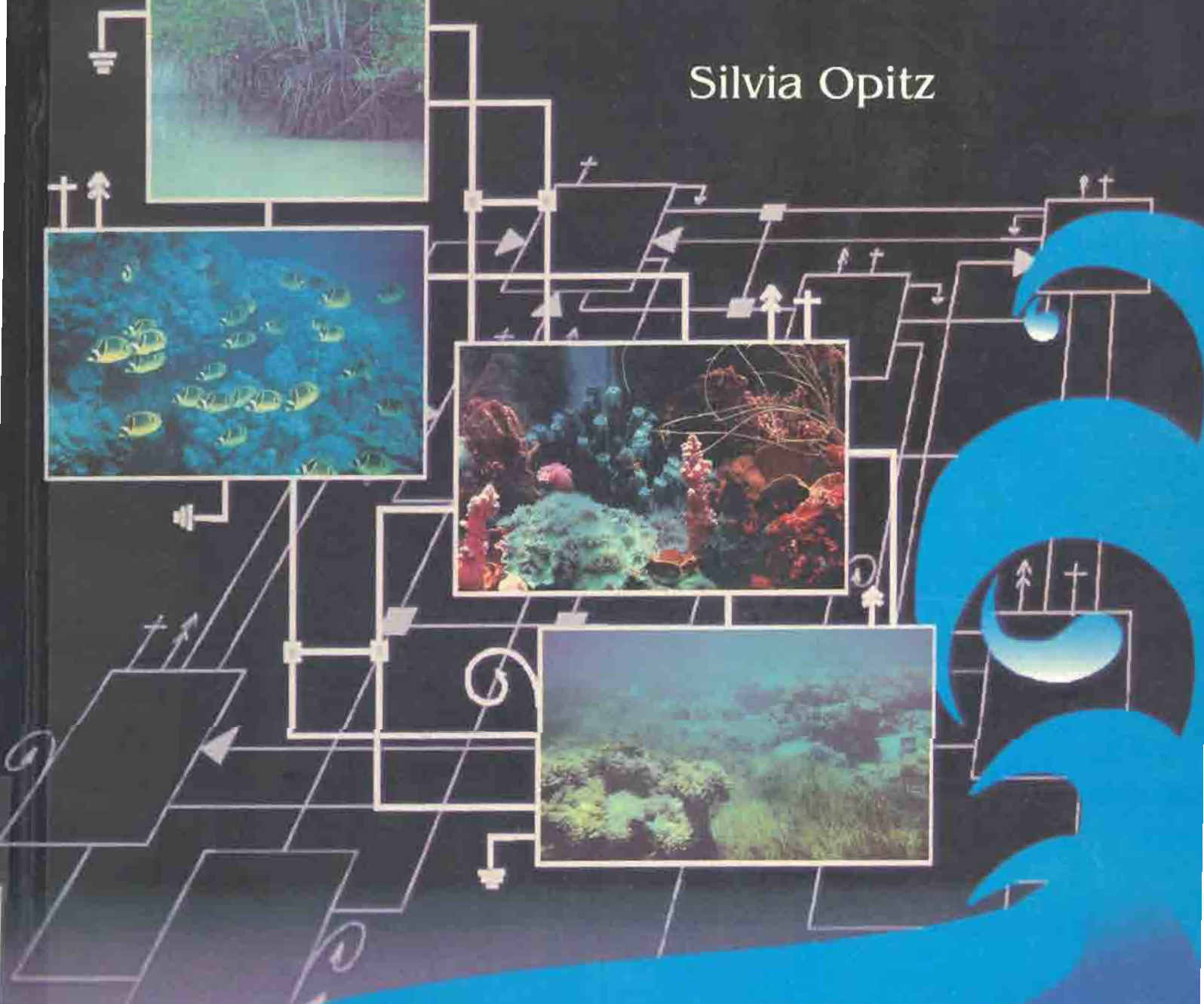


Trophic Interactions in Caribbean Coral Reefs

Silvia Opitz



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INTERNATIONAL CENTER FOR LIVING AQUATIC RESOURCES MANAGEMENT

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1996



INTERNATIONAL CENTER FOR LIVING AQUATIC RESOURCES MANAGEMENT

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APR 14 1997

SILVIA OPITZ

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Published by the International Center for Living Aquatic Resources Management, MCPO Box 2631, 0718 Makati City, Philippines.

Printed in Manila, Philippines.

Opitz, S. 1996. Trophic interactions in Caribbean coral reefs. ICLARM Tech. Rep. 43, 341 p.

Cover: Design emphasizes the systematic properties of coral reefs (central insert), inhabited by a variety of fishes (insert, left below), and often connected to other equally productive and equally threatened ecosystems, mangroves (insert, upper left) and seagrasses (insert, lower right). The wave at the lower right side symbolizes how the Sea interconnects these elements, also connected in the models (background diagram).

Copyediting by: Marie Sol M. Sadorra
Proofreading by: Marie Sol M. Sadorra, Cecille Legazpi, Francisco Torres, Jr. and Jessica A. Moya
Layout by: Ariel C. Aquisap, Albert B. Contemprate and Ovidio F. Espiritu, Jr.
Cover concept by: Silvia Opitz
Cover design by: Alan Siegfried Esquillon
Illustrations by: Albert B. Contemprate and Ovidio F. Espiritu, Jr.

ISSN 0115-5547
ISBN 971-8709-60-6

ICLARM Contribution No. 1085

13489

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FOREWORD

Coral reefs are important tropical resource systems and, through a rigorous priority setting process (ICLARM Strategic Plan 1992), were selected by ICLARM as one of the three major aquatic resource systems for research focus. The others were coastal inshore systems and inland small ponds and rice flood waters.

As one activity on coral reefs, ICLARM - in collaboration with a number of partner institutions - is developing a global database on coral reefs, ReefBase - to provide managers and scientists access to a standardized, key subset of widely scattered information on coral reefs. Comprehensive and standardized information on such complex systems as coral reefs is critical to understanding their function.

The present document - although it describes an earlier, unfished state of a small part of the world's coral reefs, the Virgin Island/Puerto Rico, area of the Caribbean - therefore presents a vast amount of standardized information. Here, the information presented was standardized through its incorporation into a mass-balance (ECOPATH) model of trophic interactions within a reef - the most detailed model of this type so far published.

The information thus standardized and validated should be useful for many other coral reef modeling efforts.

The trophic models themselves should also be of interest, as they show, also for the first time at this level of details, how groups - corals, algae, invertebrates and fish - usually studied by different scientists - interact, and impact on each other, thus providing a quantitative underpinning for a systems view.

This study*, therefore, represents a significant contribution to coral reef and aquatic systems trophodynamics research.

Meryl J. Williams
Director General
ICLARM

*Conducted by Dr. S. Opitz towards her doctorate requirements at the University of Kiel, Germany, under the guidance of Dr. D. Pauly and with advice from Dr. V. Christensen of ICLARM.

FOREWORD

The work before you is the most detailed account ever published of the energy fluxes within a coral reef ecosystem — and I am well aware that this is a strong claim to make in view of the enormous research effort which has been devoted to coral reefs to date, and of the enormous literature which presents the results of this effort.

Upon perusing that literature, however, it quickly becomes clear that only a small fraction deals with the energetics of more than two species at a time: the bulk of published coral reef studies deals with the dynamics or trophic biology of either one species, or the interactions between two species.

Studies referring to ensembles of coral reef species obviously do exist as well but they tend to concentrate on the definition, identity and maintenance of the “communities” formed by coral reef species, with little attention devoted to their energetics: few coral reef researchers followed the path opened by the pioneering study of the metabolism of Eniwetak lagoon, published in 1955 by the Odum brothers.

One reason for this may be the lack of a straightforward tool for integrating scattered knowledge on the interactive elements of coral reef ecosystems: simulation modeling, e.g., as brilliantly applied by T.R. McClanahan to coral reef systems, requires formal skills possessed by only few biologists potentially interested in using a formal modeling framework.

This situation changed in the mid-1980s, when J.J. Polovina and coworkers developed the ECOPATH approach and software, and applied this to the description of a coral reef system in the Northern Hawaiian Island, the French Frigate Shoals. This approach, fully described in the present work, does not need restating here. Suffice to say that, while based on a simple “Master Equation” with parameters that are relatively easy to estimate, it allows incorporation, into a formal framework, of the immense amount of single-species data, and on data on species interactions alluded to above.

The creators of this approach — J. Polovina and colleagues for the initial version, V. Christensen and this author for its vastly expanded successor ECOPATH II — earlier attributed the versatility of the ECOPATH approach to the “steady-state” or “equilibrium” assumption that seemed to underlie its Master Equation.

However, this assumption is not necessary for the ECOPATH approach to work. Rather, what is needed is *mass-balance*, i.e., one must not only be able to account for all flows of biomass among the different elements of an ecosystem, during the period being modeled (usually one year), but one should also account for any *difference* in biomass that may occur between the beginning and the end of that period.

Equilibrium or steady-state implies that this difference should be zero — a problem when the system being modeled is known to display interannual changes of biomass. However, recent changes incorporated on the Windows version of ECOPATH (ECOPATH 3.0) not only allow considering between-year changes of biomass (through a term for “biomass accumulated”, added to the Master Equation), but also explicitly consider seasonal changes of all biomass and fluxes during an annual production cycle.

These features, along with the explicit consideration of uncertainty on the input values (accommodated through a Monte-Carlo simulation, whose output can be interpreted in a Bayesian context), have given the ECOPATH approach dynamic features that were lacking when Dr. Opitz began the work which led to this document.

The fact that Dr. Opitz could assemble as many biomass and rate estimates as she did, and then establish mass-balance in her models — without recourse to the routines recently developed to assist users of the ECOPATH software — speaks for the depth of understanding she achieved of the energetics of Caribbean coral reef systems, and of the biology of their components.

Indeed, I assume that many future models of coral reef systems — whether constructed using the ECOPATH approach or not — will rely on the biomass and rate estimates obtained by Dr. Opitz, and validated by their “fit” to her model.

I conclude, thus, by congratulating Dr. Opitz for a work destined to be found, much thumbed, on the desk of every would-be modeler of coral reef ecosystems.

Daniel Pauly
Principal Science Adviser
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ABSTRACT

Five thermodynamically balanced models of the trophic interactions and organic matter transfer between compartments of a Caribbean coral reef system are presented. Inputs to the models were obtained from published data and from parameter estimates based on multivariate statistics. The models were analyzed using the ECOPATH II program (Version 1.0) of Daniel Pauly, Villy Christensen and coworkers at the International Center for Living Aquatic Resources Management (ICLARM) (in Manila, Philippines) which combines elements of network flow analysis with the steady-state approach of Jeffrey Polovina's original ECOPATH. A single model with 50 boxes, 2 models with 20 boxes and 2 with 11 boxes were constructed, based on two different methods of aggregation. Their features were compared.

These balanced models indicate that coral reef systems are in a "steady-state" or "flow-through equilibrium", when the appropriate spatial and temporal scale is selected. This implies that investigations on reef community structure which relied on a small spatial scale, and which suggest a high degree of stochastic variability may not address issues related to the stability of structures at larger scales.

The models' outputs indicate the existence of short cycles, effectively recycling organic matter within the reef system, with the larger part of net primary production being recycled directly to the detrital pool. Thus, transfer efficiencies between trophic levels are generally low.

Selected outputs of the models were compared with features of the original coral reef system, i.e., unfished segments of the Puerto Rico - Virgin Islands (PRVI) coral reef area and were found to be compatible.

Summary statistics of the models were compared with those of two other coral reefs also modelled using ECOPATH II. The models presented here refer to a large system in terms of biomass and throughput, compared to French Frigate Shoals, Hawaii, and to Bolinao, Philippines. These statistics indicate an intermediate stage of system maturity when compared to the other two systems. Features of the modelled system (degree of piscivory, avoidance, by piscivorous fish, of larger herbivorous reef fish as prey, sediment production by parrotfish) were compared with those of other coral reef systems and were found to be compatible.

A potential fisheries yield of $4 \text{ gm}^{-2}\text{year}^{-1}$ for unfished segments of the PRVI reef area was estimated from the models' outputs. Present overexploitation of lobsters and conchs in the PRVI area may in part be explained by the strong predation pressure already exerted on these resources in the unexploited system as indicated by the high ecotrophic efficiencies of crustaceans and gastropods.

Possible solutions could be developed or suggested for several methodological problems identified during the modelling process. Also, worthwhile areas of future theoretical and field research were identified:

1. A large number of fish species can effectively be reduced to fewer groups by cluster analysis. Appropriate aggregation parameters are those referring to energetic requirements of a species such as size, activity level and type of food.
2. The impact of predator abundance on predation pressure may be accounted for by introducing a weighting factor, which should be the product of density, mean weight and relative food requirements of a predator.

3. Density estimates from visual censuses, especially for migrating species, such as apex predators, should be divided by the time an individual fish was actually seen during the census. This would lead to more realistic values of “effective” biomass, i.e., a biomass which is related to the carrying capacity of the reef system for this species.

4. Density estimates for small and/or cryptic reef species should be obtained from repeated rotenone or similar samplings.

5. Density estimates for small schooling pelagics are urgently needed and may be obtained by hydroacoustic methods.

6. For the construction of nonfish groups one should, as was done for fishes, start from the species level and define appropriate energetic aggregation criteria by objective selection of ecological variables.

7. For future models, it will be a worthwhile theoretical task to identify objective criteria for the determination of spatial and temporal scales of the system which is to be modelled, to define appropriate system boundaries.

8. More information on diet, food consumption requirements, abundances and growth performance should be obtained for nonfish taxa in the PRVI reef area. This could strongly improve the reliability of the models' output.

9. Information on length-weight data and growth parameters for fish species without interest for fisheries (e.g., very small ones) are needed as well as catch data on a species level or for ecological guilds, i.e., on a generic or family level, for the PRVI reef area.

1. INTRODUCTION

All kinds of superlatives have been used to describe the ecosystem "coral reef", the most famous being "the oasis in the desert", which is related to the high production rates often measured for components of the coral reef community compared to rates measured in the nutrient-poor water masses surrounding coral reefs (Polovina 1984b).

There is general agreement that coral reefs are characterized by three main features: (1) high species diversity; (2) pronounced complexity of relationships within the system community; and (3) high rates of production in usually nutrient- and plankton-poor oceanic waters (see Lewis 1981).

It is hypothesized here that coral reefs can survive under such unfavorable conditions mainly because these three features are interconnected, resulting in an effective recycling and conservation of organic matter within the system. This hypothesis is based on small-scale observations such as the symbiosis between the coral polyps and their symbionts, the "zooxanthellae". Whether this hypothesis applies to an entire coral reef ecosystem unit is, to date, not known. There is further disagreement on the question whether coral reefs may be considered to be in a steady-state, i.e., whether import plus consumption are balanced by production plus respiration for the entire system.

The main objective of this report was to test whether it is possible to construct a thermodynamically balanced model of a Caribbean coral reef system, the assumption being that if such model could be constructed, this would indicate that the coral reef system in question can indeed be considered as being in some form of steady state.

A Caribbean coral reef area was selected as study site for a modelling effort because I became enthralled by these beautiful ecosystems since I have "dived" Caribbean coral reefs in 1983/1984. Another reason was that several Caribbean reef areas have been studied in such detail that enough input data appeared to exist for the creation of a model representative of the larger Caribbean area. My choice of a model of a Caribbean coral reef as a research topic was also influenced by the consideration that tools for proper management of coral reef systems, especially of fisheries resources, are urgently needed.

The whole system approach seems very promising, since it will provide information for conservationists with a holistic claim as well as for fisheries managers concerned with fisheries yields from a small number of target species. Several of these commercially interesting species are already heavily exploited and some show declines in catch/effort although they are reported to be only lightly exploited. Munro (1984), in an unpublished report to the Caribbean Fisheries Management Council (CFMC) "on assessment of the fisheries of Puerto Rico and the U.S. Virgin Islands", attempted to find out why "the reported landings for the Virgin Islands only amount to 5%-13% of the 'norm' (for other Caribbean reef areas) of 5 tonnes km⁻²". He concluded that "either the landings are greatly underestimated or the fisheries are rather lightly exploited or both". In contrast to this statement, fisheries statistics in CFMC (1984, 1985) showed a decline in landings for target fish species in the Puerto Rico - Virgin Islands (PRVI)

area, and contributions in Mahon (1987) reported on the overexploitation of target invertebrates (conchs, lobsters) along with an increasing fishing effort for the same area.

The lack of appropriate data for proper stock assessment is emphasized regularly throughout the relevant literature (see e.g., Munro 1984; Mahon 1987; PDT 1990). Many unpublished reports of investigations and surveys lead a poor life in shelves or drawers; the more lucky ones are distributed through the person-to-person method. Thus, there is a need to pool, check and quantify data scattered over a wide range of formal and informal documentation. The construction of a model is an excellent reason for such compilatory work. By logical combination of the elements of a system, one may obtain new insights in its functioning as a whole and in that of its elements as well.

Holistic steady-state models, in contrast to simulation models, can be parameterized with relatively small expenditure in input data while they may, at the same time, correctly describe properties of the system being modelled (Silvert 1981). In spite of their "simplicity", these models may reflect relevant structures of the system and contribute to its understanding; simultaneously, they may indicate gaps of knowledge and thus identify areas of future practical research. According to Silvert (1981), the "complexity of a model is not in itself a criterion of its quality. The degree of complexity is determined chiefly by its function and the observations it is supposed to describe, not by the internal structure of the system" that is being modelled.

Leontief (1951) developed economic input-output analysis aiming to quantify the amount of raw materials and industrial services required to produce a quantity of consumer goods. Augustinovic (1970) later worked out an inverse analysis where the fate of system inputs is traced to their destination through the economic flow diagram. Hannon (1973) and Finn (1976) were the first to apply these techniques to biological systems.

Network analysis today provides tools for unifying the concepts of food chains, food webs, species richness, diversity, etc., by depicting ecosystems as number of compartments, interconnected by flows of energy or matter from one compartment to another. Wulff et al. (1989) believed that "network flow analysis contains a lot of information about the dynamic structure of a whole system and how it functions".

Network analysis today includes analysis of cycles, through-flows, storages and information theory. Finn (1976) developed an index showing the proportion of the flow in a system that is recycled, relative to total flow. The magnitude of this "cycling index" depends to some extent on the substance used as currency (biomass, carbon, nitrogen, phosphorus, etc.). Ulanowicz (1980, 1986), one of the leading scientists of Working Group 59/73 of the Scientific Committee on Oceanographic Research devoted to "Mathematical Models in Biological Oceanography", (Platt et al. 1981), introduced three indices describing the state of an ecosystem: (1) Total System Throughput, the sum of all flows occurring within a system; (2) Ascendency, a measure of the mutual information content within the system; and which concerns the size of the system as well as diversity of flows; and (3) Development Capacity, a measure of the potential for growth of a system towards maturity.

In network analysis, comparison of two different elements (e.g., species) within a system may be established by defining their relative position in the food web as one plus the weighted mean trophic level of their preys. Conversely, the division of any kind of diet into compartments allows the construction of pyramids (see Lindeman 1942) representing flows and transfer efficiencies between trophic levels. This technique allows intercomparison of ecosystems or comparison of different states of a single system based on the analysis of the food web linking its components.

1.1. The ECOPATH Model

The modelling approach used here was developed by J.J. Polovina (1984a, 1984b, 1984c, 1985, 1986) and Polovina and Ow (1983), named "ECOPATH", was originally designed to estimate standing stock and production budget of an entire coral reef ecosystem, the "French Frigate Shoals", in the Northwestern Hawaiian Islands.

ECOPATH II is derived from the original ECOPATH software. It was considerably enhanced and modified by combining the steady-state approach of the original ECOPATH with techniques derived from network analysis and information theory. ECOPATH II was designed as a tool to produce straightforward equilibrium box models for any kind of aquatic system. The development and updating of ECOPATH II are part of a project funded by the Danish International Development Agency (DANIDA) on "Global Comparisons of Aquatic Ecosystems", executed by Drs. Villy Christensen and Daniel Pauly at the International Center for Living Aquatic Resources Management (ICLARM). Aquatic scientists, spread all around the world, are contributing to this project by applying ECOPATH II to "their" ecosystems. The present report is based on a thesis written as part of this project; therein, ECOPATH II was used to generate five quantitative steady-state models of the trophic interactions in a Caribbean coral reef.

In the following, a short comparative description is given of the theoretical background of inputs required and outputs provided by Polovina and Ow's original ECOPATH and by the ECOPATH II software of Pauly, Christensen and coworkers (Christensen and Pauly 1990a, 1990b; Pauly et al. 1993a). Analyses of models included here were executed with version 1.0 of ECOPATH II. Since version 2.0 of ECOPATH II is distributed together with a comprehensive user's manual (Christensen and Pauly 1991), I advise the reader to consult this manual for additional information on differences between versions 1.0 and 2.0 (see also Christensen and Pauly 1992b for version 2.1 and Christensen and Pauly 1995 and 1996 for ECOPATH 3.0).

1.1.1. Comparative Description of ECOPATH I and ECOPATH II

1.1.1.1. POLOVINA'S ORIGINAL ECOPATH

Theoretical Background

The basic assumption of the original version of the ECOPATH model (termed here ECOPATH I) is that the ecosystem which will be modelled must be under equilibrium conditions, i.e., in a "steady-state". Polovina (1984b) stated that "equilibrium conditions exist when the mean annual biomass for each species (group) does not change from year to year. This assumption results in a system of biomass budget equations which, for each species (group), can be expressed as:

Production of biomass for species (i) - all predation on species (i) - nonpredatory biomass mortality for species (i) = 0 for all (i) ...1.1)

The ECOPATH model expresses each term in the budget equation as a linear function of the unknown mean annual biomasses (B_i 's), so the resulting biomass budget equations become a system of simultaneous equations linear in the B_i 's." This system of equations can be

expressed in matrix form as $AB = 0$, where A is an $n \times n$ matrix of coefficients, and B is an n -dimensional vector.

Outputs

Outputs of ECOPATH I are: estimates of mean annual biomass, mean annual consumption and mean annual production of each species (group). Additionally, an estimate of necessary net primary production to support the system is computed which may be compared with "external" estimates to validate the model.

Inputs

The inputs required by ECOPATH I (and ECOPATH II) are few compared to, e.g., simulation models. In the following, the inputs necessary to solve the system of biomass budget equations via the ECOPATH I computer program are briefly introduced.

Species Groups

Subsequent to identifying the various components of the ecosystem under study, these may be regrouped into species groups. ECOPATH I allows a maximum input of 20 species (groups). A species group is an aggregation of species with a common physical habitat, similar diet and similar life history characteristics (Polovina 1984b).

Biomass

ECOPATH I is a "top-down model", i.e., the biomasses of the subordinated species (groups) are determined by the biomass (and food consumption) of the apex predator (group). When there is no fishery catch, a biomass estimate for at least one top predator (group) must be provided to produce a unique and nontrivial set of B_i 's (i.e., $B_i > 0$) which solves the system of equations. In an ecosystem where there is fishing mortality, the fish catches may be entered instead of an estimate of a species' (group's) biomass to insure a nontrivial solution.

Production/Biomass Ratio (P/B)

Polovina (1984b, 1985) defined production (P) over one year for a cohort of animals as:

$$P = \int_0^1 N_t \frac{d}{dt}(W_t) dt \quad \dots 1.2)$$

and mean annual biomass (B) for the cohort as:

$$B = \int_0^1 N_t W_t dt \quad \dots 1.3)$$

where N_t is the number of animals and W_t the mean individual weight at time t .

Allen (1971) showed that, under equilibrium conditions, for a number of growth and mortality functions, including negative exponential mortality and “von Bertalanffy” growth, the ratio of annual production to mean biomass for a cohort of fish is equal to the annual instantaneous total mortality (Z). When mortality is expressed by a negative exponential function, total species group production (P) is equal to the sum of the production (P_i) of n cohorts (or species) i with annual instantaneous total mortality (Z_i).

In an ecosystem under equilibrium (i.e., under steady state) and where (when) there is very little or no fishing mortality, the P/B ratio of fishes and invertebrates is thus equal to their instantaneous rate of natural mortality (M) (Christensen and Pauly 1991).

Predation Mortality

Predation mortality is the fraction of the biomass of a species group which is consumed by its predator in a given ecosystem. Polovina’s ECOPATH model computes this mortality in the same fashion as the PROBUB model by Laevastu and Larkins (1981). Here, two types of information are needed:

Diet Composition Matrix

A diet composition matrix DC_{ij} must be specified, where an entry DC_{ij} from this matrix refers to the proportion of prey j in the diet of predator (or consumer) i . The primary source of this information is the analysis of consumers’ stomach contents. Composition of stomach contents may be expressed in percent of total volume, weight or energy contents. Macdonald and Green (1983) showed that the estimation of diet composition is largely unaffected by the units used (volume, weight or energy); on the other hand, “numbers of items” or “occurrence” produce misleading diet compositions, due to the large difference in energy contents between small and large prey items (Pauly 1985, 1986).

Food Required

The second type of information needed to ascertain predation mortality is the food requirements of the predator. Food required (R_i) by a species (group) (i) is expressed as:

$$R_i = b_i B_i + a_i P_i \quad \dots 1.4)$$

where B_i is the mean annual species (group) biomass, P_i is the annual production of species (group) i , and a_i and b_i are parameters to be estimated from energetics studies. The component $b_i B_i$ is the food required to maintain the biomass B_i and the component $a_i P_i$ is the food required to support the biomass production P_i (Laevastu and Larkins 1981).

In the original ECOPATH model, the production of species (group) i is $P_i = C_i B_i$, so the food required for species (group) B_i is

$$R_i = b_i B_i + a_i P_i = b_i B_i + a_i C_i B_i = (b_i + a_i C_i) B_i \quad \dots 1.5)$$

Thus, the amount of species (group) j consumed by predator species (group) i is given as:

$$R_i DC_{ij} = (b_i + a_i C_i) B_i DC_{ij} \quad \dots 1.6)$$

Predation mortality consumes generally between 75% and 90% of the populations' production (Ricker 1969; Polovina 1984a, 1984b); a change in predation mortality has little influence on the biomasses of species groups calculated by ECOPATH I, as was shown by Polovina (1984c) by simple sensitivity analysis on the input parameters.

Nonpredation Mortality

All mortality attributable to causes other than predation, such as fishing, spawning and disease, is considered "nonpredation mortality".

Habitat Area

For each species (group) included in ECOPATH I, an estimate of the area of distribution for which the other input values are valid was required. This was not included in ECOPATH II as biomass and flows are easy to adjust to any standard area, without using a special software or routine.

Summary of inputs required for the original version of ECOPATH:

1. number and names of species (groups) considered in the model;
2. at least one biomass estimate of one of the apex predators or its annual fisheries catch;
3. annual fisheries catch (including discard) for all exploited species (groups), or nonpredation mortality ($d_i B_i$) when there is no fishing;
4. P/B ratio (usually M_i or Z_i) for each species (group);
5. food required to support biomass production ($a_i P_i$) of each species (group);
6. food required to maintain the biomass ($b_i B_i$) of each species (group);
7. diet composition DC_{ij} of each species (group); and
8. habitat area of each species (group).

1.1.1.2. ECOPATH II

Theoretical Background

ECOPATH II shares with ECOPATH I its "basic equation" (1.1). This routine balances an equilibrium model using the mass-balance equation:

$$\text{Consumption} + \text{Import} = \text{Production} + \text{Respiration}$$

where "production" is the sum of "export + mortality due to predation + flow to detritus". Both versions of ECOPATH (I and II) are structured around the feature that, in a balanced system, the consumption of a predator (group) generates the predation mortality of its prey (group/s).

New Features and Outputs

DEFINITIONS AND PARAMETER ESTIMATION ROUTINE. In ECOPATH II, the set of simultaneous linear equations, representing an ecosystem, is solved using the “generalized inverse method” (Mackay 1981); this has the advantage that a generalized inverse can be found even if the set of equations is overdetermined (more equations than unknowns) or underdetermined (less equations than unknowns). By linking the species groups into a combined system, the basic routines of ECOPATH I and II allow the estimation of one missing parameter (= unknown) of each species (group). In ECOPATH I, only biomass could be the unknowns, whereas other parameters may be unknown in ECOPATH II (see Christensen and Pauly 1992a, 1992b and further below).

For all consumers of the system, a request was incorporated into ECOPATH II for the percentages of ingestion that are lost due to egestion (feces) and to excretion (urine); values of 15% and 5%, respectively, are provided as defaults, based on Winberg (1956). The entries (or defaults) are subsequently used for the computation of net conversion efficiencies (production/[ingestion - egestion]) and of flows to the “detritus box”. Detritus is derived from nonpredatory “leaks” of organic substance from any trophic level, mainly egestion, excretion and nonpredatory mortality.

ECOPATH II was provided with checks for “i’s” that do not balance, i.e., for which production is larger or equal to the amount of food consumed. Gross efficiency (GE = production/consumption) should be < 0.5 for all species (groups). Ecotrophic efficiency (EE), the part of production consumed by predators, was, throughout, a required input in ECOPATH I. It is now either an input or output of ECOPATH II and is used to ascertain that the amount consumed by predators does not surpass production of a species (group). In a balanced system, EE must be ≤ 1.0 .

ECOPATH II calculates fractional trophic levels as suggested by Odum and Heald (1975). A routine assigns a trophic level (TL) of 1 to primary producers and detritus and a TL of 1+ (the weighted average of the prey’s TL) to consumers.

The concept of “omnivory index” was included into ECOPATH II in 1987 (see Pauly et al. 1993a). It is calculated as the variance of the trophic levels of a consumer’s prey groups. When the output value is zero, the consumers in question feed on a single discrete trophic level.

Ivlev’s electivity index E_i (Ivlev 1961) is computed by ECOPATH II for each species (group), based on biomasses, not numbers. E_i is scaled so that $E_i = -1$ corresponds to total avoidance, $E_i = 0$ to nonselective feeding, and $E_i = 1$ to exclusive feeding. Since this index is not independent of prey density a further index, the “standardized forage ratio” of Chesson (1983) was incorporated into ECOPATH II. This is independent of prey availability and initially ranged between 0 and 1, with $S_i = 0$ representing avoidance and $S_i = 1$ exclusive feeding. In ECOPATH II, this index has been reexpressed such that it can now range between -1 and 1, and thus allows direct comparison with Ivlev’s index of electivity.

NETWORK FLOW INDICES. A routine for calculating “ascendency” was included in ECOPATH II based on Ulanowicz (1986) as modified by Ulanowicz and Norden (1990). “Ascendency” is a measure of the “average mutual information in a system”. It is scaled by system throughput (sum of all flows) and expressed in flowbits (e.g., bit * gm⁻²·year⁻¹ WW). In addition to ascendency and related statistics (overhead, capacity, etc.), holistic properties of ecosystems that can be computed using ECOPATH II are total system throughput and Finn’s cycling index

(Finn 1976). The theory leading to these system properties can be found in Ulanowicz (1986) and is further discussed in Christensen and Pauly (1991).

FLows AND TRAnSFER EFFICIENCIES. Based on Lindeman (1942), Christensen and Pauly (1990b) provided ECOPATH II with routines to calculate discrete trophic levels and, since an ecosystem is treated as a thermodynamical unit, to calculate flows and transfer efficiencies between trophic levels. The combination of throughput and transfer efficiency of each trophic level allows the construction of solid "trophic pyramids" (see Christensen and Pauly 1992b) representing an entire system; these pyramids can be used for intercomparison of ecosystems.

MIXED TROPHIC IMPACTS. This routine allows the computation of direct and indirect impacts which a change in biomass of a predator group will have on other groups in the system, assuming that the diet matrix remains unchanged. This routine, based on Hannon (1973) and Hannon and Joiris (1989), may thus be viewed as a tool for sensitivity analysis.

AGGREGATION OF BOXES. The ascendancy and related features of an ecosystem are affected by the number of groups by which the system is represented. An aggregation routine was included into ECOPATH II which allows for reduction of the initial number of boxes to any desired number. This routine identifies, under the constraint of no change in throughput, the pair of "boxes" which, when pooled, least reduce the ascendancy of the ecosystem. As shown in Christensen and Pauly (1992b), "pairing" of boxes generally involves groups with low throughputs (at least initially) and having similar trophic levels.

Inputs Required by ECOPATH II

PRIMARy PRoDUCTION (PP). Net primary production (NPP, in the currency of the model, here in $\text{gm}^{-2}\cdot\text{year}^{-1}$ WW) is not an input for ECOPATH I. However, this is a required input for ECOPATH II, in order to "(1) encourage users to obtain independent, i.e., "external" estimates of a parameter which closely correlates with a number of important characteristics of a system; (2) to allow for validation of internally generated NPP, which should be smaller than or equal to the "external" input value and more importantly; (3) to allow for the difference between observed (= external) and computed (= internal) NPP to be added to the detritus box and thus quantify a linkage important in virtually all ecosystems (i.e., the transfer of uningested, dead phytoplankton and algal biomass to the detrital pool, from where it becomes available to a variety of consumers and/or for burial or export" [Pauly et al. 1993a]).

DETRITUS BIOMASS (D). Detritus biomass is not a required input in ECOPATH I, but is needed by ECOPATH II for various computations when detritivores are included in a system, e.g., to compute the "electivity" of detritivores for detritus.

BIOMASS AND/OR ECOTROPHIC EFFICIENCY. Ecotrophic efficiency (EE) is an input of ECOPATH I expressing the fraction of total production consumed by predators ($a_i P_i$). In ECOPATH II, ecotrophic efficiency must be either entered (when biomass is unknown) or is estimated by the program (when biomass is known and was entered). In ECOPATH II, the fraction $1-EE$ of production is directed toward the detritus box, from which it may be exported out of the system.

EXPORT AND/OR FISHERIES CATCH (C). Export from a species (group) consists of catches and emigration or transport out of the system. Fisheries catch (and/or the quantity killed by fishing and discarded) is required for all exploited species. This input is needed for both versions of ECOPATH in a system of which some elements are exploited.

PRODUCTION/BIOMASS RATIO (P/B). The P/B ratio is an input parameter required for ECOPATH I as well as ECOPATH II. As stated above, under certain conditions, the P/B ratio of a population can be set equal to instantaneous total mortality Z (or natural mortality M for an unfished stock). The P/B ratio of primary producers can be estimated either by direct methods (oxygen production, C_{14} , etc.) or if those data are not available, by indirect methods, e.g., empirical equations (Lafontaine and Peters 1986; Pauly et al. 1993a). Mann (1982) provides P/B estimates for various benthic algae and seagrasses.

FOOD CONSUMPTION PER UNIT BIOMASS (Q/B). The relative food consumption of a species (group) other than primary producers is required as input parameter by both versions of ECOPATH. Pauly et al. (1993a) defined Q/B as the number of times a population consumes its own weight per year. In ECOPATH I, this input corresponds to the "food required (FR)". Pauly (1986) used the symbols "Q/B" for cases when FR refers "to age-structured populations, i.e., when one must consider the fact that there are, in a natural, steady-state population of fish more young than old fish, and that the former eat (per unit weight) more than the latter" (Pauly et al. 1993a). Methods to calculate Q/B of an age-structured population are described in the "Materials and methods" section.

DIET COMPOSITION. The diet composition (DC) is also an input required by both versions of ECOPATH. Only species (groups) occurring within the system may be part of the diet^a and, at least in ECOPATH II, some species (groups) should feed on detritus, as also happens in reality.

UNASSIMILATED FOOD. Egestion and excretion refer to parts of ingestion that are not assimilated. The respective fractions can be entered as an optional input if known for a species (group). Default values of 15% and 5%, respectively, are provided by the program based on Winberg (1956).

Summary of Inputs Required for ECOPATH II:

1. one independent estimate of net primary production (NPP);
2. one independent estimate of detritus biomass (D);
3. number of species (groups) considered in the model;
4. biomass (B_i) for each species group or ecotrophic efficiency (EE) when biomass is unknown;
5. export (usually fisheries catch, including discard) for all exploited species [groups];
6. P/B ratio (usually M_i or Z_i) for each species (group);
7. food consumption per unit biomass (Q/B) for each species (group);
8. diet composition DC_{ij} of each species (group); and
9. optional inputs: egestion and excretion (defaults 15% and 5%); the defaults were accepted here throughout.

^aIn recent version of ECOPATH, consumers may also feed on "imports," i.e., items not included in the system.

1.2. Development of the Models

Fig. 1.1 shows a flow chart of the logical development and construction of the five models of a Caribbean coral reef that are presented here.

Starting from the motivation for such a modelling effort, the first step was to formulate an objective, the second to define the size and boundaries of the system which was to be modelled. The next step was the collection and definition of the elements represented in the system. Empirical methods were used to obtain values for the key input parameters of ECOPATH II. The literature was systematically searched for data useful for the composition of a diet matrix, for the estimation of preliminary biomass values and for the computation of vital statistics by empirical methods. After a preliminary set of inputs was prepared, the elements of the system were reduced to 50 species (groups) or "boxes", using a clustering technique.

The reduced set of inputs was entered into ECOPATH II and subsequently modified until the 50-box model was balanced. The final outputs were compared with the initial inputs and discrepancies discussed.

The balanced 50-box model was then reduced to 20-box and 11-box models by two different methods of aggregation. The resulting 20- and 11-box models from both methods were compared. Based on the outputs from all models, the implications of the modelling were discussed.

The number(s) of the section(s), in which the above steps are described, is (are) given in the upper right corner of the boxes in Fig. 1.1.

1.3. Objectives

Summarizing, the main objectives of this report are:

1. to test whether it is possible to construct quantitative steady-state models of the trophic interactions in a Caribbean coral reef;
2. to assess the reliability of such models by comparison of outputs with independent quantitative information on the dynamics of Caribbean and other coral reefs;
3. to possibly identify, describe and quantify presently unknown properties (qualitative and quantitative) of coral reef ecosystems, relevant to their functioning and exploitation by fishing;
4. to identify methodological problems associated with the modelling process and propose solutions to these, e.g., on how to handle the high species diversity on coral reefs, based on various taxonomical approaches, or on how to deal with migrating species that only spend a fraction of their time on the reef; and
5. to reveal information gaps where additional theoretical and field research is needed.

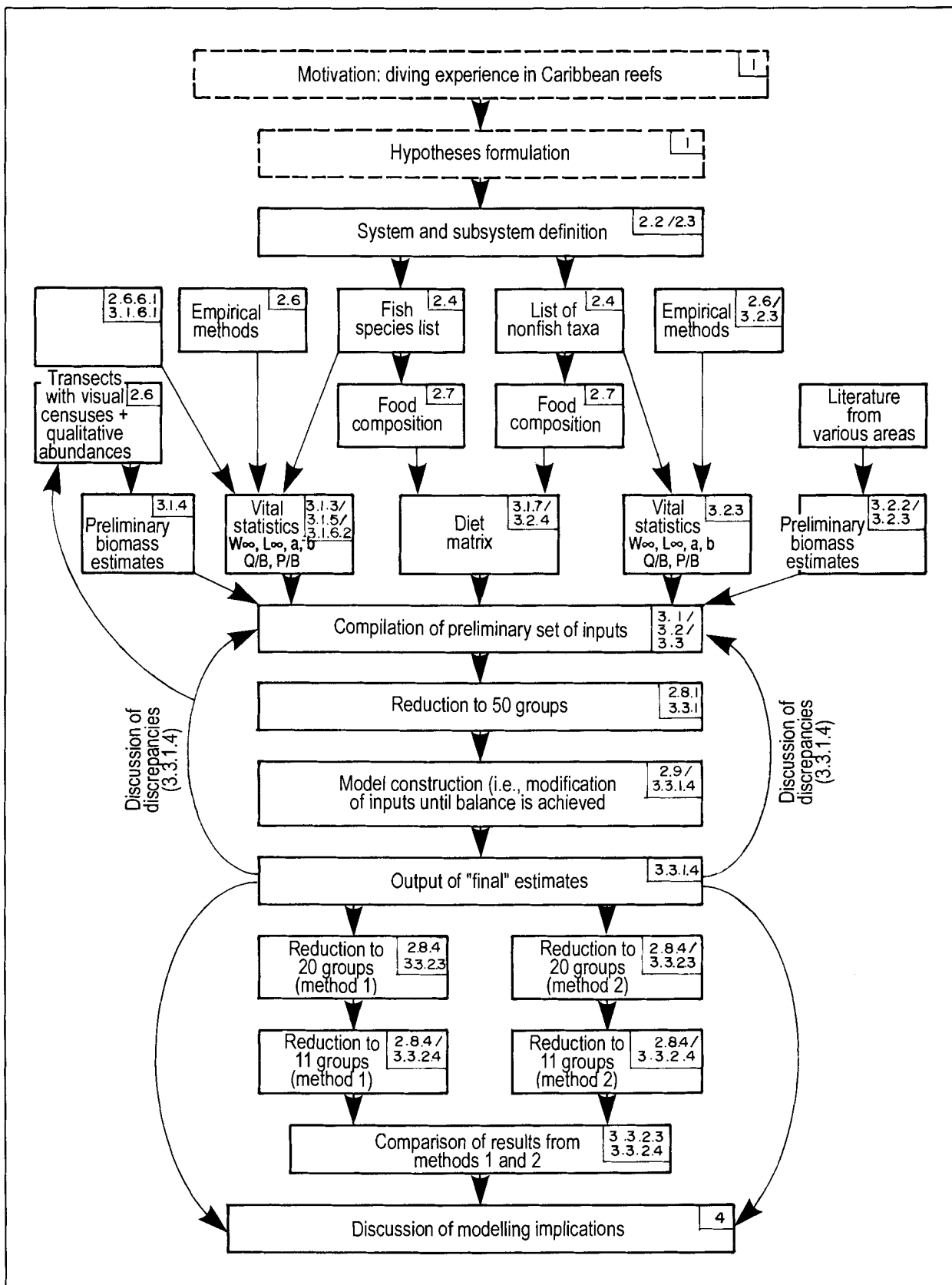


Fig. 1.1. Flow chart showing development and construction of five trophic models of a Caribbean coral reef. The numbers in the upper right corner of the boxes refer to the sections describing the steps.

2. MATERIALS AND METHODS

2.1. Sources of Data

Input data used here for the modelling of a Caribbean coral reef were obtained from published reports and papers and from parameter estimates based on multivariate statistics.

2.2. Choice of Suitable System

The decision to model a Caribbean, rather than an Indo-Pacific reef, was made on the basis of the author's personal SCUBA diving experience in the Caribbean, and because it was assumed that more literature data would be available for the former area.

Within the Caribbean, the Puerto Rico-Virgin Islands (PRVI) shelf complex appears to have been studied in greater detail than any other area, notably with regard to the food habits of fishes (see Randall 1967). Nevertheless, information from other areas had to be considered; wherever possible, data were taken from other areas of the Caribbean. When such data were lacking, literature on Indo-Pacific reefs was consulted, then that covering other tropical areas. Some literature data on various invertebrate groups were also taken from subtropical and temperate areas; in such cases, the relevant rates were always adjusted to the higher temperatures of PRVI area (see section 2.6.6).

2.3. System Boundaries and the Concept of "Subsystems"

The PRVI insular shelf extends over more than 1,000 km². The US Virgin Islands of St. Thomas, St. John and St. Croix are surrounded by about 200 km², the British Virgin Islands by 343 km² and Puerto Rico by 554 km² of shelf area. A summary description on topographical features of the PRVI shelf area may be found in Ogden and Gladfelter (1983) and in Jacobsen and Browder (1987).

The coral reefs fringing the Virgin Islands form one of several distinct ecological units that are interconnected through various components and processes. Based on our own independent literature studies, we agree with Jacobsen and Browder's (1987) concept of the shallower part of the reef consisting of distinctive, though interconnected subsystems (Fig. 2.1, Table 2.1).

Six of the seven subsystems of Jacobsen and Browder (1987) were combined here into a modelling "unit" comprising a transect from the water edge (mangrove estuaries) at 0 m depth down to the sand/mudflats at the 100-m contour with the extension of the pelagic subunit reduced to the area overlying the five demersal subunits. The shelf ("break") area below 100 m depth, which Jacobsen and Browder (1987) identified as their seventh subsystem, was not considered here, mainly because of the lack of suitable information (on, e.g., presence of nonfish taxa, abundance of species or feeding relationships), although it might be significant in both ecological and fisheries contexts.

The five demersal units were assumed to comprise each of the same area, i.e., 20% of the total area. This is a rough estimate, but the data in Table 2.1 do not suggest this to be erroneous. The inputs and outputs of biomass, production and consumption for the components of the system were all estimated for an average square meter from the five demersal subsystems and the overlaying pelagic zone.

2.4. System Components

The system was divided into two main components: fish species and nonfish taxa.

2.4.1. Fish Species

Overall, Caribbean reefs support over 500 species of fishes described in specialized

Table 2.1. Areal extent of subsystems of the PRVI shelf (based on Jacobsen and Browder 1987).

| Subsystem | Location | Area (km ²) |
|--------------------|------------------------------------------------------------------------------------|-------------------------------------------|
| Mangrove estuaries | Puerto Rico (PR) Virgin Islands (VI) | 6.405 n.s. |
| Seagrass beds | Culebra (PR) St. Croix (VI) | 0.433 2.322 |
| Coral reefs | Culebra (PR) Vieques (PR) St. Croix (VI) St. Thomas (VI) St. John (VI) | 0.923 0.245 3.584 1.901 0.855 |
| Algal plains | Puerto Rico Virgin Islands | n.s. n.s. |
| Sand/mud | Puerto Rico Virgin Islands | n.s. n.s. |
| Shelf break | PR and VI | 78.000 |
| Pelagic zone | PR and VI | 10,900.000 |

n.s. - not stated.

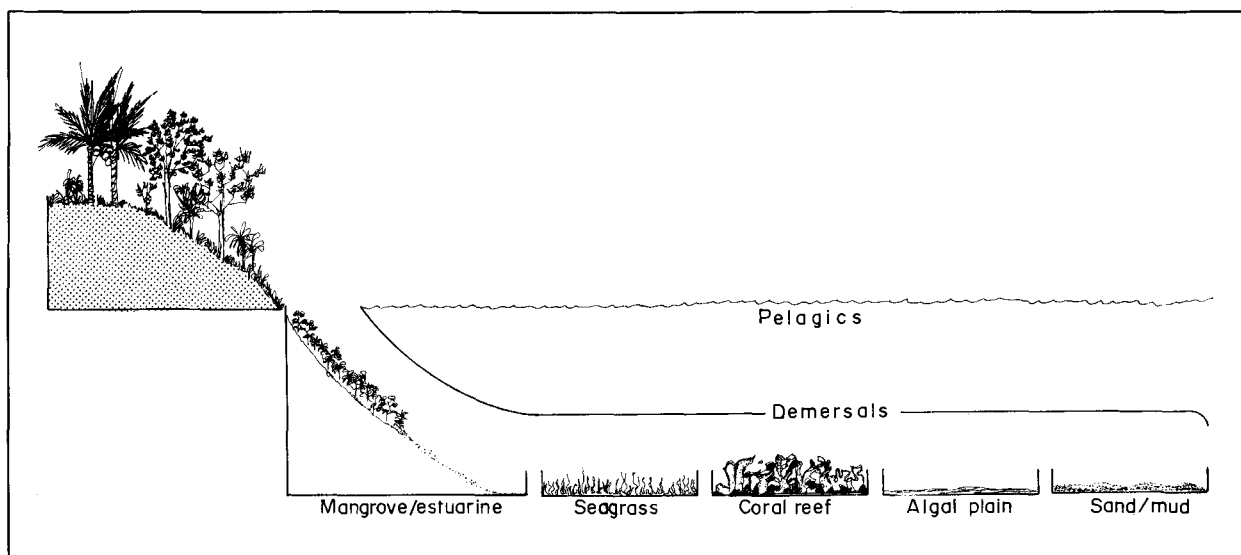


Fig. 2.1. Diagram of conceptual subsystems included in the models (modified from Jacobsen and Browder 1987).

texts (Bohlke and Chaplin 1968; Randall 1968, 1983). However, site-specific studies of the PRVI area reported fewer than 500 fish species: Parrish (1982) recorded 130 species on a reef off southwest Puerto Rico; Collette and Talbot (1972) recorded 107 species on a reef off St. John; Clavijo et al. (1980) listed 212 fish species that used reefs off St. Croix as "typical habitat" while adults. Kimmel (1985) identified around 200 fish species on reefs of La Parguera, Puerto Rico.

The more than 200 fish species included in Randall's (1967) study of the food habits of West Indian reef fishes were taken as scaffold for the model. This species list was completed by

including more fish species with affinities with coral reefs, especially sharks, from Fischer (1978). Table 2.2 lists all the fish species included in the present model.

2.4.2. Nonfish Taxa

The nonfish taxa included in the model consist mainly of groups reported by Randall from fish stomachs, as completed with information from various published sources (Table 2.3). These nonfish taxa comprise different functional groups. The energetic basis of the system is the detrital pool, with particulate organic matter (POM) and dissolved organic matter (DOM) included under the same label. This pool receives all dead organic material, excreta and egesta from the organisms in the system. Conversely, this pool serves as a food source for different detritivores of various levels of organization, e.g., bacteria, worms, crustaceans, finfish (Sorokin 1987).

Autotrophs belong to the next category of organisms. The following primary producers contribute to the reef metabolism (Mann 1982):

1. symbiotic algae, which consist of symbiotic dinoflagellates, called "zooxanthellae", and filamentous chlorophyceans (the former live inside the wall cells of the coral polyps and exchange metabolites directly with the host animal; the latter live attached to the coral skeleton and contribute roughly three times more weight than the former to total organic mass of hermatypic corals [Odum and Odum 1955]);
2. macrophytic brown and red algae;
3. coralline algae;
4. seagrasses (spermatophytes);
5. benthic diatoms; and
6. phytoplankton.

Randall (1967), in his analysis of reef fish stomachs, identified around 150 species of primary producers as food items. A variety of organisms was pooled under the label "decomposers/microfauna". The elements of this group are single-celled heterotrophic organisms such as bacteria, ciliates, foraminiferans, etc. This functional group is, like the detritus pool, of utmost importance for the entire reef metabolism. Through very high turnover rates and the ability to transform dead organic material such that it is reintroduced into the trophic cycle, the decomposer/microfauna group can be viewed as the "motor" of the reef whereas the detritus pool serves as "fuel" (note that this analogy has a problem, since some part of the "motor" is always consumed by the reef community).

As was done with "decomposers/microfauna", various organisms were pooled under the label "zooplankton" (e.g., several crustacean families, jellyfish, eggs and larvae, insects). Benthic invertebrates, sea turtles and birds form the last categories of nonfish groups.

The version of ECOPATH II used for this work did not allow the inclusion of consumers with a P/R ratio > 1.^a Therefore the symbiotic coral-zooxanthellae-filamentous algae complex was separated into a producer component (the zooxanthellae and filamentous algae, which were grouped with the benthic autotrophs) and a consumer component (the heterotrophic part of the coral polyps).

The models presented here deal exclusively with the organic part of the system, i.e., the inorganic part, such as the skeletons of corals, foraminiferans, worms, etc., are not considered.

^aRecent version of ECOPATH allows this.

Table 2.2. Fish species included in the ECOPATH II models of a Caribbean coral reef. Species are listed in alphabetical order of families. English and Spanish common names are from Fischer (1978).

*=not included in diet composition matrix.

| Family | Species | English common name | Spanish common name |
|---------------|----------------------------------------------------------------|-----------------------|-----------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> Castelnau 1855 | Ocean surgeon | <i>Navajon pardo</i> |
| | <i>Acanthurus chirurgus</i> (Bloch 1787) | Doctorfish | <i>Navajon cirujano</i> |
| | <i>Acanthurus coeruleus</i> Bloch & Schneider 1801 | Blue tang surgeonfish | <i>Navajon azul</i> |
| Albulidae | * <i>Albula vulpes</i> (Linnaeus 1758) | Bonefish | <i>Macabí</i> |
| Antennariidae | <i>Antennarius multiocellatus</i> (Cuvier & Valenciennes 1837) | Longlure frogfish | - |
| | <i>Antennarius striatus</i> (Cuvier 1817) | Splitlure frogfish | - |
| Apogonidae | <i>Apogon conklini</i> (Silvester 1915) | Freckled cardinalfish | - |
| | <i>Apogon maculatus</i> (Poey 1861) | Flamefish | - |
| Atherinidae | <i>Allanetta harringtonensis</i> (Goode 1877) | Reef silversides | - |
| | <i>Atherinomorus stipes</i> (Müller & Troschel 1848) | Hardhead silversides | - |
| Aulostomidae | <i>Aulostomus maculatus</i> Valenciennes 1842 | Trumpetfish | <i>Trompeta</i> |
| Balistidae | * <i>Balistes capriscus</i> Gmelin 1788 | Gray triggerfish | <i>Pejepuerco blanco</i> |
| | <i>Balistes vetula</i> Linnaeus 1758 | Queen triggerfish | <i>Pejepuerco cachuo</i> |
| | <i>Canthidermis sufflamen</i> (Mitchill 1815) | Ocean triggerfish | - |
| | <i>Melichthys niger</i> (Bloch 1786) | Black durgon | <i>Calafate negro</i> |
| | * <i>Xanthichthys ringens</i> (Linnaeus 1758) | Sargassum triggerfish | - |
| Belonidae | * <i>Ablennes hians</i> (Valenciennes 1846) | Flat needlefish | <i>Agujon sable</i> |
| | <i>Platybelone argalus argalus</i> (LeSueur 1821) | Keeltail needlefish | <i>Agujon de quilla</i> |
| | <i>Strongylura timucu</i> (Walbaum 1792) | Timucu needlefish | <i>Agujon timuco</i> |
| | <i>Tylosurus acus acus</i> (Lacepède 1803) | Agujon needlefish | <i>Marao ojo</i> |
| | <i>Tylosurus crocodilus crocodilus</i> (Peron & LeSueur 1821) | Hound needlefish | <i>Marao lisero</i> |
| Blenniidae | <i>Entomacrodus nigricans</i> Gill 1859 | Pearl blenny | - |
| | <i>Ophioblennius atlanticus</i> (Cuvier & Valenciennes 1836) | Redlip blenny | - |
| | <i>Parablennius marmoreus</i> (Poey 1875) | Seaweed blenny | - |
| | <i>Scartella cristata</i> (Linnaeus 1758) | Molly miller | - |
| Bothidae | <i>Bothus lunatus</i> (Linnaeus 1758) | Peacock flounder | <i>Lenguado ocelado</i> |
| | <i>Bothus ocellatus</i> (Agassiz 1831) | Eyed flounder | - |
| Carangidae | * <i>Alectis ciliaris</i> (Bloch 1788) | African pomano | <i>Pampano de hebra</i> |
| | <i>Caranx bartholomaei</i> Cuvier 1833 | Yellow jack | <i>Cojínua amarilla</i> |
| | * <i>Caranx crysos</i> (Mitchill 1815) | Blue runner | <i>Cojínua negra</i> |
| | <i>Caranx latus</i> Agassiz 1831 | Horse-eye jack | <i>Jurel ojo</i> |
| | * <i>Caranx lugubris</i> Poey 1860 | Black jack | <i>Jurel negro</i> |
| | <i>Caranx ruber</i> (Bloch 1793) | Bar jack | <i>Cojínua carbonera</i> |
| | * <i>Decapterus macarellus</i> (Cuvier 1833) | Mackerel scad | <i>Macarela caballa</i> |
| | <i>Decapterus punctatus</i> (Cuvier 1829) | Round scad | <i>Macarela chuparaco</i> |
| | * <i>Elagatis bipinnulatus</i> (Quoy & Gaimard 1824) | Rainbow runner | <i>Macarela salmon</i> |
| | <i>Oligoplites saurus</i> (Bloch & Schneider 1801) | Atlantic leatherjack | <i>Zapatero sietecueros</i> |
| | <i>Selar crumenophthalmus</i> (Bloch 1793) | Bigeye scad | <i>Chicharro ojo</i> |
| | <i>Seriola dumerili</i> (Risso 1810) | Greater amberjack | <i>Medregal coronado</i> |

continued...

Table 2.2 continued

| Family | Species | English common name | Spanish common name | |
|--------------------|-------------------------------------------------------------|-----------------------------------------------------|---------------------------------|---|
| Carcharhinidae | * <i>Seniola rivoliana</i> Cuvier 1833 | Almaco jack | <i>Medregal limon</i> | |
| | <i>Trachinotus falcatus</i> (Linnaeus 1758) | Permit | <i>Pampano palometa</i> | |
| | <i>Trachinotus goodei</i> Jordan & Evermann 1896 | Palometa pompano | <i>Pampano listado</i> | |
| | <i>Carcharhinus acronotus</i> (Poey 1861) | Blacknose shark | <i>Tiburón amarillo</i> | |
| | <i>Carcharhinus falciformis</i> (Bibron 1839) | Silky shark | <i>Tiburón jaquetón</i> | |
| | <i>Carcharhinus leucas</i> (Valenciennes 1839) | Bull shark | <i>Tiburón sarda</i> | |
| | <i>Carcharhinus limbatus</i> (Valenciennes 1839) | Blacktip shark | <i>Tiburón macuira</i> | |
| | <i>Carcharhinus longimanus</i> (Lesson 1830) | Oceanic whitetip shark | <i>Tiburón oceánico</i> | |
| | <i>Carcharhinus perezii</i> (Poey 1876) | Caribbean reef shark | <i>Tiburón coralino</i> | |
| | <i>Galeocerdo cuvier</i> (LeSueur 1822) | Tiger shark | <i>Tintorera, Alecrin</i> | |
| Chaetodontidae | <i>Negaprion brevirostris</i> (Poey 1868) | Lemon shark | <i>Tiburón galano</i> | |
| | <i>Rhizoprionodon porosus</i> (Poey 1861) | Caribbean sharpnose shark | <i>Cazon playón</i> | |
| | <i>Chaetodon aculeatus</i> (Poey 1860) | Longsnout butterflyfish | - | |
| | <i>Chaetodon capistratus</i> Linnaeus 1758 | Foureye butterflyfish | - | |
| | * <i>Chaetodon ocellatus</i> Bloch 1787 | Spotfin butterflyfish | - | |
| | <i>Chaetodon sedentarius</i> Poey 1860 | Reef butterflyfish | - | |
| | <i>Chaetodon striatus</i> Linnaeus 1758 | Banded butterflyfish | - | |
| | <i>Amblycirrhitus pinos</i> (Mowbray 1927) | Redspotted hawkfish | - | |
| | Cirrhitidae | <i>Labrisomus guppyi</i> Norman | Shadow blenny | - |
| | Clinidae | <i>Labrisomus nuchipinnis</i> (Quoy & Gaimard 1824) | Hairy blenny | - |
| Clupeidae | <i>Harengula clupeiola</i> (Cuvier 1829) | False pilchard, f. herring | <i>Sardineta escamuda</i> | |
| | <i>Harengula humeralis</i> (Cuvier 1829) | Red-ear sardine | <i>Sardineta de lay</i> ** | |
| | <i>Jenkinsia lamprotaenia</i> (Gosse 1851) | Dwarf herring | <i>Sardineta canalerita</i> | |
| | <i>Opisthonema oglinum</i> (LeSueur 1818) | Atlantic thread herring | <i>Machuelo hebra atlántico</i> | |
| | * <i>Sardinella aurita</i> Valenciennes 1847 | Round sardinella | <i>Sardinella atlántica</i> | |
| Congridae | <i>Heteroconger halis</i> Böhlke 1957 | Garden eel | - | |
| Dactylopteridae | <i>Dactylopterus volitans</i> (Linnaeus 1758) | Flying gurnard | <i>Alon</i> | |
| Dasyatidae | <i>Dasyatis americana</i> Hildebrand & Schroeder 1928 | Southern stingray | <i>Raya latigo americana</i> | |
| Diodontidae | <i>Chilomycterus antennatus</i> (Cuvier 1818) | Bridled burrfish | - | |
| | * <i>Chilomycterus antillarum</i> Jordan & Rutter 1897 | Web burrfish | - | |
| | <i>Diodon holocanthus</i> Linnaeus 1758 | Spiny puffer | - | |
| | <i>Diodon hystrix</i> Linnaeus 1758 | Porcupinefish | <i>Pejerizo común</i> | |
| Elopidae | * <i>Elops saurus</i> Linnaeus 1776 | Ladyfish | <i>Malacho</i> | |
| Emmelichthyidae | <i>Inermia vittata</i> Poey 1861 | Boga | - | |
| Engraulidae | <i>Anchoa hepsetus</i> (Linnaeus 1758) | Striped anchovy | <i>Anchoa legitima</i> | |
| | <i>Anchoa lyolepis</i> (Evermann & Marsh 1902) | Dusky anchovy | <i>Anchoa trompalarga</i> | |
| Ephippidae | <i>Chaetodipterus faber</i> (Broussonet 1782) | Atlantic spadefish | <i>Paguala</i> | |
| Fistulariidae | <i>Fistularia tabacaria</i> Linnaeus 1758 | Cornetfish | - | |
| Gerreidae | <i>Eucinostomus argenteus</i> Baird & Girard 1854 | Silver mojarra; spotfin mojarra | <i>Mojarrita plateada</i> | |
| | <i>Gerres cinereus</i> (Walbaum 1792) | Yellowfin mojarra | <i>Mojarra blanca</i> | |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> (Bonnatere 1783) | Nurse shark | <i>Gata atlántica</i> | |
| Gobiidae | * <i>Bathygobius soporator</i> (Cuvier & Valenciennes 1837) | Frillfin goby | - | |

continued...

Table 2.2 continued

| Family | Species | English common name | Spanish common name |
|---------------|---------------------------------------------------------|---------------------------------|---------------------------|
| | <i>Coryphopterus glaucofraenum</i> Gill 1863 | Bridled goby | - |
| | <i>Gnatholepis thompsoni</i> Jordan 1902 | Goldspot goby | - |
| | <i>Gobiosoma evelynae</i> Böhlke & Robins 1968 | Sharknose goby | - |
| Grammidae | <i>Gramma loreto</i> Poey 1868 | Fairy basslet | - |
| | <i>Gramma melacara</i> Böhlke & Randall 1963 | Blackcap basslet | - |
| Grammistidae | <i>Rypticus saponaceus</i> (Bloch & Schneider 1801) | Greater soapfish | <i>Jabonero</i> |
| Haemulidae | <i>Anisotremus surinamensis</i> (Bloch 1791) | Black margate | <i>Burro pompon</i> |
| | <i>Anisotremus virginicus</i> (Linnaeus 1758) | Porkfish | <i>Burro catalina</i> |
| | <i>Haemulon album</i> Cuvier 1829 | White margate | <i>Ronco blanco</i> |
| | <i>Haemulon aurolineatum</i> Cuvier 1829 | Tomtate grunt | <i>Ronco jeniguano</i> |
| | * <i>Haemulon bonariense</i> Cuvier 1829 | Black grunt | <i>Ronco rayado</i> |
| | <i>Haemulon carbonarium</i> Poey 1860 | Caesar grunt | <i>Ronco carbonero</i> |
| | <i>Haemulon chrysargyreum</i> Günther 1859 | Smallmouth grunt | <i>Ronco boquilla</i> |
| | <i>Haemulon flavolineatum</i> (Desmarest 1823) | French grunt | <i>Ronco amarillo</i> |
| | <i>Haemulon macrostomum</i> Günther 1859 | Spanish grunt | <i>Ronco caco</i> |
| | * <i>Haemulon melanurum</i> (Linnaeus 1758) | Cottonwick grunt | <i>Ronco mapurite</i> |
| | <i>Haemulon parrai</i> (Desmarest 1823) | Sailor's grunt: sailor's choice | <i>Ronco plateado</i> |
| | <i>Haemulon plumieri</i> (Lacepede 1802) | White grunt | <i>Ronco margariteno</i> |
| | <i>Haemulon sciurus</i> (Shaw 1803) | Bluestriped grunt | <i>Ronco catire</i> |
| | * <i>Pomadasys corvinaeformis</i> (Steindachner 1868) | Roughneck grunt | <i>Corocoro gris</i> |
| | * <i>Pomadasys crocro</i> (Cuvier 1830) | Burro grunt | <i>Corocoro crocro</i> |
| Hemiramphidae | <i>Hemiramphus balao</i> LeSueur 1823 | Balao halfbeak | <i>Agujeta balaju</i> |
| | <i>Hemiramphus brasiliensis</i> (Linnaeus 1758) | Ballyhoo halfbeak | <i>Agujeta brasilena</i> |
| | * <i>Hyporamphus unifasciatus</i> (Ranzani 1842) | Common halfbeak | <i>Agujeta blanca</i> |
| Holocentridae | <i>Holocentrus ascensionis</i> (Osbeck 1765) | Longjaw squirrelfish | <i>Candil gallito</i> |
| | <i>Holocentrus coruscus</i> (Poey 1860) | Reef squirrelfish | - |
| | <i>Neoniphon marianus</i> (Cuvier & Valenciennes 1829) | Squirrelfish ? | - |
| | <i>Holocentrus rufus</i> (Walbaum 1792) | Longspine squirrelfish | <i>Candil soldado</i> |
| | <i>Sargocentron vexillarium</i> (Poey 1860) | Dusky squirrelfish | - |
| | <i>Myripristis jacobus</i> Cuvier 1829 | Blackbar soldierfish | <i>Candil colorado</i> |
| | * <i>Plectyprops retrospinis</i> (Guichenot 1853) | Cardinal soldierfish | - |
| Kyphosidae | <i>Kyphosus incisor</i> (Cuvier 1831) | Yellow sea chub | <i>Chopa amarilla</i> |
| | <i>Kyphosus sectatrix</i> (Linnaeus 1758) | Bermuda sea chub | <i>Chopa blanca</i> |
| Labridae | <i>Bodianus rufus</i> (Linnaeus 1758) | Spanish hogfish | <i>Vieja colorada</i> |
| | <i>Clepticus parrae</i> (Bloch & Schneider 1801) | Creole wrasse | - |
| | <i>Halichoeres bivittatus</i> (Bloch 1791) | Slippery dick | - |
| | <i>Halichoeres garnoti</i> (Cuvier & Valenciennes 1839) | Yellowhead wrasse | - |
| | <i>Halichoeres maculipinna</i> (Müller & Troschel 1848) | Clown wrasse | - |
| | <i>Halichoeres poeyi</i> (Steindachner 1867) | Black-ear wrasse | - |
| | <i>Halichoeres radiatus</i> (Linnaeus 1758) | Pudding wife (wrasse) | <i>Doncella arco-iris</i> |
| | <i>Xyrichtys novacula</i> (Linnaeus 1758) | Pearly razorfish | <i>Doncella cuchilla</i> |
| | <i>Xyrichtys splendens</i> (Castelnau 1855) | Green razorfish | - |
| | <i>Lachnolaimus maximus</i> (Walbaum 1792) | Hogfish | <i>Doncella de pluma</i> |

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Table 2.2 continued

| Family | Species | English common name | Spanish common name |
|----------------------------------------------------------|----------------------------------------------------------|------------------------------------------|---------------------------|
| Lutjanidae | <i>Thalassoma bifasciatum</i> (Bloch 1791) | Bluehead | - |
| | <i>Lutjanus analis</i> (Cuvier 1828) | Mutton snapper | <i>Pargo criollo</i> |
| | <i>Lutjanus apodus</i> (Walbaum 1792) | Schoolmaster (snapper) | <i>Pargo amarillo</i> |
| | <i>Lutjanus cyanopterus</i> (Cuvier 1828) | Cubera snapper | <i>Pargo cubera</i> |
| | <i>Lutjanus griseus</i> (Linnaeus 1758) | Gray snapper | <i>Pargo prieto</i> |
| | <i>Lutjanus jocu</i> (Bloch & Schneider 1801) | Dog snapper | <i>Pargo jocu</i> |
| | <i>Lutjanus mahagoni</i> (Cuvier 1828) | Mahogany snapper | <i>Pargo ojo</i> |
| | <i>Lutjanus synagris</i> (Linnaeus 1758) | Lane snapper | <i>Pargo biajaiba</i> |
| | <i>Ocyurus chrysurus</i> (Bloch 1791) | Yellowtail snapper | <i>Rabirubia</i> |
| | Malacanthidae | <i>Malacanthus plumieri</i> (Bloch 1786) | Sand tilefish |
| <i>Tarpon atlanticus</i> (Valenciennes 1846) | | Tarpon | <i>Tarpon</i> |
| Megalopidae | <i>Aluterus schoepfii</i> (Walbaum 1792) | Orange filefish | <i>Cachua perra</i> |
| Monacanthidae | <i>Aluterus scripta</i> (Osbeck 1765) | Scrawled filefish | - |
| | <i>Cantherines macroceros</i> (Hollard 1854) | Whitespotted filefish | - |
| | <i>Cantherines pullus</i> (Ranzani 1842) | Orangespotted filefish | <i>Lija pintada</i> |
| | <i>Monacanthus ciliatus</i> (Mitchill 1818) | Fringed filefish | <i>Lija de clavo</i> |
| | * <i>Monacanthus tuckeri</i> Bean 1906 | Slender filefish | - |
| | * <i>Stefanolepis setifer</i> (Bennett 1830) | Pygmy filefish | <i>Lija de hebra</i> |
| | <i>Mugil curema</i> Valenciennes 1836 | White mullet | <i>Lisa criolla</i> |
| | <i>Mulloidichthys martinicus</i> (Cuvier 1829) | Yellow goatfish | <i>Salmonete amarillo</i> |
| | <i>Pseudupeneus maculatus</i> (Bloch 1793) | Spotted goatfish | <i>Salmonete manchado</i> |
| | Muraenidae | <i>Echidna catenata</i> (Bloch 1795) | Chain moray |
| * <i>Enchelycore nigricans</i> (Bonnaterre 1788) | | Viper moray | <i>Morena negra</i> |
| * <i>Gymnothorax funebris</i> (Ranzani 1840) | | Green moray | <i>Morena congrio</i> |
| <i>Lycodontis moringa</i> (Cuvier 1829) | | Spotted moray | <i>Morena pintada</i> |
| <i>Gymnothorax vicinus</i> (Castelnaud 1855) | | Purplemouth moray | <i>Morena amarilla</i> |
| * <i>Gymnothorax miliaris</i> (Kaup 1856) | | Goldentail moray | <i>Morena dorada</i> |
| <i>Aetobatus narinari</i> (Euphrasen 1790) | | Spotted eagle ray | <i>Chucho pintado</i> |
| <i>Ogcocephalus nasutus</i> (Cuvier & Valenciennes 1837) | | Redbellied batfish | - |
| <i>Myrichthys breviceps</i> (Gronow 1854) | | Sharptail eel | - |
| <i>Myrichthys ocellatus</i> (Kaup 1856) | | Goldspotted snake eel | - |
| Ophichthidae | <i>Ophichthus ophis</i> (Linnaeus 1758) | Spotted snake eel | <i>Tieso pintado</i> |
| | <i>Opisthognathus aurifrons</i> (Jordan & Thompson 1905) | Yellowhead jawfish | - |
| | * <i>Opisthognathus macrogathus</i> Poey 1860 | Longjaw jawfish | - |
| | <i>Opisthognathus maxillosus</i> Poey 1860 | Mottled jawfish | - |
| | <i>Opisthognathus whitehurstii</i> (Longley 1931) | Dusky jawfish | - |
| | <i>Acanthostracion polygonius</i> Poey 1876 | Honeycomb cowfish | - |
| | <i>Acanthostracion quadricornis</i> (Linnaeus 1758) | Scrawled cowfish | <i>Torito azul</i> |
| | <i>Lactophrys trigonus</i> (Linnaeus 1758) | Buffalo trunkfish | <i>Chapin bufalo</i> |
| | <i>Lactophrys bicaudalis</i> (Linnaeus 1758) | Spotted trunkfish | <i>Chapin pintado</i> |
| | <i>Lactophrys triqueter</i> (Linnaeus 1758) | Smooth trunkfish | <i>Chapin baqueta</i> |
| Pempheridae | <i>Pempheris schomburgki</i> Müller & Troschel 1848 | Copper sweeper | - |
| | * <i>Pempheris poeyi</i> Bean 1885 | Shortfin sweeper | - |

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Table 2.2 continued

| Family | Species | English common name | Spanish common name | |
|-------------------------------------------------|--------------------------------------------------------------------------|------------------------------------------------------------------|------------------------------|-------------------------|
| Pomacanthidae | <i>Centropyge argi</i> Woods & Kanazawa 1951 | Cherubfish | - | |
| | <i>Holacanthus ciliaris</i> (Linnaeus 1758) | Queen angelfish | <i>Isabelita patale</i> | |
| | <i>Holacanthus tricolor</i> (Bloch 1795) | Rock beauty | <i>Isabelita medioluto</i> | |
| | <i>Pomacanthus arcuatus</i> (Linnaeus 1758) | Gray angelfish | <i>Cachama blanca</i> | |
| Pomacentridae | <i>Pomacanthus paru</i> (Bloch 1787) | French angelfish | <i>Cachama negra</i> | |
| | <i>Abudefduf saxatilis</i> (Linnaeus 1758) | Sergeant major | <i>Petaca rayada</i> | |
| | <i>Abudefduf taurus</i> (Müller & Troschel 1848) | Night sergeant | <i>Petaca rezobada</i> | |
| | <i>Chromis cyanea</i> (Poey 1860) | Blue chromis | - | |
| | <i>Chromis multilineata</i> (Guichenot 1853) | Brown chromis | <i>Jaqueta parda</i> | |
| | <i>Microspathodon chrysurus</i> (Cuvier & Valenciennes 1830) | Yellowtail damselfish | <i>Jaqueta rabo amarillo</i> | |
| | <i>Stegastes fuscus</i> (Cuvier 1830) | Dusky damselfish | - | |
| | <i>Stegastes leucostictus</i> (Müller & Troschel 1848) | Beau gregory | - | |
| | <i>Stegastes planifrons</i> (Cuvier 1830) | Yellow damselfish | - | |
| | <i>Stegastes variabilis</i> (Castelnau 1855) | Cocoa damselfish | - | |
| | Priacanthidae | <i>Priacanthus arenatus</i> Cuvier in Cuvier & Valenciennes 1829 | Atlantic bigeye | <i>Catalufa toro</i> |
| | | <i>Heteropriacanthus cruentatus</i> (Lacepède 1802) | Glasseye | <i>Catalufa de roca</i> |
| Rachycentridae | * <i>Rachycentron canadum</i> (Linnaeus 1766) | Cobia | <i>Cobie</i> | |
| Scaridae | <i>Scarus coelestinus</i> Valenciennes in Cuvier & Valenciennes 1839 | Midnight parrotfish | <i>Loro negro</i> | |
| | * <i>Scarus coeruleus</i> (Bloch 1786) | Blue parrotfish | <i>Loro azul</i> | |
| | <i>Scarus iserti</i> Bloch 1790 | Striped parrotfish | <i>Loro rayado</i> | |
| | <i>Scarus guacamaia</i> Cuvier 1831 | Rainbow parrotfish | <i>Loro guacamayo</i> | |
| | <i>Scarus taeniopterus</i> Desmarest 1831 | Princess parrotfish | <i>Loro listado</i> | |
| | <i>Scarus vetula</i> Bloch & Schneider 1801 | Queen parrotfish | <i>Loro perico</i> | |
| | <i>Sparisoma aurofrenatum</i> Valenciennes in Cuvier & Valenciennes 1839 | Redband parrotfish | <i>Loro manchado</i> | |
| | <i>Sparisoma chrysopterygum</i> (Bloch & Schneider 1801) | Redtail parrotfish | <i>Loro verde</i> | |
| | <i>Sparisoma radians</i> Valenciennes in Cuvier & Valenciennes 1839 | Bucktooth parrotfish | - | |
| | <i>Sparisoma rubripinne</i> Valenciennes in Cuvier & Valenciennes 1839 | Redfin parrotfish | <i>Loro basto</i> | |
| | <i>Sparisoma viride</i> (Bonnaterre 1788) | Stoplight parrotfish | <i>Loro viejo</i> | |
| | Sciaenidae | <i>Equetus lanceolatus</i> (Linnaeus 1758) | Jackknife fish | - |
| | | <i>Equetus punctatus</i> (Bloch & Schneider 1801) | Spotted drum | - |
| | | <i>Odontoscion dentex</i> (Cuvier 1830) | Reef croaker | <i>Bombache de roca</i> |
| | Scombridae | <i>Pareques acuminatus</i> (Bloch & Schneider 1801) | Cubby | - |
| <i>Euthynnus alletteratus</i> (Rafinesque 1810) | | Little tunny | <i>Bacoreta</i> | |
| <i>Scomberomorus cavalla</i> (Cuvier 1829) | | King mackerel | <i>Carite lucio</i> | |
| <i>Scomberomorus regalis</i> (Bloch 1793) | | Cero | <i>Carite chinigua</i> | |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> Cuvier 1829 | Barbfish | - | |
| | <i>Scorpaena grandicornis</i> Cuvier 1829 | Grass scorpionfish | - | |
| | <i>Scorpaena inermis</i> Cuvier 1829 | Mushroom scorpionfish | - | |
| | <i>Scorpaena plumieri</i> Bloch 1789 | Spotted scorpionfish | <i>Rascacio negro</i> | |
| | <i>Scorpaenodes caribbaeus</i> Meek & Hildebrand 1928 | Reef scorpionfish | - | |
| Serranidae | <i>Alphestes afer</i> (Bloch 1793) | Mutton hamlet | <i>Guaseta</i> | |
| | <i>Cephalopholis cruentata</i> Lacepède 1802 | Grasby (seabass) | <i>Cherna enjambre</i> | |
| | <i>Cephalopholis fulva</i> (Linnaeus 1758) | Coney (seabass) | <i>Cherna cabrilla</i> | |

continued...

Table 2.2 continued

| Family | Species | English common name | Spanish common name |
|----------------|-------------------------------------------------------|---------------------------|---------------------------|
| | * <i>Diplectrum formosum</i> (Linnaeus 1758) | Sand seabass; sand perch | <i>Serrano arenero</i> |
| | <i>Epinephelus adscensionis</i> (Osbeck 1771) | Rock hind | <i>Mero cabrilla</i> |
| | <i>Epinephelus guttatus</i> (Linnaeus 1758) | Red hind | <i>Mero colorado</i> |
| | <i>Epinephelus itajara</i> (Lichtenstein 1822) | Giant grouper; jewfish | <i>Mero guasa</i> |
| | * <i>Epinephelus morio</i> (Valenciennes 1828) | Red grouper | <i>Mero americano</i> |
| | <i>Epinephelus striatus</i> (Bloch 1792) | Nassau grouper | <i>Cherna criolla</i> |
| | <i>Hypoplectrus aberrans</i> (Poey 1868) | Yellowbellied hamlet | - |
| | <i>Hypoplectrus chlorurus</i> (Valenciennes 1828) | Yellowtail hamlet | - |
| | <i>Hypoplectrus nigricans</i> (Poey 1852) | Black hamlet | - |
| | <i>Hypoplectrus puella</i> (Cuvier 1828) | Barred hamlet | - |
| | <i>Mycteroperca bonaci</i> (Poey 1861) | Black grouper | <i>Cuna bonaci</i> |
| | * <i>Mycteroperca interstitialis</i> (Poey 1861) | Yellowmouth grouper | <i>Cuna amarilla</i> |
| | * <i>Mycteroperca rubra</i> (Bloch 1793) | Comb grouper | <i>Cuna negra</i> |
| | <i>Mycteroperca tigris</i> (Valenciennes 1833) | Tiger grouper | <i>Cuna gata</i> |
| | <i>Mycteroperca venenosa</i> (Linnaeus 1758) | Yellowfin grouper | <i>Cuna de piedra</i> |
| | <i>Paranthias furcifer</i> (Valenciennes 1828) | Creole fish | <i>Cuna lucero</i> |
| | * <i>Serranus tabacarius</i> (Cuvier 1829) | Tobacco fish | - |
| | <i>Serranus tigrinus</i> (Bloch 1790) | Harlequin bass | - |
| | * <i>Serranus tortugarum</i> Longley 1935 | Chalk bass | - |
| Sparidae | <i>Archosargus rhomboidalis</i> (Linnaeus 1758) | Western Atlantic seabream | <i>Sargo amarillo</i> |
| | <i>Calamus bajonado</i> (Bloch & Schneider 1801) | Jolthead porgy | <i>Pluma bajonado</i> |
| | <i>Calamus calamus</i> (Valenciennes 1830) | Saucereye porgy | <i>Pluma calamo</i> |
| | <i>Calamus pennatula</i> Guichenot 1868 | Pluma porgy | <i>Pluma plumilla</i> |
| | <i>Diplodus argentus caudimacula</i> (Poey 1861) | Silver porgy | <i>Sargo fino</i> |
| Sphyraenidae | <i>Sphyraena barracuda</i> (Walbaum 1792) | Great barracuda | <i>Picuda barracuda</i> |
| | <i>Sphyraena picudilla</i> (Poey 1860) | Southern sennet | <i>Picuda china</i> |
| Sphyrnidae | <i>Sphyrna lewini</i> (Cuvier, Griffith & Smith 1834) | Scalloped hammerhead | <i>Cornuda comun</i> |
| | <i>Sphyrna tiburo</i> (Linnaeus 1758) | Bonnethead | <i>Cornuda de corona</i> |
| Synodontidae | <i>Synodus foetens</i> (Linnaeus 1766) | Galliwasp | - |
| | <i>Synodus intermedius</i> (Agassiz 18??) | Sand diver | - |
| | <i>Synodus synodus</i> (Linnaeus 1758) | Rockspear | - |
| Tetraodontidae | <i>Canthigaster rostrata</i> (Bloch 1782) | Sharpnose puffer | - |
| | <i>Sphoeroides spengleri</i> (Bloch 1785) | Bandtail puffer | <i>Tamboril collarete</i> |
| Triakidae | <i>Mustelus canis</i> (Mitchill 1815) | Smooth dogfish | <i>Musola dentuda</i> |

Table 2.3. Nonfish taxa included in the ECOPATH II models of a Caribbean coral reef (from stomach content analyses of West Indian reef fishes by Randall 1967).
D = detritus, A = autotroph, I = invertebrate, R = reptile, B = bird

| ID | | Taxon | Remarks |
|----|----|--------------------|---------------------------------------------------------------------------------------------------|
| D | 1 | Detritus | Dead particulate and dissolved organic matter |
| A | 1 | Benthic algae | See section 2.4.2 and Appendix Table 8.5.1 |
| A | 2 | Spermatophytes | See Appendix Table 8.5.1 |
| A | 3 | Phytoplankton | Pelagic primary producers |
| I | 1 | Microfauna | Bacteria, Foraminifera |
| I | 2 | Zooplankton | See Appendix Table 8.5.3 |
| I | 3 | Sponges | See Appendix Table 8.5.2 |
| I | 4 | Fire corals | Millepora, Hydrozoa (see section 2.4.2 and Appendix Table 8.5.2) |
| I | 5 | Sea fans | Gorgonacea, Octocorallia, Anthozoa (see section 2.4.2 and Appendix Table 8.5.2) |
| I | 6 | Sea anemones | Actiniaria and Zoanthidea, Hexacorallia, Anthozoa (see section 2.4.2 and Appendix Table 8.5.2) |
| I | 7 | Stony corals | Scleractinia, Octocorallia, Anthozoa (see section 2.4.2 and Appendix Table 8.5.2) |
| I | 8 | Bryozoans | See Appendix Table 8.5.2 |
| I | 9 | Sipunculids | Worms (see Appendix Table 8.5.2) |
| I | 10 | Priapuloids | Worms (see Appendix Table 8.5.2) |
| I | 11 | Chitons | Mollusca (see Appendix Table 8.5.2) |
| I | 12 | Gastropods | Mollusca (see Appendix Table 8.5.2) |
| I | 13 | Bivalves | Mollusca (see Appendix Table 8.5.2) |
| I | 14 | Scaphopods | Mollusca (see Appendix Table 8.5.2) |
| I | 15 | Squids | Mollusca (see Appendix Table 8.5.2) |
| I | 16 | Octopuses | Mollusca (see Appendix Table 8.5.2) |
| I | 17 | Polychaetes | Worms (see Appendix Table 8.5.2) |
| I | 18 | Echiuroids | Worms (see Appendix Table 8.5.2) |
| I | 19 | Pycnogonids | Arthropoda, Arachnidea (see Appendix Table 8.5.2) |
| I | 20 | Barnacles | Cirripedia, Crustacea (see Appendix Table 8.5.2) |
| I | 21 | Stomatopods | Crustacea (see Appendix Table 8.5.2) |
| I | 22 | Amphipods | Crustacea (see Appendix Table 8.5.2) |
| I | 23 | Tanaids | Crustacea (see Appendix Table 8.5.2) |
| I | 24 | Isopods | Crustacea (see Appendix Table 8.5.2) |
| I | 25 | Shrimps | Natantia, Crustacea (see Appendix Table 8.5.2) |
| I | 26 | Spiny lobsters | Panuliridae, Crustacea (see Appendix Table 8.5.2) |
| I | 27 | Scyllarid lobsters | Crustacea (see Appendix Table 8.5.2) |
| I | 28 | Hermit crabs | Diogenidae, Crustacea (see Appendix Table 8.5.2) |
| I | 29 | Crabs | Reptantia, Crustacea (see Appendix Table 8.5.2) |
| I | 30 | Hemichordates | Wormlikes (see Appendix Table 8.5.2) |
| I | 31 | Starfish | Asteroidea, Echinodermata (see Appendix Table 8.5.2) |
| I | 32 | Brittle stars | Ophiuroidea, Echinodermata (see Appendix Table 8.5.2) |
| I | 33 | Sea urchins | Echinoidea, Echinodermata (see Appendix Table 8.5.2) |
| I | 34 | Sea cucumbers | Holothuroidea, Echinodermata (see Appendix Table 8.5.2) |
| I | 35 | Ascidians | Tunicata, Chordata (see Appendix Table 8.5.2) |
| R | 1 | Sea turtles | See Appendix Table 8.5.2 |
| B | 1 | Sea birds | See Polovina (1984b) |

A complete list of reported nonfish food items from stomachs of West Indian reef fishes is to be found in Appendix 5.1 for benthic algae and spermatophytes; in Appendix 5.2 for invertebrates and sea turtles; and in Appendix 5.3 for items that are part of the “zooplankton” group.

2.5. Fishing Activities

To simplify the modelling work and to reduce data requirements, an unexploited system was assumed. Since data from a large geographical range had to be used, especially for the nonfish taxa, the resulting model largely reflects a “virtual” system the description of which would not

necessarily be improved by inclusion of fisheries catch data from a specific area. Moreover, even if this had have been attempted, it would have been extremely difficult to obtain catch data for individual species - except for certain commercial species of the Serranidae and Lutjanidae families.

The ecosystem described here can thus be seen as representing unfished areas of the PRVI shelf and may therefore be directly compared with the unexploited French Frigate Shoals reef ecosystem described by Polovina (1984a,b,c, 1985, 1986) and Polovina and Ow (1983).

2.6. Methods and Data Used for the Computation of Input Parameters Required for ECOPATH II

2.6.1. Growth Parameter Estimation

For the computation of several input parameters of ECOPATH II (e.g., Q/B, M), growth parameters had to be estimated for most of the fish and invertebrates included in the model. Growth of fish was generally assumed to follow the von Bertalanffy growth function (VBGF) expressed for growth in length as:

$$L_t = L_\infty \left(1 - e^{-K(t-t_0)} \right) \quad \dots 2.1)$$

where L_t is the length at age t , L_∞ the asymptotic length, K is the growth coefficient (year⁻¹) and t_0 the theoretical "age" at length zero (Pauly 1984). Growth in weight is defined as:

$$W_t = W_\infty \left(1 - e^{-K(t-t_0)} \right)^b \quad \dots 2.2)$$

where W_t is the weight at age t , W_∞ the asymptotic weight and b is the exponent of a length-weight relationship of the form:

$$W = aL^b \quad \dots 2.3)$$

When only one pair of L , W value was available, the value of b was set equal to 3, and the relationship was used to estimate the multiplicative factor required in equation 2.3.

$$a = W / L^3 \quad \dots 2.4)$$

2.6.1.1. ASYMPTOTIC SIZES

For the majority of fish species considered in the model, size-at-age or length-frequency data were not available from which L_∞ or W_∞ could be estimated. Therefore, asymptotic sizes were estimated from the reported maximum sizes (L_{\max} or W_{\max}) using the relationships (Pauly 1984):

$$L_{\max} / 0.95 \approx L_{\infty} \quad \dots 2.5)$$

$$W_{\max} / 0.86 \approx W_{\infty} \quad \dots 2.6)$$

2.6.1.2. GROWTH COEFFICIENT K

Estimates of the growth coefficient K of the VBGF were obtained directly from published sources or via the equation

$$\log_{10} K = \phi' - 2\log_{10} L_{\infty} \quad \dots 2.7)$$

for length and

$$\log_{10} K = \phi - 0.67\log_{10} W_{\infty} \quad \dots 2.8)$$

for weight. The required estimates of ϕ' and ϕ were generally obtained from the means of several values of

$$\phi' = \log_{10} K + 2\log_{10} L_{\infty}, \text{ and} \quad \dots 2.9)$$

$$\phi = \log_{10} K + 0.67\log_{10} W_{\infty} \quad \dots 2.10)$$

where K, L_{∞} and W_{∞} are growth parameter estimates for various populations of the same species (Pauly 1979a; Pauly and Munro 1984; Pauly 1985). This approach which, strictly speaking, should be used only within species was used here between species (generally of the same genus) when these were similar in shape and habits.

2.6.1.3. ESTIMATION OF t_0

For all cases considered here, rough estimates of t_0 were obtained via an empirical model by Pauly (1979a) of the form:

$$\log_{10}(-t_0) \approx -0.39 - 0.28\log_{10} L_{\infty} - 1.04\log_{10} K \quad \dots 2.11)$$

with L_{∞} as the asymptotic length in cm (total length) and K the growth coefficient of the species considered, expressed in year⁻¹. This parameter has little influence on the estimation of Q/B (see Pauly 1986), the only use for which it was required here.

2.6.2. Habitat and Abundances

2.6.2.1. FISH SPECIES

Ecological information on feeding areas, shelter, time of activity, schooling behavior, etc., as well as information on abundance of Caribbean reef fishes were assembled from various

published sources (e.g., Randall 1967, 1968; Jacobsen and Browder 1987). An attempt was made to classify information on spatial occurrence according to the subsystems defined in Fig. 2.1. Qualitative abundances were grouped applying the scale presented in Table 2.4.

Category "E" was considered here because some of the species for which growth data and/or other information were available, while not occurring in the PRVI area, do occur in other

Table 2.4. Qualitative abundance levels as used to estimate average density (Nm^{-2}) of fish species included in the models.^a

| Abundance level | Remarks | Nm^{-2} |
|-----------------|-------------------------------|------------------|
| V Very abundant | Usually seen in large numbers | 5.00000 |
| A Abundant | Usually seen in some numbers | 0.50000 |
| C Common | Usually seen | 0.05000 |
| F Fairly common | Seen about half of the time | 0.00500 |
| O Occasional | Usually not seen | 0.00050 |
| R Rare | Very unlikely | 0.00005 |
| E Absent | (see text) | 0.00000 |

^aAs assessed following comparison of various scales (see text).

areas of the Caribbean and thus might be useful for others who might wish to construct models of other areas of the Caribbean.

To reexpress qualitative statements about abundances into numerical values, the following procedure was applied: A series of six published sets of numerical abundances from visual

censuses and from rotenone stations carried out in the PRVI area (Jacobsen and Browder 1987) was transformed and ranked. Every list was treated separately since many fish species occurred in several lists. Furthermore, it was important to check whether the whole range of abundances was covered by the majority of the sets, and whether their respective rankings differed markedly. A parallel listing of the sets revealed, when a \log_{10} scale was applied, that the whole range of abundance values (from $1.3 \cdot 10^{-6} \text{ Nm}^{-2}$ to $8.4 \cdot 10^{-1} \text{ Nm}^{-2}$) could be covered by six levels of abundance. The last column in Table 2.4 presents an average density value in Nm^{-2} for each abundance level. This average density was computed as the mean of the corresponding subrange. Density values refer to an average m^2 for the five subsystems considered here.

Qualitative abundances from several sources (e.g., Clavijo et al. 1980; Randall 1983; Jacobsen and Browder 1987) were transferred one level downward so they would correspond to the new scale, as suggested by a comparison of visual census figures with qualitative abundances data, for a variety of species for which both estimates were available. This transfer had the disadvantage that the qualitative definitions of the different abundance levels did not completely overlap. However, for the present application, it appeared more important to obtain a consistent scale for all data sets.

When several estimates of abundance were available for the same species, a selection was made according to three hierarchical levels:

- data (individuals m^{-2}) from visual censuses or rotenone stations, by subsystems;
- qualitative abundances from tables in Jacobsen and Browder (1987) based on published visual census figures; and
- qualitative abundances derived from general descriptions of the biology of the species in question.

For species for which no indication of abundance could be found, abundance was assumed equal to the average for species with similar characteristics.

The list thus prepared, with adjusted qualitative abundances for every species and every subsystem, was used to obtain numerical values by substituting qualitative abundances by its numerical average range value in Nm^{-2} , divided by four. Then, the sum of four subsystems

(seagrass beds and algal mats were combined here in one subsystem) was computed by simple addition of estimates for a species across subsystems.

2.6.2.2. NONFISH TAXA

For nonfish taxa, even rough qualitative estimates of abundance are scarce to nonexistent in coral reef systems. Although some investigations have been carried out in the PRVI reef area, they concentrated mostly on commercially exploited species, especially conchs and lobster. The limited information on abundance of nonfish taxa that were identified in the course of this work may be found in Table 3.9 in the "Results" section.

2.6.3. Biomass

2.6.3.1. FISH SPECIES

The biomass in g per m² wet weight (gm⁻²WW) of each fish species was calculated by multiplying the average numerical abundance obtained as described above with the mean individual weight of the corresponding species. Mean individual weight (W_{mean}) was obtained through

$$W_{\text{mean}} = W_{\infty} * 0.3 * 0.86 \quad \dots 2.12)$$

with W_{∞} being asymptotic weight in gWW and assuming that the mean weight corresponds roughly to 30% of maximum weight, itself about 86% of the asymptotic weight (see equation 2.6). Note that this definition of W_{mean} implies that the bulk of the biomass is comprised by late juveniles and young adults.

2.6.3.2. NONFISH TAXA

Many of the biomass values for nonfish taxa were indicated in the corresponding references in units other than wet (=live) weight. A summary of the conversion factors used for the computation of biomass and P/B values for nonfish taxa is given in Table 2.5.

2.6.3.3. DETRITUS BIOMASS

The following empirical equation was derived by Pauly et al. (1993a, Appendix IV) to obtain rough estimates of detritus biomass (D , gCm⁻²) as a function of primary production (PP , gCm⁻² year⁻¹) and euphotic depth (E , in m):

$$\log_{10} D = -2.41 + 0.954 \log_{10} PP + 0.863 \log_{10} E \quad \dots 2.13)$$

Fifty-two percent of the variation ($R = 0.718$) in the data set was explained by equation (2.12), so the fit is not tight. However, the *biomass* of detritus (as opposed to the *flows* in and out of this box) is not an important parameter and hence the low precision of estimates for (2.13) may be acceptable here.

Table 2.5. Conversion factors used for ECOPATH II input parameter estimation of nonfish groups.

| Taxon | Ratio | % ^a | Source |
|-------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------|-----------------|-------------------------------------------------|
| Organic tissue | 1 kcal = 1 g WW | | Steele (1974) |
| Organic tissue | 1 g C = 10 g WW | 10 | Steele (1974) |
| Organic tissue | 1gDW=4.5 gWW | 20-25 | Steele (1974) |
| Plants | 1 g C = 40 kJ | - | G. Graf, IfM, Kiel (pers. comm.) |
| Plants | 1gCm ² ·day ⁻¹ = 1gO ₂ m ⁻² ·hour ⁻¹ *0.3 | | Connor and Adey (1977) |
| Primary producers | 1 g C = 3.33 g DW | 30 ^b | G. Graf and D. Schramm, IfM, Kiel (pers. comm.) |
| Algae, seagrasses, phytoplankton | 1 g C = 16.7 g WW | 6 | G. Graf and D. Schramm, IfM, Kiel (pers. comm.) |
| Algae, seagrasses, phytoplankton | 1 g DW = 5.71 g WW | 15-20 | G. Graf and D. Schramm, IfM, Kiel (pers. comm.) |
| Foraminifera | 1 g AFDW = 1.16 g WW | 86.5 | Odum and Odum (1955) |
| Animals | 1 g C = 48 kJ | | G. Graf, IfM, Kiel (pers. comm.) |
| Sponges | 1 g DW = 3.33 g WW | 30 | D. Barthel, IfM, Kiel (pers. comm.) |
| Sea anemones | 1 g DW = 5.88 g WW | 83 | Pauly et al. (1993b) |
| Gastropods (mean of several species) | 1 g DW = 5.56 g WW | 18 | Pauly et al. (1993b) |
| Gastropods (mean of several species) | 1 g DW = 18.6 kJ | | Pauly et al. (1993b) |
| Bivalves (mean of several species) | 1 g DW = 10 g WW | 10 | Pauly et al. (1993b) |
| Polychaetes (mean of several species) | 1 g DW = 5 g WW | 20 | Pauly et al. (1993b) |
| Cirripedia (two species) | 1 g DW = 5.3 kJ | | Berry (1982) |
| Amphipods (one species) | 1 g DW = 4 g WW | 26 | Pauly et al. (1993b) |
| Euphausiids (<i>E. superba</i>) | 1 g DW = 5 g WW | 19 | Pauly et al. (1993b) |
| Isopods (1 species) | 1 g DW = 4 g WW | 25 | Pauly et al. (1993b) |
| Shrimps (2 species) | 1 g DW = 3.8 g WW | 26.7 | Pauly et al. (1993b) |
| Small crustaceans (mean of euphausiids, amphipods, isopods and shrimps) | 1 g DW = 4 g WW | 25 | Pauly et al. (1993b) |
| Spiny lobster (1 species) | 1 g DW = 5.55 g WW | 18 | Smale (1978) |
| Starfish (2 species) | 1 g DW = 3.5 g WW | 29 | Pauly et al. (1993b) |
| Sea urchins (1 species) | 1 g DW = 3.1 g WW | 32 | Pauly et al. (1993b) |
| Sea urchins (1 species) | 1 kJ = 3.7 g DW | | Pauly et al. (1993b) |
| Sea urchins (1 species) | 1 kJ = 1.2 g WW | | Pauly et al. (1993b) |
| Sea cucumbers (3 species) | 1 g DW = 11.1 g WW | 9 | Pauly et al. (1993b) |
| Sea cucumbers (3 species) | 1 kJ = 1.08 g WW | | Pauly et al. (1993b) |
| Echinoderms (mean of several taxa) | 1 g DW = 4 g WW | 25 | Pauly et al. (1993b) |
| Shell organics | 1 g DW = 23.9 kJ | | Horn (1986) |

Notes:

^aC or DW expressed as percent of WW.^bC or DW expressed as percent of DW.

Source: AFDW = ash free, C = carbon, DW = dry weight, WW = wet (live) weight.

2.6.4. Environmental Temperature

Mean annual temperature was determined from hydrographical charts (DHI 1969) to be 28°C for the waters around the PRVI area. An estimate of mean annual temperature was required to compute natural mortality *M* and food consumption (*Q/B*) from empirical models (see below). This value of 28°C agrees closely with temperatures indicated by Randall (1962a) for two different study sites in the US Virgin Islands.

2.6.5. Production/Biomass Ratio

Under certain conditions mentioned above, the P/B ratio is equal to total mortality Z or natural mortality M (in case of no fishing) of a population of fish or invertebrates. Given the assumption of zero fishing mortality made here, the literature was searched for published values of M (or of Z in unexploited systems).

When natural mortality (or Z) was indicated as a percentage, or as percentage survival, the estimate was transformed into an exponential rate based on:

$$N_t = N_0 * e^{-Mt} \quad \dots 2.14)$$

where N_t is the number of fish in a given cohort at time t and N_0 is the initial number of fish.

In cases where no values of M for fishes or invertebrates with high metabolic rate could be found, this parameter was estimated by means of an empirical model derived by Pauly (1980), of the form

$$\log_{10} M = -0.0066 - 0.279 \log_{10} L_{\infty} + 0.6543 \log_{10} K + 0.463 \log_{10} T \quad \dots 2.15)$$

and

$$\log_{10} M = -0.2107 - 0.0824 \log_{10} W_{\infty} + 0.6757 \log_{10} K + 0.4687 \log_{10} T \quad \dots 2.16)$$

where L_{∞} and W_{∞} refer to asymptotic length in cm and asymptotic weight in g, respectively, K refers to year⁻¹ and T is the mean annual temperature in °C of the water in which the stock in question lives (i.e., 28°C, see above). Estimates of natural mortality M for clupeoid species obtained via equation (2.15) or (2.16) were corrected with $M * 0.6$ as suggested by Pauly (1980). In the case of the low-metabolism echinoderms (sea urchins, holothuroids), M was assumed to be lower than implied by equation (2.15) or (2.16), i.e., about equal to K (as was demonstrated to be appropriate for tropical sea urchins by Longhurst and Pauly 1987).

2.6.6. Food Consumption per Unit Biomass (Q/B)

One of the key input parameters in ECOPATH I as well as in ECOPATH II is the food consumption of a species (group). Polovina (1984a,b) called this input the “food required” (FR). Pauly (1986), instead, used the expression “food consumption per unit biomass” (Q/B) and defined it as the number of times an age-structured population of fish consumes its own weight per year. In the forthcoming, Q/B shall be used *sensu* Pauly.

Published data on food consumption of tropical reef fishes and invertebrates are scarce. Thus, Polovina (1984a,b,c), in the absence of appropriate data, used estimates of consumption by Pacific salmon for the reef fishes in the ECOPATH modelling of French Frigate Shoals, Northern Hawaiian Islands. Furthermore, all reef invertebrates, except squids, lobsters and crabs, were lumped together as “heterotrophic benthos” with one common estimate of Q/B. Menzel (1958,1960) was one of the firsts to carry out food conversion experiments on coral reef fishes using two species from Bermuda, the herbivorous *Holacanthus bermudensis* (Pomacanthidae) and the carnivorous *Epinephelus guttatus* (Serranidae) for his experiments.

To date a series of methods exists for estimating the quantity of food eaten by a fish population during a certain period of time (e.g., Bajkov 1935; Beverton and Holt 1957; Ursin 1967; Elliot and Persson 1978; Jobling 1981; Daan 1973, 1983; Andersen 1982; Armstrong et al. 1983; Durbin et al. 1983; Rice et al. 1983; Stewart et al. 1983; Majkowski and Hearn 1984; and Pennington 1984). Most of the available methods are based on combining stomach contents data obtained from the field with experimental results. These parameters then allow for the determination of transition rates for stomach contents. Such investigations require a great number of stomachs because of the generally high variabilities of stomach contents in natural fish populations. This requirement is generally difficult to meet with regard to coral reef fishes which abundances are often rather low. In addition to this, reefs are unsuited for trawl fishing for obvious reasons and, instead, fishes on reefs and in adjacent areas are caught by a wide variety of gears from traps to hook and lines. Therefore, catches and hence stomach contents are usually not representative. Fishes, when trap-caught, may consume prey usually unavailable to them. Also the mean weight of the food in the stomach of fishes caught with hook and lines, tends to be underestimated because generally "hungry" fish are caught.

Another approach to estimate the food consumption is by executing laboratory or pond feeding experiments. According to Mann (1978), extrapolations of food consumption estimates from laboratory experiments to natural conditions can be divided into direct and indirect methods. The direct methods attempt to simulate natural conditions in the laboratory by imitating environmental variables such as temperature, salinity, food type, availability of food, water movements, diel cycles, etc., as well as the growth rates of fish occurring in the field. Because of practical difficulties in meeting all of these requirements, indirect methods have evolved. Indirect estimates usually use nitrogen and/or energy budgets as a basis for the prediction of food requirements of fish under a range of conditions (Ivlev 1939, 1961; Winberg 1956; Gerking 1962). Although more assumptions than in the direct method are involved, the indirect approach leaves more space for the incorporation of the effects of environmental variables. A detailed discussion of the subject can be found in Mann (1978).

2.6.6.1. THE FOOD CONSUMPTION MODEL

As a contribution to the indirect approach, an analytical method for the estimation of Q/B has been developed by Pauly (1986) which may be applied in cases where stomach analyses from field samples cannot be carried out. The method links up estimates of gross food conversion efficiency, obtained from laboratory experiments, with field data on growth. Thus, it aims to make the best use of available data from laboratory or pond experiments without the need for extensive field sampling of stomachs. By combining experimental and field data, this model reduces the effects of two sources of bias from experimental data on food conversion: (1) experimental fish are most often given a higher ration than is available to them in nature (Windell 1978); and (2) captive fish are usually stressed and therefore have lower conversion efficiencies than in nature (Pauly 1986).

Pauly's model assumes that the population under consideration is in a steady state and that the fish (or invertebrate) grows according to the von Bertalanffy theory of growth (von Bertalanffy 1934, 1938, 1951; Pauly 1984). Note that these assumptions are also inherent to the reef ecosystem model developed here.

The model incorporates the following main features:

1. experimental data
 - a. food conversion efficiencies and/or weight-specific daily ration obtained from feeding experiments; and
 - b. a conversion parameter β (beta; see below for definition)
2. field data
 - a. the parameters (W_∞ , K and t_0) of VBGF; and
 - b. an estimate of mortality (here M).

Parameter Estimation

ESTIMATION OF β (BETA). Feeding experiments can be used to obtain estimates of gross conversion efficiency (K_1), which is defined for any time interval by (Ivlev 1939, 1966)

$$K_1 = \text{growth increment/food ingested} \quad \dots 2.17)$$

or (Paloheimo and Dickie 1966)

$$K_1 = (dw/F)dt \quad \dots 2.18)$$

dw/dt is the growth rate and F the food ingested. Growth increment and food ingested are expressed in the same units (e.g., g wet weight, dry weight or protein). Under stable laboratory or environmental conditions, food conversion efficiency and fish weight usually show a strong negative correlation. This can be expressed by the regression

$$\log_{10} K_1 = \log_{10} a + b \log_{10} W \quad \dots 2.19)$$

which leads to the conventional model

$$K_1 = aW^b \quad \dots 2.20)$$

(for a discussion of this model, see Jones 1976). Pauly (1986) derived the alternative model

$$K_1 = 1 - (W/W_\infty)^\beta \quad \dots 2.21)$$

with β as the slope of a double logarithmic plot when equation (2.21) is transformed to

$$-\log_{10}(1-K_1) = \beta \log_{10} W_\infty - \beta \log_{10} W \quad \dots 2.22)$$

W_∞ is the weight at which $K_1 = 0$ (x-intercept), and $K_1 = 1$ when $W = 0$. W_∞ is thus defined as the weight at which dw/dt is 0, independently of the amount of food ingested. This corresponds to the definition of asymptotic weight (W_∞) in the VBGF (see above).

For cases in which the available data lead to unrealistic estimates of W_{∞} , equation (2.22) can be rewritten as

$$\beta = \bar{C} / (\log_{10} W_{\infty \text{VBGF}} - \overline{\log W}) \quad \dots 2.23)$$

where \bar{C} is the mean of the $-\log_{10}(1-K_t)$ values, $\overline{\log W}$ the mean of the $\log_{10} W$ values and $W_{\infty \text{VBGF}}$ an estimate of asymptotic size suitable to describe the growth of the fish in question. The same value of W_{∞} can thus be used for a combined analysis of growth and food consumption.

Pauly (1986) also emphasized the appropriate choice of the regression model used. Thus when (1) the $\log_{10} W$ values are not controlled by the experimenter or (2) regression parameters are required rather than prediction of C-values, a "geometric mean" or Type II regression should be used instead of an "arithmetic mean" or Type I regression. Conversion from a Type I to a Type II regression can be performed through

$$b' = b / |r| \quad \dots 2.24)$$

and

$$a' = \bar{C} - b' \overline{\log_{10} W} \quad \dots 2.25)$$

where a, b are the parameters of Type I, a' and b' the parameters of Type II regression and r is the correlation coefficient linking the y (= C) and the x (= $\log_{10} W$) values (Ricker 1973).

ESTIMATION OF FOOD CONSUMPTION (Q/B). Food conversion efficiency as a function of age t of a given fish $K_1(t)$ can be obtained by inserting equation (2.2) into equation (2.21):

$$K_{1(t)} = 1 - \left(1 - e^{-K(t-t_0)}\right)^{b\beta} \quad \dots 2.26)$$

where K, t_0 and β are as defined above.

The feeding rate (dq/dt) of a fish of age t can then be expressed by rewriting equation (2.17) as follows:

$$dq/dt = (dw/dt) / K_{1(t)} \quad \dots 2.27)$$

The growth rate (dw/dt, e.g., in kg year⁻¹) is the first derivative of VBGF and has the form:

$$\frac{dw}{dt} = W_{\infty} * 3K \left(1 - e^{-K(t-t_0)}\right)^{b-1} * e^{-K(t-t_0)} \quad \dots 2.28)$$

The cumulative food consumption (Q_c) of a fish can thus be obtained by integrating equation (2.26) from age at recruitment (t_r) to age at which the fish dies (t_{max})

$$Q_c = \int_{t_r}^{t_{max}} \left(\frac{dw}{dt} \right) K_1(t) dt \quad \dots 2.29)$$

By assuming exponential decrease with instantaneous total mortality Z , viz

$$N_t = N_r e^{-Z(t-t_r)} \quad \dots 2.30)$$

with N_t the number of fish in the population at time t and N_r the number of recruits, the food consumption per unit biomass (Q/B) of an age-structured population can be estimated from:

$$\frac{Q}{B} = \frac{\int_{t_r}^{t_{max}} \frac{(dw/dt) N_t}{1 - \left(1 - e^{-K(t-t_0)}\right)^{b\beta}} dt}{\int_{t_r}^{t_{max}} W(t) * N_t * dt} \quad \dots 2.31)$$

Thus, the parameters needed for the estimation of Q/B of a fish or invertebrate population are b , β , W_∞ , Z (or M for an unfished system), K , t_0 , t_r and t_{max} . The parameter b is the exponent of the length-weight relationship and can generally be set equal to 3 (see above); the parameter β is estimated from feeding experiments; W_∞ , K and t_0 are obtained from growth data, whereas t_r and t_{max} can be set more or less arbitrarily (a sensitivity analysis of the model showed only negligible effects for changes of t_r , t_{max} and t_0 ; see Pauly 1986).

ESTIMATION OF MAINTENANCE RATION AND TROPHIC EFFICIENCY. Apart from Q/B , the above model can also be used to estimate:

1. maintenance ration (R_M);
2. trophic efficiency (E_T); and
3. respiratory and excretory losses (R_L).

Maintenance ration is defined as the amount of food necessary to maintain a given weight of a fish. R_M is usually obtained through controlled feeding experiments (Windell 1978), through starvation experiments (Jones 1976) or by measuring the oxygen consumption of a fasting animal (Ursin 1967). Pauly (1986) used the relationship between Q/B and fish weight to estimate R_M ; W_∞ being defined as the weight at which all food ingested is only used for

maintenance and not for growth, the ration at (or near) W_{∞} can be used as estimate of R_M . In practice, R_M is obtained by extrapolating to W_{∞} , using a semilog plot, estimates of Q/B for two fixed weights close to W_{∞} , e.g., $0.90 \cdot W_{\infty}$ and $0.95 \cdot W_{\infty}$ (Pauly and Palomares 1986).

Population trophic efficiency can also be estimated from the model with

$$E_T = Z \cdot (B / Q) \quad \dots 2.32)$$

where Z is total mortality (or M natural mortality for an unfished population) and B/Q the reciprocal of the output of equation (2.33). E_T expresses population production per unit food consumed. Food that has not been turned into production of biomass goes to respiratory and excretory losses; the relative amount of such losses can be calculated through

$$R_L = 1 - E_T \quad \dots 2.33)$$

Pauly's (1986) food consumption model was used to obtain estimates of Q/B for several macroinvertebrates (see Results section). All values of Q/B , R_m and E_T were calculated with a BASIC computer program called MAXIMS (Jarre et al. 1990).

The bulk of the Q/B estimates used for the development of the empirical model, described below, was calculated by applying published results of food conversion experiments and growth parameters of the fish stocks considered to the above described food consumption model (see Palomares 1987).

2.6.6.2. THE EMPIRICAL MODEL

Fish Species

Food consumption per unit biomass (Q/B) for the fish species included in the present models of a Caribbean coral reef was computed with an empirical model of the form:

$$\log_{10} Q / B = -0.0771 - 0.2018 \log_{10} W_{\infty} + 0.6121 \log_{10} T + 0.5156 \log_{10} A + 0.5471 F \quad \dots 2.34)$$

(Palomares and Pauly 1989), where Q/B is the food consumption of an age-structured population over its mean biomass, W_{∞} is the mean asymptotic (or maximum) weight (as defined by the VBGF) of the fish of a given population (live weight in g), T is the mean environmental temperature in °C, A an index of the mean activity level of the fish of a given species, derived from the shape ("aspect ratio") of its caudal fin (see below) and F is the food type, with : carnivorous = 0 and herbivorous = 1. The derivation of this model, which is modified from Palomares (1987), is given in Palomares and Pauly (1989). It is based on Q/B estimates for 33 demersal and pelagic fish stocks ranging in size from myctophids to tuna and occurring in marine waters with mean annual temperatures ranging from 10 to 28°C.

ASPECT RATIO OF THE CAUDAL FIN. The aspect ratio (A) of the caudal fin is defined as the square of the height (h^2) of the caudal fin over its surface area (s) or

$$A = h^2 / s \quad \dots 2.35)$$

(Lindsey 1978). This ratio can be used to characterize the activity level of fishes which use their caudal fin as (main) organ of propulsion. High aspect ratios indicate that the fish are fast and continuous swimmers (e.g., tunas) while low aspect ratios refer to slow and/or "burst" swimmers (e.g., groupers, see Fig. 2.2). The aspect ratio of the caudal fin is thus related to the activity level and energy requirements of the fish (Sambilay 1990). It is therefore an indirect measure for the food consumption of a fish (or a population of fish).

Based on photographs in Randall (1968) and drawings in Fischer (1978), the aspect ratios of around 230 Caribbean coral reef fish species were determined using a computerized image analyzing system. This newly available technology permits to rapidly analyze a high number of pictures, especially when combined with a special data storage program, as was the case in this study. This program was written for the present application and kindly made available by R. Froese (ICLARM). For computing Q/B, estimates of aspect ratios obtained by Randall's photographs were used and only when these were not available were values from the Food and Agriculture Organization (FAO) drawings applied. Indications of length for a variety of species were adapted from Randall (1968).

For sharks, which have heterocercal tails, and for rays, an aspect ratio of 7.0 was assumed. This value is based on the reflection that sharks and rays act as apex predators, roaming through widespread hunting grounds and thus may have an activity level similar to large scombrids and jacks, which highest aspect ratios varied between 6 and 7.

For species without well-formed caudal fins (such as moray eels), an aspect ratio of 0.7 was assumed; this corresponds to the lowest level of A among fish species with caudal fins. This assumption was made since morays and similar fishes usually spend much time hidden in crevices waiting for preys to pass by.

FOOD TYPE. The food type (F) of equation (2.34) is a dummy variable which originally took only two values: 0 for carnivorous and 1 for herbivorous fish. To be able to deal with omnivory (i.e., diets containing both animals and plants), linearity was assumed between 0 and 1, i.e., the variable F was set at values proportional to the fraction of plants in the diet when solving for Q/B.

Nonfish Taxa

The majority of estimates of Q/B for nonfish taxa were obtained by searching directly in the literature for estimates of this parameter. As pointed out above, for several taxa Q/B could be

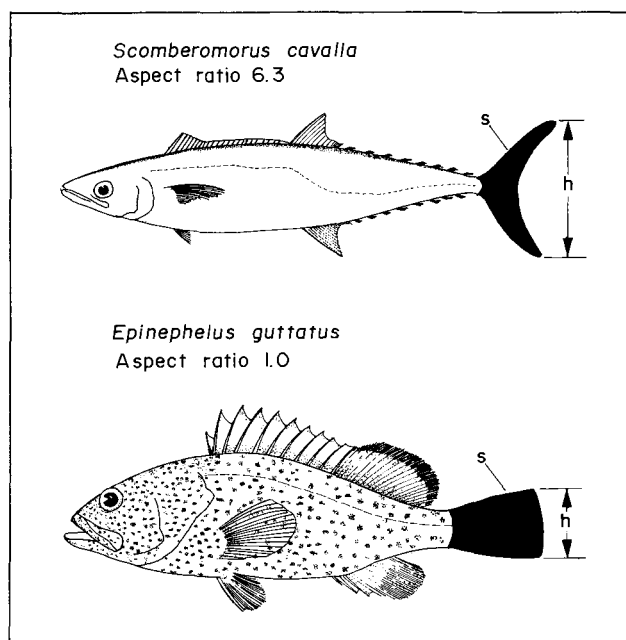


Fig. 2.2. Schematic representation of method to estimate the aspect ratio of the caudal fin of fish, given height (h) and surface area (s).

estimated through the food consumption model (Pauly 1986), i.e., via β as obtained from published estimates on food conversion or daily ration and from growth parameters.

I also attempted to estimate Q/B from data on energy metabolism, i.e., from oxygen consumption estimates. This approach yielded, for reasons not explored here, excessively high consumption estimates and was therefore abandoned.

Consumption values from the literature were adjusted to a mean annual temperature of 28°C, prevailing in the Caribbean waters around the Virgin Islands, through a reexpression of the slope associated with temperature in equation (2.34), i.e.,

$$V = (T / T')^{0.6121} \quad \dots 2.36)$$

V is the factor for temperature adjustment, T the temperature at PRVI (i.e., 28°C) and T' the original habitat or experimental temperature.

2.7. Diet Composition Matrix

It was known from a previous application of ECOPATH II to the Peruvian upwelling system (Jarre et al. 1991), that changes in the diet composition had a strong impact on model structure. Therefore, the diet matrix of the present modelling work was elaborated as carefully as possible for the species groups involved.

2.7.1. Fish Species

The data for the diet composition matrix DC_{ij} were obtained primarily from a study carried out in St. John, US Virgin Islands, by Randall (1967).

The results of Randall's stomach contents analyses were given in percent of volume of the different major groups of food organisms. This is roughly equivalent to relative weight and thus, approximates the relative amounts of energy extracted by consumers from various system components (MacDonald and Green 1983).

The taxonomic level of the organisms identified by Randall (1967) from fish stomachs varied from species to families and higher categories, with most of the invertebrates grouped under the higher categories. Fishes, on the other hand, were identified mostly to generic or specific level without indication of percentage volume of the single genus or species. For the construction of the DC matrix, the percentages of "fish" were divided into equal parts between all the genera and species listed as prey for the respective fish species. When fish were identified only to family or generic level, the respective percent of volume was distributed evenly over all species of the family or the genus represented in the matrix. Only 18 fish species appeared as prey but not as predators. Their share was distributed evenly over all the species of the same family or genus considered in the matrix. Species of families that were not represented in the matrix were considered as "unidentified" fish (F0). The F0 in the matrix contains also all cases where the share of fish in a diet was not identified to a lower taxonomic category.

Unidentified stomach contents were assigned to the identified parts according to the relative abundance of the latter. Eggs and larvae were assigned as "zooplankton", together with

a variety of other organisms (see Appendix 8.5.3). In Randall's study, "algae and organic detritus" were listed as a combined item in many cases. Their relative contribution was divided into two equal parts. The symbiotic coral-algae complex was separated into a producer and a consumer component; when a fish species' diet included the item "corals", the corresponding percentage was divided into two shares with 60% of it for plant tissue (benthic algae) and 40% for animal tissue (corals).

Only five species of sharks, four of the family Carcharhinidae and one of the family Ginglymostomatidae were included in Randall's study. These five species were represented only by few individuals, many of which had (nearly) empty stomachs. The same was true for stomach analyses of the two species of rays included in the model. Diet for cartilaginous species was generally stated as "fish", preliminarily included into the food matrix under the label F0. Randall's list of shark species was completed by including more sharks with affinities to coral reefs of the PRVI area, based on accounts in Fischer (1978). From qualitative descriptions of the diet composition given therein, quantitative shares were estimated for each food item according to the already existing DC matrix. The item "fish" or "small fish" was most difficult to translate into quantitative information. It was assumed that, when part of the diet was stated to be "fish", every fish species in the system had the same probability of being a victim of a shark's appetite. So the generalization "feeds on fishes" was translated by distributing 100% (minus the part of the diet represented by invertebrates) into equal percentages over all fish species in the DC matrix. Excluded were (1) species larger than the one under consideration and (2) reef fish species with $W_{\infty} < 100$ g. The latter restriction was adopted based on the notion that a large animal, such as a tiger shark, weighing about half a ton, would generally not manage to extract gobies, blennies, damselfishes, etc., of 10-20 g from between the coral heads and other crevices. When the diet, or part of it, was stated to be "pelagic fish", this was divided into the pelagic species in the DC matrix. When a shark fed on "small pelagic fish" the same procedure was applied, but excluding the scombrids and large jacks from the list of possible candidates. When the food or part of it consisted of "small fish", the diet was distributed evenly over all species with $W_{\infty} < 1,000$ g. The weight limits are arbitrary. Such arbitrary limits were necessary, however, since a rigorous definition of what a "small fish" is does not exist.

The relative contribution of fish versus nonfish taxa is also arbitrary. The only restriction was the statement "feeds mainly on...". In such cases, the diet part referred to was considered to represent from 60% to 80% of the total diet.

2.7.2. Nonfish Taxa

To establish a diet matrix for the nonfish taxa, the food items of fish stomachs analyzed by Randall (1967) were listed under their corresponding taxon together with the number of fish species for which an item was stated as part of the diet (see Appendix Tables 8.5.1 and 8.5.2). Based on this listing, a search of the Aquatic Sciences and Fisheries Abstracts (ASFA) database was carried out and ca. 300 publications were selected. Since quantitative information on diet composition of coral reef invertebrates is still scarce (see also Lewis 1981), the quantitative diet composition of each taxon was estimated in many cases from qualitative or rough quantitative information and/or was transferred from ecosystems other than coral reefs.

For the construction of the DC matrix, all percentages of volume were reexpressed as fraction of one.

2.8. Aggregation of Species and Taxa

2.8.1. First Reduction

ECOPATH II, as presently implemented, allows a maximum input of 50 species groups. This restriction made it necessary to reduce the initial number of fish species (over 200) and 41 nonfish taxa into 50 species groups. The procedures used for this reduction are described below.

2.8.1.1. FISH SPECIES

A set of ecologically meaningful criteria was needed by which the fish species could be aggregated into species groups, using cluster analysis. Since I was concerned with a model of energy flow, variables which relate to flows between elements of the system had to be identified. These variables must necessarily relate to the factors which determine the food consumption of a species. Referring to the section on "Food consumption (Q/B)", these variables are: (1) size, (2) activity level and (3) type of food. These variables were parametrized as in equation (2.34), using: (1) size = asymptotic weight W_{∞} in g; (2) activity level = aspect ratio A; and (3) type of food = degree of herbivory, in %. These three were treated as being of equal importance. Prior to the aggregation of fish species into species groups by cluster analysis, two of the three variables were transformed such that they evenly covered a range of values.

Asymptotic weights (originally in g wet weight) were reexpressed in logarithmic form and thus reduced to values between 0 and 6. Food type was divided by 10 and thus ranged then from 0 to 10. A value of 0 corresponded now to 100% carnivory and a value of 10-100% herbivory. Aspect ratio was not transformed since this evenly covered values between 0.7 and 7.0.

To determine whether the three variables were correlated, a factor analysis was carried out. According to Backhaus et al. (1986), prior to the execution of a cluster analysis on a given set of elements and variables, it is advisable to carry out a factor analysis to detect dependent variables and exclude them from the aggregation process. The factor analysis showed that the variables were largely independent, except for a negligible correlation between aspect ratio and size (Fig. 2.3).

Cluster analysis was performed, using the prepared matrix of 243 fish species and 3 variables. "Quadratic Euclidean distance (QED)" was selected as index of dissimilarity. By relying on the square of the distance between the variables of two elements, this index emphasizes large distances.

Three hierarchical agglomerative cluster-algorithms were applied to the matrix of input data: (1) "average linkage", (2) "centroid" and (3) "ward". All three are conservative algorithms (they have no contracting or dilating effect on the grouping process) and may be used with measures of dissimilarity (e.g., QED). "Average linkage" and "ward" are monotonous algorithms where the measure of dissimilarity (QED) increases monotonously with decreasing number of clusters. "Ward" has the exclusive feature of forming groups of similar sizes, i.e., groups with a similar number of elements. By comparison of the results (clusters) obtained by the three algorithms, "average linkage" was finally selected to be most appropriate. The species groups aggregated by this algorithm represented best the conditions in the reef, even with reference to the habitat of certain species groups (see below).

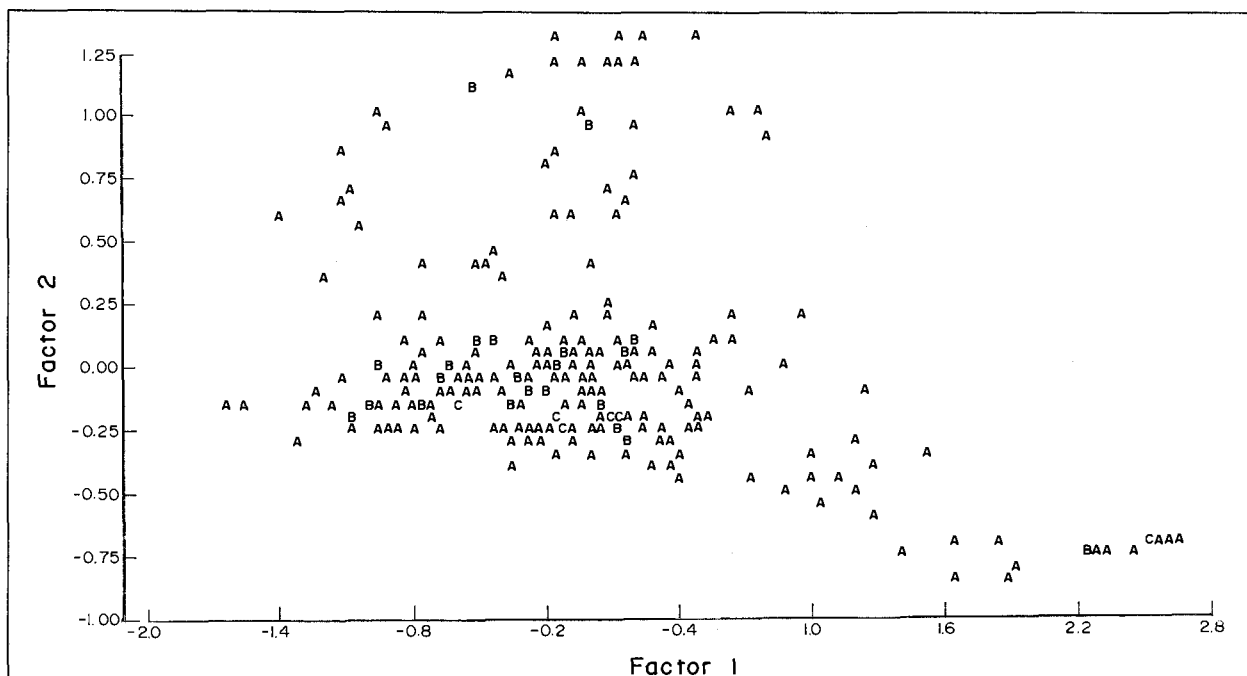


Fig. 2.3. Results of the factor analysis; relation between asymptotic weight (factor 1) and aspect ratio (factor 2).

A QED value of 1.5 was achieved when all fish species were aggregated into a single cluster. This corresponds to a cutoff level of 100%. At a cutoff level of 20% (QED = 0.3), 29 species groups were identified. Of these, 6 species, each forming a single cluster, were regarded as outliers and merged (outside the cluster analysis) with larger species groups.

An additional criterion of distinction outside the cluster analysis was applied to the prepared boxes. This criterion referred to the vertical dimension of the habitat. It was defined to be the level in the water column where a species spends most of its time. The water column was divided into two levels: demersal/off bottom and midwater/pelagic. Following this criterion, the species groups were divided into a demersal and a pelagic fraction whenever they contained species from both habitats. This division was omitted in groups of large predators because not enough information on preferred level was available, or a species could not be assigned to a distinct level. Overall, 27 fish species groups or boxes were obtained by the above-described method of aggregation.

The theoretical background for the factor analysis and cluster analyses was obtained from Backhaus et al. (1986) and Pielou (1984). The analyses were performed with the assistance of Mr. C. Klingenberg, using SAS on a VAX 750 mainframe at the Institut für Meereskunde, Kiel.

2.8.1.2. NONFISH TAXA

The aggregation of nonfish taxa could not be performed in the same manner as for fish species due to their broad taxonomic range. Values for size, activity and food type were almost entirely lacking at the species level. Activity could not be transformed into a simple parameter due to the extreme difference in lifestyle. Reduction from initially 41 to 23 groups was performed by applying the following, largely subjective aggregation criteria (in order of importance):

1. Availability of estimates of P/B and Q/B: Estimates of these two input parameters were only available for 24 of the 41 taxa. Fortunately, most of the taxa for which estimates could be obtained were definitely of great importance to reef metabolism.
2. Size: Rough indications of size for the majority of taxa were obtained from Barnes' (1963) *Invertebrate Zoology*. For some, size indications were encountered while compiling estimates of P/B and Q/B of nonfish taxa. Size was roughly scaled into four groups: small, medium, large, very large. It was assumed that, for taxa with a wide spectrum of sizes (e.g., polychaetes, gastropods), the average size contributed the bulk of biomass to the respective box.
3. Similarity of diet composition: Diet composition was reduced to six compartments: detritus, benthic producers/symbiotic algae, phytoplankton, microfauna, zooplankton, macroinvertebrates/fish.
4. Similarity of lifestyle: Sessile, semisessile, burrowing in bottom or calcareous skeletons, free living, filtering, grazing, scavenging, predatory were the categories considered.
5. Taxonomic closeness: This feature was used in cases (e.g., corals/sea anemones) where the preceding criteria were not applicable or did not allow a decision because, e.g., the lifestyles were too different, as in the case of sessile or slow-moving organisms which had no adequate group to be pooled with in terms of sizes or diet compositions.

2.8.2. Weighting Within Species Groups

To obtain a reasonable distribution of food percentages within the species groups, these had to be weighted by considering throughput of every species belonging to the respective group. A weighting factor (WF) was computed for every species through

$$WF = W_{\text{mean}} * AB * Q / B \quad \dots 2.37)$$

where W_{mean} is as defined for equation (2.12) above, AB is the abundance of a species in Nm^{-2} and Q/B the annual food consumption per unit biomass.

2.8.3. Combining Species Groups

Single fish species and nonfish taxa were combined into the respective groups according to the results of the aggregation procedure.

2.8.3.1. P/B RATIO AND FOOD CONSUMPTION (Q/B)

A group estimate of P/B ratio and Q/B was obtained from the median of the respective parameter for each group of estimates for single species or taxon.

2.8.3.2. DIET COMPOSITION MATRIX

For the condensation of the DC matrix, groups had to be combined vertically (columns) and horizontally (rows). All columns and rows belonging to one group were summed up. Thus, each row in the reduced DC matrix represented a predator group and each column a prey group. To bring the sum of a row back to unity (all values had formerly been multiplied by the

corresponding weighting factor, see above), each value in a row was divided by the sum of the row:

$$d_{(ij)} = \frac{d_{(ij)}}{\sum_{k=1}^n d_{(ik)}} \quad \dots 2.38)$$

where d_{ij} is the weighted j th value of species group i , k is the number of columns and d_{ij} ' the resulting part of 1.0 in column j of group i . This procedure was carried out for each species group and each level of aggregation.

2.8.4. Second Reduction of Species Groups

The initial 50-box model was still rather difficult to handle and visualize. For this reason, and also to find out whether information on the energy flow is lost when the system is further aggregated, the number of boxes was further reduced. Models with an equal number of boxes generated by two different methods of reduction were also compared.

2.8.4.1. METHOD 1: CLUSTER ANALYSIS + INTUITION

Fish Groups

The fish groups were further reduced by increasing the average distance (expressed by the dissimilarity index QED) between clusters. Two additional cutoff levels of 40% and 50% were selected. After separating the "pelagic-midwater" from the "demersal-off bottom" species, two sets with nine, respectively, four fish species groups were obtained. Combining groups was achieved by computing a mean (weighted by biomass) for the three key input parameters. The DC values were computed as weighted means (by throughput = biomass * consumption) from the groups of which they consisted.

Nonfish Groups

The nonfish groups were again difficult to reduce, especially because no standardized criterion could be identified for lumping nonfish groups with each other and with fish groups, a necessary procedure when reducing the boxes of the system (<20). Based on diet composition, size and lifestyle, boxes were combined intuitively with each other as well as with fish groups. Results of this "intuitive guesswork" (Ulanowicz 1986) were contrasted with those of the objective aggregation routine included in ECOPATH II, described in the Introduction and summarized in Section 2.8.4.2. The resulting total number of boxes including all fish species and nonfish taxa was 20 and 11, respectively.

2.8.4.2. METHOD 2: ECOPATH II AGGREGATION ROUTINE

As described in the Introduction, the ECOPATH II program (Pauly and Christensen 1993) includes an aggregation routine, based on Ulanowicz (1986), which allows the stepwise reduction of the number of boxes in a model from a highest possible number of 50 to any small

number (after the initial system is balanced). Boxes are linked pairwise such that ascendancy (a measure of the information content of a system, see Introduction) is reduced as little as possible. With this aggregation routine, the boxes of the present coral reef system were reduced pairwise until the 50 initial boxes of the system were reduced to 10. The intermediate models with the box numbers corresponding to those obtained by method 1 were used to compare models resulting from both methods.

2.9. ECOPATH II Parameter Estimation Routine

The databases for the 50-, 20- and 11-box models obtained by method 1 were fed into the ECOPATH II program, and the ecotrophic efficiency (EE) for each box was computed. Then, the original input values of the 50-box model were gradually modified, until all components (boxes) showed an $EE < 1$. Starting from the balanced 50-box model, the ECOPATH II aggregation routine (method 2) was then used to compute the parameter values for each aggregation step (this involved automatical transformation of the diet matrix) such that every new model resulting from a further reduction of boxes was again balanced.

Besides estimation of EE, ECOPATH II was used to compute the estimates for each box of the parameters of the "master equation" (see Introduction). With the estimates for food intake, predation mortality (= production (P) * EE), flow to detritus and assimilation (A) (all in $\text{gm}^{-2}\cdot\text{year}^{-1}$), an energy balance for every box could be calculated. Furthermore, ECOPATH II calculated trophic level, omnivory index, gross efficiency, net efficiency, R/A ratio and P/R ratio for each box of the system. Based on the diet composition matrix, ECOPATH II computed, finally, for each predator, the food intake in $\text{gm}^{-2}\cdot\text{year}^{-1}$, Ivlev's electivity index and the standardized forage ratio of each prey group (see Introduction).

2.10. Whole System Properties

In the Introduction, the various features of ECOPATH II were presented. Besides the ability to produce a balanced system, the program offers several routines for the computation of various whole system properties useful for comparing different ecosystems or models of the same ecosystem.

These summary statistics are: sum of all production, sum of all imports, sum of all respiratory flows, sum of flows into detritus, total system throughput, throughput cycled, Finn's cycling index and net primary production necessary to support the system.

The "network flow indices" routine enabled the computation of ascendancy, overhead and capacity for distinctive sources (inputs, internal and dissipations) of the system. The same routine computed flows and transfer efficiencies within the system subsequent to dividing it into discrete trophic levels. Finally, the routine called "mixed trophic impacts" allowed the computation of a predator - prey matrix in which the impact of an increase in biomass for a predator group is shown for all prey groups.

2.11. Graphical Representation of Models

The graphical representation of the box models followed suggestions made on this topic by Jarre et al. (1989) and Pauly and Christensen (1993). Thus, the surface area of each box was drawn so as to be proportional to the logarithm of the biomass in that box. Also, care was taken

to arrange the boxes along the ordinate (trophic level) such that they did not overlap and enough space was left to draw the arrows representing the flows. Furthermore, the boxes were arranged along the abscissa so as to minimize crossovers of arrows. Whenever possible, flows were combined to reduce the number of arrows between boxes. About 80-90% of flows into a box from the components of the system were included into the graphical representations of the models, the rest was omitted for clarity. The complete sets of flows for each box are shown in Appendix 8.8.

3. RESULTS

3.1. Fish Species

3.1.1 Ecology

The available information on the spatial distribution of coral reef fish species in the Puerto Rico - Virgin Islands (PRVI) area are summarized, by subsystem, in Table 3.1. While the majority of species live within the coral reef, a number of others use the reef only as shelter and forage in the adjacent areas, such as the seagrass beds and the algal flats. These species contribute to reef metabolism mainly by means of their excreted and egested matters (feces). Several species use mangrove areas and estuaries as nursery for their offspring. These few examples show that in terms of energy flow, the subsystems cannot really be separated from each other (see also Ogden and Gladfelter 1983).

The resident coral reef fishes and the regular visitors to reefs show a very high species diversity, whereas the pelagic or oceanic fishes that are only indirectly related to reef metabolism (through their foraging activity or release of excreta) contribute relatively few species.

However, Table 3.1 is only a first attempt to group into compartments what is, in reality, a gradient with smooth transitions.

3.1.2. Fisheries and Management

The fisheries in the PRVI reef area consist mainly of small-scale, commercial fishing, and the rest of recreational fishing (and perhaps some subsistence fishing). The Caribbean Fisheries Management Council (CFMC 1985) estimates that 22% of total landings originate from the recreational sector in the US Virgin Islands. In the shallow-water reef fisheries, it is estimated that recreational landings contribute 13% of total landings. The overwhelming part of domestic requirements of fish products in the PRVI area (Puerto Rico, 85%; US Virgin Islands, over 60%) is covered by imports. Table 3.2 gives the number of fishers and vessels comprising the commercial fisheries in the PRVI area.

Fish traps are by far the most important fishing gear in terms of units as well as yield in the PRVI area. The second most important gear is the handline. Table 3.3 gives the relative importance of the different gear in the fisheries of Puerto Rico and the US Virgin Islands. A local annual fisheries production of around 3,000 t for Puerto Rico and 1,700 t for the US Virgin Islands is indicated (Table 3.3).

The CFMC reports the following 10 species as important in PRVI: Nassau grouper (*Epinephelus striatus*), red hind (*E. guttatus*), coney (*Cephalopholis fulva*), lane snapper (*Lutjanus synagris*), mutton snapper (*L. analis*), yellowtail snapper (*Ocyurus chrysurus*), white grunt (*Haemulon plumieri*), queen triggerfish (*Balistes vetula*), spotted goatfish (*Pseudupeneus*

Table 3.1. Spatial distribution of Caribbean reef fishes (with emphasis on the Virgin Islands and notes on their ecology).

F=foraging, H=home, CI=cleaner, y=young, 1=bottom, 2=well above bottom, 3=midwater, 4=pelagic; + = reported to occur in subsystem.

VA=very abundant, A=abundant, C=common, F=fairly common, O=occasional, R=rare, E=absent.

R=resident, V=regular visitor, T=transient.

W=wide range from shallow to deep; V=very shallow, surface to 2 m; SH=shallow, 1-15 m; D=deep, below 15 m; +=upper depth limit SH.

* = from tables in Boulon (1986b); type of length not stated, therefore total length is assumed; average size of species estimated during census on reef; species with an estimate are of commercial importance in the Virgin Islands Biosphere Reserve; values in brackets were recorded for genus or family.

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|-----------------------------|-----------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|-----------------------------------------------------------------------------------------------|---------------------------------------------------------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | HF1y | HF1 | F1 | | | SH | | O | V | (11.4) | |
| | <i>Acanthurus chirurgus</i> | HF1y | HF1 | F1 | | | W | | O | | (11.4) | |
| | <i>Acanthurus coeruleus</i> | HF1y | HF1 | + | | | SH | | C | R | (11.4) | More closely related to coral reefs than above species. |
| Albulidae | <i>Albula vulpes</i> | | | | HF1 | | SH | | S | | | Food: worms, molluscs, crabs, shrimps, squid (FAO 1977). |
| Antennariidae | <i>Antennarius multiocellatus</i> | | HF1 | | | | SH | | S | | | Most common of family on West Indian reefs; commercially unimportant. |
| | <i>Antennarius striatus</i> | | HF1 | HF1 | HF1 | | | | O | | | In different habitats; commercially unimportant. |
| Apogonidae | <i>Apogon binotatus</i> | | HF1 | | | | | | | R | | |
| | <i>Apogon conklini</i> | | HF1 | | | | SH | | C | R | | |
| | <i>Apogon maculatus</i> | | HF1 | | | | W | | C | R | | Largest and most common inshore cardinal fish of West Indies (WI). |
| | <i>Apogon pigmentaria</i> | | HF1 | | | | | | | R | | |
| | <i>Apogon quadrisquamatus</i> | | HF1 | | | | | | | R | | |
| | <i>Apogon townsendi</i> | | HF1 | | | | | | | R | | |
| <i>Apogon xenus</i> | | HF1 | | | | | | | R | | | |
| Atherinidae | <i>Allanetta harringtonensis</i> | | HF3 | HF3 | HF3 | | | | O | | | Widely used as bait. Important forage for commercial fishes. |
| | <i>Atherinomorus stipes</i> | | HF3 | HF3 | HF3 | | | | O | | | See above. Most common inshore of family in WI. |
| Aulostomidae | <i>Aulostomus maculatus</i> | | HF2 | F2 | | | A | | O | | | Clear water. |
| Balistidae | <i>Balistes caprisicus</i> | | HF1 | HF1 | HF1 | | W | 50 | S | | | |
| | <i>Balistes vetula</i> | | HF1 | HF1 | HF1 | | W | 100 | O | | 27.9 | |
| | <i>Canthidermis sufflamen</i> | | | | | HF4 | W | | S | | | Rare inshore, but common on offshore reefs, near dropoffs to clear water. |
| | <i>Melichthys niger</i> | | HF2 | | | | W | 30 | O | | | Outer coral reefs with clear water. |
| <i>Xanthichthys ringens</i> | | HF1 | HF3y | | | D | >35 | O | | | One of most abundant WI reef fishes in >35m, above rare; young in floating <i>Sargassum</i> . | |
| Belonidae | <i>Ablennes hians</i> | | | | | HF4 | V | | S | | | Inshore occurrence around islands. |
| | <i>Platybelone argalus</i> | HF4 | HF4 | HF4 | HF4 | HF4 | | | F | | | Frequently found in inshore waters. |
| | <i>Strongylura marina</i> | HF4 | | | HF3 | | | | E | | | Coastal areas and mangrove-lined lagoons. |

continued...

Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks | |
|-----------------------------|-------------------------------------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | HF1y | HF1 | F1 | | | SH | | O | V | (11.4) | Inshore species, often in mangrove sloughs, enters fresh waters. More offshore than <i>T. crocodilus</i> but also in coastal waters. Largest of Belonidae; occurs more inshore than above species. | |
| | <i>Strongylura timucu</i> | | HF3 | | HF3 | | | | F | | | | |
| | <i>Tylosurus acus</i> | | | | | HF4 | | | O | | | | |
| | <i>Tylosurus crocodilus</i> | | | | HF4 | HF4 | | | O | | | | |
| Blenniidae | <i>Entomacrodus nigricans</i> | | HF1 | | | | ++ | | F | R | | Inshore species, bottom-bound. | |
| | <i>Hypoleurochilus</i> sp. | | HF1 | | | | SH | | A | | | One of most abundant fishes on WI reefs (0.65 individuals m ⁻² based on Randall 1968). | |
| | <i>Ophioblennius atlanticus</i> | | HF1 | | | | | | O | | | Rocky areas close to shore. | |
| | <i>Parablennius marmoratus</i> <i>Scartella cristata</i> | | HF1 | | | | ++ | | F | | | | |
| Bothidae | <i>Bothus lunatus</i> | HF1 | HF1 | + | HF1 | | 0 | 65 | F | | | Also in mangrove areas. | |
| | <i>Bothus ocellatus</i> | | HF1 | + | HF1 | | SH | | O | | | | |
| | <i>Paralichthys tropicus</i> | | | | HF1 | | ++ | 183 | F | | | | |
| Carangidae | <i>Alectis ciliaris</i> | | | | | HF2 | W | >60 | O | | | Solitary openwater species. | |
| | <i>Caranx bartholomaei</i> | | | | | H2F1 | W | | O | | | Solitary or in small groups; uncommon in inshore waters. | |
| | <i>Caranx crysos</i> | | F1 | | | HF3 | W | | O | | 27.9 | Large schools; sometimes feeding on coral reefs. | |
| | <i>Caranx hippos</i> | | | | | HF3 | | | S | | | Rare in clearwater reef areas. | |
| | <i>Caranx latus</i> | HF2 | HF2 | | | HF3 | W | | O | | | Small groups of small fishes over inshore reefs; larger fish, more offshore. | |
| | <i>Caranx lugubris</i> | | | | | HF3 | 24 | >65 | S | T | 25.4 | Free clear water. | |
| | <i>Caranx ruber</i> | HF2 | HF2 | | | | SH | | C | V-T | 22.8 | Most common jack in WI; foraging and cleaning on reef. | |
| | <i>Decapterus macarellus</i> | | | | | F4 | W | | S | | | Occasionally over outer reef areas; zooplanktonfeeder. | |
| | <i>Decapterus punctatus</i> | | F2,3 | | | F4 | W | 90 | O | | | Occasionally over outer reef areas; zooplanktonfeeder. | |
| | <i>Elagatis bipinnulatus</i> | | F3 | | | F4 | W | | S | | | Pelagic, but occasionally nearshore, also over reefs. | |
| | <i>Oligoplites saurus</i> | HF3,4 | | | | HF3,4 | | | | C | | | Small groups near surface; more turbid than clear waters. |
| | <i>Selar crumenophthalmus</i> | HF2,3 | F2 | | | HF2,3 | | SH | | O | | | Schooling fish over shallow reefs or in turbid waters. |
| | <i>Selene vomer</i> | | HF1 | | | HF1 | | ++ | | | | | Small milling schools. |
| | <i>Seriola dumerili</i> | | F1,4 | | | | HF2,4 | W | 360 | S | | 53.3 | Most common of genus in tropical West Atlantic. |
| | <i>Seriola rivoliana</i> | | F1 | | | | | W | | S | | | Rare in shallow waters. |
| <i>Trachinotus falcatus</i> | | F1,4 | F1,4 | F1,4 | F1,4 | | V | >35 | O | | | Rare in WI. | |
| <i>Trachinotus goodei</i> | | HF1 | | | HF1 | | ++ | | O | | | Surf zone and clear waters along sandy beaches; schooling. | |

continued...

Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks | |
|-------------------------------|-----------------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|---------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------|
| Carcharhinidae | <i>Carcharhinus acronotus</i> | | HF | | HF | | | | S | | | Feeds on small fishes. | |
| | <i>Carcharhinus falciformis</i> | | + | + | + | + | W | >500 | S | | | Feeds on fishes, squids, crabs. | |
| | <i>Carcharhinus leucas</i> | | | HF1,4 | HF1,4 | | W | | S | | | Feeds on fishes (mackerels, tunas, smaller sharks, rays); invertebrates (crabs, shrimps, sea urchins); and carrion. | |
| | <i>Carcharhinus limbatus</i> | | HF4 | HF4 | HF4 | HF4 | W | | S | | | Aggregating; feeds on small schooling fishes, rays, squids. | |
| | <i>Carcharhinus longimanus</i> | | HF4 | HF4 | HF4 | HF4 | | | S | | | Females aggregate; feeds on fishes (scombrids, carangids), squids, portunid crabs, turtles, carrion. | |
| | <i>Carcharhinus perezii (springeri)</i> | | HF1 | HF1 | HF1 | | 30 | | O | | | | Probably most common member of genus in inshore water of WI islands; feeds on fishes including rays. |
| | <i>Galeocerdo cuvieri</i> | | HF3 | HF3 | HF3 | HF3 | W | | O | | | | Coastal as well as offshore waters, near the surface; feeds on fishes (also sharks, rays), turtles, seabirds, squids, conchs, crabs, carrion. |
| | <i>Negaprion brevirostris</i> | HF1 | HF1 | HF1 | HF1 | | W | | O | | | | Common inshore species entering also brackish and even fresh water. |
| <i>Rhizoprionodon porosus</i> | | HF1 | HF1 | HF1 | HF1 | W | >500 | O | | | | Primarily a coastal species, but also found in offshore waters. | |
| Chaetodontidae | <i>Chaetodon aculeatus</i> | | HF1 | | | | D | >100 | S | | | Rare in shallow water, most common of family from 30 m depth. | |
| | <i>Chaetodon capistratus</i> | | HF1 | | | | SH | | C | | | Most common of family in WI reefs. | |
| | <i>Chaetodon ocellatus</i> | | HF1 | | | | W | | S | | | Found in deeper water than others of family. | |
| | <i>Chaetodon sedentarius</i> | | HF1 | | | | W | | S | | | | |
| <i>Chaetodon striatus</i> | | HF1 | | | | SH | | O | | | | | |
| Centropomidae | <i>Centropomus undecimalis</i> | | | | | | | | | | 30.4 | | |
| Cirrhitidae | <i>Amblycirrhitus pinos</i> | | HF1 | | | | W | | S-O | R | | | |

continued...

Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|----------------------------|-----------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|----------------------------------------------------------------------------------------|
| Clinidae | <i>Acanthemblemaria spinosa</i> | | HF1 | | | | | | | R | | |
| | <i>Emblemariopsis leptocirris</i> | | HF1 | | | | | | | R | | |
| | <i>Enneanectes altivelis</i> | | HF1 | | | | | | | R | | |
| | <i>Labrisomus guppyi</i> | | HF1 | | | | ++ | | C | | | |
| | <i>Labrisomus haitiensis</i> | | HF1 | | | | | | | R | | |
| | <i>Labrisomus kalisheræ</i> | | HF1 | | | | ++ | | F | | | |
| | <i>Labrisomus nuchipinnis</i> | | HF1 | | | | ++ | | C | | | Largest and most abundant inshore clinid fish in WI. |
| | <i>Pseudemblemaria signifera</i> | | HF1 | | | | | | | R | | |
| | <i>Starksia hassi</i> | | HF1 | | | | | | | R | | |
| <i>Starksia lepicoelia</i> | | HF1 | | | | | | | R | | | |
| Clupeidae | <i>Harengula clupeiola</i> | | HF4 | HF4 | HF4 | | ++ | | VA | | | Pelagic, in coastal waters, estuaries and lagoons; schooling. |
| | <i>Harengula humeralis</i> | HF4 | HF4 | HF4 | HF4 | | ++ | | VA | 10.1 | | Pelagic, in coastal waters, estuaries and lagoons; schooling. |
| | <i>Jenkinsia lamprotaenia</i> | HF4 | HF4 | HF4 | HF4 | | ++ | | VA | V | * | Often extremely abundant; forms large schools near the surface. |
| | <i>Opisthonema oglinum</i> | | HF4 | HF4 | HF4 | | ++ | | C | | | Pelagic, in coastal waters, often forming compact schools. |
| | <i>Sardinella aurita</i> | | HF4 | HF4 | HF4 | | ++ | | C | | | Pelagic, in coastal waters, often forming compact circular schools; feeds on plankton. |
| Congridae | <i>Heteroconger halis</i> | | | | HF1 | | ++ | 50 | F | | | Burrowed in sand near reefs; zooplankton feeder. |
| Dactylopteridae | <i>Dactylopterus volitans</i> | | | | HF1 | | ++ | | O | | | No commercial interest. |
| Dasyatidae | <i>Dasyatis americana</i> | | + | | HF1 | | ++ | | F | | | Most common of genus in WI, regular visitor to reefs. |
| | <i>Urolophus jamaicensis</i> | | | | | | | | | | | |
| Diodontidae | <i>Chilomycterus antennatus</i> | HF1 | + | HF1 | | | W | | S-O | | | Relatively rare; most often taken from seagrass beds. |
| | <i>Chilomycterus antillarum</i> | | + | + | | | W | 30 | S | | | |
| | <i>Diodon holocanthus</i> | HF1 | HF1 | | HF1 | | SH | | O-C | | | More often in muddy habitats (mangroves) than on reefs. |
| | <i>Diodon hystrix</i> | | HF1 | | | | SH | 30 | O | | | Most frequent of family. |
| Echeneidae | <i>Echeneis naucrates</i> | | HF | | | | | | O | | | Attached sometimes to larger reef fishes, sometimes solitary in reef. |
| | <i>Remora remora</i> | | HF | | | | | | O | | | Attached sometimes to larger reef fishes, sometimes solitary in reef. |
| Elopidae | <i>Elops saurus</i> | | HF | | HF | | ++ | | O | | | Common in WI in shallow brackish lagoons. |

continued...

Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|------------------------------|------------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------|
| Emmelichthyidae | <i>Emmelichthys atlanticus</i> | | HF2 | | | | 3 | 50 | F | R | | Small aggregations above coral heads. |
| | <i>Inermia vittata</i> | | | | | HF3 | D | | C | | | Slender schooling fish in clear blue waters. |
| Engraulidae | <i>Anchoa hepsetus</i> | | HF3 | HF3 | HF3 | | ++ | >70 | A | | | Forms large schools; feeds on small zooplankton and zoobenthos. |
| | <i>Anchoa lyolepis</i> | | | HF3 | HF3 | | ++ | >55 | A | | | Forms large, compact schools; on sandy beaches and mangrove lagoons. |
| Ephippidae | <i>Chaetodipterus faber</i> | | HF2 | | | | W | | S | | 22.8 | Schooling. |
| Fistulariidae | <i>Fistularia tabacaria</i> | | + | HF1 | | | SH | | S-O | | | |
| Gerreidae | <i>Eucinostomus argenteus</i> | | | | HF1 | | ++ | | C | | (20.3) | Also in brackish environments. |
| | <i>Gerres cinereus</i> | | HF1 | | HF1 | | ++ | | C | | (20.3) | |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | | HF1 | + | HF1 | | W | | O | | | Most common reef shark in WI; feeds mostly on invertebrates (shrimps, crabs, lobsters, squids, other molluscs, sea urchins); also on small fish. |
| Gobiidae | <i>Bathygobius soporator</i> | HF1 | HF1 | | HF1 | | 0 | | S-O | | | Most common shore fish of tropical America. |
| | <i>Coryphopterus dicrus</i> | | HF1 | | | | | | | R | | Most common goby of WI reefs. |
| | <i>Coryphopterus eidolon</i> | | HF1 | | | | | | | R | | |
| | <i>Coryphopterus glaucofraenum</i> | | HF1 | HF1 | HF1 | W | 27 | | O | R | | |
| | <i>Coryphopterus personatus</i> | | HF1 | | | | | | | R | | |
| | <i>Coryphopterus thrix</i> | | HF1 | | | | | | | R | | |
| | <i>Gnatholepis thompsoni</i> | | HF1 | | HF1 | | 33 | | C | R | | |
| | <i>Gobiosoma evelynae</i> | | HF1 | | | | SH | | F | R | | |
| | <i>Gobiosoma horsti</i> | | HF1 | | | | SH | | F | R | | |
| | <i>Gobiosoma saucrum</i> | | HF1 | | | | | | | R | | |
| | <i>Lythrypnus elasson</i> | | HF1 | | | | | | | R | | |
| | <i>Lythrypnus nesiotes</i> | | HF1 | | | | | | | R | | |
| | <i>Lythrypnus sp.</i> | | HF1 | | | | | | | R | | |
| | <i>Quisquilius hipoliti</i> | | HF1 | | | | | | | R | | |
| <i>Risor ruber</i> | | HF1 | | | | | | | R | | | |
| Grammidae (Serranidae) | <i>Gramma loreto</i> | | HF1 | | | | W | 70 | O | | | Cleaner; common in caves or beneath ledges; highly prized aquarium fish. |
| | <i>Gramma melacara</i> | | HF1 | | | | 10 | 70 | O | | | Bahamas and Belize, rare in <30 m, very common on reef fronts in <50 m in the Bahamas. |
| Grammistidae (Serranidae) | <i>Rypticus saponaceus</i> | | + | | + | | SH | 50 | O | | | |
| | <i>Rypticus subbifrenatus</i> | | + | | | | SH | | | R | | |

continued...

Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|---------------------------------|----------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|-----------------------------------------------------------------------------------------|
| Haemulidae (Pomadasyidae) | <i>Anisotremus surinamensis</i> | | HF1 | F1 | F1 | | SH | | S-O | | | Very shallow water, one of the largest grunts. |
| | <i>Anisotremus virginicus</i> | | HF1Cy | | | | SH | | S | | 17.7 | Common in Florida Keys. |
| | <i>Haemulon album</i> | | H1 | F1 | F1 | H1 | SH | 30 | O | | 25.4 | Largest of genus; clear water; active at night; good food fish. |
| | <i>Haemulon aurolineatum</i> | + | H1 | F1 | F1 | H1 | W | 25 | O | R | 10.6 | Smallest and most common of family. |
| | <i>Haemulon bonariense</i> | + | HF1 | HF1 | HF1 | | SH | | S | | | Abundant in southern Caribbean, also found in mangrove-lined lagoons. |
| | <i>Haemulon boschmae</i> | | HF1 | | | | V | 100 | S | | | Forms loose aggregates. |
| | <i>Haemulon carbonarium</i> | | + | HF1 | HF1 | | W | 25 | S | | 12.7 | Forms schools around coral reefs in clear water. |
| | <i>Haemulon chrysargyreum</i> | | | H1 | F1,3 | F1,3 | W | 25 | O | | 11.4 | Small schools hiding in caves by day. |
| | <i>Haemulon flavolineatum</i> | HF1y | HF1 | HF1 | | | W | 25 | C | R | 11.4 | Most common grunt on WI reefs; one of most abundant species in WI; commonly in schools. |
| | <i>Haemulon macrostomum</i> | | | HF1 | HF1 | | W | | S | | 13.9 | Clear water. |
| | <i>Haemulon melanurum</i> | | | HF1 | HF1 | | W | | S | | | Clear water; schooling. |
| | <i>Haemulon parrai</i> | | | HF1 | HF1 | HF1 | W | 30 | S | | 19.0 | More common in continental waters; schooling. |
| | <i>Haemulon plumieri</i> | HF1 | HF1 | HF1 | HF1 | | W | 35 | O | | 15.2 | Common reef fish. |
| | <i>Haemulon sciurus</i> | HF1 | HF1 | HF1 | | | W | 30 | O | | 17.7 | One of most common and most colorful WI grunts. |
| | <i>Haemulon steindachneri</i> | | + | | HF1 | | 0 | 25 | E | | | |
| <i>Haemulon striatum</i> | | | | | | HF1 | | 100 | S | | | Prefers deeper water than other species of genus. |
| <i>Orthopristis chrysoptera</i> | | | + | | HF1 | | ++ | | | | | Only occasionally on reefs. |
| <i>Orthopristis poeyi</i> | | | + | | HF1 | | ++ | | | | | Cuba only. |
| <i>Orthopristis ruber</i> | HF1 | | + | | HF1 | | ++ | 70 | | | | Southern Caribbean; also in brackish water. |
| <i>Pomadasy corvinaeformis</i> | | | HF1 | | HF1 | | ++ | | S | | | Cuba is northern limit of distribution. |
| <i>Pomadasy crocro</i> | HF1 | | | | HF1 | | ++ | | S | | | Also quite common in brackish, mangrove-lined lagoons; anadromous. |
| Hemiramphidae | <i>Hemiramphus balao</i> | | + | HF4 | HF4 | | | | F | | | Inshore surface dwelling fish forming sizeable schools. |
| | <i>Hemiramphus brasiliensis</i> | | + | HF4 | HF4 | | V | | O | | | Inshore surface dwelling fish forming sizeable schools. |
| | <i>Hyporhamphus unifasciatus</i> | | + | HF4 | HF4 | | | | O | | | Inshore surface schooling species, frequently entering estuaries. |
| Holocentridae | <i>Holocentrus ascensionis</i> | + | H1 | F1 | F1 | H1 | SH | 30 | O | | (13.7) | Nocturnal. |
| | <i>Holocentrus coruscus</i> | + | H1 | F1 | F1 | H1 | SH | 30 | O | R | (13.7) | Nocturnal. |
| | <i>Holocentrus rufus</i> | + | H1 | F1 | F1 | H1 | W | 32 | C | V | (13.7) | Nocturnal; more abundant on well-developed reefs than <i>H. ascensionis</i> . |
| | <i>Myripristis jacobus</i> | + | HF1 | | | HF1 | SH | 25 | O | R | (13.7) | Nocturnal; aggregating around coral reefs. |
| | <i>Neoniphon marianus</i> | + | H1 | F1 | F1 | H1 | W | 70 | O | R | (13.7) | Rare in shallow water, maybe most common of family in 35-70 m depth; nocturnal. |

continued...

Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|------------|---------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|-------------------------------------------------------------------------------------------------------------------------|
| | <i>Plectrypops retrospinis</i> | | HF1 | | | | W | | S | | (13.7) | Nocturnal; secretive. |
| | <i>Sargocentron vexillarium</i> | + | H1 | F1 | F1 | H1 | SH | | O | | (13.7) | Most common inshore squirrelfish in WI. |
| Kyphosidae | <i>Kyphosus incisor</i> | | HF1 | HF1 | | | | SH | O | | (25.4) | Sometimes in floating <i>Sargassum</i> weed. |
| | <i>Kyphosus sectatrix</i> | | HF1 | HF1 | HF1 | | | SH | O | | (25.4) | Young commonly found in floating <i>Sargassum</i> weed. |
| Labridae | <i>Bodianus rufus</i> | | HF1 | | | | W | 35 | O | | 20.3 | Moderately large. |
| | <i>Clepticus parrae</i> | | HF3 | F3 | F3 | | W | | O | | | Reef for shelter, feeding in midwater on zooplankton. |
| | <i>Halichoeres bivittatus</i> | | + | HF1 | HF1 | | SH | | O | V | | Most common of genus. |
| | <i>Halichoeres garnoti</i> | | HF1 | | HF1 | | W | | S | V | | Not uncommon. |
| | <i>Halichoeres maculipinna</i> | | HF1 | | | | SH | | O | T | | |
| | <i>Halichoeres poeyi</i> | | + | HF1 | | | SH | | C | | | |
| | <i>Halichoeres radiatus</i> | | HF1 | HF1 | HF1 | | SH | 50 | O | | | Largest of genus and second largest of family in WI. |
| | <i>Lachnolaimus maximus</i> | | HF1 | HF1 | HF1 | | SH | | S | | | Largest tropical wrasse, open areas near reefs (gorgonians). |
| | <i>Thalassoma bifasciatum</i> | | HF1 | | | | | SH | C | R | | One of most abundant WI reef fishes. |
| | <i>Xyrichtys novacula</i> | | | | HF1 | | 1 | 90 | F | | | |
| | <i>Xyrichtys splendens</i> | | | | HF1 | | SH | | O | | | Smaller than <i>X. novacula</i> . |
| Lutjanidae | <i>Apsilus dentatus</i> | | HF2 | | | | | 120-180 | | | | Mainly rocky bottoms; young may be found near surface. |
| | <i>Lutjanus analis</i> | | + | HF2 | + | | W | | O | | 55.8 | More open water over vegetated sand bottom; forms small aggregations. |
| | <i>Lutjanus apodus</i> | HF2 | HF2 | HF2 | HF2 | | | SH | C | | 19.0 | Most common of family on WI coral reefs; forms aggregations by day. |
| | <i>Lutjanus buccanella</i> | | HF2 | | HF2 | | ++ | 150 | E | | | Young in 35-50 m; adults between 80-150 m. |
| | <i>Lutjanus campechanus</i> | | HF2 | | | | 10 | 190 | E | | | Mainly in 30-130 m; juveniles in shallow waters and over sand or mud. |
| | <i>Lutjanus cyanopterus</i> | HF2y | HF2 | | HF2 | | W | 40 | S | T | | Large near rocks and reefs; small often on mangrove-lined coasts; least common of shallow water species of WI. |
| | <i>Lutjanus griseus</i> | HF2 | HF2 | HF2 | HF2 | HF2 | W | 180 | S-O | | 26.6 | Various habitats (rock-mangrove). |
| | <i>Lutjanus jocu</i> | HF2y | HF2 | | HF2y | | W | | O | | 20.3 | Adults in reefs; young in estuaries and coastal waters. |
| | <i>Lutjanus mahagoni</i> | HF2y | HF2 | + | + | | SH | | O | | 15.2 | Coral reef species in clear water. |
| | <i>Lutjanus synagris</i> | | HF2 | HF2 | HF2 | | W | 400 | F | | | Forms large aggregations, especially during the spawning period. |

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Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|----------------|----------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------|
| | <i>Lutjanus vivanus</i> | | | | | HF2 | ++ | >200 | S | | | Very common between 90 and 140 m, near shelf edge; not a real reef species. |
| | <i>Ocyurus chrysurus</i> | HF2 | HF2 | HF2y | | | 10 | 70 | C | V-T | 17.7 | One of the most common and active shallow-water reef fishes of the Caribbean. |
| Malacanthidae | <i>Malacanthus plumieri</i> | | + | | HF1 | | 10-50 | 153 | O | | | Sand and rubble near reefs; only shallow-water species of family in West Atlantic. |
| Megalopidae | <i>Tarpon atlanticus</i> | HF | | | | | | | O | | | |
| Monacanthidae | <i>Aluterus monoceros</i> | | | | | | | | F | | | |
| | <i>Aluterus schoepfii</i> | | + | HF1,3 | HF1,3 | | ++ | 50 | S | | | |
| | <i>Aluterus scripta</i> | | + | HF1 | | | SH | | S | | | |
| | <i>Cantherines macrocerus</i> | | + | | | | SH | | S | | | |
| | <i>Cantherines pullus</i> | | + | | | | SH | 50 | O | | | Most common filefish in WI. |
| | <i>Monacanthus ciliatus</i> | | HF1 | HF1 | HF1 | | SH | 50 | O | | | Fish of vegetated areas. |
| | <i>Monacanthus tuckeri</i> | | + | | | | SH | | O | V | | Smallest of WI filefishes. |
| | <i>Stephanolepis setifer</i> | | | HF2 | HF2 | | ++ | 80 | O | | | |
| Mugilidae | <i>Mugil curema</i> | HF | | | HF3 | | ++ | | O | | 25.4 | Typical in murky waters (estuaries, mangroves, lagoons) but also in clear waters close to the surface. |
| Mullidae | <i>Mulloidichthys martinicus</i> | + | H1 | HF1y | HF1 | | SH | | O | | 16.5 | Solitary or in small groups. |
| | <i>Pseudupeneus maculatus</i> | + | H1 | HF1y | HF1 | | SH | 50 | O | V | 13.7 | Solitary, young in seagrass beds. |
| Muraenidae | <i>Enchelycore nigricans</i> | | HF1 | | | | SH | >12 | S | | | Mainly in very shallow water. |
| | <i>Echidna catenata</i> | | HF1 | | | | SH | | S | | | Very shallow water beneath rocks; feeds mainly on crabs. |
| | <i>Gymnothorax funebris</i> | HF1 | HF1 | | HF1 | | SH | 5 | O | | | Also in mangroves. |
| | <i>Gymnothorax vicinus</i> | | HF1 | HF1 | | | SH | | O | | | Secretive. |
| | <i>Gymnothorax miliaris</i> | | HF1 | | | | SH | 50 | S | R | | Secretive. |
| | <i>Lycodontis moringa</i> | | HF1 | | | | SH | 50 | O | V | | Most common moray in shallow water in WI. |
| Myliobatidae | <i>Aetobatus narinari</i> | | HF3 | HF3 | HF3 | | SH | | S | | | Coastal surface waters; visitor to reefs; solitary or large schools; feeds on bivalves (cockles, oysters), snails, shrimps, worms, cephalopods, small fish. |
| Ogcocephalidae | <i>Ogcocephalus nasutus</i> | | + | HF1 | HF1 | | ++ | | O | | | Occasionally on reefs. |
| Ophichthidae | <i>Myrichthys breviceps</i> | | + | HF1 | HF1 | | SH | | O | | | May be observed in reefs; burrowing in mud or sand. |

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Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|------------------|-------------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|-------------------------------------------------------------------------------|
| | <i>Myrichthys ocellatus</i> | | + | | | | SH | | O | | | Feeds primarily on crabs; burrowing in mud or sand. |
| | <i>Ophichthus ophis</i> | | | HF1 | | | SH | 50 | F | | | Burrowing in mud or sand. |
| Opisthognathidae | <i>Opisthognathus aurifrons</i> | | HF1 | | | | ++ | | F | | | |
| | <i>Opisthognathus macrognathus</i> | | HF1 | | | | ++ | | S | | | |
| | <i>Opisthognathus maxillosus</i> | | HF1 | | | | ++ | | O-F | | | Most common of family. |
| | <i>Opisthognathus whitehurstii</i> | | HF1 | | | | ++ | | O | | | |
| Ostraciidae | <i>Acanthostracion polygonius</i> | | HF1 | | | | W | | S | | (17.7) | Least common trunk fish in WI. |
| | <i>Acanthostracion quadricornis</i> | | | HF1 | | | SH | 80 | O | | (17.7) | |
| | <i>Lactophrys bicaudalis</i> | | HF1 | HF1 | | | SH | 50 | S | | (17.7) | |
| | <i>Lactophrys trigonus</i> | | | HF1 | | | SH | 50 | S-O | | (17.7) | Largest of genus. |
| | <i>Lactophrys triqueter</i> | | HF1 | F1 | F1 | | SH | 50 | O | R | (17.7) | |
| Pempheridae | <i>Pempheris schomburgki</i> | | H1F2 | | | | SH | | O | | | Nocturnal. |
| | <i>Pempheris poeyi</i> | | H1F2 | | | | | | O | | | Nocturnal. |
| Pomacanthidae | <i>Centropyge argi</i> | | HF1 | | | | D | >30 | O | | | Not common in shallow water, moderately common in 30 m depth. |
| | <i>Holacanthus ciliaris</i> | | HF1 | | | | SH | | O | R | (20.3) | |
| | <i>Holacanthus tricolor</i> | | HF1 | | | | SH | | O | | 15.2 | Clear reef areas. |
| | <i>Pomacanthus arcuatus</i> | + | HF1 | | | | SH | | O | T | 27.9 | Usually in pairs. |
| | <i>Pomacanthus paru</i> | | HF1 | | | | SH | | O | | (20.3) | Young are often cleaners. |
| Pomacentridae | <i>Abudefduf saxatilis</i> | | HF1 | | | | SH | <15 | O | | | |
| | <i>Abudefduf taurus</i> | | HF1 | HF1 | | | ++ | <5 | O | | | Largest of WI pomacentrids. |
| | <i>Chromis cyanea</i> | | H1F3 | | | | W | | O | R | | |
| | <i>Chromis multilineata</i> | | H1F3 | | | | W | >40 | O | R | | |
| | <i>Microspathodon chrysurus</i> | | HF1 | | | | SH | 7-10 | O | | | Second in size to <i>A. taurus</i> ; one of most common reef fishes in WI. |
| | <i>Pomacentrus partitus</i> | | HF1 | | | | | | O | R | | |
| | <i>Stegastes fuscus</i> | | + | | | | SH | | C | | | One of most abundant of all fish. |
| | <i>Stegastes leucostictus</i> | | + | | | | SH | | C | | | Very common. |
| | <i>Stegastes planifrons</i> | | + | | | | SH | | S | R | | Abundant. |
| | <i>Stegastes variabilis</i> | | + | | | | SH | | O | R | | Abundant. |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | | HF2 | | | | SH | - | S | | | Nocturnal, solitary. |
| | <i>Priacanthus arenatus</i> | | HF2 | | | | SH | 75 | O | | | Nocturnal, occurs in small aggregations. |
| Rachycentridae | <i>Rachycentron canadum</i> | | HF3 | | | HF3 | W | | S | | | Clear water reef areas; mainly pelagic. |
| Scaridae | <i>Cryptotomus roseus</i> | | | + | + | | 8 | 60 | | | (13.7) | |
| | <i>Nicholsina usta</i> | | | + | | | ++ | 83 | | | (13.7) | |
| | <i>Scarus coelestinus</i> | | H1 | F1 | | | SH | <20 | F | | (13.7) | Second largest after <i>S. guacamaia</i> . |
| | <i>Scarus coeruleus</i> | | + | + | | | SH | | O | | (13.7) | |

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Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|--------------|---------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|-------------------------------------------------------------------------|
| | <i>Scarus guacamaia</i> | HF1y | HF1 | F1 | | | W | | O-F | | (13.7) | Adults mainly on reefs, young often in mangrove areas. |
| | <i>Scarus iserti</i> | | HF1 | F1 | | | SH | | F | V | (13.7) | Most common of genus in WI and smallest of genus in Atlantic. |
| | <i>Scarus taeniopterus</i> | | HF1 | | | | W | | O | T | (13.7) | |
| | <i>Scarus vetula</i> | | HF1 | HF1 | | | SH | | A | R | (13.7) | One of the most common scarids in WI reefs. |
| | <i>Sparisoma aurofrenatum</i> | | HF1 | HF1 | + | | W | | F | V | (13.7) | Young mainly in seagrass beds. |
| | <i>Sparisoma chrysopterygum</i> | | HF1 | HF1 | + | | SH | | F | | (13.7) | Second in abundance to <i>S. radians</i> in WI reefs. |
| | <i>Sparisoma radians</i> | | | HF1 | | | SH | | O | | (13.7) | Smallest of genus. |
| | <i>Sparisoma rubripinne</i> | | HF1 | HF1 | | | SH | | F | | (13.7) | Most common of genus in inshore portions of WI reefs. |
| | <i>Sparisoma viride</i> | | HF1 | HF1 | | | SH | | C | V | (13.7) | Young in vegetated areas. |
| Sciaenidae | <i>Equetus lanceolatus</i> | | H1F2 | | + | | SH | | O | | | Found in deeper water than other members of genus. |
| | <i>Equetus punctatus</i> | | HF | | | | SH | | S | | | Largest of genus in WI; secretive by day in reefs and usually solitary. |
| | <i>Odontoscion dentex</i> | + | H1F2 | | F2 | | SH | | O | | | Nocturnal, hides in caves by day. |
| | <i>Pareques acuminatus</i> | | HF | | | | SH | | O | | | Small groups; hides under rock ledges by day. |
| Scombridae | <i>Euthynnus aletteratus</i> | | | | | F4 | W | | O | | | Green shelf waters, forms compact schools. |
| | <i>Scomberomorus cavalla</i> | | F1,3 | | F1,3 | F1,3 | W | | O | | 81.2 | Largest of genus, midwater but often near bottom. |
| | <i>Scomberomorus maculatus</i> | | | | | | | | | | 38.1 | |
| | <i>Scomberomorus regalis</i> | | HF | | | | W | | O | T | | Most common member of genus; clear inshore waters. |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | | HF1 | | HF1 | | ++ | 100 | F | | | |
| | <i>Scorpaena grandicornis</i> | | | HF1 | | | SH | | S-O | | | |
| | <i>Scorpaena inermis</i> | | HF1 | HF1 | | | W | | S | | | |
| | <i>Scorpaena plumieri</i> | | HF1 | | | | SH | | O | | | Largest and most common of genus on shallow WI reefs. |
| | <i>Scorpaenodes caribbaeus</i> | | HF1 | | | | SH | | O | | | Common inshore fish but rarely seen. |
| Serranidae | <i>Alphesthes afer</i> | | HF1 | HF1 | | | SH | | S-O | | 17.7 | Much more common in seagrass beds than on reefs. |
| | <i>Cephalopholis cruentata</i> | | HF1 | | | | W | >70 | O | R | 19.0 | Most abundant on reefs with ledges and caves. |
| | <i>Cephalopholis fulva</i> | | HF1 | | | | W | 40 | O | | 19.0 | One of the most common groupers on WI reefs. |
| | <i>Dermatolepis inermis</i> | | HF1 | | | | | | O | | | Secretive. |

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Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|--------|------------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|-------------------------------------------------------------------------------------------------------------|
| | <i>Diplectrum formosum</i> | | | | HF1 | | 1 | 50 | S | | | At the bases of reefs, known in WI only from Cuba and islands of the southern Caribbean. |
| | <i>Epinephelus adscensionis</i> | | HF1 | | | | SH | >30 | O | | | Most common in shallow water on rocky bottom. |
| | <i>Epinephelus guttatus</i> | | HF1 | | | | W | | O | | 17.7 | Most common of genus in WI. |
| | <i>Epinephelus itajara</i> | | HF1 | | HF1 | | ++ | | O | | | Largest of family. |
| | <i>Epinephelus morio</i> | | HF1 | | | | 5 | 150 | S | | | Abundant in Florida and Gulf of Mexico. |
| | <i>Epinephelus striatus</i> | | HF1 | HF1 | | | SH | 90 | O | R | 31.7 | One of the most important commercial fishes of WI; young in seagrass beds. |
| | <i>Hypoplectrus aberrans</i> | | HF1 | | | | SH | | O | | | |
| | <i>Hypoplectrus chlorurus</i> | | HF1 | | | | SH | | O | | | |
| | <i>Hypoplectrus nigricans</i> | | HF1 | | | | SH | | O | | | Largest of genus. |
| | <i>Hypoplectrus puella</i> | | HF1 | | | | SH | | C | | | Most common of genus in WI. |
| | <i>Mycteroperca bonaci</i> | | HF1 | | | | W | | S | | 29.2 | Rare in Virgin Islands and Puerto Rico, common in the Bahamas, Cuba, Florida Keys and southern Caribbean. |
| | <i>Mycteroperca cidi</i> | | HF1 | | HF1y | | W | >20 | E | | | Common inshore species at coast of Venezuela and offshore islands, juveniles over sand, larger >20 m depth. |
| | <i>Mycteroperca interstitialis</i> | HF1 | HF1 | | HF1y | | W | >55 | S | | | Small- and middle-sized individuals occur in mangrove-lined lagoons. |
| | <i>Mycteroperca microlepis</i> | | HF1 | HF1 | | | 48 | 80 | E | | | Depth refers to catch on Campeche Bank. |
| | <i>Mycteroperca phenax</i> | | HF1 | | | | 64 | 90 | E | | | Depth refers to catch on Campeche Bank. |
| | <i>Mycteroperca rubra</i> | HF1y | HF1 | | HF1y | | | | S | | | Rare in the north, common to the south; also common (especially young) in mangrove-lined lagoons. |
| | <i>Mycteroperca tigris</i> | | HF1 | | | | W | >30 | O | | 30.4 | |
| | <i>Mycteroperca venenosa</i> | | HF1 | | | | W | >80 | S | | | Most common species of genus generally in WI. |
| | <i>Paranthias furcifer</i> | | | | | H2F3 | D | 70 | S | | | Feeds on zooplankton in midwater, forms small schools. |
| | <i>Serranus dewegeri</i> | | HF1 | HF1y | | | ++ | | E | | | Southern Caribbean; one of the most abundant inshore fishes of rocky bottom around islands off Venezuela. |
| | <i>Serranus tabacarius</i> | | HF1 | | | | 5 | >20 | O | | | |
| | <i>Serranus tigrinus</i> | | HF1 | | | | W | | S | | | Most common species of genus on WI reefs but nowhere abundant. |
| | <i>Serranus tortugarum</i> | | + | | HF1 | | | | O | | | Small groups, hovering over a patch of coral rubble or an old conch shell. |

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Table 3.1 continued

| Family | Species | Mangroves/ estuaries | Coral reefs | Seagrass beds/ algal mats | Sand and mudflats | Offshore reef | Average depth/ upper limit | Depth limit (m) | Abundance | Type | Average size* (cm) | Remarks |
|----------------|---------------------------------|-------------------------|----------------|---------------------------------|-------------------------|------------------|----------------------------------|-----------------------|-----------|------|--------------------------|----------------------------------------------------------------------------------------------------------------------------|
| Sparidae | <i>Archosargus rhomboidalis</i> | HF1 | + | HF1 | | | | | S-O | | 7.6 | Rare on coral reefs; more common near mangroves and on vegetated areas. |
| | <i>Calamus bajonado</i> | | HF1 | HF1 | HF1 | | 6 | 45 | S | | (17.7) | Largest of genus; clear water; large adults usually solitary. |
| | <i>Calamus calamus</i> | | HF1 | HF1y | HF1y | | 6 | 75 | O | | (17.7) | |
| | <i>Calamus penna</i> | | + | + | | | 3 | 87 | S | | (17.7) | |
| | <i>Calamus pennatula</i> | | HF1 | HF1 | HF1 | | W,Sy | 85 | S-O | T | (17.7) | Most common of genus in WI; young in shallower waters. |
| | <i>Diplodus caudimaculata</i> | | HF1 | | HF1 | | | ++ | O | | (17.7) | Shallow, calm waters; young may form aggregations. |
| Sphyraenidae | <i>Sphyraena barracuda</i> | HF3y | HF3 | HF3y | HF3y | HF3 | W | | O | | 76.2 | Openwater species; young in aggregations, larger, solitary over reefs. |
| | <i>Sphyraena picudilla</i> | | HF2 | HF2 | HF2 | | W | | S | | | Small schooling species, mainly over seagrass beds; uncommon in VI. |
| Sphyrnidae | <i>Sphyrna lewini</i> | | + | + | + | + | W | | S | | | Young occur mostly in coastal waters; feed on pelagic fish (including smaller sharks and rays), squids, crabs, sting rays. |
| | <i>Sphyrna tiburo</i> | | HF1 | | HF1 | | 10-25 | 80 | O | | | Often preyed upon by larger sharks; feeds on crabs and shrimps, less on clams, octopus and small fishes. |
| Syngnathidae | <i>Syngnathus</i> sp. | | + | | | | | | | R | | |
| Synodontidae | <i>Synodus foetens</i> | | | | HF1 | | | ++ | O | | | Not a reef fish. |
| | <i>Synodus intermedius</i> | | HF1 | | HF1 | | | ++ | F | V | | Largest and most common of family in WI. |
| | <i>Synodus synodus</i> | | | | HF1 | | | ++ | O | | | |
| Tetraodontidae | <i>Canthigaster rostrata</i> | | HF1 | HF1 | | | W | | O-C | R | | Solitary. |
| | <i>Sphoeroides spengleri</i> | | HF1 | HF1 | HF1 | | SH | 30 | O | | | |
| | <i>Sphoeroides testudineus</i> | HF1 | | | HF1 | | ++ | | S | | | One of most common species in mangrove areas and estuarine coastlines; may form huge aggregates. |
| Triakidae | <i>Mustelus canis</i> | | + | + | HF1 | | W | >150 | S | | | Coastal waters, occasionally in fresh water; on muddy bottom; feeds on crabs, lobsters, small fish and molluscs. |

Table 3.2. Number of fishers and vessels in Puerto Rico and U.S. Virgin Islands 1970-1981 (reproduced from Table 6 in CFMC 1985).

| Year | Number of fishers | | Number of vessels | |
|------|-------------------|----------------|-------------------|----------------|
| | Puerto Rico | Virgin Islands | Puerto Rico | Virgin Islands |
| 1970 | 1,082 | 400 | 869 | - |
| 1971 | 994 | - | 811 | - |
| 1972 | 968 | - | 797 | - |
| 1973 | 927 | - | 785 | - |
| 1974 | 1,182 | - | 835 | - |
| 1975 | 1,230 | 450 | 902 | - |
| 1976 | - | 509 | - | - |
| 1977 | 1,368 | 846 | 1,036 | - |
| 1978 | 1,442 | 265 | 1,073 | 231 |
| 1979 | - | 281 | - | 223 |
| 1980 | 1,447 | 355 | 1,084 | 237 |
| 1981 | - | 397 | - | - |
| 1982 | 1,872 | 578 | 1,449 | - |

-.Data not available.

Table 3.3. Relative importance of the different gear in the fisheries of the PRVI area (adapted from Table 5 in CFMC 1985).

| Gear/ product | Number of units | Puerto Rico (1980) | | US Virgin Islands (1979-1980) | |
|------------------|--------------------|--------------------|----------------|-------------------------------|----------------|
| | | Catch (t) | % ^a | Catch (t) | % ^a |
| Fish trap | 12,586 | 1,268 | 42.0 | 1,098 | 66.0 |
| Lobster pot | 2,252 | 29 | 1.0 | 109 | 6.5 |
| Fishnet | - | - | - | 215 | 13.0 |
| Beach seine | 238 | 249 | 8.2 | - | - |
| Gillnet | 870 | 264 | 8.7 | - | - |
| Castnet | 827 | 19 | 0.6 | - | - |
| Hookfish | - | - | - | 147 | 8.8 |
| Handline | 2,391 | 635 | 21.0 | - | - |
| Troll line | 2,057 | 209 | 6.9 | - | - |
| Trotline | 331 | 11 | 0.4 | - | - |
| Spear | 341 | 168 | 5.6 | 20 | 1.2 |
| Hand | - | 170 | 5.6 | - | - |
| Hand lobster | - | - | - | 11 | 0.7 |
| Conch | - | - | - | 57 | 3.4 |
| Whelk | - | - | - | 6 | 0.4 |
| Total | - | 3,022 | 100.0 | 1,663 | 100.0 |

^aPercent of total catch from all gears.

maculatus) and yellow goatfish (*Mulloidichthys martinicus*). Based on biostatistical surveys, Jacobsen and Browder (1987) calculated that these 10 species contribute approximately 40% to Virgin Islands landings, by weight, and 45% to Puerto Rico landings. Table 3.4 gives the average annual catch per trap for Puerto Rico for three years. The landings per year were divided by the total number of traps assuming that the number of hauls remained constant. There was a decline in catch/effort for all the families listed.

Boulon (1986a) listed fish species and invertebrates of commercial importance in the Virgin Islands Biosphere Reserve (Table 3.1). The only catch data available on over 50 individual species are reported in Boulon and Clavijo (1986) based on a study of three commercial fishpot fishers on St. John, US Virgin Islands, from January to June 1984.

Table 3.5. Estimated asymptotic length (L_{∞}) and weight (W_{∞}) of Caribbean reef fishes.

| Family | Species | L_{∞} (cm) | | Source | W_{∞} (g) | Source |
|------------------------------|------------------------------------|----------------------------|-----------|---------------------------------------------------------------------------------|----------------------------|---------------------------------------------------------------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 25 | FL | Pauly (1978) | 288 | Randall (1962, 1967, 1968) |
| | <i>Acanthurus chirurgus</i> | 33 | FL | Pauly (1978) | 820 | Munro (1983) |
| | <i>Acanthurus coeruleus</i> | 32 | FL | Pauly (1978) | 983 | Munro (1983) |
| Albulidae | <i>Albula vulpes</i> | 82 | | Beebe and Tee-Van (1928) | 7,150 | Beebe and Tee-Van (1928), Randall (1962, 1967, 1968) |
| Antennariidae | <i>Antennarius multiocellatus</i> | 15 | (-) | Randall (1962, 1967, 1968) | 70 | * |
| | <i>Antennarius striatus</i> | 15 | / | Figuereido and Menezes (1977); Randall (1962, 1967, 1968) | 70 | * |
| Apogonidae | <i>Apogon conklini</i> | 5 | SL/ | Randall (1962, 1967, 1968) | 3.3 | Smith and Tyler (1972) |
| | <i>Apogon maculatus</i> | 10 | / | Randall (1962, 1967, 1968) | 37 | Smith and Tyler (1972) |
| Atherinidae | <i>Allanetta harringtonensis</i> | 8 | / | Randall (1962, 1967, 1968) | 11 | Beebe and Tee-Van (1928) |
| | <i>Atherinomorus stipes</i> | 9 | + | Randall (1962, 1967, 1968) | 16 | Beebe and Tee-Van (1928) |
| Aulostomidae | <i>Aulostomus maculatus</i> | 76 | / | Randall (1962, 1967, 1968) | 777 | Beebe and Tee-Van (1928) |
| Balistidae | <i>Balistes capricus</i> | 31 | | Johnson and Solomon (1984); Randall (1962, 1967, 1968) | 611 | Polovina and Ralston (1987) |
| | <i>Balistes vetula</i> | 45 | FL | Munro (1983); Randall (1962, 1967, 1968) | 2,586 | Randall (1962, 1967, 1968) |
| | <i>Canthidermis sufflamen</i> | 56 | + | Randall (1962, 1967, 1968) | 4,783 | Munro (1983) |
| | <i>Melichthys niger</i> | 36 | | Randall (1962, 1967, 1968) | 960 | * |
| | <i>Xanthichthys ringens</i> | 25 | | Randall (1962, 1967, 1968) | 320 | * |
| Belontiidae | <i>Ablennes hians</i> | 87 | | Fischer (1978) | 751 | Beebe and Tee-Van (1928) |
| | <i>Platybelone argalus argalus</i> | 51 | / | Randall (1962, 1967, 1968) | 150 | * |
| | <i>Strongylura marina</i> | 64 | | Fischer (1978) | 300 | * |
| | <i>Strongylura timucu</i> | 46 | / | Randall (1962, 1967, 1968) | 110 | * |
| | <i>Tylosurus acus</i> | 135 | SL | Fischer (1978) | 2,800 | * |
| | <i>Tylosurus crocodilus</i> | 107 | SL | Fischer (1978) | 1,400 | * |
| Blenniidae | <i>Entomacrodus nigricans</i> | 8.5 | | Randall (1962, 1967, 1968) | 13 | * |
| | <i>Ophioblennius atlanticus</i> | 12 | / | Randall (1962, 1967, 1968) | 35 | * |
| | <i>Parablennius marmoratus</i> | 8 | | Randall (1962, 1967, 1968) | 10 | * |
| | <i>Scartelia cristata</i> | 11 | | Randall (1962, 1967, 1968) | 27 | * |
| Bothidae | <i>Bothus lunatus</i> | 46 | / | Randall (1962, 1967, 1968) | 3,000 | * |
| | <i>Bothus ocellatus</i> | 18 | | Randall (1962, 1967, 1968) | 180 | * |
| | <i>Paralichthys tropicus</i> | 54 | % | Randall (1962, 1967, 1968) | 5,000 | * |
| Carangidae | <i>Alectis ciliaris</i> | 91 | TL/ | Randall (1962, 1967, 1968) | 10,700 | Randall (1962, 1967, 1968) |
| | <i>Caranx bartholomaei</i> | 81 | TL | Munro (1983) | 9,862 | Munro (1983) |
| | <i>Caranx crysos</i> | 58 | TL | Munro (1983); Randall (1962, 1967, 1968) | 2,314 | Munro (1983) |
| | <i>Caranx hippos</i> | 95 | TL | Munro (1983) | 8,490 | Randall (1962, 1967, 1968) |
| | <i>Caranx latus</i> | 80 | % | Randall (1962, 1967, 1968) | 6,250 | Randall (1962, 1967, 1968) |
| | <i>Caranx lugubris</i> | 80 | % | Randall (1962, 1967, 1968) | 6,554 | Randall (1962, 1967, 1968) |
| | <i>Caranx ruber</i> | 56 | % | Randall (1962, 1967, 1968) | 3,160 | Munro (1983) |
| | <i>Decapterus macarellus</i> | 35 | SL | Fischer (1978) | 750 | * |
| | <i>Decapterus punctatus</i> | 32 | | Beebe and Tee-Van (1928) | 573 | Beebe and Tee-Van (1928) |
| | <i>Elagatis bipinnulatus</i> | 107 | FL | Fischer (1978) | 10,500 | Fischer (1978) |
| | <i>Oligoplites saurus</i> | 31 | FL% | Fischer (1978) | 334% | Fischer (1978) |
| | <i>Selar crumenophthalmus</i> | 40 | TL | Figuereido and Menezes (1977); Randall (1962, 1967, 1968) | 1,240 | Beebe and Tee-Van (1928) |
| | <i>Selene vomer</i> | 50 | TL | Figuereido and Menezes (1977) | 1,400 | * |
| | <i>Seriola dumerilii</i> | 194 | | Randall (1962, 1967, 1968) | 80,000 | Figuereido and Menezes (1977); Fischer (1978); Randall (1962, 1967, 1968) |
| | <i>Seriola rivoliana</i> | 97 | | Fischer (1978) | 24,000 | Fischer (1978); Randall (1962, 1967, 1968) |
| | <i>Trachinotus falcatus</i> | 105 | FL | Fischer (1978) | 36,000 | Fischer (1978) |
| | <i>Trachinotus goodii</i> | 50 | | Figuereido and Menezes (1977); Fischer (1978); Randall (1962, 1967, 1968) | 3,900 | * |
| Carcharhinidae | <i>Carcharhinus acronotus</i> | 200 | L_{max} | Fischer (1978) | 90,000 | * |
| | <i>Carcharhinus falciformis</i> | 350 | L_{max} | Fischer (1978) | 350,000 | * |
| | <i>Carcharhinus leucas</i> | 350 | L_{max} | Fischer (1978) | 350,000 | Clark and von Schmidt (1965) |
| | <i>Carcharhinus limbatus</i> | 247 | L_{max} | Fischer (1978) | 116,000 | Clark and von Schmidt (1965) |
| | <i>Carcharhinus longimanus</i> | 350 | L_{max} | Fischer (1978) | 350,000 | * |
| | <i>Carcharhinus perezii</i> | 230 | L_{max} | Fischer (1978) | 100,000 | * |
| | <i>Galeocerdo cuvier</i> | 457 | | Pauly (1978) | 672,000 | Clark and von Schmidt (1965) |
| | <i>Negaprion brevirostris</i> | 284 | | Pauly and Palomares (1986) | 110,000 | Collette and Nauen (1983) |
| | <i>Rhizoprionodon porosus</i> | 107 | | Randall (1962, 1967, 1968) | 5,020 | Randall (1962, 1967, 1968) |
| | Chaetodontidae | <i>Chaetodon aculeatus</i> | 9.5 | % | Randall (1962, 1967, 1968) | 19 |
| <i>Chaetodon capistratus</i> | | 14.2 | | Munro (1983) | 120 | * |
| <i>Chaetodon ocellatus</i> | | 16.8 | | Munro (1983) | 200 | * |
| <i>Chaetodon sedentarius</i> | | 14.7 | % | Randall (1962, 1967, 1968) | 130 | * |
| <i>Chaetodon striatus</i> | | 15.8 | | Munro (1983) | 164 | Beebe and Tee-Van (1928) |

continued...

Table 3.5 continued

| Family | Species | L _∞ (cm) | Source | W _∞ (g) | Source |
|------------------------------|------------------------------------|------------------------|----------------------------------------------------------------------------|-----------------------|------------------------------------------------------------|
| Cirrhitidae | <i>Amblycirrhitus pinos</i> | 8 (SL) | Beebe and Tee-Van (1928) | 15 | Smith and Tyler (1972) |
| Clinidae | <i>Labrisomus nuchipinnis</i> | 20 | Randall (1962, 1967, 1968) | 137 | Smith and Tyler (1972) |
| Clupeidae | <i>Harengula clupeola</i> | 18 % | Figuereido and Menezes (1977) | 240 | * |
| | <i>Harengula humeralis</i> | 22 | Randall (1962, 1967, 1968) | 440 | * |
| | <i>Jenkinsia lamprotaenia</i> | 6.8 % | Fischer (1978) | 13 | * |
| | <i>Opisthonema oglinum</i> | 26 % | Fischer (1978) | 730 | Beebe and Tee-Van (1928) |
| Congridae | <i>Heteroconger halis</i> | 48 | Randall (1962, 1967, 1968) | 150 | * |
| Coryphaenidae | <i>Coryphaena hippurus</i> | 151 | Pauly and Palomares (1986) | 147,000 | Pauly and Palomares (1986) |
| Dactylopteridae | <i>Dactylopterus volitans</i> | 45 + | Fischer (1978) | 1,825 | Beebe and Tee-Van (1928); Figuereido and Menezes (1977) |
| Dasyatidae | <i>Dasyatis americana</i> | 300 | Fischer (1978) | 122,000 | * |
| | <i>Urolophus jamaicensis</i> | 76 | Randall (1962, 1967, 1968) | | |
| Diodontidae | <i>Chilomycterus antennatus</i> | 26.3 % | Randall (1962, 1967, 1968) | 1,420 | Beebe and Tee-Van (1928) |
| | <i>Chilomycterus antillarum</i> | 26.3 % | Randall (1962, 1967, 1968) | 1,420 | * |
| | <i>Diodon holocanthus</i> | 51 | Randall (1962, 1967, 1968) | 11,300 | * |
| | <i>Diodon hystrix</i> | 60 % | Randall (1962, 1967, 1968) | 18,446 | Beebe and Tee-Van (1928) |
| Echeneidae | <i>Echeneis naucrates</i> | 85 % | Randall (1962, 1967, 1968) | 2,080 | Randall (1962, 1967, 1968) |
| | <i>Remora remora</i> | 79 | Randall (1962, 1967, 1968) | 1,700 | * |
| Elopidae | <i>Elops saurus</i> | 47 | Pauly (1978) | 444 | Beebe and Tee-Van (1928) |
| Emmelichthyidae | <i>Inermia vittata</i> | 23 (SL)/ | Randall (1962, 1967, 1968) | 282 | Smith and Tyler (1972) |
| Engraulidae | <i>Anchoa hepsetus</i> | 15 | Fischer (1978) | 50 | Beebe and Tee-Van (1928); |
| | <i>Anchoa lyolepis</i> | 8 | Fischer (1978); Randall (1962, 1967, 1968) | 8 | Figuereido and Menezes (1977) |
| | <i>Anchoa mitchilli</i> | 10 | Fischer (1978) | 15 | * |
| | <i>Anchoa parva</i> | 6 | Fischer (1978) | 3 | * |
| Ephippidae | <i>Chaetodipterus faber</i> | 90 | Fischer (1978); Randall (1962, 1967, 1968) | 9,000 | Beebe and Tee-Van (1928); Randall (1962, 1967, 1968) |
| Exocoetidae | <i>Exocoetus volitans</i> | 18 / | Fischer (1978) | 90 | * |
| Fistulariidae | <i>Fistularia tabacaria</i> | 183 | Randall (1962, 1967, 1968) | 10,800 | * |
| Gerreidae | <i>Eucinostomus argenteus</i> | 30 % | Figuereido and Menezes (1977) | 290 | * |
| | <i>Eucinostomus lefroyi</i> | | | | |
| | <i>Gerres cinereus</i> | 40 | Beebe and Tee-Van (1928); Fischer (1978); Randall (1962, 1967, 1968) | 690 | Randall (1962, 1967, 1968) |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | 430 | Fischer (1978) | 500,000 | Figuereido and Menezes (1977) |
| Gobiidae | <i>Bathygobius soporator</i> | 15 | Randall (1962, 1967, 1968) | 70 | * |
| | <i>Coryphopterus glaucofraenum</i> | 8 (SL) | Randall (1962, 1967, 1968) | 10.5 | Smith and Tyler (1972) |
| | <i>Gnatholepis thompsoni</i> | 6 SL/ | Randall (1962, 1967, 1968) | 4.1 | Smith and Tyler (1972) |
| | <i>Gobiosoma evelynae</i> | 5 (SL) | Randall (1962, 1967, 1968) | 2.2 | Smith and Tyler (1972) |
| | <i>Gobiosoma horsti</i> | 5 (SL) | Randall (1962, 1967, 1968) | 3.1 | Smith and Tyler (1972) |
| Grammidae (Serranidae) | <i>Gramma loreto</i> | 8 + | Randall (1962, 1967, 1968) | 15 | * |
| | <i>Gramma melacara</i> | 10 - | Randall (1962, 1967, 1968) | 30 | * |
| | <i>Rypticus saponaceus</i> | 29.5 (SL%) | Randall (1962, 1967, 1968) | 685 | Smith and Tyler (1972) |
| Haemulidae (Pomadasyidae) | <i>Anisotremus sunnamiensis</i> | 63 TL% | Randall (1962, 1967, 1968) | 4,770 % | Randall (1962, 1967, 1968) |
| | <i>Anisotremus virginicus</i> | 30 / | Randall (1962, 1967, 1968) | 850 | Munro (1983) |
| | <i>Haemulon album</i> | 65 FL | Pauly (1980) | 5,300 | Munro (1983) |
| | <i>Haemulon aurolineatum</i> | 23 FL | Munro (1983) | 188 | Munro (1983) |
| | <i>Haemulon bonariense</i> | 30 FL | Munro (1983) | 385 | Munro (1983) |
| | <i>Haemulon boschmae</i> | 19 / | Fischer (1978); Randall (1962, 1967, 1968) | | |
| | <i>Haemulon carbonarium</i> | 38 % | Randall (1962, 1967, 1968) | 861 | Munro (1983) |
| | <i>Haemulon chrysargyreum</i> | 24 % | Randall (1962, 1967, 1968) | 251 | Munro (1983) |
| | <i>Haemulon flavolineatum</i> | 27 / | Randall (1962, 1967, 1968) | 480 | Munro (1983) |
| | <i>Haemulon macrostomum</i> | 45 % | Randall (1962, 1967, 1968) | 1,795 | Randall (1962, 1967, 1968) |
| | <i>Haemulon melanurum</i> | 30 FL | Munro (1983) | 427 | Munro (1983) |
| | <i>Haemulon parrai</i> | 41 | Randall (1962, 1967, 1968) | 1,265 | * |
| | <i>Haemulon plumieri</i> | 42 | Munro (1983) | 1,360 | Munro (1983) |
| | <i>Haemulon sciurus</i> | 41 % | Randall (1962, 1967, 1968) | 1,185 | Randall (1962, 1967, 1968) |
| | <i>Haemulon steindachneri</i> | 27 | Randall (1962, 1967, 1968) | 360 | * |
| | <i>Haemulon striatum</i> | 28 | Randall (1962, 1967, 1968) | 400 | * |
| | <i>Orthopristis chrysoptera</i> | 46 | Darcy (1983); Fischer (1978) | 900 | Darcy (1983) |
| | <i>Orthopristis poeyi</i> | 24 | Fischer (1978) | 130 | * |
| | <i>Orthopristis ruber</i> | 33 | Randall (1962, 1967, 1968) | 330 | * |

continued...

Table 3.5 continued

| Family | Species | L_{∞} (cm) | Source | W_{∞} (g) | Source |
|--------------------------------------|----------------------------------|-----------------------------|------------------------------------------------------------------------------|----------------------------|---------------------------------------------------------|
| Hemiramphidae | <i>Pomadasys corvinaeformis</i> | 25 | Fischer (1978) | 386 | Beebe and Tee-Van (1928) |
| | <i>Pomadasys crocro</i> | 33 | Figuereido and Menezes (1977) | 890 | * |
| | <i>Hemiramphus balao</i> | 40 | Fischer (1978) | 614 | Berkeley and Houde (1978) |
| | <i>Hemiramphus brasiliensis</i> | 32.5 | Pauly and Murphy (1982) | 298 | Berkeley and Houde (1978) |
| | <i>Hyporhamphus unifasciatus</i> | 27 | Fischer (1978) | 167 | Beebe and Tee-Van (1928) |
| Holocentridae | <i>Holocentrus ascensionis</i> | 37 TL% | Randall (1962, 1967, 1968) | 838 | Munro (1983) |
| | <i>Holocentrus coruscus</i> | 13 | Randall (1962, 1967, 1968) | 57 | Beebe and Tee-Van (1928) |
| | <i>Holocentrus rufus</i> | 32 | Fischer (1978); Randall (1962, 1967, 1968) | 583 | Munro (1983) |
| | <i>Myripristis jacobus</i> | 23 % | Randall (1962, 1967, 1968) | 397 | Beebe and Tee-Van (1928) |
| | <i>Neoniphon manianus</i> | 19 % | Randall (1962, 1967, 1968) | 163 | Beebe and Tee-Van (1928) |
| | <i>Plectrypops retrospinis</i> | 13 | Randall (1962, 1967, 1968) | 57 | * |
| | <i>Sargocentron vexillarium</i> | 18 - | Randall (1962, 1967, 1968) | 181 | Beebe and Tee-Van (1928) |
| Kyphosidae | <i>Kyphosus incisor</i> | 70 % | Randall (1962, 1967, 1968) | 6,122 | Randall (1962, 1967, 1968) |
| | <i>Kyphosus sectatrix</i> | 80 % | Randall (1962, 1967, 1968) | 9,139 | Randall (1962, 1967, 1968) |
| Labridae | <i>Bodianus rufus</i> | 40 | Fischer (1978); Randall (1962, 1967, 1968) | 1,560 | Beebe and Tee-Van (1928) |
| | <i>Clepticus parrae</i> | 25 | Randall (1962, 1967, 1968) | 447 | Beebe and Tee-Van (1928) |
| | <i>Halichoeres bivittatus</i> | 23 % | Randall (1962, 1967, 1968) | 230 | * |
| | <i>Halichoeres garnoti</i> | 19 | Randall (1962, 1967, 1968) | 132 | Beebe and Tee-Van (1928) |
| | <i>Halichoeres maculipinna</i> | 11 | Randall (1962, 1967, 1968) | 25 | * |
| | <i>Halichoeres poeyi</i> | 20 | Randall (1962, 1967, 1968) | 150 | * |
| | <i>Halichoeres radiatus</i> | 46 | Randall (1962, 1967, 1968) | 2,283 | Beebe and Tee-Van (1928) |
| | <i>Lachnolaimus maximus</i> | 86 % | Randall (1962, 1967, 1968) | 7,728 | Randall (1962, 1967, 1968) |
| | <i>Thalassoma bifasciatum</i> | 17 | Pauly (1978) | 97 | Beebe and Tee-Van (1928) |
| | <i>Xyrichtys novacula</i> | 23 % | Randall (1962, 1967, 1968) | 230 | * |
| | <i>Xyrichtys splendens</i> | 14 | Randall (1962, 1967, 1968) | 50 | * |
| | Lutjanidae | <i>Apsilus dentatus</i> | 65 Tlf | Pauly (1980) | 4,000 |
| | | 67 TLm | Pauly (1980) | 4,500 | Pauly (1980) |
| <i>Lutjanus analis</i> | | 74 FL% | Randall (1962, 1967, 1968) | 5,511 | Randall (1962, 1967, 1968) |
| <i>Lutjanus apodus</i> | | 59 FL% | Randall (1962, 1967, 1968) | 3,502 | Randall (1962, 1967, 1968) |
| <i>Lutjanus buccanella</i> | | 60 Tlf | Pauly (1980) | 3,200 | Pauly (1980) |
| | | 53 TLm | Pauly (1980) | 1,890 | Pauly (1980) |
| <i>Lutjanus campechanus</i> | | 95 TL | Polovina and Ralston (1987) | 18,000 | Beebe and Tee-Van (1928) |
| <i>Lutjanus cyanopterus</i> | | 160 | Fischer (1978) | 70,000 | Randall (1962, 1967, 1968) |
| <i>Lutjanus gnseus</i> | | 55 | Pauly (1978) | 1,590 | Beebe and Tee-Van (1928) |
| <i>Lutjanus jocu</i> | | 83 % | Randall (1962, 1967, 1968) | 13,380 | Randall (1962, 1967, 1968) |
| <i>Lutjanus mahagoni</i> | | 40 % | Randall (1962, 1967, 1968) | 980 | * |
| <i>Lutjanus synagris</i> | | 43 FL% | Munro (1983) | 1,213 | Munro (1983) |
| <i>Lutjanus vivanus</i> | | 70 FL | Munro (1983) | 4,700 | * |
| <i>Ocyurus chrysurus</i> | | 75 TL% | Randall (1962, 1967, 1968) | 3,570 | Munro (1983) |
| <i>Pristipomoides macrophthalmus</i> | | 37 FL% _m | Munro (1983) | | |
| <i>Rhombopites aurorubens</i> | 63 | Polovina and Ralston (1987) | | | |
| Malacanthidae | <i>Malacanthus plumieri</i> | 61 / | Randall (1962, 1967, 1968) | 1,126 | Randall (1962, 1967, 1968) |
| Megalopidae | <i>Tarpon atlanticus</i> | 250 | Fischer (1978) | 160,000 | Beebe and Tee-Van (1928); Randall (1962, 1967, 1968) |
| Monacanthidae | <i>Aluterus monoceros</i> | 61 + | Randall (1962, 1967, 1968) | 2,500 | * |
| | <i>Aluterus schoepfii</i> | 51 / | Randall (1962, 1967, 1968) | 1,892 | Beebe and Tee-Van (1928) |
| | <i>Aluterus scripta</i> | 76 % | Randall (1962, 1967, 1968) | 2,941 | Randall (1962, 1967, 1968) |
| | <i>Cantherines macrocerus</i> | 44 % | Randall (1962, 1967, 1968) | 2,700 | * |
| | <i>Cantherines pullus</i> | 19 mean | Randall (1962, 1967, 1968) | 220 | Beebe and Tee-Van (1928) |
| | <i>Monacanthus ciliatus</i> | 20 | Fischer (1978) | 448 | Beebe and Tee-Van (1928) |
| | <i>Monacanthus tuckeri</i> | 10 % | Randall (1962, 1967, 1968) | 24 | Smith and Tyler (1972) |
| | <i>Stephanolepis setifer</i> | 19 % | Randall (1962, 1967, 1968) | 220 | * |
| | Mugilidae | <i>Mugil curema</i> | 38 | Randall (1962, 1967, 1968) | 767 |
| Mullidae | <i>Mulloidichthys martinicus</i> | 35 mean | Pauly (1980) | 440 | Munro (1983) |
| | <i>Pseudupeneus maculatus</i> | 28 / | Munro (1983); Randall (1962, 1967, 1968) | 393 | Munro (1983) |
| Muraenidae | <i>Enchelycore nigricans</i> | 100 | Fischer (1978) | 900 | * |
| | <i>Echidna catenata</i> | 71 | Randall (1962, 1967, 1968) | 600 | * |
| | <i>Gymnothorax funebris</i> | 190 | Fischer (1978); Randall (1962, 1967, 1968) | 12,000 | Randall (1962, 1967, 1968) |
| | <i>Gymnothorax miliaris</i> | 50 | Figuereido and Menezes (1977); Fischer (1987); Randall (1962, 1967, 1968) | 201 | Smith and Tyler (1972) |
| | <i>Gymnothorax vicinus</i> | 122 / | Fischer (1978); Randall (1962, 1967, 1968) | 3,000 | * |
| | <i>Lycodontis moringa</i> | 100 | Fischer (1978); Smith and Tyler (1972) | 950 mean | Beebe and Tee-Van (1928); Smith and Tyler (1972) |
| Myliobatidae | <i>Aelobatus narinari</i> | 370 Len | Fischer (1978) | 230,000 | Figuereido and Menezes (1977) |
| | | 280 Wid | Fischer (1978) | | |

continued...

Table 3.5 continued

| Family | Species | L_{∞} (cm) | | Source | W_{∞} (g) | Source |
|----------------------------|-------------------------------------|----------------------------|------|-------------------------------------------------------------|------------------------------|----------------------------------------------------|
| Ogcocephalidae | <i>Ogcocephalus nasutus</i> | 28 | / | Randall (1962, 1967, 1968) | 641 | Beebe and Tee-Van (1928) |
| Ophichthidae | <i>Myrichthys breviceps</i> | 91 | + | Randall (1962, 1967, 1968) | 640 | * |
| | <i>Myrichthys ocellatus</i> | 91 | / | Randall (1962, 1967, 1968) | 640 | * |
| | <i>Ophichthus ophis</i> | 135 | | Randall (1962, 1967, 1968) | 2,091 | Randall (1962, 1967, 1968) |
| Opisthognathidae | <i>Opisthognathus aurifrons</i> | 10 | - | Randall (1962, 1967, 1968) | 30 | * |
| | <i>Opisthognathus macrognathus</i> | 20 | - | Randall (1962, 1967, 1968) | 240 | * |
| | <i>Opisthognathus maxillosus</i> | 13 | / | Randall (1962, 1967, 1968) | 65 | * |
| | <i>Opisthognathus whitehurstii</i> | 8 | | Randall (1962, 1967, 1968) | 15 | * |
| Ostraciidae | <i>Acanthostracion polygonius</i> | 41 | % | Randall (1962, 1967, 1968) | 2,200 | * |
| | <i>Acanthostracion quadricornis</i> | 46 | / | Randall (1962, 1967, 1968) | 3,050 | * |
| | <i>Lactophrys bicaudalis</i> | 45 | | Fischer (1978) | 5,572 | Beebe and Tee-Van (1928) |
| | <i>Lactophrys trigonus</i> | 46 | + | Randall (1962, 1967, 1968) | 3,052 | Beebe and Tee-Van (1928) |
| | <i>Lactophrys triqueter</i> | 30 | | Fischer (1978); Randall (1962, 1967, 1968) | 1,394 1,755 | Beebe and Tee-Van (1928) Smith and Tyler (1972) |
| Pemppheridae | <i>Pemppheris schomburgki</i> | 15 | | Randall (1962, 1967, 1968) | 100 | * |
| | <i>Pemppheris poeyi</i> | 10 | / | Randall (1962, 1967, 1968) | 30 | * |
| Pomacanthidae | <i>Centropyge argi</i> | 7 | | Randall (1962, 1967, 1968) | 10 | * |
| | <i>Holacanthus ciliaris</i> | 46 | | Randall (1962, 1967, 1968) | 1,988 | Randall (1962, 1967, 1968) |
| | <i>Holacanthus tricolor</i> | 36 | % | Munro (1983) | 1,306 | Munro (1983) |
| | <i>Pomacanthus arcuatus</i> | 60 | | Fischer (1978); Munro (1983); Randall (1962, 1967, 1968) | 12,407 | Beebe and Tee-Van (1928) |
| | <i>Pomacanthus paru</i> | 43 | % | Randall (1962, 1967, 1968) | 2,769 | Munro (1983) |
| Pomacentridae | <i>Abudefduf saxatilis</i> | 20 | | Fischer (1978) | 483 | Beebe and Tee-Van (1928) |
| | <i>Abudefduf taurus</i> | 25 | | Fischer (1978) | 900 | * |
| | <i>Chromis cyanea</i> | 13 | / | Beebe and Tee-Van (1928); Randall (1962, 1967, 1968) | 93 | Beebe and Tee-Van (1928) |
| | <i>Chromis multilineata</i> | 20 | SL | Fischer (1978) | 339 | Smith and Tyler (1972) |
| | <i>Microspathodon chrysurus</i> | 20 | | Fischer (1978) | 412 | Beebe and Tee-Van (1928) |
| | <i>Stegastes fuscus</i> | 15 | / | Randall (1962, 1967, 1968) | 168 | Beebe and Tee-Van (1928) |
| | <i>Stegastes leucostictus</i> | 10 | | Randall (1962, 1967, 1968) | 50 | * |
| | <i>Stegastes planifrons</i> | 11 | TL | Pauly (1978) | 68 | Smith and Tyler (1972) |
| | <i>Stegastes variabilis</i> | 10 | + | Randall (1962, 1967, 1968) | 54 | Smith and Tyler (1972) |
| | Pomatomidae | <i>Pomatomus saltator</i> | 115 | | Pauly (1978) | |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | 30 | | Randall (1962, 1967, 1968) | 700 | * |
| | <i>Priacanthus arenatus</i> | 40 | | Fischer (1978); Randall (1962, 1967, 1968) | 1,653 | Beebe and Tee-Van (1928) |
| Rachycentridae | <i>Rachycentron canadum</i> | 160 | | Pauly (1978) | 33,400 | Randall (1962, 1967, 1968) |
| Scaridae | <i>Cryptotomus roseus</i> | 13 | () | Randall (1962, 1967, 1968) | | |
| | <i>Nicholsina usta</i> | 31 | % | Randall (1962, 1967, 1968) | | |
| | <i>Scarus coelestinus</i> | 80 | % | Randall (1962, 1967, 1968) | 8,556 | Randall (1962, 1967, 1968) |
| | <i>Scarus coeruleus</i> | 62 | | Pauly (1978) | 3,720 | Randall (1962, 1967, 1968) |
| | <i>Scarus guacamaia</i> | 99 | % | Randall (1962, 1967, 1968) | 23,000 | Randall (1962, 1967, 1968) |
| | <i>Scarus iserti</i> | 20 | FL | Munro (1983) | 141 | Munro (1983) |
| | <i>Scarus taeniopterus</i> | 35 | FL | Munro (1983) | 1,102 | Beebe and Tee-Van (1928) |
| | <i>Scarus velula</i> | 54 | SL% | Randall (1962, 1967, 1968) | 5,558 | Smith and Tyler (1972) |
| | <i>Sparisoma aurofrenatum</i> | 26 | FL | Munro (1983) | 324 | Munro (1983) |
| | <i>Sparisoma chrysopterygum</i> | 42 | FL | Fischer (1978); Pauly (1978); Randall (1962, 1967, 1968) | 1,510 | Munro (1983) |
| | <i>Sparisoma radians</i> | 20 | % | Randall (1962, 1967, 1968) | 150 | * |
| | <i>Sparisoma rubripinne</i> | 46.5 | | Pauly (1978) | 2,734 | Randall (1962, 1967, 1968) |
| | <i>Sparisoma viride</i> | 50 | FL | Munro (1983) | 2,430 | Munro (1983) |
| | Sciaenidae | <i>Equetus lanceolatus</i> | 25 | | Figueredo and Menezes (1977) | 325 |
| <i>Equetus punctatus</i> | | 28 | % | Randall (1962, 1967, 1968) | 460 | Beebe and Tee-Van (1928) |
| <i>Odonotoscion dentex</i> | | 25 | | Figueredo and Menezes (1977) | 318 | Beebe and Tee-Van (1928) |
| <i>Pareques acuminatus</i> | | 24 | mean | Figueredo and Menezes (1977); Randall (1962, 1967, 1968) | 280 | * |
| Scombridae | <i>Auxis rochei</i> | 50 | FL | Fischer (1978) | 1,800 | * |
| | <i>Auxis thazard</i> | 50 | FL | Fischer (1978) | 1,840 | Beebe and Tee-Van (1928) |
| | <i>Euthynnus aletterafus</i> | 93 | FL | Collette and Nauen (1983) | 12,200 | Collette and Nauen (1983) |
| | <i>Scomberomorus cavalla</i> | 137 | FL | Pauly (1978) | 34,285 | Collette and Nauen (1983) |
| | <i>Scomberomorus maculatus</i> | 77 | FL | Collette and Nauen (1983) | 4,800 | Collette and Nauen (1983) |
| | <i>Scomberomorus regalis</i> | 83.5 | FL | Collette and Nauen (1983) | 4,900 | Collette and Nauen (1983) |
| | <i>Thunnus atlanticus</i> | 90 | | Pauly (1978) | | |
| | | | | | | |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | 36 | / | Randall (1962, 1967, 1968) | 1,600 | Beebe and Tee-Van (1928) |
| | <i>Scorpaena grandicornis</i> | 18 | / | Randall (1962, 1967, 1968) | 212 | Beebe and Tee-Van (1928) |
| | <i>Scorpaena inermis</i> | >7 | | Randall (1962, 1967, 1968) | 10 | * |
| | <i>Scorpaena plumieri</i> | 43 | / | Randall (1962, 1967, 1968) | 4,330 | Beebe and Tee-Van (1928) |
| | <i>Scorpaenodes caribbaeus</i> | 13 | / | Randall (1962, 1967, 1968) | 80 | * |

continued...

Table 3.5 continued

| Family | Species | L_{∞} (cm) | Source | W_{∞} (g) | Source |
|----------------|------------------------------------|----------------------|----------------------------------------------------------------------------|---------------------|---------------------------------------------------------------|
| Serranidae | <i>Alphesthes afer</i> | 33 | Fischer (1978) | 1,202 | Beebe and Tee-Van (1928); Figuereido and Menezes (1977) |
| | <i>Cephalopholis cruentata</i> | 34 | Munro (1983) | 691 | Munro (1983) |
| | <i>Cephalopholis fulva</i> | 34 | Munro (1983) | 640 | Munro (1983) |
| | <i>Dermatolepis inermis</i> | 85 % | Randall (1962, 1967, 1968) | 10,513 | Randall (1962, 1967, 1968) |
| | <i>Diplectrum formosum</i> | 31 | Fischer (1978); Pauly (1978); Randall (1962, 1967, 1968) | 550 | * |
| | <i>Epinephelus adscensionis</i> | 50 % | Randall (1962, 1967, 1968) | 1,981 | Randall (1962, 1967, 1968) |
| | <i>Epinephelus guttatus</i> | 58 % | Randall (1962, 1967, 1968) | 2,919 | Munro (1983) |
| | <i>Epinephelus itajara</i> * | 228 % | Randall (1962, 1967, 1968) | 381,644 | Randall (1962, 1967, 1968) |
| | <i>Epinephelus morio</i> | 71 | Fischer (1978) | 5,010 | Randall (1962, 1967, 1968) |
| | <i>Epinephelus striatus</i> | 100 | Pauly (1978); Polovina and Ralston (1987) | 23,800 | Randall (1962, 1967, 1968) |
| | <i>Hypoplectrus aberrans</i> | 12 | Randall (1962, 1967, 1968) | 50 | * |
| | <i>Hypoplectrus chlorurus</i> | 13 | Randall (1962, 1967, 1968) | 64 | * |
| | <i>Hypoplectrus nigricans</i> | 15 | Randall (1962, 1967, 1968) | 90 | * |
| | <i>Hypoplectrus puella</i> | 13 SL | Randall (1962, 1967, 1968) | 64 | Randall (1962, 1967, 1968) |
| | <i>Mycteroperca bonaci</i> | 120 | Polovina and Ralston (1987) | 90,000 | Figuereido and Menezes (1977) |
| | <i>Mycteroperca cidi</i> | 114 | Randall (1962, 1967, 1968) | 77,000 | * |
| | <i>Mycteroperca interstitialis</i> | 70 | Figuereido and Menezes (1977); Randall (1962, 1967, 1968) | 4,000 | Figuereido and Menezes (1977) |
| | <i>Mycteroperca microlepis</i> | 110 mean | Pauly (1978); Polovina and Ralston (1987) | 70,000 | * |
| | <i>Mycteroperca phenax</i> | 72 + | Polovina and Ralston (1987) | 43,700 | Fischer (1978) |
| | <i>Mycteroperca rubra</i> | 70 | Randall (1962, 1967, 1968) | 10,719 | Fischer (1978) |
| | <i>Mycteroperca tigris</i> | 102 | Fischer (1978); Randall (1962, 1967, 1968) | 17,043 | Randall (1962, 1967, 1968) |
| | <i>Mycteroperca venenosa</i> | 86 FL | Munro (1983); Polovina and Ralston (1987) | 12,270 | Randall (1962, 1967, 1968) |
| | <i>Paranthias furcifer</i> | 37 | Polovina and Ralston (1987) | 1,100 | Beebe and Tee-Van (1928) |
| | <i>Serranus dewegeri</i> | 32 | Randall (1962, 1967, 1968) | 700 | * |
| | <i>Serranus tabacarius</i> | 18 | Randall (1962, 1967, 1968) | 141 | Beebe and Tee-Van (1928) |
| | <i>Serranus tigrinus</i> | 10 | Randall (1962, 1967, 1968) | 21 | Beebe and Tee-Van (1928) |
| | <i>Serranus tortugarum</i> | 8 | Randall (1962, 1967, 1968) | 12 | * |
| Sparidae | <i>Archosargus rhomboidalis</i> | 35 | Figuereido and Menezes (1977) | 1,200 | * |
| | <i>Calamus bajonado</i> | 64 | Beebe and Tee-Van (1928) | 4,500 | Beebe and Tee-Van (1928) |
| | <i>Calamus calamus</i> | 38 mean | Beebe and Tee-Van (1928); Fischer (1978); Randall (1962, 1967, 1968) | 1,541 | Beebe and Tee-Van (1928) |
| | <i>Calamus penna</i> | 50 % | Figuereido and Menezes (1977) | 2,000 | * |
| | <i>Calamus pennatula</i> | 39 % | Randall (1962, 1967, 1968) | 1,700 | * |
| | <i>Diplodus caudimacula</i> | 28 | Fischer (1978) | 600 | * |
| Sphyrinaeidae | <i>Sphyrna barracuda</i> | 178 | Pauly (1978); Randall (1962, 1967, 1968) | 57,800 | Randall (1962, 1967, 1968) |
| | <i>Sphyrna picudilla</i> | 50 | Fischer (1978) | 1,300 | * |
| Sphyrnidae | <i>Sphyrna mokarran</i> | 550 | Figuereido and Menezes (1977); Randall (1962, 1967, 1968) | 711,000 mean | Figuereido and Menezes (1977); Randall (1962, 1967, 1968) |
| | <i>Sphyrna lewini</i> | 420 | Fischer (1978) | 400,853 | Figuereido and Menezes (1977); Randall (1962, 1967, 1968) |
| | <i>Sphyrna tiburo</i> | 150 | Fischer (1978) | 18,000 | * |
| Synodontidae | <i>Synodus intermedius</i> | 46 | Randall (1962, 1967, 1968) | 1,103 | Beebe and Tee-Van (1928) |
| | <i>Synodus foetens</i> | 46 | Randall (1962, 1967, 1968) | 1,100 | * |
| | <i>Synodus synodus</i> | 33 | Randall (1962, 1967, 1968) | 400 | * |
| Tetraodontidae | <i>Canthigaster rostrata</i> | 11 (SL) | Randall (1962, 1967, 1968) | 82 | Smith and Tyler (1972) |
| | <i>Sphoeroides spengleri</i> | 15 | Fischer (1978) | 92 | Beebe and Tee-Van (1928) |
| | <i>Sphoeroides testudineus</i> | 30 | Fischer (1978) | | |
| Triakidae | <i>Mustelus canis</i> | 150 / | Fischer (1978) | 15,000 | * |

Symbols: FL = fork length.
 SL = standard length.
 TL = total length.
 Len = length.
 Wid = width.
 f = female.
 m = male.
 mean = mean from several sources or several values in one source
 / = approximate asymptotic size.
 + = rather more than recorded size.
 - = rather less than recorded size.
 () = 'reported' to reach the recorded size.
 % = L_{\max}/W_{\max} transformed into L/W by equation (2.5)/(2.6).
 * = calculated with mean 'a' from similarly sized species of genus or family.

Table 3.6. Estimated average biomass of fish species in Virgin Islands reef environments.

| Family | Species | Average weight (g/ind.) | Average density (Nm ⁻²) | Average biomass (gm ⁻²) |
|----------------|-----------------------------------|-------------------------|-------------------------------------|-------------------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 74 | 3.75E-02 | 2.7750 |
| | <i>Acanthurus chirurgus</i> | 212 | 1.5E-02 | 3.1800 |
| | <i>Acanthurus coeruleus</i> | 254 | 2.5E-02 | 6.3500 |
| Albulidae | <i>Albula vulpes</i> | 1,845 | 1.25E-05 | 0.0231 |
| Antennariidae | <i>Antennarius multiocellatus</i> | 18 | 1.25E-05 | 0.0002 |
| | <i>Antennarius striatus</i> | 18 | 1.25E-04 | 0.0023 |
| Apogonidae | <i>Apogon conklini</i> | 1 | 1.26E-02 | 0.0126 |
| | <i>Apogon maculatus</i> | 10 | 1.25E-03 | 0.0125 |
| Atherinidae | <i>Allanetta harringtonensis</i> | 3 | 1.25E-01 | 0.3754 |
| | <i>Atherinomorus stipes</i> | 4 | 1.25E-01 | 0.5000 |
| Aulostomidae | <i>Aulostomus maculatus</i> | 200 | 2.5E-03 | 0.5000 |
| Balistidae | <i>Balistes capriscus</i> | 158 | 1.25E-05 | 0.0020 |
| | <i>Balistes vetula</i> | 667 | 1.38E-03 | 0.9171 |
| | <i>Canthidermis sufflamen</i> | 1,234 | 1.25E-05 | 0.0154 |
| | <i>Melichthys niger</i> | 248 | 1.25E-04 | 0.0310 |
| | <i>Xanthichthys ringens</i> | 83 | 1.25E-04 | 0.0104 |
| Belonidae | <i>Ablennes hians</i> | 194 | 1.25E-05 | 0.0024 |
| | <i>Platybelone argalus</i> | 39 | 1.25E-03 | 0.0488 |
| | <i>Strongylura timucu</i> | 28 | 1.25E-03 | 0.0350 |
| | <i>Tylosurus acus</i> | 722 | 1.25E-04 | 0.0903 |
| | <i>Tylosurus crocodilus</i> | 361 | 1.25E-03 | 0.4513 |
| Blenniidae | <i>Entomacrodus nigricans</i> | 3 | 1.25E-03 | 0.0038 |
| | <i>Ophioblennius atlanticus</i> | 9 | 1.25E-03 | 0.0113 |
| | <i>Parablennius marmoratus</i> | 3 | 1.25E-03 | 0.0038 |
| | <i>Scartella cristata</i> | 7 | 1.25E-04 | 0.0009 |
| Bothidae | <i>Bothus lunatus</i> | 774 | 1.25E-04 | 0.0968 |
| | <i>Bothus ocellatus</i> | 46 | 1.25E-05 | 0.0006 |
| Carangidae | <i>Alectis ciliaris</i> | 2,761 | 1.25E-05 | 0.0345 |
| | <i>Caranx bartholomaei</i> | 2,544 | 1.25E-04 | 0.3180 |
| | <i>Caranx crysos (fusus)</i> | 597 | 1.38E-04 | 0.0821 |
| | <i>Caranx latus</i> | 1,613 | 2.5E-04 | 0.4033 |
| | <i>Caranx lugubris</i> | 1,691 | 1.25E-05 | 0.0211 |
| | <i>Caranx ruber</i> | 815 | 3.88E-02 | 31.5813 |
| | <i>Decapterus macarellus</i> | 194 | 1.25E-05 | 0.0024 |
| | <i>Decapterus punctatus</i> | 148 | 1.25E-04 | 0.0185 |
| | <i>Elagatis bipinnulatus</i> | 2,709 | 1.25E-05 | 0.0339 |
| | <i>Oligoplites saurus</i> | 86 | 1.25E-02 | 1.0750 |
| | <i>Selar crumenophthalmus</i> | 320 | 1.25E-04 | 0.0400 |
| | <i>Seriola dumerili</i> | 20,640 | 1.25E-05 | 0.2580 |
| | <i>Seriola rivoliana</i> | 6,192 | 1.25E-05 | 0.0774 |
| | <i>Trachinotus falcatus</i> | 9,288 | 1.25E-05 | 0.1161 |
| | <i>Trachinotus goodei</i> | 1,006 | 1.25E-04 | 0.1258 |
| Carcharhinidae | <i>Carcharhinus acronotus</i> | 23,220 | 1.25E-05 | 0.2903 |
| | <i>Carcharhinus falciformis</i> | 90,300 | 1.25E-05 | 1.1288 |
| | <i>Carcharhinus leucas</i> | 90,300 | 1.25E-05 | 1.1288 |
| | <i>Carcharhinus limbatus</i> | 29,928 | 1.25E-05 | 0.3741 |
| | <i>Carcharhinus longimanus</i> | 90,300 | 1.25E-05 | 1.1288 |
| | <i>Carcharhinus perezi</i> | 25,800 | 1.25E-04 | 3.2250 |
| | <i>Galeocerdo cuvieri</i> | 173,376 | 1.25E-04 | 21.6720 |

continued...

Table 3.6 continued

| Family | Species | Average weight (g/ind.) | Average density (Nm ⁻²) | Average biomass (gm ⁻²) |
|--------------------|------------------------------------|-------------------------|-------------------------------------|-------------------------------------|
| | <i>Negaprion brevirostris</i> | 28,380 | 1.25E-04 | 3.5475 |
| | <i>Rhizoprionodon porosus</i> | 1,295 | 1.25E-04 | 0.1619 |
| Chaetodontidae | <i>Chaetodon aculeatus</i> | 5 | 1.25E-05 | 0.0001 |
| | <i>Chaetodon capistratus</i> | 31 | 2.5E-03 | 0.0775 |
| | <i>Chaetodon ocellatus</i> | 52 | 1.25E-05 | 0.0007 |
| | <i>Chaetodon sedentarius</i> | 34 | 1.25E-05 | 0.0004 |
| | <i>Chaetodon striatus</i> | 42 | 1.25E-05 | 0.0005 |
| Cirrhitidae | <i>Amblycirrhitus pinos</i> | 4 | 1.38E-04 | 0.0006 |
| Clinidae | <i>Labrisomus guppyi</i> | 35 | 1.25E-02 | 0.4375 |
| | <i>Labrisomus nuchipinnis</i> | 35 | 1.25E-02 | 0.4375 |
| Clupeidae | <i>Harengula clupeiola</i> | 62 | 1.25E-01 | 7.7500 |
| | <i>Harengula humeralis</i> | 114 | 1.39E-01 | 15.8175 |
| | <i>Jenkinsia lamprotaenia</i> | 3 | 1.39E+00 | 4.1625 |
| | <i>Opisthomema oglinum</i> | 188 | 1.25E-02 | 2.3500 |
| | <i>Sardinella aurita</i> | 258 | 1.25E-02 | 3.2250 |
| Congridae | <i>Heteroconger halis</i> | 39 | 1.25E-02 | 0.4875 |
| Dactylopteridae | <i>Dactylopterus volitans</i> | 471 | 1.25E-02 | 5.8875 |
| Dasyatidae | <i>Dasyatis americana</i> | 31,476 | 1.25E-03 | 39.3450 |
| Diodontidae | <i>Chilomycterus antennatus</i> | 366 | 1.38E-04 | 0.0503 |
| | <i>Chilomycterus antillarum</i> | 366 | 1.25E-05 | 0.0046 |
| | <i>Diodon holocanthus</i> | 2,915 | 1.25E-02 | 36.4739 |
| | <i>Diodon hystrix</i> | 4,759 | 1.25E-04 | 0.5949 |
| Elopidae | <i>Elops saurus</i> | 115 | 1.25E-04 | 0.0144 |
| Emmelichthyidae | <i>Inermia vittata</i> | 73 | 1.25E-02 | 0.9125 |
| Engraulidae | <i>Anchoa hepsetus</i> | 13 | 1.25E-03 | 0.0163 |
| | <i>Anchoa lyolepis</i> | 2 | 1.25E-03 | 0.0025 |
| Ephippidae | <i>Chaetodipterus faber</i> | 2,322 | 2.5E-04 | 0.5805 |
| Fistulariidae | <i>Fistularia tabacaria</i> | 2,786 | 1.25E-04 | 0.3483 |
| Gerreidae | <i>Eucinostomus argenteus</i> | 75 | 1.25E-02 | 0.9394 |
| | <i>Gerres cinereus</i> | 178 | 1.39E-02 | 2.4698 |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | 129,000 | 1.25E-04 | 16.1250 |
| Gobiidae | <i>Bathygobius soporator</i> | 18 | 1.38E-04 | 0.0025 |
| | <i>Coryphopterus glaucofraenum</i> | 3 | 2.5E-02 | 0.0750 |
| | <i>Gnatholepis thompsoni</i> | 1 | 1.26E-02 | 0.0126 |
| | <i>Gobiosoma evelynae</i> | 0.6 | 1.25E-01 | 0.0751 |
| | <i>Gobiosoma horsti</i> | 0.8 | 1.25E-01 | 0.1001 |
| Grammidae | <i>Gramma loreto</i> | 4 | 1.25E-03 | 0.0050 |
| | <i>Gramma melacara</i> | 8 | 1.25E-04 | 0.0010 |
| Grammistidae | <i>Rypticus saponaceus</i> | 177 | 1.25E-02 | 2.2147 |
| Haemulidae | <i>Anisotremus surinamensis</i> | 1,231 | 1.38E-04 | 0.1693 |
| | <i>Anisotremus virginicus</i> | 219 | 1.38E-04 | 0.0301 |
| | <i>Haemulon album</i> | 1,367 | 2.63E-04 | 0.3588 |

continued...

Table 3.6 continued

| Family | Species | Average weight (g/ind.) | Average density (Nm ⁻²) | Average biomass (gm ⁻²) |
|---------------|----------------------------------|-------------------------|-------------------------------------|-------------------------------------|
| | <i>Haemulon aurolineatum</i> | 49 | 1.26E-02 | 0.6186 |
| | <i>Haemulon bonariense</i> | 99 | 1.25E-05 | 0.0012 |
| | <i>Haemulon carbonarium</i> | 222 | 1.25E-04 | 0.0278 |
| | <i>Haemulon chrysargyreum</i> | 65 | 1.38E-03 | 0.0894 |
| | <i>Haemulon flavolineatum</i> | 124 | 1.5E-01 | 18.6000 |
| | <i>Haemulon macrostomum</i> | 463 | 1.25E-05 | 0.0058 |
| | <i>Haemulon melanurum</i> | 110 | 1.25E-05 | 0.0014 |
| | <i>Haemulon parrai</i> | 326 | 1.25E-03 | 0.4075 |
| | <i>Haemulon plumieri</i> | 351 | 1.51E-02 | 5.3089 |
| | <i>Haemulon sciurus</i> | 306 | 3.88E-03 | 1.1858 |
| | <i>Pomadasys corvinaeformis</i> | 100 | 1.25E-05 | 0.0013 |
| | <i>Pomadasys croco</i> | 230 | 1.25E-05 | 0.0029 |
| Hemiramphidae | <i>Hemiramphus balao</i> | 158 | 1.25E-03 | 0.1975 |
| | <i>Hemiramphus brasiliensis</i> | 77 | 1.25E-04 | 0.0096 |
| | <i>Hyporhamphus unifasciatus</i> | 43 | 1.25E-04 | 0.0054 |
| Holocentridae | <i>Holocentrus ascensionis</i> | 216 | 2.53E-03 | 0.5454 |
| | <i>Holocentrus coruscus</i> | 15 | 1.51E-03 | 0.0227 |
| | <i>Holocentrus rufus</i> | 150 | 1.39E-02 | 2.0831 |
| | <i>Myripristis jacobus</i> | 102 | 3.88E-04 | 0.0395 |
| | <i>Neoniphon marianus</i> | 42 | 2.63E-04 | 0.0110 |
| | <i>Plectrypops retrospinis</i> | 15 | 1.5E-03 | 0.0225 |
| | <i>Sargocentron vexillarium</i> | 47 | 1.4E-03 | 0.0658 |
| Kyphosidae | <i>Kyphosus incisor</i> | 1,579 | 1.38E-03 | 2.1711 |
| | <i>Kyphosus sectatrix</i> | 2,358 | 2.5E-03 | 5.8950 |
| Labridae | <i>Bodianus rufus</i> | 402 | 2.5E-04 | 0.1005 |
| | <i>Clepticus parrae</i> | 115 | 1.25E-02 | 1.4375 |
| | <i>Halichoeres bivittatus</i> | 59 | 1.38E-02 | 0.8113 |
| | <i>Halichoeres garnoti</i> | 34 | 1.25E-03 | 0.0425 |
| | <i>Halichoeres maculipinna</i> | 6 | 1.25E-03 | 0.0075 |
| | <i>Halichoeres poeyi</i> | 39 | 1.26E-02 | 0.4924 |
| | <i>Halichoeres radiatus</i> | 589 | 1.25E-04 | 0.0736 |
| | <i>Lachnolaimus maximus</i> | 1,994 | 1.25E-05 | 0.0249 |
| | <i>Thalassoma bifasciatum</i> | 25 | 2.5E-02 | 0.6250 |
| | <i>Xyrichtys novacula</i> | 59 | 1.25E-02 | 0.7375 |
| | <i>Xyrichtys splendens</i> | 13 | 1.25E-02 | 0.1625 |
| Lutjanidae | <i>Lutjanus analis</i> | 1,422 | 2.63E-04 | 0.3733 |
| | <i>Lutjanus apodus</i> | 904 | 1.5E-02 | 13.5600 |
| | <i>Lutjanus cyanopterus</i> | 18,060 | 1.25E-05 | 0.2258 |
| | <i>Lutjanus griseus</i> | 410 | 1.5E-02 | 6.1500 |
| | <i>Lutjanus jocu</i> | 3,452 | 1.25E-04 | 0.4315 |
| | <i>Lutjanus mahagoni</i> | 253 | 1.5E-02 | 3.7950 |
| | <i>Lutjanus synagris</i> | 313 | 2.5E-03 | 0.7825 |
| | <i>Ocyurus chrysurus</i> | 921 | 2.75E-02 | 25.3275 |
| Malacanthidae | <i>Malacanthus plumieri</i> | 291 | 1.28E-02 | 3.7103 |
| Megalopidae | <i>Tarpon atlanticus</i> | 41,280 | 1.38E-04 | 5.6760 |
| Monacanthidae | <i>Aluterus schoepfii</i> | 488 | 1.25E-03 | 0.6100 |
| | <i>Aluterus scripta</i> | 759 | 1.25E-05 | 0.0095 |
| | <i>Cantherines macrocerus</i> | 697 | 1.25E-05 | 0.0087 |
| | <i>Cantherines pullus</i> | 57 | 1.25E-04 | 0.0071 |
| | <i>Monacanthus ciliatus</i> | 116 | 1.25E-02 | 1.4500 |
| | <i>Monacanthus tuckeri</i> | 6 | 1.26E-03 | 0.0076 |
| | <i>Stephanolepis setifer</i> | 57 | 1.25E-04 | 0.0071 |

continued...

Table 3.6 continued

| Family | Species | Average weight (g/ind.) | Average density (Nm ⁻²) | Average biomass (gm ⁻²) |
|------------------|-------------------------------------|-------------------------|-------------------------------------|-------------------------------------|
| Mugilidae | <i>Mugil curema</i> | 198 | 1.38E-04 | 0.0272 |
| Mullidae | <i>Mulloidichthys martinicus</i> | 114 | 1.5E-02 | 1.7100 |
| | <i>Pseudupeneus maculatus</i> | 101 | 5.E-03 | 0.5050 |
| Muraenidae | <i>Echidna catenata</i> | 155 | 1.25E-05 | 0.0019 |
| | <i>Enchelycore nigricans</i> | 232 | 1.25E-05 | 0.0029 |
| | <i>Gymnothorax funebris</i> | 3,096 | 1.25E-04 | 0.3870 |
| | <i>Gymnothorax miliaris</i> | 52 | 1.25E-05 | 0.0007 |
| | <i>Gymnothorax vicinus</i> | 774 | 1.25E-04 | 0.0968 |
| | <i>Lycodontis moringa</i> | 245 | 1.25E-05 | 0.0031 |
| Myliobatidae | <i>Aetobatus narinari</i> | 59,340 | 1.25E-05 | 0.7418 |
| Ogcocephalidae | <i>Ogcocephalus nasutus</i> | 165 | 1.25E-04 | 0.0206 |
| Ophichthidae | <i>Myrichthys breviceps</i> | 165 | 1.25E-05 | 0.0021 |
| | <i>Myrichthys ocellatus</i> | 165 | 1.25E-05 | 0.0021 |
| | <i>Ophichthus ophis</i> | 540 | 1.25E-03 | 0.6750 |
| Opisthognathidae | <i>Opisthognathus aunifrons</i> | 8 | 1.25E-03 | 0.0100 |
| | <i>Opisthognathus macrognathus</i> | 62 | 1.25E-05 | 0.0008 |
| | <i>Opisthognathus maxillosus</i> | 17 | 6.25E-04 | 0.0106 |
| | <i>Opisthognathus whitehurstii</i> | 4 | 1.25E-03 | 0.0050 |
| Ostraciidae | <i>Acanthostracion polygonius</i> | 568 | 1.28E-03 | 0.7242 |
| | <i>Acanthostracion quadricornis</i> | 787 | 1.5E-04 | 0.1181 |
| | <i>Lactophrys bicaudalis</i> | 1,438 | 1.5E-04 | 0.2157 |
| | <i>Lactophrys trigonus</i> | 787 | 1.28E-03 | 1.0034 |
| | <i>Lactophrys triqueter</i> | 360 | 2.51E-03 | 0.9045 |
| Pempheridae | <i>Pempheris poeyi</i> | 8 | 1.25E-04 | 0.0010 |
| | <i>Pempheris schomburgki</i> | 26 | 1.25E-03 | 0.0325 |
| Pomacanthidae | <i>Centropyge argi</i> | 3 | 1.25E-04 | 0.0004 |
| | <i>Holacanthus ciliaris</i> | 513 | 1.28E-02 | 6.5407 |
| | <i>Holacanthus tricolor</i> | 337 | 2.5E-04 | 0.0843 |
| | <i>Pomacanthus arcuatus</i> | 3,201 | 2.51E-03 | 8.0425 |
| | <i>Pomacanthus paru</i> | 714 | 2.5E-04 | 0.1785 |
| Pomacentridae | <i>Abudefduf saxatilis</i> | 125 | 1.25E-03 | 0.1563 |
| | <i>Abudefduf taurus</i> | 232 | 1.25E-04 | 0.0290 |
| | <i>Chromis cyanea</i> | 24 | 1.25E-02 | 0.3000 |
| | <i>Chromis multilineata</i> | 87 | 1.25E-02 | 1.0875 |
| | <i>Microspathodon chrysurus</i> | 106 | 1.25E-02 | 1.3250 |
| | <i>Stegastes fuscus</i> | 43 | 1.26E-01 | 5.4288 |
| | <i>Stegastes leucostictus</i> | 13 | 1.25E-02 | 0.1625 |
| | <i>Stegastes planifrons</i> | 18 | 1.25E-02 | 0.2250 |
| | <i>Stegastes variabilis</i> | 14 | 1.25E-02 | 0.1750 |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | 181 | 1.25E-04 | 0.0226 |
| | <i>Priacanthus arenatus</i> | 426 | 1.25E-04 | 0.0533 |
| Rachycentridae | <i>Rachycentron canadum</i> | 8,617 | 1.25E-05 | 0.1077 |
| Scaridae | <i>Scarus coelestinus</i> | 2,207 | 1.25E-04 | 0.2759 |
| | <i>Scarus coeruleus</i> | 960 | 1.25E-04 | 0.1200 |
| | <i>Scarus guacamaia</i> | 5,934 | 1.26E-02 | 74.9168 |
| | <i>Scarus iserti</i> | 36 | 2.5E-03 | 0.0900 |
| | <i>Scarus taeniopterus</i> | 284 | 1.25E-04 | 0.0355 |
| | <i>Scarus vetula</i> | 1,434 | 1.38E-02 | 19.7175 |

continued...

Table 3.6 continued

| Family | Species | Average weight (g/ind.) | Average density (Nm ⁻²) | Average biomass (gm ⁻²) |
|----------------|------------------------------------|-------------------------|-------------------------------------|-------------------------------------|
| | <i>Sparisoma aurofrenatum</i> | 84 | 1.25E-03 | 0.1050 |
| | <i>Sparisoma chrysopterum</i> | 390 | 1.38E-02 | 5.3625 |
| | <i>Sparisoma radians</i> | 39 | 1.25E-01 | 4.8799 |
| | <i>Sparisoma rubripinne</i> | 705 | 1.25E-03 | 0.8813 |
| | <i>Sparisoma viride</i> | 627 | 1.38E-03 | 0.8621 |
| Sciaenidae | <i>Equetus lanceolatus</i> | 84 | 1.25E-04 | 0.0105 |
| | <i>Equetus punctatus</i> | 119 | 1.25E-04 | 0.0149 |
| | <i>Odontoscion dentex</i> | 82 | 2.5E-04 | 0.0205 |
| | <i>Paregues acuminatus</i> | 72 | 1.25E-05 | 0.0009 |
| Scombridae | <i>Euthynnus aletteratus</i> | 3,148 | 1.25E-04 | 0.3935 |
| | <i>Scomberomorus cavalla</i> | 8,846 | 1.38E-04 | 1.2163 |
| | <i>Scomberomorus regalis</i> | 1,264 | 1.38E-03 | 1.7380 |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | 413 | 1.25E-03 | 0.5163 |
| | <i>Scorpaena grandicornis</i> | 55 | 6.25E-05 | 0.0034 |
| | <i>Scorpaena inermis</i> | 3 | 1.25E-03 | 0.0038 |
| | <i>Scorpaena plumieri</i> | 1,117 | 1.38E-04 | 0.1536 |
| | <i>Scorpaenodes caribbaeus</i> | 21 | 1.25E-04 | 0.0026 |
| Serranidae | <i>Alphestes afer</i> | 310 | 6.25E-05 | 0.0194 |
| | <i>Cephalopholis cruentata</i> | 178 | 1.25E-04 | 0.0223 |
| | <i>Cephalopholis fulva</i> | 165 | 1.38E-03 | 0.2269 |
| | <i>Diplectrum formosum</i> | 142 | 1.25E-05 | 0.0018 |
| | <i>Epinephelus adscensionis</i> | 511 | 2.5E-04 | 0.1278 |
| | <i>Epinephelus guttatus</i> | 753 | 2.5E-03 | 1.8825 |
| | <i>Epinephelus itajara</i> | 98,464 | 1.25E-04 | 12.3080 |
| | <i>Epinephelus morio</i> | 1,293 | 1.25E-05 | 0.0162 |
| | <i>Epinephelus striatus</i> | 6,140 | 2.5E-04 | 1.5350 |
| | <i>Hypoplectrus aberrans</i> | 13 | 1.25E-04 | 0.0016 |
| | <i>Hypoplectrus chlorurus</i> | 17 | 1.25E-05 | 0.0002 |
| | <i>Hypoplectrus nigricans</i> | 23 | 1.25E-04 | 0.0029 |
| | <i>Hypoplectrus puella</i> | 17 | 1.25E-03 | 0.0213 |
| | <i>Mycteroperca bonaci</i> | 23,220 | 2.5E-04 | 5.8050 |
| | <i>Mycteroperca interstitialis</i> | 1,032 | 1.25E-05 | 0.0129 |
| | <i>Mycteroperca rubra</i> | 2,766 | 1.25E-05 | 0.0346 |
| | <i>Mycteroperca tigris</i> | 4,397 | 1.25E-04 | 0.5496 |
| | <i>Mycteroperca venenosa</i> | 3,166 | 1.25E-05 | 0.0396 |
| | <i>Paranthias furcifer</i> | 284 | 1.25E-05 | 0.0036 |
| | <i>Serranus tabacarius</i> | 36 | 1.25E-05 | 0.0005 |
| | <i>Serranus tigninus</i> | 5 | 1.38E-03 | 0.0069 |
| | <i>Serranus tortugarum</i> | 3 | 1.25E-04 | 0.0004 |
| Sparidae | <i>Archosargus rhomboidalis</i> | 310 | 1.5E-04 | 0.0465 |
| | <i>Calamus bajonado</i> | 1,161 | 2.5E-05 | 0.0290 |
| | <i>Calamus calamus</i> | 398 | 2.5E-04 | 0.0995 |
| | <i>Calamus pennatula</i> | 439 | 1.38E-04 | 0.0604 |
| | <i>Diplodus caudimacula</i> | 155 | 1.38E-04 | 0.0213 |
| Sphyraenidae | <i>Sphyraena barracuda</i> | 14,912 | 1.5E-03 | 22.3680 |
| | <i>Sphyraena picudilla</i> | 335 | 1.25E-05 | 0.0042 |
| Sphyrnidae | <i>Sphyrma lewini</i> | 103,420 | 1.25E-05 | 1.2928 |
| | <i>Sphyrma tiburo</i> | 4,644 | 1.25E-04 | 0.5805 |
| Synodontidae | <i>Synodus foetens</i> | 284 | 1.25E-04 | 0.0355 |
| | <i>Synodus intermedius</i> | 285 | 1.38E-03 | 0.3919 |
| | <i>Synodus synodus</i> | 103 | 1.25E-03 | 0.1288 |
| Tetraodontidae | <i>Canthigaster rostrata</i> | 21 | 1.26E-02 | 0.2651 |
| | <i>Sphoeroides spengleri</i> | 24 | 1.39E-03 | 0.0333 |
| Triakidae | <i>Mustelus canis</i> | 3,870 | 1.25E-05 | 0.0484 |

Table 3.7. Estimated growth parameters and natural mortality (M) of Caribbean reef fishes; mean environmental temperature = 28°C.

| Family | Species | Origin | L _∞ (cm) | K year ⁻¹ | M year ⁻¹ | Source ^a | |
|----------------|----------------------------------|------------------------------|------------------------|-------------------------|-------------------------|-----------------------------------------------------------|----------------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | VI | 25 FL | 0.383 | 1.00 * | Pauly (1981)* | |
| | | Jamaica | 28 FL | 0.500 | 2.20 > | Nicholson and Hartsuijker (1982) | |
| | | Jamaica | 30 FL | 0.180 | 0.58 * | Nicholson and Hartsuijker (1982) | |
| | | <i>Acanthurus chirurgus</i> | VI | 33 FL | 0.254 | 0.71 * | Pauly (1978) |
| | | <i>Acanthurus coeruleus</i> | VI | 32 FL | 0.247 | 0.70 * | Pauly (1978) |
| | | | Jamaica | 30 FI | 0.500 | 2.35 > | Nicholson and Hartsuijker (1982) |
| | | | Jamaica | 28 FL | 0.220 | 0.68 * | Nicholson and Hartsuijker (1982) |
| Apogonidae | <i>Apogon conklini</i> | | 5 SL/ | 9.111 | 12.50 ** | Pauly (1978) | |
| | <i>Apogon cyanosoma</i> | Red Sea | 9.4 | 2.904 | 4.96 * | Pauly (1978) | |
| | <i>Apogon maculatus</i> | | 10 / | 2.278 | 4.16 ** | Pauly (1978) | |
| Atherinidae | <i>Allanetta harringtonensis</i> | | 8 / | 1.855 | 3.87 ** | Pauly (1978) | |
| | <i>Atherinomorus stipes</i> | | 9 + | 1.466 | 3.21 ** | Pauly (1978) | |
| Balistidae | <i>Balistes capriscus</i> | Gulf of Mexico | 31 | 0.383 | 0.94 * | Johnson and Salomon (1984) | |
| | <i>Balistes vetula</i> | Jamaica | 45 FL | 0.570 | 2.60 | Munro (1983) | |
| | | Jamaica | 54 FL | 0.125 | 0.39 * | Nicholson and Hartsuijker (1982) | |
| | | VI | 60 TL | 0.230 | 0.56 * | Pauly (1978) | |
| Blenniidae | <i>Parablennius marmoreus</i> | | 8 | | 2.53 | Eyberg (1984); annual P/B ratio for the respective genera | |
| | <i>Scartella cristata</i> | | 11 | | 2.84 | Eyberg (1984); annual P/B ratio for the respective genera | |
| Bothidae | <i>Bothus lunatus</i> | | 46 / | 0.321 | 0.75 ** | Pauly (1978) | |
| | <i>Bothus ocellatus</i> | | 18 | 2.101 | 3.35 ** | Pauly (1978) | |
| Carangidae | <i>Caranx ruber</i> | Jamaica | 53 | 0.235 | 0.59 * | Nicholson and Hartsuijker (1982) | |
| | | Jamaica | 56 | 0.240 | 1.40 | Munro (1983) | |
| | | Jamaica | 57 | 0.240 | 1.51 | Pauly (1980) | |
| | | <i>Decapterus macarellus</i> | | 35 SL | 0.397 | 0.93 ** | Pauly (1978) |
| | | <i>Decapterus punctatus</i> | Red Sea | 32 | 0.318 | 0.83 * | Pauly (1978) |
| | | <i>Seriola dumerili</i> | | 194 | 0.118 | 0.26 ** | Pauly (1978) |
| | | <i>Seriola rivoliana</i> | | 97 | 0.473 | 0.79 ** | Pauly (1978) |
| Carcharhinidae | <i>Carcharhinus acronotus</i> | | 200 | | | | |
| | | | 168 | 0.159 | 0.33 * | Pauly (1978) | |
| | <i>Carcharhinus leucas</i> | | 350 | | | | |
| | | 305 | 0.125 | 0.24 * | Pauly (1978) | | |
| | <i>Carcharhinus limbatus</i> | | 247 | | | | |
| | | 228 | 0.139 | 0.28 * | Pauly (1978) | | |

continued...

Table 3.7 continued

| Family | Species | Origin | L _∞ (cm) | K year ⁻¹ | M year ⁻¹ | Source ^a |
|--------------------|------------------------------------|---------|------------------------|-------------------------|-------------------------|-----------------------------------------------------------------------|
| | <i>Galeocerdo cuvier</i> | | 457 | 0.106 | 0.19 * | Pauly (1978) |
| | <i>Negaprion brevirostris</i> | | 320 + | 0.132 | 0.25 * | Pauly (1978) |
| | <i>Prionace glauca</i> | | 383 | 0.110 | 0.21 * | Pauly (1978) |
| Chaetodontidae | <i>Chaetodon aculeatus</i> | | 9.5 | 1.957 | 3.82 ** | Ralston (1976); daily K * 365; ϕ' for <i>Chaetodon miliaris</i> |
| | <i>Chaetodon capistratus</i> | | 14.2 | 0.876 | 2.02 ** | Ralston (1976); daily K * 365; ϕ' for <i>Chaetodon miliaris</i> |
| | <i>Chaetodon ocellatus</i> | | 16.8 | 0.626 | 1.55 ** | Ralston (1976); daily K * 365; ϕ' for <i>Chaetodon miliaris</i> |
| | <i>Chaetodon sedentarius</i> | | 14.7 | 0.817 | 1.91 ** | Ralston (1976); daily K * 365; ϕ' for <i>Chaetodon miliaris</i> |
| | <i>Chaetodon striatus</i> | | 15.8 | 0.708 | 1.70 ** | Ralston (1976); daily K * 365; ϕ' for <i>Chaetodon miliaris</i> |
| Clupeidae | <i>Harengula clupeola</i> | | 18 | 0.627 | 0.91 ** | Hubold and Mazzetti (1982); ϕ' computed from <i>Harengula jaguana</i> |
| | <i>Harengula humeralis</i> | | 22 | 0.420 | 0.66 ** | Hubold and Mazzetti (1982); ϕ' computed from <i>Harengula jaguana</i> |
| | <i>Sardinella aurita</i> | | 29 | 0.488 | 0.68 ** | Pauly (1978) |
| | <i>Sardinella brasiliensis</i> | | | | 0.74 | Saccardo (1983) |
| Coryphaenidae | <i>Coryphaena hippurus</i> | Florida | 135 | 0.620 | 0.86 * | Pauly (1978) |
| | | Florida | 167 | 0.530 | 0.73 * | Pauly (1978) |
| | | mean | 151 | 0.575 | | |
| Elopidae | <i>Elops saurus</i> | Cuba | 47 | 0.240 | 0.62 * | Pauly (1978) |
| Engraulidae | <i>Anchoa hepsetus</i> | | 15 | 0.585 | 1.53 ** | Pauly (1978) |
| | <i>Anchoa lyolepis</i> | | 8 | 2.055 | 4.14 ** | Pauly (1978) |
| | <i>Anchoa naso</i> | Ecuador | 8.6 | 1.780 | 3.69 * | Pauly (1978) |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | | 430 | 0.141 | 0.24 * | Pauly (1978) |
| Gobiidae | <i>Bathygobius soporator</i> | | 15 | 0.223 | 0.81 ** | Pauly (1978) |
| | <i>Coryphopterus glaucofraenum</i> | | 8 (SL) | 0.785 | 2.20 ** | Pauly (1978) |
| | <i>Gnatholepis thompsoni</i> | | 6 SL/ | 1.395 | 3.48 ** | Pauly (1978) |
| | <i>Gobiosoma evelynae</i> | | 5 (SL) | 1.103 | 3.14 ** | Pauly (1978) |
| | <i>Gobiosoma horsti</i> | | 5 (SL) | 1.103 | 3.14 ** | Pauly (1978) |
| | <i>Gobiosoma robustum</i> | Florida | 3.6 | 2.006 | 5.09 * | Pauly (1978) |
| | | Florida | 6 | 0.813 | 2.44 * | Pauly (1978) |
| Haemulidae | <i>Haemulon album</i> | Jamaica | 68 | 0.200 | 0.79 | Munro (1983) |
| (Pomadasyidae) | | Jamaica | 75 | 0.200 | 0.79 | Pauly (1980) |
| | <i>Haemulon aurolineatum</i> | Jamaica | 23 FL | 0.350 | 2.37 > | Nicholson and Hartsuijker (1982) |
| | <i>Haemulon flavolineatum</i> | Jamaica | 27 / | 0.185 | 0.90 > | Nicholson and Hartsuijker (1982) |
| | <i>Haemulon plumieri</i> | Jamaica | 42 FL | 0.345 | 1.77 | Munro (1983), Pauly (1980) |
| | | Jamaica | 42 FL | 0.260 | 0.67 * | Nicholson and Hartsuijker (1982) |

continued...

Table 3.7 continued

| Family | Species | Origin | L _∞ (cm) | K year ⁻¹ | M year ⁻¹ | Source ^a |
|---------------|---------------------------------|------------------|------------------------|-------------------------|-------------------------|----------------------------------|
| | | Jamaica | 46 FL | 0.150 | 0.46 * | Nicholson and Hartsuijker (1982) |
| | <i>Haemulon sciurus</i> | Jamaica | 41 FL | 0.260 | 0.68 * | Munro (1983) |
| Hemiramphidae | <i>Hemiramphus brasiliensis</i> | Florida | 32.5 | 0.587 | 1.23 * | Pauly and Murphy (1982) |
| Holocentridae | <i>Holocentrus ascensionis</i> | Jamaica | 30 | 1.180 | 4.32 | Munro (1983) |
| | | Jamaica | 30 | 0.260 | 0.74 * | Nicholson and Hartsuijker (1982) |
| | | Jamaica | 31 | 0.235 | 0.69 * | Nicholson and Hartsuijker (1982) |
| | <i>Holocentrus coruscus</i> | | 13 | 1.973 | 3.52 ** | Pauly (1978) |
| | <i>Holocentrus diadema</i> | Red Sea | 17 | 1.129 | 2.27 * | Pauly (1978) |
| | <i>Holocentrus rufus</i> | Jamaica | 23 | 0.940 | 4.56 | Munro (1983) |
| | <i>Neoniphon marianus</i> | | 19 | 0.924 | 1.93 ** | Pauly (1978) |
| | | Jamaica | 23 | 0.290 | 0.86 * | Nicholson and Hartsuijker (1982) |
| | <i>Sargocentron vexillarium</i> | | 18 - | 1.029 | 2.10 ** | Pauly (1978) |
| Labridae | <i>Thalassoma bifasciatum</i> | Florida | 17 | 0.750 | 1.73 * | Pauly (1978) |
| Lutjanidae | <i>Apsilus dentatus</i> | Jamaica | 65 f | 0.300 | 0.83 | Pauly (1980) |
| | | Jamaica | 67 m | 0.650 | 1.90 | Pauly (1980) |
| | <i>Lutjanus analis</i> | | 74 FL | 0.120 | 0.20 | Bannerot et al.(1987) |
| | | Cuba | 81 TL | 0.116 | 0.33 * | Pozo (1979) |
| | | Florida | 86 TL | 0.153 | 0.39 * | Polovina and Ralston (1987) |
| | <i>Lutjanus apodus</i> | Jamaica | 63 TL | 0.180 | 0.54 | Pauly (1980) |
| | <i>Lutjanus buccanella</i> | Jamaica | 53 TL | 0.700 | 2.24 | Pauly (1980) |
| | | Jamaica | 60 | 0.350 | 1.83 | Pauly (1980) |
| | <i>Lutjanus campechanus</i> | mean | 95 TL | 0.150 | 0.37 * | Polovina and Ralston (1987) |
| | <i>Lutjanus griseus</i> | Cuba | 51 FL | 0.240 | 0.61 * | Baez et al. (1980) |
| | | Florida | 55 FLm | 0.167 | 0.47 * | Pauly (1978) |
| | <i>Lutjanus synagris</i> | Cuba | 42 FL | 0.268 | 0.69 * | Pauly (1978) |
| | | Florida | 50 TL | 0.134 | 0.42 * | Polovina and Ralston (1987) |
| | | Ceara | 50 | 0.231 | 0.59 * | Pauly (1978) |
| | <i>Ocyurus chrysurus</i> | Jamaica | 75 TL | 0.250 | 0.60 | Munro (1983) |
| | | Southern Florida | 45 FL | 0.279 | 0.69 * | Johnson and Attramadal (1983) |
| | | Jamaica | 60 FL | 0.250 | 0.60 | Munro (1983) |
| | | Jamaica | 60 FL | 0.250 | 0.62 | Pauly (1980) |
| | | | 69 TL | 0.160 | 0.20 | Bannerot et al. (1987) |
| | <i>Rhomboplites aurorubens</i> | | 63 | 0.198 | 0.50 * | Polovina and Ralston (1987) |
| Malacanthidae | <i>Malacanthus plumieri</i> | | 61 / | 0.147 | 0.42 ** | Pauly (1978) |

continued...

Table 3.7 continued

| Family | Species | Origin | L _∞ (cm) | K year ⁻¹ | M year ⁻¹ | Source ^a |
|---------------|----------------------------------|------------------|------------------------|-------------------------|-------------------------|-----------------------------------------------------------------------------------------------------------|
| | <i>Lutjanus synagris</i> | Cuba | 42 FL | 0.268 | 0.69 * | Pauly (1978) |
| | | Florida | 50 TL | 0.134 | 0.42 * | Polovina and Ralston (1987) |
| | | Ceara | 50 | 0.231 | 0.59 * | Pauly (1978) |
| | <i>Ocyurus chrysurus</i> | Jamaica | 75 TL | 0.250 | 0.60 | Munro (1983) |
| | | Southern Florida | 45 FL | 0.279 | 0.69 * | Johnson and Altramadal (1983) |
| | | Jamaica | 60 FL | 0.250 | 0.60 | Munro (1983) |
| | | Jamaica | 60 FL | 0.250 | 0.62 | Pauly (1980) |
| | | | 69 TL | 0.160 | 0.20 | Bannerot et al. (1987) |
| | <i>Rhomboplites aurorubens</i> | | 63 | 0.198 | 0.50 * | Polovina and Ralston (1987) |
| Malacanthidae | <i>Malacanthus plumieri</i> | | 61 / | 0.147 | 0.42 ** | Pauly (1978) |
| Megalopidae | <i>Tarpon atlanticus</i> | Ceara | 263.3 FLf | 0.065 | 0.16 * | Ferreira de Menezes and Pinto Paiva (1966) |
| | | Ceara | 206.2 FLm | 0.084 | 0.21 * | Ferreira de Menezes and Pinto Paiva (1966) |
| Mugilidae | <i>Mugil curema</i> | | 38 | 0.341 | 0.83 ** | Pauly (1978) |
| | | | 53 | 0.1 | 0.34 * | Alvarez-Lajonchere (1981) |
| Mullidae | <i>Mulloidichthys martinicus</i> | Jamaica | 30 | 0.400 | 0.98 * | Nicholson and Hartsuijker (1982) |
| | | Jamaica | 33 | 0.190 | 0.59 * | Nicholson and Hartsuijker (1982) |
| | | Jamaica | 35 | 0.400 | 1.70 | Munro (1983) |
| | <i>Pseudupeneus maculatus</i> | Jamaica | 25 f | 0.350 | 0.95 | Munro (1983) |
| | | Jamaica | 27 m | 0.700 | 1.89 | Munro (1983) |
| | | Jamaica | 29 | 0.270 | 0.77 * | Nicholson and Hartsuijker (1983) |
| Pomacanthidae | <i>Holacanthus ciliaris</i> | | 46 | 0.136 | 0.43 ** | Pauly (1986, Table 2; ϕ for <i>Holacanthus bermudensis</i>) |
| | <i>Holacanthus tricolor</i> | | 36 | 0.180 | 0.55 ** | Pauly (1986, Table 2; ϕ for <i>Holacanthus bermudensis</i>) |
| | <i>Pomacanthus arcuatus</i> | Jamaica | 41 | 0.230 | 0.63 * | Nicholson and Hartsuijker (1983) |
| | | Jamaica | 47 | 0.214 | 0.57 * | Pauly (1978) |
| | | Jamaica | 60 | | 1.78 > | Munro (1983) |
| | <i>Pomacanthus paru</i> | | 43 | 0.231 | 0.62 ** | Based on ϕ ' of <i>Pomacanthus arcuatus</i> |
| Pomacentridae | <i>Stegastes fuscus</i> | | 15 / | | 1.60 | Meekan (1988); mortality of 24%/month for <i>Pomacentrus</i> sp. modified into an annual exponential rate |
| | <i>Stegastes leucostictus</i> | | 10 | | 1.60 | Meekan (1988); mortality of 24%/month for <i>Pomacentrus</i> sp. |
| | <i>Stegastes planifrons</i> | | 10.2 f | 0.580 | 1.69 * | Pauly (1981) |
| | | | 11.6 m | 0.330 | 1.13 * | Pauly (1981) |
| | <i>Stegastes variabilis</i> | | 10 + | | 1.60 | Meekan (1988); mortality of 24%/month for <i>Pomacentrus</i> sp. modified into an annual exponential rate |

continued...

Table 3.7 continued

| Family | Species | Origin | L _∞ (cm) | K year ⁻¹ | M year ⁻¹ | Source ^a |
|----------------|--------------------------------|----------------------------|------------------------|-------------------------|-----------------------------|----------------------------------------------------------------------|
| Pomatomidae | <i>Pomatomus saltator</i> | Cape Cod, Gulf of Maine | 115 | 0.135 | 0.33 * | Pauly (1978) |
| Rachycentridae | <i>Rachycentron canadum</i> | Chesapeake Bay | 134 | 0.210 | 0.42 * | Pauly (1978) |
| | | | 160 | 0.200 | 0.39 * | Pauly (1978) |
| Scaridae | <i>Scarus coeruleus</i> | VI | 61.5 | 0.325 | 0.70 * | Pauly (1978) |
| | <i>Scarus guacamaia</i> | VI | 80 | 0.293 | 0.61 * | Pauly (1978) |
| | <i>Scarus taeniopterus</i> | Jamaica | 35 FL | | 1.59 > | Munro (1983) |
| | | Jamaica | 31 FL | 0.305 | 0.81 * | Nicholson and Hartsuijker (1982) |
| | <i>Scarus vetula</i> | VI | 53 SL | 0.599 | 1.09 * | Pauly (1978) |
| | <i>Sparisoma aurofrenatum</i> | Jamaica | 26 FL | 0.350 | 0.94 * | Nicholson and Hartsuijker (1982) |
| | <i>Sparisoma chrysopterus</i> | VI | 42 FL | 0.782 | 1.38 * | Pauly (1978) |
| | <i>Sparisoma rubripinne</i> | VI | 46.5 | 0.584 | 1.11 * | Pauly (1978) |
| Scombridae | <i>Scomberomorus cavalla</i> | Ceara | 116 | 0.180 | 0.40 * | Pauly (1978) |
| | | Ceara | 137 | 0.150 | 0.34 * | Pauly (1978) |
| | <i>Scomberomorus maculatus</i> | Florida | 77 FL | 0.356 | 0.70 * | Pauly (1978) |
| | <i>Scomberomorus regalis</i> | | 83.5 FL | 0.303 | 0.61 ** | Pauly (1978) |
| | <i>Thunnus atlanticus</i> | Cuba | 78 | 0.330 | 0.67 | Pauly (1978) |
| | | Florida | 90 | 0.216 | 0.48 * | Pauly (1978) |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | | 36 / | 0.197 | 0.59 ** | Pauly (1978) |
| | <i>Scorpaena grandicornis</i> | | 18 / | 0.788 | 1.76 ** | Pauly (1978) |
| | <i>Scorpaena guttata</i> | California | 35 | 0.389 | 0.92 * | Pauly (1978) |
| | <i>Scorpaena plumieri</i> | | 43 / | 0.138 | 0.44 ** | Pauly (1978) |
| | <i>Scorpaenodes caribbaeus</i> | | 13 / | | | |
| Serranidae | <i>Cephalopholis cruentata</i> | Jamaica | 34 | 0.340 | 0.85 * | Polovina and Ralston (1987) |
| | | Curaçao | 42 | 0.130 | 0.43 * | Polovina and Ralston (1987) |
| | <i>Cephalopholis fulva</i> | Jamaica | 34 | 0.630 | 1.95 | Munro (1983) |
| | | Jamaica | 34 | 0.630 | 0.55 | Pauly (1980) |
| | | Jamaica | 34 | 0.440 | 1.01 * | Nicholson and Hartsuijker (1982) |
| | | Jamaica | 37.5 | 0.175 | 0.54 * | Nicholson and Hartsuijker (1982) |
| | <i>Diplectrum formosum</i> | Florida | 31 | 0.287 | 0.78 * | Pauly (1978) |
| | <i>Epinephelus guttatus</i> | Jamaica | 52 TL | 0.240 | 0.68 | Munro (1983) |
| | | Jamaica | 53 TL | 0.480 | 0.94 * | Computed with ELEFAN, based on length-frequency data in Munro (1983) |
| | <i>Epinephelus morio</i> | Florida | 67 | 0.179 | 0.46 * | Polovina and Ralston (1987) |
| | | Mexico | 93 | 0.113 | 0.31 * | Polovina and Ralston (1987) |
| | <i>Epinephelus striatus</i> | Jamaica | 90 | 0.090 | 0.24 | Munro (1983) |
| | VI | 97 | 0.183 | 0.36 | Polovina and Ralston (1987) | |
| | VI | 110 | 0.224 | 0.47 * | Pauly (1978) | |

continued...

Table 3.7 continued

| Family | Species | Origin | L_{∞} (cm) | K year ⁻¹ | M year ⁻¹ | Source ^a |
|--------------|--------------------------------|----------------|----------------------|-------------------------|-------------------------|------------------------------------------------------------|
| | <i>Mycteroperca bonaci</i> | | 120 | 0.160 | 0.37 * | Polovina and Ralston (1987) |
| | <i>Mycteroperca microlepis</i> | Florida | 101 TL | 0.156 | 0.38 * | Polovina and Ralston (1987) |
| | | Florida | 124 TL | 0.124 | 0.31 * | Pauly (1978) |
| | <i>Mycteroperca phenax</i> | Florida | 72 TL | 0.166 | 0.43 * | Polovina and Ralston (1987) |
| | <i>Mycteroperca venenosa</i> | Jamaica | 86 | 0.170 | 0.42 | Munro (1983) |
| | | Jamaica | 86 | 0.100 | 0.30 * | Polovina and Ralston (1987) |
| | <i>Paranthias furcifer</i> | Gulf of Mexico | 37 | 0.220 | 0.63 * | Polovina and Ralston (1987) |
| | <i>Serranus tabacarius</i> | | 18 | 0.086 | 0.41 ** | Bouain (1983); mean ϕ' computed from three species of |
| | <i>Serranus tigrinus</i> | | 10 | 0.279 | 1.05 ** | Bouain (1983); mean ϕ' computed from three species of |
| | | | | | | genus |
| | <i>Serranus tortugarum</i> | | 8 | 0.436 | 1.50 ** | Bouain (1983); mean ϕ' computed from three species of |
| | | | | | | genus |
| Sphyraenidae | <i>Sphyraena barracuda</i> | Florida | 156 f | 0.113 | 0.27 * | Pauly (1978) |
| | | Florida | 178 m | 0.087 | 0.22 * | Pauly (1978) |
| Sphyrnidae | <i>Sphyma tiburo</i> | | 150 | 0.11 | 0.27 * | Pauly (1978) |
| Triakidae | <i>Mustelus canis</i> | | 150 | 0.138 | 0.31 * | Pauly (1978) |

^aFor values of M and/or K [when M was computed by equations (2.9), (2.10) and (2.15)].

Symbols:

- FL = fork length.
- SL = standard length.
- TL = total length.
- f = female.
- m = male.
- VI = Virgin Islands.
- * = computed with equation (2.15).
- ** = computed via ϕ or ϕ' [equations (2.9) or (2.10)] and equation (2.15).
- > = M/K divided by K or K_{mean} from other source.
- / = approximate asymptotic size.
- + = rather more than the recorded size.
- = rather less than the recorded size.
- () = 'reported' to reach the recorded size.

Table 3.8. Estimated food consumption of Caribbean reef fishes. Q/B values represent mean population consumption of the corresponding species; mean environmental temperature = 28°C.

| Family | Species | Food type | Aspect ratio | W _∞ (g) | Q/B (%BWD) | Q/B _p (year ⁻¹) |
|----------------|------------------------------------|-----------|--------------|--------------------|------------|----------------------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 0.89 | 2.18 | 288 | 9.42 | 34.38 |
| | <i>Acanthurus chirurgus</i> | 0.88 | 1.78 | 820 | 6.78 | 23.87 |
| | <i>Acanthurus coeruleus</i> | 0.88 | 1.86 | 983 | 6.68 | 24.38 |
| Antennariidae | <i>Antennarius multiocellatus</i> | 0 | 1.22 | 70 | 3.03 | 11.06 |
| | <i>Antennarius striatus</i> | 0 | 0.80 | 70 | 2.43 | 8.87 |
| Apogonidae | <i>Apogon conklini</i> | 0 | 1.72 | 3.3 | 6.69 | 24.42 |
| | <i>Apogon maculatus</i> | 0 | 1.53 | 37 | 3.87 | 14.13 |
| Atherinidae | <i>Allanetta harringtonensis</i> | 0 | 1.71 | 11 | 5.24 | 19.12 |
| | <i>Atherinomorus stipes</i> | 0 | 1.88 | 16 | 5.10 | 18.60 |
| Aulostomidae | <i>Aulostomus maculatus</i> | 0 | 1.70 | 777 | 2.21 | 8.06 |
| Balistidae | <i>Balistes caprisus</i> | 0.50 * | 2.44 | 611 | 5.25 | 19.16 |
| | <i>Balistes vetula</i> | 0.12 | 1.98 | 2,586 | 2.18 | 7.96 |
| | <i>Canthidermis sufflamen</i> | 0.25 | 1.71 | 4,783 | 2.10 | 7.67 |
| | <i>Melichthys niger</i> | 0.75 | 2.30 | 960 | 6.36 | 23.21 |
| Belonidae | <i>Platybelone argalus argalus</i> | 0 | 2.65 | 150 | 3.87 | 14.12 |
| | <i>Strongylura timucu</i> | 0 | 1.69 | 110 | 3.26 | 11.91 |
| | <i>Tylosurus crocodilus</i> | 0 | 3.11 | 1,400 | 2.68 | 9.78 |
| Blenniidae | <i>Entomacrodus nigricans</i> | 0.84 | 1.02 | 13 | 11.16 | 40.72 |
| | <i>Ophioblennius atlanticus</i> | 0.87 | 0.92 | 35 | 9.01 | 32.88 |
| | <i>Parablennius marmoratus</i> | 0.69 | 1.26 | 10 | 10.87 | 39.67 |
| | <i>Scartella cristata</i> | 0.87 | 1.18 | 27 | 10.76 | 39.29 |
| Bothidae | <i>Bothus lunatus</i> | 0 | 1.11 | 3,000 | 1.35 | 4.94 |
| | <i>Bothus ocellatus</i> | 0 | 1.51 | 180 | 2.79 | 10.19 |
| Carangidae | <i>Alectis ciliaris</i> | 0 * | 4.82 | 10,700 | 2.23 | 8.13 |
| | <i>Caranx bartholomaei</i> | 0 | 4.06 | 9,862 | 2.07 | 7.56 |
| | <i>Caranx crysos (fusus)</i> | 0 | 4.19 | 2,314 | 2.82 | 10.31 |
| | <i>Caranx hippos</i> | 0 * | 3.76 | 8,490 | 2.05 | 7.50 |
| | <i>Caranx latus</i> | 0 | 4.48 | 6,250 | 2.39 | 8.72 |
| | <i>Caranx lugubris</i> | 0 | 5.48 | 6,554 | 2.63 | 9.59 |
| | <i>Caranx ruber</i> | 0 | 4.52 | 3,160 | 2.76 | 10.06 |
| | <i>Decapterus macarellus</i> | 0 | 3.80 | 750 | 3.37 | 12.30 |
| | <i>Decapterus punctatus</i> | 0 | 3.66 | 573 | 3.49 | 12.73 |
| | <i>Oligoplites saurus</i> | 0 | 3.12 | 334 | 3.58 | 13.08 |
| | <i>Selar crumenophthalmus</i> | 0 | 3.54 | 1,240 | 2.93 | 10.70 |
| | <i>Seriola dumerili</i> | 0 | 4.49 | 80,000 | 1.43 | 5.22 |
| | <i>Trachinotus falcatus</i> | 0 | 3.88 | 36,000 | 1.56 | 5.69 |
| | <i>Trachinotus goodei</i> | 0 | 5.01 | 3,900 | 2.79 | 10.18 |
| Carcharhinidae | <i>Carcharhinus acronotus</i> | 0 | 7 * | 90,000 | 1.76 | 6.41 |
| | <i>Carcharhinus falciformis</i> | 0 | 7 * | 350,000 | 1.34 | 4.87 |
| | <i>Carcharhinus leucas</i> | 0 | 7 * | 350,000 | 1.34 | 4.87 |
| | <i>Carchannus leucas</i> | 0 | 3.85 S | 350,000 | 0.98 | 3.58 |
| | <i>Carcharhinus limbatus</i> | 0 | 7 * | 116,000 | 1.67 | 6.09 |
| | <i>Carcharhinus longimanus</i> | 0 | 7 * | 350,000 | 1.34 | 4.87 |
| | <i>Carcharhinus perezi</i> | 0 | 7 * | 100,000 | 1.72 | 6.28 |
| | <i>Galeocerdo cuvier</i> | 0 | 7 * | 672,000 | 1.17 | 4.27 |
| | <i>Negaprion brevirostris</i> | 0 | 7 * | 110,000 | 1.69 | 6.16 |
| | <i>Negaprion brevirostris</i> | 0 | 1.56 S | 110,000 | 0.78 | 2.84 |
| | <i>Rhizoprionodon porosus</i> | 0 | 7 * | 5,020 | 3.15 | 11.48 |
| Chaetodontidae | <i>Chaetodon aculeatus</i> | 0 | 1.85 | 19 | 4.88 | 17.82 |
| | <i>Chaetodon capistratus</i> | 0 | 2.53 | 120 | 3.95 | 14.42 |

continued...

Table 3.8 continued

| Family | Species | Food type | Aspect ratio | W _∞ (g) | Q/B (%BWD) | Q/B ₁ (year ⁻¹) |
|------------------------------|------------------------------------|-----------|--------------|--------------------|------------|----------------------------------------|
| | <i>Chaetodon sedentarius</i> | 0 | 2.29 | 130 | 3.70 | 13.51 |
| | <i>Chaetodon striatus</i> | 0 | 2.37 | 164 | 3.59 | 13.09 |
| Cirrhitidae | <i>Amblycirrhitus pinos</i> | 0 | 1.52 | 15 | 4.62 | 16.86 |
| Clinidae | <i>Labrisomus nuchipinnis</i> | 0 | 1.29 | 137 | 2.72 | 9.93 |
| Clupeidae | <i>Harengula clupeiola</i> | 0 | 2.65 | 240 | 3.52 | 12.85 |
| | <i>Harengula humeralis</i> | 0.03 | 2.90 | 440 | 3.39 | 12.37 |
| | <i>Jenkinsia lamprotaenia</i> | 0 | 2.19 | 13 | 5.74 | 20.95 |
| | <i>Opisthonema oglinum</i> | 0 | 3.13 | 730 | 3.06 | 11.18 |
| Dactylopteridae | <i>Dactylopterus volitans</i> | 0 | 1.60 | 1,825 | 1.80 | 6.57 |
| Dasyatidae | <i>Dasyatis americana</i> | 0 | 7 * | 122,000 | 1.65 | 6.03 |
| Diodontidae | <i>Chilomycterus antennatus</i> | 0 | 1.40 | 1,420 | 1.77 | 6.46 |
| | <i>Diodon holocanthus</i> | 0 | 1.37 | 11,300 | 1.15 | 4.20 |
| | <i>Diodon hystrix</i> | 0 | 1.09 | 18,446 | 0.93 | 3.38 |
| Echeneidae | <i>Echeneis naucrates</i> | 0 | 1.10 | 2,080 | 1.45 | 5.29 |
| | <i>Remora remora</i> | 0 | 1.60 | 1,700 | 1.83 | 6.68 |
| Elopidae | <i>Elops saurus</i> | 0 * | 3.41 | 444 | 3.54 | 12.92 |
| Emmelichthyidae | <i>Inermia vittata</i> | 0 | 2.92 | 282 | 3.58 | 13.07 |
| Engraulidae | <i>Anchoa lyolepis</i> | 1 * | 1.77 | 8 | 19.99 | 72.96 |
| Ephippidae | <i>Chaetodipterus faber</i> | 0.07 | 3.11 | 9,000 | 2.01 | 7.34 |
| Fistulariidae | <i>Fistularia tabacaria</i> | 0 | 1.70 | 10,800 | 1.30 | 4.75 |
| Gerreidae | <i>Eucinostomus argenteus</i> | 0 | 1.94 | 290 | 2.89 | 10.55 |
| | <i>Gerres cinereus</i> | 0 | 3.91 | 690 | 3.48 | 12.69 |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | 0 | 7 * | 500,000 | 1.24 | 4.54 |
| | <i>Ginglymostoma cirratum</i> | 0 | 0.76 S | 500,000 | 0.40 | 1.44 |
| Gobiidae | <i>Coryphopterus glaucofraenum</i> | 0.44 | 1.11 | 10.5 | 7.35 | 26.83 |
| | <i>Gnatholepis thompsoni</i> | 0.65 | 0.96 | 4.1 | 10.72 | 39.13 |
| | <i>Gobiosoma evelynae</i> | 0 | 0.79 | 2.2 | 4.85 | 17.70 |
| Grammidae (Serranidae) | <i>Gramma loreto</i> | 0 | 1.13 | 15 | 3.97 | 14.49 |
| | <i>Gramma melacara</i> | 0 | 1.36 | 30 | 3.80 | 13.87 |
| Grammistidae (Serranidae) | <i>Rypticus saponaceus</i> | 0 | 1.37 | 685 | 2.03 | 7.41 |
| Haemulidae (Pomadasyidae) | <i>Anisotremus surinamensis</i> | 0 | 2.66 | 4,770 | 1.93 | 7.04 |
| | <i>Anisotremus virginicus</i> | 0 | 2.84 | 850 | 2.82 | 10.31 |
| | <i>Haemulon album</i> | 0 | 2.77 | 5,300 | 1.93 | 7.04 |
| | <i>Haemulon aurolineatum</i> | 0 | 2.44 | 188 | 3.55 | 12.95 |
| | <i>Haemulon bonariense</i> | 0 * | 2.45 | 385 | 3.07 | 11.22 |
| | <i>Haemulon carbonarium</i> | 0 | 2.84 | 861 | 2.82 | 10.29 |
| | <i>Haemulon chrysargyreum</i> | 0 | 2.59 | 251 | 3.45 | 12.58 |
| | <i>Haemulon flavolineatum</i> | 0 | 2.46 | 480 | 2.94 | 10.75 |
| | <i>Haemulon macrostomum</i> | 0 | 2.69 | 1,795 | 2.36 | 8.63 |
| | <i>Haemulon melanurum</i> | 0 * | 1.79 | 427 | 2.56 | 9.34 |
| | <i>Haemulon parrai</i> | 0 | 2.53 | 1,265 | 2.46 | 8.98 |
| | <i>Haemulon plumieri</i> | 0 | 2.86 | 1,360 | 2.58 | 9.42 |
| | <i>Haemulon sciurus</i> | 0 | 2.45 | 1,185 | 2.45 | 8.94 |

continued...

Table 3.8 continued

| Family | Species | Food type | Aspect ratio | W_{∞} (g) | Q/B (%BWD) | Q/B (year ⁻¹) |
|---------------|----------------------------------|-----------|--------------|------------------|------------|---------------------------|
| | <i>Haemulon steindachneri</i> | 0 * | 2.02 | 360 | 2.82 | 10.29 |
| | <i>Haemulon striatum</i> | 0 * | 1.83 | 400 | 2.62 | 9.58 |
| | <i>Orthopristis chrysoptera</i> | 0 * | 2.46 | 900 | 2.59 | 9.47 |
| | <i>Orthopristis ruber</i> | 0 * | 2.46 | 330 | 3.17 | 11.58 |
| Hemiramphidae | <i>Hemiramphus balao</i> | 0 | 2.73 | 614 | 2.96 | 10.79 |
| | <i>Hemiramphus brasiliensis</i> | 0.81 | 3.23 | 298 | 10.36 | 37.80 |
| Holocentridae | <i>Holocentrus ascensionis</i> | 0 | 2.64 | 838 | 2.73 | 9.97 |
| | <i>Holocentrus coruscus</i> | 0 | 3.03 | 57 | 5.05 | 18.42 |
| | <i>Holocentrus rufus</i> | 0 | 2.23 | 583 | 2.69 | 9.82 |
| | <i>Myripristis jacobus</i> | 0 | 2.46 | 397 | 3.06 | 11.16 |
| | <i>Neoniphon marianus</i> | 0 | 3.02 | 163 | 4.07 | 14.86 |
| | <i>Plectrypops retrospinis</i> | 0 | 1.73 | 57 | 3.77 | 13.77 |
| | <i>Sargocentron vexillarium</i> | 0 | 2.65 | 181 | 3.73 | 13.60 |
| Kyphosidae | <i>Kyphosus incisor</i> | 1 | 2.83 | 6,122 | 6.68 | 24.37 |
| | <i>Kyphosus sectatrix</i> | 1 | 2.90 | 9,139 | 6.23 | 22.76 |
| Labridae | <i>Bodianus rufus</i> | 0 | 1.20 | 1,560 | 1.60 | 5.85 |
| | <i>Clepticus parrae</i> | 0 | 1.84 | 447 | 2.58 | 9.40 |
| | <i>Halichoeres bivittatus</i> | 0 | 1.27 | 230 | 2.43 | 8.88 |
| | <i>Halichoeres garnoti</i> | 0 | 1.44 | 132 | 2.90 | 10.59 |
| | <i>Halichoeres maculipinna</i> | 0 | 1.25 | 25 | 3.77 | 13.77 |
| | <i>Halichoeres poeyi</i> | 0 | 1.19 | 150 | 2.56 | 9.35 |
| | <i>Halichoeres radiatus</i> | 0 | 1.58 | 2,283 | 1.71 | 6.25 |
| | <i>Lachnolaimus maximus</i> | 0 | 1.51 | 7,728 | 1.31 | 4.78 |
| | <i>Thalassoma bifasciatum</i> | 0 | 1.07 | 97 | 2.65 | 9.66 |
| | <i>Xyrichtys novacula</i> | 0 | 1.48 | 230 | 2.63 | 9.60 |
| | <i>Xyrichtys splendens</i> | 0 | 1.48 | 50 | 3.58 | 13.06 |
| Lutjanidae | <i>Lutjanus analis</i> | 0 | 2.06 | 5,511 | 1.64 | 5.99 |
| | <i>Lutjanus apodus</i> | 0 | 2.00 | 3,502 | 1.77 | 6.46 |
| | <i>Lutjanus campechanus</i> | 0 * | 1.41 | 18,000 | 1.06 | 3.88 |
| | <i>Lutjanus cyanopterus</i> | 0 | 2.18 | 70,000 | 1.01 | 3.69 |
| | <i>Lutjanus griseus</i> | 0 | 2.06 | 1,590 | 2.11 | 7.70 |
| | <i>Lutjanus jocu</i> | 0 | 2.08 | 13,380 | 1.38 | 5.04 |
| | <i>Lutjanus mahagoni</i> | 0 | 2.05 | 980 | 2.32 | 8.47 |
| | <i>Lutjanus synagris</i> | 0 | 1.97 | 1,213 | 2.18 | 7.95 |
| | <i>Ocyurus chrysurus</i> | 0 | 2.96 | 3,570 | 2.16 | 7.89 |
| Malacanthidae | <i>Malacanthus plumieri</i> | 0 | 1.42 | 1,126 | 1.87 | 6.82 |
| Megalopidae | <i>Tarpon atlanticus</i> | 0 | 2.96 | 160,000 | 1.00 | 3.66 |
| Monacanthidae | <i>Aluterus monoceros</i> | 0.50 * | 1.70 | 2,500 | 3.28 | 11.96 |
| | <i>Aluterus schoepfii</i> | 0.99 | 0.72 | 1,892 | 4.13 | 15.09 |
| | <i>Aluterus scripta</i> | 0.43 | 0.72 | 2,941 | 1.86 | 6.79 |
| | <i>Cantherines macrocerus</i> | 0.29 | 1.46 | 2,700 | 2.29 | 8.35 |
| | <i>Cantherines pullus</i> | 0.42 | 1.47 | 220 | 4.48 | 16.37 |
| | <i>Monacanthus ciliatus</i> | 0.34 | 1.24 | 448 | 3.21 | 11.73 |
| | <i>Monacanthus tuckeri</i> | 0.31 | 1.37 | 24 | 5.90 | 21.53 |
| Mugilidae | <i>Mugil curema</i> | 1 | 2.72 | 767 | 9.95 | 36.33 |
| Mullidae | <i>Mulloidichthys martinicus</i> | 0 | 2.26 | 440 | 2.87 | 10.48 |
| | <i>Pseudupeneus maculatus</i> | 0 | 2.30 | 393 | 2.96 | 10.80 |
| Muraenidae | <i>Enchelycore nigricans</i> | 0 | 0.7 * | 900 | 1.36 | 4.95 |
| | <i>Echidna catenata</i> | 0 | 0.7 * | 600 | 1.47 | 5.38 |
| | <i>Gymnothorax funebris</i> | 0 | 0.7 * | 12,000 | 0.80 | 2.94 |
| | <i>Gymnothorax miliaris</i> | 0 | 0.7 * | 201 | 1.84 | 6.70 |

continued...

Table 3.8 continued

| Family | Species | Food type | Aspect ratio | W_{∞} (g) | Q/B (%BWD) | Q/B (year ⁻¹) |
|------------------|-------------------------------------|-----------|--------------|------------------|------------|---------------------------|
| | <i>Gymnothorax vicinus</i> | 0 | 0.7 * | 3,000 | 1.06 | 3.89 |
| | <i>Lycodontis moringa</i> | 0 | 0.7 * | 950 | 1.34 | 4.90 |
| Myliobatidae | <i>Aetobatus narinari</i> | 0 | 7 * | 230,000 | 1.45 | 5.31 |
| Ophichthidae | <i>Myrichthys breviceps</i> | 0 | 0.7 * | 640 | 1.45 | 5.31 |
| | <i>Myrichthys ocellatus</i> | 0 | 0.7 * | 640 | 1.45 | 5.31 |
| | <i>Ophichthus ophis</i> | 0 | 0.7 * | 2,091 | 1.14 | 4.18 |
| Opisthognathidae | <i>Opisthognathus aurifrons</i> | 0 | 0.84 | 30 | 2.96 | 10.81 |
| | <i>Opisthognathus macrognathus</i> | 0 | 1.14 | 240 | 2.28 | 8.33 |
| | <i>Opisthognathus whitehurstii</i> | 0 | 0.82 | 65 | 2.51 | 9.15 |
| | <i>Opisthognathus maxillosus</i> | 0 | 0.76 | 15 | 3.24 | 11.81 |
| Ostraciidae | <i>Acanthostracion polygonius</i> | 0 | 1.17 | 2,200 | 1.47 | 5.38 |
| | <i>Acanthostracion quadricornis</i> | 0.1 | 1.14 | 3,050 | 1.55 | 5.65 |
| | <i>Lactophrys bicaudalis</i> | 0.15 | 1.23 | 5,572 | 1.52 | 5.55 |
| | <i>Lactophrys trigonus</i> | 0.08 | 1.87 | 3,052 | 1.95 | 7.11 |
| | <i>Lactophrys triqueter</i> | 0.02 | 1.98 | 1,394 | 2.18 | 7.96 |
| Pempheridae | <i>Pempheris schomburgki</i> | 0 | 2.28 | 100 | 3.89 | 14.19 |
| Pomacanthidae | <i>Centropyge argi</i> | 0.75 | 1.44 | 10 | 12.57 | 45.90 |
| | <i>Holacanthus ciliaris</i> | 0.02 | 1.29 | 1,988 | 1.63 | 5.95 |
| | <i>Holacanthus tricolor</i> | 0.01 | 1.65 | 1,306 | 1.99 | 7.26 |
| | <i>Pomacanthus arcuatus</i> | 0.08 | 2.89 | 12,407 | 1.84 | 6.72 |
| | <i>Pomacanthus paru</i> VI | 0.13 | 1.81 | 2,769 | 2.08 | 7.59 |
| | <i>Pomacanthus paru</i> Jamaica | 0.13 | 1.81 | 2,229 | 2.17 | 7.92 |
| Pomacentridae | <i>Abudefduf saxatilis</i> | 0.09 | 2.98 | 483 | 3.64 | 13.28 |
| | <i>Abudefduf taurus</i> | 0.94 | 2.01 | 900 | 7.64 | 27.88 |
| | <i>Chromis cyanea</i> | 0 | 1.79 | 93 | 3.49 | 12.72 |
| | <i>Chromis multilineata</i> | 0 | 1.79 | 339 | 2.68 | 9.78 |
| | <i>Microspathodon chrysurus</i> | 0.78 | 1.88 | 412 | 7.06 | 25.77 |
| | <i>Stegastes fuscus</i> | 0.51 | 1.83 | 168 | 5.94 | 21.67 |
| | <i>Stegastes leucostictus</i> | 0.2 | 1.31 | 50 | 4.32 | 15.77 |
| | <i>Stegastes planifrons</i> | 0.25 | 1.37 | 68 | 4.43 | 16.16 |
| | <i>Stegastes variabilis</i> | 0.45 | 1.54 | 54 | 6.34 | 23.13 |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | 0 | 1.79 | 700 | 2.32 | 8.46 |
| | <i>Priacanthus arenatus</i> | 0 | 1.83 | 1,653 | 1.97 | 7.20 |
| Rachycentridae | <i>Rachycentron canadum</i> | 0 | 1.56 | 33,400 | 0.99 | 3.62 |
| Scaridae | <i>Scarus coelestinus</i> | 0.99 | 1.06 | 8,556 | 3.71 | 13.53 |
| | <i>Scarus coeruleus</i> | 1 * | 1.48 | 3,720 | 5.28 | 19.27 |
| | <i>Scarus guacamaia</i> | 1 | 1.19 | 23,000 | 3.27 | 11.95 |
| | <i>Scarus iserti</i> | 1 * | 1.44 | 141 | 10.09 | 36.82 |
| | <i>Scarus taeniopterus</i> | 0.99 | 1.09 | 1,102 | 5.70 | 20.82 |
| | <i>Scarus vetula</i> | 0.97 | 1.05 | 5,558 | 3.94 | 14.37 |
| | <i>Sparisoma aurofrenatum</i> | 0.99 | 1.33 | 324 | 8.07 | 29.47 |
| | <i>Sparisoma chrysopterygum</i> | 1 | 1.44 | 1,510 | 6.25 | 22.82 |
| | <i>Sparisoma radians</i> | 1 | 1.26 | 150 | 9.28 | 33.88 |
| | <i>Sparisoma rubripinne</i> | 1 | 1.51 | 2,734 | 5.69 | 20.77 |
| | <i>Sparisoma viride</i> | 1 | 1.43 | 2,430 | 5.66 | 20.66 |
| Sciaenidae | <i>Equetus lanceolatus</i> | 0 | 0.97 | 325 | 1.98 | 7.21 |
| | <i>Equetus punctatus</i> | 0 | 1.25 | 460 | 2.10 | 7.66 |
| | <i>Odontoscion dentex</i> | 0 | 1.68 | 318 | 2.63 | 9.59 |
| | <i>Pareques acuminatus</i> | 0 | 1.34 | 280 | 2.40 | 8.77 |
| Scombridae | <i>Euthynnus aletteratus</i> | 0 | 6.29 | 12,200 | 2.49 | 9.08 |
| | <i>Scomberomorus cavalla</i> | 0 | 6.31 | 34,285 | 2.02 | 7.39 |

continued...

Table 3.8 continued

| Family | Species | Food type | Aspect ratio | W _∞ (g) | Q/B (%BWD) | Q/B (year ⁻¹) |
|----------------|------------------------------------|-----------|--------------|--------------------|------------|---------------------------|
| | <i>Scomberomorus maculatus</i> | 0 * | 4.87 | 4,800 | 2.63 | 9.61 |
| | <i>Scomberomorus regalis</i> | 0 | 6.16 | 4,900 | 2.96 | 10.80 |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | 0 | 1.03 | 1,600 | 1.47 | 5.37 |
| | <i>Scorpaena grandicornis</i> | 0 | 1.03 | 212 | 2.22 | 8.09 |
| | <i>Scorpaena plumieri</i> | 0 | 1.00 | 4,330 | 1.19 | 4.33 |
| Serranidae | <i>Alphesthes afer</i> | 0 | 1.20 | 1,202 | 1.69 | 6.17 |
| | <i>Cephalopholis cruentata</i> | 0 | 1.23 | 691 | 1.91 | 6.99 |
| | <i>Cephalopholis fulva</i> | 0 | 1.47 | 640 | 2.13 | 7.78 |
| | <i>Dermatolepis inermis</i> | 0 * | 1.51 | 10,513 | 1.23 | 4.49 |
| | <i>Diplectrum formosum</i> | 0 * | 1.49 | 550 | 2.21 | 8.08 |
| | <i>Epinephelus adscensionis</i> | 0 | 1.53 | 1,981 | 1.73 | 6.33 |
| | <i>Epinephelus guttatus</i> | 0 | 1.03 | 2,919 | 1.31 | 4.77 |
| | <i>Epinephelus itajara</i> | 0 | 1.28 | 381,644 | 0.55 | 1.99 |
| | <i>Epinephelus morio</i> | 0 | 1.46 | 5,010 | 1.40 | 5.12 |
| | <i>Epinephelus striatus</i> | 0 | 1.57 | 23,800 | 1.06 | 3.88 |
| | <i>Hypoplectrus aberrans</i> | 0 | 1.84 | 50 | 4.01 | 14.62 |
| | <i>Hypoplectrus chlorurus</i> | 0 | 1.53 | 64 | 3.46 | 12.64 |
| | <i>Hypoplectrus nigricans</i> | 0 | 1.77 | 90 | 3.48 | 12.70 |
| | <i>Hypoplectrus puella</i> | 0 | 1.58 | 64 | 3.52 | 12.86 |
| | <i>Mycteroperca bonaci</i> | 0 | 1.22 | 90,000 | 0.71 | 2.60 |
| | <i>Mycteroperca cidi</i> | 0 * | 1.13 | 77,000 | 0.71 | 2.58 |
| | <i>Mycteroperca interstitialis</i> | 0 | 1.29 | 4,000 | 1.38 | 5.03 |
| | <i>Mycteroperca phenax</i> | 0 * | 1.36 | 43,700 | 0.87 | 3.18 |
| | <i>Mycteroperca rubra</i> | 0 * | 1.20 | 10,719 | 1.09 | 3.97 |
| | <i>Mycteroperca tigris</i> | 0 | 1.81 | 17,043 | 1.22 | 4.46 |
| | <i>Mycteroperca venenosa</i> | 0 | 1.08 | 12,270 | 1.00 | 3.66 |
| | <i>Paranthias furcifer</i> | 0 | 2.27 | 1,100 | 2.39 | 8.73 |
| | <i>Serranus dewegeri</i> | 0 | 1.40 | 700 | 2.04 | 7.45 |
| | <i>Serranus tabacarius</i> | 0 | 1.24 | 141 | 2.65 | 9.67 |
| | <i>Serranus tigrinus</i> | 0 | 1.45 | 21 | 4.21 | 15.37 |
| | <i>Serranus tortugarum</i> | 0 | 1.58 | 12 | 4.94 | 18.03 |
| Sparidae | <i>Archosargus rhomboidalis</i> | 0.83 | 2.47 | 1,200 | 6.98 | 25.46 |
| | <i>Calamus bajonado</i> | 0 | 3.04 | 4,500 | 2.09 | 7.63 |
| | <i>Calamus calamus</i> | 0 | 3.18 | 1,541 | 2.66 | 9.70 |
| | <i>Calamus penna</i> | 0 | 3.11 | 2,000 | 2.49 | 9.10 |
| | <i>Calamus pennatula</i> | 0 | 3.04 | 1,700 | 2.55 | 9.29 |
| | <i>Diplodus caudimacula</i> | 0.8 | 2.92 | 600 | 8.43 | 30.76 |
| Sphyraenidae | <i>Sphyraena barracuda</i> | 0 | 1.61 | 57,800 | 0.90 | 3.28 |
| | <i>Sphyraena picudilla</i> | 0 | 2.62 | 1,300 | 2.49 | 9.08 |
| Sphyrnidae | <i>Sphyrna lewini</i> | 0 | 7 * | 400,853 | 1.30 | 4.74 |
| | <i>Sphyrna tiburo</i> | 0 | 7 * | 18,000 | 2.43 | 8.87 |
| | <i>Sphyrna tiburo</i> | 0 | 1.63 S | 18,000 | 1.15 | 4.18 |
| Synodontidae | <i>Synodus intermedius</i> | 0 | 2.38 | 1,103 | 2.45 | 8.94 |
| | <i>Synodus foetens</i> | 0 | 2.65 | 1,100 | 2.59 | 9.45 |
| | <i>Synodus synodus</i> | 0 | 2.38 | 400 | 3.01 | 10.98 |
| Tetraodontidae | <i>Canthigaster rostrata</i> | 0.18 | 1.52 | 82 | 4.12 | 15.04 |
| | <i>Sphoeroides spengleri</i> | 0.08 | 1.57 | 92 | 3.60 | 13.16 |
| Triakidae | <i>Mustelus canis</i> | 0 | 7 * | 15,000 | 2.52 | 9.20 |

Symbols: * = assumed value.

VI = Virgin Islands.

S = aspect ratio from Table 1 in Pauly et al. (1993).

3.2. Nonfish Taxa

3.2.1. Ecology, Fisheries and Management

Although I assumed a coral reef ecosystem without fishing activities, which was realistic when Randall investigated food habits of West Indian reef fishes in 1967, some information on fisheries and management were included here to give an idea of later developments.

Boulon (1986b) gave a short description of spatial occurrence of the three commercially most important invertebrates in the Virgin Islands Biosphere Reserve:

Adults of *Strombus gigas*, the queen conch, are found from grass and mud-bottom mangrove embayments out to deep (10 m+) algal plains. They are most often found in beds of seagrass (*Thalassia testudinum*, *Syringodium filiforme*, or a mixture of both), but are also found on sand flats. Juvenile conch are generally found in shallower water and in areas of less dense seagrass. Young conch may be restricted in their movements in thick beds of seagrass.

Cittarium pica, the West Indian topshell, is primarily found in rocky, coastal habitats, from slightly above the water level to several feet below. They appear to prefer coasts that are subject to at least some wave action. The smaller individuals tend to be found higher up in the habitat range of this species with larger, mature individuals being found in deeper (1-2 m) subtidal bedrock or upper fore-reef zones.

Panulirus argus, the spiny lobster, appears to be most abundant near reef-grassflat boundaries. Juveniles were found in mangroves and *Thalassia* habitats. In the reef habitat, lobsters are primarily found in dense grass beds (?) where they spend the daylight hours. At night, they forage for food either on the reef or on adjacent algal plains.

Mahon (1987) gave a summary on ecology, fisheries and management of "shared fishery resources of the Lesser Antilles region". The only invertebrates discussed therein are queen conch and spiny lobster; the former is stated to be "the second most valuable resource in the Caribbean as a whole", while "conch stocks throughout the Caribbean region are considered to be fully or overexploited".

In Puerto Rico, all conch production is consumed locally. The west coast dominates the fisheries with landings averaging 68.2 t during the early and mid-1970s, and increasing to 189 t in 1982. Price per pound rose from US\$0.5 in 1975 to US\$1.37 in 1984, with a total worth for the latter year of \$628,350, thus, making conch one of the country's most commercially important species.^a

Inshore populations were noticeably reduced by the mid-1970s, when fishing pressure on deepwater areas increased. In the population off La Parguera, Puerto Rico, fishing mortality is estimated to be 68% ($F = 0.4 \text{ year}^{-1}$). The conch fishery is classified as "overfished".

A directed, commercial conch fishery exists on St. Croix, US Virgin Islands. St. Thomas and St. John are less fished because conch occur deeper and farther from shore. In 1982, 9.7 conch·ha⁻¹ and 1.58 * 10⁶ individuals were estimated for St. Thomas/St. John, and 7.6 conch·ha⁻¹ and 0.26 * 10⁶ individuals for St. Croix. Primary fishing grounds for conch in the US Virgin Islands are the west and northwest coasts of St. Croix and the south shelf on St. Thomas. The conch resources in the US Virgin Islands are classified as "overfished".

^aAll monetary figures are in US\$, unless otherwise stated.

Queen conch are harvested primarily by free and scuba diving. The regulations that have been enacted or proposed to reduce fishing pressure include: a minimum size limit on shell length (e.g., 18 cm), shell diameter, lip thickness or meat weight (e.g., 225 g market weight), seasonal closures, total closures, prohibitions on the use of scuba and on export.

In Puerto Rico, total spiny lobster landings decreased from 133.58 t in 1983 to 112.62 t in 1985, and the market value decreased from $\$1.04 \times 10^{-6}$ to $\$0.92 \times 10^{-6}$. For 1985-1986, approximately 20% of lobsters sampled were less than 76 mm carapace length (CL) as compared to 16% for 1984-1985. Regulations include a minimum CL of 89 mm; no capture of berried females; prohibition on the use of spears, hooks and drugs, poisons or explosives; and the use of degradable panels on lobster traps.

In the US Virgin Islands, estimated domestic landings were 22.7 t in 1975; this rose to 74 t in 1979, but the 1982 estimate was 45.4 t. Only 50% of lobster catches are assumed to be reported, which hinders reliable determination of stock size. Depletion of spiny lobster stocks is indicated by declining average size and the necessity for fishers to travel longer distances to the fishing grounds. Regulations are the same as for Puerto Rico.

Indications on the commercial importance of other nonfish taxa, such as shrimps and turtles, were not found. Indications on their ecology, however, were found in general zoology textbooks and in the references assembled for construction of their diet composition matrix (see below).

3.2.2. Abundance and Mean Individual Weight

While assembling the ECOPATH II input parameters for nonfish taxa, a number of estimates of abundance and weight per individual could be identified (Table 3.9).

3.2.3. Biomass, P/B Ratio and Food Consumption

Table 3.10 shows the complete database of the three ECOPATH II key input parameters for the 41 nonfish taxa originating from stomach contents records of West Indian reef fishes (Randall 1967). Following is a brief description on the origin of every range or single value, in form of 135 numbered "statements":^b

1. To assess total system net primary productivity (NPP), a selection of published values of coral reef gross and net PP was compiled from various sources (Table 3.11). Emphasis was put on information from the Caribbean Sea. Available values for distinct parts of the community were related to their corresponding subsystem(s) (mangrove, seagrass bed, sand flat, coral reef, algal plain). A mean NPP (in $\text{gCm}^{-2}\text{year}^{-1}$) was computed for every subsystem and from these again a mean total system NPP of ca. $1,200 \text{ gCm}^{-2}\text{year}^{-1}$ (= $20,000 \text{ gm}^{-2}\text{year}^{-1}\text{WW}$, conversion: carbon = 6% of WW) was obtained. Compared to overall NPP for coral reefs worldwide (e.g., in Lewis 1981 and Hatcher 1990, see Table PP), the value obtained by the above-described procedure for a Virgin Islands reef system seemed high. This value, however, was strongly influenced by the high NPP estimates reported for parts of the algal community by Connor and Adey (1977) for the Virgin Islands. The uncertainty of the estimate of total system NPP is critical since this value is a direct measure of the carrying capacity of the system. The reliability of this parameter has a strong impact on the degree of similarity in terms of size (biomass) between the real system and the model.

^bThe number corresponding to each statement may also be found in Table 3.10.

Table 3.9. Abundance (Nm⁻²) and mean individual live weight (g) of some nonfish taxa from Caribbean reefs and other areas.

| Nonfish group | Abundance (Nm ⁻²) | Source | Mean weight (gWW/l) | Source |
|------------------------------------------------------------------------|-------------------------------|--------------------------------------|---------------------|----------------------------------------------------------------|
| Sponges | 0.76 | Beets and Lewand (1986) ^b | 86-184 7.50 | Wilkinson (1987) ^m Wilkinson (1987) ^l |
| Sea fans (Gorgonacea) (Octocorallia, Anthozoa) | 0.26 | Beets and Lewand (1986) ^b | - | |
| Sea anemones (Actiniaria, Zoanthidea, Hexacorallia and Anthozoa) | 7.26 | Beets and Lewand (1986) ^b | - | |
| Gastropods | 3.01 | Beets and Lewand (1986) ^b | - | |
| Bivalves | 0.12 | Beets and Lewand (1986) ^b | - | |
| Octopuses | 0.01 | Beets and Lewand (1986) ^b | 2,800 | Guerra (1979) ^e |
| Polychaetes | 1.54 | Beets and Lewand (1986) ^b | 0.006 | Naim and Amoureux (1982) ^h |
| Amphipods | 516 | Berry (1982) ^c | 0.005 | Berry (1982) ^c |
| Isopods (fish parasites) | | | 0.14 | Schwinghamer et al. (1986, Table 1) ^j |
| Shrimps | 1.25 | Beets and Lewand (1986) ^b | - | |
| Spiny lobsters | 0.02 | Beets and Lewand (1986) ^b | 58.7 217 | Munro (1983) ^f Munro (1983) ^g |
| Scyllarid lobsters | 1.35 | Beets and Lewand (1986) ^b | - | |
| Crabs | 0.54 | Beets and Lewand (1986) ^b | - | |
| Ophiuroids | 2.5 | Beets and Lewand (1986) ^b | - | |
| Echinoids | 2.1 | Bak et al. (1984) ^a | 57 | Sambilay et al. (1990) ⁱ |
| | 3.3 | Beets and Lewand (1986) ^b | - | |
| | 2.6-7 | Weil et al. (1984) ^k | - | |
| Holothurians | 15 | Berry (1982) ^d | 4 | Berry (1982) ^d |
| Ascidians (Tunicata) | 0.69 | Beets and Lewand (1986) ^b | | |

Sources:

^aFrom Table 1 in Bak et al. (1984), an average density of 2.1 individuals per m² was computed for *Diadema antillarum* in Curaçao, Netherlands Antilles.

^bBeets and Lewand (1986, Tables 1-4) indicated density values (individuals m⁻²) for five phototransects of a US Virgin Islands reef area. The area covered was 0.5 m on either side of the transect line. Values in Table DMWNFG represented means for five transects.

^cBerry (1982, in Table 1) reported a mean density for amphipods (several species) on the ORI reef, Natal province, South Africa, of 516 individuals per m²; live weight is 1-3.5 gm⁻² (conversion: DW=25% of WW). Mean weight per individual is 4.5 mg.

^dBerry (1982), in Table 1, gave a mean density of 15 individuals per m² and a biomass equivalent to 55.1 kJm⁻² for holothurians on the ORI reef, Natal province, South Africa. 1 kJ corresponds to 1.08 g WW; mean individual live weight is 3.96 g.

^eGuerra (1979) gave growth estimates for *Octopus vulgaris* (in the Mediterranean) which yielded a W_{∞} of 10,850 g. W_{mean} was computed from equation (2.12).

^fMunro (1983) recorded a mean live weight of 58.7 g for individual *Panulirus guttatus* around Jamaica.

^gMunro (1983) reported a biomass of 63-119 kgkm⁻² and a density of 290-550 individuals per km² for *P. argus* in Jamaican waters. Hence, mean individual live weight is 217 g.

^hNaim and Amoureux (1982) gave a maximum polychaete density of 1,900 individuals m⁻² = 11 gm⁻² for a coral reef in Moorea, French Polynesia. Mean weight per individual is 5.8 mg.

ⁱPauly et al. (1993b) for Caribbean *D. antillarum*.

^jSchwinghamer et al. (1986, Table 1).

^kWeil et al. (1984); mean density of *D. antillarum* for several coral reefs in Venezuela.

^lWilkinson (1987, Table 1) indicated density values for Caribbean reef sponges of 7.37 individuals m⁻² (0.5 km distance from land) and 7.60 individuals m⁻² (5 km distance from land).

^mWith density and biomass values from Table 1 in Wilkinson (1987), a range of mean weight of 86-184 g/individual was calculated for Caribbean reef sponges.

Table 3.10. Database of ECOPATH II key input parameters of nonfish groups for Caribbean coral reef models; Q/B = annual consumption/unit biomass, P/B = production/biomass ratio.

Net primary production (NPP) of overall system: 20,000 (15,000-25,000) g m⁻²year⁻¹WW(see text statement no. 1).
 Export of benthic producers from system: 480 gm⁻²year⁻¹WW(see text statement no. 2).

| ID | Nonfish group | Biomass (gm ⁻² WW) | Source (TSN) | Q/B (year ⁻¹) | Source (TSN) | P/B (year ⁻¹) | Source (TSN) |
|------|------------------------------------------------------------------|-------------------------------|--------------|---------------------------|--------------|---------------------------|--------------|
| D 1 | Detritus, POM, DOM | 2,000 | 3 | - | - | - | - |
| A 1 | Epilithic and endolithic algae | 360 | 4 | - | - | 30.8 | 5 |
| | Reef turf algae | 109 | 6 | - | - | - | - |
| | Benthic fleshy algae | 6,000 | 7 | - | - | - | - |
| | Macroalgae | 415 | 8 | - | - | - | - |
| | Benthic algae | 2,800 | 9 | - | - | - | - |
| | | 55-164 | 10 | - | - | - | - |
| | Spermatophytes | 163 | 11 | - | - | - | - |
| | | | | | | 12.8 | 12 |
| | | | | | | 13.25 | 13 |
| | | | | | | 20.1 | 12 |
| | | | | | | 13.5 | 15 |
| | Benthic autotrophs (excluding symbionts) | 1,500 | 14 | - | - | - | - |
| A 2 | Symbiotic algae (zooxanthellae, filaments) | 730 | 16 | - | - | 9.3 | 17 |
| | | | | | | 10.2 | 18 |
| | | | | | | 16.6 | 19 |
| | Benthic autotrophs | 3,500-4,000 | 20 | - | - | 12.5 | 21 |
| | | | | | | 13 | 22 |
| | | | | | | 70 | 26 |
| A 3 | Phytoplankton | 0.102 | 23 | - | - | - | - |
| | | 11.7-83 | 24 | - | - | - | - |
| | | 73 | 25 | - | - | - | - |
| I 1 | Bacterioplankton | 2-5 | 27 | 1,043 | 28 | 267 | 29 |
| | Benthic bacteria | 0.1 | 31 | 7,300 | 33 | 313 | 30 |
| | | 0.2-1.732 | | | | 155 | 34 |
| | Foraminifera | 1 | 35 | 30 | 36 | 10 | 37 |
| | Decomposers/microfauna | 5.5 | 38 | 2,190 | 39 | 220 | 40 |
| I 2 | Zooplankton | 5-30 | 41 | 50 | 42 | 40 | 26 |
| | | | | 280 | 26 | - | - |
| I 3 | Sponges | 30.5 | 43 | 4.02 | 45 | 1.7 | 46 |
| | | 507-1,049 | 44 | | | | |
| I 4 | Fire corals (Millepora, Hydrozoa) | | | | | | |
| I 5 | Sea fans (Gorgonacea) (Octocorallia, Anthozoa) | 14 | 47 | 3 | +52 | 0.08 | 48 |
| | | | | 15 | +52 | | |
| I 6 | Sea anemones (Actiniaria and Zoanthidea, Hexacorallia, Anthozoa) | 1.3 | 49 | 3 | +52 | | |
| | | | 15 | +52 | | | |
| I 7 | Stony corals (Scleractinia) (Octocorallia, Anthozoa) | 21-57 | 50 | 3 | +52 | | |
| | Corals | 187 | 51 | 15 | +52 | | |
| | | | 51 | 3 | 52 | 0.37 | 53 |
| | | | | 15 | 52 | 1.8 | 53 |
| I 8 | Bryozoans | 9.2 | 54 | | | | |
| I 9 | Sipunculid worms | 4.5 | 55 | | | | |
| | | 7.5-11.5 | 56 | | | | |
| I 10 | Priapuloids | | | | | | |
| I 11 | Chitons | 18 | 57 | 7.2 | 59 | 0.26 | 60 |
| | | 30 | 57 | 16.2 | 59 | 0.42 | 60 |
| | | 61 | 58 | | | | |
| I 12 | Gastropods | 39 | 61 | 4.7 | 62 | 0.1 | 66 |
| | | | | 7.7 | 63 | 2.5 | 67 |
| | | | | 15.7 | 64 | 2.1-5.9 | 68 |
| | | | | 56.4 | 65 | | |
| I 13 | Bivalves | 60-170 | 69 | 9 | 70 | 1.35 | 73 |
| | | | | 9.5 | 71 | 2.23 | 68 |
| | | | | 29.1 | 72 | 3.8 | 74 |
| I 14 | Scaphopods | | | | | | |
| | Molluscs (excluding Cephalopods) | 47-143 | 75 | | | | |
| | | 396 | 76 | | | | |
| I 15 | Squids | | | 16.6 | 77 | 0.9-1.6 | 79 |
| | | | | 20.8 | 78 | | |
| I 16 | Octopuses | 28 | 80 | 6.76 | 81 | 1.09 | 82 |
| | | | | | | 3.06 | 83 |
| I 17 | Polychaetes | 1.9 | 84 | 24.2 | 89 | 2.48 | 68 |
| | | 11 | 85 | 99 | 90 | 5.8 | 89 |
| | | 10 | 86 | | | | |
| | | 20 | 87 | | | | |
| | | 34 | 88 | | | | |
| I 18 | Echiuroids | | | | | | |
| I 19 | Pycnogonids | | | | | | |

continued...

Table 3.10 continued

| ID | Nonfish group | Biomass (gm ⁻² WW) | Source (TSN) | Q/B (year ⁻¹) | Source (TSN) | P/B (year ⁻¹) | Source (TSN) |
|------|--------------------------|----------------------------------|-----------------|------------------------------|-----------------|------------------------------|-----------------|
| I 20 | Barnacles | 35-83 | 91 | | | | |
| I 21 | Stomatopods | | | | | | |
| I 22 | Amphipods | 1.1-3.5 | 92 | 33.4 | 93 | 2.41 | 68 |
| | | | | 125.5 | 94 | | |
| | | | | 164 | 95 | | |
| I 23 | Tanaids | | | 33.4 | 93 | | |
| I 24 | Isopods (fish parasites) | | | 33.4 | 93 | 2.4 | 68 |
| | Small crustaceans | 0.16 | 96 | | | 1.4-3.9 | 68 |
| I 25 | Shrimps | 0.25-0.52 | 97 | 26.9 | 99 | 0.9-1.6 | 100 |
| | | 3.8 | 98 | | | 1.8-2.8 | 101 |
| I 26 | Spiny lobsters | 0.09 | 102 | 7.4 | 103 | 0.35 | 104 |
| | | | | | | 0.45 | 105 |
| | | | | | | 1.03 | 106 |
| I 27 | Scyllarid lobsters | | | | | | |
| I 28 | Hermit crabs | 0.1-0.2 | 107 | | | | |
| I 29 | Crabs | 2.3-4.9 | 108 | 10.6 | 109 | 0.73 | 110 |
| | | | | 17.4 | 109 | 1.61 | 111 |
| | Lobsters and crabs | 3.31 | 26 | 8.2 | 26 | 0.52 | 26 |
| | Crustaceans | 2.5-5.6 | 112 | | | | |
| | | 161 | 113 | | | | |
| I 30 | Hemichordates | | | | | | |
| I 31 | Asteroids | 14-25 | 114 | 3.24 | 115 | 0.49 | 116 |
| I 32 | Ophiuroids | 1-3 | 117 | | | | |
| I 33 | Echinoids | 67 | 118 | 2.81 | 122 | 0.8 | 68 |
| | | 40-148 | 119 | | | 2 | 123 |
| | | 120 | 120 | | | | |
| | | 148-211 | 121 | | | | |
| I 34 | Holothurians | 23-44 | 124 | 3.36 | 126 | 0.29 | 127 |
| | | 59 | 125 | | | | |
| | Echinoderms | 140-200 | 128 | | | | |
| | | 1,053 | 129 | | | | |
| I 35 | Ascidians (Tunicata) | 35 | 130 | 9-49 | 132 | 2.3 | 133 |
| | | 744 | 131 | 24 | 132 | | |
| R 1 | Sea turtles | 0.015 | 26 | 2.8 | 134 | 0.15 | 26 |
| | | | | 3.5 | 26 | | |
| | | | | 11.5 | 135 | | |
| B 1 | Sea birds | 0.015 | 26 | 80 | 26 | 5.4 | 26 |

Symbols:

A = autotrophs.

B = birds.

D = detritus.

I = invertebrates.

R = reptiles.

TSN = text statement number.

* = assumed value.

Table 3.11. Primary productivity of coral reefs and components; GPP = gross primary production, NPP = net primary production, P/R = production/respiration ratio.

| Community component | Geographical location | GPP ^a | NPP ^a | P/R | Source |
|-------------------------------------------|-----------------------------------|-------------------------|-------------------------|----------|-------------------------------------------------------|
| Overall system | Tropics | 840-2,190 ^c | -3.7-62 | 1H | Hatcher (1990, Fig.1) |
| Overall system | Tropics | - | 200-1,000 | | Lewis (1981) |
| Overall system | Tropics | - | 1,200 | >1 | Lewis (1981, Table 1); median of reefs with P/R >1 |
| Overall system | Puerto Rico, El Mario reefs | 4,450 | 350 | 1.1 | Odum et al. (1959) |
| Fore reef | Tropics | 730-2,555 ^c | -365-1,862 | 0.5-5.5 | Hatcher (1990, Fig.1) |
| Reef flat - reef crest | Tropics | 730-2,555 ^c | 110-548 | 1-4 | Hatcher (1990, Fig.1) |
| Reef flat - back reef | Tropics | 949-14,500 ^c | 621-9,855 | 0.7-3.2 | Hatcher (1990, Fig.1) |
| Sand sheet and shallow patch reefs | Tropics | 329-4,709 ^c | -183-1,241 | 0.7-1.4 | Hatcher (1990, Fig.1) |
| Water over algal plains, seagrass meadows | Tropics | 3.65-730 ^c | -475-511 | 0.1-1.4 | Hatcher (1990, Fig.1) |
| Surface pelagic waters | Virgin Islands | - | 20-267 | - | Jacobsen and Browder (1987); mean from Table 51 |
| Lagoon | Joyuda Lagoon, Puerto Rico | 1,260 ^c | 475 | 1.6 | Owen and Tilly (1985) |
| Lagoon | British West Indies, Bimini | 319 | 114 | - | Odum and Hoskin (1958) |
| Corals | Tropics | 281-3,723 ^c | 2.9-14.6 ^b | 0.5-5 | Hatcher (1990, Fig.1) |
| Gorgonacea | Florida | 2,080 | 533 | 3.9 | Kanwisher and Wainwright (1967) |
| Scleractinia | Florida | 2,385 | 769 | 3.1 | Kanwisher and Wainwright (1967) |
| <i>Acropora palmata</i> | Virgin Islands | 485-1,405 ^c | 197-708 | 1.7-2 | Rogers and Salesky (1981) |
| <i>A. palmata</i> | Virgin Islands | 880 | 551-672 | 2-2.5 | Gladfelter et al. (1977) |
| Benthic algae | PRVI area | - | 110 | - | Stanhope (1980) |
| Coralline algae | Tropics | 292-1,022 ^c | 0.22-4.27 ^b | 1-5.4 | Hatcher (1990, Fig.1) |
| Corallinaceae | Curaçao | 890 | 370 | 1.7 | Wanders (1976a) |
| Algal turfs | Tropics | 329-4,417 ^c | 6.2-102.2 ^b | 1.2-6.7 | Hatcher (1990, Fig.1) |
| Algal turfs | Virgin Islands | - | 213.5 ^c | - | Rogers and Salesky (1981) |
| Reef turf algae | PRVI area | - | 274 | - | Carpenter (1985) |
| Macroalgae - reef flat | Tropics | 840-14,381 ^c | 9.13-4.31 ^b | 1.2-6.3 | Hatcher (1990, Fig.1) |
| Macroalgae - lagoon | Tropics | -1,460 ^c | 0.073-14.6 ^b | 1.9-2.8 | Hatcher (1990, Fig.1) |
| Macroscopic algae | Virgin Islands | - | 2,010 | - | Rogers and Salesky (1981) |
| Microalgae | Tropics | 29-1,351 ^c | ?-132.5 ^b | 1.1-10.3 | Hatcher (1990, Fig.1) |
| Fleshy and filamentous algae | Curaçao | 712 | 452 | 2.7 | Wanders (1976a) |
| Fleshy and filamentous algae | Virgin Islands | 10,950 | 6,278 | 2.3 | Connor and Adey (1977) |
| Algal | Virgin Islands | 2,486-4,139 | 1,577-2,321 | 2.4 | Connor and Adey (1977) |
| <i>Sargassum platycarpum</i> | Curaçao | 3,840 | 2,550 | 2.9 | Wanders (1976b) |
| <i>Halimeda</i> | Jamaica | 1,460 | 839 | 2.3 | Hillis-Colinvaux (1974) |
| Seagrass | Tropics | 1,095-5,480 | 1.5-3.2 ^b | 1.5-2.5 | Hatcher (1990, Fig.1) |
| Turtle grassbed | Florida, Long Keys | 3,880 | 1,140 | 1.4 | Odum (1957) |
| Turtle grassbed | Puerto Rico, West La Gata reef | 980 | -310 | 0.8 | Odum et al. (1959) |
| Turtle grassbed | Puerto Rico, Isla Magueyes | 1,350 | -150 | 0.9 | Odum et al. (1959) |
| <i>Thalassia</i> | Barbados | - | 368 | - | Patriquin (1973) |
| <i>Thalassia</i> | Cuba | - | 585 | - | Buesa (1974) |
| <i>Thalassia</i> | Florida | - | 534-712 | - | Odum (1957) |
| <i>Thalassia</i> | Jamaica | - | 704 | - | Greenway (1974) |
| <i>Thalassia</i> | Puerto Rico | - | 800 | - | Burkholder et al. (1959) |

^a(gCm⁻²·year⁻¹).^b(gCg·year⁻¹DW).^cValues adjusted to annual production.

2. From Table 2 in Zieman et al. (1979), total export of primary producers from Virgin Islands seagrass meadows was calculated as follows:

| Item | Unit | Value |
|-------------------------------------|-------------------------------------------|-------------|
| Combined surface and bedload export | (g day ⁻¹ DW) | 162,000.000 |
| Export area | (m ²) | 620,000.000 |
| Daily export | (g m ⁻² day ⁻¹ DW) | 0.261 |
| Annual export | (g m ⁻² year ⁻¹ DW) | 95.370 |
| Export in live weight | (g m ⁻² year ⁻¹ WW) | 477.000 |

Conversion: DW = 17.5% of WW.

Annual export of benthic producers amounted to around 480 g live weight m⁻².

3. Detritus standing stock was calculated by applying equation (2.13). A gross PP of 3,000 gCm⁻²year⁻¹ (mean from overall system values in Table PP) and an approximate euphotic depth of 40 m (personal observation) yielded a detritus standing stock of 195 gCm⁻² which was around 2,000 gm⁻²WW. Conversion: C = 10% of WW for organic material.

4. 21.5 gCm⁻² was the average algal biomass (mean of highest and lowest value) from Fig. 5 in Klumpp and McKinnon (1989) for an epilithic algal community (EAC) on Davis Reef, Great Barrier Reef, Australia. Conversion: C=6% of WW.

5. In Klumpp and McKinnon (1989), the range of gross production of the above (no. 4) EAC was 1.705-2.511 gCm⁻²day⁻¹ for the reef flat and 1.114-1.716 gCm⁻²day⁻¹ for the reef slope. An average annual production of 662 gCm⁻² was calculated (highest + lowest value/2) which resulted in a P/B ratio of 30.8 when divided by an average biomass of 21.5 gCm⁻² (see no. 4).

6. Carpenter (1985) reported a biomass of 19.0 gm⁻²DW for reef turf algae in the PRVI area. Live weight was 109 gm⁻² with a conversion of 17.5%.

7. A biomass of 1,022.1 gm⁻²DW was obtained for benthic fleshy algae on a Caribbean *Porites furcata* reef from Table 2 in Glynn (1973). Conversion: 17.5% of WW.

8. Biomass for macroalgae was derived from a value of 71.27 gm⁻²DW in Table 3 in Dominguez and Alcolado (1990) for macroalgae in the Gulf of Batabanó, Cuba. Conversion: DW = 17.5% of WW.

9. From Tables 8-12 in Odum and Odum (1955), an average biomass for benthic algae (excluding symbiotic algae in corals) of 490 gm⁻²DW was calculated. Conversion: DW = 17.5% of WW.

10. Stanhope (1980) gave a biomass of 4.3-11.7 gm⁻²DW for benthic algae associated with coral reefs in the PRVI area. Converted into live weight, the biomass was 55-164 gm⁻² (DW = 17.5% of WW).

11. A mean standing crop of 142 gm⁻²DW was assessed by Gonzalez-Liboy (1979) for turtle grassbeds (*Thalassia testudineum*) for seven sites off La Parguera, Puerto Rico. DW-WW relationship was assumed to be 17.5%. This live weight was only valid for one subsystem, so the calculated value of 811 g was divided by 5. This procedure was applied in cases where the distribution of an organism was clearly limited to a specific subsystem.

12. Zieman et al. (1979, in Table 1) indicated a daily production of 2.7 gm⁻²DW and a standing crop of 77 gm⁻²DW for a turtle grassbed (*Thalassia testudineum*) in the US Virgin Islands, which resulted in an annual P/B ratio of 12.8. Production and standing crop for the seagrass *Syringodium filiforme* were 0.32 gm⁻²day⁻¹DW and 5.8 gm⁻²DW, respectively, which yielded an annual P/B ratio of 20.1.

13. A mean daily turnover ($= P/B * 100$) of 3.63% was assessed by Gonzalez-Liboy (1979) for the above turtle grassbeds. Annual turnover ($= P/B$) was thus 13.25 ($3.63 * 3.65$).

14. A mean standing crop of $1,513 \text{ gm}^{-2}\text{WW}$ for benthic autotrophs, excluding symbiotic algae (zooxanthellae and filamentous algae), was estimated from biomass values of different components described in nos. 4, 6, 7, 8, 9, 10 and 11.

15. A weighted (by biomass) mean annual P/B ratio of 13.5 for benthic autotrophs was estimated from nos. 5, 12 and 13, and an assumed annual P/B ratio of 12.5 for benthic algae, benthic fleshy algae and macroalgae (see no. 21).

16. From Table 4 in Odum and Odum (1955), an average dry weight for symbiotic algae (zooxanthellae and filaments) of 0.063 gcm^{-2} was obtained. Conversion DW-WW was 17.5%. Since the calculated live weight of $3,600 \text{ gm}^{-2}$ was only valid for the reef zone, and few or no corals grow in the other subsystems included in the model, the value was divided by 5. The resulting biomass of $720 \text{ gm}^{-2}\text{WW}$ was increased to $730 \text{ gm}^{-2}\text{WW}$ to account for zooxanthellae biomass in foraminiferans (see no. 35).

17. A PP for corals of $281\text{-}3,723 \text{ gCm}^{-2}\text{year}^{-1}$ was read off Fig. 1 in Hatcher (1990). A mean production of $33,366 \text{ gm}^{-2}\text{year}^{-1}\text{WW}$ (conversion: $C = 6\%$ of WW) and a biomass of $3,600 \text{ gm}^{-2}\text{WW}$ for symbionts resulted in an annual P/B of 9.3.

18. From Kanwisher and Wainwright (1967), an average gross primary productivity of $2,200 \text{ gCm}^{-2}\text{year}^{-1}$ was obtained for Gorgonacea (sea fans) and Scleractinia (stony corals). Applying a conversion of $C = 6\%$ of WW and a symbiont biomass of $3,600 \text{ gm}^{-2}\text{WW}$, the resulting P/B ratio was 10.2 year^{-1} .

19. Production of zooxanthellae was indicated with $10 \text{ kcalm}^{-2}\text{day}^{-1}$ in Fig. 2 in Sorokin (1987) which is $3,650 \text{ gm}^{-2}\text{year}^{-1}\text{WW}$. When zooxanthellae biomass is around $220 \text{ gm}^{-2}\text{WW}$ (Odum and Odum 1955, Table 4; conversion DW = 17.5% of WW), annual P/B is 16.6.

20. Odum and Odum (1955) estimated a mean standing stock of benthic producers of $703 \text{ gm}^{-2}\text{DW}$ with a low standard deviation, i.e., the biomass seemed to be evenly distributed over the reef regardless of species composition. A conversion of DW = 17.5% WW resulted in a live weight biomass of ca. $3,500\text{-}4,000 \text{ gm}^{-2}$ for benthic autotrophs.

21. Odum and Odum (1955) indicated an annual ratio of 12.5 for gross PP to standing stock of benthic autotrophs for the Eniwetok atoll reef.

22. Weighted (by biomass) mean annual P/B ratio for benthic autotrophs and symbiotic algae was 13.0.

23. Chlorophyll-a concentrations per depth range of pelagic waters in Great Cruz Bay, St. John, Virgin Islands, were given in Table 52 in Jacobsen and Browder (1987). The median was $0.37 \text{ mgChl.-am}^{-2}$. A conversion factor of Chl.-a to phytoplankton carbon of 6% was determined from Table 1 and Fig. 5 in Knoppers and Opitz (1984). Assuming a conversion of 6% for carbon: WW resulted in a biomass of $0.102 \text{ gm}^{-2}\text{WW}$ for phytoplankton.

24. Margalef (1971) gave a phytoplankton biomass range of $0.7\text{-}5.0 \text{ g Cm}^{-2}$ for the northeastern Caribbean. This, transformed into wet weight, yielded a range of $11.7\text{-}83 \text{ g}$ (Carbon = 6% of WW).

25. Jones (1963) indicated a phytoplankton production of $307 \text{ gCm}^{-2} \text{ year}^{-1} = 5,117 \text{ gm}^{-2} \text{ year}^{-1}\text{WW}$ (conversion $C = 6\%$ of WW) for a Florida patch reef. Assuming a P/B ratio of 70 (Polovina 1984b) yielded a biomass of $73 \text{ gm}^{-2}\text{WW}$.

26. All estimates with this source number in Table 3.10 were adopted from Polovina (1984b).

27. Sorokin (1987) reported 0.2-0.5 gm⁻³ as range for bacterioplankton biomass. An assumed mean water depth of 10 m resulted in a value of 2-5 gm⁻² bacterioplankton standing stock.

28. A daily consumption of 10 kcalm⁻² was read off Fig. 2 in Sorokin (1987) for bacterioplankton. A conversion of 1 kcal = 1 g live weight and a biomass of 2-5 gm⁻² yielded a daily consumption of 286% of body weight (% BWday⁻¹) and an annual Q/B of 1,043.

29. Lewis (1981, Table 3b) presented values for production and biomass of bacteria in the water column above various coral reefs. From these values, a mean annual P/B ratio of 267 can be calculated.

30. Production of bacterioplankton was 3 kcalm⁻²day⁻¹ in Fig. 2 in Sorokin (1987). Divided by a biomass of 3.5 gm⁻², this resulted in an annual P/B ratio of 313 (1 kcal = 1 gWW).

31. Odum and Odum (1955) calculated a biomass of 0.1 gm⁻² for bacteria on reef substrates.

32. Sorokin (1987) gave a biomass range of 0.2-1.7 gm⁻² for benthic bacteria as an average for coral reefs.

33. Sorokin (1987), in Fig. 2, gave a ration of 20 kcalm⁻²day⁻¹ = 20 gm⁻²WW for demersal microflora on coral reefs. Based on an average biomass of 1 gm⁻² (0.2-1.7 gm⁻², no. 32), the annual consumption was 7,300 *BW.

34. From Table 3a in Lewis (1981), a mean annual P/B ratio of 155 was calculated for bacteria in coral sediments.

35. A value for organic tissue of foraminiferans was obtained as follows:

a. 31.4 g was the mean foraminiferan biomass of a four-year sampling period in the Gulf of Batabanó, Cuba (based on Table 4 in Gomez et al. 1980).

b. Odum and Odum (1955, Table 12) reported an ash-free DW of 48 gm⁻² for foraminifera and an AFDW:WW conversion of 86.5% for this component of the decomposers on the Eniwetok reef atoll, South Pacific. Thus foraminiferan wet weight was 55.5 gm⁻².

c. From Table 2 in Glynn (1973), a biomass of 135.6 gm⁻²DW was obtained for foraminiferans on a Caribbean *P. furcata* reef. A conversion of 86.5% (Odum and Odum 1955) yielded a live weight of 157 gm⁻².

d. A mean foraminiferan biomass of 81 gm⁻²WW was computed from a, b and c. Odum and Odum (1955) gave a loss on ignition for foraminiferans of 13.5% per individual. Organic tissue of foraminiferans is assumed to approximately represent the difference between dry weight and wet weight, so that the main component of dry biomass consists of the calcareous skeleton. Like coral polyps, foraminiferans, in Caribbean coral reefs, cover their metabolic requirements largely through the activity of symbiotic microalgae (zooxanthellae). These autotroph organisms are assumed to represent 80-90% of the organic tissue of foraminiferans (Prof. Röttger, Department of Microbiology, University Kiel, pers. comm.). Hence, isolated organic tissue of foraminiferans was reduced to 1 gm⁻²WW.

36. Annual P/B ratio of foraminiferans (no. 37) is about ten times the P/B ratio of corals. Their functioning as an ecological unit resembles that of corals: both have a calcareous skeleton and symbiotic zooxanthellae which supply a considerable part of the ration - ca. 40% of photosynthesis C₁₄ is transferred to the animal (Drits et al. 1987). Considering the much smaller size of foraminiferans and therefore a higher metabolism, I assumed their Q/B to be at least ten times that of corals. Hence, annual consumption should not be < 30 *BW.

37. Hallock (1981) reported carbonate turnover rates for foraminiferans of 11-16 times per year for Palau, Western Caroline Islands and of 7-11 times for Hawaii due to slower growth rates and the absence of the family Calcarinidae. Species composition of Caribbean reef foraminiferans was not known; an average annual P/B ratio of 10 was assumed.

38. Sum of foraminifera (1 gm^{-2}), benthic bacteria (0.9 gm^{-2}) and bacterioplankton (3.5 gm^{-2})

39. Consumption, computed via the combined daily ration of $30 \text{ kcalm}^{-2}\text{day}^{-1}$, for bacterioplankton and demersal decomposers (Fig. 2 in Sorokin 1987) was $556\% * 3.65 = 2,028^*$ year⁻²BW based on a wet biomass of 5.4 gm^{-2} and a conversion of $1 \text{ kcal} = 1 \text{ gWW}$.

40. Mean weighted (by biomass) annual P/B ratio for microfauna (bacterioplankton, benthic microflora and foraminifera) was 220.

41. Sorokin (1987) reported a range of $0.5\text{-}3 \text{ gm}^{-3}$ for zooplankton biomass on coral reefs; based on an assumed average water depth of 10 m, the biomass was thus $5\text{-}30 \text{ gm}^{-2}$.

42. Sorokin (1987) in Fig. 2 gave a consumption of $2.5 \text{ kcalm}^{-2}\text{day}^{-1}$ for reef zooplankton. For an average biomass of 18 gm^{-2} , daily consumption was 13.9% BW and annual Q/B was 50.1.

43. Glynn (1973, in Table 2) reported a sponge biomass of $11.8 \text{ gm}^{-2}\text{DW}$ for a *P. furcata* reef in the Caribbean. A conversion of DW = 30% of WW yielded a wet biomass for sponges of 39.3 gm^{-2} . A 22.5% wet mass was subtracted for inorganic skeletal material (D. Barthel, IfM, Kiel, pers. comm.). The organic weight was 30.5 gm^{-2} .

44. Wilkinson (1987, in Table 1) gave a biomass value of $1,353.8 \text{ gm}^{-2}\text{WW}$ (0.5 km distance from land) and 654.2 gm^{-2} (5 km distance from land) for sponge populations around St. Croix, US Virgin Islands. Subtracting 22.5% for inorganic skeletal material yielded a range of organic wet weight of $507\text{-}1,049 \text{ gm}^{-2}$.

45. Wilkinson (1987) reported a consumption of $0.084 \text{ gCm}^{-2}\text{day}^{-1}$ for a Great Barrier Reef sponge biomass of $98.3 \text{ gm}^{-2}\text{WW} = 76.2 \text{ gm}^{-2}$ of organic wet weight. A conversion of carbon to live weight of 10% yielded a daily sponge consumption of 1.1% BW. Thus, sponges consumed 4.02 times their body weight per year. This value might underestimate consumption of Caribbean sponge populations. Sponges in the Pacific show a much higher infestation with autotrophic symbionts (Cyanophyceae) than species in the tropical Atlantic; metabolites of these symbionts are consumed directly by the sponge host, as those of symbiotic zooxanthellae by the coral polyp (see Wilkinson 1987).

46. Weight-at-age data for the Caribbean sponge *Pseudoceratina crassa* were read off Fig. 1 in Wilkinson and Cheshire (1988). The slope of a semilogarithmic plot from these data yielded an annual production rate of 2.97, indicating that an individual would treble its initial biomass in one year. When assuming a low natural mortality of $10\% \text{ year}^{-1}$ and exponential growth, an annual P/B ratio of 1.7 can be obtained as a crude preliminary estimate.

47. A biomass value of $16.52 \text{ gm}^{-2}\text{DW}$ was adopted from Table 12 in Alcolado (1990) for calculating a gorgonian standing stock. Dry weight was converted to live weight (22.5%) and yielded 73.4 gm^{-2} . From this value, 5% was subtracted for zooxanthellae biomass (see Table 4 in Odum and Odum 1955). The resulting animal biomass was divided by 5 to account for its occurrence in only one subsystem.

48. Yoshioka and Yoshioka (1990) reported an average annual survival rate of 92% for gorgonians at La Parguera, Puerto Rico. From this value, an exponential rate of total mortality $Z = M = 0.08 \text{ year}^{-1}$ can be computed. This corresponds to an estimate of P/B (see Allen 1971).

49. Glynn (1973) in Table 2 gave a biomass of $1.1 \text{ gm}^{-2}\text{DW}$ for zoanths on a Caribbean *P. furcata* reef. With a conversion of DW = 17% of WW, the live biomass was 6.5 gm^{-2} . Divided by 5 to account for the subsystems' effect led to a live weight of sea anemones of 1.3 gm^{-2} .

50. A weight range of 60.1-160 gm²DW for stony corals was taken from Fig. 8 in Martínez-Estalella and Alcolado (1990a). This value originated from a coral reef area near the Isla de la Juventud at the southern margin of the Gulf of Batabanó, Cuba. Conversion of DW = 22.5% of WW resulted in a live weight of 267-711 gm². Martínez-Estalella and Alcolado (1990a) did not mention if their coral biomass values were reduced for plant material (symbiotic zooxanthellae and filamentous algae). If not, the stony coral animal live weight decreased to 107-285 gm² (animal tissue ca. 40% of plant tissue, subpolyp zone not included; see Odum and Odum 1955, Table 4). Accounting for the subsystem effect, the final animal biomass for stony corals was 21.4-57 gm²WW.

51. Odum and Odum (1955, in Table 4) gave a biomass of animal tissue in corals of 0.021 gcm²DW = 933.3 gm²WW (DW = 22.5% of WW). Accounting for the subsystems effect, the live weight of corals for the representative area in the models was 933.3/5 = 187 gm².

52. Sorokin (1987, his Fig. 2) gave a consumption value from reef system components (excluding supply by symbiotic algae) of 8.2 kcalm⁻²day⁻¹ for coral polyps and sea anemones, and 1 kcal was assumed to equal 1 gWW. A biomass of 1,000 gm² for stony corals (calcareous skeleton and symbionts excluded) and sea anemones (see nos. 49 and 51) yielded a daily consumption of 0.825% BW and an annual Q/B of 3.0. When consumption was based on an average biomass of 200 gm² (no. 50), the annual consumption increased to 15 * BW.

53. Production of coral polyps and sea anemones was given as 1 kcalm⁻²day⁻¹ in Fig. 2 of Sorokin (1987). This value yielded an annual P/B ratio of 0.365 based on a biomass of 1,000 gm²WW and 1.825 with a biomass of 200 gm²WW (conversion 1kcal = 1 gWW).

54. Dahl (1973) in Table 2 gave a live weight of 2.3 g for 0.25 m² of a coral reef and sand habitat in Puerto Rico.

55. Glynn (1973) in Table 1 gave a sipunculid dry weight of 0.9 gm² for a Caribbean *P. furcata* reef area. Applying a conversion factor for polychaetes (DW = 20% of WW) yielded a live biomass of 4.5gm².

56. Berry (1982), in Table 1, gave for polychaetes a dry biomass range of 1.5-2.3 gm²DW which was a live weight of 7.5-11.5 gm²WW when a conversion of DW = 20% of WW was applied.

57. Horn (1986), in Table 4, gave an average dry biomass of 3.3 gm² and 5.4 gm² (sum of somatic tissue and shell organics) for two populations of *Chiton pelliserpentis* on a New Zealand high- and lowshore habitat. A conversion of DW = 18% of WW (mean of several gastropod species; for chitons, no value was available) yielded an average live biomass of 18.4 gm² (highshore) and 30.2 gm² (lowshore).

58. Based on values in Glynn (1973), an average dry biomass of 55 gm² was computed for chitons from two reef-boring polyplacophores (*Chiton tuberculatus*, Florida reef, 41 gm² and *Acanthopleura granulata*, Puerto Rico reef, 69 gm²). A live biomass of 306 gm² was obtained by an 18% DW:WW conversion for gastropods. This occurred in only one subsystem, so the average live weight was 61.1 gm².

59. In Table 7, Horn (1986) gave an average consumption of 471 (highshore) and 1,521 (lowshore) kJm⁻²year⁻¹ for *Chiton pelliserpentis*. If 18.6 kJ corresponded to 1 gDW of somatic tissue and 23.9 kJ to 1 gDW of shell organics, annual consumption for chitons was: highshore, 7.2 * BW and lowshore, 16.2 * BW (average biomass: highshore, 3.3 gm²DW and 5.4 gm²DW, lowshore; see no. 57). Both values are rather low for an herbivorous animal.

60. Horn (1986) gave an annual mortality rate of 22.9% for highshore and 34.3% for lowshore chitons. These values were transformed into exponential rates of natural mortality M = 0.26 and 0.42 year⁻¹, respectively; M was set equal to P/B (Allen 1971).

61. A dry biomass of 60-80 g 10m⁻² for gastropods from Fig. 19 in Martínez-Estalella and Alcolado (1990b) was converted into live biomass = 38.9 gm⁻² with a conversion of DW = 18% of WW.

62. Huebner and Edwards (1981) gave a daily feeding rate of 1.28% BW for the gastropod *Polinices duplicatus*. The annual rate was $1.28 * 3.65 = 4.7 * BW$.

63. Riddle et al. (1990) in Table 5 gave a consumption for infaunal gastropods (>2.0 mm) from Davies Reef Lagoon, Great Barrier Reef, Australia, of 277 kJm⁻²year⁻¹. Biomass was indicated with 0.748 gCm⁻². Q/B was then 7.7 year⁻¹ (conversion: 1 gC = 48 kJ).

64. Deslous-Paoli et al. (1985) gave an energy budget for the gastropod *Crepidula fornicata* in the basin of Marennes-Oleron, France: average biomass for an individual representative of the population was equivalent to 1.87 kJ and the annual consumption was 20.08 kJ per individual. Q/B was thus 10.74 year⁻¹. Adjusted for temperature ($V = 1.465$ when $T' = 15^{\circ}\text{C}$), the consumption increased to $Q/B = 15.7 \text{ year}^{-1}$.

65. From Table 3 in Carefoot (1970), gross conversion efficiencies (K_1) for two seahares (*Aplysia dactylomela* and *A. juliana*), fed on various species of algae, were computed. A mean value of $b = 0.0204$ (see Pauly 1986) and an annual consumption of $56.4 * BW$ was computed using growth parameters for the seahare *Dolabella auricularia* from Pauly and Calumpong (1984): $W_{\infty} = 490 \text{ g}$, $K = 0.9 \text{ year}^{-1}$, $t_0 = -0.23 \text{ year}^{-1}$ and $b = 3$. Natural mortality $M = 1.6$ was estimated through equation (2.15). The computations were executed with the MAXIMS software.

66. Berg and Olsen (1989) gave an average natural mortality rate M of 0.1 year⁻¹ for the abundant queen conch (*Strombus gigas*) in the PRVI area; this was set equal P/B.

67. Riddle et al. (1990, in Table 5) gave a P/B ratio for gastropods (>2 mm, mean weight = 50.5 mg/l) of 2.5 year⁻¹.

68. All estimates of P/B ratio with this statement number in Table 3.10 were obtained from Table 1 in Schwinghamer et al. (1986). Each P/B ratio in Table 3.10 represented the average of all estimates listed for a taxon in Schwinghamer et al. (1986); their P/B ratios were assembled from temperate and boreal areas.

69. A dry biomass of 60.1-170 g 10m⁻² for bivalves from Fig. 20 in Martínez-Estalella and Alcolado (1990b) was converted to a live biomass of 60-170 gm⁻² with a conversion of DW = 10% of WW.

70. Salzwedel (1980) gave an energy budget for an "average" *Tellina fabula* population in the German Bight, North Sea, of mean biomass = 60 kJm⁻² and consumption = 290 kJm⁻²year⁻¹, at a temperature of about 10°C. Annual Q/B for *T. fabula* was thus 4.8. This value was adjusted for temperature to a consumption of 9.5 year⁻¹*BW with a factor of 1.88. ($T = 28^{\circ}\text{C}$, $T' = 10^{\circ}\text{C}$).

71. Riddle et al. (1990, in Table 5) gave a consumption for infaunal bivalves (>2.0 mm) from Davies Reef Lagoon, Great Barrier Reef, Australia, of 657 kJm⁻²year⁻¹. Biomass was indicated with 1.443 gCm⁻². Annual Q/B was then 9.5 (conversion: 1 gC = 48 kJ).

72. The daily rations for three species of bivalves (*Mytilus edulis*, *Modiolus modiolus* and *Arctica islandica*) were calculated as follows: filtration rates were determined from equations in Moehlenberg and Riisgard (1979) and multiplied by an average organic content of 0.16 mg l⁻¹ of seawater. The parameter b and Q/B values were computed with MAXIMS (Jarre et al. 1990). The growth constant K was obtained from Appendix 4 in Vakily (1992). W_{max} in the filtration experiments of Moehlenberg and Riisgard were assumed to be 86% of W_{∞} of the three species. M was assumed to equal K , as is commonly done in bivalves. The average annual Q/B of the three species was adjusted for temperature. Table 3.12a/b shows the parameters used for the computation of annual Q/B for bivalves.

Table 3.12a. Parameters used to estimate annual Q/B of some bivalves.

| Parameter/species | <i>Mytilus edulis</i> | | <i>Modiolus modiolus</i> | | <i>Arctica islandica</i> | |
|----------------------------------------|-----------------------|--------|--------------------------|--------|--------------------------|--------|
| Size (gDW) | 0.011 | 1.361 | 0.058 | 1.555 | 0.011 | 1.310 |
| Volume filtered (l day ⁻¹) | 9.1 | 219.1 | 17.0 | 200.5 | 8.1 | 157.5 |
| Daily ration (gDWday ⁻¹) | 0.0015 | 0.0351 | 0.0027 | 0.0321 | 0.0013 | 0.0252 |
| Daily ration (%) | 13.3 | 2.6 | 4.7 | 2.1 | 11.8 | 1.9 |
| W_{∞} (gWW) | | 15.8 | | 18.1 | | 15.2 |
| $K = M$ (year ⁻¹) | | 0.36 | | 0.08 | | 0.02 |
| b | | 3 | | 3 | | 3 |
| β | | 0.02 | | 0.009 | | 0.0012 |

Table 3.12b. Consumption of some bivalves.

| Species | Q/B (% BWD) | Q/B (year ⁻¹) |
|--------------------------|----------------|------------------------------|
| <i>Mytilus edulis</i> | 5.8 | 21.0 |
| <i>Modiolus modiolus</i> | 2.8 | 10.4 |
| <i>Arctica islandica</i> | 5.3 | 19.3 |
| Mean | 4.6 | 16.9 |
| Adjusted ($V = 1.724$) | 7.9 | 29.1 |

73. In Salzwedel (1980), biomass for *T. fabula* in the German Bight was 60 kJm⁻² and production was 81 kJm⁻²year⁻¹. The resulting P/B ratio was 1.35 year⁻¹.

74. Berry (1982, in Table 1) gave a biomass equivalent of 6,702.5 kJm⁻² for organic tissue of a *Perna perna* population on the Oceanographic Research Institute (ORI) Reef, Natal Province, South Africa. Production was 25,458 kJm⁻²year⁻¹. Annual P/B was thus 3.8.

75. A dry biomass of 66-200 g 10m² for molluscs from Fig. 18 in Martínez-Estalella and Alcolado (1990b) was converted to a live biomass of 47-143 gm⁻². A mean conversion for gastropods and bivalves of DW = 14% of WW was applied.

76. Glynn (1973) in Table 2 gave a dry biomass of 59.4 gm⁻² for molluscs on a Caribbean *P. furcata* reef. This corresponded to a live biomass of 424.3 gm⁻² when DW = 14% of WW.

77. Pauly et al. (1993b) estimated an average daily consumption of 4.56% BW for squids as an ecological group. Annual Q/B was thus 16.6.

78. Abolmasova (1985) reported a daily consumption of 13.8-5.7% of body weight from feeding experiments with captive squid *Sthenoteuthis oualaniensis* of 20-70 g body weight. Experimental temperature was 29-30°C. The value of 5.7% was assumed to pertain to a 70 g animal and used to calculate an annual consumption of 20.8 * BW for tropical squids.

79. From Table 10.6 in Longhurst and Pauly (1987) and Table 8 in Pauly et al. (1993b), various annual rates of natural mortality M for squids were obtained:

| Species | M(year ⁻¹) | Source |
|-----------------------------------------------------------------|------------------------|----------------------------|
| <i>Loligo duvauceli</i> | 1.50 | Longhurst and Pauly (1987) |
| <i>Sepioteuthis lessoniana</i> | 1.10 | Longhurst and Pauly (1987) |
| <i>Loligo duvauceli</i> and <i>Loligo opalescens</i> | 1.58 | Pauly et al. (1993b) |
| <i>Sepioteuthis lessoniana</i> and <i>Illex illecebrosus</i> | 0.85 | Pauly et al. (1993b) |

M was assumed to equal P/B, therefore, a range of 0.85-1.58 year⁻¹ was accepted for squids.

80. From Beets and Lewand (1986), an octopus density for a Virgin Island reef area of 0.01 individuals m⁻² (mean of Tables 1-4) was obtained. From growth estimates in Guerra (1979) and from equation (2.12), a medium-sized individual of *Octopus vulgaris* was calculated to weigh 2,800 g. The resulting live biomass was 28 gm⁻²WW.

81. Two species of octopus (*O. vulgaris* and *O. briareus*) were identified as prey items of fish diets on West Indian reefs (see Appendix Table 8.5.2). For *O. vulgaris*, annual population consumption was computed as follows: From Table 3 in Smale and Buchan (1981), estimates for gross conversion efficiency (K_1) and b were obtained for females and males separately (see Table 3.13a). With MAXIMS (Jarre et al. 1990), annual Q/B was then computed using growth parameters from Guerra (1979) (Table 3.13b). Age at size 0 (t_0) and natural mortality M were obtained from equations (2.11) and (2.16), respectively. Annual Q/B was then adjusted for temperature (Table 3.13c).

82. From growth parameters in Guerra (1979), an annual natural mortality rate M of 1.09 was calculated for *O. vulgaris* ($W_\infty = 10,850$ g, $K = 0.72$ year⁻¹, $T = 28^\circ\text{C}$). M was assumed to equal P/B.

83. Buchan and Smale (1981) gave an annual P/B ratio of 3.33 for female and 2.78 for male *O. vulgaris* on the ORI Reef, Natal Province, South Africa. The mean was 3.06 year⁻¹.

84. Lewis (1981) calculated an average biomass of 0.19 gCm⁻² (= 1.9 gm⁻²WW) for polychaetes in coral clumps on reefs in the Bahamas and off South Florida from data in Vittor and Johnson (1977).

85. Naim and Amoureux (1982) gave a maximum polychaete density of 1,900 individuals = 11 gm⁻² for a coral reef in Moorea, French Polynesia.

Table 3.13a. Conversion efficiencies of *O. vulgaris* (Smale and Buchan, 1981 modified).

| No. ind. | Days | Diet | Initial weight (g) | Final weight (g) | Mean weight (g) | Change in weight (g) | Food intake (g) | K_1 | β |
|--------------------|------|------|--------------------|------------------|-----------------|----------------------|-----------------|--------|---------|
| Females | | | | | | | | | |
| 1 | 84 | R+M | 74 | 1,013 | 544 | 939 | 3,481 | 0.2698 | 0.1051 |
| 2 | 77 | M | 77 | 744 | 411 | 667 | 2,644 | 0.2523 | 0.0888 |
| 3 | 97 | M | 80 | 1,292 | 686 | 1,212 | 6,236 | 0.1944 | 0.0783 |
| 4 | 70 | R+M | 146 | 1,129 | 638 | 983 | 2,810 | 0.3498 | 0.1519 |
| 5 | 194 | R+M | 197 | 4,353 | 2,275 | 4,156 | 8,741 | 0.4755 | 0.4131 |
| 6 | 138 | R+M | 238 | 4,332 | 2,285 | 4,094 | 8,398 | 0.4875 | 0.4291 |
| 7 | 81 | R+M | 247 | 1,646 | 947 | 1,399 | 3,939 | 0.3552 | 0.1799 |
| 8 | 53 | R+M | 282 | 1,728 | 1,005 | 1,446 | 2,490 | 0.5807 | 0.3653 |
| 9 | 55 | R+M | 350 | 866 | 608 | 516 | 1,493 | 0.3456 | 0.1474 |
| 10 | 58 | R+M | 351 | 1,251 | 801 | 900 | 2,377 | 0.3786 | 0.1826 |
| 11 | 25 | M | 497 | 881 | 689 | 384 | 1,718 | 0.2235 | 0.0918 |
| 12 | 196 | R+M | 545 | 4,542 | 2,544 | 3,997 | 9,903 | 0.4036 | 0.3563 |
| 13 | 31 | M | 580 | 1,113 | 847 | 533 | 1,910 | 0.2791 | 0.1283 |
| 14 | 98 | R+M | 952 | 3,111 | 2,032 | 2,159 | 5,620 | 0.3842 | 0.2894 |
| Mean females | | | | | | | | | 0.2148 |
| Males | | | | | | | | | |
| 1 | 83 | M | 13 | 277 | 145 | 214 | 908 | 0.2357 | 0.0623 |
| 2 | 278 | R+M | 284 | 5,685 | 2,985 | 5,401 | 9,062 | 0.5960 | 0.7023 |
| 3 | 91 | R+M | 291 | 1,815 | 1,053 | 1,524 | 2,764 | 0.5514 | 0.3437 |
| 4 | 94 | R+M | 320 | 1,709 | 1,015 | 1,389 | 3,435 | 0.4044 | 0.2187 |
| 5 | 158 | R+M | 393 | 2,259 | 1,326 | 1,866 | 5,988 | 0.3116 | 0.1776 |
| 6 | 127 | R+M | 486 | 1,480 | 983 | 994 | 4,149 | 0.2396 | 0.1141 |
| 7 | 139 | M | 528 | 1,768 | 1,148 | 1,240 | 5,413 | 0.2291 | 0.1158 |
| 8 | 36 | M | 540 | 1,142 | 841 | 602 | 1,715 | 0.3510 | 0.1691 |
| 9 | 112 | M | 542 | 1,100 | 821 | 558 | 3,207 | 0.1740 | 0.0741 |
| 10 | 106 | M | 635 | 1,492 | 1,064 | 857 | 3,562 | 0.2406 | 0.1185 |
| 11 | 112 | M | 889 | 1,680 | 1,285 | 791 | 4,455 | 0.1776 | 0.0917 |
| Mean males | | | | | | | | | 0.1989 |
| Mean of all values | | | | | | | | | 0.2078 |

M = mussels, R = rock lobster.

Table 3.13b. Parameters used to estimate consumption of *O. vulgaris*.

| Parameter (unit) | | Source |
|-----------------------------|--------|-----------------------------------------------|
| L_{∞} (cm) | 30 | Guerra (1979) |
| W_{∞} (g) | 10,850 | Guerra (1979) |
| W_{max} (g) | 4,000 | Largest individual of Smale and Buchan (1981) |
| W_r (g) | 1 | Set value |
| K (year ⁻¹) | 0.72 | Guerra (1979) |
| t_0 (year ⁻¹) | -0.22 | Equation (2.11) |
| b | 2.987 | Guerra (1979) |
| β | 0.2078 | Table 3.13a |
| M (year ⁻¹) | 1.09 | Equation (2.16) |
| T (°C) | 28 | DHI (1967) |
| T' (°C) | 23 | Smale and Buchan (1981) |
| V | 1.128 | Pauly et al. (1993b) |

Table 3.13c. Consumption and maintenance ration of *O. vulgaris* at 28°C.

| Temperature | Q/B (% BWD) | R_m (% BWD) | Q/B (year ⁻¹) | R_m (*BWyear ⁻¹) |
|-------------|-------------|---------------|---------------------------|--------------------------------|
| 23°C | 1.6415 | 0.9528 | 5.9916 | 3.4778 |
| 28°C | 1.85 | 1.08 | 6.76 | 3.92 |

86. Ibarzábal (1990, in Fig. 15) gave a biomass value of > 2 gm⁻²DW for polychaetes in an area near the Isla de la Juventud, Cuba. Applying a conversion of DW = 20% of WW, the minimum live weight was 10 gm⁻².

87. Kohn and Lloyd (1973) estimated biomass of polychaetes on coral reefs in the Eastern Indian Ocean of 2-6 gm⁻²DW (ca. 20 gm⁻²WW).

88. Glynn (1973, in Table 2) gave a dry biomass of 6.7 gm⁻² for polychaetes in a Caribbean *P. furcata* reef area. Allowing for DW = 20% of WW yielded a live weight of 33.5 gm⁻².

89. Riddle et al. (1990, in Table 5) reported the following consumption rates and P/B ratios for polychaetes in Davies Reef Lagoon, Great Barrier Reef, Australia:

| Feeding type | Size | Biomass (mgCm ⁻²) | Consumption (kJm ⁻² year ⁻¹) | P/B (year ⁻¹) | Q/B ^a (year ⁻¹) |
|--------------|---------|-------------------------------|-----------------------------------------------------|---------------------------|----------------------------------------|
| Macrophagous | > 2 mm | 124 | 73 | 2.6 | 12.3 |
| | 0.5-2mm | 71 | 126 | 10.3 | 40.0 |
| Microphagous | > 2 mm | 217 | 157 | 3.3 | 15.1 |
| | 0.5-2mm | 224 | 373 | 8.5 | 34.7 |

^aConsumption/48/biomass (1gC = 48 kJ).

A weighted (by biomass) mean annual Q/B of 24.2 with a P/B ratio of 5.8 could thus be calculated for polychaetes as an ecological group.

90. From Table 1 in Pamatmat and Findlay (1983), a mean daily ration for the polychaete *Capitella capitata* of 27% BW was determined. Annual Q/B for animals of 45-167 mgWW was 99.

91. Berry (1982, in Table 1) gave a dry biomass range for two species of Cirripedia on the ORI Reef, Natal Province, South Africa, of 8.74 to 20.46 gm⁻² (calcareous shells excluded). Allowing for a conversion DW = 25% of WW (mean for various small crustaceans) resulted in a live biomass range of 35.4-82.8 gm⁻².

92. Berry (1982, in Table 1) gave a dry biomass range for amphipods (several species) of 0.28-0.92 gm⁻² which corresponded to a live biomass of 1.1-3.5 gm⁻² (conversion 1:4).

93. Riddle et al. (1990, in Table 5) gave a biomass of 0.086 gCm⁻² and a consumption of 69 kJm⁻²year⁻¹ for crustaceans in Davies Reef Lagoon, Great Barrier Reef, Australia, of 5.32 mg (>2 mm length) mean body weight. Annual Q/B was thus 16.7. Consumption for crustaceans of a mean weight of 0.072 mg (0.5-2 mm length) was 46.3*BW year⁻¹ (249 kJm⁻²year⁻¹/48/0.112 gCm⁻²). Weighted (by biomass) mean annual Q/B of small crustaceans was then 33.4.

94. Applying an equation by Cammen (1980):

$$C = 0.381 * W^{0.742}$$

...3.3)

where C = daily ingestion rate in mgday⁻¹DW, W = body weight in mgDW, for deposit feeders and detritivores yielded a daily ingestion rate of 0.515 mg = 34.3% BW for amphipods of mean individual dry body weight of 1.5 mg (Berry 1982). Annual Q/B was 125.25.

95. Hargrave (1985) gave an average daily consumption rate of 30-60% BW for lysianassid amphipods (size not given) of the Nares Abyssal Plain (5,830 m depth) in the northwest Atlantic Ocean, where temperature (not given) can be presumed to be very low. A mean consumption rate of 45% BW yielded an annual Q/B of 164. This already high estimate of Q/B was not adjusted to the higher temperature of waters around PRVI.

96. Odum and Odum (1955, p. 309) gave a biomass value for "microcrustacea" in algal encrusting mats of 0.022 gm⁻²DW which is 0.17 gm⁻²WW (wet volume assumed to have a specific density of 1). The authors did not comment on species composition and size of this ecological group.

97. In Fig. 32 in Martínez-Iglesias and Alcolado (1990), a dry biomass range of 66-140 g 100m⁻² was indicated for decapod crustaceans in the Gulf of Batabanó, Cuba, north of the Isla de la Juventud. In Fig. 34 (same source), it was shown that penaeidean and caridean shrimps comprise an average 10% of the decapod biomass of the samples taken during the study. Allowing for DW = 26.7% of WW yielded a shrimp wet biomass of 0.25-0.52 gm⁻².

98. Odum and Odum (1955, in Table 8, quadrat A), gave a dry biomass for predominantly herbivorous gammarids and other small crustacean of 4.7 gm⁻². A mean live biomass of 18.8 gm⁻² was computed with DW = 25% of WW. This ecological group was not listed for quadrats B,C,D and E by Odum and Odum (1955, Tables 9-12); hence, the average reef area contained 18.8/5 = 3.76 gm⁻²WW.

99. Pauly et al. (1993b) computed a daily consumption of 7.93% BW for "shrimps" as an ecological group; annual Q/B was thus 26.9.

100. Pauly et al. (1993b) estimated a natural mortality rate for *Metapenaeus monoceros* of M = 1.6 year⁻¹ and for *Penaeus monodon* of M = 0.9 year⁻¹. M was assumed to equal P/B. Hence, an annual range of P/B for shrimps of 0.9-1.6 was obtained.

101. Garcia (1985), in Table 1, gave estimates of annual natural mortality M (= P/B) for fully exploited penaeid shrimp stocks in the Gulf of Mexico of 1.8-2.8 year⁻¹. Penaeids comprised ca. 25% of occurrence of "shrimps" in West Indian fish stomachs analyzed by Randall (1967). M was set equal P/B.

102. Munro (1983) reported a biomass of 63-119 kgkm⁻² = 0.063-0.119 gm⁻² for *Panulirus argus* in Jamaican waters.

103. Food consumption for spiny lobsters was computed as follows: Table 3.14a shows food conversions and corresponding weights for *Panulirus homarus*, read off Fig. 17 in Smale (1978) for an experimental temperature of 28°C. Dry weight was converted into wet weight with

DW = 18% of WW. These data were used to estimate b through equation (2.23), applying a mean W_{∞} of 833 gWW from females and males (see Table 3.14b). From growth parameters in Table 3.14b, the annual consumption (Table 3.14c) of spiny lobster was computed with MAXIMS (Jarre et al. 1990).

104. Malkov (1978) gave a natural mortality estimate M of 0.35 year^{-1} for *P. argus* in Cuban waters.

105. Buesa (1969) reported an annual natural mortality of 56-78% for *P. argus* in Cuban waters; from this, a mean exponential rate of $M = 0.45 \text{ year}^{-1}$ was computed; M was assumed to equal P/B .

Table 3.14a. Food conversions of *P. homarus* (Smale 1978, Fig. 17, 28°C).

| DW(g) | $K_1 \cdot 100$ | WW(g) | $\text{Log}_{10} \text{WW}$ | K_1 | $-\text{Log}_{10}(1-K_1)$ |
|-------|-----------------|--------|-----------------------------|-------|---------------------------|
| 1.5 | 31.3 | 8.33 | 0.920818 | 0.313 | 0.163043 |
| 3.8 | 19.8 | 21.11 | 1.324511 | 0.198 | 0.095825 |
| 5.7 | 14.0 | 31.67 | 1.500602 | 0.140 | 0.065501 |
| 7.2 | 12.2 | 40.00 | 1.602059 | 0.122 | 0.056500 |
| 11.2 | 17.2 | 62.22 | 1.793945 | 0.172 | 0.081969 |
| 13.1 | 15.4 | 72.78 | 1.861998 | 0.154 | 0.072629 |
| 14.9 | 10.9 | 82.78 | 1.917913 | 0.109 | 0.050122 |
| 16.8 | 9.2 | 93.33 | 1.970036 | 0.092 | 0.041914 |
| 18.3 | 9.9 | 101.67 | 2.007178 | 0.099 | 0.045275 |
| 26.2 | 16.7 | 145.56 | 2.163028 | 0.167 | 0.079354 |
| 27.9 | 9.0 | 155.00 | 2.190331 | 0.090 | 0.040958 |
| 31.2 | 8.7 | 173.33 | 2.238882 | 0.087 | 0.039529 |
| 32.8 | 1.8 | 182.22 | 2.260601 | 0.018 | 0.007888 |
| 33.9 | 5.0 | 188.33 | 2.274927 | 0.050 | 0.022276 |
| 35.1 | 5.9 | 195.00 | 2.290034 | 0.059 | 0.026410 |
| 36.4 | 4.0 | 202.22 | 2.305828 | 0.040 | 0.017728 |
| 37.5 | 2.0 | 208.33 | 2.318758 | 0.020 | 0.008773 |
| Mean | | | 1.937732 | | 0.053865 |

Table 3.14b. Parameters used to estimate consumption of *P. homarus*.

| Parameter (unit) | Female | Male | Source |
|----------------------------|---------|---------|---------------------------------------------|
| Cl_{∞} (mm) | 94.2 | 120.0 | Smale (1978) |
| Cl_{\max} (mm) | 76.0 | 81.0 | Smale (1978, Table 3) |
| a | 0.00069 | 0.00126 | Munro (1983, for <i>P. guttatus</i>) |
| b | 3.01 | 2.85 | Munro (1983, for <i>P. guttatus</i>) |
| W_{∞} (g) | 604.0 | 1,062.0 | - |
| W_{\max} (g) | 316.0 | 346.0 | - |
| W_r (g) | 1.0 | 1.0 | Set value |
| K (year^{-1}) | 0.337 | 0.177 | Smale (1978, K (day^{-1})*365) |
| t_0 | 0.15 | -0.01 | Equation (2.11) |
| M (year^{-1}) | 0.7 | 0.7 | Mean of nos. 95, 96 and 97 |
| β | 0.05484 | 0.05484 | See text |

Table 3.14c. Consumption and maintenance ration of *P. homarus*.

| Sex | Q/B (% BWD) | R_m (% BWD) | Q/B (year^{-1}) | R_m (*BW year^{-1}) |
|--------|----------------|------------------|-------------------------------|------------------------------------|
| Female | 2.4138 | 1.6773 | 8.8105 | 6.1222 |
| Male | 1.6173 | 0.9305 | 5.9031 | 3.3963 |
| F + M | 2.02 | 1.30 | 7.36 | 4.76 |

106. Munro (1983, in Table 16.2) estimated M for female and male *P. argus* from unexploited parts of Pedro Bank, Jamaica, to be 0.81 and 1.25 year⁻¹, respectively. Hence, mean P/B for *P. argus* was 1.03 year⁻¹.

107. A biomass range of 66-149 g 100m⁻²DW for decapod crustaceans was read off Fig. 32 in Martínez-Iglesias and Alcolado (1990). Fig. 34 (same source) indicated a percentage of anomuran biomass of 3.2. Allowing for a DW/WW conversion of 1:4 yielded a live biomass for hermit crabs of 0.09-0.18 gm⁻².

108. Brachyuran biomass in Martínez-Iglesias and Alcolado (1990) was 87.2% of decapod biomass (Fig. 34). A dry biomass range of 66-140 g100m⁻² (Fig. 32) and a conversion DW/WW of 1:4 resulted in a live biomass for crabs of 2.3-4.9 gm⁻².

109. Consumption for the majid crab *Mithrax spinosissimus* was estimated as follows: From Tables 2 and 4 in Wilber and Wilber (1989), length increments ($\delta l/\delta t$) in relation to CL of *M. spinosissimus* were prepared for a "forced" Gulland and Holt plot (Pauly 1984). An asymptotic carapace width (CW) of 150 mm (Munro 1983: for *M. spinosissimus* around Jamaica) was used as "external" $CL(\infty)$. The slope of the plot yielded an estimate of $K = 0.00087 \text{ day}^{-1}$, i.e., $K = 0.32 \text{ year}^{-1}$. Estimates of daily ration for *M. spinosissimus* were read off Fig. 1 in Winfree and Weinstein (1989). Two different kinds of diets were provided which yielded two differing values for β and hence of consumption. Table 3.15a shows the data used for the estimation of Q/B of *M. spinosissimus*. Table 3.15b presents the population consumption computed using MAXIMS (Jarre et al. 1990).

110. Based on $W_\infty = 1,326 \text{ g}$ for female and 2,820 g for male *M. spinosissimus*, a value of $K = 0.32 \text{ year}^{-1}$ (see no. 109) and a mean environmental temperature of 28°C, values of M of 0.75 for females and 0.71 for males were computed from equation (2.16). Mean P/B for *M. spinosissimus* was thus 0.73 year⁻¹.

111. Erhardt and Restrepo (1989) reported a natural mortality rate M of 1.60 year⁻¹ for the Florida stone crab *Menippe mercenaria* (Majidae). Majid crabs comprised ca. 30% of occurrence of crab species in West Indian fish stomachs analyzed by Randall (1967).

112. A biomass range of 66-140 g 100m⁻²DW for decapod crustaceans in the Gulf of Batabanó, Cuba, was obtained from Fig. 32 in Martínez-Iglesias and Alcolado (1990). Allowing for a DW/WW conversion of 1:4 yielded a live biomass of 2.6-5.6 gm⁻².

113. Glynn (1973), in Table 2, gave a dry biomass for crustaceans on a Caribbean *P. furcata* reef of 40.3 gm⁻². When an average conversion of 25% is applied, live biomass was 161.2 gm⁻².

114. A biomass range of 41-71 g 10m⁻²DW for asteroids in the Gulf of Batabanó, Cuba, was obtained from Fig. 38A in Corvea et al. (1990). Live weight of starfish was 14.1-24.5 gm⁻² when DW is 29% of WW.

Table 3.15a. Parameters used to estimate consumption of *M. spinosissimus*.

| Parameter (unit) | Female | Male | Source |
|-----------------------------|---------|---------|----------------------|
| CW_{\max} (mm) | 154 | 175 | Munro (1983) |
| a | 0.00028 | 0.00018 | Munro (1983) |
| b | 3.0515 | 3.1654 | Munro (1983) |
| W_r (g) | 651 | 727 | Munro (1983) |
| W_{\max} (g) | 1,140 | 2,425 | Munro (1983) |
| W_∞ (g) | 1,326 | 2,820 | Equation (2.6) |
| K (year ⁻¹) | 0.32 | 0.32 | See text |
| t_0 (year ⁻¹) | 0.32 | 0.31 | Equation (2.11) |
| M (year ⁻¹) | 0.75 | 0.71 | Equation (2.16) |
| β_1 | 0.0179 | 0.0223 | Computed with MAXIMS |
| β_2 | 0.0285 | 0.0385 | Computed with MAXIMS |

1 diet: wild reef algae; 2 diet: fish - *Gracilaria* sp. - *Ulva* sp.

Table 3.15b. Consumption of a Caribbean population of *M. spinosissimus*.

| Sex | | Q/B (% BWD) | Q/B (year ⁻¹) |
|--------|---|----------------|------------------------------|
| Female | 1 | 5.2372 | 19.1157 |
| Male | 1 | 4.2936 | 15.6715 |
| Female | 2 | 3.2977 | 12.0366 |
| Male | 2 | 2.5036 | 9.1318 |
| F + M | 1 | 4.765 | 17.394 |
| F + M | 2 | 2.901 | 10.587 |

1 and 2: see above.

115. From data in Wurzian (1984) on *Astropecten aranciatus* in the Northern Adriatic Sea, a rough estimate of food consumption for starfish was computed as follows: density of *A. aranciatus* was 3.67 individuals 100 m⁻²; this was 0.037 individuals m⁻². Ingestion was 9.5 kJm⁻²year⁻¹. A mean radius (R) of 12.3 cm was computed from R-frequency data in Fig. 1 (same source) and converted to a live weight of 130 g applying the equation in Fig. 2:

$$WW = 0.1609 * R^{2.8293}$$

...3.4)

where WW is wet weight in g and R is the radius of the starfish. With a density of 0.037 individuals m⁻² and an individual weight of 130 g, an average live biomass of 4.76 gm⁻² was computed. Ingestion was converted into wet weight with 1 kJ = 1.2 gWW (conversion for sea urchins) to 11.37 gm⁻²year⁻¹. Hence, annual food consumption for an average individual of *A. aranciatus* was 2.39 *BW. Adjusted for temperature, annual Q/B increased to 3.24 (V = 1.36; T' = 17°C: mean annual temperature for the Northern Adriatic was determined from charts in KNMI 1957).

116. Production of *A. aranciatus* in the Northern Adriatic was indicated with 1.958 kJm⁻²year⁻¹ (Wurzian 1984). Converted to live weight, production was 2.35 gm⁻²year⁻¹. Based on an average biomass of 4.76 gm⁻²WW (see no. 115), the annual P/B ratio was 0.49.

117. A biomass range of 2-6 g10m⁻²DW for ophiuroids in the Gulf of Batabanó, Cuba, was obtained from Fig. 41A in Corvea et al. (1990). Live weight of ophiuroids was 0.7-2.1 g when DW is 29% of WW.

118. McClanahan (1988), in Table 1, gave density and biomass values for three species of sea urchins on a Kenyan coral reef. Mean biomass was 67.4 gm⁻²WW.

119. A biomass range of 127-472 g10m⁻²DW for echinoids in the Gulf of Batabanó, Cuba, was obtained from Fig. 39A in Corvea et al. (1990). Live weight of sea urchins was thus 39.7-147.5 gm⁻² based on a conversion DW/WW of 32%.

120. From Table 1 in Bak et al. (1984), an average density of 2.1 individuals m² was computed for *Diadema antillarum* in Curaçao, Netherlands Antilles. Pauly et al. (1993b) computed an average weight per individual of the same species of 57 gWW. Based on these values, the biomass was 119.7 gm⁻². *D. antillarum* is the most common sea urchin in PRVI reef areas.

121. Weil et al. (1984) reported a mean *D. antillarum* density of 2.6-3.7 individuals m² for several coral reefs in Venezuela. Based on an average individual weight of 57 g (see no. 120), the mean biomass range was 148.2-210.9 gm⁻²WW.

122. Pauly et al. (1993b) estimated food consumptions for various species of sea urchins. When abundance of the respective genera in stomach analyses of reef fishes was considered (Randall 1967, see Appendix 8.5.2), the weighted mean consumption for sea urchins as an ecological group is 0.77% BWD or 2.81 * BWyear⁻¹.

123. Hawkins (1985) gave a P/B estimate of 2.0 year⁻¹ for the sea urchin *D. antillarum* in Barbados.

124. A biomass range of 21-40 g 10m⁻²DW for holothurians in the Gulf of Batabanó, Cuba, was obtained from Fig. 40A in Corvea et al. (1990). Live weight of sea cucumbers was 23.3-44.4 gm⁻² when DW is 9% of WW.

125. Berry (1982), in Table 1, gave a caloric equivalent of biomass of 55.1 kJm⁻² for a holothurian species on the ORI Reef, Natal Province, South Africa. Allowing for a conversion of 1 kJ = 1.08 gWW, the live biomass was 59.4 gm⁻².

126. Pauly et al. (1993b) estimated the consumption of several species of tropical holothurians from published data. From this source, consumption estimates for *Holothuria* sp. (Q/B = 0.77% BWD) and *Opheodesoma spectabilis* (Q/B = 1.07% BWD) were used to compute an average consumption of 3.36 * BWyear⁻¹ for the ecological group "holothurians" of the present Caribbean reef model.

127. In Pauly et al. (1993b), estimates of the growth constant K (year⁻¹) were given for various species of holothurians. The mean annual K of species similar to those occurring in the PRVI area was 0.29, assumed to equal M (= P/B).

128. Corvea et al. (1990, in Fig. 36) gave a dry biomass range for echinoderms in the Gulf of Batabanó, Cuba, of 351-500 g10m⁻². This corresponded to a live weight of 140.4-200 gm⁻².

129. Glynn (1973, in Table 2) gave a dry biomass for echinoderms on a Caribbean *P. furcata* reef of 263.3 gm⁻². This corresponded to a live weight of 1,053.2 gm⁻².

130. Dahl (1973, in Table 1) gave a wet biomass of 8.8 g 0.25m⁻² which corresponded to 35.2 gm⁻² for tunicates on a Puerto Rican algal plain.

131. Berry (1982, in Table 1) gave a biomass of 148.8 gm⁻²DW (flesh only) for the ascidian *Pyura stolonifera* on the ORI Reef, Natal Province, South Africa. Allowing for DW = 20% of WW (organic tissue) yielded a live weight of 744 gm⁻².

132. Consumption of ascidians (sessile tunicates) was estimated as follows: Mullin (1983) provided equations to compute filtering rates of the planktonic living salp *Thalia democratica* on phytoplankton and bacteria. Benthic tunicates also filter phytoplankton and microfauna (besides zooplankton and detritus particles); hence, the suggested filtering rates were assumed to be valid also for sessile ascidians. From stomach analyses of West Indian reef fishes (Randall 1967) and from a collection of Caribbean reef invertebrates and plants (Colin 1978), species composition and rough estimates of abundance were obtained. Meinkoth (1981) provided information on size of solitary and colonial ascidians. Weight estimates were obtained by computing the volume of a cylinder whose height corresponded to the length of an individual ascidian and whose radius corresponded to its width/2, and by setting the specific weight of the cylinder equal to that of water (1 cm³ = 1 g). Body carbon of ascidians was computed via the equations in Mullin (1983):

$$\text{for solitaries } C = 1.03 * L^{1.775} \quad \dots 3.5)$$

$$\text{for aggregates } C = 0.515 * L^{2.1} \quad \dots 3.6)$$

where carbon is organic carbon content of body in mg and L is body length in mm. Mullin (1983) reported median total particulate organic carbon in his experimental filtering media as 142 µgCl⁻¹ (range 69-200 µgCl⁻¹). I assumed from diet composition percentages for ascidians (see Appendix Table 8.6.2) that 60% of the volume filtered (VF) was grazed at rates for bacteria and 40% at rates for phytoplankton. The VF of every food item was computed by multiplying the corresponding grazing rates with the body carbon of the respective species. The consumption of a food item was obtained by dividing the total organic carbon content (142 µgCl⁻¹) into the respective percentages (60% for bacteria, 40% for phytoplankton) and by then multiplying the shares by VF. Rations per hour were converted into consumption of % of body weight applying mean carbon-weight (C/W) ratios of 0.0112% for solitary and 0.0013% for colonial species. The range of consumption values included in Table 3.10 does not report the results for the colonial species as they are unrealistically

low. An average annual Q/B of 24 is the median for solitary species. Table 3.16 shows the data used for the estimation of food consumption for ascidians as an ecological group in the present Caribbean reef model.

133. Berry (1982), in Table 1, gave a caloric equivalent of biomass of 2,300 kJm⁻² for organic tissue of the detritivorous ascidian *Pyura stolonifera* on the ORI Reef, Natal Province, South Africa. Production of organic tissue was given as 5,213 kJm⁻²year⁻¹. Hence, the P/B ratio was 2.3 year⁻¹.

134. U. Richter (Hagenbecks Tierpark, Hamburg) reported the following personal observations: (a) juvenile *Emydura albertinii* (marine turtle, New Guinea, piscivorous) of 65 g live weight consumed around 0.5 g fish or shellfish flesh per day which was 0.77% BWD and 2.8*BWyear⁻¹, (b) a "giant turtle" (species name not available) of 300 kg consumed around 2 kg of plant food per day which was 0.67% BWD or 2.5*BWyear⁻¹. These values of turtle consumption were somewhat lower than that of 3.5*BWyear⁻¹ estimated by Polovina (1984b) in his coral reef model of French Frigate Shoals, Hawaii. The low values are surprising since, in the first case, we deal with a very young organism with high metabolic requirements. In the second case, the consumption rate appears to be low for an herbivorous animal. However, annual P/B ratio was also very low (0.15).

135. From Fig. 1 in Nuijta and Uchida (1982), maximum daily growth for juvenile loggerhead turtles (*Caretta caretta*) was obtained for a feeding rate of 3.1% BW. It is not known if the growth potential is reached at that feeding rate. This feeding rate yielded an annual consumption estimate for juvenile carnivorous loggerhead turtles of 11.5*BW. Experimental temperature was 20.5°C. For this turtle, the amount of food required to maintain basal metabolism was less for higher temperatures than for lower ones (see Nuijta and Uchida 1982). Therefore, the Q/B estimate was not corrected for temperature.

The estimates presented in points 1-135 were summarized; a range and an average input value were determined for each of the three input parameters and for each taxon for which estimates were available (Table 3.17).

3.2.4. Diet Composition Matrix

In the diet composition (DC) matrix of nonfish taxa (Appendix Table 8.6.2), constituents are listed in rough systematical order. The matrix was assembled from a wide variety of sources (see "Materials and methods" section). The references on which the matrix was based are listed in Appendix Table 8.6.3 by author(s) and year of publication.

3.3. The Models

3.3.1. The 50-Box Model

3.3.1.1. AGGREGATION OF FISH SPECIES

The results from the cluster analysis, aggregating 243 fish species stepwise into 1 species group, are depicted in Fig. 3.1. This dendrogram includes the names of fish species combined into the respective clusters. The first cutoff level at a quadratic Euclidean distance (QED) of 0.3 (20% of the total 1.5) yielded 29 fish species groups (see "Materials and methods" section).

Overall, the cluster analysis produced a distinct separation among carnivorous, omnivorous and herbivorous species. Furthermore, apex predators, such as large sharks and rays, as well

Table 3.16. Food consumption of ascidians (class: Tunicata).

| Species | N | Type | Height L(mm) | Width (mm) | Volume ^a (cm ³) | W (g) | C-Body (μ g) | C/W-ratio (%) | GR-bact (ml μ gC ⁻¹ h ⁻¹) | GR Phyto (ml μ gC ⁻¹ h ⁻¹) |
|--------------------------------|---|------|-----------------|---------------|-------------------------------------------|----------|----------------------|------------------|---------------------------------------------------------|----------------------------------------------------------|
| <i>Ascidia nigra</i> | 2 | Is | 102 | 60 | 288.4 | 288.4 | 3,785 | 0.001312 | 5.646 ^b | 8.998 ^d |
| <i>Boltenia echinata</i> | 1 | I | 25 | 35 | 24.1 | 24.1 | 312 | 0.001294 | 1.425 ^b | 2.056 ^d |
| <i>B. ovifera</i> | 1 | I | 76 | 51 | 155.3 | 155.3 | 2,245 | 0.001445 | 4.233 ^b | 6.606 ^d |
| <i>B. villosa</i> | 1 | I | 41 | 32 | 33.0 | 33.0 | 751 | 0.002275 | 2.313 ^b | 3.456 ^d |
| <i>Clavelina huntsmani</i> | ? | I | 51 | 10 | 4.0 | 4.0 | 1,106 | 0.027651 | 2.864 ^b | 4.346 ^d |
| <i>Clavelina picta</i> | ? | I | 19 | 6 | 0.5 | 0.5 | 192 | 0.038340 | 1.090 ^b | 1.541 ^d |
| <i>Didemnum albidum</i> | 2 | C | 3 | 102 | 24.5 | 24.5 | 5 | 0.000021 | 0.248 ^c | 0.314 ^e |
| <i>D. carnulentum</i> | 2 | C | 3 | 121 | 34.5 | 34.5 | 5 | 0.000014 | 0.248 ^c | 0.314 ^e |
| <i>Distaplia stylifer</i> | ? | C | 16 | 19 | 4.5 | 4.5 | 174 | 0.003865 | 0.742 ^c | 1.061 ^e |
| <i>Ecteinascidia turbinata</i> | 1 | I | 25 | 10 | 2.0 | 2.0 | 312 | 0.015600 | 1.425 ^b | 2.056 ^d |
| <i>Polycarpa obtecta</i> | 1 | Is | 51 | 44 | 77.5 | 77.5 | 1,106 | 0.001425 | 2.864 ^b | 4.346 ^d |

I = individuals, Is = solitary individuals, C = colonies.

| Species | VFbact (mlh ⁻¹) | VFphyto (mlh ⁻¹) | Cons. B. (μ gh ⁻¹) | Cons. P. (μ gh ⁻¹) | Q/B-bac (%BCh ⁻¹) | Q/B-phy (%BCh ⁻¹) | Q/B (%BCh ⁻¹) | Q/B (%BCday ⁻¹) | Q/B (%BWD) | Q/B (year ⁻¹) |
|--------------------------------|--------------------------------|---------------------------------|----------------------------------------|----------------------------------------|----------------------------------|----------------------------------|------------------------------|--------------------------------|---------------|------------------------------|
| <i>Ascidia nigra</i> | 21,372.33 | 34,058.91 | 1,816.65 | 1,941.36 | 47.99 | 51.29 | 49.64 | 1,191.34 | 13.31 | 48.57 |
| <i>Boltenia echinata</i> | 444.73 | 641.38 | 37.80 | 36.56 | 12.12 | 11.72 | 11.92 | 285.98 | 3.19 | 11.66 |
| <i>B. ovifera</i> | 9,504.47 | 14,833.16 | 807.88 | 845.49 | 35.98 | 37.66 | 36.82 | 883.63 | 9.87 | 36.03 |
| <i>B. villosa</i> | 1,736.90 | 2,549.48 | 147.64 | 147.89 | 19.66 | 19.70 | 19.68 | 472.33 | 5.28 | 19.26 |
| <i>Clavelina huntsmani</i> | 3,168.22 | 4,806.41 | 269.30 | 273.97 | 24.35 | 24.77 | 24.56 | 589.41 | 6.58 | 24.03 |
| <i>Clavelina picta</i> | 208.86 | 295.40 | 17.75 | 16.48 | 9.26 | 8.78 | 9.02 | 216.53 | 2.42 | 8.83 |
| <i>Didemnum albidum</i> | 1.28 | 1.62 | 0.11 | 0.09 | 2.11 | 1.79 | 1.95 | 46.78 | 0.06 | 0.22 |
| <i>D. carnulentum</i> | 1.28 | 1.62 | 0.11 | 0.09 | 2.11 | 1.79 | 1.95 | 46.78 | 0.06 | 0.22 |
| <i>Distaplia stylifer</i> | 129.04 | 184.62 | 10.97 | 10.52 | 6.31 | 6.05 | 6.18 | 148.25 | 0.19 | 0.70 |
| <i>Ecteinascidia turbinata</i> | 444.73 | 641.38 | 37.8 | 36.56 | 12.12 | 11.72 | 11.92 | 285.98 | 3.19 | 11.66 |
| <i>Polycarpa obtecta</i> | 3,168.22 | 4,806.41 | 269.30 | 273.97 | 24.35 | 24.77 | 24.56 | 589.41 | 6.58 | 24.03 |

Annual consumption of ascidians (median of all values except colonies).

24.00

^aVolume of cylinder: $\pi \cdot r^2 \cdot h$ ($r = W/2$).

^bGrazing rate for solitaries on bacteria: $0.061 \cdot L^{0.979}$.

^cGrazing rate for aggregates on bacteria: $0.121 \cdot L^{0.854}$.

^dGrazing rate for solitaries on phytoplankton: $0.07 \cdot L^{1.05}$.

^eGrazing rate for aggregates on phytoplankton: $0.141 \cdot L^{0.728}$.

Table 3.17. Key input parameters of 41 nonfish groups for Caribbean coral reef models; average values = mean of two values or median of > two values; P/B = production/biomass ratio, Q/B = annual consumption per unit biomass.

Net primary production (PP) of overall system: 20,000 g·m⁻²·year⁻¹ WW.

Export of benthic producers from system: 480 g·m⁻²·year⁻¹ WW.

| ID | Nonfish group | Range of biomass (g·m ⁻² WW) | Average biomass (g·m ⁻² WW) | Range of P/B (year ⁻¹) | Average P/B (year ⁻¹) | Range of Q/B (year ⁻¹) | Average Q/B (year ⁻¹) |
|------|------------------------------------------|-----------------------------------------|----------------------------------------|------------------------------------|-----------------------------------|------------------------------------|-----------------------------------|
| D 1 | Detritus, POM, DOM | 1,000-3,000 | 2,000.000 | | | | |
| A 1 | Benthic autotrophs (excluding symbionts) | 109-3,300 | 1,500.000 | 12.8-30.8 | 13.25 | | |
| A 2 | Symbiotic algae | | 730.000 | 9.3-16.6 | 10.20 | | |
| A 3 | Phytoplankton | 0.1-83 | 42.000 | | 70.00 | | |
| I 1 | Decomposers/microfauna | 3.1-7.7 | 5.400 | 170-220 | 195.00 | 1,900-2,200 | 2,050.00 |
| I 2 | Zooplankton | 5-30 | 17.500 | | 40.00 | 50-280 | 165.00 |
| I 3 | Sponges | 30.5-1,049 | 800.000 | | 1.70 | 4.02 | |
| I 4 | Fire corals | | | | | 3-15 | 9.00 |
| I 5 | Sea fans | | 14.000 | | 0.08 | 3-15 | 9.00 |
| I 6 | Sea anemones | | 1.300 | | | 3-15 | 9.00 |
| I 7 | Stony corals | 21-57 | 39.000 | | | 3-15 | 9.00 |
| I 8 | Corals | | 187.000 | 0.37-1.8 | 1.09 | 3-15 | 9.00 |
| I 9 | Bryozoans | | 9.200 | | | | |
| I 10 | Sipunculid worms | 4.5-11.5 | 7.000 | | | | |
| I 11 | Priapuloids | | | | | | |
| I 12 | Chitons | 18-61 | 36.000 | 0.26-0.42 | 0.34 | 7.2-16.2 | 11.70 |
| I 13 | Gastropods | | 39.000 | 0.1-5.9 | 2.50 | 4.7-56.4 | 14.00 |
| I 14 | Bivalves | 60-170 | 115.000 | 1.35-3.8 | 2.23 | 9-29.1 | 9.50 |
| I 15 | Scaphopods | | | | | | |
| I 16 | Shelled molluscs | 47-396 | 246.000 | | | | |
| I 17 | Squids | | | 0.9-1.6 | 1.25 | 16.6-20.8 | 18.70 |
| I 18 | Octopuses | | 28.000 | 1.09-3.06 | 2.08 | | 6.76 |
| I 19 | Polychaetes | 1.9-34 | 11.000 | 2.48-5.8 | 4.14 | 24.2-99 | 61.60 |
| I 20 | Echiuroids | | | | | | |
| I 21 | Pycnogonids | | | | | | |
| I 22 | Barnacles | 35-83 | 59.000 | | | | |
| I 23 | Stomatopods | | | | | | |
| I 24 | Amphipods | 1.1-3.5 | 2.300 | | 2.41 | 33.4-164 | 125.50 |
| I 25 | Tanaids | | | | | | 125.50 |
| I 26 | Isopods (fish parasites) | | | | 2.40 | | 125.50 |
| I 27 | Small crustaceans | | 0.160 | 1.4-3.9 | 2.65 | | |
| I 28 | Shrimps | 0.25-3.8 | 2.100 | 0.9-2.8 | 1.80 | | 26.90 |
| I 29 | Spiny lobsters | | 0.090 | 0.35-1.03 | 0.45 | | 7.40 |
| I 30 | Scyllarid lobsters | | | | | | |
| I 31 | Hermit crabs | 0.1-0.2 | 0.150 | | | | |
| I 32 | Crabs | 2.3-4.9 | 3.600 | 0.73-1.61 | 1.17 | 10.6-17.4 | 14.00 |
| I 33 | Lobsters and crabs | | 3.310 | | 0.52 | | 8.20 |
| I 34 | Crustaceans | 2.5-161 | 82.500 | | | | |
| I 35 | Hemichordates | | | | | | |
| I 36 | Asteroids | 14-25 | 19.500 | | 0.49 | | 3.24 |
| I 37 | Ophiuroids | 1-3 | 2.000 | | | | |
| I 38 | Echinoids | 40-211 | 107.000 | 0.8-2 | 1.40 | | 2.81 |
| I 39 | Holothurians | 23-59 | 46.200 | | 0.29 | | 3.36 |
| I 40 | Echinoderms | 140-1,053 | 612.000 | | | | |
| I 41 | Ascidians (Tunicata) | 35-744 | 390.000 | | 2.30 | 9-49 | 24.00 |
| R 1 | Sea turtles | | 0.015 | | 0.15 | 2.8-11.5 | 3.50 |
| B 1 | Sea birds | | 0.015 | | 5.40 | | 80.00 |

D = detritus, A = autotroph, I = invertebrate, R = reptile, B = bird.

as scombrids and large jacks were jointly aggregated into clusters due to their high aspect ratio and high asymptotic weight. The vast majority of (predominantly) carnivorous reef species and small pelagics were clustered according to the resulting mixture of asymptotic weight and aspect ratio. These basic features of aggregation were observed as well by the results of the “centroid” agglomeration algorithm. The “centroid” algorithm produced more single species clusters than “average linkage”, an undesirable feature for the present application. Besides, “average linkage” and “centroid” differed in several cases in the way they combined omnivorous species with carnivores or herbivores. Slight differences could also be observed for species with intermediate sizes and aspect ratios. Since the reef fish community can be viewed as an assemblage of species with smooth gradations in terms of size, activity and even DC than a set of separate compartments (see above), it is not surprising that species with intermediate parameter values were grouped differently by two different aggregation algorithms. What was surprising, rather, was that the results from both methods were largely compatible.

3.3.1.2. ASSEMBLAGE CHARACTERISTICS OF FISH SPECIES GROUPS

Table 3.18 presents the characteristics of 27 fish species groups which resulted from a further treatment of the 29 fish species groups from the cluster analysis. These 27 groups are part of the 50-box model, which was completed by adding 23 nonfish groups. An extended version of Table 3.18, where the species are only grouped but not condensed may be found in Appendix Table 8.7.1.

3.3.1.3. INPUT VALUES

Tables 3.19 and 3.20 present the estimates of the three key parameters: biomass, P/B ratio and Q/B, which served as input values for the groups of the 50-box model. The model was balanced based on these inputs and the corresponding food matrix, presented in Table 3.21, with all ecotrophic efficiencies (EEs) < 1 and input NPP $>$ output NPP.

3.3.1.4. MODIFICATION STRATEGY OF INPUT VALUES

The inputs were ranked intuitively according to the accuracy of the methods by which the respective inputs were obtained (see “Materials and methods”). The biomass values of fish and nonfish groups were considered to be the least accurate of all inputs. The P/B and Q/B values of nonfish groups were considered to be less accurate than those of fish groups. The diet matrix of nonfish groups was considered less accurate than that of fish groups. Within the fish groups, the DCs of large, wide-ranging top predators, like sharks, scombrids and jacks, were considered less accurate than that of the other fish groups. The inputs were gradually modified following the above sequence of accuracy.

3.3.1.5. PARAMETER ESTIMATION

Output values

In Table 3.22, the changes of the three key input parameters necessary to balance the 50-box model are documented together with the resulting change in ecotrophic efficiency. Table 3.23 documents necessary changes within the diet composition matrix. Table 3.24 shows

Table 3.18. Characteristics of 27 fish species groups defined by cluster analysis.

L_∞, W_∞ = asymptotic length/live weight.

H = habitat (a feature not included in cluster analysis), P = pelagic, D = demersal.

| Group no. | Fish species group | No. of species | Range of L _∞ (cm) | Range of W _∞ (g) | Average aspect ratio ^a | Average food type ^a | H | Most important species in group (by biomass) |
|-----------------|-----------------------------------------------|----------------|------------------------------|-----------------------------|-----------------------------------|--------------------------------|-----|---------------------------------------------------------------------|
| 1 | Large sharks/rays, carnivorous | 13 | 200-457 | 90,000-672,000 | 7 | 0 | P/D | <i>Dasyatis americana</i> <i>Galeocerdo cuvier</i> |
| 2 | Sharks/scombrids, carnivorous | 6 | 83.5-150 | 4,900-34,300 | 6.65 | 0 | P/D | <i>Scomberomorus regalis</i> <i>Scomberomorus cavalla</i> |
| 3 | Large jacks, carnivorous | 3 | 97-194 | 24,000-80,000 | 4.5 | 0 | P/D | <i>Seriola dumerili</i> <i>Trachinotus falcatus</i> |
| 4 | Intermediate jacks, carnivorous | 8 | 56-107 | 2,300-10,700 | 4.5 | 0 | P/D | <i>Caranx ruber</i> <i>Caranx latus</i> |
| 5 | Small jacks, carnivorous | 4 | 32-47 | 444-1,240 | 3.6 | 0 | P/D | <i>Selar crumenophtalmus</i> <i>Decapterus punctatus</i> |
| 6 | Intermediate reef fish, carnivorous 1 | 12 | 45-90 | 2,586-12,407 | 2.45 | 0 | D | <i>Ocyurus chrysurus</i> <i>Lutjanus apodus</i> |
| 7 ^b | Large to intermediate schooling fish, pelagic | 11 | 18-135 | 150-2,800 | 3.0 | 0 | P | <i>Harengula humeralis</i> <i>Harengula clupeiola</i> |
| 8 | Intermediate reef fish, carnivorous 2 | 39 | 13-55 | 57-1,795 | 2.5 | 0 | D | <i>Haemulon flavolineatum</i> <i>Lutjanus griseus</i> |
| 9 | Hemiramphidae, herbivorous | 2 | 27-32.5 | 167-298 | 3.1 | 0.8 | P | <i>Hemiramphus brasiliensis</i> <i>Hyporhamphus unifasciatus</i> |
| 10 | Kyphosidae, herbivorous | 2 | 70-80 | 6,122-9,139 | 2.85 | 1 | D | <i>Kyphosus sectatrix</i> <i>Kyphosus incisor</i> |
| 11 | Intermediate reef fish, herbivorous | 10 | 20-38 | 288-1,200 | 2.25 | 0.85 | D | <i>Acanthurus coeruleus</i> <i>Acanthurus chirurgus</i> |
| 12 | Large reef fish, carnivorous | 13 | 51-190 | 7,728-70,000 | 1.6 | 0 | D | <i>Diodon holocanthus</i> <i>Sphyræna barracuda</i> |
| 13 | Intermediate reef fish, carnivorous 3 | 36 | 20-135 | 290-5,010 | 1.4 | 0 | D | <i>Dactylopterus volitans</i> <i>Malacanthus plumieri</i> |
| 14 | Small reef fish, carnivorous 1 | 29 | 10-50 | 30-325 | 1.3 | 0 | D | <i>Halichoeres bivittatus</i> <i>Xyrichtys novacula</i> |
| 15 ^a | Small schooling fish, pelagic | 4 | 6.8-48 | 11-110 | 1.8 | 0 | P | <i>Jenkinsia lamprotaenia</i> <i>Atherinomorus stipes</i> |
| 16 | Engraulidae, herbivorous | 2 | 8-15 | 8-50 | 1.9 | 0.7 | P | <i>Anchoa hepsetus</i> <i>Anchoa lyolepis</i> |
| 17 | Small reef fish, carnivorous 2 | 12 | 5-11 | 3.3-37 | 1.43 | 0 | D | <i>Apogon conklini</i> <i>Apogon maculatus</i> |
| 19 | Intermediate reef fish, carnivorous 4 | 7 | 36-76 | 1,306-5,572 | 1.3 | 0.1 | D | <i>Holacanthus ciliaris</i> <i>Lactophrys bicaudalis</i> |
| 20 | Small reef fish, omnivorous 1 | 5 | 10-20 | 54-448 | 1.5 | 0.4 | D | <i>Stegastes fuscus</i> <i>Monacanthus ciliatus</i> |
| 21 | Small reef fish, omnivorous 2 | 4 | 10-11 | 24-82 | 1.35 | 0.2 | D | <i>Canthigaster rostrata</i> <i>Stegastes planifrons</i> |
| 22 | Small reef fish, omnivorous 3 | 5 | 7-8.5 | 4.1-13 | 1.1 | 0.7 | D | <i>Coryphopterus glaucofraenum</i> <i>Gnatholepis thompsoni</i> |
| 23 | Large Scaridae, herbivorous | 3 | 54-99 | 5,558-23,000 | 1.1 | 1 | D | <i>Scarus guacamaia</i> <i>Scarus vetula</i> |
| 24 | Intermediate Scaridae, herbivorous | 6 | 35-62 | 1,102-3,720 | 1.4 | 1 | D | <i>Sparisoma chrysopteron</i> <i>Sparisoma rubripinne</i> |
| 25 | Small Scaridae, herbivorous | 3 | 20-26 | 141-324 | 1.3 | 1 | D | <i>Sparisoma radians</i> <i>Sparisoma aurofrenatum</i> |
| 26 | Blenniidae, herbivorous | 2 | 11-12 | 27-35 | 1.05 | 0.9 | D | <i>Ophioblennius atlanticus</i> <i>Scartella cristata</i> |
| 27 | Small Gobiidae, carnivorous | 2 | 5 | 2.2-3.1 | 0.8 | 0 | D | <i>Gobiosoma horsti</i> <i>Gobiosoma evelynae</i> |

^aMedian of species-specific values.^bGrouped by habitat "midwater-pelagic" outside cluster-analysis.

Table 3.19. Key input parameters of 27 fish species groups prepared for a 50- box model of a Caribbean coral reef; average values = mean of 2 values or median of > 2 values.

| Group no. | Fish species group | No. of species | Biomass (g·m ⁻² WW) | Average P/B (year ⁻¹) | Average Q/B (year ⁻¹) |
|-----------------|--------------------------------------------|----------------|--------------------------------|-----------------------------------|-----------------------------------|
| 1 | Large sharks/rays, carnivorous | 13 | 95.676 | 0.240 | 4.900 |
| 2 | Sharks/scombrids, carnivorous | 6 | 4.139 | 0.340 | 9.150 |
| 3 | Large jacks, carnivorous | 3 | 0.452 | 0.525 | 5.700 |
| 4 | Intermediate jacks, carnivorous | 8 | 32.600 | 1.170 | 9.150 |
| 5 | Small jacks, carnivorous | 4 | 0.075 | 0.830 | 12.500 |
| 6 | Intermediate reef fish, carnivorous 1 | 12 | 50.400 | 0.580 | 6.850 |
| 7 ^a | Large-intermediate schooling fish, pelagic | 11 | 31.920 | 0.680 | 12.400 |
| 8 | Intermediate reef fish, carnivorous 2 | 39 | 46.376 | 1.265 | 10.300 |
| 9 | Hemiramphidae, herbivorous | 2 | 0.015 | 1.230 | 39.100 |
| 10 | Kyphosidae, herbivorous | 2 | 8.066 | 0.600 ^b | 23.600 |
| 11 | Intermediate reef fish, herbivorous | 10 | 13.787 | 0.710 | 25.650 |
| 12 | Large reef fish, carnivorous | 13 | 63.121 | 0.380 | 3.900 |
| 13 | Intermediate reef fish, carnivorous 3 | 36 | 20.713 | 0.640 | 6.400 |
| 14 | Small reef fish, carnivorous 1 | 29 | 4.652 | 1.665 | 9.800 |
| 15 ^a | Small schooling fish, pelagic | 4 | 5.073 | 3.540 | 18.850 |
| 16 | Engraulidae, herbivorous | 2 | 0.019 | 2.835 | 43.400 |
| 17 | Small reef fish, carnivorous 2 | 12 | 0.065 | 3.820 | 14.650 |
| 18 | Large groupers, carnivorous | 2 | 18.113 | 0.370 | 2.300 |
| 19 | Intermediate reef fish, carnivorous 4 | 7 | 7.156 | 0.550 | 6.000 |
| 20 | Small reef fish, omnivorous 1 | 5 | 7.068 | 1.600 | 16.400 |
| 21 | Small reef fish, omnivorous 2 | 4 | 0.660 | 1.505 | 16.000 |
| 22 | Small reef fish, omnivorous 3 | 5 | 0.096 | 2.530 | 39.700 |
| 23 | Large Scaridae, herbivorous | 3 | 94.910 | 0.850 | 13.500 |
| 24 | Intermediate Scaridae, herbivorous | 6 | 7.871 | 1.155 | 20.750 |
| 25 | Small Scaridae, herbivorous | 3 | 5.075 | 0.940 | 33.900 |
| 26 | Blenniidae, herbivorous | 2 | 0.012 | 2.840 | 36.100 |
| 27 | Small Gobiidae, carnivorous | 2 | 0.175 | 3.140 | 17.450 |

^aGrouped by habitat "midwater-pelagic" outside cluster analysis.

^bAssumed value, estimated from groups with similar ecology.

Table 3.20. Key input parameters of 23 nonfish groups prepared for a 50-box model of a Caribbean coral reef; average values = mean of 2 values or median of > 2 values.

NPP of overall system: 20,000 (15,000-25,000) g·m⁻²·year⁻¹ WW.

Export of benthic producers from system: 480 g·m⁻²·year⁻¹ WW.

| | Nonfish group | Range of biomass (g·m ⁻² WW) | Average biomass (g·m ⁻² WW) | Range of P/B (year ⁻¹) | Average P/B (year ⁻¹) | Range of Q/B (year ⁻¹) | Average Q/B (year ⁻¹) |
|----|-------------------------------------------------------|-----------------------------------------|----------------------------------------|------------------------------------|-----------------------------------|------------------------------------|-----------------------------------|
| 1 | Sea birds | | 0.015 | | 5.40 | | 80.00 |
| 2 | Squids | | *1.000 | 0.9-1.6 | 1.25 | 16.6-20.8 | 18.70 |
| 3 | Sea turtles | | 0.015 | | 0.15 | 2.8-11.5 | 3.50 |
| 4 | Octopuses | | 28.000 | 1.09-3.06 | 2.08 | | 6.76 |
| 5 | Lobsters | | 1.090 | 0.35-1.03 | 0.45 | | 7.40 |
| 6 | Crabs | 2.3-4.9 | 3.600 | 0.73-1.61 | 1.17 | 10.6-17.4 | 14.00 |
| 7 | Shrimps/hermit crabs/ stomatopods | 0.1-3.8 | 3.250 | 0.9-2.8 | 1.80 | | 26.90 |
| 8 | Small benthic arthropods | 4.1-6.1 | 5.300 | 2.4-2.41 | 2.40 | 33.4-164 | 125.50 |
| 9 | Asteroids | 14-25 | 19.500 | | 0.49 | | 3.24 |
| 10 | Echinoids | 40-211 | 107.000 | 0.8-2 | 1.40 | | 2.81 |
| 11 | Gastropods | | 39.000 | 0.1-5.9 | 2.50 | 4.7-56.4 | 14.00 |
| 12 | Chitons/scaphopods | 19-62 | 37.000 | 0.26-0.42 | 0.34 | 7.2-16.2 | 11.70 |
| 13 | Polychaetes/priapuloids/ ophiurids | 3.9-38 | 14.000 | 2.48-5.8 | 4.14 | 24.2-99 | 61.60 |
| 14 | Holothuroids/sipunculids/ echiuroids/hemichordates | 29.5-72.5 | 55.200 | | 0.29 | | 3.36 |
| 15 | Bivalves | 60-170 | 115.000 | 1.35-3.8 | 2.23 | 9-29.1 | 9.50 |
| 16 | Ascidians/barnacles/ bryozoans | 79-836 | 458.000 | | 2.30 | 9-49 | 24.00 |
| 17 | Sponges | 30.5-1,049 | 800.000 | | 1.70 | | 4.02 |
| 18 | Corals/sea anemones | 54-188 | 121.000 | 0.37-1.8 | 1.09 | 3-15 | 9.00 |
| 19 | Zooplankton | 5-30 | 17.500 | | 40.00 | 50-280 | 165.00 |
| 20 | Decomposers/microfauna | 3.1-7.7 | 5.400 | 170-220 | 195.00 | 1,900-2,200 | 2,050.00 |
| 21 | Phytoplankton | 0.1-73 | 26.000 | | 70.00 | - | - |
| 22 | Benthic autotrophs | 840-4,030 | 2,230.000 | 12.8-30.8 | 13.25 | - | - |
| 23 | Detritus, POM, DOM | 1,000-3,000 | 2,000.000 | - | - | - | - |

* An arbitrary minimum biomass of 1 g·m⁻² has been assigned to all taxa for which an estimate was lacking.

Table 3.21 . Diet composition matrix prepared for the construction of a 50-box model of a Caribbean coral reef.

| Group no. | Predator group | No. of original groups or species | Prey group | | | | | | | | | | |
|-----------------|---------------------------------------------------|-----------------------------------|------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| | | | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 |
| 1 | Large sharks/rays, carnivorous | 13 | 0.049 | 0.017 | 0.003 | 0.007 | 0.004 | 0.009 | 0.019 | 0.054 | 0.002 | 0.002 | 0.042 |
| 2 | Sharks/scombrids, carnivorous | 6 | - | 0.013 | - | 0.031 | 0.031 | 0.015 | 0.176 | 0.04 | 0.115 | - | 0.016 |
| 3 | Large jacks, carnivorous | 3(2) ^b | - | - | - | 0.134 | - | 0.135 | - | 0.223 | - | - | 0.044 |
| 4 | Intermediate jacks, carnivorous | 8(4) | - | - | - | - | - | 0.001 | 0.056 | 0.11 | - | - | 0.107 |
| 5 | Small jacks, carnivorous | 4(2) | - | - | - | 0.002 | 0.001 | - | 0.014 | - | - | - | 0.02 |
| 6 | Intermediate reef fish, carnivorous 1 | 13(12) | - | - | - | - | - | 0.001 | - | 0.009 | - | - | - |
| 7 ^a | Large to intermediate schooling fish, pelagic | 11(9) | - | - | - | - | - | - | 0.003 | - | - | - | 0.006 |
| 8 | Intermediate reef fish, carnivorous 2 | 39(35) | - | - | - | 0.001 | - | - | 0.001 | 0.04 | - | - | - |
| 9 | Hemiramphidae, herbivorous | 2(1) | - | - | - | - | - | - | - | - | - | - | - |
| 10 | Kyphosidae, herbivorous | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 11 | Intermediate reef fish, herbivorous | 10(9) | - | - | - | - | - | - | - | - | - | - | - |
| 12 | Large reef fish, carnivorous | 13(10) | - | - | 0.004 | 0.024 | 0.02 | 0.019 | 0.043 | 0.058 | - | - | 0.019 |
| 13 | Intermediate reef fish, carnivorous 3 | 36(29) | - | - | - | - | 0.001 | - | 0.002 | 0.021 | - | - | 0.005 |
| 14 | Small reef fish, carnivorous 1 | 29(23) | - | - | - | - | - | - | 0.002 | 0.002 | 0.002 | - | - |
| 15 ^a | Small schooling fish, pelagic | 4 | - | - | - | - | - | - | - | - | - | - | - |
| 16 | Engraulidae, herbivorous | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 17 | Small reef fish, carnivorous 2 | 12(11) | - | - | - | - | - | - | - | 0.002 | - | - | 0.001 |
| 18 | Large groupers, carnivorous | 2 | 0.041 | - | - | - | - | - | - | 0.191 | - | - | - |
| 19 | Intermediate reef fish, carnivorous 4 | 7 | - | - | - | - | - | - | - | - | - | - | - |
| 20 | Small reef fish, omnivorous 1 | 5(4) | - | - | - | - | - | - | 0.003 | 0.003 | 0.003 | - | - |
| 21 | Small reef fish, omnivorous 2 | 4(3) | - | - | - | - | - | - | - | - | - | - | - |
| 22 | Small reef fish, omnivorous 3 | 5 | - | - | - | - | - | - | - | - | - | - | - |
| 23 | Large Scaridae, herbivorous | 3 | - | - | - | - | - | - | - | - | - | - | - |
| 24 | Intermediate Scaridae, herbivorous | 6(5) | - | - | - | - | - | - | - | - | - | - | - |
| 25 | Small Scaridae, herbivorous | 3 | - | - | - | - | - | - | - | - | - | - | - |
| 26 | Blenniidae, herbivorous | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 27 | Small Gobiidae, carnivorous | 2(1) | - | - | - | - | - | - | - | - | - | - | - |
| 28 | Sea birds | 1 | - | 0.02 | 0.01 | 0.06 | 0.03 | 0.01 | 0.28 | 0.04 | 0.1 | - | 0.01 |
| 29 | Squids | 1 | - | - | - | - | - | - | 0.12 | - | 0.03 | - | - |
| 30 | Sea turtles | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 31 | Octopuses | 1 | - | - | - | - | - | - | - | 0.002 | - | - | 0.002 |
| 32 | Lobsters | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 33 | Crabs | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 34 | Shrimps/hermit crabs/stomatopods | 3 | - | - | - | - | - | - | - | 0.01 | - | - | - |
| 35 | Amphipods/isopods/tanaids/pycnogonids | 4 | - | - | - | - | 0.005 | - | 0.01 | 0.03 | 0.005 | - | 0.01 |
| 36 | Asteroids | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 37 | Echinoids | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 38 | Gastropods | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 39 | Chitons/scaphopods | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 40 | Polychaetes/priapuloids/ophiuroids | 3 | - | - | - | - | - | - | - | - | - | - | - |
| 41 | Holothuroids/sipunculids/echiuroids/hemichordates | 4 | - | - | - | - | - | - | - | - | - | - | - |
| 42 | Bivalves | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 43 | Ascidians/barnacles/bryozoans | 3 | - | - | - | - | - | - | - | - | - | - | - |
| 44 | Sponges | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 45 | Sea anemones/corals | 4 | - | - | - | - | - | - | - | - | - | - | - |
| 46 | Zooplankton | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 47 | Microfauna | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 48 | Phytoplankton | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 49 | Benthic autotrophs | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 50 | Detritus, POM, DOM | 1 | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 3.21 continued

| Group no. | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | 23 | 24 | 25 | 26 | 27 |
|-----------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| 1 | 0.009 | 0.067 | 0.043 | 0.049 | 0.028 | 0.03 | 0.001 | 0.006 | 0.007 | 0.004 | 0.001 | 0.002 | 0.004 | 0.006 | 0.003 | - |
| 2 | 0.014 | 0.084 | 0.061 | 0.136 | 0.04 | 0.009 | - | - | 0.003 | 0.001 | - | - | 0.001 | 0.002 | 0.001 | - |
| 3 | - | 0.134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 4 | - | - | - | 0.11 | 0.107 | 0.001 | - | - | 0.053 | 0.053 | 0.053 | 0.016 | 0.073 | 0.122 | 0.053 | - |
| 5 | - | - | - | 0.006 | 0.019 | - | - | - | - | - | - | - | - | - | - | - |
| 6 | - | 0.057 | 0.009 | 0.062 | - | 0.003 | - | - | 0.02 | - | - | 0.006 | 0.008 | 0.016 | - | - |
| 7 ^a | - | - | - | 0.047 | 0.006 | - | - | - | - | - | - | - | - | - | - | - |
| 8 | 0.001 | 0.001 | - | 0.067 | 0.002 | 0.001 | - | - | - | - | - | - | - | - | - | - |
| 9 | - | - | - | 0.2 | - | - | - | - | - | - | - | - | - | - | - | - |
| 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 12 | 0.017 | 0.021 | 0.002 | 0.059 | 0.001 | - | - | - | 0.006 | 0.017 | - | 0.006 | 0.008 | 0.006 | 0.001 | - |
| 13 | - | 0.009 | 0.068 | 0.002 | 0.002 | 0.005 | - | - | 0.006 | 0.001 | 0.015 | 0.002 | 0.002 | 0.001 | 0.002 | 0.007 |
| 14 | - | - | 0.005 | 0.002 | 0.002 | 0.002 | - | - | 0.002 | 0.002 | 0.006 | - | - | 0.002 | 0.006 | 0.002 |
| 15 ^a | - | - | - | 0.002 | 0.002 | - | - | - | - | - | - | - | - | - | - | - |
| 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 17 | - | - | 0.012 | - | - | 0.001 | - | - | 0.001 | - | 0.004 | - | - | - | 0.001 | 0.002 |
| 18 | 0.232 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 20 | - | - | 0.008 | 0.004 | 0.004 | 0.004 | - | - | 0.004 | 0.004 | 0.004 | - | - | 0.004 | 0.004 | 0.004 |
| 21 | - | - | 0.003 | 0.003 | 0.003 | 0.003 | - | - | - | 0.003 | 0.003 | - | - | - | 0.003 | 0.002 |
| 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 28 | 0.005 | 0.03 | 0.025 | 0.2 | 0.1 | 0.008 | - | 0.005 | 0.005 | 0.003 | - | 0.002 | 0.005 | 0.002 | - | - |
| 29 | - | - | - | 0.12 | 0.03 | - | - | - | - | - | - | - | - | - | - | - |
| 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 31 | 0.001 | 0.003 | 0.007 | - | - | 0.001 | - | - | 0.002 | 0.002 | - | - | - | - | - | - |
| 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 33 | - | - | 0.013 | - | - | 0.007 | - | - | - | - | - | - | - | - | - | - |
| 34 | - | - | 0.01 | - | 0.01 | 0.005 | - | - | - | - | - | - | - | - | - | - |
| 35 | 0.01 | 0.03 | 0.02 | 0.005 | - | 0.01 | - | - | 0.005 | - | - | - | - | - | - | - |
| 36 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 38 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 39 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 40 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 41 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 3.21 continued

| Group no. | 28 | 29 | 30 | 31 | 32 | 33 | 34 | 35 | 36 | 37 | 38 | 39 | 40 | 41 | 42 | 43 |
|-----------------|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| 1 | 0.01 | 0.021 | 0.02 | 0.003 | 0.015 | 0.109 | 0.059 | - | - | 0.008 | 0.015 | 0.006 | 0.082 | 0.11 | 0.058 | - |
| 2 | - | 0.034 | - | 0.045 | 0.005 | 0.036 | 0.069 | - | - | - | 0.008 | 0.001 | 0.006 | - | 0.007 | - |
| 3 | - | - | - | - | - | 0.013 | 0.02 | - | - | 0.082 | 0.158 | - | - | - | 0.057 | - |
| 4 | - | 0.014 | - | - | - | 0.004 | 0.041 | - | - | - | 0.004 | - | - | - | - | - |
| 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 6 | - | 0.005 | - | 0.014 | - | 0.189 | 0.072 | - | 0.001 | 0.018 | 0.017 | - | 0.006 | 0.003 | 0.01 | 0.017 |
| 7 [†] | - | - | - | - | - | - | 0.007 | - | - | - | - | - | 0.195 | - | - | - |
| 8 | - | - | - | 0.021 | 0.001 | 0.232 | 0.091 | 0.031 | - | 0.023 | 0.035 | 0.033 | 0.255 | 0.096 | 0.043 | 0.001 |
| 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 11 | - | - | - | - | - | - | 0.004 | - | - | - | 0.001 | - | 0.002 | - | - | - |
| 12 | - | 0.001 | - | 0.009 | 0.002 | 0.032 | 0.03 | - | - | 0.076 | 0.43 | - | - | - | 0.083 | - |
| 13 | - | - | - | 0.015 | - | 0.259 | 0.202 | 0.042 | - | 0.006 | 0.004 | 0.01 | 0.072 | 0.021 | 0.023 | 0.01 |
| 14 | - | - | - | - | - | 0.189 | 0.056 | 0.042 | - | 0.051 | 0.144 | 0.04 | 0.112 | 0.008 | 0.06 | - |
| 15 [*] | - | - | - | - | - | - | - | 0.015 | - | - | - | - | - | - | - | 0.006 |
| 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 17 | - | - | - | 0.004 | - | 0.063 | 0.23 | 0.123 | - | - | 0.005 | 0.002 | 0.079 | 0.001 | 0.002 | - |
| 18 | - | - | 0.035 | - | 0.426 | 0.075 | - | - | - | - | - | - | - | - | - | - |
| 19 | - | - | - | - | - | 0.001 | 0.002 | - | 0.002 | 0.003 | - | - | 0.006 | 0.006 | - | 0.023 |
| 20 | - | - | - | - | - | 0.025 | 0.024 | 0.056 | - | - | 0.038 | - | 0.071 | - | 0.008 | 0.002 |
| 21 | - | - | - | - | - | 0.07 | 0.057 | 0.051 | 0.021 | 0.021 | 0.035 | - | 0.129 | 0.005 | 0.023 | 0.012 |
| 22 | - | - | - | - | - | - | - | 0.008 | - | - | - | - | 0.083 | - | 0.071 | - |
| 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 26 | - | - | - | - | - | - | - | - | - | - | 0.001 | - | - | - | - | - |
| 27 | - | - | - | - | - | - | - | 1 | - | - | - | - | - | - | - | - |
| 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 29 | - | 0.3 | - | - | - | - | 0.1 | - | - | - | - | - | - | - | - | - |
| 30 | - | - | - | - | 0.06 | 0.025 | 0.04 | - | 0.02 | 0.02 | 0.025 | - | - | - | 0.02 | 0.02 |
| 31 | - | - | - | 0.01 | - | 0.035 | 0.03 | - | - | - | 0.83 | 0.01 | 0.005 | - | 0.06 | - |
| 32 | - | - | - | - | - | - | - | - | - | - | 0.15 | - | 0.1 | - | 0.5 | - |
| 33 | - | - | - | - | - | 0.04 | 0.07 | 0.01 | - | 0.002 | 0.08 | - | 0.004 | 0.001 | 0.03 | 0.003 |
| 34 | - | 0.015 | - | 0.02 | 0.015 | 0.005 | 0.065 | 0.23 | 0.005 | 0.005 | 0.095 | 0.02 | 0.12 | 0.005 | 0.125 | - |
| 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.02 |
| 36 | - | - | - | - | - | 0.02 | - | - | 0.12 | 0.02 | 0.2 | 0.01 | 0.06 | 0.01 | 0.09 | 0.005 |
| 37 | - | - | - | - | - | - | - | - | 0.01 | 0.01 | - | - | 0.01 | 0.01 | 0.01 | - |
| 38 | - | - | - | - | 0.004 | 0.002 | 0.006 | 0.006 | - | - | 0.04 | 0.06 | 0.1 | 0.018 | 0.03 | 0.048 |
| 39 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.04 |
| 40 | - | - | - | - | - | - | - | 0.04 | - | - | 0.01 | - | 0.095 | - | 0.01 | 0.01 |
| 41 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 3.21 continued

| Group no. | 44 | 45 | 46 | 47 | 48 | 49 | 50 | Sum |
|-----------------|-------|-------|-------|-------|-------|-------|-------|-----|
| 1 | - | - | - | - | - | - | 0.016 | 1 |
| 2 | - | - | - | - | - | - | - | 1 |
| 3 | - | - | - | - | - | - | - | 1 |
| 4 | - | - | 0.022 | - | - | - | - | 1 |
| 5 | - | - | 0.938 | - | - | - | - | 1 |
| 6 | 0.111 | 0.01 | 0.318 | - | - | 0.018 | - | 1 |
| 7 ^a | - | - | 0.722 | - | - | 0.014 | - | 1 |
| 8 | 0.001 | 0.004 | 0.018 | - | - | 0.001 | - | 1 |
| 9 | - | - | - | - | - | 0.8 | - | 1 |
| 10 | - | - | - | - | - | 1 | - | 1 |
| 11 | - | - | 0.002 | - | - | 0.533 | 0.458 | 1 |
| 12 | - | - | 0.006 | - | - | - | - | 1 |
| 13 | 0.004 | 0.003 | 0.173 | - | - | 0.005 | - | 1 |
| 14 | - | - | 0.26 | - | - | 0.001 | - | 1 |
| 15 ^a | - | - | 0.972 | 0.003 | - | - | - | 1 |
| 16 | - | - | 0.2 | - | 0.7 | - | 0.1 | 1 |
| 17 | - | - | 0.467 | - | - | - | - | 1 |
| 18 | - | - | - | - | - | - | - | 1 |
| 19 | 0.921 | 0.007 | 0.001 | - | - | 0.028 | - | 1 |
| 20 | 0.001 | 0.032 | 0.134 | - | - | 0.297 | 0.259 | 1 |
| 21 | 0.065 | 0.118 | 0.109 | 0.005 | - | 0.18 | 0.076 | 1 |
| 22 | - | - | 0.252 | - | - | 0.294 | 0.292 | 1 |
| 23 | 0.002 | 0.001 | - | - | - | 0.997 | - | 1 |
| 24 | 0.001 | - | - | - | - | 0.999 | - | 1 |
| 25 | - | - | - | - | - | 1 | - | 1 |
| 26 | - | - | 0.005 | - | - | 0.497 | 0.497 | 1 |
| 27 | - | - | - | - | - | - | - | 1 |
| 28 | - | - | 0.05 | - | - | - | - | 1 |
| 29 | - | - | 0.3 | - | - | - | - | 1 |
| 30 | 0.34 | 0.016 | - | - | - | 0.414 | - | 1 |
| 31 | - | - | - | - | - | - | - | 1 |
| 32 | - | - | - | - | - | 0.15 | 0.1 | 1 |
| 33 | 0.02 | 0.016 | 0.04 | 0.03 | 0.03 | 0.574 | 0.03 | 1 |
| 34 | - | - | 0.15 | 0.055 | - | 0.005 | 0.03 | 1 |
| 35 | 0.03 | 0.058 | 0.1 | 0.08 | - | 0.447 | 0.125 | 1 |
| 36 | 0.005 | 0.012 | - | 0.04 | - | 0.058 | 0.35 | 1 |
| 37 | 0.05 | 0.08 | - | - | - | 0.82 | - | 1 |
| 38 | 0.04 | 0.018 | - | 0.064 | - | 0.364 | 0.2 | 1 |
| 39 | 0.02 | 0.008 | - | 0.42 | - | 0.512 | - | 1 |
| 40 | 0.025 | 0.02 | 0.11 | 0.095 | 0.15 | 0.15 | 0.285 | 1 |
| 41 | - | - | - | 0.2 | 0.015 | 0.125 | 0.66 | 1 |
| 42 | - | - | - | 0.12 | 0.16 | - | 0.72 | 1 |
| 43 | - | - | 0.225 | 0.2 | 0.235 | - | 0.34 | 1 |
| 44 | - | - | - | 0.001 | - | - | 0.999 | 1 |
| 45 | - | - | 0.15 | 0.1 | - | 0.65 | 0.1 | 1 |
| 46 | - | - | - | 0.75 | 0.2 | - | 0.05 | 1 |
| 47 | - | - | - | - | - | 0.05 | 0.95 | 1 |
| 48 | - | - | - | - | - | - | - | 0 |
| 49 | - | - | - | - | - | - | - | 0 |
| 50 | - | - | - | - | - | - | - | 0 |

^aGrouped by habitat "midwater-pelagic" outside cluster analysis.

^bValue outside parentheses: no. of species indicated by cluster analysis.

Value inside parentheses: no. of species with DC values available in DC matrix.

Table 3.22. Results of the ECOPATH II parameter estimation for the construction of a 50-box model of a Caribbean coral reef.

Original inputs which had to be modified are shown in parentheses.

P/B = annual production/biomass ratio; Q/B = annual consumption /unit biomass; EE = ecotrophic efficiency.

| Group no. | Species group | Biomass (g·m ⁻²) | Biomass change (%) | P/B (year ⁻¹) | P/B change (%) | Q/B (year ⁻¹) | Q/B change (%) | Start EE | End EE | |
|-----------|-----------------------------------------------|------------------------------|------------------------|---------------------------|----------------|---------------------------|----------------|-----------|--------|-------|
| 1 | Large sharks/rays, carnivorous | 0.300 | (95.676) | - 99.7 | 0.240 | - | 4.90 | 1.053 | 0.320 | |
| 2 | Sharks/scombrids, carnivorous | 0.414 | (4.139) | - 90 | 0.340 | - | 9.15 | 6.030 | 0.626 | |
| 3 | Large jacks, carnivorous | 0.181 | (0.452) | - 60 | 0.525 | - | 5.70 | 10.120 | 0.604 | |
| 4 | Intermediate jacks, carnivorous | 1.630 | (32.600) | - 95 | 1.170 | - | 9.15 | 0.295 | 0.411 | |
| 5 | Small jacks, carnivorous | 1.000 | (0.075) | + 1,233 | 0.830 | - | 12.50 | 184.200 | 0.725 | |
| 6 | Intermediate reef fish, carnivorous 1 | 3.024 | (50.400) | - 94 | 0.580 | - | 6.85 | 0.358 | 0.776 | |
| 7 | Large to intermediate schooling fish, pelagic | 11.172 | (31.920) | - 65 | 0.680 | - | 12.40 | 2.510 | 0.953 | |
| 8 | Intermediate reef fish, carnivorous 2 | 6.493 | (46.376) | - 86 | 1.265 | - | 10.30 | 2.201 | 0.933 | |
| 9 | Hemiramphidae, herbivorous | 1.125 | (0.015) | +7,400 | 1.230 | - | 39.10 | 527.800 | 0.888 | |
| 10 | Kyphosidae, herbivorous | 2.420 | (8.066) | - 70 | 0.600 | - | 23.60 | 0.194 | 0.587 | |
| 11 | Intermediate reef fish, herbivorous | 9.651 | (13.787) | - 30 | 0.710 | - | 25.65 | 6.855 | 0.672 | |
| 12 | Large reef fish, carnivorous | 2.525 | (63.121) | - 96 | 0.380 | - | 3.90 | 1.081 | 0.692 | |
| 13 | Intermediate reef fish, carnivorous 3 | 7.250 | (20.713) | - 65 | 0.640 | - | 6.40 | 6.187 | 0.825 | |
| 14 | Small reef fish, carnivorous 1 | 4.419 | (4.652) | - 5 | 1.665 | - | 9.80 | 6.774 | 0.955 | |
| 15 | Small schooling fish, pelagic | 10.146 | (5.073) | + 100 | 3.540 | - | 18.85 | 8.600 | 0.793 | |
| 16 | Engraulidae, herbivorous | 3.325 | (0.019) | + 17,400 | 2.835 | - | 43.40 | 979.300 | 0.791 | |
| 17 | Small reef fish, carnivorous 2 | 1.300 | (0.065) | + 1,900 | 3.820 | - | 14.65 | 101.100 | 0.884 | |
| 18 | Large groupers, carnivorous | 0.725 | (18.113) | - 96 | 0.370 | - | 2.30 | 0.070 | 0.510 | |
| 19 | Intermediate reef fish, carnivorous 4 | 1.431 | (7.156) | - 80 | 0.550 | - | 6.00 | 0.716 | 0.739 | |
| 20 | Small reef fish, omnivorous 1 | 3.534 | (7.068) | - 50 | 1.600 | - | 16.40 | 2.887 | 0.711 | |
| 21 | Small reef fish, omnivorous 2 | 0.990 | (0.660) | + 50 | 1.505 | - | 16.00 | 23.160 | 0.932 | |
| 22 | Small reef fish, omnivorous 3 | 0.960 | (0.096) | + 900 | 2.530 | - | 39.70 | 78.390 | 0.866 | |
| 23 | Large Scaridae, herbivorous | 18.982 | (94.910) | - 80 | 0.850 | - | 13.50 | 0.118 | 0.336 | |
| 24 | Intermediate Scaridae, herbivorous | 5.510 | (7.871) | - 30 | 1.155 | - | 20.75 | 3.156 | 0.607 | |
| 25 | Small Scaridae, herbivorous | 5.075 | | - | 0.940 | - | 33.90 | 9.864 | 0.735 | |
| 26 | Blenniidae, herbivorous | 0.600 | (0.012) | + 4,900 | 2.840 | - | 36.10 | 543.800 | 0.854 | |
| 27 | Small Gobiidae, carnivorous | 0.175 | | - | 3.140 | - | 17.45 | 2.740 | 0.939 | |
| 28 | Sea birds | 0.017 | (0.015) ^b | + 20 | 5.400 | - | 80.00 | 57.870 | 0.320 | |
| 29 | Squids | 1.500 | (1.000) ^b | + 50 | 1.300 | (1.25) ^a | + 4 | 17.5 | 19.360 | 0.894 |
| 30 | Sea turtles | 0.500 | (0.015) ^b | + 1,900 | 0.150 | | 3.50 | 4,815.000 | 0.876 | |
| 31 | Octopuses | 8.400 | (28.000) ^b | - 70 | 1.900 | (2.08) ^a | - 9 | 6.76 | 0.443 | 0.849 |
| 32 | Lobsters | 3.270 | (1.090) ^b | + 200 | 1.000 | (0.45) ^a | + 22 | 7.40 | 60.010 | 0.960 |
| 33 | Crabs | 19.000 | (3.600) ^c | + 428 | 1.600 | (1.17) ^a | + 37 | 14.00 | 70.960 | 0.938 |
| 34 | Shrimps/hermit crabs/stomatopods | 10.000 | (3.250) ^c | + 208 | 2.800 | (1.80) ^a | + 56 | 26.90 | 30.240 | 0.949 |
| 35 | Amphipods/isopods/tanaids/ pycnogonids | 13.250 | (5.300) ^c | + 150 | 5.000 | (2.40) ^c | + 108 | 125.50 | 7.147 | 0.927 |
| 36 | Asteroids | 25.000 | (19.500) ^a | + 28 | 0.490 | | 3.24 | 1.222 | 0.963 | |
| 37 | Echinoids | 100.000 | (107.000) ^a | - 6 | 1.100 | (1.40) ^a | - 21 | 3.70 | 0.321 | 0.648 |
| 38 | Gastropods | 46.800 | (39.000) ^a | + 20 | 2.800 | (2.50) ^a | + 12 | 14.00 | 3.727 | 0.974 |

continued...

Table 3.22 continued

Original inputs which had to be modified are shown in parentheses.

P/B = annual production/biomass ratio; Q/B = annual consumption /unit biomass; EE = ecotrophic efficiency.

| Group no. | Species group | Biomass (g·m ⁻²) | Biomass change (%) | P/B (year ⁻¹) | P/B change (%) | Q/B (year ⁻¹) | Q/B change (%) | Start EE | End EE | | | |
|-----------|-------------------------------------------------------|------------------------------|--------------------------|---------------------------|----------------|---------------------------|----------------|----------|-------------------------|-------|-------|-------|
| 39 | Chitons/scaphopods | 62.000 | (37.000) ^a | + 68 | 0.360 | (0.34) ^a | + 6 | 11.70 | - | 4.674 | 0.911 | |
| 40 | Polychaetes/priapuloids/ophiuroids | 33.000 | (14.000) ^a | + 136 | 5.200 | (4.14) ^a | + 26 | 61.60 | - | 7.265 | 0.919 | |
| 41 | Holothuroids/sipunculids/ echiuroids/hemichordates | 66.240 | (55.200) ^a | + 20 | 0.310 | (0.29) ^b | + 7 | 3.36 | - | 7.239 | 0.916 | |
| 42 | Bivalves | 109.250 | (115.000) ^a | - 5 | 2.230 | - | - | 9.50 | - | 0.549 | 0.913 | |
| 43 | Ascidians/barnacles/bryozoans | 137.400 | (458.000) ^a | - 70 | 2.300 | - | - | 20.00 | (24.00) ^a | - 17 | 0.072 | 0.798 |
| 44 | Sponges | 800.000 | - | - | 1.50 | (1.70) ^b | - 25 | 5.00 | (4.02) ^b | + 24 | 0.095 | 0.249 |
| 45 | Corals/sea anemones | 121.000 | - | - | 1.090 | - | - | 9.00 | - | - | 0.817 | 0.977 |
| 46 | Zooplankton | 32.000 | (17.500) ^c | + 83 | 40.000 | - | - | 165.00 | - | - | 4.825 | 0.921 |
| 47 | Decomposers/microfauna | 15.000 | (5.400) ^c | + 178 | 280.00 | (195.00) ^c | + 44 | 1,900.00 | (2,050.00) ^a | - 7 | 4.754 | 0.979 |
| 48 | Phytoplankton | 40.000 | (42.000) ^a | - 5 | 70.000 | - | - | - | - | - | 1.906 | 0.740 |
| 49 | Benthic autotrophs | 1,300.000 | (2,230.000) ^a | - 42 | 13.250 | - | - | - | - | - | 0.167 | 0.360 |
| 50 | Detritus, POM, DOM | 2,000.000 | - | - | - | - | - | - | - | - | - | - |

^aWithin available range.^bNo range available.^cOut of available range.

Table 3. 23. Results of the ECOPATH II parameter estimation for the construction of a 50-box model of a Caribbean coral reef diet composition matrix. Values in parentheses refer to original inputs that were modified during the parameter estimation routine.

| Group no. | Predator group | No. of original groups or species | Prey group | | | | | | | | | | |
|-----------|-----------------------------------------------|-----------------------------------|------------------|-----------------|-------|------------------|-----------------|-----------------|-----------------|-----------------|------------------|-----------------|------------------|
| | | | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 |
| 1 | Large sharks/rays, carnivorous | 13 | 0.01 (0.049) | 0.01 (0.017) | 0.003 | 0.05 (0.007) | 0.004 | 0.1 (0.009) | 0.01 (0.019) | 0.01 (0.054) | 0.002 | 0.06 (0.002) | 0.052 (0.042) |
| 2 | Sharks/scombrids, carnivorous | 6 | - | 0.01 (0.013) | - | 0.034 (0.031) | 0.05 (0.031) | 0.015 | 0.176 | 0.04 | 0.115 | 0.02 (-) | 0.02 (0.016) |
| 3 | Large jacks, carnivorous | 3(2) | - | - | - | 0.15 (0.134) | - | 0.2 (0.135) | - | 0.16 (0.223) | - | - | 0.044 |
| 4 | Intermediate jacks, carnivorous | 8(4) | - | - | - | - | - | 0.001 | 0.056 | 0.11 | - | 0.015 | 0.107 |
| 5 | Small jacks, carnivorous | 4(2) | - | - | - | 0.002 | 0.001 | - | 0.014 | - | - | - | 0.02 |
| 6 | Intermediate reef fish, carnivorous 1 | 13(12) | - | - | - | - | - | 0.001 | - | 0.009 | - | 0.003 (-) | 0.004 (-) |
| 7 | Large to intermediate schooling fish, pelagic | 11(9) | - | - | - | - | - | - | 0.003 | - | - | - | 0.01 (0.006) |
| 8 | Intermediate reef fish, carnivorous 2 | 39(35) | - | - | - | 0.001 | - | - | 0.001 | 0.02 (0.040) | - | 0.003 (-) | 0.004 (-) |
| 9 | Hemiramphidae, herbivorous | 2(1) | - | - | - | - | - | - | - | - | - | - | - |
| 10 | Kyphosidae, herbivorous | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 11 | Intermediate reef fish, herbivorous | 10(9) | - | - | - | - | - | - | - | - | - | - | - |
| 12 | Large reef fish, carnivorous | 13(10) | - | - | 0.004 | 0.024 | 0.02 | 0.03 (0.019) | 0.04 (0.043) | 0.04 (0.058) | - | 0.003 | 0.023 (0.019) |
| 13 | Intermediate reef fish, carnivorous 3 | 36(29) | - | - | - | - | 0.003 (-) | 0.01 (0.001) | 0.002 (-) | 0.01 (0.020) | - | 0.003 (-) | 0.009 (0.005) |
| 14 | Small reef fish, carnivorous 1 | 29(23) | - | - | - | - | - | - | 0.002 | 0.002 | 0.002 | - | - |
| 15 | Small schooling fish, pelagic | 4 | - | - | - | - | - | - | - | - | - | - | - |
| 16 | Engraulidae, herbivorous | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 17 | Small reef fish, carnivorous 2 | 12(11) | - | - | - | - | - | - | - | 0.002 | - | - | 0.001 |
| 18 | Large groupers, carnivorous | 2 | 0.005 (0.041) | 0.005 (-) | - | 0.01 (-) | 0.01 (-) | 0.085 (-) | - | 0.09 (0.191) | - | 0.02 (-) | 0.02 (-) |
| 19 | Intermediate reef fish, carnivorous 4 | 7 | - | - | - | - | - | - | - | - | - | - | - |
| 20 | Small reef fish, omnivorous 1 | 5(4) | - | - | - | - | - | - | 0.003 | 0.003 | 0.003 | - | - |
| 21 | Small reef fish, omnivorous 2 | 4(3) | - | - | - | - | - | - | - | - | - | - | - |
| 22 | Small reef fish, omnivorous 3 | 5 | - | - | - | - | - | - | - | - | - | - | - |
| 23 | Large Scaridae, herbivorous | 3 | - | - | - | - | - | - | - | - | - | - | - |
| 24 | Intermediate Scaridae, herbivorous | 6(5) | - | - | - | - | - | - | - | - | - | - | - |
| 25 | Small Scaridae, herbivorous | 3 | - | - | - | - | - | - | - | - | - | - | - |
| 26 | Blenniidae, herbivorous | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 27 | Small Gobiidae, carnivorous | 2(1) | - | - | - | - | - | - | - | - | - | - | - |
| 28 | Sea birds | 1 | - | 0.02 | 0.01 | 0.06 | 0.03 | 0.01 | 0.28 | 0.04 | 0.1 | - | 0.01 |
| 29 | Squids | 1 | - | - | - | - | - | - | 0.15 (0.120) | - | 0.015 (0.030) | - | - |
| 30 | Sea turtles | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 31 | Octopuses | 1 | - | - | - | - | - | - | - | 0.002 | - | - | 0.002 |
| 32 | Lobsters | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 33 | Crabs | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 34 | Shrimps/hermit crabs/ stomatopods | 3 | - | - | - | - | - | - | - | 0.01 | - | - | - |
| 35 | Amphipods/isopods/tanaids/pycnogonids | 4 | - | - | - | - | - | - | - | - | - | - | - |
| | | | | | | | | (0.005) | (0.010) | (0.030) | (0.005) | | (0.010) |
| 36 | Asteroids | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 37 | Echinoids | 1 | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 3. 23 continued

| Group no. | Predator group | No. of original groups or species | Prey group | | | | | | | | | | | |
|-----------|---------------------------------------------------|-----------------------------------|------------|---|---|---|---|---|-------|-------|---|-------|-------|---|
| | | | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | |
| 38 | Gastropods | 1 | - | - | - | - | - | - | - | - | - | - | - | - |
| 39 | Chitons/scaphopods | 2 | - | - | - | - | - | - | - | - | - | - | - | - |
| 40 | Polychaetes/priapuloids/ophiuroids | 3 | - | - | - | - | - | - | - | - | - | - | - | - |
| 41 | Holothuroids/sipunculids/echiuroids/hemichordates | 4 | - | - | - | - | - | - | - | - | - | - | - | - |
| 42 | Bivalves | 1 | - | - | - | - | - | - | - | - | - | - | - | - |
| 43 | Ascidians/barnacles/bryozoans | 3 | - | - | - | - | - | - | - | - | - | - | - | - |
| 44 | Sponges | 1 | - | - | - | - | - | - | - | - | - | - | - | - |
| 45 | Sea anemones/corals | 4 | - | - | - | - | - | - | - | - | - | - | - | - |
| 46 | Zooplankton | 1 | - | - | - | - | - | - | - | - | - | - | - | - |
| 47 | Microfauna | 1 | - | - | - | - | - | - | - | - | - | - | - | - |
| 48 | Phytoplankton | 1 | - | - | - | - | - | - | - | - | - | - | - | - |
| 49 | Benthic autotrophs | 2 | - | - | - | - | - | - | - | - | - | - | - | - |
| 50 | Contribution to detritus | 1 | - | - | - | - | - | - | 0.001 | 0.001 | - | 0.001 | 0.002 | - |

continued...

Table 3.23 continued

| Group no. | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | 23 | 24 | 25 | 26 | 27 |
|-----------|------------------|------------------|------------------|-----------------|-------|------------------|------------------|-----------------|------------------|------------------|-------|------------------|------------------|------------------|-------|------------------|
| 1 | 0.03 (0.009) | 0.025 (0.067) | 0.02 (0.043) | 0.049 | 0.028 | 0.006 (0.030) | 0.093 (0.001) | 0.02 (0.006) | 0.025 (0.007) | 0.004 | 0.001 | 0.025 (0.002) | 0.002 (0.004) | 0.001 (0.006) | - | - |
| 2 | 0.014 | 0.084 | 0.02 (0.061) | 0.136 | 0.04 | 0.02 (0.009) | - | 0.006 (-) | 0.014 (0.003) | 0.001 | - | 0.03 (-) | 0.001 | 0.002 | 0.001 | - |
| 3 | - | 0.124 (0.134) | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 4 | - | - | - | 0.11 | 0.107 | 0.001 | - | 0.005 (-) | 0.053 | 0.053 | 0.053 | 0.016 | 0.073 | 0.122 | 0.053 | - |
| 5 | - | - | - | 0.006 | 0.019 | - | - | - | - | - | - | - | - | - | - | - |
| 6 | - | 0.057 | 0.005 (0.009) | 0.062 | - | 0.003 | - | 0.003 | 0.02 | - | - | 0.03 (0.006) | 0.016 (0.008) | 0.016 | - | - |
| 7 | - | - | - | 0.047 | 0.006 | - | - | - | - | - | - | - | - | - | - | - |
| 8 | 0.001 | 0.015 (0.001) | - | 0.067 | 0.002 | 0.001 | - | 0.003 (-) | - | - | - | 0.02 (-) | 0.03 (-) | 0.01 (-) | - | - |
| 9 | - | - | - | 0.2 | - | - | - | - | - | - | - | - | - | - | - | - |
| 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 12 | 0.005 (0.017) | 0.04 (0.021) | 0.001 (0.002) | 0.059 | 0.001 | - | - | 0.003 | 0.017 (0.006) | 0.006 (0.017) | - | 0.01 (0.006) | 0.01 (0.008) | 0.008 (0.006) | 0.001 | - |
| 13 | - | 0.012 (0.009) | 0.01 (0.068) | 0.002 | 0.002 | 0.01 (0.005) | - | 0.003 (-) | 0.04 (0.006) | 0.001 | 0.015 | 0.055 (0.002) | 0.007 (0.002) | 0.006 (0.001) | 0.002 | 0.004 (0.007) |
| 14 | - | - | 0.003 (0.005) | 0.002 | 0.002 | 0.002 | - | - | 0.005 (0.002) | 0.002 | 0.006 | - | - | 0.002 | 0.006 | 0.002 |
| 15 | - | - | - | 0.002 | 0.002 | - | - | - | - | - | - | - | - | - | - | - |
| 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 17 | - | - | 0.008 (0.012) | - | - | 0.003 (0.001) | - | - | 0.001 | - | 0.004 | - | - | - | 0.001 | 0.002 |
| 18 | 0.232 | - | - | - | - | - | - | 0.01 (-) | - | - | - | 0.248 (-) | - | - | - | - |
| 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 20 | - | - | 0.004 (0.008) | 0.004 | 0.004 | 0.004 | - | - | 0.006 (0.004) | 0.004 | 0.004 | - | - | 0.004 | 0.004 | 0.003 (0.004) |
| 21 | - | - | 0.003 | 0.003 | 0.003 | 0.003 | - | - | - | 0.003 | 0.003 | - | - | - | 0.003 | 0.002 |
| 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 28 | 0.005 | 0.03 | 0.025 | 0.2 | 0.1 | 0.008 | - | 0.005 | 0.005 | 0.003 | - | 0.002 | 0.005 | 0.002 | - | - |
| 29 | - | - | - | 0.13 (0.120) | 0.03 | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 3.23 continued

| Group no. | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | 23 | 24 | 25 | 26 | 27 |
|-----------|---------|---------|-----------------|---------|-------|---------|----|----|---------|-------|----|-------|-------|-------|----|----|
| 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 31 | 0.001 | 0.003 | 0.007 | - | - | 0.001 | - | - | 0.002 | 0.002 | - | - | - | - | - | - |
| 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 33 | - | - | 0.01 (0.013) | - | - | 0.007 | - | - | - | - | - | - | - | - | - | - |
| 34 | - | - | 0.01 | - | 0.01 | 0.005 | - | - | - | - | - | - | - | - | - | - |
| 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| | (0.010) | (0.030) | (0.020) | (0.005) | - | (0.010) | - | - | (0.005) | - | - | - | - | - | - | - |
| 36 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 38 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 39 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 40 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 41 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 50 | - | - | - | 0.002 | 0.001 | - | - | - | 0.001 | - | - | 0.003 | 0.001 | 0.002 | - | - |

continued...

Table 3.23 continued

| Group no. | 28 | 29 | 30 | 31 | 32 | 33 | 34 | 35 | 36 | 37 | 38 | 39 | 40 | 41 | 42 | 43 |
|-----------|-----------------|------------------|------------------|-----------------|------------------|-----------------|------------------|------------------|-------|-----------------|-----------------|-----------------|------------------|-----------------|------------------|-----------------|
| 1 | 0.02 (0.010) | 0.005 (0.021) | 0.005 (0.020) | 0.02 (0.003) | 0.005 (0.015) | 0.05 (0.109) | 0.05 (0.059) | - | - | 0.008 | 0.01 (0.015) | - | 0.05 (0.006) | 0.05 (0.082) | 0.067 (0.110) | - |
| 2 | - | 0.01 (0.034) | - | 0.045 | 0.005 | 0.02 (0.036) | 0.05 (0.069) | - | - | - | 0.008 | - | 0.006 (0.001) | - | 0.007 | - |
| 3 | - | - | - | - | - | 0.01 (0.013) | 0.015 (0.020) | - | - | 0.082 | 0.158 | - | - | - | 0.057 | - |
| 4 | - | 0.004 (0.014) | - | - | - | 0.004 | 0.031 (0.041) | - | - | - | 0.004 | - | - | - | - | - |
| 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 6 | - | 0.001 (0.005) | - | 0.014 | - | 0.09 (0.189) | 0.03 (0.072) | - | 0.001 | 0.05 (0.018) | 0.017 | - | 0.006 | 0.003 | 0.01 | 0.06 (0.017) |
| 7 | - | - | - | - | - | - | 0.004 (0.007) | - | - | - | - | - | 0.12 (0.195) | - | - | - |
| 8 | - | - | - | 0.05 (0.021) | 0.001 | 0.1 (0.252) | 0.02 (0.091) | 0.031 | - | 0.1 (0.023) | 0.035 | 0.02 (0.033) | 0.18 (0.255) | 0.06 (0.096) | 0.073 (0.043) | 0.05 (0.001) |
| 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 11 | - | - | - | - | - | - | 0.004 | - | - | - | 0.001 | - | 0.002 | - | - | - |
| 12 | - | 0.001 | - | 0.009 | 0.002 | 0.032 | 0.03 | - | - | 0.1 (0.076) | 0.4 (0.430) | - | - | - | 0.075 (0.083) | - |
| 13 | - | - | - | 0.03 (0.015) | - | 0.15 (0.259) | 0.1 (0.202) | 0.042 | - | 0.06 (0.006) | 0.004 | 0.01 | 0.05 (0.072) | 0.01 (0.021) | 0.023 | 0.04 (0.01) |
| 14 | - | - | - | - | - | 0.1 (0.189) | 0.04 (0.056) | 0.042 | - | 0.11 (0.051) | 0.144 | 0.04 | 0.1 (0.112) | 0.008 | 0.06 | - |
| 15 | - | - | - | - | - | - | - | 0.01 (0.015) | - | - | - | - | - | - | - | 0.03 (0.006) |
| 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 17 | - | - | - | 0.02 (0.004) | - | 0.063 | 0.15 (0.230) | 0.1 (0.123) | - | - | 0.005 | 0.002 | 0.079 | 0.001 | 0.03 (0.002) | - |
| 18 | - | - | 0.035 | - | 0.16 (0.426) | 0.07 (0.075) | - | - | - | - | - | - | - | - | - | - |
| 19 | - | - | - | - | - | 0.001 | 0.002 | - | 0.002 | 0.003 | - | - | 0.006 | 0.006 | - | 0.023 |
| 20 | - | - | - | - | - | 0.01 (0.025) | 0.01 (0.024) | 0.056 | - | - | 0.038 | - | 0.05 (0.071) | - | 0.02 (0.008) | 0.02 (0.002) |
| 21 | - | - | - | - | - | 0.05 (0.070) | 0.04 (0.057) | 0.035 (0.051) | 0.021 | 0.05 (0.021) | 0.035 | - | 0.1 (0.129) | 0.005 | 0.027 (0.023) | 0.02 (0.012) |
| 22 | - | - | - | - | - | - | - | 0.008 | - | - | - | - | 0.05 (0.083) | - | 0.071 | - |
| 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 26 | - | - | - | - | - | - | - | - | - | - | 0.001 | - | - | - | - | - |
| 27 | - | - | - | - | - | - | - | 1 | - | - | - | - | - | - | - | - |
| 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 29 | - | 0.01 | - | - | - | - | 0.1 (0.300) | - | - | - | - | - | - | - | - | - |

continued...

Table 3.23 continued

| Group no. | 28 | 29 | 30 | 31 | 32 | 33 | 34 | 35 | 36 | 37 | 38 | 39 | 40 | 41 | 42 | 43 | |
|-----------|----|------------------|----|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|-----------------|
| 30 | - | - | - | - | 0.06 | 0.025 | 0.04 | - | 0.02 | 0.02 | 0.025 | - | - | - | - | 0.02 | 0.02 |
| 31 | - | - | - | 0.02 (0.010) | - | 0.025 (0.035) | 0.02 (0.030) | - | - | - | 0.5 (0.830) | 0.01 | 0.005 | - | - | 0.4 (0.060) | - |
| 32 | - | - | - | - | - | - | - | - | - | - | 0.15 | - | 0.05 (0.100) | - | - | 0.5 | - |
| 33 | - | - | - | - | - | 0.005 (0.040) | 0.01 (0.070) | 0.005 (0.010) | - | 0.05 (0.002) | 0.05 (0.080) | - | 0.004 | 0.001 | 0.05 | 0.02 (0.030) | 0.02 (0.003) |
| 34 | - | 0.005 (0.015) | - | 0.025 (0.020) | 0.005 (0.015) | 0.002 (0.005) | 0.002 (0.065) | 0.04 (0.230) | 0.01 (0.005) | 0.11 (0.005) | 0.08 (0.095) | 0.01 (0.020) | 0.05 (0.120) | 0.005 | 0.005 | 0.212 (0.125) | - |
| 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.05 (0.020) |
| 36 | - | - | - | - | - | 0.01 (0.02) | - | - | 0.08 (0.120) | 0.07 (0.020) | 0.135 (0.200) | 0.005 (0.010) | 0.03 (0.060) | 0.005 (0.010) | 0.005 (0.010) | 0.145 (0.090) | 0.02 (0.005) |
| 37 | - | - | - | - | - | - | - | - | 0.006 (0.010) | 0.015 (0.010) | - | - | 0.004 (0.010) | 0.005 (0.010) | 0.01 | - | - |
| 38 | - | - | - | - | 0.002 (0.004) | 0.002 | 0.004 (0.006) | 0.006 | - | - | 0.02 (0.040) | 0.02 (0.060) | 0.05 (0.100) | 0.015 (0.018) | 0.04 (0.030) | 0.058 (0.048) | - |
| 39 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.04 |
| 40 | - | - | - | - | - | - | - | 0.015 (0.040) | - | - | 0.01 | - | 0.03 (0.095) | - | - | 0.03 (0.010) | 0.04 (0.010) |
| 41 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| 50 | - | - | - | 0.001 | - | 0.002 | 0.002 | 0.015 | 0.001 | 0.005 | 0.006 | 0.006 | 0.018 | 0.002 | 0.01 | 0.026 | - |

continued...

Table 3.23 continued

| Group no. | 44 | 45 | 46 | 47 | 48 | 49 | 50 | Sum |
|-----------|-----------------|-----------------|------------------|------------------|-----|------------------|------------------|-----|
| 1 | - | - | - | - | - | - | 0.02 (0.016) | 1 |
| 2 | - | - | - | - | - | - | - | 1 |
| 3 | - | - | - | - | - | - | - | 1 |
| 4 | - | - | 0.022 | - | - | - | - | 1 |
| 5 | - | - | 0.938 | - | - | - | - | 1 |
| 6 | 0.2 (0.111) | 0.01 | 0.25 (0.318) | - | - | 0.029 (0.018) | - | 1 |
| 7 | - | - | 0.722 | - | - | 0.088 (0.014) | - | 1 |
| 8 | 0.07 (0.001) | 0.004 | 0.018 | - | - | 0.01 (0.001) | - | 1 |
| 9 | - | - | - | - | - | 0.8 | - | 1 |
| 10 | - | - | - | - | - | 1 | - | 1 |
| 11 | - | - | 0.002 | - | - | 0.533 | 0.458 | 1 |
| 12 | - | - | 0.006 | - | - | - | - | 1 |
| 13 | 0.07 (0.004) | 0.003 | 0.15 (0.173) | - | - | 0.052 (0.005) | - | 1 |
| 14 | - | - | 0.26 | - | - | 0.058 (0.001) | - | 1 |
| 15 | - | - | 0.95 (0.972) | 0.006 (0.003) | - | - | - | 1 |
| 16 | - | - | 0.2 | - | 0.7 | - | 0.1 | 1 |
| 17 | - | - | 0.5 (0.467) | - | - | 0.028 (-) | - | 1 |
| 18 | - | - | - | - | - | - | - | 1 |
| 19 | 0.921 | 0.007 | 0.001 | - | - | 0.028 | - | 1 |
| 20 | 0.02 (0.001) | 0.02 (0.032) | 0.1 (0.134) | - | - | 0.326 (0.297) | 0.28 (0.259) | 1 |
| 21 | 0.08 (0.065) | 0.1 (0.118) | 0.109 | 0.005 | - | 0.2 (0.180) | 0.1 (0.076) | 1 |
| 22 | - | - | 0.2 (0.252) | - | - | 0.335 (0.294) | 0.336 (0.292) | 1 |
| 23 | 0.002 | 0.001 | - | - | - | 0.997 | - | 1 |
| 24 | 0.001 | - | - | - | - | 0.999 | - | 1 |
| 25 | - | - | - | - | - | 1 | - | 1 |
| 26 | - | - | 0.005 | - | - | 0.497 | 0.497 | 1 |
| 27 | - | - | - | - | - | - | - | 1 |
| 28 | - | - | 0.05 | - | - | - | - | 1 |
| 29 | - | - | 0.565 (0.300) | - | - | - | - | 1 |

continued...

Table 3.23 continued

| Group no. | 44 | 45 | 46 | 47 | 48 | 49 | 50 | Sum |
|-----------|---------|---------|---------|---------|---------|---------|---------|-----|
| 30 | 0.34 | 0.016 | - | - | - | 0.414 | - | 1 |
| 31 | - | - | - | - | - | - | - | 1 |
| 32 | - | - | - | - | - | 0.18 | 0.12 | 1 |
| | | | | | | (0.150) | (0.100) | |
| 33 | 0.05 | 0.01 | 0.03 | 0.03 | 0.02 | 0.6 | 0.048 | 1 |
| | (0.020) | (0.016) | (0.040) | | (0.030) | (0.574) | (0.030) | |
| 34 | - | - | 0.15 | 0.055 | - | 0.084 | 0.12 | 1 |
| | | | | | | (0.005) | (0.030) | |
| 35 | 0.06 | 0.04 | 0.04 | 0.04 | - | 0.6 | 0.17 | 1 |
| | (0.030) | (0.058) | (0.100) | (0.080) | | (0.447) | (0.125) | |
| 36 | 0.02 | 0.012 | - | 0.02 | - | 0.088 | 0.36 | 1 |
| | (0.005) | | | (0.040) | | (0.058) | (0.350) | |
| 37 | 0.05 | 0.05 | - | - | - | 0.86 | - | 1 |
| | | (0.080) | | | | (0.820) | | |
| 38 | 0.04 | 0.015 | - | 0.03 | - | 0.4 | 0.298 | 1 |
| | | (0.018) | | (0.064) | | (0.364) | (0.200) | |
| 39 | 0.02 | 0.008 | - | 0.3 | - | 0.632 | - | 1 |
| | | | | (0.42) | | (0.512) | | |
| 40 | 0.05 | 0.01 | 0.05 | 0.05 | 0.1 | 0.245 | 0.37 | 1 |
| | (0.025) | (0.020) | (0.110) | (0.095) | (0.150) | (0.150) | (0.285) | |
| 41 | - | - | - | 0.1 | 0.01 | 0.13 | 0.76 | 1 |
| | | | | (0.200) | (0.015) | (0.125) | (0.660) | |
| 42 | - | - | - | 0.1 | 0.15 | - | 0.75 | 1 |
| | | | | (0.120) | (0.160) | | (0.720) | |
| 43 | - | - | 0.15 | 0.1 | 0.2 | - | 0.55 | 1 |
| | | | (0.225) | (0.200) | (0.235) | | (0.340) | |
| 44 | - | - | - | 0.001 | - | - | 0.999 | 1 |
| 45 | - | - | 0.15 | 0.1 | - | 0.65 | 0.1 | 1 |
| 46 | - | - | - | 0.6 | 0.2 | - | 0.2 | 1 |
| | | | | (0.750) | | | (0.050) | |
| 47 | - | - | - | - | - | 0.05 | 0.95 | 1 |
| 48 | - | - | - | - | - | - | (1) | (1) |
| 49 | - | - | - | - | - | - | (1) | (1) |
| 50 | 0.073 | 0.01 | 0.05 | 0.25 | 0.033 | 0.475 | - | (1) |

Table 3.24. Results of the ECOPATH II parameter estimation for the construction of a 50-box model of a Caribbean coral reef. Additional parameters. Import to detritus: 475 g·m⁻², other imports to boxes are 0; catches for all boxes are 0. Excretion = 5% and egestion = 15% (default values) of the food intake of each box.

| Group no. | Species group | Food intake ^a | Respiration ^a | Predation mortality (production *EE) ^a | Flow to detritus ^a | Assimilation ^a | Net production (g·m ⁻² ·year ⁻¹) | Trophic level |
|-----------|--------------------------------------------|--------------------------|--------------------------|---------------------------------------------------|-------------------------------|---------------------------|---------------------------------------------------------|---------------|
| 1 | Large sharks/rays, carnivorous | 1.47 | 1.104 | 0.023 | 0.343 | 1.176 | 0.072 | 3.920 |
| 2 | Sharks/scombrids, carnivorous | 3.79 | 2.890 | 0.088 | 0.810 | 3.030 | 0.141 | 4.155 |
| 3 | Large jacks, carnivorous | 1.03 | 0.730 | 0.057 | 0.244 | 0.825 | 0.094 | 4.050 |
| 4 | Intermediate jacks, carnivorous | 14.91 | 10.024 | 0.784 | 4.106 | 11.932 | 1.908 | 3.630 |
| 5 | Small jacks, carnivorous | 12.50 | 9.170 | 0.601 | 2.729 | 10.000 | 0.829 | 3.600 |
| 6 | Intermediate reef fish, carnivorous 1 | 20.71 | 14.818 | 1.361 | 4.536 | 16.572 | 1.754 | 3.480 |
| 7 | Large intermediate schooling fish, pelagic | 138.53 | 103.229 | 7.240 | 28.064 | 110.826 | 7.597 | 3.470 |
| 8 | Intermediate reef fish, carnivorous 2 | 66.88 | 45.289 | 7.660 | 13.929 | 53.502 | 8.210 | 3.440 |
| 9 | Hemiramphidae, herbivorous | 43.99 | 33.806 | 1.229 | 8.952 | 35.190 | 1.384 | 2.520 |
| 10 | Kyphosidae, herbivorous | 57.11 | 44.238 | 0.853 | 12.022 | 45.690 | 1.453 | 2.000 |
| 11 | Intermediate reef fish, herbivorous | 247.55 | 191.186 | 4.603 | 51.759 | 198.038 | 6.850 | 2.020 |
| 12 | Large reef fish, carnivorous | 9.85 | 6.919 | 0.664 | 2.265 | 7.878 | 0.960 | 3.660 |
| 13 | Intermediate reef fish, carnivorous 3 | 46.40 | 32.480 | 3.829 | 10.091 | 37.120 | 4.641 | 3.440 |
| 14 | Small reef fish, carnivorous 1 | 43.31 | 27.287 | 7.026 | 8.993 | 34.645 | 7.357 | 3.355 |
| 15 | Small schooling fish, pelagic | 191.25 | 117.085 | 28.483 | 45.684 | 153.002 | 35.918 | 3.595 |
| 16 | Engraulidae, herbivorous | 144.31 | 106.018 | 7.456 | 30.832 | 115.444 | 9.426 | 2.320 |
| 17 | Small reef fish, carnivorous 2 | 19.04 | 10.270 | 4.390 | 4.385 | 15.236 | 4.966 | 3.570 |
| 18 | Large groupers, carnivorous | 1.67 | 1.066 | 0.137 | 0.465 | 1.334 | 0.269 | 3.890 |
| 19 | Intermediate reef fish, omnivorous | 8.59 | 6.082 | 0.582 | 1.923 | 6.869 | 0.788 | 2.990 |
| 20 | Small reef fish, omnivorous 1 | 57.96 | 40.712 | 4.022 | 13.224 | 46.366 | 5.657 | 2.580 |
| 21 | Small reef fish, omnivorous 2 | 15.84 | 11.182 | 1.389 | 3.269 | 12.672 | 1.490 | 2.980 |
| 22 | Small reef fish, omnivorous 3 | 38.11 | 28.061 | 2.103 | 7.948 | 30.490 | 2.428 | 2.480 |
| 23 | Large Scaridae, herbivorous | 256.26 | 188.871 | 5.415 | 61.971 | 205.006 | 16.116 | 2.000 |
| 24 | Intermediate Scaridae, herbivorous | 114.33 | 85.102 | 3.863 | 25.367 | 91.466 | 6.364 | 2.000 |
| 25 | Small Scaridae, herbivorous | 172.04 | 132.863 | 3.507 | 35.672 | 137.634 | 4.771 | 2.000 |
| 26 | Blenniidae, herbivorous | 21.66 | 15.624 | 1.455 | 4.581 | 17.328 | 1.704 | 2.010 |
| 27 | Small Gobiidae, carnivorous | 3.05 | 1.894 | 0.516 | 0.644 | 2.443 | 0.550 | 3.285 |
| 28 | Sea birds | 1.36 | 0.996 | 0.029 | 0.334 | 1.088 | 0.091 | 4.240 |
| 29 | Squids | 26.25 | 19.050 | 1.743 | 5.457 | 21.000 | 1.950 | 3.910 |
| 30 | Sea turtles | 1.75 | 1.325 | 0.066 | 0.359 | 1.400 | 0.075 | 2.730 |
| 31 | Octopuses | 56.78 | 29.467 | 13.556 | 13.761 | 45.427 | 15.967 | 3.310 |
| 32 | Lobsters | 24.20 | 16.088 | 3.140 | 4.970 | 19.359 | 3.271 | 2.820 |
| 33 | Crabs | 266.00 | 182.400 | 28.526 | 55.074 | 212.800 | 30.412 | 2.430 |
| 34 | Shrimps/hermit crabs/stomatopods | 269.00 | 187.200 | 26.561 | 55.239 | 215.200 | 27.988 | 3.085 |
| 35 | Amphipods/isopods/tanaids/pycnogonids | 1,662.88 | 1,264.050 | 61.417 | 337.408 | 1,330.300 | 66.254 | 2.285 |
| 36 | Asteroids | 81.00 | 52.550 | 11.796 | 16.654 | 64.800 | 12.249 | 2.710 |
| 37 | Echinoids | 370.00 | 186.000 | 71.315 | 112.685 | 296.000 | 110.054 | 2.170 |
| 38 | Gastropods | 655.20 | 393.120 | 127.695 | 134.385 | 524.160 | 131.104 | 2.370 |
| 39 | Chitons/scaphopods | 725.40 | 558.000 | 20.339 | 147.061 | 580.320 | 22.326 | 2.380 |

continued...

Table 3.24 continued

| Group no. | Species group | Food intake ^a | Respiration ^a | Predation mortality (production *EE) ^a | Flow to detritus ^a | Assimilation ^a | Net production (g·m ⁻² ·year ⁻¹) | Trophic level |
|-----------|-------------------------------------------------------|--------------------------|--------------------------|---------------------------------------------------|-------------------------------|---------------------------|---------------------------------------------------------|---------------|
| 40 | Polychaetes/priapuloids/ophiuroids | 2,032.80 | 1,454.640 | 157.634 | 420.526 | 1,626.240 | 171.528 | 2.350 |
| 41 | Holothuroids/sipunculids/ echiuroids/hemichordates | 222.57 | 157.519 | 18.803 | 46.245 | 178.053 | 20.527 | 2.100 |
| 42 | Bivalves | 1,037.88 | 586.672 | 222.354 | 228.849 | 830.300 | 243.542 | 2.100 |
| 43 | Ascidians/barnacles/bryozoans | 2,748.00 | 1,882.380 | 252.302 | 613.318 | 2,198.400 | 316.168 | 2.340 |
| 44 | Sponges | 4,000.00 | 2,000.000 | 299.177 | 1,700.823 | 3,200.000 | 1,201.514 | 2.000 |
| 45 | Sea anemones/corals | 1,089.00 | 739.310 | 128.803 | 220.887 | 871.200 | 131.835 | 2.340 |
| 46 | Zooplankton | 5,280.00 | 2,944.000 | 1,179.004 | 1,156.996 | 4,224.000 | 1,280.135 | 2.600 |
| 47 | Microfauna | 28,500.00 | 18,600.000 | 4,112.797 | 5,787.203 | 22,800.000 | 4,201.018 | 2.000 |
| 48 | Phytoplankton | - | - | 2,073.120 | 726.880 | - | 2,801.514 | 1.000 |
| 49 | Benthic autotrophs | - | - | 6,201.429 | 11,023.570 | - | - | 1.000 |
| 50 | Detritus, POM, DOM | - | - | - | - | - | - | 1.000 |

^a(g·m⁻²·year⁻¹ WW).

continued...

Table 3.24 continued.

| Group no. | Omnivory index | Gross efficiency | Net efficiency | Respiration/ assimilation | Production/ respiration |
|-----------|----------------|------------------|----------------|---------------------------|-------------------------|
| 1 | 0.5950 | 0.049 | 0.058 | 0.939 | 0.065 |
| 2 | 0.3031 | 0.037 | 0.044 | 0.954 | 0.049 |
| 3 | 0.3684 | 0.092 | 0.108 | 0.885 | 0.129 |
| 4 | 0.3988 | 0.128 | 0.151 | 0.840 | 0.190 |
| 5 | 0.0280 | 0.066 | 0.078 | 0.917 | 0.090 |
| 6 | 0.3076 | 0.085 | 0.100 | 0.894 | 0.118 |
| 7 | 0.2702 | 0.055 | 0.065 | 0.931 | 0.074 |
| 8 | 0.2505 | 0.123 | 0.144 | 0.846 | 0.181 |
| 9 | 1.0773 | 0.031 | 0.037 | 0.961 | 0.041 |
| 10 | 0.0000 | 0.025 | 0.030 | 0.968 | 0.033 |
| 11 | 0.0263 | 0.028 | 0.033 | 0.965 | 0.036 |
| 12 | 0.3181 | 0.097 | 0.115 | 0.878 | 0.139 |
| 13 | 0.2831 | 0.100 | 0.118 | 0.875 | 0.143 |
| 14 | 0.1704 | 0.170 | 0.200 | 0.788 | 0.270 |
| 15 | 0.0086 | 0.188 | 0.221 | 0.765 | 0.307 |
| 16 | 0.4096 | 0.065 | 0.077 | 0.918 | 0.089 |
| 17 | 0.1515 | 0.261 | 0.307 | 0.674 | 0.484 |
| 18 | 0.4604 | 0.161 | 0.189 | 0.799 | 0.252 |
| 19 | 0.0362 | 0.092 | 0.108 | 0.885 | 0.129 |
| 20 | 0.5494 | 0.098 | 0.115 | 0.878 | 0.139 |
| 21 | 0.4694 | 0.094 | 0.111 | 0.882 | 0.133 |
| 22 | 0.4761 | 0.064 | 0.075 | 0.920 | 0.087 |
| 23 | 0.0038 | 0.063 | 0.074 | 0.921 | 0.085 |
| 24 | 0.0010 | 0.056 | 0.065 | 0.930 | 0.075 |
| 25 | 0.0000 | 0.028 | 0.033 | 0.965 | 0.036 |

continued...

Table 3.24 continued.

| Group no. | Omnivory index | Gross efficiency | Net efficiency | Respiration/ assimilation | Production/ respiration |
|-----------|----------------|------------------|----------------|---------------------------|-------------------------|
| 26 | 0.0146 | 0.079 | 0.093 | 0.902 | 0.109 |
| 27 | 0.0000 | 0.180 | 0.212 | 0.775 | 0.290 |
| 28 | 0.2708 | 0.067 | 0.078 | 0.915 | 0.091 |
| 29 | 0.1887 | 0.074 | 0.087 | 0.907 | 0.102 |
| 30 | 0.4528 | 0.043 | 0.051 | 0.946 | 0.057 |
| 31 | 0.0709 | 0.281 | 0.331 | 0.649 | 0.542 |
| 32 | 0.3011 | 0.135 | 0.159 | 0.831 | 0.203 |
| 33 | 0.4187 | 0.114 | 0.135 | 0.857 | 0.167 |
| 34 | 0.4034 | 0.104 | 0.122 | 0.870 | 0.150 |
| 35 | 0.2831 | 0.040 | 0.047 | 0.950 | 0.052 |
| 36 | 0.4394 | 0.151 | 0.178 | 0.811 | 0.233 |
| 37 | 0.1756 | 0.297 | 0.350 | 0.628 | 0.592 |
| 38 | 0.3337 | 0.200 | 0.235 | 0.750 | 0.333 |
| 39 | 0.2585 | 0.031 | 0.036 | 0.962 | 0.040 |
| 40 | 0.3277 | 0.084 | 0.099 | 0.894 | 0.118 |
| 41 | 0.0900 | 0.092 | 0.109 | 0.885 | 0.130 |
| 42 | 0.0900 | 0.235 | 0.276 | 0.707 | 0.415 |
| 43 | 0.3684 | 0.115 | 0.135 | 0.856 | 0.168 |
| 44 | 0.0010 | 0.300 | 0.353 | 0.625 | 0.601 |
| 45 | 0.3684 | 0.121 | 0.142 | 0.849 | 0.178 |
| 46 | 0.2400 | 0.242 | 0.285 | 0.697 | 0.435 |
| 47 | 0.0000 | 0.147 | 0.173 | 0.816 | 0.226 |
| 48 | 0.0000 | - | - | - | - |
| 49 | 0.0000 | - | - | - | - |
| 50 | 0.3669 | - | - | - | - |

additional parameters and ratios computed by ECOPATH II for the balanced 50-box model. These outputs permit to estimate flow rates (in $\text{gm}^{-2}\text{year}^{-1}$) in and out of each box of the system.

Modifications Required to Balance the 50-Box Model

The input value of net PP had to be modified slightly, from $20,000 \text{ gm}^{-2}\text{year}^{-1}$ to $20,500 \text{ gm}^{-2}\text{year}^{-1}$. This was still well within the established range of NPP estimates ($15,000\text{-}25,000 \text{ gm}^{-2}\text{year}^{-1}$).

The strongest changes were applied to the biomass estimates since they were considered to be least accurate of all input values. The biomass of detritus was left unchanged, as well as biomass of phytoplankton, whereas that of benthic primary producers had to be lowered by 42% from $2,230$ to $1,300 \text{ gm}^{-2}$. This value was still within the range estimated for an average m^2 from five subsystems.

It was observed during the balancing process that the input biomass values of fish groups were generally too high, while those of the majority of nonfish taxa were too low. This was especially true for strongly carnivorous groups. Another aspect of the initial biomass estimates of fish groups was that values for large predators were too high, while those for small ones were too low.

The density estimates for fishes, which are generally based on visual censuses, appeared to have provided unrealistically high abundances for groups of large predatory fish. For example, the biomass of large sharks and rays had to be decreased from 96 gm^{-2} to 0.3 gm^{-2} , a change of 99.7%. Obviously, the density estimates for relatively rare, but large and wide-ranging predators depended strongly on their presence during a census. This point will be discussed in more detail below.

The majority of biomass values for sharks and rays was computed via qualitative indications of density. Since a common definition in terms of numbers does not exist for qualitative indications of abundance, it is quite possible that expressions such as "abundant" or "common" refer to a comparison between species of the same ecological group, so that, e.g., nurse sharks are abundant on the reef compared to tiger sharks but that this abundance has nothing in common with the expression "abundant" used, e.g., for doctorfish, parrotfish, etc., or even small pelagics such as dwarf herring or sardine.

Densities of small and/or cryptic reef fish species are obviously strongly underestimated by visual census techniques. Thus, for example "Blenniidae" had to be raised from 0.012 gm^{-2} to 0.6 gm^{-2} , a change of biomass of 4,900%.

The density of small schooling pelagic fish was also strongly underestimated. Thus, for example, the initial estimate of the biomass of Engraulidae had to be raised from 0.019 gm^{-2} to 3.325 gm^{-2} (17,400% change). This can be explained by the fact that only the highest density level obtained from visual censuses for reef fishes was assigned to small schooling pelagics. In all censuses in which small pelagics were explicitly included, they were stated as "too numerous to count". Still higher levels of abundance might have to be added to the ones presented in "Materials and methods" in order to properly account for densities of small pelagics.

Groups of large to intermediate herbivorous reef fish (e.g., families Scaridae or Kyphosidae) showed generally low EEs. Their input biomasses were reduced in line with reduction of biomass of 60-99% performed on groups containing similarly sized carnivorous species. An 80% reduction of biomass is the average for fishes of this size range. The biomass of large herbivorous reef fish could have been reduced further to raise their EE, but there was

no good reason to do so, except that the relatively high EE of benthic autotrophs would have slightly decreased.

The biomass values of nonfish groups had to be raised because of the diet preferences of fish groups. Conspicuous was the discrepancy between “offer” and “demand” in crustaceans, which were the most difficult groups to balance. This is because these groups feed on a relatively high trophic level and their consumption is rather high (up to 125 times their body weight per year). This makes it difficult to raise their biomass without serious implications for the whole system.

Initial biomasses of crustaceans were generally too low to satisfy the food requirements of the fish groups. Two opposite tendencies might be responsible for this discrepancy: (1) the comparably high percentages of crustaceans in the diet composition of fish groups might be an artefact, i.e., due to the high amount of undigestible exoskeletons, the percentage of crustaceans in the food of fishes might be overestimated; and (2) the biomass estimates for crustaceans are derived mainly from the Gulf of Batabanó, Cuba, from an area characterized by sand and mud. The combined biomass estimates for crustaceans from this area, of 5-10 gm⁻²WWV, do not compare well with an overall estimate of crustacean biomass of around 160 gm⁻²WWV recorded by Glynn (1973) for a Caribbean *Porites furcata* reef (see also Table 3.17).

The effect of erroneous biomass estimates on weighting factors within species groups is, on the other hand, of minor importance since these groups are composed of species with similar ecological features; the above problems concerning the input biomasses are thus valid for all species of a group.

After the biomasses, the strongest modifications were performed on the DC matrix, particularly of elasmobranchs and on invertebrates. This was not unexpected, since these parts of the matrix were most heterogenous in the first place. Raising the biomass of crustaceans was not enough to correct for the unrealistically high initial EE values for these groups. Thus, the diet of groups which fed on crustaceans was shifted, in part, to ecological groups with low EEs (sea urchins, ascidians, sponges, etc.). This implied a shift to lower trophic levels.

The diet of large carnivorous fish groups was similarly modified, often from carnivorous to herbivorous fishes. The latter showed in general low EEs, especially the groups containing large species, which also had high biomasses. For example, large parrotfish (Scaridae) with the highest output biomass for a single fish family in the 50-box model (almost 19 gm⁻²) have a very low EE (EE = 0.336). The original DC matrix of fishes did not identify large herbivorous reef fish as a preferred food item of carnivorous species. It might be an artefact from Randall's stomach analyses, but this is unlikely since it can be assumed that herbivorous fish should have been identified as reliably as carnivorous fish. This suggests that large predatory fish species tend to avoid herbivorous reef fish as prey. Parrotfish produce a skin mucus which is reported to protect them from predation while they sleep. If the low predation mortality of herbivorous reef fish is true, and not an artefact, this would mean that predator biomass would have to be lowered further to lower the EE of prey groups below 1. This would also mean that large herbivorous reef fish serve mainly as energy supply for the detrital pool. This matter will be discussed further below.

The initial P/B values, especially of fish groups, were considered to be more reliable than the initial biomass estimates; thus, the overwhelming majority of P/B outputs were kept within the initial range of input P/B estimated for the respective groups.

The consumption estimates for nonfish groups were slightly modified in a few groups only. For sea urchins and sponges, Q/B was raised in order to lower their gross efficiency below 0.4 and to lower their P/R ratio to 0.6 year⁻¹.

Output gross efficiencies (GE = production/consumption) for all boxes were generally between 0.1 and 0.5. High GEs are characteristic of small-sized and/or young animals. Low GEs are characteristic of large-sized and/or old animals. The GE of apex predators should be close to zero when they are not exploited by fisheries. GE output values from the 50-box model for fish groups were somewhat low, never exceeding 0.3. To raise these estimates, the P/B ratios could be increased and/or the Q/B reduced, thus decreasing the EEs of prey groups. These modifications would result in an increase of the biomass of a number of prey groups.

Graphical Representation of the 50-Box Model

Fig. 3.2 shows the mass-balanced 50-box model of an unexploited segment of coral reef in the PRVI area. This graph demonstrates that it is possible to balance a quantitative steady-state model of the energy flow within a coral reef system based on literature data and parameter estimations derived from multivariate statistics.

3.3.1.6. SUMMARY STATISTICS

Summary statistics computed by ECOPATH II for the 50-box model are presented in Table 3.25.

3.3.1.7. NETWORK FLOW INDICES

Network flow indices computed by ECOPATH II for the 50-box model are listed in Table 3.26.

3.3.1.8. FLOWS AND TRANSFER EFFICIENCIES

Table 3.27 presents flows and transfer efficiencies for discrete trophic levels computed by ECOPATH II for the 50-box model.

3.3.1.9. MIXED TROPHIC IMPACTS

Fig. 3.3 shows the direct and indirect impacts an increase in the biomass of the groups to the left of the histograms would have on the groups positioned above them. The bars pointing upwards show positive impacts while those pointing downwards, negative impacts. The impacts are relative, not absolute ones but are comparable between histograms. Note that the routine assumes the food matrix to remain stable, which would not happen in the long term (but may be reasonable for the short term). The mixed trophic impacts can also be viewed as a simple sensitivity analysis *sensu* Majkowski (1982).

A biomass increase of benthic autotrophs (group 49) has a direct positive effect on all groups feeding on benthic producers. The impact is greatest for groups which feed almost exclusively on benthic autotrophs, such as Kyphosidae (group 10), intermediate herbivorous reef fish (11), parrotfish (family Scaridae, groups 23, 24, 25), herbivorous blennies (26), echinoids (37) and corals (45). The impact on detritus is also positive since a larger amount of biomass would be directed towards this box.

An increase in phytoplankton biomass would have a strong direct positive impact on engraulids (16) since they feed largely on phytoplankton. It has a moderate direct positive

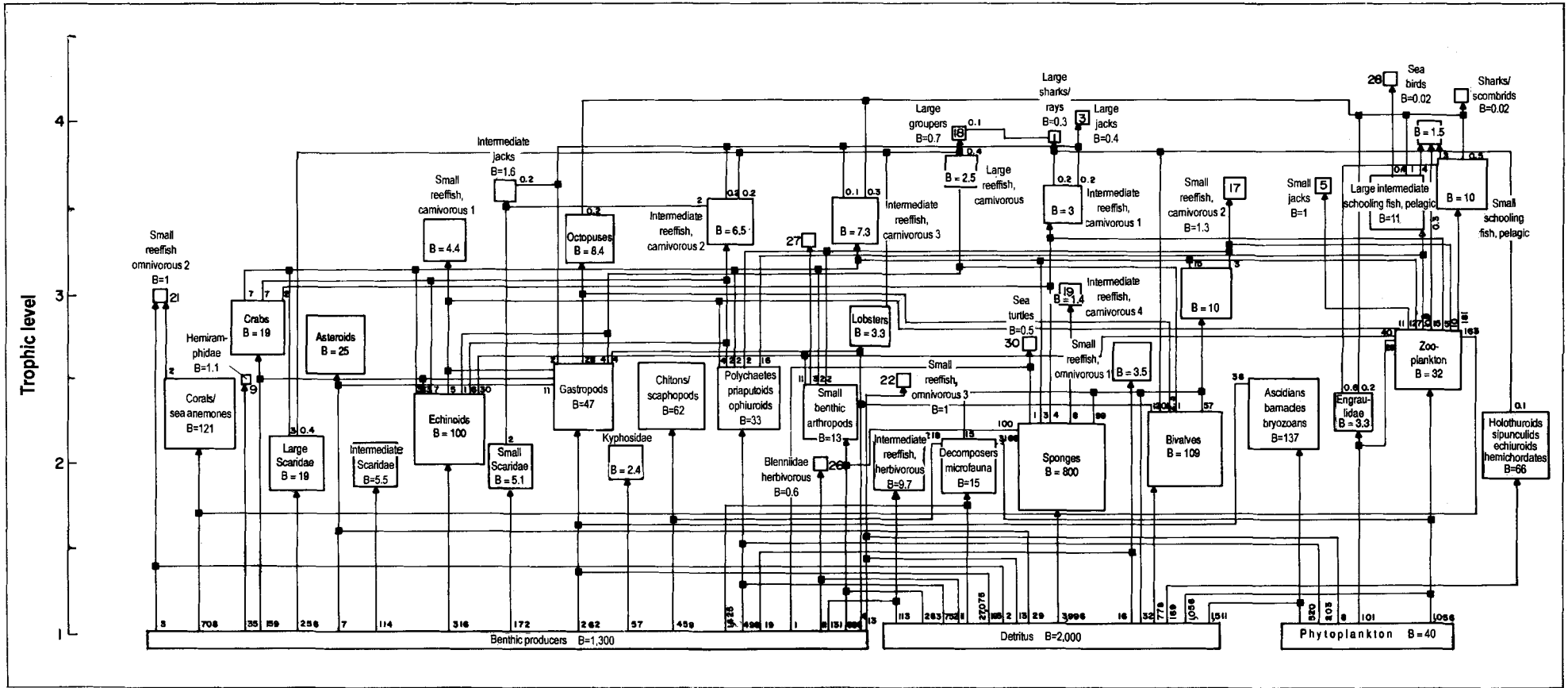


Fig. 3.2. Graphical representation of the 50-box model of a Caribbean coral reef. The area of each box is proportional to the logarithm of the biomass (B, in gm⁻²WW) of each group. Values represent the average for all subsystems in Fig. 2.1. Flows are in gm⁻²year⁻¹ WW. Incoming flows (bottom half of a box) represent at least 80% of the diet of a group.

Table 3.25. Summary statistics computed by ECOPATH II for a 50-box model of a Caribbean coral reef. PP = primary production.

| | | | |
|--------------------------------|---|---------|------------------------------------|
| Sum of all production | = | 51,282 | gm ² year ⁻¹ |
| Sum of all imports* | = | 475 | gm ² year ⁻¹ |
| Sum of all respiratory flows | = | 32,523 | gm ² year ⁻¹ |
| Sum of all flows into detritus | = | 23,193 | gm ² year ⁻¹ |
| <hr/> | | | |
| Total system throughput | = | 107,473 | gm ² year ⁻¹ |
| Throughput cycled | = | 19,686 | gm ² year ⁻¹ |
| Finn's cycling index | = | 18.3% | of throughput |
| <hr/> | | | |
| Input total net PP | = | 20,500 | gm ² year ⁻¹ |
| Calculated total net PP | = | 20,025 | gm ² year ⁻¹ |

*Difference between input total net PP and calculated total net PP.

Table 3.26. Network flow indices computed by ECOPATH II for a 50-box model of a Caribbean coral reef.

| Source | Ascendency (flowbits) | % | Overhead (flowbits) | % | Capacity (flowbits) | % |
|--------------|--------------------------|------|------------------------|------|------------------------|-------|
| Inputs | 952 | 0.2 | 2,678 | 0.6 | 3,630 | 0.8 |
| Internal | 94,210 | 21.5 | 226,071 | 51.6 | 320,281 | 73.1 |
| Dissipations | 52,362 | 12.0 | 61,940 | 14.1 | 114,302 | 26.1 |
| Totals | 147,524 | 33.7 | 290,689 | 66.3 | 438,213 | 100.0 |

Table 3.27. Flows and transfer efficiencies for discrete trophic levels computed by ECOPATH II for a 50-box model of a Caribbean coral reef.

| Trophic level | Flows originating from the primary producers | | | | | Flow to detritus | | |
|--------------------------------------------|----------------------------------------------|---------|-------------|------------|------|------------------|------|------|
| | Import | Export | Respiration | Throughput | | | | |
| I | 0.000 | 480.000 | 0.000 | 0.000 | | 11,750.450 | | |
| II | 0.000 | 0.000 | 5,252.860 | 7,794.549 | | 1,665.249 | | |
| III | 0.000 | 0.000 | 578.620 | 876.441 | | 188.912 | | |
| IV | 0.000 | 0.000 | 74.367 | 108.909 | | 23.422 | | |
| V | 0.000 | 0.000 | 7.595 | 11.120 | | 2.393 | | |
| VI | 0.000 | 0.000 | 0.763 | 1.133 | | 0.246 | | |
| VII | 0.000 | 0.000 | 0.083 | 0.124 | | 0.027 | | |
| VIII | 0.000 | 0.000 | 0.009 | 0.013 | | 0.003 | | |
| IX | 0.000 | 0.000 | 0.001 | 0.001 | | 0.000 | | |
| Trophic level | Flows originating from the detritus | | | | | Flow to detritus | | |
| | Import | Export | Respiration | Throughput | | | | |
| I | 475.000 | 0.000 | 0.000 | 23,668.000 | | 0.000 | | |
| II | 0.000 | 0.000 | 22,975.980 | 36,171.262 | | 8,299.198 | | |
| III | 0.000 | 0.000 | 2,985.809 | 4,896.084 | | 1,060.269 | | |
| IV | 0.000 | 0.000 | 582.137 | 850.005 | | 183.624 | | |
| V | 0.000 | 0.000 | 58.936 | 84.245 | | 17.906 | | |
| VI | 0.000 | 0.000 | 4.997 | 7.404 | | 1.599 | | |
| VII | 0.000 | 0.000 | 0.543 | 0.808 | | 0.176 | | |
| VIII | 0.000 | 0.000 | 0.060 | 0.089 | | 0.020 | | |
| IX | 0.000 | 0.000 | 0.007 | 0.010 | | 0.002 | | |
| Transfer efficiencies (%) by trophic level | | | | | | | | |
| Source | I | II | III | IV | V | VI | VII | VIII |
| Producers | - | 11.2 | 12.4 | 10.2 | 10.2 | 10.9 | 10.8 | - |
| Detritus | - | 13.5 | 17.4 | 9.9 | 8.8 | 10.9 | 11.1 | - |
| All flows | - | 13.1 | 16.6 | 9.9 | 9.0 | 10.9 | 11.0 | - |

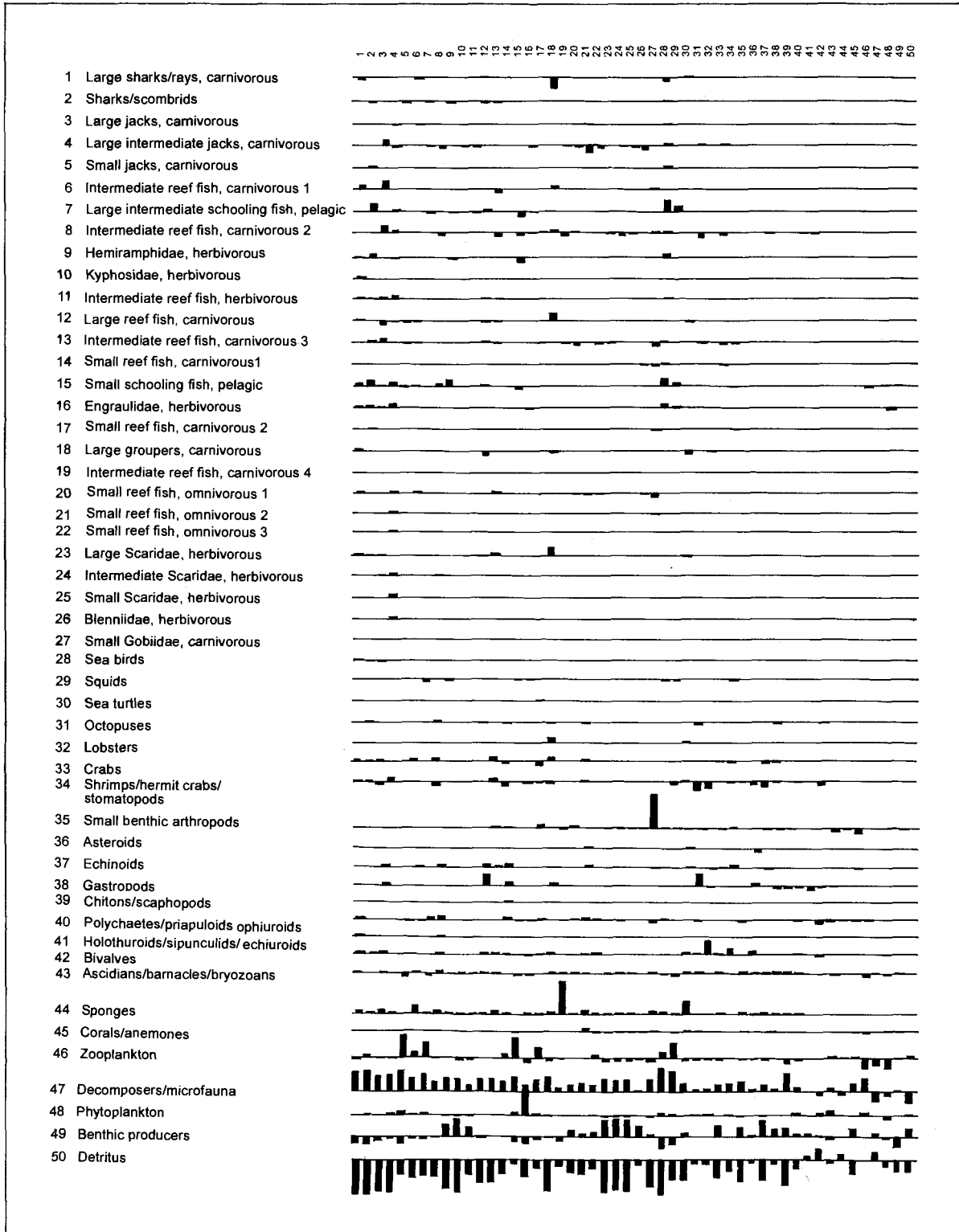


Fig. 3.3. Mixed trophic impacts of a 50-box model of a Caribbean coral reef showing the direct and indirect impacts an increase in the biomass of the groups to the left of the histograms will have on the groups positioned above the histograms.

impact on bivalves (42), ascidians, barnacles and bryozoans (43) and zooplankton (46) since these groups feed to some extent on phytoplankton. A slight indirect positive impact is observed for groups feeding on groups which feed on phytoplankton, such as sharks and scombrids, which feed on phytoplanktivorous engraulids.

An increase in biomass of decomposers/microfauna (group 47) has a large indirect positive impact on almost all other groups (recall the analogy with the system's motor), except for a small indirect negative impact on groups, which feed to a large extent on detritus. These groups may be regarded as competitors for the same food source since group 47 feeds almost exclusively on detritus. Therefore an increase of decomposers/microfauna biomass would have a direct negative impact on the detritus box.

Increasing the biomass of the apex predators (birds, sharks, scombrids, large jacks and groupers) would have a relatively small negative impact on their prey groups. This is due to the small biomass of these groups combined with moderate relative food requirements. An increase in the biomass of apex predators would have a slight direct positive effect on groups which themselves feed, albeit, to a small extent on these predators, such as the seabirds.

Overall, an increase in biomass of a given group has a direct negative impact on its prey groups and an indirect negative impact on groups competing for the same food source. Further, such increase has a direct positive impact on the predators of that group and an indirect positive impact on groups feeding, in turn, on those predators.

3.3.2. Comparison of Models Based on Two Methods of Aggregation

3.3.2.1. FURTHER AGGREGATION OF FISH GROUPS BY CLUSTER ANALYSIS

The results of the cluster analysis used to pass from 29 to 1 fish species group are depicted in Fig. 3.1.

The strong separation of herbivores and carnivores is maintained up to a cutoff level of 80% (QED 1.2). The strongest separation between large and fast-swimming apex predators and the vast majority of reef fish is maintained up to a cutoff level of 99.9% (QED 1.49). The reef fish fauna is thus characterized by three principal energetic groups: (1) the large and fast-swimming apex predators with a predominantly pelagic habitat and wide-ranging migrations (exceptions are not well identified due to set values used here for the aspect ratios of sharks and rays; some of them, especially rays and the nurse shark, probably belong to the large reef fishes); (2) the predominantly carnivorous; and (3) the predominantly herbivorous reef fishes. (The omnivores were pooled with either group depending on the percentage of plant or animal food in their diet).

3.3.2.2. AGGREGATION BY THE ECOPATH II AGGREGATION ROUTINE

Fig. 3.4 shows the results of the ECOPATH II aggregation routine which allowed a stepwise reduction of the number of boxes from 50 to 1. Species groups with a similar trophic level were aggregated first as in the case of the strictly herbivorous fish groups and the strictly carnivorous groups which belong largely to the apex predators; within each trophic set, groups with a low biomass were aggregated first in order to reduce ascendancy as little as possible. It may also be observed from Fig. 3.4 that the aggregation routine tends to form two main trophic groups by combining groups with a similar trophic level: one carnivorous and one herbivorous. This tendency becomes more obvious when the number of boxes is further reduced. Due to their

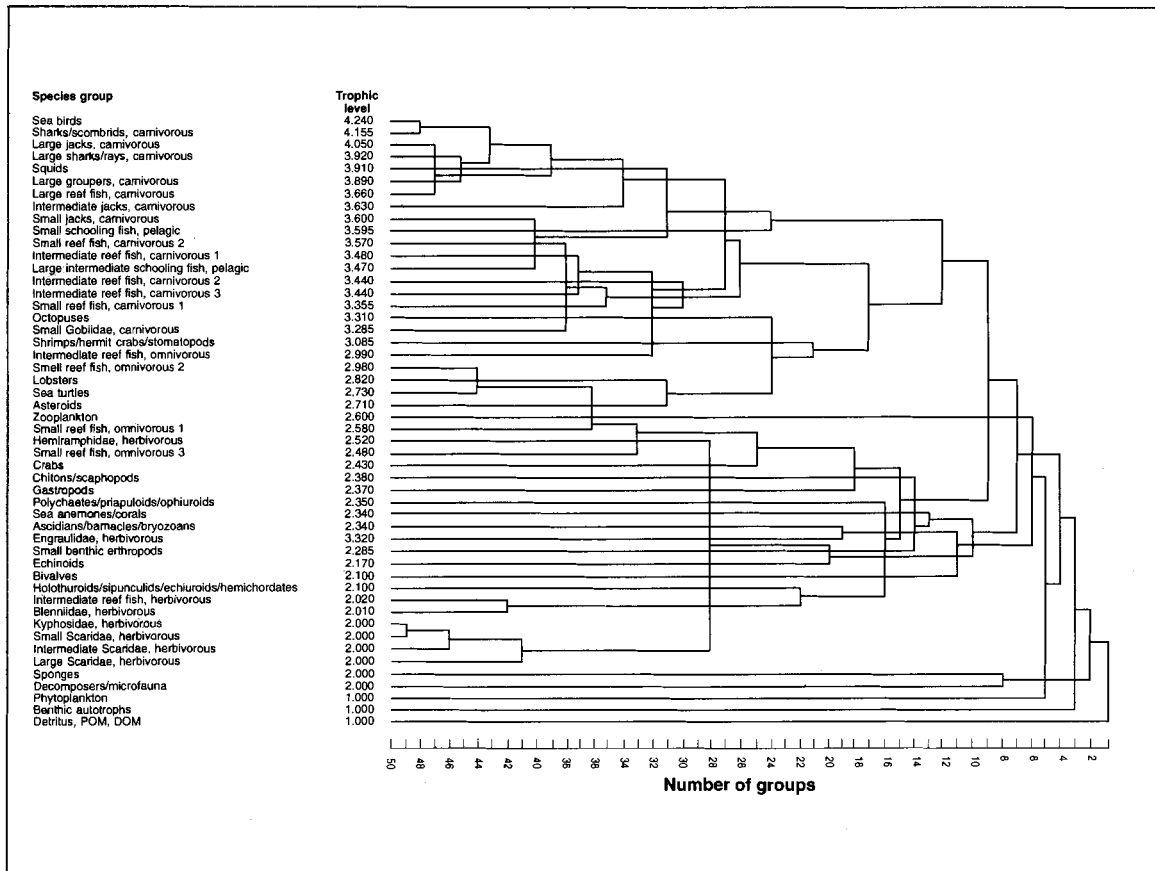


Fig. 3.4. Graphical representation of the results of the ECOPATH II aggregation routine showing the stepwise reduction of the number of boxes from 50 to 1 for models of a Caribbean coral reef.

high biomass and/or degree of detritivory, sponges and microfauna are kept as single groups until the system is reduced to 8 boxes.

Benthic producers and detritus are also kept separate until the system is reduced to a minimum number of boxes (5, 3 and 1). This dendrogram cannot be compared directly with the results of the cluster analysis, since the latter was applied exclusively to fish species. Nevertheless, it should be noticed that both figures show a separation into main trophic groups (recall the separation into herbivorous and carnivorous reef fish groups which was maintained up to a cutoff level of 80% in Fig. 3.1). The separation into large apex predators with a mainly pelagic habitat and reef fish with a demersal habitat was not repeated by the ECOPATH II aggregation routine.

3.3.2.3. THE 20-BOX MODELS

Method 1: Cluster Analysis + Intuition

COMPOSITION OF GROUPS. Table 3.28 shows the composition of groups for the first of the two 20-box models. As pointed out in "Materials and methods", the group composition resulted from an aggregation of fish species by cluster analysis and "intuitive guesswork" for the nonfish groups based on the criteria relative to DC, body size and lifestyle.

In contrast to the 50-box model, some of the boxes of the 20-box model include both fish and nonfish groups: groups are now classified by a combination of diet, size and lifestyle, leaving apart the separation into fish and nonfish. This is a first step away from taxonomics, toward a purely energetic approach (within the fish community, steps in this direction were taken by combining species and groups with similar energy requirements, also irrespective of taxonomics).

PARAMETER ESTIMATION. The condensed input values for the key parameters and the inputs for the DC matrix may be found in Tables 3.29 and 3.30, respectively. Output values are the same as input values with one exception referring to box 8, small benthic carnivores. In the 50-box model, due to their low biomass (1.3 gm^{-2}), the sea anemones were merged with the corals, even though they are fully heterotrophic (feeding mainly on zooplankton). Within the 20-box model, they were merged with small benthic carnivores. Their excessive predation mortality was shifted to benthic autotrophs; this was unproblematic since the predators of sea anemones include omnivorous and herbivorous groups. Therefore, the EE of benthic autotrophs (box 19) increased from 0.360 to 0.362, while the EE of "corals" decreased from 0.977 in the 50-box model to 0.713 in the 20-box model.

Table 3.31 presents additional parameters and ratios computed by ECOPATH II. These parameter estimates allow, as already mentioned for the 50-box model, computation of the energy balance, following the "master equation", of each box of the system.

Fig. 3.5 presents the 20-box model of an unexploited segment of the PRVI coral reefs system, aggregated following method 1 (see above).

SYSTEM PROPERTIES. System properties, such as summary statistics, network flow indices, flows and transfer efficiencies and mixed trophic impacts, were estimated by ECOPATH II for the 20-box model as obtained by method 1. The results showed only negligible differences in relation to the 50-box model. Summary statistics, network flow indices and flows for the 20-box model, obtained by method 1, are listed in Appendix Tables 8.9.1 to 8.9.3. Table 3.32 presents the transfer efficiencies between discrete trophic levels.

Table 3.28. Composition of groups of a 20-box model of a Caribbean coral reef obtained by method 1 (cluster analysis + intuition); names of subgroups refer to original groups in the 50-box model.

| Group no. | Predator/prey group | Group no. | Predator/prey group |
|-----------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------|
| 1 | B Sea birds | 9 | Benthic invertebrates, omnivorous |
| 2 | Sharks/rays/scombrids | I I | Gastropods + Polychaetes/priapuloids/ ophiuroids |
| | F Large sharks/rays, carnivorous + F Sharks/scombrids, carnivorous | 10 | Large to intermediate reef fish, herbivorous (>50% plant food) |
| 3 | Large to intermediate jacks, carnivorous | F F F | F Kyphosidae, herbivorous + Intermediate reef fish, herbivorous + Large Scaridae, herbivorous + Intermediate Scaridae, herbivorous |
| | F Large jacks, carnivorous + F Intermediate jacks, carnivorous | | |
| 4 | Large to intermediate schooling pelagics | 11 | Small reef fish, herbivorous (>50% plant food) |
| | F Small jacks, carnivorous + F* Large to intermediate schooling fish, pelagic + F Hemiramphidae, herbivorous + I Squids | F F F | F Small reef fish, omnivorous 3 + Small Scaridae, herbivorous + Blenniidae, herbivorous |
| 5 | Small schooling fish, pelagic | 12 | Benthic invertebrates, herbivorous (>50% plant food) |
| | F Small schooling fish, pelagic + F Engraulidae, herbivorous | I I I I | Crabs Amphipods/isopods/ tanaids/pycnogonids + Echinoids + Chitons/scaphopods |
| 6 | Large groupers, carnivorous | 13 | Sessile detritivores (>50% detrital food) |
| 7 | Large to intermediate benthic carnivores (>50% live animal food) | I I | Holothuroids/sipunculids/ echiuroids/ hemichordates + Bivalves |
| | F Intermediate reef fish, carnivorous 1 + F Large reef fish, carnivorous + F Intermediate reef fish, carnivorous 2 + F Intermediate reef fish, carnivorous 3 + F Intermediate reef fish, carnivorous 4 + R Sea turtles + F Octopuses + F Lobsters | 14 | Sessile detritivores (>50% detrital food) |
| 8 | Small benthic carnivores (> 50 % live animal food) | I I | Ascidians/barnacles/bryozoans + Sponges |
| | F Small reef fish, carnivorous 1 + F Small reef fish, carnivorous 2 + F Small reef fish, omnivorous 1 + F Small reef fish, omnivorous 2 + F Small Gobiidae, carnivorous + I Shrimps/hermit crabs/stomatopods + I Asteroids + I Sea anemones | 15 16 17 18 19 20 | Corals Zooplankton Decomposers/microfauna Phytoplankton Benthic autotrophs Detritus |

B = bird, F = fish, I = invertebrate, R = reptile.

Table 3.29. Results of the ECOPATH II parameter estimation for a 20-box model of a Caribbean coral reef (method 1: cluster analysis + intuition; average values are weighted by biomass); P/B = production/biomass ratio, Q/B = consumption/unit biomass, EE = ecotrophic efficiency (part of production consumed by predators).

| Group no. | Predator group | No. of groups ^a | Biomass (g·m ⁻²) | Average P/B (year ⁻¹) | Average Q/B (year ⁻¹) | End EE |
|-----------|------------------------------------------------------------------|----------------------------|------------------------------|-----------------------------------|-----------------------------------|------------------|
| 1 | Sea birds | 1 | 0.017 | 5.400 | 80.000 | 0.344 |
| 2 | Sharks/rays/scombrids, carnivorous | 2 | 0.714 | 0.298 | 7.364 | 0.527 |
| 3 | Large to intermediate jacks, carnivorous | 2 | 1.811 | 1.106 | 8.805 | 0.355 |
| 4 | Large to intermediate schooling pelagics | 4 | 14.797 | 0.795 | 14.954 | 0.923 |
| 5 | Small schooling pelagic fish | 2 | 13.471 | 3.366 | 24.910 | 0.789 |
| 6 | Large groupers, carnivorous | 1 | 0.725 | 0.370 | 2.300 | 0.51 |
| 7 | Large to intermediate benthic carnivores (>50% live animal food) | 8 | 32.893 | 1.084 | 7.149 | 0.831 |
| 8 | Small benthic carnivores (>50% live animal food) | 8 | 46.718 | 1.320 | 10.722 | 0.942 (1.423) |
| 9 | Benthic invertebrates, omnivorous | 2 | 79.800 | 3.792 | 33.684 | 0.937 |
| 10 | Large to intermediate reef fish, herbivorous (>50% plant food) | 4 | 36.563 | 0.843 | 18.468 | 0.475 |
| 11 | Small reef fish, herbivorous (>50% plant food) | 3 | 6.635 | 1.342 | 34.938 | 0.788 |
| 12 | Benthic invertebrates, herbivorous (>50% plant food) | 4 | 194.250 | 1.179 | 15.569 | 0.798 |
| 13 | Semisessile detritivores (>50% detrital food) | 2 | 175.490 | 1.505 | 7.182 | 0.909 |
| 14 | Sessile detritivores (>50% detrital food) | 2 | 937.400 | 1.617 | 7.199 | 0.364 |
| 15 | Corals | 1 | 120.000 | 1.090 | 9.000 | 0.713 |
| 16 | Zooplankton | 1 | 32.000 | 40.000 | 165.000 | 0.928 |
| 17 | Decomposers/microfauna | 1 | 15.000 | 280.000 | 1,900.000 | 0.979 |
| 18 | Phytoplankton | 1 | 40.000 | 70.000 | - | 0.74 |
| 19 | Benthic autotrophs | 1 | 1,300.000 | 13.250 | - | 0.362 (0.360) |
| 20 | Detritus, POM, DOM | 1 | 2,000.000 | - | - | - |

^a Number of groups refers to original groups from the 50-box model.

Table 3.30. Diet composition matrix for a 20-box model of a Caribbean coral reef (method 1: cluster analysis + intuition). Values in parentheses refer to original values which had to be modified to balance the model.

| Group no. | Predator group | No. of groups ^a | Prey group | | | | | | | | | | |
|-----------|------------------------------------------------------------------|----------------------------|------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| | | | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 |
| 1 | Sea birds | 1 | - | 0.02 | 0.07 | 0.41 | 0.3 | - | 0.09 | 0.041 | - | 0.017 | 0.002 |
| 2 | Sharks/rays/scombrids, carnivorous | 2 | 0.006 | 0.013 | 0.039 | 0.258 | 0.148 | 0.026 | 0.21 | 0.105 | 0.027 | 0.09 | 0.003 |
| 3 | Large to intermediate jacks, carnivorous | 2 | - | - | 0.01 | 0.056 | 0.203 | - | 0.14 | 0.13 | 0.014 | 0.2 | 0.212 |
| 4 | Large to intermediate schooling pelagics | 4 | - | - | - | 0.023 | 0.093 | - | - | 0.014 | 0.075 | 0.007 | - |
| 5 | Small schooling pelagic fish | 2 | - | - | - | - | 0.002 | - | - | - | - | - | - |
| 6 | Large groupers, carnivorous | 1 | - | 0.01 | 0.01 | 0.01 | - | - | 0.612 | - | - | 0.288 | - |
| 7 | Large to intermediate benthic carnivores (>50% live animal food) | 8 | - | - | 0.001 | 0.004 | 0.028 | - | 0.058 | 0.05 | 0.234 | 0.038 | 0.009 |
| 8 | Small benthic carnivores (>50% live animal food) | 8 | - | - | - | 0.004 | 0.007 | - | 0.023 | 0.04 | 0.135 | - | 0.003 |
| 9 | Benthic invertebrates, omnivorous | 2 | - | - | - | - | - | - | - | 0.004 | 0.047 | - | - |
| 10 | Large to intermediate reef fish, herbivorous (>50% plant food) | 4 | - | - | - | - | - | - | - | 0.001 | 0.001 | - | - |
| 11 | Small reef fish, herbivorous (>50% plant food) | 3 | - | - | - | - | - | - | - | - | 0.008 | - | - |
| 12 | Benthic invertebrates, herbivorous (>50% plant food) | 4 | - | - | - | - | - | - | - | 0.003 | 0.005 | - | - |
| 13 | Semisessile detritivores (>50% detrital food) | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 14 | Sessile detritivores (>50% detrital food) | 2 | - | - | - | - | - | - | - | - | - | - | - |
| 15 | Corals | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 16 | Zooplankton | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 17 | Microfauna | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 18 | Phytoplankton | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 19 | Benthic autotrophs | 1 | - | - | - | - | - | - | - | - | - | - | - |
| 20 | Detritus | 1 | - | - | - | - | - | - | - | - | - | - | - |

^aNumber of groups refers to original groups from the 50-box model.

continued...

Table 3.30 continued

| Group no. | Prey group 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | Sum |
|-----------|---------------|-------|-------|-------|-------|-------|-------|------------------|-------|-----|
| 1 | - | - | - | - | 0.05 | - | - | - | - | 1 |
| 2 | 0.031 | 0.038 | - | - | - | - | - | - | 0.006 | 1 |
| 3 | 0.01 | 0.004 | - | - | 0.021 | - | - | - | - | 1 |
| 4 | - | - | - | - | 0.574 | - | - | 0.214 | - | 1 |
| 5 | 0.006 | - | 0.017 | - | 0.628 | 0.003 | 0.301 | - | 0.043 | 1 |
| 6 | 0.07 | - | - | - | - | - | - | - | - | 1 |
| 7 | 0.15 | 0.198 | 0.116 | 0.002 | 0.057 | - | - | 0.043 | 0.012 | 1 |
| 8 | 0.15 | 0.151 | 0.014 | 0.002 | 0.153 | 0.038 | - | (0.038) 0.119 | 0.161 | 1 |
| 9 | 0.018 | 0.036 | 0.092 | 0.006 | 0.038 | 0.045 | 0.076 | (0.109) 0.286 | 0.352 | 1 |
| 10 | - | - | 0.001 | - | 0.001 | - | - | (0.284) 0.828 | 0.168 | 1 |
| 11 | 0.001 | 0.012 | - | - | 0.033 | - | - | 0.844 | 0.102 | 1 |
| 12 | 0.007 | 0.006 | 0.087 | 0.025 | 0.025 | 0.097 | 0.002 | 0.645 | 0.098 | 1 |
| 13 | - | - | - | - | - | 0.1 | 0.125 | (0.639) 0.023 | 0.752 | 1 |
| 14 | - | - | - | - | 0.061 | 0.041 | 0.081 | - | 0.817 | 1 |
| 15 | - | - | - | - | 0.15 | 0.1 | - | 0.65 | 0.1 | 1 |
| 16 | - | - | - | - | - | 0.6 | 0.2 | - | 0.2 | 1 |
| 17 | - | - | - | - | - | - | - | 0.05 | 0.95 | 1 |
| 18 | - | - | - | - | - | - | - | - | (1) | (1) |
| 19 | - | - | - | - | - | - | - | - | (1) | (1) |
| 20 | - | - | - | - | - | - | - | - | - | 0 |

Table 3.31. Results of the ECOPATH II parameter estimation for a 20-box model of a Caribbean coral reef, method 1 (cluster analysis + intuition). Additional parameters. Import to detritus: 475 g·m⁻², other imports to boxes are 0; catches for all boxes are 0. Excretion = 5% and egestion = 15% (nonassimilated food, default values) of the food intake of each box.

| Group no. | Predator group | Food intake ^a | Respiration ^a | Predation mortality (production *EE) ^a | Flow to detritus ^a | Assimilation ^a | Trophic level |
|-----------|------------------------------------------------------------------|--------------------------|--------------------------|---------------------------------------------------|-------------------------------|---------------------------|---------------|
| 1 | Sea birds | 1.36 | 0.996 | 0.032 | 0.332 | 1.088 | 4.189 |
| 2 | Sharks/rays/scombrids, carnivorous | 5.26 | 3.994 | 0.112 | 1.152 | 4.206 | 4.038 |
| 3 | Large to intermediate jacks, carnivorous | 15.95 | 10.754 | 0.712 | 4.481 | 12.757 | 3.652 |
| 4 | Large to intermediate schooling pelagics | 221.27 | 165.256 | 10.857 | 45.161 | 177.019 | 3.297 |
| 5 | Small schooling pelagic fish | 335.56 | 223.107 | 35.763 | 76.692 | 268.450 | 3.039 |
| 6 | Large groupers, carnivorous | 1.67 | 1.066 | 0.137 | 0.465 | 1.334 | 3.872 |
| 7 | Large to intermediate benthic carnivores (>50% live animal food) | 235.15 | 152.466 | 29.639 | 53.047 | 188.122 | 3.306 |
| 8 | Small benthic carnivores (>50% live animal food) | 500.91 | 339.061 | 58.073 | 103.777 | 400.728 | 3.005 |
| 9 | Benthic invertebrates, omnivorous | 2,687.98 | 1,847.785 | 283.596 | 556.603 | 2,150.386 | 2.353 |
| 10 | Large to intermediate reef fish, herbivorous (>50% plant food) | 675.25 | 509.374 | 14.650 | 151.221 | 540.196 | 2.006 |
| 11 | Small reef fish, herbivorous (>50% plant food) | 231.81 | 176.547 | 7.018 | 48.249 | 185.451 | 2.078 |
| 12 | Benthic invertebrates, herbivorous (>50% plant food) | 3,024.28 | 2,190.402 | 182.647 | 651.229 | 2,419.423 | 2.298 |
| 13 | Semisessile detritivores (>50% detrital food) | 1,260.37 | 744.183 | 240.156 | 276.030 | 1,008.295 | 2.100 |
| 14 | Sessile detritivores (>50% detrital food) | 6,748.34 | 3,882.898 | 551.077 | 2,314.367 | 5,398.674 | 2.139 |
| 15 | Corals | 1,080.00 | 733.200 | 93.207 | 253.593 | 864.000 | 2.340 |
| 16 | Zooplankton | 5,280.00 | 2,944.000 | 1,187.915 | 1,148.085 | 4,224.000 | 2.600 |
| 17 | Microfauna | 28,500.00 | 18,600.000 | 4,113.075 | 5,786.925 | 22,800.000 | 2.000 |
| 18 | Phytoplankton | - | 0.000 | 2,071.501 | 728.498 | - | 1.000 |
| 19 | Benthic autotrophs | - | 0.000 | 6,227.238 | 10,997.762 | - | 1.000 |
| 20 | Detritus, POM, DOM | - | 0.000 | - | - | - | 1.000 |

^a(g·m⁻²·year⁻¹ WW).

continued....

Table 3.31 continued

| Group no. | Omnivory index | Gross efficiency | Net efficiency | Respiration/assimilation | Production/respiration |
|-----------|----------------|------------------|----------------|--------------------------|------------------------|
| 1 | 0.0872 | 0.068 | 0.085 | 0.915 | 0.091 |
| 2 | 0.2729 | 0.040 | 0.051 | 0.949 | 0.053 |
| 3 | 0.2968 | 0.126 | 0.157 | 0.843 | 0.186 |
| 4 | 0.4948 | 0.053 | 0.066 | 0.934 | 0.071 |
| 5 | 0.5716 | 0.135 | 0.169 | 0.831 | 0.203 |
| 6 | 0.3758 | 0.161 | 0.201 | 0.799 | 0.252 |
| 7 | 0.2180 | 0.152 | 0.190 | 0.810 | 0.234 |
| 8 | 0.4612 | 0.123 | 0.154 | 0.846 | 0.182 |
| 9 | 0.3236 | 0.113 | 0.141 | 0.859 | 0.164 |
| 10 | 0.0097 | 0.046 | 0.057 | 0.943 | 0.061 |
| 11 | 0.1092 | 0.038 | 0.048 | 0.952 | 0.050 |
| 12 | 0.2701 | 0.076 | 0.095 | 0.905 | 0.105 |
| 13 | 0.0900 | 0.210 | 0.262 | 0.738 | 0.355 |
| 14 | 0.1780 | 0.225 | 0.280 | 0.719 | 0.390 |
| 15 | 0.3684 | 0.121 | 0.151 | 0.849 | 0.178 |
| 16 | 0.2400 | 0.242 | 0.303 | 0.697 | 0.435 |
| 17 | 0.0000 | 0.147 | 0.184 | 0.816 | 0.226 |
| 18 | 0.0000 | - | - | - | - |
| 19 | 0.0000 | - | - | - | - |
| 20 | 0.3667 | - | - | - | - |

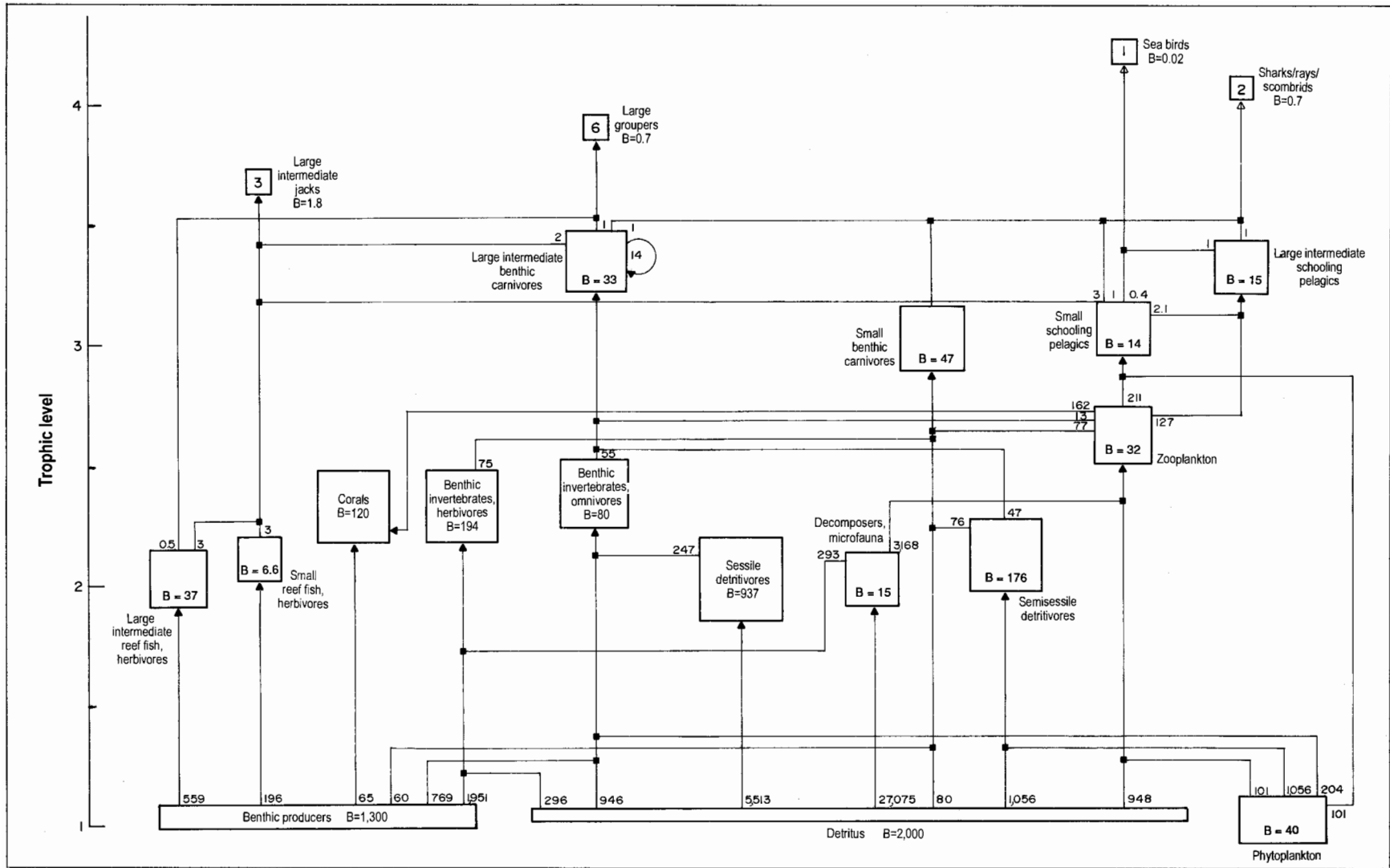


Fig. 3.5. Graphical representation of a 20-box model of a Caribbean coral reef as obtained by method 1 (cluster analysis + intuition). The area of each box is proportional to the logarithm of the biomass (B , in $\text{gm}^2 \text{WW}$) of each group. Values represent the average for all subsystems in Fig. 2.1. Flows are in $\text{gm}^2 \text{year}^{-1} \text{WW}$. Incoming flows (entering bottom half of a box) represent at least 90% of the diet of a group.

Table 3.32. Transfer efficiencies (%) by discrete trophic level for a 20-box model (method 1).

| Source | I | II | III | IV | V | VI | VII | VIII |
|-----------|---|------|------|-----|-----|------|------|------|
| Producers | - | 10.8 | 11.8 | 9.3 | 9.6 | 10.1 | - | |
| Detritus | - | 13.6 | 17.4 | 8.9 | 8.8 | 10.1 | 10.1 | - |
| All flows | - | 13.1 | 16.6 | 8.9 | 8.9 | 10.1 | 11.1 | - |

Transfer efficiencies by discrete trophic levels for primary producers were somewhat lower (4% to 9%) than in the 50-box model. For detritus, transfer efficiencies were lower for levels IV, VI and VII. Throughput per trophic level for detritus is around 7-10 times that for primary producers. Thus, resulting transfer efficiencies for combined flows were mainly affected by efficiencies computed for detritus. The differences in transfer efficiencies for models with a different number of boxes may be because the efficiencies are computed after the flows involved in cycles are removed, and there are more flows involved in cycles after aggregating formerly distinct boxes. Since detritus is involved in more cycles than any other food source, the differences in transfer efficiencies for detritus must also be more pronounced when comparing models with a different number of boxes.

Trophic level VII for primary producers had, in contrast to the 50-box model, no estimate of transfer efficiency. This may be explained by the effect of aggregating groups of differing food types, where the fraction of plant food in the resulting groups was reduced to nearly zero due to weighting of DC by the biomass and consumption requirements of the groups included.

MIXED TROPHIC IMPACTS. Fig. 3.6 shows the direct and indirect impacts on the biomass of prey groups by an increased biomass of the predator group for the 20-box model obtained by method 1. Relationships and impacts between groups correspond basically to those discussed for the 50-box model. The figure was included here for a better visualization of impacts, when the system is reduced to 20 boxes and for comparison with the 20-box model obtained by method 2 (see below).

Method 2: ECOPATH II - Aggregation Routine

COMPOSITION OF GROUPS. Table 3.33 shows the group composition of the second of the two 20-box models. The group composition is the result of a combination of groups from the balanced 50-box model by the ECOPATH II aggregation routine. In contrast to method 1, the combination of groups is based on a single criterion, the least reduction (by pairing boxes) of average shared information (i.e., ascendancy). Interestingly, this routine tends to group boxes of similar trophic levels, i.e., the aggregated boxes in the reduced models consist of carnivorous, omnivorous and herbivorous groups without reference criteria, such as size. Therefore, a large carnivorous fish group is produced which includes species ranging from tiger shark to gobies (Group 1, aggregation 26).

Also, because a trophic level of 1 is assigned to autotrophs as well as to detritus (see Introduction), nonfish taxa or groups feeding to some or large extent on detritus are combined with groups feeding on benthic autotrophs (group 4, aggregation 22).

The omnivorous group 6 (aggregation 25) includes fish and nonfish groups with a wide spectrum of omnivory. The groups with similar trophic levels are more likely to be aggregated when their biomass values are low because the associated flows are less, and hence have little impact on ascendancy. This might also be the reason why so many nonfish groups from the 50-

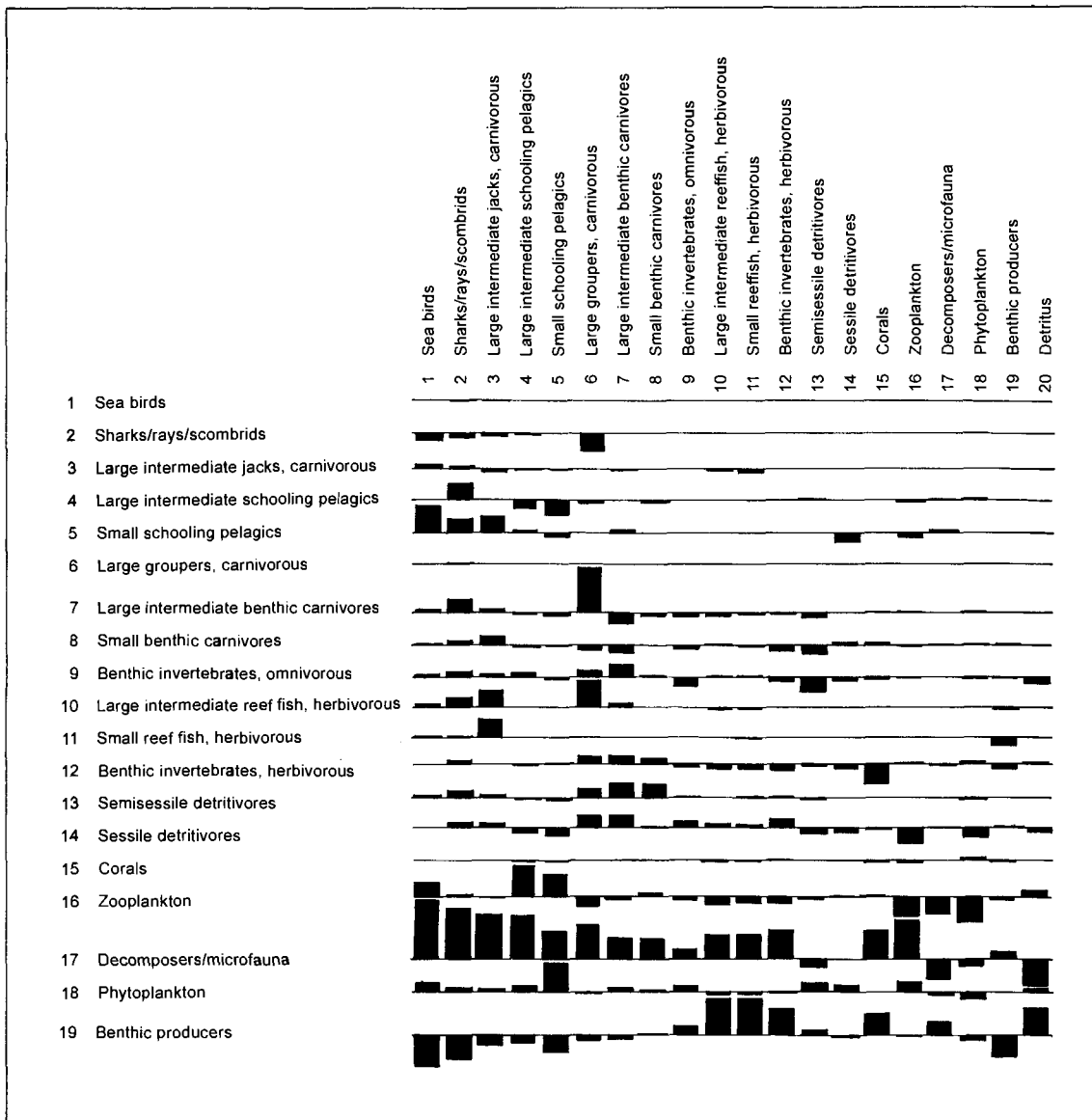


Fig. 3.6. Mixed trophic impacts of a 20-box model (method 1) of a Caribbean coral reef showing the direct and indirect impacts an increase in the biomass of the groups to the left of the histograms will have on the groups above the histograms.

Table 3.33. Composition of groups of a 20-box model of a Caribbean coral reef as obtained by method 2 (ECOPATH II aggregation routine); names of subgroups refer to original groups in the 50-box model.

| Group no. | Predator/prey group | Group no. | Predator/prey group |
|-----------|-------------------------------------------------------|-----------|---------------------------------------|
| 1 | Aggregation 26 | 6 | Aggregation 25 |
| F | Large sharks/rays, carnivorous | F | Small reef fish, omnivorous 1 |
| F | Sharks/scombrids, carnivorous | F | Small reef fish, omnivorous 2 |
| F | Large jacks, carnivorous | F | Small reef fish, omnivorous 3 |
| F | Intermediate jacks, carnivorous | R | Sea turtles |
| F | Intermediate reef fish, carnivorous 1 | I | Crabs |
| F | Intermediate reef fish, carnivorous 2 | | |
| F | Large reef fish, carnivorous | | |
| F | Intermediate reef fish, carnivorous 3 | 7 | Aggregation 21 |
| F | Small reef fish, carnivorous 1 | | |
| F | Small reef fish, carnivorous 2 | I | Octopuses |
| F | Large groupers, carnivorous | I | Lobsters |
| F | Intermediate reef fish, carnivorous 4 | I | Shrimps/hermit crabs/stomatopods |
| F | Small Gobiidae, carnivorous | I | Asteroids |
| B | Sea birds | | |
| 2 | Aggregation 24 | 8 | Amphipods/isopods/tanaids/pycnogonids |
| F | Small jacks, carnivorous | 9 | Gastropods |
| F | Large to intermediate schooling fish, pelagic | | |
| F | Small schooling fish, pelagic | 10 | Chitons/scaphopods |
| I | Squids | | |
| | | 11 | Polychaetes/priapuloids/ophiuroids |
| 3 | Aggregation 20 | | |
| F | Hemiramphidae, herbivorous | 12 | Bivalves |
| F | Kyphosidae, herbivorous | | |
| F | Large Scaridae, herbivorous | 13 | Ascidians/barnacles/bryozoans |
| F | Intermediate Scaridae, herbivorous | | |
| F | Small Scaridae, herbivorous | | |
| I | Echinoids | 14 | Sponges |
| 4 | Aggregation 22 | 15 | Sea anemones/corals |
| F | Intermediate reef fish, herbivorous | 16 | Zooplankton |
| F | Blenniidae, herbivorous | | |
| I | Holothuroids/sipunculids/ echiuroids/hemichordates | 17 | Microfauna |
| | | 18 | Phytoplankton |
| 5 | Engraulidae, herbivorous | 19 | Benthic autotrophs |
| | | 20 | Detritus, POM, DOM |

B = bird, F = fish, I = invertebrate, R = reptile.

box model continued to exist as separate entities within the 20-box model. This is certainly a positive feature, since differences in DC, consumption requirements and size are pronounced between nonfish groups.

When comparing the group composition produced by method 1 with that by method 2, the food type is apparently the principal aggregation criterion. The basic producer groups of the system are left unchanged in both 20-box models (groups 15-20, corals to detritus). The combination of fish with nonfish and the distinction of the food type in a group depend primarily on the contrast of (1) intuitive use of more than one aggregation criterion, and (2) objective combination of groups by a single aggregation criterion. The choice of method depends on the objective of the modelling effort. Method 1 produces a more detailed view of the system where ecological criteria, such as size, lifestyle and, to some extent, habitat (e.g., pelagic vs. demersal) are explicitly considered. If one aims, though, to produce an energy pyramid of the system considered, where energy transfer efficiencies and flow rates between distinct trophic levels are of interest, method 2 is certainly the better choice.

PARAMETER ESTIMATION. Tables 3.34 and 3.35 list the outputs for the three key parameters, as well as EEs and additional parameters. The percent food intake of prey for 20 predator groups (database of a food matrix) may be found in Appendix Table 8.8.3. Fig. 3.7 shows the 20-box model of the PRVI coral reef as obtained by method 2.

Table 3.34. Results of the ECOPATH II parameter estimation for a 20-box model of a Caribbean coral reef (method 2: ECOPATH II aggregation routine); P/B = production/biomass ratio, Q/B = consumption/unit biomass, EE = ecotrophic efficiency (part of production consumed by predators).

| Group no. | Predator group | No. of groups ^a | Biomass (g·m ⁻²) | Average P/B (year ⁻¹) | Average Q/B (year ⁻¹) | End EE |
|-----------|---------------------------------------|----------------------------|------------------------------|-----------------------------------|-----------------------------------|--------|
| 1 | Aggregation 26 | 14 | 29.884 | 1.064 | 8.100 | 0.854 |
| 2 | Aggregation 24 | 4 | 23.818 | 1.944 | 15.473 | 0.822 |
| 3 | Aggregation 20 | 6 | 133.112 | 1.053 | 7.616 | 0.615 |
| 4 | Aggregation 22 | 3 | 76.491 | 0.380 | 6.429 | 0.855 |
| 5 | Engraulidae, herbivorous | 1 | 3.325 | 2.835 | 43.400 | 0.791 |
| 6 | Aggregation 25 | 5 | 24.984 | 1.603 | 15.196 | 0.902 |
| 7 | Aggregation 21 | 4 | 46.670 | 1.274 | 9.235 | 0.926 |
| 8 | Amphipods/isopods/tanaids/pycnogonids | 1 | 13.250 | 5.000 | 125.500 | 0.927 |
| 9 | Gastropods | 1 | 46.800 | 2.800 | 14.000 | 0.974 |
| 10 | Chitons/scaphopods | 1 | 62.000 | 0.360 | 11.700 | 0.911 |
| 11 | Polychaetes/priapuloids/ophiuroids | 1 | 33.000 | 5.200 | 61.600 | 0.919 |
| 12 | Bivalves | 1 | 109.500 | 2.230 | 9.500 | 0.913 |
| 13 | Ascidians/barnacles/bryozoans | 1 | 137.400 | 2.300 | 20.000 | 0.798 |
| 14 | Sponges | 1 | 800.000 | 1.500 | 5.000 | 0.249 |
| 15 | Corals/sea anemones | 1 | 121.000 | 1.090 | 9.000 | 0.977 |
| 16 | Zooplankton | 1 | 32.000 | 40.000 | 165.000 | 0.921 |
| 17 | Decomposers/microfauna | 1 | 15.000 | 280.000 | 1,900.000 | 0.979 |
| 18 | Phytoplankton | 1 | 40.000 | 70.000 | - | 0.74 |
| 19 | Benthic autotrophs | 1 | 1,300.000 | 13.250 | - | 0.36 |
| 20 | Detritus, DOM, POM | 1 | 2,000.000 | - | - | - |

^a Number of groups refers to original groups from the 50-box model.

Table 3.35. Results of the ECOPATH II parameter estimation for a 20-box model of a Caribbean coral reef method 2 (ECOPATH II aggregation routine). Additional parameters.

Import to detritus: 475 g·m⁻², other imports to boxes are 0; catches for all boxes are 0. Excretion = 5% and egestion = 15% (nonassimilated food, default values) of the food intake of each box.

| Group no. | Predator group | Food intake ^a | Respiration ^a | Predation mortality (production *EE) ^a | Flow to detritus ^a | Assimilation ^a | Trophic level |
|-----------|---------------------------------------|--------------------------|--------------------------|---------------------------------------------------|-------------------------------|---------------------------|---------------|
| 1 | Aggregation 26 | 242.06 | 161.848 | 27.145 | 53.070 | 193.650 | 3.45 |
| 2 | Aggregation 24 | 368.53 | 248.534 | 38.067 | 81.934 | 294.828 | 3.57 |
| 3 | Aggregation 20 | 1,013.73 | 670.880 | 86.182 | 256.670 | 810.985 | 2.08 |
| 4 | Aggregation 22 | 491.77 | 364.329 | 24.860 | 102.585 | 393.420 | 2.05 |
| 5 | Engraulidae, herbivorous | 144.31 | 106.018 | 7.456 | 30.832 | 115.444 | 2.32 |
| 6 | Aggregation 25 | 379.66 | 263.680 | 36.105 | 79.875 | 303.728 | 2.48 |
| 7 | Aggregation 21 | 430.98 | 285.306 | 55.052 | 90.624 | 344.786 | 3.02 |
| 8 | Amphipods/isopods/tanaids/pycnogonids | 1,662.88 | 1,264.050 | 61.417 | 337.408 | 1,330.300 | 2.28 |
| 9 | Gastropods | 655.20 | 393.120 | 127.695 | 134.385 | 524.160 | 2.37 |
| 10 | Chitons/scaphopods | 725.40 | 558.000 | 20.339 | 147.061 | 580.320 | 2.38 |
| 11 | Polychaetes/priapuloids/ophiuroids | 2,032.80 | 1,454.640 | 157.634 | 420.526 | 1,626.240 | 2.35 |
| 12 | Bivalves | 1,037.88 | 586.672 | 222.354 | 228.849 | 830.300 | 2.10 |
| 13 | Ascidians/barnacles/bryozoans | 2,748.00 | 1,882.380 | 252.302 | 613.318 | 2,198.400 | 2.34 |
| 14 | Sponges | 4,000.00 | 2,000.000 | 299.177 | 1,700.823 | 3,200.000 | 2.00 |
| 15 | Corals/sea anemones | 1,089.00 | 739.310 | 128.803 | 220.887 | 871.200 | 2.34 |
| 16 | Zooplankton | 5,280.00 | 2,944.000 | 1,179.004 | 1,156.996 | 4,224.000 | 2.60 |
| 17 | Microfauna | 28,500.00 | 18,600.000 | 4,112.797 | 5,787.203 | 22,800.000 | 2.00 |
| 18 | Phytoplankton | - | 0.000 | 2,073.120 | 726.880 | - | 1.00 |
| 19 | Benthic autotrophs | - | 0.000 | 6,201.429 | 11,023.570 | - | 1.00 |
| 20 | Detritus, POM, DOM | - | - | - | - | - | 1.00 |

^a(g·m⁻²·year⁻¹ WW).

continued...

SYSTEM PROPERTIES. System properties, such as summary statistics, network flow indices and size of flows from primary producers, detritus and all flows combined for a 20-box model as obtained by method 2, are shown in Appendix Tables 8.9.4 to 8.9.6. These properties showed only small differences in relation to the 50-box model and to the 20-box model as obtained by method 1. This is mainly due to the fact that both methods of aggregation maintain a given throughput during the aggregation process. Table 3.36 presents the transfer efficiencies between discrete trophic levels.

Transfer efficiencies by discrete trophic levels for primary producers were somewhat higher (1% to 6.5%) than in the 50-box model. Compared to the 20-box model obtained by method 1, differences varied from 5% to 14%. For detritus, transfer efficiencies showed differences to the 50-box model of -1% to +4% and to the 20-box model, obtained by method 1, of -1% to +14%. For combined flows, differences of transfer efficiencies varied from -1% to +5% compared to the 50-box model, and from -1% to +14% for the 20-box model as obtained by method 1. Differences in transfer efficiencies were more pronounced for levels V, VI and VII when compared to the 50-box model (+5% to +9%), and for levels IV to VII when compared to the 20-box model as obtained by method 1 (+10% to +14%). These more pronounced differences between the 20-box models, obtained by two different aggregation methods, may as well be due to efficiencies computed after removing flows involved in cycles. Since the two methods produced somewhat differing groups, the efficiencies should also be different.

MIXED TROPHIC IMPACTS. Fig. 3.8 shows, as its predecessors of the 50-box model and the 20-box model obtained by method 1, the direct and indirect impacts on the biomass of prey groups by a raise of biomass of the predator group. As for the 20-box model obtained by method 1, the present figure was included for a better visualization of impacts, when the system is reduced to 20 boxes by the ECOPATH II aggregation routine and for comparison with the 20-box model as obtained by method 1.

3.3.2.2. THE 11-BOX MODELS

Method 1 : Cluster Analysis + Intuition

COMPOSITION OF GROUPS. Table 3.37 shows the group composition for the 11-box model of a Caribbean coral reef as obtained by method 1. Main aggregation criterion was now the food type, and the aim was to reduce the system to a very low number of boxes. The division into two main habitats (pelagic-midwater, demersal-off bottom) is still visible, mainly because the group of apex predators was maintained by the cluster analysis. The zooplanktivorous schooling pelagics (group 2) correspond exactly to group 2 (aggregation 24) in the 20-box model obtained by method 2.

The omnivore group (4) is rather small, consisting of only two invertebrate taxa, because a group or taxon was identified as omnivorous only when none of the three main food sources (live animal, plant, detrital food) outweighed the others, i.e., when one element could not be classified by the criterion ">50% of a distinct food source".

For this method of aggregation and with increasing level of aggregation, the number of aggregation criteria is reduced. This means that distinctness of main features of a system is paid for in terms of loss in detail. Thus, it is up to the modeller or user of the model to decide which element is more appropriate for the respective objective and from this, to select an appropriate number of boxes for the final model.

Table 3.36. Transfer efficiencies (%) by discrete trophic level for a 20-box model (method 2).

| Source | I | II | III | IV | V | VI | VII | VIII |
|-----------|---|------|------|------|------|------|------|------|
| Producers | - | 11.3 | 12.2 | 10.3 | 10.8 | 11.5 | 11.5 | - |
| Detritus | - | 13.5 | 17.3 | 9.7 | 9.6 | 11.4 | 11.5 | - |
| All flows | - | 13.1 | 16.5 | 9.8 | 9.8 | 11.4 | 11.5 | - |

Table 3.37. Composition of groups of an 11-box model of a Caribbean coral reef obtained by method 1 (cluster analysis + intuition); names of subgroups refer to original groups in the 50-box model.

| Group no. | Predator/prey group | Group no. | Predator/prey group |
|-----------|---------------------------------------------------|-----------|-------------------------------------------|
| 1 | Apex predators, mainly pelagic | 5 | Benthic herbivores (>50% plant food) |
| | F Large sharks/rays, carnivorous + | F | Kyphosidae, herbivorous + |
| | F Sharks/scombrids, carnivorous + | F | Intermediate reef fish, herbivorous + |
| | F Large jacks, carnivorous + | F | Large Scaridae, herbivorous + |
| | F Intermediate jacks, carnivorous + | F | Intermediate Scaridae, herbivorous + |
| | B Sea birds | F | Small reef fish, omnivorous 3 + |
| 2 | Schooling pelagics, zooplanktivorous | F | Small Scaridae, herbivorous + |
| | F Small jacks, carnivorous + | F | Blenniidae, herbivorous + |
| | F Large to intermediate schooling fish, pelagic + | I | Crabs + |
| | F Hemiramphidae, herbivorous + | I | Amphipods/isopods/tanaids/pycnogonids + |
| | F Small schooling fish, pelagic + | I | Echinoids + |
| | F Engraulidae, herbivorous + | I | Chitons/scaphopods + |
| | I Squids | I | Corals |
| 3 | Benthic carnivores (>50% live animal food) | 6 | Benthic detritivores (>50% detrital food) |
| | F Large groupers, carnivorous + | I | Holothuroids/sipunculids |
| | F Intermediate reef fish, carnivorous 1 + | I | echiuroids/hemichordates + |
| | F Large reef fish, carnivorous + | I | Bivalves + |
| | F Intermediate reef fish, carnivorous 2 + | I | Ascidians/barnacles/bryozoans + |
| | F Intermediate reef fish, carnivorous 3 + | I | Sponges |
| | F Intermediate reef fish, carnivorous 4 + | 7 | Zooplankton |
| | F Small reef fish, carnivorous 1 + | 8 | Decomposers/microfauna |
| | F Small reef fish, carnivorous 2 + | 9 | Phytoplankton |
| | F Small reef fish, omnivorous 1 + | 10 | Benthic autotrophs |
| | F Small reef fish, omnivorous 2 + | 11 | Detritus |
| | F Small Gobiidae, carnivorous + | | |
| | R Sea turtles + | | |
| | I Octopuses + | | |
| | I Lobsters + | | |
| | I Asteroids + | | |
| | I Sea anemones + | | |
| 4 | Benthic omnivores | | |
| | I Gastropods + | | |
| | I Polychaetes/priapuloids/ophiuroids | | |

B = bird, F = fish, I = invertebrate, R = reptile.

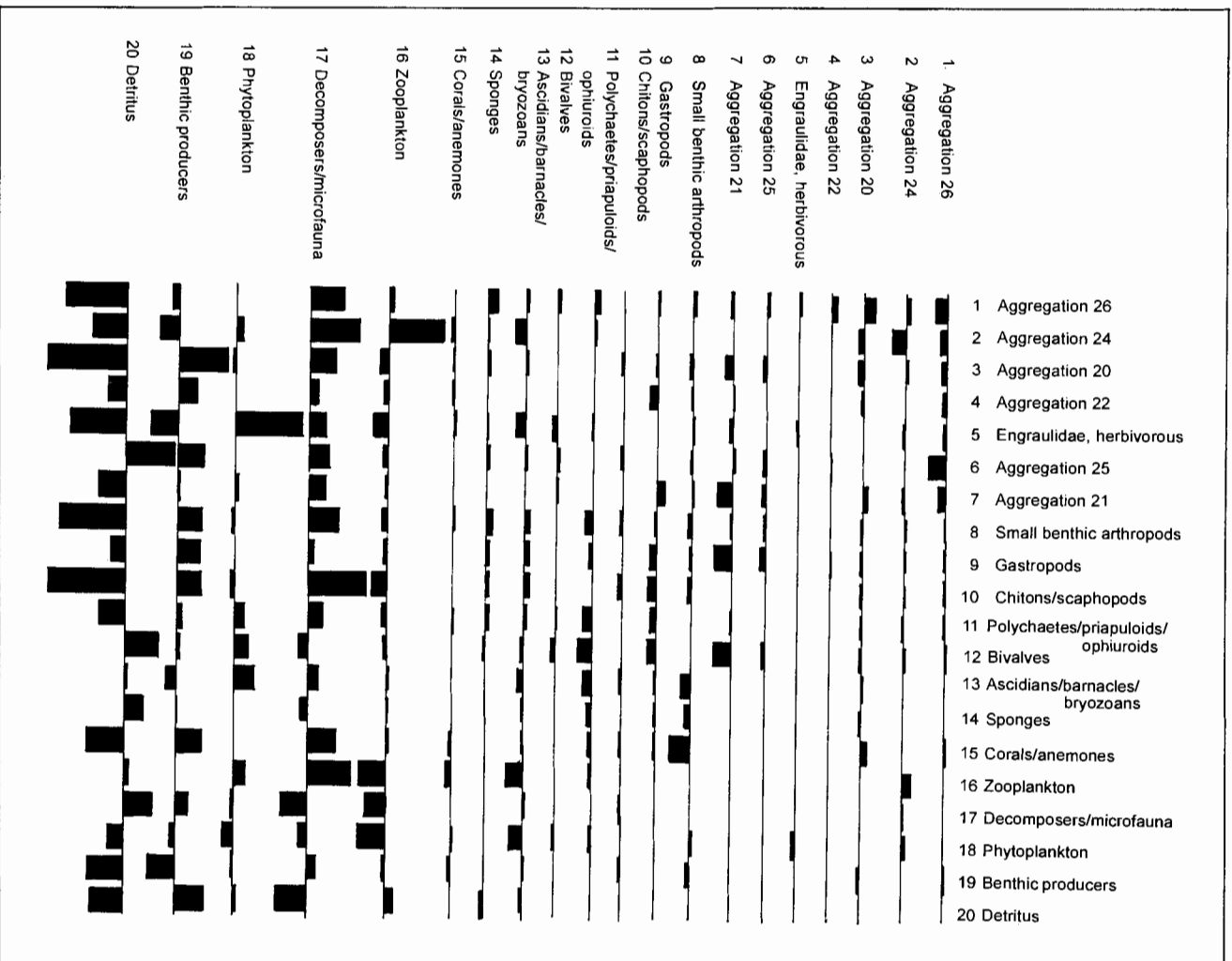


Fig. 3.8. Mixed trophic impacts of a 20-box model (method 2) of a Caribbean coral reef showing the direct and indirect impacts an increase in the biomass of the groups to the left of the histograms will have on the groups above the histograms.

PARAMETER ESTIMATION. Tables 3.38 and 3.40 present input/output estimates for the three key parameters as well as EEs and additional parameters and ratios. Input values did not have to be modified to equilibrate the system, therefore the same tables are valid for input as well as for output. The same is true for the diet matrix (Table 3.39).

Fig. 3.9 shows the 11-box model as produced by method 1.

SYSTEM PROPERTIES. Summary statistics, network flow indices and size of flows for primary producers, detritus and all flows combined for an 11-box model, obtained by method 1, are presented in Appendix Tables 8.9.7 to 8.9.9. The results, again, showed only small differences when compared to the previous models. Table 3.41 presents the transfer efficiencies between discrete trophic levels.

Overall, transfer efficiencies for primary producers showed only negligible differences when compared to the 20-box model obtained by the same method. The only more pronounced differences could be observed in level V for detritus (+8%) and consequently for all flows combined (+7%) and in level VII for all flows combined (-10%). The differences may be because before computing the transfer efficiencies, ECOPATH II removes the flows involved in cycles. A difference in group composition between the two methods must therefore result in differing transfer efficiencies.

MIXED TROPHIC IMPACTS. Fig. 3.10 shows the direct and indirect impacts on the biomass of prey groups by a raise of biomass of the predator group for an 11-box model obtained by method 1. As for the 20-box models, the figure was included here for comparison with the previous models.

Method 2: ECOPATH II - Aggregation Routine

COMPOSITION OF GROUPS. Table 3.42 shows the composition of groups for an 11-box model resulting from a further reduction of the number of boxes by the ECOPATH II aggregation routine. At this level of reduction, fish groups are strongly fused with nonfish ones. The tendency to produce groups of similar food type is maintained. Aggregation 12 (group 1) represents a very large group (large in the sense of containing the highest number of groups from the 50-box model) of carnivores. Group 2 (aggregation 20) holds the herbivores consisting of few groups, then group 3 (aggregation 15), the omnivores. Group 4 (aggregation 11) is a mixture of phytoplankton-feeding engraulids with detritivorous and phytoplanktivorous bivalves, ascidians, barnacles and bryozoans. Group 5 (aggregation 13) consists of invertebrates feeding predominantly on autotrophs. Groups 6-11 were maintained as in the 50-box model. Except for the sponges (group 6), which were pooled with other detritivores in the 20- and 11-box models of method 1, these groups are represented as single boxes in the 11-box models from both methods. Both 11-box models contain similar qualitative groupings although, as stated above, the detritivores are not separated from the herbivores due to the assignment of the same trophic level to both food types in method 2.

PARAMETER ESTIMATION. Tables 3.43, 3.44 and Appendix Table 8.8.5 show the ECOPATH II outputs for the 11-box model. Fig. 3.11 presents the second of the two 11-box models of a Caribbean coral reef.

Table 3.38. Results of the ECOPATH II parameter estimation for an 11-box model of a Caribbean coral reef (method 1: cluster analysis + intuition); average values are weighted by biomass; P/B = production/biomass ratio, Q/B = consumption/unit biomass, EE = ecotrophic efficiency (part of production consumed by predators).

| Group no. | Predator group | No. of groups ^a | Biomass (g·m ⁻²) | Average P/B (year ⁻¹) | Average Q/B (year ⁻¹) | End EE |
|-----------|--------------------------------------------|----------------------------|------------------------------|-----------------------------------|-----------------------------------|--------|
| 1 | Apex predators, mainly pelagic | 5 | 2.542 | 0.908 | 8.876 | 0.254 |
| 2 | Schooling pelagics, zooplanktivorous | 6 | 28.268 | 2.02 | 19.689 | 0.817 |
| 3 | Benthic carnivores (>50% live animal food) | 17 | 80.336 | 1.215 | 9.183 | 0.894 |
| 4 | Benthic invertebrates, omnivorous | 2 | 79.8 | 3.792 | 33.684 | 0.947 |
| 5 | Benthic herbivores (>50% plant food) | 11 | 357.448 | 1.118 | 14.02 | 0.739 |
| 6 | Benthic detritivores (>50% detrital food) | 4 | 1,112.89 | 1.599 | 7.196 | 0.445 |
| 7 | Zooplankton | 1 | 32 | 40 | 165 | 0.925 |
| 8 | Decomposers/microfauna | 1 | 15 | 280 | 1,900 | 0.979 |
| 9 | Phytoplankton | 1 | 40 | 70 | - | 0.74 |
| 10 | Benthic autotrophs | 1 | 1,300 | 13.25 | - | 0.362 |
| 11 | Detritus, POM, DOM | 1 | 2,000 | - | - | - |

^a Number of groups refers to original groups from the 50-box model.

Table 3.39. Diet composition matrix for an 11-box model of a Caribbean coral reef, method 1 (cluster analysis + intuition).

| Group no. | Predator group | No. of groups ^a | Prey group | | | | | | | | | | | Sum |
|-----------|--------------------------------------------|----------------------------|------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-----|
| | | | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | |
| 1 | Apex predators, mainly pelagic | 5 | 0.026 | 0.32 | 0.279 | 0.016 | 0.328 | 0.012 | 0.018 | - | - | - | 0.001 | 1 |
| 2 | Schooling pelagics, zooplanktivorous | 6 | - | 0.047 | 0.006 | 0.03 | 0.006 | 0.01 | 0.607 | 0.002 | 0.181 | 0.085 | 0.026 | 1 |
| 3 | Benthic carnivores (>50% live animal food) | 17 | - | 0.018 | 0.077 | 0.167 | 0.169 | 0.213 | 0.122 | 0.026 | - | 0.095 | 0.113 | 1 |
| 4 | Benthic invertebrates, omnivorous | 2 | - | - | 0.004 | 0.047 | 0.024 | 0.128 | 0.038 | 0.045 | 0.076 | 0.286 | 0.352 | 1 |
| 5 | Benthic herbivores (>50% plant food) | 12 | - | - | 0.002 | 0.004 | 0.019 | 0.057 | 0.049 | 0.08 | 0.001 | 0.68 | 0.108 | 1 |
| 6 | Benthic detritivores (>50% detrital food) | 4 | - | - | - | - | - | - | 0.051 | 0.05 | 0.088 | 0.004 | 0.807 | 1 |
| 7 | Zooplankton | 1 | - | - | - | - | - | - | - | 0.6 | 0.2 | - | 0.2 | 1 |
| 8 | Microfauna | 1 | - | - | - | - | - | - | - | - | - | 0.05 | 0.95 | 1 |
| 9 | Phytoplankton | 1 | - | - | - | - | - | - | - | - | - | - | (1) | (1) |
| 10 | Benthic autotrophs | 1 | - | - | - | - | - | - | - | - | - | - | (1) | (1) |
| 11 | Detritus, POM, DOM | 1 | - | - | - | - | - | - | - | - | - | - | - | 0 |

^a Number of groups refers to original groups from the 50-box model.

Table 3.40. Results of the ECOPATH II parameter estimation for an 11-box model of a Caribbean coral reef, method 1 (cluster analysis + intuition). Additional parameters. Import to detritus: 475 g·m⁻², other imports to boxes are 0; catches for all boxes are 0. Excretion = 5% and egestion = 15% (nonassimilated food, default values) of the food intake of each box.

| Group no. | Predator group | Food intake ^a | Respiration ^a | Predation mortality (production *EE) ^a | Flow to detritus ^a | Assimilation ^a | Trophic level |
|-----------|--------------------------------------------|--------------------------|--------------------------|---------------------------------------------------|-------------------------------|---------------------------|---------------|
| 1 | Apex predators, mainly pelagic | 22.56 | 15.742 | 0.587 | 6.234 | 18.050 | 4.320 |
| 2 | Schooling pelagics, zooplanktivorous | 556.57 | 388.154 | 46.658 | 121.757 | 445.255 | 3.574 |
| 3 | Benthic carnivores (>50% live animal food) | 737.73 | 492.572 | 87.214 | 157.939 | 590.180 | 3.469 |
| 4 | Benthic invertebrates, omnivorous | 2,687.98 | 1,847.785 | 286.639 | 553.559 | 2,150.386 | 2.774 |
| 5 | Benthic herbivores (>50% plant food) | 5,011.42 | 3,609.510 | 295.144 | 1,106.767 | 4,009.137 | 2.969 |
| 6 | Benthic detritivores (>50% detrital food) | 8,008.36 | 4,627.174 | 792.685 | 2,588.498 | 6,406.685 | 2.234 |
| 7 | Zooplankton | 5,280.00 | 2,944.000 | 1,184.375 | 1,151.625 | 4,224.000 | 2.800 |
| 8 | Decomposers/microfauna | 28,500.00 | 18,600.000 | 4,110.585 | 5,789.415 | 22,800.000 | 2.000 |
| 9 | Phytoplankton | - | 0.000 | 2,070.772 | 729.228 | - | 1.000 |
| 10 | Benthic autotrophs | - | 0.000 | 6,230.956 | 12,419.045 | - | 1.000 |
| 11 | Detritus, POM, DOM | - | 0.000 | - | - | - | 1.000 |

^a(g·m⁻²·year⁻¹ WWV).

continued....

Table 3.40 continued

| Group no. | Omnivory index | Gross efficiency | Net efficiency | Respiration/assimilation | Production/respiration |
|-----------|----------------|------------------|----------------|--------------------------|------------------------|
| 1 | 0.1224 | 0.102 | 0.128 | 0.872 | 0.147 |
| 2 | 0.2388 | 0.103 | 0.128 | 0.872 | 0.147 |
| 3 | 0.4523 | 0.132 | 0.165 | 0.835 | 0.198 |
| 4 | 0.3915 | 0.113 | 0.141 | 0.859 | 0.164 |
| 5 | 0.1661 | 0.080 | 0.100 | 0.900 | 0.111 |
| 6 | 0.2526 | 0.222 | 0.278 | 0.722 | 0.385 |
| 7 | 0.1600 | 0.242 | 0.303 | 0.697 | 0.435 |
| 8 | 0.0000 | 0.147 | 0.184 | 0.816 | 0.226 |
| 9 | 0.0000 | - | - | - | - |
| 10 | 0.0000 | - | - | - | - |
| 11 | 0.3656 | - | - | - | - |

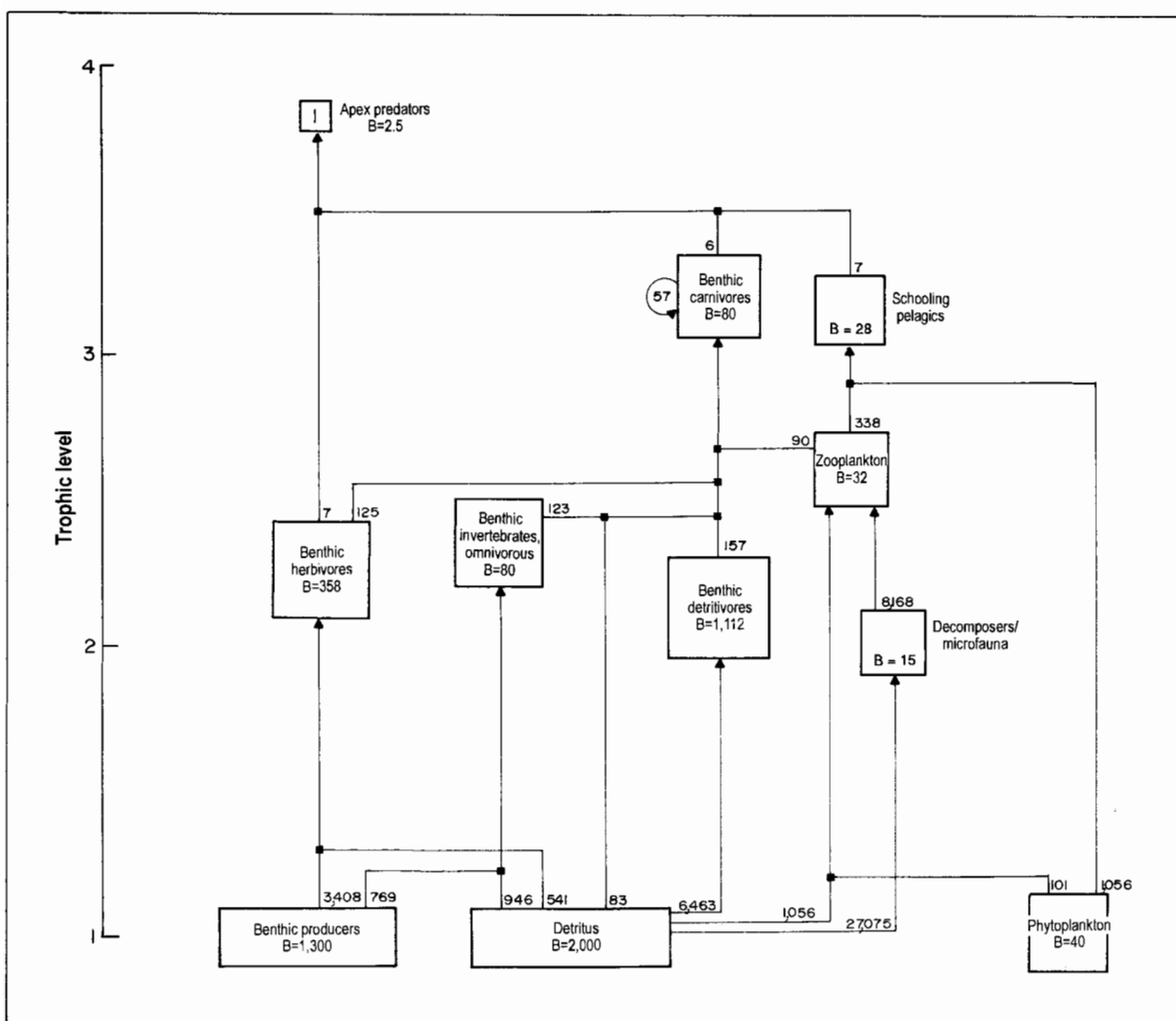


Fig. 3.9. Graphical representation of an 11-box model as obtained by method 1 (cluster analysis + intuition). The area of each box is proportional to the logarithm of the biomass (B, in gm⁻²WW) of each group. Values represent the average for all subsystems in Fig. 2.1. Flows are in gm⁻²year⁻¹WW. Incoming flows (entering bottom half of a box) represent at least 95% of the diet of a group.

Table 3.41. Transfer efficiencies (%) by discrete trophic level for an 11-box model (method 1).

| Source | I | II | III | IV | V | VI | VII | VIII |
|-----------|---|------|------|-----|-----|------|------|------|
| Producers | - | 10.7 | 11.8 | 9.5 | 9.8 | 10.0 | - | - |
| Detritus | - | 13.6 | 17.1 | 9.0 | 9.5 | 9.9 | 10.0 | - |
| All flows | - | 13.1 | 16.4 | 9.1 | 9.5 | 9.9 | 10.0 | - |

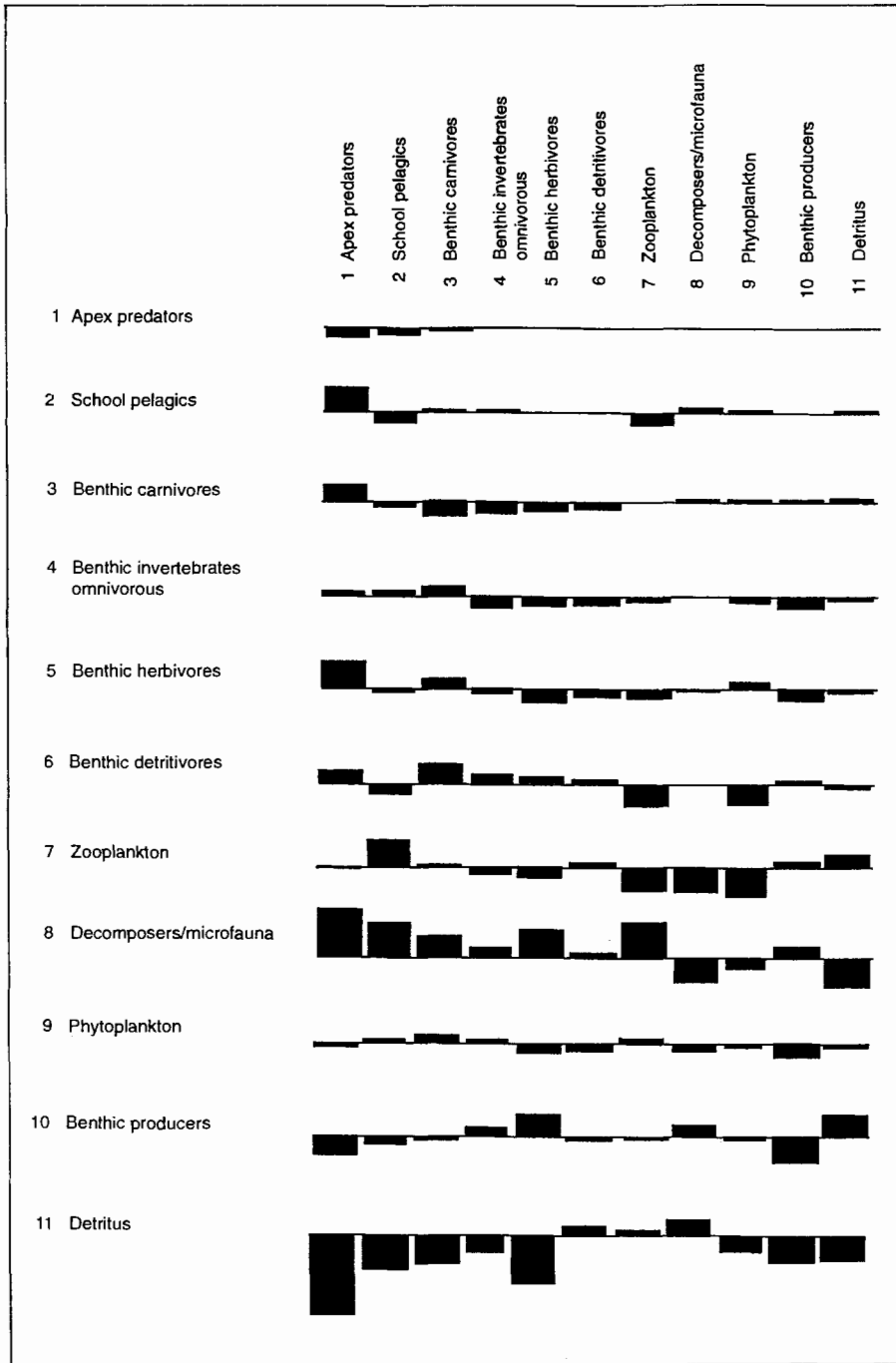


Fig. 3.10. Mixed trophic impacts of an 11-box model (method 1) of a Caribbean coral reef showing the direct and indirect impacts an increase in the biomass of the groups to the left of the histograms will have on the groups positioned above the histograms.

Table 3.42. Composition of groups for an 11-box model of a Caribbean coral reef obtained by method 2 (ECOPATH II aggregation routine); names of subgroups refer to original groups in the 50-box model.

| Group no. | Predator/prey group | Group no. | Predator/prey group |
|-----------|-------------------------------------------------|-----------|---------------------------------------------------|
| 1 | Aggregation 12 | 3 | Aggregation 15 |
| F | Large sharks/rays, carnivorous + | F | Intermediate reef fish, herbivorous + |
| F | Sharks/scombrids, carnivorous + | F | Small reef fish, omnivorous 1 + |
| F | Large jacks, carnivorous + | F | Small reef fish, omnivorous 2 + |
| F | Intermediate jacks, carnivorous + | F | Small reef fish, omnivorous 3 + |
| F | Small jacks, carnivorous + | F | Blenniidae, herbivorous + |
| F | Intermediate reef fish, carnivorous 1 + | R | Sea turtles + |
| F | Large to intermediate schooling fish, pelagic + | I | Crabs + |
| F | Intermediate reef fish, carnivorous 2 + | I | Gastropods + |
| F | Large reef fish, carnivorous + | I | Polychaetes/priapuloids/ophiuroids + |
| F | Intermediate reef fish, carnivorous 3 + | I | Holothuroids/sipunculids/echiuroids/hemichordates |
| F | Small reef fish, carnivorous 1 + | | |
| F | Small schooling fish, pelagic + | 4 | Aggregation 11 |
| F | Small reef fish, carnivorous 2 + | F | Engraulidae, herbivorous + |
| F | Large groupers, carnivorous + | I | Bivalves + |
| F | Intermediate reef fish, carnivorous 4 + | I | Ascidians/barnacles/bryozoans |
| F | Small Gobiidae, carnivorous + | | |
| B | Sea birds + | 5 | Aggregation 13 |
| I | Squids + | I | Amphipods/isopods/tanaids/pycnogonids + |
| I | Octopuses + | I | Chitons/scaphopods + |
| I | Lobsters + | I | Sea anemones/corals |
| I | Shrimps/hermit crabs/stomatopods + | | |
| I | Asteroids + | | |
| | | 6 | Sponges |
| 2 | Aggregation 20 | 7 | Zooplankton |
| F | Hemiramphidae, herbivorous + | 8 | Microfauna |
| F | Kyphosidae, herbivorous + | 9 | Phytoplankton |
| F | Large Scaridae, herbivorous + | | |
| F | Intermediate Scaridae, herbivorous + | 10 | Benthic autotrophs |
| F | Small Scaridae, herbivorous + | | |
| I | Echinoids + | 11 | Detritus, POM, DOM |

B = bird, F = fish, I = invertebrate, R = reptile.

Table 3.43. Results of the ECOPATH II parameter estimation for an 11-box model of a Caribbean coral reef (method 2: ECOPATH II aggregation routine); P/B = production/biomass ratio, Q/B = consumption/unit biomass, EE = ecotrophic efficiency (part of production consumed by predators).

| Group no. | Predator group | No. of groups ^a | Biomass (g·m ⁻²) | Average P/B (year ⁻¹) | Average Q/B (year ⁻¹) | End EE |
|-----------|------------------------|----------------------------|------------------------------|-----------------------------------|-----------------------------------|--------|
| 1 | Aggregation 12 | 22 | 100.372 | 1.371 | 10.377 | 0.874 |
| 2 | Aggregation 20 | 6 | 133.112 | 1.053 | 7.616 | 0.615 |
| 3 | Aggregation 15 | 10 | 181.275 | 2.051 | 19.636 | 0.931 |
| 4 | Aggregation 11 | 3 | 249.975 | 2.277 | 15.722 | 0.847 |
| 5 | Aggregation 13 | 3 | 196.250 | 1.123 | 17.719 | 0.955 |
| 6 | Sponges | 1 | 800.000 | 1.500 | 5.000 | 0.249 |
| 7 | Zooplankton | 1 | 32.000 | 40.000 | 165.000 | 0.921 |
| 8 | Decomposers/microfauna | 1 | 15.000 | 280.000 | 1,900.000 | 0.979 |
| 9 | Phytoplankton | 1 | 40.000 | 70.000 | - | 0.740 |
| 10 | Benthic autotrophs | 1 | 1,300.000 | 13.250 | - | 0.360 |
| 11 | Detritus, DOM, POM | 1 | 2,000.000 | - | - | - |

^a Number of groups refers to original groups from the 50-box model.

Table 3.44. Results of the ECOPATH II parameter estimation for an 11-box model of a Caribbean coral reef, method 2 (ECOPATH II aggregation routine). Additional parameters. Import to detritus: 475 g·m⁻², other imports to boxes are 0; catches for all boxes are 0. Excretion = 5% and egestion = 15% (nonassimilated food, default values) of the food intake of each box.

| Group no. | Predator group | Food intake ^a | Respiration ^a | Predation mortality (production *EE) ^a | Flow to detritus ^a | Assimilation ^a | Trophic level |
|-----------|------------------------|--------------------------|--------------------------|---------------------------------------------------|-------------------------------|---------------------------|---------------|
| 1 | Aggregation 12 | 1,041.58 | 695.688 | 120.265 | 225.627 | 833.263 | 3.33 |
| 2 | Aggregation 20 | 1,013.73 | 670.880 | 86.182 | 256.670 | 810.985 | 2.08 |
| 3 | Aggregation 15 | 3,559.43 | 2,475.769 | 346.294 | 737.371 | 2,847.547 | 2.33 |
| 4 | Aggregation 11 | 3,930.18 | 2,575.070 | 482.112 | 872.998 | 3,144.143 | 2.28 |
| 5 | Aggregation 13 | 3,477.28 | 2,561.360 | 210.559 | 705.356 | 2,781.820 | 2.32 |
| 6 | Sponges | 4,000.00 | 2,000.000 | 299.177 | 1,700.823 | 3,200.000 | 2.00 |
| 7 | Zooplankton | 5,280.00 | 2,944.000 | 1,179.004 | 1,156.996 | 4,224.000 | 2.60 |
| 8 | Decomposers/microfauna | 28,500.00 | 18,600.000 | 4,112.797 | 5,787.203 | 22,800.000 | 2.00 |
| 9 | Phytoplankton | - | - | 2,073.121 | 726.880 | - | 1.00 |
| 10 | Benthic autotrophs | - | - | 6,201.430 | 11,023.570 | - | 1.00 |
| 11 | Detritus, POM, DOM | - | - | - | - | - | 1.00 |

^a(g·m⁻²·year⁻¹ WW).

| Group no. | Omnivory index | Gross efficiency | Net efficiency | Respiration/assimilation | Production/respiration |
|-----------|----------------|------------------|----------------|--------------------------|------------------------|
| 1 | 0.3260 | 0.132 | 0.170 | 0.835 | 0.198 |
| 2 | 0.1216 | 0.138 | 0.170 | 0.827 | 0.209 |
| 3 | 0.3234 | 0.104 | 0.130 | 0.869 | 0.150 |
| 4 | 0.3075 | 0.145 | 0.180 | 0.819 | 0.221 |
| 5 | 0.3014 | 0.063 | 0.080 | 0.921 | 0.086 |
| 6 | 0.0010 | 0.300 | 0.380 | 0.625 | 0.600 |
| 7 | 0.2400 | 0.242 | 0.300 | 0.697 | 0.435 |
| 8 | 0.0000 | 0.147 | 0.180 | 0.816 | 0.226 |
| 9 | 0.0000 | - | - | - | - |
| 10 | 0.0000 | - | - | - | - |
| 11 | 0.3648 | - | - | - | - |

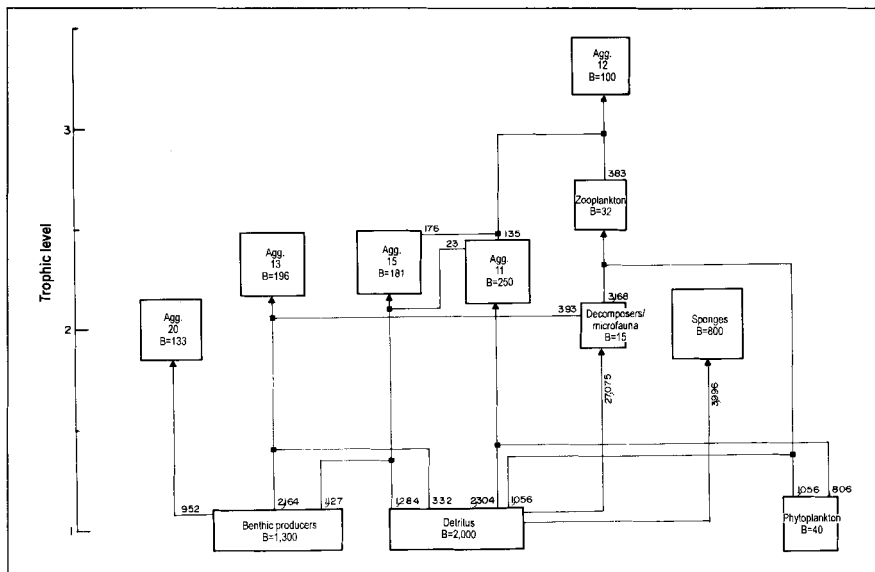


Fig. 3.11. Graphical representation of an 11-box model as obtained by method 2 (ECOPATH II aggregation routine). The area of each box is proportional to the logarithm of the biomass (B, gm⁻²WW) of each group. Values represent the average for all subsystems in Fig. 2.1. Flows are in gm⁻²·year⁻¹·WW. Incoming flows (entering bottom half of a box) represent at least 95% of the diet of a group.

SYSTEM PROPERTIES. System properties such as summary statistics, network flow indices and size of flows for primary producers, detritus and all flows combined are listed in Appendix Tables 8.9.10 to 8.9.12. Table 3.45 presents the transfer efficiencies between discrete trophic levels.

Table 3.45. Transfer efficiencies (%) by discrete trophic level for an 11-box model (method 2).

| Source | I | II | III | IV | V | VI | VII | VIII |
|-----------|---|------|------|------|------|------|-----|------|
| Producers | - | 11.4 | 12.2 | 10.4 | 10.5 | 10.9 | - | - |
| Detritus | - | 13.5 | 17.4 | 10.4 | 10.1 | 10.7 | - | - |
| All flows | - | 13.1 | 16.6 | 10.4 | 10.2 | 10.8 | - | - |

Transfer efficiencies of levels II to IV for primary producers showed only negligible differences when compared to those of the 20-box model obtained by the same method (2). Transfer efficiencies of levels V and VI showed differences of -3% and -5%, respectively. Level VII had no estimate. When compared to transfer efficiencies of the 11-box model obtained by method 1, differences were more pronounced, ranging from +3% to +9% for primary producers. For detritus, estimates for levels IV to VI differed from -6% to +7% when compared to those of the 20-box model obtained by method 2 and from +6% to +16% (level IV) when compared to the 11-box model obtained by method 1. No estimate of transfer efficiency of level VII was computed for detritus. Resulting transfer efficiencies for all flows of levels IV to VI showed differences from -5% to +6% when compared to the 20-box model obtained by the same method and from +7% to +14% (level IV) when compared to the 11-box model obtained by method 1. The 11-box model obtained by method 2 includes one discrete trophic level less than the previous models. This is due to the differing composition of carnivores which in the 20- and 11-box models obtained by method 1 discriminate between reef carnivores, schooling pelagics and large pelagic apex predators. In the 11-box model obtained by method 2, these groups are combined into a single group of carnivores.

MIXED TROPHIC IMPACTS. Fig. 3.12, showing the direct and indirect impacts on the biomass of prey groups by a raise of biomass of the predator group, was included for comparison of the 11-box model obtained by method 2 with the previous models.

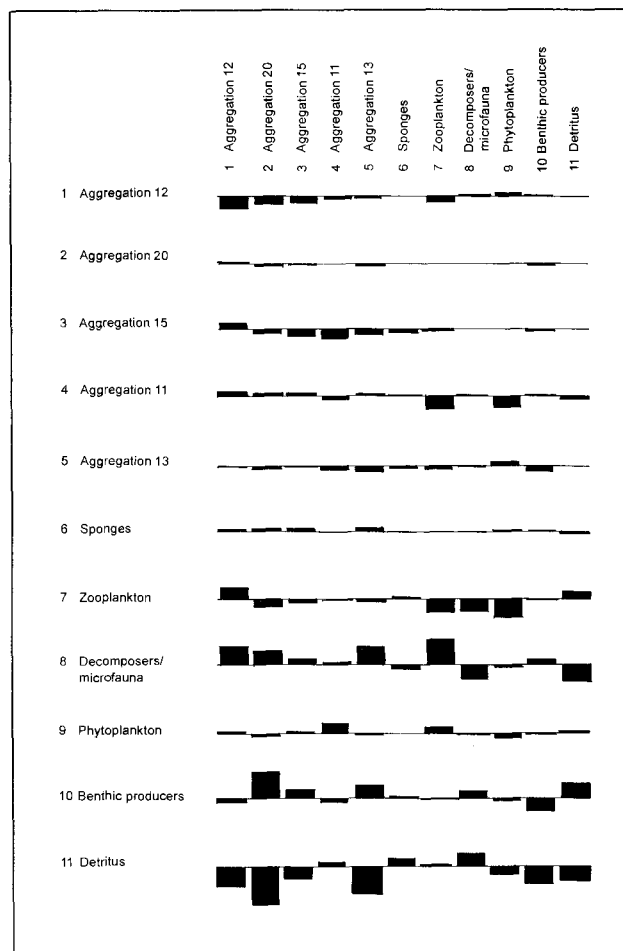


Fig. 3.12. Mixed trophic impacts of an 11-box model (method 2) of a Caribbean coral reef showing the direct and indirect impacts an increase in the biomass of the groups to the left of the histograms will have on the groups above the histograms.

4. DISCUSSION

The models presented in the "Results" section show that it is possible to construct a quantitative steady-state model of a coral reef system from published data and parameter estimates based largely on multivariate statistics.

4.1. Compatibility of the Models with the PRVI Coral Reef System

In the following, selected outputs will be compared to "external" estimates, i.e., values not used to produce the models, to assess the "match" between the models and real coral reef ecosystems (unexploited segments of the Puerto Rico-Virgin Islands [PRVI] reef area and coral reef ecosystems in general).

Benthic autotrophs showed a relatively high ecotrophic efficiency (EE) of 0.36 in the five models. According to Jacobsen and Browder (1987), only 5%-10% of net productivity of *Thalassia* is consumed directly. This corresponds to an EE of 0.05-0.1. Seagrasses formed a sizable fraction of benthic autotrophs but it is well possible that a considerably stronger grazing pressure was exerted on the algae, the other elements of this box. The reef contains a specialized fish fauna feeding directly on benthic plants: mixed aggregations of acanthurids, scarids, chaetodontids, for example, graze on coral-algal reef flats like sheep on a meadow (see e.g., Earle 1972).

Based on rotenone sampling, Randall (1963b) obtained a reef fish biomass of 160 gm⁻² and 158 gm⁻² for two fringing reef areas in the Virgin Islands. Small pelagics as well as large apex predators (sharks, rays, scombrids, jacks) were not included in this figure. These values were not used as biomass inputs for the models presented here because (1) the estimates referred only to a single subsystem and, more important, (2) an independent estimate of reef fish biomass was needed for validation of the models' outputs (subsequent comparison of results from the model with "external" estimates). Only by using independent estimates can the compatibility of both be checked and thus the results confirmed or rejected.

The models' output of total fish biomass amounted to 104 gm⁻², including small pelagics and apex predators for an average system composed of the five subsystems in Fig. 2.1. Demersal reef fish biomass was 75 gm⁻². This value was only 47% of Randall's (1963b) estimate. Although his figure was based on a rotenone sampling from the subsystem "coral reef" where biomasses for typical reef fishes usually are higher than for an average square meter from five subsystems, his figure was probably still an underestimate of the real values. Apparently, the two reefs were sampled by fish poison only once each. Even sampling by poison will underestimate fish biomass on a reef probably due to escape of larger individuals or, in case of small cryptic species, by flight into hiding places, where they subsequently die undetected (Pauly 1984).

Smith (1973) showed that not all individuals present in a specific reef area are killed by the first application of poison. For a more realistic estimate of population size, sampling has to be

repeated. A “Leslie” plot of catch/effort (catch per application of poison) versus cumulative catch can be used to obtain an estimate of real population size from the intersection of the resulting regression line with the abscissa, as was shown by Pauly (1984) for two coral reef fish species (see Fig. 4.1).

From Fig. 4.1, a ratio between the first sample and the initial population size of 0.31 for reef eels and of 0.32 for bluehead wrasse can be estimated. Applying the mean ratio of 0.315 to Randall's fish biomass estimate of 160 gm⁻² yielded an initial fish biomass of 533 gm⁻².

Another evaluation of reef fish standing stock, carried out by Dammann (1969, cited in Lewis 1981) yielded a fish biomass for a Virgin Islands fringing reef of 38 gm⁻². Since this paper was not available to me, I cannot discuss the methodology involved. Note, however, that the fish biomass estimated here is intermediate between Randall's and Dammann's estimates of fish biomass.

Fast and Pagan-Font (1973, cited in Jacobsen and Browder 1987) obtained a biomass estimate for a natural reef in Puerto Rico of 23.3 gm⁻². Fishing pressure on reef fish in Puerto Rico is more pronounced than in the Virgin Islands, and this is sufficient to explain the lower biomasses.

Jacobsen and Browder (1987, p. 34) stated that “herbivorous fishes dominate tropical reef faunas throughout the world”. They do not specify in what sense, if in numbers or in biomass. I assume they are referring to biomass.

The strictly herbivorous reef fish of the families Scaridae and Acanthuridae in Randall's study made up between 20.7% (33.1:160.1 gm⁻²) and 28.7% (45.4:157.8 gm⁻²) of reported fish biomass. The biomass estimate of herbivores for the first reef area of 33.1 gm⁻² compares very well with the models' estimate of 32.6 gm⁻² (demersals only), although this figure contains additional species from other families. Herbivorous fish biomass of the model amounts to 43.4% (32.6:75.1 gm⁻²) of total reef fish biomass (demersals only) whereas herbivorous fish contribute to total number of demersal reef species only with 12% (30 out of 243). The model's output for herbivorous reef fish biomass is thus compatible with “external” estimates.

4.2. Comparison of the Models with Other Coral Reef Areas

In the following, features of our PRVI coral reef model will be compared to those of other coral reefs. Table 4.1 presents a comparison of selected outputs of three coral reef models,

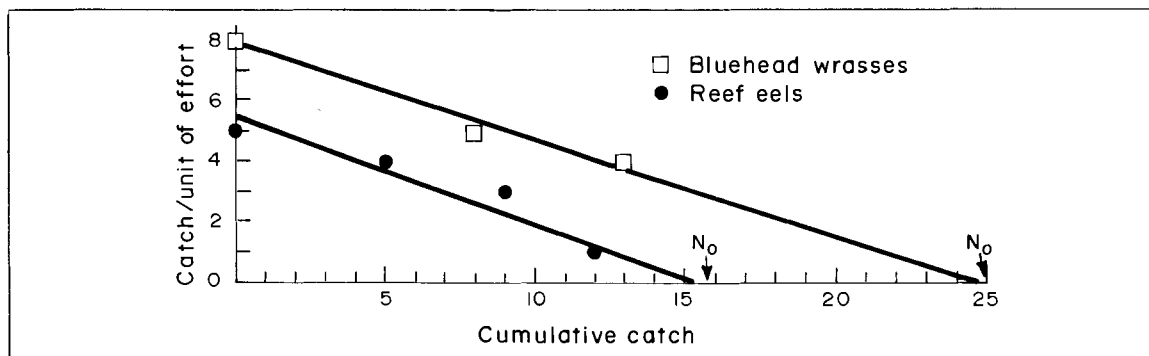


Fig. 4.1. Leslie plots for reef eels (*Kaupichthys hyoproroides*) and bluehead wrasses (*Thalassoma bifasciatum*) from an isolated Bahamian reef patch, with estimates of unexploited population size (reproduced from Fig. 6.1 in Pauly 1984).

Table 4.1. Comparison of properties of three coral reef models, analyzed by ECOPATH II.

| Properties/models | PRVI | Bolinao | FFS |
|-------------------------------------------------------------------------|-------------------------|-------------------|-----------|
| Number of boxes | 50, 20, 11 ^a | 23 | 16 |
| Total system throughput (g·m ⁻² ·year ⁻¹ WW) | 107,473 | 39,307 | 4,973 |
| Cycling index (% of throughput) | 18.3 | - ^b | 11.0 |
| Capacity (flowbits) | 438,213 | 86,774 | 8,893,465 |
| Ascendency (% of capacity) | 34 | 44 | 14.2 |
| Overhead (% of capacity) | 66 | (56) ^c | 74.8 |
| Sum of all production (g·m ⁻² ·year ⁻¹) | 51,282 | 1,938 | - |
| Sum of all respiratory flows (g·m ⁻² ·year ⁻¹) | 32,523 | 18,965 | - |
| Sum of all flows into detritus (g·m ⁻² ·year ⁻¹) | 23,193 | 18,405 | - |
| Total biomass excluding detritus (D) (gm ⁻²) | 3,048 | 1,879 | 175 |
| | (2,000) | (10) | - |
| Total fish biomass (gm ⁻²) | 104 | 6.8 | 17.5 |
| Ratio fish/total biomass | 0.03 | 0.004 | 0.1 |

^aEstimates in this column refer to 50 boxes. Values for 20 and 11 boxes differ only slightly.

^bNo estimate available.

^cComputed here as capacity (100%) - ascendency (44%).

analyzed by ECOPATH II, such as (1) our PRVI coral reef system; (2) the French Frigate Shoals (FFS) in Hawaii (Pauly et al. 1993a); and (3) segments of the Bolinao coral reef system (Aliño et al. 1993) in the Philippines.

Table 4.1 shows that the PRVI system is large in terms of throughput when compared to Bolinao and FFS. Total biomass and fish biomass are highest in the PRVI system, although the ratio of fish biomass to total biomass is highest for FFS. It is surprising that such a high fish biomass is maintained by such a small biomass of nonfish groups. Estimates of P/B and Q/B differ to some degree for those ecological groups from both models which can be compared directly (see tables in Pauly et al. 1993a). If P/B of prey groups is higher and Q/B is lower than in the PRVI model, the ecological efficiency (production/consumption) is increased and thus, the same biomass of predators can be maintained by a smaller biomass of prey groups. The inverse seems to be the case for the Bolinao model in which a very low fish biomass is maintained by a large nonfish biomass. This very low fish biomass is the result of the strong overfishing of the fish resources of the Bolinao reef system (McManus et al. 1992).

Ascendency is, as stated before, a measure of average mutual information in a system. Ulanowicz and Norden (1990) supposed systems, in the absence of major perturbations, to mature in the direction of increasing ascendency. This would imply that the Bolinao system is the most mature, the PRVI area intermediate and the FFS system the most immature. On the other hand, FFS is situated close to the northern limit for coral reefs and hence, production, respiration, etc. and, in consequence, ascendency are low compared to PRVI and Bolinao. This indicates that caution must be taken when using ascendency as an index of maturity (Christensen 1995).

Parrish et al. (1986), in their study on "piscivory in a Hawaiian coral reef fish community", found that 12%-31% of the biomass of the reef fish community of the four stations sampled consisted of piscivores. Piscivory is not considered as a separate feeding type in the model of a Caribbean coral reef presented here; rather as part of carnivory. Our findings can thus not be directly compared. However, one can estimate from the output diet composition (DC) matrix of the 50-box model (Table 3.23) that about one-third of the diet of the carnivorous-omnivorous fish (referring only to demersal groups) consists of fish. Piscivory amounts thus to 18.7% of demersal reef fish biomass. This figure is well within the range reported by Parrish et al. (1986).

As stated, (large) herbivorous reef fish are not a preferred food item of piscivorous reef fish, as demonstrated by the respective input and output percentages of this prey item in the DC matrix of the 50-box model (see "Results" section). This is further reflected by the "standardized forage ratios" (S_i) computed by ECOPATH II for each predator in relation to its prey groups. This ratio indicates a preference or avoidance of a potential prey group. In contrast to Ivlev's electivity index, this ratio is independent of prey density. Table 4.2 shows the ratios for all groups of herbivorous fish in the 50-box model in relation to their predator groups.

The majority of values in Table 4.2 are negative, and several values are positive but close to zero. Many of these were increased (as shown by the "+" sign in Table 4.2) due to a raise of the respective diet fraction performed to reduce predation pressure on carnivorous fish groups (see "Results"). Since a value of -1 for S_i corresponds to total avoidance of a prey group and $S_i = 0$ to nonselective feeding, this confirms that herbivorous fishes are indeed not a preferred food item for the majority of carnivorous predators.

Interestingly, Parrish et al. (1986) stated, with reference to Hawaiian reefs, that a number of herbivorous and omnivorous species from the families Scaridae, Acanthuridae, Chaetodontidae, etc. "are not eaten in significant quantity by abundant predators"! They stated further that "these non-piscivorous fishes consume considerable food resources; the route by which these resources pass on through the trophic system remains in doubt". Parrish et al.'s findings confirmed the model's results on low EEs for larger herbivorous fish groups.

If not esteemed as food source by higher trophic levels, herbivorous reef fish seem to be more strongly involved in other processes on the reef. Gygi (1975) estimated the erosion of coral reefs (i.e., sediment production), generated by the stoplight parrotfish *Sparisoma viride* on two reef types in Bermuda, as a side-effect of its feeding habits: "The fish scrape away algae and some of the rock below with strong, beak-like jaws, and crush the mouthful in the pharyngeal mill. Swallowed rock bits are used as a milling agent to rip up algal cells". The excreted sediment is then deposited as sand. For an algal-vermetid rock flat with coral-algal patch reefs, Gygi determined a parrotfish standing stock of 20 gm⁻²WW. The resulting erosion from this fish biomass was computed to be 5,814 gha⁻¹day⁻¹. This corresponded to an erosion of 0.58 gm⁻²day⁻¹ and to 212 gm⁻²year⁻¹. The model's output biomass for scarids was ca. 30 gm⁻².

Table 4.2. Standardized forage ratios for herbivorous reef fish groups as computed by ECOPATH II for a 50-box model of a Caribbean coral reef.

| Predator | Herbivorous prey groups ^a | | | | | |
|--------------------------------------|--------------------------------------|---------|---------|---------|---------|--------|
| | KY | IH | LS | IS | SS | BH |
| Large sharks/rays | -0.121+ | -0.712+ | -0.921+ | -0.978- | -0.988- | - |
| Sharks/scombrids | 0.115+ | -0.527+ | -0.618+ | -0.947 | -0.889 | -0.602 |
| Intermediate jacks | -0.053 | 0.240 | -0.785 | 0.325 | 0.573 | 0.889 |
| Intermediate reef fish carnivorous 1 | 0.051+ | -0.465+ | 0.174+ | 0.456+ | 0.490 | - |
| Intermediate reef fish carnivorous 2 | 0.085+ | -0.439+ | 0.002+ | 0.700+ | 0.313+ | - |
| Large reef fish carnivorous | -0.359 | -0.044+ | -0.667+ | -0.181+ | -0.249+ | -0.222 |
| Intermediate reef fish carnivorous 3 | -0.310+ | -0.434+ | 0.110+ | -0.229+ | -0.332+ | 0.181 |
| Small reef fish carnivorous 1 | - | - | - | - | -0.545 | 0.795 |
| Large groupers | 0.112+ | -0.529+ | 0.335+ | - | - | - |
| Small reef fish omnivorous 1 | - | - | - | - | -0.173 | 0.739 |
| Sea birds | - | -0.760 | -0.973 | -0.787 | -0.902 | - |

^aBH = herbivorous Blenniidae

H = intermediate reef fish herbivorous

IS = intermediate Scaridae

KY = Kyphosidae

LS = large Scaridae

SS = small Scaridae

Thus, sediment production or reef erosion due to scarid feeding activities would amount to 318 $\text{gm}^{-2}\text{year}^{-1}$. Parrotfish in the model do not feed exclusively on encrusting algae or corals. From the algal composition in stomachs of West Indian parrotfish, given by Randall (1967, see also Appendix Table 8.5.1), it was estimated, that at least 50% of the diet consisted of encrusting algae and corals. The sediment production by parrotfish would thus amount to at least 160 $\text{gm}^{-2}\text{year}^{-1}$. Parrotfish can therefore be considered important sediment producers in coral reefs of the PRVI area.

4.3. New Insights into Structure and Functioning of a Coral Reef as an Ecosystem Unit

The overwhelming part of production of large- to intermediate-sized herbivorous reef fish, which provide the bulk of herbivorous fish biomass, is not transferred to higher trophic levels. The models' outputs indicate that much of their production is recycled directly into the detrital pool. However, the fish must die before they can be decomposed by bacteria and other microfauna. The large- and intermediate-sized parrotfish (groups 23 and 24 in the 50-box model) showed, indeed, higher natural mortalities (= P/B values) than similar-sized carnivorous reef fish groups (Table 4.3).

The nutrients from the decomposed fish are available to reef autotrophs. These autotrophs (mainly reef algae), once more, are consumed by the herbivorous reef fish population. This represents, thus, a very short loop for effective recycling of matter. This shortcut is also reflected in the difference in transfer efficiencies between discrete trophic levels. In the case of the 50-box model, highest efficiencies of 13% to 16% were computed by the corresponding ECOPATH II routine for the transfer of matter between levels I to II and II to III for all flows and of 11.2% to 12.4%, respectively, for primary producers (see Table 3.27). Following is a sharp decline in transfer efficiencies between levels III to IV to 9.9% for all flows and to 10.2% for primary producers.

In the "Introduction", I pointed out three features of general agreement characterizing coral reefs: high species diversity, pronounced complexity of trophic relationships and high rates of production in spite of being surrounded by nutrient-poor oceanic waters.

A fourth feature should be added to these three: the existence of shortcuts for the effective recycling of matter. Throughputs of 59% for all flows and of 89% for primary producers of the respective total throughputs occur on trophic level II, the highest observed. Second highest throughputs occur on trophic level III with 7.7% for all flows and 10% for primary producers of the respective total throughputs. An 86% of total flow from primary producers to detritus is

Table 4.3. Comparison of natural mortalities M (= P/B values) of selected groups of herbivorous and carnivorous reef fish from the outputs of the balanced 50-box model (Tables 3.18 and 3.22).

| Group no. | Group name | Size range (g) | M (year ⁻¹) |
|-----------|---------------------------------------|----------------|-------------------------|
| 6 | Intermediate reef fish, carnivorous 1 | 2,586 - 12,407 | 0.580 |
| 12 | Large reef fish, carnivorous | 7,728 - 70,000 | 0.380 |
| 13 | Intermediate reef fish, carnivorous 3 | 290 - 5,010 | 0.640 |
| 11 | Intermediate reef fish, herbivorous | 288 - 1,200 | 0.710 |
| 23 | Large Scaridae, herbivorous | 5,558 - 23,000 | 0.850 |
| 24 | Intermediate Scaridae, herbivorous | 1,102 - 3,720 | 1.155 |

recycled directly without entering the higher levels of the system. Thus, for primary producers, the corresponding backflows to detritus are second highest for trophic level II with 12.2% and third for III with 1.4% of total backflow. For combined flows, as well, they are second highest for trophic level II with 43% and third for III with 5.4%.

Coral reef ecosystems are often viewed as “mature” systems, in which structures and processes are maintained at some optimum level, i.e., with minimum leaks (see Odum and Odum 1955). Thus, the reef, as a thermodynamical unit, may not benefit from sharks or tunas which are not an integral part of the actual reef community. “Feeding” these occasional visitors might mean a net loss of energy, since sharks and tunas and other large apex predators, feeding in part on the reef community, tend to migrate away from these systems and take the energy ingested with them.

The reef, as a thermodynamical unit, may also not benefit from passing matter up through many trophic levels. With every additional trophic level through which matter is passed, about 80%-90% of its energy is lost. Thus, direct recycling of matter may be the best way for the reef to conserve energy and to build up biomass. Other reef components with a high biomass for a single taxon, such as the strongly herbivorous echinoids (80% plant food) or the detritivorous sponges, are additional examples for this aspect of reef metabolism. They show rather low exploitation rates as food source for other elements of the system, as expressed by relatively low EEs compared to the majority of nonfish taxa. This feature is furthermore supported by the fact that transfer efficiencies between discrete trophic levels stay on the same rather low level of 9%-11% up to the highest discrete trophic levels in all models. Pauly and Christensen (1995) found, when comparing trophic efficiencies in 48 different ecosystems, all analyzed using ECOPATH II, an average trophic efficiency of 10%, with a range of 2%-34%.

Summarizing, it can be hypothesized that the reef’s “strategy” is not to achieve high transfer efficiencies between trophic levels but to build up biomass through maintenance of short cycles for an effective recycling of matter back to its base, the detrital pool. Or else: only ecosystems where those features had developed during evolution could survive in nutrient-poor waters.

During a long-term study on community structure of an artificial reef in the Virgin Islands, Ogden and Ebersole (1981) found an astonishing stability in species composition. They discussed critically results from the Pacific (Great Barrier Reef, Australia) where this stability was never reported. Sale (1975, 1977) and Talbot et al. (1978) suggested that high diversity in coral reef fish communities is maintained by the chance arrival—“lottery”—of larval colonists to unpredictably available habitat space. They cited as evidence the stochastic composition of coral reef fish communities on small natural and artificial structures. Ogden and Ebersole (1981) attributed this stochastic community structure found by some workers in the Pacific to the small structures—natural and artificial—that these workers have studied. Small coral heads are, according to Ogden and Ebersole (1981), too small for detection of “real” differences in community structure. They argued that, when the appropriate scale is selected, coral reef systems become stable (i.e., display flow-through equilibrium) and show an orderly functioning. Moreover, the predictability of the community seems to increase with the size of the reef. From the results of the models presented here, it can be concluded that a coral reef system can straightforwardly be made to reflect a steady-state or flow-through equilibrium, when a whole system approach, such as in our models, is applied.

This does not prevent stochastic events such as predation, disturbance, incidental recruitment of species, etc., from taking place on a smaller scale within that system. As pointed out earlier, for the present models, fish groups were aggregated, based on similarity of energetic

requirements and not on taxonomical closeness. Thus, individuals of a certain fish species within a “box” may easily be replaced by those of another species with similar energetic features. When considering the flow of energy through the reef, the only important feature of a species group or “box” is its capacity to maintain its biomass and production rate, so that the same amount of energy remains available to other boxes of the system. The species composition within that box may be viewed as a stochastic feature, depending on the above-described “lottery”. The results of investigations of community structures, which emphasize their stochastic feature, cannot express anything about overall performance of reef metabolism, such as stability or instability of structures and processes at higher scales (e.g., relationships between boxes, trophic levels, etc.). Thus, from these experiments, it cannot be concluded if the reef as an entity is in a steady state or not.

Hall et al. (1990), in their study on the usefulness of power analysis for the experimental design of energy flow studies, discussed the compatibility of results from energy flow models with those from field experiments. They concluded that field experiments could not be substituted by modelling. They did not specify to what scale of model or experiment they referred. From their examples, it must be assumed that the scale is relatively small, e.g., one predator and a few prey items. I do not doubt the necessity of field experiments on energy flow, on the contrary, they provide urgently needed estimates for modelling energy flows on a larger scale.

Jones (1986, 1987) discussed the hypothesis of recruitment limitation to coral reefs. In general, space and food are assumed to be major limiting factors. Opinions on this subject are divided. Jones (1987) found indications from his experiments with a zooplankton-feeding pomacentrid species at One Tree Island (Great Barrier Reef, Australia) that recruitment of larvae or postlarval 0+ cohorts of this species was not limited by the reef community, it was accidental. Mortality did not seem to be density-dependent, but time of maturation of juveniles was influenced by population structure. When individuals from older cohorts were present, maturation time of young fish was longer, suggesting a possible food limitation for recruitment. If this kind of “stock-recruitment interaction” is valid for the majority of species within a box, it would substantiate the hypothesis that a “box” tends to maintain its average standing stock biomass and food requirements.

Most fisheries can be viewed as an apex predator within a system. First, they remove the predators, then take the production these took. When the apex predators of our PRVI reef models, such as sharks, rays, large scombrids, jacks and groupers, are removed, the production of the reef fish community consumed by these groups becomes available to the fisheries. Excluding the schooling pelagic fish (the production of this group consumed by apex predators is $1.9 \text{ gm}^{-2}\cdot\text{year}^{-1}$), the reef fish production then available to the fisheries amounts to about $4 \text{ gm}^{-2}\cdot\text{year}^{-1}$ (excluding small blennies and gobies). The fish production available to the fisheries may be increased by extending these to lower trophic levels, such as the large carnivorous reef fish (group 12 in the 50-box model). Recalling the “norm” of 5 t km^{-2} fisheries yield for Caribbean coral reefs (Munro 1984), it seems possible to attain this yield as well for the unfished parts of the PRVI reef area, without destructive effects for the reef fish community as a whole. Overall, it is advisable to fish large individuals from higher trophic levels. This has a positive effect on prey density and should lead, at least when overall fishing pressure remains low, to increase of recruits to the adult guilds. This, again, may lead to sustainable fisheries yields.

The overfishing of invertebrates, presently occurring in PRVI, can also be explained, if to a small extent, by the results of the model. A strong predation pressure is already exerted in the

unexploited system on, e.g., crustaceans and gastropods (see high EEs for these groups). There is not much surplus production for the fisheries.

The deductions on fisheries potential for the area are very preliminary. The shortcomings of input data for nonfish groups were already pointed out. Actual standing stocks might be higher than outputs from models imply, but the “true” relations among biomass, production and consumption for nonfish groups are to date not known with sufficient accuracy.

4.4. Identification and Solution of Methodological Problems During the Modelling Process

4.4.1. Aggregation of Fish Groups

The high fish species diversity within the PRVI coral reef system was successfully reduced into fewer groups with similar energetic requirements. It is difficult to parametrize and combine ecological information of various dimensions, e.g., spatial, temporal, energetic in a given model. Emphasis should focus on one aspect, depending on the objectives of the model. In the present case, the emphasis was on energy flow and trophic relations. The model reflects to some extent the spatial aspect by the separation of pelagics and demersals and by the explicit consideration of subsystems. The division in two distinct horizontal levels is also reflected by consumption requirements and prey items.

The aggregation of nonfish taxa was achieved by considering as well as possible the energetic aspect, leading to more or less distinct trophic groups. Taxa were aggregated intuitively based on criteria known to be related to energetic requirements. This was only achieved for higher taxonomic categories, since estimates at the species level were generally unavailable. A first attempt to leave the taxonomic level and proceed to an ataxonomic approach for combining elements, including fish and nonfish, was realized in the 20- and 10-box models of methods 1 and 2.

4.4.2. Weighting Factors

Parrish et al. (1986) emphasized the impact of predator abundance on predation pressure for a Hawaiian coral reef system. I agree with these authors. Indeed, it is not only abundance (in weight) of a predator which is of importance, but also its food requirements (Q/B). Within species groups, a weighting factor could be calculated for every species of the model from average biomasses and Q/B estimates to determine the “throughput” of every species in a group. The percentages of respective prey items (DC matrix) were then weighted by these factors (see “Materials and methods”). Between species groups, this relationship is produced by the key input parameters for each group (biomass, mortality = P/B ratio, and Q/B).

Thus, ECOPATH II and some of the methods developed in the course of the work leading to this report provide possible solutions for a variety of problems related to ecosystem analysis.

4.4.3. Input Estimates for Nonfish Taxa

Input values for nonfish taxa had to be assembled from a wide geographical range for a wide variety of taxa. Preparation of input values was based on various qualitative and quantitative assumptions. The match between these data and the actual rates and states on the ground is not necessarily close. However, balancing of the model forced the various rates for

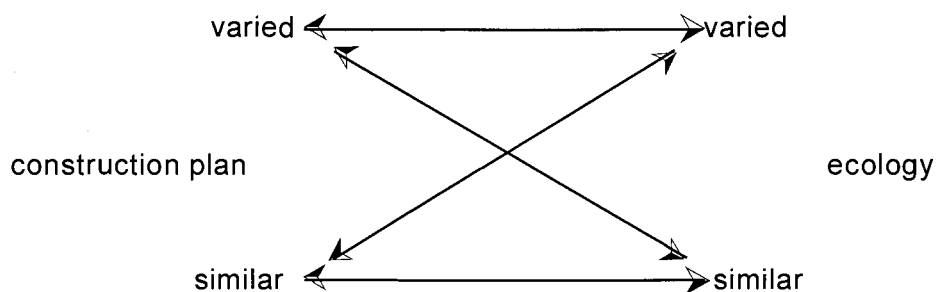
these groups to become mutually compatible. It is assumed that this process also rendered them more similar to the values on the ground.

4.5. Gaps of Knowledge and Suggestions for Future Research

4.5.1. Theoretical Research

From the above considerations, the following theoretical research topics may be deducted:

1. Determination of appropriate scales: How can a suitable ecological system unit be determined for modelling, and how does one deal with migrations? The system unit should not be too small because of stochastic effects, and not too large because of possible loss of genuine system identity. Is it possible to identify objective criteria?
2. Construction of groups: Objective methods for assembling of nonfish groups should be developed. Ataxonomical aggregation of reef nonfish taxa should be initiated at the specific level, as was done for the fishes. Nonfish taxa tend to strongly differ in their anatomy (they represent a much larger systematical range than fishes). If one aims to assemble ecologically similar groups, one must base the aggregation process on ecological features, not on morphological ones. The figure below shows possible combinations between construction plan and ecology by connecting all features.



An objective procedure for the construction of groups would be as follows: (1) determination of relevant ecological variables (depending on one's objective); (2) application of a factor analysis to identify independent variables; (3) cluster analysis to aggregate species, and eventually; (4) discriminant analysis to determine if clusters differ significantly.

4.5.2. Field Research

Several times in the foregoing, it was pointed out that the input data for nonfish groups were too scattered, often based on risky assumptions and therefore quite unreliable. To improve this situation, the database for inputs must be improved. This opens a wide area for future practical research. In the first place, the existing literature could be searched more thoroughly for estimates and data sets on the species level.

Needed are, furthermore, analyses of diet, food consumption requirements, abundances and biomass, growth performance, etc., of reef invertebrates and other nonfish taxa, preferably at the species level.

Other areas requiring work are length-weight data and growth parameters for reef fish species which may not be of interest for fisheries, and rigorous studies of the DC of apex predators.

The initial biomasses of fish groups had to be changed considerably to balance the 50-box model. Biomasses of groups, including small pelagic or cryptic fish species, were strongly increased whereas biomasses of groups, including large- to intermediate-sized fish species (both carnivores and herbivores), were decreased, more or less proportional to the size of the fishes. The input biomasses of the large, highly motile predators had to be decreased the most.

Densities, used to compute biomasses of small pelagic fish species, were assumed to correspond to the highest abundance level encountered for reef fish species. More realistic estimates of abundance of small pelagic stocks in a given area should be obtained by the application of, e.g., hydroacoustic techniques (Thorne 1979).

The need to increase input biomasses of groups, including cryptic species, is not surprising: very small, cryptic species tend to be overlooked during visual censuses, and other methods should be used for biomass estimation (e.g., successive rotenone sampling and a Leslie plot, see above).

The case of large, highly motile predators is as straightforward. It could be that the SCUBA divers performing the censuses actually attract faraway apex predators (whose sense of smell and hearing is geared, indeed, to detect faraway, potential preys).

It could also be that census surveys in general, or at least those whose results were used in the present study, do not account appropriately for potential effect on estimated biomass of a brief sighting of a large or very large fish, compared with much longer sightings of smaller fish.

If so, one could imagine a correction factor, proportional to the inverse of the sighting time of any recorded fish, which would lead to estimates of "effective" biomass, i.e., to estimates of the biomass relying on a given reef for its food.

Last but not least, catch data must be considered, since there is fishing activity in the area. If it is not possible to obtain such data at the species level, attempts should be made to report catch data for ecological guilds, i.e., at the generic or family level. The fish species are divided into three main groups: (1) apex predators, loosely related to the reef; (2) a mainly carnivorous group comprising around 70%-80% of reef fish species; and (3) a mainly herbivorous group comprising a little more than 10% of species but around 40% of total fish biomass. This division could be utilized for other purposes. The species-rich cluster of (predominately) carnivores could be grouped by species of the same family or genus and thus better be matched with species clusters used in fisheries statistics. Appeldoorn and Lindeman (1985) referred to species clusters known from fisheries statistics "in many respects to behave similar to a unispecies stock with size selective mortality". They identified species clusters to "approximating an ecological guild". Many of the model's fish groups contain several species of a family or genus. Examples are members of the families Haemulidae, Muraenidae, Blenniidae, Scaridae, Serranidae, etc. Note that, for fisheries management purposes, it is generally advisable to separate fish groups from nonfish groups.

What are the mechanisms by which stock size of, e.g., herbivorous reef fish is controlled, since predation pressure on these groups is rather low? Are they food- or recruitment-limited?

In future models, the ecological position of the different developmental stages of a fish species could be modelled by treating each stage as a different species with different inputs. These inputs could be prepared and stages clustered according to, e.g., energetic requirements.

Seasonal differences (e.g., rainy and dry seasons) could be modelled by producing one model for each season or month (as done by Jarre-Teichmann and Pauly 1993 for the Peru system).

New insights into structure and functioning of a coral reef as an ecosystem unit were achieved by the modelling process. By modifying inputs deducted from published information on the system (these changes are fully documented in the input-output tables in the "Results" section), a hypothetical reef could be constructed. Overall, the model presented here can be assumed to represent the specific coral reef system from which the majority of information was gathered. This hypothetical reef can now be tested and improved.

5. CONCLUSIONS

1. Quantitative steady-state models of a Caribbean coral reef system could be constructed from published data and estimates, based largely on multivariate statistics.

2. Calculated net primary production necessary to support the system modelled therein, of 20,025 gm⁻²year⁻¹WW, was well within the range of a rough estimate of 15,000-25,000 gm⁻²year⁻¹WW previously available for the Puerto Rico-Virgin Islands (PRVI) coral reef area.

3. Total fish biomass of 104 gm⁻²WW rendered an intermediate value when compared to model-independent estimates of coral reef fish biomass (23-160 gm⁻²WW) for several segments in the PRVI coral reef area.

4. The estimated total biomass, total fish biomass and total system throughput of the models indicate a large system compared to two other coral reef models analyzed by ECOPATH II (French Frigate Shoals [FFS], Hawaii, and Bolinao reef area, Philippines). The ratio of fish biomass/total biomass was intermediate for PRVI with a higher ratio for FFS and a very low ratio for Bolinao due to high exploitation by fisheries.

5. The degree of piscivory for the Caribbean reef models presented here of 19% of reef fish biomass is well within the range reported for a Hawaiian coral reef fish community.

6. Large- to intermediate-sized herbivorous reef fish are not a preferred food item of larger piscivorous reef fish. This finding is also in agreement with observations from a Hawaiian coral reef fish community.

7. It was estimated that, based on their output biomass, parrotfish produce at least 160 g sand m⁻²year⁻¹ and may thus be considered as important sediment producers of the PRVI reef area.

8. The models' outputs indicate the existence of short cycles for an effective recycling of organic matter within the reef system. Most of the net primary production is recycled directly to the detrital pool. Transfer efficiencies between trophic levels are low.

9. The fact that the model could be balanced straightforwardly can be interpreted as showing that coral reef systems are in a "steady-state" or "flow-through equilibrium", when the appropriate spatial and temporal scale is selected. Investigations on small scales of reef community structure, which emphasize the stochastic variability of coral reefs, do not necessarily yield information on stability or instability of structures and processes at higher scales.

10. A very preliminary potential fisheries yield of 4 gm⁻²year⁻¹WW for unfished segments of the PRVI reef area was estimated from one of the models' outputs, i.e., the amount consumed by the models' apex predators. When these predators are removed first, their consumption becomes available to the fisheries.

11. A strong predation pressure is exerted on lobsters and conchs in unexploited parts of the PRVI coral reef system as indicated by the high ecotrophic efficiencies for crustaceans and gastropods. This shows how vulnerable these resources may be to an additional exploitation by fisheries.

12. The large number of fish species occurring in the PRVI reefs can effectively be reduced to a smaller number of groups by cluster analysis. Appropriate aggregation parameters are those referring to energetic requirements of a species, such as size, activity level and type of food.

13. For assembling nonfish groups, one should start from the species level and define appropriate energetic aggregation criteria by objective selection of ecological variables.

14. The contribution of predator abundance to predation pressure can be accounted for by weighting the fractions of prey of the diet composition by applying a weighting factor. This should be the product of density, mean weight and relative food consumption (Q/B) of a predator.

15. A theoretical task for future models will be to identify objective criteria for the determination of spatial and temporal scales of the system which is to be modelled and define appropriate system boundaries.

16. Information on diet, food consumption requirements, abundances and growth performance should be obtained for nonfish taxa in the PRVI reef area. This would strongly improve the reliability of the models' outputs.

17. Information on length-weight data and growth parameters for fish species without interest for fisheries (e.g., very small ones) are needed.

18. Density estimates from visual censuses, especially for highly motile species, such as apex predators, should account for factors which may lead to overestimation of biomass. Density estimates for small and/or cryptic reef species should be obtained from repeated rotenone or similar sampling.

Density estimates for small schooling pelagics are urgently needed; appropriate estimates could be obtained by hydroacoustic techniques.

19. Catch data, preferably at the species level, should be obtained and included into the models. If it is impossible to obtain such data, attempts should be made to report catch data for ecological guilds, i.e., at the generic or family level.

20. The models, as they are now, represent a more constraining hypothesis than in the beginning. This new hypothesis can again be tested and improved.

6. ACKNOWLEDGEMENTS

Among all the people who helped and supported me in many ways during the preparation of the thesis upon which this work is based, I particularly thank: Dr. D. Pauly (ICLARM, Manila) for accepting me as his doctoral student, for his professional advice and support throughout the course of the investigation, for valuable critical comments and suggestions on contents and style of earlier drafts of the thesis which led to this report; Prof. Dr. Schnack for being my host during the preparation of this report and for his general support, advice and suggestions; Prof. Dr. Chapman, for critical comments and suggestions on contents and style of an earlier draft of this thesis; Dr. Villy Christensen, for advice on various aspects of the ECOPATH II software; the technical staff of the Fishery Biology Department, K. Burkert, B. Rohloff, H. Worthmann, D. Jarosch, for their friendly and general technical support; C. Klingenberg for his help with the performance of the cluster analyses on the VAX computer at the Institut für Meereskunde; Dr. R. Froese for the preparation of a PC program to estimate and store the aspect ratios of the caudal fin of fishes and for putting an image-analyzing system at my disposal to measure these aspect ratios; the colleagues at the Institut für Meereskunde, Kiel, the Kiel University, and the Alfred Wegener Institute for Polar and Marine Research, Bremerhaven, especially D. Barthel, T. Brey, C. Clemmesen, G. Graf, T. Grünkorn, V. Isaac, A. Jarre, F. Koester, A. Morales, M. Prein, Prof. Røttger, M. Ruth, L. Sarmiento, M. Vakily, U. Waller and W. Welsch for their readiness to discuss critical matters and for providing valuable information; I. Achenbach for helping me with the preparation of the figures and for her friendship; and Mrs. Wollweber, the Chief Librarian of the Institut für Meereskunde and Dr. Kinzer for providing me with important literature.

My very special thanks go to my husband, H. Kausch. During the many years of the preparation of this report it was him who supported me financially, took care of the technical aspects of daily life, looked after our son and kept up my good spirit.

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8. APPENDICES

Appendix 8.1. Length-weight data of Caribbean reef fishes by author.

Table 8.1.1. Length-weight data of Caribbean reef fishes after Beebe and Tee-Van (1928).

| Family | Length Species | Observed | | Estimated | | (g) ^b | a ^{b,c} |
|--------------------|---------------------------------|-----------------------------|-----------------------|------------------------|------------------|------------------|------------------|
| | | Weight (cm) ^a | L _∞ (g) | W _∞ (cm) | (g) ^b | | |
| Acanthuridae | <i>Acanthurus bahianus</i> | 12.5 | 67 | 32 / | | | |
| | <i>Acanthurus coeruleus</i> | 9.5 | 40 | 32 / | | | |
| Albulidae | <i>Albula vulpes</i> | 45.0 | 1,361 | 82 | | 6,300 | 0.01494 |
| Atherinidae | <i>Atherinomorus stipes</i> | 4.5 | 2 | 8 - | | | 0.022 |
| Aulostomidae | <i>Aulostomus maculatus</i> | 57.8 | 341 | | | | 0.00177 |
| Balistidae | <i>Balistes vetula</i> | 16.6 | 171 | 40 | | | |
| | <i>Canthidermis sufflamen</i> | | | 64 | | | |
| Belonidae | <i>Ablennes hians</i> | 65.4 | 320 | 95 | | | |
| Canthigasteridae | <i>Canthigaster rostrata</i> | 5.0 | 7 | | | | |
| Carangidae | <i>Caranx bartholomaei</i> | 6.5 | 5 | 40 | | | |
| | <i>Caranx crysos</i> | 9.5 | 17 | 32 | | | |
| | <i>Caranx latus</i> | 11.5 | 42 | 58 | | | |
| | | 15.8 | 103 | | | | |
| | | 23.8 | 320 | | | | |
| | <i>Decapterus punctatus</i> | 9.5 | 15 | 32 | | | 0.0175 |
| | <i>Oligoplites saurus</i> | 23.3 | 132 | 32 | | | 0.0104 |
| | <i>Selar crumenophthalmus</i> | 17.5 | 104 | 64 | | | 0.01935 ma |
| | | 17.8 | 109 | | | | |
| | <i>Trachinotus falcatus</i> | 18.0 | 206 | 32 | | 1,400 | |
| Chaetodontidae | <i>Chaetodon striatus</i> | 10.2 | 44 | 13 | | | 0.0415 |
| Clupeidae | <i>Opisthonema oglinum</i> | 8.5 | 10 | | | | 0.0173 ma |
| | | 12.7 | 31 | | | | |
| | | 21.4 | 201 | | | | |
| Coryphaenidae | <i>Coryphaena hippurus</i> | 32.5 | 1,474 | 191 | | 15,000 | |
| | | 64.5 | 2,268 | | | | |
| Dactylopteridae | <i>Dactylopterus volitans</i> | 7.0 | 6 | 32 / | | | 0.02 |
| | | 18.2 | 136 | | | | |
| Diodontidae | <i>Chilomycterus antennatus</i> | 12.7 | 160 | 21 | | | 0.0781 |
| | <i>Diodon hystrix</i> | 5.6 | 15 | | | | 0.0854 |
| Echeneidae | <i>Echeneis naucrates</i> | 29.7 | 80 | 101 | | | 0.00305 |
| Elopidae | <i>Elops saurus</i> | 27.8 | 92 | | | | 0.00428 |
| Engraulidae | <i>Anchoa hepsetus</i> | 3.9 | 1 | 16 | | | 0.0147 |
| | | 5.4 | 2 | | | | |
| | | 7.0 | 5 | | | | |
| Ephippidae | <i>Chaetodipterus faber</i> | | | 95 | | 9,100 | |
| Gerreidae | <i>Gerres cinereus</i> | 30.0 | 1,247 | 40 | | | 0.04619 |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | 72.2 | 1,928 | 318 | | | |

continued...

Table 8.1.1 continued

| Family | Length Species | Observed | | Estimated | | (g) ^b | a ^{b,c} |
|------------------------------|----------------------------------|-----------------------------|-----------------------|------------------------|-------------------------|------------------|------------------|
| | | Weight (cm) ^a | L _∞ (g) | W _∞ (cm) | | | |
| Gobiidae | <i>Bathygobius soporator</i> | | | 16 | | | |
| | <i>Gobiosoma horsti</i> | | | 7 | | | |
| Haemulidae (Pomadasyidae) | <i>Anisotremus virginicus</i> | 20.5 | 330 | 32 | | 900 | |
| | <i>Haemulon bonariense</i> | 18.5 | 147 | | | | |
| | | 26.4 | 401 | | | | |
| | <i>Haemulon plumieri</i> | 23.5 | 316 | 48 | | 1,800 | 0.0244 |
| | <i>Haemulon sciurus</i> | 21.1 | 270 | 48 | | | |
| | <i>Pomadasys corvinaeformis</i> | 12.2 | 45 | | | | 0.0247 ma |
| | 18.4 | 153 | | | | | |
| Hemiramphidae | <i>Hemiramphus brasiliensis</i> | | | 40 | | | |
| | <i>Hyporhamphus unifasciatus</i> | 14.7 | 27 | 32 | | | 0.0085 |
| Holocentridae | <i>Holocentrus ascensionis</i> | 15.6 | 85 | 64 | | | 0.0224 |
| | <i>Holocentrus coruscus</i> | 8.5 | 16 | | | | 0.0261 |
| | <i>Myripristis jacobus</i> | 13.2 | 75 | 32 | | | 0.0326 |
| | <i>Neoniphon marianus</i> | 13.0 | 52 | 17 + | | | 0.0237 |
| | <i>Sargocentron vexillarium</i> | 10.0 | 31 | | | | 0.031 |
| Kyphosidae | <i>Kyphosus sectatrix</i> | 7.8 | 13 | 48 | | | 0.0274 |
| Labridae | <i>Bodianus rufus</i> | 17.3 | 136 | 64 | | | 0.0263 |
| | <i>Clepticus parrae</i> | 16.3 | 124 | | | | 0.0286 |
| | <i>Halichoeres gamoti</i> | 14.0 | 53 | 21 | | | 0.0193 |
| | <i>Halichoeres radiatus</i> | 6.8 | 4 | | | | 0.02345 ma |
| | | 13.9 | 55 | | | | |
| | | 32.5 | 907 | | | | |
| | <i>Lachnolaimus maximus</i> | 34.1 | 1,361 | 64 | | 6,800 | 0.0301 |
| | 11.5 | 30 | | | | 0.0197 | |
| Lutjanidae | <i>Lutjanus analis</i> | 54.2 | 4,536 | | | 11,000 | |
| | <i>Lutjanus apodus</i> | 11.5 | 42 | | | 3,600 | |
| | | 22.5 | 308 | | | | |
| | <i>Lutjanus campechanus</i> | 21.7 | 252 | 95 | | 18,000 | 0.02466 |
| | <i>Lutjanus griseus</i> | 9.0 | 14 | 95 | | 8,200 | 0.00956 |
| | <i>Lutjanus jocu</i> | 15.0 | 107 | | | | |
| | <i>Lutjanus synagris</i> | 12.8 | 50 | 32 | | 1,800 | 0.03932 ma |
| | | 28.5 | 907 | | | | |
| | | | 64 | | | | |
| Malacanthidae | <i>Malacanthus plumieri</i> | 29.7 | 190 | 40 | | | 0.00725 |
| Megalopidae | <i>Tarpon atlanticus</i> | 7.8 | 5 | 260 | | 159,000 | |
| | | 18.8 | 65 | | | | |
| | | 106.0 | 16,329 | | | | |
| Monacanthidae | <i>Aluterus schoepfii</i> | 2.2 | 1 | | | | 0.01426 |
| | | 43.0 | 1,134 | | | | |
| | <i>Aluterus scripta</i> | 8.3 | 7 | 95 | | | |
| | <i>Cantherines pullus</i> | 13.1 | 72 | | | 2,700 | 0.032 |
| | 5.0 | 7 | 21 | | | 0.056 | |
| Mugilidae | <i>Mugil curema</i> | 41.8 | 1,021 | | | | 0.01398 |
| Mullidae | <i>Mulloidichthys martinicus</i> | 16.5 | 98 | 28 | | | 0.02182 |
| | <i>Pseudupeneus maculatus</i> | 19.0 | 145 | 22 | | | 0.02114 |
| Muraenidae | <i>Gymnothorax funebris</i> | 108.6 | 1,588 | | | | 0.00124 |
| | <i>Lycodontis moringa</i> | 63.6 | 227 | | | | 0.00088 |
| Myiobatidae | <i>Aetobatus narinari</i> | | | | 382 Length 239 Width | 204,000 | |
| Ogcocephalidae | <i>Ogcocephalus nasutus</i> | 18.5 | 185 | | | | 0.0292 |
| Ostraciidae | <i>Lactophrys bicaudalis</i> | 10.1 | 63 | 42 | | | 0.06115 |
| | <i>Lactophrys trigonus</i> | 21.6 | 316 | 24 | | | 0.03136 |
| | <i>Lactophrys triqueter</i> | 13.5 | 127 | 27 + | | | 0.05162 |

continued...

Table 8.1.1 continued

| Family | Length Species | Observed | | Estimated | | a ^{b,c} |
|--------------------------|---------------------------------|-----------------------------|-----------------------|------------------------|------------------|------------------|
| | | Weight (cm) ^a | L _∞ (g) | W _∞ (cm) | (g) ^b | |
| Pomacanthidae | <i>Holacanthus ciliaris</i> | | | 64 | | |
| | <i>Holacanthus tricolor</i> | 11.5 | 84 | 32 | | 0.05523 |
| | <i>Pomacanthus arcuatus</i> | 24.0 | 794 | 64 | | 0.05744 |
| | <i>Pomacanthus paru</i> | 23.5 | 680 | 32 + | | 0.0524 |
| Pomacentridae | <i>Abudefduf saxatilis</i> | 10.8 | 76 | 16 | | 0.06033 |
| | <i>Chromis cyanea</i> | 8.5 | 26 | 13 / | | 0.04234 |
| | <i>Microspathodon chrysurus</i> | 10.0 | 53 | 16 | | 0.05149 |
| | | 11.5 | 76 | | | |
| | <i>Stegastes fuscus</i> | 7.5 | 21 | 16 / | | 0.04978 |
| Priacanthidae | <i>Priacanthus arenatus</i> | 24.0 | 357 | | | 0.02583 |
| Scaridae | <i>Scarus coelestinus</i> | 46.0 | 3,175 | 95 | | 0.03262 |
| | <i>Scarus guacamaia</i> | 50.0 | 3,856 | 95 | | 0.03085 |
| | <i>Scarus iserti</i> | | | 19 | | |
| | <i>Scarus taeniopterus</i> | 18.0 | 150 | | | 0.02572 |
| | <i>Sparisoma aurofrenatum</i> | 12.8 | 54 | 23 / | | 0.02653 ma |
| | | 13.0 | 60 | | | |
| | <i>Sparisoma chrysopterygum</i> | | | 32 / | | |
| Sciaenidae | <i>Equetus lanceolatus</i> | 14.0 | 57 | | | 0.02077 |
| | <i>Equetus punctatus</i> | 18.7 | 137 | | | 0.02095 |
| | <i>Odontoscion dentex</i> | 9.5 | 18 | 32 | | 0.02035 |
| | | 12.6 | 41 | | | |
| | | 13.1 | 44 | | | |
| Scombridae | <i>Auxis thazard</i> | 39.5 | 907 | | | 0.01472 |
| | <i>Euthynnus aletteratus</i> | 49.6 | 1,701 | | | |
| | <i>Scomberomorus maculatus</i> | 28.0 | 234 | 111 | 4,500 | |
| | | 74.5 | 4,309 | | | |
| | <i>Scomberomorus regalis</i> | | | 191 | 9,100 | |
| | <i>Thunnus obesus</i> | 62.0 | 5,443 | 159 | | |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | 8.7 | 24 | | | 0.03427 ma |
| | | 14.2 | 92 | | | |
| | <i>Scorpaena grandicornis</i> | 10.9 | 47 | | | 0.03629 |
| | <i>Scorpaena plumieri</i> | 11.6 | 85 | 32 / | | 0.05446 |
| Serranidae | <i>Alphesthes afer</i> | 14.5 | 102 | 32 + | | 0.03346 |
| | <i>Cephalopholis cruentata</i> | 5.9 | 6 | 32 | | |
| | | 16.6 | 114 | | | |
| | <i>Cephalopholis fulva</i> | 17.7 | 143 | 32 | | 0.02579 |
| | <i>Epinephelus guttatus</i> | 13.8 | 59 | 48 | | 0.0229 |
| | | 23.5 | 303 | | | |
| | <i>Epinephelus itajara</i> | 12.2 | 45 | 254 | 314,000 | |
| | <i>Epinephelus morio</i> | 26.5 | 460 | 95 | | 0.02472 |
| | <i>Epinephelus striatus</i> | | | 95 | 23,000 | |
| | <i>Mycteroperca venenosa</i> | 40.6 | 1,814 | 95 - | | 0.02711 |
| | <i>Paranthias furcifer</i> | 16.4 | 96 | 27 | | 0.02176 |
| | <i>Serranus tabacarius</i> | 10.5 | 26 | | | 0.02416 m |
| | | 10.8 | 29 | | | |
| | | 10.8 | 34 | | | |
| <i>Serranus tigrinus</i> | 7.3 | 8 | | | 0.02056 | |
| Sparidae | <i>Calamus bajonado</i> | 14.3 | 94 | 64 | 4,500 | |
| | <i>Calamus calamus</i> | 21.9 | 295 | 40 | | 0.02809 |
| Sphyraenidae | <i>Sphyraena barracuda</i> | 70.5 | 2,495 | 167 | 25,000 | 0.00712 |
| Synodontidae | <i>Synodus intermedius</i> | 22.5 | 129 | 40 | | 0.01133 |
| Tetraodontidae | <i>Sphoeroides spengleri</i> | 8.0 | 14 | | | 0.02734 |

^aType of length not stated.

^b'a' computed with equation (2.4); 'b' assumed equal to 3.0.

^cCalculated only when an estimate of 'a' or W_∞ was needed for combined asymptotic sizes from various authors (Table 3.5).

Symbols:

ma = mean 'a' from several L/W pairs.

/ = approximate maximum size.

+ = rather more than the recorded maximum size.

- = rather less than the recorded maximum size.

Table 8.1.2 Length-weight data of Caribbean reef fishes after Randall (1962, 1967, 1968).

| Family | Species | Length (cm) | Weight (g) | L _{max} (cm) | W _{max} (g)* | a* | b |
|-------------------------------|------------------------------------|-------------|------------|-----------------------|-----------------------|------------|----------|
| Acanthuridae | <i>Acanthurus bahianus</i> | | | 36 ! | | 0.0541 | 2.665 FL |
| | <i>Acanthurus chirurgus</i> | | | 34 ! | | | |
| | <i>Acanthurus coeruleus</i> | | | 37 ! | | | |
| Albulidae | <i>Albula vulpes</i> | 49.5 | 1,180 | 102 * | 8,600 * | 0.011 | 3 as |
| Antennariidae | <i>Antennarius multiocellatus</i> | | | 15 (-) | | | |
| | <i>Phrynelox scaber</i> | | | 15 / | | | |
| Apogonidae | <i>Apogon conklini</i> | | | 5 / | | | |
| | <i>Apogon maculatus</i> | | | 10 / | | | |
| Atherinidae | <i>Allanetta harringtonensis</i> | | | 8 / | | | |
| | <i>Atherinomorus stipes</i> | | | 9 + | | | |
| Aulostomidae | <i>Aulostomus maculatus</i> | | | 76 / | | | |
| Balistidae | <i>Balistes capriscus</i> | | | 30 (+) | | 0.0291 | 2.993 FL |
| | <i>Balistes vetula</i> | | | 57 / | | | |
| | <i>Canthidermis sufflamen</i> | | | 56 + | 4,500 - | | |
| | <i>Melichthys niger</i> | | | 36 | | | |
| | <i>Xanthichthys ringens</i> | | | 25 | | | |
| Belonidae | <i>Platybelone argalus argalus</i> | | | 51 / | | | |
| | <i>Strongylura timucu</i> | | | 46 / | | | |
| | <i>Tylosurus crocodilus</i> | | | 152 | | | |
| Blenniidae | <i>Entomacrodus nigricans</i> | | | 8+ | | | |
| | <i>Ophioblennius atlanticus</i> | | | 12/ | | | |
| | <i>Parablennius marmoreus</i> | | | 8 | | | |
| | <i>Scartella cristata</i> | | | 11 | | | |
| Bothidae | <i>Bothus lunatus</i> | | | 46 / | | | |
| | <i>Bothus ocellatus</i> | | | 18 | | | |
| | <i>Paralichthys tropicus</i> | | | 51 + | | | |
| Carangidae | <i>Alectis ciliaris</i> | 40.6 | 950 | 91 / | | 0.0142 | 3 as |
| | <i>Caranx bartholomaei</i> | | | 76 / | | | |
| | <i>Caranx crysos</i> | | | 58 ! | | | |
| | <i>Caranx hippos</i> | 85.6 | 6,210 | | 18,000 / | 0.0099 | 3 as |
| | <i>Caranx latus</i> | 71.9 | 3,860 | 76 ! | | 0.0122 | 3 as |
| | | 33.8 | 540 | | | | |
| | <i>Caranx lugubris</i> | 62.2 | 3,080 | 76 ! | | 0.0128 | 3 as |
| | <i>Caranx ruber</i> | | | 61 ! | | | |
| | <i>Decapterus macarellus</i> | | | 30 + | | | |
| | <i>Decapterus punctatus</i> | | | 30 (-) | | | |
| | <i>Elegatis bipinnulatus</i> | 41.9 | 450 | 119 | 13,600 | | |
| | <i>Oligoplites saurus</i> | | | 30 + | | | |
| | <i>Selar crumenophthalmus</i> | | | 41 / | | | |
| | <i>Seriola dumerili</i> | 111.8 | 14,060 | | 80,000 | 0.0108 ma | 3 as |
| | | 180.3 | 67,585 | | | | |
| <i>Seriola rivoliana</i> | 66.0 | 2,860 | | 23,000 (+) | | | |
| <i>Trachinotus falcatus</i> | 31.2 | 450 | 107 | 23,000 | | | |
| SL | 50.0 | 3,620 | | | | | |
| SL | 81.0 | 14,950 | | | | | |
| | 104.1 | 15,195 | | | | | |
| | | | | 37 | | | |
| | | | | 51 () | | | |
| Carcharhinidae | <i>Carcharhinus perezii</i> | | | 274 + | | | |
| | <i>Galeocerdo cuvier</i> | 279.4 | 117,930 | 549 | | 0.00804 ma | 3 as |
| | | 422.9 | 807,390 | 914 () | | | |
| | <i>Negaprion brevirostris</i> | 63.5 | 1,220 | 335 | | 0.0048 | 3 as |
| <i>Rhizoprionodon porosus</i> | 88.9 | 2,860 | 107 | | 0.0041 | 3 as | |
| Chaetodontidae | <i>Chaetodon aculeatus</i> | | | 9 ! | | | |
| | <i>Chaetodon capistratus</i> | | | 10 | | | |
| | | | | 15 () | | | |
| | <i>Chaetodon sedentarius</i> | | | 14 ! | | | |
| | <i>Chaetodon striatus</i> | | | 15 | | | |

continued...

Table 8.1.2 continued

| Family | Species | Length (cm) | Weight (g) | L _{max} (cm) | W _{max} (g) ^a | a ^a | b |
|---------------------------|------------------------------------|-------------|------------|-----------------------|-----------------------------------|----------------|------|
| Cirrhitidae | <i>Amblycirrhitus pinos</i> | | | 8 | | | |
| Clinidae | <i>Labrisomus nuchipinnis</i> | | | 20 / | | | |
| Clupeidae | <i>Harengula clupeola</i> | | | 14 | | | |
| | <i>Harengula humeralis</i> | | | 22 | | | |
| | <i>Jenkinsia lamprotaenia</i> | | | 6 | | | |
| | <i>Opisthonema oglinum</i> | | | 30 | | | |
| Congridae | <i>Heteroconger halis</i> | | | 48 | | | |
| Dactylopteridae | <i>Dactylopterus volitans</i> | | | 33 ! | | | |
| Dasyatidae | <i>Dasyatis americana</i> | | | 152 | Width | | |
| | <i>Urolophus jamaicensis</i> | | | 76 | | | |
| Diodontidae | <i>Chilomycterus antennatus</i> | | | 25 ! | | | |
| | <i>Chilomycterus antillarum</i> | | | 25 ! | | | |
| | <i>Diodon holocanthus</i> | | | 51 | | | |
| | <i>Diodon hystrix</i> | | | 57 ! | | | |
| Echeneidae | <i>Echeneis naucrates</i> | | | 81 ! | 1,800 ! | | |
| | <i>Remora remora</i> | | | 79 | | | |
| Emmelichthyidae | <i>Inermia vittata</i> | | | 23 / | | | |
| Engraulidae | <i>Anchoa lyolepis</i> | | | 8 | | | |
| Ephippidae | <i>Chaetodipterus faber</i> | 35.1 | 1,360 | 90 | 9,000 | | |
| Fistulariidae | <i>Fistularia tabacaria</i> | | | 183 | | | |
| Gerreidae | <i>Eucinostomus argenteus</i> | | | 19 ! | | | |
| | <i>Gerres cinereus</i> | 36.8 | 540 | 39 ! | | 0.0108 | 3 as |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | 160.0 | 23,590 | 427 () | | | |
| Gobiidae | <i>Bathygobius soporator</i> | | | 15 | | | |
| | <i>Coryphopterus glaucofraenum</i> | | | 8 | | | |
| | <i>Gobiosoma sp. (evelynae)</i> | | | 4 / | | | |
| | <i>Gnatholepis thompsoni</i> | | | 6 / | | | |
| Grammidae (Serranidae) | <i>Gramma loreto</i> | | | 8 + | | | |
| | <i>Gramma melacara</i> | | | 10 - | | | |
| Grammistidae (Serranidae) | <i>Rypticus saponaceus</i> | | | 28 ! | | | |
| Haemulidae (Pomadasyidae) | <i>Anisotremus surinamensis</i> | | | 60 ! | 4,100 ! | | |
| | <i>Anisotremus virginicus</i> | | | 30 / | | | |
| | <i>Haemulon album</i> | 46.7 | 1,540 | 65 ! | | 0.0151 | 3 as |
| | <i>Haemulon aurolineatum</i> | | | 25 - | | | |
| | <i>Haemulon bonariense</i> | | | 28 / | | | |
| | <i>Haemulon boschmae</i> | | | 19 / | | | |
| | <i>Haemulon carbonarium</i> | | | 36 ! | | | |
| | <i>Haemulon chrysargyreum</i> | | | 23 ! | | | |
| | <i>Haemulon flavolineatum</i> | | | 27 / | | 0.0197 | 3 as |
| | <i>Haemulon macrostomum</i> | 38.6 | 1,130 | 43 ! | | | |
| | <i>Haemulon melanurum</i> | | | 33 | | | |
| | <i>Haemulon parrai</i> | | | 41 | | | |
| | <i>Haemulon plumieri</i> | 26.4 | 450 | 41 | | 0.0245 | 3 as |
| | <i>Haemulon sciurus</i> | 29.7 | 450 | 39 ! | | 0.0172 | 3 as |
| | | | | 46 | | | |
| | <i>Haemulon steindachneri</i> | | | 27 | | | |
| | <i>Haemulon striatum</i> | | | 28 | | | |
| | <i>Orthopristis ruber</i> | | | 33 | | | |
| Hemiramphidae | <i>Hemiramphus balao</i> | | | 38 / | | | |
| | <i>Hemiramphus brasiliensis</i> | | | 38 / | | | |

continued...

Table 8.1.2 continued

| Family | Species | Length (cm) | Weight (g) | L _{max} (cm) | W _{max} (g) ^a | a ^a | b |
|----------------|----------------------------------|-------------|------------|-----------------------|-----------------------------------|----------------|----------|
| Holocentridae | <i>Holocentrus ascensionis</i> | | | 35 ! | | | |
| | <i>Holocentrus coruscus</i> | | | 13 | | | |
| | <i>Holocentrus rufus</i> | | | 32 + | | | |
| | <i>Myripristis jacobus</i> | | | 22 ! | | | |
| | <i>Neoniphon marianus</i> | | | 18 ! | | | |
| | <i>Plectrypops retrospinis</i> | | | 13 | | | |
| | <i>Sargocentron vexillarium</i> | | | 18 - | | | |
| Kyphosidae | <i>Kyphosus incisor</i> | 67.3 | 5,440 | 76 () | | 0.01785 | 3 as |
| | <i>Kyphosus sectatrix</i> | | | 76 ! | | | |
| Labridae | <i>Bodianus rufus</i> | | | 39 | | | |
| | | | | 61 () | | | |
| | <i>Clepticus parrae</i> | | | 25 | | | |
| | | | | 30 () | | | |
| | <i>Halichoeres bivittatus</i> | | | 22 ! | | | |
| | <i>Halichoeres garnoti</i> | | | 19 | | | |
| | <i>Halichoeres maculipinna</i> | | | 11 | | | |
| | <i>Halichoeres poeyi</i> | | | 20 | | | |
| | <i>Halichoeres radiatus</i> | | | 46 | | | |
| | <i>Lachnolaimus maximus</i> | 81.3 | 6,530 | | 11,000 () | 0.0121 | 3 as |
| | <i>Thalassoma bifasciatum</i> | | | 15 () | | | |
| | | | 22 ! | | | | |
| | <i>Xyrichtys novacula</i> | | | 14 | | | |
| | <i>Xyrichtys splendens</i> | | | | | | |
| Lutjanidae | <i>Lutjanus analis</i> | 38.1 | 770 | | 11,000 () | 0.0136 m | 3 as |
| | | 69.9 | 4,540 | | | | |
| | <i>Lutjanus apodus</i> | 53.3 | 2,495 | 56 ! | 3,600 () | 0.0218 | 2.939 FL |
| | <i>Lutjanus cyanopterus</i> | 90.2 | 12,380 | | 45,000 + | 0.0168 | 3 as |
| | <i>Lutjanus griseus</i> | 39.6 | 1,040 | | 3,600 | 0.0167 | 3 as |
| | <i>Lutjanus jocu</i> | 39.4 | 950 | 79 ! | 9,000 (+) | 0.0234 | 3 as |
| | SL | 63.0 | 7,825 | | | | |
| | <i>Lutjanus mahagoni</i> | | | 38 ! | | | |
| | <i>Lutjanus synagris</i> | | | 36 | | | |
| | <i>Ocyurus chrysurus</i> | 71.1 | 2,270 | 76 / | | 0.0063 | 3 as |
| Malacanthidae | <i>Malacanthus plumieri</i> | 52.8 | 730 | 61 / | | 0.0049 | 3 as |
| Megalopidae | <i>Tarpon atlanticus</i> | 36.6 | 1,590 | 244 + | 160,000 | | |
| Monacanthidae | <i>Aluterus monoceros</i> | | | 61 + | | | |
| | <i>Aluterus schoepfii</i> | | | 51 | | | |
| | <i>Aluterus scripta</i> | | | 72 ! | 2,500 ! | 0.0067 | 3 as |
| | | | | 91 | | | |
| | <i>Cantherines macrocerus</i> | | | 42 ! | | | |
| | <i>Cantherines pullus</i> | | | 18 ♀ | | | |
| | | | | 20 ♂ | | | |
| | <i>Monacanthus ciliatus</i> | | | 13 ! | | | |
| | <i>Monacanthus tuckeri</i> | | | 9 ! | | | |
| | <i>Stephanolepis setifer</i> | | | 18 ! | | | |
| Mugilidae | <i>Mugil curema</i> | | | 38 | | | |
| Mullidae | <i>Mulloidichthys martinicus</i> | 32.8 | 500 | 39 | | 0.0141 | 3 as |
| | <i>Pseudupeneus maculatus</i> | | | 28 / | | | |
| Muraenidae | <i>Enchelycore nigricans</i> | | | 76 + | | | |
| | <i>Echidna catenata</i> | | | 71 | | | |
| | <i>Gymnothorax funebris</i> | 83.8 | 1,500 | 189 | 12,000 | 0.0025 | 3 as |
| | <i>Gymnothorax miliaris</i> | | | 51 / | | | |
| | <i>Gymnothorax vicinus</i> | | | 122 / | | | |
| | <i>Lycodontis moringa</i> | | | 122 / | | | |
| Myliobatidae | <i>Aetobatus narinari</i> | 91.4 | 34,470 | 230 Width | | 0.0451 | 3 as |
| Ogcocephalidae | <i>Ogcocephalus nasutus</i> | | | 28 / | | | |
| Ophichthidae | <i>Myrichthys breviceps</i> | | | 91 + | | | |
| | <i>Myrichthys ocellatus</i> | | | 91 / | | | |
| | <i>Ophichthus ophis</i> | 122.0 | 1,400 | 135 | | 0.0008 ma | 3 as |
| | | 116.3 | 1,470 | | | | |

continued...

Table 8.1.2 continued

| Family | Species | Length (cm) | Weight (g) | L _{max} (cm) | W _{max} (g) ^a | a ^a | b |
|-----------------------------|-------------------------------------|----------------|---------------|--------------------------|--------------------------------------|----------------|------|
| Opisthognathidae | <i>Opisthognathus aurifrons</i> | | | 10 - | | | |
| | <i>Opisthognathus macrognathus</i> | | | 20 - | | | |
| | <i>Opisthognathus maxillosus</i> | | | 13 / | | | |
| | <i>Opisthognathus whitehurstii</i> | | | 8 ! | | | |
| Ostraciidae | <i>Acanthostracion polygonus</i> | 770 | | 39 ! | | | |
| | <i>Acanthostracion quadricomis</i> | 4,540 | | 46 / | | | |
| | <i>Lactophrys bicaudalis</i> | 12,380 | | 41 + | | | |
| | <i>Lactophrys trigonus</i> | 2,495 | | 46 + | | | |
| | <i>Lactophrys triqueter</i> | | | 30 / | | | |
| Pempheridae | <i>Pempheris poeyi</i> | | | 10 / | | | |
| | <i>Pempheris schomburgki</i> | | | 15 | | | |
| Pomacanthidae | <i>Centropyge argi</i> | | | 7 | | | |
| | <i>Holacanthus ciliaris</i> | 42.7 | 1,590 | 46 | | 0.0204 | 3 as |
| | <i>Holacanthus tricolor</i> | | | 25 | | | |
| | <i>Pomacanthus arcuatus</i> | | | 30 | | | |
| | <i>Pomacanthus paru</i> | | | 43 | | | |
| Pomacentridae | <i>Abudefduf saxatilis</i> | | | 18 + | | | |
| | <i>Abudefduf taurus</i> | | | 20 / | | | |
| | <i>Chromis cyanea</i> | | | 13 / | | | |
| | <i>Chromis multilineata</i> | | | 16 ! | | | |
| | <i>Microspathodon chrysurus</i> | | | 19 / | | | |
| | <i>Stegastes fuscus</i> | | | 13 ! | | | |
| | | | | 15 / | | | |
| | <i>Stegastes leucostictus</i> | | | 10 | | | |
| | <i>Stegastes planifrons</i> | | | 13 / | | | |
| | <i>Stegastes variabilis</i> | | | 10 + | | | |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | | | 30 / | | | |
| | <i>Priacanthus arenatus</i> | | | 41 / | | | |
| Rachycentridae | <i>Rachycentron canadum</i> | 56.6 | 1,720 | 178 | 46,000 | 0.0088 ma | 3 as |
| Scaridae | <i>Cryptotomus roseus</i> | | | 13 () | | | |
| | <i>Nicholsina usta</i> | | | 29 ! | | | |
| | <i>Scarus coelestinus</i> | 51.6 | 2,400 | 76 ! | 7,000 ! | 0.0167 ma | 3 as |
| | <i>Scarus coeruleus</i> | 62.5 | 3,360 | 58 ! ♂ | 3,400 ! ♂ | 0.0156 ma | 3 as |
| | | | | 91 | | | |
| | <i>Scarus guacamaia</i> | 57.2 | 3,220 | 94 ! | 20,000 ! | | |
| | <i>Scarus iserti</i> | | | 25 | | | |
| | <i>Scarus taeniopterus</i> | | | 30 / | | | |
| | <i>Scarus vetula</i> | | | 51 ! | | | |
| | <i>Sparisoma aurofrenatum</i> | | | 25 + | | | |
| | <i>Sparisoma chrysopteron</i> | | | 41 ! ♀ | | | |
| | | | | 51 / ♂ | | | |
| | <i>Sparisoma radians</i> | | | 19 ! | | | |
| <i>Sparisoma rubripinne</i> | | | 34 ! ♀ | | 0.0031 | 3.562 FL | |
| | | | 44 ! ♂ | | | | |
| <i>Sparisoma viride</i> | | | 51 + | 1,600 + | | | |
| Sciaenidae | <i>Equetus lanceolatus</i> | | | 23 / | | | |
| | <i>Equetus punctatus</i> | | | 27 ! | | | |
| | <i>Odontoscion dentex</i> | | | 18 + | | | |
| | <i>Pareques acuminatus</i> | | | 23 () | | | |
| Scombridae | <i>Euthynnus aletteratus</i> | 32.5 | 450 | 91 / | 9,100 + | | |
| | <i>Scomberomorus cavalla</i> | 110.7 | 10,660 | 165 + | 36,000 / | | |
| | <i>Scomberomorus maculatus</i> | 53.3 | 860 | 122 () | 9,100 () | | |
| | <i>Scomberomorus regalis</i> | 41.4 | 450 | 81 ! | 12,000 () | | |
| | | | 122 () | | | | |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | | | 36 / | | | |
| | <i>Scorpaena grandicomis</i> | | | 18 / | | | |
| | <i>Scorpaena plumieri</i> | | | 43 / | | | |
| | <i>Scorpaenodes caribbaeus</i> | | | 13 / | | | |

continued...

Table 8.1.2 continued

| Family | Species | Length (cm) | Weight (g) | L _{max} (cm) | W _{max} (g) ^a | a ^a | b | |
|----------------------------|------------------------------------|-------------|------------|-----------------------|-----------------------------------|----------------|----------|----------|
| Serranidae | <i>Alphesthes afer</i> | | | 30 / | | | | |
| | <i>Cephalopholis cruentata</i> | | | 30 / | | | | |
| | <i>Cephalopholis fulva</i> | | | 41 - | | | | |
| | <i>Dermatolepis inermis</i> | | | 81 ! | 9,100 ! | 0.0171 | 3 as | |
| | <i>Diplectrum formosum</i> | | | 30 () | | | | |
| | <i>Epinephelus adscensionis</i> | 33.8 | 590 | 61 / | | 0.0158 m | 3 as | |
| | <i>Epinephelus guttatus</i> | | | 55 | | | | |
| | <i>Epinephelus itajara</i> | 165.1 | 154,220 | 217 ! | 308,000 ! | 0.0322 m | 3 as | |
| | <i>Epinephelus morio</i> | 42.7 | 1,090 | | 23,000 (+) | 0.0140 | 3 as | |
| | <i>Epinephelus striatus</i> | 40.4 | 950 | | 25,000 () | 0.0033 | 3.425 FL | |
| | <i>Hypoplectrus aberrans</i> | | | 12 | | | | |
| | <i>Hypoplectrus chlorurus</i> | | | 13 | | | | |
| | <i>Hypoplectrus nigricans</i> | | | 15 | | | | |
| | <i>Hypoplectrus puella</i> | | | 13-VI | | | | |
| | <i>Mycteroperca bonaci</i> | 39.6 | 910 | | 15-Florida | 82,000 | | |
| | <i>Mycteroperca cidi</i> | | | 114 | | | | |
| | <i>Mycteroperca interstitialis</i> | | | 69 | | | | |
| | <i>Mycteroperca phenax</i> | | | 61 (-) | | | | |
| | <i>Mycteroperca rubra</i> | | | 70 | | | | |
| | <i>Mycteroperca tigris</i> | 35.6 | 635 | 102 | | | 0.0160 m | 3 as |
| | | 71.1 | 5,900 | | | | | |
| | <i>Mycteroperca venenosa</i> | | | 91 + | | | 0.0045 | 3.325 FL |
| | <i>Paranthias furcifer</i> | | | 27 + | | | | |
| <i>Serranus dewegeri</i> | | | 32 | | | | | |
| <i>Serranus tabacarius</i> | | | 18 | | | | | |
| <i>Serranus tigrinus</i> | | | 10 | | | | | |
| <i>Serranus tortugarum</i> | | | 8 | | | | | |
| Sparidae | <i>Archosargus rhomboidalis</i> | | | 33 | | | | |
| | <i>Calamus bajonado</i> | | | 56 | | | | |
| | <i>Calamus calamus</i> | | | 37 | | | | |
| | <i>Calamus penna</i> | | | 41 + | | | | |
| | <i>Calamus pennatula</i> | | | 37 | | | | |
| | <i>Diplodus caudimacula</i> | | | 30 / | | | | |
| Sphyraenidae | <i>Sphyraena barracuda</i> | 167.6 | 46,720 | 183 (-) | | 0.0099 | 3 as | |
| | <i>Sphyraena picudilla</i> | | | 46 (+) | | | | |
| Sphyrnidae | <i>Sphyrna lewini</i> | 276.9 | 108,860 | 305 | | 0.0051 | 3 as | |
| | <i>Sphyrna mokarran</i> | 80.0 | 1,950 | 549 + | | 0.0038 | 3 as | |
| Synodontidae | <i>Synodus foetens</i> | | | 46 | | | | |
| | <i>Synodus intermedius</i> | | | 46 | | | | |
| | <i>Synodus synodus</i> | | | 33 | | | | |
| Tetraodontidae | <i>Canthigaster rostrata</i> | | | 11 | | | | |
| | <i>Sphoeroides spengleri</i> | | | 30 (+) | | | | |
| | <i>Sphoeroides testudineus</i> | | | | | | | |

^aCalculated only when an estimate of 'a' or W_∞ was needed for combined asymptotic sizes from various authors (Table 3.5).

Symbols:

! = specimen measured by Randall.

* = world record (hook and line).

/ = approximate maximum size.

+ = rather more than the recorded size.

- = rather less than the recorded size.

() = 'said to reach' the recorded size.

FL = fork length.

SL = standard length.

ma = mean 'a' from several L/W pairs.

as = 'b' from equation (2.3), assumed equal to 3.0.

VI = Virgin Islands.

♀ = female.

♂ = male.

Table 8.1.3. Length-weight data of Caribbean reef fishes after Smith and Tyler (1972).

| Fish | Species | Length (cm) ^a | Weight (g) | a ^b |
|------------------------------|------------------------------------|--------------------------|------------|----------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 2.3 juvenile | 0.50 | 0.0411 |
| | <i>Acanthurus coeruleus</i> | 2.6 | 0.90 | 0.0512 |
| Apogonidae | <i>Apogon binotatus</i> | 3.6 ♂ | 1.50 | 0.0322 |
| | | 3.8 ♀ | 1.80 | 0.0328 |
| | | 4.0 ♀ | 1.90 | 0.0297 |
| | | 4.0 ♀ | 2.00 | 0.0313 |
| | <i>Apogon conklini</i> | 2.5 ♂ | 0.40 | 0.0256 |
| | | 3.2 ♀ | 0.90 | 0.0275 |
| | <i>Apogon maculatus</i> | 4.6 ♂ | 3.70 | 0.0380 |
| | | 5.0 ♀ | 4.40 | 0.0352 |
| | | 5.1 ♀ | 5.00 | 0.0377 |
| | <i>Apogon pigmentaria</i> | 1.8 ♂ | 0.20 | 0.0343 |
| | | 2.6 ♀ | 0.50 | 0.0284 |
| | | 3.0 ♂ | 0.70 | 0.0259 |
| | <i>Apogon quadrisquamatus</i> | 1.7 ♂ | 0.30 | 0.0611 |
| | | 3.1 ♀ | 1.10 | 0.0369 |
| | <i>Apogon townsendi</i> | 3.6 ♂ | 1.30 | 0.0279 |
| | | 4.0 ♀ | 1.70 | 0.0266 |
| | <i>Apogon xenus</i> | 2.4 ♂ | 0.40 | 0.0289 |
| 3.5 | | 1.20 | 0.0280 | |
| 3.6 | | 1.20 | 0.0257 | |
| | 4.4 ♀ | 2.40 | 0.0282 | |
| Blenniidae | <i>Hypleurocheilus</i> sp. | 4.0 ? | 1.00 | 0.0156 |
| Canthigasteridae | <i>Canthigaster rostrata</i> | 1.3 ♂ | 0.10 | 0.0455 |
| | | 2.6 | 1.50 | 0.0853 |
| | | 5.8 ♂ | 10.50 | 0.0538 |
| Cirrihitidae | <i>Amblycirrhitus pinos</i> | 2.3 | 0.40 | 0.0329 |
| | | 3.4 ♀ | 1.00 | 0.0254 |
| Clinidae | <i>Acanthemblemaria spinosa</i> | 2.0 | 0.10 | 0.0125 |
| | | 2.0 | 0.10 | 0.0125 |
| | <i>Emblemaria leptocirris</i> | 2.0 ♂ | 0.40 | 0.0500 |
| | | 2.3 ♀ | 0.50 | 0.0411 |
| | <i>Labrisomus haitiensis</i> | 3.6 | 0.80 | 0.0171 |
| | | 1.4 ♀ | 0.10 | 0.0364 |
| | <i>Pseudemblemaria signifera</i> | 1.5 ♂ | 0.10 | 0.0296 |
| | | 1.6 ♀ | 0.10 | 0.0244 |
| | | 1.7 ♂ | 0.10 | 0.0204 |
| | | 2.0 | 0.20 | 0.0250 |
| <i>Starksia hassi</i> | 2.1 | 0.20 | 0.0216 | |
| | 2.2 | 0.20 | 0.0188 | |
| Starksia lepicoelia | | | | |
| Emmelichthyidae | <i>Inermia vittata</i> | 4.1 | 1.60 | 0.0232 |
| Gobiidae | <i>Coryphopterus dicrus</i> | 1.7 ♀ | 0.10 | 0.0204 |
| | | 2.4 ♀ | 0.30 | 0.0217 |
| | | 2.5 ♂ | 0.30 | 0.0192 |
| | <i>Coryphopterus eidolon</i> | 1.8 | 0.10 | 0.0171 |
| | | 3.2 ♂ | 0.60 | 0.0183 |
| | | 3.9 ♀ | 1.10 | 0.0185 |
| | | 4.4 | 1.40 | 0.0164 |
| | <i>Coryphopterus glaucofraenum</i> | 3.4 ♂ | 0.80 | 0.0204 |
| | | 3.4 ♀ | 0.90 | 0.0229 |
| | | 3.8 ♀ | 1.00 | 0.0182 |
| | <i>Coryphopterus personatus</i> | 1.7 ♀ | 0.10 | 0.0204 |
| | | 2.3 ♀ | 0.30 | 0.0247 |
| | | 2.4 ♀ | 0.30 | 0.0217 |
| | | 2.6 ♂ | 0.30 | 0.0171 |
| | | 2.8 ♀ | 0.30 | 0.0137 |
| | | 2.9 ♂ | 0.50 | 0.0205 |
| | <i>Coryphopterus thrix</i> | 2.7 ♀ | 0.30 | 0.0152 |
| | | 2.8 ♀ | 0.40 | 0.0182 |
| | | 4.0 ♂ | 1.00 | 0.0156 |
| <i>Gnatholepis thompsoni</i> | 4.0 ♂ | 1.00 | 0.0156 | |
| | 5.2 ♀ | 3.20 | 0.0228 | |

continued...

Table 8.1.3 continued

| Fish | Species | Length (cm) ^a | Weight (g) | a ^b |
|------------------------------|-------------------------------|--------------------------|------------|----------------|
| | <i>Gobiosoma evelynae</i> | 1.8 ? | 0.10 | 0.0171 |
| | | 2.2 ?♂ | 0.20 | 0.0188 |
| | | 2.6 ?♀ | 0.30 | 0.0171 |
| | <i>Gobiosoma horsti</i> | 2.3 ? | 0.30 | 0.0247 |
| | <i>Gobiosoma saucrum</i> | 1.4 ♀ | 0.10 | 0.0364 |
| | | 1.6 ♀ | 0.10 | 0.0244 |
| | | 1.7 ♂ | 0.10 | 0.0204 |
| | | 2.0 ♂ | 0.10 | 0.0125 |
| | <i>Lythrypnus elasson</i> | 0.8 | 0.02 | 0.0391 |
| | | L _{max} 1.3 | 0.02 | 0.0091 |
| | <i>Lythrypnus nesiotis</i> | 0.7 | 0.02 | 0.0583 |
| | | 1.2 | 0.02 | 0.0116 |
| | | L _{max} 1.4 | | |
| | <i>Lythrypnus</i> sp. | 0.9 | 0.10 | 0.1372 |
| | <i>Quisquilius hipoliti</i> | 2.0 | 0.20 | 0.0250 |
| | | 2.2 ♂ | 0.30 | 0.0282 |
| | <i>Risor ruber</i> | 1.0 ♂ | 0.10 | 0.1000 |
| | | 1.1 ♀ | 0.10 | 0.0751 |
| | | 1.2 ♂ | 0.10 | 0.0579 |
| Grammistidae (Serranidae) | <i>Rypticus subbifrenatus</i> | 6.4 | 7.00 | 0.0267 |
| Haemulidae (Pomadasyidae) | <i>Haemulon aurolineatum</i> | 3.0 | 0.50 | 0.0185 |
| | | 4.9 | 2.50 | 0.0212 |
| | <i>Haemulon flavolineatum</i> | 3.0 | 0.90 | 0.0333 |
| Holocentridae | <i>Holocentrus coruscus</i> | 4.9 | 3.40 | 0.0289 |
| | <i>Holocentrus rufus</i> | 6.1 | 4.80 | 0.0211 |
| | <i>Myripristis jacobus</i> | 3.7 | 2.00 | 0.0395 |
| | | 3.9 | 2.50 | 0.0421 |
| | <i>Neoniphon marianus</i> | 4.5 | 2.60 | 0.0285 |
| | | 6.4 | 7.40 | 0.0282 |
| Labridae | <i>Halichoeres garnoti</i> | 4.2 ♀ | 1.50 | 0.0202 |
| | | 6.0 ♂ | 4.10 | 0.0190 |
| | | 7.3 ♀ | 6.70 | 0.0172 |
| | <i>Thalassoma bifasciatum</i> | 1.7 | 0.10 | 0.0204 |
| | | 2.1 | 0.20 | 0.0216 |
| | | 2.8 ♂ | 0.40 | 0.0182 |
| | | 2.8 ♀ | 0.50 | 0.0228 |
| | | 3.0 ♂ | 0.80 | 0.0296 |
| | | 4.5 ♀ | 1.80 | 0.0198 |
| | | 5.0 ♂ | 2.40 | 0.0192 |
| | | 7.5 ♂ | 7.50 | 0.0178 |
| Monacanthidae | <i>Monacanthus tuckeri</i> | 1.6 ?juvenile | 0.10 | 0.0244 |
| Muraenidae | <i>Gymnothorax miliaris</i> | 32.5 TL♀ | 55.40 | 0.0016 |
| | <i>Lycodontis moringa</i> | 25.9 TL | 17.90 | 0.0010 |
| Ostraciidae | <i>Lactophrys triqueter</i> | 10.0 | 65.00 | 0.0650 |
| Pomacanthidae | <i>Holacanthus ciliaris</i> | 2.3 | 0.50 | 0.0411 |
| | <i>Pomacanthus arcuatus</i> | 1.8 | 0.30 | 0.0514 |
| Pomacentridae | <i>Chromis cyanea</i> | 3.7 ♂ | 2.40 | 0.0474 |
| | | 8.1 ♀ | 22.20 | 0.0418 |
| | <i>Chromis multilineata</i> | 5.0 | 5.30 | 0.0424 |
| | <i>Pomacentrus partitus</i> | 3.0 | 1.20 | 0.0444 |
| | | 4.1 | 2.20 | 0.0319 |
| | | 4.4 | 3.80 | 0.0446 |
| | <i>Stegastes planifrons</i> | 5.7 ♀ | 9.50 | 0.0513 |
| | <i>Stegastes variabilis</i> | 2.4 | 0.80 | 0.0579 |
| | | 3.5 | 2.10 | 0.0490 |
| | | 3.8 ♀ | 3.30 | 0.0601 |
| | | 5.1 ♂ | 6.30 | 0.0475 |
| Scaridae | <i>Scarus vetula</i> | 23.0 ♂ | 430.00 | 0.0353 |

continued...

Table 8.1.3 continued

| Fish | Species | Length (cm) ^a | Weight (g) | a ^b |
|--------------|--------------------------------|--------------------------|------------|----------------|
| Serranidae | <i>Cephalopholis cruentata</i> | 25.0 | 425.00 | 0.0272 |
| | | 30.0 | 600.00 | 0.0222 |
| | <i>Epinephelus striatus</i> | 30.0 | 400.00 | 0.0148 |
| | <i>Hypoplectrus puella</i> | 2.2 | 0.30 | 0.0282 |
| | | 3.5 | 1.30 | 0.0303 |
| Syngnathidae | <i>Syngnathus</i> sp. | 3.6 TL | 0.10 | 0.0021 |
| Synodontidae | <i>Synodus intermedius</i> | 17.0 | 82.00 | 0.0167 |

^a'Length' refers to standard length unless stated otherwise.

^b'a' computed with equation (2.4); 'b' assumed equal to 3.0.

Symbols:

M = mean length from female and male.

♀ = female.

TL = total length.

♂ = male.

? = type of length not recorded.

Table 8.1.4. Length-weight data of fishes from the southeastern Atlantic coast of Brazil after Figueredo and Menezes (1977).

| Family | Species | Length (cm) | Weight (g) | L _∞ (cm) | W _∞ (g) ^a | a ^{a,b} |
|------------------------------|---------------------------------|-------------|------------|---------------------|---------------------------------|------------------|
| Albulidae | <i>Albula vulpes</i> | 35 ! | | 100 | | |
| Antennariidae | <i>Antennarius striatus</i> | | | 15 | | |
| Carangidae | <i>Alectis ciliaris</i> | | | 100 + | | |
| | <i>Caranx bartholomaei</i> | | | 90 + | 7,500 | |
| | <i>Caranx crysos</i> | | | 70 - | | |
| | <i>Caranx hippos</i> | | | 100 + | 25,000 | |
| | <i>Caranx latus</i> | | | 80 / | | |
| | <i>Caranx lugubris</i> | | | 100 - | 7,000 + | |
| | <i>Caranx ruber</i> | | | 60 + | | |
| | <i>Decapterus macarellus</i> | | | 40 / | | |
| | <i>Decapterus punctatus</i> | | | 30 / | | |
| | <i>Oligoplites saurus</i> | | | 30 / | | |
| | <i>Selar crumenophthalmus</i> | | | 40 / | | |
| | <i>Selene vomer</i> | | | 50 / | | |
| | <i>Seriola dumerili</i> | | | 170 | 80,000 | |
| | <i>Seriola rivoliana</i> | | | 100 | 20,000 | |
| | <i>Trachinotus falcatus</i> | | | 120 | 40,000 | |
| <i>Trachinotus goodei</i> | | | 50 | | | |
| Carcharhinidae | <i>Galeocерdo cuvieri</i> | 400 | 525,000 | 550 / | | 0.0082 |
| | <i>Negaprion brevirostris</i> | 250 | 80,000 | 300 / | | |
| | <i>Prionace glauca</i> | 270 | 70,000 | 400 / | | 0.0036 |
| | <i>Rhizoprionodon porosus</i> | | | 100 + | 4,500 | |
| Clupeidae | <i>Harengula clupeola</i> | 17 ! | | | | |
| | <i>Opisthonema oglinum</i> | | | 30 / | | |
| Coryphaenidae | <i>Coryphaena hippurus</i> | 118 ! | | 200 | | |
| Dactylopteridae | <i>Dactylopterus volitans</i> | 36 ! | | 45 / | | |
| Dasyatidae | <i>Dasyatis americana</i> | | | 150 Width | | |
| Echeneidae | <i>Echeneis naucrates</i> | 65 ! | | 100 / | | |
| | <i>Remora remora</i> | | | 60 + | | |
| Elopidae | <i>Elops saurus</i> | | | 90 / | | |
| Engraulidae | <i>Anchoa lyolepis</i> | 8.3 ! | | | | |
| Fistulariidae | <i>Fistularia tabacaria</i> | | | 180 | | |
| Gerreidae | <i>Eucinostomus argenteus</i> | 28 ! | | 30 / | | |
| Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | 250 | 250,000 | 450 | | 0.0160 |
| Haemulidae (Pomadasyidae) | <i>Anisotremus surinamensis</i> | 35 | | 50 + | | |
| | <i>Anisotremus virginicus</i> | 31.5 ! | | 35 / | | |
| | <i>Haemulon aurolineatum</i> | 23.5 ! | | 25 / | | |
| | <i>Haemulon plumieri</i> | 28.5 ! | | 40 | | |
| | <i>Haemulon steindachneri</i> | 28.3 ! | | 30 / | | |
| | <i>Orthopristis ruber</i> | 28.8 ! | | 40 / | | |
| | <i>Pomadasys corvinaeformis</i> | 24.8 ! | | 25 / | | |
| | <i>Pomadasys crocro</i> | 21.5 ! | | 30 / | | |
| Holocentridae | <i>Holocentrus ascensionis</i> | | | 34 + | | |
| | <i>Myripristis jacobus</i> | 22.5 ! | | | | |
| Lutjanidae | <i>Lutjanus analis</i> | 35 ! | | 80 / | | |
| | <i>Lutjanus cyanopterus</i> | 120 ! | | | | |
| | <i>Lutjanus griseus</i> | 143 ! | | | | |
| | <i>Lutjanus jocu</i> | 49 ! | | 70 / | | |
| | <i>Lutjanus synagris</i> | 40.5 ! | | 40 / | | |
| | <i>Lutjanus vivanus</i> | 34.3 ! | | 50 + | | |
| | <i>Ocyurus chrysurus</i> | 42.5 ! | | 70 / | | |
| | <i>Rhomboplites aurorubens</i> | 44.5 ! | | 50 + | | |
| Malacanthidae | <i>Malacanthus plumieri</i> | 46.5 ! | | 60 | | |

continued...

Table 8.1.4 continued

| Family | Species | Length (cm) | Weight (g) | L _∞ (cm) | W _∞ (g) ^a | a ^{a,b} |
|----------------------------|-------------------------------------|-------------|------------|---------------------|---------------------------------|------------------|
| Megalopidae | <i>Tarpon atlanticus</i> | | | 250 | 150,000 | |
| Muraenidae | <i>Echidna catenata</i> | | | 100 - | | |
| | <i>Gymnothorax funebris</i> | | | 200 / | | |
| | <i>Gymnothorax miliaris</i> | 50 ! | | | | |
| | <i>Gymnothorax vicinus</i> | | | 100 | | |
| | <i>Lycodontis moringa</i> | | | 100 | | |
| Myliobatidae | <i>Aetobatus narinari</i> | | | 200 Width | 200,000 | |
| Ophichthidae | <i>Ophichthus ophis</i> | | | 150 ? | | |
| Pomatomidae | <i>Pomatomus saltator</i> | | | 100 + | | |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | | | 30 | | |
| | <i>Priacanthus arenatus</i> | | | 40 | | |
| Rachycentridae | <i>Rachycentron canadum</i> | | | 200 / | | |
| Sciaenidae | <i>Equetus lanceolatus</i> | | | 25 / | | |
| | <i>Odontoscion dentex</i> | 22 ! | | 25 / | | |
| | <i>Pareques acuminatus</i> | 17 ! | | 25 / | | |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | 21.5 ! | | | | |
| | <i>Scorpaena plumieri</i> | | | 35 / | | |
| Serranidae | <i>Alphestes afer</i> | | | 33 | | |
| | <i>Diplectrum formosum</i> | | | 30 / | | |
| | <i>Epinephelus adscensionis</i> | | | 60 / | | |
| | <i>Epinephelus itajara</i> | 270 | 375,000 | | 400,000 | |
| | <i>Epinephelus morio</i> | | | 70 + | 12,000+ | |
| | <i>Mycteroperca bonaci</i> | | | 100 + | 90,000 | |
| | <i>Mycteroperca interstitialis</i> | | | 70 + | 4,000 | |
| | <i>Mycteroperca microlepis</i> | | | 70 / | | |
| | <i>Mycteroperca rubra</i> | | | 80 / | | |
| | <i>Mycteroperca tigris</i> | 75 | 6,000 | 100 / | | 0.0142 |
| | <i>Mycteroperca venenosa</i> | | | 100 - | 12,000 | |
| <i>Paranthias furcifer</i> | | | 20 / | | | |
| Sparidae | <i>Archosargus rhomboidalis</i> | 35.5 ! | | 35 | | |
| | <i>Calamus penna</i> | 50 ! | | | | |
| | <i>Calamus pennatula</i> | 35 ! | | | | |
| Sphyrnidae | <i>Sphyrna lewini</i> | 280 | 125,000 | 300 + | | 0.0057 |
| | <i>Sphyrna mokarran</i> | 370 ♀ | 240,000 | 550 | | 0.0047 |
| Synodontidae | <i>Synodus foetens</i> | 36.5 ! | | | | |
| | <i>Synodus intermedius</i> | | | 50 | | |

^aCalculated only when an estimate of 'a' or W_∞ was needed for combined asymptotic sizes from various authors (Table 3.5).

^b'a' computed with equation (2.4); 'b' assumed equal to 3.0.

Symbols:

! = specimen measured by the authors.

/ = approximate maximum size.

+ = rather more than the recorded size.

- = rather less than the recorded size.

Table 8.1.5. Length-weight data of Caribbean reef fishes after FAO species identification sheets (Fischer 1978).

| Family | Species | Common length (cm) | Common weight (g) | Maximum length (cm) | Maximum weight (g) |
|-------------------------------|------------------------------------|---------------------------|-------------------|-----------------------------|--------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 25.0 | | 36.0 | |
| | <i>Acanthurus chirurgus</i> | 25.0 | | 35.0 | |
| | <i>Acanthurus coeruleus</i> | 25.0 | | 36.0 | |
| Albulidae | <i>Albula vulpes</i> | 35.0 | | 77.0 | |
| Balistidae | <i>Balistes capriscus</i> | 20.0 | | 30.0 | |
| | <i>Balistes vetula</i> | 30.0 | | 50.0 | |
| | <i>Melichthys niger</i> | 30.0 | | 50.0 | |
| Belonidae | <i>Ablennes hians</i> | 70.0 SL | | 82.5 SL+ | |
| | <i>Platybelone argalus argalus</i> | 30.0 SL | | 38.2 SL+ | |
| | <i>Strongylura marina</i> | 50.0 SL | | 64.0 SL | |
| | <i>Strongylura timucu</i> | 30.0 SL | | 42.0 SL | |
| | <i>Tylosurus acus</i> | 90.0 SL | | 128.5 SL | |
| | <i>Tylosurus crocodilus</i> | 90.0 SL | | 101.3 SL+ | |
| Bothidae | <i>Bothus lunatus</i> | 35.0 | | 45.0 | |
| | <i>Paralichthys tropicus</i> | 30.0 | | 50.0 + | |
| Carangidae | <i>Alectis ciliaris</i> | 90.0 FL | | 150.0 | |
| | | 84.5 FL | 7,900 | | |
| | | 109.0 FL | 16,500 | | |
| | <i>Caranx bartholomaei</i> | 45.0 FL | | 100.0 | |
| | | 89.5 | 7,600 | | |
| | <i>Caranx crysos</i> | 35.0 FL | | 68.0 | |
| | <i>Caranx hippos</i> | 60.0 FL | | 101.0 | 25,000 |
| | <i>Caranx latus</i> | 50.0 FL | | 80.0 + | 16,000 |
| | <i>Caranx lugubris</i> | 70.0 FL | | 99.0 | 7,000 |
| | <i>Caranx ruber</i> | 40.0 FL | | 50.0 + | 8,200 () |
| | <i>Decapterus macarellus</i> | 20.0 SL | | 35.0 SL | |
| | <i>Decapterus punctatus</i> | 15.0 FL | | 30.0 SL | |
| | <i>Elagatis bipinnulatus</i> | 80.0 FL | | 107.0 FL | 10,500 |
| | <i>Oligoplites saurus</i> | 27.0 FL | | 29.7 FL | 287 |
| | <i>Selar crumenophthalmus</i> | 24.0 FL | | 27.0 SL | |
| | <i>Seriola dumerilii</i> | 70.0 FL | 2,000 | | 80,300 |
| | | 110.0 FL | 8,000 | | |
| 150.0 FL | | 67,600 | | | |
| <i>Seriola rivoliana</i> | 55.0 FL | 2,500 | 97.0 | 24,000 | |
| | 80.0 FL | 3,400 | | | |
| | 83.0 FL | 9,800 | | | |
| <i>Trachinotus falcatus</i> | 94.0 | 17,000 | 105.5 FL | 36,000 | |
| | 114.0 | 22,900 | | | |
| <i>Trachinotus goodei</i> | 31.0 FL | | 49.3 | | |
| Carcharhinidae | <i>Carcharhinus acronotus</i> | 100.0 | | 200.0 | |
| | <i>Carcharhinus falciformis</i> | 250.0 | | 350.0 | |
| | <i>Carcharhinus leucas</i> | 260.0 | | 350.0 | |
| | <i>Carcharhinus limbatus</i> | 150.0 | | 247.0 | |
| | <i>Carcharhinus longimanus</i> | 270.0 | | 350.0 | |
| | <i>Carcharhinus perezi</i> | 150.0 | | 230.0 | |
| | <i>Galeocerdo cuvier</i> | 400.0 | | 650.0 + | |
| | <i>Negaprion brevirostris</i> | 240.0 | | 320.0 | |
| | Clupeidae | <i>Harengula clupeola</i> | 9.0 | | 10.0 |
| <i>Jenkinsia lamprotaenia</i> | | 5.0 | | 6.5 | |
| <i>Opisthonema oglinum</i> | | 20.0 | | 25.0 | |
| Coryphaenidae | <i>Coryphaena hippurus</i> | 100.0 | | 200.0 | |
| Dactylopteridae | <i>Dactylopterus volitans</i> | 20.0 | | 45.0 + | |
| Dasyatidae | <i>Dasyatis americana</i> | 90.0 Width | | 300.0 Length 200.0 Width | |
| Diodontidae | <i>Diodon hystrix</i> | 25.0 | | 65.0 | |
| Elopidae | <i>Elops saurus</i> | 60.0 | | 100.0 | |

continued...

Table 8.1.5 continued

| Family | Species | Common length (cm) | Common weight (g) | Maximum length (cm) | Maximum weight (g) |
|------------------------------|----------------------------------|--------------------|-------------------|---------------------|--------------------|
| Engraulidae | <i>Anchoa hepsetus</i> | 10.0 | | 15.0 | |
| | <i>Anchoa lyolepis</i> | 6.0 | | 7.5 | |
| Ephippidae | <i>Chaetodipterus faber</i> | 50.0 | | 90.0 | |
| Gerreidae | <i>Eucinostomus argenteus</i> | 15.0 | | 20.0 | |
| | <i>Gerres cinereus</i> | 28.0 | | 39.0 | |
| Tnglymostomatidae | <i>Ginglymostoma cirratum</i> | 340.0 | | 430.0 | |
| Grammistidae (Serranidae) | <i>Rypticus saponaceus</i> | 25.0 | | 32.0 | |
| Haemulidae (Pomadasyidae) | <i>Anisotremus surinamensis</i> | 45.0 | | 60.0 | |
| | <i>Anisotremus virginicus</i> | 25.0 | | 35.0 | |
| | <i>Haemulon album</i> | 35.0 | | 42.0 | |
| | <i>Haemulon aurolineatum</i> | 17.0 | | 25.0 | |
| | <i>Haemulon bonariense</i> | 30.0 | | 40.0 | |
| | <i>Haemulon boschmae</i> | 15.0 | | 19.0 | |
| | <i>Haemulon carbonarium</i> | 20.0 | | 35.0 | |
| | <i>Haemulon chrysargyreum</i> | 17.0 | | 23.0 | |
| | <i>Haemulon flavolineatum</i> | 17.0 | | 22.0 | |
| | <i>Haemulon macrostomum</i> | 24.0 | | 28.0 | |
| | <i>Haemulon melanurum</i> | 25.0 | | 33.0 | |
| | <i>Haemulon parrai</i> | 30.0 | | 41.0 | |
| | <i>Haemulon plumieri</i> | 30.0 | | 40.0 | |
| | <i>Haemulon sciurus</i> | 25.0 | | 37.0 | |
| | <i>Haemulon steindachneri</i> | 20.0 | | 27.0 | |
| | <i>Haemulon striatum</i> | 17.0 | | 22.0 | |
| | <i>Orthopristis chrysoptera</i> | 30.0 | | 46.0 | |
| | <i>Orthopristis poeyi</i> | 20.0 | | 24.0 | |
| | <i>Orthopristis ruber</i> | 30.0 | | 40.0 | |
| | <i>Pomadasys corvinaeformis</i> | 18.0 | | 25.0 | |
| <i>Pomadasys crocro</i> | 25.0 | | 33.0 | | |
| Hemiramphidae | <i>Hemiramphus balao</i> | 35.0 | | 40.0 | |
| | <i>Hemiramphus brasiliensis</i> | 35.0 | | 40.5 | |
| | <i>Hyporamphus unifasciatus</i> | 20.0 | | 27.0 | |
| Kyphosidae | <i>Kyphosus incisor</i> | 45.0 | | 67.0 | |
| | <i>Kyphosus sectatrix</i> | 50.0 | | 76.0 | |
| Labridae | <i>Bodianus rufus</i> | 28.0 | | 40.0 | |
| | <i>Lachnolaimus maximus</i> | 35.0 | | 70.0 | |
| Lutjanidae | <i>Lutjanus analis</i> | 50.0 | | 76.0 | |
| | <i>Lutjanus apodus</i> | 35.0 | | 62.0 | |
| | <i>Lutjanus campechanus</i> | 60.0 | | 80.0 | |
| | <i>Lutjanus cyanopterus</i> | 90.0 | | 160.0 | |
| | <i>Lutjanus griseus</i> | 35.0 | | 65.0 | |
| | <i>Lutjanus jocu</i> | 60.0 | | 74.0 | |
| | <i>Lutjanus mahagoni</i> | 38.0 | | 48.0 | |
| | <i>Lutjanus synagris</i> | 25.0 | | 35.0 | |
| | <i>Ocyurus chrysurus</i> | 40.0 | | 61.0 | |
| Malacanthidae | <i>Malacanthus plumieri</i> | 50.0 | | 60.0 | |
| Megalopidae | <i>Tarpon atlanticus</i> | 130.0 | | 250.0 | |
| Monacanthidae | <i>Aluterus schoepfii</i> | 40.0 | | 60.0 | |
| | <i>Cantherines pullus</i> | 12.0 | | 20.0 | |
| | <i>Monacanthus ciliatus</i> | 10.0 | | 20.0 | |
| | <i>Stephanolepis setifer</i> | 10.0 | | 20.0 | |
| Mugilidae | <i>Mugil curema</i> | 30.0 | | 45.0 | |
| Mullidae | <i>Mulloidichthys martinicus</i> | 28.0 | | 40.0 | |
| | <i>Pseudupeneus maculatus</i> | 22.0 | | 30.0 | |

continued...

Table 8.1.5 continued

| Family | Species | Common length (cm) | Common weight (g) | Maximum length (cm) | Maximum weight (g) | |
|--------------------------------|-------------------------------------|----------------------|-------------------|-----------------------------|--------------------|--|
| Muraenidae | <i>Enchelycore nigricans</i> | | | 100.0 | | |
| | <i>Gymnothorax funebris</i> | | | 190.0 | | |
| | <i>Gymnothorax miliaris</i> | | | 50.0 | | |
| | <i>Gymnothorax vicinus</i> | | | 122.0 | | |
| | <i>Lycodontis moringa</i> | | | 100.0 | | |
| Myliobatidae | <i>Aetobatus narinari</i> | 140.0 Width | | 370.0 Length 280.0 Width | | |
| Ophichthidae | <i>Ophichthus ophis</i> | | | 140.0 | | |
| Ostraciidae | <i>Acanthostracion quadricornis</i> | 20.0 | | 35.0 | | |
| | <i>Lactophrys bicaudalis</i> | 20.0 | | 45.0 | | |
| | <i>Lactophrys trigonus</i> | 20.0 | | 45.0 | | |
| | <i>Lactophrys triqueter</i> | 15.0 | | 30.0 | | |
| Pomacanthidae | <i>Holacanthus ciliaris</i> | 30.0 | | 45.0 | | |
| | <i>Holacanthus tricolor</i> | 30.0 | | 35.0 | | |
| | <i>Pomacanthus arcuatus</i> | 45.0 | | 60.0 | | |
| | <i>Pomacanthus paru</i> | 28.0 | | 37.0 | | |
| Pomacentridae | <i>Abudefduf saxatilis</i> | 15.0 | | 20.0 | | |
| | <i>Abudefduf taurus</i> | 20.0 | | 25.0 | | |
| | <i>Chromis multilineata</i> | 12.0 | | 20.0 | | |
| | <i>Microspathodon chrysurus</i> | 15.0 | | 20.0 | | |
| Pomatomidae | <i>Pomatomus saltator</i> | 60.0 | | 110.0 | | |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | 20.0 | | 30.0 | | |
| | <i>Priacanthus arenatus</i> | 35.0 | | 40.0 | | |
| Rachycentridae | <i>Rachycentron canadum</i> | 100.0 | | 200.0 | | |
| Scaridae | <i>Nicholsina usta</i> | 18.0 | | 29.0 | | |
| | <i>Scarus coelestinus</i> | 50.0 | | 76.0 | | |
| | <i>Scarus coeruleus</i> | 35.0 | | 59.0 + | | |
| | <i>Scarus guacamaia</i> | 70.0 | | 94.0 | | |
| | <i>Scarus iserti</i> | 18.0 | | 27.0 | | |
| | <i>Scarus taeniopterus</i> | 22.0 | | 33.0 | | |
| | <i>Scarus vetula</i> | 32.0 | | 51.0 | | |
| | <i>Sparisoma aurofrenatum</i> | 20.0 | | 28.0 | | |
| | <i>Sparisoma chrysopterus</i> | 25.0 | | 45.0 | | |
| | <i>Sparisoma rubripinne</i> | 30.0 | | 45.0 | | |
| | <i>Sparisoma viride</i> | 38.0 | | 64.0 | | |
| | Scombridae | <i>Auxis rochei</i> | 18.0 | | 25.0 | |
| | | <i>Auxis thazard</i> | 40.0 | | 50.0 | |
| <i>Euthynnus aletteratus</i> | | 35.0 | 6,000 | 40.0 FL | | |
| <i>Scomberomorus cavalla</i> | | 70.0 | | 150.0 FL | 45,000 | |
| <i>Scomberomorus maculatus</i> | | 50.0 | | 70.0 FL+ | | |
| <i>Scomberomorus regalis</i> | | 45.0 | | 80.0 FL+ | | |
| <i>Thunnus obesus</i> | | 180.0 | | 236.0 | | |
| Scorpaenidae | <i>Scorpaena plumieri</i> | 30.0 | | 43.0 | | |
| Serranidae | <i>Alphesthes afer</i> | 25.0 | | 33.0 | | |
| | <i>Cephalopholis cruentata</i> | 20.0 | | 32.0 | | |
| | <i>Cephalopholis fulva</i> | 25.0 | | 30.0 | | |
| | <i>Dermatolepis inermis</i> | | | 90.0 | | |
| | <i>Diplectrum formosum</i> | | | 30.0 | | |
| | <i>Epinephelus adscensionis</i> | | | 40.0 | | |
| | <i>Epinephelus guttatus</i> | 40.0 | | 60.0 | | |
| | <i>Epinephelus itajara</i> | | 40,000 | 240.0 | 260,000 | |
| | <i>Epinephelus morio</i> | 70.0 - | | 70.0 ♀ 72.0 ♂ | | |
| | <i>Epinephelus striatus</i> | | 10,000 - | 100.0 | 20,000 | |
| | <i>Mycteroperca bonaci</i> | 70.0 | 26,000 | 100.0 + | 65,000 | |
| | <i>Mycteroperca cidi</i> | 50.0 | | 100.0 | | |
| | <i>Mycteroperca interstitialis</i> | 40.0 | | 50.0 | | |

continued...

Table 8.1.5 continued

| Family | Species | Common length (cm) | Common weight (g) | Maximum length (cm) | Maximum weight (g) |
|----------------|---------------------------------|--------------------|-------------------|---------------------|--------------------|
| | <i>Mycteroperca microlepis</i> | 50.0 | | 70.0 | |
| | <i>Mycteroperca phenax</i> | 30.0 | | 40.0 + | 7,500 |
| | <i>Mycteroperca rubra</i> | 35.0 | 1,340 | 49.0 + | |
| | <i>Mycteroperca tigris</i> | 40.0 | | 50.0 | |
| | <i>Mycteroperca venenosa</i> | 50.0 - | | 70.0 | |
| | <i>Paranthias furcifer</i> | 20.0 | | 35.0 | |
| | <i>Serranus dewegeri</i> | | | 30.0 + | |
| Sparidae | <i>Archosargus rhomboidalis</i> | 20.0 | | 32.0 | |
| | <i>Calamus bajonado</i> | 54.0 | | 68.0 | |
| | <i>Calamus calamus</i> | 30.0 | | 36.0 | |
| | <i>Calamus penna</i> | 28.0 | | 36.0 | |
| | <i>Calamus pennatula</i> | 30.0 | | 36.0 | |
| | <i>Diplodus caudimacula</i> | 22.0 | | 28.0 | |
| Sphyraenidae | <i>Sphyraena barracuda</i> | 130.0 | | 200.0 | |
| | <i>Sphyraena picudilla</i> | 36.0 | | 50.0 | |
| Sphyrnidae | <i>Sphyrna lewini</i> | 360.0 | | 420.0 | |
| Tetraodontidae | <i>Sphoeroides spengleri</i> | 12.0 | | 15.0 | |
| | <i>Sphoeroides testudineus</i> | 20.0 | | 30.0 | |

Symbols:

+ = rather more than the recorded size.

- = rather less than the recorded size.

() = 'reported' to reach the recorded size.

SL = standard length.

FL = fork length.

♀ = female.

♂ = male.

Table 8.1.6. Length-weight data of Caribbean reef fishes after Munro (1983).

| Family | Species | L _m (cm) * | W _m (g) | M a | F b | Z K | M | F | |
|------------------------------|---------------------------------|--------------------------------------|-----------------------|--------|-------------|------------|---------|------|--|
| Acanthuridae | <i>Acanthurus bahianus</i> | 28 30 TL | 547 | 0.0191 | 3.080 | 4.4 M/K | | | |
| | <i>Acanthurus chirurgus</i> | 35 | 980 | 0.0221 | 3.009 | 4.9 M/K | | | |
| | <i>Acanthurus coeruleus</i> | 30 33 TL | 809 | 0.0278 | 3.022 | 4.7 M/K | | | |
| Balistidae | <i>Balistes capriscus</i> | 31 | | | | | | | |
| | <i>Balistes vetula</i> | 45 | 2,920 | 0.0516 | 2.875 | 0.57 | 2.6 | | |
| | <i>Canthidermis sufflamen</i> | 55 | 4,783 | 0.0599 | 2.817 | | | | |
| | <i>Melichthys niger</i> | 32 * | | | | | | | |
| | <i>Xanthichthys ringens</i> | 25 | | | | | | | |
| Carangidae | <i>Alectis ciliaris</i> | 76 * | | | | | | | |
| | <i>Caranx bartholomaei</i> | 69 81 TL | 9,862 | 0.0063 | 3.368 | | | | |
| | <i>Caranx crysos</i> | 48 58 TL | 2,314 | 0.0065 | 3.302 | | | | |
| | <i>Caranx hippos</i> | 79 * | | | | | | | |
| | <i>Caranx latus</i> | 73 85 TL | | | | (2.46) | | | |
| | <i>Caranx lugubris</i> | 69 * | | | | | | | |
| | <i>Caranx ruber</i> | 52 60 TL | 2,494 | 0.0083 | 3.191 | 0.24 | 1.3-1.5 | | |
| | <i>Elagatis bipinnulatus</i> | 65 * | | | | | | | |
| | <i>Selar crumenophthalmus</i> | 22 * | | | | | | | |
| | <i>Selene vomer</i> | 21 * | | | | | | | |
| | <i>Seriola dumerili</i> | 124 * | | | | | | | |
| | <i>Seriola rivoliana</i> | 53 * | | | | | | | |
| | Chaetodontidae | <i>Chaetodon aculeatus</i> | 8.8 ?* | 15.2 * | 0.0223 | 3.000 | | | |
| | | <i>Chaetodon capistratus</i> | 13 ?Q* | | | | | | |
| | | 14 ?Q* | | | | | | | |
| <i>Chaetodon ocellatus</i> | | 16 ?* | | | | | | | |
| <i>Chaetodon sedentarius</i> | | 13 ?* | | | | | | | |
| <i>Chaetodon striatus</i> | | 15 ?* | | | | | | | |
| Haemulidae (Pomadasyidae) | <i>Anisotremus surinamensis</i> | | | 0.0233 | 3.010 | | | | |
| | <i>Anisotremus virginicus</i> | 30 | 846 | 0.0102 | 3.330 | | | | |
| | <i>Haemulon album</i> | 65 | 5,297 | 0.0144 | 3.070 | 0.2 | | | |
| | <i>Haemulon aurolineatum</i> | 23 | 188 | 0.0298 | 2.790 | | | | |
| | <i>Haemulon bonariense</i> | 30 | 385 | 0.0485 | 2.640 | | | | |
| | <i>Haemulon carbonarium</i> | | | 0.0404 | 2.740 | | | | |
| | <i>Haemulon chrysargyreum</i> | | | 0.0141 | 3.080 | | | | |
| | <i>Haemulon flavolineatum</i> | 25 | 374 | 0.0107 | 3.250 | 4.88 M/K | | | |
| | <i>Haemulon macrostomum</i> | | | 0.0176 | 3.060 | | | | |
| | <i>Haemulon melanurum</i> | 30 | 427 | 0.0557 | 2.630 | | | | |
| | <i>Haemulon plumieri</i> | 42 | 1,357 | 0.0238 | 2.930 | 0.345 | 1.77 | 5.13 | |
| <i>Haemulon sciurus</i> | 40 | 1,114 | 0.0314 | 2.840 | 0.26 | | 1.6 | | |
| Holocentridae | <i>Holocentrus ascensionis</i> | 26.5 | 345 | 0.0565 | 2.660 | 3.66 M/K | | | |
| | <i>Holocentrus rufus</i> | 23.5 | 231 | 0.0178 | 3.000 | 4.85 M/K | | | |
| Lutjanidae | <i>Apsilis dentatus</i> | 56 ♀ 58 ♂ | | | 0.3 0.65 | 0.8 1.9 | | | |
| | <i>Lutjanus analis</i> | 60 ♂* 64 TL♂* | | | | | | | |
| | <i>Lutjanus apodus</i> | 57 ♀* 60 TL♀* | 3,800 | 0.0089 | 3.204 | (0.18) | | | |
| | <i>Lutjanus buccanella</i> | 46 ♀ 49.9 TL♀ 54 ♂ 58.6 TL♂ | | | 0.35 0.7 | 1.8 2.2 | | | |
| | <i>Lutjanus griseus</i> | 45 ♂* | | | | | | | |
| | <i>Lutjanus jocu</i> | 72 ♂* | | | | | | | |
| | <i>Lutjanus mahagoni</i> | 26 ♂* | | | | | | | |
| | <i>Lutjanus synagris</i> | 41 ♂* 44.6 TL♂* | 1,050 g | 0.0189 | 2.943 g | | | | |
| | <i>Lutjanus vivanus</i> | 70 () 76.5 (TL) | | | | | | | |

continued...

Table 8.1.6 continued

| Family | Species | L _{max} (cm) * | W _{max} (g) | M _a | F _b | Z _K | M | F |
|---------------|--------------------------------------|----------------------------|-------------------------|------------------|----------------|----------------|-------|---|
| | <i>Ocyurus chrysurus</i> | 60 75 TL | 3,600 | 0.0145 | 3.032 | 0.25 | 0.6 | |
| | <i>Pristipomoides macrophthalmus</i> | 35 ♂ * | | | | | | |
| | <i>Rhomboplites aurorubens</i> | 39 ♀ * | | | | | | |
| Mullidae | <i>Mulloidichthys martinicus</i> | 30 ♀ 29 ♂ | 513 363 | 0.0089 0.0820 | 3.223 2.493 | | | |
| | <i>Pseudupeneus maculatus</i> | 25 ♀ 26.5 ♂ | 259 367 | 0.0552 0.0099 | 2.626 3.210 | | | |
| Pomacanthidae | <i>Holacanthus ciliaris</i> | 33.3 ? * | | 0.0280 | 3.000 | | | |
| | <i>Holacanthus tricolor</i> | 34.4 ? * | 1,140 * | | | | | |
| | <i>Pomacanthus arcuatus</i> | 60 (?) | | | | | | |
| | <i>Pomacanthus paru</i> | 36 ? * 37.5 ? * | 1,625 * | 0.0348 | 3.000 | | | |
| Scaridae | <i>Scarus coelestinus</i> | 32.1 ♀ * 29.5 ♂ * | | | | | | |
| | <i>Scarus coeruleus</i> | 44.5 * | | | | | | |
| | <i>Scarus guacamaia</i> | 45.7 ♀ * | | | | | | |
| | <i>Scarus iserti</i> | 19.7 ♀ * 20.8 ♂ * | | 0.0166 | 3.020 | | | |
| | <i>Scarus taeniopterus</i> | 35 | | | | 6.5 M/K | | |
| | <i>Scarus vetula</i> | 24.8 ♀ * 28.6 ♂ * | | | | | | |
| | <i>Sparisoma aurofrenatum</i> | 26 | 324 | 0.0129 | 3.110 | 5.9 Z/K | | |
| | <i>Sparisoma chrysopterygum</i> | 40 | 1,290 | 0.0093 | 3.210 | 11.7 Z/K | | |
| | <i>Sparisoma rubripinne</i> | 22.7 ♀ * 34.5 ♂ * | | 0.0031 | 3.562 | | | |
| | <i>Sparisoma viride</i> | 50 | 2,430 | 0.0537 | 2.740 | 10.5 Z/K | | |
| Serranidae | <i>Cephalopholis cruentata</i> | 34 ? | 691 | 0.0076 | 3.237 | 0.34 | | |
| | <i>Cephalopholis fulva</i> | 34 ? | 638 | 0.0729 | 2.574 | 0.63 | 1.95 | |
| | <i>Epinephelus guttatus</i> | 52 ? | 2,113 | 0.0176 | 2.960 | 0.48 | 0.68 | |
| | <i>Epinephelus striatus</i> | 90 ? | 12,900 | 0.0107 | 3.112 | 0.09 | 0.235 | |
| | <i>Mycteroperca venenosa</i> | 86 ? | | | | 0.17 | 0.41 | |

*Length' refers to fork length unless stated otherwise.

Symbols:

* = L_{max}, W_{max}
() = 'reported' to reach the recorded size.
g = gutted weight.

TL = total length.

? = type of length not stated.

♀ = female.

♂ = male.

Appendix 8.2

Table 8.2.1. Density (ind.m⁻³) and crop (gm⁻³) of fishes of a coral reef in the US Virgin Islands after Smith and Tyler (1972).

| Family | Species | ind.m ⁻³ | gm ⁻³ |
|------------------------------|------------------------------------|-------------------------------|------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 0.01 | 0.005 |
| | <i>Acanthurus coeruleus</i> | 0.03 | 0.01 |
| Apogonidae | <i>Apogon binotatus</i> | 0.70 | 1.02 |
| | <i>Apogon conklini</i> | 0.11 | 0.05 |
| | <i>Apogon maculatus</i> | 0.09 | 0.37 |
| | <i>Apogon pigmentaria</i> | 0.03 | 0.014 |
| | <i>Apogon quadrisquamatus</i> | 0.12 | 0.03 |
| | <i>Apogon townsendi</i> | 1.18 | 1.55 |
| | <i>Apogon xenus</i> | 0.03 | 0.28 |
| Blenniidae | <i>Hypoleurochilus</i> sp. | 0.02 | 0.002 |
| Canthigasteridae | <i>Canthigaster rostrata</i> | 0.07 | 0.280 |
| Cirrhitidae | <i>Amblycirrhites pinos</i> | 0.02 | 0.010 |
| Clinidae | <i>Acanthemblemaria spinosa</i> | 0.01 | 0.001 |
| | <i>Emblemariopsis leptocirris</i> | 0.01 | 0.001 |
| | <i>Enneanectes altivelis</i> | 0.02 | 0.001 |
| | <i>Labrisomus haitiensis</i> | 0.10 | 0.08 |
| | <i>Pseudemblemaria signifera</i> | 0.05 | 0.005 |
| | <i>Starksia hassi</i> | 0.02 | 0.004 |
| | <i>Starksia lepicoelia</i> | 0.04 | 0.008 |
| Emmelichthyidae | <i>Emmelichthys atlanticus</i> | 0.1 | 0.16 |
| Gobiidae | <i>Coryphopterus dicrus</i> | 0.06 | 0.01 |
| | <i>Coryphopterus eidolon</i> | 0.25 | 0.14 |
| | <i>Coryphopterus glaucofraenum</i> | 0.27 | 0.22 |
| | <i>Coryphopterus personatus</i> | 0.23 | 0.07 |
| | <i>Coryphopterus thrix</i> | 0.04 | 0.02 |
| | <i>Gnatholepis thompsoni</i> | 0.06 | 0.06 |
| | <i>Gobiosoma evelynae</i> | 0.10 | 0.02 |
| | <i>Gobiosoma horsti</i> | 0.01 | 0.003 |
| | <i>Gobiosoma saucrum</i> | 0.06 | 0.005 |
| | <i>Lythrypnus elasson</i> | 0.03 | 0.002 |
| | <i>Lythrypnus nesiotis</i> | 0.23 | 0.004 |
| | <i>Lythrypnus</i> sp. | 0.02 | 0.001 |
| | <i>Quisquilius hipoliti</i> | 0.04 | 0.009 |
| | <i>Risor ruber</i> | 0.03 | 0.003 |
| | Grammistidae (Serranidae) | <i>Rypticus subbifrenatus</i> | 0.01 |
| Haemulidae (Pomadasyidae) | <i>Haemulon aurolineatum</i> | 0.55 | 0.62 |
| | <i>Haemulon flavolineatum</i> | 0.40 | 0.35 |
| Holocentridae | <i>Holocentrus coruscus</i> | 0.08 | 0.27 |
| | <i>Holocentrus rufus</i> | 0.01 | 0.05 |
| | <i>Myripristis jacobus</i> | 0.10 | 0.20 |
| | <i>Neoniphon marianus</i> | 0.30 | 1.63 |
| Labridae | <i>Halichoeres bivittatus</i> | 0.01 | 0.12 |
| | <i>Halichoeres garnoti</i> | 0.05 | 0.13 |
| | <i>Thalassoma bifasciatum</i> | 0.16 | 0.26 |
| Lutjanidae | <i>Ocyurus chrysurus</i> | 0.10 | 0.70 |
| Monacanthidae | <i>Monacanthus tocken</i> | 0.01 | 0.001 |
| Mullidae | <i>Pseudupeneus maculatus</i> | 0.01 | 1.10 |
| Muraenidae | <i>Lycodontis moringa</i> | 0.01 | 0.18 |
| | <i>Gymnothorax miliaris</i> | 0.02 | 0.03 |

continued...

Table 8.2.1 continued

| Family | Species | ind.m ⁻³ | gm ⁻³ |
|------------------|--------------------------------|---------------------|------------------|
| Ostraciidae | <i>Lactophrys triqueter</i> | 0.01 | 0.65 |
| Pomacanthidae | <i>Holacanthus ciliaris</i> | 0.03 | 0.012 |
| | <i>Chromis cyanea</i> | 0.35 | 1.23 |
| | <i>Chromis multilineata</i> | 0.18 | 0.96 |
| | <i>Pomacentrus partitus</i> | 0.06 | 0.15 |
| | <i>Stegastes planifrons</i> | 0.10 | 0.60 |
| | <i>Stegastes variabilis</i> | 0.09 | 0.26 |
| Scaridae | <i>Scarus iserti</i> | 0.02 | 0.30 |
| | <i>Scarus vetula</i> | 0.01 | 2.00 |
| | <i>Sparisoma viride</i> | 0.01 | 0.001 |
| Serranidae | <i>Cephalopholis cruentata</i> | 0.02 | 4.00 |
| | <i>Epinephelus striatus</i> | 0.01 | 4.00 |
| | <i>Hypoplectrus puella</i> | 0.04 | 0.032 |
| Syngnathidae | <i>Syngnathus</i> sp. | 0.01 | 0.001 |
| Synodontidae | <i>Synodus intermedius</i> | 0.01 | 0.82 |
| Sum (64 species) | | | 25.17 |

Appendix 8.3

Table 8.3.1. Aspect ratio (A) of Caribbean reef fishes based on drawings in Fischer (1978) and photographs in Randall (1968); length and sex of specimen from photographs in Randall (1968).

| Family | Species | A (Fischer) | Length (cm) | Sex | A (Randall) |
|----------------|------------------------------------|----------------|----------------|-----|----------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 2.22 | 20 | - | 2.18 |
| | <i>Acanthurus chirurgus</i> | 1.87 | 22 | - | 1.78 |
| | <i>Acanthurus coeruleus</i> | 1.98 | 17 | - | 1.86 |
| Antennariidae | <i>Antennarius multiocellatus</i> | | 12 | | 1.14 |
| | | | 19 | - | 1.22 |
| | <i>Antennarius striatus</i> | | 7 | - | 1.11 |
| | | | 15 | | 0.80 |
| Apogonidae | <i>Apogon conklini</i> | | 6 | - | 1.72 |
| | <i>Apogon maculatus</i> | | 6 | - | 1.53 |
| Atherinidae | <i>Allanetta harringtonensis</i> | | 5 | - | 1.71 |
| | <i>Atherinomorus stipes</i> | | 7 | - | 1.88 |
| Aulostomidae | <i>Aulostomus maculatus</i> | 1.19 | 59 | - | 1.70 |
| Balistidae | <i>Balistes capriscus</i> | 2.47 | 13 | - | 2.44 |
| | <i>Balistes vetula</i> | 1.99 | 33 | - | 1.98 |
| | <i>Canthidermis sufflamen</i> | | 41 | - | 1.71 |
| | <i>Melichthys niger</i> | 2.05 | 33 | - | 2.30 |
| | <i>Xanthichthys ringens</i> | | 12 | | 1.92 |
| Belonidae | <i>Ablennes hians</i> | 3.27 | | - | |
| | <i>Platybelone argalus argalus</i> | 2.44 | 34 | - | 2.65 |
| | <i>Strongylura marina</i> | 1.52 | | - | |
| | <i>Strongylura notata</i> | | 25 | - | 1.64 |
| | <i>Strongylura timucu</i> | 1.50 | 32 | - | 1.69 |
| | <i>Tylosurus acus</i> | 1.83 | | - | |
| | <i>Tylosurus crocodilus</i> | 2.16 | 58 | - | 3.11 |
| Blenniidae | <i>Entomacrodus nigricans</i> | | 6 | - | 1.02 |
| | <i>Ophioblennius atlanticus</i> | | 11 | - | 0.92 |
| | <i>Parablennius mammoreus</i> | | 5 | - | 1.26 |
| | <i>Scartella cristata</i> | | 8 | - | 1.18 |
| Bothidae | <i>Bothus lunatus</i> | 1.40 | 28 | - | 1.11 |
| | <i>Bothus ocellatus</i> | | 7 | ♀ | 1.18 |
| | | | 10 | ♂ | 1.51 |
| | <i>Paralichthys tropicus</i> | 1.71 | 24 | - | 1.38 |
| Carangidae | <i>Alectis ciliaris</i> | | 41 | - | 4.82 |
| | <i>Caranx bartholomaei</i> | 4.14 | 37 | - | 4.06 |
| | <i>Caranx crysos (fusus)</i> | 3.52 | 29 | - | 4.19 |
| | <i>Caranx hippos</i> | 3.40 | 33 | - | 3.76 |
| | <i>Caranx latus</i> | 4.32 | 34 | - | 4.48 |
| | <i>Caranx lugubris</i> | 3.96 | 62 | - | 5.48 |
| | <i>Caranx ruber</i> | 2.40 | 27 | - | 4.52 |
| | <i>Decapterus macarellus</i> | 2.98 | 18 | - | 3.80 |
| | <i>Decapterus punctatus</i> | 2.80 | 16 | - | 3.66 |
| | <i>Oligoplites saurus</i> | 3.43 | 25 | - | 3.12 |
| | <i>Selar crumenophthalmus</i> | 3.15 | 17 | - | 3.54 |
| | <i>Seriola dumerili</i> | 3.30 | 112 | - | 4.49 |
| | <i>Trachinotus falcatus</i> | 3.97 | 31 | - | 5.79 |
| | | | 104 | - | 3.88 |
| | <i>Trachinotus goodei</i> | 2.69 | 21 | - | 5.01 |
| Chaetodontidae | <i>Chaetodon aculeatus</i> | | 8 | - | 1.85 |
| | <i>Chaetodon capistratus</i> | | 10 | - | 2.53 |
| | <i>Chaetodon sedentarius</i> | | 11 | - | 2.29 |
| | <i>Chaetodon striatus</i> | | 14 | - | 2.37 |
| Cirrhitidae | <i>Amblycirrhitis pinos</i> | | 7 | - | 1.52 |
| Clinidae | <i>Labrisomus nuchipinnis</i> | | 11 | - | 1.29 |

continued...

Table 8.3.1 continued

| Family | Species | A (Fischer) | Length (cm) | Sex | A (Randall) |
|------------------------------|------------------------------------|----------------|----------------|----------|----------------|
| Clupeidae | <i>Harengula clupeiola</i> | 1.72 | 11 | - | 2.65 |
| | <i>Harengula humeralis</i> | | 11 | - | 2.90 |
| | <i>Jenkinsia lamprotaenia</i> | 1.82 | 5 | - | 2.19 |
| | <i>Opisthonema oglinum</i> | 2.53 | 13 | - | 3.13 |
| Dactylopteridae | <i>Dactylopterus volitans</i> | 1.63 | 33 | - | 1.60 |
| Diodontidae | <i>Chilomycterus antennatus</i> | | 14 | - | 1.40 |
| | <i>Diodon holocanthus</i> | | 14 | - | 1.37 |
| | <i>Diodon hystrix</i> | 0.92 | 38 | - | 1.09 |
| Echeneidae | <i>Echeneis naucrates</i> | | 32 | - | 1.10 |
| | <i>Remora remora</i> | | 20 | - | 1.60 |
| Elopidae | <i>Elops saurus</i> | 4.53 | 33 | - | 3.41 |
| Emmelichthyidae | <i>Emmelichthys atlanticus</i> | | 10 | - | 2.67 |
| | <i>Inermia vittata</i> | | 15 | - | 2.92 |
| Engraulidae | <i>Anchoa hepsetus</i> | 2.50 | | - | |
| | <i>Anchoa lyolepis</i> | 1.96 | 6 | - | 1.77 |
| | <i>Anchoa mitchilli</i> | 2.95 | | - | |
| | <i>Anchoa parva</i> | 2.24 | | - | |
| Ephippidae | <i>Chaetodipterus faber</i> | 3.04 | 35 | - | 3.11 |
| Exocoetidae | <i>Exocoetus volitans</i> | 1.76 | | - | |
| Fistulariidae | <i>Fistularia tabacaria</i> | | 39 | - | 1.70 |
| Gerreidae | <i>Eucinostomus argenteus</i> | 2.98 | 12 | - | 1.94 |
| | <i>Eucinostomus lefroyi</i> | | 12 | - | 2.64 |
| | <i>Gerres cinereus</i> | 3.68 | 37 | - | 3.91 |
| Gobiidae | <i>Bathygobius soporator</i> | | 16 | - | 0.92 |
| | <i>Coryphopterus glaucofraenum</i> | | 6 | - | 1.11 |
| | <i>Coryphopterus personatus</i> | | 3 | - | 1.06 |
| | <i>Gnatholepis thompsoni</i> | | 5 | - | 0.96 |
| | <i>Gobiosoma evelynae</i> | | 4 | - | 0.79 |
| | <i>Ioglossus helenae</i> | | 11 | - | 0.84 |
| | <i>Quisquilius hipoliti</i> | | 3 | - | 1.09 |
| Grammidae (Serranidae) | <i>Gramma loreto</i> | | 6 | - | 1.13 |
| | <i>Gramma melacara</i> | | 6 | - | 1.36 |
| Grammistidae (Serranidae) | <i>Rypticus saponaceus</i> | 1.21 | 17 | - | 1.37 |
| Haemulidae (Pomadasyidae) | <i>Anisotremus surinamensis</i> | | 26 | - | 2.66 |
| | <i>Anisotremus virginicus</i> | | 6 | juvenile | 1.87 |
| | | | 32 | adult | 2.84 |
| | <i>Haemulon album</i> | | 47 | - | 2.77 |
| | <i>Haemulon aurolineatum</i> | | 18 | - | 2.44 |
| | <i>Haemulon bonariense</i> | | 19 | - | 2.45 |
| | <i>Haemulon boschmae</i> | | 11 | - | 1.96 |
| | <i>Haemulon carbonarium</i> | | 33 | - | 2.84 |
| | <i>Haemulon chrysargyreum</i> | | 19 | - | 2.59 |
| | <i>Haemulon flavolineatum</i> | | 19 | - | 2.46 |
| | <i>Haemulon macrostomum</i> | | 39 | - | 2.69 |
| | <i>Haemulon melanurum</i> | | 18 | - | 1.79 |
| | <i>Haemulon parrai</i> | | 33 | - | 2.53 |
| | <i>Haemulon plumieri</i> | | 26 | - | 2.86 |
| | <i>Haemulon sciurus</i> | | 30 | - | 2.45 |
| | <i>Haemulon steindachneri</i> | | 14 | - | 2.02 |
| | <i>Haemulon striatum</i> | | 18 | - | 1.83 |
| <i>Orthopristis ruber</i> | | 25 | - | 2.46 | |
| Hemiramphidae | <i>Hemiramphus balao</i> | 2.87 | 25 | - | 2.73 |
| | <i>Hemiramphus bermudensis</i> | 3.15 | | - | |

continued...

Table 8.3.1 continued

| Family | Species | A (Fischer) | Length (cm) | Sex | A (Randall) |
|------------------|-------------------------------------|----------------|----------------|----------|----------------|
| | <i>Hemiramphus brasiliensis</i> | 3.04 | 28 | - | 3.23 |
| | <i>Hyporamphus unifasciatus</i> | 2.45 | | - | |
| Holocentridae | <i>Holocentrus ascensionis</i> | 2.56 | 27 | - | 2.64 |
| | <i>Holocentrus coruscus</i> | | 11 | - | 3.03 |
| | <i>Holocentrus rufus</i> | 2.24 | 25 | - | 2.23 |
| | <i>Myripristis jacobus</i> | 2.61 | 13 | - | 2.46 |
| | <i>Neoniphon marianus</i> | | 12 | - | 3.02 |
| | <i>Plectrypops retrospinis</i> | | 10 | - | 1.73 |
| | <i>Sargocentron vexillarium</i> | | 9 | - | 2.65 |
| Kyphosidae | <i>Kyphosus incisor</i> | 2.97 | 25 | - | 2.83 |
| | <i>Kyphosus sectatrix</i> | 2.83 | 32 | - | 2.90 |
| Labridae | <i>Bodianus pulchellus</i> | 1.12 | 13 | - | 1.79 |
| | <i>Bodianus rufus</i> | 1.37 | 26 | - | 1.20 |
| | <i>Clepticus parrae</i> | | 24 | - | 1.84 |
| | <i>Halichoeres bivittatus</i> | | 14 | - | 1.27 |
| | <i>Halichoeres garnoti</i> | | 6 | juvenile | 1.48 |
| | | | 9 | ♀ | 1.09 |
| | | | 12 | ♂ | 1.44 |
| | <i>Halichoeres maculipinna</i> | | 6 | juvenile | 1.44 |
| | | | 13 | ♂ | 1.25 |
| | <i>Halichoeres pictus</i> | | 9 | ♀ | 1.39 |
| | | | 11 | ♂ | 1.50 |
| | <i>Halichoeres poeyi</i> | | 10 | - | 1.17 |
| | | | 16 | ♂ | 1.19 |
| | <i>Halichoeres radiatus</i> | 1.63 | 12 | juvenile | 0.87 |
| | | | 24 | ♀ | 1.58 |
| | <i>Xyrichtys novacula</i> | 1.40 | 18 | - | 1.48 |
| | <i>Xyrichtys splendens</i> | | 5 | - | 1.54 |
| | | | 12 | - | 1.48 |
| | <i>Hemipteronotus martinicensis</i> | | 8 | ♀ | 1.09 |
| | | | 13 | ♂ | 1.18 |
| | <i>Lachnolaimus maximus</i> | 1.66 | 22 | - | 1.51 |
| | <i>Thalassoma bifasciatum</i> | | 9 | ♀ | 1.32 |
| | | | 12 | ♂ | 1.07 |
| Lutjanidae | <i>Lutjanus analis</i> | 2.66 | 38 | - | 2.06 |
| | <i>Lutjanus apodus</i> | 1.83 | 25 | - | 2.00 |
| | <i>Lutjanus campechanus</i> | 1.23 | | - | 1.41 |
| | <i>Lutjanus cyanopterus</i> | 2.08 | 90 | - | 2.18 |
| | <i>Lutjanus griseus</i> | 1.90 | 40 | - | 2.06 |
| | <i>Lutjanus jocu</i> | 2.38 | 39 | - | 2.08 |
| | <i>Lutjanus mahagoni</i> | 2.01 | 24 | - | 2.05 |
| | <i>Lutjanus synagris</i> | 1.80 | 18 | - | 1.97 |
| | <i>Ocyurus chrysurus</i> | 3.56 | 16 | - | 2.96 |
| Malacanthidae | <i>Malacanthus plumieri</i> | 1.80 | 53 | - | 1.42 |
| Megalopidae | <i>Tarpon atlanticus</i> | | 37 | - | 2.96 |
| Monacanthidae | <i>Aluterus monoceros</i> | | 47 | - | 1.70 |
| | <i>Aluterus schoepfii</i> | 0.87 | 43 | - | 0.72 |
| | <i>Aluterus scripta</i> | | 50 | - | 0.72 |
| | <i>Cantherines macrocerus</i> | | 37 | ♀ | 1.46 |
| | <i>Cantherines pullus</i> | 1.68 | 17 | - | 1.47 |
| | <i>Monacanthus ciliatus</i> | 1.36 | 10 | - | 1.24 |
| | <i>Monacanthus tuckeri</i> | | 9 | - | 1.37 |
| | <i>Stephanolepis setifer</i> | 1.13 | 18 | - | 1.25 |
| Mugilidae | <i>Mugil curema</i> | 1.92 | 26 | - | 2.72 |
| Mullidae | <i>Mulloidichthys martinicus</i> | 4.85 | 33 | - | 2.26 |
| | <i>Pseudupeneus maculatus</i> | 2.85 | 24 | - | 2.30 |
| Opisthognathidae | <i>Opisthognathus aurifrons</i> | | 6 | - | 0.84 |
| | <i>Opisthognathus macrognathus</i> | | 12 | - | 1.14 |

continued...

Table 8.3.1 continued

| Family | Species | A (Fischer) | Length (cm) | Sex | A (Randall) |
|----------------|-------------------------------------|----------------|----------------|----------|----------------|
| | <i>Opisthognathus maxillosus</i> | | 8 | - | 0.76 |
| | <i>Opisthognathus whitehurstii</i> | | 5 | - | 0.82 |
| Ostraciidae | <i>Acanthostracion polygonius</i> | | 25 | - | 1.17 |
| | <i>Acanthostracion quadricornis</i> | 1.82 | 24 | - | 1.14 |
| | <i>Lactophrys bicaudalis</i> | 1.89 | 37 | - | 1.23 |
| | <i>Lactophrys trigonus</i> | 1.77 | 29 | - | 1.89 |
| | | | 33 | - | 1.87 |
| | <i>Lactophrys triqueter</i> | 2.18 | 14 | - | 1.98 |
| Pempheridae | <i>Pempheris schomburgki</i> | | 11 | - | 2.28 |
| Pomacanthidae | <i>Centropyge argi</i> | | 5 | - | 1.44 |
| | <i>Holacanthus ciliaris</i> | | 6 | juvenile | 1.44 |
| | | | 21 | adult | 1.29 |
| | <i>Holacanthus tricolor</i> | | 17 | - | 1.65 |
| | <i>Pomacanthus arcuatus</i> | | 30 | - | 2.89 |
| | <i>Pomacanthus paru</i> | | 4 | juvenile | 1.39 |
| | | | 24 | adult | 1.81 |
| Pomacentridae | <i>Abudefduf saxatilis</i> | | 13 | - | 2.98 |
| | <i>Abudefduf taurus</i> | | 17 | - | 2.01 |
| | <i>Chromis cyanea</i> | | 12 | - | 1.79 |
| | <i>Chromis multilineata</i> | | 13 | - | 1.79 |
| | <i>Microspathodon chrysurus</i> | | 18 | - | 1.88 |
| | <i>Pomacentrus partitus</i> | | 6 | - | 1.38 |
| | <i>Stegastes fuscus</i> | | 9 | - | 1.83 |
| | <i>Stegastes leucostictus</i> | | 5 | - | 1.31 |
| | <i>Stegastes planifrons</i> | | 6 | - | 1.37 |
| | <i>Stegastes variabilis</i> | | 7 | - | 1.54 |
| Pomatomidae | <i>Pomatomus saltator</i> | 2.16 | | - | |
| Priacanthidae | <i>Heteropriacanthus cruentatus</i> | | 25 | - | 1.79 |
| | <i>Priacanthus arenatus</i> | | 27 | - | 1.83 |
| Rachycentridae | <i>Rachycentron canadus</i> | | 57 | - | 1.56 |
| Scaridae | <i>Cryptotomus roseus</i> | | 10 | - | 1.21 |
| | <i>Nicholsina usta</i> | | 15 | - | 1.12 |
| | <i>Scarus coelestinus</i> | | 52 | - | 1.06 |
| | <i>Scarus coeruleus</i> | | 33 | - | 0.95 |
| | | | 62 | ♂ | 1.48 |
| | <i>Scarus guacamaia</i> | | 57 | - | 1.19 |
| | <i>Scarus iserti</i> | | 21 | ♀ | 1.24 |
| | | | 22 | ♂ | 1.44 |
| | <i>Scarus taeniopterus</i> | | 19 | ♀ | 1.23 |
| | | | 24 | ♂ | 1.09 |
| | <i>Scarus vetula</i> | | 26 | ♀ | 1.49 |
| | | | 34 | ♂ | 1.05 |
| | <i>Sparisoma aurofrenatum</i> | | 16 | ♀ | 1.17 |
| | | | 19 | ♂ | 1.33 |
| | <i>Sparisoma chrysopterygum</i> | | 26 | ♀ | 1.39 |
| | | | 27 | ♂ | 1.44 |
| | <i>Sparisoma radians</i> | | 9 | ♀ | 1.26 |
| | | | 8 | ♂ | 1.18 |
| | <i>Sparisoma rubripinne</i> | | 18 | - | 1.12 |
| | | | 41 | ♂ | 1.51 |
| | <i>Sparisoma viride</i> | | 26 | ♀ | 1.66 |
| | | | 31 | ♂ | 1.43 |
| Sciaenidae | <i>Equetus lanceolatus</i> | | 18 | - | 0.97 |
| | <i>Equetus punctatus</i> | | 18 | - | 1.25 |
| | <i>Odontoscion dentex</i> | | 14 | - | 1.68 |
| | <i>Pareques acuminatus</i> | | 12 | - | 1.34 |
| Scombridae | <i>Euthynnus aletteratus</i> | | 33 | - | 6.29 |
| | <i>Scomberomorus cavalla</i> | | 111 | - | 6.31 |
| | <i>Scomberomorus maculatus</i> | | 53 | - | 4.87 |
| | <i>Scomberomorus regalis</i> | | 41 | - | 6.16 |

continued...

Table 8.3.1 continued

| Family | Species | A (Fischer) | Length (cm) | Sex | A (Randall) |
|----------------|------------------------------------|----------------|----------------|-----|----------------|
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | | 20 | - | 1.03 |
| | <i>Scorpaena grandicornis</i> | | 12 | - | 1.03 |
| | <i>Scorpaena plumieri</i> | | 16 | - | 1.00 |
| | <i>Scorpaenodes canibbaeus</i> | | 7 | - | 1.01 |
| Serranidae | <i>Alphesthes afer</i> | | 23 | - | 1.20 |
| | <i>Cephalopholis cruentata</i> | | 19 | - | 1.23 |
| | <i>Cephalopholis fulva</i> | | 23 | - | 1.47 |
| | <i>Dermatolepis inermis</i> | | 15 | - | 1.51 |
| | <i>Diplectrum formosum</i> | 1.63 | 18 | - | 1.49 |
| | <i>Epinephelus adscensionis</i> | | 34 | - | 1.53 |
| | <i>Epinephelus guttatus</i> | | 25 | - | 1.03 |
| | <i>Epinephelus itajara*</i> | | 165 | - | 1.28 |
| | <i>Epinephelus morio</i> | | 43 | - | 1.46 |
| | <i>Epinephelus striatus</i> | | 40 | - | 1.57 |
| | <i>Hypoplectrus aberrans</i> | | 12 | - | 1.84 |
| | <i>Hypoplectrus chlorurus</i> | | 9 | - | 1.53 |
| | <i>Hypoplectrus nigricans</i> | | 11 | - | 1.77 |
| | <i>Hypoplectrus puella</i> | | 12 | - | 1.58 |
| | <i>Mycteroperca bonaci</i> | | 40 | - | 1.22 |
| | <i>Mycteroperca cidi</i> | | 24 | - | 1.13 |
| | <i>Mycteroperca interstitialis</i> | | 32 | - | 1.29 |
| | <i>Mycteroperca phenax</i> | | 27 | - | 1.36 |
| | <i>Mycteroperca rubra</i> | | 18 | - | 1.20 |
| | <i>Mycteroperca tigris</i> | | 36 | - | 1.81 |
| | <i>Mycteroperca venenosa</i> | | 22 | - | 1.08 |
| | <i>Paranthias furcifer</i> | | 24 | - | 2.27 |
| | <i>Serranus dewegeri</i> | | 22 | - | 1.40 |
| | <i>Serranus tabacarius</i> | | 11 | - | 1.24 |
| | <i>Serranus tigrinus</i> | | 9 | - | 1.45 |
| | <i>Serranus tortugarum</i> | | 5 | - | 1.58 |
| Sparidae | <i>Archosargus rhomboidalis</i> | | 17 | - | 2.47 |
| | <i>Calamus bajonado</i> | | 34 | - | 3.04 |
| | <i>Calamus calamus</i> | | 26 | - | 3.18 |
| | <i>Calamus penna</i> | | 24 | - | 3.11 |
| | <i>Calamus pennatula</i> | | 31 | - | 3.04 |
| | <i>Diplodus caudimacula</i> | | 27 | - | 2.92 |
| Sphyraenidae | <i>Sphyraena barracuda</i> | | 49 | - | 1.61 |
| | <i>Sphyraena picudilla</i> | | 36 | - | 2.62 |
| Synodontidae | <i>Synodus foetens</i> | | 31 | - | 2.65 |
| | <i>Synodus intermedius</i> | | 38 | - | 2.38 |
| | <i>Synodus synodus</i> | | 12 | - | 2.38 |
| Tetraodontidae | <i>Canthigaster rostrata</i> | | 7 | - | 1.52 |
| | <i>Sphoeroides spengleri</i> | | 12 | - | 1.57 |

Symbols:

♀ = female.

♂ = male.

Appendix 8.4

Table 8.4.1. Food consumption of Caribbean reef fishes (without accounting for omnivory); Q/B values represent mean population consumption of the corresponding species; mean environmental temperature = 28°C.

| Family | Species | Food type | Aspect ratio | W _m (g) | Q/B (%BWD) | Q/B (year ⁻¹) |
|----------------|------------------------------------------------------------|-----------|--------------|--------------------|--------------|---------------------------|
| Acanthuridae | <i>Acanthurus bahianus</i> | 1 | 2.18 | 288 | 10.82 | 39.49 |
| | <i>Acanthurus chirurgus</i> | 1 | 1.78 | 820 | 7.89 | 28.79 |
| | <i>Acanthurus coeruleus</i> | 1 | 1.86 | 983 | 7.77 | 28.36 |
| Balistidae | <i>Balistes caprisus</i> | 0 * | 2.44 | 611 | 2.80 | 10.20 |
| | <i>Balistes vetula</i> | 0 | 1.98 | 2,586 | 1.87 | 6.84 |
| | <i>Canthidermis sufflamen</i> | 0 | 1.71 | 4,783 | 1.53 | 5.60 |
| | <i>Melichthys niger</i> | 1 | 2.30 | 960 | 8.71 | 31.79 |
| Blenniidae | <i>Entomacrodus nigricans</i> | 1 | 1.02 | 13 | 13.65 | 49.81 |
| | <i>Ophioblennius atlanticus</i> | 1 | 0.92 | 35 | 10.61 | 38.73 |
| | <i>Parablennius mammoreus</i> | 1 | 1.26 | 10 | 16.06 | 58.62 |
| | <i>Scartella cristata</i> | 1 | 1.18 | 27 | 12.68 | 46.28 |
| Bothidae | <i>Bothus lunatus</i> | 0 | 1.11 | 3,000 | 1.35 | 4.94 |
| | <i>Bothus ocellatus</i> | 0 | 1.51 | 180 | 2.79 | 10.19 |
| Clupeidae | <i>Harengula humeralis</i> | 0 | 2.90 | 440 | 3.27 | 11.92 |
| Ephippidae | <i>Chaetodipterus faber</i> | 0 | 3.11 | 9,000 | 1.84 | 6.72 |
| Gobiidae | <i>Coryphopterus glaucofraenum</i> | 0 | 1.11 | 10.5 | 4.22 | 15.41 |
| | <i>Gnatholepis thompsoni</i> | 1 | 0.96 | 4.1 | 16.67 | 60.83 |
| Hemiramphidae | <i>Hemiramphus brasiliensis</i> | 1 | 3.23 | 298 | 13.16 | 48.02 |
| Monacanthidae | <i>Aluterus monoceros</i> | 0 * | 1.70 | 2,500 | 1.75 | 6.37 |
| | <i>Aluterus schoepfii</i> | 1 | 0.72 | 1,892 | 4.19 | 15.28 |
| | <i>Aluterus scripta</i> | 0 | 0.72 | 2,941 | 1.08 | 3.95 |
| | <i>Cantherines macrocerus</i> | 0 | 1.46 | 2,700 | 1.59 | 5.79 |
| | <i>Cantherines pullus</i> | 0 | 1.47 | 220 | 2.64 | 9.64 |
| | <i>Monacanthus ciliatus</i> | 0 | 1.24 | 448 | 2.09 | 7.65 |
| | <i>Monacanthus tuckeri</i> | 0 | 1.37 | 24 | 3.99 | 14.57 |
| Ostraciidae | <i>Acanthostracion quadricornis</i> | 0 | 1.14 | 3,050 | 1.37 | 4.98 |
| | <i>Lactophrys trigonus</i> | 0 | 1.87 | 3,052 | 1.76 | 6.43 |
| | <i>Lactophrys bicaudalis</i> | 0 | 1.23 | 5,572 | 1.26 | 4.59 |
| | <i>Lactophrys triqueter</i> | 0 | 1.98 | 1,394 | 2.13 | 7.76 |
| Pomacanthidae | <i>Centropyge argi</i> | 1 | 1.44 | 10 | 17.23 | 62.89 |
| | <i>Holacanthus ciliaris</i> | 0 | 1.29 | 1,988 | 1.59 | 5.80 |
| | <i>Holacanthus tricolor</i> | 0 | 1.65 | 1,306 | 1.96 | 7.15 |
| | <i>Pomacanthus arcuatus</i> | 0 | 2.89 | 12,407 | 1.66 | 6.06 |
| | <i>Pomacanthus paru</i> Virgin Islands Jamaica | 0 0 | 1.81 1.81 | 2,769 2,229 | 1.77 1.84 | 6.45 6.73 |
| Pomacentridae | <i>Abudefduf saxatilis</i> | 0 | 2.98 | 483 | 3.25 | 11.86 |
| | <i>Abudefduf taurus</i> | 1 | 2.01 | 900 | 8.24 | 30.07 |
| | <i>Microspathodon chrysurus</i> | 1 | 1.88 | 412 | 9.31 | 33.99 |
| | <i>Stegastes fuscus</i> | 1 | 1.83 | 168 | 11.00 | 40.17 |
| | <i>Stegastes leucostictus</i> | 0 | 1.31 | 50 | 3.36 | 12.26 |
| | <i>Stegastes planifrons</i> <i>Stegastes variabilis</i> | 0 0 | 1.37 1.54 | 68 54 | 3.23 3.59 | 11.79 13.12 |
| Scaridae | <i>Scarus coelestinus</i> | 1 | 1.06 | 8,556 | 3.75 | 13.70 |
| | <i>Scarus taeniopterus</i> | 1 | 1.09 | 1,102 | 5.78 | 21.08 |
| | <i>Scarus vetula</i> | 1 | 1.05 | 5,558 | 4.09 | 14.93 |
| | <i>Sparisoma aurofrenatum</i> | 1 | 1.33 | 324 | 8.18 | 29.84 |
| Sparidae | <i>Archosargus rhomboidalis</i> | 1 | 2.47 | 1,200 | 8.64 | 31.54 |
| | <i>Diplodus caudimacula</i> | 1 | 2.92 | 600 | 10.84 | 39.57 |
| Tetraodontidae | <i>Canthigaster rostrata</i> | 0 | 1.52 | 82 | 3.28 | 11.98 |
| | <i>Sphoeroides spengleri</i> | 0 | 1.57 | 92 | 3.26 | 11.90 |

Symbol:

* = Assumed (derived from other members of genus or family).

Appendix 8.5. List of species of nonfish taxa.

Table 8.5.1. Benthic primary producers in fish stomachs of West Indian reef fishes (based on Randall 1967); occurrence refers to number of fish species with respective item in stomach.

| Taxon and class | Species | Occurrence | Sum | | |
|------------------------------------------|----------------------------------|-------------------------------|-------|--------------|--|
| | | | Genus | Higher taxon | |
| Spermatophytes (including seagrasses) | <i>Cymodocea manatorum</i> | 16 | | | |
| | <i>Cymodocea</i> sp. | 1 | 17 | | |
| | <i>Halophila baillonis</i> | 7 | 7 | | |
| | <i>Ruppia maritima</i> | 1 | 1 | | |
| | <i>Thalassia</i> sp. | 1 | | | |
| | <i>Thalassia testudineum</i> | 23 | 24 | | |
| | <i>Vaucheria</i> sp. | 1 | 1 | 50 | |
| Algae | Algae * | 3 | | 3 | |
| Rhodophyta ^a | <i>Acanthophora spicifera</i> | 8 | 8 | | |
| | <i>Acrochaetium</i> sp. | 2 | 2 | | |
| | <i>Amphiroa fragilissima</i> | 14 | 14 | | |
| | <i>Anacystis marina</i> | 3 | | | |
| | <i>Anacystis</i> sp. | 2 | 5 | | |
| | <i>Asterocystis ramosa</i> | 2 | 2 | | |
| | <i>Avrainvillea nigricans</i> | 1 | | | |
| Chlorophyta ^b | <i>Avrainvillea rawsonii</i> | 1 | | | |
| | <i>Avrainvillea</i> sp. | 1 | 3 | | |
| Cyanophyta ^c | Bluegreens | 1 | 1 | | |
| | <i>Botryocladia occidentalis</i> | 1 | 1 | | |
| Chlorophyta | <i>Bryopsis pennata</i> | 4 | | | |
| | <i>Bryopsis</i> sp. | 2 | 6 | | |
| | <i>Bryothamnion triquetrum</i> | 2 | 2 | | |
| Rhodophyta | <i>Callithamnion</i> sp. | 1 | 1 | | |
| | <i>Calothrix crustacea</i> | 2 | | | |
| | <i>Calothrix</i> sp. | 2 | 4 | | |
| Chlorophyta | <i>Caulerpa cupressoides</i> | 2 | | | |
| | <i>Caulerpa mexicana</i> | 1 | | | |
| | <i>Caulerpa prolifera</i> | 1 | | | |
| | <i>Caulerpa racemosa</i> | 5 | | | |
| | <i>Caulerpa sertularioides</i> | 5 | | | |
| | <i>Caulerpa</i> sp. | 1 | | | |
| | <i>Caulerpa taxifolia</i> | 2 | | | |
| | <i>Caulerpa vickersiae</i> | 1 | 18 | | |
| | Rhodophyta | <i>Centroceras clavulatum</i> | 14 | | |
| | | <i>Centroceras</i> sp. | 7 | 21 | |
| <i>Ceramium byssoideum</i> | | 12 | | | |
| <i>Ceramium fastigiatum</i> | | 1 | | | |
| <i>Ceramium nitens</i> | | 6 | | | |
| Chlorophyta | <i>Ceramium</i> sp. | 18 | 37 | | |
| | <i>Chaetomorpha</i> sp. | 1 | 1 | | |
| Rhodophyta | <i>Champia parvula</i> | 2 | 2 | | |
| | <i>Chondria littoralis</i> | 1 | | | |
| | <i>Chondria</i> sp. | 1 | 2 | | |
| Chlorophyta | <i>Cladophora delicatula</i> | 3 | | | |
| | <i>Cladophora</i> sp. | 8 | 11 | | |
| | <i>Codium intertextum</i> | 1 | | | |
| | <i>Codium isthmocladum</i> | 1 | 2 | | |
| | <i>Coelothrix irregularis</i> | 12 | 12 | | |
| | <i>Colpomenia sinuosa</i> | 1 | 1 | | |
| | Rhodophyta | <i>Corallina cubensis</i> | 3 | | |
| | | <i>Corallina subulata</i> | 1 | 4 | |
| <i>Cryptonemia crenulata</i> | | 1 | 1 | | |
| Diatoms | | 17 | 17 | | |
| Phaeophyta ^d | <i>Dichothrix penicillata</i> | 1 | | | |
| | <i>Dichothrix</i> sp. | 1 | 2 | | |
| | <i>Dictyopteris delicatula</i> | 10 | | | |
| | <i>Dictyopteris justii</i> | 1 | | | |
| | <i>Dictyopteris plagiogramma</i> | 2 | | | |
| | <i>Dictyopteris</i> sp. | 1 | 14 | | |
| | <i>Dictyosphaeria favulosa</i> | 2 | 2 | | |
| Chlorophyta Phaeophyta | <i>Dictyota bartayresii</i> | 4 | | | |
| | <i>Dictyota ciliata</i> | 1 | | | |
| | <i>Dictyota dentata</i> | 3 | | | |
| | <i>Dictyota divaricata</i> | 13 | | | |
| | <i>Dictyota linearis</i> | 1 | | | |
| | <i>Dictyota</i> sp. | 10 | 32 | | |

continued...

Table 8.5.1 continued

| Taxon and class | Species | Occurrence | Sum | |
|-----------------|------------------------------------|------------|-------|--------------|
| | | | Genus | Higher taxon |
| Rhodophyta | <i>Digenia simplex</i> | 1 | 1 | |
| | <i>Dilophus guineensis</i> | 1 | 1 | |
| Phaeophyta | <i>Ectocarpus breviarticulatus</i> | 1 | | |
| | <i>Ectocarpus</i> sp. | 2 | | 3 |
| | <i>Endesme zosteræ</i> | 1 | | 1 |
| | <i>Enteromorpha flexuosa</i> | 6 | | |
| | <i>Enteromorpha</i> sp. | 18 | | 24 |
| | <i>Eucheuma acanthocladum</i> | 1 | | |
| | <i>Eucheuma echinocarpium</i> | 2 | | 3 |
| Rhodophyta | <i>Galaxura lapidescens</i> | 2 | | |
| | <i>Galaxura</i> sp. | 1 | | |
| | <i>Galaxura squalida</i> | 1 | | 4 |
| | <i>Gelidiella acerosa</i> | 1 | | 1 |
| | <i>Gelidium corneum</i> | 6 | | |
| | <i>Gelidium pusillum</i> | 5 | | |
| | <i>Gelidium rigidulum</i> | 3 | | |
| | <i>Gelidium</i> sp. | 8 | | 22 |
| Phaeophyta | <i>Giffordia mitchellæ</i> | 1 | | 1 |
| Rhodophyta | <i>Gracilaria debilis</i> | 1 | | |
| | <i>Gracilaria mamillaris</i> | 1 | | |
| | <i>Gracilaria</i> sp. | 2 | | |
| | <i>Gracilaria verrucosa</i> | 1 | | 5 |
| | <i>Grateloupia cuneifolia</i> | 1 | | 1 |
| Chlorophyta | <i>Halimeda incrassata</i> | 2 | | |
| | <i>Halimeda monile</i> | 2 | | |
| | <i>Halimeda opuntia</i> | 8 | | |
| | <i>Halimeda</i> sp. | 5 | | 17 |
| Rhodophyta | <i>Halymenia floresia</i> | 1 | | 1 |
| | <i>Herposiphonia secunda</i> | 4 | | |
| | <i>Herposiphonia</i> sp. | 13 | | |
| | <i>Herposiphonia tenella</i> | 1 | | 18 |
| | <i>Heterosiphonia wurdemanni</i> | 2 | | 2 |
| | <i>Hypnea musciformis</i> | 7 | | |
| | <i>Hypnea spinella</i> | 5 | | 12 |
| | <i>Jania capillacea</i> | 5 | | |
| | <i>Jania rubens</i> | 7 | | |
| | <i>Jania</i> sp. | 1 | | 13 |
| | <i>Laurencia corallopsis</i> | 1 | | |
| | <i>Laurencia gemnifera</i> | 1 | | |
| | <i>Laurencia obtusa</i> | 11 | | |
| | <i>Laurencia papillosa</i> | 6 | | |
| | <i>Laurencia</i> sp. | 3 | | 22 |
| | <i>Liagora ceranoides</i> | 3 | | 3 |
| | <i>Lithothamnion</i> sp. | 1 | | 1 |
| | <i>Lomentaria uncinata</i> | 1 | | 1 |
| | <i>Lyngbya majuscula</i> | 14 | | |
| | <i>Lyngbya</i> sp. | 18 | | 32 |
| | <i>Microcoleus chthonoplastes</i> | 3 | | |
| | <i>Microcoleus</i> sp. | 6 | | 9 |
| | <i>Nitophyllum</i> sp. | 1 | | 1 |
| Cyanophyta | <i>Oscillatoria</i> sp. | 4 | | 4 |
| Phaeophyta | <i>Padina gymnospora</i> | 2 | | |
| | <i>Padina sanctæ-crucis</i> | 2 | | |
| | <i>Padina</i> sp. | 1 | | 5 |
| Chlorophyta | <i>Penicillus pyriformis</i> | 1 | | 1 |
| Rhodophyta | <i>Peyssonnelia</i> sp. | 1 | | 1 |
| | <i>Phormidium penicellatum</i> | 1 | | |
| | <i>Phormidium</i> sp. | 2 | | 3 |
| | <i>Plectonema nostocorum</i> | 2 | | 2 |
| | <i>Pocockiella variegata</i> | 5 | | |
| | <i>Pocockiella</i> sp. | 1 | | 6 |
| | <i>Polysiphonia binneyi</i> | 1 | | |
| | <i>Polysiphonia ferrulacea</i> | 13 | | |
| | <i>Polysiphonia</i> sp. | 21 | | 35 |
| | <i>Rhizoclonium ripanum</i> | 6 | | |
| | <i>Rhizoclonium</i> sp. | 4 | | 10 |
| Phaeophyta | <i>Sargassum fluitans</i> | 3 | | |
| | <i>Sargassum hystrix</i> | 2 | | |
| | <i>Sargassum natans</i> | 3 | | |
| | <i>Sargassum platycarpum</i> | 3 | | |

continued...

Table 8.5.1 continued

| Taxon and class | Species | Occurrence | Sum | |
|-----------------|----------------------------------|------------|-------|--------------|
| | | | Genus | Higher taxon |
| | <i>Sargassum polyceratum</i> | 1 | | |
| | <i>Sargassum pteropleuron</i> | 1 | | |
| | <i>Sargassum rigidulum</i> | 1 | | |
| | <i>Sargassum</i> sp. | 4 | 18 | |
| | <i>Spatoglossum schroderi</i> | 1 | 1 | |
| Rhodophyta | <i>Spermothamnion investiens</i> | 2 | 2 | |
| Phaeophyta | <i>Sphacelaria tribuloides</i> | 2 | | |
| | <i>Sphacelaria</i> sp. | 1 | 3 | |
| | <i>Spirulina</i> sp. | 2 | 2 | |
| Rhodophyta | <i>Spyridia filamentosa</i> | 7 | 7 | |
| Phaeophyta | <i>Turbinaria turbinata</i> | 2 | 2 | |
| Chlorophyta | <i>Udotea flabellum</i> | 2 | 2 | |
| | <i>Ulva lactuca</i> | 4 | | |
| | <i>Ulva</i> sp. | 3 | 7 | |
| | <i>Valonia utricularis</i> | 1 | 2 | |
| | <i>Valonia ventricosa</i> | 1 | | |
| | <i>Vidalia obtusiloba</i> | 1 | 1 | |
| Rhodophyta | <i>Wrangelia argus</i> | 2 | 2 | 543 |

^aRhodophyta (red algae).

^bChlorophyta (green algae).

^cCyanophyta (blue-green algae).

^dPhaeophyta (brown algae).

Table 8.5.2. Invertebrates in stomachs of West Indian reef fishes (based on Randall 1967); items are listed in systematical order; "occurrence" refers to number of fish species with respective item in stomach.

| Taxon | Species | Occurrence | Genus | Sum Higher taxon |
|----------------------------------|-------------------------------------|------------|-------|---------------------|
| Foraminifera | - | 6 | - | 6 |
| Sponges | - | 20 | - | 21 |
| | <i>Mycale</i> sp. | 1 | 1 | |
| Hydrozoans | <i>Hydroids</i> | 12 | - | 16 |
| | <i>Millepora alvicornis</i> | 2 | 2 | |
| | <i>Pennaria</i> sp. | 1 | 1 | |
| | <i>Sertularia</i> sp. | 1 | 1 | |
| Gorgonians (Sea fans) | - | 6 | - | 14 |
| | <i>Eunicea calyculata</i> | 1 | 1 | |
| | <i>Gorgonia flabellum</i> | 2 | 2 | |
| | <i>Muricea atlantica</i> | 1 | | |
| | <i>Muricea laxa</i> | 1 | | |
| | <i>Muricea</i> sp. | 1 | 3 | |
| | <i>Plexaura flexuosa</i> | 1 | 1 | |
| | <i>Pterogorgia</i> sp. | 1 | 1 | |
| Alcyonarians (Leather corals) | - | 1 | - | 1 |
| Zoanthids (Sea anemones) | - | 1 | - | 22 |
| | <i>Bunodactis stelloides</i> | 1 | 1 | |
| | <i>Bunodosoma granulifera</i> | 1 | 1 | |
| | <i>Palythoa</i> sp. | 1 | 1 | |
| | <i>Phyllactis flosculifera</i> | 2 | 2 | |
| | <i>Rhodactis sanctithomae</i> | 1 | 1 | |
| | <i>Zoanthus pulchellus</i> | 2 | | |
| | <i>Zoanthus sociatus</i> | 5 | | |
| | <i>Zoanthus</i> sp. | 8 | 15 | |
| Scleractinia (Stony corals) | - | 7 | - | 10 |
| | <i>Cladocora arbuscula</i> | 1 | 1 | |
| | <i>Colpophyllia</i> sp. | 1 | 1 | |
| | <i>Oculina diffusa</i> | 1 | 1 | |
| Bryozoans | - | 6 | - | 6 |
| Sipunculids | - | 8 | - | 22 |
| | <i>Aspidosiphon cumingi</i> | 1 | | |
| | <i>Aspidosiphon</i> sp. | 9 | | |
| | <i>Aspidosiphon speciosus</i> | 1 | | |
| | <i>Aspidosiphon spinosocutatus</i> | 1 | 12 | |
| | <i>Siphonosoma cumanensis</i> | 2 | 2 | |
| Priapuloids | - | 1 | - | 1 |
| Chitons | - | 7 | - | 20 |
| | <i>Acanthochiton hemphilli</i> | 1 | 1 | |
| | <i>Acanthochitona pygmaea</i> | 3 | | |
| | <i>Acanthochitona</i> sp. | 1 | 4 | |
| | <i>Calloplax janeirensis</i> | 1 | 1 | |
| | <i>Choneplax lata</i> | 2 | 2 | |
| | <i>Ischnochiton papillosus</i> | 4 | | |
| | <i>Ischnochiton</i> sp. | 1 | 5 | |
| Gastropods | - | 15 | - | 328 |
| | acmaeids (limpets) | 8 | - | |
| | acteonids | 1 | - | |
| | alcidids | 1 | - | |
| | buccinids | 1 | - | |
| | columbellids | 1 | - | |
| | muricids | 1 | - | |
| | turbinids | 2 | - | |
| | turrids | 4 | - | |
| | vermetids | 3 | - | |
| | <i>Acmaea antillarum</i> (limpets) | 4 | | |
| | <i>Acmaea leucopleura</i> (limpets) | 3 | | |

continued...

Table 8.5.2 continued

| Taxon | Species | Occurrence | Genus | Sum Higher taxon |
|-------|-----------------------------------------|------------|-------|---------------------|
| | <i>Acmaea pustulata</i> (limpets) | 10 | | |
| | <i>Acmaea</i> sp. | 6 | 23 | |
| | <i>Alvania auberiana</i> | 2 | 2 | |
| | <i>Anachis sparsa</i> | 1 | | |
| | <i>Anachis</i> sp. | 5 | 6 | |
| | <i>Arene tricarinata</i> | 2 | | |
| | <i>Arene</i> sp. | 5 | 7 | |
| | <i>Astraea caelata</i> | 1 | | |
| | <i>Astraea longispina</i> | 2 | | |
| | <i>Astraea ruber</i> | 2 | | |
| | <i>Astraea</i> sp. | 4 | 9 | |
| | <i>Atys riiseana</i> | 1 | | |
| | <i>Atys</i> sp. | 1 | 2 | |
| | <i>Bailya</i> sp. | 1 | 1 | |
| | <i>Balcis intermedia</i> (in Holoth.?) | 1 | 1 | |
| | <i>Batillaria</i> sp. | 1 | 1 | |
| | <i>Bittium varium</i> | 4 | 1 | |
| | <i>Bulla</i> sp. | 1 | | |
| | <i>Bulla striata</i> | 1 | 2 | |
| | <i>Bullata ovuliformis</i> | 2 | 2 | |
| | <i>Caecum nitidum</i> | 1 | | |
| | <i>Caecum pulchellum</i> | 3 | | |
| | <i>Caecum</i> sp. | 1 | 5 | |
| | <i>Calliostoma javanicum</i> | 1 | | |
| | <i>Calliostoma</i> sp. | 3 | 4 | |
| | <i>Cerithiopsis emersoni</i> | 1 | 1 | |
| | <i>Cerithium algicola</i> | 3 | | |
| | <i>Cerithium eburneum</i> | 1 | | |
| | <i>Cerithium litteratum</i> | 3 | | |
| | <i>Cerithium</i> sp. | 5 | 12 | |
| | <i>Cittarium pica</i> | 2 | 2 | |
| | <i>Columbella mercatoria</i> | 9 | 9 | |
| | <i>Conus juliae</i> | 1 | 1 | |
| | <i>Coralliophila caribea</i> | 1 | 1 | |
| | <i>Crassispira fuscescens</i> | 1 | | |
| | <i>Crassispira nigrescens</i> | 1 | 2 | |
| | <i>Crassostrea rhizophorae</i> | 2 | 2 | |
| | <i>Crucibulum auricula</i> | 1 | 1 | |
| | <i>Cyclostremiscus ornatus</i> | 1 | 1 | |
| | <i>Cymatium labiosum</i> | 1 | 1 | |
| | <i>Daphnella lymneiformis</i> | 1 | 1 | |
| | <i>Diodora cayensis</i> | 1 | | |
| | <i>Diodora listeri</i> | 3 | | |
| | <i>Diodora</i> sp. | 3 | | |
| | <i>Diodora variegata</i> | 1 | | |
| | <i>Diodora viridula</i> | 1 | 9 | |
| | <i>Drillia</i> sp. | 1 | 1 | |
| | <i>Emarginula pumila</i> | 2 | 2 | |
| | <i>Epitonium</i> sp. | 2 | | |
| | <i>Epitonium turritellulum</i> | 1 | 3 | |
| | <i>Fasciolaria tulipa</i> | 2 | 2 | |
| | <i>Fissurella angusta</i> (limpets) | 1 | | |
| | <i>Fissurella barbadensis</i> (limpets) | 6 | | |
| | <i>Fissurella nodosa</i> (limpets) | 1 | | |
| | <i>Fissurella</i> sp. (limpets) | 6 | 14 | |
| | <i>Haminoea elegans</i> | 3 | 3 | |
| | <i>Hemitoma octoradiata</i> | 5 | 5 | |
| | <i>Hipponix subrufus</i> | 1 | 1 | |
| | <i>Hyalina albolineata</i> | 3 | | |
| | <i>Hyalina avena</i> | 1 | | |
| | <i>Hyalina</i> sp. | 6 | 10 | |
| | <i>Litiopa melanostoma</i> | 1 | 1 | |
| | <i>Littorina angulifera</i> | 1 | | |
| | <i>Littorina meleagris</i> | 1 | | |
| | <i>Littorina ziczac</i> | 1 | 3 | |
| | <i>Mangelia</i> sp. | 2 | 2 | |
| | <i>Marginella</i> sp. | 1 | 1 | |
| | <i>Melampus coffeus</i> | 2 | 2 | |
| | <i>Mitra barbadensis</i> | 1 | | |
| | <i>Mitra</i> sp. | 1 | | |

continued...

Table 8.5.2 continued

| Taxon | Species | Occurrence | Sum | |
|----------|---------------------------------------|------------|-------|--------------|
| | | | Genus | Higher taxon |
| | <i>Mitrella lunata</i> | 3 | | |
| | <i>Mitrella nitens</i> | 1 | 6 | |
| | <i>Modulus modiolus</i> | 9 | 9 | |
| | <i>Murex pomum</i> | 2 | | |
| | <i>Murex</i> sp. | 1 | 3 | |
| | <i>Muricopsis hexagonus</i> | 1 | 1 | |
| | <i>Nassarius albus</i> | 2 | | |
| | <i>Nassarius</i> sp. | 5 | 7 | |
| | <i>Natica canrena</i> | 1 | | |
| | <i>Natica</i> sp. (juvenile) | 1 | 2 | |
| | <i>Nerita peloronta</i> | 1 | 1 | |
| | <i>Nitidella laevigata</i> | 1 | | |
| | <i>Nitidella nitida</i> | 2 | | |
| | <i>Nitidella ocellata</i> | 3 | | |
| | <i>Nitidella</i> sp. | 3 | 9 | |
| | <i>Oliva caribaeensis</i> | 1 | | |
| | <i>Oliva reticularis</i> | 1 | | |
| | <i>Oliva</i> sp. | 2 | 4 | |
| | <i>Olivella nivea</i> | 1 | | |
| | <i>Olivella petiolita</i> | 1 | | |
| | <i>Olivella</i> sp. | 8 | 10 | |
| | <i>Persicula lavalleeana</i> | 4 | 4 | |
| | <i>Petalococonchus</i> sp. (vermetid) | 1 | 1 | |
| | <i>Planaxis lineatus</i> | 2 | 2 | |
| | <i>Polinices lacteus</i> | 1 | 1 | |
| | <i>Pseudostomatella coccinea</i> | 3 | | |
| | <i>Pseudostomatella erythrocoma</i> | 1 | 4 | |
| | <i>Purpura patula</i> (thaid) | 1 | 1 | |
| | <i>Pusia</i> sp. | 1 | 1 | |
| | <i>Pyramidella candida</i> | 1 | 1 | |
| | <i>Pyrgocythara</i> sp. | 1 | 1 | |
| | <i>Retusa</i> sp. | 1 | 1 | |
| | <i>Risomurex roseus</i> | 1 | 1 | |
| | <i>Rissoina cancellata</i> | 1 | | |
| | <i>Rissoina fischeri</i> | 1 | | |
| | <i>Rissoina</i> sp. | 1 | 3 | |
| | <i>Schismope</i> sp. | 1 | 1 | |
| | <i>Seila adamsi</i> | 1 | 1 | |
| | <i>Sinum perspectivum</i> | 1 | 1 | |
| | <i>Siphonaria</i> sp. (limpets) | 2 | 2 | |
| | <i>Smaragdia viridis</i> | 3 | 3 | |
| | <i>Strombus gallus</i> | 1 | | |
| | <i>Strombus gigas</i> | 12 | | |
| | <i>Strombus</i> sp. | 2 | 15 | |
| | <i>Synaptocochlea picta</i> | 2 | 2 | |
| | <i>Tegula excavata</i> | 1 | | |
| | <i>Tegula fasciata</i> | 6 | | |
| | <i>Tegula lividomaculata</i> | 2 | | |
| | <i>Tegula</i> sp. | 2 | 11 | |
| | <i>Tricolia adamsi</i> | 3 | | |
| | <i>Tricolia bella</i> | 1 | | |
| | <i>Tricolia tessellata</i> | 3 | 7 | |
| | <i>Tridachia crispata</i> | 1 | 1 | |
| | <i>Trivia</i> sp. | 1 | 1 | |
| | <i>Turbo castanea</i> | 7 | | |
| | <i>Turbo</i> sp. | 1 | 8 | |
| | <i>Turbonilla</i> sp. | 1 | 1 | |
| | <i>Turritella</i> sp. | 1 | 1 | |
| | <i>Vasum muricatum</i> | 1 | 1 | |
| | <i>Zebina browniana</i> | 3 | 3 | |
| Bivalves | - | 13 | - | 143 |
| | arcids | 1 | - | |
| | ostreids | 1 | - | |
| | pinnids | 1 | - | |
| | <i>Aequipecten gibbus</i> | 2 | | |
| | <i>Aequipecten lineolaris</i> | 1 | 3 | |
| | <i>Americardia guppyi</i> | 2 | | |
| | <i>Americardia media</i> | 1 | 3 | |
| | <i>Antigona listeri</i> | 1 | 1 | |

continued...

Table 8.5.2 continued

| Taxon | Species | Occurrence | Genus | Sum Higher taxon |
|------------|----------------------------------|------------|-------|---------------------|
| | <i>Arca imbricata</i> | 1 | | |
| | <i>Arca</i> sp. | 1 | | |
| | <i>Arca umbonata</i> | 1 | | |
| | <i>Arca zebra</i> | 3 | 6 | |
| | <i>Arcopsis adamsi</i> | 2 | 2 | |
| | <i>Asaphis deflorata</i> | 1 | 1 | |
| | <i>Atrina rigida</i> | 1 | | |
| | <i>Atrina seminuda</i> | 1 | 2 | |
| | <i>Barbatia cancellaria</i> | 3 | | |
| | <i>Babartia domingensis</i> | 2 | 5 | |
| | <i>Brachidontes citrinus</i> | 1 | | |
| | <i>Brachidontes exustus</i> | 3 | | |
| | <i>Brachidontes</i> sp. | 1 | 5 | |
| | <i>Chama sarda</i> | 1 | | |
| | <i>Chama</i> sp. | 1 | 2 | |
| | <i>Chione cancellata</i> | 1 | | |
| | <i>Chione paphia</i> | 1 | | |
| | <i>Chione</i> sp. | 1 | 3 | |
| | <i>Chlamys</i> sp. | 1 | 1 | |
| | <i>Codakia costata</i> | 1 | | |
| | <i>Codakia orbicularis</i> | 1 | | |
| | <i>Codakia pectinella</i> | 1 | 3 | |
| | <i>Corbula contracta</i> | 1 | 1 | |
| | <i>Crassinella</i> sp. | 1 | 1 | |
| | <i>Crassostrea rhizophorae</i> | 2 | 2 | |
| | <i>Crenella divaricata</i> | 1 | 1 | |
| | <i>Cumingia antillarum</i> | 4 | 4 | |
| | <i>Divaricella quadrisulcata</i> | 1 | 1 | |
| | <i>Ervilia nitens</i> | 2 | 2 | |
| | <i>Glycymeris decussata</i> | 1 | | |
| | <i>Glycymeris pectinata</i> | 1 | | |
| | <i>Glycymeris</i> sp. | 1 | 3 | |
| | <i>Gouldia cerina</i> | 2 | 2 | |
| | <i>Isognomon alatus</i> | 1 | | |
| | <i>Isognomon radiatus</i> | 3 | | |
| | <i>Isognomon</i> sp. | 2 | 6 | |
| | <i>Laevicardium</i> sp. | 7 | 7 | |
| | <i>Lima pellucida</i> | 1 | 1 | |
| | <i>Lithophaga bisulcata</i> | 1 | 1 | |
| | <i>Lyropecten antillarum</i> | 1 | 1 | |
| | <i>Macoma cerina</i> | 2 | 2 | |
| | <i>Microcardium</i> sp. | 1 | 1 | |
| | <i>Musculus lateralis</i> | 5 | 5 | |
| | <i>Nuculana</i> sp. | 1 | 1 | |
| | <i>Ostrea</i> sp. | 3 | 3 | |
| | <i>Papyridea semisulcata</i> | 2 | | |
| | <i>Papyridea soleniformis</i> | 1 | | |
| | <i>Papyridea</i> sp. | 1 | 4 | |
| | <i>Pecten ziczac</i> | 1 | | |
| | <i>Pecten</i> sp. | 4 | 5 | |
| | <i>Pinctada radiata</i> | 2 | | |
| | <i>Pinctada</i> sp. | 2 | 4 | |
| | <i>Pinna carnea</i> | 1 | 1 | |
| | <i>Pitar fulminata</i> | 1 | | |
| | <i>Pitar</i> sp. | 3 | 4 | |
| | <i>Pseudochama</i> sp. | 1 | 1 | |
| | <i>Pteria colymbus</i> | 1 | 1 | |
| | <i>Solemya occidentalis</i> | 3 | 3 | |
| | <i>Spondylus</i> sp. | 1 | 1 | |
| | <i>Tellina caribaea</i> | 2 | | |
| | <i>Tellina listeri</i> | 1 | | |
| | <i>Tellina</i> sp. | 11 | 14 | |
| | <i>Trachycardium magnum</i> | 2 | | |
| | <i>Trachycardium muricatum</i> | 1 | | |
| | <i>Trachycardium</i> sp. | 2 | 5 | |
| | <i>Volsella</i> sp. | 1 | 1 | |
| | <i>Yoldia perprotracta</i> | 1 | 1 | |
| Scaphopods | - | 3 | - | 13 |
| | <i>Cadulus acus</i> | 1 | | |

continued...

Table 8.5.2 continued

| Taxon | Species | Occurrence | Sum | |
|-------------|----------------------------------------------|------------|-------|--------------|
| | | | Genus | Higher taxon |
| | <i>Cadulus</i> sp. | 3 | 4 | |
| | <i>Dentalium semistriolatum</i> | 1 | | |
| | <i>Dentalium</i> sp. | 5 | 6 | |
| Squids | - | 11 | - | 14 |
| | <i>Dorytheutis plei</i> | 1 | | |
| | <i>Dorytheutis</i> sp. | 1 | 2 | |
| | <i>Sepiateuthis sepioidea</i> | 1 | 1 | |
| Octopuses | - | 18 | - | 20 |
| | <i>Octopus briareus</i> | 1 | | |
| | <i>Octopus vulgaris</i> | 1 | 2 | |
| Polychaetes | - | 51 | - | 86 |
| | ampharetids | 1 | - | |
| | arabellids | 1 | - | |
| | capitellids | 1 | - | |
| | chaetopterids | 1 | - | |
| | eunicids | 2 | - | |
| | flabelligerids | 1 | - | |
| | glycerids | 1 | - | |
| | lumbrinerids | 1 | - | |
| | maldanids | 1 | - | |
| | neroids | 1 | - | |
| | onuphids | 2 | - | |
| | opheliids | 1 | - | |
| | pectinariids | 1 | - | |
| | polynoids | 1 | - | |
| | sabellids | 3 | - | |
| | serpulids | 4 | - | |
| | syllids | 1 | - | |
| | terebellids | 3 | - | |
| | <i>Cariboa</i> sp. | 1 | 1 | |
| | <i>Chloeia</i> sp. | 2 | 2 | |
| | <i>Eunice</i> sp. (fire worm) | 3 | 3 | |
| | <i>Sabellastarte magnifica</i> (palolo worm) | 1 | 1 | |
| | <i>Semiodera</i> sp. | 1 | 1 | |
| Echiuroids | - | 2 | - | 2 |
| Pycnogonids | - | 2 | - | 2 |
| Barnacles | appendages | 5 | - | 5 |
| Stomatopods | - | 26 | - | 53 |
| | <i>Gonodactylus curacaoensis</i> | 2 | | |
| | <i>Gonodactylus oerstedii</i> | 6 | | |
| | <i>Gonodactylus</i> sp. | 7 | 15 | |
| | <i>Lysiosquilla glabriuscula</i> | 2 | 2 | |
| | <i>Nannosquilla</i> | 1 | 1 | |
| | <i>Pseudosquilla ciliata</i> | 6 | 6 | |
| | <i>Squilla</i> | 1 | | |
| | <i>Squilla tricarinata</i> | 1 | 2 | |
| | <i>Squilla</i> sp. | 1 | 1 | |
| Amphipods | - | 29 | - | 64 |
| | caprellids | 5 | - | |
| | gammarids | 3 | - | |
| | hyperiids | 8 | - | |
| | isaeids | 1 | - | |
| | lysianassids (gammarid) | 1 | - | |
| | metopids | 1 | - | |
| | photids | 1 | - | |
| | <i>Ampelisca</i> sp. (gammarid) | 1 | 1 | |
| | <i>Colomastix</i> sp. | 1 | 1 | |
| | <i>Corophium</i> sp. | 1 | 1 | |
| | <i>Cymadusa filosa</i> | 1 | | |
| | <i>Cymadusa</i> sp. | 1 | 2 | |
| | <i>Elasmopus</i> sp. | 2 | 2 | |

continued.

Table 8.5.2 continued

| Taxon | Species | Occurrence | Genus | Sum Higher taxon |
|--------------------|--------------------------------------------------------|------------|-------|---------------------|
| | <i>Eurystheus</i> sp. | 1 | 1 | |
| | <i>Hyale</i> sp. | 1 | 1 | |
| | <i>Leucothoe</i> sp. | 1 | 1 | |
| | <i>Magamphopus</i> sp. | 1 | 1 | |
| | <i>Neomicrodentopus</i> sp. | 1 | 1 | |
| | <i>Paraphoxus</i> sp. | 1 | 1 | |
| | <i>Photis</i> sp. | 2 | 2 | |
| Tanaids | - | 14 | - | 15 |
| | apseudids | 1 | - | |
| Isopods | - | 34 | - | 51 |
| | anthurids | 1 | - | |
| | cymothoids | 1 | - | |
| | excorallanids | 2 | - | |
| | flabelliferans | 1 | - | |
| | gnathiids (larvae) | 2 | - | |
| | sphaeromids | 2 | - | |
| | <i>Asellote</i> sp. | 1 | 1 | |
| | <i>Cymodoce</i> sp. | 1 | 1 | |
| | <i>Excorallana antillensis</i> | 1 | | |
| | <i>Excorallana</i> sp. | 1 | 2 | |
| | <i>Paracerceis caudata</i> | 3 | 3 | |
| | <i>Stenetrium</i> sp. | 1 | 1 | |
| Shrimps | - | 32 | - | 166 |
| | alpheids | 40 | - | |
| | axiids | 2 | - | |
| | carideans | 24 | - | |
| | gnathophyllids | 6 | - | |
| | hippolytids | 1 | - | |
| | palaemonids | 7 | - | |
| | penaeids | 23 | - | |
| | stenopids | 1 | - | |
| | stenopodids | 1 | - | |
| | <i>Alpheus</i> sp. | 1 | 1 | |
| | <i>Brachycarpus biunguiculatus</i> | 1 | 1 | |
| | <i>Calianassa</i> sp. | 2 | 2 | |
| | <i>Gnathophylloides mineri</i> | 1 | 1 | |
| | <i>Lucifer faxani</i> | 1 | | |
| | <i>Lucifer</i> sp. | 2 | 3 | |
| | <i>Lysmata moorei</i> (cleaner shrimp) | 1 | 1 | |
| | <i>Penaeopsis goodei</i> | 1 | | |
| | <i>Penaeopsis</i> sp. | 1 | 2 | |
| | <i>Periclimenes</i> sp. (cleaner shrimp) | 3 | 3 | |
| | <i>Processa</i> sp. | 1 | 1 | |
| | <i>Sicyonia laevigata</i> | 1 | 1 | |
| | <i>Sicyonia</i> sp. | 1 | 1 | |
| | <i>Solenocera</i> sp. | 1 | 1 | |
| | <i>Stenopus hispidus</i> (cleaner shrimp) | 2 | 2 | |
| | <i>Synalpheus</i> sp. | 2 | 2 | |
| | <i>Tozeuma</i> sp. | 3 | 3 | |
| | <i>Trachycaris restrictus</i> | 1 | 1 | |
| | <i>Trachypenaeus</i> sp. | 2 | 2 | |
| | <i>Upogebia</i> sp. (Thalassinids) | 1 | 1 | |
| Spiny lobsters | - | 1 | - | 7 |
| | <i>Panulirus argus</i> | 4 | | |
| | <i>Panulirus guttatus</i> | 2 | 6 | |
| Scyllarid lobsters | - | 3 | - | 5 |
| | <i>Scyllarides aequinoctialis</i> (slipper lobster) | 2 | 2 | |
| Hermit crabs | - | 21 | - | 49 |
| | diogenids | 3 | - | |
| | pagurids | 3 | - | |
| | <i>Calcinus fibiceus</i> | 1 | | |
| | <i>Calcinus sulcatus</i> | 1 | 2 | |
| | <i>Clibanarius</i> sp. | 1 | | |

continued...

Table 8.5.2 continued

| Taxon | Species | Occurrence | Genus | Sum Higher taxon |
|---------------|------------------------------------|------------|-------|---------------------|
| | <i>Clibanarius tricolor</i> | 2 | 3 | |
| | <i>Dardanus venosus</i> | 2 | 2 | |
| | <i>Paguristes anomalus</i> | 2 | | |
| | <i>Paguristes depressus</i> | 3 | | |
| | <i>Paguristes grayi</i> | 2 | | |
| | <i>Paguristes</i> sp. | 2 | | |
| | <i>Paguristes wassi</i> | 1 | 10 | |
| | <i>Pagurus bonairensis</i> | 1 | | |
| | <i>Pagurus miamensis</i> | 1 | 2 | |
| | <i>Petrochirus diogenes</i> | 2 | 2 | |
| | <i>Spiropagurus</i> sp. | 1 | 1 | |
| Crabs | - | 33 | - | 258 |
| | calappids | 9 | - | |
| | gonoplacid (<i>Speocarcinus</i>) | 3 | - | |
| | grapsids | 3 | - | |
| | hippids | 4 | - | |
| | leucosids | 1 | - | |
| | majids | 29 | - | |
| | oxystomes | 1 | - | |
| | parthenopids | 2 | - | |
| | pinotherids | 1 | - | |
| | porcellanids | 6 | - | |
| | portunids | 24 | - | |
| | raninids | 2 | - | |
| | xanthids | 17 | - | |
| Majidae | <i>Acanthonyx petiverii</i> | 1 | 1 | |
| | <i>Actaea acantha</i> | 1 | 1 | |
| Albuneidae | <i>Albunea gibbesii</i> | 2 | 2 | |
| | <i>Arenaeus cribarius</i> | 1 | 1 | |
| Calappidae | <i>Calappa flammea</i> | 1 | | |
| | <i>Calappa gallus</i> | 2 | | |
| | <i>Calappa ocellata</i> | 2 | | |
| | <i>Calappa</i> sp. | 2 | 7 | |
| | <i>Callinectes</i> sp. | 1 | 1 | |
| Xanthidae | <i>Carpilius corallinus</i> | 2 | 2 | |
| | <i>Chorinus heros</i> | 1 | 1 | |
| | <i>Cronius ruber</i> | 5 | | |
| | <i>Cronius tumidulus</i> | 2 | 7 | |
| | <i>Domecia hispida</i> | 4 | 4 | |
| Leucosidae | <i>Ebalia stimpsonii</i> | 1 | 1 | |
| Hippidae | <i>Emerita portoricensis</i> | 1 | | |
| | <i>Emerita</i> sp. | 1 | 2 | |
| | <i>Euryplax nitida</i> | 1 | 1 | |
| Majidae | <i>Herbstia</i> sp. | | | |
| | <i>Leiolambrus nitidus</i> | 1 | 1 | |
| | <i>Leptodius floridanus</i> | 1 | 1 | |
| | <i>Macrocoelema</i> sp. | 1 | 1 | |
| | <i>Microphrys bicornutus</i> | 2 | 2 | |
| | <i>Mithrax cinctimanus</i> | 1 | | |
| | <i>Mithrax coryphe</i> | 3 | | |
| | <i>Mithrax forceps</i> | 1 | | |
| | <i>Mithrax sculptus</i> | 7 | | |
| | <i>Mithrax</i> sp. | 16 | | |
| | <i>Mithrax verrucosus</i> | 1 | 29 | |
| | <i>Ocypode albicans</i> | 1 | 1 | |
| | <i>Panopeus</i> sp. | 2 | 2 | |
| Parthenopidae | <i>Parthenope serrata</i> | 1 | 1 | |
| | <i>Percnon gibbesi</i> | 7 | | |
| | <i>Percnon</i> sp. | 1 | 8 | |
| | <i>Persephona punctata</i> | 1 | 1 | |
| Porcellanidae | <i>Petrolisthes galathinus</i> | 7 | | |
| | <i>Petrolisthes ponta</i> | 1 | | |

continued...

Table 8.5.2 continued

| Taxon | Species | Occurrence | Genus | Sum Higher taxon |
|---------------|---------------------------------------|------------|-------|---------------------|
| | <i>Petrolisthes</i> sp. | 4 | 12 | |
| | <i>Pilumnus</i> sp. | 1 | 1 | |
| | <i>Pisosoma</i> sp. | 1 | 1 | |
| Majidae | <i>Pitho aculeata</i> | 1 | | |
| | <i>Pitho lherminieri</i> | 2 | | |
| | <i>Pitho</i> sp. | 3 | 6 | |
| Xanthidae | <i>Platypodia spectabilis</i> | 1 | 1 | |
| Porcellanidae | <i>Porcellana sayana</i> | 1 | 1 | |
| Portunidae | <i>Portunus anaps</i> | 1 | | |
| | <i>Portunus bahamensis</i> | 1 | | |
| | <i>Portunus ordwayi</i> | 2 | | |
| | <i>Portunus sebae</i> | 5 | | |
| | <i>Portunus</i> sp. | 6 | 15 | |
| Raninidae | <i>Ranilia muricata</i> | 1 | 1 | |
| Grapsidae | <i>Sesarma</i> sp. | 1 | 1 | |
| Majidae | <i>Stenorynchus seticornis</i> | 4 | 4 | |
| | <i>Xanthodius denticulatus</i> | 1 | 1 | |
| Hemichordates | - | 6 | - | 6 |
| Asteroids | <i>Astropecten</i> sp. | 1 | 1 | 6 |
| | <i>Linckia guildingii</i> | 1 | 1 | |
| | <i>Ophidiaster</i> sp. | 1 | 1 | |
| | <i>Oreaster reticulatus</i> | 3 | 3 | |
| Ophiuroids | - | 6 | - | 55 |
| | amphiurid | 1 | - | |
| | <i>Amphipholis gracillima</i> | 1 | 1 | |
| | <i>Amphiura fibulata</i> | 1 | 1 | |
| | <i>Ophiactis</i> sp. | 1 | 1 | |
| | <i>Ophiocoma echinata</i> | 7 | | |
| | <i>Ophiocoma riisei</i> | 3 | | |
| | <i>Ophiocoma</i> sp. | 3 | 13 | |
| | <i>Ophioderma brevispinum</i> | 1 | | |
| | <i>Ophioderma rubicundum</i> | 3 | | |
| | <i>Ophioderma</i> sp. | 2 | | |
| | <i>Ophioderma squamosissima</i> | 1 | 7 | |
| | <i>Ophionereis</i> sp. | 3 | 3 | |
| | <i>Ophiothrix lineata</i> | 1 | | |
| | <i>Ophiothrix</i> sp. | 21 | 22 | |
| Echinoids | - | 5 | - | 68 |
| | spatangoids | 8 | - | |
| | <i>Brissus brissus</i> (heart urchin) | 1 | 1 | |
| | <i>Clypeaster</i> sp. | 1 | 1 | |
| | <i>Diadema antillarum</i> | 22 | 22 | |
| | <i>Echinometra lacunter</i> | 3 | | |
| | <i>Echinometra</i> sp. | 7 | | |
| | <i>Echinometra viridis</i> | 6 | 16 | |
| | <i>Eucidaris tribuloides</i> | 6 | 6 | |
| | <i>Lytechinus variegatus</i> | 7 | 7 | |
| | <i>Moira atropus</i> | 1 | 1 | |
| | <i>Tripneustes esculentus</i> | 1 | 1 | |
| Holothurians | - | 8 | - | 10 |
| | <i>Holothuria arenicola</i> | 1 | 1 | |
| | <i>Thyone pseudofusus</i> | 1 | 1 | |
| Tunicates | - | 9 | - | 36 |
| | appendicularians | 8 | - | |
| | didemnids | 4 | - | |
| | salps | 4 | - | |
| | <i>Ascidia nigra</i> | 2 | 2 | |

continued...

Table 8.5.2 continued

| Taxon | Species | Occurrence | Sum | |
|-------------|--------------------------------|------------|-------|--------------|
| | | | Genus | Higher taxon |
| | <i>Boltenia</i> sp. | 1 | 1 | |
| | <i>Ecteinascidia turbinata</i> | 1 | 1 | |
| | <i>Polycarpa insulsa</i> | 1 | 1 | |
| | <i>Microcosmos exasperatus</i> | 1 | 1 | |
| | <i>Trididemnum savignii</i> | 4 | | |
| | <i>Trididemnum</i> sp. | 1 | 5 | |
| Sea turtles | <i>Caretta caretta</i> | 1 | 1 | 2 |
| | <i>Eretmochelys imbricata</i> | 1 | 1 | |

Table 8.5.3. Composition of "zooplankton" in the diet composition matrix of a Caribbean coral reef (US Virgin Islands, based on Randall 1967); "occurrence" refers to the number of fish species with respective item in stomach.

| Taxon | Item | Occurrence |
|--------------|-------------------------|------------|
| Arthropoda | Insects | 3 |
| Annelida | Polychaete larvae | 1 |
| Cnidaria | Scyphozoans (medusae) | 2 |
| | Siphonophores (medusae) | 11 |
| Crustacea | Barnacle larvae | 5 |
| | Copepods | 47 |
| | Crab larvae | 26 |
| | Euphausiids | 1 |
| | Hermit crab larvae | 4 |
| | Isopod larvae | 1 |
| | Mysids | 12 |
| | Ostracods | 19 |
| | Scyllarid larvae | 5 |
| | Shrimp larvae | 37 |
| | Stomatopod larvae | 11 |
| Ctenophora | Ctenophores (medusae) | 1 |
| Mollusca | Bivalve larvae | 2 |
| | Cephalopod larvae | 4 |
| | Gastropod larvae | 11 |
| | Heteropods | 2 |
| | Mollusc eggs | 6 |
| | Pteropods | 13 |
| Tunicata | Appendicularians | 11 |
| | Salps | 5 |
| Unidentified | Fish eggs | 14 |
| | Fish larvae | 11 |

Appendix 8.6. Diet composition matrix.

Table 8.6.1. Taxa and species included in the diet composition matrix of the ECOPATH II models of a Caribbean coral reef. Species are listed in alphabetical order of families.

| Family or higher taxon | Code | Species |
|------------------------|------|-------------------------------------|
| Organic detritus | D 1 | |
| Benthic algae | A 1 | |
| Spermatophytes | A 1 | |
| Symbiotic algae | A 2 | |
| Phytoplankton | A 3 | |
| Decomposers/microfauna | I 1 | |
| Zooplankton | I 2 | |
| Sponges | I 3 | |
| Hydrozoans | I 4 | |
| Sea fans | I 5 | |
| Sea anemones | I 6 | |
| Stony corals | I 7 | |
| Bryozoans | I 8 | |
| Sipunculid worms | I 9 | |
| Priapuloids | I 10 | |
| Chitons | I 11 | |
| Gastropods | I 12 | |
| Bivalves | I 13 | |
| Scaphopods | I 14 | |
| Squids | I 15 | |
| Octopuses | I 16 | |
| Polychaetes | I 17 | |
| Echiuroids | I 18 | |
| Pycnogonids | I 19 | |
| Barnacles | I 20 | |
| Stomatopods | I 21 | |
| Amphipods | I 22 | |
| Tanaids | I 23 | |
| Isopods | I 24 | |
| Shrimps | I 25 | |
| Spiny lobsters | I 26 | |
| Scyllarid lobsters | I 27 | |
| Hermit crabs | I 28 | |
| Crabs | I 29 | |
| Hemichordates | I 30 | |
| Asteroids | I 31 | |
| Ophiuroids | I 32 | |
| Echinoids | I 33 | |
| Holothurians | I 34 | |
| Tunicates | I 35 | |
| Sea turtles | R 1 | |
| Sea birds | B 1 | |
| | F 0 | Unidentified fish |
| Acanthuridae | F 1 | <i>Acanthurus bahianus</i> |
| | F 2 | <i>Acanthurus chirurgus</i> |
| | F 3 | <i>Acanthurus coeruleus</i> |
| Antennariidae | F 4 | <i>Antennarius multiocellatus</i> |
| | F 5 | <i>Antennarius striatus</i> |
| Apogonidae | F 6 | <i>Apogon conklini</i> |
| | F 7 | <i>Apogon maculatus</i> |
| Atherinidae | F 8 | <i>Allanetta harringtonensis</i> |
| | F 9 | <i>Atherinomorus stipes</i> |
| Aulostomidae | F 10 | <i>Aulostomus maculatus</i> |
| Balistidae | F 11 | <i>Balistes vetula</i> |
| | F 12 | <i>Canthidermis sufflamen</i> |
| | F 13 | <i>Melichthys niger</i> |
| Belonidae | F 14 | <i>Platybelone argalus argalus</i> |
| | F 15 | <i>Strongylura timucu</i> |
| | F 16 | <i>Tylosurus acus acus</i> |
| | F 17 | <i>Tylosurus crocodilus crocod.</i> |
| Blenniidae | F 18 | <i>Entomacrodus nigricans</i> |
| | F 19 | <i>Ophioblennius atlanticus</i> |
| | F 20 | <i>Parablennius marmoratus</i> |
| | F 21 | <i>Scartella cristata</i> |
| Bothidae | F 22 | <i>Bothus lunatus</i> |
| | F 23 | <i>Bothus ocellatus</i> |

continued...

Table 8.6.1 continued

| Family or higher taxon | Code | Species | |
|------------------------|----------------|------------------------------------|---------------------------------|
| Carangidae | F 24 | <i>Caranx bartholomaei</i> | |
| | F 25 | <i>Caranx latus</i> | |
| | F 26 | <i>Caranx ruber</i> | |
| | F 27 | <i>Decapterus punctatus</i> | |
| | F 28 | <i>Oligoplites saurus</i> | |
| | F 29 | <i>Selar crumenophthalmus</i> | |
| | F 30 | <i>Seriola dumerili</i> | |
| | F 31 | <i>Trachinotus falcatus</i> | |
| | F 32 | <i>Trachinotus goodei</i> | |
| | Carcharhinidae | F 33 | <i>Carcharhinus acronotus</i> |
| F 34 | | <i>Carcharhinus falciformis</i> | |
| F 35 | | <i>Carcharhinus leucas</i> | |
| F 36 | | <i>Carcharhinus limbatus</i> | |
| F 37 | | <i>Carcharhinus longimanus</i> | |
| F 38 | | <i>Carcharhinus perezii</i> | |
| F 39 | | <i>Galeocerdo cuvier</i> | |
| F 40 | | <i>Negaprion brevirostris</i> | |
| Chaetodontidae | F 41 | <i>Rhizoprionodon porosus</i> | |
| | F 42 | <i>Chaetodon aculeatus</i> | |
| | F 43 | <i>Chaetodon capistratus</i> | |
| | F 44 | <i>Chaetodon sedentarius</i> | |
| | F 45 | <i>Chaetodon striatus</i> | |
| Cirrihidae | F 46 | <i>Amblycirrhites pinos</i> | |
| Clinidae | F 47 | <i>Labrisomus guppyi</i> | |
| | F 48 | <i>Labrisomus nuchipinnis</i> | |
| Clupeidae | F 49 | <i>Harengula clupeola</i> | |
| | F 50 | <i>Harengula humeralis</i> | |
| | F 51 | <i>Jenkinsia lamprotaenia</i> | |
| | F 52 | <i>Opisthonema oglinum</i> | |
| Congridae | F 53 | <i>Heteroconger halis</i> | |
| Dactylopteridae | F 54 | <i>Dactylopterus volitans</i> | |
| Dasyatidae | F 55 | <i>Dasyatis americana</i> | |
| Diodontidae | F 56 | <i>Chilomycterus antennatus</i> | |
| | F 57 | <i>Diodon holocanthus</i> | |
| | F 58 | <i>Diodon hystrix</i> | |
| Emmelichthyidae | F 59 | <i>Inermia vittata</i> | |
| Engraulidae | F 60 | <i>Anchoa hepsetus</i> | |
| | F 61 | <i>Anchoa lyolepis</i> | |
| Ephippidae | F 62 | <i>Chaetodipterus faber</i> | |
| Fistulariidae | F 63 | <i>Fistularia tabacaria</i> | |
| Gerreidae | F 64 | <i>Eucinostomus argenteus</i> | |
| | F 65 | <i>Gerres cinereus</i> | |
| Ginglymostomatidae | F 66 | <i>Ginglymostoma cirratum</i> | |
| Gobiidae | F 67 | <i>Coryphopterus glaucofraenum</i> | |
| | F 68 | <i>Gnatholepis thompsoni</i> | |
| | F 69 | <i>Gobiosoma evelynae</i> | |
| | F 70 | <i>Gramma loreto</i> | |
| Grammidae | F 71 | <i>Gramma melacara</i> | |
| | F 72 | <i>Rypticus saponaceus</i> | |
| Grammistidae | F 73 | <i>Anisotremus surinamensis</i> | |
| | F 74 | <i>Anisotremus virginicus</i> | |
| | F 75 | <i>Haemulon album</i> | |
| | F 76 | <i>Haemulon aurolineatum</i> | |
| | F 77 | <i>Haemulon carbonarium</i> | |
| | F 78 | <i>Haemulon chrysargyreum</i> | |
| | F 79 | <i>Haemulon flavolineatum</i> | |
| | F 80 | <i>Haemulon macrostomum</i> | |
| | F 81 | <i>Haemulon parrai</i> | |
| | F 82 | <i>Haemulon plumieri</i> | |
| | F 83 | <i>Haemulon sciurus</i> | |
| | Hemiramphidae | F 84 | <i>Hemiramphus balao</i> |
| | | F 85 | <i>Hemiramphus brasiliensis</i> |
| Holocentridae | F 86 | <i>Holocentrus ascensionis</i> | |
| | F 87 | <i>Holocentrus coruscus</i> | |
| | F 88 | <i>Neoniphon marianus</i> | |
| | F 89 | <i>Holocentrus rufus</i> | |
| | F 90 | <i>Sargocentron vexillarium</i> | |
| | F 91 | <i>Myripristis jacobus</i> | |
| Kyphosidae | F 92 | <i>Kyphosus incisor</i> | |
| | F 93 | <i>Kyphosus sectatrix</i> | |
| Labridae | F 94 | <i>Bodianus rufus</i> | |
| | F 95 | <i>Clepticus parrae</i> | |

continued...

Table 8.6.1 continued

| Family or higher taxon | Code | Species |
|------------------------|-------|-------------------------------------|
| | F 96 | <i>Halichoeres bivittatus</i> |
| | F 97 | <i>Halichoeres gamoti</i> |
| | F 98 | <i>Halichoeres maculipinna</i> |
| | F 99 | <i>Halichoeres poeyi</i> |
| | F 100 | <i>Halichoeres radiatus</i> |
| | F 101 | <i>Xyrichtys novacula</i> |
| | F 102 | <i>Xyrichtys splendens</i> |
| | F 103 | <i>Lachnolaimus maximus</i> |
| | F 104 | <i>Thalassoma bifasciatum</i> |
| Lutjanidae | F 105 | <i>Lutjanus analis</i> |
| | F 106 | <i>Lutjanus apodus</i> |
| | F 107 | <i>Lutjanus cyanopterus</i> |
| | F 108 | <i>Lutjanus griseus</i> |
| | F 109 | <i>Lutjanus jocu</i> |
| | F 110 | <i>Lutjanus mahagoni</i> |
| | F 111 | <i>Lutjanus synagris</i> |
| | F 112 | <i>Ocyurus chrysurus</i> |
| Malacanthidae | F 113 | <i>Malacanthus plumieri</i> |
| Megalopidae | F 114 | <i>Tarpon atlanticus</i> |
| Monacanthidae | F 115 | <i>Aluterus schoepfii</i> |
| | F 116 | <i>Aluterus scripta</i> |
| | F 117 | <i>Cantherines macroceros</i> |
| | F 118 | <i>Cantherines pullus</i> |
| | F 119 | <i>Monacanthus ciliatus</i> |
| Mugilidae | F 120 | <i>Mugil curema</i> |
| Mullidae | F 121 | <i>Mulloidichthys martinicus</i> |
| | F 122 | <i>Pseudupeneus maculatus</i> |
| Muraenidae | F 123 | <i>Echidna catenata</i> |
| | F 124 | <i>Lycodontis moina</i> |
| | F 125 | <i>Gymnothorax vicinus</i> |
| Myliobatidae | F 126 | <i>Aetobatus narinari</i> |
| Ogcocephalidae | F 127 | <i>Ogcocephalus nasutus</i> |
| Ophichthidae | F 128 | <i>Myrichthys breviceps</i> |
| | F 129 | <i>Myrichthys ocellatus</i> |
| | F 130 | <i>Ophichthys ophis</i> |
| Opisthognathidae | F 131 | <i>Opisthognathus aurifrons</i> |
| | F 132 | <i>Opisthognathus maxillosus</i> |
| | F 133 | <i>Opisthognathus whitehurstii</i> |
| Ostraciidae | F 134 | <i>Acanthostracion polygonius</i> |
| | F 135 | <i>Acanthostracion quadricomis</i> |
| | F 136 | <i>Lactophrys trigonus</i> |
| | F 137 | <i>Lactophrys bicaudalis</i> |
| | F 138 | <i>Lactophrys triqueter</i> |
| Pempheridae | F 139 | <i>Pempheris schomburgki</i> |
| Pomacanthidae | F 140 | <i>Centropyge argi</i> |
| | F 141 | <i>Holacanthus ciliaris</i> |
| | F 142 | <i>Holacanthus tricolor</i> |
| | F 143 | <i>Pomacanthus arcuatus</i> |
| | F 144 | <i>Pomacanthus paru</i> |
| Pomacentridae | F 145 | <i>Abudefduf saxatilis</i> |
| | F 146 | <i>Abudefduf taurus</i> |
| | F 147 | <i>Chromis cyanea</i> |
| | F 148 | <i>Chromis multilineata</i> |
| | F 149 | <i>Microspathodon chrysurus</i> |
| | F 150 | <i>Stegastes fuscus</i> |
| | F 151 | <i>Stegastes leucostictus</i> |
| | F 152 | <i>Stegastes planifrons</i> |
| | F 153 | <i>Stegastes variabilis</i> |
| Priacanthidae | F 154 | <i>Priacanthus arenatus</i> |
| | F 155 | <i>Heteropriacanthus cruentatus</i> |
| Scaridae | F 156 | <i>Scarus coelestinus</i> |
| | F 157 | <i>Scarus iserti</i> |
| | F 158 | <i>Scarus guacamaia</i> |
| | F 159 | <i>Scarus taeniopterus</i> |
| | F 160 | <i>Scarus vetula</i> |
| | F 161 | <i>Sparisoma aurofrenatum</i> |
| | F 162 | <i>Sparisoma chrysopterygum</i> |
| | F 163 | <i>Sparisoma radians</i> |
| | F 164 | <i>Sparisoma rubripinne</i> |
| | F 165 | <i>Sparisoma viride</i> |
| Sciaenidae | F 166 | <i>Equetus lanceolatus</i> |
| | F 167 | <i>Equetus punctatus</i> |

continued...

Table 8.6.1 continued

| Family or higher taxon | Code | Species |
|------------------------|-------|---------------------------------------|
| | F 168 | <i>Odontoscion dentex</i> |
| | F 169 | <i>Pareques acuminatus</i> |
| Scombridae | F 170 | <i>Euthynnus alletteratus</i> |
| | F 171 | <i>Scomberomorus cavalla</i> |
| | F 172 | <i>Scomberomorus regalis</i> |
| Scorpaenidae | F 173 | <i>Scorpaena brasiliensis</i> |
| | F 174 | <i>Scorpaena grandicornis</i> |
| | F 175 | <i>Scorpaena inermis</i> |
| | F 176 | <i>Scorpaena plumieri</i> |
| | F 177 | <i>Scorpaenodes caribbaeus</i> |
| Serranidae | F 178 | <i>Alphestes afer</i> |
| | F 179 | <i>Cephalopholis cruentata</i> |
| | F 180 | <i>Cephalopholis fulva</i> |
| | F 181 | <i>Epinephelus adscensionis</i> |
| | F 182 | <i>Epinephelus guttatus</i> |
| | F 183 | <i>Epinephelus itajara</i> |
| | F 184 | <i>Epinephelus striatus</i> |
| | F 185 | <i>Hypoplectrus aberrans</i> |
| | F 186 | <i>Hypoplectrus chlorurus</i> |
| | F 187 | <i>Hypoplectrus nigricans</i> |
| | F 188 | <i>Hypoplectrus puella</i> |
| | F 189 | <i>Mycteroperca bonaci</i> |
| | F 190 | <i>Mycteroperca tigris</i> |
| | F 191 | <i>Mycteroperca venenosa</i> |
| | F 192 | <i>Paranthias furcifer</i> |
| | F 193 | <i>Serranus tigrinus</i> |
| Sparidae | F 194 | <i>Archosargus rhomboidalis</i> |
| | F 195 | <i>Calamus bajonado</i> |
| | F 196 | <i>Calamus calamus</i> |
| | F 197 | <i>Calamus pennatula</i> |
| | F 198 | <i>Diplodus argenteus caudimacula</i> |
| Sphyracidae | F 199 | <i>Sphyracna barracuda</i> |
| | F 200 | <i>Sphyracna picudilla</i> |
| Sphyrnidae | F 201 | <i>Sphyrna lewini</i> |
| | F 202 | <i>Sphyrna tiburo</i> |
| Synodontidae | F 203 | <i>Synodus foetens</i> |
| | F 204 | <i>Synodus intermedius</i> |
| | F 205 | <i>Synodus synodus</i> |
| Tetraodontidae | F 206 | <i>Canthigaster rostrata</i> |
| | F 207 | <i>Sphoeroides spengleri</i> |
| Triakidae | F 208 | <i>Mustelus canis</i> |

Symbols:

| | | | | | |
|---|---|-----------|---|---|--------------|
| A | = | autotroph | F | = | fish |
| B | = | bird | I | = | invertebrate |
| D | = | detritus | R | = | reptile |

Table 8.6.2. The complete diet composition matrix for the construction of steady-state models of a Caribbean coral reef area, including 41 nonfish taxa and 208 fish species.

| Predator/prey | D1 | A1 | A2 | A3 | I1 | I2 | I3 | I4 | I5 | I6 | I7 |
|-------------------------|------|------|------|------|------|-------|-------|-------|-------|-------|-------|
| D 1 Detritus, POM, DOM | - | - | - | - | - | - | - | - | - | - | - |
| A 1 Benthic autotrophs | - | - | - | - | - | - | - | - | - | - | - |
| A 2 Symbiotic algae | - | - | - | - | - | - | - | - | - | - | - |
| A 3 Phytoplankton | - | - | - | - | - | - | - | - | - | - | - |
| I 1 Zooplankton | - | - | - | 0.2 | - | 0.8 | - | - | - | - | - |
| I 2 Microfauna | 1 | - | - | - | - | - | - | - | - | - | - |
| I 3 Sponges | 1 | - | - | - | - | - | - | - | - | - | - |
| I 4 Fire corals | 0.1 | - | 0.65 | - | 0.15 | 0.1 | - | - | - | - | - |
| I 5 Sea fans | 0.1 | - | 0.65 | - | 0.15 | 0.1 | - | - | - | - | - |
| I 6 Sea anemones | 0.2 | - | - | - | 0.7 | 0.1 | - | - | - | - | - |
| I 7 Stony corals | 0.1 | - | 0.65 | - | 0.15 | 0.1 | - | - | - | - | - |
| I 8 Bryozoans | 0.1 | - | - | 0.7 | 0.09 | 0.1 | - | - | - | - | - |
| I 9 Sipunculid worms | 0.5 | 0.25 | - | - | - | 0.25 | - | - | - | - | - |
| I 10 Priapuloids | - | - | - | - | - | - | - | - | - | - | - |
| I 11 Chitons | - | 0.8 | - | - | - | 0.04 | 0.04 | 0.04 | - | - | - |
| I 12 Gastropods | 0.2 | 0.35 | - | - | - | 0.064 | 0.04 | 0.006 | 0.012 | 0.006 | 0.008 |
| I 13 Bivalves | 0.72 | - | - | 0.16 | - | 0.12 | - | - | - | - | - |
| I 14 Scaphopods | - | - | - | - | - | 1 | - | - | - | - | - |
| I 15 Squids | - | - | - | - | 0.3 | - | - | - | - | - | - |
| I 16 Octopuses | - | - | - | - | - | - | - | - | - | - | - |
| I 17 Polychaetes | 0.3 | 0.15 | - | 0.14 | 0.1 | 0.1 | 0.03 | 0.009 | 0.018 | 0.009 | 0.009 |
| I 18 Echiuroids | 0.45 | 0.5 | - | - | - | 0.05 | - | - | - | - | - |
| I 19 Pycnogonids | - | 0.1 | - | - | 0.1 | - | 0.15 | 0.25 | 0.2 | 0.1 | - |
| I 20 Barnacles | - | - | - | 0.4 | 0.4 | 0.2 | - | - | - | - | - |
| I 21 Stomatopods | - | - | - | - | 0.05 | - | - | - | - | - | - |
| I 22 Amphipods | 0.2 | 0.5 | - | - | 0.2 | 0.1 | - | - | - | - | - |
| I 23 Tanaids | - | 0.8 | - | - | - | 0.2 | - | - | - | - | - |
| I 24 Isopods | 0.2 | - | - | - | - | - | - | - | - | - | - |
| I 25 Shrimps | 0.04 | 0.01 | - | - | 0.2 | 0.08 | - | - | - | - | - |
| I 26 Spiny lobsters | - | 0.3 | - | - | - | - | - | - | - | - | - |
| I 27 Scyllarid lobsters | 0.2 | - | - | - | - | - | - | - | - | - | - |
| I 28 Hermit crabs | 0.05 | - | - | 0.05 | 0.05 | 0.05 | - | - | - | - | - |
| I 29 Crabs | 0.03 | 0.55 | - | 0.03 | 0.04 | 0.03 | 0.02 | - | - | - | 0.04 |
| I 30 Hemichordates | 0.1 | - | - | 0.8 | - | 0.1 | - | - | - | - | - |
| I 31 Asteroids | 0.35 | 0.05 | - | - | - | 0.04 | 0.005 | - | 0.01 | 0.005 | 0.005 |
| I 32 Ophiuroids | 0.3 | - | - | 0.3 | 0.3 | 0.1 | - | - | - | - | - |
| I 33 Echinoids | - | 0.7 | - | - | - | - | 0.05 | - | - | - | 0.2 |
| I 34 Holothurians | 0.7 | 0.1 | - | - | - | 0.2 | - | - | - | - | - |
| I 35 Tunicates | 0.4 | - | - | 0.2 | 0.2 | 0.2 | - | - | - | - | - |
| R 1 Sea turtles | - | 0.39 | - | - | - | - | 0.34 | - | - | - | 0.04 |
| B 1 Sea birds | - | - | - | - | 0.05 | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I8 | I9 | I10 | I11 | I12 | I13 | I14 | I15 | I16 | I17 | I18 | I19 | I20 | I21 |
|------|------|-------|-----|------|-------|------|------|------|------|-------|-----|-----|-------|-------|
| D 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| A 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| A 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| A 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 8 | - | - | - | - | - | - | - | - | - | 0.01 | - | - | - | - |
| I 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 10 | - | - | - | - | - | - | - | - | - | 1 | - | - | - | - |
| I 11 | 0.04 | - | - | - | - | - | - | - | - | - | - | - | 0.04 | - |
| I 12 | - | - | - | 0.03 | 0.04 | 0.03 | 0.03 | - | - | 0.1 | - | - | 0.004 | 0.002 |
| I 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 15 | - | - | - | - | - | - | - | 0.3 | - | - | - | - | - | - |
| I 16 | - | - | - | 0.01 | 0.83 | 0.06 | - | - | 0.01 | 0.005 | - | - | - | - |
| I 17 | 0.01 | - | - | - | 0.01 | 0.01 | - | - | - | 0.045 | - | - | - | - |
| I 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 19 | 0.1 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 21 | - | - | - | - | 0.2 | 0.2 | - | 0.05 | - | 0.03 | - | - | - | - |
| I 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 25 | - | - | - | - | 0.05 | 0.04 | 0.03 | - | 0.03 | 0.1 | - | - | - | 0.03 |
| I 26 | - | - | - | - | 0.3 | 0.4 | - | - | - | - | - | - | - | - |
| I 27 | - | - | - | - | - | 0.6 | - | - | - | 0.2 | - | - | - | - |
| I 28 | - | - | - | - | - | 0.8 | - | - | - | - | - | - | - | - |
| I 29 | - | 0.001 | - | - | 0.08 | 0.03 | - | - | - | 0.002 | - | - | 0.003 | - |
| I 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 31 | - | - | - | - | 0.2 | 0.09 | 0.01 | - | - | 0.01 | - | - | - | - |
| I 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 33 | - | - | - | - | - | 0.01 | - | - | - | - | - | - | - | - |
| I 34 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| R 1 | - | - | - | - | 0.025 | 0.02 | - | - | - | - | - | - | 0.02 | 0.02 |
| B 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I22 | I23 | I24 | I25 | I26 | I27 | I28 | I29 | I30 | I31 | I32 | I33 | I34 | I35 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|------|-------|-------|------|-------|
| D 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| A 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| A 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| A 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 12 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.018 | - | - | - | - | 0.004 |
| I 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 15 | - | - | - | 0.1 | - | - | - | - | - | - | - | - | - | - |
| I 16 | - | - | - | 0.01 | - | - | 0.02 | 0.035 | - | - | - | - | - | - |
| I 17 | 0.045 | - | - | - | - | - | - | - | - | - | 0.01 | - | - | 0.005 |
| I 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 21 | 0.12 | 0.05 | 0.05 | 0.09 | - | - | - | - | - | 0.02 | 0.02 | 0.02 | 0.02 | - |
| I 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 25 | 0.11 | 0.09 | 0.05 | 0.02 | 0.01 | 0.01 | 0.01 | 0.01 | - | - | 0.06 | - | - | - |
| I 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 29 | 0.01 | - | - | - | - | - | 0.07 | 0.04 | - | - | 0.002 | 0.002 | - | - |
| I 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 31 | - | - | - | - | - | - | - | 0.02 | - | 0.12 | 0.05 | 0.02 | 0.01 | 0.005 |
| I 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 33 | - | - | - | - | - | - | - | - | - | 0.01 | 0.01 | 0.01 | 0.01 | - |
| I 34 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| I 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| R 1 | - | - | - | - | 0.06 | - | 0.02 | 0.025 | - | 0.02 | - | 0.02 | - | - |
| B 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | R1 | B1 | FO | Sum | Qualitative indications on fish |
|------|----|----|------|-----|---------------------------------------|
| D 1 | - | - | - | 0 | |
| A 1 | - | - | - | 0 | |
| A 2 | - | - | - | 0 | |
| A 3 | - | - | - | 0 | |
| I 1 | - | - | - | 1 | |
| I 2 | - | - | - | 1 | |
| I 3 | - | - | - | 1 | |
| I 4 | - | - | - | 1 | |
| I 5 | - | - | - | 1 | |
| I 6 | - | - | - | 1 | |
| I 7 | - | - | - | 1 | |
| I 8 | - | - | - | 1 | |
| I 9 | - | - | - | 1 | |
| I 10 | - | - | - | 1 | |
| I 11 | - | - | - | 1 | |
| I 12 | - | - | - | 1 | |
| I 13 | - | - | - | 1 | |
| I 14 | - | - | - | 1 | |
| I 15 | - | - | 0.3 | 1 | small pelagics |
| I 16 | - | - | 0.02 | 1 | Labridae, Pomacentridae, Myliobatidae |
| I 17 | - | - | - | 1 | |
| I 18 | - | - | - | 1 | |
| I 19 | - | - | - | 1 | |
| I 20 | - | - | - | 1 | |
| I 21 | - | - | 0.08 | 1 | Engraulidae, Mullidae |
| I 22 | - | - | - | 1 | |
| I 23 | - | - | - | 1 | |
| I 24 | - | - | 0.8 | 1 | Fish (parasite) |
| I 25 | - | - | 0.02 | 1 | |
| I 26 | - | - | - | 1 | |
| I 27 | - | - | - | 1 | |
| I 28 | - | - | - | 1 | |
| I 29 | - | - | 0.02 | 1 | |
| I 30 | - | - | - | 1 | |
| I 31 | - | - | - | 1 | |
| I 32 | - | - | - | 1 | |
| I 33 | - | - | - | 1 | |
| I 34 | - | - | - | 1 | |
| I 35 | - | - | - | 1 | |
| R 1 | - | - | - | 1 | |
| B 1 | - | - | 0.95 | 1 | |

continued...

Table 8.6.2 continued

| | Predator/prey | D1 | A1 | A2 | A3 | I1 | I2 | I3 | I4 | I5 |
|------|------------------------------------|-------|-------|-------|-----|-------|-------|----|-------|-------|
| F 1 | <i>Acanthurus bahianus</i> | 0.459 | 0.541 | - | - | - | - | - | - | - |
| F 2 | <i>Acanthurus chirurgus</i> | 0.469 | 0.527 | - | - | - | - | - | - | - |
| F 3 | <i>Acanthurus coeruleus</i> | 0.464 | 0.532 | 0.001 | - | - | 0.003 | - | - | - |
| F 4 | <i>Antennarius multiocellatus</i> | - | - | - | - | - | - | - | - | - |
| F 5 | <i>Antennarius striatus</i> | - | - | - | - | - | - | - | - | - |
| F 6 | <i>Apogon conklini</i> | - | - | - | - | - | 0.6 | - | - | - |
| F 7 | <i>Apogon maculatus</i> | - | - | - | - | - | 0.387 | - | - | - |
| F 8 | <i>Allanetta harringtonensis</i> | - | - | - | - | - | 1 | - | - | - |
| F 9 | <i>Atherinomorus stipes</i> | - | - | - | - | 0.028 | 0.91 | - | - | - |
| F 10 | <i>Aulostomus maculatus</i> | - | - | - | - | - | - | - | - | - |
| F 11 | <i>Balistes vetula</i> | - | 0.012 | 0.001 | - | - | - | - | - | - |
| F 12 | <i>Canthidermis sufflamen</i> | - | 0.025 | - | - | - | 0.609 | - | - | - |
| F 13 | <i>Melichthys niger</i> | - | 0.752 | 0.004 | - | 0.001 | 0.153 | - | - | - |
| F 14 | <i>Platybelone argalus argalus</i> | - | - | - | - | - | 0.03 | - | - | - |
| F 15 | <i>Strongylura timucu</i> | - | - | - | - | - | - | - | - | - |
| F 16 | <i>Tylosurus acus</i> | - | - | - | - | - | 0.037 | - | - | - |
| F 17 | <i>Tylosurus crocodilus</i> | - | - | - | - | - | - | - | - | - |
| F 18 | <i>Entomacrodus nigricans</i> | 0.482 | 0.482 | - | - | - | - | - | - | - |
| F 19 | <i>Ophioblennius atlanticus</i> | 0.498 | 0.497 | - | - | - | 0.005 | - | - | - |
| F 20 | <i>Parablennius marmoreus</i> | 0.396 | 0.396 | 0.014 | - | - | - | - | 0.01 | - |
| F 21 | <i>Scartella cristata</i> | 0.496 | 0.496 | - | - | - | - | - | - | - |
| F 22 | <i>Bothus lunatus</i> | - | - | - | - | - | - | - | - | - |
| F 23 | <i>Bothus ocellatus</i> | - | - | - | - | - | - | - | - | - |
| F 24 | <i>Caranx bartholomaei</i> | - | - | - | - | - | - | - | - | - |
| F 25 | <i>Caranx latus</i> | - | - | - | - | - | 0.084 | - | - | - |
| F 26 | <i>Caranx ruber</i> | - | - | - | - | - | 0.022 | - | - | - |
| F 27 | <i>Decapterus punctatus</i> | - | - | - | - | - | 1 | - | - | - |
| F 28 | <i>Oligoplites saurus</i> | - | - | - | - | - | 0.013 | - | - | - |
| F 29 | <i>Selar crumenophthalmus</i> | - | - | - | - | - | 0.904 | - | - | - |
| F 30 | <i>Seriola dumerilii</i> | - | - | - | - | - | - | - | - | - |
| F 31 | <i>Trachinotus falcatulus</i> | - | - | - | - | - | - | - | - | - |
| F 32 | <i>Trachinotus goodei</i> | - | - | - | - | - | - | - | - | - |
| F 33 | <i>Carcharhinus acronotus</i> | - | - | - | - | - | - | - | - | - |
| F 34 | <i>Carcharhinus falciformis</i> | - | - | - | - | - | - | - | - | - |
| F 35 | <i>Carcharhinus leucas</i> | 0.1 | - | - | - | - | - | - | - | - |
| F 36 | <i>Carcharhinus limbatus</i> | - | - | - | - | - | - | - | - | - |
| F 37 | <i>Carcharhinus longimanus</i> | 0.1 | - | - | - | - | - | - | - | - |
| F 38 | <i>Carcharhinus perezii</i> | - | - | - | - | - | - | - | - | - |
| F 39 | <i>Galeocerdo cuvier</i> | 0.05 | - | - | - | - | - | - | - | - |
| F 40 | <i>Negaprion brevirostris</i> | 0.1 | - | - | - | - | - | - | - | - |
| F 41 | <i>Rhizoprionodon porosus</i> | - | - | - | - | - | - | - | - | - |
| F 42 | <i>Chaetodon aculeatus</i> | - | - | - | - | - | 0.173 | - | - | - |
| F 43 | <i>Chaetodon capistratus</i> | - | - | 0.039 | - | - | 0.064 | - | - | 0.026 |
| F 44 | <i>Chaetodon sedentarius</i> | - | - | 0.124 | - | - | - | - | 0.082 | - |
| F 45 | <i>Chaetodon striatus</i> | - | - | - | - | - | 0.025 | - | - | - |
| F 46 | <i>Amblycirrhitus pinos</i> | - | - | - | - | - | 0.64 | - | - | - |
| F 47 | <i>Labrisomus guppyi</i> | - | - | - | - | - | - | - | - | - |
| F 48 | <i>Labrisomus nuchipinnis</i> | - | - | - | - | - | - | - | - | - |
| F 49 | <i>Harengula clupeiola</i> | - | - | - | - | - | 0.95 | - | - | - |
| F 50 | <i>Harengula humeralis</i> | - | 0.025 | - | - | - | 0.679 | - | - | - |
| F 51 | <i>Jenkinsia lamprotaenia</i> | - | - | - | - | - | 0.982 | - | - | - |
| F 52 | <i>Opisthonema oglinum</i> | - | - | - | - | - | 0.592 | - | - | - |
| F 53 | <i>Heteroconger halis</i> | - | - | - | - | - | 1 | - | - | - |
| F 54 | <i>Dactylopterus volitans</i> | - | - | - | - | - | - | - | - | - |
| F 55 | <i>Dasyatis americana</i> | - | - | - | - | - | - | - | - | - |
| F 56 | <i>Chilomycterus antennatus</i> | - | - | - | - | - | - | - | - | - |
| F 57 | <i>Diodon holocanthus</i> | - | - | - | - | - | - | - | - | - |
| F 58 | <i>Diodon hystrix</i> | - | - | - | - | - | - | - | - | - |
| F 59 | <i>Inermia vittata</i> | - | - | - | - | - | 0.987 | - | - | - |
| F 60 | <i>Anchoa hepsetus</i> | 0.1 | - | - | 0.7 | - | 0.2 | - | - | - |

continued...

Table 8.6.2 continued

| | I6 | I7 | I8 | I9 | I10 | I11 | I12 | I13 | I14 | I15 | I16 | I17 | I18 | I19 |
|------|-------|-------|----|-------|-----|-------|-------|-------|-----|-------|-------|-------|-----|-----|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | 0.002 | - | - | - | - | 0.002 | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | 0.059 | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | 0.037 | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 11 | - | 0.001 | - | 0.009 | - | 0.001 | 0.017 | 0.047 | - | - | - | 0.022 | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | 0.002 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | 0.036 | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | 0.092 | - | - |
| F 21 | - | - | - | - | - | - | 0.008 | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | 0.029 | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | - | - | - | - | - | - | 0.004 | - | - | 0.014 | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | 0.478 | 0.172 | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | 0.083 | 0.039 | - | - | - | - | - | - |
| F 33 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 34 | - | - | - | - | - | - | - | - | - | 0.1 | - | - | - | - |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 36 | - | - | - | - | - | - | - | - | - | 0.1 | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | 0.1 | - | - | - | - |
| F 38 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 39 | - | - | - | - | - | - | 0.05 | - | - | 0.05 | - | - | - | - |
| F 40 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 41 | - | - | - | - | - | - | 0.15 | - | - | - | - | - | - | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | 0.525 | - | - |
| F 43 | 0.536 | - | - | - | - | - | - | - | - | - | - | 0.314 | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | 0.295 | - | - |
| F 45 | 0.325 | - | - | - | - | - | - | - | - | - | - | 0.587 | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | 0.122 | - | - |
| F 47 | - | - | - | - | - | 0.2 | 0.05 | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | 0.165 | - | - | - | - | 0.1 | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | 0.05 | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | 0.296 | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | 0.224 | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | 0.072 | - | - | - | - | - | - |
| F 55 | - | - | - | 0.206 | - | - | - | 0.108 | - | - | - | 0.173 | - | - |
| F 56 | - | - | - | - | - | - | 0.566 | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | 0.677 | 0.128 | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | 0.314 | 0.131 | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | 0.013 | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I20 | I21 | I22 | I23 | I24 | I25 | I26 | I27 | I28 | I29 | I30 | I31 | I32 | I33 |
|------|-------|-------|-------|-------|-------|-------|-----|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | 0.125 | - | - | - | - | - | - | - | 0.125 | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | 0.239 | 0.035 | 0.067 | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | 0.008 | - | 0.013 | 0.282 | - | - | - | 0.273 | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | 0.062 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | 0.265 | - | - | - | - | - | - | - | - |
| F 11 | - | 0.001 | - | - | - | 0.008 | - | 0.001 | 0.02 | 0.057 | - | 0.014 | 0.034 | 0.747 |
| F 12 | - | - | 0.07 | - | - | - | - | - | - | - | - | - | - | 0.296 |
| F 13 | - | - | - | - | - | 0.005 | - | - | 0.017 | 0.035 | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | 0.04 | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | 0.091 | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | 0.092 | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | 0.114 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | 0.033 | 0.177 | - | - | 0.211 | - | - | - | 0.296 | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | 0.007 | 0.038 | - | - | - | - | - | - | - | - |
| F 26 | - | 0.006 | - | - | - | 0.035 | - | - | - | 0.004 | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | 0.061 | 0.039 | - | - | - | 0.25 |
| F 32 | - | - | - | - | 0.017 | - | - | - | 0.009 | - | - | - | - | - |
| F 33 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 34 | - | - | - | - | - | - | - | - | - | 0.1 | - | - | - | - |
| F 35 | - | - | - | - | - | 0.1 | - | - | - | 0.1 | - | - | - | 0.1 |
| F 36 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | 0.1 | - | - | - | - |
| F 38 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 39 | - | - | - | - | - | - | - | - | - | 0.05 | - | - | - | - |
| F 40 | - | - | - | - | - | 0.1 | - | - | - | 0.1 | - | - | - | - |
| F 41 | - | - | - | - | - | 0.15 | - | - | - | - | - | - | - | - |
| F 42 | - | - | - | 0.019 | - | 0.129 | - | - | - | - | - | - | - | 0.154 |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | 0.234 | - | - | 0.265 | - | - | - | - | - | - | - | - |
| F 45 | - | - | 0.021 | 0.021 | - | 0.021 | - | - | - | - | - | - | - | - |
| F 46 | - | - | 0.021 | 0.014 | 0.025 | 0.106 | - | - | - | 0.072 | - | - | - | - |
| F 47 | - | - | - | - | 0.025 | - | - | - | - | 0.725 | - | - | - | - |
| F 48 | - | - | 0.053 | - | - | 0.077 | - | - | - | 0.272 | - | - | 0.123 | 0.105 |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | 0.018 | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | 0.086 | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | 0.193 | 0.014 | - | - | 0.054 | - | - | - | 0.617 | - | - | - | - |
| F 55 | - | 0.02 | - | - | - | 0.076 | - | - | - | 0.176 | 0.023 | - | - | - |
| F 56 | - | - | - | - | 0.206 | 0.004 | - | - | 0.214 | 0.01 | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | 0.042 | 0.037 | - | - | - | 0.116 |
| F 58 | - | - | - | - | - | - | - | - | 0.059 | 0.149 | - | - | - | 0.347 |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I34 | I35 | R1 | B1 |
|------|-----|-------|-----|------|
| F 1 | - | - | - | - |
| F 2 | - | - | - | - |
| F 3 | - | - | - | - |
| F 4 | - | - | - | - |
| F 5 | - | - | - | - |
| F 6 | - | - | - | - |
| F 7 | - | - | - | - |
| F 8 | - | - | - | - |
| F 9 | - | - | - | - |
| F 10 | - | - | - | - |
| F 11 | - | 0.006 | - | - |
| F 12 | - | - | - | - |
| F 13 | - | - | - | - |
| F 14 | - | - | - | - |
| F 15 | - | - | - | - |
| F 16 | - | - | - | - |
| F 17 | - | - | - | - |
| F 18 | - | - | - | - |
| F 19 | - | - | - | - |
| F 20 | - | - | - | - |
| F 21 | - | - | - | - |
| F 22 | - | - | - | - |
| F 23 | - | - | - | - |
| F 24 | - | - | - | - |
| F 25 | - | - | - | - |
| F 26 | - | - | - | - |
| F 27 | - | - | - | - |
| F 28 | - | - | - | - |
| F 29 | - | - | - | - |
| F 30 | - | - | - | - |
| F 31 | - | - | - | - |
| F 32 | - | - | - | - |
| F 33 | - | - | - | - |
| F 34 | - | - | - | - |
| F 35 | - | - | - | - |
| F 36 | - | - | - | - |
| F 37 | - | - | 0.1 | - |
| F 38 | - | - | - | - |
| F 39 | - | - | 0.1 | 0.05 |
| F 40 | - | - | - | - |
| F 41 | - | - | - | - |
| F 42 | - | - | - | - |
| F 43 | - | 0.021 | - | - |
| F 44 | - | - | - | - |
| F 45 | - | - | - | - |
| F 46 | - | - | - | - |
| F 47 | - | - | - | - |
| F 48 | - | - | - | - |
| F 49 | - | - | - | - |
| F 50 | - | - | - | - |
| F 51 | - | - | - | - |
| F 52 | - | - | - | - |
| F 53 | - | - | - | - |
| F 54 | - | - | - | - |
| F 55 | - | - | - | - |
| F 56 | - | - | - | - |
| F 57 | - | - | - | - |
| F 58 | - | - | - | - |
| F 59 | - | - | - | - |
| F 60 | - | - | - | - |

Table 8.6.2 continued

| | Predator/prey | F0 | F1 | F2 | F3 | F4 | F5 | F6 | F7 | F8 |
|------|------------------------------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | <i>Acanthurus bahianus</i> | - | - | - | - | - | - | - | - | - |
| F 2 | <i>Acanthurus chirurgus</i> | - | - | - | - | - | - | - | - | - |
| F 3 | <i>Acanthurus coeruleus</i> | - | - | - | - | - | - | - | - | - |
| F 4 | <i>Antennarius multiocellatus</i> | - | - | - | - | - | - | - | - | - |
| F 5 | <i>Antennarius striatus</i> | 0.5 | - | - | - | - | - | - | - | - |
| F 6 | <i>Apogon conklini</i> | - | - | - | - | - | - | - | - | - |
| F 7 | <i>Apogon maculatus</i> | - | - | - | - | - | - | - | - | - |
| F 8 | <i>Allanetta harringtonensis</i> | - | - | - | - | - | - | - | - | - |
| F 9 | <i>Atherinomorus stipes</i> | - | - | - | - | - | - | - | - | - |
| F 10 | <i>Aulostomus maculatus</i> | - | 0.049 | 0.012 | 0.012 | - | - | 0.019 | 0.019 | - |
| F 11 | <i>Balistes vetula</i> | 0.002 | - | - | - | - | - | - | - | - |
| F 12 | <i>Canthidermis sufflamen</i> | - | - | - | - | - | - | - | - | - |
| F 13 | <i>Melichthys niger</i> | 0.031 | - | - | - | - | - | - | - | - |
| F 14 | <i>Platybelone argalus argalus</i> | - | - | - | - | - | - | - | - | - |
| F 15 | <i>Strongylura timucu</i> | - | - | - | - | - | - | - | - | - |
| F 16 | <i>Tylosurus acus</i> | - | - | - | - | - | - | - | - | - |
| F 17 | <i>Tylosurus crocodilus</i> | - | 0.182 | - | - | - | - | - | - | - |
| F 18 | <i>Entomacrodus nigricans</i> | - | - | - | - | - | - | - | - | - |
| F 19 | <i>Ophioblennius atlanticus</i> | - | - | - | - | - | - | - | - | - |
| F 20 | <i>Parablennius marmoratus</i> | - | - | - | - | - | - | - | - | - |
| F 21 | <i>Scartella cristata</i> | - | - | - | - | - | - | - | - | - |
| F 22 | <i>Bothus lunatus</i> | - | - | - | - | - | - | - | - | - |
| F 23 | <i>Bothus ocellatus</i> | - | - | - | - | - | - | - | - | - |
| F 24 | <i>Caranx bartholomaei</i> | 0.4 | - | - | - | - | - | - | - | - |
| F 25 | <i>Caranx latus</i> | - | - | - | - | - | - | - | - | 0.145 |
| F 26 | <i>Caranx ruber</i> | 0.051 | 0.017 | 0.017 | 0.068 | - | - | - | - | 0.025 |
| F 27 | <i>Decapterus punctatus</i> | - | - | - | - | - | - | - | - | - |
| F 28 | <i>Oligoplites saurus</i> | - | - | - | - | - | - | - | - | - |
| F 29 | <i>Selar crumenophthalmus</i> | 0.096 | - | - | - | - | - | - | - | - |
| F 30 | <i>Seriola dumerili</i> | - | - | - | - | - | - | - | - | - |
| F 31 | <i>Trachinotus falcatus</i> | - | - | - | - | - | - | - | - | - |
| F 32 | <i>Trachinotus goodei</i> | - | - | - | - | - | - | - | - | - |
| F 33 | <i>Carcharhinus acronotus</i> | - | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 |
| F 34 | <i>Carcharhinus falciformis</i> | - | 0.005 | 0.005 | - | - | - | - | - | - |
| F 35 | <i>Carcharhinus leucas</i> | - | - | - | - | - | - | - | - | - |
| F 36 | <i>Carcharhinus limbatus</i> | - | - | - | - | - | - | - | - | 0.05 |
| F 37 | <i>Carcharhinus longimanus</i> | - | - | - | - | - | - | - | - | - |
| F 38 | <i>Carcharhinus perezi</i> | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | <i>Galeocerdo cuvier</i> | - | 0.003 | 0.003 | 0.003 | - | - | - | - | - |
| F 40 | <i>Negaprion brevirostris</i> | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 41 | <i>Rhizoprionodon porosus</i> | - | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 |
| F 42 | <i>Chaetodon aculeatus</i> | - | - | - | - | - | - | - | - | - |
| F 43 | <i>Chaetodon capistratus</i> | - | - | - | - | - | - | - | - | - |
| F 44 | <i>Chaetodon sedentarius</i> | - | - | - | - | - | - | - | - | - |
| F 45 | <i>Chaetodon striatus</i> | - | - | - | - | - | - | - | - | - |
| F 46 | <i>Amblycirrhilus pinos</i> | - | - | - | - | - | - | - | - | - |
| F 47 | <i>Labrisomus guppyi</i> | - | - | - | - | - | - | - | - | - |
| F 48 | <i>Labrisomus nuchipinnis</i> | 0.105 | - | - | - | - | - | - | - | - |
| F 49 | <i>Harengula clupeiola</i> | - | - | - | - | - | - | - | - | - |
| F 50 | <i>Harengula humeralis</i> | - | - | - | - | - | - | - | - | - |
| F 51 | <i>Jenkinsia lamprotaenia</i> | - | - | - | - | - | - | - | - | - |
| F 52 | <i>Opisthonema oglinum</i> | 0.098 | - | - | - | - | - | - | - | - |
| F 53 | <i>Heteroconger halis</i> | - | - | - | - | - | - | - | - | - |
| F 54 | <i>Dactylopterus volitans</i> | 0.05 | - | - | - | - | - | - | - | - |
| F 55 | <i>Dasyatis americana</i> | - | 0.018 | 0.018 | 0.018 | - | - | - | - | - |
| F 56 | <i>Chilomycterus antennatus</i> | - | - | - | - | - | - | - | - | - |
| F 57 | <i>Diodon holocanthus</i> | - | - | - | - | - | - | - | - | - |
| F 58 | <i>Diodon hystrix</i> | - | - | - | - | - | - | - | - | - |
| F 59 | <i>Inermia vittata</i> | - | - | - | - | - | - | - | - | - |
| F 60 | <i>Anchoa hepsetus</i> | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F9 | F10 | F11 | F12 | F13 | F14 | F15 | F16 | F17 | F18 | F19 | F20 | F21 | F22 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | 0.009 | 0.046 | 0.009 | 0.009 | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | 0.24 | 0.24 | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | 0.145 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | 0.025 | - | - | - | - | - | - | - | - | 0.051 | 0.051 | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | 0.067 | 0.067 | 0.066 | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | 0.01 | 0.01 | - | - | 0.01 | 0.01 | 0.01 | - | - | - | 0.01 | - | 0.01 | - |
| F 34 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.006 | 0.006 | 0.006 | 0.006 | - | - | - | - | 0.005 |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 36 | 0.05 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | - | - | - | 0.003 |
| F 40 | 0.005 | 0.005 | - | - | 0.005 | 0.005 | 0.005 | 0.005 | - | - | 0.005 | - | 0.005 | - |
| F 41 | 0.007 | 0.007 | - | - | 0.007 | 0.007 | 0.007 | - | - | - | 0.007 | - | 0.007 | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F23 | F24 | F25 | F26 | F27 | F28 | F29 | F30 | F31 | F32 | F33 | F34 | F35 | F36 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|------|------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | 0.054 | 0.054 | 0.054 | 0.054 | 0.054 | 0.054 | 0.053 | 0.053 | 0.053 | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | 0.286 | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | 0.2 | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | - | - | - | - | - | 0.01 | - | - | - | - | - | - | - | - |
| F 34 | 0.005 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | - | - | 0.006 |
| F 35 | - | - | - | - | - | - | - | - | - | - | 0.05 | - | - | 0.05 |
| F 36 | - | - | - | - | - | 0.05 | - | - | - | - | - | - | - | - |
| F 37 | - | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | - | - |
| F 39 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.01 | 0.01 | 0.01 | 0.01 |
| F 40 | - | - | - | - | - | 0.005 | - | - | - | - | - | - | - | - |
| F 41 | - | - | - | - | - | 0.007 | - | - | - | - | - | - | - | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F37 | F38 | F39 | F40 | F41 | F42 | F43 | F44 | F45 | F46 | F47 | F48 | F49 | F50 |
|------|------|-------|-----|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | - | 0.073 | 0.037 | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.182 |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | 0.145 | 0.145 |
| F 26 | - | - | - | - | - | - | - | - | - | - | - | - | 0.051 | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | - | - | - | - | - | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | - | 0.01 | 0.01 | 0.01 |
| F 34 | - | 0.006 | - | 0.006 | 0.006 | - | - | - | - | - | - | - | 0.005 | 0.005 |
| F 35 | - | 0.05 | - | 0.05 | 0.05 | - | - | - | - | - | - | - | - | - |
| F 36 | - | - | - | - | - | - | - | - | - | - | - | - | 0.05 | 0.05 |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | - | - | - | 0.005 | 0.01 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | 0.01 | 0.01 | - | 0.01 | 0.01 | - | 0.003 | 0.003 | 0.003 | - | - | - | 0.003 | 0.003 |
| F 40 | - | - | - | - | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.005 | 0.005 | 0.005 |
| F 41 | - | - | - | - | - | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | - | - | 0.007 | 0.007 |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.2.6 continued

| | F51 | F52 | F53 | F54 | F55 | F56 | F57 | F58 | F59 | F60 | F61 | F62 | F63 | F64 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | 0.019 | 0.018 | - | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | 0.97 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | 0.48 | - | - | - | - | - | - | - | - | 0.24 | 0.24 | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | 0.182 | 0.181 | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | 0.286 | - | - | - | - | - | - | - | - | 0.143 | 0.142 | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | 0.051 | - | - | - | - | - | - | - | - | 0.051 | 0.051 | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | 0.987 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | 0.426 | - | - | - | - | - | - | - | - | 0.213 | 0.213 | - | - | - |
| F 33 | 0.01 | 0.01 | 0.01 | - | - | - | - | - | 0.01 | 0.005 | 0.005 | - | - | 0.01 |
| F 34 | - | 0.005 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | - | 0.005 | 0.005 | 0.005 |
| F 35 | - | - | - | - | 0.05 | - | - | - | - | - | - | - | - | - |
| F 36 | 0.05 | 0.05 | - | - | 0.05 | - | - | - | 0.05 | 0.025 | 0.025 | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.003 | 0.002 | 0.005 | 0.005 | 0.005 |
| F 39 | - | 0.003 | 0.003 | 0.003 | 0.04 | 0.003 | 0.003 | 0.003 | 0.003 | - | - | 0.003 | 0.003 | 0.003 |
| F 40 | 0.005 | 0.005 | 0.005 | - | 0.1 | - | - | - | 0.005 | 0.003 | 0.002 | - | - | 0.005 |
| F 41 | 0.007 | 0.014 | 0.007 | - | - | - | - | - | 0.007 | 0.004 | 0.003 | - | - | 0.007 |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | 0.028 | 0.027 | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F65 | F66 | F67 | F68 | F69 | F70 | F71 | F72 | F73 | F74 | F75 | F76 | F77 | F78 |
|------|-------|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | 0.75 | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | 0.012 | 0.012 | 0.012 | - | - | - | - | - | - | 0.037 | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | 0.283 | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | 0.2 | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | 0.01 | - | - | - | - | 0.01 | 0.01 | 0.01 | - | 0.01 | - | 0.01 | 0.01 | 0.01 |
| F 34 | 0.005 | - | - | - | - | - | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 36 | - | - | - | - | - | - | - | - | - | - | - | - | 0.05 | 0.05 |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | 0.003 | 0.01 | - | - | - | - | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 |
| F 40 | 0.005 | - | - | - | - | 0.005 | 0.005 | 0.005 | - | 0.005 | - | 0.005 | 0.005 | 0.005 |
| F 41 | 0.007 | - | - | - | - | 0.007 | 0.007 | 0.007 | - | 0.007 | - | 0.007 | 0.007 | 0.007 |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F79 | F80 | F81 | F82 | F83 | F84 | F85 | F86 | F87 | F88 | F89 | F90 | F91 | F92 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | 0.037 | - | - | - | - | - | - | - | 0.037 | - | 0.037 | - | 0.037 | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | 0.291 | - |
| F 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | 0.01 | - | - | - | - | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | - |
| F 34 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 36 | 0.05 | - | - | - | - | 0.05 | 0.05 | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 |
| F 40 | 0.005 | - | - | - | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - |
| F 41 | 0.007 | - | - | - | - | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F121 | F122 | F123 | F124 | F125 | F126 | F127 | F128 | F129 | F130 | F131 | F132 | F133 | F134 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | 0.037 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | 0.051 | 0.051 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | 0.01 | 0.01 | 0.01 | 0.01 | - | - | - | 0.01 | 0.01 | - | 0.01 | 0.01 | - | - |
| F 34 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.006 | 0.005 | 0.005 | 0.005 | 0.005 | - | - | - | 0.005 |
| F 35 | - | - | - | - | - | - | 0.05 | - | - | - | - | - | - | - |
| F 36 | - | - | - | - | - | - | 0.05 | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.04 | 0.003 | 0.003 | 0.003 | 0.003 | - | - | - | 0.003 |
| F 40 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.1 | - | 0.005 | 0.005 | - | 0.005 | 0.005 | - | - |
| F 41 | 0.007 | 0.007 | 0.007 | 0.007 | - | - | - | 0.007 | 0.007 | - | 0.007 | 0.007 | - | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | 0.018 | 0.018 | 0.018 | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F135 | F136 | F137 | F138 | F139 | F140 | F141 | F142 | F143 | F144 | F145 | F146 | F147 | F148 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | - | - | - | 0.037 | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | - | - | - | - | 0.01 | - | - | - | - | - | 0.01 | 0.01 | 0.01 | 0.01 |
| F 34 | 0.005 | 0.005 | 0.005 | 0.005 | - | - | 0.005 | 0.005 | 0.006 | 0.006 | 0.005 | 0.005 | - | 0.005 |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 36 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | 0.003 |
| F 40 | - | - | - | - | 0.005 | - | - | - | - | - | 0.005 | 0.005 | 0.005 | 0.005 |
| F 41 | - | - | - | - | 0.007 | - | - | - | - | - | 0.007 | 0.007 | 0.007 | 0.007 |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F149 | F150 | F151 | F152 | F153 | F154 | F155 | F156 | F157 | F158 | F159 | F160 | F161 | F162 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | 0.018 | 0.018 | 0.018 | 0.018 | - | - | - | - | - | - | - | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | 0.04 | 0.04 |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | - | - | - | 0.051 | - | - | - | 0.005 | 0.056 | 0.005 | 0.005 | 0.005 | 0.056 | 0.005 |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | 0.2 | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | - | 0.01 | - | 0.01 | - | 0.01 | - | 0.01 | - |
| F 34 | 0.005 | 0.005 | - | - | - | 0.005 | 0.005 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.005 | 0.005 |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 36 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | 0.003 | 0.003 | - | - | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 |
| F 40 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.005 | - | 0.005 | - | 0.005 | - | 0.005 | - |
| F 41 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | - | 0.007 | - | 0.007 | - | 0.007 | - | 0.007 | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F163 | F164 | F165 | F166 | F167 | F168 | F169 | F170 | F171 | F172 | F173 | F174 | F175 | F176 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | 0.5 | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | 0.04 | 0.04 | 0.04 | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | 0.005 | 0.005 | 0.005 | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | 0.01 | - | - | 0.01 | 0.01 | 0.01 | 0.01 | - | - | - | - | 0.01 | - | - |
| F 34 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.005 |
| F 35 | - | - | - | - | - | - | - | 0.05 | 0.05 | 0.05 | - | - | - | - |
| F 36 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | 0.05 | 0.05 | 0.05 | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.01 | 0.005 | 0.01 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | 0.003 |
| F 40 | 0.005 | - | - | 0.005 | 0.005 | 0.005 | 0.005 | - | - | - | - | 0.005 | - | - |
| F 41 | 0.007 | - | - | 0.007 | 0.007 | 0.007 | 0.007 | - | - | - | - | 0.007 | - | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.055 |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F177 | F178 | F179 | F180 | F181 | F182 | F183 | F184 | F185 | F186 | F187 | F188 | F189 | F190 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | 0.01 | - | 0.01 | 0.01 | - | - | - | - | 0.01 | 0.01 | 0.01 | 0.01 | - | - |
| F 34 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.006 | 0.006 | - | - | - | - | 0.006 | 0.006 |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 36 | 0.05 | - | - | - | - | - | - | - | - | - | 0.05 | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 39 | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | - | - | - | 0.003 | 0.003 |
| F 40 | 0.005 | - | 0.005 | 0.005 | - | - | - | - | 0.005 | 0.005 | 0.005 | 0.005 | - | - |
| F 41 | 0.007 | - | 0.007 | 0.007 | - | - | - | - | 0.007 | 0.007 | 0.007 | 0.007 | - | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F191 | F192 | F193 | F194 | F195 | F196 | F197 | F198 | F199 | F200 | F201 | F202 | F203 | F204 |
|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|------|-------|-------|-------|
| F 1 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 2 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 3 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 4 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 5 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 6 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 7 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 8 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 9 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 10 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 11 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 12 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 13 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 14 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 15 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 16 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 17 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 18 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 19 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 20 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 21 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 22 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 23 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 24 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 25 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 26 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 27 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 28 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 29 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 30 | - | - | - | - | 0.067 | 0.067 | 0.066 | - | - | - | - | - | - | - |
| F 31 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 32 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 33 | - | 0.01 | 0.01 | - | - | - | - | 0.01 | - | - | - | - | 0.01 | - |
| F 34 | 0.006 | 0.006 | 0.005 | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.006 | 0.005 | 0.005 |
| F 35 | - | - | - | - | - | - | - | - | - | - | - | 0.05 | - | - |
| F 36 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 37 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 38 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | - | 0.01 | 0.005 | 0.005 |
| F 39 | 0.003 | 0.003 | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.01 | 0.01 | 0.003 | 0.003 |
| F 40 | - | 0.005 | 0.005 | - | - | - | - | 0.005 | - | - | - | - | 0.005 | - |
| F 41 | - | 0.007 | 0.007 | - | - | - | - | 0.007 | - | - | - | - | 0.007 | - |
| F 42 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 43 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 44 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 45 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 46 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 47 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 48 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 49 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 50 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 51 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 52 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 53 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 54 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 55 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 56 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 57 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 58 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 59 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 60 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F205 | F206 | F207 | F208 | Sum |
|------|-------|-------|-------|-------|-----|
| F 1 | - | - | - | - | 1 |
| F 2 | - | - | - | - | 1 |
| F 3 | - | - | - | - | 1 |
| F 4 | - | - | - | - | 1 |
| F 5 | - | - | - | - | 1 |
| F 6 | - | - | - | - | 1 |
| F 7 | - | - | - | - | 1 |
| F 8 | - | - | - | - | 1 |
| F 9 | - | - | - | - | 1 |
| F 10 | - | - | - | - | 1 |
| F 11 | - | - | - | - | 1 |
| F 12 | - | - | - | - | 1 |
| F 13 | - | - | - | - | 1 |
| F 14 | - | - | - | - | 1 |
| F 15 | - | - | - | - | 1 |
| F 16 | - | - | - | - | 1 |
| F 17 | - | - | - | - | 1 |
| F 18 | - | - | - | - | 1 |
| F 19 | - | - | - | - | 1 |
| F 20 | - | - | - | - | 1 |
| F 21 | - | - | - | - | 1 |
| F 22 | - | - | - | - | 1 |
| F 23 | - | - | - | - | 1 |
| F 24 | - | - | - | - | 1 |
| F 25 | - | - | - | - | 1 |
| F 26 | - | - | - | - | 1 |
| F 27 | - | - | - | - | 1 |
| F 28 | - | - | - | - | 1 |
| F 29 | - | - | - | - | 1 |
| F 30 | - | - | - | - | 1 |
| F 31 | - | - | - | - | 1 |
| F 32 | - | - | - | - | 1 |
| F 33 | 0.01 | - | 0.01 | - | 1 |
| F 34 | 0.005 | - | - | 0.006 | 1 |
| F 35 | - | - | - | 0.05 | 1 |
| F 36 | - | - | - | - | 1 |
| F 37 | - | - | - | - | 1 |
| F 38 | 0.005 | 0.005 | 0.005 | 0.01 | 1 |
| F 39 | 0.003 | - | - | 0.01 | 1 |
| F 40 | 0.005 | - | 0.005 | - | 1 |
| F 41 | 0.007 | - | 0.007 | - | 1 |
| F 42 | - | - | - | - | 1 |
| F 43 | - | - | - | - | 1 |
| F 44 | - | - | - | - | 1 |
| F 45 | - | - | - | - | 1 |
| F 46 | - | - | - | - | 1 |
| F 47 | - | - | - | - | 1 |
| F 48 | - | - | - | - | 1 |
| F 49 | - | - | - | - | 1 |
| F 50 | - | - | - | - | 1 |
| F 51 | - | - | - | - | 1 |
| F 52 | - | - | - | - | 1 |
| F 53 | - | - | - | - | 1 |
| F 54 | - | - | - | - | 1 |
| F 55 | - | - | - | - | 1 |
| F 56 | - | - | - | - | 1 |
| F 57 | - | - | - | - | 1 |
| F 58 | - | - | - | - | 1 |
| F 59 | - | - | - | - | 1 |
| F 60 | - | - | - | - | 1 |

continued...

Table 8.6.2 continued

| | Predator/prey | D1 | A1 | A2 | A3 | I1 | I2 | I3 | I4 | I5 |
|-------|------------------------------------|-------|-------|-------|-----|-------|-------|-------|-------|-------|
| F 61 | <i>Anchoa lyolepis</i> | 0.1 | - | - | 0.7 | - | 0.2 | - | - | - |
| F 62 | <i>Chaetodipterus faber</i> | - | 0.073 | 0.052 | - | - | 0.168 | 0.327 | - | 0.025 |
| F 63 | <i>Fistularia tabacaria</i> | - | - | - | - | - | - | - | - | - |
| F 64 | <i>Eucinostomus argenteus</i> | - | - | - | - | - | 0.021 | - | - | - |
| F 65 | <i>Gerres cinereus</i> | - | - | - | - | - | - | - | - | - |
| F 66 | <i>Ginglymostoma cirratum</i> | - | - | - | - | - | - | - | - | - |
| F 67 | <i>Coryphopterus glaucofraenum</i> | 0.25 | 0.25 | - | - | - | 0.3 | - | - | - |
| F 68 | <i>Gnatholepis thompsoni</i> | 0.37 | 0.37 | - | - | - | 0.217 | - | - | - |
| F 69 | <i>Gobiosoma evelynae</i> | - | - | - | - | - | - | - | - | - |
| F 70 | <i>Gramma loreto</i> | - | - | - | - | - | 1 | - | - | - |
| F 71 | <i>Gramma melacara</i> | - | - | - | - | - | 1 | - | - | - |
| F 72 | <i>Rypticus saponaceus</i> | - | - | - | - | - | - | - | - | - |
| F 73 | <i>Anisotremus surinamensis</i> | - | - | - | - | - | - | - | - | - |
| F 74 | <i>Anisotremus virginicus</i> | - | - | - | - | 0.005 | 0.05 | - | - | - |
| F 75 | <i>Haemulon album</i> | - | - | - | - | - | - | 0.001 | - | - |
| F 76 | <i>Haemulon aurolineatum</i> | - | - | - | - | - | 0.35 | - | - | - |
| F 77 | <i>Haemulon carbonarium</i> | - | - | - | - | - | 0.001 | - | - | - |
| F 78 | <i>Haemulon chrysargyreum</i> | - | - | - | - | - | 0.23 | - | - | - |
| F 79 | <i>Haemulon flavolineatum</i> | - | - | - | - | - | - | - | - | - |
| F 80 | <i>Haemulon macrostomum</i> | - | - | - | - | - | - | - | - | - |
| F 81 | <i>Haemulon parrai</i> | - | - | - | - | - | - | - | - | - |
| F 82 | <i>Haemulon plumieri</i> | - | - | - | - | - | - | - | - | - |
| F 83 | <i>Haemulon sciurus</i> | - | - | - | - | - | 0.016 | - | - | - |
| F 84 | <i>Hemiramphus balao</i> | - | - | - | - | - | 0.678 | - | - | - |
| F 85 | <i>Hemiramphus brasiliensis</i> | - | 0.81 | - | - | - | - | - | - | - |
| F 86 | <i>Holocentrus ascensionis</i> | - | - | - | - | - | - | - | - | - |
| F 87 | <i>Holocentrus coruscus</i> | - | - | - | - | - | - | - | - | - |
| F 88 | <i>Neoniphon marianus</i> | - | - | - | - | - | 0.051 | - | - | - |
| F 89 | <i>Holocentrus rufus</i> | - | - | - | - | - | 0.026 | - | - | - |
| F 90 | <i>Sargocentron vexillarium</i> | - | - | - | - | - | 0.275 | - | - | - |
| F 91 | <i>Myrpristis jacobus</i> | - | - | - | - | - | 0.785 | - | - | - |
| F 92 | <i>Kyphosus incisor</i> | - | 1 | - | - | - | - | - | - | - |
| F 93 | <i>Kyphosus sectatrix</i> | - | 1 | - | - | - | - | - | - | - |
| F 94 | <i>Bodianus rufus</i> | - | - | - | - | - | - | - | - | - |
| F 95 | <i>Clepticus parrae</i> | - | - | - | - | - | 1 | - | - | - |
| F 96 | <i>Halichoeres bivittatus</i> | - | - | - | - | 0.001 | - | - | - | - |
| F 97 | <i>Halichoeres garnoti</i> | - | - | - | - | - | - | - | - | - |
| F 98 | <i>Halichoeres maculipinna</i> | - | - | - | - | - | 0.188 | - | - | - |
| F 99 | <i>Halichoeres poeyi</i> | - | - | - | - | - | - | - | - | - |
| F 100 | <i>Halichoeres radiatus</i> | - | - | - | - | - | - | - | - | - |
| F 101 | <i>Xyrichtys novacula</i> | - | - | - | - | - | - | - | - | - |
| F 102 | <i>Xyrichtys splendens</i> | - | - | - | - | - | 0.775 | - | - | - |
| F 103 | <i>Lachnolaimus maximus</i> | - | - | - | - | - | - | - | - | - |
| F 104 | <i>Thalassoma bifasciatum</i> | - | - | - | - | - | 0.439 | - | - | - |
| F 105 | <i>Lutjanus analis</i> | - | - | - | - | - | - | - | - | - |
| F 106 | <i>Lutjanus apodus</i> | - | - | - | - | - | - | - | - | - |
| F 107 | <i>Lutjanus cyanopterus</i> | - | - | - | - | - | - | - | - | - |
| F 108 | <i>Lutjanus griseus</i> | - | - | - | - | - | - | - | - | - |
| F 109 | <i>Lutjanus jocu</i> | - | - | - | - | - | 0.018 | - | - | - |
| F 110 | <i>Lutjanus mahagoni</i> | - | - | - | - | - | - | - | - | - |
| F 111 | <i>Lutjanus synagris</i> | - | 0.027 | - | - | - | - | 0.014 | - | - |
| F 112 | <i>Ocyurus chrysurus</i> | - | - | - | - | - | 0.571 | - | - | - |
| F 113 | <i>Malacanthus plumieri</i> | - | - | - | - | - | - | - | - | - |
| F 114 | <i>Tarpon atlanticus</i> | - | - | - | - | - | - | - | - | - |
| F 115 | <i>Aluterus schoepfi</i> | - | 0.988 | - | - | - | - | - | - | - |
| F 116 | <i>Aluterus scripta</i> | - | 0.432 | 0.312 | - | - | - | 0.004 | 0.158 | 0.05 |
| F 117 | <i>Cantherines macroceros</i> | - | 0.029 | 0.059 | - | - | - | 0.865 | 0.02 | 0.019 |
| F 118 | <i>Cantherines pullus</i> | 0.214 | 0.26 | 0.035 | - | - | 0.001 | 0.327 | 0.013 | 0.007 |
| F 119 | <i>Monacanthus ciliatus</i> | 0.107 | 0.26 | 0.002 | - | - | 0.306 | - | 0.001 | - |
| F 120 | <i>Mugil curema</i> | 0.5 | 0.5 | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I6 | I7 | I8 | I9 | I10 | I11 | I12 | I13 | I14 | I15 | I16 | I17 | I18 | I19 |
|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | 0.185 | 0.009 | - | - | - | - | - | - | - | - | - | 0.137 | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | 0.027 | - | - | 0.017 | 0.035 | - | - | - | 0.199 | - | - |
| F 65 | - | - | - | 0.061 | - | - | 0.147 | 0.233 | - | - | - | 0.128 | - | - |
| F 66 | - | - | - | - | - | 0.02 | 0.02 | 0.02 | 0.02 | 0.05 | 0.02 | - | - | - |
| F 67 | - | - | - | - | - | - | - | 0.1 | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | 0.168 | 0.015 | - | - | - | - | - | - |
| F 74 | - | - | - | 0.004 | - | 0.008 | 0.038 | 0.055 | 0.002 | - | - | 0.14 | - | - |
| F 75 | - | - | 0.001 | 0.257 | 0.008 | 0.008 | 0.028 | 0.055 | 0.002 | - | - | 0.143 | 0.01 | - |
| F 76 | - | - | - | - | - | - | 0.021 | 0.016 | 0.004 | - | - | 0.31 | - | - |
| F 77 | - | - | - | 0.048 | - | 0.098 | 0.152 | - | - | - | - | 0.08 | - | - |
| F 78 | - | - | - | 0.012 | - | - | 0.018 | 0.147 | - | 0.001 | 0.001 | 0.191 | - | - |
| F 79 | - | - | - | 0.102 | - | 0.057 | 0.005 | 0.032 | 0.008 | - | 0.033 | 0.398 | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | 0.034 | - | - | - | - | - | 0.051 | 0.028 | 0.002 | - | - | 0.029 | - | - |
| F 82 | - | - | - | 0.086 | - | 0.007 | 0.071 | 0.013 | - | - | - | 0.15 | - | - |
| F 83 | - | - | 0.003 | 0.032 | - | - | 0.048 | 0.158 | 0.002 | - | 0.016 | 0.053 | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | 0.101 | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | 0.01 | - | - | - | - | 0.039 | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | 0.006 | 0.078 | - | - | - | - | 0.045 | - | - |
| F 90 | - | - | - | - | - | 0.103 | 0.251 | - | - | - | - | 0.04 | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | 0.07 | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | 0.007 | 0.105 | 0.081 | - | - | - | 0.002 | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | 0.034 | 0.129 | 0.039 | - | - | - | 0.181 | - | - |
| F 97 | - | - | - | 0.005 | - | 0.021 | 0.146 | 0.076 | - | - | - | 0.06 | - | - |
| F 98 | - | - | - | 0.009 | - | 0.015 | 0.053 | 0.024 | 0.002 | - | 0.043 | 0.475 | - | - |
| F 99 | - | - | - | 0.054 | - | 0.046 | 0.222 | 0.23 | - | - | - | 0.019 | - | - |
| F 100 | - | - | - | - | - | 0.002 | 0.213 | 0.251 | - | - | - | 0.014 | - | - |
| F 101 | - | - | - | - | - | - | 0.411 | 0.298 | 0.06 | - | - | 0.08 | - | - |
| F 102 | - | - | - | - | - | - | 0.051 | 0.034 | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | 0.397 | 0.426 | 0.006 | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | 0.063 | - | - | - | - | 0.056 | - | 0.022 |
| F 105 | - | - | - | - | - | - | 0.133 | - | - | - | 0.031 | - | - | - |
| F 106 | - | - | - | - | - | - | 0.005 | - | - | - | 0.035 | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | 0.068 | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | 0.036 | - | - | 0.022 | 0.07 | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | 0.094 | - | - | - |
| F 111 | - | - | - | - | - | 0.002 | 0.003 | 0.002 | 0.002 | 0.002 | 0.002 | 0.041 | - | - |
| F 112 | - | - | - | - | - | - | 0.025 | 0.012 | - | 0.01 | 0.01 | - | - | - |
| F 113 | - | - | - | 0.104 | - | 0.057 | - | - | - | - | - | 0.104 | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | 0.006 | - | - | - | - | - | - | - |
| F 116 | 0.024 | - | - | - | - | - | 0.006 | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | 0.018 | 0.003 | 0.024 | - | - | - | - | 0.002 | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | 0.016 | 0.024 | - | - | - | 0.043 | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I20 | I21 | I22 | I23 | I24 | I25 | I26 | I27 | I28 | I29 | I30 | I31 | I32 | I33 |
|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | 0.001 | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | 0.443 | 0.062 | - | 0.081 | - | - | - | 0.115 | - | - | - | - |
| F 65 | - | 0.032 | 0.004 | - | - | 0.05 | - | - | - | 0.307 | 0.019 | - | 0.019 | - |
| F 66 | - | - | - | - | - | 0.05 | 0.05 | 0.05 | - | 0.05 | - | - | - | 0.05 |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | 0.1 | - |
| F 68 | - | - | 0.043 | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | 1 | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | 0.083 | - | - | - | 0.342 | - | - | - | 0.096 | - | - | - | - |
| F 73 | - | 0.025 | - | - | - | 0.025 | - | - | 0.033 | 0.105 | - | 0.018 | 0.055 | 0.541 |
| F 74 | - | 0.052 | 0.033 | 0.012 | 0.09 | 0.162 | - | - | 0.006 | 0.178 | - | - | 0.165 | - |
| F 75 | - | 0.017 | 0.006 | - | - | 0.049 | - | - | 0.006 | 0.097 | 0.036 | - | 0.053 | 0.203 |
| F 76 | - | - | 0.04 | 0.011 | 0.003 | 0.197 | - | - | 0.026 | 0.022 | - | - | - | - |
| F 77 | 0.019 | - | 0.002 | - | - | 0.041 | - | - | - | 0.383 | - | - | 0.06 | 0.109 |
| F 78 | - | 0.065 | 0.063 | - | 0.052 | 0.088 | - | - | 0.02 | 0.108 | - | - | - | - |
| F 79 | - | 0.008 | 0.006 | - | 0.041 | 0.036 | - | - | 0.009 | 0.169 | - | - | 0.031 | 0.015 |
| F 80 | - | - | - | - | - | - | - | - | - | 0.132 | - | - | - | 0.868 |
| F 81 | - | 0.005 | 0.072 | - | 0.005 | 0.38 | - | - | - | 0.337 | - | - | 0.024 | - |
| F 82 | - | - | 0.006 | 0.004 | - | 0.069 | - | - | - | 0.311 | 0.034 | - | 0.059 | 0.128 |
| F 83 | - | 0.048 | 0.021 | 0.001 | 0.015 | 0.11 | - | - | - | 0.295 | - | - | 0.059 | 0.092 |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | 0.007 | 0.119 | - | - | - | 0.825 | - | - | - | - |
| F 87 | - | - | - | - | - | 0.719 | - | - | - | 0.281 | - | - | - | - |
| F 88 | - | - | - | - | - | 0.596 | - | - | - | 0.353 | - | - | - | - |
| F 89 | - | 0.004 | - | - | 0.016 | 0.155 | - | - | - | 0.587 | - | - | 0.071 | - |
| F 90 | - | - | - | - | 0.059 | 0.107 | - | - | - | 0.139 | - | - | - | - |
| F 91 | - | - | 0.033 | - | 0.018 | 0.063 | - | - | - | 0.031 | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | 0.013 | - | - | 0.007 | 0.063 | - | - | 0.039 | 0.342 | - | - | 0.196 | 0.145 |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | 0.032 | - | - | - | 0.038 | - | - | 0.009 | 0.241 | - | - | 0.076 | 0.187 |
| F 97 | - | - | - | - | - | 0.081 | - | - | 0.048 | 0.331 | - | - | 0.168 | 0.032 |
| F 98 | - | - | 0.055 | 0.015 | 0.018 | 0.07 | - | - | - | 0.033 | - | - | - | - |
| F 99 | - | 0.002 | - | - | 0.004 | - | - | - | 0.053 | 0.331 | - | - | 0.106 | 0.071 |
| F 100 | - | 0.009 | - | - | - | - | - | - | 0.047 | 0.197 | - | - | 0.068 | 0.199 |
| F 101 | - | - | 0.008 | - | 0.047 | 0.096 | - | - | - | - | - | - | - | - |
| F 102 | - | - | 0.132 | - | 0.008 | - | - | - | - | - | - | - | - | - |
| F 103 | 0.005 | - | 0.01 | - | - | - | - | - | 0.049 | 0.061 | - | - | - | 0.046 |
| F 104 | - | - | - | 0.025 | 0.077 | 0.042 | - | - | - | 0.083 | - | - | 0.082 | 0.016 |
| F 105 | - | 0.008 | - | - | - | 0.023 | 0.019 | - | 0.029 | 0.453 | - | - | - | - |
| F 106 | - | 0.045 | - | - | - | 0.041 | - | - | - | 0.267 | - | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | 0.132 | - | 0.009 | - | 0.4 | - | - | - | - |
| F 109 | - | - | - | - | - | - | 0.069 | 0.018 | - | 0.16 | - | - | - | - |
| F 110 | - | - | - | - | - | 0.125 | - | - | - | 0.031 | - | - | - | - |
| F 111 | - | 0.027 | 0.007 | 0.007 | 0.007 | 0.206 | 0.007 | 0.007 | 0.007 | 0.206 | - | - | - | - |
| F 112 | - | 0.025 | - | - | - | 0.066 | - | - | - | 0.203 | - | - | - | - |
| F 113 | - | 0.15 | 0.016 | - | - | 0.014 | - | - | - | 0.185 | - | - | 0.219 | 0.027 |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | 0.006 | - | - | - | - | - |
| F 116 | - | - | - | - | - | 0.003 | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | 0.006 | - | - | 0.005 | - | - | - | - | - | - | 0.02 | 0.001 |
| F 119 | - | - | 0.067 | 0.056 | 0.038 | 0.08 | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | | I34 | I35 | R1 | B1 |
|-------|-------|-----|-------|----|----|
| F 61 | - | - | - | - | - |
| F 62 | 0.023 | - | - | - | - |
| F 63 | - | - | - | - | - |
| F 64 | - | - | - | - | - |
| F 65 | - | - | - | - | - |
| F 66 | - | - | - | - | - |
| F 67 | - | - | - | - | - |
| F 68 | - | - | - | - | - |
| F 69 | - | - | - | - | - |
| F 70 | - | - | - | - | - |
| F 71 | - | - | - | - | - |
| F 72 | - | - | - | - | - |
| F 73 | - | - | - | - | - |
| F 74 | - | - | - | - | - |
| F 75 | - | - | - | - | - |
| F 76 | - | - | - | - | - |
| F 77 | - | - | - | - | - |
| F 78 | - | - | - | - | - |
| F 79 | 0.05 | - | - | - | - |
| F 80 | - | - | - | - | - |
| F 81 | 0.033 | - | - | - | - |
| F 82 | 0.028 | - | - | - | - |
| F 83 | - | - | - | - | - |
| F 84 | - | - | - | - | - |
| F 85 | - | - | - | - | - |
| F 86 | - | - | - | - | - |
| F 87 | - | - | - | - | - |
| F 88 | - | - | - | - | - |
| F 89 | - | - | - | - | - |
| F 90 | - | - | - | - | - |
| F 91 | - | - | - | - | - |
| F 92 | - | - | - | - | - |
| F 93 | - | - | - | - | - |
| F 94 | - | - | - | - | - |
| F 95 | - | - | - | - | - |
| F 96 | - | - | - | - | - |
| F 97 | - | - | - | - | - |
| F 98 | - | - | - | - | - |
| F 99 | 0.017 | - | - | - | - |
| F 100 | - | - | - | - | - |
| F 101 | - | - | - | - | - |
| F 102 | - | - | - | - | - |
| F 103 | - | - | - | - | - |
| F 104 | - | - | - | - | - |
| F 105 | - | - | - | - | - |
| F 106 | - | - | - | - | - |
| F 107 | - | - | - | - | - |
| F 108 | - | - | - | - | - |
| F 109 | - | - | - | - | - |
| F 110 | - | - | - | - | - |
| F 111 | 0.014 | - | - | - | - |
| F 112 | - | - | - | - | - |
| F 113 | - | - | - | - | - |
| F 114 | - | - | - | - | - |
| F 115 | - | - | - | - | - |
| F 116 | - | - | 0.011 | - | - |
| F 117 | 0.008 | - | - | - | - |
| F 118 | - | - | 0.064 | - | - |
| F 119 | - | - | - | - | - |
| F 120 | - | - | - | - | - |

continued...

Table 8.6.2 continued

| Predator/prey | F0 | F1 | F2 | F3 | F4 | F5 | F6 | F7 | F8 |
|-----------------------------------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 61 <i>Anchoa lyolepis</i> | - | - | - | - | - | - | - | - | - |
| F 62 <i>Chaetodipterus faber</i> | - | - | - | - | - | - | - | - | - |
| F 63 <i>Fistularia tabacaria</i> | - | - | - | - | - | - | - | - | - |
| F 64 <i>Eucinostomus argenteus</i> | - | - | - | - | - | - | - | - | - |
| F 65 <i>Gerres cinereus</i> | - | - | - | - | - | - | - | - | - |
| F 66 <i>Ginglymostoma cirratum</i> | - | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 |
| F 67 <i>Coryphopterus glaucofraenum</i> | - | - | - | - | - | - | - | - | - |
| F 68 <i>Gnatholepis thompsoni</i> | - | - | - | - | - | - | - | - | - |
| F 69 <i>Gobiosoma evelynae</i> | - | - | - | - | - | - | - | - | - |
| F 70 <i>Gramma loreto</i> | - | - | - | - | - | - | - | - | - |
| F 71 <i>Gramma melacara</i> | - | - | - | - | - | - | - | - | - |
| F 72 <i>Rypticus saponaceus</i> | - | - | - | - | - | - | - | - | - |
| F 73 <i>Anisotremus surinamensis</i> | 0.015 | - | - | - | - | - | - | - | - |
| F 74 <i>Anisotremus virginicus</i> | - | - | - | - | - | - | - | - | - |
| F 75 <i>Haemulon album</i> | - | - | - | - | - | - | - | - | - |
| F 76 <i>Haemulon aurolineatum</i> | - | - | - | - | - | - | - | - | - |
| F 77 <i>Haemulon carbonarium</i> | - | - | - | - | - | - | - | - | - |
| F 78 <i>Haemulon chrysargyreum</i> | 0.004 | - | - | - | - | - | - | - | - |
| F 79 <i>Haemulon flavolineatum</i> | - | - | - | - | - | - | - | - | - |
| F 80 <i>Haemulon macrostomum</i> | - | - | - | - | - | - | - | - | - |
| F 81 <i>Haemulon parrai</i> | - | - | - | - | - | - | - | - | - |
| F 82 <i>Haemulon plumieri</i> | 0.034 | - | - | - | - | - | - | - | - |
| F 83 <i>Haemulon sciurus</i> | 0.031 | - | - | - | - | - | - | - | - |
| F 84 <i>Hemiramphus balao</i> | - | - | - | - | - | - | - | - | - |
| F 85 <i>Hemiramphus brasiliensis</i> | - | - | - | - | - | - | - | - | - |
| F 86 <i>Holocentrus ascensionis</i> | - | - | - | - | - | - | - | - | - |
| F 87 <i>Holocentrus coruscus</i> | - | - | - | - | - | - | - | - | - |
| F 88 <i>Neoniphon marianus</i> | - | - | - | - | - | - | - | - | - |
| F 89 <i>Holocentrus rufus</i> | 0.012 | - | - | - | - | - | - | - | - |
| F 90 <i>Sargocentron vexillarium</i> | - | 0.009 | 0.009 | 0.008 | - | - | - | - | - |
| F 91 <i>Myripristis jacobus</i> | - | - | - | - | - | - | - | - | - |
| F 92 <i>Kyphosus incisor</i> | - | - | - | - | - | - | - | - | - |
| F 93 <i>Kyphosus sectatrix</i> | - | - | - | - | - | - | - | - | - |
| F 94 <i>Bodianus rufus</i> | - | - | - | - | - | - | - | - | - |
| F 95 <i>Clepticus parrae</i> | - | - | - | - | - | - | - | - | - |
| F 96 <i>Halichoeres bivittatus</i> | 0.033 | - | - | - | - | - | - | - | - |
| F 97 <i>Halichoeres garnoti</i> | 0.032 | - | - | - | - | - | - | - | - |
| F 98 <i>Halichoeres maculipinna</i> | - | - | - | - | - | - | - | - | - |
| F 99 <i>Halichoeres poeyi</i> | 0.052 | - | - | - | - | - | - | - | - |
| F 100 <i>Halichoeres radiatus</i> | - | - | - | - | - | - | - | - | - |
| F 101 <i>Xyrichtys novacula</i> | - | - | - | - | - | - | - | - | - |
| F 102 <i>Xyrichtys splendens</i> | - | - | - | - | - | - | - | - | - |
| F 103 <i>Lachnolaimus maximus</i> | - | - | - | - | - | - | - | - | - |
| F 104 <i>Thalassoma bifasciatum</i> | 0.095 | - | - | - | - | - | - | - | - |
| F 105 <i>Lutjanus analis</i> | - | 0.02 | - | - | - | - | - | - | - |
| F 106 <i>Lutjanus apodus</i> | - | - | - | - | - | - | - | - | 0.02 |
| F 107 <i>Lutjanus cyanopterus</i> | - | - | - | - | - | - | - | - | - |
| F 108 <i>Lutjanus griseus</i> | - | - | - | - | - | - | - | - | - |
| F 109 <i>Lutjanus jocu</i> | - | - | - | - | - | - | - | - | 0.014 |
| F 110 <i>Lutjanus mahagoni</i> | - | - | - | - | - | - | - | - | 0.125 |
| F 111 <i>Lutjanus synagris</i> | 0.109 | - | - | - | - | - | 0.014 | - | 0.035 |
| F 112 <i>Ocyurus chrysurus</i> | - | - | - | - | - | - | - | - | - |
| F 113 <i>Malacanthus plumieri</i> | - | - | - | - | - | - | - | - | - |
| F 114 <i>Tarpon atlanticus</i> | - | - | - | - | - | - | - | - | 0.5 |
| F 115 <i>Aluterus schoepfi</i> | - | - | - | - | - | - | - | - | - |
| F 116 <i>Aluterus scripta</i> | - | - | - | - | - | - | - | - | - |
| F 117 <i>Cantherines macroceros</i> | - | - | - | - | - | - | - | - | - |
| F 118 <i>Cantherines pullus</i> | - | - | - | - | - | - | - | - | - |
| F 119 <i>Monacanthus ciliatus</i> | - | - | - | - | - | - | - | - | - |
| F 120 <i>Mugil curema</i> | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F9 | F10 | F11 | F12 | F13 | F14 | F15 | F16 | F17 | F18 | F19 | F20 | F21 | F22 |
|-------|-------|-------|-----|-----|-------|-------|-------|-----|-----|-------|-------|-------|-------|-----|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | 0.006 | - | - | 0.006 | 0.006 | 0.006 | - | - | - | 0.006 | - | 0.006 | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | 0.002 | 0.002 | 0.002 | 0.001 | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 106 | 0.02 | 0.041 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | 0.014 | 0.03 | - | - | 0.029 | - | - | - | - | - | - | - | - | - |
| F 110 | 0.125 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | 0.035 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | 0.5 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F23 | F24 | F25 | F26 | F27 | F28 | F29 | F30 | F31 | F32 | F33 | F34 | F35 | F36 |
|-------|-------|-----|-----|-----|-------|-----|-------|-----|-----|-----|-----|-----|-----|-----|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | - | - | - | 0.006 | - | 0.006 | - | - | - | - | - | - | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 106 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F37 | F38 | F39 | F40 | F41 | F42 | F43 | F44 | F45 | F46 | F47 | F48 | F49 | F50 |
|-------|-----|-----|-----|-----|-----|-------|-------|-------|-------|-------|-----|-------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | - | - | - | - | - | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | - | 0.006 | 0.006 | 0.006 |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 106 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | 0.042 | 0.042 |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F51 | F52 | F53 | F54 | F55 | F56 | F57 | F58 | F59 | F60 | F61 | F62 | F63 | F64 |
|-------|-------|-------|-------|-----|-----|-------|-------|-------|-------|-------|-------|-----|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | 0.006 | 0.006 | - | - | - | - | - | 0.006 | 0.003 | 0.003 | - | - | 0.006 |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | 0.221 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | 0.19 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | 0.01 | 0.01 | - | - | - | - | 0.021 | - |
| F 106 | 0.04 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 107 | 0.042 | 0.042 | - | - | - | - | 0.083 | 0.083 | - | - | - | - | - | - |
| F 108 | 0.391 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | 0.029 | 0.029 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | 0.026 | - | - | 0.045 | 0.045 | 0.045 | - | 0.035 | 0.035 | - | - | - |
| F 112 | 0.078 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | 0.062 | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F65 | F66 | F67 | F68 | F69 | F70 | F71 | F72 | F73 | F74 | F75 | F76 | F77 | F78 |
|-------|-------|-----|-------|-------|-------|-------|-------|-------|-----|-------|-------|-------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | - | - | - | - | 0.006 | 0.006 | 0.006 | - | 0.006 | - | 0.006 | 0.006 | 0.006 |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | 0.053 | 0.053 | 0.053 | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | 0.007 | 0.007 | 0.007 | - | - | - | - | - | - | 0.02 | - | - |
| F 106 | - | - | - | - | - | - | - | - | - | - | 0.005 | 0.004 | 0.004 | 0.005 |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | 0.163 | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | - | - | - | - | 0.003 | 0.032 | 0.003 | 0.003 |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F79 | F80 | F81 | F82 | F83 | F84 | F85 | F86 | F87 | F88 | F89 | F90 | F91 | F92 |
|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-----|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | - | - | - | - | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | 0.01 | - | - | 0.01 | - | - | - | 0.02 | - | - | - | - | - | - |
| F 106 | 0.004 | 0.004 | 0.005 | 0.004 | 0.005 | - | - | - | - | - | - | - | - | - |
| F 107 | - | - | - | - | 0.163 | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | 0.003 | 0.003 | 0.003 | 0.032 | 0.003 | - | - | 0.005 | 0.005 | 0.005 | 0.034 | 0.005 | 0.005 | - |
| F 110 | - | - | - | - | - | - | - | 0.042 | 0.042 | 0.041 | 0.042 | 0.042 | 0.041 | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F93 | F94 | F95 | F96 | F97 | F98 | F99 | F100 | F101 | F102 | F103 | F104 | F105 | F106 |
|-------|-----|-------|-------|-------|-------|-------|-------|-------|-------|-------|------|-------|------|------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | - | - | 0.006 | 0.006 | 0.006 | 0.006 | - | 0.006 | 0.006 | 0.006 | - | 0.006 | - | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | 0.16 | - | - | - | - | 0.16 | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | 0.02 | - | - | - | - | - | - | - | - | - |
| F 106 | - | 0.041 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | 0.029 | - | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | 0.013 | 0.012 | 0.012 | 0.012 | 0.013 | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F107 | F108 | F109 | F110 | F111 | F112 | F113 | F114 | F115 | F116 | F117 | F118 | F119 | F120 |
|-------|------|------|------|-------|------|------|-------|------|------|------|------|-------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | 0.5 | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | - | - | - | 0.006 | - | - | - | - | - | - | - | 0.006 | 0.006 | 0.006 |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | 0.02 | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | 0.021 | - | - | - | - | - | 0.04 | - |
| F 106 | - | - | - | - | - | - | - | - | - | - | - | 0.041 | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F121 | F122 | F123 | F124 | F125 | F126 | F127 | F128 | F129 | F130 | F131 | F132 | F133 | F134 |
|-------|-------|-------|-------|-------|------|------|------|-------|-------|-------|-------|-------|------|------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | 0.5 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | 0.006 | 0.006 | 0.006 | - | - | - | 0.006 | 0.006 | - | 0.006 | 0.006 | - | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | 0.021 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 106 | - | - | - | 0.04 | - | - | - | - | - | - | - | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | 0.029 | - | 0.029 | - | - | - | 0.024 | 0.024 | 0.009 | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F135 | F136 | F137 | F138 | F139 | F140 | F141 | F142 | F143 | F144 | F145 | F146 | F147 | F148 |
|-------|------|------|------|------|-------|------|------|-------|------|------|-------|-------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | - | - | - | - | 0.006 | - | - | - | - | - | 0.006 | 0.006 | 0.006 | 0.006 |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 106 | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.04 |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | - | 0.029 | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F149 | F150 | F151 | F152 | F153 | F154 | F155 | F156 | F157 | F158 | F159 | F160 | F161 | F162 |
|-------|-------|-------|-------|-------|-------|------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | - | 0.006 | - | 0.006 | - | - | - | 0.006 | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | 0.02 | - | - |
| F 106 | - | 0.041 | - | - | - | - | - | 0.008 | 0.008 | 0.008 | 0.008 | 0.008 | 0.049 | 0.008 |
| F 107 | - | - | - | - | - | - | - | 0.051 | 0.051 | 0.051 | 0.051 | 0.051 | 0.017 | 0.017 |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | - | 0.009 | 0.009 | 0.009 | 0.009 | 0.009 | 0.009 | 0.009 |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F163 | F164 | F165 | F166 | F167 | F168 | F169 | F170 | F171 | F172 | F173 | F174 | F175 | F176 |
|-------|-------|-------|-------|-------|-------|-------|-------|------|------|------|-------|-------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | - | - | 0.006 | 0.006 | 0.006 | 0.006 | - | - | - | - | 0.006 | - | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | - | - | 0.02 |
| F 106 | 0.008 | 0.008 | 0.008 | - | - | - | - | - | - | - | 0.008 | 0.008 | 0.008 | 0.049 |
| F 107 | 0.017 | 0.017 | 0.017 | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | 0.009 | 0.009 | 0.036 | - | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F177 | F178 | F179 | F180 | F181 | F182 | F183 | F184 | F185 | F186 | F187 | F188 | F189 | F190 |
|-------|-------|-------|-------|-------|------|------|------|------|-------|-------|-------|-------|------|------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | 0.006 | - | 0.006 | 0.006 | - | - | - | - | 0.006 | 0.006 | 0.006 | 0.006 | - | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 106 | 0.008 | 0.005 | 0.005 | 0.005 | - | - | - | - | 0.005 | 0.005 | 0.005 | 0.005 | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | 0.014 | 0.014 | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F191 | F192 | F193 | F194 | F195 | F196 | F197 | F198 | F199 | F200 | F201 | F202 | F203 | F204 |
|-------|------|------|-------|------|------|------|------|-------|------|------|------|------|-------|-------|
| F 61 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 62 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 63 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 64 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 65 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 66 | - | - | 0.006 | - | - | - | - | 0.006 | - | - | - | - | - | - |
| F 67 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 68 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 69 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 70 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 71 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 72 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 73 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 74 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 75 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 76 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 77 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 78 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 79 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 80 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 81 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 82 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 83 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 84 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 85 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 86 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 87 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 88 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 89 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 90 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 91 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 92 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 93 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 94 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 95 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 96 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 97 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 98 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 99 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 100 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 101 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 102 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 103 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 104 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 105 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 106 | - | - | 0.005 | - | - | - | - | - | - | - | - | - | - | - |
| F 107 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 108 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 109 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 110 | - | - | - | - | - | - | - | - | - | - | - | - | 0.083 | 0.083 |
| F 111 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 112 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 113 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 114 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 115 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 116 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 117 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 118 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 119 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 120 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F205 | F206 | F207 | F208 | Sum |
|-------|-------|-------|-------|------|-----|
| F 61 | - | - | - | - | 1 |
| F 62 | - | - | - | - | 1 |
| F 63 | - | - | - | - | 1 |
| F 64 | - | - | - | - | 1 |
| F 65 | - | - | - | - | 1 |
| F 66 | 0.006 | 0.006 | 0.006 | - | 1 |
| F 67 | - | - | - | - | 1 |
| F 68 | - | - | - | - | 1 |
| F 69 | - | - | - | - | 1 |
| F 70 | - | - | - | - | 1 |
| F 71 | - | - | - | - | 1 |
| F 72 | - | - | - | - | 1 |
| F 73 | - | - | - | - | 1 |
| F 74 | - | - | - | - | 1 |
| F 75 | - | - | - | - | 1 |
| F 76 | - | - | - | - | 1 |
| F 77 | - | - | - | - | 1 |
| F 78 | - | - | - | - | 1 |
| F 79 | - | - | - | - | 1 |
| F 80 | - | - | - | - | 1 |
| F 81 | - | - | - | - | 1 |
| F 82 | - | - | - | - | 1 |
| F 83 | - | - | - | - | 1 |
| F 84 | - | - | - | - | 1 |
| F 85 | - | - | - | - | 1 |
| F 86 | - | - | - | - | 1 |
| F 87 | - | - | - | - | 1 |
| F 88 | - | - | - | - | 1 |
| F 89 | - | - | - | - | 1 |
| F 90 | - | - | - | - | 1 |
| F 91 | - | - | - | - | 1 |
| F 92 | - | - | - | - | 1 |
| F 93 | - | - | - | - | 1 |
| F 94 | - | - | - | - | 1 |
| F 95 | - | - | - | - | 1 |
| F 96 | - | - | - | - | 1 |
| F 97 | - | - | - | - | 1 |
| F 98 | - | - | - | - | 1 |
| F 99 | - | - | - | - | 1 |
| F 100 | - | - | - | - | 1 |
| F 101 | - | - | - | - | 1 |
| F 102 | - | - | - | - | 1 |
| F 103 | - | - | - | - | 1 |
| F 104 | - | - | - | - | 1 |
| F 105 | - | - | 0.02 | - | 1 |
| F 106 | - | - | - | - | 1 |
| F 107 | - | - | - | - | 1 |
| F 108 | - | - | - | - | 1 |
| F 109 | - | - | - | - | 1 |
| F 110 | 0.084 | - | - | - | 1 |
| F 111 | - | - | - | - | 1 |
| F 112 | - | - | - | - | 1 |
| F 113 | - | - | - | - | 1 |
| F 114 | - | - | - | - | 1 |
| F 115 | - | - | - | - | 1 |
| F 116 | - | - | - | - | 1 |
| F 117 | - | - | - | - | 1 |
| F 118 | - | - | - | - | 1 |
| F 119 | - | - | - | - | 1 |
| F 120 | - | - | - | - | 1 |

continued...

Table 8.6.2 continued

| | Predator/prey | D1 | A1 | A2 | A3 | I1 | I2 | I3 | I4 | I5 |
|-------|-------------------------------------|-------|-------|-------|----|-------|-------|-------|-------|-------|
| F 121 | <i>Mulloidichthys martinicus</i> | - | - | - | - | - | 0.161 | - | - | - |
| F 122 | <i>Pseudupeneus maculatus</i> | - | - | - | - | - | 0.007 | - | - | - |
| F 123 | <i>Echidna catenata</i> | - | - | - | - | - | - | - | - | - |
| F 124 | <i>Lycodontis moringa</i> | - | - | - | - | - | - | - | - | - |
| F 125 | <i>Lycodontis vicinus</i> | - | - | - | - | - | - | - | - | - |
| F 126 | <i>Aetobatus narinari</i> | - | - | - | - | - | - | - | - | - |
| F 127 | <i>Ogcocephalus nasutus</i> | - | 0.111 | - | - | - | - | - | - | - |
| F 128 | <i>Myrichthys acuminatus</i> | - | - | - | - | - | - | - | - | - |
| F 129 | <i>Myrichthys oculatus</i> | - | - | - | - | - | - | - | - | - |
| F 130 | <i>Ophichthus ophis</i> | - | - | - | - | - | - | - | - | - |
| F 131 | <i>Opisthognatus aurifrons</i> | - | - | - | - | - | 1 | - | - | - |
| F 132 | <i>Opisthognatus maxillosus</i> | - | - | - | - | - | 0.058 | - | - | - |
| F 133 | <i>Opisthognatus whitehurstii</i> | - | - | - | - | - | - | - | - | - |
| F 134 | <i>Acanthostracion polygonius</i> | - | - | 0.187 | - | - | - | 0.156 | - | 0.124 |
| F 135 | <i>Acanthostracion quadricornis</i> | - | 0.102 | 0.029 | - | - | 0.038 | 0.352 | - | 0.019 |
| F 136 | <i>Lactophrys trigonus</i> | - | 0.079 | - | - | - | 0.01 | - | - | - |
| F 137 | <i>Rhinesomus bicaudalis</i> | - | 0.15 | - | - | - | - | - | - | - |
| F 138 | <i>Rhinesomus triqueter</i> | - | 0.021 | - | - | - | 0.005 | 0.066 | - | - |
| F 139 | <i>Pempheris schomburgki</i> | - | - | - | - | - | 0.715 | - | - | - |
| F 140 | <i>Centropyge argi</i> | 0.5 | 0.5 | - | - | - | - | - | - | - |
| F 141 | <i>Holacanthus ciliaris</i> | - | 0.014 | 0.003 | - | - | - | 0.968 | 0.002 | - |
| F 142 | <i>Holacanthus tricolor</i> | - | 0.008 | - | - | - | - | 0.971 | - | - |
| F 143 | <i>Pomacanthus arcuatus</i> | - | 0.084 | 0.012 | - | - | 0.016 | 0.728 | 0.002 | 0.006 |
| F 144 | <i>Pomacanthus paru</i> | - | 0.135 | 0.01 | - | - | 0.001 | 0.748 | 0.001 | 0.006 |
| F 145 | <i>Abudefduf saxatilis</i> | - | 0.088 | - | - | - | 0.319 | - | - | - |
| F 146 | <i>Abudefduf taurus</i> | - | 0.94 | 0.001 | - | - | - | - | - | - |
| F 147 | <i>Chromis cyanea</i> | - | - | - | - | - | 1 | - | - | - |
| F 148 | <i>Chromis multilineata</i> | - | - | - | - | - | 1 | - | - | - |
| F 149 | <i>Microspathodon chrysurus</i> | 0.446 | 0.447 | 0.021 | - | - | 0.008 | 0.003 | - | - |
| F 150 | <i>Stegastes fuscus</i> | 0.281 | 0.3 | 0.001 | - | - | 0.114 | - | 0.001 | - |
| F 151 | <i>Stegastes leucostictus</i> | 0.113 | 0.113 | 0.052 | - | 0.02 | 0.283 | - | 0.027 | - |
| F 152 | <i>Stegastes planifrons</i> | 0.121 | 0.161 | 0.005 | - | - | 0.091 | - | - | - |
| F 153 | <i>Stegastes variabilis</i> | 0.258 | 0.257 | 0.034 | - | - | - | 0.021 | 0.014 | - |
| F 154 | <i>Priacanthus arenatus</i> | - | - | - | - | - | 0.267 | - | - | - |
| F 155 | <i>Priacanthus cruentatus</i> | - | - | - | - | - | 0.52 | - | - | - |
| F 156 | <i>Scarus coelestinus</i> | - | 0.986 | 0.001 | - | 0.002 | - | 0.001 | - | - |
| F 157 | <i>Scarus iserti</i> | - | 1 | - | - | - | - | - | - | - |
| F 158 | <i>Scarus guacamaia</i> | - | 1 | - | - | - | - | - | - | - |
| F 159 | <i>Scarus taeniopterus</i> | - | 0.985 | - | - | - | - | 0.015 | - | - |
| F 160 | <i>Scarus vetula</i> | - | 0.972 | 0.011 | - | - | - | 0.01 | - | 0.007 |
| F 161 | <i>Sparisoma aurofrenatum</i> | - | 0.991 | 0.005 | - | - | - | - | - | 0.003 |
| F 162 | <i>Sparisoma chrysopterum</i> | - | 1 | - | - | - | - | - | - | - |
| F 163 | <i>Sparisoma radians</i> | - | 1 | - | - | - | - | - | - | - |
| F 164 | <i>Sparisoma rubripinne</i> | - | 0.996 | - | - | - | - | 0.004 | - | - |
| F 165 | <i>Sparisoma viride</i> | - | 0.998 | 0.001 | - | - | - | 0.001 | - | - |
| F 166 | <i>Equetus lanceolatus</i> | - | - | - | - | - | - | - | - | - |
| F 167 | <i>Equetus punctatus</i> | - | - | - | - | - | 0.141 | - | - | - |
| F 168 | <i>Odontoscion dentex</i> | - | - | - | - | - | 0.391 | - | - | - |
| F 169 | <i>Pareques acuminatus</i> | - | - | - | - | - | 0.472 | - | - | - |
| F 170 | <i>Euthynnus alletteratus</i> | - | - | - | - | - | - | - | - | - |
| F 171 | <i>Scomberomorus cavalla</i> | - | - | - | - | - | - | - | - | - |
| F 172 | <i>Scomberomorus regalis</i> | - | - | - | - | - | - | - | - | - |
| F 173 | <i>Scorpaena brasiliensis</i> | - | - | - | - | - | - | - | - | - |
| F 174 | <i>Scorpaena grandicornis</i> | - | - | - | - | - | - | - | - | - |
| F 175 | <i>Scorpaena inermis</i> | - | - | - | - | - | - | - | - | - |
| F 176 | <i>Scorpaena plumieri</i> | - | - | - | - | - | - | - | - | - |
| F 177 | <i>Scorpaenodes caribbaeus</i> | - | - | - | - | - | - | - | - | - |
| F 178 | <i>Alphesthes afer</i> | - | - | - | - | - | - | - | - | - |
| F 179 | <i>Cephalopholis cruentata</i> | - | - | - | - | - | - | - | - | - |
| F 180 | <i>Cephalopholis fulva</i> | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I6 | I7 | I8 | I9 | I10 | I11 | I12 | I13 | I14 | I15 | I16 | I17 | I18 | I19 |
|-------|-------|-------|-------|-------|-----|-------|-------|-------|-------|-------|-------|-------|------|-----|
| F 121 | - | - | - | 0.069 | - | 0.08 | 0.007 | 0.133 | 0.004 | - | - | 0.19 | - | - |
| F 122 | - | - | - | 0.045 | - | - | 0.008 | 0.069 | - | - | - | 0.137 | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | 0.125 | - | - | - |
| F 126 | - | - | - | 0.02 | - | - | 0.3 | 0.45 | - | 0.02 | 0.02 | 0.02 | 0.02 | - |
| F 127 | - | - | - | - | - | - | 0.259 | 0.017 | - | - | - | 0.044 | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | 0.012 | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | 0.5 | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | 0.143 | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | 0.023 | - | - | - | - | - | - | - |
| F 135 | 0.124 | - | - | - | - | - | - | 0.004 | - | - | - | 0.021 | - | - |
| F 136 | - | - | - | - | - | 0.008 | 0.022 | 0.104 | - | - | - | 0.099 | - | - |
| F 137 | - | - | - | - | - | - | 0.011 | - | - | - | - | 0.065 | - | - |
| F 138 | - | - | - | 0.175 | - | 0.007 | 0.032 | 0.019 | - | - | - | 0.327 | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | 0.273 | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | 0.004 | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | 0.021 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | 0.046 | - | 0.001 | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | 0.049 | - | 0.002 | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | 0.45 | - | - | - | - | - | 0.055 | - | - | - | - | 0.01 | - | - |
| F 146 | 0.053 | - | - | - | - | 0.001 | 0.002 | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | 0.014 | - | - | - | - | 0.001 | - | - | - | - | - | - | - |
| F 150 | 0.038 | - | - | - | - | - | 0.042 | 0.006 | - | - | - | 0.071 | - | - |
| F 151 | - | 0.008 | - | - | - | - | 0.003 | - | - | - | - | 0.165 | - | - |
| F 152 | 0.303 | 0.004 | - | - | - | - | - | - | - | - | - | 0.088 | - | - |
| F 153 | - | 0.008 | - | - | - | - | 0.014 | 0.007 | - | - | - | 0.193 | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | 0.011 | 0.011 | 0.011 | - | - |
| F 155 | - | - | - | - | - | - | 0.013 | - | - | 0.022 | 0.022 | 0.17 | - | - |
| F 156 | - | 0.001 | - | - | - | 0.001 | 0.001 | 0.001 | 0.001 | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | 0.001 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | 0.313 | - | - |
| F 167 | - | - | - | - | - | - | 0.032 | 0.006 | - | - | - | 0.054 | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | 0.366 | - | 0.067 | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | 0.002 | - | - | - |
| F 172 | - | - | - | - | - | - | - | - | - | - | 0.023 | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | 0.062 | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | 0.025 | - | - | - |
| F 179 | - | - | - | - | - | - | 0.038 | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | I20 | I21 | I22 | I23 | I24 | I25 | I26 | I27 | I28 | I29 | I30 | I31 | I32 | I33 |
|-------|-------|-------|-------|-------|-------|-------|-----|------|-------|-------|-------|-------|-------|-------|
| F 121 | - | 0.008 | 0.045 | 0.009 | 0.058 | 0.061 | - | - | - | 0.081 | - | - | 0.09 | 0.004 |
| F 122 | - | 0.037 | 0.021 | 0.002 | 0.022 | 0.252 | - | - | - | 0.348 | - | - | 0.008 | - |
| F 123 | - | - | - | - | - | 0.037 | - | - | - | 0.963 | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | 0.25 | - | - | - | - |
| F 126 | - | - | - | - | - | 0.05 | - | - | - | - | - | - | - | - |
| F 127 | 0.006 | - | - | - | - | - | - | - | - | 0.475 | - | - | - | - |
| F 128 | - | 0.07 | - | - | - | - | - | - | - | 0.86 | - | - | - | 0.07 |
| F 129 | - | 0.155 | - | - | - | 0.04 | - | - | - | 0.761 | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | 0.286 | 0.286 | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | 0.11 | 0.54 | - | - | - | 0.05 | - | - | 0.08 | - |
| F 134 | - | - | - | - | - | 0.133 | - | - | - | - | - | - | - | - |
| F 135 | - | - | 0.003 | - | - | 0.003 | - | - | 0.095 | - | - | - | - | - |
| F 136 | - | - | 0.019 | - | - | 0.003 | - | - | 0.004 | 0.464 | - | 0.025 | 0.016 | 0.073 |
| F 137 | - | - | - | - | - | - | - | - | - | 0.044 | - | 0.065 | 0.136 | 0.109 |
| F 138 | - | - | 0.013 | - | - | 0.078 | - | - | 0.025 | 0.098 | 0.036 | - | - | 0.025 |
| F 139 | - | - | 0.012 | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | 0.031 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | 0.003 | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | 0.003 | - | - | 0.041 | - | - | - | - | - | - | 0.016 | - |
| F 150 | - | - | 0.016 | - | 0.021 | 0.012 | - | - | 0.004 | 0.029 | - | - | - | - |
| F 151 | - | - | 0.021 | - | - | 0.013 | - | - | 0.013 | 0.038 | - | - | - | - |
| F 152 | - | - | 0.035 | - | 0.035 | 0.035 | - | - | 0.035 | 0.035 | - | 0.013 | 0.013 | 0.013 |
| F 153 | - | - | 0.011 | 0.009 | 0.15 | 0.014 | - | - | - | - | - | - | 0.01 | - |
| F 154 | - | - | - | - | 0.016 | 0.347 | - | - | - | 0.049 | - | - | - | - |
| F 155 | - | - | 0.007 | - | 0.05 | 0.05 | - | - | - | 0.073 | - | - | - | - |
| F 156 | - | - | 0.001 | - | 0.001 | 0.001 | - | - | - | 0.001 | - | - | - | 0.001 |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | 0.625 | - | - | - | 0.062 | - | - | - | - |
| F 167 | - | 0.004 | - | - | 0.013 | 0.137 | - | - | 0.15 | 0.459 | - | - | - | - |
| F 168 | - | - | - | - | 0.178 | 0.19 | - | - | - | 0.052 | - | - | - | - |
| F 169 | - | - | 0.003 | - | 0.035 | 0.411 | - | - | - | 0.06 | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | 0.92 | - | 0.01 | - | - | - | - | - | - |
| F 172 | - | - | - | - | - | 0.016 | - | - | - | - | - | - | - | - |
| F 173 | - | 0.215 | - | - | - | 0.535 | - | - | - | 0.107 | - | - | - | - |
| F 174 | - | - | - | - | - | 0.875 | - | - | - | - | - | - | - | - |
| F 175 | - | 0.08 | - | - | - | 0.91 | - | - | - | 0.007 | - | - | - | - |
| F 176 | - | - | - | - | - | 0.187 | - | - | 0.016 | 0.313 | - | - | - | - |
| F 177 | - | - | - | - | - | 0.731 | - | - | - | 0.193 | - | - | - | - |
| F 178 | - | - | - | - | - | 0.073 | - | - | - | 0.832 | - | - | - | - |
| F 179 | - | 0.089 | - | - | - | 0.173 | - | - | - | 0.038 | - | - | - | - |
| F 180 | - | 0.133 | - | - | - | 0.222 | - | - | - | 0.185 | - | - | - | - |

continued...

Table 8.6.2 continued

| | I34 | I35 | R1 | B1 |
|-------|-------|-------|----|----|
| F 121 | - | - | - | - |
| F 122 | - | - | - | - |
| F 123 | - | - | - | - |
| F 124 | - | - | - | - |
| F 125 | - | - | - | - |
| F 126 | - | - | - | - |
| F 127 | - | - | - | - |
| F 128 | - | - | - | - |
| F 129 | - | - | - | - |
| F 130 | - | - | - | - |
| F 131 | - | - | - | - |
| F 132 | - | - | - | - |
| F 133 | - | - | - | - |
| F 134 | - | 0.377 | - | - |
| F 135 | - | 0.21 | - | - |
| F 136 | 0.027 | 0.047 | - | - |
| F 137 | 0.207 | 0.213 | - | - |
| F 138 | - | 0.073 | - | - |
| F 139 | - | - | - | - |
| F 140 | - | - | - | - |
| F 141 | - | 0.009 | - | - |
| F 142 | - | - | - | - |
| F 143 | - | 0.105 | - | - |
| F 144 | - | 0.048 | - | - |
| F 145 | - | - | - | - |
| F 146 | - | - | - | - |
| F 147 | - | - | - | - |
| F 148 | - | - | - | - |
| F 149 | - | - | - | - |
| F 150 | - | 0.002 | - | - |
| F 151 | - | 0.038 | - | - |
| F 152 | 0.013 | - | - | - |
| F 153 | - | - | - | - |
| F 154 | - | - | - | - |
| F 155 | - | - | - | - |
| F 156 | - | - | - | - |
| F 157 | - | - | - | - |
| F 158 | - | - | - | - |
| F 159 | - | - | - | - |
| F 160 | - | - | - | - |
| F 161 | - | - | - | - |
| F 162 | - | - | - | - |
| F 163 | - | - | - | - |
| F 164 | - | - | - | - |
| F 165 | - | - | - | - |
| F 166 | - | - | - | - |
| F 167 | - | - | - | - |
| F 168 | - | - | - | - |
| F 169 | - | - | - | - |
| F 170 | - | - | - | - |
| F 171 | - | - | - | - |
| F 172 | - | - | - | - |
| F 173 | - | - | - | - |
| F 174 | - | - | - | - |
| F 175 | - | - | - | - |
| F 176 | - | - | - | - |
| F 177 | - | - | - | - |
| F 178 | - | - | - | - |
| F 179 | - | - | - | - |
| F 180 | - | - | - | - |

continued...

Table 8.6.2 continued

| | Predator/Prey | F0 | F1 | F2 | F3 | F4 | F5 | F6 | F7 | F8 |
|-------|-------------------------------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 121 | <i>Mulloidichthys martinicus</i> | - | - | - | - | - | - | - | - | - |
| F 122 | <i>Pseudupeneus maculatus</i> | 0.022 | - | - | - | - | - | - | - | - |
| F 123 | <i>Echidna catenata</i> | - | - | - | - | - | - | - | - | - |
| F 124 | <i>Lycodontis moringa</i> | - | - | - | - | - | - | - | - | - |
| F 125 | <i>Lycodontis vicinus</i> | - | - | - | - | - | - | - | - | - |
| F 126 | <i>Aetobatus narinari</i> | - | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 |
| F 127 | <i>Ogcocephalus nasutus</i> | - | - | - | - | - | - | - | - | - |
| F 128 | <i>Myrichthys acuminatus</i> | - | - | - | - | - | - | - | - | - |
| F 129 | <i>Myrichthys ocellatus</i> | 0.032 | - | - | - | - | - | - | - | - |
| F 130 | <i>Ophichthus ophis</i> | - | - | - | - | - | - | - | - | - |
| F 131 | <i>Opisthognathus aurifrons</i> | - | - | - | - | - | - | - | - | - |
| F 132 | <i>Opisthognathus maxillosus</i> | 0.227 | - | - | - | - | - | - | - | - |
| F 133 | <i>Opisthognathus whitehurstii</i> | 0.22 | - | - | - | - | - | - | - | - |
| F 134 | <i>Acanthostracion polygonius</i> | - | - | - | - | - | - | - | - | - |
| F 135 | <i>Acanthostracion quadricornis</i> | - | - | - | - | - | - | - | - | - |
| F 136 | <i>Lactophrys trigonus</i> | - | - | - | - | - | - | - | - | - |
| F 137 | <i>Rhinesomus bicaudalis</i> | - | - | - | - | - | - | - | - | - |
| F 138 | <i>Rhinesomus triquetter</i> | - | - | - | - | - | - | - | - | - |
| F 139 | <i>Pempheris schomburgki</i> | - | - | - | - | - | - | - | - | - |
| F 140 | <i>Centropyge argi</i> | - | - | - | - | - | - | - | - | - |
| F 141 | <i>Holacanthus ciliaris</i> | - | - | - | - | - | - | - | - | - |
| F 142 | <i>Holacanthus tricolor</i> | - | - | - | - | - | - | - | - | - |
| F 143 | <i>Pomacanthus arcuatus</i> | - | - | - | - | - | - | - | - | - |
| F 144 | <i>Pomacanthus paru</i> | - | - | - | - | - | - | - | - | - |
| F 145 | <i>Abudefduf saxatilis</i> | - | - | - | - | - | - | - | - | - |
| F 146 | <i>Abudefduf taurus</i> | - | - | - | - | - | - | - | - | - |
| F 147 | <i>Chromis cyanea</i> | - | - | - | - | - | - | - | - | - |
| F 148 | <i>Chromis multilineata</i> | - | - | - | - | - | - | - | - | - |
| F 149 | <i>Microspathodon chrysurus</i> | - | - | - | - | - | - | - | - | - |
| F 150 | <i>Pomacentrus fuscus</i> | 0.062 | - | - | - | - | - | - | - | - |
| F 151 | <i>Pomacentrus leucostictus</i> | 0.093 | - | - | - | - | - | - | - | - |
| F 152 | <i>Pomacentrus planifrons</i> | - | - | - | - | - | - | - | - | - |
| F 153 | <i>Pomacentrus variabilis</i> | - | - | - | - | - | - | - | - | - |
| F 154 | <i>Priacanthus arenatus</i> | - | - | - | - | - | - | - | - | 0.094 |
| F 155 | <i>Priacanthus cruentatus</i> | - | - | - | - | - | - | - | - | - |
| F 156 | <i>Scarus coelestinus</i> | - | - | - | - | - | - | - | - | - |
| F 157 | <i>Scarus croicensis</i> | - | - | - | - | - | - | - | - | - |
| F 158 | <i>Scarus guacamaia</i> | - | - | - | - | - | - | - | - | - |
| F 159 | <i>Scarus taeniopterus</i> | - | - | - | - | - | - | - | - | - |
| F 160 | <i>Scarus vetula</i> | - | - | - | - | - | - | - | - | - |
| F 161 | <i>Sparisoma aurofrenatum</i> | - | - | - | - | - | - | - | - | 0.003 |
| F 162 | <i>Sparisoma chrysopterum</i> | - | - | - | - | - | - | - | - | - |
| F 163 | <i>Sparisoma radians</i> | - | - | - | - | - | - | - | - | - |
| F 164 | <i>Sparisoma rubripinne</i> | - | - | - | - | - | - | - | - | - |
| F 165 | <i>Sparisoma viride</i> | - | - | - | - | - | - | - | - | - |
| F 166 | <i>Equetus lanceolatus</i> | - | - | - | - | - | - | - | - | - |
| F 167 | <i>Equetus punctatus</i> | 0.004 | - | - | - | - | - | - | - | - |
| F 168 | <i>Odontoscion dentex</i> | 0.189 | - | - | - | - | - | - | - | - |
| F 169 | <i>Pareques acuminatus</i> | 0.019 | - | - | - | - | - | - | - | - |
| F 170 | <i>Euthynnus alletteratus</i> | - | - | - | - | - | - | - | - | 0.142 |
| F 171 | <i>Scomberomorus cavalla</i> | - | - | - | - | - | - | - | - | - |
| F 172 | <i>Scomberomorus regalis</i> | - | - | - | - | - | - | - | - | 0.096 |
| F 173 | <i>Scorpaena brasiliensis</i> | 0.143 | - | - | - | - | - | - | - | - |
| F 174 | <i>Scorpaena grandicornis</i> | 0.125 | - | - | - | - | - | - | - | - |
| F 175 | <i>Scorpaena inermis</i> | 0.003 | - | - | - | - | - | - | - | - |
| F 176 | <i>Scorpaena plumieri</i> | - | 0.047 | 0.047 | 0.047 | - | - | - | - | - |
| F 177 | <i>Scorpaenodes caribbaeus</i> | - | - | - | - | - | - | - | - | - |
| F 178 | <i>Alphestes afer</i> | - | 0.012 | 0.012 | 0.012 | - | - | - | - | - |
| F 179 | <i>Cephalopholis cruentata</i> | - | - | - | - | - | - | 0.047 | 0.047 | - |
| F 180 | <i>Cephalopholis fulva</i> | - | 0.022 | 0.022 | 0.022 | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F9 | F10 | F11 | F12 | F13 | F14 | F15 | F16 | F17 | F18 | F19 | F20 | F21 | F22 |
|-------|-------|-------|-----|-----|-------|-------|-------|-------|-------|-----|-------|-----|-------|-----|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | 0.001 | - | - | 0.001 | 0.001 | 0.001 | - | - | - | 0.001 | - | 0.001 | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | 0.094 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | 0.047 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 172 | 0.032 | - | - | - | - | 0.016 | 0.016 | 0.016 | 0.016 | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | 0.065 | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F23 | F24 | F25 | F26 | F27 | F28 | F29 | F30 | F31 | F32 | F33 | F34 | F35 | F36 |
|-------|-------|-----|-----|-------|-------|-------|-------|-----|-----|-----|-----|-----|-----|-----|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | - | - | - | 0.001 | 0.001 | - | - | - | - | - | - | - | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 172 | - | - | - | 0.064 | - | - | 0.064 | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F37 | F38 | F39 | F40 | F41 | F42 | F43 | F44 | F45 | F46 | F47 | F48 | F49 | F50 |
|-------|-----|-----|-----|-----|-----|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | - | - | - | - | - | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | - | 0.001 | 0.001 | 0.001 |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | 0.023 | 0.023 |
| F 171 | - | - | - | - | - | - | - | - | - | - | - | - | 0.017 | 0.016 |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | 0.096 | 0.032 |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | 0.024 | 0.024 | 0.024 | 0.024 | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | 0.033 | 0.033 | - | - | - |

continued...

Table 8.6.2 continued

| | F51 | F52 | F53 | F54 | F55 | F56 | F57 | F58 | F59 | F60 | F61 | F62 | F63 | F64 |
|-------|-------|-------|-------|-----|-----|-----|-----|-----|-------|-------|-------|-----|-----|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | 0.001 | 0.001 | - | - | - | - | - | 0.001 | 0.001 | - | - | - | 0.001 |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | 0.047 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | 0.073 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | 0.12 | 0.023 | - | - | - | - | - | - | - | 0.047 | 0.047 | - | - | - |
| F 171 | 0.017 | 0.016 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 172 | 0.064 | 0.064 | - | - | - | - | - | - | - | 0.032 | 0.032 | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | 0.141 | - | 0.14 | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | 0.034 | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F65 | F66 | F67 | F68 | F69 | F70 | F71 | F72 | F73 | F74 | F75 | F76 | F77 | F78 |
|-------|-------|-----|-------|-----|-----|-------|-------|-------|-----|-------|-------|-------|-------|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | 0.022 | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | 0.5 | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | - | - | - | - | 0.001 | 0.001 | 0.001 | - | 0.001 | - | 0.001 | 0.001 | 0.001 |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | 0.5 | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | 0.001 | 0.001 | 0.001 | 0.001 |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | 0.189 | - | - | - | - | - | - | - | 0.011 | 0.011 | 0.011 | 0.011 |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F79 | F80 | F81 | F82 | F83 | F84 | F85 | F86 | F87 | F88 | F89 | F90 | F91 | F92 |
|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-----|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | - | - | - | - | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | 0.162 | 0.362 | - | - | - | - | - | - | - |
| F 172 | - | - | - | - | - | - | 0.064 | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | 0.011 | 0.011 | 0.011 | 0.011 | 0.011 | - | - | - | 0.092 | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F93 | F94 | F95 | F96 | F97 | F98 | F99 | F100 | F101 | F102 | F103 | F104 | F105 | F106 |
|-------|-----|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|------|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | - | - | 0.001 | 0.001 | 0.001 | 0.001 | - | 0.001 | 0.001 | 0.001 | - | 0.001 | - | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | - | - | 0.02 | 0.021 |
| F 172 | - | 0.006 | 0.069 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F107 | F108 | F109 | F110 | F111 | F112 | F113 | F114 | F115 | F116 | F117 | F118 | F119 | F120 |
|-------|------|-------|------|-------|------|-------|------|------|------|------|------|-------|-------|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | 0.5 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | - | - | - | 0.001 | - | - | - | - | - | - | - | 0.001 | 0.001 | 0.001 |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | 0.02 | 0.021 | 0.02 | 0.021 | 0.02 | 0.021 | - | - | - | - | - | - | - | 0.049 |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | 0.065 | 0.065 | - |

continued...

Table 8.6.2 continued

| | F121 | F122 | F123 | F124 | F125 | F126 | F127 | F128 | F129 | F130 | F131 | F132 | F133 | F134 |
|-------|-------|-------|-------|-------|------|------|------|-------|-------|------|-------|-------|------|------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | 0.001 | 0.001 | 0.001 | - | - | - | 0.001 | 0.001 | - | 0.001 | 0.001 | - | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | 0.095 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F135 | F136 | F137 | F138 | F139 | F140 | F141 | F142 | F143 | F144 | F145 | F146 | F147 | F148 |
|-------|------|------|------|------|-------|------|-------|-------|------|------|-------|-------|-------|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | - | - | - | - | 0.001 | - | - | - | - | - | 0.001 | 0.001 | 0.001 | 0.001 |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | 0.064 | 0.064 |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | 0.038 | 0.038 | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | 0.092 | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F149 | F150 | F151 | F152 | F153 | F154 | F155 | F156 | F157 | F158 | F159 | F160 | F161 | F162 |
|-------|-------|-------|-------|-------|-------|------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | 0.125 | 0.125 | 0.125 | 0.125 | 0.125 | - | - |
| F 126 | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 | - | 0.001 | - | 0.001 | - | - | - | 0.001 | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 | 0.007 |

continued...

Table 8.6.2 continued

| | F163 | F164 | F165 | F166 | F167 | F168 | F169 | F170 | F171 | F172 | F173 | F174 | F175 | F176 |
|-------|-------|-------|-------|-------|-------|-------|-------|------|-------|-------|------|-------|------|------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | - | - | 0.001 | 0.001 | 0.001 | 0.001 | - | - | - | - | 0.001 | - | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | - | - | - | - | - | - | 0.04 | 0.008 | 0.008 | - | - | - | - |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | 0.007 | 0.007 | 0.007 | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F177 | F178 | F179 | F180 | F181 | F182 | F183 | F184 | F185 | F186 | F187 | F188 | F189 | F190 |
|-------|-------|-------|-------|-------|-------|-------|------|-------|-------|-------|-------|-------|------|-------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | 0.001 | - | 0.001 | 0.001 | - | - | - | - | 0.001 | 0.001 | 0.001 | 0.001 | - | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | - | 0.001 | 0.001 | 0.001 | 0.007 | 0.007 | - | 0.007 | - | - | - | - | - | 0.001 |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F191 | F192 | F193 | F194 | F195 | F196 | F197 | F198 | F199 | F200 | F201 | F202 | F203 | F204 |
|-------|-------|-------|-------|------|------|------|------|-------|------|------|------|------|------|------|
| F 121 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 122 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 123 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 124 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 125 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 126 | - | - | 0.001 | - | - | - | - | 0.001 | - | - | - | - | - | - |
| F 127 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 128 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 129 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 130 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 131 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 132 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 133 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 134 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 135 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 136 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 137 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 138 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 139 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 140 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 141 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 142 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 143 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 144 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 145 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 146 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 147 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 148 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 149 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 150 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 151 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 152 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 153 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 154 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 155 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 156 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 157 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 158 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 159 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 160 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 161 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 162 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 163 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 164 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 165 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 166 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 167 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 168 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 169 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 170 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 171 | 0.001 | 0.001 | 0.001 | - | - | - | - | - | - | - | - | - | - | - |
| F 172 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 173 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 174 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 175 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 176 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 177 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 178 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 179 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 180 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |

continued...

Table 8.6.2 continued

| | F205 | F206 | F207 | F208 | Sum |
|-------|-------|-------|-------|------|-----|
| F 121 | - | - | - | - | 1 |
| F 122 | - | - | - | - | 1 |
| F 123 | - | - | - | - | 1 |
| F 124 | - | - | - | - | 1 |
| F 125 | - | - | - | - | 1 |
| F 126 | 0.001 | 0.001 | 0.001 | - | 1 |
| F 127 | - | - | 0.088 | - | 1 |
| F 128 | - | - | - | - | 1 |
| F 129 | - | - | - | - | 1 |
| F 130 | - | - | - | - | 1 |
| F 131 | - | - | - | - | 1 |
| F 132 | - | - | - | - | 1 |
| F 133 | - | - | - | - | 1 |
| F 134 | - | - | - | - | 1 |
| F 135 | - | - | - | - | 1 |
| F 136 | - | - | - | - | 1 |
| F 137 | - | - | - | - | 1 |
| F 138 | - | - | - | - | 1 |
| F 139 | - | - | - | - | 1 |
| F 140 | - | - | - | - | 1 |
| F 141 | - | - | - | - | 1 |
| F 142 | - | - | - | - | 1 |
| F 143 | - | - | - | - | 1 |
| F 144 | - | - | - | - | 1 |
| F 145 | - | - | - | - | 1 |
| F 146 | - | - | - | - | 1 |
| F 147 | - | - | - | - | 1 |
| F 148 | - | - | - | - | 1 |
| F 149 | - | - | - | - | 1 |
| F 150 | - | - | - | - | 1 |
| F 151 | - | - | - | - | 1 |
| F 152 | - | - | - | - | 1 |
| F 153 | - | - | - | - | 1 |
| F 154 | - | - | - | - | 1 |
| F 155 | - | - | - | - | 1 |
| F 156 | - | - | - | - | 1 |
| F 157 | - | - | - | - | 1 |
| F 158 | - | - | - | - | 1 |
| F 159 | - | - | - | - | 1 |
| F 160 | - | - | - | - | 1 |
| F 161 | - | - | - | - | 1 |
| F 162 | - | - | - | - | 1 |
| F 163 | - | - | - | - | 1 |
| F 164 | - | - | - | - | 1 |
| F 165 | - | - | - | - | 1 |
| F 166 | - | - | - | - | 1 |
| F 167 | - | - | - | - | 1 |
| F 168 | - | - | - | - | 1 |
| F 169 | - | - | - | - | 1 |
| F 170 | - | - | - | - | 1 |
| F 171 | - | - | - | - | 1 |
| F 172 | - | - | - | - | 1 |
| F 173 | - | - | - | - | 1 |
| F 174 | - | - | - | - | 1 |
| F 175 | - | - | - | - | 1 |
| F 176 | - | - | - | - | 1 |
| F 177 | - | - | - | - | 1 |
| F 178 | - | - | - | - | 1 |
| F 179 | - | - | - | - | 1 |
| F 180 | - | - | 0.063 | - | 1 |

continued...

Table 8.6.2 continued

| Predator/prey | D1 | A1 | A2 | A3 | I1 | I2 | I3 | I4 | I5 |
|---------------------------------------|-------|-------|-------|----|----|-------|-------|-------|----|
| F 181 <i>Epinephelus adscensionis</i> | - | - | - | - | - | - | - | - | - |
| F 182 <i>Epinephelus guttatus</i> | - | - | - | - | - | - | - | - | - |
| F 183 <i>Epinephelus itajara</i> | - | - | - | - | - | - | - | - | - |
| F 184 <i>Epinephelus striatus</i> | - | - | - | - | - | - | - | - | - |
| F 185 <i>Hypoplectrus aberrans</i> | - | - | - | - | - | 0.008 | - | - | - |
| F 186 <i>Hypoplectrus chlorurus</i> | - | - | - | - | - | - | - | - | - |
| F 187 <i>Hypoplectrus nigricans</i> | - | - | - | - | - | 0.059 | - | - | - |
| F 188 <i>Hypoplectrus puella</i> | - | - | - | - | - | 0.089 | - | - | - |
| F 189 <i>Mycteroperca bonaci</i> | - | - | - | - | - | - | - | - | - |
| F 190 <i>Mycteroperca tigris</i> | - | - | - | - | - | - | - | - | - |
| F 191 <i>Mycteroperca venenosa</i> | - | - | - | - | - | - | - | - | - |
| F 192 <i>Paranthias furcifer</i> | - | - | - | - | - | 0.902 | - | - | - |
| F 193 <i>Serranus tigrinus</i> | - | - | - | - | - | - | - | - | - |
| F 194 <i>Archosargus rhomboidalis</i> | - | 0.834 | - | - | - | 0.035 | - | - | - |
| F 195 <i>Calamus bajonado</i> | - | - | - | - | - | - | - | - | - |
| F 196 <i>Calamus calamus</i> | - | - | - | - | - | - | - | - | - |
| F 197 <i>Calamus pennatula</i> | - | - | - | - | - | - | - | - | - |
| F 198 <i>Diplodus caudimacula</i> | - | 0.8 | - | - | - | - | - | - | - |
| F 199 <i>Sphyræna barracuda</i> | - | - | - | - | - | 0.019 | - | - | - |
| F 200 <i>Sphyræna picudilla</i> | - | - | - | - | - | - | - | - | - |
| F 201 <i>Sphyræna lewini</i> | - | - | - | - | - | - | - | - | - |
| F 202 <i>Sphyræna tiburo</i> | - | - | - | - | - | - | - | - | - |
| F 203 <i>Synodus foetens</i> | - | - | - | - | - | - | - | - | - |
| F 204 <i>Synodus intermedius</i> | - | - | - | - | - | - | - | - | - |
| F 205 <i>Synodus synodus</i> | - | - | - | - | - | - | - | - | - |
| F 206 <i>Canthigaster rostrata</i> | 0.011 | 0.172 | 0.018 | - | - | 0.013 | 0.167 | 0.012 | - |
| F 207 <i>Sphoeroides spengleri</i> | 0.018 | 0.07 | - | - | - | 0.031 | - | - | - |
| F 208 <i>Mustelus canis</i> | - | - | - | - | - | - | - | - | - |

continued...

| | I6 | I7 | I8 | I9 | I10 | I11 | I12 | I13 | I14 | I15 | I16 | I17 | I18 | I19 |
|-------|----|----|----|-------|-----|-------|-------|-------|------|-------|-------|-------|------|-------|
| F 181 | - | - | - | - | - | 0.016 | 0.032 | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | 0.07 | - | 0.02 | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | - | - | - | - | - | 0.016 | 0.007 | - | 0.026 | 0.026 | - | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | - | - | - | - | - | - | 0.039 | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | 0.043 | 0.034 | - | - | - | 0.004 | - | - |
| F 195 | - | - | - | - | - | - | 0.111 | 0.135 | - | - | - | 0.067 | - | - |
| F 196 | - | - | - | 0.006 | - | 0.019 | 0.084 | 0.151 | - | - | - | 0.194 | - | - |
| F 197 | - | - | - | 0.05 | - | - | 0.082 | 0.123 | - | - | - | 0.14 | - | - |
| F 198 | - | - | - | - | - | 0.02 | 0.16 | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | - | 0.026 | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | 0.179 | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | - | 0.1 | - | - | - | - |
| F 202 | - | - | - | - | - | - | - | 0.05 | - | - | 0.25 | - | - | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | 0.05 | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | 0.087 | 0.058 | - | - | - | 0.131 | - | 0.013 |
| F 207 | - | - | - | - | - | 0.015 | 0.098 | 0.164 | - | - | - | 0.078 | - | - |
| F 208 | - | - | - | - | - | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | - | - | - |

continued...

Table 8.6.2 continued

| | I20 | I21 | I22 | I23 | I24 | I25 | I26 | I27 | I28 | I29 | I30 | I31 | I32 | I33 |
|-------|-----|-------|-------|-----|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 181 | - | - | - | - | - | 0.046 | - | - | - | 0.705 | - | - | - | - |
| F 182 | - | 0.174 | - | - | - | 0.11 | - | - | - | 0.415 | - | - | - | - |
| F 183 | - | - | - | - | - | - | 0.456 | 0.233 | - | 0.122 | - | - | - | - |
| F 184 | - | 0.056 | - | - | 0.002 | 0.051 | 0.035 | - | 0.012 | 0.229 | - | - | - | - |
| F 185 | - | 0.08 | - | - | - | 0.556 | - | - | - | 0.236 | - | - | - | - |
| F 186 | - | - | - | - | - | 0.562 | - | - | - | 0.19 | - | - | - | - |
| F 187 | - | 0.029 | - | - | - | 0.294 | - | - | - | 0.178 | - | - | - | - |
| F 188 | - | 0.053 | - | - | 0.037 | 0.51 | - | - | - | 0.211 | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | - | - | 0.008 | - | - | - | - | - | - | - | - |
| F 192 | - | - | 0.034 | - | - | 0.064 | - | - | - | - | - | - | - | - |
| F 193 | - | 0.091 | - | - | - | 0.733 | - | - | - | 0.079 | - | - | - | - |
| F 194 | - | - | 0.002 | - | - | - | - | - | - | 0.048 | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | 0.011 | 0.223 | - | - | - | 0.453 |
| F 196 | - | - | - | - | - | - | - | - | 0.153 | 0.147 | - | - | 0.156 | 0.09 |
| F 197 | - | 0.03 | - | - | - | 0.058 | - | - | 0.08 | 0.215 | - | - | 0.142 | 0.04 |
| F 198 | - | - | - | - | - | - | - | - | - | 0.02 | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | - | 0.1 | - | - | - | - |
| F 202 | - | - | - | - | - | 0.25 | - | - | - | 0.25 | - | - | - | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | 0.005 | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | 0.039 | - | - | 0.037 | - | - | 0.029 | 0.122 | - | 0.042 | - | 0.042 |
| F 207 | - | - | 0.058 | - | 0.011 | 0.056 | - | - | 0.009 | 0.209 | 0.035 | - | 0.061 | 0.071 |
| F 208 | - | - | - | - | - | - | 0.1 | 0.1 | - | 0.2 | - | - | - | - |

continued...

| | I34 | I35 | R1 | B1 |
|-------|------|-------|-------|----|
| F 181 | - | - | - | - |
| F 182 | - | - | - | - |
| F 183 | - | - | 0.056 | - |
| F 184 | - | - | - | - |
| F 185 | - | - | - | - |
| F 186 | - | - | - | - |
| F 187 | - | - | - | - |
| F 188 | - | - | - | - |
| F 189 | - | - | - | - |
| F 190 | - | - | - | - |
| F 191 | - | - | - | - |
| F 192 | - | - | - | - |
| F 193 | - | - | - | - |
| F 194 | - | - | - | - |
| F 195 | - | - | - | - |
| F 196 | - | - | - | - |
| F 197 | 0.04 | - | - | - |
| F 198 | - | - | - | - |
| F 199 | - | - | - | - |
| F 200 | - | - | - | - |
| F 201 | - | - | - | - |
| F 202 | - | - | - | - |
| F 203 | - | - | - | - |
| F 204 | - | - | - | - |
| F 205 | - | - | - | - |
| F 206 | - | 0.007 | - | - |
| F 207 | - | 0.009 | - | - |
| F 208 | - | - | - | - |

continued...

Table 8.6.2. continued.

| Predator/prey | F0 | F1 | F2 | F3 | F4 | F5 | F6 | F7 | F8 |
|---------------------------------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 181 <i>Epinephelus adscensionis</i> | - | - | - | - | - | - | - | - | - |
| F 182 <i>Epinephelus guttatus</i> | - | - | - | - | - | - | - | - | - |
| F 183 <i>Epinephelus itajara</i> | - | - | - | - | - | - | - | - | - |
| F 184 <i>Epinephelus striatus</i> | - | 0.005 | 0.005 | 0.005 | - | - | - | - | 0.007 |
| F 185 <i>Hypoplectrus aberrans</i> | - | - | - | - | - | - | - | - | - |
| F 186 <i>Hypoplectrus chlorurus</i> | - | - | - | - | - | - | - | - | - |
| F 187 <i>Hypoplectrus nigricans</i> | - | - | - | - | - | - | - | - | - |
| F 188 <i>Hypoplectrus puella</i> | - | - | - | - | - | - | - | - | - |
| F 189 <i>Mycteroperca bonaci</i> | - | - | - | - | - | - | - | - | - |
| F 190 <i>Mycteroperca tigris</i> | - | 0.095 | 0.024 | 0.095 | - | - | - | - | 0.036 |
| F 191 <i>Mycteroperca venenosa</i> | - | 0.048 | - | - | - | - | - | - | 0.024 |
| F 192 <i>Paranthias furcifer</i> | - | - | - | - | - | - | - | - | - |
| F 193 <i>Serranus tigrinus</i> | 0.097 | - | - | - | - | - | - | - | - |
| F 194 <i>Archosargus rhomboidalis</i> | - | - | - | - | - | - | - | - | - |
| F 195 <i>Calamus bajonado</i> | - | - | - | - | - | - | - | - | - |
| F 196 <i>Calamus calamus</i> | - | - | - | - | - | - | - | - | - |
| F 197 <i>Calamus pennatula</i> | - | - | - | - | - | - | - | - | - |
| F 198 <i>Diplodus caudimacula</i> | - | - | - | - | - | - | - | - | - |
| F 199 <i>Sphyrna barracuda</i> | - | 0.053 | - | - | - | - | - | - | - |
| F 200 <i>Sphyrna picudilla</i> | 0.821 | - | - | - | - | - | - | - | 0.08 |
| F 201 <i>Sphyrna lewini</i> | - | - | - | - | - | - | - | - | 0.02 |
| F 202 <i>Sphyrna tiburo</i> | - | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 |
| F 203 <i>Synodus foetens</i> | - | - | - | - | - | - | - | - | - |
| F 204 <i>Synodus intermedius</i> | - | - | - | - | - | - | - | - | 0.079 |
| F 205 <i>Synodus synodus</i> | 1 | - | - | - | - | - | - | - | - |
| F 206 <i>Canthigaster rostrata</i> | - | - | - | - | - | - | - | - | - |
| F 207 <i>Sphoeroides spengleri</i> | 0.007 | - | - | - | - | - | - | - | - |
| F 208 <i>Mustelus canis</i> | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 |

continued...

| | F9 | F10 | F11 | F12 | F13 | F14 | F15 | F16 | F17 | F18 | F19 | F20 | F21 | F22 |
|-------|-------|-------|-----|-----|-------|-------|-------|-------|-------|-------|-------|-------|-------|-----|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | 0.037 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | 0.007 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | 0.03 | 0.03 | 0.03 | 0.03 | - |
| F 186 | - | - | - | - | - | - | - | - | - | 0.062 | 0.062 | 0.062 | 0.062 | - |
| F 187 | - | - | - | - | - | - | - | - | - | 0.11 | 0.11 | 0.11 | 0.11 | - |
| F 188 | - | - | - | - | - | - | - | - | - | 0.025 | 0.025 | 0.025 | 0.025 | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | 0.036 | - | - | - | - | - | - | - | - | - | 0.072 | - | - | - |
| F 191 | 0.024 | 0.047 | - | - | - | - | - | - | - | - | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | 0.027 | - | - | - | - | 0.013 | 0.013 | 0.013 | 0.013 | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | 0.02 | - | - | - | - | 0.02 | 0.02 | 0.02 | 0.02 | - | - | - | - | - |
| F 202 | 0.002 | 0.002 | - | - | 0.002 | 0.002 | 0.002 | - | - | - | 0.002 | - | 0.002 | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | 0.079 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | 0.003 | - | - | 0.003 | 0.003 | 0.003 | - | - | - | 0.003 | - | 0.003 | - |

continued...

Table 8.6.2 continued

| | F23 | F24 | F25 | F26 | F27 | F28 | F29 | F30 | F31 | F32 | F33 | F34 | F35 | F36 |
|-------|-----|-------|-------|-------|-------|-------|-------|-------|-------|-------|------|------|------|------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | 0.048 | - | - | - | - | - | - | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | 0.024 | 0.024 | 0.024 | 0.059 | 0.006 | 0.006 | 0.006 | 0.006 | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 |
| F 202 | - | - | - | - | - | 0.002 | - | - | - | - | - | - | - | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | 0.018 | 0.018 | 0.018 | 0.018 | 0.018 | 0.018 | 0.018 | 0.018 | 0.018 | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | - | - | - | - | - | 0.003 | - | - | - | - | - | - | - | - |

continued...

| | F37 | F38 | F39 | F40 | F41 | F42 | F43 | F44 | F45 | F46 | F47 | F48 | F49 | F50 |
|-------|------|------|-----|------|------|-------|-------|-------|-------|-------|-----|-------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | - | - | - | - | - | - | - | - | - | - | - | 0.015 | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | - | - | - | 0.066 | 0.013 |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | 0.02 | 0.02 | - | 0.02 | 0.02 | - | - | - | - | - | - | - | 0.02 | 0.02 |
| F 202 | - | - | - | - | - | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | - | 0.002 | 0.002 | 0.002 |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | 0.079 | 0.079 |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | - | - | - | - | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | 0.003 | 0.003 | 0.003 |

continued...

Table 8.6.2 continued

| | F51 | F52 | F53 | F54 | F55 | F56 | F57 | F58 | F59 | F60 | F61 | F62 | F63 | F64 |
|-------|-------|-------|-------|-----|-------|-----|-------|-------|-------|-------|-------|-----|-----|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | 0.036 | - | - | - | - | - |
| F 183 | - | - | - | - | 0.067 | - | 0.033 | 0.033 | - | - | - | - | - | - |
| F 184 | 0.015 | - | - | - | - | - | - | - | - | 0.015 | 0.014 | - | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | 0.5 | - |
| F 190 | 0.072 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | 0.066 | 0.013 | - | - | - | - | 0.027 | 0.027 | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | 0.02 | 0.02 | - | - | 0.02 | - | - | - | - | 0.01 | 0.01 | - | - | - |
| F 202 | 0.002 | 0.002 | 0.002 | - | - | - | - | - | 0.002 | 0.001 | 0.001 | - | - | 0.002 |
| F 203 | - | - | - | - | - | - | - | - | - | 0.5 | 0.5 | - | - | - |
| F 204 | 0.156 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | 0.003 | 0.003 | - | - | - | - | - | 0.003 | 0.002 | 0.001 | - | - | 0.003 |

continued...

Table 8.6.2 continued

| | F65 | F66 | F67 | F68 | F69 | F70 | F71 | F72 | F73 | F74 | F75 | F76 | F77 | F78 |
|-------|-------|-----|-----|-----|-----|-------|-------|-------|-----|-------|-------|-------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | - | - | - | - | - | - | - | - | - | - | 0.015 | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | 0.008 | 0.008 | 0.008 | 0.008 |
| F 191 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | - | 0.006 | 0.006 | 0.006 | 0.006 |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 202 | 0.002 | - | - | - | - | 0.002 | 0.002 | 0.002 | - | 0.002 | - | 0.002 | 0.002 | 0.002 |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | 0.017 | 0.017 | 0.017 | 0.017 |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | - | - | - | - | 0.003 | 0.003 | 0.003 | - | 0.003 | - | 0.003 | 0.003 | 0.003 |

continued...

Table 8.6.2 continued

| | F79 | F80 | F81 | F82 | F83 | F84 | F85 | F86 | F87 | F88 | F89 | F90 | F91 | F92 |
|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-----|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | 0.014 | - | - | - | - | - | - | 0.003 | 0.003 | 0.003 | 0.017 | 0.003 | 0.015 | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | 0.5 | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | 0.078 | 0.008 | 0.008 | 0.008 | 0.008 | - | - | - | - | - | - | - | - | - |
| F 191 | 0.047 | - | - | - | - | - | - | 0.017 | 0.017 | 0.017 | 0.017 | 0.017 | 0.007 | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 | - | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | 0.02 | 0.02 | - | - | - | - | - | - | - |
| F 202 | 0.002 | - | - | - | - | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | 0.017 | 0.017 | 0.017 | 0.017 | 0.017 | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | - | - | - | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - |

continued...

| | F93 | F94 | F95 | F96 | F97 | F98 | F99 | F100 | F101 | F102 | F103 | F104 | F105 | F106 |
|-------|-----|-----|-------|-------|-------|-------|-----|-------|-------|-------|------|-------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | - | 0.014 | 0.015 | 0.015 | - | - | - | - | - | - | - | 0.002 | 0.002 |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | - | - | - | - | - | - | - | - | 0.048 | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 202 | - | - | 0.002 | 0.002 | 0.002 | 0.002 | - | 0.002 | 0.002 | 0.002 | - | 0.002 | - | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | - | - | 0.003 | 0.003 | 0.003 | 0.003 | - | 0.003 | 0.003 | 0.003 | - | 0.003 | - | - |

continued...

Table 8.6.2 continued

| | F107 | F108 | F109 | F110 | F111 | F112 | F113 | F114 | F115 | F116 | F117 | F118 | F119 | F120 |
|-------|-------|-------|-------|-------|-------|-------|------|------|------|------|------|-------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | 0.101 | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | 0.036 | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | 0.015 | - | - | - | - | - | 0.015 | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | 0.072 | - | - |
| F 191 | - | - | - | - | - | - | - | - | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | 0.053 | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | 0.02 | - | - | - | - | - | - |
| F 202 | - | - | - | 0.002 | - | - | - | - | - | - | - | 0.002 | 0.002 | 0.002 |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | - | - | - | 0.003 | - | - | - | - | - | - | - | 0.003 | 0.003 | 0.003 |

continued...

| | F121 | F122 | F123 | F124 | F125 | F126 | F127 | F128 | F129 | F130 | F131 | F132 | F133 | F134 |
|-------|-------|-------|-------|-------|-------|------|------|-------|-------|------|-------|-------|------|------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | 0.036 | - | - | 0.018 | 0.018 | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | 0.015 | 0.015 | 0.028 | 0.015 | - | - | - | - | - | - | - | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | 0.053 | - | - | - | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | 0.02 | - | - | - | - | - | - | - | - |
| F 202 | 0.002 | 0.002 | 0.002 | 0.002 | - | - | - | 0.002 | 0.002 | - | 0.002 | 0.002 | - | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | 0.003 | 0.003 | 0.003 | - | - | - | 0.003 | 0.003 | - | 0.003 | 0.003 | - | - |

continued...

Table 8.6.2 continued

| | F135 | F136 | F137 | F138 | F139 | F140 | F141 | F142 | F143 | F144 | F145 | F146 | F147 | F148 |
|-------|------|-------|------|------|-------|------|------|------|------|------|-------|-------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | 0.015 | - | - | - | - | - | - | - | - | - | - | 0.017 | 0.016 |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | - | - | - | - | - | - | - | - | - | 0.047 | - | 0.048 | 0.047 |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 202 | - | - | - | - | 0.002 | - | - | - | - | - | 0.002 | 0.002 | 0.002 | 0.002 |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | - | - | - | - | 0.003 | - | - | - | - | - | 0.003 | 0.003 | 0.003 | 0.003 |

continued...

| | F149 | F150 | F151 | F152 | F153 | F154 | F155 | F156 | F157 | F158 | F159 | F160 | F161 | F162 |
|-------|-------|-------|-------|-------|-------|------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | 0.02 | 0.02 |
| F 182 | - | - | - | - | - | - | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | 0.017 | 0.016 | 0.002 | 0.002 | 0.002 | - | 0.015 | 0.004 | 0.004 | 0.004 | 0.004 | 0.02 | 0.017 | 0.017 |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | 0.072 | - | 0.072 | - | - | - | 0.009 | 0.009 | 0.009 | 0.009 | 0.009 | 0.019 | 0.019 |
| F 191 | - | 0.059 | 0.012 | 0.059 | 0.012 | - | - | 0.005 | 0.053 | 0.005 | 0.005 | 0.052 | 0.005 | 0.005 |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 202 | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 | - | 0.002 | - | 0.002 | - | 0.002 | - | 0.002 | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | - | - | 0.003 | - | 0.003 | - | 0.003 | - | 0.003 |

continued...

Table 8.6.2 continued

| | F163 | F164 | F165 | F166 | F167 | F168 | F169 | F170 | F171 | F172 | F173 | F174 | F175 | F176 |
|-------|-------|-------|-------|-------|-------|-------|-------|------|------|------|------|-------|------|------|
| F 181 | 0.02 | 0.02 | 0.02 | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | 0.003 | 0.003 | 0.003 | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | 0.001 | 0.001 | 0.001 | - | - | - | - | - | - | - | - | - | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | 0.019 | 0.019 | 0.019 | - | - | 0.072 | - | - | - | - | - | - | - | - |
| F 191 | 0.005 | 0.005 | 0.005 | - | - | - | - | - | - | - | - | - | - | - |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | 0.005 | 0.005 | 0.005 | - | - | - | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | 0.02 | 0.02 | 0.02 | - | - | - | - |
| F 202 | 0.002 | - | - | 0.002 | 0.002 | 0.002 | 0.002 | - | - | - | - | 0.002 | - | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | - | - | 0.003 | 0.003 | 0.003 | 0.003 | - | - | - | - | 0.003 | - | - |

continued...

| | F177 | F178 | F179 | F180 | F181 | F182 | F183 | F184 | F185 | F186 | F187 | F188 | F189 | F190 |
|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | - | - | 0.015 | - | - | - | - | - | - | - | 0.014 | - | - |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | - | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 202 | 0.002 | - | 0.002 | 0.002 | - | - | - | - | 0.002 | 0.002 | 0.002 | 0.002 | - | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | 0.003 | - | 0.003 | 0.003 | - | - | - | - | 0.003 | 0.003 | 0.003 | 0.003 | - | - |

continued...

Table 8.6.2 continued

| | F191 | F192 | F193 | F194 | F195 | F196 | F197 | F198 | F199 | F200 | F201 | F202 | F203 | F204 |
|-------|-------|-------|-------|------|------|------|------|-------|------|-------|------|------|-------|-------|
| F 181 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 182 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 183 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 184 | - | - | - | - | - | - | - | - | - | - | - | - | 0.005 | 0.02 |
| F 185 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 186 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 187 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 188 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 189 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 190 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 191 | 0.003 | 0.003 | 0.003 | - | - | - | - | - | - | - | - | - | - | 0.048 |
| F 192 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 193 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 194 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 195 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 196 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 197 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 198 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 199 | - | - | - | - | - | - | - | - | - | 0.053 | - | - | 0.018 | 0.018 |
| F 200 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 201 | - | - | - | - | - | - | - | - | 0.02 | 0.02 | - | 0.02 | - | - |
| F 202 | - | 0.002 | 0.002 | - | - | - | - | 0.002 | - | - | - | - | 0.002 | - |
| F 203 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 204 | - | - | 0.158 | - | - | - | - | - | - | - | - | - | - | - |
| F 205 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 206 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 207 | - | - | - | - | - | - | - | - | - | - | - | - | - | - |
| F 208 | - | 0.003 | 0.003 | - | - | - | - | 0.003 | - | - | - | - | 0.003 | - |

continued...

Table 8.6.2 continued

| | F205 | F206 | F207 | F208 | Sum |
|-------|-------|-------|-------|------|-----|
| F 181 | - | - | - | - | 1 |
| F 182 | - | - | - | - | 1 |
| F 183 | - | - | - | - | 1 |
| F 184 | 0.005 | - | - | - | 1 |
| F 185 | - | - | - | - | 1 |
| F 186 | - | - | - | - | 1 |
| F 187 | - | - | - | - | 1 |
| F 188 | - | - | - | - | 1 |
| F 189 | - | - | - | - | 1 |
| F 190 | - | - | - | - | 1 |
| F 191 | - | - | - | - | 1 |
| F 192 | - | - | - | - | 1 |
| F 193 | - | - | - | - | 1 |
| F 194 | - | - | - | - | 1 |
| F 195 | - | - | - | - | 1 |
| F 196 | - | - | - | - | 1 |
| F 197 | - | - | - | - | 1 |
| F 198 | - | - | - | - | 1 |
| F 199 | 0.018 | 0.053 | - | - | 1 |
| F 200 | - | - | - | - | 1 |
| F 201 | - | - | - | 0.02 | 1 |
| F 202 | 0.002 | - | 0.002 | - | 1 |
| F 203 | - | - | - | - | 1 |
| F 204 | - | - | - | - | 1 |
| F 205 | - | - | - | - | 1 |
| F 206 | - | - | - | - | 1 |
| F 207 | - | - | - | - | 1 |
| F 208 | 0.003 | - | 0.003 | - | 1 |

Table 8.6.3. Summary of publications used for the construction of the diet composition matrix of nonfish taxa.

| ID | Species group | Publication |
|-----|----------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| D1 | Detritus | |
| A1 | Benthic algae/ spermatophytes | |
| A2 | Symbiotic algae | |
| A3 | Phytoplankton | |
| I1 | Microfauna | Sorokin (1987) |
| I2 | Zooplankton | Gottfried and Roman (1983), Sorokin (1987), Jarre et al. (1989) |
| I3 | Sponges | Reiswig (1971), Wilkinson (1987) |
| I4 | Hydrozoans | Lewis (1982), Minchin (1983), Sorokin (1987) |
| I5 | Gorgonians | Lewis (1982), Minchin (1983), Sebens and Koehl (1984), Sorokin (1987) |
| I6 | Zoantharians | Lewis (1982), Minchin (1983), Sebens and Koehl (1984), Sorokin (1987) |
| I7 | Scleractinians | Meinkoth (1981), Lewis (1982), Minchin (1983), Sorokin (1987) |
| I8 | Bryozoans | Meinkoth (1981), Kaminski (1984), Winston (1986) |
| I9 | Sipunculid worms | Meinkoth (1981) |
| I10 | Priapuloids | Meinkoth (1981) |
| I11 | Chitons | Lewis (1982), Steneck and Watling (1982), Nishihama et al. (1986) |
| I12 | Gastropods | Kasinathan et al. (1975), Engstrom (1977), Hoffman et al. (1978), Milera and Cortes (1979), Kohn (1980), Broom (1981), Garrity and Levings (1981), Griffiths (1981), Behrens-Yamada (1982), Brawley and Adey (1982), Creese and Underwood (1982), Lewis (1982), Maes (1982), Rietsma et al. (1982), Salvat and Denizot (1982), Steneck (1982), Taylor (1982), Tsikhon-Lukanina (1982), VanMontfrans et al. (1982), Anon. (1983), Broom (1983), Fujioka and Yamazato (1983), McKillup and Butler (1983), Penney and Griffiths (1983), Thangavelu and Muthiah (1983), Berry (1984), Boggs et al. (1984), Morgan and Kitting (1984), Robertson and Schutt (1984), Tomita and Mizushima (1984), Underwood (1984), Watanabe (1984), Hardison and Kitting (1985), Hawkins (1985), Hughes (1985a, 1985b), Kitting (1985), Kotaki et al. (1985), Perron et al. (1985), Steinberg (1985), Winter and Hamilton (1985), Borja (1986), DSouza (1986), Kohlmeyer and Bebout (1986), Mook (1986), Taylor (1987), Berry (1988) |
| I13 | Bivalves | Berg and Alatalo (1981), Schmid and Schaerer (1981), Pohlo (1982), Tsikhon-Lukanina (1982), Fehlbeck (1983), Shafir and Loya (1983), Graham (1984), Reid (1985), Chang et al. (1988), Davenport (1988), Scott (1988) |
| I14 | Scaphopods | Poon (1987) |
| I15 | Squids | Nigmatullin and Toporova (1982), OSullivan and Cullen (1983), Nemoto et al. (1984), Maurer and Bowman (1985), Nicol and ODor (1985), Vovk (1985), Guo-Xin et al. (1986), Yang et al. (1986), Chesalin (1987), Lipinski and Linkowski (1988) |
| I16 | Octopuses | Smale and Buchan (1981), Ambrose and Nelson (1983), Ambrose (1984), Nixon and Budelmann (1984), Lam and Chiu (1985), Kobayashi (1986) |
| I17 | Polychaetes | Fauchald and Jumars (1979), Vreeland and Lasker (1989) |
| I18 | Echiuroids | Schembri and Jaccarini (1978), Ohta (1984) |
| I19 | Pycnogonids | Meinkoth (1981), Bamber and Davis (1982), Child and Harbison (1986), Staples and Watson (1987) |
| I20 | Barnacles | Lewis (1981), Meinkoth (1981), Tsikhon-Lukanina et al. (1986), Sorokin (1987) |
| I21 | Stomatopods | Giovanardi and Piccinetti (1984), Dominguez and Reaka (1988) |
| I22 | Amphipods | Zimmerman et al. (1977), Zimmerman (1979), Brawley and Adey (1981), Howard (1982), Nielsen and Kofeod (1982), Gunnill (1983), Hendler and Miller (1984), Miller (1984), Stuart et al. (1985), Yamashita et al. (1985), SainteMarie (1986), Brawley and Fei (1987), Buschmann and Santelices (1987), Baerlocher et al. (1988), Morrisey (1988) |
| I23 | Tanaids | Johnson and Attramadai (1983), Delille et al. (1985) |
| I24 | Isopods | Waegele (1982), Sandifer and Kerby (1983), Segal (1987) |
| I25 | Shrimps | Corredor and Criales (1977), Balasubramanian et al. (1979), Guitart and Hondares (1980), Chong and Sasekumar (1981), Moriarty and Barclay (1981), Reed et al. (1982), Beal (1983), |

continued...

Table 8.6.3 continued

| ID | Species group | Publication |
|-----|--------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| | | Glynn (1983), Kim et al. (1983), Criales (1984), Howard (1984), Motos (1985), Primavera and Gacutan (1985), Cockcroft and Mclachlan (1986), Figueras (1986), Oya (1987), Scott et al. (1988), Stoner and Zimmerman (1988) |
| I26 | Spiny lobsters | Brito Perez and Diaz Iglesia (1983), Joll and Phillips (1984) |
| I27 | Scyllarid lobsters | Meinkoth (1981), Suthers and Anderson (1981), Rudloe (1983), Lau (1987) |
| I28 | Hermit crabs | Gibbons (1984a, 1984b), Wicksten (1988) |
| I29 | Crabs | Prahl (1978), Patel et al. (1979), Kropp (1981), Meinkoth (1981), Nelson (1981), Paul (1981), Robertson and Pfeiffer (1981), Schembri (1981), Drummond-Davis et al. (1982), Laughlin (1982), McDonald (1982), Seiple and Salmon (1982), Williams (1982), Klumpp and Nichols (1983), McLean (1983), Morales and Antezana (1983), Saisho et al. (1983), DuPreez (1984), Kotaki et al. (1985), Lawton and Hughes (1985), Signor (1985), Chilton and Bull (1986), Choy (1986), Kilar and Lou (1986), Matsui et al. (1986), Kurihara and Okamoto (1987), Wear and Haddon (1987), Coen (1988), Perez and Bellwood (1988), Winfree and Weinstein (1989) |
| I30 | Hemichordates | Dilly (1985), Lester (1985) |
| I31 | Asteroids | Schembri and Jaccarini (1978), Bitter et al. (1980), Town (1981), Freitas (1982), Harrold (1982), Scheibling (1982), Bitter and Penchaszadeh (1983), Penchaszadeh and Molinet (1983), Shivji et al. (1983), Bitter Soto (1984), Chiu et al. (1985), Dearborn and Edwards (1985), McClintock and Lawrence (1985) |
| I32 | Ophiuroids | Chartock (1983), Lee et al. (1983), Clements and Stancyk (1984), Hensen (1984), Pearson and Gage (1984) |
| I33 | Echinoids | Roushdy and Hansen (1960), Carpenter (1981), Drifmeyer (1981), McClintock et al. (1982), Regis and Thomassin (1982), Vadas et al. (1982), Gomez et al. (1983), Hughes et al. (1985), Andrew and Stocker (1986), Coyer et al. (1987), Downing and El-Zahr (1987), Foster (1987), Jafari and Mahasneh (1987), Briscoe and Sebens (1988), Dean et al. (1988), Stevenson (1988) |
| I34 | Holothurians | Akhmeteva et al. (1982), Moriarty (1982), Hammond (1983), Hammond and Wilkinson (1985), Massin and Doumen (1986), Sisak and Sander (1987) |
| I35 | Tunicates | Meinkoth (1981), Stuart and Klumpp (1984) |
| R1 | Sea turtles | Witzell (1983), Acevedo et al. (1984), Meylan (1988) |
| B1 | Sea birds | Polovina (1984) |

Appendix 8.7

Table 8.7.1. Characteristics of 27 fish species groups (extended version).

| Group no. | Family | Species | L (cm) | W (g) | W _{mean} (g) | Density (N·m ⁻²) | Biomass (g·m ⁻²) | Aspect ratio | Food type | Q/B% day ⁻¹ | Q/B year ⁻¹ | Natural mortality | Weighing factor | Level | |
|----------------|--------------------|-----------------------------------------|------------------------------|-----------|-----------------------|------------------------------|------------------------------|--------------|-----------|------------------------|------------------------|-------------------|-----------------|---------|---|
| 1 | Carcharhinidae | <i>Galeocerdo cuvier</i> | 457 | 672,000 | 173,376 | 1.25E-04 | 21.6720 | 7 * | 0 | 1.2 | 4.4 | 0.19 | 95.357 | 4 | |
| | Ginglymostomatidae | <i>Ginglymostoma cirratum</i> | 430 | 500,000 * | 129,000 | 1.25E-04 | 16.1250 | 7 * | 0 | 1.2 | 4.5 | 0.24 | 73.144 | 1 | |
| | Sphyrnidae | <i>Sphyrna lewini</i> | 420 | 400,853 | 103,420 | 1.25E-05 | 1.2928 | 7 * | 0 | 1.3 | 4.7 | | 6.131 | 4 1 | |
| | Carcharhinidae | <i>Carcharhinus falciformis</i> | 350 Lmax | 350,000 | 90,300 | 1.25E-05 | 1.1288 | 7 * | 0 | 1.3 | 4.9 | | 5.502 | 4 | |
| | Carcharhinidae | <i>Carcharhinus longimanus</i> | 350 Lmax | 350,000 | 90,300 | 1.25E-05 | 1.1288 | 7 * | 0 | 1.3 | 4.9 | | 5.502 | 4 | |
| | Carcharhinidae | <i>Carcharhinus leucas</i> | 350 Lmax | 350,000 | 90,300 | 1.25E-05 | 1.1288 | 7 * | 0 | 1.3 | 4.9 | 0.24 | 5.502 | 4 1 | |
| | Myliobatidae | <i>Aetobatus narinari</i> | 370 Len | 230,000 | 59,340 | 1.25E-05 | 0.7418 | 7 * | 0 | 1.5 | 5.3 | | 3.935 | 4 3 2 1 | |
| | Dasyatidae | <i>Dasyatis americana</i> | 300 | 122,000 * | 31,476 | 1.25E-03 | 39.3450 | 7 * | 0 | 1.7 | 6.0 | | 237.243 | 1 | |
| | Carcharhinidae | <i>Carcharhinus limbatus</i> | 247 Lmax | 116,000 | 29,928 | 1.25E-05 | 0.3741 | 7 * | 0 | 1.7 | 6.1 | 0.28 | 2.279 | 4 | |
| | Carcharhinidae | <i>Negaprion brevirostris</i> | 284 | 110,000 | 28,380 | 1.25E-04 | 3.5475 | 7 * | 0 | 1.7 | 6.2 | 0.25 | 21.842 | 2 1 | |
| | Carcharhinidae | <i>Carcharhinus perezii</i> (springeri) | 230 Lmax | 100,000 | 25,800 | 1.25E-04 | 3.2250 | 7 * | 0 | 1.7 | 6.3 | | 20.242 | 1 | |
| | Carcharhinidae | <i>Carcharhinus acronotus</i> | 200 Lmax | 90,000 | 23,220 | 1.25E-05 | 0.2903 | 7 * | 0 | 1.8 | 6.4 | 0.33 | 1.861 | 1 * | |
| | # | Megalopidae | <i>Tarpon atlanticus</i> | 250 | 160,000 | 41,280 | 1.38E-04 | 5.6760 | 3 | 0 | 1.0 | 3.7 | 0.19 | 20.772 | 4 |
| | 2 | Sphyrnidae | <i>Sphyrna tiburo</i> | 150 | 18,000 * | 4,644 | 1.25E-04 | 0.5805 | 7 * | 0 | 2.4 | 8.9 | 0.27 | 5.150 | 1 |
| Triakidae | | <i>Mustelus canis</i> | 150 | 15,000 * | 3,870 | 1.25E-05 | 0.0484 | 7 * | 0 | 2.5 | 9.2 | 0.31 | 0.445 | 1 | |
| Carcharhinidae | | <i>Rhizoprionodon porosus</i> | 107 | 5,020 | 1,295 | 1.25E-04 | 0.1619 | 7 * | 0 | 3.1 | 11.5 | | 1.858 | 1 2 3 4 | |
| Scombridae | | <i>Scomberomorus cavalla</i> | 137 FL | 34,285 | 8,846 | 1.38E-04 | 1.2163 | 6.3 | 0 | 2.0 | 7.4 | 0.37 | 8.985 | 4 | |
| Scombridae | | <i>Euthynnus aletteratus</i> | 93 FL | 12,200 | 3,148 | 1.25E-04 | 0.3935 | 6.3 | 0 | 2.5 | 9.1 | | 3.573 | 4 | |
| Scombridae | | <i>Scomberomorus regalis</i> | 83.5 FL | 4,900 | 1,264 | 1.37E-03 | 1.7380 | 6.2 | 0 | 3.0 | 10.8 | 0.47 | 18.763 | 4 | |
| 3 | Carangidae | <i>Seriola dumerilii</i> | 194 | 80,000 | 20,640 | 1.25E-05 | 0.2580 | 4.5 | 0 | 1.4 | 5.2 | 0.26 | 1.347 | 4 2 1 | |
| | Carangidae | <i>Seriola rivoliana</i> | 97 | 24,000 | 6,192 | 1.25E-05 | 0.0774 | 4.5 * | 0 * | 1.8 | 6.7 | 0.79 | 0.516 | 4 1 | |
| | Carangidae | <i>Trachinotus falcatus</i> | 105 FL | 36,000 | 9,288 | 1.25E-05 | 0.1161 | 3.9 | 0 | 1.6 | 5.7 | | 0.661 | 4 1 | |
| 4 | Carangidae | <i>Caranx lugubris</i> | 80 | 6,554 | 1,691 | 1.25E-05 | 0.0211 | 5.5 | 0 | 2.6 | 9.6 | | 0.203 | 3 | |
| | Carangidae | <i>Trachinotus goodei</i> | 50 | 3,900 * | 1,006 | 1.25E-04 | 0.1258 | 5 | 0 | 2.8 | 10.2 | | 1.279 | 3 * | |
| | Carangidae | <i>Alectis ciliaris</i> | 91 TL/ | 10,700 | 2,761 | 1.25E-05 | 0.0345 | 4.8 | 0 * | 2.2 | 8.1 | | 0.281 | 2 4 | |
| | Carangidae | <i>Elagatis bipinnulatus</i> | 107 FL | 10,500 | 2,709 | 1.25E-05 | 0.0339 | 4.5 * | 0 * | 2.2 | 7.9 | | 0.267 | 4 3 | |
| | Carangidae | <i>Caranx latus</i> | 80 | 6,250 | 1,613 | 2.50E-04 | 0.4033 | 4.5 | 0 | 2.4 | 8.7 | | 3.517 | 2 3 | |
| | Carangidae | <i>Caranx ruber</i> | 56 | 3,160 | 815 | 3.87E-02 | 31.5813 | 4.5 | 0 | 2.8 | 10.1 | 1.17 | 317.673 | 2 | |
| | Carangidae | <i>Caranx crysos</i> | 58 TL | 2,314 | 597 | 1.38E-04 | 0.0821 | 4.2 | 0 | 2.8 | 10.3 | | 0.846 | 3 1 | |
| | Carangidae | <i>Caranx bartholomaei</i> | 81 TL | 9,862 | 2,544 | 1.25E-04 | 0.3180 | 4.1 | 0 | 2.1 | 7.6 | | 2.405 | 1 2 | |
| | 5 | Carangidae | <i>Decapterus macarellus</i> | 35 SL | 750 * | 194 | 1.25E-05 | 0.0024 | 3.8 | 0 | 3.4 | 12.3 | 0.93 | 0.030 | 4 |
| Carangidae | | <i>Decapterus punctatus</i> | 32 | 573 | 148 | 1.25E-04 | 0.0185 | 3.7 | 0 | 3.5 | 12.7 | 0.83 | 0.235 | 4 | |
| Carangidae | | <i>Selar crumenophthalmus</i> | 40 TL | 1,240 | 320 | 1.25E-04 | 0.0400 | 3.5 | 0 | 2.9 | 10.7 | | 0.428 | 3 2 | |
| Elopidae | | <i>Elops saurus</i> | 47 | 444 | 115 | 1.25E-04 | 0.0144 | 3.4 | 0 * | 3.5 | 12.9 | 0.62 | 0.186 | 4 | |
| 6 | | Sparidae | <i>Calamus bajonado</i> | 64 | 4,500 | 1,161 | 2.50E-05 | 0.0290 | 3 | 0 | 2.1 | 7.6 | 0.221 | 1 | |
| | Lutjanidae | <i>Ocyurus chrysurus</i> | 75 TL | 3,570 | 921 | 2.75E-02 | 25.3275 | 3 | 0 | 2.2 | 7.9 | 0.60 | 199.830 | 2 | |
| | Haemulidae | <i>Haemulon album</i> | 65 FL | 5,300 | 1,367 | 2.62E-04 | 0.3588 | 2.8 | 0 | 1.9 | 7.0 | 0.79 | 2.527 | 1 | |
| | Haemulidae | <i>Anisotremus surinamensis</i> | 63 TL | 4,770 | 1,231 | 1.38E-04 | 0.1693 | 2.7 | 0 | 1.9 | 7.0 | | 1.192 | 1 | |
| | Ephippidae | <i>Chaetodipterus faber</i> | 90 | 9,000 | 2,322 | 2.50E-04 | 0.5805 | 3.1 | 0.07 | 2.0 | 7.3 | | 4.261 | 2 | |
| | Pomacanthidae | <i>Pomacanthus arcuatus</i> | 60 | 12,407 | 3,201 | 2.51E-03 | 8.0425 | 2.9 | 0.08 | 1.8 | 6.7 | 0.63 | 53.932 | 1 | |
| | Balistidae | <i>Canthidermis sufflamen</i> | 56 + | 4,783 | 1,234 | 1.25E-05 | 0.0154 | 1.7 | 0.025 | 1.6 | 5.8 | | 0.089 | 4 | |
| | Albulidae | <i>Albula vulpes</i> | 82 | 7,150 | 1,845 | 1.25E-05 | 0.0231 | 2.2 * | 0 * | 1.6 | 5.9 | | 0.136 | 1 | |
| | Lutjanidae | <i>Lutjanus analis</i> | 74 FL | 5,511 | 1,422 | 2.62E-04 | 0.3733 | 2.1 | 0 | 1.6 | 6.0 | 0.33 | 2.236 | 2 | |
| | Lutjanidae | <i>Lutjanus apodus</i> | 59 FL | 3,502 | 904 | 1.50E-02 | 13.5600 | 2 | 0 | 1.8 | 6.5 | 0.54 | 87.655 | 2 | |
| | Balistidae | <i>Balistes vetula</i> | 45 FL | 2,586 | 667 | 1.37E-03 | 0.9171 | 2 | 0.01 | 1.9 | 6.9 | 0.56 | 6.356 | 1 | |
| | Ostraciidae | <i>Lactophrys trigonus</i> | 46 + | 3,052 | 787 | 1.28E-03 | 1.0034 | 1.9 | 0.05 | 1.9 | 6.8 | | 6.871 | 1 | |
| | 7 | Belonidae | <i>Tylosurus acus</i> | 135 SL | 2,800 * | 722 | 1.25E-04 | 0.0903 | 3 * | 0 | 2.3 | 8.3 | | 0.753 | 4 |
| Belonidae | | <i>Tylosurus crocodilus</i> | 107 SL | 1,400 * | 361 | 1.25E-03 | 0.4513 | 3.1 | 0 | 2.7 | 9.8 | | 4.415 | 4 | |
| Clupeidae | | <i>Opisthonema oglinum</i> | 26 | 730 | 188 | 1.25E-02 | 2.3500 | 3.1 | 0 | 3.1 | 11.2 | | 26.283 | 4 | |

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Table 8.7.1 continued

| Group no. | Family | Species | L (cm) | W (g) | W _{max} (g) | Density (N·m ⁻²) | Biomass (g·m ⁻²) | Aspect ratio | Food type | Q/B% day ⁻¹ | Q/B year ⁻¹ | Natural mortality | Weighing factor | Level |
|-----------|-----------------|----------------------------------|---------|---------|----------------------|------------------------------|------------------------------|--------------|-----------|------------------------|------------------------|-------------------|-----------------|-------|
| | Belonidae | <i>Ablennes hians</i> | 87 | 751 | 194 | 1.25E-05 | 0.0024 | 3 * | 0 | 3.0 | 10.9 | | 0.026 | 4 * |
| | Hemiramphidae | <i>Hemiramphus balao</i> | 40 | 614 | 158 | 1.25E-03 | 0.1975 | 2.7 | 0 | 3.0 | 10.8 | | 2.130 | 4 |
| | Clupeidae | <i>Sardinella aurita</i> | 29 | 1,000 * | 258 | 1.25E-02 | 3.2250 | 3 * | 0.2 * | 3.6 | 13.2 | 0.68 | 42.614 | 4 |
| | Carangidae | <i>Oligoplites saurus</i> | 31 FL | 334 | 86 | 1.25E-02 | 1.0750 | 3.1 | 0 | 3.6 | 13.1 | | 14.060 | 3 * |
| | Emmelichthyidae | <i>Inermia vittata</i> | 23 (SL) | 282 | 73 | 1.25E-02 | 0.9125 | 2.9 | 0 | 3.6 | 13.1 | | 11.924 | 3 |
| | Belonidae | <i>Platybelone argalus</i> | 51 / | 150 * | 39 | 1.25E-03 | 0.0488 | 2.65 | 0 | 3.9 | 14.1 | | 0.688 | 4 |
| | Clupeidae | <i>Harengula clupeiola</i> | 18 | 240 * | 62 | 1.25E-01 | 7.7500 | 2.7 | 0 | 3.5 | 12.8 | 0.91 | 99.489 | 4 |
| | Clupeidae | <i>Harengula humeralis</i> | 22 | 440 * | 114 | 1.39E-01 | 15.8175 | 2.9 | 0.03 | 3.4 | 12.4 | 0.66 | 195.804 | 4 |
| 8 # | Gerreidae | <i>Gerres cinereus</i> | 40 | 690 | 178 | 1.39E-02 | 2.4698 | 3.9 | 0 | 3.5 | 12.7 | | 31.336 | 1 |
| | Sparidae | <i>Calamus calamus</i> | 38 | 1,541 | 398 | 2.50E-04 | 0.0995 | 3.2 | 0 | 2.7 | 9.7 | | 0.965 | 1 |
| | Sparidae | <i>Calamus pennatula</i> | 39 | 1,700 * | 439 | 1.38E-04 | 0.0604 | 3 | 0 | 2.5 | 9.3 | | 0.561 | 1 |
| | Haemulidae | <i>Haemulon plumieri</i> | 42 | 1,360 | 351 | 1.51E-02 | 5.3089 | 2.9 | 0 | 2.6 | 9.4 | 0.67 | 50.025 | 1 |
| | Haemulidae | <i>Haemulon carbonarium</i> | 38 | 861 | 222 | 1.25E-04 | 0.0278 | 2.8 | 0 | 2.8 | 10.3 | | 0.286 | 1 |
| | Haemulidae | <i>Anisotremus virginicus</i> | 30 / | 850 | 219 | 1.38E-04 | 0.0301 | 2.8 | 0 | 2.8 | 10.3 | | 0.310 | 1 |
| | Haemulidae | <i>Haemulon macrostomum</i> | 45 | 1,795 | 463 | 1.25E-05 | 0.0058 | 2.7 | 0 | 2.4 | 8.6 | | 0.050 | 1 |
| | Synodontidae | <i>Synodus foetens</i> | 46 | 1,100 * | 284 | 1.25E-04 | 0.0355 | 2.65 | 0 | 2.6 | 9.5 | | 0.336 | 1 |
| | Sphyraenidae | <i>Sphyraena picudilla</i> | 50 | 1,300 * | 335 | 1.25E-05 | 0.0042 | 2.6 | 0 | 2.5 | 9.1 | | 0.038 | 2 |
| | Holocentridae | <i>Holocentrus ascensionis</i> | 37 TL | 838 | 216 | 2.53E-03 | 0.5454 | 2.6 | 0 | 2.7 | 10 | 0.74 | 5.437 | 1 |
| | Haemulidae | <i>Haemulon parrai</i> | 41 | 1,265 * | 326 | 1.25E-03 | 0.4075 | 2.5 | 0 | 2.5 | 9.0 | | 3.655 | 1 |
| | Haemulidae | <i>Haemulon sciurus</i> | 41 | 1,185 | 306 | 3.87E-03 | 1.1858 | 2.5 | 0 | 2.5 | 8.9 | 0.68 | 10.606 | 1 |
| | Haemulidae | <i>Pomadasys crocro</i> | 33 | 890 * | 230 | 1.25E-05 | 0.0029 | 2.5 * | 0 * | 2.6 | 9.6 | | 0.028 | 1 |
| | Synodontidae | <i>Synodus intermedius</i> | 46 | 1,103 | 285 | 1.37E-03 | 0.3919 | 2.4 | 0 | 2.4 | 8.9 | | 3.504 | 1 |
| | Serranidae | <i>Paranthias furcifer</i> | 37 | 1,100 | 284 | 1.25E-05 | 0.0036 | 2.3 | 0 | 2.4 | 8.7 | 0.63 | 0.031 | 2 3 |
| | Lutjanidae | <i>Lutjanus griseus</i> | 55 | 1,590 | 410 | 1.50E-02 | 6.1500 | 2.1 | 0 | 2.1 | 7.7 | 0.54 | 47.340 | 2 |
| | Lutjanidae | <i>Lutjanus mahagoni</i> | 40 | 980 * | 253 | 1.50E-02 | 3.7950 | 2.05 | 0 | 2.3 | 8.5 | | 32.136 | 2 |
| | Lutjanidae | <i>Lutjanus synagris</i> | 43 FL | 1,213 | 313 | 2.50E-03 | 0.7825 | 2 | 0.03 | 2.3 | 8.3 | 0.59 | 6.461 | 2 |
| | Ostraciidae | <i>Lactophrys triqueter</i> | 30 | 1,394 | 360 | 2.51E-03 | 0.9045 | 2 | 0.02 | 2.2 | 8.0 | | 7.199 | 1 |
| | Ogcocephalidae | <i>Ogcocephalus nasutus</i> | 28 / | 641 | 165 | 1.25E-04 | 0.0206 | | 0.1 | - | - | | 0.185 | 1 |
| | Holocentridae | <i>Neoniphon marianus</i> | 19 | 163 | 42 | 2.62E-04 | 0.0110 | 3 | 0 | 4.1 | 14.9 | 1.93 | 0.164 | 1 |
| # | Holocentridae | <i>Holocentrus coruscus</i> | 13 | 57 | 15 | 1.51E-03 | 0.0227 | 3 | 0 | 5.0 | 18.4 | 3.52 | 0.418 | 1 |
| | Holocentridae | <i>Sargocentron vexillarium</i> | 18 - | 181 | 47 | 1.40E-03 | 0.0658 | 2.7 | 0 | 3.7 | 13.6 | 2.1 | 0.895 | 1 |
| | Haemulidae | <i>Haemulon chrysargyreum</i> | 24 | 251 | 65 | 1.37E-03 | 0.0894 | 2.6 | 0 | 3.4 | 12.6 | | 1.125 | 1 3 |
| | Haemulidae | <i>Haemulon flavolineatum</i> | 27 / | 480 | 124 | 1.50E-01 | 18.6000 | 2.5 | 0 | 2.9 | 10.7 | 0.90 | 199.904 | 1 |
| | Holocentridae | <i>Myripristis jacobus</i> | 23 | 397 | 102 | 3.87E-04 | 0.0395 | 2.5 | 0 | 3.1 | 11.2 | | 0.441 | 1 |
| | Haemulidae | <i>Pomadasys corvinaeformis</i> | 25 | 386 | 100 | 1.25E-05 | 0.0013 | 2.5 * | 0 * | 3.1 | 11.3 | | 0.014 | 1 |
| | Haemulidae | <i>Haemulon bonariense</i> | 30 FL | 385 | 99 | 1.25E-05 | 0.0012 | 2.5 | 0 * | 3.1 | 11.2 | | 0.014 | 1 |
| | Chaetodontidae | <i>Chaetodon ocellatus</i> | 16.8 | 200 * | 52 | 1.25E-05 | 0.0007 | 2.5 * | 0 * | 3.5 | 12.9 | 1.55 | 0.008 | 1 |
| | Chaetodontidae | <i>Chaetodon capistratus</i> | 14.2 | 120 * | 31 | 2.50E-03 | 0.0775 | 2.5 | 0 | 3.9 | 14.4 | 2.02 | 1.117 | 1 |
| | Synodontidae | <i>Synodus synodus</i> | 33 | 400 * | 103 | 1.25E-03 | 0.1288 | 2.4 | 0 | 3.0 | 11.0 | | 1.413 | 1 |
| | Haemulidae | <i>Haemulon aurolineatum</i> | 23 FL | 188 | 49 | 1.26E-02 | 0.6186 | 2.4 | 0 | 3.5 | 12.9 | 2.37 | 8.010 | 1 |
| | Chaetodontidae | <i>Chaetodon striatus</i> | 15.8 | 164 | 42 | 1.25E-05 | 0.0005 | 2.4 | 0 | 3.6 | 13.1 | 1.70 | 0.007 | 1 |
| | Mullidae | <i>Mulloidichthys martinicus</i> | 35 | 440 | 114 | 1.5E-02 | 1.7100 | 2.3 | 0 | 2.9 | 10.5 | 0.98 | 17.915 | 1 |
| | Mullidae | <i>Pseudupeneus maculatus</i> | 28 / | 393 | 101 | 5.00E-03 | 0.5050 | 2.3 | 0 | 3.0 | 10.8 | 0.95 | 5.455 | 1 |
| | Chaetodontidae | <i>Chaetodon sedentarius</i> | 14.7 | 130 * | 34 | 1.25E-05 | 0.0004 | 2.3 | 0 | 3.7 | 13.5 | 1.91 | 0.006 | 1 |
| | Pempheridae | <i>Pempheris schomburgki</i> | 15 | 100 * | 26 | 1.25E-03 | 0.0325 | 2.3 | 0 | 3.9 | 14.2 | | 0.461 | 1 2 |
| | Holocentridae | <i>Holocentrus rufus</i> | 32 | 583 | 150 | 1.39E-02 | 2.0831 | 2.2 | 0 | 2.7 | 9.8 | 2.71 | 20.459 | 1 |
| | Pomacentridae | <i>Abudefduf saxatilis</i> | 20 | 483 | 125 | 1.25E-03 | 0.1563 | 3 | 0.1 | 3.6 | 13.3 | | 2.075 | 1 |
| 9 | Hemiramphidae | <i>Hemiramphus brasiliensis</i> | 32.5 | 298 | 77 | 1.25E-04 | 0.0096 | 3.2 | 0.8 | 10.4 | 37.8 | 1.23 | 0.364 | 4 |
| | Hemiramphidae | <i>Hyporhamphus unifasciatus</i> | 27 | 167 | 43 | 1.25E-04 | 0.0054 | 3 * | 0.8 * | 11.1 | 40.4 | | 0.217 | 4 |
| 10 | Kyphosidae | <i>Kyphosus sectatrix</i> | 80 | 9,139 | 1,579 | 1.37E-03 | 2.1711 | 2.9 | 1 | 6.2 | 22.8 | | 52.919 | 1 |
| | Kyphosidae | <i>Kyphosus incisor</i> | 70 | 6,122 | 2,358 | 2.50E-03 | 5.8950 | 2.8 | 1 | 6.7 | 24.4 | | 134.155 | 1 |

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Table 8.7.1 continued

| Group no. | Family | Species | L (cm) | W (g) | W _{mean} (g) | Density (N·m ⁻²) | Biomass (g·m ⁻²) | Aspect ratio | Food type | Q/B% day ⁻¹ | Q/B year ⁻¹ | Natural mortality | Weighing factor | Level |
|-----------------|-------------------------------|-------------------------------------|-----------------------------|----------|-----------------------|------------------------------|------------------------------|--------------|-----------|------------------------|------------------------|-------------------|-----------------|-------|
| 11 # | Balistidae | <i>Balistes capriscus</i> | 31 | 611 | 158 | 1.25E-05 | 0.0020 | 2.4 | 0.5 * | 5.2 | 19.2 | 0.94 | 0.038 | 1 |
| | Balistidae | <i>Melichthys niger</i> | 36 | 960 * | 248 | 1.25E-04 | 0.0310 | 2.3 | 0.75 | 6.4 | 23.2 | | 0.719 | 2 |
| | Sparidae | <i>Diplodus caudimaculata</i> | 28 | 600 * | 155 | 1.38E-04 | 0.0213 | 2.9 | 0.8 | 8.4 | 30.8 | | 0.655 | 1 |
| | Sparidae | <i>Archosargus rhomboidalis</i> | 35 | 1,200 * | 310 | 1.50E-04 | 0.0465 | 2.5 | 0.8 | 7.0 | 25.5 | | 1.184 | 1 |
| | Mugilidae | <i>Mugil curema</i> | 38 | 767 | 198 | 1.38E-04 | 0.0272 | 2.7 | 1 | 9.9 | 36.3 | 0.59 | 0.989 | 1 |
| | Pomacentridae | <i>Microspathodon chrysurus</i> | 20 | 412 | 106 | 1.25E-02 | 1.3250 | 1.9 | 0.8 | 7.1 | 25.8 | | 34.140 | 1 |
| | Acanthuridae | <i>Acanthurus bahianus</i> | 25 FL | 288 | 74 | 3.75E-02 | 2.7750 | 2.2 | 0.9 | 9.4 | 34.4 | 1.00 | 95.409 | 1 |
| | Pomacentridae | <i>Abudefduf taurus</i> | 25 | 900 * | 232 | 1.25E-04 | 0.0290 | 2 | 0.9 | 7.6 | 27.9 | | 0.809 | 1 |
| | Acanthuridae | <i>Acanthurus coeruleus</i> | 32 FL | 983 | 254 | 2.50E-02 | 6.3500 | 1.9 | 0.9 | 6.7 | 24.4 | 0.70 | 154.811 | 1 |
| Acanthuridae | <i>Acanthurus chirurgus</i> | 33 FL | 820 | 212 | 1.50E-02 | 3.1800 | 1.8 | 0.9 | 6.8 | 24.7 | 0.71 | 78.701 | 1 | |
| 12 # | Lutjanidae | <i>Lutjanus cyanopterus</i> | 160 | 70,000 | 18,060 | 1.25E-05 | 0.2258 | 2.2 | 0 | 1.0 | 3.7 | | 0.834 | 2 |
| | Lutjanidae | <i>Lutjanus jocu</i> | 83 | 13,380 | 3,452 | 1.25E-04 | 0.4315 | 2.1 | 0 | 1.4 | 5.0 | | 2.173 | 2 |
| | Serranidae | <i>Mycteroperca tigris</i> | 102 | 17,043 | 4,397 | 1.25E-04 | 0.5496 | 1.8 | 0 | 1.2 | 4.5 | 0.42 | 2.451 | 1 |
| | Sphyraenidae | <i>Sphyraena barracuda</i> | 178 | 57,800 | 14,912 | 1.50E-03 | 22.3680 | 1.6 | 0 | 0.9 | 3.3 | 0.25 | 73.386 | 2 3 |
| | Serranidae | <i>Epinephelus striatus</i> | 100 | 23,800 | 6,140 | 2.50E-04 | 1.5350 | 1.6 | 0 | 1.1 | 3.9 | | 5.956 | 1 |
| | Rachycentridae | <i>Rachycentron canadum</i> | 160 | 33,400 | 8,617 | 1.25E-05 | 0.1077 | 1.6 | 0 | 1.0 | 3.6 | 0.40 | 0.390 | 2 3 4 |
| | Fistulariidae | <i>Fistularia tabacaria</i> | 183 | 10,800 * | 2,786 | 1.25E-04 | 0.3483 | 1.7 | 0 | 1.3 | 4.7 | | 1.649 | 1 |
| | Labridae | <i>Lachnolaimus maximus</i> | 86 | 7,728 | 1,994 | 1.25E-05 | 0.0249 | 1.5 | 0 | 1.3 | 4.8 | | 0.119 | 1 |
| | Diodontidae | <i>Diodon holocanthus</i> | 51 | 11,300 * | 2,915 | 1.25E-02 | 36.4739 | 1.4 | 0 | 1.2 | 4.2 | | 153.250 | 1 |
| | Serranidae | <i>Mycteroperca rubra</i> | 70 | 10,719 | 2,766 | 1.25E-05 | 0.0346 | 1.2 | 0 * | 1.1 | 4.0 | | 0.137 | 1 |
| | Diodontidae | <i>Diodon hystrix</i> | 60 | 18,446 | 4,759 | 1.25E-04 | 0.5949 | 1.1 | 0 | 0.9 | 3.4 | | 2.012 | 1 |
| | Serranidae | <i>Mycteroperca venenosa</i> | 86 FL | 12,270 | 3,166 | 1.25E-05 | 0.0396 | 1.1 | 0 | 1.0 | 3.7 | 0.36 | 0.145 | 1 |
| | Muraenidae | <i>Gymnothorax funebris</i> | 190 | 12,000 | 3,096 | 1.25E-04 | 0.3870 | 0.7 * | 0 | 0.8 | 2.9 | | 1.137 | 1 |
| | 13 | Balistidae | <i>Xanthichthys ringens</i> | 25 | 320 * | 83 | 1.25E-04 | 0.0104 | 1.9 | 0 * | 2.8 | 10.3 | | 0.107 |
| Gerreidae | | <i>Eucinostomus argenteus</i> | 30 | 290 * | 75 | 1.25E-02 | 0.9394 | 1.9 | 0 | 2.9 | 10.5 | | 9.893 | 1 |
| Priacanthidae | | <i>Heteropriacanthus cruentatus</i> | 30 | 700 * | 181 | 1.25E-04 | 0.0226 | 1.8 | 0 | 2.3 | 8.5 | | 0.191 | 2 |
| Labridae | | <i>Clepticus parrae</i> | 25 | 447 | 115 | 1.25E-02 | 1.4375 | 1.8 | 0 | 2.6 | 9.4 | | 13.515 | 3 |
| Haemulidae | | <i>Haemulon melanurum</i> | 30 FL | 427 | 110 | 1.25E-05 | 0.0014 | 1.8 | 0 * | 2.6 | 9.3 | | 0.013 | 1 |
| Pomacentridae | | <i>Chromis multilineata</i> | 20 SL | 339 | 87 | 1.25E-02 | 1.0875 | 1.8 | 0 | 2.7 | 9.8 | | 10.631 | 1 3 |
| Aulostomidae | | <i>Aulostomus maculatus</i> | 76 / | 777 | 200 | 2.50E-03 | 0.5000 | 1.7 | 0 | 2.2 | 8.1 | | 4.032 | 2 |
| Sciaenidae | | <i>Odontoscion dentex</i> | 25 | 318 | 82 | 2.50E-04 | 0.0205 | 1.7 | 0 | 2.6 | 9.6 | | 0.197 | 1 2 |
| Serranidae | | <i>Cephalopholis fulva</i> | 34 | 640 | 165 | 1.37E-03 | 0.2269 | 1.5 | 0 | 2.1 | 7.8 | 0.78 | 1.765 | 1 |
| Serranidae | | <i>Dipllectrum formosum</i> | 31 | 550 * | 142 | 1.25E-05 | 0.0018 | 1.5 | 0 * | 2.2 | 8.1 | 0.78 | 0.014 | 1 |
| Grammistidae | | <i>Rypticus saponaceus</i> | 29.5 (SL) | 685 | 177 | 1.25E-02 | 2.2147 | 1.4 | 0 | 2.0 | 7.4 | | 16.408 | 1 |
| Sciaenidae | | <i>Equetus punctatus</i> | 28 | 460 | 119 | 1.25E-04 | 0.0149 | 1.25 | 0 | 2.1 | 7.7 | | 0.114 | 1 |
| Serranidae | | <i>Cephalopholis cruentata</i> | 34 | 691 | 178 | 1.25E-04 | 0.0223 | 1.2 | 0 | 1.9 | 7.0 | 0.64 | 0.155 | 1 |
| Priacanthidae | | <i>Priacanthus arenatus</i> | 40 | 1,653 | 426 | 1.25E-04 | 0.0533 | 1.8 | 0 | 2.0 | 7.2 | | 0.383 | 2 |
| Labridae | | <i>Halichoeres radiatus</i> | 46 | 2,283 | 589 | 1.25E-04 | 0.0736 | 1.6 | 0 | 1.7 | 6.3 | | 0.460 | 1 |
| Dactylopteridae | | <i>Dactylopterus volitans</i> | 45 + | 1,825 | 471 | 1.25E-02 | 5.8875 | 1.6 | 0 | 1.8 | 6.6 | | 38.668 | 1 |
| Serranidae | | <i>Epinephelus morio</i> | 71 | 5,010 | 1,293 | 1.25E-05 | 0.0162 | 1.5 | 0 | 1.4 | 5.1 | 0.39 | 0.083 | 1 |
| Serranidae | | <i>Epinephelus adscensionis</i> | 50 | 1,981 | 511 | 2.50E-04 | 0.1278 | 1.5 | 0 | 1.7 | 6.3 | | 0.808 | 1 |
| Diodontidae | | <i>Chilomycterus antennatus</i> | 26.3 | 1,420 | 366 | 1.38E-04 | 0.0503 | 1.4 | 0 | 1.8 | 6.5 | | 0.325 | 1 * |
| Diodontidae | | <i>Chilomycterus antillarum</i> | 26.3 | 1,420 * | 366 | 1.25E-05 | 0.0046 | 1.4 * | 0 * | 1.8 | 6.5 | | 0.030 | 1 * |
| Malacanthidae | | <i>Malacanthus plumieri</i> | 61 / | 1,126 | 291 | 1.27E-02 | 3.7103 | 1.4 | 0 | 1.9 | 6.8 | 0.42 | 25.309 | 1 |
| Serranidae | | <i>Mycteroperca interstitialis</i> | 70 | 4,000 | 1,032 | 1.25E-05 | 0.0129 | 1.3 | 0 | 1.4 | 5.0 | | 0.065 | 1 |
| Ostraciidae | | <i>Acanthostracion polygonius</i> | 41 | 2,200 * | 568 | 1.28E-03 | 0.7242 | 1.2 | 0 | 1.5 | 5.4 | | 3.895 | 1 |
| Labridae | <i>Bodianus rufus</i> | 40 | 1,560 | 402 | 2.50E-04 | 0.1005 | 1.2 | 0 | 1.6 | 5.9 | | 0.588 | 1 | |
| Serranidae | <i>Alphesites afer</i> | 33 | 1,202 | 310 | 6.25E-05 | 0.0194 | 1.2 | 0 | 1.7 | 6.2 | | 0.119 | 1 | |
| Bothidae | <i>Bothus lunatus</i> | 46 / | 3,000 * | 774 | 1.25E-04 | 0.0968 | 1.1 | 0 | 1.4 | 4.9 | 0.75 | 0.478 | 1 | |
| Scorpaenidae | <i>Scorpaena plumieri</i> | 43 / | 4,330 | 1,117 | 1.38E-04 | 0.1536 | 1 | 0 | 1.2 | 4.3 | 0.44 | 0.665 | 1 | |
| Serranidae | <i>Epinephelus guttatus</i> | 58 | 2,919 | 753 | 2.50E-03 | 1.8825 | 1 | 0 | 1.3 | 4.8 | 0.68 | 8.976 | 1 | |
| Scorpaenidae | <i>Scorpaena brasiliensis</i> | 36 / | 1,600 | 413 | 1.25E-03 | 0.5163 | 1 | 0 | 1.5 | 5.4 | 0.59 | 2.775 | 1 | |
| Muraenidae | <i>Gymnothorax vicinus</i> | 122 / | 3,000 * | 774 | 1.25E-04 | 0.0968 | 0.7 * | 0 | 1.1 | 3.9 | | 0.376 | 1 | |
| Ophichthidae | <i>Ophichthus ophis</i> | 135 | 2,091 | 540 | 1.25E-03 | 0.6750 | 0.7 * | 0 | 1.1 | 4.2 | | 2.821 | 1 | |
| Muraenidae | <i>Lycodontis moringa</i> | 100 | 950 | 245 | 1.25E-05 | 0.0031 | 0.7 * | 0 | 1.3 | 4.9 | | 0.015 | 1 | |

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Table 8.7.1 continued

| Group no. | Family | Species | L (cm) | W (g) | W _{mean} (g) | Density (N·m ⁻²) | Biomass (g·m ⁻²) | Aspect ratio | Food type | Q/B% day ⁻¹ | Q/B year ⁻¹ | Natural mortality | Weighing factor | Level |
|-----------|------------------|------------------------------------|--------|---------|-----------------------|------------------------------|------------------------------|--------------|-----------|------------------------|------------------------|-------------------|-----------------|-------|
| | Muraenidae | <i>Enchelycore nigricans</i> | 100 | 900 * | 232 | 1.25E-05 | 0.0029 | 0.7 * | 0 * | 1.4 | 5.0 | | 0.014 | 1 |
| | Ophichthidae | <i>Mynichthys ocellatus</i> | 91 / | 640 * | 165 | 1.25E-05 | 0.0021 | 0.7 * | 0 * | 1.5 | 5.3 | | 0.011 | |
| | Ophichthidae | <i>Myrichthys breviceps</i> | 91 + | 640 * | 165 | 1.25E-05 | 0.0021 | 0.7 * | 0 | 1.5 | 5.3 | | 0.011 | 1 |
| | Muraenidae | <i>Echidna catenata</i> | 71 | 600 * | 155 | 1.25E-05 | 0.0019 | 0.7 * | 0 | 1.5 | 5.4 | | 0.010 | 1 |
| 14 | Pempheridae | <i>Pempheris poeyi</i> | 10 / | 30 * | 8 | 1.25E-04 | 0.0010 | 2 * | 0 * | 4.6 | 16.9 | | 0.017 | 12 |
| | Pomacentridae | <i>Chromis cyanea</i> | 13 / | 93 | 24 | 1.25E-02 | 0.3000 | 1.8 | 0 | 3.5 | 12.7 | 1.6 | 3.817 | 13 |
| | Serranidae | <i>Hypoplectrus nigricans</i> | 15 | 90 * | 23 | 1.25E-04 | 0.0029 | 1.8 | 0 | 3.5 | 12.7 | | 0.037 | 1 |
| | Serranidae | <i>Hypoplectrus aberrans</i> | 12 | 50 * | 13 | 1.25E-04 | 0.0016 | 1.8 | 0 | 4.0 | 14.6 | | 0.024 | 1 |
| | Holocentridae | <i>Plectrypops retrospinis</i> | 13 | 57 * | 15 | 1.50E-03 | 0.0225 | 1.7 | 0 | 3.8 | 13.8 | | 0.310 | 1 |
| | Serranidae | <i>Hypoplectrus puella</i> | 13 SL | 64 | 17 | 1.25E-03 | 0.0213 | 1.6 | 0 | 3.5 | 12.9 | | 0.273 | 1 |
| | Serranidae | <i>Hypoplectrus chlorurus</i> | 13 | 64 * | 17 | 1.25E-05 | 0.0002 | 1.5 | 0 | 3.5 | 12.6 | | 0.003 | 1 |
| | Labridae | <i>Xyrichtys splendens</i> | 14 | 50 * | 13 | 1.25E-02 | 0.1625 | 1.5 | 0 | 3.6 | 13.1 | | 2.122 | 15 |
| | Tetraodontidae | <i>Sphaeroides spengleri</i> | 15 | 92 | 24 | 1.39E-03 | 0.0333 | 1.6 | 0.1 | 3.6 | 13.2 | | 0.438 | 1 |
| | Labridae | <i>Xyrichtys novacula</i> | 23 | 230 * | 59 | 1.25E-02 | 0.7375 | 1.5 | 0 | 2.6 | 9.6 | | 7.084 | 1 |
| | Bothidae | <i>Bothus ocellatus</i> | 18 | 180 * | 46 | 1.25E-05 | 0.0006 | 1.5 | 0 | 2.8 | 10.2 | 3.35 | 0.006 | 1 |
| | Labridae | <i>Halichoeres gamoti</i> | 19 | 132 | 34 | 1.25E-03 | 0.0425 | 1.4 | 0 | 2.9 | 10.6 | | 0.450 | 1 |
| | Sciaenidae | <i>Pareques acuminatus</i> | 24 | 280 * | 72 | 1.25E-05 | 0.0009 | 1.3 | 0 | 2.4 | 8.8 | | 0.008 | |
| | Labridae | <i>Halichoeres bivittatus</i> | 23 | 230 * | 59 | 1.38E-02 | 0.8113 | 1.3 | 0 | 2.4 | 8.9 | | 7.207 | 1 |
| | Clinidae | <i>Labrisomus nuchipinnis</i> | 20 | 137 | 35 | 1.25E-02 | 0.4375 | 1.3 | 0 | 2.7 | 9.9 | | 4.347 | 1 |
| | Clinidae | <i>Labrisomus guppyi</i> | 20 * | 137 * | 35 | 1.25E-02 | 0.4375 | 1.3 * | 0 * | 2.7 | 9.9 | | 4.347 | 1 |
| | Labridae | <i>Halichoeres poeyi</i> | 20 | 150 * | 39 | 1.26E-02 | 0.4924 | 1.2 | 0 | 2.6 | 9.4 | | 4.604 | 1 |
| | Serranidae | <i>Serranus tabacarius</i> | 18 | 141 | 36 | 1.25E-05 | 0.0005 | 1.2 | 0 | 2.6 | 9.7 | 0.41 | 0.004 | 1 |
| | Antennariidae | <i>Antennarius multicephalus</i> | 15 (-) | 70 * | 18 | 1.25E-05 | 0.0002 | 1.2 | 0 | 3.0 | 11.0 | | 0.002 | 1 |
| | Opisthognathidae | <i>Opisthognathus macrogathus</i> | 20 - | 240 * | 62 | 1.25E-05 | 0.0008 | 1.1 | 0 | 2.3 | 8.3 | | 0.006 | 1 |
| | Labridae | <i>Thalassoma bifasciatum</i> | 17 | 97 | 25 | 2.50E-02 | 0.6250 | 1.1 | 0 | 2.6 | 9.7 | 1.73 | 6.038 | 1 |
| | Sciaenidae | <i>Equetus lanceolatus</i> | 25 | 325 | 84 | 1.25E-04 | 0.0105 | 1 | 0 | 2.0 | 7.2 | | 0.076 | 12 |
| | Scorpaenidae | <i>Scorpaena grandicomis</i> | 18 / | 212 | 55 | 6.25E-05 | 0.0034 | 1 | 0 | 2.2 | 8.1 | 1.76 | 0.028 | 1 |
| | Scorpaenidae | <i>Scorpaenodes caribbaeus</i> | 13 / | 80 * | 21 | 1.25E-04 | 0.0026 | 1 | 0 | 2.7 | 9.8 | | 0.026 | 1 |
| | Gobiidae | <i>Bathygobius soporator</i> | 15 | 70 * | 18 | 1.38E-04 | 0.0025 | 0.9 | 0 * | 2.6 | 9.5 | 0.81 | 0.024 | 1 |
| | Antennariidae | <i>Antennarius striatus</i> | 15 / | 70 * | 18 | 1.25E-04 | 0.0023 | 0.8 | 0 | 2.4 | 8.9 | | 0.020 | 1 |
| | Opisthognathidae | <i>Opisthognathus maxillosus</i> | 13 / | 65 * | 17 | 6.25E-04 | 0.0106 | 0.8 | 0 | 2.4 | 8.8 | | 0.093 | 1 |
| | Muraenidae | <i>Gymnothorax miliaris</i> | 50 | 201 | 52 | 1.25E-05 | 0.0007 | 0.7 * | 0 * | 1.8 | 6.7 | | 0.004 | 1 |
| | Congridae | <i>Heteroconger halis</i> | 48 | 150 * | 39 | 1.25E-02 | 0.4875 | 0.7 * | 0 | 1.9 | 7.1 | | 3.467 | 1 |
| 15 | Belontiidae | <i>Strongylura timucu</i> | 46 / | 110 * | 28 | 1.25E-03 | 0.0350 | 1.7 | 0 | 3.3 | 11.9 | | 0.417 | 3 |
| | Clupeidae | <i>Jenkinsia lamprotaenia</i> | 6.8 | 13 * | 3 | 1.39E+00 | 4.1625 | 2.2 | 0 * | 5.7 | 21.0 | | 87.213 | 4 |
| | Atherinidae | <i>Atherinomorus stipes</i> | 9 + | 16 | 4 | 1.25E-01 | 0.5000 | 1.9 | 0 | 5.1 | 18.6 | 3.21 | 9.299 | 3 |
| | Atherinidae | <i>Allanetta harringtonensis</i> | 8 / | 11 | 3 | 1.25E-01 | 0.3754 | 1.7 | 0 | 5.2 | 19.1 | 3.87 | 7.177 | 3 |
| 16 | Engraulidae | <i>Anchoa hepsetus</i> | 15 | 50 | 13 | 1.25E-03 | 0.0163 | 2 * | 0.7 | 10.1 | 36.8 | 1.53 | 0.599 | 3 |
| | Engraulidae | <i>Anchoa lyolepis</i> | 8 | 8 * | 2 | 1.25E-03 | 0.0025 | 1.8 | 0.7 * | 13.7 | 50.0 | 4.14 | 0.125 | 3 |
| 17 | Chaetodontidae | <i>Chaetodon aculeatus</i> | 9.5 | 19 | 5 | 1.25E-05 | 0.0001 | 1.9 | 0 | 4.9 | 17.8 | 3.82 | 0.001 | 1 |
| # | Apogonidae | <i>Apogon conklini</i> | 5 SL/ | 3.3 | 1 | 1.26E-02 | 0.0126 | 1.7 | 0 | 6.7 | 24.4 | 12.5 | 0.308 | 1 |
| | Serranidae | <i>Serranus tortugarum</i> | 8 | 12 * | 3 | 1.25E-04 | 0.0004 | 1.6 | 0 | 4.9 | 18.0 | 1.5 | 0.007 | |
| | Apogonidae | <i>Apogon maculatus</i> | 10 / | 37 | 10 | 1.25E-03 | 0.0125 | 1.5 | 0 | 3.9 | 14.1 | 4.16 | 0.177 | 1 |
| | Cirrhitidae | <i>Amblycirrhitus pinos</i> | 8 (SL) | 15 | 4 | 1.38E-04 | 0.0006 | 1.5 | 0 | 4.6 | 16.9 | | 0.009 | 1 |
| | Serranidae | <i>Serranus tigrinus</i> | 10 | 21 | 5 | 1.37E-03 | 0.0069 | 1.45 | 0 | 4.2 | 15.4 | 1.05 | 0.106 | 1 |
| | (Serranidae) | <i>Gramma melacara</i> | 10 - | 30 * | 8 | 1.25E-04 | 0.0010 | 1.4 | 0 | 3.8 | 13.9 | | 0.014 | 1 |
| | Labridae | <i>Halichoeres maculipinna</i> | 11 | 25 * | 6 | 1.25E-03 | 0.0075 | 1.3 | 0 | 3.8 | 13.8 | | 0.103 | 1 |
| | Grammidae | <i>Gramma loreto</i> | 8 + | 15 * | 4 | 1.25E-03 | 0.0050 | 1.1 | 0 | 4.0 | 14.5 | | 0.072 | 1 |
| | Scorpaenidae | <i>Scorpaena inermis</i> | >7 | 10 * | 3 | 1.25E-03 | 0.0038 | 1 * | 0 | 4.0 | 14.8 | | 0.055 | 1 |
| | Opisthognathidae | <i>Opisthognathus aurifrons</i> | 10 - | 30 * | 8 | 1.25E-03 | 0.0100 | 0.8 | 0 | 3.0 | 10.8 | | 0.108 | 1 |
| | Opisthognathidae | <i>Opisthognathus whitehurstii</i> | 8 | 15 * | 4 | 1.25E-03 | 0.0050 | 0.8 | 0 | 3.4 | 12.3 | | 0.061 | 1 |
| 18 | Serranidae | <i>Epinephelus itajara</i> | 228 | 381,644 | 98,464 | 1.25E-04 | 12.3080 | 1.3 | 0 | 0.5 | 2.0 | | 24.502 | 1 |
| | Serranidae | <i>Mycteroperca bonaci</i> | 120 | 90,000 | 23,220 | 2.50E-04 | 5.8050 | 1.2 | 0 | 0.7 | 2.6 | 0.37 | 15.114 | 1 |

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Table 8.7.1 continued

| Group no. | Family | Species | L (cm) | W (g) | W _{mean} (g) | Density (N·m ⁻²) | Biomass (g·m ⁻²) | Aspect ratio | Food type | Q/B% day ⁻¹ | Q/B year ⁻¹ | Natural mortality | Weighing factor | Level |
|-----------|------------------|-------------------------------------|---------|---------|-----------------------|------------------------------|------------------------------|--------------|-----------|------------------------|------------------------|-------------------|-----------------|-------|
| 19 | Ostraciidae | <i>Acanthostracion quadricornis</i> | 46 / | 3,050 * | 787 | 1.50E-04 | 0.1181 | 1.1 | 0.1 | 1.5 | 5.7 | | 0.667 | 1 |
| | Ostraciidae | <i>Lactophrys bicaudalis</i> | 45 | 5,572 | 1,438 | 1.50E-04 | 0.2157 | 1.2 | 0.15 | 1.5 | 5.5 | | 1.196 | 1 |
| | Monacanthidae | <i>Cantherines macrocerus</i> | 44 | 2,700 * | 697 | 1.25E-05 | 0.0087 | 1.5 | 0.03 | 1.6 | 6.0 | | 0.052 | 1 * |
| | Monacanthidae | <i>Aluterus scripta</i> | 76 | 2,941 | 759 | 1.25E-05 | 0.0095 | 0.7 | 0.4 | 1.9 | 6.8 | | 0.064 | 1 |
| | Pomacanthidae | <i>Pomacanthus paru</i> | 43 | 2,769 | 714 | 2.50E-04 | 0.1785 | 1.8 | 0.135 | 2.1 | 7.6 | 0.62 | 1.365 | 1 |
| | Pomacanthidae | <i>Holacanthus tricolor</i> | 36 | 1,306 | 337 | 2.50E-04 | 0.0843 | 1.65 | 0.008 | 2.0 | 7.2 | 0.55 | 0.609 | 1 |
| | Pomacanthidae | <i>Holacanthus ciliaris</i> | 46 | 1,988 | 513 | 1.27E-02 | 6.5407 | 1.3 | 0.014 | 1.6 | 5.9 | 0.43 | 38.581 | 1 |
| 20 | Monacanthidae | <i>Monacanthus ciliatus</i> | 20 | 448 | 116 | 1.25E-02 | 1.4500 | 1.2 | 0.3 | 3.2 | 11.7 | | 17.014 | 1 |
| | Monacanthidae | <i>Cantherines pullus</i> | 19 | 220 | 57 | 1.25E-04 | 0.0071 | 1.5 | 0.4 | 4.5 | 16.4 | | 0.117 | 2 * |
| | Monacanthidae | <i>Stephanolepis setifer</i> | 19 | 220 * | 57 | 1.25E-04 | 0.0071 | 1.25 | 0.4 * | 4.0 | 14.7 | | 0.105 | 2 |
| | Pomacentridae | <i>Stegastes variabilis</i> | 10 + | 54 | 14 | 1.25E-02 | 0.1750 | 1.5 | 0.45 | 6.3 | 23.1 | 1.6 | 4.047 | 2 * |
| | Pomacentridae | <i>Stegastes fuscus</i> | 15 / | 168 | 43 | 1.26E-01 | 5.4288 | 1.8 | 0.5 | 5.9 | 21.7 | 1.6 | 117.620 | 2 * |
| 21 | Canthigasteridae | <i>Canthigaster rostrata</i> | 11 (SL) | 82 | 21 | 1.26E-02 | 0.2651 | 1.5 | 0.2 | 4.1 | 15.0 | | 3.985 | 1 |
| | Pomacentridae | <i>Stegastes leucostictus</i> | 10 | 50 * | 13 | 1.25E-02 | 0.1625 | 1.3 | 0.2 | 4.3 | 15.8 | 1.6 | 2.563 | 1 * |
| | Pomacentridae | <i>Stegastes planifrons</i> | 11 TL | 68 | 18 | 1.25E-02 | 0.2250 | 1.4 | 0.25 | 4.4 | 16.2 | 1.41 | 3.636 | 1 * |
| | Monacanthidae | <i>Monacanthus tuckeri</i> | 10 | 24 | 6 | 1.26E-03 | 0.0076 | 1.4 | 0.3 | 5.9 | 21.5 | | 0.163 | 1 * |
| 22 # | Gobiidae | <i>Coryphopterus glaucofraenum</i> | 8 (SL) | 10.5 | 3 | 2.50E-02 | 0.0750 | 1.1 | 0.4 | 7.4 | 26.8 | 2.2 | 2.012 | 1 |
| | Gobiidae | <i>Gnatholepis thompsoni</i> | 6 SL/ | 4.1 | 1 | 1.26E-02 | 0.0126 | 1 | 0.65 | 10.7 | 39.1 | 3.48 | 0.494 | 1 |
| | Blenniidae | <i>Parablennius marmoratus</i> | 8 | 10 * | 3 | 1.25E-03 | 0.0038 | 1.3 | 0.7 | 10.9 | 39.7 | 2.53 | 0.149 | 1 |
| | Pomacanthidae | <i>Centropyge argi</i> | 7 | 10 * | 3 | 1.25E-04 | 0.0004 | 1.4 | 0.75 | 12.6 | 45.9 | | 0.017 | 1 |
| | Blenniidae | <i>Entomacrodus nigricans</i> | 8.5 | 13 * | 3 | 1.25E-03 | 0.0038 | 1 | 0.8 | 11.2 | 40.7 | | 0.153 | 1 |
| 23 | Scaridae | <i>Scarus guacamaia</i> | 99 | 23,000 | 5,934 | 1.26E-02 | 74.9168 | 1.2 | 1 | 3.3 | 12.0 | 0.61 | 895.357 | 1 |
| | Scaridae | <i>Scarus coelestinus</i> | 80 | 8,556 | 2,207 | 1.25E-04 | 0.2759 | 1.1 | 1 | 3.7 | 13.5 | | 3.733 | 1 |
| | Scaridae | <i>Scarus vetula</i> | 54 SL | 5,558 | 1,434 | 1.38E-02 | 19.7175 | 1.05 | 1 | 3.9 | 14.4 | 1.09 | 283.385 | 1 |
| 24 | Scaridae | <i>Scarus coeruleus</i> | 62 | 3,720 | 960 | 1.25E-04 | 0.1200 | 1.5 | 1 * | 5.3 | 19.3 | 0.70 | 2.312 | 1 * |
| | Scaridae | <i>Sparisoma rubripinne</i> | 46.5 | 2,734 | 705 | 1.25E-03 | 0.8813 | 1.5 | 1 | 5.7 | 20.8 | 1.11 | 18.303 | 1 |
| | Scaridae | <i>Sparisoma viride</i> | 50 FL | 2,430 | 627 | 1.37E-03 | 0.8621 | 1.4 | 1 | 5.7 | 20.7 | | 17.812 | 1 |
| | Scaridae | <i>Sparisoma chrysopterum</i> | 42 FL | 1,510 | 390 | 1.38E-02 | 5.3625 | 1.4 | 1 | 6.3 | 22.8 | 1.38 | 122.349 | 1 |
| | Scaridae | <i>Scarus taeniopterus</i> | 35 FL | 1,102 | 284 | 1.25E-04 | 0.0355 | 1.1 | 1 | 5.7 | 20.8 | 1.20 | 0.739 | 1 |
| | Monacanthidae | <i>Aluterus schoepfii</i> | 51 / | 1,892 | 488 | 1.25E-03 | 0.6100 | 0.7 | 1 | 4.1 | 15.1 | | 9.205 | 1 3 |
| 25 | Scaridae | <i>Scarus iserti</i> | 20 FL | 141 | 36 | 2.50E-03 | 0.0900 | 1.4 | 1 | 10.1 | 36.8 | | 3.313 | 1 |
| | Scaridae | <i>Sparisoma aurofrenatum</i> | 26 FL | 324 | 84 | 1.25E-03 | 0.1050 | 1.3 | 1 | 8.1 | 29.5 | 0.94 | 3.094 | 1 |
| | Scaridae | <i>Sparisoma radians</i> | 20 | 150 * | 39 | 1.25E-01 | 4.8799 | 1.3 | 1 | 9.3 | 33.9 | | 165.353 | 1 |
| 26 | Blenniidae | <i>Scartella cristata</i> | 11 | 27 * | 7 | 1.25E-04 | 0.0009 | 1.2 | 0.9 | 10.8 | 39.3 | | 0.034 | 1 |
| | Blenniidae | <i>Ophioblennius atlanticus</i> | 12 / | 35 * | 9 | 1.25E-03 | 0.0113 | 0.9 | 0.9 | 9.0 | 32.9 | 2.84 | 0.370 | 1 |
| 27 | Gobiidae | <i>Gobiosoma horsti</i> | 5 (SL) | 3.1 | 0.8 | 1.25E-01 | 0.1001 | 0.85 * | 0 * | 4.7 | 17.2 | 3.14 | 1.721 | 1 |
| | Gobiidae | <i>Gobiosoma evelynae</i> (sp.) | 5 (SL) | 2.2 | 0.6 | 1.25E-01 | 0.0751 | 0.8 | 0 | 4.8 | 17.7 | 3.14 | 1.329 | 1 |

Symbol:

= single species grouped outside cluster analysis.

* = estimated values.

1 = demersal, 2 = off bottom, 3 = midwater, 4 = pelagic.

Appendix 8.8. Food intake and electivity indices for each consumer and all models.

Table 8.8.1. The 50-box model.

| | | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|-----------|---------------------|------------------|-------------|--------------------------|---------------------------|
| Predator: | (1) LGSHARKS/RAYSC | | | | |
| | (0' Cycle) | | | | |
| 1. | LGSHARKS/RAYSC | 0.010 | 0.01 | 0.988 | 0.029 |
| 2. | SHARKS/SCOMBRID | 0.010 | 0.01 | 0.984 | -0.134 |
| 3. | LGJACKSC | 0.003 | 0.00 | 0.976 | -0.315 |
| 4. | IMJACKSC | 0.050 | 0.07 | 0.987 | -0.013 |
| 5. | SMJACKSC | 0.004 | 0.01 | 0.906 | -0.778 |
| 6. | IMREEF FISHC1 | 0.100 | 0.15 | 0.988 | 0.025 |
| 7. | LGIMSCHOOLFISHP | 0.010 | 0.01 | 0.638 | -0.946 |
| 8. | IMREEF FISHC2 | 0.010 | 0.01 | 0.722 | -0.908 |
| 9. | HEMIRAMPHIDAEH | 0.002 | 0.00 | 0.799 | -0.895 |
| 10. | KYPHOSIDAEH | 0.060 | 0.09 | 0.984 | -0.121 |
| 11. | IMREEF FISHH | 0.052 | 0.08 | 0.929 | -0.712 |
| 12. | LGREEF FISHC | 0.030 | 0.04 | 0.967 | -0.457 |
| 13. | IMREEF FISHC3 | 0.025 | 0.04 | 0.891 | -0.806 |
| 14. | SMREEF FISHC1 | 0.020 | 0.03 | 0.916 | -0.752 |
| 15. | SMSCHOOLFISHP | 0.049 | 0.07 | 0.921 | -0.738 |
| 16. | ENGRAULIDAEH | 0.028 | 0.04 | 0.954 | -0.583 |
| 17. | SMREEF FISHC2 | 0.006 | 0.01 | 0.918 | -0.748 |
| 18. | LGGROUPERSC | 0.093 | 0.14 | 0.997 | 0.626 |
| 19. | IMREEF FISHC4 | 0.020 | 0.03 | 0.972 | -0.390 |
| 20. | SMREEF FISHO1 | 0.025 | 0.04 | 0.946 | -0.638 |
| 21. | SMREEF FISHO2 | 0.004 | 0.01 | 0.907 | -0.776 |
| 22. | SMREEF FISHO3 | 0.001 | 0.00 | 0.680 | -0.937 |
| 23. | LGSCARIDAEH | 0.025 | 0.04 | 0.739 | -0.921 |
| 24. | IMSCARIDAEH | 0.002 | 0.00 | 0.294 | -0.978 |
| 25. | SMSCARIDAEH | 0.001 | 0.00 | -0.003 | -0.988 |
| 28. | SEABIRDS | 0.020 | 0.03 | 1.000 | 0.986 |
| 29. | SQUIDS | 0.005 | 0.01 | 0.888 | -0.812 |
| 30. | SEATURTLES | 0.005 | 0.01 | 0.961 | -0.523 |
| 31. | OCTOPUSES | 0.020 | 0.03 | 0.846 | -0.862 |
| 32. | LOBSTERS | 0.005 | 0.01 | 0.771 | -0.909 |
| 33. | CRABS | 0.050 | 0.07 | 0.860 | -0.848 |
| 34. | SHRMP/HCRB/STOM | 0.050 | 0.07 | 0.924 | -0.730 |
| 37. | ECHINIDS | 0.008 | 0.01 | -0.425 | -0.995 |
| 38. | GASTROPODS | 0.010 | 0.01 | 0.038 | -0.987 |
| 40. | POLY/PRIA/OPHIU | 0.050 | 0.07 | 0.769 | -0.910 |
| 41. | HOL/SIP/ECH/HEM | 0.050 | 0.07 | 0.584 | -0.954 |
| 42. | BIVALVES | 0.067 | 0.10 | 0.512 | -0.963 |
| 50. | Detritus | 0.020 | 0.03 | -0.904 | -0.999 |
| TOTAL | | 1.000 | 1.47 | | |
| Predator: | (2) SHARKS/SCOMBRID | | | | |
| | (0' Cycle) | | | | |
| 2. | SHARKS/SCOMBRID | 0.010 | 0.04 | 0.984 | 0.589 |
| 4. | IMJACKSC | 0.034 | 0.13 | 0.981 | 0.536 |
| 5. | SMJACKSC | 0.050 | 0.19 | 0.992 | 0.795 |
| 6. | IMREEF FISHC1 | 0.015 | 0.06 | 0.923 | -0.144 |
| 7. | LGIMSCHOOLFISHP | 0.176 | 0.67 | 0.975 | 0.421 |
| 8. | IMREEF FISHC2 | 0.040 | 0.15 | 0.938 | -0.035 |
| 9. | HEMIRAMPHIDAEH | 0.115 | 0.44 | 0.996 | 0.913 |
| 10. | KYPHOSIDAEH | 0.020 | 0.08 | 0.953 | 0.115 |
| 11. | IMREEF FISHH | 0.020 | 0.08 | 0.825 | -0.527 |
| 12. | LGREEF FISHC | 0.014 | 0.05 | 0.931 | -0.088 |
| 13. | IMREEF FISHC3 | 0.084 | 0.32 | 0.966 | 0.281 |
| 14. | SMREEF FISHC1 | 0.020 | 0.08 | 0.916 | -0.189 |
| 15. | SMSCHOOLFISHP | 0.136 | 0.52 | 0.971 | 0.350 |
| 16. | ENGRAULIDAEH | 0.040 | 0.15 | 0.968 | 0.299 |
| 17. | SMREEF FISHC2 | 0.020 | 0.08 | 0.975 | 0.411 |
| 19. | IMREEF FISHC4 | 0.006 | 0.02 | 0.910 | -0.226 |
| 20. | SMREEF FISHO1 | 0.014 | 0.05 | 0.905 | -0.254 |
| 21. | SMREEF FISHO2 | 0.001 | 0.00 | 0.672 | -0.738 |
| 23. | LGSCARIDAEH | 0.030 | 0.11 | 0.777 | -0.618 |
| 24. | IMSCARIDAEH | 0.001 | 0.00 | -0.044 | -0.947 |
| 25. | SMSCARIDAEH | 0.002 | 0.01 | 0.331 | -0.889 |
| 26. | BLENNIIDAEH | 0.001 | 0.00 | 0.788 | -0.602 |

continued...

Table 8.8.1 continued

| | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|----------------------------------------|------------------|-------------|--------------------------|---------------------------|
| 29. SQUIDS | 0.010 | 0.04 | 0.942 | 0.005 |
| 31. OCTOPUSES | 0.045 | 0.17 | 0.929 | -0.106 |
| 32. LOBSTERS | 0.005 | 0.02 | 0.771 | -0.628 |
| 33. CRABS | 0.020 | 0.08 | 0.683 | -0.729 |
| 34. SHRMP/HCRB/STOM | 0.050 | 0.19 | 0.924 | -0.140 |
| 38. GASTROPODS | 0.008 | 0.03 | -0.074 | -0.950 |
| 40. POLY/PRIA/OPHIU | 0.006 | 0.02 | -0.043 | -0.947 |
| 42. BIVALVES | 0.007 | 0.03 | -0.511 | -0.981 |
| TOTAL | 1.000 | 3.79 | | |
| Predator: (3) LGJACKSC | | | | |
| 4. IMJACKSC | 0.150 | 0.15 | 0.996 | 0.948 |
| 6. IMREEF FISHC1 | 0.200 | 0.21 | 0.994 | 0.914 |
| 8. IMREEF FISHC2 | 0.160 | 0.17 | 0.984 | 0.732 |
| 11. IMREEF FISHH | 0.044 | 0.05 | 0.917 | 0.039 |
| 13. IMREEF FISHC3 | 0.124 | 0.13 | 0.977 | 0.624 |
| 33. CRABS | 0.010 | 0.01 | 0.453 | -0.782 |
| 34. SHRMP/HCRB/STOM | 0.015 | 0.02 | 0.767 | -0.481 |
| 37. ECHINOIDS | 0.082 | 0.08 | 0.611 | -0.679 |
| 38. GASTROPODS | 0.158 | 0.16 | 0.889 | -0.114 |
| 42. BIVALVES | 0.057 | 0.06 | 0.450 | -0.784 |
| TOTAL | 1.000 | 1.03 | | |
| Predator: (4) IMJACKSC | | | | |
| 6. IMREEF FISHC1 | 0.001 | 0.01 | 0.251 | -0.910 |
| 7. LGMSCHOOLFISHP | 0.056 | 0.84 | 0.924 | -0.159 |
| 8. IMREEF FISHC2 | 0.110 | 1.64 | 0.977 | 0.435 |
| 10. KYPHOSIDAEH | 0.015 | 0.22 | 0.938 | -0.053 |
| 11. IMREEF FISHH | 0.107 | 1.60 | 0.965 | 0.240 |
| 15. SMSCHOOLFISHPEL | 0.110 | 1.64 | 0.964 | 0.229 |
| 16. ENGRAULIDAEH | 0.107 | 1.60 | 0.988 | 0.670 |
| 17. SMREEF FISHC2 | 0.001 | 0.01 | 0.590 | -0.802 |
| 19. IMREEF FISHC4 | 0.005 | 0.07 | 0.893 | -0.331 |
| 20. SMREEF FISHO1 | 0.053 | 0.79 | 0.974 | 0.382 |
| 21. SMREEF FISHO2 | 0.053 | 0.79 | 0.993 | 0.801 |
| 22. SMREEF FISHO3 | 0.053 | 0.79 | 0.993 | 0.807 |
| 23. LGSCARIDAEH | 0.016 | 0.24 | 0.619 | -0.785 |
| 24. IMSCARIDAEH | 0.073 | 1.09 | 0.971 | 0.325 |
| 25. SMSCARIDAEH | 0.122 | 1.82 | 0.984 | 0.573 |
| 26. BLENNIIDAEH | 0.053 | 0.79 | 0.996 | 0.889 |
| 29. SQUIDS | 0.004 | 0.06 | 0.862 | -0.446 |
| 33. CRABS | 0.004 | 0.06 | 0.030 | -0.942 |
| 34. SHRMP/HCRB/STOM | 0.031 | 0.46 | 0.880 | -0.383 |
| 38. GASTROPODS | 0.004 | 0.06 | -0.397 | -0.976 |
| 46. ZOOPLANKTON | 0.022 | 0.33 | 0.553 | -0.821 |
| TOTAL | 1.000 | 14.91 | | |
| Predator: (6) IMREEF FISHC1 (0' Cycle) | | | | |
| 6. IMREEF FISHC1 | 0.001 | 0.02 | 0.251 | -0.550 |
| 8. IMREEF FISHC2 | 0.009 | 0.19 | 0.750 | 0.108 |
| 10. KYPHOSIDAEH | 0.003 | 0.06 | 0.724 | 0.051 |
| 11. IMREEF FISHH | 0.004 | 0.08 | 0.353 | -0.465 |
| 13. IMREEF FISHC3 | 0.057 | 1.18 | 0.951 | 0.778 |
| 14. SMREEF FISHC1 | 0.005 | 0.10 | 0.702 | 0.005 |
| 15. SMSCHOOLFISHPEL | 0.062 | 1.28 | 0.937 | 0.714 |
| 17. SMREEF FISHC2 | 0.003 | 0.06 | 0.842 | 0.356 |
| 19. IMREEF FISHC4 | 0.003 | 0.06 | 0.827 | 0.311 |
| 20. SMREEF FISHO1 | 0.020 | 0.41 | 0.932 | 0.692 |
| 23. LGSCARIDAEH | 0.030 | 0.62 | 0.777 | 0.174 |
| 24. IMSCARIDAEH | 0.016 | 0.33 | 0.872 | 0.456 |
| 25. SMSCARIDAEH | 0.016 | 0.33 | 0.882 | 0.490 |
| 29. SQUIDS | 0.001 | 0.02 | 0.542 | -0.258 |
| 31. OCTOPUSES | 0.014 | 0.29 | 0.788 | 0.200 |
| 33. CRABS | 0.090 | 1.86 | 0.920 | 0.638 |
| 34. SHRMP/HCRB/STOM | 0.030 | 0.62 | 0.876 | 0.469 |
| 36. ASTEROIDS | 0.001 | 0.02 | -0.664 | -0.932 |
| 37. ECHINOIDS | 0.050 | 1.04 | 0.432 | -0.388 |

continued...

Table 8.8.1 continued

| | | Diet composition | Food intake | Ivlev's elec- tivity index | Standardized forage ratio |
|------------------------------------------|---------------------|---------------------|----------------|-------------------------------|------------------------------|
| 38. | GASTROPODS | 0.017 | 0.35 | 0.294 | -0.516 |
| 40. | POLY/PRIA/OPHIU | 0.006 | 0.12 | -0.043 | -0.725 |
| 41. | HOL/SIP/ECH/HEM | 0.003 | 0.06 | -0.628 | -0.924 |
| 42. | BIVALVES | 0.010 | 0.21 | -0.368 | -0.852 |
| 43. | ASCI/BARN/BRYOZ | 0.060 | 1.24 | 0.376 | -0.444 |
| 44. | SPONGES | 0.200 | 4.14 | 0.116 | -0.640 |
| 45. | CORALS/ANEMONES | 0.010 | 0.21 | -0.411 | -0.865 |
| 46. | ZOOPLANKTON | 0.250 | 5.18 | 0.951 | 0.776 |
| 49. | Benthic prod. | 0.029 | 0.60 | -0.798 | -0.962 |
| TOTAL | | 1.000 | 20.71 | | |
| Predator: (5) SMJACKSC | | | | | |
| 4. | IMJACKSC | 0.002 | 0.03 | 0.722 | 0.202 |
| 5. | SMJACKSC (0' Cycle) | 0.001 | 0.01 | 0.669 | 0.099 |
| 7. | LGIMSCHOOLFISHP | 0.014 | 0.18 | 0.727 | 0.212 |
| 11. | IMREEF FISHH | 0.020 | 0.25 | 0.825 | 0.444 |
| 15. | SMSCHOOLFISHPEL | 0.006 | 0.08 | 0.498 | -0.167 |
| 16. | ENGRAULIDAEH | 0.019 | 0.24 | 0.933 | 0.775 |
| 46. | ZOOPLANKTON | 0.938 | 11.73 | 0.987 | 0.984 |
| TOTAL | | 1.000 | 12.50 | | |
| Predator: (7) LGIMSCHOOLFISHP (0' Cycle) | | | | | |
| 7. | LGIMSCHOOLFISHP | 0.003 | 0.42 | 0.151 | -0.444 |
| 11. | IMREEF FISHH | 0.010 | 1.39 | 0.679 | 0.207 |
| 15. | SMSCHOOLFISHPEL | 0.047 | 6.51 | 0.918 | 0.768 |
| 16. | ENGRAULIDAEH | 0.006 | 0.83 | 0.802 | 0.461 |
| 34. | SHRMP/HCRB/STOM | 0.004 | 0.55 | 0.338 | -0.269 |
| 40. | POLY/PRIA/OPHIU | 0.120 | 16.62 | 0.897 | 0.705 |
| 46. | ZOOPLANKTON | 0.722 | 100.02 | 0.983 | 0.979 |
| 49. | Benthic prod. | 0.088 | 12.19 | -0.491 | -0.824 |
| TOTAL | | 1.000 | 138.53 | | |
| Predator: (8) IMREEF FISHC2 (0' Cycle) | | | | | |
| 4. | IMJACKSC | 0.001 | 0.07 | 0.512 | -0.266 |
| 7. | LGIMSCHOOLFISHP | 0.001 | 0.07 | -0.378 | -0.845 |
| 8. | IMREEF FISHC2 | 0.020 | 1.34 | 0.879 | 0.507 |
| 10. | KYPHOSIDAEH | 0.003 | 0.20 | 0.724 | 0.085 |
| 11. | IMREEF FISHH | 0.004 | 0.27 | 0.353 | -0.439 |
| 12. | LGREEF FISHC | 0.001 | 0.07 | 0.333 | -0.457 |
| 13. | IMREEF FISHC3 | 0.015 | 1.00 | 0.825 | 0.336 |
| 15. | SMSCHOOLFISHPEL | 0.067 | 4.48 | 0.942 | 0.752 |
| 16. | ENGRAULIDAEH | 0.002 | 0.13 | 0.505 | -0.275 |
| 17. | SMREEF FISHC2 | 0.001 | 0.07 | 0.590 | -0.157 |
| 19. | IMREEF FISHC4 | 0.003 | 0.20 | 0.827 | 0.342 |
| 23. | LGSCARIDAEH | 0.020 | 1.34 | 0.683 | 0.002 |
| 24. | IMSCARIDAEH | 0.030 | 2.01 | 0.930 | 0.700 |
| 25. | SMSCARIDAEH | 0.010 | 0.67 | 0.817 | 0.313 |
| 31. | OCTOPUSES | 0.050 | 3.34 | 0.936 | 0.725 |
| 32. | LOBSTERS | 0.001 | 0.07 | 0.214 | -0.554 |
| 33. | CRABS | 0.100 | 6.69 | 0.927 | 0.691 |
| 34. | SHRMP/HCRB/STOM | 0.020 | 1.34 | 0.820 | 0.320 |
| 35. | SMBENTHARTHROPO | 0.031 | 2.07 | 0.844 | 0.392 |
| 37. | ECHINOIDS | 0.100 | 6.69 | 0.669 | -0.024 |
| 38. | GASTROPODS | 0.035 | 2.34 | 0.581 | -0.170 |
| 39. | CHITONS/SCAPHOP | 0.020 | 1.34 | 0.239 | -0.535 |
| 40. | POLY/PRIA/OPHIU | 0.180 | 12.04 | 0.930 | 0.701 |
| 41. | HOL/SIP/ECH/HEM | 0.060 | 4.01 | 0.641 | -0.075 |
| 42. | BIVALVES | 0.073 | 4.88 | 0.543 | -0.225 |
| 43. | ASCI/BARN/BRYOZ | 0.050 | 3.34 | 0.295 | -0.490 |
| 44. | SPONGES | 0.070 | 4.68 | -0.387 | -0.849 |
| 45. | CORALS/ANEMONES | 0.004 | 0.27 | -0.714 | -0.940 |
| 46. | ZOOPLANKTON | 0.018 | 1.20 | 0.479 | -0.306 |
| 49. | Benthic prod. | 0.010 | 0.67 | -0.925 | -0.986 |
| TOTAL | | 1.000 | 66.88 | | |

continued...

Table 8.8.1 continued

| | | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|---------------------|--------------------|------------------|-------------|--------------------------|---------------------------|
| Predator: | (9) HEMIRAMPHIDAEH | | | | |
| 15. SMSCHOOLFISHPEL | | 0.200 | 8.80 | 0.980 | 0.999 |
| 49. Benthic prod. | | 0.800 | 35.19 | 0.513 | 0.209 |
| TOTAL | | 1.000 | 43.99 | | |
| Predator: | (10) KYPHOSIDAEH | | | | |
| 49. Benthic prod. | | 1.000 | 57.11 | 0.590 | 1.000 |
| TOTAL | | 1.000 | 57.11 | | |
| Predator: | (11) IMREEF FISHH | | | | |
| 34. SHRMP/HCRB/StOM | | 0.004 | 0.99 | 0.338 | 0.927 |
| 38. GASTROPODS | | 0.001 | 0.25 | -0.805 | -0.032 |
| 40. POLY/PRIA/OPHIU | | 0.002 | 0.50 | -0.531 | 0.468 |
| 45. CORALS/ANEMONES | | 0.002 | 0.50 | -0.846 | -0.161 |
| 49. Benthic prod. | | 0.533 | 131.94 | 0.348 | 0.930 |
| 50. Detritus | | 0.458 | 113.38 | 0.072 | 0.850 |
| TOTAL | | 1.000 | 247.55 | | |
| Predator: | (12) LGREEF FISHC | | | | |
| 3. LGJACKSC | | 0.004 | 0.04 | 0.982 | 0.819 |
| 4. IMJACKSC | | 0.024 | 0.24 | 0.973 | 0.725 |
| 5. SMJACKSC | | 0.020 | 0.20 | 0.980 | 0.798 |
| 6. IMREEF FISHC1 | | 0.030 | 0.30 | 0.961 | 0.604 |
| 7. LGIMSCHOOLFISHP | | 0.040 | 0.39 | 0.895 | 0.163 |
| 8. IMREEF FISHC2 | | 0.040 | 0.39 | 0.938 | 0.418 |
| 10. KYPHOSIDAEH | | 0.003 | 0.03 | 0.724 | -0.359 |
| 11. IMREEF FISHH | | 0.023 | 0.23 | 0.847 | -0.044 |
| 12. LGREEF FISHC | (0' Cycle) | 0.005 | 0.05 | 0.818 | -0.138 |
| 13. IMREEF FISHC3 | | 0.040 | 0.39 | 0.931 | 0.370 |
| 14. SMREEF FISHC1 | | 0.001 | 0.01 | 0.066 | -0.843 |
| 15. SMSCHOOLFISHPEL | | 0.059 | 0.58 | 0.934 | 0.393 |
| 16. ENGRAULIDAEH | | 0.001 | 0.01 | 0.206 | -0.796 |
| 19. IMREEF FISHC4 | | 0.003 | 0.03 | 0.827 | -0.109 |
| 20. SMREEF FISHO1 | | 0.017 | 0.17 | 0.921 | 0.306 |
| 21. SMREEF FISHO2 | | 0.006 | 0.06 | 0.937 | 0.411 |
| 23. LGSCARIDAEH | | 0.010 | 0.10 | 0.453 | -0.667 |
| 24. IMSCARIDAEH | | 0.010 | 0.10 | 0.803 | -0.181 |
| 25. SMSCARIDAEH | | 0.008 | 0.08 | 0.777 | -0.249 |
| 26. BLENNIIDAEH | | 0.001 | 0.01 | 0.788 | -0.222 |
| 29. SQUIDS | | 0.001 | 0.01 | 0.542 | -0.597 |
| 31. OCTOPUSES | | 0.009 | 0.09 | 0.688 | -0.421 |
| 32. LOBSTERS | | 0.002 | 0.02 | 0.511 | -0.624 |
| 33. CRABS | | 0.032 | 0.32 | 0.790 | -0.217 |
| 34. SHRMP/HCRB/StOM | | 0.030 | 0.30 | 0.876 | 0.073 |
| 37. ECHINOIDS | | 0.100 | 0.98 | 0.669 | -0.449 |
| 38. GASTROPODS | | 0.400 | 3.94 | 0.955 | 0.551 |
| 42. BIVALVES | | 0.075 | 0.74 | 0.552 | -0.587 |
| 46. ZOOPLANKTON | | 0.006 | 0.06 | -0.027 | -0.868 |
| TOTAL | | 1.000 | 9.85 | | |
| Predator: | (13) IMREEF FISHC3 | | | | |
| 5. SMJACKSC | | 0.003 | 0.14 | 0.876 | 0.128 |
| 6. IMREEF FISHC1 | | 0.010 | 0.46 | 0.887 | 0.177 |
| 7. LGIMSCHOOLFISHP | | 0.002 | 0.09 | -0.051 | -0.860 |
| 8. IMREEF FISHC2 | | 0.010 | 0.46 | 0.772 | -0.208 |
| 10. KYPHOSIDAEH | | 0.003 | 0.14 | 0.724 | -0.310 |
| 11. IMREEF FISHH | | 0.009 | 0.42 | 0.650 | -0.434 |
| 13. IMREEF FISHC3 | (0' Cycle) | 0.012 | 0.56 | 0.786 | -0.173 |
| 14. SMREEF FISHC1 | | 0.010 | 0.46 | 0.839 | -0.016 |
| 15. SMSCHOOLFISHPEL | | 0.002 | 0.09 | -0.002 | -0.847 |
| 16. ENGRAULIDAEH | | 0.002 | 0.09 | 0.505 | -0.595 |
| 17. SMREEF FISHC2 | | 0.010 | 0.46 | 0.950 | 0.551 |
| 19. IMREEF FISHC4 | | 0.003 | 0.14 | 0.827 | -0.055 |
| 20. SMREEF FISHO1 | | 0.040 | 1.86 | 0.966 | 0.681 |
| 21. SMREEF FISHO2 | | 0.001 | 0.05 | 0.672 | -0.401 |
| 22. SMREEF FISHO3 | | 0.015 | 0.70 | 0.975 | 0.767 |

continued...

Table 8.8.1 continued

| | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|-------------------------------------------|------------------|-------------|--------------------------|---------------------------|
| 23. LGSCARIDAEH | 0.055 | 2.55 | 0.872 | 0.110 |
| 24. IMSCARIDAEH | 0.007 | 0.32 | 0.730 | -0.299 |
| 25. SMSCARIDAEH | 0.006 | 0.28 | 0.713 | -0.332 |
| 26. BLENNIIDAEH | 0.002 | 0.09 | 0.888 | 0.181 |
| 27. SMGOBIIDAEH | 0.004 | 0.19 | 0.983 | 0.845 |
| 31. OCTOPUSES | 0.030 | 1.39 | 0.895 | 0.215 |
| 33. CRABS | 0.150 | 6.96 | 0.951 | 0.561 |
| 34. SHRMP/HCRB/STOM | 0.100 | 4.64 | 0.961 | 0.642 |
| 35. SMBENTHARTHROPO | 0.042 | 1.95 | 0.882 | 0.155 |
| 37. ECHINOIDS | 0.060 | 2.78 | 0.504 | -0.596 |
| 38. GASTROPODS | 0.004 | 0.19 | -0.397 | -0.931 |
| 39. CHITONS/SCAPHOP | 0.010 | 0.46 | -0.102 | -0.873 |
| 40. POLY/PRIA/OPHIU | 0.050 | 2.32 | 0.769 | -0.216 |
| 41. HOL/SIP/ECH/HEM | 0.010 | 0.46 | -0.135 | -0.881 |
| 42. BIVALVES | 0.023 | 1.07 | 0.030 | -0.837 |
| 43. ASCI/BARN/BRYOZ | 0.040 | 1.86 | 0.190 | -0.782 |
| 44. SPONGES | 0.070 | 3.25 | -0.387 | -0.929 |
| 45. CORALS/ANEMONES | 0.003 | 0.14 | -0.778 | -0.979 |
| 46. ZOOPLANKTON | 0.150 | 6.96 | 0.919 | 0.344 |
| 49. Benthic prod. | 0.052 | 2.41 | -0.664 | -0.967 |
| TOTAL | 1.000 | 46.40 | | |
| Predator: (14) SMREEF FISHC1 | | | | |
| 7. LGIMSCHOOLFISHP | 0.002 | 0.09 | -0.051 | -0.765 |
| 8. IMREEF FISHC2 | 0.002 | 0.09 | 0.217 | -0.626 |
| 9. HEMIRAMPHIDAEH | 0.002 | 0.09 | 0.799 | 0.152 |
| 14. SMREEF FISHC1 (0' Cycle) | 0.003 | 0.13 | 0.548 | -0.325 |
| 15. SMSCHOOLFISHPEL | 0.002 | 0.09 | -0.002 | -0.744 |
| 16. ENGRAULIDAEH | 0.002 | 0.09 | 0.505 | -0.378 |
| 17. SMREEF FISHC2 | 0.002 | 0.09 | 0.772 | 0.079 |
| 20. SMREEF FISHO1 | 0.005 | 0.22 | 0.754 | 0.036 |
| 21. SMREEF FISHO2 | 0.002 | 0.09 | 0.821 | 0.215 |
| 22. SMREEF FISHO3 | 0.006 | 0.26 | 0.939 | 0.674 |
| 25. SMSCARIDAEH | 0.002 | 0.09 | 0.331 | -0.545 |
| 26. BLENNIIDAEH | 0.006 | 0.26 | 0.961 | 0.795 |
| 27. SMGOBIIDAEH | 0.002 | 0.09 | 0.966 | 0.823 |
| 33. CRABS | 0.100 | 4.33 | 0.927 | 0.619 |
| 34. SHRMP/HCRB/STOM | 0.040 | 1.73 | 0.906 | 0.520 |
| 35. SMBENTHARTHROPO | 0.042 | 1.82 | 0.882 | 0.425 |
| 37. ECHINOIDS | 0.110 | 4.76 | 0.695 | -0.092 |
| 38. GASTROPODS | 0.144 | 6.24 | 0.879 | 0.412 |
| 39. CHITONS/SCAPHOP | 0.040 | 1.73 | 0.530 | -0.347 |
| 40. POLY/PRIA/OPHIU | 0.100 | 4.33 | 0.877 | 0.405 |
| 41. HOL/SIP/ECH/HEM | 0.008 | 0.35 | -0.242 | -0.835 |
| 42. BIVALVES | 0.060 | 2.60 | 0.470 | -0.417 |
| 46. ZOOPLANKTON | 0.260 | 11.26 | 0.952 | 0.746 |
| 49. Benthic prod. | 0.058 | 2.51 | -0.632 | -0.936 |
| TOTAL | 1.000 | 43.31 | | |
| Predator: (15) SMSCHOOLFISHPEL (0' Cycle) | | | | |
| 15. SMSCHOOLFISHPEL | 0.002 | 0.38 | -0.002 | -0.535 |
| 16. ENGRAULIDAEH | 0.002 | 0.38 | 0.505 | -0.033 |
| 34. SHRMP/HCRB/STOM | 0.010 | 1.91 | 0.669 | 0.223 |
| 43. ASCI/BARN/BRYOZ | 0.030 | 5.74 | 0.049 | -0.498 |
| 46. ZOOPLANKTON | 0.950 | 181.69 | 0.987 | 0.997 |
| 47. DECOMP/MICROFAU | 0.006 | 1.15 | 0.338 | -0.236 |
| TOTAL | 1.000 | 191.25 | | |
| Predator: (16) ENGRAULIDAEH | | | | |
| 46. ZOOPLANKTON | 0.200 | 28.86 | 0.939 | 0.892 |
| 48. Phytoplankton | 0.700 | 101.01 | 0.978 | 0.985 |
| 50. Detritus | 0.100 | 14.43 | -0.597 | -0.813 |
| TOTAL | 1.000 | 144.31 | | |
| Predator: (17) SMREEF FISHC2 | | | | |
| 8. IMREEF FISHC2 | 0.002 | 0.04 | 0.217 | -0.639 |
| 11. IMREEF FISHH | 0.001 | 0.02 | -0.313 | -0.862 |

continued...

Table 8.8.1 continued

| | | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|------------------------------|---------------------------|------------------|-------------|--------------------------|---------------------------|
| 14. | SMREEF FISHC1 | 0.008 | 0.15 | 0.803 | 0.140 |
| 17. | SMREEF FISHC2 | 0.003 | 0.06 | 0.842 | 0.260 |
| 20. | SMREEF FISHO1 | 0.001 | 0.02 | 0.176 | -0.663 |
| 22. | SMREEF FISHO ³ | 0.004 | 0.08 | 0.909 | 0.519 |
| 26. | BLENNIIDAEH | 0.001 | 0.02 | 0.788 | 0.098 |
| 27. | SMGOBIIDAEC | 0.002 | 0.04 | 0.966 | 0.814 |
| 31. | OCTOPUSES | 0.020 | 0.38 | 0.846 | 0.275 |
| 33. | CRABS | 0.063 | 1.20 | 0.887 | 0.426 |
| 34. | SHRMP/HCRB/StOM | 0.150 | 2.86 | 0.974 | 0.864 |
| 35. | SMBENTHARTHROPO | 0.100 | 1.90 | 0.949 | 0.716 |
| 38. | GASTROPODS | 0.005 | 0.10 | -0.299 | -0.858 |
| 39. | CHITONS/SCAPHOP | 0.002 | 0.04 | -0.720 | -0.955 |
| 40. | POLY/PRIA/OPHIU | 0.079 | 1.50 | 0.847 | 0.277 |
| 41. | HOL/SIP/ECH/HEM | 0.001 | 0.02 | -0.858 | -0.979 |
| 42. | BIVALVES | 0.030 | 0.57 | 0.162 | -0.672 |
| 46. | ZOOPLANKTON | 0.500 | 9.52 | 0.975 | 0.870 |
| 49. | Benthic prod. | 0.028 | 0.53 | -0.804 | -0.970 |
| TOTAL | | 1.000 | 19.04 | | |
| Predator: (18) LGGROUPERSC | | | | | |
| 1. | LGS SHARKS/RAYSC | 0.005 | 0.01 | 0.977 | 0.443 |
| 2. | SHARKS/SCOMBRID | 0.005 | 0.01 | 0.968 | 0.299 |
| 4. | IMJACKSC | 0.010 | 0.02 | 0.937 | -0.040 |
| 5. | SMJACKSC | 0.010 | 0.02 | 0.961 | 0.207 |
| 6. | IMREEF FISHC1 | 0.085 | 0.14 | 0.986 | 0.639 |
| 8. | IMREEF FISHC2 | 0.090 | 0.15 | 0.972 | 0.362 |
| 10. | KYPHOSIDAEH | 0.020 | 0.03 | 0.953 | 0.112 |
| 11. | IMREEF FISHH | 0.020 | 0.03 | 0.825 | -0.529 |
| 12. | LGREEF FISHC | 0.232 | 0.39 | 0.996 | 0.899 |
| 19. | IMREEF FISHC4 | 0.010 | 0.02 | 0.945 | 0.026 |
| 23. | LGSCARIDAEH | 0.248 | 0.41 | 0.970 | 0.335 |
| 30. | SEATURTLES | 0.035 | 0.06 | 0.994 | 0.858 |
| 32. | LOBSTERS | 0.160 | 0.27 | 0.992 | 0.789 |
| 33. | CRABS | 0.070 | 0.12 | 0.898 | -0.290 |
| TOTAL | | 1.000 | 1.67 | | |
| Predator: (19) IMREEF FISHC4 | | | | | |
| 33. | CRABS | 0.001 | 0.01 | -0.580 | 0.124 |
| 34. | SHRMP/HCRB/StOM | 0.002 | 0.02 | 0.005 | 0.680 |
| 36. | ASTEROIDS | 0.002 | 0.02 | -0.425 | 0.328 |
| 37. | ECHINOIDS | 0.003 | 0.03 | -0.737 | -0.161 |
| 40. | POLY/PRIA/OPHIU | 0.006 | 0.05 | -0.043 | 0.651 |
| 41. | HOL/SIP/ECH/HEM | 0.006 | 0.05 | -0.372 | 0.384 |
| 43. | ASCI/BARN/BRYOZ | 0.023 | 0.20 | -0.084 | 0.624 |
| 44. | SPONGES | 0.921 | 7.91 | 0.706 | 0.968 |
| 45. | CORALS/ANEMONES | 0.007 | 0.06 | -0.548 | 0.171 |
| 46. | ZOOPLANKTON | 0.001 | 0.01 | -0.727 | -0.141 |
| 49. | Benthic prod. | 0.028 | 0.24 | -0.804 | -0.319 |
| TOTAL | | 1.000 | 8.59 | | |
| Predator: (20) SMREEF FISHO1 | | | | | |
| 7. | LGIMSCHOOLFISHP | 0.003 | 0.17 | 0.151 | -0.616 |
| 8. | IMREEF FISHC2 | 0.003 | 0.17 | 0.400 | -0.418 |
| 9. | HEMIRAMPHIDAEH | 0.003 | 0.17 | 0.862 | 0.423 |
| 14. | SMREEF FISHC1 | 0.004 | 0.23 | 0.641 | -0.104 |
| 15. | SMSCHOOLFISHPEL | 0.004 | 0.23 | 0.331 | -0.482 |
| 16. | ENGRAULIDAEH | 0.004 | 0.23 | 0.717 | 0.040 |
| 17. | SMREEF FISHC2 | 0.004 | 0.23 | 0.879 | 0.483 |
| 20. | SMREEF FISHO1 | 0.006 | 0.35 | 0.791 | 0.214 |
| 21. | SMREEF FISHO2 | 0.004 | 0.23 | 0.907 | 0.587 |
| 22. | SMREEF FISHO3 | 0.004 | 0.23 | 0.909 | 0.598 |
| 25. | SMSCARIDAEH | 0.004 | 0.23 | 0.598 | -0.173 |
| 26. | BLENNIIDAEH | 0.004 | 0.23 | 0.942 | 0.739 |
| 27. | SMGOBIIDAEC | 0.003 | 0.17 | 0.977 | 0.912 |
| 33. | CRABS | 0.010 | 0.58 | 0.453 | -0.362 |
| 34. | SHRMP/HCRB/StOM | 0.010 | 0.58 | 0.669 | -0.054 |
| 35. | SMBENTHARTHROPO | 0.056 | 3.25 | 0.910 | 0.603 |

continued...

Table 8.8.1 continued

| | Diet composition | Food intake | Ilev's electivity index | Standardized forage ratio |
|------------------------------|------------------|-------------|-------------------------|---------------------------|
| 38. GASTROPODS | 0.038 | 2.20 | 0.608 | -0.159 |
| 40. POLY/PRIA/OPHIU | 0.050 | 2.90 | 0.769 | 0.157 |
| 42. BIVALVES | 0.020 | 1.16 | -0.039 | -0.721 |
| 43. ASCI/BARN/BRYOZ | 0.020 | 1.16 | -0.153 | -0.772 |
| 44. SPONGES | 0.020 | 1.16 | -0.776 | -0.957 |
| 45. CORALS/ANEMONES | 0.020 | 1.16 | -0.090 | -0.745 |
| 46. ZOOPLANKTON | 0.100 | 5.80 | 0.881 | 0.490 |
| 49. Benthic prod. | 0.326 | 18.89 | 0.117 | -0.637 |
| 50. Detritus | 0.280 | 16.23 | -0.172 | -0.780 |
| TOTAL | 1.000 | 57.96 | | |
| Predator: (21) SMREEF FISHO2 | | | | |
| 14. SMREEF FISHC1 | 0.003 | 0.05 | 0.548 | -0.159 |
| 15. SMSCHOOLFISHPEL | 0.003 | 0.05 | 0.198 | -0.523 |
| 16. ENGRAULIDAEH | 0.003 | 0.05 | 0.640 | -0.015 |
| 17. SMREEF FISHC2 | 0.003 | 0.05 | 0.842 | 0.438 |
| 21. SMREEF FISHO2 (0' Cycle) | 0.003 | 0.05 | 0.877 | 0.547 |
| 22. SMREEF FISHO3 | 0.003 | 0.05 | 0.881 | 0.559 |
| 26. BLENNIIDAEH | 0.003 | 0.05 | 0.924 | 0.710 |
| 27. SMGOBIIDAEH | 0.002 | 0.03 | 0.966 | 0.882 |
| 33. CRABS | 0.050 | 0.79 | 0.860 | 0.492 |
| 34. SHRMP/HCRB/STOM | 0.040 | 0.63 | 0.906 | 0.644 |
| 35. SMBENTHARTHROPO | 0.035 | 0.55 | 0.860 | 0.494 |
| 36. ASTEROIDS | 0.021 | 0.33 | 0.618 | -0.052 |
| 37. ECHINOIDS | 0.050 | 0.79 | 0.432 | -0.305 |
| 38. GASTROPODS | 0.035 | 0.55 | 0.581 | -0.110 |
| 40. POLY/PRIA/OPHIU | 0.100 | 1.58 | 0.877 | 0.547 |
| 41. HOL/SIP/ECH/HEM | 0.005 | 0.08 | -0.448 | -0.852 |
| 42. BIVALVES | 0.027 | 0.43 | 0.110 | -0.585 |
| 43. ASCI/BARN/BRYOZ | 0.020 | 0.32 | -0.153 | -0.733 |
| 44. SPONGES | 0.080 | 1.27 | -0.329 | -0.809 |
| 45. CORALS/ANEMONES | 0.100 | 1.58 | 0.613 | -0.060 |
| 46. ZOOPLANKTON | 0.109 | 1.73 | 0.890 | 0.590 |
| 47. DECOMP/MICROFAU | 0.005 | 0.08 | 0.254 | -0.477 |
| 49. Benthic prod. | 0.200 | 3.17 | -0.126 | -0.720 |
| 50. Detritus | 0.100 | 1.58 | -0.597 | -0.900 |
| TOTAL | 1.000 | 15.84 | | |
| Predator: (22) SMREEF FISHO3 | | | | |
| 35. SMBENTHARTHROPO | 0.008 | 0.30 | 0.506 | 0.540 |
| 40. POLY/PRIA/OPHIU | 0.050 | 1.91 | 0.769 | 0.807 |
| 42. BIVALVES | 0.071 | 2.71 | 0.533 | 0.567 |
| 46. ZOOPLANKTON | 0.200 | 7.62 | 0.939 | 0.979 |
| 49. Benthic prod. | 0.335 | 12.77 | 0.131 | 0.158 |
| 50. Detritus | 0.336 | 12.81 | -0.082 | -0.060 |
| TOTAL | 1.000 | 38.11 | | |
| Predator: (23) LGSCARIDAEH | | | | |
| 44. SPONGES | 0.002 | 0.51 | -0.975 | -0.727 |
| 45. CORALS/ANEMONES | 0.001 | 0.26 | -0.920 | -0.310 |
| 49. Benthic prod. | 0.997 | 255.49 | 0.589 | 0.999 |
| TOTAL | 1.000 | 256.26 | | |
| Predator: (24) IMSCARIDAEH | | | | |
| 44. SPONGES | 0.001 | 0.11 | -0.987 | -0.852 |
| 49. Benthic prod. | 0.999 | 114.22 | 0.590 | 1.000 |
| TOTAL | 1.000 | 114.33 | | |
| Predator: (25) SMSCARIDAEH | | | | |
| 49. Benthic prod. | 1.000 | 172.04 | 0.590 | 1.000 |
| TOTAL | 1.000 | 172.04 | | |

continued...

Table 8.8.1 continued

| | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|----------------------------|------------------|-------------|--------------------------|---------------------------|
| Predator: (26) BLENNIIDAEH | | | | |
| 38. GASTROPODS | 0.001 | 0.02 | -0.805 | 0.142 |
| 46. ZOOPLANKTON | 0.005 | 0.11 | -0.118 | 0.843 |
| 49. Benthic prod. | 0.497 | 10.77 | 0.317 | 0.956 |
| 50. Detritus | 0.497 | 10.77 | 0.113 | 0.912 |
| TOTAL | 1.000 | 21.66 | | |
| Predator: (27) SMGOBIIDAEC | | | | |
| 35. SMBENTHARTHROPO | 1.000 | 3.05 | 0.995 | 1.000 |
| TOTAL | 1.000 | 3.05 | | |
| Predator: (28) SEABIRDS | | | | |
| 2. SHARKS/SCOMBRID | 0.020 | 0.03 | 0.992 | 0.758 |
| 3. LGJACKSC | 0.010 | 0.01 | 0.993 | 0.790 |
| 4. IMJACKSC | 0.060 | 0.08 | 0.989 | 0.685 |
| 5. SMJACKSC | 0.030 | 0.04 | 0.987 | 0.621 |
| 6. IMREEF FISHC1 | 0.010 | 0.01 | 0.887 | -0.391 |
| 7. LGIMSCHOOOLFISHP | 0.280 | 0.38 | 0.984 | 0.558 |
| 8. IMREEF FISHC2 | 0.040 | 0.05 | 0.938 | -0.098 |
| 9. HEMIRAMPHIDAEH | 0.100 | 0.14 | 0.996 | 0.877 |
| 11. IMREEF FISHH | 0.010 | 0.01 | 0.679 | -0.760 |
| 12. LGREEF FISHC | 0.005 | 0.01 | 0.818 | -0.586 |
| 13. IMREEF FISHC3 | 0.030 | 0.04 | 0.909 | -0.291 |
| 14. SMREEF FISHC1 | 0.025 | 0.03 | 0.932 | -0.140 |
| 15. SMSCHOOOLFISHPEL | 0.200 | 0.27 | 0.980 | 0.464 |
| 16. ENGRAULIDAEH | 0.100 | 0.14 | 0.987 | 0.622 |
| 17. SMREEF FISHC2 | 0.008 | 0.01 | 0.938 | -0.098 |
| 19. IMREEF FISHC4 | 0.005 | 0.01 | 0.893 | -0.367 |
| 20. SMREEF FISHO1 | 0.005 | 0.01 | 0.754 | -0.686 |
| 21. SMREEF FISHO2 | 0.003 | 0.00 | 0.877 | -0.428 |
| 23. LGSCARIDAEH | 0.002 | 0.00 | -0.306 | -0.973 |
| 24. IMSCARIDAEH | 0.005 | 0.01 | 0.642 | -0.787 |
| 25. SMSCARIDAEH | 0.002 | 0.00 | 0.331 | -0.902 |
| 46. ZOOPLANKTON | 0.050 | 0.07 | 0.775 | -0.659 |
| TOTAL | 1.000 | 1.36 | | |
| Predator: (29) SQUIDS | | | | |
| 7. LGIMSCHOOOLFISHP | 0.150 | 3.94 | 0.971 | 0.809 |
| 9. HEMIRAMPHIDAEH | 0.015 | 0.39 | 0.971 | 0.807 |
| 15. SMSCHOOOLFISHPEL | 0.130 | 3.41 | 0.970 | 0.799 |
| 16. ENGRAULIDAEH | 0.030 | 0.79 | 0.957 | 0.714 |
| 29. SQUIDS | 0.010 | 0.26 | 0.942 | 0.622 |
| 34. SHRMP/HCRB/STOM | 0.100 | 2.63 | 0.961 | 0.741 |
| 46. ZOOPLANKTON | 0.565 | 14.83 | 0.978 | 0.860 |
| TOTAL | 1.000 | 26.25 | | |
| Predator: (30) SEATURTLES | | | | |
| 32. LOBSTERS | 0.060 | 0.10 | 0.979 | 0.982 |
| 33. CRABS | 0.025 | 0.04 | 0.738 | 0.440 |
| 34. SHRMP/HCRB/STOM | 0.040 | 0.07 | 0.906 | 0.795 |
| 36. ASTEROIDS | 0.020 | 0.04 | 0.603 | 0.210 |
| 37. ECHINOIDS | 0.020 | 0.04 | 0.005 | -0.456 |
| 38. GASTROPODS | 0.025 | 0.04 | 0.459 | 0.006 |
| 42. BIVALVES | 0.020 | 0.04 | -0.039 | -0.490 |
| 43. ASCI/BARN/BRYOZ | 0.020 | 0.04 | -0.153 | -0.573 |
| 44. SPONGES | 0.340 | 0.60 | 0.364 | -0.110 |
| 45. CORALS/ANEMONES | 0.016 | 0.03 | -0.199 | -0.604 |
| 49. Benthic prod. | 0.414 | 0.72 | 0.233 | -0.251 |
| TOTAL | 1.000 | 1.75 | | |
| Predator: (31) OCTOPUSES | | | | |
| 8. IMREEF FISHC2 | 0.002 | 0.11 | 0.217 | -0.271 |
| 11. IMREEF FISHH | 0.002 | 0.11 | 0.023 | -0.445 |
| 12. LGREEF FISHC | 0.001 | 0.06 | 0.333 | -0.149 |
| 13. IMREEF FISHC3 | 0.003 | 0.17 | 0.353 | -0.128 |

continued...

Table 8.8.1 continued

| | Diet composition | Food intake | Ivlev's elec- tivity index | Standardized forage ratio |
|--------------------------------|------------------|-------------|-------------------------------|------------------------------|
| 14. SMREEF FISHC1 | 0.007 | 0.40 | 0.778 | 0.512 |
| 17. SMREEF FISHC2 | 0.001 | 0.06 | 0.590 | 0.186 |
| 20. SMREEF FISHO1 | 0.002 | 0.11 | 0.481 | 0.031 |
| 21. SMREEF FISHO2 | 0.002 | 0.11 | 0.821 | 0.602 |
| 31. OCTOPUSES (0' Cycle) | 0.020 | 1.14 | 0.846 | 0.656 |
| 33. CRABS | 0.025 | 1.42 | 0.738 | 0.436 |
| 34. SHRMP/HCRB/STOM | 0.020 | 1.14 | 0.820 | 0.598 |
| 38. GASTROPODS | 0.500 | 28.39 | 0.964 | 0.941 |
| 39. CHITONS/SCAPHOP | 0.010 | 0.57 | -0.102 | -0.540 |
| 40. POLY/PRIA/OPHIU | 0.005 | 0.28 | -0.133 | -0.562 |
| 42. BIVALVES | 0.400 | 22.71 | 0.897 | 0.773 |
| TOTAL | 1.000 | 56.78 | | |
| Predator: (32) LOBSTERS | | | | |
| 38. GASTROPODS | 0.150 | 3.63 | 0.884 | 0.923 |
| 40. POLY/PRIA/OPHIU | 0.050 | 1.21 | 0.769 | 0.806 |
| 42. BIVALVES | 0.500 | 12.10 | 0.917 | 0.957 |
| 49. Benthic prod. | 0.180 | 4.36 | -0.177 | -0.159 |
| 50. Detritus | 0.120 | 2.90 | -0.535 | -0.525 |
| TOTAL | 1.000 | 24.20 | | |
| Predator: (33) CRABS | | | | |
| 14. SMREEF FISHC1 | 0.010 | 2.66 | 0.839 | 0.784 |
| 17. SMREEF FISHC2 | 0.007 | 1.86 | 0.929 | 0.925 |
| 33. CRABS (0' Cycle) | 0.005 | 1.33 | 0.141 | -0.089 |
| 34. SHRMP/HCRB/STOM | 0.010 | 2.66 | 0.669 | 0.539 |
| 35. SMBENTHARTHROPO | 0.005 | 1.33 | 0.312 | 0.095 |
| 37. ECHINOIDS | 0.050 | 13.30 | 0.432 | 0.235 |
| 38. GASTROPODS | 0.050 | 13.30 | 0.687 | 0.564 |
| 40. POLY/PRIA/OPHIU | 0.004 | 1.06 | -0.241 | -0.447 |
| 41. HOL/SIP/ECH/HEM | 0.001 | 0.27 | -0.858 | -0.910 |
| 42. BIVALVES | 0.050 | 13.30 | 0.396 | 0.192 |
| 43. ASCI/BARN/BRYOZ | 0.020 | 5.32 | -0.153 | -0.370 |
| 44. SPONGES | 0.050 | 13.30 | -0.520 | -0.672 |
| 45. CORALS/ANEMONES | 0.010 | 2.66 | -0.411 | -0.587 |
| 46. ZOOPLANKTON | 0.030 | 7.98 | 0.651 | 0.514 |
| 47. DECOMP/MICROFAU | 0.030 | 7.98 | 0.820 | 0.755 |
| 48. Phytoplankton | 0.020 | 5.32 | 0.432 | 0.235 |
| 49. Benthic prod. | 0.600 | 159.60 | 0.399 | 0.196 |
| 50. Detritus | 0.048 | 12.77 | -0.784 | -0.860 |
| TOTAL | 1.000 | 266.00 | | |
| Predator: (34) SHRMP/HCRB/STOM | | | | |
| 8. IMREEF FISHC2 | 0.010 | 2.69 | 0.772 | 0.358 |
| 14. SMREEF FISHC1 | 0.010 | 2.69 | 0.839 | 0.521 |
| 16. ENGRAULIDAEH | 0.010 | 2.69 | 0.876 | 0.623 |
| 17. SMREEF FISHC2 | 0.005 | 1.35 | 0.902 | 0.699 |
| 29. SQUIDS | 0.005 | 1.35 | 0.888 | 0.657 |
| 31. OCTOPUSES | 0.025 | 6.72 | 0.875 | 0.620 |
| 32. LOBSTERS | 0.005 | 1.35 | 0.771 | 0.355 |
| 33. CRABS | 0.002 | 0.54 | -0.306 | -0.756 |
| 34. SHRMP/HCRB/STOM (0' Cycle) | 0.002 | 0.54 | 0.005 | -0.581 |
| 35. SMBENTHARTHROPO | 0.040 | 10.76 | 0.877 | 0.625 |
| 36. ASTEROIDS | 0.010 | 2.69 | 0.338 | -0.305 |
| 37. ECHINOIDS | 0.110 | 29.59 | 0.695 | 0.198 |
| 38. GASTROPODS | 0.080 | 21.52 | 0.792 | 0.405 |
| 39. CHITONS/SCAPHOP | 0.010 | 2.69 | -0.102 | -0.648 |
| 40. POLY/PRIA/OPHIU | 0.050 | 13.45 | 0.769 | 0.351 |
| 41. HOL/SIP/ECH/HEM | 0.005 | 1.35 | -0.448 | -0.819 |
| 42. BIVALVES | 0.212 | 57.03 | 0.815 | 0.459 |
| 46. ZOOPLANKTON | 0.150 | 40.35 | 0.919 | 0.752 |
| 47. DECOMP/MICROFAU | 0.055 | 14.80 | 0.897 | 0.685 |
| 49. Benthic prod. | 0.084 | 22.60 | -0.508 | -0.843 |
| 50. Detritus | 0.120 | 32.28 | -0.535 | -0.853 |
| TOTAL | 1.000 | 269.00 | | |

continued...

Table 8.8.1 continued

| | | Diet composition | Food intake | Ivlev's elec- tivity index | Standardized forage ratio |
|-----------|-----------------------|---------------------|----------------|-------------------------------|------------------------------|
| Predator: | (35) SMBENTHARTHROPO | | | | |
| 43. | ASCI/BARN/BRYOZ | 0.050 | 83.14 | 0.295 | 0.571 |
| 44. | SPONGES | 0.060 | 99.77 | -0.451 | -0.168 |
| 45. | CORALS/ANEMONES | 0.040 | 66.51 | 0.251 | 0.535 |
| 46. | ZOOPLANKTON | 0.040 | 66.51 | 0.726 | 0.535 |
| 47. | DECOMP/MICROFAU | 0.040 | 66.51 | 0.862 | 0.961 |
| 49. | Benthic prod. | 0.600 | 997.73 | 0.399 | 0.652 |
| 50. | Detritus | 0.170 | 282.69 | -0.400 | -0.106 |
| | TOTAL | 1.000 | 1,662.88 | | |
| Predator: | (36) ASTEROIDS | | | | |
| 33. | CRABS | 0.010 | 0.81 | 0.453 | 0.401 |
| 36. | ASTEROIDS | 0.080 | 6.48 | 0.883 | 0.899 |
| 37. | ECHINOIDS (0' Cycle) | 0.070 | 5.67 | 0.559 | 0.519 |
| 38. | GASTROPODS | 0.135 | 10.94 | 0.871 | 0.884 |
| 39. | CHITONS/SCAPHOP | 0.005 | 0.41 | -0.421 | -0.488 |
| 40. | POLY/PRIA/OPHIU | 0.030 | 2.43 | 0.642 | 0.614 |
| 41. | HOL/SIP/ECH/HEM | 0.005 | 0.41 | -0.448 | -0.513 |
| 42. | BIVALVES | 0.145 | 11.74 | 0.740 | 0.728 |
| 43. | ASCI/BARN/BRYOZ | 0.020 | 1.62 | -0.153 | -0.231 |
| 44. | SPONGES | 0.020 | 1.62 | -0.776 | -0.808 |
| 45. | CORALS/ANEMONES | 0.012 | 0.97 | -0.333 | -0.404 |
| 47. | DECOMP/MICROFAU | 0.020 | 1.62 | 0.741 | 0.729 |
| 49. | Benthic prod. | 0.088 | 7.13 | -0.491 | -0.552 |
| 50. | Detritus | 0.360 | 29.16 | -0.048 | -0.126 |
| | TOTAL | 1.000 | 81.00 | | |
| Predator: | (37) ECHINOIDS | | | | |
| 36. | ASTEROIDS | 0.006 | 2.22 | 0.096 | 0.764 |
| 37. | ECHINOIDS (0' Cycle) | 0.015 | 5.55 | -0.138 | 0.631 |
| 40. | POLY/PRIA/OPHIU | 0.004 | 1.48 | -0.241 | 0.556 |
| 41. | HOL/SIP/ECH/HEM | 0.005 | 1.85 | -0.448 | 0.360 |
| 42. | BIVALVES | 0.010 | 3.70 | -0.368 | 0.445 |
| 44. | SPONGES | 0.050 | 18.50 | -0.520 | 0.272 |
| 45. | CORALS/ANEMONES | 0.050 | 18.50 | 0.352 | 0.870 |
| 49. | Benthic prod. | 0.860 | 318.20 | 0.539 | 0.931 |
| | TOTAL | 1.000 | 370.00 | | |
| Predator: | (38) GASTROPODS | | | | |
| 32. | LOBSTERS | 0.002 | 1.31 | 0.511 | 0.627 |
| 33. | CRABS | 0.002 | 1.31 | -0.306 | -0.177 |
| 34. | SHRMP/HCRB/STOM | 0.004 | 2.62 | 0.338 | 0.469 |
| 35. | SMBENTHARTHROPO | 0.006 | 3.93 | 0.391 | 0.519 |
| 38. | GASTROPODS (0' Cycle) | 0.020 | 13.10 | 0.367 | 0.496 |
| 39. | CHITONS/SCAPHOP | 0.020 | 13.10 | 0.239 | 0.377 |
| 40. | POLY/PRIA/OPHIU | 0.050 | 32.76 | 0.769 | 0.851 |
| 41. | HOL/SIP/ECH/HEM | 0.015 | 9.83 | 0.067 | 0.209 |
| 42. | BIVALVES | 0.040 | 26.21 | 0.298 | 0.432 |
| 43. | ASCI/BARN/BRYOZ | 0.058 | 38.00 | 0.361 | 0.491 |
| 44. | SPONGES | 0.040 | 26.21 | -0.597 | -0.504 |
| 45. | CORALS/ANEMONES | 0.015 | 9.83 | -0.230 | -0.095 |
| 47. | DECOMP/MICROFAU | 0.030 | 19.66 | 0.820 | 0.894 |
| 49. | Benthic prod. | 0.400 | 262.08 | 0.217 | 0.355 |
| 50. | Detritus | 0.298 | 195.25 | -0.141 | -0.002 |
| | TOTAL | 1.000 | 655.20 | | |
| Predator: | (39) CHITONS/SCAPHOP | | | | |
| 43. | ASCI/BARN/BRYOZ | 0.040 | 29.02 | 0.190 | -0.181 |
| 44. | SPONGES | 0.020 | 14.51 | -0.776 | -0.889 |
| 45. | CORALS/ANEMONES | 0.008 | 5.80 | -0.500 | -0.730 |
| 47. | DECOMP/MICROFAU | 0.300 | 217.62 | 0.980 | 0.998 |
| 49. | Benthic prod. | 0.632 | 458.45 | 0.421 | 0.078 |
| | TOTAL | 1.000 | 725.40 | | |

continued...

Table 8.8.1 continued

| | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|---------------------|----------------------|-------------|--------------------------|---------------------------|
| Predator: | (40) POLY/PRIA/OPHIU | | | |
| 35. SMBENTHARTHROPO | 0.015 | 30.49 | 0.702 | 0.705 |
| 38. GASTROPODS | 0.010 | 20.33 | 0.038 | -0.002 |
| 40. POLY/PRIA/OPHIU | 0.030 | 60.98 | 0.642 | 0.639 |
| 42. BIVALVES | 0.030 | 60.98 | 0.162 | 0.125 |
| 43. ASCI/BARN/BRYOZ | 0.040 | 81.31 | 0.190 | 0.155 |
| 44. SPONGES | 0.050 | 101.64 | -0.520 | -0.554 |
| 45. CORALS/ANEMONES | 0.010 | 20.33 | -0.411 | -0.449 |
| 46. ZOOPLANKTON | 0.050 | 101.64 | 0.775 | 0.786 |
| 47. DECOMP/MICROFAU | 0.050 | 101.64 | 0.888 | 0.913 |
| 48. Phytoplankton | 0.100 | 203.28 | 0.853 | 0.874 |
| 49. Benthic prod. | 0.245 | 498.04 | -0.025 | -0.066 |
| 50. Detritus | 0.370 | 752.14 | -0.034 | -0.076 |
| TOTAL | 1.000 | 2,032.80 | | |
| Predator: | (41) HOL/SIP/ECH/HEM | | | |
| 47. DECOMP/MICROFAU | 0.100 | 22.26 | 0.942 | 0.996 |
| 48. Phytoplankton | 0.010 | 2.23 | 0.116 | 0.263 |
| 49. Benthic prod. | 0.130 | 28.93 | -0.329 | -0.197 |
| 50. Detritus | 0.760 | 169.15 | 0.315 | 0.453 |
| TOTAL | 1.000 | 222.57 | | |
| Predator: | (42) BIVALVES | | | |
| 47. DECOMP/MICROFAU | 0.100 | 103.79 | 0.942 | 0.975 |
| 48. Phytoplankton | 0.150 | 155.68 | 0.900 | 0.926 |
| 50. Detritus | 0.750 | 778.41 | 0.309 | 0.276 |
| TOTAL | 1.000 | 1,037.88 | | |
| Predator: | (43) ASCI/BARN/BRYOZ | | | |
| 46. ZOOPLANKTON | 0.150 | 412.20 | 0.919 | 0.901 |
| 47. DECOMP/MICROFAU | 0.100 | 274.80 | 0.942 | 0.941 |
| 48. Phytoplankton | 0.200 | 549.60 | 0.924 | 0.909 |
| 50. Detritus | 0.550 | 1,511.40 | 0.163 | -0.097 |
| TOTAL | 1.000 | 2,748.00 | | |
| Predator: | (44) SPONGES | | | |
| 47. DECOMP/MICROFAU | 0.001 | 4.00 | -0.496 | 0.735 |
| 50. Detritus | 0.999 | 3,996.00 | 0.432 | 0.995 |
| TOTAL | 1.000 | 4,000.00 | | |
| Predator: | (45) CORALS/ANEMONES | | | |
| 46. ZOOPLANKTON | 0.150 | 163.35 | 0.919 | 0.939 |
| 47. DECOMP/MICROFAU | 0.100 | 108.90 | 0.942 | 0.968 |
| 49. Benthic prod. | 0.650 | 707.85 | 0.432 | 0.365 |
| 50. Detritus | 0.100 | 108.90 | -0.597 | -0.657 |
| TOTAL | 1.000 | 1,089.00 | | |
| Predator: | (46) ZOOPLANKTON | | | |
| 47. DECOMP/MICROFAU | 0.600 | 3,168.00 | 0.990 | 0.995 |
| 48. Phytoplankton | 0.200 | 1,056.00 | 0.924 | 0.719 |
| 50. Detritus | 0.200 | 1,056.00 | -0.329 | -0.804 |
| TOTAL | 1.000 | 5,280.00 | | |
| Predator: | (47) DECOMP/MICROFAU | | | |
| 49. Benthic prod. | 0.050 | 1,425.00 | -0.675 | 0.597 |
| 50. Detritus | 0.950 | 27,075.00 | 0.411 | 0.997 |
| TOTAL | 1.000 | 28,500.00 | | |
| Predator: | (50) Detritus | | | |
| 1. LGSHARKS/RAYSC | 0.000 | 0.34 | -0.602 | 0.002 |
| 2. SHARKS/SCOMBRID | 0.000 | 0.81 | -0.403 | 0.003 |
| 3. LGJACKSC | 0.000 | 0.24 | -0.546 | 0.002 |

continued...

Table 8.8.1 continued

| | Diet composition | Food intake | Ivlev's electricity index | Standardized forage ratio |
|---------------------|------------------|-------------|---------------------------|---------------------------|
| 4. IMJACKSC | 0.000 | 4.11 | -0.292 | 0.004 |
| 5. SMJACKSC | 0.000 | 2.73 | 0.255 | 0.004 |
| 6. IMREEF FISHC1 | 0.000 | 4.54 | -0.508 | 0.002 |
| 7. LGIMSCHOOLFISHP | 0.001 | 28.06 | -0.293 | 0.004 |
| 8. IMREEF FISHC2 | 0.001 | 13.93 | -0.363 | 0.003 |
| 9. HEMIRAMPHIDAEH | 0.000 | 8.95 | 0.268 | 0.012 |
| 10. KYPHOSIDAEH | 0.001 | 12.02 | 0.039 | 0.008 |
| 11. IMREEF FISHH | 0.002 | 51.76 | 0.077 | 0.008 |
| 12. LGREEF FISHC | 0.000 | 2.27 | -0.673 | 0.001 |
| 13. IMREEF FISHC3 | 0.000 | 10.09 | -0.535 | 0.002 |
| 14. SMREEF FISHC1 | 0.000 | 8.99 | -0.386 | 0.003 |
| 15. SMSCHOOLFISHPEL | 0.002 | 45.68 | -0.010 | 0.007 |
| 16. ENGRAULIDAEH | 0.001 | 30.83 | 0.337 | 0.014 |
| 17. SMREEF FISHC2 | 0.000 | 4.38 | -0.153 | 0.005 |
| 18. LGGROUPERS | 0.000 | 0.47 | -0.755 | 0.001 |
| 19. IMREEF FISHC4 | 0.000 | 1.92 | -0.547 | 0.002 |
| 20. SMREEF FISHO1 | 0.001 | 13.22 | -0.102 | 0.006 |
| 21. SMREEF FISHO2 | 0.000 | 3.27 | -0.164 | 0.005 |
| 22. SMREEF FISHO3 | 0.000 | 7.95 | 0.286 | 0.013 |
| 23. LGSCARIDAEH | 0.003 | 61.97 | -0.169 | 0.005 |
| 24. IMSCARIDAEH | 0.001 | 25.37 | 0.001 | 0.007 |
| 25. SMSCARIDAEH | 0.002 | 35.67 | 0.209 | 0.011 |
| 26. BLENNIIDAEH | 0.000 | 4.58 | 0.249 | 0.012 |
| 27. SMGOBIIDAE | 0.000 | 0.64 | -0.110 | 0.006 |
| 28. SEABIRDS | 0.000 | 0.33 | 0.621 | 0.031 |
| 29. SQUIDS | 0.000 | 5.46 | -0.116 | 0.006 |
| 30. SEATURTLES | 0.000 | 0.36 | -0.730 | 0.001 |
| 31. OCTOPUSES | 0.001 | 13.76 | -0.474 | 0.003 |
| 32. LOBSTERS | 0.000 | 4.97 | -0.503 | 0.002 |
| 33. CRABS | 0.002 | 55.07 | -0.226 | 0.005 |
| 34. SHRMP/HCRB/STOM | 0.002 | 55.24 | 0.092 | 0.009 |
| 35. SMBENTHARTHROPO | 0.015 | 337.41 | 0.694 | 0.040 |
| 36. ASTEROIDS | 0.001 | 16.65 | -0.747 | 0.001 |
| 37. ECHINIDS | 0.005 | 112.68 | -0.606 | 0.002 |
| 38. GASTROPODS | 0.006 | 134.39 | -0.231 | 0.004 |
| 39. CHITONS/SCAPHOP | 0.006 | 147.06 | -0.319 | 0.004 |
| 40. POLY/PRIA/OPHIU | 0.018 | 420.53 | 0.470 | 0.020 |
| 41. HOL/SIP/ECH/HEM | 0.002 | 46.25 | -0.736 | 0.001 |
| 42. BIVALVES | 0.010 | 228.85 | -0.374 | 0.003 |
| 43. ASCI/BARN/BRYOZ | 0.026 | 613.32 | -0.014 | 0.007 |
| 44. SPONGES | 0.073 | 1,700.82 | -0.367 | 0.003 |
| 45. CORALS/ANEMONES | 0.010 | 220.89 | -0.431 | 0.003 |
| 46. ZOOPLANKTON | 0.050 | 1,157.00 | 0.775 | 0.057 |
| 47. DECOMP/MICROFAU | 0.250 | 5,787.20 | 0.976 | 0.603 |
| 48. Phytoplankton | 0.031 | 726.88 | 0.596 | 0.028 |
| 49. Benthic prod. | 0.475 | 11,023.57 | 0.297 | 0.013 |
| TOTAL | 1.000 | 23,193.50 | | |

Table 8.8.2. The 20-box model as obtained by method 1 (cluster analysis + intuition).

| | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|-------------------------------|------------------|-------------|--------------------------|---------------------------|
| Predator: (1) SEA BIRDS | | | | |
| 2. SHARK/RAY/SCOMB | 0.020 | 0.03 | 0.986 | 0.698 |
| 3. L-IMJACKSCARN | 0.070 | 0.10 | 0.990 | 0.795 |
| 4. L-IMSCHOOLPEL | 0.410 | 0.56 | 0.986 | 0.695 |
| 5. SMALLSCHOOLPEL | 0.300 | 0.41 | 0.982 | 0.617 |
| 7. L-IMBENTHCARN | 0.090 | 0.12 | 0.865 | -0.395 |
| 8. SMALLBENTHCARN | 0.041 | 0.06 | 0.632 | -0.759 |
| 10. L-IMREEFFHERB | 0.017 | 0.02 | 0.402 | -0.865 |
| 11. SMALLREEFFHERB | 0.002 | 0.00 | 0.207 | -0.911 |
| 16. ZOOPLANKTON | 0.050 | 0.07 | 0.775 | -0.606 |
| TOTAL | 1.000 | 1.36 | | |
| Predator: (2) SHARK/RAY/SCOMB | | | | |
| 1. SEA BIRDS | 0.006 | 0.03 | 0.999 | 0.966 |
| 2. SHARK/RAY/SCOMB (0' Cycle) | 0.013 | 0.07 | 0.978 | -0.132 |
| 3. L-IMJACKSCARN | 0.039 | 0.21 | 0.982 | -0.045 |
| 4. L-IMSCHOOLPEL | 0.258 | 1.36 | 0.978 | -0.154 |
| 5. SMALLSCHOOLPEL | 0.148 | 0.78 | 0.965 | -0.374 |
| 6. LGGROUPERSCARN | 0.026 | 0.14 | 0.989 | 0.222 |
| 7. L-IMBENTHCARN | 0.210 | 1.10 | 0.940 | -0.585 |
| 8. SMALLBENTHCARN | 0.105 | 0.55 | 0.838 | -0.832 |
| 9. BENTHINVOMNI | 0.027 | 0.14 | 0.261 | -0.973 |
| 10. L-IMREEFFHERB | 0.090 | 0.47 | 0.851 | -0.818 |
| 11. SMALLREEFFHERB | 0.003 | 0.02 | 0.391 | -0.964 |
| 12. BENTHINVHERB | 0.031 | 0.16 | -0.108 | -0.987 |
| 13. SEMISESSDETRIV | 0.038 | 0.20 | 0.044 | -0.983 |
| 20. Detritus | 0.006 | 0.03 | -0.970 | -1.000 |
| TOTAL | 1.000 | 5.26 | | |
| Predator: (3) L-IMJACKSCARN | | | | |
| 3. L-IMJACKSCARN (0' Cycle) | 0.010 | 0.16 | 0.931 | 0.241 |
| 4. L-IMSCHOOLPEL | 0.056 | 0.89 | 0.901 | 0.043 |
| 5. SMALLSCHOOLPEL | 0.203 | 3.24 | 0.974 | 0.679 |
| 7. L-IMBENTHCARN | 0.140 | 2.23 | 0.911 | 0.105 |
| 8. SMALLBENTHCARN | 0.130 | 2.07 | 0.867 | -0.118 |
| 9. BENTHINVOMNI | 0.014 | 0.22 | -0.061 | -0.909 |
| 10. L-IMREEFFHERB | 0.200 | 3.19 | 0.930 | 0.236 |
| 11. SMALLREEFFHERB | 0.212 | 3.38 | 0.988 | 0.883 |
| 12. BENTHINVHERB | 0.010 | 0.16 | -0.587 | -0.972 |
| 13. SEMISESSDETRIV | 0.004 | 0.06 | -0.794 | -0.988 |
| 16. ZOOPLANKTON | 0.021 | 0.33 | 0.536 | -0.694 |
| TOTAL | 1.000 | 15.95 | | |
| Predator: (4) L-IMSCHOOLPEL | | | | |
| 4. L-IMSCHOOLPEL (0' Cycle) | 0.023 | 5.09 | 0.774 | 0.055 |
| 5. SMALLSCHOOLPEL | -0.093 | 20.58 | 0.944 | 0.723 |
| 8. SMALLBENTHCARN | 0.014 | 3.10 | 0.204 | -0.659 |
| 9. BENTHINVOMNI | 0.075 | 16.60 | 0.652 | -0.205 |
| 10. L-IMREEFFHERB | 0.007 | 1.55 | -0.017 | -0.769 |
| 16. ZOOPLANKTON | 0.574 | 127.01 | 0.978 | 0.943 |
| 19. Benthic prod. | 0.214 | 47.35 | -0.092 | -0.798 |
| TOTAL | 1.000 | 221.27 | | |
| Predator: (5) SMALLSCHOOLPEL | | | | |
| 5. SMALLSCHOOLPEL (0' Cycle) | 0.002 | 0.67 | -0.143 | -0.813 |
| 12. BENTHINVHERB | 0.006 | 2.01 | -0.730 | -0.958 |
| 14. SESSILEDTRIV | 0.017 | 5.70 | -0.832 | -0.975 |
| 16. ZOOPLANKTON | 0.628 | 210.73 | 0.980 | 0.958 |
| 17. DECOMP/MICROF | 0.003 | 1.01 | 0.005 | -0.756 |
| 18. Phytoplankton | 0.301 | 101.00 | 0.949 | 0.754 |
| 20. Detritus | -0.043 | 14.43 | -0.804 | -0.971 |
| TOTAL | 1.000 | 335.56 | | |
| Predator: (6) LGGROUPERSCARN | | | | |
| 2. SHARK/RAY/SCOMB | 0.010 | 0.02 | 0.972 | 0.779 |
| 3. L-IMJACKSCARN | 0.010 | 0.02 | 0.931 | 0.433 |

continued...

Table 8.8.2 continued

| | | Diet composition | Food intake | lvlev's elec- tivity index | Standardized forage ratio |
|------------------------------|---------------------------|------------------|-------------|-------------------------------|------------------------------|
| 4. | L-IMSCHOOLPEL | 0.010 | 0.02 | 0.547 | -0.566 |
| 7. | L-IMBENTHCARN | 0.612 | 1.02 | 0.979 | 0.851 |
| 10. | L-IMREEFFHERB | 0.288 | 0.48 | 0.951 | 0.585 |
| 12. | BENTHINVHERB | 0.070 | 0.12 | 0.291 | -0.744 |
| TOTAL | | 1.000 | 1.67 | | |
| Predator: (7) L-IMBENTHCARN | | | | | |
| 3. | L-IMJACKSCARN | 0.001 | 0.24 | 0.472 | -0.156 |
| 4. | L-IMSCHOOLPEL | 0.004 | 0.94 | 0.154 | -0.481 |
| 5. | SMALLSCHOOLPEL | 0.028 | 6.58 | 0.826 | 0.509 |
| 7. | L-IMBENTHCARN (0' Cycle) | 0.058 | 13.64 | 0.798 | 0.436 |
| 8. | SMALLBENTHCARN | 0.050 | 11.76 | 0.688 | 0.190 |
| 9. | BENTHINVOMNI | 0.234 | 55.03 | 0.873 | 0.646 |
| 10. | L-IMREEFFHERB | 0.038 | 8.94 | 0.680 | 0.174 |
| 11. | SMALLREEFFHERB | 0.009 | 2.12 | 0.745 | 0.310 |
| 12. | BENTHINVHERB | 0.150 | 35.27 | 0.592 | 0.018 |
| 13. | SEMISESDETRIV | 0.198 | 46.56 | 0.701 | 0.217 |
| 14. | SESSILEDETRIV | 0.116 | 27.28 | -0.231 | -0.726 |
| 15. | CORALS | 0.002 | 0.47 | -0.845 | -0.958 |
| 16. | ZOOPLANKTON | 0.057 | 13.40 | 0.800 | 0.441 |
| 19. | Benthic prod. | 0.043 | 10.11 | -0.714 | -0.919 |
| 20. | Detritus | 0.012 | 2.82 | -0.941 | -0.985 |
| TOTAL | | 1.000 | 235.15 | | |
| Predator: (8) SMALLBENTHCARN | | | | | |
| 4. | L-IMSCHOOLPEL | 0.004 | 2.00 | 0.154 | -0.445 |
| 5. | SMALLSCHOOLPEL | 0.007 | 3.51 | 0.448 | -0.141 |
| 7. | L-IMBENTHCARN | 0.023 | 11.52 | 0.558 | 0.013 |
| 8. | SMALLBENTHCARN (0' Cycle) | 0.040 | 20.04 | 0.624 | 0.120 |
| 9. | BENTHINVOMNI | 0.135 | 67.62 | 0.790 | 0.458 |
| 11. | SMALLREEFFHERB | 0.003 | 1.50 | 0.391 | -0.211 |
| 12. | BENTHINVHERB | 0.150 | 75.14 | 0.592 | 0.066 |
| 13. | SEMISESDETRIV | 0.151 | 75.64 | 0.626 | 0.123 |
| 14. | SESSILEDETRIV | 0.014 | 7.01 | -0.860 | -0.959 |
| 15. | CORALS | 0.002 | 1.00 | -0.845 | -0.955 |
| 16. | ZOOPLANKTON | 0.153 | 76.64 | 0.920 | 0.822 |
| 17. | DECOMP/MICROF | 0.038 | 19.03 | 0.855 | 0.625 |
| 19. | Benthic prod. | 0.119 | 59.61 | -0.368 | -0.772 |
| 20. | Detritus | 0.161 | 80.65 | -0.422 | -0.797 |
| TOTAL | | 1.000 | 500.91 | | |
| Predator: (9) BENTHINVOMNI | | | | | |
| 8. | SMALLBENTHCARN | 0.004 | 10.75 | -0.396 | -0.644 |
| 9. | BENTHINVOMNI (0' Cycle) | 0.047 | 126.34 | 0.497 | 0.229 |
| 12. | BENTHINVHERB | 0.018 | 48.38 | -0.363 | -0.620 |
| 13. | SEMISESDETRIV | 0.036 | 96.77 | 0.017 | -0.310 |
| 14. | SESSILEDETRIV | 0.092 | 247.29 | -0.337 | -0.602 |
| 15. | CORALS | 0.006 | 16.13 | -0.597 | -0.777 |
| 16. | ZOOPLANKTON | 0.038 | 102.14 | 0.714 | 0.557 |
| 17. | DECOMP/MICROF | 0.045 | 120.96 | 0.876 | 0.851 |
| 18. | Phytoplankton | 0.076 | 204.29 | 0.811 | 0.727 |
| 19. | Benthic prod. | 0.286 | 768.73 | 0.052 | -0.277 |
| 20. | Detritus | 0.352 | 946.17 | -0.059 | -0.379 |
| TOTAL | | 1.000 | 2,687.98 | | |
| Predator: (10) L-IMREEFFHERB | | | | | |
| 8. | SMALLBENTHCARN | 0.001 | 0.68 | -0.805 | -0.306 |
| 9. | BENTHINVOMNI | 0.001 | 0.68 | -0.881 | -0.530 |
| 14. | SESSILEDETRIV | 0.001 | 0.68 | -0.989 | -0.950 |
| 16. | ZOOPLANKTON | 0.001 | 0.68 | -0.727 | -0.120 |
| 19. | Benthic prod. | 0.828 | 559.10 | 0.526 | 0.975 |
| 20. | Detritus | 0.168 | 113.44 | -0.404 | 0.388 |
| TOTAL | | 1.000 | 675.25 | | |

continued...

Table 8.8.2 continued

| | | Diet composition | Food intake | Ilev's elec- tivity index | Standardized forage ratio |
|--------------------|---------------------|---------------------|----------------|------------------------------|------------------------------|
| Predator: | (11) SMALLREEFFHERB | | | | |
| 9. BENTHINVOMNI | | 0.008 | 1.85 | -0.328 | 0.027 |
| 12. BENTHINVHERB | | 0.001 | 0.23 | -0.949 | -0.902 |
| 13. SEMISESSDETRIV | | 0.012 | 2.78 | -0.487 | -0.171 |
| 16. ZOOPLANKTON | | 0.033 | 7.65 | 0.678 | 0.915 |
| 19. Benthic prod. | | 0.844 | 195.65 | 0.532 | 0.815 |
| 20. Detritus | | 0.102 | 23.64 | -0.591 | -0.314 |
| TOTAL | | 1.000 | 231.81 | | |
| Predator: | (12) BENTHINVHERB | | | | |
| 8. SMALLBENTHCARN | | 0.003 | 9.07 | -0.510 | -0.743 |
| 9. BENTHINVOMNI | | 0.005 | 15.12 | -0.519 | -0.749 |
| 12. BENTHINVHERB | (0' Cycle) | 0.007 | 21.17 | -0.692 | -0.848 |
| 13. SEMISESSDETRIV | | 0.006 | 18.15 | -0.706 | -0.855 |
| 14. SESSILEDETRIV | | 0.087 | 263.11 | -0.362 | -0.648 |
| 15. CORALS | | 0.025 | 75.61 | 0.025 | -0.345 |
| 16. ZOOPLANKTON | | 0.025 | 75.61 | 0.595 | 0.325 |
| 17. DECOMP/MICROF | | 0.097 | 293.36 | 0.941 | 0.970 |
| 18. Phytoplankton | | 0.002 | 6.05 | -0.597 | -0.794 |
| 19. Benthic prod. | | 0.645 | 1,950.66 | 0.429 | 0.092 |
| 20. Detritus | | 0.098 | 296.38 | -0.603 | -0.798 |
| TOTAL | | 1.000 | 3,024.28 | | |
| Predator: | (13) SEMISESSDETRIV | | | | |
| 17. DECOMP/MICROF | | 0.100 | 126.04 | 0.942 | 0.946 |
| 18. Phytoplankton | | 0.125 | 157.55 | 0.881 | 0.787 |
| 19. Benthic prod. | | 0.023 | 28.99 | -0.836 | -0.936 |
| 20. Detritus | | 0.752 | 947.80 | 0.310 | -0.157 |
| TOTAL | | 1.000 | 1,260.37 | | |
| Predator: | (14) SESSILEDETRIV | | | | |
| 16. ZOOPLANKTON | | 0.061 | 411.65 | 0.812 | 0.750 |
| 17. DECOMP/MICROF | | 0.041 | 276.68 | 0.865 | 0.846 |
| 18. Phytoplankton | | 0.081 | 546.62 | 0.822 | 0.768 |
| 20. Detritus | | 0.817 | 5,513.40 | 0.347 | 0.076 |
| TOTAL | | 1.000 | 6,748.34 | | |
| Predator: | (15) CORALS | | | | |
| 16. ZOOPLANKTON | | 0.150 | 162.00 | 0.919 | 0.850 |
| 17. DECOMP/MICROF | | 0.100 | 108.00 | 0.942 | 0.921 |
| 19. Benthic prod. | | 0.650 | 702.00 | 0.432 | -0.091 |
| 20. Detritus | | 0.100 | 108.00 | -0.597 | -0.852 |
| TOTAL | | 1.000 | 1,080.00 | | |
| Predator: | (16) ZOOPLANKTON | | | | |
| 17. DECOMP/MICROF | | 0.600 | 3,168.00 | 0.990 | 0.987 |
| 18. Phytoplankton | | 0.200 | 1,056.00 | 0.924 | 0.406 |
| 20. Detritus | | 0.200 | 1,056.00 | -0.329 | -0.919 |
| TOTAL | | 1.000 | 5,280.00 | | |
| Predator: | (17) DECOMP/MICROF | | | | |
| 19. Benthic prod. | | 0.050 | 1,425.00 | -0.675 | 0.212 |
| 20. Detritus | | 0.950 | 27,075.00 | 0.411 | 0.992 |
| TOTAL | | 1.000 | 28,500.00 | | |
| Predator: | (20) Detritus | | | | |
| 1. SEA BIRDS | | 0.000 | 0.33 | 0.619 | 0.038 |
| 2. SHARK/RAY/SCOMB | | 0.000 | 1.15 | -0.480 | 0.003 |
| 3. L-IMJACKSCARN | | 0.000 | 4.48 | -0.300 | 0.005 |
| 4. L-IMSCHOOLPEL | | 0.002 | 45.16 | -0.202 | 0.006 |
| 5. SMALLSCHOOLPEL | | 0.003 | 76.69 | 0.107 | 0.011 |
| 6. LGGROUPERS-CARN | | 0.000 | 0.47 | -0.755 | 0.001 |
| 7. L-IMBENTHCARN | | 0.002 | 53.05 | -0.480 | 0.003 |

continued...

Table 8.8.2 continued

| | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|--------------------|------------------|-------------|--------------------------|---------------------------|
| 8. SMALLBENTHCARN | 0.004 | 103.78 | -0.348 | 0.004 |
| 9. BENTHINVOMNI | 0.024 | 556.60 | 0.206 | 0.014 |
| 10. L-IMREEFFHERB | 0.007 | 151.22 | -0.053 | 0.008 |
| 11. SMALLREEFFHERB | 0.002 | 48.25 | 0.226 | 0.014 |
| 12. BENTHINVHERB | 0.028 | 651.23 | -0.156 | 0.007 |
| 13. SEMISESSDETRIV | 0.012 | 276.03 | -0.490 | 0.003 |
| 14. SESSILEDETRIV | 0.100 | 2,314.37 | -0.301 | 0.005 |
| 15. CORALS | 0.011 | 253.59 | -0.370 | 0.004 |
| 16. ZOOPLANKTON | 0.049 | 1,148.09 | 0.773 | 0.070 |
| 17. DECOMP/MICROF | 0.249 | 5,786.93 | 0.976 | 0.752 |
| 18. Phytoplankton | 0.031 | 728.50 | 0.597 | 0.035 |
| 19. Benthic prod. | 0.474 | 10,997.76 | 0.296 | 0.016 |
| TOTAL | 1.000 | 23,197.67 | | |

Table 8.8.3. The 20-box model as obtained by method 2 (ECOPATH II aggregation routine).

| | | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|---------------------|------------------|------------------|-------------|--------------------------|---------------------------|
| Predator: | (1) Agg. 26 | | | | |
| | (0' Cycle) | | | | |
| 1. Agg. 26 | | 0.058 | 14.14 | 0.816 | 0.192 |
| 2. Agg. 24 | | 0.051 | 12.29 | 0.830 | 0.236 |
| 3. Agg. 20 | | 0.126 | 30.45 | 0.653 | -0.187 |
| 4. Agg. 22 | | 0.037 | 9.06 | 0.424 | -0.483 |
| 5. ENGRAULIDAEH | | 0.009 | 2.25 | 0.868 | 0.371 |
| 6. Agg. 25 | | 0.116 | 28.15 | 0.918 | 0.595 |
| 7. Agg. 21 | | 0.076 | 18.35 | 0.783 | 0.095 |
| 8. SMBENTHARTHROPO | | 0.045 | 10.80 | 0.889 | 0.458 |
| 9. GASTROPODS | | 0.055 | 13.42 | 0.713 | -0.071 |
| 10. CHITONS/SCAPHOP | | 0.015 | 3.57 | 0.091 | -0.712 |
| 11. POLY/PRIA/OPHIU | | 0.085 | 20.47 | 0.856 | 0.329 |
| 12. BIVALVES | | 0.042 | 10.25 | 0.323 | -0.568 |
| 13. ASCI/BARN/BRYOZ | | 0.027 | 6.64 | 0.004 | -0.753 |
| 14. SPONGES | | 0.083 | 19.98 | -0.315 | -0.865 |
| 15. CORALS/ANEMONES | | 0.003 | 0.67 | -0.792 | -0.968 |
| 16. ZOOPLANKTON | | 0.143 | 34.59 | 0.915 | 0.578 |
| 19. Benthic prod. | | 0.029 | 6.97 | -0.799 | -0.969 |
| 20. Detritus | | 0.000 | 0.03 | -0.999 | -1.000 |
| TOTAL | | 1.000 | 242.06 | | |
| Predator: | (2) Agg. 24 | | | | |
| | (0' Cycle) | | | | |
| 1. Agg. 26 | | 0.000 | 0.03 | -0.977 | -0.997 |
| 2. Agg. 24 | | 0.041 | 15.18 | 0.794 | 0.045 |
| 3. Agg. 20 | | 0.001 | 0.39 | -0.922 | -0.990 |
| 4. Agg. 22 | | 0.004 | 1.64 | -0.547 | -0.933 |
| 5. ENGRAULIDAEH | | 0.006 | 2.24 | 0.804 | 0.074 |
| 7. Agg. 21 | | 0.014 | 5.09 | 0.198 | -0.697 |
| 11. POLY/PRIA/OPHIU | | 0.045 | 16.62 | 0.747 | -0.079 |
| 13. ASCI/BARN/BRYOZ | | 0.016 | 5.74 | -0.272 | -0.873 |
| 16. ZOOPLANKTON | | 0.836 | 308.27 | 0.985 | 0.978 |
| 17. DECOMP/MICROFAU | | 0.003 | 1.15 | 0.023 | -0.778 |
| 19. Benthic prod. | | 0.033 | 12.19 | -0.772 | -0.970 |
| TOTAL | | 1.000 | 368.53 | | |
| Predator: | (3) Agg. 20 | | | | |
| | (0' Cycle) | | | | |
| 2. Agg. 24 | | 0.009 | 8.80 | 0.296 | 0.728 |
| 3. Agg. 20 | | 0.005 | 5.55 | -0.656 | -0.287 |
| 4. Agg. 22 | | 0.002 | 1.85 | -0.785 | -0.518 |
| 7. Agg. 21 | | 0.002 | 2.22 | -0.617 | -0.224 |
| 11. POLY/PRIA/OPHIU | | 0.001 | 1.48 | -0.635 | -0.253 |
| 12. BIVALVES | | 0.004 | 3.70 | -0.711 | -0.382 |
| 14. SPONGES | | 0.019 | 19.13 | -0.787 | -0.523 |
| 15. CORALS/ANEMONES | | 0.019 | 18.76 | -0.129 | 0.382 |
| 19. Benthic prod. | | 0.939 | 952.25 | 0.570 | 0.899 |
| TOTAL | | 1.000 | 1,013.73 | | |
| Predator: | (4) Agg. 22 | | | | |
| 7. Agg. 21 | | 0.002 | 0.99 | -0.642 | -0.642 |
| 9. GASTROPODS | | 0.001 | 0.27 | -0.888 | -0.889 |
| 11. POLY/PRIA/OPHIU | | 0.001 | 0.50 | -0.733 | -0.733 |
| 15. CORALS/ANEMONES | | 0.001 | 0.50 | -0.919 | -0.920 |
| 16. ZOOPLANKTON | | 0.000 | 0.11 | -0.933 | -0.933 |
| 17. DECOMP/MICROFAU | | 0.045 | 22.26 | 0.877 | 0.973 |
| 18. Phytoplankton | | 0.005 | 2.23 | -0.273 | -0.263 |
| 19. Benthic prod. | | 0.349 | 171.64 | 0.151 | 0.182 |
| 20. Detritus | | 0.596 | 293.29 | 0.202 | 0.236 |
| TOTAL | | 1.000 | 491.77 | | |
| Predator: | (5) ENGRAULIDAEH | | | | |
| 16. ZOOPLANKTON | | 0.200 | 28.86 | 0.939 | 0.742 |
| 18. Phytoplankton | | 0.700 | 101.01 | 0.978 | 0.963 |
| 20. Detritus | | 0.100 | 14.43 | -0.597 | -0.923 |
| TOTAL | | 1.000 | 144.31 | | |

continued...

Table 8.8.3 continued

| | | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|-----------|---------------------|------------------|-------------|--------------------------|---------------------------|
| Predator: | (6) Agg. 25 | | | | |
| 1. | Agg. 26 | 0.014 | 5.46 | 0.417 | 0.028 |
| 2. | Agg. 24 | 0.001 | 0.45 | -0.596 | -0.810 |
| 3. | Agg. 20 | 0.038 | 14.53 | 0.184 | -0.236 |
| 4. | Agg. 22 | 0.022 | 0.62 | -0.804 | -0.914 |
| 5. | ENGRAULIDAEH | 0.001 | 0.28 | 0.055 | -0.359 |
| 6. | Agg. 25 | 0.010 | 3.65 | 0.321 | -0.089 |
| 7. | Agg. 21 | 0.012 | 4.42 | 0.114 | -0.304 |
| 8. | SMBENTHARTHROPO | 0.014 | 5.43 | 0.690 | 0.437 |
| 9. | GASTROPODS | 0.042 | 16.10 | 0.641 | 0.354 |
| 11. | POLY/PRIA/OPHIU | 0.020 | 7.45 | 0.500 | 0.140 |
| 12. | BIVALVES | 0.046 | 17.63 | 0.364 | -0.037 |
| 13. | ASCI/BARN/BRYOZ | 0.018 | 6.83 | -0.204 | -0.567 |
| 14. | SPONGES | 0.043 | 16.32 | -0.573 | -0.798 |
| 15. | CORALS/ANEMONES | 0.014 | 5.43 | -0.253 | -0.601 |
| 16. | ZOOPLANKTON | 0.061 | 23.12 | 0.811 | 0.667 |
| 17. | DECOMP/MICROFAU | 0.021 | 8.06 | 0.754 | 0.554 |
| 18. | Phytoplankton | 0.014 | 5.32 | 0.278 | -0.137 |
| 19. | Benthic prod. | 0.514 | 195.15 | 0.332 | -0.075 |
| 20. | Detritus | 0.114 | 43.39 | -0.552 | -0.786 |
| TOTAL | | 1.000 | 379.66 | | |
| Predator: | (7) Agg. 21 | | | | |
| 1. | Agg. 26 | 0.017 | 7.52 | 0.493 | -0.244 |
| 2. | Agg. 24 | 0.003 | 1.34 | -0.204 | -0.765 |
| 3. | Agg. 20 | 0.082 | 35.26 | 0.513 | -0.219 |
| 4. | Agg. 22 | 0.004 | 1.86 | -0.556 | -0.892 |
| 5. | ENGRAULIDAEH | 0.006 | 2.69 | 0.809 | 0.356 |
| 6. | Agg. 25 | 0.007 | 2.99 | 0.168 | -0.557 |
| 7. | Agg. 21 | 0.047 | 20.05 | 0.668 | 0.030 |
| 8. | SMBENTHARTHROPO | 0.025 | 10.76 | 0.810 | 0.357 |
| 9. | GASTROPODS | 0.150 | 64.48 | 0.883 | 0.591 |
| 10. | CHITONS/SCAPHOP | 0.008 | 3.66 | -0.182 | -0.755 |
| 11. | POLY/PRIA/OPHIU | 0.040 | 17.37 | 0.721 | 0.137 |
| 12. | BIVALVES | 0.240 | 103.59 | 0.835 | 0.431 |
| 13. | ASCI/BARN/BRYOZ | 0.004 | 1.62 | -0.757 | -0.946 |
| 14. | SPONGES | 0.004 | 1.62 | -0.954 | -0.991 |
| 15. | CORALS/ANEMONES | 0.002 | 0.97 | -0.828 | -0.963 |
| 16. | ZOOPLANKTON | 0.094 | 40.35 | 0.873 | 0.555 |
| 17. | DECOMP/MICROFAU | 0.038 | 16.41 | 0.855 | 0.495 |
| 19. | Benthic prod. | 0.079 | 34.08 | -0.530 | -0.884 |
| 20. | Detritus | 0.149 | 64.34 | -0.453 | -0.859 |
| TOTAL | | 1.000 | 430.98 | | |
| Predator: | (8) SMBENTHARTHROPO | | | | |
| 13. | ASCI/BARN/BRYOZ | 0.050 | 83.14 | 0.295 | 0.174 |
| 14. | SPONGES | 0.060 | 99.77 | -0.451 | -0.567 |
| 15. | CORALS/ANEMONES | 0.040 | 66.51 | 0.251 | 0.123 |
| 16. | ZOOPLANKTON | 0.040 | 66.51 | 0.726 | 0.713 |
| 17. | DECOMP/MICROFAU | 0.040 | 66.51 | 0.862 | 0.904 |
| 19. | Benthic prod. | 0.600 | 997.73 | 0.399 | 0.295 |
| 20. | Detritus | 0.170 | 282.69 | -0.400 | -0.522 |
| TOTAL | | 1.000 | 1,662.88 | | |
| Predator: | (9) GASTROPODS | | | | |
| 4. | Agg. 22 | 0.015 | 9.83 | -0.005 | -0.260 |
| 6. | Agg. 25 | 0.002 | 1.31 | -0.424 | -0.619 |
| 7. | Agg. 21 | 0.006 | 3.93 | -0.213 | -0.448 |
| 8. | SMBENTHARTHROPO | 0.006 | 3.93 | 0.391 | 0.171 |
| 9. | GASTROPODS | 0.020 | 13.10 | 0.367 | 0.141 |
| 10. | CHITONS/SCAPHOP | 0.020 | 13.10 | 0.239 | -0.007 |
| 11. | POLY/PRIA/OPHIU | 0.050 | 32.76 | 0.769 | 0.703 |
| 12. | BIVALVES | 0.040 | 26.21 | 0.298 | 0.059 |
| 13. | ASCI/BARN/BRYOZ | 0.058 | 38.00 | 0.361 | 0.134 |
| 14. | SPONGES | 0.040 | 26.21 | -0.597 | -0.745 |
| 15. | CORALS/ANEMONES | 0.015 | 9.83 | -0.230 | -0.463 |
| 17. | DECOMP/MICROFAU | 0.030 | 19.66 | 0.820 | 0.786 |

continued...

Table 8.8.3 continued

| | Diet composition | Food intake | Ilev's electivity index | Standardized forage ratio |
|--------------------------------|------------------|-----------------|-------------------------|---------------------------|
| 19. Benthic prod. | 0.400 | 262.08 | 0.217 | -0.032 |
| 20. Detritus | 0.298 | 195.25 | -0.141 | -0.386 |
| TOTAL | 1.000 | 655.20 | | |
| Predator: (10) CHITONS/SCAPHOP | | | | |
| 13. ASCI/BARN/BRYOZ | 0.040 | 29.02 | 0.190 | -0.576 |
| 14. SPONGES | 0.020 | 14.51 | -0.776 | -0.955 |
| 15. CORALS/ANEMONES | 0.008 | 5.80 | -0.500 | -0.886 |
| 17. DECOMP/MICROFAU | 0.300 | 217.62 | 0.980 | 0.995 |
| 19. Benthic prod. | 0.632 | 458.45 | 0.421 | -0.376 |
| TOTAL | 1.000 | 725.40 | | |
| Predator: (11) POLY/PRIA/OPHIU | | | | |
| 8. SMBENTHARTHROPO | 0.015 | 30.49 | 0.702 | 0.383 |
| 9. GASTROPODS | 0.010 | 20.33 | 0.038 | -0.443 |
| 11. POLY/PRIA/OPHIU (0' Cycle) | 0.030 | 60.98 | 0.642 | 0.275 |
| 12. BIVALVES | 0.030 | 60.98 | 0.162 | -0.334 |
| 13. ASCI/BARN/BRYOZ | 0.040 | 81.31 | 0.190 | -0.308 |
| 14. SPONGES | 0.050 | 101.64 | -0.520 | -0.800 |
| 15. CORALS/ANEMONES | 0.010 | 20.33 | -0.411 | -0.743 |
| 16. ZOOPLANKTON | 0.050 | 101.64 | 0.775 | 0.528 |
| 17. DECOMP/MICROFAU | 0.050 | 101.64 | 0.888 | 0.791 |
| 18. Phytoplankton | 0.100 | 203.28 | 0.853 | 0.704 |
| 19. Benthic prod. | 0.245 | 498.04 | -0.025 | -0.493 |
| 20. Detritus | 0.370 | 752.14 | -0.034 | -0.500 |
| TOTAL | 1.000 | 2,032.80 | | |
| Predator: (12) BIVALVES | | | | |
| 17. DECOMP/MICROFAU | 0.100 | 103.79 | 0.942 | 0.937 |
| 18. Phytoplankton | 0.150 | 155.68 | 0.900 | 0.820 |
| 20. Detritus | 0.750 | 778.41 | 0.309 | -0.188 |
| TOTAL | 1.000 | 1,037.88 | | |
| Predator: (13) ASCI/BARN/BRYOZ | | | | |
| 16. ZOOPLANKTON | 0.150 | 412.20 | 0.919 | 0.764 |
| 17. DECOMP/MICROFAU | 0.100 | 274.80 | 0.942 | 0.854 |
| 18. Phytoplankton | 0.200 | 549.60 | 0.924 | 0.782 |
| 20. Detritus | 0.550 | 1,511.40 | 0.163 | -0.516 |
| TOTAL | 1.000 | 2,748.00 | | |
| Predator: (14) SPONGES | | | | |
| 17. DECOMP/MICROFAU | 0.001 | 4.00 | -0.496 | 0.434 |
| 20. Detritus | 0.999 | 3,996.00 | 0.432 | 0.986 |
| TOTAL | 1.000 | 4,000.00 | | |
| Predator: (15) CORALS/ANEMONES | | | | |
| 16. ZOOPLANKTON | 0.150 | 163.35 | 0.919 | 0.850 |
| 17. DECOMP/MICROFAU | 0.100 | 108.90 | 0.942 | 0.921 |
| 19. Benthic prod. | 0.650 | 707.85 | 0.432 | -0.091 |
| 20. Detritus | 0.100 | 108.90 | -0.597 | -0.852 |
| TOTAL | 1.000 | 1,089.00 | | |
| Predator: (16) ZOOPLANKTON | | | | |
| 17. DECOMP/MICROFAU | 0.600 | 3,168.00 | 0.990 | 0.987 |
| 18. Phytoplankton | 0.200 | 1,056.00 | 0.924 | 0.406 |
| 20. Detritus | 0.200 | 1,056.00 | -0.329 | -0.919 |
| TOTAL | 1.000 | 5,280.00 | | |

continued...

Table 8.8.3 continued

| | | Diet composition | Food intake | Ivlev's elec- tivity index | Standardized forage ratio |
|---------------------|----------------------|---------------------|----------------|-------------------------------|------------------------------|
| Predator: | (17) DECOMP/MICROFAU | | | | |
| 19. Benthic prod. | | 0.050 | 1,425.00 | -0.675 | 0.212 |
| 20. Detritus | | 0.950 | 27,075.00 | 0.411 | 0.992 |
| TOTAL | | 1.000 | 28,500.00 | | |
| Predator: | (20) Detritus | | | | |
| 1. Agg. 26 | | 0.002 | 53.07 | -0.442 | 0.003 |
| 2. Agg. 24 | | 0.004 | 81.93 | -0.144 | 0.007 |
| 3. Agg. 20 | | 0.011 | 256.67 | -0.409 | 0.004 |
| 4. Agg. 22 | | 0.004 | 102.59 | -0.548 | 0.003 |
| 5. ENGRAULIDAEH | | 0.001 | 30.83 | 0.337 | 0.018 |
| 6. Agg. 25 | | 0.003 | 79.87 | -0.179 | 0.006 |
| 7. Agg. 21 | | 0.004 | 90.62 | -0.406 | 0.004 |
| 8. SMBENTHARTHROPO | | 0.015 | 337.41 | 0.694 | 0.048 |
| 9. GASTROPODS | | 0.006 | 134.39 | -0.231 | 0.005 |
| 10. CHITONS/SCAPHOP | | 0.006 | 147.06 | -0.319 | 0.005 |
| 11. POLY/PRIA/OPHIU | | 0.018 | 420.53 | 0.470 | 0.024 |
| 12. BIVALVES | | 0.010 | 228.85 | -0.374 | 0.004 |
| 13. ASCI/BARN/BRYOZ | | 0.026 | 613.32 | -0.014 | 0.008 |
| 14. SPONGES | | 0.073 | 1,700.82 | -0.367 | 0.004 |
| 15. CORALS/ANEMONES | | 0.010 | 220.89 | -0.431 | 0.003 |
| 16. ZOOPLANKTON | | 0.050 | 1,157.00 | 0.775 | 0.069 |
| 17. DECOMP/MICROFAU | | 0.250 | 5,787.20 | 0.976 | 0.734 |
| 18. Phytoplankton | | 0.031 | 726.88 | 0.596 | 0.035 |
| 19. Benthic prod. | | 0.475 | 11,023.57 | 0.297 | 0.016 |
| TOTAL | | 1.000 | 23,193.50 | | |

Table 8.8.4. The 11-box model as obtained by method 1 (cluster analysis + intuition).

| | | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|--------------------|---------------------|------------------|-------------|--------------------------|---------------------------|
| Predator: | (1) APEX PREDATORS | | | | |
| | (0' Cycle) | | | | |
| 1. APEX PREDATORS | | 0.026 | 0.59 | 0.962 | 0.722 |
| 2. SCHOOLPELAGICS | | 0.320 | 7.22 | 0.966 | 0.761 |
| 3. BENTHCARNIVORES | | 0.279 | 6.30 | 0.892 | 0.198 |
| 4. BENTHINVOMNI | | 0.016 | 0.36 | 0.006 | -0.859 |
| 5. BENTHHERBIVORES | | 0.328 | 7.40 | 0.645 | -0.475 |
| 6. BENTHDETRITIVOR | | 0.012 | 0.27 | -0.897 | -0.992 |
| 7. ZOOPLANKTON | | 0.018 | 0.41 | 0.479 | -0.646 |
| 11. Detritus | | 0.001 | 0.02 | -0.995 | -1.000 |
| TOTAL | | 1.000 | 22.56 | | |
| Predator: | (2) SCHOOLPELAGICS | | | | |
| | (0' Cycle) | | | | |
| 2. SCHOOLPELAGICS | | 0.047 | 26.16 | 0.787 | -0.185 |
| 3. BENTHCARNIVORES | | 0.006 | 3.34 | -0.452 | -0.944 |
| 4. BENTHINVOMNI | | 0.030 | 16.70 | 0.310 | -0.743 |
| 5. BENTHHERBIVORES | | 0.006 | 3.34 | -0.844 | -0.987 |
| 6. BENTHDETRITIVOR | | 0.010 | 5.57 | -0.913 | -0.993 |
| 7. ZOOPLANKTON | | 0.607 | 337.84 | 0.979 | 0.930 |
| 8. DECOMP/MICROFAU | | 0.002 | 1.11 | -0.195 | -0.901 |
| 9. Phytoplankton | | 0.181 | 100.74 | 0.916 | 0.359 |
| 10. Benthic prod. | | 0.085 | 47.31 | -0.504 | -0.951 |
| 11. Detritus | | 0.026 | 14.47 | -0.877 | -0.990 |
| TOTAL | | 1.000 | 556.57 | | |
| Predator: | (3) BENTHCARNIVORES | | | | |
| | (0' Cycle) | | | | |
| 2. SCHOOLPELAGICS | | 0.018 | 13.28 | 0.525 | -0.192 |
| 3. BENTHCARNIVORES | | 0.077 | 56.80 | 0.657 | 0.028 |
| 4. BENTHINVOMNI | | 0.167 | 123.20 | 0.827 | 0.450 |
| 5. BENTHHERBIVORES | | 0.169 | 124.68 | 0.409 | -0.338 |
| 6. BENTHDETRITIVOR | | 0.213 | 157.14 | -0.017 | -0.674 |
| 7. ZOOPLANKTON | | 0.122 | 90.00 | 0.901 | 0.720 |
| 8. DECOMP/MICROFAU | | 0.026 | 19.18 | 0.795 | 0.353 |
| 10. Benthic prod. | | 0.095 | 70.08 | -0.461 | -0.863 |
| 11. Detritus | | 0.113 | 83.36 | -0.556 | -0.893 |
| TOTAL | | 1.000 | 737.73 | | |
| Predator: | (4) BENTHINVOMNI | | | | |
| | (0' Cycle) | | | | |
| 3. BENTHCARNIVORES | | 0.004 | 10.75 | -0.598 | -0.872 |
| 4. BENTHINVOMNI | | 0.0047 | 126.34 | 0.497 | -0.066 |
| 5. BENTHHERBIVORES | | 0.024 | 64.51 | -0.494 | -0.830 |
| 6. BENTHDETRITIVOR | | 0.128 | 344.06 | -0.265 | -0.724 |
| 7. ZOOPLANKTON | | 0.038 | 102.14 | 0.714 | 0.320 |
| 8. DECOMP/MICROFAU | | 0.045 | 120.96 | 0.876 | 0.749 |
| 9. Phytoplankton | | 0.076 | 204.29 | 0.811 | 0.557 |
| 10. Benthic prod. | | 0.286 | 768.76 | 0.052 | -0.526 |
| 11. Detritus | | 0.352 | 946.17 | -0.059 | -0.604 |
| TOTAL | | 1.000 | 2,687.98 | | |
| Predator: | (5) BENTHHERBIVORES | | | | |
| | (0' Cycle) | | | | |
| 3. BENTHCARNIVORES | | 0.002 | 10.02 | -0.777 | -0.937 |
| 4. BENTHINVOMNI | | 0.004 | 20.05 | -0.596 | -0.876 |
| 5. BENTHHERBIVORES | | 0.019 | 95.22 | -0.577 | -0.869 |
| 6. BENTHDETRITIVOR | | 0.057 | 285.65 | -0.589 | -0.874 |
| 7. ZOOPLANKTON | | 0.049 | 245.56 | 0.771 | 0.429 |
| 8. DECOMP/MICROFAU | | 0.080 | 400.91 | 0.928 | 0.917 |
| 9. Phytoplankton | | 0.001 | 5.01 | -0.776 | -0.936 |
| 10. Benthic prod. | | 0.680 | 3,407.77 | 0.451 | -0.153 |
| 11. Detritus | | 0.108 | 541.23 | -0.572 | -0.867 |
| TOTAL | | 1.000 | 5,011.42 | | |
| Predator: | (6) BENTHDETRITIVOR | | | | |
| 7. ZOOPLANKTON | | 0.051 | 408.43 | 0.779 | 0.457 |
| 8. DECOMP/MICROFAU | | 0.050 | 400.42 | 0.888 | 0.776 |

continued...

Table 8.8.4 continued

| | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|---------------------|------------------|-------------|--------------------------|---------------------------|
| 9. Phytoplankton | 0.088 | 704.74 | 0.835 | 0.610 |
| 10. Benthic prod. | 0.004 | 32.03 | -0.969 | -0.992 |
| 11. Detritus | 0.807 | 6,462.74 | 0.341 | -0.277 |
| TOTAL | 1.000 | 8,008.36 | | |
| Predator: | | | | |
| (7) ZOOPLANKTON | | | | |
| 8. DECOMP/MICROFAU | 0.600 | 3,168.00 | 0.990 | 0.975 |
| 9. Phytoplankton | 0.200 | 1,056.00 | 0.924 | 0.110 |
| 11. Detritus | 0.200 | 1,056.00 | -0.329 | -0.957 |
| TOTAL | 1.000 | 5,280.00 | | |
| Predator: | | | | |
| (8) DECOMP/MICROFAU | | | | |
| 10. Benthic prod. | 0.050 | 1,425.00 | -0.675 | -0.105 |
| 11. Detritus | 0.950 | 27,075.00 | 0.411 | 0.984 |
| TOTAL | 1.000 | 28,500.00 | | |
| Predator: | | | | |
| (11) Detritus | | | | |
| 1. APEX PREDATORS | 0.000 | 6.23 | -0.304 | 0.005 |
| 2. SCHOOLPELAGICS | 0.005 | 121.76 | -0.032 | 0.009 |
| 3. BENTHCARNIVORES | 0.007 | 157.94 | -0.401 | 0.004 |
| 4. BENTHINVOMNI | 0.024 | 553.56 | 0.203 | 0.015 |
| 5. BENTHHERBIVORES | 0.048 | 1,106.77 | -0.195 | 0.007 |
| 6. BENTHDETRITIVOR | 0.112 | 2,588.50 | -0.328 | 0.005 |
| 7. ZOOPLANKTON | 0.050 | 1,151.63 | 0.774 | 0.077 |
| 8. DECOMP/MICROFAU | 0.250 | 5,789.42 | 0.976 | 0.822 |
| 9. Phytoplankton | 0.031 | 729.23 | 0.597 | 0.039 |
| 10. Benthic prod. | 0.474 | 10,994.04 | 0.296 | 0.018 |
| TOTAL | 1.000 | 23,199.07 | | |

Table 8.8.5. The 11-box model as obtained by method 2 (ECOPATH II aggregation routine).

| | | Diet composition | Food intake | Ivlev's electivity index | Standardized forage ratio |
|--------------------|-----------------|------------------|-------------|--------------------------|---------------------------|
| Predator: | (1) Agg. 12 | | | | |
| | (0' Cycle) | | | | |
| 1. Agg. 12 | | 0.090 | 94.00 | 0.639 | -0.244 |
| 2. Agg. 20 | | 0.063 | 66.10 | 0.413 | -0.523 |
| 3. Agg. 15 | | 0.169 | 176.06 | 0.650 | -0.226 |
| 4. Agg. 11 | | 0.130 | 135.01 | 0.447 | -0.491 |
| 5. Agg. 13 | | 0.029 | 30.44 | -0.142 | -0.825 |
| 6. SPONGES | | 0.021 | 21.60 | -0.769 | -0.967 |
| 7. ZOOPLANKTON | | 0.368 | 383.20 | 0.966 | 0.930 |
| 8. DECOMP/MICROFAU | | 0.017 | 17.56 | 0.700 | -0.129 |
| 10. Benthic prod. | | 0.051 | 53.24 | -0.669 | -0.951 |
| 11. Detritus | | 0.062 | 64.37 | -0.730 | -0.961 |
| TOTAL | | 1.000 | 1,041.58 | | |
| Predator: | (2) Agg. 20 | | | | |
| | (0' Cycle) | | | | |
| 1. Agg. 12 | | 0.011 | 11.02 | -0.293 | 0.084 |
| 2. Agg. 20 | | 0.005 | 5.55 | -0.656 | -0.409 |
| 3. Agg. 15 | | 0.003 | 3.33 | -0.832 | -0.694 |
| 4. Agg. 11 | | 0.004 | 3.70 | -0.863 | -0.747 |
| 5. Agg. 13 | | 0.019 | 18.76 | -0.355 | 0.008 |
| 6. SPONGES | | 0.019 | 19.13 | -0.787 | -0.618 |
| 10. Benthic prod. | | 0.939 | 952.25 | 0.570 | 0.920 |
| TOTAL | | 1.000 | 1,013.73 | | |
| Predator: | (3) Agg. 15 | | | | |
| | (0' Cycle) | | | | |
| 1. Agg. 12 | | 0.004 | 15.25 | -0.645 | -0.878 |
| 2. Agg. 20 | | 0.004 | 14.53 | -0.732 | -0.911 |
| 3. Agg. 15 | | 0.047 | 166.91 | 0.133 | -0.421 |
| 4. Agg. 11 | | 0.065 | 231.24 | 0.135 | -0.419 |
| 5. Agg. 13 | | 0.025 | 89.04 | -0.217 | -0.671 |
| 6. SPONGES | | 0.041 | 144.17 | -0.593 | -0.857 |
| 7. ZOOPLANKTON | | 0.035 | 124.87 | 0.694 | 0.331 |
| 8. DECOMP/MICROFAU | | 0.043 | 151.61 | 0.870 | 0.766 |
| 9. Phytoplankton | | 0.059 | 210.83 | 0.764 | 0.485 |
| 10. Benthic prod. | | 0.317 | 1,126.91 | 0.103 | -0.447 |
| 11. Detritus | | 0.361 | 1,284.06 | -0.047 | -0.562 |
| TOTAL | | 1.000 | 3,559.43 | | |
| Predator: | (4) Agg. 11 | | | | |
| 7. ZOOPLANKTON | | 0.112 | 441.06 | 0.893 | 0.495 |
| 8. DECOMP/MICROFAU | | 0.096 | 378.59 | 0.940 | 0.756 |
| 9. Phytoplankton | | 0.205 | 806.29 | 0.926 | 0.668 |
| 11. Detritus | | 0.586 | 2,304.24 | 0.193 | -0.674 |
| TOTAL | | 1.000 | 3,930.18 | | |
| Predator: | (5) Agg. 13 | | | | |
| | (0' Cycle) | | | | |
| 4. Agg. 11 | | 0.032 | 112.16 | -0.211 | -0.777 |
| 5. Agg. 13 | | 0.021 | 72.32 | -0.303 | -0.814 |
| 6. SPONGES | | 0.033 | 114.28 | -0.656 | -0.924 |
| 7. ZOOPLANKTON | | 0.066 | 229.87 | 0.825 | 0.424 |
| 8. DECOMP/MICROFAU | | 0.113 | 393.04 | 0.949 | 0.926 |
| 10. Benthic prod. | | 0.622 | 2,164.03 | 0.415 | -0.350 |
| 11. Detritus | | 0.113 | 391.59 | -0.557 | -0.897 |
| TOTAL | | 1.000 | 3,477.28 | | |
| Predator: | (6) SPONGES | | | | |
| 8. DECOMP/MICROFAU | | 0.001 | 4.00 | -0.496 | 0.143 |
| 11. Detritus | | 0.999 | 3,996.00 | 0.432 | 0.974 |
| TOTAL | | 1.000 | 4,000.00 | | |
| Predator: | (7) ZOOPLANKTON | | | | |
| 8. DECOMP/MICROFAU | | 0.600 | 3,168.00 | 0.990 | 0.975 |
| 9. Phytoplankton | | 0.200 | 1,056.00 | 0.924 | 0.110 |
| 11. Detritus | | 0.200 | 1,056.00 | -0.329 | -0.957 |
| TOTAL | | 1.000 | 5,280.00 | | |

continued...

Table 8.8.5 continued

| | | Diet composition | Food intake | Ivlev's elec- tivity index | Standardized forage ratio |
|--------------------|---------------------|---------------------|----------------|-------------------------------|------------------------------|
| Predator: | (8) DECOMP/MICROFAU | | | | |
| 10. Benthic prod. | | 0.050 | 1,425.00 | -0.675 | -0.105 |
| 11. Detritus | | 0.950 | 27,075.00 | 0.411 | 0.984 |
| TOTAL | | 1.000 | 28,500.00 | | |
| Predator: | (11) Detritus | | | | |
| 1. Agg. 12 | | 0.010 | 225.63 | -0.343 | 0.005 |
| 2. Agg. 20 | | 0.011 | 256.67 | -0.409 | 0.004 |
| 3. Agg. 15 | | 0.032 | 737.37 | -0.061 | 0.009 |
| 4. Agg. 11 | | 0.038 | 873.00 | -0.136 | 0.007 |
| 5. Agg. 13 | | 0.030 | 705.36 | -0.122 | 0.008 |
| 6. SPONGES | | 0.073 | 1,700.82 | -0.367 | 0.005 |
| 7. ZOOPLANKTON | | 0.050 | 1,157.00 | 0.775 | 0.078 |
| 8. DECOMP/MICROFAU | | 0.250 | 5,787.20 | 0.976 | 0.828 |
| 9. Phytoplankton | | 0.031 | 726.88 | 0.596 | 0.039 |
| 10. Benthic prod. | | 0.475 | 11,023.57 | 0.297 | 0.018 |
| TOTAL | | 1.000 | 23,193.50 | | |

Appendix 8.9. Summary statistics and flows between discrete trophic levels for the 20- and 11-box models from two methods.

Table 8.9.1. Summary statistics for a 20-box model of a Caribbean coral reef (method 1). PP = primary production.

| | | | |
|---------------------------------|---|---------|---------------------------------------|
| Sum of all production | = | 51,285 | g·m ⁻² ·year ⁻¹ |
| Sum of all imports ^a | = | 475 | g·m ⁻² ·year ⁻¹ |
| Sum of all respiratory flows | = | 32,525 | g·m ⁻² ·year ⁻¹ |
| Sum of all flows into detritus | = | 23,198 | g·m ⁻² ·year ⁻¹ |
| Total system throughput | = | 107,483 | g·m ⁻² ·year ⁻¹ |
| Throughput cycled | = | 20,459 | g·m ⁻² ·year ⁻¹ |
| Finn's cycling index | = | 19.03% | of throughput |
| Input total net PP | = | 20,500 | g·m ⁻² ·year ⁻¹ |
| Calculated total net PP | = | 20,025 | g·m ⁻² ·year ⁻¹ |

^aDifference between input total net PP and calculated total net PP.

Table 8.9.2. Network flow indices for a 20-box model of a Caribbean coral reef (method 1).

| Source | Ascendency | | Overhead | | Capacity | |
|-------------|------------|------|----------|------|----------|-------|
| | flowbits | % | flowbits | % | flowbits | % |
| Input | 0 | 0.0 | 0 | 0.0 | 0 | 0.0 |
| Internal | 90,203 | 22.4 | 209,148 | 51.9 | 299,351 | 74.3 |
| Dissipation | 52,951 | 13.1 | 50,589 | 12.6 | 103,540 | 25.7 |
| Total | 143,154 | 35.5 | 259,737 | 64.5 | 402,891 | 100.0 |

Table 8.9.3 Flows and transfer efficiencies for discrete trophic levels of a 20-box model (method 1).

| Flows originating from the primary producers | | | | | |
|----------------------------------------------|--------|---------|-------------|------------|------------------|
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 0.000 | 480.000 | 0.000 | 0.000 | 11,726.258 |
| II | 0.000 | 0.000 | 5,226.313 | 7,818.739 | 1,748.263 |
| III | 0.000 | 0.000 | 550.389 | 844.163 | 194.366 |
| IV | 0.000 | 0.000 | 67.132 | 99.408 | 23.043 |
| V | 0.000 | 0.000 | 6.358 | 9.233 | 1.992 |
| VI | 0.000 | 0.000 | 0.602 | 0.883 | 0.192 |
| VII | 0.000 | 0.000 | 0.061 | 0.089 | 0.019 |
| VIII | 0.000 | 0.000 | 0.006 | 0.009 | 0.002 |
| IX | 0.000 | 0.000 | 0.001 | 0.001 | 0.000 |

| Flows originating from the detritus | | | | | |
|-------------------------------------|---------|--------|-------------|------------|------------------|
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 475.000 | 0.000 | 0.000 | 23,172.668 | 0.000 |
| II | 0.000 | 0.000 | 23,098.771 | 36,177.758 | 8,161.688 |
| III | 0.000 | 0.000 | 2,954.389 | 4,917.297 | 1,108.677 |
| IV | 0.000 | 0.000 | 563.005 | 854.232 | 215.283 |
| V | 0.000 | 0.000 | 53.000 | 75.944 | 16.284 |
| VI | 0.000 | 0.000 | 4.550 | 6.660 | 1.438 |
| VII | 0.000 | 0.000 | 0.457 | 0.671 | 0.146 |
| VIII | 0.000 | 0.000 | 0.046 | 0.068 | 0.015 |
| IX | 0.000 | 0.000 | 0.005 | 0.007 | 0.002 |

Table 8.9.4. Summary statistics for a 20-box model of a Caribbean coral reef (method 2). PP = primary production.

| | | | |
|--------------------------------|---|----------------------|---------------------------------------|
| Sum of all production | = | 51,282 | g·m ⁻² ·year ⁻¹ |
| Sum of all imports* | = | 475 | g·m ⁻² ·year ⁻¹ |
| Sum of all respiratory flows | = | 32,523 | g·m ⁻² ·year ⁻¹ |
| Sum of all flows into detritus | = | 23,193 | g·m ⁻² ·year ⁻¹ |
| <hr/> | | | |
| Total system throughput | = | 107,473 | g·m ⁻² ·year ⁻¹ |
| Throughput cycled | = | 19,726 | g·m ⁻² ·year ⁻¹ |
| Finn's cycling index | = | 18.35% of throughput | |
| <hr/> | | | |
| Input total net PP | = | 20,500 | g·m ⁻² ·year ⁻¹ |
| Calculated total net PP | = | 20,025 | g·m ⁻² ·year ⁻¹ |

*Difference between input total net PP and calculated total net PP.

Table 8.9.5. Network flow indices for a 20-box model of a Caribbean coral reef (method 2).

| Source | Ascendency | | Overhead | | Capacity | |
|-------------|------------|------|----------|------|----------|-------|
| | flowbits | % | flowbits | % | flowbits | % |
| Input | 952 | 0.2 | 2,678 | 0.6 | 3,630 | 0.9 |
| Internal | 93,243 | 21.8 | 221,083 | 51.6 | 314,326 | 73.3 |
| Dissipation | 52,340 | 12.2 | 58,334 | 13.6 | 110,674 | 25.8 |
| <hr/> | | | | | | |
| Total | 146,535 | 34.2 | 282,095 | 65.8 | 428,630 | 100.0 |

Table 8.9.6. Flows and transfer efficiencies for discrete trophic levels of a 20-box model (method 2).

| Flows originating from the primary producers | | | | | |
|----------------------------------------------|---------|---------|-------------|------------|------------------|
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 0.000 | 480.000 | 0.000 | 0.000 | 11,750.450 |
| II | 0.000 | 0.000 | 5,241.900 | 7,794.550 | 1,668.085 |
| III | 0.000 | 0.000 | 587.240 | 884.564 | 189.563 |
| IV | 0.000 | 0.000 | 73.484 | 107.762 | 23.132 |
| V | 0.000 | 0.000 | 7.547 | 11.145 | 2.394 |
| VI | 0.000 | 0.000 | 0.806 | 1.204 | 0.259 |
| VII | 0.000 | 0.000 | 0.093 | 0.139 | 0.030 |
| VIII | 0.000 | 0.000 | 0.011 | 0.016 | 0.003 |
| Flows originating from the detritus | | | | | |
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 475.000 | 0.000 | 0.000 | 23,668.496 | 0.000 |
| II | 0.000 | 0.000 | 22,976.932 | 36,171.262 | 8,299.344 |
| III | 0.000 | 0.000 | 2,991.129 | 4,894.986 | 1,058.519 |
| IV | 0.000 | 0.000 | 580.908 | 845.338 | 182.146 |
| V | 0.000 | 0.000 | 56.716 | 82.284 | 17.643 |
| VI | 0.000 | 0.000 | 5.313 | 7.925 | 1.705 |
| VII | 0.000 | 0.000 | 0.606 | 0.906 | 0.195 |
| VIII | 0.000 | 0.000 | 0.070 | 0.105 | 0.023 |

Table 8.9.7. Summary statistics for an 11-box model of a Caribbean reef (method 1). PP = primary production.

| | | | |
|--------------------------------|---|----------------------|---------------------------------------|
| Sum of all production | = | 51,285 | g·m ⁻² ·year ⁻¹ |
| Sum of all imports* | = | 475 | g·m ⁻² ·year ⁻¹ |
| Sum of all respiratory flows | = | 32,525 | g·m ⁻² ·year ⁻¹ |
| Sum of all flows into detritus | = | 23,199 | g·m ⁻² ·year ⁻¹ |
| <hr/> | | | |
| Total system throughput | = | 107,484 | g·m ⁻² ·year ⁻¹ |
| Throughput cycled | = | 20,900 | g·m ⁻² ·year ⁻¹ |
| Finn's cycling index | = | 19.45% of throughput | |
| <hr/> | | | |
| Input total net PP | = | 20,500 | g·m ⁻² ·year ⁻¹ |
| Calculated total net PP | = | 20,025 | g·m ⁻² ·year ⁻¹ |

*Difference between input total net PP and calculated total net PP.

Table 8.9.8. Network flow indices for an 11-box model of a Caribbean coral reef (method 1).

| Source | Ascendency | | Overhead | | Capacity | |
|--------------|----------------|-------------|----------------|-------------|----------------|--------------|
| | flowbits | % | flowbits | % | flowbits | % |
| Input | 0 | 0.0 | 0 | 0.0 | 0 | 0.0 |
| Internal | 88,466 | 23.5 | 194,187 | 51.5 | 282,653 | 75.0 |
| Dissipation | 52,947 | 14.0 | 41,325 | 11.0 | 94,272 | 25.0 |
| Total | 141,413 | 37.5 | 235,512 | 62.5 | 376,925 | 100.0 |

Table 8.9.9. Flows and transfer efficiencies for discrete trophic levels of an 11-box model (method 1).

| Flows originating from the primary producers | | | | | |
|----------------------------------------------|---------|---------|-------------|------------|------------------|
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 0.000 | 480.000 | 0.000 | 0.000 | 11,723.268 |
| II | 0.000 | 0.000 | 5,221.518 | 7,821.728 | 1,759.426 |
| III | 0.000 | 0.000 | 549.314 | 840.784 | 192.542 |
| IV | 0.000 | 0.000 | 66.822 | 98.928 | 22.744 |
| V | 0.000 | 0.000 | 6.411 | 9.363 | 2.037 |
| VI | 0.000 | 0.000 | 0.624 | 0.915 | 0.200 |
| VII | 0.000 | 0.000 | 0.062 | 0.091 | 0.020 |
| VIII | 0.000 | 0.000 | 0.006 | 0.009 | 0.002 |
| Flows originating from the detritus | | | | | |
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 475.000 | 0.000 | 0.000 | 23,174.063 | 0.000 |
| II | 0.000 | 0.000 | 23,098.945 | 36,179.004 | 8,154.590 |
| III | 0.000 | 0.000 | 2,963.081 | 4,925.468 | 1,118.325 |
| IV | 0.000 | 0.000 | 560.205 | 844.063 | 207.598 |
| V | 0.000 | 0.000 | 52.484 | 76.260 | 16.565 |
| VI | 0.000 | 0.000 | 4.923 | 7.210 | 1.572 |
| VII | 0.000 | 0.000 | 0.487 | 0.715 | 0.156 |
| VIII | 0.000 | 0.000 | 0.049 | 0.071 | 0.016 |

Table 8.9.10. Summary statistics for an 11-box model of a Caribbean coral reef (method 2). PP = primary production.

| | | | |
|--------------------------------|---|---------|---------------------------------------------------|
| Sum of all production | = | 51,282 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |
| Sum of all imports* | = | 475 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |
| Sum of all respiratory flows | = | 32,523 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |
| Sum of all flows into detritus | = | 23,193 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |
| | | | |
| Total system throughput | = | 107,473 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |
| Throughput cycled | = | 20,117 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |
| Finn's cycling index | = | 18.70% | of throughput |
| | | | |
| Input total net PP | = | 20,500 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |
| Calculated total net PP | = | 20,025 | $\text{g}\cdot\text{m}^{-2}\cdot\text{year}^{-1}$ |

*Difference between input total net PP and calculated total net PP.

Table 8.9.11. Network flow indices for an 11-box model of a Caribbean coral reef (method 2).

| Source | Ascendency | | Overhead | | Capacity | |
|--------------|----------------|-------------|----------------|-------------|----------------|--------------|
| | flowbits | % | flowbits | % | flowbits | % |
| Input | 952 | 0.2 | 2,678 | 0.7 | 3,630 | 0.9 |
| Internal | 90,229 | 22.7 | 205,030 | 51.5 | 295,259 | 74.2 |
| Dissipation | 52,313 | 13.1 | 46,835 | 11.8 | 99,148 | 24.9 |
| Total | 143,494 | 36.0 | 254,543 | 64.0 | 398,037 | 100.0 |

Table 8.9.12. Flows and transfer efficiencies for discrete trophic levels of an 11-box model (method 2).

| Flows originating from the primary producers | | | | | |
|----------------------------------------------|---------|---------|-------------|------------|------------------|
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 0.000 | 480.000 | 0.000 | 0.000 | 11,750.449 |
| II | 0.000 | 0.000 | 5,237.327 | 7,794.550 | 1,668.588 |
| III | 0.000 | 0.000 | 589.595 | 888.635 | 190.385 |
| IV | 0.000 | 0.000 | 74.016 | 108.655 | 23.315 |
| V | 0.000 | 0.000 | 7.692 | 11.325 | 2.439 |
| VI | 0.000 | 0.000 | 0.805 | 1.194 | 0.259 |
| VII | 0.000 | 0.000 | 0.087 | 0.130 | 0.028 |
| Flows originating from detritus | | | | | |
| Trophic level | Import | Export | Respiration | Throughput | Flow to detritus |
| I | 475.000 | 0.000 | 0.000 | 23,668.494 | 0.000 |
| II | 0.000 | 0.000 | 22,991.117 | 36,171.262 | 8,299.580 |
| III | 0.000 | 0.000 | 2,977.325 | 4,880.564 | 1,055.676 |
| IV | 0.000 | 0.000 | 577.533 | 847.562 | 181.747 |
| V | 0.000 | 0.000 | 60.478 | 88.282 | 18.856 |
| VI | 0.000 | 0.000 | 6.053 | 8.947 | 1.934 |
| VII | 0.000 | 0.000 | 0.647 | 0.961 | 0.209 |



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