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Dataset Paper

Detailed Food Web Networks of Three Greater Antillean Coral Reef Systems: The Cayman Islands, Cuba, and Jamaica

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Food webs represent one of the most complex aspects of community biotic interactions. Complex food webs are represented as networks of interspecific interactions, where nodes represent species or groups of species, and links are predator-prey interactions. This paper presents reconstructions of coral reef food webs in three Greater Antillean regions of the Caribbean: the Cayman Islands, Cuba, and Jamaica. Though not taxonomically comprehensive, each food web nevertheless comprises producers and consumers, single-celled and multicellular organisms, and species foraging on reefs and adjacent seagrass beds. Species are grouped into trophic guilds if their prey and predator links are indistinguishable. The data list guilds, taxonomic composition, prey guilds/species, and predators. Primary producer and invertebrate richness are regionally uniform, but vertebrate richness varies on the basis of more detailed occurrence data. Each region comprises 169 primary producers, 513 protistan and invertebrate consumer species, and 159, 178, and 170 vertebrate species in the Cayman Islands, Cuba, and Jamaica, respectively. Caribbean coral reefs are among the world's most endangered by anthropogenic activities. The datasets presented here will facilitate comparisons of historical and regional variation, the assessment of impacts of species loss and invasion, and the application of food webs to ecosystem analyses.

1. Introduction

Coral reef communities of the Greater Antilles of the Caribbean Sea have a long history of anthropogenic disturbance, driven by the exploitation for food of both vertebrate and invertebrate species [1, 2]. More recently, coral bleaching, storm effects, coral disease, coastal development, pollution, invasive species, and a reduction of herbivorous control of algae in spatial competition with coral [3] have resulted in dramatic declines of diversity and abundance on reefs throughout the region [4-6]. The ongoing and predicted increases of seawater temperature and acidification as consequences of anthropogenic global warming make coral reefs among the most endangered ocean ecosystems, and Greater Antillean reefs may be particularly vulnerable because of their past and recent histories of perturbation [7]. It is therefore increasingly important to integrate species-level data on systematics, ecology, and biogeography into systems-level data that are informative to regional conservation and management efforts. Here we use food web networks, in close regional proximity, to represent one important aspect of species ecology and present regional variation of detailed ecological networks.

Food web networks seek to capture the complexity of patterns of trophic interaction in biological communities [8]. These networks serve at least two purposes from the perspective of ecosystem protection. First, the basic topology of a food web network is a measure of the robustness or resistance of that community to species removal [9]. Such removals represent local or global extinctions. Network representation of communities can be used to discover interactions that would be lost as a consequence of extinction, and chains or pathways that would be disrupted [10]. They may be used additionally to infer possible secondary extinctions as a result of lost or disrupted paths [11]. The simplest such inference, topological secondary extinction [12], predicts the



FIGURE 1: Map of the northern Caribbean Basin showing regions covered in this paper. Outlines around each region, the Cayman Islands, Cuba, and Jamaica, trace the 100-meter depth contour around each island or group of islands, within which all reef sites used in this study are included. Cyan colour: Cayman Islands; green colour: Cuba; red colour: Jamaica. The region outlined in red, but disconnected from the island of Jamaica, represents Pedro Bank (Map courtesy of The World Marble Globe, version 1.3.3, http://edu.kde.org/marble/).

secondary extinction of a species if it loses all its incoming paths, that is, its prey resources, as a consequence of an initial extinction. Topological extinction thus measures the structural robustness of a food web [13], but it is limited to a minimum estimate of secondary extinction because it functions only in the direction of energy flow (bottom-up) [14]. A further limitation is imposed by the typical lack of demographic parameters in complex food web networks (but see [15]), forcing one to ignore demographic changes that could result from the initial extinction(s), such as top-down trophic cascades and Allee effects. Nevertheless, parameter modeling can indicate the potential for demographic instability and tipping points in the community.

A second purpose is the assessment of temporal and geographic variance of interactions within communities. Temporal variation will arise under conditions of varying or changing environments, such as those caused by climate change or biotic invasions, and the possible microevolutionary adaptation to such changes. Geographic variation is expected on the basis of varying composition between communities (β diversity), even those of the same type, whether driven by local environmental differences or stochastic processes, as well as different histories and regimes of anthropogenic disturbance. This measurement of temporal and geographic variation, however, is challenged by the level of ecological detail required to construct a food web network.

The food webs presented in this paper capture, to the greatest extent currently possible, the patterns of trophic interactions in coral reef communities of the Cayman Islands, Cuba, and Jamaica. In constructing the food webs we strove to include as many species as possible, their trophic interactions, and at least for vertebrates whether those species foraged preferentially on coral reefs, or seagrass beds, or both. The results are three highly detailed food webs that should both

serve as useful sources of ecological data within each region and facilitate comparisons across that area of the Greater Antilles and beyond as the number of similar regional datasets increases. The datasets will also be useful tools for assessing the robustness of these ecologically and economically important communities, and the potential outcomes of various types of disturbance and conservation measures.

2. Methodology

This study considers coral reef systems in three national regions: the Cayman Islands, Cuba, and Jamaica. Though the systems are not contiguous within each region, for example, offshore islands and shallows such as Pedro Bank off Jamaica, we consider the reefs within each region to be subject to uniform political administration and subject to similar anthropogenic policies and activities (Figure 1). We therefore refer in the study to three individual regions, namely, the Cayman Islands, Cuba, and Jamaica. Regions are certainly heterogeneous in terms of reef physiography and oceanographic conditions as well as anthropogenic factors. For example, each region has established marineprotected areas (41, 42, and 12 in number for the Cayman Islands, Cuba, and Jamaica, resp.; see World Bank Data Catalog, http://data.worldbank.org/data-catalog) where one could expect local species richness and abundance of exploited species to be greater than areas where harvesting is permitted [16–18]. Nevertheless, our data are pooled across localities in each region for two reasons. First, the absence of a species from a locality because of anthropogenic impacts could be intermittent and dependent on the timescales of replenishment by immigration and extirpation. Second, food webs vary in time and space in terms of both species composition as well as the nature (strength and even direction) of trophic interactions [12]. It is therefore useful to present data that are integrated across fine-scale spatial and temporal variation as representative of a region's typical or expected food webs. Those data in turn can serve as starting points for further examination of the effects of finer-scale variation, such as differences between exploited, protected, and pristine areas. In effect, food webs at finer geographic or temporal scales will be subsets of our food webs, with the latter greatly facilitating construction of finer-scale webs.

The compositions of coral reef and associated seagrass communities of the three regions were assessed by extensive review of the available literature and databases [19–124]. Taxa present elsewhere in the Caribbean Sea or Gulf of Mexico, but not recorded explicitly from one of our Greater Antillean locations, were omitted from the dataset. We included only species for which reasonable detail of prey and predator species as well as foraging habitat (see the following) could be obtained; anecdotal records of trophic interactions were not used. This resulted in the underrepresentation of several potentially important taxonomic groups, for example, Asteroidea and Cirripedia. Nevertheless, the dataset includes primary producer species representative of coccolithophores, diatoms (including epiphytic species), filamentous algae, coarsely branched algae, jointed calcareous

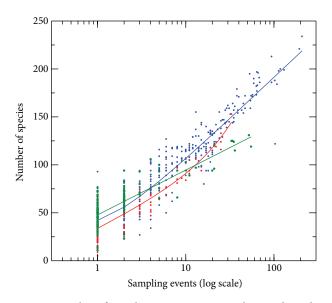


FIGURE 2: Number of sampling events per regional site and number of species discovered, as recorded in the REEF database [125]. Blue: Cayman Islands; green: Cuba; red: Jamaica. Lines are fractional polynomial regressions fit to each dataset. Functions are as follows: Cayman Islands: $24.6x^{-1} + 37.7 \ln x + 17.4$; Cuba: $20.3 \ln x + 47.8$; Jamaica: $1.3x + 19.8 \ln x + 32.5$.

algae, thick leathery algae, coralline encrusting algae, and seagrasses. Planktonic bacteria are represented as a single entry, or node, in the networks. Consumers are represented by dinoflagellates, planktonic and benthic foraminifera, tintinnids, epibenthic sponges, endolithic sponges, ahermatypic and hermatypic corals, copepods, amphipods, appendicularians, siphonophores, isopods, chaetognaths, scyphozoans, gorgonians, corallimorphs, zoanthids, echinoids, bivalves, gastropods (including pteropods and nudibranchs), polychaetes, holothurians, malacostracan crustaceans, cartilaginous and bony fish, and sea turtles.

Nonvertebrate taxa were treated as occurring uniformly among all three locations because of a general lack of biogeographic specificity at this scale available in published data. More precise data are available for vertebrate species, however, and we took advantage of this to differentiate the three food webs. Vertebrate occurrences were obtained from REEF (http://www.reef.org/), GBIF (http://www.gbif.org/), and Fishbase (http://www.fishbase.org/) databases and utilized to conservatively minimize apparent differences among the locations. We searched the REEF Volunteer Survey database [125] for records of every vertebrate species in our literature dataset. Those species that are not recorded in the survey, but are present in our dataset, were automatically considered to be present. All REEF sites in each region were queried individually, but the results were combined to construct the regional species lists. We included both Expert and Novice observations, again to be conservative in our estimate of the compositional differences among the regions. Our queries covered the period from 1999 to 2010. We considered the presence of a species in at least one site within a region to count for the entire region. This allows for the

possibilities that species were overlooked during surveys and that individuals at one site could relocate to or reproductively seed another site. Furthermore, we ignored demographic data such as estimates of population densities, which are extremely low for some species, and recorded a species as present even if only a single individual was observed, because the REEF data bear a significant sampling effect signal. The impact of sampling effort and number of reports was assessed by comparing the number of reports per site and the number of species recorded for each location. Results indicate clearly that Cuba and Jamaica have been sampled far fewer times than have the Cayman Islands and that lower species richnesses of the former locations in the REEF databases could very well be reversed if sampling was intensified (Figure 2). Moreover, given the much greater area of Cuba's coastal region, important refuges and relatively pristine areas such as Los Jardines de Reina are not captured effectively in the REEF database.

We therefore augmented the data for each location with occurrences recorded in the GBIF and Fishbase databases. While the REEF data were constrained to reports from 1999 to 2010, all GBIF and Fishbase occurrences in the Twentieth century were incorporated into our dataset. This results in a very conservative estimate of present species composition, since the historic databases include species from each location that are not only absent in the REEF surveys but may in fact now be extirpated from a locality. Finally, maximum body length was recorded for all fish species as listed in Fishbase.

Food web assembly was as follows. We included species both restricted to coral reefs and those commonly found in adjacent seagrass beds. All eukaryotic clades resolvable to the species level were considered for inclusion in the food webs, including protistan producers and consumers, but we included only species for which diet could be specified. Most primary producer and invertebrate species were then organized into trophic guilds [11] unless consumer diets are highly specialized and known with specificity; such is the case for cephalopod and stomatopod species. Each guild comprises data on taxonomic composition, trophic ecologies (species prey and predators), and primary habitat. Given uncertainty in the full array of trophic interactions for most of these species, a trophic guild is an aggregation of species where members of a guild potentially share prey drawn from the same guild(s), likewise for predators, and share the same habitat. One hundred and sixty-nine primary producer species were subsequently aggregated into nine guilds while 513 protistan and invertebrate consumers were aggregated into 99 guilds (Figure 3).

Cartilaginous and bony fish diets were obtained from the detailed reports of Randall [39] and Opitz [126]. The Opitz ECOPATH data proved to be particularly valuable, being based on extensive gut analyses, though primarily from the US Virgin Islands. Many fish dietary items, however, are present in very small proportions of total diet diversity and may therefore represent only occasional feeding rather than major dietary components or are possibly ingested incidentally. We thus included only those prey items which comprised 1% or more of a species diet. All vertebrate species were then categorized into three foraging groups: species

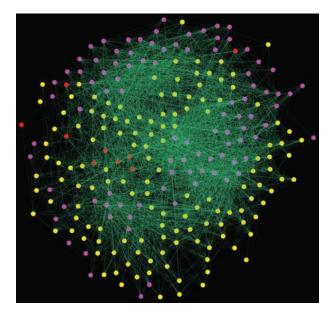


FIGURE 3: Food web illustration of trophic guilds and trophic species, and trophic interactions between them. This web represents the Cuban dataset. All nodes and interactions in the data are illustrated. There are 266 nodes and 3899 interactions (edges) in the network. The obvious complexity of the network emphasizes the improbability of any easy simplification of the data. Node colours are as follows: red: primary producers; magenta: protistan and invertebrate guilds; yellow: vertebrate species and trophic species.

that forage primarily on coral reefs (reef foragers), those that forage primarily in seagrass beds (seagrass foragers), and those that forage frequently in both habitats (reef-seagrass foragers) [126]. Our documentation, however, does not preclude the presence of these species in other coastal habitats not covered in our study, for example, mangrove environments. Vertebrate species were subsequently aggregated if they shared exactly the same prey and predators, thereby representing trophic species [127]. Trophic species are conceptually a subset of trophic guilds [14], and hence the latter term only is used from here on. Ultimately, 184 vertebrate species were aggregated into 162 trophic species (Figure 3).

3. Dataset Description

The dataset associated with this Dataset Paper consists of 7 items which are described as follows.

Dataset Item 1 (Table). Trophic data for the Cayman Islands. Data specify trophic guilds, the number of prey per guild, a list of those prey guilds, and the major foraging habitat of the guild. In the first column is given the Guild Number. Guild numbers are standard among all the datasets; for example, guild number 1 represents planktonic bacteria in the Cayman, Cuban, and Jamaican datasets. In the second column (Guild Description) is given the common language guild descriptions. The descriptions indicate the major trophic niche of the guild in the case of multispecies guilds (e.g.,

Macroplanktonic carnivores I), the major taxon of the guild (e.g., *Eucidaris*), or in the case of vertebrates, a common name of the species (e.g., scorpionfish). In the third column (Foraging Habitat) is given the major foraging habitat of vertebrate guild members. In this column, r means reef; rg, reef and seagrass beds; and g, seagrass beds. In the fourth column (Number of Prey) is given the total number of guilds that contain species that are preyed upon by the guild in question. In the fifth column (Prey) is given a list of prey guilds. Missing data are indicated by a period (.).

Column 1: Guild Number
Column 2: Guild Description

Column 3: Foraging Habitat

Column 4: Number of Prey

Column 5: Prey

Dataset Item 2 (Table). Trophic data for Cuba. Data specify trophic guilds, the number of prey per guild, a list of those prey guilds, and the major foraging habitat of the guild. In the first column is given the Guild Number. Guild numbers are standard among all the datasets; for example, guild number 1 represents planktonic bacteria in the Cayman, Cuban, and Jamaican datasets. In the second column (Guild Description) is given the common language guild descriptions. The descriptions indicate the major trophic niche of the guild in the case of multispecies guilds (e.g., Macroplanktonic carnivores I), the major taxon of the guild (e.g., Eucidaris), or in the case of vertebrates, a common name of the species (e.g., scorpionfish). In the third column (Foraging Habitat) is given the major foraging habitat of vertebrate guild members. In this column, r means reef; rg, reef and seagrass beds; and g, seagrass beds. In the fourth column (Number of Prey) is given the total number of guilds that contain species that are preyed upon by the guild in question. In the fifth column (Prey) is given a list of prey guilds. Missing data are indicated by a period (.).

Column 1: Guild Number

Column 2: Guild Description

Column 3: Foraging Habitat

Column 4: Number of Prey

Column 5: Prey

Dataset Item 3 (Table). Trophic data for Jamaica. Data specify trophic guilds, the number of prey per guild, a list of those prey guilds, and the major foraging habitat of the guild. In the first column is given the Guild Number. Guild numbers are standard among all the datasets; for example, guild number 1 represents planktonic bacteria in the Cayman, Cuban, and Jamaican datasets. In the second column (Guild Description) is given the common language guild descriptions. The descriptions indicate the major trophic niche of the guild in the case of multispecies guilds (e.g., Macroplanktonic carnivores I), the major taxon of the guild (e.g., Eucidaris),

or in the case of vertebrates, a common name of the species (e.g., scorpionfish). In the third column (Foraging Habitat) is given the major foraging habitat of vertebrate guild members. In this column, r means reef; rg, reef and seagrass beds; and g, seagrass beds. In the fourth column (Number of Prey) is given the total number of guilds that contain species that are preyed upon by the guild in question. In the fifth column (Prey) is given a list of prey guilds. Missing data are indicated by a period (.).

Column 1: Guild Number Column 2: Guild Description Column 3: Foraging Habitat Column 4: Number of Prey Column 5: Prey

Dataset Item 4 (Table). Guild key. This is a list of all guilds present in Dataset Items 1-3 (Tables) and the taxa assigned to those guilds. There is a total of 265 guilds, though none of the regions described contains all those guilds. There are also several guilds or species that are absent from all the regions, but have nevertheless been recorded in the northern Caribbean region, and are likely to be present in undescribed refuges or occasional members of the regions, for example, the tiger shark *Galeocerdo cuvieri*. In the first column is given the Guild Number; in the second (Taxa), the taxa assigned to guild; in the third (Fish Body Length), the average body size (fork length, cm) of bony and cartilaginous fish species. Maximum body size is recorded if average size is not available. Missing data are indicated by a period (.). In columns 4-6 is shown the presence of vertebrate species in the Cayman Islands, Cuba, or Jamaica indicated by "x" sign.

Column 1: Guild Number

Column 2: Taxa

Column 3: Fish Body Length (cm)

Column 4: Cayman Islands

Column 5: Cuba
Column 6: Jamaica

Dataset Item 5 (Binary Matrix). Binary adjacency matrix of the Cayman Islands food web. Rows are predatory guilds and columns are prey. The *ij*th element of the matrix is 1 if guild *i* preys upon species in guild *j*, and 0 otherwise. Note that the matrices are therefore asymmetric about the diagonal, and that there are 265 rows representing each guild in the dataset. Taxa that are missing from the food web, for example, the tiger shark Galeocerdo cuvieri, are included as disconnected nodes, that is, rows and columns comprising zero elements only. This is for consistency with future datasets of related regions in which the species might be present.

Dataset Item 6 (Binary Matrix). Binary adjacency matrix of the Cuban food web. Rows are predatory guilds and columns are prey. The ijth element of the matrix is 1 if guild i preys

upon species in guild *j*, and 0 otherwise. Note that the matrices are therefore asymmetric about the diagonal and that there are 265 rows representing each guild in the dataset. Taxa that are missing from the food web, for example, the tiger shark *Galeocerdo cuvieri*, are included as disconnected nodes, that is, rows and columns comprising zero elements only. This is for consistency with future datasets of related regions in which the species might be present.

Dataset Item 7 (Binary Matrix). Binary adjacency matrix of the Jamaican food web. Rows are predatory guilds and columns are prey. The *ij*th element of the matrix is 1 if guild *i* preys upon species in guild *j*, and 0 otherwise. Note that the matrices are therefore asymmetric about the diagonal and that there are 265 rows representing each guild in the dataset. Taxa that are missing from the food web, for example, the tiger shark *Galeocerdo cuvieri*, are included as disconnected nodes, that is, rows and columns comprising zero elements only. This is for consistency with future datasets of related regions in which the species might be present.

4. Concluding Remarks

The datasets presented here are syntheses of decades of work by multiple ecologists and systematists; yet they are far from being taxonomically comprehensive. There is a pressing need for additional data of the ecological roles of organisms in important communities such as coral reefs. Nevertheless, the current data should prove to be helpful in both understanding the long-term and large-scale dynamics of these communities and in formulating strategies for the protection of the communities and species within. Ecosystem dynamics play out over multiple scales of time and space [128] and multiple levels of biological organization, from organisms to ecosystems themselves [129]. Organizing data relevant to those scales, such as shifting historical baselines, multiple disturbances that act on different spatial scales, and variance of community composition across those scales, presents significant challenges to the formulation of coherent strategies for the conservation, restoration, and sustainable economic use of coral reef ecosystems in the Caribbean region and elsewhere [6, 130]. At the same time, it is understood that the nature of a species interactions, for example, number or strength, can be important determinants of its probability of extinction (or survival) under different circumstances. It is also understood that the configurations of those interactions for all species in a community are important determinants of the communities resilience to change or robustness against disturbances. Complex network food webs present a basic scaffolding for tying together data on an ecosystem's species richness, biotic interactions, and functional diversity. Much of the work relating food web properties to ecosystem health has dealt so far with model or relatively small communities [131]. The datasets presented here make it possible to extend this work to an ecosystem that is both one of the ocean's richest and most complex and one of its most critically endangered [7].

Finally, these datasets are both starting points and works in progress. They may be used by workers to examine regionwide phenomena such as the trophic impacts of coral bleaching or invasive species, and likewise they can be subsampled to reflect within-region variation of community composition and anthropogenic factors. They are works in progress because they will be revised as data on those already included, as well as omitted species become available, and as the state of Caribbean coral reefs continues to evolve.

Dataset Availability

The dataset associated with this Dataset Paper is dedicated to the public domain using the CC0 waiver and is available at http://dx.doi.org/10.7167/2013/857470/dataset/.

Conflict of Interests

The authors declare that they have no competing interests or conflict of interests.

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References

- [1] J. B. C. Jackson, "Reefs since Columbus," *Coral Reefs*, vol. 16, no. 1, pp. S23–S32, 1997.
- [2] M. J. Hardt, "Lessons from the past: the collapse of Jamaican coral reefs," Fish and Fisheries, vol. 10, no. 2, pp. 143–158, 2009.
- [3] P. J. Mumby, C. P. Dahlgren, A. R. Harborne et al., "Fishing, trophic cascades, and the process of grazing on coral reefs," *Science*, vol. 311, no. 5757, pp. 98–101, 2006.
- [4] K. P. Sebens, "Biodiversity of coral reefs: what are we losing and why?" *American Zoologist*, vol. 34, no. 1, pp. 115–133, 1994.
- [5] T. A. Gardner, I. M. Côté, J. A. Gill, A. Grant, and A. R. Watkinson, "Long-term region-wide declines in Caribbean corals," *Science*, vol. 301, no. 5635, pp. 958–960, 2003.
- [6] C. Mora, "A clear human footprint in the coral reefs of the Caribbean," *Proceedings of the Royal Society B*, vol. 275, no. 1636, pp. 767–773, 2008.
- [7] L. Burke, K. Reytar, M. Spalding, and A. Perry, Reefs at Risk Revisited, World Resources Institute, Washington, DC, USA, 2011.
- [8] J. M. Montoya, S. L. Pimm, and R. V. Solé, "Ecological networks and their fragility," *Nature*, vol. 442, no. 7100, pp. 259–264, 2006.
- [9] J. A. Dunne, R. J. Williams, and N. D. Martinez, "Network structure and biodiversity loss in food webs: robustness increases with connectance," *Ecology Letters*, vol. 5, no. 4, pp. 558–567, 2002.
- [10] S. Allesina and A. Bodini, "Who dominates whom in the ecosystem? Energy flow bottlenecks and cascading extinctions," *Journal of Theoretical Biology*, vol. 230, no. 3, pp. 351–358, 2004.

- [11] P. D. Roopnarine, "Extinction cascades and catastrophe in ancient food webs," *Paleobiology*, vol. 32, no. 1, pp. 1–19, 2006.
- [12] P. D. Roopnarine, "Ecological modeling of paleocommunity food webs," in *Conservation Paleobiology*, G. Dietl and K. Flessa, Eds., vol. 15 of *Paleontological Society Papers*, pp. 195–220, The Paleontological Society, 2009.
- [13] J. A. Dunne and R. J. Williams, "Cascading extinctions and community collapse in model food webs," *Philosophical Transactions of the Royal Society B*, vol. 364, no. 1524, pp. 1711–1723, 2009.
- [14] P. D. Roopnarine, "Networks, extinction and paleocommunity food webs," in *Quantitative Methods in Paleobiology*, J. Alroy and G. Hunt, Eds., vol. 16 of *Paleontological Society Papers*, The Paleontological Society, 2010.
- [15] J. Bascompte, C. J. Melián, and E. Sala, "Interaction strength combinations and the overfishing of a Marine food web," *Proceedings of the National Academy of Sciences of the United States of America*, vol. 102, no. 15, pp. 5443–5447, 2005.
- [16] M. J. H. Newman, G. A. Paredes, E. Sala, and J. B. C. Jackson, "Structure of Caribbean coral reef communities across a large gradient of fish biomass," *Ecology Letters*, vol. 9, no. 11, pp. 1216–1227, 2006.
- [17] N. A. J. Graham, T. R. McClanahan, M. A. MacNeil et al., "Climate warming, Marine protected areas and the ocean-scale integrity of coral reef ecosystems," *PLoS ONE*, vol. 3, no. 8, Article ID e3039, 2008.
- [18] E. R. Selig and J. F. Bruno, "A global analysis of the effectiveness of Marine protected areas in preventing coral loss," *PLoS ONE*, vol. 5, no. 2, Article ID e9278, 2010.
- [19] H. B. Owre and M. Foyo, "Studies on Caribbean zooplankton. Description of the program and results of the first cruise," *Bulletin of Marine Science*, vol. 22, pp. 483–521, 1972.
- [20] A. Acosta, M. Casas, C. A. Vargas, and J. Camacho, "Lista de zoantharia (Cnidaria: Anthozoa) del caribe y Colombia," *Biota Colombiana*, vol. 6, pp. 147–162, 2005.
- [21] S. Zea and E. Weil, "Taxonomy of the Caribbean excavating sponge species complex *Cliona caribbaea—C. aprica—C. langae* (Porifera, Hadromerida, Clionaidae)," *Caribbean Journal of Science*, vol. 39, no. 3, pp. 348–370, 2003.
- [22] J. L. Wulff, "Rapid diversity and abundance decline in a Caribbean coral reef sponge community," *Biological Conservation*, vol. 127, no. 2, pp. 167–176, 2006.
- [23] K. P. Sebens, S. P. Grace, B. Helmuth, E. J. Maney Jr., and J. S. Miles, "Water flow and prey capture by three scleractinian corals, *Madracis mirabilis*, Montastrea *cavernoss* and *Porites porites* in a field enclosure," *Marine Biology*, vol. 131, no. 2, pp. 347–360, 1998.
- [24] T. E. Bowman, "Pelagic amphipods of the genus Hyperia and closely related genera (Hyperiidea: Hyperiidae)," Smithsonian Contributions to Zoology, vol. 136, 76 pages, 1973.
- [25] M. A. Buzas, R. K. Smith, and K. A. Beem, "Ecology and systematics of Foraminifera in two *Thalassia* habitats, Jamaica, West Indies," *Smithsonian Contributions to Paleobiology*, vol. 31, 139 pages, 1977.
- [26] C. S. Rogers, T. H. Suchanek, and F. A. Pecora, "Effects of Hurricanes David and Frederic (1979) on shallow Acropora palmata reef communities: st. Croix, U.S. Virgin Islands," Bulletin of Marine Science, vol. 32, no. 2, pp. 532–548, 1982.
- [27] C. Birkeland and S. Neudecker, "Foraging behavior of two Caribbean chaetodontids: *Chaetodon capistratus* and *C. aculeatus*," *Copeia*, no. 1, pp. 169–178, 1981.

[28] K. P. Sebens, "Intertidal distribution of zoanthids on the Caribbean coast of Panama: effects of predation and desiccation," *Bulletin of Marine Science*, vol. 32, pp. 316–335, 1982.

- [29] T. F. Goreau and J. W. Wells, "The shallow-water scleractinia of Jamaica: revised list of species and their vertical distribution range," *Bulletin of Marine Science*, vol. 17, pp. 442–453, 1967.
- [30] T. S. Park, "Calanoid copepods from the Caribbean Sea and Gulf of Mexico. 2. New species and new records from pankton samples," *Bulletin of Marine Science*, vol. 20, pp. 472–546, 1970.
- [31] N. Knowlton and B. D. Keller, "A new, sibling species of snapping shrimp associated with the Caribbean sea anemone *Bartholomea annulata* (Alpheus armatus/immaculatus)," *Bulletin of Marine Science*, vol. 33, no. 2, pp. 353–362, 1983.
- [32] H. M. Reiswig, "Population dynamics of three Jamaican Demospongiae," Bulletin of Marine Science, vol. 23, pp. 191–226, 1973.
- [33] D. L. Meyer, "Distribution and living habits of comatulid crinoids near discovery bay, Jamaica," *Bulletin of Marine Science*, vol. 23, pp. 245–259, 1973.
- [34] E. M. Preston and J. L. Preston, "Ecological structure in a West Indian gorgonian fauna," *Bulletin of Marine Science*, vol. 25, pp. 248–258, 1975.
- [35] J. B. C. Jackson, "The ecology of molluscs of *Thalassia* communities, Jamaica, West Indies. I. Distribution, environmental physiology, and ecology of common shallow-water species," *Bulletin of Marine Science*, vol. 23, pp. 313–350, 1973.
- [36] D. A. West, "Symbiotic zoanthids (Anthozoa: Cnidaria) of Puerto Rico," Bulletin of Marine Science, no. 29, pp. 253–271, 1979
- [37] P. S. Lobel, "Herbivory by damselfishes and their role in coral reef community ecology," *Bulletin of Marine Science*, vol. 30, pp. 273–289, 1979.
- [38] B. A. Hazlett, "Biotic aspects of the distribution of the crabs *Panopeus herbstii* and Mithrax sculptus," *Bulletin of Marine Science*, vol. 29, pp. 576–580, 1979.
- [39] J. E. Randall, "Food Habits of Reef Fishes of the West Indies," Studies in Tropical Oceanography, vol. 5, pp. 665–847, 1967.
- [40] C. P. Santos, A. B. Coutinho, and E. Hajdu, "Spongivory by Eucidaris tribuloides from Salvador, Bahia (Echinodermata: Echinoidea)," Journal of the Marine Biological Association of the United Kingdom, vol. 82, no. 2, pp. 295–297, 2002.
- [41] D. T. Dy, F. T. Uy, and C. A. Coralles, "Feeding, respiration, and excretion by the tropical sea urchin *Tripneustes gratilla* (Echinodermata: Echinoidea) from the Philippines," *Journal of the Marine Biological Association of the United Kingdom*, vol. 82, pp. 299–302, 2002.
- [42] R. H. Gore, L. E. Scotto, and L. J. Becker, "Community composition: stability, and trophic partitioning in decapod crustaceans inhabiting some subtropical sabellariid worm reefs," *Bulletin of Marine Science*, vol. 28, pp. 221–248, 1978.
- [43] W. M. Goldberg, "The ecology of the coral-octocoral communities off the southeast Florida coast: geomorphology, species composition, and zonation," *Bulletin of Marine Science*, vol. 23, pp. 465–488, 1973.
- [44] R. Wolcott and C. Messing, "A comparison of diets and water agitation methods for larval culture of the edible sea urchin," *Bulletin of Marine Science*, vol. 77, pp. 177–190, 2005.
- [45] G. L. Voss and N. Voss, "An ecological survey of soldier key, biscayne bay, Florida," *Bulletin of Marine Science of the Gulf and Caribbean*, vol. 5, pp. 203–229, 1955.
- [46] G. S. Kleppel, "On the diets of calanoid copepods," *Marine Ecology Progress Series*, vol. 99, no. 1-2, pp. 183–195, 1993.

[47] J. Cobb and J. M. Lawrence, "Diets and coexistence of the sea urchins *Lytechinus variegatus* and *Arbacia punctulata* (Echinodermata) along the central Florida gulf coast," *Marine Ecology Progress Series*, vol. 295, pp. 171–182, 2005.

- [48] R. B. Manning and D. K. Camp, "A new genus of stomatopod from the caribbean sea (Stomatopoda: Squillidae)," *Journal of Crustacean Biology*, vol. 21, no. 1, pp. 202–204, 2001.
- [49] P. E. Tschudin, "Shell morphology, shell texture and species discrimination of Caribbean Tucetona (Bivalvia, Glycymeridae)," *Journal of Paleontology*, vol. 75, no. 3, pp. 658–679, 2001.
- [50] Y. Nakamura and J. T. Turner, "Predation and respiration by the small cyclopoid copepod *Oithona similis*: how important is feeding on ciliates and heterotrophic flagellates?" *Journal of Plankton Research*, vol. 19, no. 9, pp. 1275–1288, 1997.
- [51] F. L. M. Mantelatto and M. Petracco, "Natural diet of the crab Hepatus pudibundus (Brachyura: Calappidae) in Fortaleza bay, Ubatuba (SP), Brazil," Journal of Crustacean Biology, vol. 17, no. 3, pp. 440–446, 1997.
- [52] F. Sander and E. Moore, "A comparative study of inshore and offshore copepod populations at barbados, West Indes," *Crustaceana*, vol. 35, pp. 225–240, 1978.
- [53] M. S. Hill, "Spongivory on Caribbean reefs releases corals from competition with sponges," *Oecologia*, vol. 117, no. 1-2, pp. 143–150, 1998.
- [54] C. R. Wilkinson and A. C. Cheshire, "Growth rate of Jamaican coral reef sponges after hurricane allen," *Biological Bulletin*, vol. 175, pp. 175–179, 1998.
- [55] B. Kensley and R. H. Gore, "Coralaxius Abelei, new genus and new species (Crustacea: Decapoda: Thalassinidea: Axiidae): a coral-inhabiting shrimp from the Florida keys and the Western Caribbean sea," Proceedings of the Biological Society of Washington, vol. 93, pp. 1277–1294, 1980.
- [56] C. A. Góes and J. E. Lins-Oliveira, "Natural diet of the spiny lobster, *Panulirus echinatus* Smith, 1869 (Crustacea: Decapoda: Palinuridae), from São Pedro and São Paulo Archipelago, Brazil," *Brazilian Journal of Biology*, vol. 69, no. 1, pp. 143–148, 2009.
- [57] T. F. Goreau, "The ecology of Jamaican Coral Reefs I. Species composition and zonation," *Ecology*, vol. 40, pp. 67–90, 1959.
- [58] J. W. Porter, "Patterns of species diversity in Caribbean Reef Corals," *Ecology*, vol. 53, pp. 745–748, 1972.
- [59] I. Goodbody, "Continuous breeding in populations of two tropical crustaceans, *Mysidium columbiae* (Zimmer) and *Emerita portoricensis* schmidt," *Ecology*, vol. 46, pp. 195–197, 1965.
- [60] J. E. Randall, "Grazing effect on sea grasses by herbivorous reef fishes in the West Indies," *Ecology*, vol. 46, pp. 255–260, 1965.
- [61] M. M. Littler, D. S. Littler, and P. R. Taylor, "Selective herbivore increases biomass of its prey: a Chiton-Coralline reef-building association," *Ecology*, vol. 76, no. 5, pp. 1666–1681, 1995.
- [62] D. R. Robertson, "Cohabitation of competing territorial damselfishes on a Caribbean coral reef," *Ecology*, vol. 65, no. 4, pp. 1121–1135, 1984.
- [63] M. E. Hay, "Patterns of fish and urchin grazing on Caribbean coral reefs: are previous results typical?" *Ecology*, vol. 65, no. 2, pp. 446–454, 1984.
- [64] J. C. Martínez Iglesias and J. E. García Raso, "The crustacean decapod communities of three coral reefs from the southwestern Caribbean Sea of Cuba: species composition, abundance and structure of the communities," *Bulletin of Marine Science*, vol. 65, no. 2, pp. 539–557, 1999.

- [65] H. Nomura, K. Aihara, and T. Ishimaru, "Feeding of the chaetognath Sagitta crassa Tokioka in heavily eutrophicated Tokyo bay, Japan," *Plankton and Benthos Research*, vol. 2, no. 3, pp. 120–127, 2007.
- [66] R. Gasca and C. T. Shih, "Hyperiid amphipods from surface waters of the western Caribbean Sea (1991)," *Crustaceana*, vol. 74, no. 5, pp. 489–499, 2001.
- [67] J. L. Rueda and C. Salas, "Trophic dependence of the emerald neritid Smaragdia viridis (Linnaeus, 1758) on two seagrasses from European coasts," *Journal of Molluscan Studies*, vol. 73, no. 2, pp. 211–214, 2007.
- [68] E. Moore and F. Sander, "A comparative study of zooplankton from oceanic, shelf, and harbor waters of Jamaica," *Biotropica*, vol. 11, pp. 196–206, 1979.
- [69] D. F. Webber, M. K. Webber, and J. C. Roff, "Effects of flood waters on the planktonic community of the Hellshire coast, southeast Jamaica," *Biotropica*, vol. 24, no. 3, pp. 362–374, 1992.
- [70] J. W. Porter, "Autotrophy, heterotrophy, and resource partitioning in Caribbean reef-building corals," *The American Naturalist*, vol. 110, pp. 731–742, 1979.
- [71] R. B. Aronson and W. F. Precht, "Herbivory and algal dynamics on the coral reef at discovery bay, Jamaica," *Limnology and Oceanography*, vol. 45, no. 1, pp. 251–255, 2000.
- [72] A. R. Emery, "Preliminary observations on coral reef plankton," Limnology and Oceanography, vol. 13, pp. 293–303, 1968.
- [73] M. Ribes, R. Coma, and J. M. Gili, "Heterotrophic feeding by gorgonian corals with symbiotic zooxanthella," *Limnology and Oceanography*, vol. 43, no. 6, pp. 1170–1179, 1998.
- [74] S. M. Lewis, "The role of herbivorous fishes in the organization of a Caribbean coral reef community," *Ecological Monographs*, vol. 56, pp. 184–200, 1986.
- [75] B. F. McPherson, "Feeding and Oxygen Uptake of the Tropical Sea Urchin *Eucidaris tribuloides* (Lamark)," *Biological Bulletin*, vol. 135, pp. 308–321, 1968.
- [76] P. Francour, "Predation on holothurians: a literature review," *Invertebrate Biology*, vol. 116, no. 1, pp. 52–60, 1997.
- [77] K. E. Holmes, "Effects of eutrophication on bioeroding sponge communities with the description of new West Indian sponges, *Cliona spp.* (Porifera: Hadromerida: Clionidae)," *Invertebrate Biology*, vol. 119, no. 2, pp. 125–138, 2000.
- [78] R. E. Martin and W. D. Liddell, "Foraminiferal biofacies on a North Coast fringing reef (1–75m), discovery bay, Jamaica," *Palaios*, vol. 3, no. 3, pp. 298–314, 1988.
- [79] T. F. Goreau, "Mass expulsion of zooxanthellae from Jamaican reef communities after hurricane flora," *Science*, vol. 145, no. 3630, pp. 383–386, 1964.
- [80] T. J. Goreau, "Bleaching and reef community change in Jamaica: 1951–1991," American Zoologist, vol. 32, no. 6, pp. 683–695, 1992.
- [81] S. Rowley, "A critical evaluation of the symbiotic association between tropical tube-dwelling Polychaetes and their Hermatypic coral hosts, with a focus on *Spirobranchus giganteus* (Pallas, 1766)," *The Plymouth Student Scientist*, vol. 1, pp. 335–353, 2008.
- [82] T. Ishimaru, S. Nishida, and R. Marumo, "Food size selectivity of zooplankton evaluated from the occurence of coccolithophorids in the guts," *Bulletin of the Plankton Society of Japan*, vol. 35, pp. 101–114, 1988.
- [83] J. B. C. Jackson, "Overgrowth competition between encrusting cheilostome ectoprocts in a Jamaican crytic reef environment," *Journal of Animal Ecology*, vol. 48, no. 3, pp. 805–823, 1979.

- [84] B. J. Godley, S. M. Smith, P. F. Clark, and J. D. Taylor, "Molluscan and crustacean items in the diet of the loggerhead turtle, *Caretta caretta* (Linnaeus, 1758) [Testudines: Chelonidae] in the eastern Mediterranean," *Journal of Molluscan Studies*, vol. 63, no. 3, pp. 474–476, 1997.
- [85] H. M. Reiswig, "Particle feeding in natural populations of three Marine demosponges," *Biological Bulletin*, vol. 141, pp. 568–591, 1971.
- [86] P. Knight-Jones and A. S. Y. Mackie, "A revision of *Sabellastarte* (Polychaeta: Sabellidae)," *Journal of Natural History*, vol. 37, no. 19, pp. 2269–2301, 2003.
- [87] J. D. Woodley, E. A. Chornesky, P. A. Clifford et al., "Hurricane Allen's impact on Jamaican coral reefs," *Science*, vol. 214, no. 4522, pp. 749–755, 1981.
- [88] C. M. Wahle, "Regeneration of injuries among Jamaican gorgonians: the roles of colony physiology and environment," *Biological Bulletin*, vol. 165, pp. 778–790, 1983.
- [89] T. T. Noji, U. V. Bathmann, B. Von Bodungen et al., "Clearance of picoplankton-sized particles and formation of rapidly sinking aggregates by the pteropod, *Limacina retroversa*," *Journal of Plankton Research*, vol. 19, no. 7, pp. 863–875, 1997.
- [90] L. Q. Yokoyama and A. C. Z. Amaral, "The diet of Ophionereis reticulata (Echinodermata: Ophiuroidea) in southeastern Brazil," Revista Brasileira de Zoologia, vol. 25, no. 3, pp. 576–578, 2008.
- [91] E. Broglio, S. H. Jónasdóttir, A. Calbet, H. H. Jakobsen, and E. Saiz, "Effect of heterotrophic versus autotrophic food on feeding and reproduction of the calanoid copepod Acartia tonsa: relationship with prey fatty acid composition," *Aquatic Microbial Ecology*, vol. 31, no. 3, pp. 267–278, 2003.
- [92] L. G. Abele and W. Kim, "An illustrated guide to the Marine decapod crustaceans of Florida," Tech. Rep., State of Florida Department of Environmental Regulation, 1986.
- [93] M. Dunlap and J. R. Pawlik, "Video-monitored predation by Caribbean reef fishes on an array of mangrove and reef sponges," *Marine Biology*, vol. 126, no. 1, pp. 117–123, 1996.
- [94] E. I. Gilbert, Juvenile Green Turtle (Chelonia mydas) foraging ecology: feeding selectivity and forage nutrient analysis [M.S. thesis], University of Central Florida, 1998.
- [95] M. A. Faust, "Dinoflagellate associations in a coral reefmangrove ecosystem: pelican and associated Cays, Belize," *Atoll Research Bulletin*, no. 466–480, pp. 135–149, 2000.
- [96] C. T. Perry, "Macroborers within coral framework at discovery bay, north Jamaica: species distribution and abundance, and effects on coral preservation," *Coral Reefs*, vol. 17, no. 3, pp. 277–287, 1998.
- [97] T. R. McClanahan, "Predation and the control of the sea urchin *Echinometra viridis* and fleshy algae in the patch reefs of Glovers Reef, Belize," *Ecosystems*, vol. 2, no. 6, pp. 511–523, 1999.
- [98] R. J. Livingston, "The relationship of physical factors and biological response in coastal seagrass meadows," *Estuaries*, vol. 7, no. 4, pp. 377–390, 1984.
- [99] M. A. E. Malaquias, E. Berecibar, and D. G. Reid, "Reassessment of the trophic position of Bullidae (Gastropoda: Cephalaspidea) and the importance of diet in the evolution of cephalaspidean gastropods," *Journal of Zoology*, vol. 277, no. 1, pp. 88–97, 2009.
- [100] P. W. Glynn, "Ecology of a Caribbean coral reef. The Porites reef-flat biotope: part II. Plankton community with evidence for depletion," *Marine Biology*, vol. 22, no. 1, pp. 1–21, 1973.

- [101] M. G. Frick, "Lepidochelys kempi (Kemp's Ridley) Caretta caretta (Loggerhead), and Malaclemys terrapin centrata (Carolina Diamondback Terrapin) diet and predation," Herpetological Review, vol. 28, 149 pages, 1997.
- [102] M. G. Frick, "Dermochelys coriacea (Leatherback Sea Turtle) Lepidochelys kempi (Kemp's Ridley Sea Turtle, and Caretta caretta (Loggerhead Sea Turtle) pelagic feeding," Herpetological Review, vol. 30, 165 pages, 1999.
- [103] A. G. Payne, C. Smith, and A. C. Campbell, "Interactions between ophiuroids and beaugregory damselfish," *Journal of the Marine Biological Association of the United Kingdom*, vol. 83, no. 3, pp. 625–632, 2003.
- [104] Y. M. León and K. A. Bjorndal, "Selective feeding in the hawksbill turtle, an important predator in coral reef ecosystems," *Marine Ecology Progress Series*, vol. 245, pp. 249–258, 2002.
- [105] D. Lirman and P. Biber, "Seasonal dynamics of macroalgal communities of the Northern Florida Reef Tract," *Botanica Marina*, vol. 43, no. 4, pp. 305–314, 2000.
- [106] M. M. Littler, P. R. Taylor, and D. S. Littler, "Algal resistance to herbivory on a Caribbean barrier reef," *Coral Reefs*, vol. 2, no. 2, pp. 111–118, 1983.
- [107] H. R. Lasker, "A comparison of the particulate feeding abilities of three species of gorgonian soft coral," *Marine Ecology Progress Species*, vol. 5, pp. 61–67, 1981.
- [108] L. S. Hammond, "Analysis of grain-size selection by deposit-feeding holothurians and echinoids (Echinodermata) from a shallow reef lagoon, discovery bay, Jamaica," *Marine Ecology Progress Series*, vol. 8, pp. 25–36, 1982.
- [109] L. S. Hammond, "Nutrition of deposit-feeding holothuroids and echinoids from a shallow reef lagoon, discovery bay, Jamaica," *Marine Ecology Progress Series*, vol. 10, pp. 297–305, 1983.
- [110] K. W. Rylaarsdam, "Life histories and abundance patterns of colonial corals on Jamaican reefs," *Marine Ecology Progress Series*, vol. 13, pp. 249–260, 1983.
- [111] H. R. Lasker, "Prey preferences and browsing pressure of the butterflyfish Chaetodon capistratus on Caribbean gorgonians," *Marine Ecology Progress Series*, vol. 21, pp. 213–220, 1985.
- [112] D. J. Gerhart, "Gregariousness in the gorgonian-eating gastro-pod *Cyphoma gibbosum*: tests of several possible causes," *Marine Ecology Progress Series*, vol. 31, pp. 255–263, 1986.
- [113] J. L. Ruesink and C. Drew Harvell, "Specialist predation on the Caribbean gorgonian *Plexaurella* spp. by *Cyphoma signatum* (Gastropoda)," *Marine Ecology Progress Series*, vol. 65, pp. 265–272, 1990.
- [114] J. B. Lewis, "Heterotrophy in corals: zooplankton predation by the hydrocoral Millepora complanata," *Marine Ecology Progress Series*, vol. 90, no. 3, pp. 251–256, 1992.
- [115] J. Cobb and J. M. Lawrence, "Diets and coexistence of the sea urchins Lytechinus variegatus and Arbacia punctulata (Echinodermata) along the central Florida gulf coast," Marine Ecology Progress Series, vol. 295, pp. 171–182, 2005.
- [116] M. W. Miller and C. L. Gerstner, "Reefs of an uninhabited Caribbean island: fishes, benthic habitat, and opportunities to discern reef fishery impact," *Biological Conservation*, vol. 106, no. 1, pp. 37–44, 2002.
- [117] J. A. Mortimer, "The feeding ecology of the West Caribbean green turtle (*Chelonia mydas*) in Nicaragua," *Biotropica*, vol. 13, pp. 49–58, 1981.
- [118] D. F. Amorocho and R. D. Reina, "Feeding ecology of the East Pacific green sea turtle *Chelonia mydas agassizii* at Gorgona National Park, Colombia," *Endangered Species Research*, vol. 3, pp. 43–51, 2007.

[119] M. E. Hay, T. Colburn, and D. Downing, "Spatial and temporal patterns in herbivory on a Caribbean fringing reef: the effects on plant distribution," *Oecologia*, vol. 58, no. 3, pp. 299–308, 1983.

- [120] D. K. Padilla, "Algal structural defenses: form and calcification in resistance to tropical limpets," *Ecology*, vol. 70, no. 4, pp. 835–842, 1989.
- [121] D. P. Báez and N. E. Ardila, "Poliquetos (Annelida: Polychaeta) del Mar Caribe colombiano," *Biota Colombiana*, vol. 4, pp. 89–109, 2003.
- [122] J. E. Randall and W. D. Hartman, "Sponge-feeding fishes of the West Indies," *Marine Biology*, vol. 1, no. 3, pp. 216–225, 1968.
- [123] C. J. Berg Jr., "Behavior and ecology of conch (Superfamily Strombacea) on a deep subtidal algal plain," *Bulletin of Marine Science*, vol. 25, pp. 307–317, 1975.
- [124] J. T. Turner, "The feeding ecology of some zooplankters that are important prey items of larval fish," Tech. Rep. 7, NOAA Technical Report NMFS, 1984.
- [125] Reef Environmental Education Foundation, "Reef environmental education foundation survey data," Tech. Rep., Reef Environmental Education Foundation, 2010, http://www.reef.org/.
- [126] S. Opitz, Trophic Interactions in Caribbean Coral Reefs, International Center for Living Aquatic Resources Management, Manila, Philippines, 1996.
- [127] F. Briand and J. E. Cohen, "Community food webs have scale-invariant structure," *Nature*, vol. 307, no. 5948, pp. 264–267, 1984.
- [128] J. B. C. Jackson, M. X. Kirby, W. H. Berger et al., "Historical overfishing and the recent collapse of coastal ecosystems," *Science*, vol. 293, no. 5530, pp. 629–638, 2001.
- [129] P. D. Roopnarine and K. D. Angielczyk, "The evolutionary palaeoecology of species and the tragedy of the commons," *Biology Letters*, vol. 8, no. 1, pp. 147–150, 2012.
- [130] P. J. Mumby and A. Hastings, "The impact of ecosystem connectivity on coral reef resilience," *Journal of Applied Ecology*, vol. 45, no. 3, pp. 854–862, 2008.
- [131] R. M. May, Stability and Complexity in Model Ecosystems, Princeton University Press, Princeton, NJ, USA, 2nd edition, 2001.

















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