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Author: Ainsworth, C. H.

Source: Marine and Coastal Fisheries: Dynamics, Management, and

Ecosystem Science, 3(1): 190-218

Published By: American Fisheries Society

URL: https://doi.org/10.1080/19425120.2010.549047

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Marine and Coastal Fisheries: Dynamics, Management, and Ecosystem Science 3:190-218, 2011

American Fisheries Society 2011 ISSN: 1942-5120 online

DOI: 10.1080/19425120.2010.549047

ARTICLE

Quantifying Species Abundance Trends in the Northern Gulf of California Using Local Ecological Knowledge

C. H. Ainsworth*1

National Oceanic and Atmospheric Administration, Northwest Fisheries Science Center, 2725 Montlake Boulevard East, Seattle, Washington 98112, USA

Abstract

Ecosystem-based fisheries management requires data on all parts of the ecosystem, and this can be a barrier in data-poor systems. Marine ecologists need a means of drawing together diverse information to reconstruct species abundance trends for a variety of purposes. This article uses a fuzzy logic approach to integrate information from multiple data sources and describe biomass trends for marine species groups in the northern Gulf of California, Mexico. Forty-two species groups were analyzed, comprising fish, invertebrates, birds, mammals, turtles, and algae. The most important new data series comes from recent interviews with fishers in the northern part of the gulf. Respondents were asked to classify the abundance of various targeted and untargeted marine species groups from 1950 to the present. The fuzzy logic method integrates their responses with catch-per-unit-effort series, intrinsic vulnerability to fishing determined from life history parameters, biomass predicted by a Schaefer harvest model, and other simple indices. The output of the fuzzy logic routine is a time series of abundance for each species group that can be compared with known trends. The results suggest a general decline in species abundance across fished and unfished taxa, with a few exceptions. Information gathered from interviews indicated that older fishers tended to recognize a greater relative decrease in species abundance since 1970 than did younger fishers, providing another example of Pauly's (1995) shifting cognitive baselines.

Resource managers face an expanding array of challenges in the northern Gulf of California, Mexico. The area has ecological significance as a biodiversity "hotspot" with a high degree of endemism in fish (Gilligan 1980; Enriquez-Andrade et al. 2005), invertebrates (Brusca 2006; Hendrickx 2007), and plants (Felger 2000), and it contains critical habitat for migratory and endangered species (Velarde and Anderson 1994; Jaramillo-Legorreta et al. 2007; Lercari and Chávez 2007). Unfortunately, marine conservation is often at odds with the fisheries that are so critical to the economic and food security of coastal communities (Guerroero-Ruíz et al. 2006; Lluch-Cota et al. 2007). Agriculture, aquaculture, ecotourism, and other marine-use sectors also continue to grow and compete with fisheries for space and resources.

Some have advocated ecosystem-based fisheries management (EBFM) as an integrated approach to managing human activities and a means of reconciling human needs with those of the natural system (Garcia et al. 2003; Pikitch et al. 2004). However, EBFM is broad by definition, and quantitative tools and analyses meant to support EBFM decisions can have large data requirements; this has proved to be a barrier to implementation of management decisions (Frid et al. 2006). Deficiency in scientific survey information is most evident in developing tropical and subtropical nations like Mexico, where species diversity is high and food web dynamics are complex, yet resources for stock assessment and monitoring are scarce (Silvestre and Pauly 1997). Here, there is a need for systematized information on species abundance (Lluch-Cota et al. 2007),

Subject editor: Kenneth Rose, Louisiana State University, Baton Rouge, Louisiana, USA

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^{*}Corresponding author: cameron.ainsworth@noaa.gov

¹Present address: Marine Resources Assessment Group Americas, 2725 Montlake Boulevard East, Seattle, Washington 98112, USA. Received February 18, 2010; accepted August 21, 2010

particularly time series data, to support ecological modeling and other endeavors.

To patch together the history of marine populations it is sometimes necessary to draw on unconventional sources of information. One useful and largely untapped resource is the local ecological knowledge (LEK) held by fishers and community members (Johannes et al. 2000). Although there are many examples in which LEK has been collected and used for study and management purposes, there are few examples in which such data are used quantitatively. Previous publications have used LEK to define species ranges (Gerhardinger et al. 2009). Some have applied statistical models to identify critical population declines (Turvey et al. 2009) or even estimate species abundance in terrestrial (Anadón et al. 2009) and marine systems (Ainsworth et al. 2008). Local ecological knowledge offers some important advantages over other types of scientific data: it is inexpensive to collect, it can be pertinent to a wide range of species, and it can inform our understanding of ecosystem changes over long time periods and wide geographic ranges. The data also reveal how the fishing industry perceives the resource and resource supply. This is a useful perspective for understanding fisher's motives and using the LEK data in applications.

This article presents the results from a series of LEK interviews recently conducted in the northern Gulf of California. The interviews help identify changes in the marine ecosystem from 1950 to the present as perceived by artisanal fishers. Considered in this study is whether the information conveyed by fishers is affected by their reliance on stocks for food and livelihood and whether younger and older fishers perceive ecosystem changes differently. Specifically, the analysis looks for evidence of the psychosocial phenomenon called shifting cognitive baselines (Pauly 1995), in which each generation of resource users accepts a lower standard as normal and so does not have an accurate appreciation of the true magnitude of historical resource decline.

The abundance trends from the LEK data are combined with five other data series developed here (simple stock size and fishery indicators) to produce composite trends detailing the likely changes in relative abundance for 42 species groups in the northern Gulf of California. The intention is that these time series for fish, invertebrates, mammals, birds, and reptiles will be useful in a variety of quantitative EBFM applications, including ecosystem modeling. A fuzzy logic algorithm (Zadeh 1965; Bellman and Zadeh 1970) is used here as a method of deriving numerical trends from ordinal LEK data, standardizing the various data streams, and combining them into a composite time series while resolving disagreements according to a transparent-rule matrix.

METHODS

Six data series are developed here and then integrated using a fuzzy logic system in order to reconstruct species abundance trends. Three series can be considered directly representative of species abundance, and the other three relate to the exploitation trends of fisheries. The abundance indicators are (1) species abundance trends from LEK interviews, (2) a catch-per-unit-effort (CPUE) series compiled from published and unpublished literature that represents relative species abundance over time, and (3) a separate yes/no indicator from LEK interviews corresponding to whether fishers had noticed a stock depletion during their careers. The exploitation indicators include (4) biomass predictions from a simple Schaefer (logistic growth) harvest model, (5) a measure of species groups' vulnerability to fishing, and (6) a yes/no indicator from LEK interviews corresponding to whether fishers had noticed a reduction in animal body size. Two parallel analyses are conducted and compared, one processing the abundance indicators and one processing the exploitation indicators. The next section explains the development of these indicators. The fuzzy logic implementation is described later.

Indicators from LEK Interviews

Researchers interviewed 81 fishers in the towns of Puerto Peñasco, Golfo de Santa Clara, Rodolfo Campodónico, Bahia Kino, Desemboque, and Puerto Libertad (Sonora) between April and June 2008 and September 2008 and February 2009. This represents around 2% of the estimated 3,800 artisanal fishers in the northern Gulf of California (P. Turk-Boyer, Centro Intercultural de Estudios de Desiertos y Océanos [CEDO], unpublished data). Interviewees ranged in age from 20 to 89 years and had expertise in the following gear types: gill net, cast net, shrimp and finfish trawls, longline, hand line, harpoon, compressor diving, and traps. On average, respondents had 28.5 years of fishing experience. The interview forms used are available from C. H. Ainsworth.

Interviewees were asked to characterize the abundance of fish, invertebrates, mammals, birds, and reptiles into one of three categories (low, medium, or high) for each of the six decades since 1950. Species were aggregated into groups; the structure was chosen to be compatible with species-aggregated, trophicmodeling approaches for fisheries research. The format of the groups generally aggregates invertebrate species, and it represents species of commercial interest and ecological importance, such as keystone species, in more detail. A statistical test is used here to show whether fishers' perceptions of species abundance varies depending on whether or not they rely on the stock for their livelihoods. For this, respondents were divided into two pools, those that targeted a given species group and those that did not. Relative abundance trends from LEK interviews were developed for 24 species groups (Table A.1 in the appendix) for each decade and used as input in the fuzzy logic procedure.

In addition to the abundance trends, simple indicators were collected in the form of yes/no questions. The questions for each species group were whether the respondents had noticed localized depletions or extirpations of the species and whether they had noticed a reduction in average body size. The yes/no responses were obtained for 36 species groups (Table A.1). They were not recorded by decade; a single response was assumed to represent the net change over a respondent's career.

Catch per Unit Effort

Catch-per-unit-effort trends were determined based on CPUE or catch-and-effort data from Mexican government statistics (Diario Oficial de la Federación 2004, 2006 [see Table A.2]; CONAPESCA 2009). Additional information was obtained from unpublished statistics (V. M. Valdéz-Ornelas and E. Torreblanca, Pronatura Noroeste). Data from a concurrent study to estimate unreported catch provided port-level information (S. Perez-Valencia, CEDO, unpublished data), while a study of fishery logbooks also provided high-quality information for a small number of vessels (J. Torre and M. Rojo, Comunidad y Biodiversidad, unpublished data). Finally, the remaining data gaps were filled by other literature sources listed in Table A.2.

Catch data for major commercial species were taken from Diario Oficial de la Federación (2004, 2006) and CONAPESCA (2009), while the unpublished information, surveys, logbooks, and literature sources provided estimates of catch for minor targets and bycatch species. Where statistics were available by state, total catch in the northern gulf was assumed equal to that of Sonora and Baja California combined. The catch values also include estimates of unreported catch and discards, which are largely based on expert opinion (L. E. Calderón-Aguilera, Centro de Investigación Científica y de Educación Superior de Ensenada, personal communication). Effort trends in artisanal fisheries were based on the number of vessels operating (Galindo-Bect 2003); this approach has the advantage that it can capture trends for unregistered vessels, which may constitute as much as one-third of the fleet, judging by recent aerial surveys (Rodríguez-Valencia et al. 2008). Other effort series were located for lobsters (Vega-Velásquez 2006), small pelagic organisms (Arreguín-Sánchez et al. 2006), groupers (Arreguín-Sánchez et al. 2006), totoaba Totoaba macdonaldi (Lercari and Chávez 2007), and shrimp (Galindo-Bect 2003). As a last resort, effort was based on human population growth rate in two Sonoran cities, Puerto Peñasco and San Felipe (INEGI and Government of Sonora 2008a, 2008b). Catch and effort references are listed in Table A.2. Full documentation of development of catch, effort, and CPUE series is available from the author (cameron.ainsworth@noaa.gov). The CPUE series were developed for 34 species groups (Table A.1) as annual trends (units vary) and averaged to the decade level for input into the fuzzy logic procedure.

Harvest Model

A logistic growth model with harvests provides biomass (*B*) estimates at each year *t*, as in equation (1) (Schaefer 1954; Hilborn and Walters 1992),

$$B_{t} = B_{t-1} + rB_{t-1} \left(1 - \frac{B_{t-1}}{K} \right) - C_{t-1}.$$
 (1)

Catch (C) was obtained from the assembled catch series (Table A.2). The carrying capacity of the ecosystem (K) was assumed to be the unfished biomass (B_0) estimated by Lozano-Montes (2006) and Lozano-Montes et al. (2008). The B_0 values

are highly uncertain, but they cover a large number of data-poor species and represent the best available estimates. The intrinsic rate of population increase (r) was estimated according to the equation $r = 4MSY/B_0$. Where possible, the maximum sustainable yield (MSY) was taken from Mexican stock assessments (Diario Oficial de la Federación 2004, 2006; Table A.2). In the absence of species distribution data, the MSY for the northern Gulf of California was assumed to be equal to the MSY values for Sonora and Baja California combined. In some cases, only values for Sonora were available, and these were inflated 20% to account for missing areas. Where the MSYs referred to Pacific Ocean stocks, the MSYs for gulf stocks were assumed to be similar on a per-area basis (the stock area was assumed to be 10% for pelagic species and 20% for other species in the gulf). If MSY estimates were not available, r was estimated using the empirical formula of Pauly (1984), that is,

$$r \approx 9.13 \bar{W}^{-0.26}$$
,

where \bar{W} is mean body weight, approximated as $\bar{W} = (W_{\text{max}} + W_m)/2$ (Pauly 1984), with $W_{\text{max}} = \text{maximum}$ weight and $W_{\text{m}} = \text{weight}$ at maturity. These weight data were collected from FishBase (references provided in Tables A.3, A.4).

The model was initialized at 1950 using B_0 and run to 2008, calculating annual biomass estimates B_t in t/km^2 . These were then averaged to produce decadal values ($B_{1950}, B_{1960}, \ldots, B_{2000}$) and used as inputs to the fuzzy logic algorithm. Series were developed for 26 species groups (Table A.1). Between averaging these biomass values over decades, using fuzzy sets to describe them, and combining these estimates with other abundance series, the method presented here should be insensitive to the large uncertainties involved in these calculations.

Vulnerability Index

As a final indicator of species abundance, the relative vulnerability to fishing of each exploited species group is estimated based on the method of Cheung et al. (2005). Those authors combined life history characteristics, including maximum length, the Von Bertalanffy growth constant, maximum age, fecundity, age at first maturity, and natural mortality, in a fuzzy logic framework to produce a composite vulnerability-to-fishing score. The index was shown empirically to predict species status better than common alternative proxies. Their method was recreated here. Life history information for Gulf of California species was taken from FishBase and other sources (see Tables A.3, A.4); then, using the published membership functions and expert rule set, the data were collated to produce a final vulnerability score for each species in a procedure analogous to the one described in this article. The only notable difference between this treatment and the one by Cheung et al. (2005) is that the geographic range of a species was not considered as an indicator of vulnerability because it is not relevant when considering a localized area like the Gulf of California. The vulnerability scores were calculated for individual species and averaged to the level of species groups. The available life history data allowed the calculation for 21 species groups (Table A.1). The scores are time independent, and so the same score is used for each decade in the analysis.

Fuzzy Logic Overview

Fuzzy set theory, also called fuzzy logic (Zadeh 1965; Bellman and Zadeh 1970), emulates an expert's judgment by combining inputs through a heuristic IF–THEN rule matrix to reach a conclusion regarding the data. It is a means of computing with words (Zadeh 1999), where "linguistic variables," representing a wide range of possible data types, combine according to relationships similar to "rules of thumb" contained in a rule matrix. However, where classical logic requires that a variable be categorizable into a single class (e.g., a Boolean variable belonging exclusively to a yes or no category), the linguistic variables in a fuzzy set can hold varying degrees of membership in multiple classes. For example, if "purple" is a fuzzy set that describes all colours composed of red and blue light, then indigo, violet, and fuchsia could be said to hold increasing membership in the "red" category relative to the "blue."

A Worked Example

Membership in the linguistic variable categories determines what rules operate (or "fire") in the rule matrix. In this paper, the strength at which the rules fire is determined by the algebraic sum of the intersecting memberships (Figure 1C). Many conclusions may be reached simultaneously (Figure 1D) with varying degrees of belief (firing strengths). Each time a cell in the rule matrix fires, it strengthens our belief in the corresponding conclusion. There are 50 different possible conclusions. Numbered 1 to 50, each conclusion represents a linear interpretation of abundance, so that a conclusion of 40 indicates twice the abundance of one numbered 20. Whenever a cell is fired, the partial membership that elicited that action is added to a running total for the corresponding conclusion category (Figure 1E). In this way, conclusions reached repeatedly (or fired with large partial memberships) will accrue a high score.

After all the information is passed through the matrix for a certain group-period combination, we are left with an array of

Repeat for each group (g) and period (p) combination

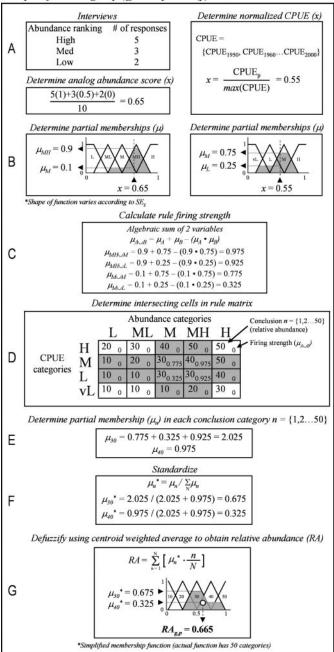


FIGURE 1. A worked example of the fuzzy logic method. A two-dimensional analysis is presented for clarity; it processes two data streams, abundance from interviews and CPUE (values are hypothetical). The algorithm presented here is repeated for each species group and time period.

50 elements; the partial memberships in each category represent our relative confidence, or degree of belief, in that abundance conclusion. The partial memberships are normalized as in Figure 1F, after which they are passed as inputs to the defuzzification membership function (Figure 1G). Defuzzification is the process by which partial memberships are converted to a single

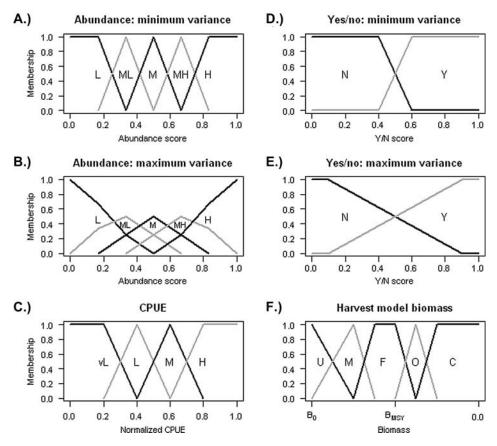


FIGURE 2. Fuzzy set membership functions. These relationships convert an analog indicator of abundance (*x*-axis) to partial memberships in linguistic abundance categories (*y*-axis). The partial memberships sum to 1 and represent the degree of belief in the abundance categories. Panels (**A**) and (**B**) show the membership in abundance categories under minimum and maximum variance between interview scores (low [L], medium–low [ML], medium [M], medium—high [MH], and high [H]); panel (**C**) shows the membership in CPUE categories (very low [vL], low [L], medium [M], and high [H]); panels (**D**) and (**E**) show the membership in yes/no (Y/N) categories under minimum and maximum variance between interview scores; and panel (**F**) shows the membership in biomass categories predicted by the harvest model (underexploited [U], moderately exploited [M], fully exploited [F], overexploited [O], and critically overfished [C]).

number representing relative abundance (i.e., a "crisp" number that is no longer part of a fuzzy set). It is therefore the reverse of the procedure described earlier to calculate memberships (i.e., Figure 1B). The defuzzification function in Figure 1G is a simplification; the actual one employed has 50 categories with triangular functions. The centroid-weighted average method of Cox (1999) is used, in which we multiply the centroid of each category (0.02, 0.04, 0.06, ..., 1.00) by the relative weighting (or confidence) in that conclusion to obtain a weighted average between 0 and 1 that represents the relative abundance for that species group and period.

Fuzzy logic implementation.—In the worked example, the rule matrix uses two dimensions (i.e., abundance and CPUE in Figure 1D). Scaling up, the implementation for both abundance and exploitation indicators uses three dimensional matrices. The next section describes the membership functions used to characterize the six data series involved. Later, combinatorial rules, the rule matrices, and defuzzification are discussed.

Membership functions.—The abundance information obtained in interviews was evaluated according to the membership

functions in Figure 2A, B. The *x*-axis in Figure 2A, B represents the analog abundance (*x*) score obtained from the interviews. It is averaged across respondents, "low" responses being assigned a value of 0, "medium" ones being assigned 0.5, and "high" ones being assigned 1.0. An average abundance score of 0.5 could be achieved through (at least) two scenarios: one in which every fisher reported medium abundance and one in which half of the fishers reported high abundance and half reported low abundance. To account for agreement between respondents, a dynamic membership function was used in which the angle subtended by the triangular functions increases from a minimum (Figure 2A) to a maximum (Figure 2B) if there was high or low agreement between respondents, respectively.

As a measure of agreement, the standard error of the mean was determined for each group–period combination as $SE_{\overline{X}} = \sigma^2/n$, where n is the number of respondents. Variance (σ^2) was calculated assuming a binomial distribution in which the majority response category (low, medium, or high) was considered "correct" and all other responses were considered "incorrect"; thus $\sigma^2 = np(1-p)$, where p is the fraction of correct responses.

Using the $SE_{\overline{\chi}}$ in this way corrects for the varying number of respondents per decade (e.g., fewer people contributed to the 1950s estimates than to the 2000–2009 estimates). The $SE_{\overline{\chi}}$ was next standardized so that the group–decade combination that had the best agreement (lowest error) adopted the membership function in Figure 2A, that with the poorest agreement adopted the function in Figure 2B, and other responses adopted intermediate functions, where slopes and intercepts were scaled linearly between the extremes.

Membership was evaluated in each of five fuzzy set abundance categories: low (L), medium—low (ML), medium (M), medium—high (MH), and high (H). Categories L and H use trapezoidal forms: beyond a certain threshold, full membership was assigned to these extreme categories. This allowed us to ignore the influence of a small number of responses that contradict the majority belief. Although the level of fishing effort, fishing skill, gear efficiency, catchability, and other factors will affect the amount of catch a fisher obtains for any given level of fish abundance, this methodology trusts that fishers can integrate over a wide range of externalities and thus have valid cognitive models of resource supply. Averaging their responses to obtain *x* should also eliminate many possible biases. For this reason, we restricted the analysis to consider only group—period combinations that had at least eight respondents.

The membership function used to categorize (normalized) CPUE per species group and decade is provided in Figure 2C. It categorizes the analog CPUE score into four linguistic categories: very low (vL), low (L), medium (M), and high (H). The membership functions used to categorize "yes/no" depletion and body size indicators into yes (Y) and no (N) categories again use a dynamic membership function to reflect the level of agreement between respondents (Figure 2D, E). As with the abundance indicators from interviews, the form of the membership function varies according to $SE_{\overline{x}}$, which was calculated assuming a binomial distribution of yes and no answers. The membership function used to evaluate biomass predictions from the logistic harvest model is provided in Figure 2F. Here, membership is evaluated in five linguistic categories: overexploited (O) or critically overfished (C) if the decadal biomass value (e.g., B_{1950} , $B_{1960}, \ldots, B_{2000}$) was below B_{MSY} , or underexploited (U), moderately exploited (M), or fully exploited (F) if biomass was above $B_{\rm MSY}$. In the case of the Cheung et al. (2005) vulnerability index, our input membership function is identical to their output (defuzzification) membership function; it is equivalent to Figure 2C with four linguistic categories: low (L), medium (M), high (H), and very high (vH).

Combining the data series.—Having determined the partial memberships in linguistic categories through the use of membership functions, we next consult the rule matrices to combine the information; Tables 1A and 1B show the abundance and exploitation matrices, respectively.

Membership in the three indicators (on X, Y, and Z axes of each matrix) leads us to a conclusion regarding the abundance for each species in each time period. The conclusions are

found inside the matrix cells (color coded in Table 1). Each cell fires with a strength proportional to the algebraic sum of the intersecting memberships. All axes are assumed to be independent, and the strength of memberships is combined using the probability-OR operator for three variables, namely,

$$\mu_{A \cup B \cup C} = \mu_A + \mu_B + \mu_C - (\mu_A \cdot \mu_B) - (\mu_A \cdot \mu_C) - (\mu_B \cdot \mu_C) + (\mu_A \cdot \mu_B \cdot \mu_C), \qquad (2)$$

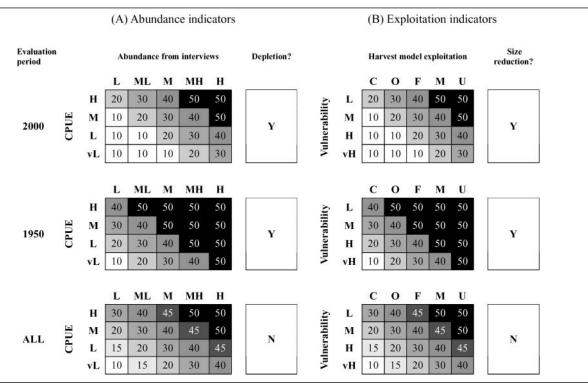
where $\mu_{A\cup B\cup C}$ is representative of our degree of belief in the corresponding conclusion. This fuzzy union operator was used based on the algebraic sum rather than the alternative union operator (μ_A OR μ_B OR $\mu_C = max[\mu_A, \mu_B, \mu_C]$) or fuzzy intersection operator (μ_A AND μ_B AND $\mu_C = min[\mu_A, \mu_B, \mu_C]$) used by previous authors (Cheung et al. 2005; Ainsworth et al. 2008) so that all operands contribute something to the output; the algorithm is thus useful for a wider range of data availabilities (see Table A.1).

Various parts of the rule matrix were accessed for each time period (1950, 1960, 1970, 1980, 1990, and 2000). If the value of the depletion indicators (i.e., stock depletion or body size reduction) is "yes," this has the effect of lowering the relative abundance score of the conclusion for recent periods (1980 to 2000) but increasing the abundance score of the conclusion for older periods (1950 to 1970) (i.e., the slope between 1950 and 2000 becomes more negative; Table 1). Matrices for the intermediate periods (1960, 1970, 1980, and 1990) are not shown, but they apply a smooth linear transition between the extreme values in 2000 (top row of Table 1) and 1950 (middle row of Table 1). If the value of the depletion indicator is "no," then the bottom row of Table 1 is accessed regardless of decade.

After all data series pass through the matrix, we are left with 50 different abundance conclusions with varying degrees of belief (partial memberships), as described in the worked example. The partial memberships are combined through defuzzification and are converted to a single crisp number representing the relative abundance. This process is repeated for each species group and period to produce time series of relative abundance that can be compared with observational series (Table 2).

Summary.—The data series used here for abundance indicators consisted of decadal abundance trends from LEK interviews, annual CPUE data averaged to decades, and a yes/no population-level depletion indicator from LEK interviews that referred to all decades. The data series used for the exploitation indicators consisted of annual stock biomass from a logistic growth harvest model averaged to decades, a vulnerability-to-fishing index that referred to all decades, and a yes/no body size change indicator from LEK interviews that referred to all decades. These numerical indicators were translated into memberships in ordinal linguistic categories (e.g., low, medium, or high) using membership functions, where membership in each category represents our relative belief in that category's being true. The memberships in each category determined the firing

TABLE 1. Heuristic rule matrices used by the fuzzy logic algorithm to combine data sources. Two parallel matrices deal with abundance and exploitation indicators, respectively. Abundance indicators include abundance estimated by interviews, catch per unit effort (CPUE) estimated from literature values, and stock depletion suggested by interviews. Exploitation indicators include exploitation status suggested by the Schaefer harvest model, vulnerability to fishing based on life history characteristics, and body size reduction suggested by interviews. The conclusions resulting from these X-Y-Z linguistic variables are identified in the cells. They represent a linear abundance index that ranges from 1 to 50. The presence of depletion or size-reduction indicators upgrades the abundance conclusion for past periods (e.g., 1950) and downgrades it for recent periods (e.g., 2000). Matrices for the intermediate periods (1990, 1980, 1970, and 1960; not shown) apply a smooth transition between the extremes 2000 and 1950. Abbreviations are as follows: L = low, ML = medium-low, M = medium, MH = medium-high, H = high, vL = very low, vH = very high, C = critically overfished, O = overexploited, F = fully exploited, M = moderately exploited, u = underexploited, Y = yes, and N = no.



strength of cells in a rule matrix, where each cell leads to a particular conclusion regarding species abundance. Finally, a crisp abundance score was determined through defuzzification, the reverse of the process that created the membership scores from the numerical indicators. Abundance per decade is plotted for each species group, representing its stock status since 1950.

RESULTS

For many species groups, the respondents in the LEK interviews were likely to indicate high biomass in the 1950s and low biomass in 2000–2009 (Figure 3). The trends are not often monotonic and there are conspicuous exceptions, like pinnipeds and seabirds. For these groups, the respondents were more likely to indicate high biomass in recent years. For pinnipeds, this conflicts with known population trends in the Gulf of California as a whole (Szteren et al. 2006; Wielgus et al. 2008). However, census data suggest that California sea lion *Zalophus californianus* rookeries in the north, where interviews occurred, may have seen population increases since the early 1990s (Szteren et al. 2006). The status of seabird populations in the gulf

is poorly known from scientific studies (Palacios and Alfaro 2005).

Comparing the abundance scores offered by people who depend on the resource economically with the scores from those who do not revealed no significant difference for any species in responses for the 2000 period (two-tailed Mann–Whitney U-test: P > 0.05). This held true for all 15 species groups tested (i.e., all exploited fish and invertebrate species; minimum sample size = 6). This suggests that fishers offered an unbiased view regardless of whether or not they depend on the stock for their livelihood, and being specialized in catching a certain type of animal did not improve or alter their assessment relative to that of a "nonexpert" fisher.

When asked to characterize the abundance of species groups for the decades between 1950 and the present, fishers showed the most agreement for the earliest decade, the 1950s (Figure 4). They generally agreed that abundance was high during this period (irrespective of the species group). Each subsequent decade had less agreement, except for the most recent decade, 2000, in which interviewees tended to agree that abundances were low. This pattern was consistent across species groups, with a few exceptions.

TABLE 2. Taxa included in the current study and those in previous studies estimating abundance and biomass trends in the northern Gulf of California.

Functional group	Species in source material	Type of information	Reference	
Blue crab	Arched swimming crab <i>Callinectes</i> arcuatus, blue swimming crab <i>C.</i> bellicosus	Relative biomass	INP (2006)	
Blue shrimp	Blue shrimp <i>Litopenaeus stylirostris</i>	CPUE	Galindo-Bect et al. (2000)	
Crabs and lobsters	Spiny lobsters (<i>Panulirus interruptus</i> , <i>P. inflatus</i> , <i>P. gracilis</i>)	CPUE	INP (2006)	
Groupers and snappers	Snapper <i>Lutjanus</i> spp., barred pargo <i>Hoplopagrus guntherii</i>	CPUE	INP (2006)	
Herbivorous echinoderms	Red urchin Strongylocentrotus franciscanus	Biomass	INP (2006)	
Large pelagic sharks	Large sharks <i>Carcharhinus</i> spp., <i>Alopias</i> spp., scalloped hammerhead <i>Sphyrna lewini</i> , whitenose shark <i>Nasolamia velox</i>	CPUE	INP (2006)	
Penaeid shrimp	Brown shrimp <i>Penaeus californiensis</i> , blue shrimp <i>P. stylirostris</i> , and white shrimp <i>P. vannamei</i>	CPUE	Magallon-Barajas (1987)	
Pinnipeds	California sea lion Zalophus californianus californianus	Relative biomass	Szteren et al. (2006)	
Small pelagics	Small pelagics (Pacific sardine Sardinops sagax caeruleus, herrings Opisthonema spp., Pacific chub mackerel Scomber japonicus, anchoveta Cetengraulis mysticetus, round herring Etrumeus teres, leatherjacks oligoplites spp., northern anchovy Engraulis mordax)	CPUE	INP (2006)	
Squid	Jumbo squid <i>Dosidicus gigas</i>	Relative biomass	Nevárez-Martínez et al. (2006)	
Totoaba	Totoaba Totoaba macdonaldi	Biomass	Larcari and Chavez (2007)	

Analyzing only the remarks from the interviewees concerning species abundance, we found that older fishers tended to recognize a greater degree of population decline since 1970 than did younger fishers (Figure 5). We considered the time since 1970 rather than that since 1950 in order to include a greater number of respondents. All species groups tested are aggregated into the six categories shown in Figure 5. The relationship between fisher age and reported abundance change is significant for mammals, other fish, turtles, and invertebrates, weakly significant for reef fish, and nonsignificant for birds (F-test). However, the results are not significant when we contrast the perceived decline against the number of years of fishing experience rather than fishers' ages: reef fish (P = 0.087), mammals (P = 0.35), birds (P = 0.31), other fish (P = 0.18), turtles (P = 0.012), and invertebrates (P = 0.51). Trusting the cognitive model of stock abundance held by older fishers (those above the median age of 43) produces a much different result in the hindcasted abundance trends resulting from the fuzzy logic routine than trusting that of the younger fishers (Figure A.1).

However, the differing opinions of these groups provide a range of plausible trajectories for relative abundance over time. The discrepancy is most noticeable in targeted and charismatic species.

The method of Cheung et al. (2005) provides an estimate of vulnerability to fishing based on life history patterns (Figure 6). Elasmobranchs, which tend to be long-lived and late maturing and have low fecundity, show the greatest overall vulnerability. This is consistent with their known biology (Stevens et al. 2000; Sadovy 2001) and the history of exploitation in the northern Gulf of California (Bizzarro et al. 2009). Also vulnerable are reef fish species, in particular, large-bodied piscivores that aggregate during breeding, such as the species included in the "groupers and snappers" group and the "large reef fish" group (Musick et al. 2000).

Crisp outputs of the fuzzy logic algorithm suggest that many species groups have suffered some degree of depletion since 1950 (Figure 7). There is satisfactory agreement between outputs from the abundance indicators and the exploitation

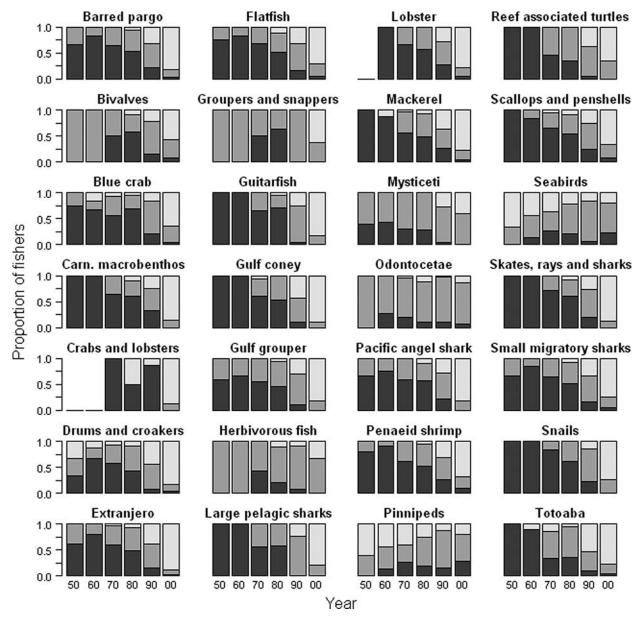


FIGURE 3. Actual interview results, presented as the proportions of fishers reporting high (dark gray), medium (medium gray), and low biomass (light gray) for each species group and time period (50 = the 1950s and so forth). Species groups that had at least eight respondents are shown. The composition of the groups is given in Table A.5 in the appendix.

indicators. Groups that show serious discrepancy between these two series are pinnipeds and blue shrimp; all other groups achieved at least a qualitative similarity. The trend based on exploitation indicators suggests that pinniped numbers have decreased; this conclusion is largely based on a perceived body size decrease by interviewees. However, the trend based on abundance indicators suggests an increase for these animals, reflecting the source interview results mentioned earlier (see Figure 3). In the case of blue shrimp, the exploitation indicators suggest a steady depletion of the stock, a result also suggested by the harvest model. However, the population trend signified by

the abundance indicators suggests that the abundance increased from 1950 to 1980 and subsequently declined. This shows the influence of the CPUE series. The initial increase in CPUE may reveal population changes, or it may reflect the introduction of modern fishing methods that increased fishing efficiency. For both pinnipeds and blue shrimp, previously published abundance series support the apparent increase in abundance proffered by the abundance indicators and discredit the decrease in abundance proffered by the exploitation indicators. The abundance indicators consistently agree with the direction of change suggested by the observational data.

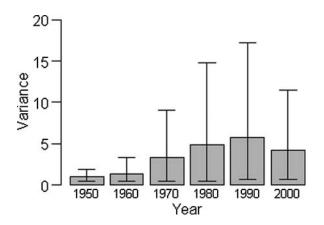


FIGURE 4. Variance among fishers' abundance scores from interviews. Variance is calculated assuming a binomial distribution (see text). Bars show the average score for all species groups analyzed; error bars show total range.

DISCUSSION

The results from interviews consistently indicated a downward trend for most populations, although this could reflect a spatial bias in reporting if fishers are referring to localized depletions in fishing areas. The interviews also were concentrated in the most heavily populated and exploited region of the gulf, which is in the northeast; this also adds a potential bias. Nevertheless, the abundance and exploitation series usually agree, which lends credence to the overall trend. In other words, the

LEK data that informed the abundance trends are consistent with the catch and life history information that informed the exploitation trends. Estimates of relative biomass and abundance from previous studies tend to corroborate the fuzzy logic outputs despite a slight mismatch in species groupings and study area used. However, decreases in relative abundance (or body size) do not necessarily indicate overexploitation, as these are natural consequences of harvesting even in well-managed stocks and may be desirable (Hilborn 2007). Although the trends determined here are relative, being comparable only across species groups and between time periods, it would be possible to develop absolute trends by scaling the fuzzy logic output to match the available partial time series from scientific sampling (as in Ainsworth et al. 2008). As yet, it is not feasible to do this in the northern Gulf of California since so few groups have reliable survey information specific to the study area.

One noteworthy result is that even most untargeted species are reported by fishers to have declined. There could be psychological factors influencing this perception among fishers (Ainsworth et al. 2008); fishers' perceptions factor into both the abundance and exploitation indicators. Unfortunately, the suggestion that both predator and prey are declining may be plausible given the major ecological changes in the Gulf of California over the last century. Regulation of the Colorado River flow by numerous dams and increased freshwater consumption in the southwestern United States and Mexico has altered the marine assemblage (Rodríguez et al. 2001; Lozano-Montes

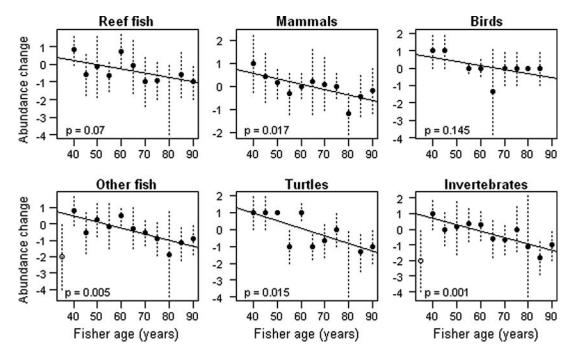


FIGURE 5. Evidence of shifting baselines in the northern Gulf of California. The *y*-axis shows the change in average perceived abundance between 1970 and 2000, the *x*-axis the age of fishers. A change of -1.0 corresponds to a reduction from high to medium or medium to low in terms of the average interview abundance score. All relationships are significant at $\alpha = 0.05$ (*F*-test) except those for reef fish and birds. Outliers (open circles) were removed from the data for invertebrates and other fish; error bars = SDs.

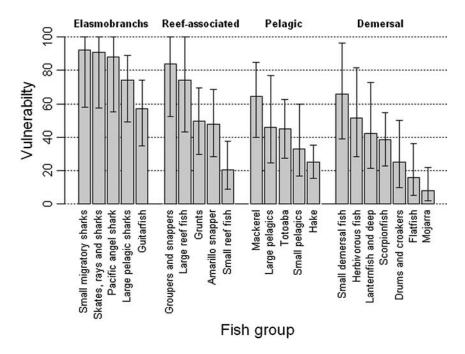


FIGURE 6. Vulnerability to fishing as predicted by the algorithm of Cheung et al. (2005) based on life history characteristics. The error bars show the upper and lower bounds of the conclusion fuzzy membership function designed by Cheung et al. and reflect the precision of their estimates based on the width of their output membership functions. The composition of the groups is given in Table A.5.

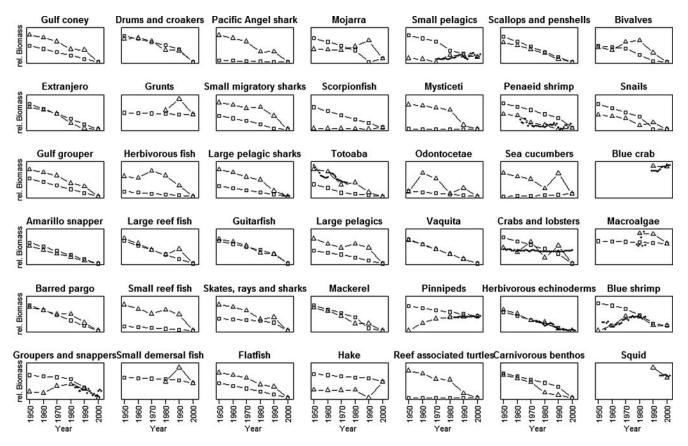


FIGURE 7. Abundance of species groups predicted by the fuzzy logic algorithm, as suggested by abundance indicators (triangles) and exploitation indicators (squares). Trends have been scaled to agree in the year 2000. Bullet points show the relative abundance trends of representative species from previous studies (Table 2); residuals are minimized with respect to abundance indicators. See Figure A.1 for the effect of respondent's age on abundance indicators. The composition of the groups is given in Table A.5.

2006), affecting the salinity gradient, estuarine habitat conditions, nutrient loading, and other factors important to fish and invertebrate production (Lavín and Sanchez 1999; Galindo-Bect et al. 2000). Therefore, management of this area may consider whether forcing factors unrelated to fishing effort have had a major influence on ecosystem structure.

Even in a region like the northern Gulf of California, which is poor in scientific data, it is possible to identify which species are likely to require special attention by fishery managers with a minimal investment in effort and no regional data using Cheung et al.'s (2005) vulnerability method. In this application, life history values were averaged across all constituent species in the aggregated species groups, so the vulnerability scores in Figure 6 represent the "average" fish in each of these groups. Individual species within groups will vary in life history parameters. We may expect the most vulnerable species to decline first under fishing pressure and perhaps even to be extirpated before the group has become seriously depleted. Methods exist to predict the most vulnerable species within a species group (Cheung and Pitcher 2004). Aggregating similar species into groups is a useful convenience for generalizing fishers' perceptions, and it is necessary for many EBFM ecosystem modeling approaches in order to simplify the food web and manage data gaps (although aggregating species carries a strong set of assumptions) (Chalcraft and Resetarits 2003).

Analysis of interview results suggests that older fishers perceive a greater decline in abundance since 1970 than do younger fishers. This confirms the results of two earlier studies in the Gulf of California suggesting that the true magnitude of stock decline is not well appreciated by those who rely on the resource (Sáenz-Arroyo et al. 2005; Lozano-Montes et al. 2008). These findings add to a growing body of literature on the subject of shifting cognitive baselines. Similar studies have been conducted in many areas of the world, with consistent results that validate Pauly's (1995) premise (e.g., North America [Baum and Myers 2004], Asia [Ainsworth et al. 2008], Africa [Bunce et al. 2008], and Europe [Airoldi and Beck 2007]).

The quantitative aspects of EBFM require some ability to forecast ecosystem-level population effects, but the data requirements of EBFM models are a barrier to their use, especially in regions like the northern Gulf of California where sufficient sampling data are unavailable. The fuzzy logic technique presented here, which was adapted and improved from Ainsworth et al. (2008), can provide numerical abundance trends that substitute for formal stock assessment. The method relies heavily on qualitative information. Nevertheless, it offers a transparent, replicable, and flexible tool that can be updated as new information becomes available. The quality of outputs is still limited by the available data. For example, difficulties in applying LEK information (Brook and McLachlan 2005) or CPUE data (Beverton and Holt 1957; Hilborn and Walters 1992) still apply. In other ecosystems, available data may support the use of a more sophisticated harvest model. However, when several sources of data are combined, reliance on any one source is reduced. This work represents the first attempt to describe abundance trends for many of the species it considers. The fuzzy logic approach is flexible enough to accommodate a range of data, and it can provide marine ecologists with time series information on abundance for a variety of EBFM applications in data-limited situations.

ACKNOWLEDGMENTS

I thank the following researchers at the Centro Intercultural de Estudios de Desiertos y Océanos (CEDO, Puerto Peñasco): Mabilia Urquidi, Sandra Reyes, Hem Nalini Morzaria Luna, Abigail Iris, and Eleazar López, along with Nabor Encinas of Comunidad y Biodiversidad (Guaymas) for carrying out the interviews. The following people provided helpful discussions and review of the manuscript: Isaac Kaplan, Phil Levin, Marc Mangel, Hem Nalini Morzaria Luna, Nick Tolimieri, Jameal Samhouri (Northwest Fisheries Science Center), and William Cheung (University of East Anglia). I also thank Kenneth Rose and two anonymous referees for their careful reviews, which greatly improved the quality of the manuscript. The David and Lucille Packard Foundation provided funding for this study. The Mia J. Tegner Memorial Research Grants Program provided funding for CEDO community interviews.

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Appendix: Supplemental Material for Quantifying Species Abundance Trends

TABLE A.1. Availability of data by functional group. The composition of the groups is given in Table A.5.

	Abundance indicator			Exploitation indicator		
Functional group	Abundance	CPUE	Depletion	Harvest model exploitation	Vulnerability	Size reduction
Gulf coney	√	√	\checkmark			\checkmark
Extranjero	\checkmark	\checkmark	\checkmark			\checkmark
Gulf grouper	\checkmark	\checkmark	\checkmark			\checkmark
Amarillo snapper			\checkmark		\checkmark	\checkmark
Barred pargo	\checkmark	\checkmark	\checkmark			\checkmark
Groupers and snappers	\checkmark	\checkmark	\checkmark	\checkmark	\checkmark	\checkmark
Drums and croakers	$\sqrt{}$	$\sqrt{}$				$\sqrt{}$
Grunts		$\sqrt{}$				
Herbivorous fish		$\sqrt{}$	\checkmark	·	$\sqrt{}$	\checkmark
Large reef fish		\checkmark	$\stackrel{\cdot}{\checkmark}$		$\sqrt{}$	$\stackrel{\cdot}{\checkmark}$
Small reef fish		\checkmark	$\sqrt{}$		\checkmark	$\sqrt{}$
Small demersal fish		√	•	\checkmark	√	•
Pacific angel shark	√	√	√	•	v /	√
Small migratory sharks	v /	1 /	2/	2/	2/	1 /
Large pelagic sharks	v /	•	2/	1/	2/	1 /
Guitarfish	v /	2/	v /	v /	2/	v /
Skates, rays and sharks	v /	V	v /	V	2/	V
Flatfish	• /	•/	•/	•/	•/	•/
Mojarra	• V	•/	•/	•/	•/	•/
Scorpionfish		·V	•/	•/	•/	•/
Totoaba	./	./	•/	./	./	./
Large pelagics	• V	./	•/	·V	./	v
Mackerel Mackerel	./	./	./	./	./	./
Hake	V	./	./	./	./	./
Small pelagics		./	·/	·/	./	·/
Mysticeti	/	V	~/	~/	V	~/
Odontocetae	v /		~/	~/		~/
Vaquita	V		v /	v /		v /
Pinnipeds	/		v /	v /		~ /
Reef-associated turtles	v /	/	v /	V		~ /
Scallops and penshells	v /	v /	~ /			~ /
Penaeid shrimp	~ /	~ /	~ /	/		~ /
Sea cucumbers	V	~ /	~ /	\checkmark		√ /
Crabs and lobsters	/	~ /	√ /	/		√ /
Herbivorous echinoderms	\checkmark	\checkmark	√ ,	\checkmark		√ /
	/	,	√ ,	/		√
Carnivorous macrobenthos	√ ,	\checkmark	√ ,	\checkmark		√ ,
Bivalves	\checkmark	√ ,	\checkmark	\checkmark		√ ,
Snails	\checkmark	\checkmark	\checkmark	\checkmark		\checkmark
Adult blue crab		\checkmark		,		
Macroalgae		\checkmark		\checkmark		
Adult blue shrimp		\checkmark		\checkmark		
Squid		\checkmark				

Catch

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Effort

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TABLE A.2. Continued.

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TABLE A.3. Life history parameter references. Numbers preceded by the letters FB are Fishbase reference numbers searchable at www.fishbase.org; all other numbers refer to the references in Table A.4. The composition of the groups is given in Table A.5.

Functional group	References
Gulf coney	1,29,45,50,80
Extranjero	1,8,29,52,FB9342
Leopard grouper	1,29,32,35,52,84,92
Gulf grouper	1,85,FB6323
Amarillo snapper	25,28,81,82,96,FB6323,FB30872
Barred pargo	3,29,FB6323
Groupers and snappers	7,14,29,46,50,52,FB55,FB2850,FB3083,FB3090,FB3094,FB3244,FB4841,FB5227,FB5592, FB6323,FB6852,FB7185,FB9342,FB26176,FB27020,FB39376,FB40592,FB46367,FB47696
Drums and croakers	6,18,19,29,39,52,59,60,64,71,73,79,88,98,FB1164,FB1166,FB1180,FB6323,FB6997,FB26176
Grunts	2,36,39,46,60,62,67,FB2850,FB9114,FB11035,FB26176,FB26585,FB40926
Herbivorous fish	43,46,51,59,62,71,93,94,95,FB268,FB273,FB1048,FB1049,FB1238,FB1396,FB1836,FB2334, FB2850,FB3678,FB3807,FB4617,FB5533,FB5592,FB5760,FB6380,FB6997,FB7253,FB8540, FB8571,FB9072,FB9267,FB9310,FB9338,FB11824,FB12360,FB12544,FB13715,FB26177, FB26587,FB28725,FB29555,FB30504,FB42001,FB48600
Large reef fish	10,14,16,19,26,37,38,52,59,60,71,101,FB1263,FB1602,FB1751,FB2850,FB3669,FB3678, FB3807,FB4525,FB4604,FB5227,FB5525,FB5592,FB6323,FB6390,FB6490,FB7142,FB8571, FB9276,FB9292,FB9311,FB9312,FB9328,FB9329,FB9341,FB11482,FB11824,FB12260,FB26112, FB26585,FB26587,FB27803,FB28688,FB37992,FB39266,FB39376,FB40789,FB40870,FB48600
Small reef fish	23,24,31,33,38,39,46,52,55,57,58,59,60,68,75,86,FB273,FB559,FB583,FB1602,FB1661,FB2272, FB2850,FB3141,FB3678,FB3807,FB5227,FB5525,FB5590,FB5592,FB6113,FB6323,FB6937, FB7247,FB9072,FB9269,FB9286,FB9289,FB9299,FB9307,FB9324,FB9333,FB9334,FB9348, FB9349,FB9710,FB11482,FB11824,FB26176,FB26177,FB26585,FB26587,FB28023,FB30504, FB30573,FB37955,FB50710

TABLE A.3. Continued.

Functional group	References
	2,14,27,52,54,59,60,72,75,100,FB273,FB1371,FB2272,FB2850,FB3669,FB3678,FB4423, FB4461,FB4525,FB4925,FB5778,FB6323,FB6347,FB8571,FB8991,FB9269,FB9271,FB9277, FB9279,FB9284,FB9301,FB9307,FB9324,FB9348,FB9992,FB10887,FB11482,FB11824,FB26176 FB26585,FB26587,FB26773,FB27327,FB28023,FB31276,FB32350,FB34120,FB34613,FB35910, FB37955,FB38374,FB38398,FB39877,FB43481
Pacific angel shark	13,43,60,FB6147
Small migratory sharks	14,102,FB244,FB6097,FB6098,FB9253
Large pelagic sharks	14,43,60,66,FB244,FB273,FB1661,FB1671,FB3213,FB3222,FB3678,FB6084,FB6088,FB6090, FB6390,FB6937,FB7200,FB8571,FB9012,FB9165,FB11824,FB12489,FB13713,FB26587, FB27093,FB27603,FB27605,FB27971,FB31395,FB31509,FB32047,FB32086,FB32407,FB35388, FB36559,FB41862,FB42004
Guitarfish	14,83,FB2850,FB6323,FB9262
Skates, rays, and sharks	14,15,52,63,71,78,89,FB244,FB247,FB2850,FB3678,FB5255,FB6145,FB6323,FB6871,FB9255,FB9257,FB9259,FB9261,FB9265,FB28023,FB43028
Flatfish	14,34,41,52,56,59,60,76,77,FB2272,FB2850,FB6997,FB9047,FB9281,FB9294,FB9330,FB9331, FB11035,FB36656,FB41284,FB42865,FB43299,FB47359,FB47696
Mojarra	14,46,FB6937,FB6997,FB8540,FB8571,FB9303,FB11824,FB26176
Scorpionfish	59,60,61,71,FB2850,FB5592,FB5760,FB6937,FB7055,FB9341,FB9352,FB11824,FB26176, FB27129,FB41274
Lanternfish and deep	69,FB2850,FB4525,FB28499,FB40826
Totoaba	5,19,20,40,71,FB796
Large pelagics	4,11,12,14,39,43,44,49,52,59,60,70,71,74,91,99,FB14,FB26,FB168,FB238,FB268,FB273,FB515, FB1139,FB1263,FB1314,FB1333,FB1345,FB1374,FB1386,FB1392,FB1414,FB1447,FB1449, FB1462,FB1467,FB1475,FB1498,FB1656,FB2850,FB2885,FB3555,FB3605,FB3669,FB3678, FB3786,FB4332,FB4525,FB4560,FB4838,FB4972,FB5337,FB5338,FB5340,FB5525,FB5730, FB5760,FB5964,FB6323,FB6390,FB6814,FB6937,FB7161,FB7173,FB7253,FB8571,FB9283, FB9319,FB9345,FB9346,FB9898,FB9987,FB11482,FB11824,FB12193,FB12260,FB12451, FB12757,FB13304,FB26319,FB26340,FB26370,FB26587,FB26849,FB27030,FB28023,FB28050, FB28958,FB32349,FB33193,FB34133,FB34137,FB34148,FB35465,FB36276,FB36645,FB36658, FB36794,FB37040,FB37813,FB41559,FB41779,FB43794,FB46593
Mackerel	14,43,43,49,52,53,62,65,71,97,FB766,FB796,FB1662,FB3578,FB3730,FB3733,FB4332, FB4525,FB4530,FB5760,FB5960,FB5964,FB6014,FB6323,FB7032,FB7193,FB12393, FB13305,FB28200,FB33255,FB35185,FB39376,FB40755,FB42455
Hake	9,17,71,FB1139
Small pelagics	14,21,22,30,42,46,47,48,49,59,60,71,87,90,FB188,FB189,FB796,FB831,FB833,FB835, FB839,FB840,FB850,FB905,FB907,FB908,FB909,FB917,FB1139,FB1836,FB2197,FB2850, FB3231,FB3669,FB3678,FB3730,FB4525,FB5760,FB5769,FB5888,FB6997,FB9273,FB9291, FB9298,FB9300,FB9336,FB10851,FB11192,FB11482,FB26420,FB27758,FB33192,FB33520, FB34034,FB37813,FB39882,FB41293

TABLE A.4. Full references for life history parameters.

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TABLE A.5. Species comprising the functional groups. Group names are in bold. The groups are based on an Atlantis ecosystem model by Ainsworth et al. (in press).

Gulf coney

Epinephelus acanthistius

Extranjero

Paralabrax loro

Paralabrax auroguttatus

Leopard grouper

Mycteroperca rosacea

Gulf grouper

Mycteroperca jordani

Amarillo snapper

Lutjanus argentiventris

Barred pargo

Hoplopagrus guentherii

Groupers and snappers

Pronotogrammus multifasciatus

Hemanthias peruanus

Cephalopholis panamensis

Diplectrum labarum

Pronotogrammus eos

Serranus aequidens

Liopropoma longilepis

Alphestes immaculatus

Diplectrum euryplectrum

Diplectrum macropoma

Diplectrum pacificum

Diplectrum sciurus

Alphestes afer

Epinephelus analogus

Dermatolepis dermatolepis

Epinephelus itajara

Epinephelus labriformis

Alphestes multiguttatus

Epinephelus niphobles

Epinephelus niveatus

Lutjanus aratus

Lutjanus colorado

Lutjanus guttatus

Lutjanus novemfasciatus

Lutjanus peru

Lutjanus viridis

Mycteroperca prionura

Mycteroperca xenarcha

Paralabrax maculatofasciatus

Paranthias colonus

Pseudogramma thaumasium

Rypticus bicolor

Rypticus nigripinnis

Serranus fasciatus Serranus psittacinus Drums and croakers

Atractoscion nobilis

Bairdiella icistia

Cheilotrema saturnum

Cynoscion parvipinnis

Umbrina roncador

Ophioscion scierus

Odontoscion xanthops

Micropogonias megalops

Micropogonias ectenes

Micropogonias altipinnis

Menticirrhus nasus

Larimus effulgens

Larimus argenteus

Larimus acclivis

Isopisthus remifer

Elattarchus archidium

Cynoscion othonopterus

Cynoscion nannus

Ophioscion strabo

Stellifer illecebrosus

Umbrina wintersteeni

Umbrina xanti

Corvula macrops

Cynoscion squamipinnis

Bairdiella armata

Cynoscion reticulatus

Cynoscion xanthulus

Larimus pacificus

Menticirrhus undulatus

Grunts

Anisotremus interruptus

Conodon serrifer

Anisotremus caesius

Anisotremus davidsonii

Anisotremus dovii

Anisotremus taeniatus

Haemulon flaviguttatum Haemulon maculicauda

Haemulon scudderii

Haemulon sexfasciatum

Haemulon steindachneri

Microlepidotus inornatus

Orthopristis chalceus

Orthopristis reddingi

Pomadasys panamensis

Xenistius californiensis

Herbivorous fish

Acanthurus xanthopterus

Prionurus punctatus

Acanthurus achilles

Acanthurus nigricans

Calotomus spinidens

Girella nigricans

Hermosilla azurea Mugil cephalus

Mugil curema

Nicholsina denticulata

Scarus compressus

Scarus ghobban

Scarus perrico

Scarus rubroviolaceus

Acanthurus triostegus

Balistes polylepis

Large reef fish

Pseudobalistes naufragium

Sufflamen verres

Chlopsis kazuko

Gorgasia punctata

Heteroconger canabus

Heteroconger digueti

Ariosoma gilberti

Paraconger similis Paraconger californiensis

Bathycongrus macrurus

Bathycongrus varidens

Gnathophis cinctus

Rhynchoconger nitens

Thalassoma lucasanum

Nemichthys larseni

Myrichthys maculosus

Myrophis vafer

Ophichthus triserialis

Ophichthus zophochir

Callechelys cliffi

Callechelys eristigma

Ethadophis byrnei

Apterichtus gymnocelus

Brotula clarkae

Lepophidium microlepis

Lepophidium negropinna

Lepophidium pardale

Lepophidium prorates Lepophidium stigmatistium

Petrotyx hopkinsi

Neobythites stelliferoides Regalecus glesne

TABLE A.5. Continued.

Sebastes diploproa Sebastes macdonaldi Sebastes sinensis Sebastes cortezi Sebastes exsul Trichiurus lepturus Bodianus diplotaenia Decodon melasma Halichoeres aestuaricola Halichoeres chierchiae Halichoeres dispilus Halichoeres nicholsi Halichoeres notospilus Halichoeres semicinctus Lobotes pacificus Nematistius pectoralis Pareques fuscovittatus Pareques viola

Pseudojuloides inornatus Halichoeres melanotis Semicossyphus pulcher Stereolepis gigas Thalassoma lutescens Thalassoma purpureum

Small reef fish

Antennarius sanguineus Apogon retrosella Ablennes hians

Hypsoblennius brevipinnis
Entomacrodus chiostictus
Grammonus diagrammus
Acanthemblemaria balanorum
Acanthemblemaria crockeri
Coralliozetus angelicus
Coralliozetus micropes
Cirriemblemaria lucasana
Protemblemaria bicirrus
Prognathodes falcifer
Forcipiger flavissimus
Chaetodon humeralis
Johnrandallia nigrirostris

Oxycirrhites typus Cirrhitus rivulatus Diodon hystrix Diodon holocanthus Chaetodipterus zonatus Fistularia commersonii

Cirrhitichthys oxycephalus

Arcos erythrops
Gobiesox adustus
Tomicodon humeralis
Tomicodon zebra
Gobiosoma paradoxum
Kyphosus elegans
Paraclinus altivelis
Xenomedea rhodopyga
Uropterygius macrocephalus
Gymnomuraena zebra
Muraena lentiginosa
Echidna nocturna
Enchelycore octaviana
Gymnothorax castaneus
Gymnothorax dovii

Gymnothorax panamensis Muraena argus Muraena clepsydra Holacanthus passer Pomacanthus zonipectus Abudefduf troschelii Chromis atrilobata Abudefduf concolor Chromis alta Chromis limbaughi Microspathodon bairdii

Microspathodon dorsalis Stegastes leucorus Stegastes rectifraenum Abudefduf declivifrons Synodus lucioceps Sphoeroides annulatus Arothron meleagris

Canthigaster punctatissima Enneanectes carminalis

Zanclus cornutus

Aluterus scriptus

Acanthemblemaria macrospilus

Apogon atricaudus
Apogon dovii
Apogon pacificus
Arothron hispidus
Bathygobius ramosus
Cosmocampus arctus arctus
Cantherhines dumerilii
Caulolatilus princeps
Chilomycterus affinis
Coralliozetus rosenblatti

Doryrhamphus excisus excisus

Elacatinus digueti
Elacatinus puncticulatus
Stegastes flavilatus
Stegastes redemptus
Gobiosoma chiquita
Gobiosoma nudum
Gobulus hancocki
Gymneleotris seminuda
Hippocampus ingens
Holacanthus clarionensis
Hypsypops rubicundus

Kuhlia mugil

Labrisomus multiporosus Labrisomus xanti

Lactoria diaphana
Lythrypnus pulchellus
Malacoctenus gigas
Medialuna californiensis
Mnierpes macrocephalus
Mulloidichthys dentatus
Myripristis leiognathus

Ogilbia ventralis

Ophioblennius steindachneri

Opistognathus scops Paraclinus mexicanus Parapsettus panamensis Plagiotremus azaleus

Heteropriacanthus cruentatus

Pristigenys serrula

Pycnomma semisquamatum Uropterygius polystictus

Scuticaria tigrina

Malacoctenus tetranemus Ostracion meleagris

Small demersal fish

Albula vulpes

Bajacalifornia burragei Phthanophaneron harveyi Antennarius avalonis Antennatus strigatus Notarius planiceps

Hexanematichthys platypogon

Bagre pinnimaculatus Notarius troschelii Aulopus bajacali Porichthys analis Porichthys ephippiatus

TABLE A.5. Continued.

Porichthys mimeticus Strongylura exilis Platybelone argalus pterura Tylosurus pacificus

Tylosurus crocodilus fodiator Hypsoblennius gentilis Hypsoblennius jenkinsi Hypsoblennius digueti Bregmaceros bathymaster Cataetyx rubrirostris Carapus dubius Echiodon exsilium Centropomus nigrescens Centropomus medius Centropomus viridis

Centropomus robalito Ekemblemaria myersi Emblemaria hypacanthus Emblemaria walkeri

Stathmonotus sinuscalifornici

Chaenopsis coheni Coralliozetus boehlkei Dactyloscopus pectoralis Gillellus semicinctus Dactylagnus mundus Dactyloscopus byersi Gillellus ornatus

Myxodagnus macrognathus Chilomycterus reticulatus Zalembius rosaceus Fistularia corneta

Rimicola eigenmanni Rimicola dimorpha Tomicodon eos Tomicodon boehlkei Gobiesox pinniger Gobiesox schultzi Tomicodon myersi Rimicola sila

Aruma histrio

Barbulifer mexicanus Barbulifer pantherinus Bollmannia stigmatura Bollmannia macropoma Bollmannia longipinnis Bollmannia ocellata Bollmannia pawneea Evermannia longipinnis

Gillichthys seta Elacatinus limbaughi Ilypnus luculentus Ophiodon elongatus Oxylebius pictus

Zaniolepis frenata Sargocentron suborbitalis

Exerpes asper

Paraclinus tanygnathus Starksia spinipenis Cryptotrema seftoni Labrisomus wigginsi Paraclinus beebei Starksia cremnobates Paraliparis ulochir Psednos pallidus Psednos griseus

Caelorinchus scaphopsis Neoconger vermiformis Gymnothorax mordax Anarchias galapagensis Gymnothorax equatorialis Gymnothorax verrilli Gymnothorax eurygnathos Eptatretus mcconnaugheyi

Eptatretus sinus Barbantus curvifrons Mentodus eubranchus Synodus sechurae

Lagocephalus lagocephalus Sphoeroides sechurae Sphoeroides lispus Lycenchelys folletti Axoclinus nigricaudus Bollmannia marginalis Calamus brachysomus Caulolatilus hubbsi

Chaenopsis alepidota alepidota

Chriolepis minutillus Chriolepis zebra Coryphopterus urospilus Crocodilichthys gracilis Ctenogobius sagittula Dactyloscopus lunaticus Enypnias seminudus Gillichthys mirabilis Girella simplicidens

Gobiesox papillifer

Gobulus crescentalis Kathetostoma averruncus Kyphosus analogus Labrisomus striatus Lythrypnus dalli Malacoctenus ebisui Malacoctenus hubbsi Malacoctenus margaritae Malacoctenus zacae Malacoctenus zonifer Microgobius cyclolepis Myxodagnus opercularis Opistognathus punctatus Opistognathus rhomaleus

Paraclinus sini Pherallodiscus funebris Quietula y-cauda Sectator ocyurus Sphoeroides lobatus Starksia grammilaga Starksia hoesei Synodus scituliceps

Pacific angel shark Squatina californica

Small migratory sharks

Mustelus dorsalis Mustelus henlei Mustelus lunulatus Mustelus californicus

Large pelagic sharks

Alopias superciliosus Carcharhinus limbatus Carcharhinus porosus Ginglymostoma cirratum Alopias pelagicus Alopias vulpinus Carcharhinus leucas Carcharhinus obscurus Carcharodon carcharias Isurus oxyrinchus Rhizoprionodon longurio Sphyrna lewini

Sphyrna mokarran Sphyrna zygaena

Guitarfish

Platyrhinoidis triseriata Zapteryx exasperata Rhinobatos leucorhynchus

TABLE A.5. Continued.

Rhinobatos glaucostigma Rhinobatos productus

Skates, rays and sharks

Hydrolagus colliei Dasyatis brevis Gymnura marmorata Gymnura crebripunctata Heterodontus francisci Heterodontus mexicanus Aetobatus narinari Myliobatis californica Mobula thurstoni Narcine entemedor Diplobatis ommata

Cephaloscyllium ventriosum

Apristurus kampae

Triakis semifasciata Urobatis halleri Urotrygon chilensis Urobatis concentricus Urobatis maculatus Urotrygon rogersi Dasyatis dipterura Myliobatis longirostris Raja equatorialis Rhinoptera steindachneri

Flatfish

Achirus mazatlanus Achirus scutum Trinectes fonsecensis Bothus constellatus Bothus leopardinus Perissias taeniopterus Symphurus atricaudus Symphurus atramentatus Symphurus callopterus Symphurus chabanaudi Symphurus fasciolaris Symphurus gorgonae Symphurus leei Symphurus melanurus Symphurus oligomerus Symphurus prolatinaris Symphurus williamsi Symphurus elongatus Paralichthys aestuarius Citharichthys mariajorisae

Ancylopsetta dendritica

Citharichthys fragilis Citharichthys stigmaeus Citharichthys xanthostigma

Hippoglossina bollmani Hippoglossina stomata Paralichthys californicus Paralichthys woolmani Pleuronichthys ocellatus

Pleuronichthys verticalis

Syacium ovale *Xystreurys liolepis*

Etropus crossotus

Mojarra

Gerres cinereus Diapterus peruvianus Eugerres axillaris Eucinostomus currani Eucinostomus entomelas Diapterus aureolus

Scorpionfish

Pontinus sierra Scorpaena sonorae Bellator xenisma Prionotus stephanophrys

Bellator loxias Prionotus albirostris Prionotus birostratus Prionotus ruscarius Scorpaena guttata Scorpaena histrio Scorpaena mystes Scorpaena plumieri Scorpaena russula Scorpaenodes xyris

Lanternfish and deep

Benthosema panamense Lampanyctus parvicauda Zalieutes elater Dibranchus hystrix Dibranchus spinosus Chauliodus macouni Stomias atriventer Bathophilus filifer

Triphoturus mexicanus

Totoaba

Totoaba macdonaldi

Large pelagics

Naucrates ductor

Caranx caballus Caranx caninus Carangoides otrynter Caranx sexfasciatus Caranx vinctus

Chloroscombrus orqueta Decapterus macrosoma Hemicaranx leucurus Hemicaranx zelotes Trachinotus kennedyi Trachinotus paitensis Trachinotus rhodopus Gnathanodon speciosus Coryphaena hippurus Remora remora Elops affinis

Lampris guttatus Mola mola Ranzania laevis Physiculus nematopus Rhincodon typus

Oncorhynchus tshawytscha Katsuwonus pelamis Auxis rochei eudorax Auxis thazard brachydorax

Sphyraena lucasana Peprilus ovatus Peprilus snyderi Xiphias gladius Euthynnus lineatus Oligoplites altus

Sarda chiliensis chiliensis

Seriola lalandi Seriola rivoliana Sphyraena ensis Elagatis bipinnulata

Mackerel

Trachurus symmetricus Gempylus serpens Scomberomorus concolor Scomber japonicus Scomberomorus sierra

Hake

Merluccius angustimanus Merluccius hernandezi

Small pelagics

Argentina sialis Atherinops affinis

TABLE A.5. Continued.

Colpichthys regis Atherinella eriarcha Atherinella nepenthe Colpichthys hubbsi Leuresthes sardina Bathylagus stilbius Sardinops sagax Harengula thrissina Opisthonema libertate

Lile stolifera

Pliosteostoma lutipinnis Opisthopterus dovi Anchoa curta Anchoa walkeri

Anchovia macrolepidota Cetengraulis mysticetus Anchoa mundeoloides Anchoa mundeola Anchoa ischana Anchoa exigua Anchoa nasus Anchoa argentivittata

Anchoa helleri

Cheilopogon pinnatibarbatus Cypselurus callopterus

Mysticeti

Balaenoptera edeni Balaenoptera physalus Balaenoptera musculus Eschrichtius robustus Megaptera novaeangliae

Odontocetae

Delphinus capensis Delphinus delphis Tursiops truncatus

Lagenorhynchus obliquidens

Berardius bairdii

Physeter macrocephalus Globicephala macrorhynchus Pseudorca crassidens

Mesoplodon peruvianus

Vaquita

Phocoena sinus

Pinnipeds

Mirounga angustirostris Zalophus californianus

Reef associated turtles

Caretta caretta

Eretmochelys imbricata

Chelonia mydas

Sea birds

Sula nebouxii Sula leucogaster Larus heermanni Larus californicus

Larus livens Thalasseus elegans Thalasseus maximus Sterna forsteri

Oceanodroma melania Halocyptena microsoma Phalacrocorax penicillatus Phalacrocorax auritus Fregata magnificens Haematopus palliatus

Egretta thula Egretta rufescens Ardea alba Ardea herodias Numenius phaeopus Calidris minutilla Arenaria melanocephala Pelecanus occidentalis Puffinus griseus Puffinus carneipes Podiceps nigrocollis

Squid

Dosidicus gigas Argonauta argo Histioteuthis hoylei Loliolopsis diomedeae Ommastrephes bartramii Pterygiotheuthis giardigiardi Sthenoteuthis oualaniensis Thysanoteuthis rhombus

Synthliboramphus craveri

Blue crab

Callinectes arcuatus Callinectes toxotes Scallops and penshells Spondylus calcifer Spondylus princeps Pteria sterna

Pinna rugosa Atrina tuberculosa

Callinectes bellicosus

Pinctada mazatlanica

Penaeid shrimp

Penaeus aztecus Penaeus duorarum Penaeus setiferus Sicyonia penicillata

Lobster

Panulirus interruptus Panulirus inflatus Panulirus gracilis Crabs and lobsters

Callinectes spp.a Panulirus spp.a Carnivorous macrobenthos

Class Polyplacophora

Class Asteroidea Class Ophiuroidea

Class Cephalopoda, order Octopoda Class Gastropoda, order Nudibranchia

Sea cucumbers Class Holothuroidea

Bivalves

Class Pelecypoda

Snails

Class Gastropoda

^aSpecies not included elsewhere.

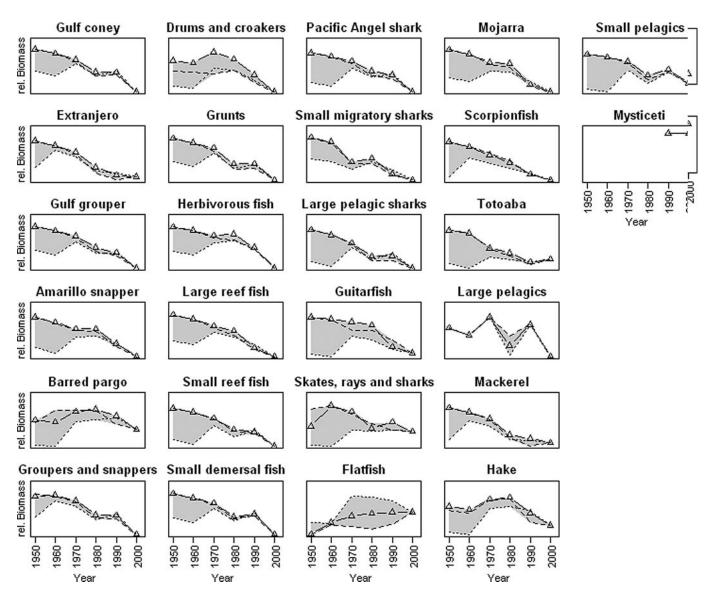


FIGURE A.1. Abundance of functional groups predicted by the fuzzy logic algorithm based on abundance indicators only (see text). Triangles show the results when the responses from all fishers are included in the algorithm, dotted lines the results when only young (below the median age of 43) fishers' responses are included, dashed lines the results when only older fishers' responses are included. The shaded areas show the total potential range (minimum–maximum) of all series. Trends have been scaled to agree in the year 2000. Species groups that had at least eight respondents are shown.