# Parameterization of an Ecosystem Model and Application for Assessing the Utility of Gulf of Mexico Pelagic Longline Spatial Closures 

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## UNIVERSITY OF MIAMI

 FOR ASSESSING THE UTILITY OF GULF OF MEXICO PELAGIC LONGLINE SPATIAL CLOSURESBy

Holly A. Perryman

## A DISSERTATION

Submitted to the Faculty of the University of Miami in partial fulfillment of the requirements for the degree of Doctor of Philosophy

Coral Gables, Florida
May 2017
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## UNIVERSITY OF MIAMI

A dissertation submitted in partial fulfillment of the requirements for the degree of

Doctor of Philosophy

# PARAMETERIZATION OF AN ECOSYSTEM MODEL AND APPLICATION FOR ASSESSING THE UTILITY OF GULF OF MEXICO PELAGIC LONGLINE SPATIAL CLOSURES 

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Parameterization of an Ecosystem Model
and Application for Assessing the
Utility of Gulf of Mexico Pelagic Longline
Spatial Closures
(Ph.D., Marine Biology and Fisheries)
(May 2017)

Abstract of a dissertation at the University of Miami.

Dissertation supervised by Professor Elizabeth Babcock.
No. of pages in text. (329)

Many highly migratory predator stocks that occupy the Gulf of Mexico are at risk, and the collapse of stocks could harm fisheries and ecosystems. Two pelagic longline spatial closures within the pelagic waters of the Gulf of Mexico have been established to protect pelagic species. In 2000, a permanent closure was established around DeSoto Canyon, with the management objectives of reducing catch and rebuilding biomass of bycatch and incidental catch species while minimizing impact to catch of target species. In 2015, a seasonal closure was established off the Louisiana shelf (Spring Closure), with the management objectives of reducing catch and rebuilding biomass of bluefin tuna (Thunnus thynnus). Pelagic spatial closures are relatively untested management tools. Science-driven analysis, including the investigation of ecosystem impacts through mathematical modeling, is necessary to address their utility. This dissertation presents research used to parameterize an ecosystem model, Atlantis, for the Gulf of Mexico marine ecosystem, followed by a study that used the Gulf of Mexico Atlantis model to conduct a policy exploration of the utility of Gulf of Mexico pelagic longline spatial closures.

Chapter 2 described the collection of Gulf of Mexico historical, species-specific landings data for the calibration of the Gulf of Mexico Atlantis model, and invest-
igated areas of uncertainty and bias, focusing on outputs from the Gulf of Mexico Atlantis model and landings-based indicators, due to unidentified landings and lack of data. U.S. landings not identified to species did not appear to bias landings-based indicators, nor does the aggregation of landings into Gulf of Mexico Atlantis functional groups. Chapter 3 described Gulf-wide spatial distributions of pelagic predatory functional groups. Distributions were estimated with generalized additive models fitted with U.S. bottom longline survey catch data (coastal models), and U.S. pelagic longline commercial catch data (pelagic models). This work advanced our knowledge on the correlations between the spatial distribution of pelagic predators within the Gulf of Mexico and the environment, and improved upon the spatial distributions previously used for the Gulf of Mexico Atlantis model. Finally, Chapter 4 described a policy exploration assessing if current pelagic longline spatial closures within the Gulf of Mexico, DeSoto Canyon and Spring Closure, could meet management objectives and evaluated possible ecosystem impacts. DeSoto Canyon was more successful at achieving management objectives and had more influence to ecosystem performance metrics than Spring Closure. Closures reduced Gulf-wide catches of bycatch and incidental groups with little reduction to catches of target groups. Rebuilding biomass of particular stocks may require additional reductions in fishing mortality.

The Atlantis framework allowed for the detailed, spatially-explicit representation of biota, fleets and spatial closures, and provided a means to explore broad-scale ecosystem impacts. This dissertation found that pelagic spatial closures could be viable means to achieve management objectives for protecting highly mobile pelagic predators from fishing pressure.

This work is dedicated to the pursuit to sustainable oceans.

## Acknowledgements

This research was funded by NOAA/SEAGRANT agreement number NA11OAR4170185 and the Cooperative Institute for Marine and Atmospheric Studies (CIMAS), a Cooperative Institute of the University of Miami and the National Oceanic and Atmospheric Administration (NOAA), cooperative agreement number NA10OAR4320143. Data were provided from: NOAA Fisheries, NOAA's Southeast Fisheries Science Center (SEFSC), NOAA's National Centers for Environmental Information (NCEI), the Texas Parks \& Wildlife Department (TPWD), the Comisión Nacional de Acuacultura y Pesca (CONAPESCA), Food and Agriculture Organization (FAO), the International Commission for the Conservation of Atlantic Tunas (ICCAT), the Archiving, Validation and Interpretation of Satellite Oceanographic (AVISO), and the National Aeronautics and Space Administration (NASA) - Moderate Resolution Imaging Spectroradiometer (MODIS).

I would like to thank my committee: Dr. Elizabeth Babcock, Dr. Cameron Ainsworth, Dr. David Die, Dr. Donald Olson, Dr. Joseph Serafy, and Dr. Michael Schirripa. Their constructive feedback strengthened this research and helped me become a better scientist. I am especially thankful for my adviser, Dr. Elizabeth Babcock. She provided me with so much support and encouragement, while also giving me the freedom to be an independent researcher. I am so grateful to work with such a wonderful group of scientists, and look forward to continue to work with them in the future.

The development of the Gulf of Mexico Integrated Ecosystem Assessment, and the Gulf of Mexico Atlantis Model were possible due to the network of researchers involved. I am thankful to colleagues at CISRO (including Dr. Elizabeth Fulton, and Dr. Bec Gorton), NOAA's NWFSC (including Dr. Isaac Kaplan, and Dr. Hem Nalini Morzaria Luna), NOAA's NEFSC (including Dr. Jason Link and Dr. Gavin Fay),

NOAA's SEFSC (including Dr. Mandy Karnauskas and Dr. Skyler Sagarese), the University of South Florida (including Micheal Drexler, Dr. Michelle Masi, and Lindsey Dornberger), CIMAS (including Dr. Arnaud Gruss), and all the others involved in these projects. I also want to thank the Atlantis Modeling Community, as well as all of the individuals who assisted me with data or data processing with ArcGIS, including Dr. Maria Estevanez, Dr. David Gloeckner, Dr. Lawrence Beerkircher, Dr. Greta Wells, Dr. Charles Weber, Dr. Lori Hale, Dr. Vivan Matter, Dr. Laura Elena Vidal, Dr. Arietta Venizelos, and Dr. Mark Fisher.

I am thankful to the faculty and staff of RSMAS, especially the librarians Annie Campbell and Angela Clark-Hughes. I am thankful to my lab (Dr. William Harford, Mathew Nuttall, La Treese Denson, and Halie O'Farrell) as well as the other fisheries labs, including the Die lab (Dr. Elizabeth Councill, Dr. Francesca Forrestal, Dr. Michelle Sculley, and Lily Hoenig), the Ehrhardt lab (Bruce Pohlot, Julie Brown, and Dr. Mark Fitchett), and the Ault lab (Molly Stevens, Christine Harvey, and Dr. Nathan Vaughan). I am grateful to all of the RSMAS students that I have worked with over the years, especially my cohort - for all their support and encouragement from the very beginning. I am especially thankful to Sharein El-Tourky.

None of this would have been possible without the love and support from my friends and family, most of which reside within the Pacific northwest and my home state: Montana. They gave me the strength and confidence to keep going. I am especially grateful to Margaret Perryman and my parents, Jim and Debbie Perryman. Words can not describe how much I love them, how thankful I am to them, and blessed I am that they are a part of my world. Lastly, I would like to thank my fur-baby, Muchacho, for always reminding me that sometimes all you need is to snuggle.

Holly A. Perryman

## University of Miami

May 2017

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## CHAPTER 1

## Introduction

The Gulf of Mexico is a large marine ecosystem bordered by the United States, Mexico and Cuba. Due to the Gulf's complex network of habitats, the ecosystem supports a high level of biological diversity: from microbial communities to highly migratory predators (e.g., sharks, tunas, and billfish). Highly migratory predators are particularly common in the Gulf's pelagic environment with its highly complex physical dynamics consisting of strong currents and eddy networks.

As the Yucatán Current moves through the Yucatán Channel, features are constricted increasing surface water flow as it moves into the Gulf of Mexico to become the Loop Current (Badan et al., 2005). The Gulf's topology causes the current to loop clockwise before exiting through the Straits of Florida. The penetration of the Loop Current into the Gulf varies, but it eventually becomes great enough to produce large, anticyclonic rings known as Loop Current eddies (Leben, 2005). All of these physical forces generate a Gulf-wide network of fronts and eddies (Wiseman et al., 1999; Oey et al., 2005), which create favorable foraging and/or breeding environments for pelagic organisms by upwelling nutrients as well as retaining and concentrating
particles (Olson et al., 1994; Bakun, 1996; Wiseman et al., 1999; Bakun and Broad, 2003).

Gulf fisheries contribute significantly to the economies of the surrounding countries. Coastal communities in particular depend heavily on the fisheries sector. Highly mobile predators are targeted with hook and line gears; either a vertical line consisting of no more than two hooks (handlines), or a horizontal mainline consisting of many hooks (longlines). U.S. commercial handliners harvest all across the Gulf shelf retaining reef fish (e.g., groupers and snappers) and pelagic fish (e.g., tunas and jacks). Longline operations consist of bottom longliners, which set hooks on or near the sea bottom, and pelagic longliners, which set hooks within the water column. U.S. commercial bottom longliners operate along the shelf and the start of the slope catching reef-based benthic species (e.g., groupers) and some highly migratory predators (i.e., sharks). U.S. commercial pelagic longliners operate in the open ocean targeting highly migratory species (e.g., tunas, swordfish and dolphinfish). Landings from U.S. commercial handline and U.S. commercial bottom longline are mostly reported in Florida, while landings from U.S. commercial pelagic longline are mostly reported in Louisiana (National Oceanic and Atmospheric Administration, 2012a). U.S. recreational handlines (i.e., tournaments, for-hire charters, and personal vessel activities) retain many different organisms but mostly target reef and pelagic fish (i.e., groupers and billfish). Recreational fishing plays an important role in the biological dynamics and coastal economy (National Oceanic and Atmospheric Administration, 2012b; Adams et al., 2004), because for some stocks recreational landings can match or surpass commercial landings.

Pelagic predators are particularly vulnerable to overfishing. Some pelagic organisms tend to be found in dense schools as they aggregate around patches of productivity in an otherwise oligotrophic environment. Advances in knowledge and technology have made it easier for fishers to locate fish schools. Thus, it is easier to locate and target large portions of the stock. Some pelagic predators tend to have slow-growing life history, meaning it can take several years for organisms to become sexually mature, which means juveniles can be subjected to fishing pressure before having an opportunity to reproduce. Because of these characteristics, and historically high fishing pressure which some species continue to experience, the sustainability of many highly migratory predator stocks are at risk. This includes some large sharks (Stevens et al., 2000; Baum et al., 2003b; Baum and Myers, 2004; Baum et al., 2005; Burgess et al., 2005; de Mutsert et al., 2008; Baum and Blanchard, 2010), Atlantic bluefin tuna, Thunnus thynnus, (Fromentin and Powers, 2005; ICCAT, 2014b), Atlantic marlins, Makaira nigricans and Kajikia albidus, (Peel et al., 2003; ICCAT, 2011, 2012), and sailfish, Istiophorus albicans (ICCAT, 2016c).

Shepherd and Myers (2005) found that large coastal sharks appear to cause strong top-down effects and their removal has lead to changes in community structure in the northern Gulf of Mexico. Thus, not only would the collapse of highly migratory stocks be devastating for local fisheries and economies, but research across terrestrial and marine ecosystems suggest that the removal of top predators could alter the structure and function of a marine ecosystem. This includes opening a niche which could be filled by organisms that are potentially harmful to the ecosystem (Parsons, 1992; Whitfield et al., 2007), reducing carbon flow to the benthic community (Parsons, 1992), or causing a trophic cascade (e.g., Parsons, 1992; Terborgh et al., 2001;

Heithaus et al., 2008; Baum and Worm, 2009; Casini et al., 2009; Bornatowski et al., 2014). A trophic cascade occurs when the removal of apex predators releases their prey groups (mesoconsumers) from predation, causing increased predation on the prey of mesoconsumers (resource species) (Heithaus et al., 2008). Trophic cascades may have negative impacts on an ecosystem, such as reduced fisheries due to the increase in natural mortality on resource species (Myers et al., 2007), a loss in goods and services due to shifts in underlining processes (Bakun and Weeks, 2006), and reducing ecosystem resistance and resilience (Britten et al., 2014).

Management of pelagic predatory stocks is both a domestic and international effort as these species are highly mobile. The International Commission for the Conservation of Atlantic Tunas (ICCAT) is an inter-governmental fishery organization responsible for the conservation of tunas and tuna-like species in the Atlantic Ocean and its adjacent seas. In the United States, the National Oceanic and Atmospheric Administration (NOAA) through the Highly Migratory Species Devision (HMSD) has primary authority for developing and implementing Fishery Management Plans (FMPs) for highly mobile species (HMS) in Atlantic federal waters, including the Gulf of Mexico. Such FMPs have enacted various input and output controls to ensure the ecological sustainability of pelagic predators (National Oceanic and Atmospheric Administration, 2016a). This includes establishing two pelagic longline spatial closures within the pelagic waters of the Gulf of Mexico. In 2000, a permanent pelagic longline spatial closure was established around the northern West Florida Slope (DeSoto Canyon) to reduce the interaction between non-targeted pelagic fish and longline fisheries. DeSoto Canyon is an area many pelagic predators frequent due to the increased productivity generated by oceanographic characteristics. In 2015,
a seasonal pelagic longline spatial closure was established off the Louisiana shelf to reduce the interactions between bluefin tuna and longline fisheries. This area experiences an increase in bluefin tuna abundance during the spring because it is part of the spawning grounds of the western stock.

Fishery spatial closures are a type of marine protected area (MPA) within which fishing is limited and/or prohibited. MPAs are a tool for ocean conservation (Agardy, 1997). They can protect marine biodiversity by conserving habitat and landscape (Gray, 1997), as well as areas of connectivity (Almany et al., 2009). In addition, MPAs can provide protection to essential habitats and species of concern by protecting areas of aggregation, such as spawning areas, foraging areas, nurseries, and migration stopovers (Norse, 1993). Spatial closures can benefit fisheries by providing biomass through spillover (e.g., McClanahan and Mangi, 2000; Kelly et al., 2002; Guidetti, 2007; Januchowski-Hartley et al., 2013), and increase the size of individuals (e.g., Babcock et al., 1999; Lester et al., 2009).

Much of the current work on MPAs focuses on coastal environments and sedentary organisms because it was originally thought that MPAs would provide little benefit to pelagic predators due to their high mobility and weak site fidelity (Roberts, 1997; Boersma and Parrish, 1999). However, Hyrenbach et al. (2000) argued that pelagic closures could be feasible tools for protecting highly migratory predators since they tend to aggregate around predictable oceanographic features. The advancing knowledge in life histories of pelagic predators, oceanography, and fisheries science suggest that pelagic MPAs have the potential to be viable management tools for protecting pelagic organisms (Game et al., 2009). MPAs for the conservation of pelagic fish are now being recommended by management agencies and stakeholders (Musick et al.,

2000a,b; ICCAT, 2007, 2009, 2010, 2014a; Highly Migratory Species Division, 2008). However, considering that there is a lack of empirical understanding of the direct and indirect impacts of pelagic MPAs, and that often MPAs can fail to meet management objectives (Jameson et al., 2002), it is imperative that science-driven analysis, including the investigation of ecosystem impacts through mathematical modeling, is done to address the utility of pelagic MPAs (Kaplan et al., 2010; Game et al., 2010; Grüss, 2014).

Ecosystem mathematical models are being developed for the Gulf of Mexico Integrated Ecosystem Assessment (IEA) program (Schirripa et al., 2013; Samhouri et al., 2014). An IEA is a framework to guide the process of synthesizing and analyzing relevant scientific information supporting Ecosystem-Based Fisheries Management (EBFM) (National Marine Fisheries Service, 1998, 2012; Levin et al., 2009; Foley et al., 2013). There has been a movement towards EBFM over the last few decades, under which scientists and managers aim to manage fisheries in an ecosystem context rather than a single-species context (Ecosystem Principles Advisory Panel, 1999; Pomeroy et al., 2010). One of the primary purposes of the Gulf of Mexico IEA is to manage the Gulf of Mexico from a broader perspective (e.g., Grüss et al., 2016b). A key component to an IEA is using ecosystem models to evaluate how different management strategies influence the status of indicators. One of the ecosystem models being developed for the Gulf IEA is Atlantis.

Atlantis is a biogeochemical and biophysical simulation framework (Fulton et al., 2004c,b; Fulton, 2010; Fulton et al., 2011). It models the turnover of chemical substances through the biotic and abiotic compartments of an ecosystem, and there are detailed routines for coupling the biological and physical components. Atlantis is an
"end-to-end" model, meaning it represents biota from bacteria up to top predators. Biota can be represented as age-structured groups, or biomass pools. There is a detailed fisheries exploitation routine that allows the simulation of individual fleets, as well as routines for simulating a range of management measures, including fishery spatial closures. Interactions between species and fisheries are spatially explicit, and the spatial domain is composed of a 3-dimensional polygon network that reflects key geographic features, habitats, and essential management jurisdictions. The Atlantis framework has been used to investigate the spatial management of fisheries, including the use of spatial closures, e.g. Ainsworth et al. (2012); Kaplan and Leonard (2012); Morzaria-Luna et al. (2013). Although Atlantis is argued to be one of the best operating models for ecosystem simulation (Plagányi, 2007), one of the disadvantages is that Atlantis requires more data than other ecosystem models, including historical landings data, and seasonal spatial distributions of simulated functional groups (i.e., groups of species with similar life histories and ecosystem function).

One method for parameterizing an Atlantis model for forecasting involves first calibrating a historical Atlantis model with landings time series data. Values of dynamic parameters in the calibrated historical model are transferred to a present day model for forward simulations. A critical component of this methodology is the collection of historical, species-specific landings (organized by gear, season, and state if possible). Data need to be aggregated based on the Atlantis-defined functional groups, which could incorporate bias into the historical landings trends. This would impact the calibration of the Gulf of Mexico Atlantis model and forecasting simulation studies. Thus, Chapter 2 of this dissertation describes the collection of Gulf of Mexico historical, species-specific landings data for the calibration of the Gulf of Mexico Atlantis
model, and investigates areas of uncertainty and bias due to unidentified landings and lack of data. This investigation provides a detailed picture of the historical development of fisheries in the Gulf, and is informative for the Gulf of Mexico Atlantis model, as well as other ecosystem models and metrics for the Gulf of Mexico.

To investigate the utility of pelagic fishery closures of the Gulf of Mexico, it is imperative that the forecasting Gulf of Mexico Atlantis model is parameterized with reasonable seasonal spatial distributions for pelagic functional groups. Spatial distributions can be inferred from predictive statistical models (Guisan and Zimmermann, 2000; Austin, 2002, 2007; Elith and Leathwick, 2009). Statistical models for predicting the spatial abundance of marine fishes depends on the fundamental relationship between catch rate (catch per unit effort) and density, and the shortcomings of using catch rate as an index of abundance have been long-studied in fisheries literature (e.g., Gulland, 1956; Beverton and Holt, 1957; Robson, 1966; Honma, 1973; Seber, 1982; Cooke and Beddington, 1984; Beddington and Cooke, 1984; Hilborn et al., 1992; Harley et al., 2001). However, advances in statistical methodologies (e.g., generalized linear modeling) address many of the shortcomings of fisheries data (Maunder and Punt, 2004).

In Chapter 3 of this dissertation, generalized additive models (GAMs) (Hastie and Tibshirani, 1986, 1990) were developed to describe the spatial distribution of pelagic functional groups within the Gulf of Mexico. Species-specific catch records were grouped according to pelagic functional groups identified for the Gulf of Mexico Atlantis model. Two types of GAMs were fitted: coastal (covering areas $0-200 \mathrm{~m}$ deep), and pelagic (covering areas greater than 200 m deep). Coastal models were fitted using NOAA's Bottom Longline Survey data, and pelagic models fitted using

NOAA's Pelagic Longline Observer Program data. A delta approach was followed to account for the zero-inflated catch data. This consisted of fitting a Bernoulli GAM with binomial data, and a Gamma GAM with zero-truncated catch rate data. Model descriptors (independent variables) considered for coastal models included year, sea bottom depth, altimetry, minimum distance from a front, as well as sea surface and sea bottom temperature, dissolved oxygen, oxygen saturation, and salinity. Descriptors considered for pelagic models included season, year, sea bottom depth, altimetry, minimum distance from a front, and sea surface temperature. Fitted models and data series describing seasonal environmental conditions were used to predict Gulfwide seasonal, spatial distributions of pelagic predator groups.

With the Gulf of Mexico Atlantis forecasting model parameterized using historical landings data, and spatial distributions of pelagic predator functional groups generated from the statistical models, it was ready to be used to explore the utility of the Gulf of Mexico pelagic longline spatial closures. Chapter 4 describes a simulation test for investigating i) if Gulf of Mexico pelagic longline fishery spatial closures are likely to achieve management objectives, and ii) potential ecosystem impacts from pelagic longline closures. The Gulf of Mexico Atlantis model was used to simulate scenarios and calculate performance measures (indicators) corresponding to management objectives of the pelagic longline fishery spatial closures, as well as broader ecological objectives. Performance metrics were compared to examine potential long-term impacts of Gulf of Mexico pelagic longline spatial closures.

In summary, this dissertation consists of three components. First, Chapter 2 describes the collection of Gulf of Mexico historical, species-specific landings data for the calibration of the Gulf of Mexico Atlantis model, and investigates areas of
uncertainty and bias due to unidentified landings and lack of data. Next, Chapter 3 describes the development of delta generalized additive models for estimating the Gulf-wide spatial distribution of pelagic predator functional groups as described for the Gulf of Mexico Atlantis model. Lastly, Chapter 4 describes a simulation test using the Gulf of Mexico Atlantis model for investigating whether Gulf of Mexico pelagic longline spatial closures could achieve management objectives, as well as their potential ecosystem impacts. This dissertation advanced our understanding regarding the strengths and weaknesses of some of the data currently available for the Gulf of Mexico IEA. In addition, it advanced our understanding of the drivers and patterns pertaining to spatial distributions of pelagic predators within the Gulf of Mexico. This work provided insight with respect to possible benefits from pelagic longline spatial closures, and improved our understanding and modeling of the Gulf of Mexico ecosystem.

## CHAPTER 2

## Landings Data for Ecosystem Fisheries Science: Lessons Learned from the Gulf of Mexico

### 2.1 Summary

Historical landings data are crucial for ecosystem based fisheries management in that they i) are needed for the calibration of ecosystem modeling tools, and ii) allow for the assessment of landings-based indicators. Such methodologies require landings data on species. Neglecting data not identified to species (ambiguous landings) could potentially bias results. This work considers Gulf of Mexico landings data to discuss potential uncertainties in the development of ecosystem based fisheries management tools, like the Gulf of Mexico Atlantis model, as well as landings-based indicators. Gulf of Mexico landings data (1980-2011) were described for the United States, Mexico, and Cuba. Landings were classified by species, then allocated into functional groups identified for the Gulf of Mexico Atlantis model. U.S. landings, both species-specific and functional group-specific, were used to compute qualitative landings-based indicators relating to stock assessment coverage, and quantitative landings-based indicators relating to system ecology (pelgic:demersal ration, and
mean trophic level). Commercial landings data have meaningful portions not identified to species, especially data from Mexico and Cuba ( $29.2 \%$ and $48.9 \%$, respectively, are unidentified). U.S. recreational data have few ambiguous landings ( $0.4 \%$ are not identified to species), but there is a lot of variation in landings data from MRIP, at least some of which is estimation error. Ambiguous landings did not appear to be adding bias to investigated indicators pertaining to U.S. waters. In addition, the aggregation of landings into Gulf of Mexico Atlantis functional groups do not appear to biasing the computation of trends. Qualitative indicators show that a majority of U.S. commercial landings are of species that are not overfished, but the majority of U.S. recreational landings are of species of unknown overfished status. Although ecosystem based fisheries management of the Gulf of Mexico would benefit from more precise landings, current data is sufficient for the development of ecosystem models.

### 2.2 Motivation

Under ecosystem-based fisheries management (EBFM), scientists and managers aim to manage fisheries in an ecosystem context rather than a single-species context (Ecosystem Principles Advisory Panel, 1999; Link, 2002; Brodziak and Link, 2002; Pikitch et al., 2004; Link, 2010). There has been a shift towards EBFM (Pomeroy et al., 2010) due to the perception that fishing operations have the power to alter the structure and function of marine ecosystems (Marasco et al., 2007), and that healthy ecosystems are needed to sustain fished populations. Hilborn (2011) argues that even if single-species management was executed well, EBFM is still necessary because pure single-species management does not consider impacts on non-target species, trophic interactions among species, and habitat-destroying fishing practices.

Integrated Ecosystem Assessment (IEA) is an assessment methodology that supports EBFM (Foley et al., 2013). Originally described by Levin et al. (2008, 2009), an IEA is a cyclic process made up of five steps. First, ecosystem objectives and threats are identified. An important part of EBFM is the capability to monitor progress toward objectives, and this is achieved with indicators (Pikitch et al., 2004). Thus, the second step of an IEA is to identify and validate indicators for assessing the state of the ecosystem. EBFM requires a suite of indicators that provide insight into the state of the ecosystem, particularly in relation to the impact of fishing (Dale and Beyeler, 2001; Rochet and Trenkel, 2003; Fulton et al., 2005; Shin and Shannon, 2010; Powers and Monk, 2010; Link et al., 2010b). This includes indicators based on landings data, which can provide information regarding changes in the assessment and fisheries management coverage of the system (e.g., Piet et al., 2010; Gascuel et al., 2012; Karnauskas et al., 2013), as well as the system's fisheries ecology (e.g., Rochet and Trenkel, 2003; Fulton et al., 2005; Shin et al., 2010). Next, there is an evaluation of the risk posed by human activities and natural processes. This is followed by the use of ecosystem models to evaluate how different management strategies influence the status of indicators. This is a process referred to as a Management Strategy Evaluation (Smith, 1994; Sainsbury, 1998; Cooke, 1999; Sainsbury et al., 2000; Butterworth et al., 2010; Punt et al., 2016). Lastly, ecosystem indicators are monitored and assessed to determine the effectiveness of management strategies. Ideally, this process is repeated to support adaptive management (Dickey-Collas, 2014) and monitoring (Uychiaoco et al., 2005).

In the USA, the National Oceanic and Atmospheric Administration (NOAA) has been developing IEAs for marine ecosystems (Samhouri et al., 2014), including the

Gulf of Mexico (Karnauskas et al., 2013; Schirripa et al., 2013). The Gulf of Mexico is a large marine ecosystem that supports the livelihoods of people in coastal communities of the United States, Mexico, and Cuba (Adams et al., 2009; Yoskowitz, 2009), as well as the many populations of marine fauna (Giattina and Altsman, 1999; Landry Jr and Costa, 1999; Mullin and J, 1999). Some of the stressors facing the Gulf include habitat modification and unsustainable exploitation of living resources (Yáñez-Arancibia and Day, 2004), both of which threaten the sustainability of fisheries stocks (Coleman et al., 1996; Baum and Myers, 2004; Ault et al., 2005; Heithaus et al., 2007a; MacKenzie et al., 2009; Beck et al., 2011). The need for a holistic approach to meet these threats has influenced a movement towards an ecosystem approach to the management of the Gulf of Mexico ecosystem and fisheries (YáñezArancibia and Day, 2004; Nugent and Cantral, 2005; Arreguín-Sánchez et al., 2008; Carollo and Reed, 2010; Day and Yáñez-Arancibia, 2013; Yáñez-Arancibia et al., 2013).

To support the Gulf of Mexico IEA, several ecosystem modeling frameworks are being developed (Schirripa et al., 2013), including Atlantis - a dynamic biogeochemical ecosystem model that simulates physical, chemical, biological, and fisheries components within a three-dimensional spatial domain (Fulton et al., 2004b,c, 2007). Atlantis has been used to investigate ecological indicators for detecting ecosystem impacts due to fisheries, investigate cumulative impacts, explore ecosystem dynamics, and test management approaches (Fulton et al., 2004a, 2005; Link et al., 2010a; Fulton et al., 2011; Kaplan et al., 2012; Ainsworth et al., 2012; Masi et al., 2017). To support the Gulf of Mexico IEA, a model for the entire Gulf of Mexico marine ecosystem was developed using the Atlantis framework (Ainsworth et al., 2015).

Initializing of the Gulf of Mexico Atlantis model for forecasting included calibrating a historical model to fit landings time series (Ainsworth et al., 2015). To do this, landings time series from 1980 to 2011 were collected and categorized into functional groups simulated in the Gulf of Mexico Atlantis model (Perryman et al., 2015). In addition, landings were partitioned across seasons and fishing fleets. Commercially important species often have complete landings profiles, however species that are not commercially important are often grouped into ambiguous categories. This includes records identified to a higher taxonomic classification (e.g., family), or no taxonomic classification (e.g., "unidentified"). Ambiguous landings may be negligible for some species, but other species can have significant portions of their landings not appropriately identified. Excluding ambiguous landings from EBFM tools could bias the computation of landings based indicators, as well as the calibration of ecosystem models (i.e., misrepresent the magnitude of fisheries on stocks).

Landings data pertaining to the Gulf of Mexico were presented and discussed in detail in the Gulf of Mexico Ecosystem Status Report (Karnauskas et al., 2013). Karnauskas et al. focused on identifying trends in Gulf-wide indicators. The following research builds on their findings. This study used Gulf of Mexico landings data that was somewhat different than data presented by Karnauskas et al. (2013) to discuss how data uncertainties, including landings not identified to species, aggregation of landings by Atlantis functional groups, and allocation of landings to season, state, and gear, could bias the Gulf of Mexico Atlantis model. Lastly, landings-based indicators were computed to discuss trends and possible bias on how data were grouped for the purposes of ecosystem modeling, including functional groups, recreational versus commercial fisheries, and seasonal and regional divisions. This study aims to gain
insight into some of the uncertainties concerning landings data available for the Gulf of Mexico for use in ecosystem models.

### 2.3 Methods

### 2.3.1 Landings Data

Landings data from Gulf of Mexico waters are available for U.S. commercial, U.S. recreational, Mexico commercial, and Cuba commercial fleets. The National Oceanic and Atmospheric Administration (NOAA) Fisheries, Fisheries Statistics Division provides summaries of U.S. commercial fisheries landings, in weight (lbs), as annual landings, or annual landings itemized by state, season, or gear (National Oceanic and Atmospheric Administration, 2012a). These landings come from a cooperative State-Federal fishery data collection system that obtains landings data from statemandated trip-tickets (which are filled out at the conclusion of every fishing trip), landing weigh-out reports provided by seafood dealers, federal logbooks of fishery catch and effort, and shipboard / portside interviews. Most states get their landings data from seafood dealers who submit monthly reports of the weight and value of landings by vessel; however, more states are switching to mandatory trip-tickets to gather landings data (National Oceanic and Atmospheric Administration, 2016b). U.S. commercial landings are dominated by menhaden, Brevoortia spp., (Figure 2.1). This analysis excludes U.S. commercial menhaden landings in order to identify underlying trends in the rest of the fisheries (de Mutsert et al., 2008; Karnauskas et al., 2013).

The NOAA Marine Recreational Information Program (MRIP) provided the bulk of the U.S. recreational data (National Oceanic and Atmospheric Administration, 2012b). MRIP is a compilation of regionally-based data collection programs that collects data from a subsample of anglers and captains, which is then expanded to all anglers based on a telephone survey to estimate effort (National Oceanic and Atmospheric Administration, 2014a). Texas is not part of MRIP surveys and instead the state conducts its own survey on recreational landings, which are then provided to NOAA (Gulf of Mexico Fishery Management Council, 2005). Lastly, NOAA's Southeast Fisheries Science Center (SEFSC) Recreational Billfish Survey System (RBS) provided data on recreational billfish tournaments within the Gulf of Mexico (A. Venizelos at NOAA, personal communication, May 8, 2013). The RBS has been collecting data on recreational billfish tournaments in the western North Atlantic, Gulf of Mexico and U.S. territories in the Caribbean since 1972, and is the primary source of U.S. recreational billfish catch and effort statistics (National Oceanic and Atmospheric Administration, 2014b). The Gulf of Mexico Ecosystem Status Report (Karnauskas et al., 2013) did not indicate if the RBS data were considered. Recreational data were extracted in numbers. Originally, MRIP data in weight were extracted but further analysis showed that MRIP data in weight had about half as many records as MRIP data in numbers. The Gulf of Mexico Ecosystem Status Report (Karnauskas et al., 2013) reports MRIP data in weight and does not discuss data in numbers.

Annual reports from the Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA) through the Comisión Nacional de Acuacultura y Pesca (CONAPESCA) provided landings data for Mexican commercial fisheries
(SAGARPA, 2016). Data provided in the reports were collected by the SAGARPA as well as the Órganos Centrales de la Secretaría from the various agencies active in the fisheries sector (SAGARPA, 1980-2011). Data, in weight (kgs), were extracted for the coastal Mexican states of Tamaulipas, Veracruz, Tabasco, Campeche, and Yucatán. Quintana Roo data were not included since landings from the Gulf of Mexico and landings from the Caribbean could not be separated, and all the major fishing ports of Quintana Roo are on the Caribbean coast. Data describing Mexican recreational landings from the Gulf of Mexico were not considered in this study (i.e., assumed to be zero) since information could not be found. These commercial landings are directly from Mexico. Landings considered in the Gulf of Mexico Ecosystem Status Report (Karnauskas et al., 2013) were from FAO.

The Food and Agriculture Organization of the United Nations (FAO) provided landings data for Cuban commercial fisheries (FAO, 2013a). Data from FAO describes total Cuban landings, in weight (tonnes). Claro et al. (2001) provided a regional breakdown (i.e., southeast, southwest, northwest and northeast) of common groups of species identified in Cuban commercial landings (1959-1998). It was assumed that the northwest region represents landings solely from the Gulf of Mexico. Thus, the data provided by Claro et al. (2001) was used to calculate average proportions of Cuban landings that were in the northwest region, which were applied to the FAO data on total Cuban landings to infer Gulf of Mexico landings. Data describing Cuban recreational landings from the Gulf of Mexico were not considered in this study (i.e., assumed to be zero) since information could not be found. The Gulf of Mexico Ecosystem Status Report (Karnauskas et al., 2013) did not discuss landings data from Cuba.

To evaluate the amount of landings associated to ambiguous groups for each region, landings data were categorized by species, genus, family+ (which includes landings identified to family or any higher Taxonomic Classification), or unidentified. This could be easily accomplished for U.S. and Cuban datasets, but many of the identifications used in Mexican data refer to general groups of organisms and not species. Thus, Mexico landings are instead categorized by "taxonomic classification", meaning that they were identified to some taxonomic group such as "snappers" or "large sharks", or "unidentified".

To evaluate whether uncertainties in species identification in the U.S. data varied over state, season, or gear, time series of total landings and fraction ambiguous were generated for each of these classifications. Both NOAA and MRIP provide landings itemized by state (i.e., the state of the port where catch was landed). Seasonal data from NOAA itemized commercial landings by month. Data were aggregated into the four seasons simulated in the Gulf of Mexico Atlantis model (winter, Jan. - Mar.; spring, Apr. - Jun.; summer, Jul. - Sep.; fall, Oct. - Dec). Seasonal data from MRIP itemized landings by six bimonthly intervals. Landings-by-gear data from NOAA were aggregated according to fleets described for the Gulf of Mexico Atlantis model (Appendix A) to simplify results while relating the analysis to EBFM tools. Three miscellaneous gear types could not be directly allocated into a fleet so each were left to stand alone for this analysis: "Combined Gears", "Not Coded", and "Unspecified Gear".

### 2.3.1.1 Functional Group-Specific Landings

Data described in section 2.3.1 were used to construct landings time series for the functional groups of the Gulf of Mexico Atlantis model. First, data were allocated into functional groups. To do this the taxonomic classification of landings were determined using the Integrated Taxonomic Information System (ITIS, 2012), FishBase (Froese and Pauly, 2016), SeaLifeBase (Palomares and Pauly, 2016), Salas et al. (2011), or the Universal Biological Indexer and Organizer (Norton et al., 2013). Speciesspecific data were directly allocated into functional groups, while ambiguous data were split amongst appropriate functional group(s). In some cases, a higher taxonomic level of identification, such as family, was sufficient to determine the appropriate functional group. In other cases, ambiguous landings were allocated to functional groups based on information from the literature, or assumptions made about the species composition. Second, data were converted to tonnes. Commercial landings were recorded by weight but recreational data (numbers) were converted to tonnes using length-weight relationships and the length information included in the datasets. This entire process was described in detail by Perryman et al. (2015).

### 2.3.2 Landings-based Indicators

This study considers two types of landings-based indicators: qualitative stock assessment coverage indicators, and quantitative community indicators. The computation of indicators were restricted to U.S. landings (section 2.3.1) because more data on species identification and status was available in the U.S. Landings-based indicators relating to ecological status were computed with i) landings itemized by season and state, as well as ii) functional group-specific landings constructed for the

Gulf of Mexico Atlantis model (section 2.3.1.1). The former allowed the assessment of indicator trends over finer temporal and spatial scales of the fisheries, and the latter allowed the analysis of impacts to indicators when data are aggregated into functional groups. Indicators are computed separately for commercial and recreational landings as to assess differences in trends between the two sectors. U.S. recreational species-specific landings data were in numbers, while the U.S. commercial speciesspecific landings data, and all functional group-specific landings data were in weight. For details on the species composition of recreational and commercial landings, see Appendix A.

### 2.3.2.1 Stock Assessment Indicators

Karnauskas et al. (2013) found that, for landings of federally managed stocks in the U.S. Gulf of Mexico, the ratio of overfished to not overfished stocks has decreased. To expand on this analysis, I included all landed species, not just those in federal fishery management plans, and evaluated overfished status and jurisdiction of i) number of landed species (combining commercial and recreational data), ii) U.S. commercial landings, and iii) U.S. recreational landings. Information on status was provided by annual Congressional Stock Status (CSS) Reports. Since 1997, NOAA has been submitting reports to Congress describing the state of the nation's marine fisheries and the effectiveness of fisheries management under the Magnuson-Stevens Fishery Conservation and Management Act as amended in 1996 by the Sustainable Fisheries Act (National Marine Fisheries Service, 1998-2012). These CSS Reports indicate the status of federally managed stocks. Species that are managed federally are identified under fishery management plans (FMPs) from the Gulf of Mexico Fish-
ery Management Council (GMFMC). Stocks primarily retained within state waters (to 16.2 km from the coast in Texas and the west coast of Florida, to 5.6 km in the other Gulf of Mexico states) are generally managed by the individual states. The Interjurisdictional Fisheries Program of the Gulf States Marine Fisheries Commission (GSMFC) provides the Gulf States with information and recommendations for interstate FMPs.

First U.S. commercial and U.S. recreational landings data were categorized based on species-specific overfished status in each year's CSS report. A species landings were categorized as unknown if the overfished status was not reported. Then, for species of unknown overfished status, landings were classified according to FMP jurisdiction (i.e., GSMFC, GMFMC, or neither). Ambiguous landings were categorized as unknown overfished status with no FMP jurisdiction, because the overfished status and FMP jurisdiction cannot be determined. The same categorization was made for landings associated with species managed by individual states and not associated with the GSMFC. Spanish mackerel (Scomberomorus maculatus) which has FMPs under both GSMFC and GMFMC, were allocated to the GMFMC.

### 2.3.2.2 Pelagic:Demersal Ratio

Landings pelagic:demersal ratio is the ratio of landings of pelagic organisms to landings of demersal/benthic organisms. To calculate the ratio, information regarding the life history of adults organisms were used to classify landings as pelagic or demersal, then total landings of pelagic species were divided by total landings of demersal species. The pelagic:demersal ratio may be an informative for the ecosystem management of the Gulf of Mexico since the metric is primarily linked to the eu-
trophication (Caddy, 1993; Caddy and Bakun, 1994; Caddy et al., 1998a; Caddy and Garibaldi, 2000; Caddy, 2000; de Leiva Moreno et al., 2000), and the Gulf of Mexico experiences periodic large-scale eutrophication which has meaningful ecosystem impacts (Malakoff, 1998; Rabalais et al., 2002b,a). Karnauskas et al. (2013) discussed the pelagic:demersal ratio with respect to fishery independent trawl survey data, but do no discuss the metric in terms of landings data. These metrics have a different interpretation because the fishery-independent pelagic:demersal ratio is tracking changes in the ecosystem, while the pelagic:demersal ratio of landings can show shifts in fishery targets.

In this study, the pelagic:demersal ratio was computed with U.S. commercial, and U.S. recreational landings time series. This was done for both species-specific and functional group-specific datasets. Species and functional groups were classified as pelagic or demersal using life history information from FishBase (Froese and Pauly, 2016) and SeaLifeBase (Palomares and Pauly, 2016). Due to the configuration of functional groups for the Gulf of Mexico Atlantis model, most groups contained species that were either all pelagic or all demersal. The exceptions were the skates and rays functional group, which was assumed to be demersal, and the large sharks functional group, which was assumed to be pelagic, when calculating functional group specific pelagic:demersal ratio (Appendix A).

### 2.3.2.3 Mean Trophic Level

Landings mean trophic level is the sum of the product of species trophic level and species landings divided by landings summed across all species. This indicator has been proposed as evidence that there has been a gradual transition in landings from
long-lived, high trophic level, piscivorous fish toward short-lived, low trophic level, invertebrates and planktivorous fish caused by sequential depletion of upper trophic level species to lower trophic level species - a phenomenon called "fishing down the foodweb" (Pauly et al., 1998; Pauly and Palomares, 2005; Pauly and Watson, 2005; Pauly et al., 2005). The Convention on Biological Diversity has identified landings mean trophic level as one of eight indicators to be tested to measure progress towards achieving a significant reduction in the current rate of biodiversity loss (Convention on Biological Diversity, 2004). Although, this indicator may be influenced by changes in fleet targeting, advancing harvesting technology, and fisheries management, rather than fisheries impact on an ecosystem (Caddy et al., 1998b; Caddy and Garibaldi, 2000; Essington et al., 2006; de Mutsert et al., 2008; Branch et al., 2010; Powers, 2010; Sethi et al., 2010), it can still be informative with respect to the targets and composition of fisheries (Shin et al., 2010).

Karnauskas et al. (2013) found the average trophic level of both Mexican and U.S. landings has increased since the 1950's. To evaluate whether this conclusion would change when the data were combined into functional groups, or with the additional datasets considered in this study, landings mean trophic level was calculated for U.S. commercial and U.S. recreational species-specific data and functional group-specific data. First, FishBase (Froese and Pauly, 2016) and SeaLifeBase (Palomares and Pauly, 2016) were used to get species-specific estimates of trophic level. If an estimate was not provided, then a value from a similar species of the same genus was assumed. Functional groups in the Gulf of Mexico Atlantis model were assigned trophic level by averaging the corresponding species-specific trophic levels.

### 2.4 Results

### 2.4.1 Landings Data

The amount of ambiguous landings varies across Gulf of Mexico countries (Figure 2.2). A majority of U.S. landings are identified to species. On average $94.8 \%$ of the commercial landings, excluding menhaden, (Figure 2.2 a ), and $95.2 \%$ of the recreational landings (Figure 2.2b). U.S. ambiguous landings mostly consist of records identified to a taxonomic classification higher than species. On average $68.6 \%$ of the commercial ambiguous landings, and $92.1 \%$ of the recreational ambiguous landings. Family is the most common taxonomic classification used other than species. Family is given for on average $57.1 \%$ of the commercial ambiguous landings, and $64.3 \%$ of the recreational ambiguous landings. Generally, the proportion of U.S landings that are of ambiguous groups fluctuates throughout the data series. After 1986, the proportion of U.S landings that are of ambiguous groups has decreased because the landings of ambiguous groups (e.g., miscellaneous finfish, and sharks) decreased while the landings of species-specific groups remained stable (Appendix A). The trend in proportion of U.S. recreational landings that are of ambiguous groups is generally decreasing due to a large decline in ambiguous landings early in the dataset.

Data from Mexico and Cuba have more landings associated to ambiguous groups. On average, $70.8 \%$ of the Mexican commercial landings are identified to taxonomic classifications while $29.2 \%$ are unidentified (Figure 2.2c). A majority of Cuban commercial landings (on average 66.8\%) are of ambiguous groups (Figure 2.2d). Most of the Cuban ambiguous landings (on average 73.0\%) are of unidentified groups. The proportion of Mexican commercial landings identified to a taxonomic classification
increased over time due a gradual decline in unidentified landings (Appendix A). The proportion of Cuban commercial landings of miscellaneous/unidentified groups decreased over time, and the proportion of Cuban commercial landings of groups with a taxonomic classification other than species increased over time (Appendix A).

On average, more U.S. commercial and U.S. recreational landings occur in the summer months (Figure 2.3). Commercial ambiguous landings are more common during the spring, while recreational ambiguous landings are more common in the summer. The percent of commercial landings that are of ambiguous groups has decreased for every season except spring (Figure 2.3c). The percent of recreational landings that are of ambiguous groups has increased for summer months, and decreased for winter months (Figure 2.3d).
U.S. commercial landings (Figure 2.4a) are predominantly landed in Louisiana, Texas, and Florida. Over time, landings from Florida have decreased and landings from Louisiana have increased. U.S. recreational landings (Figure 2.4b) are predominantly from Florida. Over time, the proportion of landings from Florida and Texas have increased while the proportion of landings from Louisiana and Mississippi have decreased. On average, for both commercial and recreational data (Figure 2.4c, 2.4d), Mississippi has a higher fraction of landings that are of ambiguous groups, followed by Florida. However, the higher average of commercial landings from Mississippi is due to a large spike in ambiguous landings in the mid-90's. This was caused by a sudden reporting of unidentified shrimp (see Appendix A). In addition, the higher average of recreational landings from Mississippi is due to a large spike in ambiguous landings in 2010. This was caused by a sudden reporting of Carcharhinidae landings (see Appendix A). Not considering these sudden spikes, Florida has the higher frac-
tion of landings that are of ambiguous groups for both commercial and recreational data.
U.S. commercial landings itemized by gear, which have been aggregated by Gulf of Mexico Atlantis fleets, are highly variable until the late 1990's (Figure 2.5). From 1980 to 2001, on average half of the commercial landings are not identified to a specific gear, thus could not be directly allocated to a fleet identified for the Gulf of Mexico Atlantis model (Figure 2.5a). From 1980 to 1996, over half of the ambiguous landings (55\%) are from gear-types that could not be directly associated to an Atlantis fleet (Figure 2.5b). The percentage of landings that are of ambiguous groups varies amongst U.S. gear-types. Trends for hook-and-line gear-types stabilize and some are generally lower in recent years than in the 1980's and 1990's (Figure 2.5c). Trends for net gears, both those operated within estuaries (Figure 2.5 d ) and those operated within the shelf Figure 2.5e), vary, with some increasing over time. Trends for miscellaneous geartypes are highly variable (Figure 2.5f).

### 2.4.2 Landings-based Indicators

### 2.4.2.1 Stock Assessment Indicators

On average, about half of the federally managed species harvested by U.S. fleets are of an unknown overfished status (Figure 2.6a). In addition, on average $62.6 \%$ of the species harvested by U.S. fleets are not identified in a GMFMC or GSMFC FMP. The number of overfished species, and the number of species of unknown overfished status has decreased over time. Many of the landed species of unknown overfished status are from the U.S. recreational data, so it's possible that this trend is driven by improvements to the MRIP dataset (e.g., improved identification of landings).

The number of not overfished species landed has increased over time. This is possibly driven by improvements and expansion of stock assessments rather than improvements in fisheries sustainability.

On average, most (60.6\%) of U.S. commercial landings (lbs) are of federally assessed stocks declared not overfished (Figure 2.6b), while most (84.0\%) of U.S. recreational landings (numbers) are of species with unknown status (Figure 2.6c). Most of the U.S. commercial landings of unknown status (on average $67.7 \%$ ) correspond to species associated to GSMFC FMPs. A majority of these landings are of blue crab (Callinectes sapidus), eastern oyster (Crassostrea virginica), and striped mullet (Mugil cephalus). Some of these species have been assessed by individual state agencies for part of the Gulf of Mexico, so U.S. commercial landings of species declared not overfished may be larger. U.S. recreational landings of unknown status are mostly (on average $59 \%$ ) of species not associated to either GSMFC or GMFMC FMPs. A majority of these landings consist of scaled sardine (Harengula jaguana), pinfish (Lagodon rhomboides), white grunt (Haemulon plumieri), and Atlantic thread herring (Opisthonema oglinum). Many of these species are used in the bait industry and they may not be at much risk of being overfished because of their short lived and fast growing life history.

Trends from U.S. commercial data and U.S. recreational data should not be compared since commercial data were in weight and recreational data were in numbers. Since the U.S. recreational landings are in numbers, and the data are highly variable, it is difficult to discern the magnitudes of the resulting trends. For instance, U.S. recreational landings of unknown status are mostly smaller bait fish, while not
overfished landings (mostly composed of Spanish mackerel) and overfished landings (mostly composed of red snapper, Lutjanus campechanus) consist of larger finfish.

### 2.4.2.2 Pelagic:Demersal Ratio

Landings pelagic:demersal ratio trends computed from U.S. commercial and U.S. recreational data are shown in Figure 2.7. There are no obvious seasonal trends of the pelagic:demersal ratio for U.S. commercial data (Figure 2.7a) or U.S. recreational data (Figure 2.7d). Seasonal pelagic:demersal ratio from U.S. recreational data are highly variable, so it is difficult to discern statistically meaningful trends. The commercial pelagic:demersal ratio trends decreased for all four seasons, while the recreational pelagic:demersal ratio increased for all six bimonthly intervals. The certainty of recreational trends is questionable since data are highly variable. Landings pelagic:demersal ratio trends differ amongst individual states for both U.S. commercial, and U.S. recreational data (Figure 2.7b and (Figure 2.7e), respectively). For both U.S. commercial, and U.S. recreational data, landings pelagic:demersal ratio is much larger for Florida landings. Florida landings had significant contributions from pelagic finfish groups (e.g., scaled sardine) while landings in the other states tend to be dominated by demersal groups (e.g., seatrout, shrimp, oysters) (Appendix A). Much of the U.S. commercial ambiguous landings consists of groups that are or have the potential to be demersal (e.g. shrimp, shellfish, flatfish, finfish), so ambiguous landings could influence U.S. trends from U.S. commercial data.

The pelagic:demersal ratio computed with functional group-specific landings tends to be similar to that computed with species-specific landings for both U.S. commercial (Figure 2.7c). The recreational pelagic:demersal ratio (Figure 2.7f) increased for both
species-specific, and functional group-specific data up until 2000, after which while the trend from species-specific data continued to increase, the trend from functional group-specific data decreased. The divergence in trends is not due to the aggregation of data into functional groups as most species are assigned to functional groups with the appropriate pelagic/demersal classification. Species composition of recreational data suggests the divergence is due to the fact that species-specific data are in numbers while functional group-specific data are in weight - meaning the species-specific ratio does not account for weight differences between organisms. Landings of some small, pelagics (e.g., scaled sardine, Harengula jaguana) increased since 2000 influencing an increase in the species-specific ratio, while landings of several relatively larger demersal species (e.g., pigfish, Orthopristis chrysoptera, red porgy, Pagrus pagrus, and yellowtail snapper, Ocyurus chrysurus) have also increased influencing a decrease in the functional group-specific ratio.

There are more concerns regarding trends from U.S. recreational data then trends from U.S. commercial data. First, seasonal landings in pelagic:demersal ratio from U.S. recreational data are highly variable, so it is difficult to discern trends. Second, U.S. recreational data landed in Florida appear to be governing the overall trend for U.S. recreational pelagic:demersal ratio. Lastly, the U.S. recreational pelagic:demersal trend computed in numbers data has a different ecological meaning than trends computed with data in weight. Thus, caution should be used when interpreting and comparing the trend to others.

### 2.4.2.3 Mean Trophic Level

Landings mean trophic level trends computed from U.S. commercial and U.S. recreational data are shown in Figure 2.8. There are no obvious seasonal trends of landings mean trophic level for U.S. commercial data (Figure 2.8a) or U.S. recreational data (Figure 2.8d). For both commercial and recreational data, summer landings mean trophic level is the only significant trend and it is declining. In addition, there are no obvious differences in landings mean trophic level amongst States for U.S. commercial data (Figure 2.8b) or U.S. recreational data (Figure 2.8e). For both commercial and recreational data, Texas landings mean trophic level is the only significant trend and it is decreasing. Landings mean trophic level computed with functional group-specific data tends to be similar to that computed with speciesspecific data for both U.S. commercial (Figure 2.8c) and U.S. recreational (Figure 2.8f) data. Values from species-specific data tend to be smaller than values from functional group-specific data

Similar to the seasonal pelagic:demersal ratios from U.S. recreational data, seasonal landings mean trophic level from U.S. recreational data are highly variable, so it is difficult to discern trends. In addition, U.S. recreational data landed in Florida appear to be governing the overall trend for U.S. recreational landings mean trophic level. U.S. commercial ambiguous landings may have some influence on landings mean trophic level, especially since the computation of landings mean trophic level seems to be particularly sensitive to values used for trophic level (Appendix A). Ambiguous landings would likely reduce the U.S. commercial metric, especially in the late 80 's to early 90 's, since much of the U.S. commercial ambiguous landings are attributed
to groups with lower trophic levels (e.g., miscellaneous shrimp, and unidentified bait finfish).

### 2.5 Discussion

Although a relatively small portion of the commercial landings from NOAA are of ambiguous groups, allocating ambiguous landings to the appropriate functional groups is important in order to i) maintain the magnitude of biomass extraction in ecosystem models, and ii) account for landings of organisms of concern that are not often identified to species. An important example of the latter is sharks. There have been improvements in the identification of shark species in the Gulf of Mexico U.S. commercial landings (Appendix A), but commercial landings of ambiguous shark groups may still represent significant amounts of biomass for some species, and ignoring these landings could bias the representation of harvesting pressure in ecosystem models like Atlantis. Mexico and Cuba landings have large portions that are of ambiguous groups, particularly miscellaneous, unidentified groups. Omitting these landings from EBFM tools (i.e., ecosystem models, indicators) would introduce bias and could lead to inappropriate management advice. Thus, it is essential to associate these ambiguous landings to taxonomic classifications.

Associating ambiguous landings to inappropriate taxonomic classifications could also introduce bias into EBFM tools. Considering Atlantis, it could shift fishing pressure and biomass loss from one group to another. This could create a situation where one group is being represented as more influenced by fishing than it is in reality, and representing another group to be less influenced by fishing than it is in reality. This, too, could lead to inappropriate management advice, like suggesting that increased
fishing pressure is sustainable for one group (when it may not be), and/or that decreased fishing pressure is necessary for the sustainability of another group (when it may not be). This is less of a concern for ambiguous landings associated to a taxonomic classification higher than species (e.g., genus, family) because these landings are more likely to be associated to appropriate functional groups as functional groups often aggregate species of similar taxonomic classifications. Fortunately, much of the U.S. ambiguous landings are associated to a taxonomic classification so much of these landings are associated to appropriate functional group(s). However, distributing ambiguous landings from Cuba and Mexico across functional groups required making additional assumptions about the data as these landings were predominantly of miscellaneous/unidentified groups. Thus, there is more uncertainty concerning the Mexican and Cuban fisheries and ecosystem model outputs may not be accurate. The allocation of ambiguous catches to functional groups could be potentially be improved by incorporating knowledge of fish distributions, gear selectivity and seasonality, as the Sea Around Us Project has done in mapping global catches (Pauly, 2007).

Representing biomass loss due to recreational fishing is important for the development of EBFM tools for the Gulf of Mexico. Recreational activities are significant to the overall fishing pressure in the Gulf of Mexico (Coleman et al., 2004b), and changes in recreational information can impact management recommendations (Griffiths and Fay, 2015). MRIP, currently the best available data on U.S. recreational landings, is necessary when reconstructing historical landings profiles for the Gulf. Fortunately, most of the MRIP data are identified to species, so recreational ambiguous landings have little impact on the historical landings time series for the Atlantis Gulf of Mexico ecosystem model, and the computation of landings-based ecosystem
indicators. However, MRIP data are highly uncertain as they are estimates based on surveys expanded across the whole fishery, unlike NOAA commercial data which are based on fishermen log books and trip tickets that cover the majority of the commercial fisheries. Thus, ecosystem model results concerning recreational fleets should be interpreted with caution. Specifically, the magnitude of landings of groups not commonly harvested by recreational activities as data from those groups tend to have more variability. Efforts are under way to improve MRIP estimates (Breidt et al., 2010), and EBFM tools would benefit from considering updated data as it becomes available. Unfortunately, information pertaining to recreational activities within the southern Gulf could not be found, and recreational activities are important sources of fishing mortality for Cuba (Claro et al., 2009), and Mexico (FAO, 2003). This is also true for illegal, unreported, and unregulated (IUU) fishing, which was also not considered in this study. Thus, landings data presented here from the southern Gulf are likely under-representing activities from Mexico and Cuba, and IUU catches could be taking place anywhere in the Gulf.
U.S. landings datasets itemized by season/state/gear are informative for EBFM, but need to be considered cautiously as they can introduce uncertainty into ecosystem model results. Small portions of the seasonal commercial landings are allocated to ambiguous groups so any uncertainty in the corresponding seasonal functional group composition or landings distribution is small. Data itemized by state were not used to calibrate the Gulf of Mexico Atlantis model, but data itemized by state are informative for other EBFM tools (i.e., indicators). Most of the ambiguous landings (both commercial and recreational) are from Florida, and the proportion of Florida landings allocated to ambiguous groups is increasing over time. This is an important
area of uncertainty to be aware of for indicator assessment. Prior to 2000 a significant amount of the commercial landings could not be directly allocated to an Atlantis fleet because these data were not specified to a gear-type (e.g., combined gears, not coded). This does not bias the historical Gulf of Mexico Atlantis model because it simulates total harvest by functional group, space and time, but does not partition landings amongst fleets. However, this data will add uncertainty to the average proportions computed to distribute landings across fleets for forecast simulations.

Some U.S. commercial gears seem to be improving species identification of landings while others seem to be getting worse. Since 1997, gears targeting sharks show the most improvement towards identifying landings to species. By the end of the series, U.S. commercial ambiguous landings are dominated by the single identification used for hammerhead shark species (Appendix A), which is likely used because identifying hammerheads to species can be difficult (FAO, 2013b). However, gill netting gears show an increase in ambiguous landings, specifically of king/cero mackerel, and sharks. In terms of management, particularly for sharks, improved species identification of landings from these gears may be needed.

Trawl and purse seine gears are not associated to much of the ambiguous landings, which is not surprising as this analysis is restricted to landings and did not consider bycatch. Bycatch refers to unwanted catch that is often discarded at sea, and it is an important source of fishery induced mortality especially for trawl and seine gears (e.g., de Silva et al., 1996; Gallaway and Cole, 1999; Diamond et al., 2000; de Silva et al., 2001; Baum et al., 2003a; Harrington et al., 2005; Finkbeiner et al., 2011). The Gulf of Mexico trawl fisheries generate 19,000 tonnes of discards, generating a discard rate of 46.2 percent (Bojorquez, 1998). More recently, the U.S. National

Bycatch Report stated the Gulf of Mexico shrimp trawl had the highest ratio for trawl fisheries (0.75), and the HMS pelagic longline the smallest amongst longline fisheries (0.23) (National Marine Fisheries Service, 2013). Bycatch can impact assessment of stocks. For instance, Cortés (2002b) determined that the Atlantic blacknose shark stock was not overfished nor experiencing overfishing. Those results were influenced by catch series that did not include bycatch. When the stock was re-assessed by SEDAR (2007) which included estimates of bycatch, SEDAR concluded that the stock was overfished and experiencing overfishing. Although landings data are essential for the construction of EBFM tools and the assessment of ecosystem indicators, so are bycatch data. Neglecting bycatch can introduce bias into EBFM tools and the ecosystem indicators, particularly for species for which bycatch is an important source of mortality.

Landings-based indicator trends computed with functional group-specific data are similar to those computed with species-specific data. Thus, in the case of the Gulf of Mexico Atlantis model i) U.S. ambiguous landings have a negligible influence on landings-based indicators trends, and ii) the aggregation into functional groups has negligible influence on landings-based indicators trends. Indicator values from functional group-specific data tend to be slightly larger than indicator values from speciesspecific data. This is likely because of the aggregation of landings data into functional groups. Thus, landings-based indicator values from the Atlantis Gulf of Mexico model may not reflect values from raw data, but the differences observed here are relatively small. The largest difference observed was between the pelagic:demersal ratio values from U.S. recreational data, but this is due to functional group data being in weight and species-specific data being in numbers. Thus, the functional groups defined for
the Gulf of Mexico Atlantis model seem to appropriately represent the hierarchy of species across the Gulf of Mexico. To fully investigate this would entail a study of different hierarchical species compositions of functional groups (e.g., Fulton, 2002).

Recreational indicators presented here should be interpreted with caution since data were in numbers, not weight, and trends from numbers data have a different meaning than trends in weight. The recreational landings pelagic:demersal ratio with species data (numbers) and functional group data (lbs) showed how number data and weight data can produce different trends. Weight data may be producing metrics with a more appropriate ecological interpretation of the ecosystem since the impact of a species on an ecosystem may be more related to the biomass of a stock than number of individuals. However, MRIP data collected in weight contains half as many records as MRIP data collected in numbers, thus the weight data do not represent as much of the U.S. recreational sector as the numbers data. For the Gulf of Mexico Atlantis model, data collected in numbers were converted to weight. Although this introduces uncertainty regarding the magnitude of functional group-specific landings in weight due to the simple assumptions made about average length-weight relationships of landed species (Hayes et al., 1995), it was the best estimate that could be made.

Quantitative community indicators reveal interesting trends and data considerations. Landings mean trophic level of U.S. recreational data are higher than landings mean trophic level of U.S. commercial data, and since 1980 both trends are relatively stable. This agrees with trends presented in the Gulf of Mexico Ecosystem Status Report (Karnauskas et al., 2013), except values in the report are somewhat larger than the values presented here. First, the U.S. commercial trend presented in the report is computed with finfish data only while this study excluded only menhaden landings.

Thus, popular demersal fisheries (e.g., shrimp, crabs, oysters) are driving down the trend computed in this study. In addition, the computation of landings mean trophic level seems to be sensitive with respect to the values of trophic levels assumed for each species, and this study and the Gulf of Mexico Ecosystem Status Report may have used somewhat different values for some species. Landings pelagic:demersal ratio was not presented by Karnauskas et al. (2013). The U.S. commercial (lbs) pelagic:demersal ratio quickly increased in the 1990's and has been slowing decreasing. The U.S. recreational pelagic:demersal ratio (functional-group trend) steadily increased until 2000. Indicators computed with data series itemized by season and state show that general trends from recreational data are dominated by data from Florida, and that trends in the western Gulf are different from trends in Florida. Computing indicators with landings datasets itemized by season and state revealed interesting trends within the ecosystem for both commercial and recreational data which may be reflecting differences in historical exploitation patterns and management (Blanchard et al., 2010).

Stock assessment indicators presented here support statements made in the Gulf of Mexico Ecosystem Status report (Karnauskas et al., 2013): a majority of landed stocks have an unknown overfished status, and are not identified under a FMP. This study found that there has been little change in the number of stocks not identified under a FMP as a decrease is likely due to the decrease in the overall number of landed stocks. Also, commercial landings are mostly of assessed stocks that are not overfished, and landings of overfished stocks decreased over time due to stocks being re-assessed and recovering from their overfished status. Commercial landings of stocks with unknown status are predominantly species under GSMFC's Interjuris-
dictional Fisheries Program, some of which have been assessed under the GSMFC or by individual states. For example, blue crabs (Callinectes sapidus) make up much of these landings, and blue crabs have a GSMFC regional management plan (Perry and VanderKooy, 2015) as well as a recent stock assessment that found the stock to be not overfished and not experiencing overfishing (VanderKooy, 2013). Lastly, recreational landings are predominantly stocks of unknown status - particularly stocks not identified under a GSMFC or GMFMC FMP. This is concerning, and there should be an effort to assess stocks of unknown status to determine species of concern. Assessment at the State level may be more appropriate for some species, in addition to low data stock assessment methods, such as Carruthers et al. (2014, 2016), and Southeast Data, Assessment, and Review (2016). In addition, the Atlantis model provides a means to qualitatively investigate fisheries impact on stocks in order to identify functional groups of concern.

Landings data are important for many aspects of fisheries science and management, and it would be advantageous of EBFM of the Gulf of Mexico if some of the landings data discussed here were improved. Historical landings from Gulf of Mexico waters off the coasts of Mexico and Cuba are uncertain due to the lack of recreational data, and large portions of landings allocated to ambiguous groups. To reduce uncertainty around Mexico and Cuba landings for the purposes of Gulf-wide EBFM tools like Atlantis, practitioners should continue working with management agencies from these regions to improve species-specific catch/landing information and develop estimates of recreational landings from Gulf waters. This could include using commercial data to indirectly estimate recreational catch series (Zeller et al., 2008). Although the MRIP dataset provides crucial information regarding recreational landings from
the northern Gulf, EBFM tools of the Gulf would benefit from the incorporation of improved estimates of recreational landings. Until such efforts can be made, data discussed here are sufficient for the construction and calibration of EBFM tools like the Atlantis Gulf of Mexico model, and for calculating ecosystem indicators for the Gulf of Mexico IEA, keeping data uncertainties in mind.


Figure 2.1: Species Composition of United States Commercial Landings Over Time. Legend shows only the seven most common species.


Figure 2.2: Regional Landings Categorized by Taxonomic Classification. Landing data are of United States commercial - excluding menhaden (a), United States recreational (b), Mexico commercial (c), and Cuban commercial (d) fleets.


Figure 2.3: Seasonal Proportion of U.S. Landings and Percent of Seasonal U.S. Landings Allocated to Ambiguous Groups. U.S. commercial landings data from NOAA are itemized into four seasons, and U.S. recreational landings data from MRIP are itemized into six bimonthly intervals. Panels (a) and (b) show the seasonal proportions for U.S. commercial landings (excluding menhaden) and U.S. recreational landings, respectively. Panels (c) and (d) show the percentage of seasonal landings allocated to ambiguous groups for U.S. commercial data (excluding menhaden) and U.S. recreational data, respectively.


Figure 2.4: State Proportion of U.S. Landings and Percent of State U.S. Landings Allocated to Ambiguous Groups. Panels (a) and (b) show the state proportions for U.S. commercial (excluding menhaden), and U.S. recreational landings, respectively. Panels (c) and (d) show the percentage of state landings allocated to ambiguous groups for U.S. commercial (excluding menhaden), and U.S. recreational data, respectively.


Figure 2.5: See following page for caption.

Figure 2.5: U.S. Commercial Landings and Ambiguous Landings Itemized by Gear. U.S. commercial gears are categorized by the fleets represented in the Gulf of Mexico Atlantis model. Panel (a) shows the proportion each fleet contributes to U.S. commercial landings (excluding the fleet targeting menhaden). Panel (b) shows the proportion each fleet contributes to U.S. commercial ambiguous landings. Panels (c-f) show the percentage of commercial landings allocated to ambiguous groups for hook-and-line fleets (c), estuary fleets (d), shelf fleets (e), and miscellaneous / unidentified gears (f).


Figure 2.6: Number of Landed Species and U.S. Landings Classified by Overfished Status in Management Jurisdiction. Panel (a) shows the number of species indicated in U.S. landings categorized by overfished status described in the U.S. Congressional Stock Status Reports, and Fishery Management Plan jurisdiction. Panels (b, c) show the U.S commercial (excluding menhaden) and U.S. recreational landings, respectively, categorized by overfished status described in the U.S. Congressional Stock Status Reports, and Fishery Management Plan jurisdiction.


Figure 2.7: Trends in Landings-Based Indicator Pelagic:Demersal Ratio. Trends were computed from U.S. commercial data (a-c) and U.S. recreational data (d-f). Panels a and d show seasonal trends. NOAA's commercial data itemized by months were aggregated into four seasons, and MRIP's recreational data were itemized by bimonthly intervals. Panels b and e show trends for Gulf States. Panels c and f show trends from the annual summaries of species-specific data (solid line) and functional group-specific data (dashed line).


Figure 2.8: Trends in Landings-Based Indicator Mean Trophic Level. Trends were computed from U.S. commercial data (a-c) and U.S. recreational data (d-f). Panels a and d show seasonal trends. NOAA's commercial data itemized by months were aggregated into four seasons, and MRIP's recreational data were itemized by bimonthly intervals. Panels b and e show trends for Gulf States. Panels c and f show trends from the annual summaries of species-specific data (solid line) and functional group-specific data (dashed line).

## CHAPTER 3

# Predicting the Biomass Distributions of Pelagic Species Across the Gulf of Mexico Using Generalized Additive Models 

### 3.1 Summary

Generalized Additive Models (GAMs) were fitted for 16 pelagic functional groups to predict spatial distributions within the Gulf of Mexico. Since data were zeroinflated a delta approach was followed, which consisted of fitting a Bernoulli GAM with binomial data and a Gamma GAM with zero-truncated catch rates [number of organisms per 100 hooks]. Delta GAMs were either coastal (covering areas 0 200 m deep) or pelagic (covering areas greater than 200 m deep). Species-specific catch records were collated based on the functional groups identified for the Gulf of Mexico Atlantis model. Coastal models were developed for 4 functional groups using NOAA's Bottom Longline Survey data, and pelagic models were developed for 15 functional groups using NOAA's Pelagic Longline Observer Program data. Descriptors considered for coastal models include year, sea bottom depth, altimetry, minimum distance from a front, as well as both sea surface and sea bottom measurements of temperature, dissolved oxygen, oxygen saturation, and salinity. Descriptors
considered for pelagic models include year, season, sea bottom depth, altimetry, sea surface temperature, and minimum distance from a front. Forward selection was used to select model descriptors. Basis dimensions of smoothing splines were iteratively adjusted based on smoother fits. Model residual diagnostics and performance were evaluated, which showed that many models seem to be underestimating catch rates. Models were used to develop seasonal distribution profiles by predicting across environmental data collected from Archiving, Validation and Interpretation of Satellite Oceanographic (AVISO) and the National Centers for Environmental Information (NCEI). Model fits and predictions for the large, predatory sharks group are discussed in detail. Fitted models for large, predatory sharks have some of the better fits, diagnostics, and performance. Fitted models are influenced by known environmental drivers as well as minimum distance from a front, and there is little research identifying the influence fronts have on the distribution of predatory sharks. Model prediction profiles successfully identify areas known to have higher catch rates of sharks within the Gulf of Mexico, thus predicted seasonal distribution profiles could help identify areas where stocks have increased vulnerability. This work advances our knowledge on the environmental cues and spatial distribution of pelagic groups within the Gulf of Mexico, suggests areas of future research, and could aid the investigation of spatial fisheries management within the Gulf of Mexico.

### 3.2 Motivation

In the Gulf of Mexico, the biomass levels of many pelagic predators are currently less than historic levels primarily due to overfishing (Pauly et al., 1998; Stevens et al., 2000; Baum et al., 2003b; Myers and Worm, 2003; Christensen et al., 2003; Peel et al.,

2003; Baum and Myers, 2004; Baum et al., 2005; Safina and Klinger, 2008). Many of these stocks continue to be subjected to fishing pressure because they support economically important fisheries (e.g., Prince et al., 1989; Weidner et al., 2001; Fromentin and Powers, 2005; Barker and Schluessel, 2005; Arreguín-Sánchez and Arcos-Huitrón, 2011; Aguilar et al., 2014) or they are caught as bycatch (e.g., de Silva et al., 2001; Serafy et al., 2004; Mandelman et al., 2008). Because top predators are known to influence the structure and function of marine ecosystems (Paine, 1980; Duffy, 2002), and the decline of top predators may reduce ecosystem sustainability (Myers et al., 2007; Heithaus et al., 2008; Baum and Worm, 2009).

The 1996 Magnuson-Stevens Act includes requirements to identify essential fish habitats, i.e. waters and substrate necessary to fish for spawning, breeding, feeding or growth to maturity (U.S. Congress, 1996). The marine environment is heterogeneous, creating patchy fish populations driven by physical and biotic forcing. When marine fauna, like pelagic predators, aggregate within essential habits they are vulnerable to fisheries, so these areas should be targeted for conservation efforts. This is especially true for bycatch species as bycatch is not only harmful to affected marine fauna but also a waste of fisheries resources. Fisheries management regulations such as spatial fishery closures offer a means to protect essential habitats, but first these areas need to be defined for each species. This can be done by determining the spatial distribution of species.

Understanding the spatial distribution of fish stocks (one aspect of spatial fisheries ecology) can not only provide information regarding essential habitats, but can also lead to a better understanding of how species abundance changes over time (Ciannelli et al., 2008). The spatial distribution of an organism is often estimated using stat-
istical methods relating abundance to measurable environmental conditions. Fishery independent catch data (i.e., survey data) are collected for statistical analysis, but sample sizes tend to be low. Fisheries dependent catch data are more abundant, but often have undesirable features that make them unsuitable to linear modeling (e.g., non-random sampling, lack of coverage of an organisms whole spatial range). However, the advancement of statistical methodologies (e.g., generalized linear modeling) provides a means to address such issues with fishery dependent catch data (Guisan et al., 2002; Venables and Dichmont, 2004; Ciannelli et al., 2008). Generalized Additive Models (GAMs) offer a particularly flexible form of statistical modeling. GAMs (Hastie and Tibshirani, 1986, 1990) can address non-linear relationships between the response and explanatory variables with smoothing splines. Such models can provide information on environmental drivers influencing stock abundance (e.g., Wall et al., 2009), and identify geographic areas of increased abundance (e.g., Saul et al., 2013).

Spatial distribution models support ecosystem based fisheries management, EBFM (Brodziak and Link, 2002; Pikitch et al., 2004) by identifying essential habitats, and provides a means to parameterize the spatial distribution of marine stocks for spatially explicit ecosystem models (e.g., Atlantis). There is a growing need to understand and predict the ecosystem effects of changing predator abundances as well as the interactions with intensifying anthropogenic stressors (Baum and Worm, 2009). This can be accomplished with spatially explicit ecosystem models, but it is important that these models are appropriately representing the temporal changes in spatial abundance.

Drexler and Ainsworth (2013) presented GAMs for predicting the spatial biomass distributions of organisms retained by Southeast Area Monitoring and Assessment Program (SEAMAP) trawls. SEAMAP trawls operate within the northern Gulf of

Mexico shelf, but models were used to predict the spatial distributions of organisms across the entire Gulf of Mexico shelf. The developed GAMs successfully predicted known areas of high abundance for some organisms (e.g., pink shrimp, Farfantepenaeus duroarum). Grüss et al. (2014) used survey data spanning the West Florida Shelf to fit delta GAMs, and found that predictions from fitted models for the different lifestage groups and seasons correctly predicted known qualitative differences between low- and high-abundance areas. Both these studies encouraged using fitted models for generating distribution maps for ecosystem models. Grüss et al. (2014) focused on the OSMOSE West Florida Shelf model, and Drexler and Ainsworth (2013) focused on the Atlantis Gulf of Mexico model. Grüss et al. did not discuss the distribution of pelagic predators, but Drexler and Ainsworth observed that models for groups less vulnerable to benthic trawling gear, such as pelagic fish, performed poorly. The authors suggested that their results for these groups may be unreliable and that the analysis of different data may be necessary.

The following presents a series of GAMs for describing the spatial biomass distributions of pelagic organisms across the Gulf of Mexico. Model fits were assessed using residual diagnostics and the performance of models were tested. Seasonal, Gulf-wide distribution profiles were developed using the fitted models to predict across grids of geographic coordinates representing seasonal averages of environmental conditions. While spatial distribution profiles are developed for several functional groups, the results for large, predatory sharks are presented and discussed in detail.

### 3.3 Materials and Methods

### 3.3.1 Data For Model Fitting

Datasets were provided by NOAA's Southeast Fisheries Science Center (SEFSC) Bottom Longline Survey, and Pelagic Observer Program (Table 3.1). The Bottom Longline Survey (Grace and Henwood, 1997; Ingram et al., 2005; Henwood et al., 2006) collects catch and environmental data along the U.S. continental shelf, operating in waters with a bottom depth between $9-366 \mathrm{~m}$. Baited hooks are suspended near the benthos, and the gear used is similar to commercial longlines. The Pelagic Observer Program (Beerkircher et al., 2002, 2004; Brooke, 2012) records catch data from observers aboard vessels in the U.S. commercial pelagic longline fleet. Vessels suspend longline gear mid-depth (approximately $33-66 \mathrm{~m}$, but the actual fishing depth is unknown due to the influences by currents and environmental conditions (Beerkircher et al., 2004) throughout the Gulf's pelagic waters.

Data were collected from the Bottom Longline Survey between 2005-2012, and Pelagic Longline Observer Program between 2005-2010. Catch records were collated based on the functional groups identified for the Gulf of Mexico Atlantis model (Ainsworth et al., 2015). Tables 3.2-3.3 show the species identification and functional group classification for data from the Bottom Longline Survey and Pelagic Longline Observer Program, respectively. While some of the functional groups identified for the Gulf of Mexico Atlantis model consist of a single species (e.g. yellowfin tuna), others consist of many species (e.g., large pelagic fish). Thus, it is likely that longline catch data attributed to a multi-species functional group will not include all of the species identified in the functional group. This study does not consider longline catch data
attributed to functional groups that i) are not associated to pelagic environments, or ii) lack sufficient data to fit a GAM.

Fishery-independent and fishery-dependent datasets can be combined to analyze spatial distributions of marine organisms if the datasets share similar spatial and temporal ranges (Pecquerie et al., 2004). Except for a small area off Louisiana, the Bottom Longline Survey and Pelagic Observer Program operate in different areas (Figure 3.1), so this study keeps the two datasets separate and constructs two series of statistical models: models fit with Bottom Longline Survey data (referred to as coastal models) and models fit with Pelagic Longline Observer data (referred to as pelagic models). Survey datasets, which are fishery-independent, are designed specifically for statistical analysis and are believed to provide more accurate information regarding catch rates, but Fox and Starr (1996) found that data from commercial operations, which are fishery-dependent, can be comparable.

In regards to large, predatory sharks, Grace and Henwood (1997) found that sharks caught by the Bottom Longline Survey tend to be a size similar to or larger than the minimum size at maturity, and that they are similar in age and size to sharks caught by commercial operations (like ones sampled by the Pelagic Longline Observer Program). Older juveniles and adults tend to spend their time in coastal/offshore waters (Hueter and Tyminski, 2007), making them more accessible to longline activities. Thus, models presented here for large sharks relate to older, likely sexually mature, organisms.

Environmental and temporal variables measured at the time of each catch observation in the datasets were used as model descriptors. Temporal variables included year and season. Season was broken down into four categories: 1 (Jan. - Mar.), 2
(Apr. - Jun.), 3 (Jul. - Sep.), or 4 (Oct. - Dec.). These seasonal categories may not line up exactly with the warming and cooling of the Gulf of Mexico, since temperatures tend to reach a minimum in early March and warm summer temperatures peak in October. However, these categories correspond to the seasonality represented in the Gulf of Mexico Atlantis model. Environmental variables required some reorganization in order to fit models, which included i) generating a single estimate for the environmental variables that have multiple measurements taken during a longline set, ii) generating estimates of key environmental variables for longline sets missing the information, and iii) associating longline sets in both datasets with additional environmental data directly related to the physical environment. Measurements of several variables (e.g., sea surface temperature, bottom depth, latitude, longitude, etc) were collected at various points during the setting and hauling of each longline. The mean of all observations in each set were used to characterize that set. Estimates of these variables were generated for all of the records in both catch datasets, however approximately $25 \%$ of the bottom longline records and $5 \%$ of the pelagic longline records were missing estimates of sea surface temperature. The Interpolate PO.DAAC MODIS L3 SST at Points tool from the Marine Geospatial Ecology Tools (MGET) toolbox in ArcGIS was used to get estimates of sea surface temperature for these records (see Appendix B). Approximately $0.05 \%$ of the bottom longline records and $9 \%$ of the pelagic longline records were dropped from this analysis because necessary information was missing and could not be recovered (e.g., date, species).

Commercial longlines often set gear based on the target species' expected position in the water column (Highly Migratory Species Division, 2000). Day sets tend to target yellowfin tuna (Thunnus albacares), when the fish tend to dive deeper into
the water column (Weng et al., 2009). Night sets tend to target swordfish (Xiphias gladius), to take advantage of their nocturnal, near-surface feeding habits (Takahashi et al., 2003). Thus, time of day is an important temporal variable to consider when using commercial data. However, time of day relates less to the local stock horizontal density and more to the vertical distribution and foraging (i.e., the catchability of a functional group). Preliminary model fits were investigated using time of day, a binomial variable indicating if catch occurred at day or night, as a model descriptor (Appendix B). This work showed that spatial distribution profiles of pelagic model changed quantitatively (i.e., the magnitude of the predicted catch rates), but not qualitatively (i.e., observed trends in distribution profiles and resulting proportion maps). Since this work is more interested in horizontal (not vertical) distribution, time of day was excluded from further model fitting.

Descriptors corresponding to altimetry and ocean fronts were incorporated into the catch datasets because top marine predators are known to aggregate around oceanographic features (Olson et al., 1994; Kleisner, 2008; Kleisner et al., 2010). Altimetry data was provided by the Archiving, Validation and Interpretation of Satellite Oceanographic (AVISO) dataset (Ducet et al., 2000). Estimates of altimetry were derived by iteratively subsetting AVISO data by catch date and averaging the four AVISO records nearest to the catch location. The relationship between catch and fronts was represented by calculating the minimum distance between the catch location and a frontal feature. Frontal features were derived using the Cayula-Cornillon Fronts in ArcGIS Raster MGET tool from the MGET toolbox. This tool uses the Cayula and Cornillon (1992) edge detection algorithm for the identification and extraction of fronts. The Cayula and Cornillon edge detection algorithm is commonly used with
sea surface temperature data, but the Gulf of Mexico is known to have weak sea surface temperature gradients (Legeckis, 1978). Features within the Gulf of Mexico are largely influenced by the physical oceanography, so AVISO altimetry data were used to derive frontal features (Appendix B). ArcGIS's Model Builder was used to develop a routine that systematically estimates minimum distance from a front for catch records (Figure 3.2). For each date represented in the AVISO dataset, frontal features are derived based on the AVISO data subsetted by date, then the minimum distance between the features and catch records are calculated.

### 3.3.2 Model Description

Generalized additive models (GAMs) estimating the abundance index of individual functional groups in coastal waters (models fit with survey data) and pelagic waters (models fit with observer data) were developed using the statistical software $R$ (R Core Team, 2014; Wood, 2006a, 2011; Wood and Wood, 2015). Catch-per-unit-effort (CPUE) is the metric commonly used as an abundance index in fisheries ecology (Hilborn et al., 1992). For this study, the calculation of longline CPUE followed an industry standard: the total numbers of individuals caught per 100 hooks. There was some debate in the literature on whether hook soak time impacts catch rates calculated from longline datasets. Ward et al. (2004) concluded that whether or not soak time effects catch rates will depend on the species, and that hook soak time does effect the catch rates of sharks and billfish. Watson et al. (2005) found soak time to have a meaningful effect on blue shark catch rates, but not on swordfish catch rates. However, recently Carruthers et al. (2011), who also studied swordfish and blue sharks, found that the method of calculating soak time will determine soak
time impact on catch rates, and ended up concluding that although hook soak time effects mortality rates it does not effect catch rates. For this study, total numbers were calculated rather than total biomass because size data, which would be necessary to convert numbers to biomass, were not available for many of the records.

A Delta approach was followed to account for the zero-inflated nature of both longline datasets. The Delta method calls for fitting two statistical models: one predicting the probability of positive catch of binomial data $(0,1)$, the other predicting the CPUE [number of organisms per 100 hooks] of zero truncated catch data. Determining an appropriate error structure for generalized models is an important aspect of model construction (Maunder and Punt, 2004). By convention, the Bernoulli distribution with a logit link function was used to model the error structure of the binomial data. Preliminary analyses suggested that the catch rate data were best supported by a gamma distribution with an inverse link function (Appendix B). Other studies fitting generalized linear models to estimate catch rates from longline data have achieved comparable, if not improved, model fits using the gamma distribution rather than the more commonly used lognormal (Punt et al., 2000; Ortiz and Arocha, 2004).

The construction of GAMs requires developing robust smoothing splines for each numerical descriptor. This study used penalized regression splines, which incorporate penalties to the least squares fitting objective based on the flexibility of a smoother (Wood and Augustin, 2002). Penalized regression splines have a smoothing parameter $(\lambda)$, which controls the tradeoff between the model's fit and smoothness, and a basis dimension $(k)$, which defines the maximum possible degrees of freedom (Guisan et al., 2002; Wood, 2006b). The $\operatorname{gam}()$ function in $R$ calculates smoothing parameters using a smoothness selection criterion, either the generalized cross-validation (GCV)
criterion or the Un-Biased Risk Estimator (UBRE) scores (Wood, 2006b). An additional penalty was incorporated to allow for the removal of a numerical descriptor if the smoothing parameter equals zero (i.e., if the smoother does not improve model fit).

Adjusting the basis dimension makes a spline more flexible (Ruppert, 2002; Li and Ruppert, 2008; Wood and Wood, 2015) but this is often not investigated (Kauermann and Opsomer, 2011; Pya and Wood, 2016). Pya and Wood (2016) concluded that the exact setting of the basis dimension is not crucial as long as it is large enough to avoid over-smoothing / under-fitting, and that the simple routine presented by Wood and Wood (2015) performs as well as complex, time expensive approaches. For this study, the routine described by Wood and Wood (2015), which follows a hypothesis testing approach, was used to check the adequacy of basis dimensions. The routine uses the k-index statistic to test if the basis dimension is large enough for the smoothing spline. The k-index is the ratio of the residual variance estimated by differencing residuals that are neighbors according to the covariates of the spline, and the residual variance estimate from the whole model fit (formulas described by Pya and Wood (2016)). The k-index should be close to one if the basis dimension is large enough. A k-index less than one indicates the possibility of a missed pattern in the residuals that could be addressed if the basis dimension is increased.

To adjust basis dimensions, first a GAM is fitted with each smoother's basis dimension set to three (the minimum accepted setting). The adequacy of basis dimensions are assessed individually for each numerical descriptor, in sequential order. If the computed k -index is less than one the basis dimension is adjusted, but if the sum of all basis dimensions in the GAM are less than three-fourths of the data's sample size
the routine stops - this is to prevent overfitting. The GAM is re-fitted with the basis dimension set to a default value suggested by Kim and $\mathrm{Gu}(2004)\left(10 n^{2 / 9}\right.$, where $n$ is the sample size), which is typically much larger than three. Thus, if the resulting k-index remains less than one then adjusting the basis dimension will likely not assist reducing residual variance. In this situation, the routine sets the basis dimension to $\max (3,\lceil e d f\rceil+1)$, where $e d f$ is the effective degrees of freedom. The latter value accounts for some improvement in smoother fit. If the resulting k-index is greater than one then the routine iteratively searches for the smallest basis dimension value that still produces a k-index greater than one. This is to find a balance between improved smoother fit and the preservation of degrees of freedom. The assessed basis dimension for each numerical descriptor is set to the determined value before assessment of the basis dimension of the next numerical descriptor commences.

The general form of the fitted GAMs is as follows:

$$
\begin{equation*}
g(\eta)=\sum_{i=1}^{n} s\left(d_{i}, k_{i}\right)+\sum_{j=1}^{m} f\left(d_{j}\right) \tag{3.1}
\end{equation*}
$$

where $\eta$ is either the probability of positive catch $\left(\eta_{B}\right)$ or the abundance index $\left(\eta_{Z}\right)$ according to the link function $g$, and $d$ represents model descriptors. Models are the summation of functions of $i$ numerical descriptors and $j$ categorical descriptors. Numerical descriptors are processed with penalized regression splines (indicated by $s()$ with $k$ being the spline's basis dimension, and categorical descriptors are treated as factors (indicated by $f())$. The bottom longline dataset contains measurements of environmental variables because a conductivity, temperature, and depth recorder (CTD) was deployed at each station. Although measurements were collected at incremental depths, this study only considers measurements recorded at the sea surface and sea
bottom as they often represent the extremes. Environmental variables considered as potential descriptors for models fit with bottom longline data include: bottom depth (m), temperature $\left({ }^{\circ} \mathrm{C}\right)$, dissolved oxygen ( ppm ), oxygen saturation (\%), and salinity (ppt). Commercial longline observers record limited environmental information. Environmental variables considered as potential descriptors for models fit with pelagic longline data include: bottom depth (m) and sea surface temperature $\left({ }^{\circ} \mathrm{C}\right)$. Year, altimetry (m), and minimum distance from a front (m) were considered as potential descriptors for all models.

Model integrity is jeopardized if a model overfits data, which can occur if the model contains an excessive number of model descriptors. Over-parameterization is particularly a concern for coastal models because fitting datasets have relatively small sample sizes and there are several variables being considered for model descriptors. Over-parameterization is less of a concern for pelagic models because fitting datasets tend to have larger sample sizes and only a few model descriptors were being considered. Forward model selection was followed to select model descriptors for all GAMs. Forward model selection was preferable over other methods (i.e., backward or stepwise) because it ensured that the most influential numerical descriptors had priority in the routine adjusting basis dimension. For coastal models, forward model selection ceased (i.e., descriptors were no longer added to a model) once no single variable improved a model's explained deviance by more than $5 \%$. Forward selection for coastal models was conducted with the entire bottom longline dataset. For pelagic models, forward model selection ceased once no single variable provided any improvement to a model's deviance explained or AIC. Forward selection for pelagic models was conducted with data subsetted for cross validation (i.e., training datasets). Basis
dimensions were not adjusted (remained set equal to three) during forward selection to ensure that a variable was selected as a model descriptor because the variable improved model fit and not because of the spline's flexibility. The presence of highly correlated descriptors can detract from a model's descriptive abilities and worsen its predictability, so forward selection ignored a variable if it was highly correlated (i.e. correlation greater than 0.80 ) with any of the selected model descriptors.

Model performance and fit was evaluated for logistic models, and delta models. The receiver operating characteristic area under the curve (AUC) metric is commonly used as a global indicator of performance for logistic models (Greiner et al., 2000). The AUC is equal to the probability that the model will correctly identify a randomly chosen pair of one positive event and one negative event (Hanley and McNeil, 1982). This study followed the arbitrary AUC guidelines suggested by Swets (1988) (i.e., 0.5 is non-informative, $0.5-0.7$ are less accurate, $0.7-0.9$ are moderately accurate, 0.9-1 are highly accurate, 1 is perfect). Cross validation was used to evaluate the performance of delta models. Functional group-specific catch records from both longline datasets were split into training and testing datasets. Training datasets contained three-fourths of a functional group's catch records and testing datasets received the remaining one-fourth. Because pelagic GAMs have factors for year and season, testing and training datasets from pelagic longline data were created based on each combination of year and season to ensure training datasets contained all of the sampled years and seasons. Results are presented by plotting observed catch rates from the testing dataset against predicted catch rates from the model fitted to the training dataset (Piñeiro et al., 2008). Model fits were assessed using Pearson residuals. Residual analysis for logistic regression is complicated because a model
produces predicted probabilities for a binary response variable, however an adequate Bernoulli model should have residuals that produce a lowess curve centered along the horizontal line with a zero intercept.

### 3.3.3 Spatial Abundance Distribution Profiles

Seasonal, spatial abundance distribution profiles spanning the entire Gulf of Mexico were developed for each functional group based on predicted abundance indices generated by the fitted statistical models. First, grids describing hypothetical seasonal conditions in the Gulf of Mexico were developed. In ArcGIS a $0.1^{\circ}$ latitude by $0.1^{\circ}$ longitude grid of geographic coordinates spanning the entire Gulf of Mexico was created using the Fishnet tool. Four versions of the grid were generated, one for each season. Next, coordinates within the grids were assigned estimates for all model descriptors. Seasonal data collected from NOAA (Table 3.1) were interpolated into rasters using the Kriging tool. Then, the Extract Values to Points tool (set to bilinear interpolation) was used to assign seasonal estimates of the environmental data to each coordinate in the appropriate seasonal grid. A single bathymetry raster was used for all four seasons. Fitted models were restricted to only predict abundance at geographic coordinates having a depth estimate within the bathymetric range of the data used to fit the model. Thus, spatial grids were divided into two groups based on the 250 m isobath - creating seperate seasonal grids for coastal waters and pelagic waters. Lastly, the abundance indices for each functional group were predicted at each geographic coordinate in the grids using the fitted GAMS (i.e., coastal models were used to predict across the coastal grids, and pelagic models were used to predict across pelagic grids).

Model predictions were used to estimate Gulf-wide abundance distribution profiles for the purpose of assigning proportion of biomass across the spatial map of the Gulf of Mexico Atlantis model. Many of the functional groups are represented in only one of the two longline datasets, thus general assumptions were made to extrapolate abundance indices into Gulf-wide abundance distribution profiles (see Appendix B). Although the large pelagic fish functional group has coastal and pelagic abundance index profiles, the two profiles could not be merged because the catchability could not be standardized. Thus, Gulf-wide abundance profiles for large pelagic fish were generated using general assumptions to extrapolate pelagic predictions across coastal waters (see Appendix B). For the large sharks and skates and rays functional groups, predictions from the coastal and pelagic models were merged by standardizing predictions to account for differences in functional group catchability between the two longline operations. Catch data from an area where the two longline datasets intersect, an area off the Louisiana coast (Figure 3.1), were used to fit simple statistical models for each of these two functional groups. Statistical models solving for CPUE were of the form shown in Equation 3.1. Model descriptors included bottom depth $(\mathrm{m})$, sea surface temperature $\left({ }^{\circ} \mathrm{C}\right)$, altimetry $(\mathrm{m})$, minimum distance from a front (m), year (2005-2010), season (1-4), and longline type (bottom or pelagic). Standardization factors were calculated by dividing the median fitted pelagic CPUE by the median fitted coastal CPUE. The coastal and pelagic profiles of predicted abundance indices were averaged across the spatial map of the Gulf of Mexico Atlantis model, and averages corresponding the pelagic predictions were standardized with the computed factor. This calculation of coastal:pelagic ratios for catchability standardization assumes the ratios to be constant across space and time. Coastal and pelagic abundance
averages were then merged to span the spatial map of the Gulf of Mexico Atlantis model. Standardization factors were used for all seasonal profiles. Additional details of the methods can be found in Appendix B.

### 3.4 Results

GAMs developed from the forward selection process are summarized in Table 3.4. The median model deviance explained by coastal Bernoulli models is $32.23 \%$ (ranging from $18.55 \%$ to $46.54 \%$ ). Based on the AUC values, half of these Bernoulli models are classified as moderately accurate while other half are classified as highly accurate. The large sharks model produced one of the best fits ( $46.54 \%$ deviance explained; 0.91 AUC). Coastal gamma models described between $40.22 \%$ and $60.45 \%$ model deviance, with a median of $46.64 \%$. The weakest fit belonged to the large sharks model. Generally, fits of the pelagic Bernoulli models are weaker than the coastal Bernouli models or the coastal or pelagic gamma models, describing on average $9.12 \%$ model deviance (ranging from $3.79 \%$ to $23.27 \%$ ). Based on the AUC values, most of these Bernoulli models are classified as moderately accurate while the few remaining Bernoulli models are classified as less accurate. The large sharks model is one of the weaker fitting models (explaining $6.14 \%$ model deviance; 0.73 AUC). Fits of pelagic gamma models vary, describing between $8.33 \%$ and $71.03 \%$ model deviance with a median of $33.51 \%$. The large sharks model is one of the better fitting models explaining $70.25 \%$ of the deviance.

Residuals from the Bernoulli models can be divided into a few general trends (Figure 3.3). Most of the Bernoulli models produce residuals that create lowess curves which are mostly or entirely within the negative region, and have positive residuals
with heavier tails than the negative residuals. Thus, a majority of Bernoulli models are successful at estimating low probabilities of catch for non-catch events and unsuccessful at estimating high probabilities of catch for catch events. Some of these Bernoulli models have residuals that produce linear lowess curves and have heavytailed positive residuals (Figure 3.3a), while others have residuals that produce parabolic lowess curves and have light-tailed positive residuals (Figure 3.3b). The latter trend is from Bernoulli models which are more successful at estimating high probabilities of catch for catch events than Bernoulli models producing the former residual trend. The remaining Bernoulli models produce residuals with the opposite behavior (i.e., lowess curves mostly or entirely within the positive region, and negative residuals with heavier tails than positive residuals), indicating that these Bernoulli models are successful at estimating high probabilities of catch for catch events and are unsuccessful at estimating low probabilities of catch for non-catch events. Residuals from these Bernoulli models either produce a linear lowess curve with heavy-tailed positive residuals (Figure 3.3c), or a parabolic lowess curve with light-tailed positive residuals (Figure 3.3d). The latter trend is from Bernoulli models which are more successful at estimating low probabilities of catch for non-catch events than Bernoulli models producing the former residual trend. Residuals from the spanish mackerel and large sharks pelagic Bernoulli models produce lowess curves that have a tendency to be flat, featureless and more centered around the horizontal axis (Figure 3.3e), suggesting that there is no left-over pattern found in the residuals.

Residual diagnostic plots for gamma models are very similar to one another, so residuals for the large sharks pelagic gamma model are shown as a general example (Figure 3.4). The Q-Q plots often show deviance residuals in a U-shape (Figure 3.4a).

The weight of the tails differ among functional groups, ranging from light-tailed to heavy-tailed residuals. Error variances have a right skew (Figure 3.4b), with the severity of the skew differing among functional groups. Error variance is not constant and tends to decrease with an increasing linear predictor (Figure 3.4c). This is not immediately obvious for all functional groups, especially those with larger sample sizes, but the standardized residuals below the zero residual horizontal line tend to suggest the trend. There is no obvious trend in residuals over time (Figure 3.4d). Box-plots showed that all datasets have outliers (Figure 3.4b), some datasets with more severe outliers than others.

Cross validation results for large sharks are shown as a general example (Figure 3.5), results from the remaining models are presented in Appendix B. Many of the results resemble that of the coastal large sharks model (Figure 3.5a), however many of them have a poor r-squared value (e.g., deep water fish have a slope $=1.05$ and an $r^{2}=0.02$, filter feeding sharks, small sharks have a slope $=0.5$ and $\left.r^{2}=0.6\right)$. The large sharks pelagic model (Figure 3.5b) produced the best cross validation results out of all of the fitted models (slope $=0.82 ; r^{2}=0.457$ ). Most of the models show a tendency to predict low catch rates for larger observed catch rates. The larger cross validation residuals from the coastal model tend to be off the Mississippi river outlet (Figure 3.5c). The same is true for larger cross validation residuals from the pelagic model, with the addition that larger residuals are closer to the slope (Figure 3.5d).

Bottom depth has a significant impact on the probability of catching large sharks for both the coastal $\left(p<2.0 E^{-16}\right)$ and pelagic ( $p<2.0 E^{-16}$ ) models. Descriptor Fits for coastal and pelagic GAMs are shown in Figure (3.6) and Figure (3.7), respectively. The probability of catching large sharks increases moving away from the
shoreline until approximately 40 m , beyond which the probability of catching large sharks decreases (Figure 3.6a). Depths greater than 1200m influence an increase in the probability of catching large sharks (Figure 3.7b). Both bottom depth smoother fits become unreliable in the deeper ranges (exceeding approximately 200 m for the coastal model and 1700 m for the pelagic model) due to sparse data.

The binomial pelagic model for large sharks is also significantly influenced by minimum distance from a front, season, and year. Minimum distance from a front significantly influenced the probability of catching large sharks in pelagic waters ( $p=$ $\left.4.6 E^{-5}\right)$. The probability of catching large sharks decreases as minimum distance from a front increases, then the probability of catching large sharks increases (Figure 3.7a). The smoother fit throughout farther distances reflect increasing uncertainty due to sparse data. The probability of catching large sharks in pelagic waters has significant differences among seasons. Figure 3.7 e shows a decrease in the probability of catching large sharks in pelagic waters, which is significantly different from winter months for summer months $(p=0.002)$ and fall months $\left(p=4.56 E^{-6}\right)$. Figure 3.7 f shows a decrease in the probability of catching large sharks in pelagic waters, which is significantly different from the value in 2005 for all years $(2006, p=0.000516 ; 2007$, $\left.p=3.96 E^{-5} ; 2008, p=0.012 ; 2009, p=8.74 E^{-6} ; 2010, p=1.6 E^{-11}\right)$.

The numerical descriptors driving the large sharks coastal gamma model are sea bottom temperature $\left(p=1.24 E^{-13}\right)$ and altimetry $\left(p=2.56 E^{-15}\right)$. Higher levels of abundance of large sharks in coastal waters are encouraged by lower sea bottom temperatures (Figure 3.6b) and altimetry between 0.20 and 0.35 (Figure 3.6c). Smoother fits become less certain in sparse data ranges. Estimates of the yearly contributions to mean catch rates tend to increase over time (Figure3.6d), but only 2012 is signific-
antly different than the reference year $(p=0.023)$. There is a decrease in large sharks abundance within coastal waters for 2011 . This may be a response to changes in environmental influences or fishing effort (e.g., the 2010 Deepwater Horizon oil spill, and/or the shifting of effort due to the dynamic network of fishing closures that followed the oil spill). The abundance of large sharks in pelagic waters does not dramatically increase or decrease over time, but in 2007 their abundance was significantly larger than the reference year $(p=0.0009)$. Seasonal changes in the abundance of large sharks in pelagic waters are significant: spring $\left(p=9.29 E^{-5}\right)$, summer $(p=0.0005)$, and fall ( $p=0.0036$ ). The numerical descriptors in the large sharks pelagic gamma model are sea surface temperature ( $p=0.01$ ), bottom depth $\left(p<2.0 E^{-16}\right)$, altimetry $\left(p=2.43 E^{-10}\right)$, and minimum distance from front $\left(p<2.0 E^{-16}\right)$.

Before predicting with fitted GAMs, densities of data used to fit GAMs were compared to densities of data used to make seasonal predictions with GAMs. Data densities pertaining to environmental data of large sharks GAMs are shown in Figure 3.8. For much of the environmental data for all functional group GAMs, data densities are similar to Figure 3.8a. This is an good because i) prediction data are within the range of fitting data, ii) there is seasonality in the prediction data, and iii) the prediction data are within plausible ranges. However, data densities for bottom oxygen saturation and sea bottom dissolved oxygen indicate no seasonality in the prediction data, and that the prediction data doesn't span the entire range of the fitting data (e.g., see Figure 3.8b). In addition, data densities indicate that for coastal models some of the environmental data used for fitting doesn't span the seasonality represented in the prediction grids (e.g., see Figure 3.8c). This is caused by the lack of seasonal coverage of the bottom longline survey, discussed earlier.

Predictions across seasonal grids for both coastal and pelagic large sharks GAMs are shown in Figure 3.9. The abundance of large sharks within coastal waters is relatively low during the winter (Figure 3.9a), increases during the spring (Figure 3.9b) and summer (Figure 3.9c), and is the highest during the fall (Figure 3.9d). The abundance of large sharks within pelagic waters does not have a strong seasonal signal (Figure 3.9e-3.9h). The borders of the pelagic maps often indicate higher levels of abundance, which seems to be an indication that large sharks abundance increases moving into shallower waters. The maps reveals several areas where large sharks may be aggregating (i.e., hotspots). Coastal predictions indicate hotspots in the southwest Florida shelf, the Mississippi River outlet, the Texas coast, and Campeche Bank. Pelagic predictions indicate hotspots off the northwest Florida slope, and in the area connecting De Soto Canyon, Mississippi Canyon, and Mississippi Fan. Predictions across seasonal grids for all fitted GAMs are shown in Appendix B.

Standard errors of spring predictions for both coastal and pelagic large sharks GAMs are shown in Figure (3.10). The coastal Bernoulli model is often the most uncertain along the edge of the grid - areas that are the shallowest or deepest (Figure 3.10a). The spring predictions from the coastal Gamma model have much error (Figure 3.10b). This is because the altimetry estimates for spring are below the range of the altimetry data used for model fitting (Figure 3.8c), and smaller altimetry values are highly uncertain (Figure 3.6c). Thus, this error is largely due to the lack of seasonal coverage of the data used for fitting. The pelagic Bernoulli model has more error around the deep-edge of the slope (Figure 3.10c). The same can be said for the pelagic gamma model, with the addition that there is commonly more error in areas within the eastern Gulf basin (Figure 3.10d). Sea bottom depth data used for fitting
does not span the greater depths represented in the seasonal grids (e.g., Figure 3.8c), which is likely contributing to the error in the two GAMs. Predictions associated to areas with high standard error should be interpreted with caution. Standard errors of seasonal predictions for all fitted GAMs are shown in Appendix B.

The coastal:pelagic ratios produced by the catchability standardization (10:5.22 for large sharks, and 100:6.1 for skates and rays) allowed merging coastal and pelagic predictions to estimate the seasonal proportion of large sharks across the spatial map of the Gulf of Mexico Atlantis (Figure 3.11). Proportions were not corrected by polygon area. For instance, the relatively large proportion of large sharks in the polygon corresponding to Campeche Bank is influenced by the large size of the polygon.

### 3.5 Discussion

### 3.5.1 Model Findings

Models fitted with bottom longline survey data (coastal models) and models fitted with pelagic longline observer data (pelagic models) range in their capabilities to adequately represent data. The AUC values ranked coastal Bernoulli models between moderately and highly accurate and pelagic Bernoulli models between moderately and less accurate. Pearson residuals for coastal Bernoulli models tended to span a smaller range than Pearson residuals from pelagic Bernoulli models (i.e., pelagic Bernoulli models have more extreme residuals than coastal Bernoulli models). Thus, coastal Bernoulli models tend to be more adequate than pelagic Bernoulli models. Considering all of the Bernoulli models fitted here, most fail to produce residuals evenly
scattered around the horizontal axis when residuals are plotted against fitted values. Instead, residuals show that the Bernoulli models can be divided into two general groups: models better at predicting low probabilities of catch for non-catch events, and models better at predicting high probabilities of catch for catch events. Most Bernoulli models fitted in this study are in the former group, which may be causing the underestimation of delta-model catch rates observed in the cross validation results.

Residuals from coastal gamma models are often less extreme compared to residuals from pelagic gamma models, however residuals from all of the fitted gamma models have similar trends. Gamma models residuals appear to be independent, but they are often not normal nor are they evenly distributed across the linear predictors. The magnitude of the skew differs between gamma models so, to various degrees, many of the gamma models do not adequately represent the data. Transforming the response variable may improve these diagnostics. Log-transforming the response variable was investigated in preliminary modeling, but was not pursued as it often failed to normalize the data (Appendix B). Gamma model diagnostics may improve with the consideration of other transformations like the Basic or Box-Cox, which were suggested by Mateu (1997) in reference to normalizing environmental data. All gamma models appear to have extreme residuals which may be leverage points (i.e., extreme predictor values), and outliers (i.e., extreme response values). Removing outliers could improve the diagnostics and fits for gamma models and possibly smoothing splines as well. However, because extreme values may be correct measurements representing important variations in the system, an analysis of the extreme values might be informative.

Cross validation results showed that all delta models have less than ideal prediction performance, and some have very poor performance (e.g., large pelagic fish, filter feeding sharks, small sharks, and other turtles). Models for large sharks produced the best performance results out of all models. Cross validation results showed all models have a tendency to underestimate higher observed catch rates. This suggests a systematic bias. This may be driven by the Bernoulli models which tend to have weaker fits than Gamma models with residuals often being negative. Pelagic Bernoulli models are particularly poor. This is likely happening since pelagic models are fitted with fishery dependent data, which doesn't appropriately represent where fish are, or are not, since fishers operate in areas where fish tend to be found. This may also influencing stronger fits for the pelagic Gamma models. When assessing model predictions to detect population hotspots (i.e., areas of increased catch rates), it is important to remember that predictions may be under-representing or missing aggregations considering that many of these models seem to be underestimating higher catch rates. This may be especially true if aggregations are occurring at time-scales smaller than season.

This work executed various recommendations from the literature in order to ensure improved model fits while not violating statistical assumptions. This includes verifying the appropriate error distribution for catch rate data, determining the link functions that improves model fits, checking for correlated descriptors, buffering against overfitting, and adjusting of a spline's basis dimension of to improve model fits. Spline basis dimensions ended up being adjusted for many of the fitted models, and a couple of interesting observations emerged. First, basis dimensions were adjusted more often for splines in coastal models than for splines in pelagic models. This may be related
to the fact that coastal models have fewer descriptors than pelagic models, because the forward modeling process restricted what variables could be used as descriptors in coastal models. Also, some descriptors became statistically important after the basis dimension was adjusted. This point is important to consider when using a model selection routine along with adjusting spline basis dimensions, because selected models may be different after adjustment of a spline's basis dimension.

Extrapolating estimates for altimetry and minimum distance from a front into catch datasets helped improve model fits, especially for pelagic models. This is understandable as frontal features offer crucial zones of productivity in the open ocean because the open ocean is oligotrophic compared to nearshore waters. Minimum distance from a front improved fits for a few of the fitted models. Similarly, Podestá et al. (1993) found an association between swordfish catch rates and distance to the nearest front, as did Kleisner et al. (2010), who modeled spatial autocorrelation of fish species and temperatures at an appropriate range of depths. In addition, minimum distance from a front explained more model deviance for pelagic models of large sharks than the other numerical descriptors. There is growing evidence that filter feeding sharks orient to fronts (Sims and Quayle, 1998; Sims et al., 2000; Priede and Miller, 2009; Miller et al., 2015), also the pelagic models presented here for filter feeding sharks are largely driven by minimum distance from a front, but there is little information on predatory sharks aggregating near/around frontal boundaries (Queiroz et al., 2012). Queiroz et al. (2012) found that blue sharks (Prionace glauca) in the northeast Atlantic ocean displayed site fidelity correlating with local frontal areas, and that the temporal and spatial pattern overlapped that of pelagic longlining activities. Queiroz et al. (2016), used movement modeling to find that sharks (i.e.,
blue shark, shortfin mako; Isurus oxyrinchus, longfin mako; Isurus paucus, tiger; Galeocerdo cuvier, great hammerhead; Sphyrna mokarran, and scalloped hammerhead; Sphyrna lewini) across the Atlantic ocean prefer habitats characterized by strong sea surface-temperature gradients (fronts). Large, predatory sharks could be drawn to frontal zones as these areas could be concentrating food sources. Thus, metrics relating to the oceans physical dynamics, like minimum distance from a front, have the potential to be critical model descriptors when estimating the distribution of pelagic organisms.

### 3.5.2 Limitations

Coastal models tend to have more adequate diagnostics and tend to explain more model deviance than pelagic models, and this could be because coastal models are fitted with fishery independent data while pelagic models are fitted with fishery dependent data. Fishery dependent data (e.g., the Pelagic Longline Observer data) represent skilled fishers sampling areas known to have increased abundance of targeted organisms. Thus, the catchability represented in the dataset is higher and can lead to a overestimate of populations. Also, fishery dependent data have limited information regarding environmental conditions during catches. Fishery independent surveys often measure a variety of environmental variables at each site with devices like a CTD. Descriptors explaining the most deviance for coastal models often are variables measured by the CTD (e.g., oxygen saturation and salinity). Similar environmental information is not present in fishery dependent datasets. Thus, fishery independent data may offer more contrast of environmental conditions for target species - allowing the flexibility to capture more environmental drivers in the model.

Many pelagic stocks have seasonal migrations that cover large areas of the Atlantic Ocean and include moving into and out of the Gulf of Mexico. Some examples include Prionace glauca (Matsunaga, 2009), Thunnus thynnus (Block et al., 2001), and Xiphias gladius (Abascal et al., 2015). Thus, it is no surprise that season often explained more deviance in pelagic models than other descriptors as season relates directly to stock density. For the coastal models, season could not be included as a predictor because data were not available in winter. The lack of a season predictor may be biasing predictions from some of the coastal models. A seasonal bias is not easily detected with predictions of large sharks or blacktip sharks because shark populations are less abundant in the Gulf of Mexico during the winter months due to their southward migration (Hueter and Tyminski, 2007; Carlson et al., 2010b). A seasonal bias is obvious in the predictions form the large pelagic fish coastal model. Coastal predictions reflect much higher catch rates in the summer than other seasons (Appendix B), suggesting that large pelagic fish are either more abundant or have higher catchability in summer. There is a seasonal signal in pelagic predictions suggesting increased abundance in summer months, but not as extreme as the coastal predictions (Appendix B). Seasonal catch records developed in Chapter 2 show that the magnitude of large pelagic fish catches do not change seasonally. This suggests that large pelagic fish species are present and can be caught in the Gulf of Mexico all year. Incorporating additional datasets so that coastal models are fitted with data covering the entire seasonal range could reduce the bias observed in coastal predictions.

Incorporating other catch datasets into model fitting could improve the taxonomic coverage of this study. First, models for other important pelagic groups could not be fitted in this study due to insufficient catch data (e.g., king mackerel, small sharks,
and small pelagic fish). Also, some of the models for multi-species functional groups were not fitted with catch information pertaining to all of the species categorized into the functional group. For example, large pelagic fish catches from the bottom longline survey primarily consist of Remora and Sphyraena spp., but the group consists of many other genera (Ainsworth et al., 2015). Also, for functional groups with a coastal, and pelagic model, there is often different taxa represented between the two catch datasets. Thus, the fitted models are missing the behavior of the other species in the group. This is important to be aware of when using these results for the parameterization of the Gulf of Mexico Atlantis ecosystem model. These issues could relate to the fact that the two operations considered in this study do not select all pelagic predators throughout the water column, so considering additional catch datasets with different harvesting strategies could be informative.

Foraging behavior of pelagic piscivores governs how they are exposed to gear like baited hooks. Humphries et al. (2010) found that many open-ocean predators (e.g., sharks, tunas, billfish and ocean sunfish) exhibit vertical movement through the water column to detect food (bait) via sound, movement, and/or oder plumes. Some of these predators feed at various depth levels throughout the entire water column - exposing them to both bottom and pelagic longline activities. Medved and Marshall (1981) investigated the feeding behavior of young sandbar sharks (Carcharhinus plumbeus) and were able to catch individuals at surface, mid-depth, and bottom depth. Lowe et al. (1996) found that larger tiger sharks (Galeocerdo cuvier) move through the water column to feed at the bottom during the night and near the surface during the day. However, other pelagic predators like Bluefin tuna (Thunnus thynnus) (Lawson et al., 2009), dolphinfish (Coryphaena hippurus) (Oxenford and Hunte, 1999), and
blue marlin (Makaira nigricans) (Goodyear et al., 2008) traverse the near-surface layer of the water column, so they are only exposed to shallow-set pelagic longline operations. If species like these are retained in bottom longline operations it is likely incidental catch occurring during the setting/hauling process while the hooks are moving through the water column.

Gear design and bait can impact an organism's susceptibility to harvesting activities. Hook size, shape, and offset all have species-specific effects on catch rates. Larger hooks have resulted in decreased catch rates of the pelagic stingray (Pteroplatytrygon violacea) (Piovano et al., 2010; Coelho et al., 2012). Circle hooks, compared to J-hooks, can increase the catch rates of yellowfin tuna (Thunnus albacares) (Falterman and Graves, 2002; Kerstetter and Graves, 2006), bigeye tuna (Thunnus obseus) (Pacheco et al., 2011), and blue sharks (Prionace glauca) (Amorim et al., 2015), and decrease the catch rates of swordfish (Xiphias gladius) (Coelho et al., 2012; Amorim et al., 2015), sailfish (Istiophorus platypterus) (Pacheco et al., 2011), as well as loggerheads (Caretta caretta) and leatherbacks (Dermochelys coriacea) (Foster et al., 2012). Increasing the circle hook offset can reduce the catch rates of swordfish (Rice et al., 2012). Mackerel bait, compared to squid bait, can result in decreased catch rates of tuna (i.e., Thunnus obesus and Thunnus alalunga) (Foster et al., 2012) and swordfish (Coelho et al., 2012; Amorim et al., 2015), in addition to increased catch rates of some mackerel sharks (i.e., Lamna nasus and Isurus oxyrinchus) (Foster et al., 2012) and blue sharks (Coelho et al., 2012; Amorim et al., 2015).

A comprehensive list of fishery independent and fishery dependent surveys conducted within U.S. Gulf of Mexico waters was presented at the 2016 Gulf of Mexico Ecosystem Modeling Workshop, GOMEMOw (Grüss et al., 2016a), and highlights
additional catch datasets possibly worth integrating with the datasets considered in this study for future model fitting efforts. Like the NMFS Expanded Annual Stock Assessment (EASA) Survey, which sampled with both vertical line and longline gear from April to October in 2011 (Fitzhugh et al., 2012; Campbell et al., 2012), or the the NMFS Small Pelagics Survey, which samples the northern Gulf of Mexico from the fall through the winter using a trawl (Ingram Jr., 2008; Pollack and Ingram Jr., 2014). These two examples are fishery independent surveys but there are also fishery dependent surveys worth considering, like the NMFS Southeast Region Headboat Observer Program (O'Hop and Sauls, 2012), the Marine Recreational Fisheries Statistics Survey (MRFSS) At-Sea Observer Program (O'Hop and Sauls, 2012), and the NMFS Shark Bottom Longline Observer Program (Hale and Carlson, 2007). All of these examples are conducted within the northern Gulf of Mexico. Considering similar catch datasets from the southern Gulf of Mexico, if they exist, would greatly benefit model fits.

Minimum distance from a front was selected for several models fitted in this study. This metric depends on adequately estimating frontal zones. Estimating frontal zones by processing altimetry data with the Cayula-Cornillon Fronts in ArcGIS Raster MGET tool in $\operatorname{ArcGIS}$ seems to sufficiently capture macroscale and large-mesoscale eddies, but struggles to represent sub-mesoscale eddies. These features are also known to also support pelagic fish (Godø et al., 2012). Thus, some of the catch records considered in this study may be closer to a frontal edge than estimated. To capture these features and potentially improve estimates of the minimum distance from a front - which may improve model fits and predictions - future work should investigate
incorporating methods that are capable of estimating finer-scaled fronts (e.g., Luo et al., 2015).

There are descriptors that were not considered in this study that have the potential to improve fitted models. Chlorophyll-A has been found to influence the movement of pelagic predators Brill and Lutcavage (2001); Drymon et al. (2013), but this is likely species specific as some studies found chlorophyll-A did not improve model fits (Su et al., 2008; Grémillet et al., 2008). Prey dynamics could also inform distribution models. Schick and Lutcavage (2009) and Benoit-Bird et al. (2013) found that including data pertaining to prey groups improved predictions of bluefin tuna distributions. Drymon et al. (2013) found the CPUE of blacktip sharks was related positively with crustacean biomass. Also, some studies standardizing the catch rates of sharks found bait type to be a significant descriptor for models (Carlson et al., 2010a; Carlson and Gulak, 2013). When interpreting variables to use to model catch rates it is important to consider not only the ability to explain model deviance but also determining if the descriptor relates to local density and/or catchability. Variables relating to local density directly describe changes in stock density, while variables relating to catchability describe how susceptible an organism is to harvesting gear and methods. It is important to know an organism's life history and to make this distinction because a variable that influences catchability and varies randomly may help explain model deviance without improving the accuracy of predicting stock density. On the other hand, if a variable influencing catchability changes over time or space, it may be necessary to include this variable in the model to avoid bias in the estimates of density.

Predictions of spatial abundance using model forecasts have some limitations. First, models were fit with in situ environmental observations, which represent instantaneous conditions, but models were forecasted with data representing time- and space-integrated means. This assumes that models trained with data representing short-term behavioral responses can predict long-term habitat suitability. Thus, results may not detect acclimation, or estimate a quick population response within a functional group's range of environmental tolerance. Second, for some model descriptors, data used to forecast with fitted models do not to properly represent the seasonality of the ecosystem. This may be driving some of the differences between model predictions and information presented in literature and other data sources. Some variables could not be considered as descriptors because seasonal Gulf-wide estimates could not be developed. For instance, preliminary model fits showed beam transmission (\%), a measurement of the penetration of light through the water column, to be a statistically important descriptor (Appendix B). However, beam transmission depends on dynamic environmental processes (e.g., cloud cover, sediment, etc), so averages spanning large temporal and spatial scales would not be meaningful.

Predictions within the southern Gulf of Mexico should be interpreted with caution as they are extrapolated (i.e., there are no data from Mexican nor Cuban waters in the datasets used for model fitting). Thus, these predictions assume that the relationship between environmental drivers and functional groups densities in the southern Gulf are the same as those in the northern Gulf, which may or may not be valid. To produce more robust predictions within the southern Gulf of Mexico models should be re-fitted with datasets that include hook and line catch data from these waters.

### 3.5.3 Large Sharks Models

Temperature and depth are driving both the coastal, and pelagic larges sharks models. Water temperature is the most influential on abundance (catch rates), and bottom depth influences where sharks are (probability of positive catch) as well as abundance. Movement of shark species often corresponds with changing sea temperatures (Morrissey and Gruber, 1993; Bigelow et al., 1999; Parsons et al., 2005; Hueter and Tyminski, 2007; Ortega et al., 2009; Carlson et al., 2010b; Baum and Blanchard, 2010), and depth influences habitat selection (Morrissey and Gruber, 1993; Heithaus et al., 2007b; Carlson et al., 2010b; Hoffmayer et al., 2014) and catch rates (Carlson et al., 2010a; Baum and Blanchard, 2010; Drymon et al., 2010; Carlson et al., 2012). Factors time and season were also significant for larges sharks pelagic models, which has been observed in studies standardizing catch rates of sharks (Carlson et al., 2010a, 2012; Carlson and Osborne, 2013; Carlson and Gulak, 2013). Sea bottom habitat could also drive the distribution of some shark species (Hannan et al., 2012). Salinity can also influence the habitat selection for sharks (Heupel and Simpfendorfer, 2008; Ubeda et al., 2009; Bethea et al., 2015), but models fitted with salinity (large sharks coastal models) did not select salinity as a model descriptor. This is because studies relating to salinity tend to focus on young, juvenile sharks and localized inshore systems, but this research aimed to describe the distribution of older juveniles and adults, which often spend most of their time in coastal/offshore waters (Hueter and Tyminski, 2007). Constructing models to describe the distribution profiles for younger organisms would be beneficial, but would require catch data that retains those individuals and should focus more on small scale, inshore studies.

Major features of predictions made with the fitted large sharks coastal model seem to be supported by observations from the literature. First, predictions indicate a seasonal signal. Catch rates are very low in the winter, but increase in the spring, summer, and fall. This corresponds to the theory of a general southward migration of sharks (Hueter and Tyminski, 2007; Carlson et al., 2010b). Second, areas of increased catch rates in the northern Gulf are predicted off the coasts of Texas, Louisiana, Mississippi and south Florida. Hotspots off of Texas, Louisiana, and south Florida appear in the spring while the hotspot off of Mississippi appears in the summer. Hueter and Tyminski (2007) concluded that 16 different species of sharks use areas off of Florida and Texas as primary and/or secondary nurseries. Their results for Texas show more older individuals were observed off Corpus Christi, and their results for Florida indicate that many older individuals were observed off the Florida Keys. These areas are approximately where the corresponding hotspots are occurring in the results presented here. Bethea et al. (2015) determined some areas in the northeast Gulf to be important nursery grounds. Results presented here did not show increased abundance in these areas, but Bethea et al. (2015) studied young of the year and juveniles while this study only studied adults.

Results presented in this study suggest a connection between large, predatory sharks and the Mississippi River outlet, particularly the dead zone. First, for both the coastal and pelagic large sharks models, extreme outliers from the cross validation tend to be located off the Mississippi River outlet. Although removing extreme outliers could improve model fits, it appears these outliers represent important instances of variation in the ecosystem - specifically how variable shark catches are in this area. Second, the predictions from the large sharks coastal model show increased
abundance of sharks along the Gulf's northeastern shelf in the summer months. This temporal and spatial domain matches that of the Gulf of Mexico hypoxic zone (a.k.a., dead zone). Studies of hypoxic conditions within the Gulf of Mexico have observed fish aggregating along the edge, and/or immediately above, the hypoxic areas since many marine teleost can not inhabit hypoxic waters (Stanley and Wilson, 2004; Zhang et al., 2009). However, experimental studies have found that some sharks are capable of altering their physiology and swimming behavior to tolerate hypoxic conditions (Metcalfe and Butler, 1984; Wise et al., 1998; Carlson and Parsons, 2001). Also, Heithaus et al. (2009) found dissolved oxygen to drive the distribution of bull sharks (Carcharhinus leucas) within Everglades National Park even in the absence of hypoxia. Thus some species of sharks might take advantage of the Gulf of Mexico hypoxic zone to forage for benthic organisms (e.g., crabs) and/or locate aggregations of fish occurring around the zone-edge. Prince and Goodyear (2006) analyzed tag data from individuals exposed to the eastern tropical Pacific hypoxic zone and suggested that the larger sizes of sailfish observed may be due to the enhanced foraging opportunities afforded by the closer proximity of predator and prey in compressed habitat.

Predictions of large sharks across the pelagic environment do not have noticeable seasonality, but there are two distinct features of these results that stand out. First, the continental slope has higher catch rates than deep, pelagic waters. Many studies have found sharks in the Gulf to have an affinity for the shelf and coast, but there have been individuals observed making trips into the deep, pelagic waters. For instance, Hoffmayer et al. (2014) tagged several dusky sharks (Carcharhinus obscurus) in the northern Gulf, and observed sharks spending most of their time along the shallow edges of the slope. However, in the fall one shark swam south to spend time at depths
greater than 300 m . Second, the slope has areas of increased catch rates. Higher catch rates are estimated around DeSoto Canyon, Mississippi Canyon, and Mississippi Fan. These increased estimates are consistent through winter to summer but decrease in the fall. Coincidentally, this spatial range and temporal pattern is similar to that of the intrusion and eddy shedding of the Loop Current (Leben, 2005). Considering the influence minimum distance from a front and altimetry have on the delta model, it is possible that these features may be connected to the physical dynamics generated from the Loop Current. Some research have suggested these areas to be important for various shark species. For instance, Etnoyer and Warrenchuk (2007) suggested the Mississippi Canyon may be a nursery for catsharks (Scyliorhinidae spp.). Also, Hueter and Tyminski (2007) found offshore coastal nurseries off Texas, Louisiana, and Mississippi through longline surveys in the months of July and August. The importance of these areas for predatory shark populations should be an topic of future research in order to improve our understanding of shark habitats in the Gulf of Mexico.

Predictions show that sharks may appregate in areas of the southern Gulf of Mexico - primarily within Campeche Bank, some coastal estuaries, and the continental slope. These results should be interpreted with caution since forecasts across the southern Gulf of Mexico were extrapolated, but some of these trends are supported by the literature. In Cuba, sharks are commonly caught as bycatch in the pelagic longline fisheries (Guitart, 1975; Aguilar et al., 2014). In Mexico, some of the indicated areas correspond to shark fisheries Castillo-Géniz et al. (1998); Pérez-Jiménez and MendezLoeza (2015), and possible nurseries (Montiel, 1988; Bonfil, 1997).

### 3.5.4 Moving Forward

Spatial distributions of pelagic fish within the Gulf of Mexico can be quite different from one another. Some of the species investigated here are not properly represented by longline datasets alone, and more catch and environmental information is needed to get a better understanding of their Gulf-wide distribution. Models for large sharks produced some of the best fits with few diagnostic issues which resulted in good performance, thus these models could aid conservation efforts for large, predatory sharks inhabiting Gulf of Mexico waters. However, The large sharks functional group consists of twenty-six different shark species, all of which have slightly different life histories, behaviors, and habitat preferences. As our knowledge and data-banks grow, species-specific investigations should be pursued so conservation plans can aim at species of concern in addition to the large sharks complex. Seasonal predictions from all of the models presented here can provide some improvement to the representation of pelagic functional groups in the Gulf of Mexico Atlantis ecosystem model, which will aid ecosystem based fisheries management efforts in the Gulf of Mexico. Considering that many of these highly migratory stocks inhabit waters far beyond the Gulf, ecosystem models like Atlantis are critical in gaining insight on how conservation efforts covering very small areas of their spatial range impact the overall status of these stocks.

Table 3.1: Data Used For Model Fitting and Predicting. Catch and effort data were provided by the Southeast Fisheries Science Center (SEFSC), and were supplemented with sea surface temperature and altimetry data from the Moderate Resolution Imaging Spectroradiometer (MODIS) and Archiving, Validation and Interpretation of Satellite Oceanographic (AVISO) datasets, respectively. AVISO data were collected from NOAA's Easier Access To Scientific Data database, while MODIS data were assigned to catch records using the Interpolate PO.DAAC MODIS L3 SST at Points tool from the Marine Geospatial Ecology Tools toolbox in ArcGIS. Data used to forecast with fitted models all came from NOAA. The National Centers for Environmental Information (NCEI) provided bathymetry data, and the NOAA's National Centers for Environmental Information (NCEI) provided climatological seasonal averages of sea temperature, dissolved oxygen, oxygen saturation, and salinity.

| Data | Source | Spatial/Temporal Resolution | Use |
| :---: | :---: | :---: | :---: |
| Coastal catch and effort | SEFSC | Northern Gulf of Mexico (9-366m isobaths) | Fitting |
| Pelagic catch and effort | SEFSC | Northern Gulf of Mexico (200m isobath - EEZ ) | Fitting |
| Sea surface temperature | MODIS | $\operatorname{grid}(4 \mathrm{~km} / 9 \mathrm{~km}) /(\text { Daily } / 8 \mathrm{Day} / \text { Monthly })^{\ddagger}$ | Fitting |
| Altimetry | AVISO | $0.25^{\circ} \times 0.25^{\circ}$ grid / weekly averages | Both |
| Bathymetry | NCEI | Gulf-wide ( $5-4000 \mathrm{~m}$ isobaths) | Forecast |
| Sea temperature ${ }^{\dagger}$ | NCEI | $0.25^{\circ} \times 0.25^{\circ} \mathrm{grid} /$ seasonal averages | Forecast |
| Dissolved oxygen ${ }^{\dagger}$ | NCEI | $1^{\circ} \mathrm{x} 1^{\circ}$ grid / seasonal averages | Forecast |
| Oxygen saturation ${ }^{\dagger}$ | NCEI | $1^{\circ} \times 1^{\circ}$ grid / seasonal averages | Forecast |
| Salinity ${ }^{\dagger}$ | NCEI | $0.25^{\circ} \times 0.25^{\circ} \mathrm{grid} /$ seasonal averages | Forecast |

[^0]Table 3.2: Species identified in NOAA's Bottom Longline Survey dataset collated by functional groups defined for the Gulf of Mexico Atlantis Model (Ainsworth et al., 2015).

| Greater amberjack (AMB)* | Large reef fish (LRF) ${ }^{\dagger}$ | Sciaenidae (SCI) ${ }^{\dagger}$ |
| :---: | :---: | :---: |
| Seriola dumerili | Brotula barbata | Menticirrhus americanus |
|  | Conger oceanicus | Micropogonias undulatus |
| Black drum (BDR) ${ }^{\dagger}$ | Echiophis punctifer |  |
| Pogonias cromis | Rhynchoconger flavus | Scamp (SCM) ${ }^{\dagger}$ |
|  | Ophichthidae | Mycteroperca phenax |
| Benthic feeding sharks (BEN)* | Ophichthus Gulf of Mexicoesi |  |
| Heptranchias perlo | Ophichthus puncticeps | Small demersal fish (SDF) ${ }^{\dagger}$ |
| Hexanchus vitulus | Ophichthus rex | Scorpaena agassizii |
|  | Trichiurus lepturus |  |
| Deep serranidae (DSR) ${ }^{\dagger}$ |  | Seatrout (SEA) ${ }^{\dagger}$ |
| Centropristis striata | Little tunny (LTN)* | Cynoscion arenarius |
| Epinephelus drummondhayi | Euthynnus alletteratus | Cynoscion nothus |
| Epinephelus flavolimbatus |  |  |
| Epinephelus nigritus | Lutjanidae (LUT) ${ }^{\dagger}$ | Small sharks (SMS)* |
| Epinephelus niveatus | Etelis oculatus | Centrophorus granulosus |
|  | Lutjanus analis | Squalus |
| Flatfish (FLT) ${ }^{\dagger}$ | Lutjanus griseus | Squalus cubensis |
| Syacium papillosum | Pristipomoides aquilonaris |  |
|  |  | Small pelagic fish (SPL)* |
| Gag grouper (GAG) ${ }^{\dagger}$ | Other demersal fish (ODF) ${ }^{\dagger}$ | Merluccius bilinearis |
| Mycteroperca microlepis | Arius felis |  |
|  | Bagre marinus | Small reef fish (SRF) ${ }^{\dagger}$ |
| Jacks (JCK)* | Gymnothorax kolpos | Caulolatilus microps |
| Carangidae | Gymnothorax nigromarginatus | Lopholatilus chamaeleonticeps |
| Seriola zonata | Haemulon plumieri | Rachycentron canadum |
|  | Muraena retifera | Synodus foetens |
| King mackerel (KMK)* | Opsanus pardus | Trachinocephalus myops |
| Scomberomorus cavalla | Pagrus pagrus |  |
|  | Prionotus tribulus | Swordfish (SWD)* |
| Large sharks (LGS) | Urophycis | Xiphias gladius |
| Carcharhinidae | Urophycis cirrata |  |
| Carcharhinus | Urophycis floridana | Blacktip sharks (TIP) |
| Carcharhinus acronotus |  | Carcharhinus limbatus |
| Carcharhinus altimus | Skates and Rays (RAY) |  |
| Carcharhinus brevipinna | Dasyatidae | Vermilion snapper (VSN) ${ }^{\dagger}$ |
| Carcharhinus falciformis | Dasyatis | Rhomboplites aurorubens |
| Carcharhinus isodon | Dasyatis americana |  |
| Carcharhinus leucas | Dasyatis centroura | Not Assigned ${ }^{\dagger}$ |
| Carcharhinus plumbeus | Dasyatis sabina | Gadidae |
| Carcharhinus signatus | Mustelus | Unidentified |
| Galeocerdo cuvier | Mustelus canis |  |
| Ginglymostoma cirratum | Mustelus norrisi |  |
| Negaprion brevirostris | Mustelus sinusmexicanus |  |
| Rhizoprionodon terraenovae | Raja eglanteria |  |
| Sphyrna | Raja garricki |  |
| Sphyrna lewini | Rhinoptera bonasus |  |
| Sphyrna mokarran | Scyliorhinus retifer |  |
| Sphyrna tiburo | Triakidae |  |
| Sphyrnidae |  |  |
|  | Red drum (RDR) ${ }^{\dagger}$ |  |
| Loggerhead (LOG)* | Sciaenops ocellatus |  |
| Caretta caretta |  |  |
|  | Red grouper (RGR) ${ }^{\dagger}$ |  |
| Large pelagic fish (LPL) | Epinephelus morio |  |
| Acanthocybium solandri |  |  |
| Coryphaena hippurus | Red snapper (RSN) ${ }^{\dagger}$ |  |
| Echeneis naucrates | Lutjanus campechanus |  |
| Echeneis neucratoides |  |  |
| Remora remora |  |  |
| Sphyraena barracuda |  |  |

Table 3.3: Species identified in the NOAA's Pelagic Longline Observer dataset collated by functional groups defined for the Gulf of Mexico Atlantis Model (Ainsworth et al., 2015).


[^1]

Figure 3.1: Geographic Distribution of Effort from NOAA's Bottom Longline Survey and NOAA's Pelagic Longline Observer Program. Effort [\# of hooks per km²] from NOAA's Bottom Longline Survey is shown in green, and effort from NOAA's Pelagic Longline Observer Program is shown in blue. This map was calculated and created in ArcGIS via the Point Density tool.


Figure 3.2: Conceptual Routine for Calculating Minimum Distance from A Front. The routine to calculate minimum distance from front (MDF) was constructed in ArcGIS's Model Builder. Oval boxes indicate data files, square boxes indicate ArcGIS tools, and oblong boxes indicate tool outputs. The diamond box indicates a feature for iterating through unique dates identified in the altimetry dataset. A front profile is constructed for an individual date ( $n$ ), which is used to calculate the minimum distance from a front for catch records associated to longline sets that occurred on date $n$.

Table 3.4: Generalized Additive Models Fitted for Seasonal Predictions. Results from the forward model selection procedure for all fitted models. Models were determined for both parts of the delta model predicting CPUE: the probability of positive catch $\left(\eta_{B}\right)$ and the abundance index $\left(\eta_{Z}\right)$. Numerical descriptors include bottom depth $(B D)$, sea surface temperature $(S S T)$, sea bottom temperature $(S B T)$, sea surface height $(S S H)$, oxygen saturation $(O S)$, dissolved oxygen $(D O)$, salinity $(S A L)$, and minimum distance from front ( $M D F$ ) - subscripts indicate whether the descriptor is measured from the sea surface $(S)$ or sea bottom $(B)$. All of which were fitted with penalized regression splines $(s())$. Year (yr) and season $(s n)$ are treated as factors $(f())$. Deviance explained (D.E., \%) is displayed for each model, and the receiver operating characteristic area under the curve (AUC) is displayed for Bernoulli models.

Table 3.4: Continued.

| Models Fit with NOAA's Bottom Longline Survey Data |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Functional Group |  | Model | D.E. | AUC |
| large sharks | $\eta_{B}$ | $s(B D, 7)$ | 46.5 | 0.91 |
|  | $\eta_{Z}$ | $s(S B T, 13)+s(S S H, 12)+f(y r)$ | 40.2 |  |
| large pelagic fish | $\eta_{B}$ | $s\left(O S_{B}, 4\right)+s(S S T, 7)+s\left(D O_{S}, 4\right)+s(B D, 9)+s\left(S A L_{B}, 7\right)$ | 22.9 | 0.86 |
|  | $\eta_{Z}$ | $s\left(D O_{S}, 3\right)+s(S S H, 3)+f(y r)+s\left(O S_{B}, 3\right)$ | 60.4 |  |
| skates and rays | $\eta_{B}$ | $s(S B T, 11)+s\left(O S_{B}, 3\right)$ | 18.5 | 0.79 |
|  | $\eta_{Z}$ | $s(S B T, 5)+s(S S T, 10)+s\left(S A L_{S}, 4\right)$ | 50.9 |  |
| blacktip sharks | $\eta_{B}$ | $s(S B T, 6)+s\left(S A L_{S}, 3\right)+s(S S H, 3)+s\left(O S_{B}, 3\right)$ | 41.6 | 0.91 |
|  | $\eta_{Z}$ | $s\left(S A L_{B}, 3\right)+s\left(D O_{B}, 15\right)+s(B D, 3)$ | $42.4$ |  |
| Models Fit with Pelagic Longline Data |  |  |  |  |
| Functional Group |  | Model | D.E. | AUC |
| other billfish | $\eta_{B}$ | $s(S S T, 3)+f(y r)+f(s n)+s(M D F, 3)+s(B D, 3)+s(S S H, 3)$ | 13.6 | 0.75 |
|  | $\eta_{Z}$ | $f(y r)+s(B D, 22)+s(S S T, 28)+s(M D F, 3)+s(S S H, 3)$ | 37.5 |  |
| blue marlin | $\eta_{B}$ | $s(S S T, 3)+f(s n)+f(y r)+s(S S H, 3)+s(M D F, 3)+s(B D, 3)$ | 9.5 | 0.71 |
|  | $\eta_{Z}$ | $s(S S T, 3)+s(S S H, 3)+f(y r)+s(B D, 3)+s(M D F, 21)+f(s n)$ | 31.8 |  |
| bluefin tuna | $\eta_{B}$ | $f(s n)+s(S S T, 3)+f(y r)+s(S S H, 3)+s(M D F, 3)+s(B D, 3)$ | 18.2 | 0.78 |
|  | $\eta_{Z}$ | $f(y r)+s(S S H, 3)+f(s n)+s(M D F, 3)+s(B D, 3)$ | 8.3 |  |
| deep water fish | $\eta_{B}$ | $f(s n)+s(S S T, 3)+s(M D F, 3)+s(B D, 3)+s(S S H, 3)$ | 11.1 | 0.77 |
|  | $\eta_{Z}$ | $f(s n)+s(S S T, 3)+s(M D F, 3)+f(y r)+s(S S H, 3)+s(B D, 3)$ | 55.0 |  |
| filter feeding sharks | $\eta_{B}$ | $f(s n)+f(y r)+s(M D F, 3)+s(B D, 3)+s(S S H, 3)$ | 7.0 | 0.72 |
|  | $\eta_{Z}$ | $f(y r)+s(B D, 5)+s(S S H, 3)+s(M D F, 3)+s(S S T, 3)$ | 33.4 |  |
| large sharks | $\eta_{B}$ | $f(s n)+f(y r)+s(M D F, 3)+s(B D, 3)+s(S S T, 3)+s(S S H, 3)$ | 6.14 | 0.73 |
|  | $\eta_{Z}$ | $s(S S T, 3)+s(B D, 3)+f(y r)+f(s n)+s(S S H, 29)+s(M D F, 33)$ | 70.3 |  |
| large pelagic fish | $\eta_{B}$ | $f(s n)+f(y r)+s(S S H, 3)+s(B D, 3)+s(S S T, 3)$ | 8.7 | 0.65 |
|  | $\eta_{Z}$ | $f(y r)+s(B D, 46)+s(S S T, 12)+s(S S H, 37)+f(s n)+s(M D F, 8)$ | 33.6 |  |
| medium pelagic fish | $\eta_{B}$ | $s(S S H, 3)+f(y r)+f(s n)+s(S S T, 3)+s(B D, 3)+s(M D F, 11)$ | 6.5 | 0.68 |
|  | $\eta_{Z}$ | $f(s n)+f(y r)+s(S S H, 3)+s(S S T, 3)+s(M D F, 3)+s(B D, 3)$ | 16.6 |  |
| skates and rays | $\eta_{B}$ | $f(y r)+f(s n)+s(S S T, 3)+s(M D F, 3)+s(B D, 3)$ | 16.1 | 0.76 |
|  | $\eta_{Z}$ | $f(s n)+f(y r)+s(S S T, 3)+s(S S H, 3)+s(B D, 3)+s(M D F, 13)$ | 36.5 |  |
| spanish mackerel | $\eta_{B}$ | $s(S S T, 3)+f(y r)+f(s n)+s(M D F, 3)+s(S S H, 8)$ | 3.8 | 0.63 |
|  | $\eta_{Z}$ | $f(s n)+s(B D, 41)+f(y r)+s(S S T, 19)+s(S S H, 9)+s(M D F, 43)$ | 25.5 |  |
| small sharks | $\eta_{B}$ | $f(s n)+f(y r)+s(S S T, 3)+s(B D, 3)+s(M D F, 3)+s(S S H, 3)$ | 4.3 | 0.65 |
|  | $\eta_{Z}$ | $s(B D, 35)+s(M D F, 3)+s(S S H, 19)+s(S S T, 3)+f(y r)+f(s n)$ | 58.5 |  |
| swordfish | $\eta_{B}$ | $f(s n)+f(y r)+s(S S H, 3)+s(S S T, 9)+s(M D F, 3)+s(B D, 3)$ | 14.9 | 0.76 |
|  | $\eta_{Z}$ | $s(B D, 49)+s(M D F, 3)+s(S S T, 3)+s(S S H, 3)+f(y r)+f(s n)$ | 48.7 |  |
| other turtles | $\eta_{B}$ | $f(s n)+f(y r)+s(B D, 3)+s(S S H, 3)+s(S S T, 3)+s(M D F, 11)$ | 4.7 | 0.67 |
|  | $\eta_{Z}$ | $f(y r)+s(M D F, 4)+s(S S T, 3)+s(S S H, 3)+s(B D, 4)$ | 71.0 |  |
| white marlin | $\eta_{B}$ | $s(S S T, 3)+f(y r)+s(S S H, 3)+f(s n)+s(B D, 3)+s(M D F, 10)$ | 18.7 | 0.82 |
|  | $\eta_{Z}$ | $f(y r)+s(S S T, 3)+s(S S H, 3)+s(M D F, 3)+s(B D, 3)$ | 25.1 |  |
| yellowfin tuna | $\eta_{B}$ | $f(s n)+f(y r)+s(S S H, 3)+s(M D F, 3)+s(S S T, 3)+s(B D, 3)$ | 23.3 | 0.80 |
|  | $\eta_{Z}$ | $s(S S T, 3)+s(B D, 3)+f(y r)+f(s n)+s(S S H, 28)+s(M D F, 4)$ | 14.7 |  |



Figure 3.3: General Trends of Residual Diagnostics for Fitted Bernoulli Models. Examples of the general trends of residuals from fitted Bernoulli models are shown for indicated functional groups. Red lines indicate the lowess smooth.


Figure 3.4: Residual Diagnostics from the Large Sharks Gamma Model Fitted with Pelagic Longline Observer Data. Residual diagnostics include: the Q-Q plot (a), box plot (b), residuals against linear predictor (c), which is the predicted value for each data point in the scale of the link function, and residuals against time (d).


Figure 3.5: Cross Validation Results from the Large Sharks Delta Generalized Additive Models. Results are presented as observed against predicted catch rates for the coastal model (a) and pelagic model (b). Results from a linear regression on the points (solid line) are shown: intercept (int), slope (s), and adjusted r-squared value (r2). The dashed line indicates the 1:1 ratio between observed and predicted values. Cross validation residuals are presented based on the corresponding geographic coordinates for the coastal model (c) and pelagic model (d). Point size indicates if the residual is larger than the $95 \%$ quantile.


Figure 3.6: Model Descriptor Fits from the Large Sharks Generalized Additive Model Fitted with Bottom Longline Survey Data. Panel (a) displays the model descriptor fit for the binomial data model, and panels (b) - (d) display the model descriptor fits for the zerotruncated data model. Solid lines indicate the fit, dashed lines indicate the $95 \%$ confidence interval, and the black dashes along the horizontal axis display the rug plot. The estimated degrees of freedom for smooth fits are included in the vertical axis label.


Figure 3.7: Model Descriptor Fits from the Large Sharks Generalized Additive Model Fitted with Pelagic Longline Data. Panels (a) - (f) display descriptor fits for the binomial data model, and panels (g) - (l) display descriptor fits for the zero-truncated data model. Solid line indicates the fit, dashed lines indicate the $95 \%$ confidence interval, and black dashes along the horizontal axis display the rug plot. The estimated degrees of freedom for smooth fits are included in the vertical axis label.


Figure 3.8: Examples of Density Plots Comparing Fitting and Predicting Data. The density curve for data used for model fitting is plotted in black, and the density curves for data used for seasonal predictions are plotted individually in the indicated shades of grey. Panel (a) displays an ideal situation: prediction data within the range of fitting data, and seasonality amongst prediction data. Panel (b) displays a less ideal situation: prediction data failing to span the range of fitting data, and no seasonality amongst prediction data. Panel (c) displays a less ideal situation: fitting data failing to span the range of prediction data.


Figure 3.9: Seasonal Predictions of Large Sharks Catch Rates. Panels (a) - (d) display the catch rates predicted from the large shark GAM fit with bottom longline survey data for season 1 (a), season 2 (b), season 3 (c), and season 4 (d). Panels (e) - (h) display the catch rates estimated when predicting the large shark GAM fit with pelagic longline observer data across for season 1 (e), season 2 (f), season 3 (g), and season 4 (h).


Figure 3.10: Example of Standard Errors of Predictions from Large Sharks Generalized Additive Models. Standard error of season 2 predictions for the large sharks coastal Bernoulli model (a), coastal Gamma model (b), pelagic Bernoulli model (c), and pelagic Gamma model (d).


Figure 3.11: Proportion of Large Sharks Abundance Aggregated by Gulf of Mexico Atlantis Polygon Map. Seasonal predictions for the coastal and pelagic large sharks models were merged using a standardized relative catchability coastal:pelagic ratio (10:5.22), which was computed with data collected from an area where the two longline catch datasets overlapped. Figures are partitioned seasonally: season 1 (a), season 2 (b), season 3 (c), and season 4 (d).

## CHAPTER 4

## Can Gulf of Mexico Pelagic Longline Fishery Closures Meet Management Objectives?

### 4.1 Summary

The Gulf of Mexico has two pelagic longline closures, a permanent closure (DeSoto Canyon), and a seasonal closure (Spring Closure), which span pelagic waters where highly migratory predators aggregate to spawn and/or forage. Management objectives of these closures include reducing the catch and rebuilding the biomass of bycatch groups (i.e., Atlantic billfish, bigeye tuna, some pelagic sharks, prohibited sharks, and sea turtles) and incidental species (i.e., bluefin tuna), without impacting catch of target species (i.e., swordfish, yellowfin tuna, bigeye tuna, skipjack tuna, albacore, dolphin fish, wahoo, and some coastal sharks). Ecosystem modeling tools like Atlantis can be used to address the utility of pelagic MPAs for mitigating fishing pressure experienced by highly migratory predators, as well as broader ecosystem impacts. A policy exploration was conducted with the Gulf of Mexico Atlantis model to investigate if Gulf of Mexico pelagic longline spatial closures could achieve management objectives, as well as potential ecosystem impacts. Performance measures
corresponding to management objectives were monitored, as well as the ecosystem performance measures average individual weight, proportion mature, pelagic:demersal ratio (for catch and the ecosystem), and ecosystem biodiversity. DeSoto Canyon was more successful at achieving management objectives than Spring Closure. Both closures reduced Gulf-wide catches of some bycatch and incidental groups (especially green turtles and miscellaneous tunas) with little reduction to total catch of target groups. Gulf-wide catch of targeted yellowfin tuna actually increased. Neither closure caused meaningful increases in biomass of bycatch or incidental groups, but there were meaningful increases in biomass of some targeted groups (especially yellowfin tuna). DeSoto Canyon changed ecosystem performance metrics while Spring Closure did not. In particular, DeSoto Canyon reduced catch pelagic:demersal ratio which increased ecosystem pelagic:demersal ratio. This study suggests that DeSoto Canyon could be meeting most of the management objectives, and that Spring Closure may not meet long-term management objectives.

### 4.2 Motivation

Many pelagic predators around the globe have historically low biomass (Pauly et al., 1998; Myers and Worm, 2003; Christensen et al., 2003; Baum and Worm, 2009). Reduction in top-down pressures can restructure marine communities (Parsons, 1992; Heithaus et al., 2008; Baum and Worm, 2009) and initiate impacts, including shifting mortality to stocks that cannot sustain such pressure (Myers et al., 2007), shifting ecosystem functionality (Casini et al., 2009), and reducing ecosystem resistance and resilience (Britten et al., 2014). Although there are stocks of Atlantic of pelagic predators (e.g., billfish and tuna) that are not overfished nor suffering overfishing (Die,

2006; Collette et al., 2011; Juan-Jordá et al., 2011), there are other stocks that are of concern. Some species of large sharks are particularly depleted (Stevens et al., 2000; Baum et al., 2003b; Baum and Myers, 2004; Baum et al., 2005; Burgess et al., 2005; de Mutsert et al., 2008; Baum and Blanchard, 2010). Around the world sharks are targeted by large-scale, artisanal, and sport fisheries (Castillo-Géniz et al., 1998; Smale, 2008; Morgan et al., 2009; Pérez-Jiménez and Mendez-Loeza, 2015; Fahmi and Dharmadi, 2015), in addition to being caught as bycatch (McKinnell and Seki, 1998; de Silva et al., 2001; Beerkircher et al., 2002; Rogan and Mackey, 2007; Mandelman et al., 2008; Petersen et al., 2009; Belcher and Jennings, 2011). The intense fishing pressure combined with their slow growing life history makes sharks particularly vulnerable to overfishing and extinction (Monte-Luna et al., 2007; Dulvy et al., 2008; García et al., 2008; Field et al., 2009; Worm et al., 2013; Ceccarelli et al., 2014). Atlantic yellowfin tuna (Thunnus albacares) are overfished but not experiencing overfishing (ICCAT, 2016b). Recently, the western stock of Atlantic bluefin tuna (Thunnus thynnus) were declared to not be experiencing overfishing and may no longer be overfished (ICCAT, 2016a). Previously, the stock was considered overfished and experiencing overfishing (Fromentin and Powers, 2005; ICCAT, 2014b), and possibly at risk of collapsing (Bjørndal and Brasão, 2006; Safina and Klinger, 2008; MacKenzie et al., 2009). Atlantic marlin (Makaira nigricans, and Kajikia albidus) are considered overfished, and possibly risk extinction, due to mortality experienced from recreational fisheries, which target marlin, pelagic longline commercial fisheries, which incidentally catch marlin, and artisanal fleets, which target marlin (Peel et al., 2003; ICCAT, 2011, 2012).

Management of pelagic predatory stocks is both a domestic and international effort as these species are highly mobile. In the United States, the National Oceanic and Atmospheric Administration (NOAA) Fisheries Program, known as the National Marine Fisheries Service (NMFS), has primary authority for developing and implementing a Fishery Management Plan (FMP) for highly mobile species (HMS) in federal waters of the Atlantic. Such FMPs have enacted input and output controls (National Oceanic and Atmospheric Administration, 2016a), including fishery area closures. Fishery time and area closures, a type of marine protected area (MPA) within which fishing is limited and/or prohibited, have been recommended by management agencies and stakeholders as a viable management option for protecting pelagic predators. The American Fisheries Society (AFS) and the International Commission for the Conservation of Atlantic Tunas (ICCAT) recommend the development, use, and evaluation of large time and area closures to protect and rebuild shark populations (Musick et al., 2000a,b; ICCAT, 2007, 2009, 2010). In 2008 NMFS implemented time/area closures proposed by the South Atlantic Fishery Management Council (SAFMC) to protect and rebuild shark stocks (Highly Migratory Species Division, 2008). ICCAT recommends the consideration of area and/or time restrictions to prevent directed fishing on the bluefin tuna spawning stock within the western Atlantic spawning grounds (i.e., the Gulf of Mexico) (ICCAT, 2014a). Peel et al. (2003) concluded that coupling time and area closures with some restraint on targeted effort may help rebuild some billfish stocks.

The Gulf of Mexico is a key area in which to consider fishery closures because the physical dynamics and topology make for a productive system for Atlantic pelagic predators. The pelagic waters consist of a dynamic network of cyclonic and anticyc-
lonic features (both filaments and eddies) primarily due to the physical forcing of the Loop Current and the episodic shedding of a warm-core, anti-cyclonic Loop Current eddy (Wiseman et al., 1999; Oey et al., 2005). Cyclonic features, which create patches of nutrient upwelling, retention, and concentration, are intensified by these dynamics (Schmitz, 2005), enhancing favorable environments for foraging planktonic larvae (Bakun, 1996). Bluefin tuna migrate into the northwest Gulf of Mexico between April and June to spawn (Dicenta et al., 1980; Richards, 1990; Mather et al., 1995; Block et al., 2005). Yellowfin tuna (Thunnus albacares) spawn in the northern Gulf of Mexico between July and August (Lang et al., 1994). Billfish, particularly sailfish (Istiophorus albicans) and blue marlin (Makaira nigricans), are believed to use the northern Gulf of Mexico as spawning grounds and early life habitat (Rooker et al., 2012). Spawning season can vary amongst billfish species (de Sylva and Breder, 1997). Although some shark species are more common within the Gulf's expansive shelf, which is used for migration (Branstetter, 1981, 1987), foraging (Hoffmayer and Parsons, 2003; Hammerschlag et al., 2012), and to access nurseries and pupping grounds (Bethea et al., 2006; Hueter and Tyminski, 2007), shark species are also encountered in the Gulf's pelagic environment (Branstetter, 1981; Beerkircher et al., 2002; Cortés, 2002a), and it is possible that these species aggregate around fronts (Queiroz et al., 2012, Chapter 3 of this dissertation).

The Gulf of Mexico's U.S. pelagic longline fleet targets swordfish, yellowfin tuna, skipjack tuna, albacore, dolphin fish, wahoo, and some coastal sharks, while incidentally catching bluefin tuna, billfish, miscellaneous tunas (e.g., some bigeye tuna), some pelagic sharks (e.g., blue sharks), prohibited sharks (e.g., hammerheads), sea turtles, seabirds, and mammals (Highly Migratory Species Division, 2000). To mitigate the
incidental catch of non-targeted species (a.k.a, bycatch), two pelagic longline fishery closures are currently established in the Gulf of Mexico. The first is a permanent closure around DeSoto Canyon with the primary management objectives to reduce the catch and rebuild the biomass of groups caught incidentally while having little impact on the catch of targeted species (Highly Migratory Species Division, 2000). The second is a seasonal closure off the Louisiana shelf with the primary management objectives to reduce the catch and rebuild the biomass of bluefin tuna (Highly Migratory Species Division, 2014). Although it was originally thought that MPAs would provide little benefit to pelagics due to their high mobility and weak site fidelity (Roberts, 1997; Boersma and Parrish, 1999), more recent research supports spatial closures as viable tools for mitigating bycatch, including bycatch of pelagic species (Goodyear, 1998; Grantham et al., 2008; Dunn et al., 2011), but there is uncertainty regarding the success of pelagic spatial closures.

Hyrenbach et al. (2000) suggested pelagic spatial closures may be feasible tools for pelagic conservation since the physical habitats highly mobile predators aggregate around tend to be spatially and temporally predictable. More recent research supports this and argues that pelagic MPAs are defensible tools for pelagic conservation due to the advances in conservation, oceanography, and fisheries science (Game et al., 2009, 2010). However, there are still concerns regarding the utility and feasibility of pelagic MPAs. First, areas must be identified within which pelagics of concern have high site fidelity, and practical enforcement plans must be developed (Kaplan et al., 2010). Second, it is not known if pelagic spatial closures will provide the same benefits as some coastal spatial closures, like providing biomass to the fisheries through spillover (e.g., McClanahan and Mangi, 2000; Kelly et al., 2002; Guidetti, 2007; Januchowski-

Hartley et al., 2013) or increasing the size of individuals (e.g., Babcock et al., 1999; Lester et al., 2009). Lastly, MPAs often fail to meet management objectives (Jameson et al., 2002). Thus, science-driven analysis, including the investigation of ecosystem impacts through mathematical modeling, should be done to address the utility and feasibility of pelagic MPAs (Kaplan et al., 2010; Game et al., 2010; Grüss, 2014).

For this study, a policy exploration was conducted to investigate i) if Gulf of Mexico pelagic longline fishery spatial closures are likely to achieve management objectives, and ii) potential ecosystem impacts from pelagic longline closures. An ecosystem model of the Gulf of Mexico was used to simulate scenarios and calculate performance measures (indicators) corresponding to management objectives of the pelagic longline fishery spatial closures, as well as broader ecological objectives. Performance metrics were then compared to evaluate potential long-term impacts of Gulf of Mexico pelagic longline spatial closures.

### 4.3 Methods

### 4.3.1 The Simulation Framework

Atlantis is a biogeochemical and biophysical modeling framework (Fulton et al., 2004c, b, 2011). It models the turnover of chemical substances through the biotic and abiotic compartments of an ecosystem, in addition to the biological and physical components. The Atlantis framework is appropriate for this study in many ways. First, Atlantis was developed with the intention to evaluate performance measures (i.e., indicators) for use in ecosystem-based fisheries management (Fulton et al., 2004a; Plagányi, 2007; Fulton et al., 2011). Second, Atlantis is an 'end-to-end' model (Fulton,
2010), meaning it represents biota from bacteria up to top predators in addition to human activities (e.g., fisheries). Third, species and fisheries interactions are spatially explicit. Fourth, the spatial domain is represented by a network of polygons that reflect the ecosystems geographic features, habitats, and essential management jurisdictions. Lastly, Atlantis contains a detailed exploitation routine which allows for the simulation of individual fleets, as well as a management routine to simulate a range of fishery management measures, including spatial closures.

The Gulf of Mexico Atlantis Model (GoMAM) is described in detail by Ainsworth et al. (2015), so the following will be a summary of the model. The spatial domain of GoMAM consists of a polygon network spanning the entire Gulf of Mexico marine ecosystem (Figure 4.2), which was developed based on bathymetry, habitat, physical oceanography, and management boundaries. The simulated biology consists of 91 functional groups of finfish, invertebrates, seabirds, mammals, plankton, and bacteria/detritus. Vertebrate functional groups have 10 age classes and all remaining groups are represented as biomass pools. The flux of nitrogen for primary producers, biomass pools, and age-structured groups are modeled differently.

Spawning and recruitment are only explicitly modeled for age-structured groups. Reproduction of simple biomass pools is included in the growth terms. For agestructured groups, nitrogen produced as spawn is temporarily removed from the system and returned as recruits after a defined larval period. In GoMAM, for tuna, billfish, and sharks this period is about a month. The stock recruitment curve assumed is the Beverton-Holt with recruitment based only on species biomass. This is for all functional groups except mammals, birds, and large sharks, which are assumed to have a constant recruit production per adult. The assumed vertical distribution
of recruits is the same as the daytime vertical distribution of juveniles. The assumed horizontal distribution of recruits is the same as the horizontal distribution of adults in the first season.

GoMAM was parameterized for forecasting based on the parameterization of a historical version of GoMAM, which was driven by historical landings time series (summarized in Chapter 1 of this dissertation, and discussed in detail by Perryman et al. (2015)), calibrated to fit historical abundance trends from 1980 to 2010. Allocating the biomass of functional groups across space is important for spatially explicit models like Atlantis (Grüss et al., 2016a). Biomass of demersal functional groups were spatially allocated using statistical models presented by Drexler and Ainsworth (2013). Statistical models presented in Chapter 2 of this dissertation were used to refine the spatial allocation of pelagic fish groups as they were originally based on somewhat homogenous assumptions.

Sedentary functional groups in GoMAM do not have horizontal movement amongst polygons. For mobile functional groups, GoMAM is set-up to simulate "prescribed movement". This means density dependent movement is not allowed and instead mobile functional groups move based on distributions that define the horizontal shift a functional groups throughout the year. GoMAM prescribes quarterly shifts, and quarterly shifts differ between adults and juveniles. Atlantis calculates the abundance ( $A$, in biomass for biomass pool groups and numbers for age structured groups) of a functional group in a polygon $(p)$ at any given time-step as

$$
\begin{equation*}
A_{p}=A_{\text {total }}\left(\varepsilon\left(F_{(Q+1), p}-F_{Q, p}\right)+F_{Q, p}\right) \tag{4.1}
\end{equation*}
$$

where $\varepsilon$ is the proportion of the quarter of the year that has passed (12 hour time steps), and $F_{Q, p}$ is the proportion of biomass in polygon $p$ during quarter $Q$ provided by the prescribed parameters. If $Q$ is the last quarter of the year then $Q+1$ is the first quarter of the next year. Atlantis can also simulate density dependent movement, which means that the spatial distribution will be determined by the food availability, but GoMAM has density dependent movement deactivated for all functional groups. Thus, any changes in catch outside fishery closures are because of changes in total biomass and not changes to local biomass due to groups responding to changes in food availability.

Simulated functional groups are allowed to migrate out of and into the Gulf of Mexico modeling domain. Abundance of functional groups migrating out of the Gulf of Mexico system are stored in a boundary polygon until the time is reached which the abundance starts to migrate back into the model domain. The abundance of functional groups while outside of the modeling domain is allowed to change (i.e., this biomass is subjected to mortality, growth, etc.). In GoMAM, migration inputs have been parameterized for functional groups that correspond to mammals, birds, sea turtles, large sharks, mackerels, billfish, yellowfin tuna, and bluefin tuna (Ainsworth et al., 2015). This is particularly important for bluefin tuna as they are a species of concern for pelagic longline fishery closures.

In GoMAM, migration inputs for adult bluefin tuna (age-classes 2-10) differ from those for juvenile bluefin tuna (age-class 1). Bluefin tuna are outside the modeling domain throughout most of the year. Migration of mobile functional groups occurs gradually rather than all at once. For bluefin tuna, the migration of juveniles spans a couple of months, while the migration of adults spans a couple of days. Some mortality
is applied to juveniles while outside the system in that $4 \%$ of the abundance does not return. Mortality outside of the system is currently not applied to adults. The abundance of juveniles and adults increases while groups are outside of the modeling domain, which adds nitrogen to the system. Preliminary work was done to adjust some of the migration inputs of bluefin tuna to reflect observations in the literature, but these alterations were not used for this study because they caused bluefin tuna to quickly collapse (Appendix C).

### 4.3.2 Gulf of Mexico Pelagic Longline Closures

The Gulf of Mexico has two fishery closures pertaining to pelagic longline operations: the DeSoto Canyon Pelagic Longline Closure, and the Spring Gear Restricted Areas (Figure 4.1).

### 4.3.2.1 DeSoto Canyon Pelagic Longline Closure

DeSoto Canyon is a valley that cuts through the broad continental shelf in the northeast Gulf of Mexico. The area's bathymetry and physical forces driven by the Loop Current interact to form cyclonic eddys that upwell cool, nutrient-rich water. This causes relatively high primary productivity (Vukovich and Maul, 1985; Hamilton et al., 2000a,b; Yuan, 2002). During a comment period for a proposed highly migratory species bycatch rule, NMFS received many comments indicating that the DeSoto Canyon area should be closed to pelagic longlining due to the high occurrence of undersized swordfish (Highly Migratory Species Division, 2000). Based on an assessment of logbook data, two areas were identified and approved as a year-round pelagic longline fishery closure that went into effect November 1, 2000. A formal assessment
of the effectiveness of DeSoto Canyon Pelagic Longline Closure, from here on referred to as DeSoto Canyon, has yet to be conducted. Landings data presented in Chapter 2 (Appendix A) suggests some evidence of decreased landings of some indicator species after DeSoto Canyon was enacted.

### 4.3.2.2 Spring Gear Restricted Areas

To protect the Atlantic bluefin tuna stock while spawning in the Gulf of Mexico, NMFS established a seasonal pelagic longline fishery closure (Highly Migratory Species Division, 2014). Several configurations were considered, but the final amendment establishing the Spring Gulf of Mexico Gear Restricted Areas consisted of two areas: one large area spanning the northwestern Gulf, and another smaller area bordering the DeSoto Canyon closure. Starting in 2015, from April 1 to May 31, pelagic longline operations are prohibited within Spring Gulf of Mexico Gear Restricted Areas, from here on referred to as Spring Closure.

### 4.3.3 Simulated Scenarios

Simulations start at the conditions from the end of the model fitted to historical data (2010) and spanned 30 years, under constant fishing mortality rates, because this was long enough to capture significant changes in the simulated ecosystem, and short enough to save on computation time. Results consist of comparing 30 year projections under the status quo scenario to the projections under other modeled scenarios. The study considers two types of scenarios: (1) fishing mortality sensitivity scenarios (the increasing and decreasing of fishing mortality rates), and (2) pelagic longline fishery closure scenarios (evaluation of pelagic longline fishing closures DeSoto Canyon and

Spring Closure). All scenarios have the same parameterized biology, ecology, and oceanography. The only variations were changes to fisheries.

### 4.3.3.1 Status Quo

This scenario allows the evaluation of system dynamics under a baseline representation of current fisheries. This includes DeSoto Canyon and Spring Closure. Fishery closures are modeled by reducing fishing mortality rate(s) within the polygon(s) corresponding to the closure's location. Often, polygons in the simulation map do not match a closure's geometry. The Intersect tool in ArcGIS was used to determine the polygon(s) covering the spatial range of a closure, and the percentage of the polygon corresponding to the closure. This information was used to develop the polygon-specific input files for simulating fishing closures, which, once a designated time step is reached in the simulation, reduce a fleet's fishing mortality within the polygon based on the regulation and the percentage of closure within the polygon. For instance, if a MPA occupies half of a polygon and the regulation closes pelagic longline fishing, then the pelagic longline fleet's fishing mortality is reduced by half within that polygon. This method does not effect fishing mortality rates in polygons that do not correspond to a closure, thus the underlining assumption here is that closed areas remove the effort that would have been in the closure. Input files were updated to include Spring Closure as it was not represented in the original calibration of GoMAM because the closure wasn't activated before 2010.

### 4.3.3.2 Fishing Mortality Sensitivity

Longline fishing mortality sensitivity scenarios are to i) evaluate the overall impact of longline fishing on the Gulf ecosystem in order to put the impact of closures into context, and ii) make sure the GoMAM behaves reasonably under large perturbations (Kaplan et al., 2012). Sensitivity scenarios consisted of two types: all longline, and pelagic longline. The former focused on all longline fleets in GoMAM, including shelf longline (reef fish), shelf longline (shark), and pelagic longline. The latter only focused on the pelagic longline fleet. These two types of sensitivity scenarios quantify the pressure pelagic longline fleets exert on pelagic predators compared to all longline fleets. Both of these two types of simulations consisted of 3 scenarios in which the fishing mortality of the indicated fleet(s) was multiplied by $0,0.5$, and 2 . Thus, in total, there are 6 fishing mortality sensitivity scenarios.

Some aspects of GoMAM's fisheries were not adjusted for this study. GoMAM contains 22 fleets (16 U.S. fleets, 5 Mexican fleets, and 1 Cuban fleet), and assumes the selectivity curve is logistic for all functional groups. Fishing mortality rates were altered for this study. Removal of biomass due to fishing is simulated with a series of constant, daily fishing mortality rates reflecting the pressure each fleet exerts on each functional group. Fishing mortality rates were originally computed using 2010 landings data (Perryman et al., 2015), but these did not include data on bycatch. To represent pelagic longline bycatch, GoMAM fishing mortality rates were updated using data describing the 2010 bycatch (National Marine Fisheries Service, 2013). In addition, these updated rates were iteratively adjusted to match 2010 simulated catch to 2010 catch data. For full details see Appendix C.

### 4.3.3.3 Pelagic Longline Closures

Because pelagic longline spatial closures DeSoto Canyon and Spring Closure are represented in the status quo scenario, to investigate their impacts to the system scenarios were simulated in which the two closures were removed from the status quo version of GoMAM. This includes a scenario in which DeSoto Canyon was removed, a scenario in which Spring Closure was removed, and a scenario in which both DeSoto Canyon and Spring Closure were removed. This allowed the assessment of individual impacts, as well as the possibility of compounding impacts. In addition, another scenario was investigated in which Spring Closure was altered to span the entire U.S. Gulf - simulating a seasonal, Gulf-wide closure of the U.S. pelagic longline fleet. This was an alternative measure considered instead of Spring Closure, but ultimately Spring Closure was preferred due to the estimated ecological gains with low fisheries impact (Highly Migratory Species Division, 2014). Thus, in total, there are 4 pelagic longline closure scenarios.

### 4.3.4 Management Objectives and Performance Measures

The primary management objectives for DeSoto Canyon are to i) reduce bycatch and incidental catch, ii) minimize the reduction in target catch, and iii) optimize survival of bycatch and incidental catch species (Highly Migratory Species Division, 2000). Performance measures for these objectives were tracked for the appropriate GoMAM functional groups. U.S. pelagic longliners target swordfish, yellowfin tuna, bigeye tuna, skipjack tuna, albacore, dolphin fish, wahoo, and some coastal sharks (Highly Migratory Species Division, 2000). The corresponding GoMAM functional groups are: swordfish, yellowfin tuna, other tuna, large pelagic fish, and large sharks.
U.S. pelagic longline incidental catch (organisms that are not targeted but may be retained if caught) primarily consists of bluefin tuna (Highly Migratory Species Division, 2000), which has its own GoMAM functional group. Catches of marine mammals and sea birds, which are always either released alive or discarded dead, were not evaluated here since more work is necessary for GoMAM to appropriately represent their spatial heterogeneity and bycatch. U.S. pelagic longline discarded bycatch (organisms caught but not retained) includes: Atlantic billfish, undersized swordfish, bigeye tuna, pelagic sharks (e.g., blue sharks), prohibited sharks, and sea turtles (Highly Migratory Species Division, 2000; National Marine Fisheries Service, 2013). The corresponding GoMAM functional groups are: white marlin, blue marlin, billfish, swordfish, other tuna, large sharks, loggerhead, Kemp's ridley, and other sea turtles.

Reducing the catch of undersized swordfish caught by U.S. pelagic longlines was a specific concern (Highly Migratory Species Division, 2000). Undersized swordfish caught by U.S. pelagic longlines are individuals weighing less than 25 kg whole weight (Cramer, 1996). In GoMAM, vertebrate functional groups, including swordfish, are represented with age structured models. The 10 age-groups were adjusted to cover the whole lifespan of the fish. In GoMAM, swordfish are first selected by pelagic longline gears by age-group 3. The median weight of swordfish of age-group 3 is 23.2 kg , so the parameterization of GoMAM is consistent with the fishery data, and catch of swordfish of age-group 3 was tracked to represent catch of undersized swordfish. The parameterization of some model inputs can be different between adult age-groups and juveniles age-groups (e.g., seasonal spatial distribution, migration, diet, etc.). GoMAM assumes swordfish become sexual maturity at age-group 3, thus
all of the age-groups selected by the fishery are considered adults and have the same parameterization.

Performance measures tracked for the reduction of bycatch and incidental catch included: catch of bluefin tuna, total catch of all bycatch groups, catch of individual bycatch groups, and proportion of swordfish catch being age-group 3 (i.e., age-at-firstcapture). Performance measures tracked to minimize the reduction in target catch included: the total catch of all target groups, and catch of individual target groups. Performance measures tracked for optimizing the survival of bycatch and incidental catch species included: the total biomass of all bycatch groups, biomass of individual bycatch groups, and biomass of bluefin tuna. For consistency, total biomass of target groups, and biomass of individual target groups were also monitored. See Table 4.1 for details.

The FMP amendment proposing Spring Closure reports a variety of socioeconomic, fisheries quota, and biological objectives (Highly Migratory Species Division, 2014), the primary objective being to prevent overfishing and rebuild bluefin tuna. The primary objective for the gear restricted areas was to reduce bluefin tuna interactions with pelagic longliners, thereby decreasing dead discards (bycatch). Thus, this work tracked the reduction of incidental catch of bluefin tuna using bluefin tuna catch as a performance measure, and the rebuilding of bluefin tuna using bluefin tuna biomass as a performance measure.
U.S. fleets targeting highly migratory pelagics, in addition to the pelagic longline fleet, include U.S. recreational groups, and U.S. hook-and-line fleets not using pelagic longling gear (e.g., fleets using vertical lines, or bottom longlines). It is possible that changes to the U.S. pelagic longline fishing may change catches from these fleets.

To evaluate this, total catches of tuna billfish, and large shark functional groups from simulated U.S. recreational fleets and non-pelagic, U.S. hook-and-line fleets (i.e., handline, shelf longline (reef fish), and shelf longline (shark)) were compared across the investigated scenarios.

Ecological objectives were also monitored to identify if closures could have broad scale ecosystem impacts, some of which have been observed for coastal spatial closures. See Table 4.1 for a summary of indicators and equations. In many fished ecosystems, the size of individuals, particularly top predators, has decreased over time due to heavy exploitation and fishing practices such as minimum size regulations focusing fishing pressure on larger organisms (Bianchi et al., 2000; Swain et al., 2007; Darimont et al., 2009). This could impact food webs and trophic structure (Woodward et al., 2005; Brose et al., 2006; Shackell et al., 2010). The reduced number of larger individuals also means a reduced number of sexually mature individuals, which can have a negative impact on spawning (Hutchings, 2000). As discussed in the introduction, spatial closures can mitigate and potentially reverse these impacts. The metric proportion of mature fish (biomass) was used to track shifts in the amount of sexually mature stock. The metric average individual size was used to track the size structure of functional groups. Changes to the average weight of multi-species functional groups should be interpreted with caution as they are composed of different species with different growth trajectories. The proportion of mature fish and size related metrics, such as average individual size, are important indicators of overfishing (Froese, 2004; Shin et al., 2005). These two stock-specific metrics were computed for individual pelagic predator functional groups, and functional group assemblages (e.g., billfishes, tunas, sharks, etc.). Results discussed in the main text will focus on
billfishes and tunas, and the corresponding functional groups. Additional results of other functional groups are shown in Appendix C.

Objectives relating to the ecosystem community were also monitored. The pelagic:demersal ratio was tracked for the catch and for the ecosystem. The pelagic:demersal ratio is primarily linked to the eutrophication (Caddy, 1993; Caddy and Bakun, 1994; Caddy et al., 1998a; Caddy and Garibaldi, 2000; Caddy, 2000; de Leiva Moreno et al., 2000), and the Gulf of Mexico experiences periodic, large-scale eutrophication which has meaningful ecosystem impacts (Malakoff, 1998; Rabalais et al., 2002b,a). In addition, the pelagic:demersal ratio can show large shifts in fishery targeting, and Fulton et al. (2005) found it to be strongly correlated with a marine ecosystems population and community attributes. Functional groups were categorized as pelagic or demersal/benthic based on life history of adults. Information from FishBase (Froese and Pauly, 2016) and SealifeBase (Palomares and Pauly, 2016) was used to classify individual species and Atlantis functional groups. The ratio was calculated as the total biomass of pelagic groups divided by the total biomass of demersal groups.

Marine ecosystem biodiversity was also monitored. Marine protected areas are used as a means to restore and/or preserve biodiversity since anthropogenic impacts (i.e., overfishing) can diminish biodiversity (Coleman and Williams, 2002; Jones et al., 2007) which can reduce ecosystem resources, resilience, and water quality (Duffy, 2002, 2003; Worm et al., 2006; Stachowicz et al., 2007; Cardinale et al., 2012). Much of the research on using MPAs to protect biodiversity is from coastal systems, but there is growing discussion regarding the use of pelagic MPAs to protect biodiversity (Worm et al., 2003; Game et al., 2009; Morato et al., 2010; Grantham et al., 2011). The Q90 biodiversity statistic was used to monitor marine ecosystem biodiversity.

Q90 is the Kempton and Taylor (1976) species diversity statistic adapted for use with ecosystem models (Ainsworth and Pitcher, 2006), which aggregate individual species into functional groups. Q90 is the interdecile slope of the cumulative log-abundance curve:

$$
\begin{equation*}
Q 90=\frac{0.8 S}{\log \left(R_{2} / R 1\right)} \tag{4.2}
\end{equation*}
$$

where $S$ is the total number of functional groups in the model, and $R_{1}$ and $R_{2}$ are the biomass values of the 10th and 90th percentiles in the cumulative abundance distribution across all functional groups.

Performance measures were calculated as the average of the values spanning the last four time steps, which are the four seasons of the last year of the simulation. All performance metrics were computed at multiple spatial scales: Gulf-wide (the entire spatial map), U.S. Gulf (all polygons within the U.S. EEZ), U.S. pelagic (all polygons within the U.S. EEZ, and a max depth greater than 200m), DeSoto Canyon waters (all polygons intersecting DeSoto Canyon), and Spring Closure waters (all polygons intersecting Spring Closure). This is allows the assessment of the spatial extent of the impacts of management measures.

### 4.4 Results

Performance measures for management objectives are summarized in Table 4.2. DeSoto Canyon caused meaningful reductions on catches within the Desoto Canyon region. Within the DeSoto Canyon polygons, removing the DeSoto Canyon closure increased catch of bycatch tunas by $53.6 \%$, bycatch billfishes by $37.2 \%$, and incidental bluefin tuna by $63.0 \%$, compared to doubling pelagic longline fishing mortality which
increased catch of bycatch tunas by $89.5 \%$, bycatch billfishes by $88.2 \%$, and incidental bluefin tuna by $52.0 \%$ within the DeSoto Canyon polygons. Within the polygons around Spring Closure, removing Spring Closure increased catch of bycatch tunas by $6.4 \%$, bycatch billfishes by $7.9 \%$, and incidental bluefin tuna by $1.5 \%$, compared to doubling pelagic longline fishing mortality which increased catch of bycatch tunas by $89.5 \%$, bycatch billfishes by $89.7 \%$, and incidental bluefin tuna by $19.7 \%$.

At the scale of the whole U.S. Gulf, removing DeSoto Canyon increased catch of bycatch tunas by $18.8 \%$, bycatch billfishes by $3.1 \%$, and incidental bluefin tuna by $10.2 \%$. Removing Spring Closure increased catch of bycatch tunas by $3.4 \%$, bycatch billfishes by $0.7 \%$, and incidental bluefin tuna by $0.6 \%$. When compared to doubling pelagic longline fishing mortality, which increased catch of bycatch tunas by $89.9 \%$, bycatch billfishes by $15.7 \%$, and incidental bluefin tuna by $39.9 \%$, it suggests that DeSoto Canyon is more effective than the Spring Closure at reducing U.S. Gulf catch of bycatch groups and incidental bluefin tuna. Removing DeSoto Canyon increased U.S. Gulf catch of target groups by $0.7 \%$, while removing Spring Closure increased U.S. Gulf catch of target groups by $0.1 \%$. When compared to doubling pelagic longline fishing mortality, which increased U.S. Gulf catch of target groups by $3.8 \%$, it is apparent that DeSoto Canyon and Spring Closure did not have meaningful impacts on U.S. Gulf catch of target groups.

DeSoto Canyon and Spring Closure had smaller impacts on U.S. Gulf biomass. Doubling pelagic longline fishing mortality decreased U.S. Gulf biomass of bycatch tunas by $4.2 \%$, bycatch billfishes by $0.7 \%$, incidental bluefin tuna by $1.7 \%$, and target groups by $0.8 \%$. Removing DeSoto Canyon decreased U.S. Gulf biomass of bycatch tunas by $0.9 \%$, bycatch billfishes by less than $0.01 \%$, incidental bluefin tuna by $0.4 \%$,
and target groups by less than $0.01 \%$. Removing Spring Closure decreased U.S. Gulf biomass of bycatch tunas by less than $0.01 \%$. The biomass of incidental bluefin tuna was more affected by simulated scenarios within Spring Closure polygons than DeSoto Canyon polygons or U.S. Gulf polygons. For instance, eliminating pelagic longline fishing increased the U.S. Gulf biomass of bluefin tuna $4.4 \%$, the DeSoto Canyon biomass of bluefin tuna $4.0 \%$, and the Spring Closure biomass of bluefin tuna $13.3 \%$. This is likely related to the fact that Spring Closure waters include a hot spot for bluefin tuna (see spatial distribution maps in Appendix B).

The impact DeSoto Canyon had on management objective performance measures varied amongst species (Figure 4.3). Within DeSoto Canyon waters, the presence of the DeSoto Canyon closure increased the biomass of miscellaneous tunas, green turtles, yellowfin tuna, and swordfish (Figure 4.3a, 4.3b). Increases in biomass were visible across spatial scales of the Gulf, especially for green turtles, miscellaneous tunas, and swordfish. Also within DeSoto Canyon waters the closure was associated with reductions in catches of billfish groups, miscellaneous tunas, bluefin tuna, green turtles, yellowfin tuna, and swordfish (Figure 4.3c, 4.3d). Gulf-wide catch of yellowfin tuna increased with the presence of DeSoto Canyon (Figure 4.3d).

Within Spring Closure waters, Spring Closure increased the biomass of yellowfin tuna but had very little effect on the biomass of other functional groups (Figure 4.4a, 4.4b). The biomass of bluefin tuna did not change within Spring Closure waters, but there was a slight increase across the Gulf ecosystem (Figure 4.4a). Within Spring Closure waters, catches of billfish groups, miscellaneous tunas, green turtles, swordfish, and yellowfin tuna were reduced (Figure 4.4c, 4.4d). Gulf-wide catches of
miscellaneous tunas, green turtles, and swordfish decreased, but catches of yellowfin tuna increased within Gulf-wide polygons as well as Spring Closure polygons.
U.S. pelagic longline effort and U.S. pelagic longline spatial closures impact catches of U.S. hook-and-line fleets other than pelagic longliners (Figure 4.5). Spring Closure had little impact on the catches from non-pelagic U.S. commercial hook-and-line fleets, but DeSoto Canyon increased the catches of billfishes from non-pelagic U.S. commercial hook-and-line fleets by $2.8 \%$, compared to an increase of $4.6 \%$ when U.S. pelagic longline fishing is reduced by half, or $9.6 \%$ when U.S. pelagic longline fishing is eliminated. In addition, DeSoto Canyon increased recreational catches of billfishes by $1.4 \%$ and tunas by $1.6 \%$. Eliminating U.S. pelagic longline fishing caused a $5.3 \%$ increase in recreational billfish catch, and a $14.1 \%$ increase in recreational tuna catch. None of the closures reduced the catch of non-pelagic U.S. hook-and-line fleets, but increasing U.S. pelagic longline fishing mortality did. In particular, there was a $8.2 \%$ reduction in billfish catch from non-pelagic U.S. commercial hook-and-line fleets, and a $7.7 \%$ reduction in tuna catch from recreational fleets when U.S. pelagic longline fishing mortality was doubled.

Results for ecosystem performance measures are shown in Table 4.3. Spring Closure had less influence on ecosystem metrics than DeSoto Canyon. Removing DeSoto Canyon decreased billfish U.S. Gulf average individual weight by less than $0.01 \%$, compared to a $0.3 \%$ reduction when U.S. pelagic longline fishing mortality is doubled. In addition, removing Desoto Canyon decreased tuna U.S. Gulf average individual weight by $0.2 \%$, compared to a $0.8 \%$ reduction when U.S. pelagic longline fishing mortality is doubled. Impacts to average individual weight and proportion mature are species specific (Figure 4.6). A reduction in pelagic longline fishing mortality had
some influence increasing U.S. Gulf average individual weight for swordfish, but no influence on other billfish groups. Reducing pelagic longline fishing mortality had the most influence on the U.S. Gulf average individual weight of tunas: increasing it for miscellaneous tunas and yellowfin tuna, and decreasing it for bluefin tuna. Reducing U.S. pelagic longline fishing mortality increased U.S. Gulf proportion mature for all billfish and tuna groups, with bluefin tuna experiencing the largest increase (53.4\%). Removing DeSoto Canyon increased U.S. Gulf catch pelagic:demersal ratio by $0.6 \%$, compared to a $2.8 \%$ reduction when U.S. pelagic longline fishing mortality was doubled. In addition, removing DeSoto Canyon decreased U.S. Gulf ecosystem pelagic:demersal ratio by $0.2 \%$, compared to a $1.1 \%$ reduction when U.S. pelagic longline fishing mortality is doubled. None of the scenarios changed the ecosystem's Q90 biodiversity metric.

### 4.5 Discussion

### 4.5.1 Model Findings

Model simulations show that the Gulf of Mexico pelagic longline spatial closure DeSoto Canyon, which has been in affect since 2000, could be achieving some management objectives. Considering U.S. Gulf metrics, first, DeSoto Canyon reduced catch of the incidental group, bluefin tuna. Total catch of bycatch groups did not change, but there were meaningful changes on a species-specific level. Specifically, catch of functional groups representing bycatch tunas, and green sea turtles were reduced. In addition, DeSoto Canyon reduced the proportion of swordfish catch being of ageclass 3 (age-at-first-capture). Second, total catch of target groups was reduced only
slightly. Thus, DeSoto Canyon did not negatively impact the catch of target species. In fact, DeSoto Canyon increased the Gulf-wide catch of yellowfin tuna, suggesting that spill over is occurring, and increased the biomass of yellowfin tuna. It would be expected to have increased catch or increased biomass, but having both is peculiar. This could be explained by the possibility of the stock being overfished in the model, which would be consistent with the latest stock assessment (ICCAT, 2016b). DeSoto Canyon also increased the biomass swordfish and miscellaneous tunas. Lastly, biomass of the incidental group bluefin tuna increased slightly, and, although total biomass of bycatch groups did not change, the biomass of groups representing bycatch tunas and green sea turtles increased slightly.

DeSoto Canyon had some influence on ecosystem objective performance metrics. Considering U.S. Gulf metrics, DeSoto Canyon increased the average weight of tunas and billfishes, particularly yellowfin tuna, miscellaneous tunas, and swordfish. Both density and biomass of these groups increased but biomass increased more, suggesting that individuals could be getting larger. The exception is bluefin tuna, which decreased in average individual weight because density increased more than biomass. DeSoto Canyon slightly increased the proportion of billfishes and tunas sexually mature. Fishing mortality sensitivity scenarios showed that bluefin tuna are especially sensitive - their proportion mature increased more than any other group. DeSoto Canyon reduced catch pelagic:demersal ratio because pelagic catches were reduced with little influence on demersal catches. In response, the ecosystem pelagic:demersal ratio increased.

None of the scenarios had a meaningful influence on ecosystem biodiversity. It is possible that these scenarios may not have enough impact on simulated biota to cause
an impact to biodiversity since pelagic longline fleets interact with a small portion of the total functional groups. While scenarios have an impact on some functional groups in the pelagic waters, benthic and microbial functional groups in the same areas are not impacted. These functional groups are abundant, and could be buffering the Q90 calculation. Also, the Q90 metric is being computed over large spatial ranges, which may be masking a signal if one is there. The Q90 calculation can be made more sensitive by passing a filter over the calculation to omit the biomass of functional groups if their biomass falls below a reference value (Ainsworth and Pitcher, 2006). This would require defining reference values for each functional group, as well as a threshold. More research is needed to fill the gap of our understanding of how pelagic MPAs may preserve biodiversity (Game et al., 2009; Grantham et al., 2011)

Spring Closure had very little influence on performance measures corresponding to its management objectives. Considering U.S. Gulf metrics, as well as those from Spring Closure waters, Spring Closure only slightly reduced catch of bluefin tuna, and did not change bluefin tuna biomass. These results suggest that Spring Closure, which has only been in affect since 2015, may not meet biological management objectives over the next couple of decades. In addition, Spring Closure had little to no influence on ecological objective performance measures. These results could be a realistic reflection of the Gulf of Mexico due to the state and dynamics of the bluefin tuna stock as a whole, or this could reflect possible model limitations. It is worth noting that Spring Closure influenced performance metrics relating to DeSoto Canyon management objectives. Specifically, Spring Closure reduced the U.S. catch of bycatch and incidental groups, and increased the biomass and catch of yellowfin tuna. This shows that DeSoto Canyon and Spring Closure had compounding im-
pacts on performance measures. Thus, objectives were more attainable when both closures were active. Networks of marine protected areas have shown to be successful management strategies (Balmford et al., 2004; Russ et al., 2008; Gaines et al., 2010).

Fishing mortality sensitivity scenarios reveal some important dynamics. First, pelagic fishery closures are not providing enough reduction in fishing mortality to cause meaningful increases in biomass. For instance, removing Spring Closure decreased bluefin tuna biomass around Spring Closure by 0.1\%, compared to a $5.3 \%$ reduction when pelagic longlining is doubled. Removing DeSoto Canyon decreased biomass miscellaneous tunas around DeSoto Canyon by $0.9 \%$, compared to a $4.4 \%$ reduction when pelagic longlining is doubled. Thus, increasing the biomass of indicator species may be dependent on complimentary management efforts further reducing fishing mortality (Allison et al., 1998; Myers and Worm, 2005). Second, alterations in longline fishing mortality influence bycatch and incidental groups more than target groups. For instance, decreasing longline fishing mortality increased the biomass of bycatch billfishes ( $0.7 \%$ ) and tunas (4.5\%), and incidental bluefin tuna (4.4\%), compared to a $1.4 \%$ increase to biomass of target groups. Thus, increasing longline effort had more negative impacts on bycatch and incidental groups than a gain from target catch. In addition, decreasing longline effort had more positive impacts on bycatch and incidental groups than loss of target catch. Target catch amongst sensitivity scenarios remained relatively stable due to catches from U.S. recreational fleets and U.S. non-longline, hook-and-line fleets since these fleets also target these groups (e.g., landings of recreational and non-longline, hook-and-line fleets increased with reductions in pelagic longline effort).

### 4.5.2 Limitations

Diagnostics of GoMAM show potential issues with the current calibration of bluefin tuna biology. First, during the simulation there is a sudden loss of bluefin tuna beyond the 3rd age-class. Although the western Atlantic bluefin tuna stock is severely depleted and it is possible very few older organisms are in the stock, the loss of these organisms in the simulated system was very sudden. Body weight diagnostics suggested it could be due to unbalanced consumption dynamics for bluefin tuna. However, adjusting parameters relating to bluefin tuna consumption caused additional problems to model dynamics (Appendix C). Second, GoMAM's modeling of the bluefin tuna seasonal migration could be improved. For instance, adults are in the simulated ecosystem longer than what is currently suggested in the literature (e.g. Block et al., 2005; Teo et al., 2007). Also, adult bluefin tuna are not subjected to additional mortality outside the system, although the Atlantic Ocean is where bluefin tuna experience significant mortality (ICCAT, 2014b, 2016a). Adjusting bluefin tuna migration, like making the seasonality stronger, improved some diagnostics but made other diagnostics worse (Appendix C). If future projects using the GoMAM model intend to focus specifically on bluefin tuna, it is recommended to adjust the treatment of the functional group in order to improve model diagnostics while still representing their known ecology. This will not be a trivial task since model tuning can directly and indirectly impact various components.

GoMAM's fisheries module could be advanced. For instance, this study reduced pelagic longline fishing mortality proportional to the area of the closure when, in reality, spatial closures may displace fishing effort (Kellner et al., 2007). The decision
to not redistribute fishing effort is an important limitation. This means indicator values for bycatch and incidental groups could be more optimistic, and indicator values for target groups could more pessimistic, than in reality since effort outside closures were not increased. Parameterizing GoMAM to model spatial closures with displaced fishing effort would require some work, but would benefit future fishery management investigations with GoMAM. It would also be beneficial to parameterize dynamic fisheries. Dynamic fisheries would be ideal especially for investigating Spring Closure, because spatial effort would respond to higher seasonal concentrations of stocks (e.g., bluefin tuna).

The fisheries module in GoMAM should also be advanced in order to improve the representation of bycatch. First, future fishery management studies with GoMAM would benefit if the model distinguished retained catches from discarded catches. Management objectives for pelagic fishery closures are not simply to reduce bycatch because this could be accomplished with lower quotas. In reality, these closures aim to reduce discards while preserving target catch, that is, increase the ratio between target catches and discarded catches. Currently, this can not be tracked in GoMAM since retained catches and discarded catches are not distinguished. Second, simulating the bycatch of sea turtles, birds, and mammals can be improved. Fairly homogenous spatial distributions were assumed for these groups, which could be improved with statistical models. Also, simple assumptions were made to parameterize fleet-specific bycatch of these groups, which could be improved by analyzing additional bycatch data.

### 4.5.3 Management Recommendations and Conclusions

Atlantic bluefin tuna is a indicator species for both DeSoto Canyon and Seasonal Closure. Both closures aim to reduce the bycatch of bluefin tuna, especially Spring Closure which aims to protect the mature stock whilst spawning in the Gulf. Gulf spawners are known to be within the Gulf of Mexico for a relatively short period of time before migrating back to their foraging grounds in the north Atlantic (Block et al., 2005). The Northern Atlantic is where the western stock spends most of their time, and it is also where most of the fishing mortality is exerted on the stock (ICCAT, 2014b). Although this study found Gulf pelagic longline closures, especially Desoto Canyon, had some influence on reducing bluefin catch these closures alone may not be providing enough protection to the spawning stock of bluefin tuna. Further reduction to the fishing mortality within the Atlantic Ocean may be necessary despite the fact the stock is already highly regulated due to its economic importance in international fisheries and the concern regarding the stock's ecological sustainability.

Replenishing the bluefin tuna stock depends on the survival and recruitment of larvae, which depends on a variety of environmental factors, including sea surface temperature (Teo et al., 2007; Muhling et al., 2010). Climate change could diminish spawning habitat within the next 50 years within the Gulf of Mexico (Muhling et al., 2011, 2014), which could have a negative impact on recruitment. This could be bad for the western bluefin stock as ICCAT (2014b) has shown that the stock is unlikely to rebuild under poor recruitment. However, there is potential for a shift in aggregation locations as ocean conditions continue to change (Martell et al., 2005). GoMAM should be considered to investigate impacts to bluefin given changes to
physical and environmental conditions due to inter-annual variability and/or climate change. This could be done by building upon the current hydrology sub-model, which would also allow advanced investigations pertaining to pelagic spatial closures within the Gulf of Mexico. The success of pelagic spatial closures will likely depend on inter-annual variability and changing ocean conditions, which can shift aggregation locations (Martell et al., 2005). Dynamic fishery spatial closures, those not fixed in space or time, could be designated to protect hydrographic features where pelagics aggregate. Fronts, being predictable, could be the basis for setting the closure's boundaries (Hyrenbach et al., 2000).

For many pelagic predators (i.e., tunas and billfishes) pelagic longlining is the dominant form of commercial longline fishing mortality, as pelagic longline fishing mortality sensitivity scenarios often produced similar results as all longline fishing mortality sensitivity scenarios, however this is was not the case for shark groups. GoMAM combines many of the large shark indicator species into one functional group, so there is no distinction between coastal species and pelagic species. The functional group's dynamics are driven more by coastal species than by pelagic species since most of the species in the group are coastal. Coastal species are more susceptible to benthic gears, e.g., bottom longlining (Ingram et al., 2005; Henwood et al., 2006; Hale and Carlson, 2007) than pelagic gears, so it is not surprising that this study saw that pelagic longline fishing and spatial closures had little influence on the large sharks functional group. Efforts to reduce the bycatch and improve the biomass of large shark species need to also incorporate coastal fleets (see Appendix C). This could include adjusting current coastal longline restriction zones, like the Reef fish longline and buoy gear restricted area (Gulf of Mexico Fishery Management Council, 1989;

Coleman et al., 2004a; Gulf of Mexico Fishery Management Council, 2016) and/or the Seasonal prohibition applicable to bottom longline fishing for Reef fish (Gulf of Mexico Fishery Management Council, 2016), to include restrictions/bans directed at shark species.

Pelagic fishery spatial closures can be a useful tool to achieve management objectives pertaining to the protection and rebuilding of highly migratory pelagic predators, but it is imperative that science-driven analysis via mathematical modeling is done to address their utility and feasibility. Although there are uncertainties regarding results from the Gulf of Mexico Atlantis model, the tool remains useful for investigating broader impacts from fisheries regulations within the Gulf. Gulf of Mexico pelagic longline spatial closures are likely reducing the bycatch of some pelagic predators, DeSoto Canyon possibly being more successful than Spring Closure. However, the impacts of the closures are likely limited since closures tend to shift fishing pressure, and the closures constitute a small part of the range of many pelagics. Rebuilding overfished populations such as bluefin tuna and the billfishes will be contingent on a suite of management strategies aiming to reduce fishing mortality inside and outside of the Gulf of Mexico, both through ICCAT and in national fisheries management plans. Future studies should consider the use of an updated hydrology sub-model in order to investigate more advanced spatial closures (e.g., rotating MPAs) and impacts due to changing ocean conditions.


Figure 4.1: United States Gulf of Mexico Pelagic Longline Fishing Closures. The shapefile for DeSoto Canyon was provided by Frick (2011), and the coordinates for Spring Closure were provided by National Oceanic and Atmospheric Administration (2016c). Figure generated in ArcGIS.


Figure 4.2: Gulf of Mexico Atlantis Model Spatial Map. Figure generated in ArcGIS.

Table 4.1: Management and Ecosystem Goals and Performance Metrics. This table summarizes U.S. pelagic longline spatial closure management objectives, investigated ecological objectives, performance metrics corresponding to objectives, and computations of performance metrics. Computation variables include the time step ( $t$ ), functional group (fid), number of functional groups in assemblage $(X)$, catch $(C)$, swordfish $(S W D)$, age-at-first-capture ( $a f c$ ), biomass ( $B$ ), bluefin tuna ( $B T N$ ), density $(D)$, mature age-classes (mature), pelagic functional groups (pelagic), and demersal functional groups (demersal).

| Objective | Performance Measures | Computation |
| :---: | :---: | :---: |
| DeSoto Canyon Management |  |  |
| Reduce catch of bycatch groups | Catch of bycatch groups | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t m a x}\left(\sum_{f i d=1}^{X} C_{f i d}^{t}\right)$ |
| Reduce catch of undersized swordfish | Proportion of swordfish catch being age-at-first-capture | $\frac{1}{4} \sum_{t=t m a x-4}^{t m a x}\left(C_{S W D_{\text {afc }}}^{t} / C_{S W D}^{t}\right)$ |
| Reduce catch of incidental groups | Catch of bluefin tuna | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t m a x}\left(C_{B T N}^{t}\right)$ |
| Minimize reduction of target catch | Catch of target groups | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t m a x}\left(\sum_{f i d=1}^{X} C_{f i d}^{t}\right)$ |
| Optimize survival of bycatch and incidental | Biomass of bycatch and incidental groups | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t \max }\left(\sum_{f i d=1}^{X} B_{f i d}^{t}\right)$ |
| Spring Closure Management |  |  |
| Promote rebuilding of bluefin tuna | Biomass of bluefin tuna |  |
| Reduce catch of incidental bluefin tuna | Catch of bluefin tuna | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t m a x}\left(C_{B T N}^{t}\right)$ |
| Ecosystem |  |  |
| Change in weight of individuals | Average weight of individuals in functional groups | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t m a x}\left(\sum_{\text {fid=1 }}^{X} B_{\text {fid }}^{t} / D_{\text {fid }}^{t}\right)$ |
| Change in stock age structure | Proportion mature of functional groups | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t m a x}\left(\sum_{f i d=1}^{X} B_{\text {fid }}^{t}{ }_{\text {mature }} / B_{\text {fid }}^{t}\right)$ |
| Change in pelagic predator biomass | pelagic:demersal ratio of ecosystem | $\frac{1}{4} \sum_{t=\text { tmax }-4}^{t m a x}\left(B_{\text {pelagic }}^{t} / B_{\text {demersal }}^{t}\right)$ |
| Change in ecosystem biodiversity | Q90 biodiversity metric of ecosystem biomass | see Eqn. (4.2) in main text |
| Shift in fishing pressure | pelagic:demersal ratio of catch; | $\frac{1}{4} \sum_{t=t \text { max }-4}^{t \max }\left(C_{\text {pelagic }}^{t} / C_{\text {demersal }}^{t}\right)$ |

Table 4.2: Summary of Results for Management Objectives Performance Metrics. Management objective performance metrics relative to the status quo. Biomass and catch metrics for bycatch groups are the sum of all bycatch groups (total), the sum of only billfish bycatch groups (billfishes), and the sum of only tuna bycatch groups (tunas). Incidental biomass and catch metrics refer to bluefin tuna. Biomass and catch metrics for target groups are the sum of all target groups. The age structure of swordfish (SWD) catch represents the proportion of swordfish catch being age-at-first-capture.
Table 4.2: Continued.

| Scenarios | Biomass of Bycatch |  |  | Biomass of Incidental | Biomass of Target | Catch of Bycatch |  |  | Catch of Incidental | Catch of Target | Age structure SWD catch |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Total | billfishes | tunas |  |  | Total | billfishes | tunas |  |  |  |
| Values computed within US Gulf |  |  |  |  |  |  |  |  |  |  |  |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All longlining $\mathrm{F} * 0$ | 1.000 | 1.007 | 1.045 | 1.044 | 1.014 | 0.972 | 0.839 | 0.000 | 0.004 | 0.942 | 0.957 |
| All longlining $\mathrm{F} * 0.5$ | 1.000 | 1.003 | 1.022 | 1.018 | 1.007 | 0.986 | 0.920 | 0.514 | 0.641 | 0.972 | 0.979 |
| All longlining $\mathrm{F} * 2$ | 1.000 | 0.993 | 0.957 | 0.983 | 0.987 | 1.026 | 1.157 | 1.894 | 1.399 | 1.050 | 1.042 |
| Pelagic longlining $\mathrm{F} * 0$ | 1.000 | 1.007 | 1.044 | 1.044 | 1.009 | 0.978 | 0.839 | 0.000 | 0.004 | 0.954 | 0.957 |
| Pelagic longlining $\mathrm{F} * 0.5$ | 1.000 | 1.004 | 1.022 | 1.018 | 1.004 | 0.989 | 0.920 | 0.514 | 0.641 | 0.978 | 0.979 |
| Pelagic longlining $\mathrm{F} * 2$ | 1.000 | 0.993 | 0.958 | 0.983 | 0.992 | 1.020 | 1.157 | 1.895 | 1.399 | 1.038 | 1.042 |
| No DeSoto Canyon | 1.000 | 0.999 | 0.991 | 0.996 | 0.999 | 1.004 | 1.031 | 1.188 | 1.102 | 1.007 | 1.013 |
| No Spring Closure | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.001 | 1.007 | 1.034 | 1.006 | 1.001 | 1.001 |
| No PLL Spatial Closures | 1.000 | 0.998 | 0.990 | 0.996 | 0.998 | 1.004 | 1.038 | 1.221 | 1.106 | 1.009 | 1.014 |
| Seasonal PLL Closure | 1.000 | 1.001 | 1.006 | 1.001 | 1.001 | 0.997 | 0.967 | 0.822 | 0.984 | 0.994 | 0.995 |
| Values computed within DeSoto Canyon |  |  |  |  |  |  |  |  |  |  |  |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All longlining $\mathrm{F} * 0$ | 1.000 | 1.008 | 1.047 | 1.040 | 1.016 | 0.978 | 0.095 | 0.000 | 0.000 | 0.944 | 0.957 |
| All longlining $\mathrm{F} * 0.5$ | 1.000 | 1.004 | 1.023 | 1.016 | 1.008 | 0.989 | 0.551 | 0.514 | 0.615 | 0.973 | 0.979 |
| All longlining $\mathrm{F} * 2$ | 1.000 | 0.992 | 0.956 | 0.985 | 0.985 | 1.020 | 1.882 | 1.894 | 1.520 | 1.048 | 1.042 |
| Pelagic longlining F * 0 | 1.000 | 1.008 | 1.046 | 1.040 | 1.012 | 0.982 | 0.095 | 0.000 | 0.000 | 0.954 | 0.957 |
| Pelagic longlining $\mathrm{F} * 0.5$ | 1.000 | 1.004 | 1.023 | 1.016 | 1.006 | 0.991 | 0.551 | 0.514 | 0.615 | 0.978 | 0.979 |
| Pelagic longlining $\mathrm{F} * 2$ | 1.000 | 0.992 | 0.956 | 0.985 | 0.989 | 1.017 | 1.882 | 1.895 | 1.520 | 1.038 | 1.042 |
| No DeSoto Canyon | 1.000 | 0.998 | 0.991 | 0.997 | 0.998 | 1.010 | 1.372 | 1.536 | 1.630 | 1.034 | 1.013 |
| No Spring Closure | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.020 | 1.018 | 0.995 | 1.000 | 1.001 |
| No PLL Spatial Closures | 1.000 | 0.998 | 0.990 | 0.996 | 0.998 | 1.010 | 1.392 | 1.553 | 1.624 | 1.034 | 1.014 |
| Seasonal PLL Closure | 1.000 | 1.001 | 1.006 | 1.001 | 1.001 | 0.997 | 0.804 | 0.809 | 0.983 | 0.992 | 0.995 |
| Values computed within Spring Closure |  |  |  |  |  |  |  |  |  |  |  |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All longlining $\mathrm{F} * 0$ | 1.000 | 1.010 | 1.052 | 1.133 | 1.028 | 0.976 | 0.079 | 0.000 | 0.000 | 0.921 | 0.957 |
| All longlining $\mathrm{F} * 0.5$ | 1.000 | 1.005 | 1.026 | 1.050 | 1.014 | 0.988 | 0.543 | 0.514 | 0.686 | 0.963 | 0.979 |
| All longlining $\mathrm{F} * 2$ | 1.000 | 0.990 | 0.950 | 0.947 | 0.975 | 1.022 | 1.897 | 1.895 | 1.197 | 1.063 | 1.042 |
| Pelagic longlining $\mathrm{F} * 0$ | 1.000 | 1.010 | 1.052 | 1.133 | 1.026 | 0.976 | 0.079 | 0.000 | 0.000 | 0.925 | 0.957 |
| Pelagic longlining $\mathrm{F} * 0.5$ | 1.000 | 1.005 | 1.026 | 1.050 | 1.012 | 0.989 | 0.543 | 0.514 | 0.686 | 0.965 | 0.979 |
| Pelagic longlining $\mathrm{F} * 2$ | 1.000 | 0.990 | 0.951 | 0.947 | 0.977 | 1.021 | 1.897 | 1.895 | 1.197 | 1.059 | 1.042 |
| No DeSoto Canyon | 1.000 | 0.998 | 0.990 | 0.987 | 0.995 | 1.004 | 1.102 | 1.156 | 0.982 | 1.011 | 1.013 |
| No Spring Closure | 1.000 | 1.000 | 0.999 | 0.999 | 0.999 | 1.002 | 1.079 | 1.064 | 1.015 | 1.006 | 1.001 |
| No PLL Spatial Closures | 1.000 | 0.998 | 0.989 | 0.986 | 0.995 | 1.005 | 1.181 | 1.219 | 0.994 | 1.016 | 1.014 |
| Seasonal PLL Closure | 1.000 | 1.002 | 1.007 | 1.002 | 1.003 | 0.996 | 0.816 | 0.846 | 0.974 | 0.990 | 0.995 |



Figure 4.3: Evaluation of Functional Group Specific Management Objective Performance Measures Upon the Establishment of DeSoto Canyon. Functional group-specific impacts for white marlin (WMR), blue marlin (BMR), other billish (BIL), miscellaneous tunas (OTN), bluefin tuna (BTN), large sharks (LGS), loggerhead sea turtles (LOG), kemp's ridley sea turtles (KMP), other sea turtles (OTN), swordfish (SWD), yellowfin tuna (YTN), and large pelagic fish (LPL). Performance metrics are compared across the whole Gulf (black), U.S. Gulf (red), U.S. Gulf open ocean (blue), and DeSoto Canyon (green). Axis values are from the status quo relative to no DeSoto Canyon.


Figure 4.4: Evaluation of Functional Group Specific Management Objective Performance Measures Upon the Establishment of Spring Closure. Functional group-specific impacts for white marlin (WMR), blue marlin (BMR), other billfish (BIL), miscellaneous tunas (OTN), bluefin tuna (BTN), large sharks (LGS), loggerhead sea turtles (LOG), kemp's ridley sea turtles (KMP), other sea turtles (OTN), swordfish (SWD), yellowfin tuna (YTN), and large pelagic fish (LPL). Performance metrics are compared across the whole Gulf (black), U.S. Gulf (red), U.S. Gulf open ocean (blue), and Spring Closure (green). Axis values are from the status quo relative to no Spring Closure.


Figure 4.5: Changes in Catch from Non-pelagic, U.S. Hook-and-Line Fleets Resulting from Pelagic Longline Fishing Mortality Sensitivity and Spatial Closure Scenarios. Changes in catch of tunas (dark grey), billfishes (grey), and large sharks (light grey) from hook-and-line shelf fleets (dashed) and sport fleets (solid) amongst indicated scenarios. Values corresponding to pelagic longline fishing mortality sensitivity scenarios (PLLF) are relative to the status quo. All other values are from the status quo relative to pelagic longline spatial closure scenarios.

Table 4.3: Summary of Results for Ecological Objectives Performance Metrics. Ecological objective performance metrics are relative to the status quo. Average individual weight, and proportion mature is shown for all billfish groups (billfishes), and all tuna groups (tunas). The pelagic:demersal ratio ( $\mathrm{P}: \mathrm{D}$ ) was computed based on biomass in the marine environment (ecosystem), and biomass caught by the fisheries (catch). The ecosystem Q90 biodiversity metric is based on biomass in the marine environment.

| Scenarios | Average Ind. Weight |  | Proportion Mature |  | P:D ratio |  | $\begin{gathered} \text { Q90 } \\ \text { ecosystem } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | billfishes | tunas | billfishes | tunas | ecosystem | catch |  |
| Values computed within US Gulf waters |  |  |  |  |  |  |  |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All Longlining F * 0 | 1.003 | 1.011 | 1.007 | 1.004 | 1.014 | 0.974 | 1.000 |
| All Longlining $\mathrm{F}^{*} 0.5$ | 1.001 | 1.005 | 1.004 | 1.002 | 1.007 | 0.988 | 1.000 |
| All Longlining F * 2 | 0.997 | 0.992 | 0.993 | 0.996 | 0.988 | 1.019 | 1.000 |
| Pelagic longlining $\mathrm{F} * 0$ | 1.003 | 1.011 | 1.007 | 1.004 | 1.013 | 0.966 | 1.000 |
| Pelagic longlining $\mathrm{F} * 1$ | 1.001 | 1.005 | 1.004 | 1.002 | 1.006 | 0.984 | 1.000 |
| Pelagic longlining $\mathrm{F} * 2$ | 0.997 | 0.992 | 0.993 | 0.996 | 0.989 | 1.028 | 1.000 |
| No DeSoto Canyon | 0.999 | 0.998 | 0.998 | 0.999 | 0.998 | 1.006 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.001 | 1.000 |
| No DC or SC | 0.999 | 0.998 | 0.998 | 0.999 | 0.997 | 1.007 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.001 | 1.001 | 1.001 | 1.002 | 0.995 | 1.000 |
| Values computed within DeSoto Canyon |  |  |  |  |  |  |  |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All Longlining F * 0 | 1.003 | 1.009 | 1.007 | 1.004 | 1.017 | 0.972 | 1.000 |
| All Longlining F * 0.5 | 1.001 | 1.004 | 1.003 | 1.002 | 1.008 | 0.988 | 1.000 |
| All Longlining F * 2 | 0.998 | 0.994 | 0.994 | 0.996 | 0.985 | 1.020 | 1.005 |
| Pelagic longlining $\mathrm{F} * 0$ | 1.003 | 1.009 | 1.007 | 1.004 | 1.015 | 0.960 | 1.000 |
| Pelagic longlining $\mathrm{F} * 1$ | 1.001 | 1.004 | 1.003 | 1.002 | 1.007 | 0.981 | 1.000 |
| Pelagic longlining $\mathrm{F} * 2$ | 0.998 | 0.993 | 0.994 | 0.996 | 0.986 | 1.032 | 1.005 |
| No DeSoto Canyon | 0.999 | 0.999 | 0.998 | 0.999 | 0.997 | 1.028 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 |
| No DC or SC | 0.999 | 0.998 | 0.998 | 0.999 | 0.997 | 1.028 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.001 | 1.001 | 1.000 | 1.002 | 0.994 | 1.000 |
| Values computed within Spring Closure |  |  |  |  |  |  |  |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All Longlining F * 0 | 1.001 | 1.006 | 1.005 | 1.002 | 1.022 | 0.938 | 1.000 |
| All Longlining F * 0.5 | 1.000 | 1.003 | 1.002 | 1.001 | 1.011 | 0.972 | 1.000 |
| All Longlining F * 2 | 0.999 | 0.997 | 0.995 | 0.998 | 0.980 | 1.047 | 1.004 |
| Pelagic longlining $\mathrm{F} * 0$ | 1.001 | 1.006 | 1.005 | 1.002 | 1.021 | 0.932 | 1.000 |
| Pelagic longlining $\mathrm{F} * 1$ | 1.001 | 1.003 | 1.002 | 1.001 | 1.010 | 0.968 | 1.000 |
| Pelagic longlining $\mathrm{F} * 2$ | 0.999 | 0.997 | 0.995 | 0.998 | 0.981 | 1.055 | 1.004 |
| No DeSoto Canyon | 1.000 | 0.999 | 0.999 | 1.000 | 0.996 | 1.009 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 0.999 | 1.004 | 1.000 |
| No DC or SC | 1.000 | 0.999 | 0.998 | 1.000 | 0.995 | 1.013 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.001 | 1.001 | 1.000 | 1.003 | 0.994 | 1.000 |



Figure 4.6: Evaluation of Ecosystem Objective Performance Metrics for Billfish and Tuna Functional Groups. Ecosystem objective performance metrics average individual weight (a), and proportion mature (b) for billfish and tuna functional groups: yellowfin tuna (YTN), bluefin tuna (BTN), little tunny (LTN), miscellaneous tunas (OTN), swordfish (SWD), white marlin (WMR), blue marlin (BMR), other billfish (BIL). U.S. Gulf metrics are compared amongst longline fishing mortality sensitivity scenarios in which fishing mortality for all longline fleets were multiplied by 0 (green), 0.5 (yellow), and 2 (red). Axis values are relative to the status quo.

## CHAPTER 5

## Conclusion

### 5.1 Summary of Dissertation

The Gulf of Mexico is an important ecosystem for Atlantic predatory pelagics (e.g., sharks, tunas, billfish) due to the environmental and physical dynamics which drive areas of productivity. The sustainability of fisheries for Atlantic pelagic predators is a concern, and the reduction of top-predators could have negative socioeconomic and biological impacts. Management efforts have included the establishment of two pelagic longline spatial closures within the pelagic waters of the Gulf of Mexico. Considering the lack of empirical data regarding the direct and indirect impacts of pelagic fisheries closures, the parameterization of a mathematical ecosystem model was necessary to assess the utility of these closures. This dissertation aided the parameterization of the Gulf of Mexico Atlantis model (i.e., chapter 2 and chapter 3), and used the model to assess if Gulf of Mexico pelagic longline closures could meet management objectives, and identify potential ecosystem impacts (chapter 4).

The work presented in Chapter 2, Landings Data for Ecosystem Fisheries Science: Lessons Learned from the Gulf of Mexico, considered Gulf of Mexico landings data to evaluate potential uncertainties in ecosystem based fisheries management metrics,
focusing on inputs to the Gulf of Mexico Atlantis model, and landings-based indicators. Meaningful portions of landings from commercial fisheries are ambiguous (not identified to species), especially in Mexico and Cuba which have large portions of unidentified landings ( $29.2 \%$ and $48.9 \%$, respectively). U. S. recreational data have minimal ambiguous landings ( $0.4 \%$ ), but landings are highly variable in part due to sampling error. U. S. ambiguous landings do not appear to be biasing the indicators. In addition, the aggregation of landings into Gulf of Mexico Atlantis functional groups do not appear to be biasing trends of landings-based indicators. While seasonspecific and State-specific U. S. landings do not introduce significant bias, much of the fleet-specific landings early in the time series could not be allocated to an Atlantis fleet due to the lack of detail in the landings dataset. This will not impact historical calibration because landings are not partitioned across fleets, but this should be considered when computing fleet-specific landings proportions for forecasting. Current landings time series appear to be sufficient for the development of ecosystem models, but ecosystem based fisheries management of the Gulf of Mexico would benefit from more precise landings data.

The work presented in Chapter 3, Predicting the Biomass Distributions of Pelagic Species Across the Gulf of Mexico Using Generalized Additive Models, Delta GAMs were fitted for describing the Gulf-wide spatial distributions of pelagic predatory functional groups using Gulf of Mexico U. S. bottom longline survey catch data (coastal models), and U. S. pelagic longline commercial catch data (pelagic models). This work advanced our knowledge on the environmental cues and spatial distribution of pelagic groups within the Gulf of Mexico, which can aid fisheries management by identifying areas of increased vulnerability. Fitted models for large, predatory sharks had some of
the better fits, diagnostics, and performance. The large, predatory sharks model was influenced by minimum distance from a front, in addition to environmental drivers. There is little research linking the spatial patterns of predatory shark species to fronts, so this suggests further studies investigating possible connections between physical features and the presence of predatory shark species. The fitted model presented in this dissertation successfully identified areas of the Gulf known to have higher catch rates of sharks, which could aid spatial management and conservation of predatory sharks.

Although many of the fitted GAMs presented in this dissertation had poor diagnostics and performances, these models improved the representation of the spatial distribution of pelagics in spatially-explicit ecosystem models. In the case of the Gulf of Mexico Atlantis model, spatial distributions were previously assumed to be fairly homogenous, lacking variability across space and season. Thus, the heterogeneous profiles produced from the statistical models improved the characterization of food web dynamics, fisheries, and spatial regulations. Also, estimates are being aggregated into large polygons, so inaccuracies at the predicted level are probably averaged out. However, distribution profiles are only used in the initial setup of Atlantis and a new stable spatial distribution is formed based on a number of simulated factors (e.g., habitat affinity, predator/prey dynamics, fisheries, etc.). Thus, it is uncertain how the spatial distributions mesh with the rest of the model. For example, if the distribution of prey groups do not correspond to the distributions estimated by the GAMs to some extent, then Atlantis will not allow the spatial distribution of predator groups to persist because they can not find food. The spatial distributions estimated by GAMs should be compared to spatial distributions GoMAM stabilize to, and if
there are significant differences than there are likely issues with the distributions of large pelagic functional groups, or their prey groups.

The work presented in Chapter 4, Can Gulf of Mexico Pelagic Longline Fishery Closures Meet Management Objectives?, conducted a policy exploration investigating if current pelagic longline spatial closures could meet management objectives, and possible ecosystem impacts. Current closures include a permanent closure over DeSoto Canyon (DeSoto Canyon), and seasonal closure off the Louisiana shelf (Spring Closure). Closures are intended to reduce bycatch and promote the rebuilding of biomass for pelagic predators. The Gulf of Mexico Atlantis model was used to simulate management scenarios altering spatial closures and pelagic longline fishing mortality, and tracked changes to performance measures. Closures reduced Gulf-wide catches of bycatch and incidental groups with little reduction to catches of target groups, but the reduction in fishing mortality was not enough to cause meaningful increases in biomass of bycatch and incidental groups. DeSoto Canyon was more successful at achieving management objectives, and had more influence to ecosystem metrics, than Spring Closure. Pelagic spatial closures can be useful tools to achieve management objectives pertaining to the protection of pelagic predators within areas where they are known to aggregate and be particularly vulnerable, but rebuilding the biomass of particular stocks may require additional reductions in fishing mortality across the Atlantic Ocean. In addition, the pelagic spatial closures may not reduce fishing mortality enough to cause meaningful impacts to ecosystem performance measures, but the pelagic:demersal ratio indicated that pelagic spatial closures can adjust the catch composition which may impact community structure.

### 5.2 Limitations

Data are a recurring limitation to this work. Gulf of Mexico landings time series from Mexico and Cuba used to calibrate the Atlantis historical model are likely an underrepresentation as data did not include recreational fisheries and historical reporting of artisanal landings are often incomplete, particularly for Mexico. In addition, there is uncertainty pertaining to U. S. recreational landings as data are estimates based on surveys of fishers that are expanded to the whole fishery.

Catch data used to fit GAMs were the primary limitation to describing the spatial distribution of pelagic predators. First, pelagic models were fitted with fisheriesdependent data. Using fisheries-independent data is preferable as it adheres to the statistical assumption of data independence (i.e. random sampling has a better representation of where species are and are not common). In addition, fisheries-independent data often contains more environmental data allowing for the consideration of a variety of model descriptors. While incorporating estimates of some model descriptors into fisheries-dependent data allowed the improvement of some model fits, this also introduces uncertainty. Second, coastal models only considered bottom longline data and lacked mid-depth hook-and-line data. Thus, organisms feeding within the water column and not at bottom (e.g., tunas, billfish) were not considered. Incorporating mid-depth hook-and-line data into this study would expand the species coverage and improve fits for coastal models. Lastly, catch data only spanned the northern Gulf, and extrapolating across the southern Gulf produced highly uncertain predictions.

GAMs were an appropriate method for modeling the spatial abundance of pelagic predatory groups. Smoothing splines allowed adequate flexibility when fitting the
environmental data, and the relaxed statistical assumptions of GAMs allowed the use of fisheries-dependent data. There are ongoing efforts for developing a GAM framework, using carefully chosen environmental predictors and a blending of available fisheries-independent and fisheries-dependent survey data, for enhancing Gulf of Mexico ecosystem models (Grüss et al., 2016a). Residual analysis of the GAMs presented in this dissertation showed that many of the gamma GAMs did not represent the data very well. This could possibly be addressed in future research by transforming the response variable using transformations like the Basic or Box-Cox (as recommended by Mateu (1997) when normalizing environmental data).

This dissertation found some parameterizations of the Gulf of Mexico Atlantis model worth adjusting to benefit future studies investigating the fisheries management of pelagic predatory groups. First, the harvesting module can be altered to distinguish bycatch from landings. This will allow more nuanced investigations of management strategies aiming to reduce bycatch while maintaining landings. In addition, the representation of fleets can be advanced to allow the redistribution of fishing mortality around MPAs, rather than reducing mortality. Also, diagnostics for some of the pelagic predatory functional groups, specifically bluefin tuna, could be improved. Model diagnostics show a sudden loss of adults seemingly due to unbalanced consumption, and bluefin diagnostics got worse when bluefin migration patterns are adjusted to better represent migration described in current literature. These may be related, as bluefin tuna spawning in the Gulf do most of their foraging in the north Atlantic. Conservation of bluefin tuna while the stock is breeding in the Gulf of Mexico is a pressing issue. To conduct this investigation strictly focusing on bluefin tuna impacts, not only would the above need to be addressed, but methods should
include different scenarios modeling possible fisher behavior regarding the pelagic longline spatial closures (especially Spring Closure as it was specifically established for spawning bluefin tuna). In addition, metrics describing the overlap between migrating bluefin tuna and the spatial closures should be analyzed, as well as catch rates in areas outside of the closures.

This dissertation project has identified several areas where the Gulf of Mexico Atlantis model could be refined to advance its functionality, and improve its modeling of the system (e.g., partitioning the large coastal sharks functional group into two functional groups: coastal sharks, and pelagic sharks), but the Gulf of Mexico Atlantis model was a very useful tool for study. The Atlantis framework allowed for the detailed, spatially-explicit representation of biota, fleets and spatial closures, and provided a means to explore broad-scale, ecosystem impacts. To take this study one step further, it would be interesting to simulate the investigated scenarios with another spatially-explicit ecosystem model to compare and contrast results. However, currently the other spatially-explicit ecosystem models of the Gulf of Mexico do not span the whole ecosystem.

### 5.3 Future Research

Qualitative assessment indicators shown in Chapter 2 reveal that a majority of U . S. commercial landings are of species that are not overfished, but the majority of $U$. S. recreational landings are of species of unknown overfished status. Because U. S. recreational fleets are important sources of fishing mortality in Gulf of Mexico waters, expanding assessment coverage to incorporate these species would be informative for fisheries management. Data-limited methods may be desirable as these species are not
commercially important and data are probably not adequate for the usual assessment methodologies. In addition, methods for reducing the uncertainty in U. S. recreational data would help landings-based investigations of the Gulf of Mexico.

Because fisheries-independent datasets are preferable to fisheries dependent datasets for statistical investigations, developing an annual pelagic longline fisheriesindependent survey within the Gulf of Mexico would be beneficial for tracking and monitoring the status of pelagic predatory stocks within the Gulf of Mexico. Getting funding for such a project may be difficult since the open ocean is expansive and oligotrophic. Considering the importance of physical metrics (e.g., fronts) when fitting GAMs, these efforts should collect metrics relating to physical oceanography (e.g., altimetry, currents) in addition to at-depth data from CTDs. Such surveys have been previously done in the Gulf but they have been discontinued(Fitzhugh et al., 2012; Campbell et al., 2012). Such a survey may become more necessary in the future as changing ocean conditions might impact recruitment and shift productive areas were pelagics aggregate.

Because foraging pelagic predators are actively seeking out food, incorporating model descriptors relating to predator-prey dynamics into GAMs could improve model fits and performance, and could be informative for the spatial ecology of some species. Schick and Lutcavage (2009) found that the fit of a generalized model predicting the distribution of bluefin tuna improved with the inclusion of prey dynamics. However, Benoit-Bird et al. (2013) found that many studies have found weak or ephemeral spatial associations between predators and prey within pelagic environments, and reported that their statistical models were unable to find a spatial relationship between predators and their prey. Instead, authors found that habitat use by predator groups
considered in their study were most strongly predicted by prey patch characteristics (i.e., depth and local density within spatial aggregations).

Mitigating the bycatch of sea turtles, birds, and mammals are also objectives for the management of pelagic longline gears. Mapping the Gulf-wide spatial distribution of these groups would benefit the parameterization of spatially explicit ecosystem models, and be informative for spatial management of the Gulf of Mexico. Considering the Gulf of Mexico Atlantis model, data should be located or collected in order to appropriately parameterize the bycatch modeling of these groups.

## APPENDIX A

## Additional Methodology and Results for Chapter 2

## A. 1 Additional Methods for Section 2.3.1

Table A.1: NOAA Commercial Gear-types Assigned to Atlantis Fleets.

| GillnetEst | LLPelgc | PotCrbEst | OtherUS |
| :---: | :---: | :---: | :---: |
| Entangling Nets (Gill) Unspc | Lines Troll, Other | Brush Trap | By Hand, Other |
| Gill Nets, Drift, Runaround | Lines Long Set With Hooks | Pots And Traps, Crab, Blue | By Hand, Oyster |
| Trammel Nets | Lines Long Drift With Hooks | Pots And Traps, Crab, Other | Cast Nets |
| Gill Nets, Stake |  | Pots And Traps, Eel | Diving Outfits, Other |
| Gill Nets, Sink/Anchor, Other | OytEst |  | Dip Nets, Drop |
| PotLbtShf | Rakes, Other | LLReefShf | Dip Nets, Common |
| Pots And Traps, Spiny Lobster | Dredge Other | Lines Long, Reef Fish | Fyke And Hoop Nets |
|  | Tongs and Grabs, Oyster | Lines Trot With Baits | Haul Seines, Beach |
| LLShkShf <br> Lines Long, Shark | Tongs and Grabs, Other |  | Haul Seines, Long |
|  |  | PotCrbShf | Hooks, Sponge |
|  | HLReefShf | Pots And Traps, Fish | Lampara \& Ring Nets |
| TwlShpEst | Reel, Electric or Hydraulic | Pots And Traps, Other | Spears |
| Beam Trawls, Butterfly Nets | Rod and Reel | Pots And Traps, Shrimp |  |
| Skimmer Net | Lines Hand, Other |  | Not Assigned |
|  | Lines Long, Vertical | SeineMenShf | Not Coded |
| TwlShpShf | RoyalRed | Encircling Nets (Purse) | Combined Gears |
| Otter Trawl Bottom, Shrimp | Otter Trawl Bottom, Shrimp | Purse Seines, Other | Unspecified Gear |
| Otter Trawl Bottom, Fish | Skimmer Net | Purse Seines, Menhaden | Troll \& Hand Lines Cmb |
| Otter Trawl Bottom, Scallop Trawls, Unspecified |  |  | Slat Traps (Virginia) |

## A.1.1 Landings Data Discrepancies

To determine if there are any discrepancies between the annual and itemized datasets from either NOAA and MRIP, plots were made to describe time series of i) the ratio between total landings in the itemized dataset against total landings in the annual dataset, and ii) the ratio between the total number of groups identified in the itemized dataset against the total number of groups identified in the annual datasets.

## A. 2 Additional Methods for Section 2.3.1.1

## A.2.1 Spatial Distribution

Seasonal landings time series were distributed across space (i.e., Atlantis polygons) using the seasonal biomass distributions of the functional groups. The construction of the seasonal biomass distributions are described above under Biomass distributions. First, the polygons that make up the region where each of the territorial fleets (i.e., U.S. commercial, U.S. recreational, Mexican commercial, and Cuban commercial) operate within were determined. Commercial landings are harvested from polygons that lie within the appropriate EEZ boundaries (note, all commercial fleets can harvest in international waters at the center of the Gulf, in the area called the donut hole). Polygons within the U.S. EEZ that do not exceed 200 m in depth were designated to contain U.S. recreational harvesting. However, U.S. recreational landings for the functional group crabs and lobsters were restricted to polygons 27 and 28 (SEDAR, 2005). Since boundary polygons 0 and 65 are reserved for flux characteristics, they were not included in the spatial distribution of landings.

Landings were only partitioned for the polygons described above; since the seasonal biomass distributions of the functional groups consider the entire polygon grid, they were partitioned in the same manner. Then, for each season and each territorial fleet, seasonal biomass distributions of the functional groups for the polygon subsets were adjusted so the distributions summed to 1 . We utilized these adjusted seasonal distributions to allocate the corresponding seasonal landings across the appropriate polygons for U.S. commercial, U.S. recreational, Mexican commercial, and Cuban commercial landings.

## A. 3 Additional Methods for Section 2.3.2

Table A.2: Functional Group Metrics for Ecological Indicator Computations.

| FID | Functional group | Trophic Level | Trophic Level Standard Error | Pelagic/Demersal |
| :---: | :---: | :---: | :---: | :---: |
| GAG | Gag grouper | 3.6 | 0 | D |
| RGR | Red grouper | 3.6 | 0 | D |
| SCM | Scamp | 4.4 | 0 | D |
| SSR | Shallow serranidae | 3.9 | 0.5 | D |
| DSR | Deep serranidae | 4.1 | 0.3 | D |
| RSN | Red snapper | 4 | 0 | D |
| VSN | Vermilion snapper | 4.3 | 0 | D |
| LUT | Lutjanidae | 4 | 0.3 | D |
| BIO | Bioeroding fish | 2.5 | 0 | D |
| LRF | Large reef fish | 3.4 | 0.6 | D |
| SRF | Small reef fish | 3.3 | 0.7 | D |
| BDR | Black drum | 3.9 | 0 | D |
| RDR | Red drum | 4.1 | 0 | D |
| SEA | Seatrout | 4.1 | 0.3 | D |
| SCI | Sciaenidae | 3.4 | 0.3 | D |
| LDY | Ladyfish | 3.9 | 0 | D |
| MUL | Mullets | 2.4 | 0.1 | D |
| POM | Pompano | 3.7 | 0.6 | D |
| SHP | Sheepshead | 3.5 | 0 | D |
| SNK | Snook | 4.2 | 0 | D |
| FLT | Flatfish | 3.4 | 0.3 | D |
| ODF | Other demersal fish | 3.7 | 0.6 | D |
| SDF | Small demersal fish | 3.7 | 0.5 | D |
| YTN | Yellowfin tuna | 4.3 | 0 | P |
| BTN | Bluefin tuna | 4.5 | 0 | P |
| LTN | Little tunny | 4.4 | 0 | P |
| OTN | Other tuna | 4.2 | 0 | P |
| SWD | Swordfish | 4.5 | 0 | P |
| WMR | White marlin | 4.5 | 0 | P |
| BMR | Blue marlin | 4.5 | 0 | P |
| BIL | Other billfish | 4.4 | 0.1 | P |
| AMB | Greater amberjack | 4.5 | 0 | P |
| JCK | Jacks | 4.1 | 0.4 | P |
| KMK | King mackerel | 4.5 | 0 | P |
| SMK | Spanish mackerel | 4.2 | 0.3 | P |
| SAR | Spanish sardine | 3.1 | 0 | P |
| LPL | Large pelagic fish | 4 | 0.5 | P |
| DWF | Deep water fish | 3.6 | 0 | P |
| MEN | Menhaden | 2.3 | 0.1 | P |
| PIN | Pinfish | 3.6 | 0 | P |
| MPL | Medium pelagic fish | 3.5 | 0.7 | P |
| SPL | Small pelagic fish | 3.3 | 0.4 | P |
| TIP | Blacktip shark | 4.4 | 0 | P |
| BEN | Benthic feeding sharks | 4.3 | 0 | P |
| LGS | Large sharks | 4.3 | 0.2 | P |
| FIL | Filter feeding sharks | 3.4 | 0 | P |
| SMS | Small sharks | 4.3 | 0 | P |
| RAY | Skates and rays | 3.7 | 0.5 | P |
| BSH | Brown shrimp | 2.5 | 0 | D |
| WSH | White shrimp | 2.5 | 0 | D |
| PSH | Pink shrimp | 2.5 | 0 | D |
| OSH | Other shrimp | 2.5 | 0 | D |
| DBR | Diving birds | 3.6 | 0 | D |
| SBR | Surface feeding birds | 3.6 | 0 | D |
| MAN | Manatee | 4.5 | 0 | D |
| MYS | Mysticeti | 3.2 | 0 | D |
| DOL | Dolphins and porpoises | 4.7 | 0 | D |
| DDO | Deep diving odontocetae | 4.7 | 0 | D |
| LOG | Loggerhead | 3.4 | 0 | P |
| KMP | Kemps ridley | 3.3 | 0 | P |
| TUR | Other turtles | 3.3 | 0 | P |
| BCR | Blue crab | 2.7 | 0 | D |
| SCR | Stone crab | 2.7 | 0 | D |
| LOB | Crabs and lobsters | 2.7 | 0 | D |
| COR | Stony corals | 2.3 | 0 | D |
| CCA | Crustose coralline algae | 2.3 | 0 | D |
| OCT | Octocorals | 2.3 | 0 | D |
| SPG | Sponges | 2.3 | 0 | D |
| CMB | Carnivorous macrobenthos | 2.2 | 0 | D |
| INF | Infaunal meiobenthos | 2 | 0 | D |
| ECH | Herbivorous echinoderms | 2 | 0 | D |
| OYS | Oysters | 2 | 0 | D |
| BIV | Bivalves | 2 | 0 | D |
| SES | Sessile filter feeders | 2 | 0 | D |
| EPI | Epiphytes | 1 | 0 | D |
| GRS | Sea grass | 1 | 0 | D |
| ALG | Macroalgae | 1 | 0 | D |
| MPB | Microphytobenthos | 2.1 | 0 | D |
| LPP | Large phytoplankton | 1 | 0 | D |
| SPP | Small phytoplankton | 1 | 0 | D |
| DIN | Toxic dinoflagellates | 1 | 0 | D |
| PRO | Protists | 1 | 0 | D |
| JEL | Jellyfish | 3.1 | 0 | D |
| SQU | Squid | 3.2 | 0 | D |
| LZP | Large zooplankton | 2.1 | 0 | D |
| SZP | Small zooplankton | 2.1 | 0 | D |
| PB | Bacteria | 1 | 0 | D |
| BB | Sediment bacteria | 1 | 0 | D |
| DC | Carrion detritus | 1 | 0 | D |
| DL | Labile detritus | 1 | 0 | D |
| DR | Refractory detritus | 1 | 0 | D |

## A. 4 Additional Results for Section 2.3.1



Figure A.1: Species Composition Time Series of United States Commercial Data. Panel (a) shows the species composition for total landings. Panel (b) shows the species composition for total landings, excluding menhaden. Panel (c) shows the species composition of ambiguous landings. Legend shows only the seven most common species.


Figure A.2: Species Composition Time Series of United States Recreational Data. Panel (a) shows the species composition for total landings. Panel (b) shows the species composition for ambiguous landings. Legend shows only the seven most common species.


Figure A.3: Species Composition Time Series of Mexican Commercial Data. Panel (a) shows the species composition for total landings. Panel (b) shows the species composition for ambiguous landings. Legend shows only the seven most common species.


Figure A.4: Species Composition Time Series of Cuban Commercial Data. Panel (a) shows the species composition for total landings. Panel (b) shows the species composition for ambiguous landings. Legend shows only the seven most common species.


Figure A.5: Species Composition Time Series of Seasonal U.S. Commercial Data. Species compositions are shown for winter, Jan. - Mar. (a), spring, Apr. - Jun. (b), summer, Jul. - Sep. (c), and fall, Oct. - Dec. (d). Legend shows only the seven most common species.


Figure A.6: Species Composition Time Series of Seasonal U.S. Commercial Ambiguous Landings. Species compositions are shown for winter, Jan. - Mar. (a), spring, Apr. - Jun. (b), summer, Jul. - Sep. (c), and fall, Oct. - Dec. (d). Legend shows only the seven most common species.


Figure A.7: Species Composition Time Series of Seasonal U.S. Recreational Landings. Species compositions are shown for wave 1, Jan. - Feb. (a), wave 2, Mar. - Apr. (b), wave 3, May. - Jun. (c), wave 4, Jul. - Aug. (d), wave 5, Sep. - Oct. (e), and wave 6, Nov. Dec. (f). Legend shows only the seven most common species.


Figure A.8: Species Composition Time Series of Seasonal U.S. Recreational Ambiguous Landings. Species compositions are shown for wave 1, Jan. - Feb. (a), wave 2, Mar. - Apr. (b), wave 3, May. - Jun. (c), wave 4, Jul. - Aug. (d), wave 5, Sep. - Oct. (e), and wave 6, Nov. - Dec. (f). Legend shows only the seven most common species.


Figure A.9: Species Composition Time Series of United States Commercial Data by State. Menhaden landings were excluded from the figures. Legend shows only the seven most common species.


Figure A.10: Species Composition Time Series of United States Ambiguous Commercial Data by State. Legend shows only the seven most common species.


Figure A.11: Species Composition Time Series of United States Total Recreational Data by State. Legend shows only the seven most common species.


Figure A.12: Species Composition Time Series of United States Ambiguous Recreational Data by State. Legend shows only the seven most common species.


Figure A.13: Species Composition Time Series of United States Total Commercial Data by Gear. Plot legends are restricted to only show seven identifications with with the most landings data associated to the plot.


Figure A.13: Continued.


Figure A.13: Continued.


Figure A.14: Species Composition Time Series of United States Ambiguous Commercial Data by Gear. Plot legends are restricted to only show seven identifications with with the most landings data associated to the plot.


Figure A.14: Continued.


Figure A.14: Continued.

## A.4.0.1 Landings Data Discrepancies

To evaluate discrepancies between annual landings datasets and itemized landings datasets, plots were constructed showing the changes in the ratios comparing i) total landings from itemized datasets and total landings from annual datasets, and ii) total number of landed groups from itemized datasets and total number of landed groups from annual datasets.
U.S. commercial data series have some discrepancies between itemized data series and the annual data series (Figure A.15). The landings dataset itemized by season tends to have more landings represented than the annual landings dataset (Figure A.15a), but less identified taxonomic groups (Figure A.15d). The landings dataset itemized by state and the annual landings dataset tend to have the same amount of landings (Figure A.15b), as well as the same number of identified groups (Figure A.15e). The landings dataset itemized by gear appears to have less landings represented than the annual dataset (Figure A.15c), as well as fewer identified groups (Figure A.15f).
U.S. recreational data series have more discrepancies between itemized data series and the annual data series (Figure A.16). The landings dataset itemized by season tends to have more landings represented than the annual landings dataset (Figure A.16a). In the earlier years, the seasonal dataset tends to have more identified groups, but since 1990 the annual dataset tends to have more identified groups (Figure A.16c). The landings dataset itemized by state tends to have more landings represented than the annual landings dataset (Figure A.16b). In the earlier years, the dataset itemized


Figure A.15: Data discrepancies between NOAA's landings data itemized by season (a,d), state (b,e), and gear (c,f) and annual landings data. Panels (a-c) show the ratio between landings from itemized data series and landings from annual data series. Panels (d-f) show the ratio between the number of groups identified in itemized data series and the number of groups identified in annual data series.
by state tends to have more identified groups, but since 1990 the annual dataset tends to have more identified groups (Figure A.16d).


Figure A.16: Data discrepancies between MRIP's landings data itemized by season (a,c), and state ( $\mathrm{b}, \mathrm{d}$ ), and annual landings data. Panels ( $\mathrm{a}-\mathrm{b}$ ) show the ratio between landings from itemized data series and landings from annual data series. Panels (c - d) show the ratio between the number of groups identified in itemized data series and the number of groups identified in annual data series.

## A. 5 Additional Results for Section 2.3.1.1

## A.5.1 Historical Landings Time Series for Atlantis

Information was developed for the calibration and implementation of the Gulf of Mexico Atlantis model. Historical time series of U.S. commercial, U.S. recreational, Mexican commercial, and Cuban commercial landings for model calibration are shown in Tables A.3, A.4, A.5, and A. 6 (respectively). Average seasonal distributions for functional groups identified in the U.S. commercial, U.S. recreational, and Mexican historical time series are shown in Table A.7, A.8, and A. 11 (respectively). The proportions for distributing landings across fleets are presented in Table A.10.

Table A.3: United States Historical Commercial Landings by Atlantis Functional Group (tonnes)

| Group | 1980 | 1981 | 1982 | 1983 | 1984 | 1985 | 1986 | 1987 | 1988 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GAG | 643.7 | 898.8 | 1069.9 | 788.4 | 703 | 819.7 | 771.8 | 697.7 | 551.8 |
| RGR | 1316.8 | 1542.5 | 1792.9 | 2715 | 2466.6 | 2598.3 | 2863.5 | 3047.2 | 2151.2 |
| SCM | 0 | 0 | 0 | 0 | 0 | 14.4 | 174 | 164.4 | 125.3 |
| SSR | 218.9 | 256.4 | 259 | 269.3 | 340 | 279.1 | 993.6 | 900.7 | 715.7 |
| DSR | 223.2 | 372.8 | 406.5 | 280.7 | 294.9 | 403.5 | 703 | 711.9 | 1002.2 |
| RSN | 2273.4 | 2706.1 | 2907.1 | 3302.6 | 2604.5 | 2013 | 1798.5 | 1522.7 | 1841.6 |
| VSN | 139.9 | 164.1 | 180.4 | 258.8 | 652.4 | 670.7 | 793.5 | 728.2 | 705.1 |
| LUT | 715.8 | 767.7 | 1194.2 | 1002.6 | 939.3 | 797.4 | 960.6 | 1226.2 | 1085.6 |
| BIO | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LRF | 72.6 | 84.8 | 76.7 | 85 | 142.5 | 110.8 | 87.2 | 152.9 | 119.5 |
| SRF | 267 | 509.9 | 490.2 | 475.7 | 727.3 | 593 | 475.1 | 818.3 | 878.1 |
| BDR | 2691.9 | 2954.4 | 1932.2 | 2389.5 | 2691.5 | 3180.2 | 3455.5 | 4828.2 | 4748.5 |
| RDR | 1240.2 | 1249.7 | 1103.3 | 1422.1 | 1972.6 | 2881.8 | 6410.6 | 2223.9 | 136.5 |
| SEA | 2234.7 | 2112.4 | 1847 | 1921.3 | 1684.3 | 1491.8 | 1862 | 1878 | 1599 |
| SCI | 3474.5 | 3612.1 | 2147.5 | 1282 | 1121.7 | 939.6 | 844.1 | 1120.5 | 827.2 |
| LDY | 612.5 | 1814.4 | 1494.3 | 1888 | 1560.3 | 1342.9 | 2032.6 | 2322.7 | 1881.5 |
| MUL | 13896.6 | 15270.3 | 12211.1 | 11718.3 | 10292.9 | 9006.8 | 11899.1 | 10758.6 | 11602.5 |
| POM | 300.8 | 247.4 | 320.7 | 274.4 | 247 | 213.1 | 240.2 | 250.1 | 263.1 |
| SHP | 539.9 | 474.8 | 558.5 | 760.6 | 683.8 | 749.5 | 791.7 | 1518.3 | 1439.1 |
| SNK | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| FLT | 697.6 | 713.2 | 990.2 | 931.3 | 937.2 | 987 | 1034.5 | 1207.7 | 724.4 |
| ODF | 569.9 | 694.1 | 728.4 | 792.3 | 1292.8 | 1046.6 | 740.5 | 1377.7 | 1155.4 |
| SDF | 42.1 | 55.7 | 55.9 | 62.4 | 112.1 | 81.5 | 45.7 | 96.4 | 70 |
| YTN | 33 | 18.2 | 63.6 | 100.2 | 376.7 | 1505.9 | 3393.6 | 4179.4 | 7815.8 |
| BTN | 5.1 | 12.1 | 16.4 | 38.8 | 70 | 69.1 | 108.5 | 175 | 138.6 |
| LTN | 0.7 | 0.5 | 2.9 | 2.3 | 1.9 | 2.3 | 0.1 | 1.5 | 108.7 |
| OTN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SWD | 837.8 | 532.3 | 587.9 | 327.7 | 307.2 | 511.7 | 320.9 | 666.3 | 970.1 |
| WMR | 0 | 0 | 0 | 0 | 4.1 | 9.4 | 39.1 | 24.6 | 0.2 |
| BMR | 0 | 0 | 0 | 0 | 0.9 | 5 | 16.2 | 16.4 | 3.2 |
| BIL | 0 | 0 | 4.3 | 1.2 | 5 | 10.2 | 11.2 | 18.1 | 0 |
| AMB | 81.6 | 107.5 | 102.3 | 127.5 | 240.9 | 346.2 | 506.4 | 705.7 | 932.4 |
| JCK | 2424.6 | 2125.5 | 2335.1 | 2261.8 | 1679.1 | 1499.2 | 1414.9 | 2034 | 2868 |
| KMK | 1543.2 | 2399.6 | 662.4 | 1437.7 | 978.8 | 826.5 | 917 | 1109.1 | 865 |
| SMK | 887.9 | 1670.2 | 1524.1 | 1031.5 | 1596.8 | 1375.8 | 1244.5 | 1300.6 | 1054.4 |
| SAR | 1348.2 | 1264.6 | 1268.3 | 1217.6 | 1601.9 | 2069.4 | 2774.4 | 2925.9 | 1594.2 |
| LPL | 1226.1 | 860 | 869.7 | 705.8 | 641.8 | 588.2 | 631.3 | 684.2 | 1115 |
| DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| MEN | 702081 | 553684.4 | 861426.8 | 962982 | 985411.7 | 884189.2 | 830743.7 | 911642.5 | 639787.1 |
| PIN | 34.6 | 45.9 | 46 | 51.4 | 92.3 | 67.2 | 37.6 | 79.4 | 57.6 |
| MPL | 71.8 | 91.8 | 115 | 125.5 | 180.5 | 132 | 68.5 | 151.8 | 128.3 |
| SPL | 158.2 | 334.2 | 824.5 | 939.2 | 415.7 | 406.2 | 1417.6 | 825.8 | 1678.8 |
| TIP | 60.6 | 60.6 | 60.6 | 65.5 | 89.5 | 83.1 | 596.5 | 825.1 | 1460 |
| BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LGS | 155.6 | 258.3 | 304.5 | 352.9 | 306.7 | 417.6 | 855.9 | 2052.8 | 3781 |
| FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 0 | 0 | 0 | 0 | 0 | 0 | 40.2 | 0 | 77.7 |
| BSH | 49861.9 | 72678.6 | 56121.7 | 48553 | 62929.2 | 63706 | 72075.9 | 67305.6 | 59535.4 |
| WSH | 29835.2 | 32212.8 | 27443.7 | 29559.9 | 39196.7 | 41175.1 | 49417 | 37339.1 | 31600.1 |
| PSH | 9345.8 | 13625.5 | 8454.8 | 9196.7 | 10632.4 | 11526.7 | 8502.1 | 7557.5 | 6583.4 |
| OSH | 6276 | 4700.6 | 3073.6 | 3772.8 | 3546.5 | 3491.3 | 8644.1 | 5572.3 | 4855.6 |
| TUR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| BCR | 19444.9 | 19248.4 | 16761.2 | 18362.6 | 25558 | 25335.4 | 24047.6 | 35592.7 | 35992.1 |
| SCR | 1710.6 | 1894 | 2583.3 | 2173.4 | 1798.3 | 1847.2 | 1834.9 | 2160.4 | 2367.2 |
| LOB | 2594 | 2292.7 | 2594 | 1708.4 | 2772.1 | 2787.6 | 2010.5 | 2513.2 | 2648.5 |
| SPG | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 43.8 |
| CMB | 0 | 0 | 0.2 | 0.9 | 0 | 0 | 0.6 | 0.5 | 1.5 |
| OYS | 7039.7 | 8785.8 | 11408.8 | 13229 | 12520.5 | 12024.8 | 10220.3 | 8515.2 | 8103.4 |
| BIV | 7.9 | 21.3 | 5.9 | 17 | 44 | 55.6 | 8 | 8.9 | 33.5 |
| SQU | 26.9 | 43.9 | 35 | 32.1 | 55.4 | 67.2 | 80.1 | 75.3 | 104.7 |

Table A.3: Continued

| Group | 1989 | 1990 | 1991 | 1992 | 1993 | 1994 | 1995 | 1996 | 1997 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GAG | 767.9 | 814.4 | 710.1 | 754.8 | 847.6 | 734.4 | 751.1 | 712.6 | 727.4 |
| RGR | 3342 | 2181.5 | 2310.8 | 2024.5 | 2893.8 | 2223.9 | 2152.8 | 2020.4 | 2199.2 |
| SCM | 137.5 | 131.7 | 163 | 146.8 | 164 | 112 | 123 | 122.5 | 153.4 |
| SSR | 967.2 | 919.1 | 715.1 | 625.5 | 1049.3 | 747.5 | 545.1 | 646.6 | 572.8 |
| DSR | 548.1 | 817.3 | 741.2 | 813.7 | 679.9 | 939.7 | 698.7 | 513.4 | 610.4 |
| RSN | 1406.1 | 1207.5 | 1016.5 | 1380.3 | 1544.5 | 1475.1 | 1339.9 | 1973.6 | 2187.7 |
| VSN | 752.4 | 1113.1 | 814.1 | 1028.7 | 1233.6 | 1197.1 | 987.9 | 828.8 | 964.3 |
| LUT | 1348 | 1213.6 | 1396.2 | 1324.1 | 1700.4 | 1500.9 | 1267.2 | 1070.3 | 1202.3 |
| BIO | 0 | 0 | 3.6 | 1.5 | 0.3 | 0.5 | 0.5 | 0.3 | 0.4 |
| LRF | 170.2 | 198.6 | 198.3 | 202.7 | 465.3 | 361.7 | 250.5 | 273.5 | 266.2 |
| SRF | 812.9 | 829.3 | 812.2 | 831.5 | 1846.9 | 1396.3 | 1026 | 1128.4 | 1200.1 |
| BDR | 2490 | 1722.4 | 1215.7 | 1801.4 | 1890.9 | 2596.1 | 2737 | 2729.9 | 2569.6 |
| RDR | 81.6 | 9.5 | 17.3 | 35.2 | 58.7 | 32.6 | 19.3 | 25.9 | 21.7 |
| SEA | 1485 | 1010.3 | 1312.8 | 1131.5 | 1552.5 | 1446.2 | 882.3 | 806.6 | 686.5 |
| SCI | 1071.1 | 1171.8 | 1154.2 | 1258.8 | 2232.4 | 1683.1 | 1765.3 | 1294 | 1277.8 |
| LDY | 2073.1 | 2629.7 | 2058.3 | 2073 | 1819 | 1935.4 | 1245.1 | 844 | 858.1 |
| MUL | 12818.3 | 13307.5 | 11624.9 | 11675.4 | 13804.4 | 12392.1 | 9624.6 | 7023.6 | 7941.1 |
| POM | 245.9 | 327.6 | 278.5 | 253.6 | 253.1 | 266.4 | 179 | 120.6 | 261.9 |
| SHP | 1927.4 | 1796.3 | 1472.6 | 1880.7 | 2113 | 1964.9 | 1818.6 | 1558.8 | 1685.3 |
| SNK | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| FLT | 813.8 | 831 | 1062.9 | 1022.7 | 1712.5 | 1399.9 | 995.8 | 847.3 | 841 |
| ODF | 1550.8 | 1765.6 | 1859.3 | 1666 | 3971.5 | 2841.7 | 1932.8 | 2338.9 | 2254.6 |
| SDF | 105.3 | 121.7 | 123.7 | 119.2 | 338.8 | 221.4 | 144 | 192.2 | 186.7 |
| YTN | 5786.2 | 3665 | 2567.7 | 4229.4 | 2910.3 | 2105.4 | 1588 | 2125.2 | 2407.4 |
| BTN | 66.4 | 102.1 | 120.4 | 81.4 | 47 | 34.2 | 26.9 | 22.5 | 16.9 |
| LTN | 49.2 | 51.6 | 51 | 460.7 | 263.7 | 28.4 | 29.5 | 89.1 | 171.9 |
| OTN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SWD | 957.1 | 445.8 | 632.5 | 590.1 | 466.9 | 339 | 583.6 | 752.9 | 593.8 |
| WMR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| BMR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| BIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| AMB | 886.9 | 555.2 | 817.3 | 461.2 | 729.5 | 578.2 | 573.1 | 576.6 | 507.4 |
| JCK | 3101.6 | 2981.6 | 3031.7 | 3062.6 | 3497.9 | 2499.3 | 1403.1 | 661.1 | 766.1 |
| KMK | 788.7 | 912.4 | 881.4 | 1045.9 | 1352.3 | 1140 | 1007.7 | 1363 | 1367.4 |
| SMK | 1448.2 | 1212.7 | 1646.6 | 1816.2 | 1303.6 | 1310 | 769.1 | 412.6 | 367.3 |
| SAR | 1079.1 | 974.4 | 760.8 | 848.4 | 772.2 | 984.2 | 173 | 498.6 | 413.3 |
| LPL | 1177.4 | 1258.3 | 1881.7 | 1814.9 | 1728.6 | 1958.5 | 1387.8 | 1211.4 | 1287.6 |
| DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| MEN | 583185.8 | 539421.6 | 552946.5 | 432763.4 | 551534.6 | 774825.8 | 472059.7 | 491689.1 | 622013.7 |
| PIN | 85.8 | 99.5 | 106.6 | 112.9 | 296.5 | 221.3 | 128 | 166.5 | 168.3 |
| MPL | 234.3 | 243.7 | 250.3 | 290.6 | 635.3 | 487.2 | 354.7 | 383.7 | 333.6 |
| SPL | 4339.6 | 2160.8 | 2475.8 | 2676 | 3216.9 | 3059.7 | 2082.6 | 2449.5 | 2425.9 |
| TIP | 1594.7 | 960.3 | 388.2 | 444.6 | 476.8 | 1002.3 | 708.7 | 489.9 | 377.6 |
| BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LGS | 5543.5 | 4005 | 3505.5 | 3237 | 1757.1 | 2000.1 | 1914.3 | 1827.7 | 1727 |
| FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 224.9 | 280.9 | 132.7 | 122.9 | 73.3 | 33.8 | 45.3 | 8.7 | 0.1 |
| BSH | 69213.4 | 76343.7 | 64545.4 | 50898.6 | 50212.5 | 49684.1 | 57209.3 | 55140.8 | 49559.2 |
| WSH | 25567.2 | 30912.2 | 32000.3 | 33681.8 | 27581.2 | 32467.8 | 34996.8 | 25561.6 | 27993.3 |
| PSH | 6274.6 | 5470.3 | 4921.5 | 4651.9 | 6937.8 | 7313.9 | 10356.9 | 13969.2 | 9227.5 |
| OSH | 4362.6 | 3183.9 | 3311.3 | 10745.2 | 8642.4 | 5862.2 | 4241 | 7716.2 | 9957 |
| TUR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| BCR | 25240.2 | 26467.1 | 29866.1 | 31664.8 | 29781.1 | 24164.4 | 24800.9 | 28331.3 | 29095.9 |
| SCR | 2337.2 | 2853.7 | 2843.4 | 3013.5 | 3021.5 | 2976.1 | 2713 | 2924.7 | 2897.2 |
| LOB | 3265.1 | 2467.1 | 2762.1 | 1840.7 | 2066.8 | 2885.6 | 3537.6 | 3386.7 | 3276.1 |
| SPG | 277.5 | 360.5 | 381.2 | 338.9 | 338.7 | 387.8 | 357.4 | 324.3 | 236.4 |
| CMB | 0 | 0 | 0 | 0.7 | 1.1 | 1.3 | 0 | 1.7 | 1.4 |
| OYS | 7177.7 | 5600 | 5607.1 | 7414.2 | 8252.6 | 9219.5 | 10016.8 | 10571 | 10881.1 |
| BIV | 1322.2 | 28.7 | 0.4 | 2.7 | 2206.9 | 868.9 | 23.8 | 94.2 | 180.8 |
| SQU | 61.4 | 58.4 | 38.7 | 69.2 | 54.2 | 64.5 | 70.6 | 97.2 | 68.7 |

Table A.3: Continued

| Group | 1998 | 1999 | 2000 | 2001 | 2002 | 2003 | 2004 | 2005 | 2006 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GAG | 1151.8 | 952.3 | 1053.9 | 1470.5 | 1385.6 | 1239.8 | 1369.9 | 1233.8 | 661.3 |
| RGR | 1799.5 | 2705.3 | 2646.6 | 2697.4 | 2678.4 | 2239.5 | 2605.2 | 2454.4 | 2333.2 |
| SCM | 115.9 | 138 | 104.4 | 143.3 | 163.4 | 168.3 | 172.6 | 164.9 | 117.3 |
| SSR | 506.8 | 530.8 | 521.9 | 370.6 | 343.9 | 364.9 | 347.2 | 221 | 217.9 |
| DSR | 513.8 | 763.6 | 899 | 737.5 | 756 | 976.4 | 767.6 | 664.1 | 614.8 |
| RSN | 2129.2 | 2212.2 | 2197.2 | 2116.5 | 2187.7 | 2016.2 | 2121.4 | 1863.7 | 2103.5 |
| VSN | 785.9 | 899.2 | 662.2 | 778 | 911.1 | 1095.8 | 968.1 | 847.4 | 800.3 |
| LUT | 1056.5 | 1178.2 | 1108.9 | 977.7 | 996.5 | 936.3 | 1009.8 | 856.1 | 836.8 |
| BIO | 0.2 | 0.3 | 0.2 | 0.1 | 0.3 | 1.1 | 0.4 | 0.7 | 0.4 |
| LRF | 203 | 239 | 246.5 | 133.2 | 166.2 | 142.8 | 141.4 | 82.1 | 87.8 |
| SRF | 940.3 | 950.2 | 916 | 602.1 | 648.6 | 588.7 | 614.4 | 542 | 437.6 |
| BDR | 2066.2 | 2308.3 | 2630.1 | 2560.1 | 2510.1 | 2379.4 | 2526.1 | 2060.4 | 1909.5 |
| RDR | 24.4 | 26.2 | 24.6 | 13.8 | 11.4 | 13.3 | 11.2 | 15.5 | 12.2 |
| SEA | 393.2 | 403 | 342 | 239.2 | 219.3 | 177 | 145.3 | 113.8 | 107.7 |
| SCI | 992.9 | 972 | 856.8 | 477.1 | 423.3 | 402.8 | 378.9 | 246.4 | 280.8 |
| LDY | 970.4 | 1917.3 | 154.8 | 544.2 | 760.5 | 866.2 | 665.1 | 870.8 | 795.8 |
| MUL | 7153.1 | 9092.1 | 7625.7 | 7295.6 | 5742.8 | 5877.3 | 6237.1 | 4092.6 | 5772.9 |
| POM | 305.8 | 210.5 | 222.6 | 166.1 | 135.9 | 130.1 | 108.8 | 102.8 | 160.9 |
| SHP | 1299.6 | 1661.8 | 1433 | 1186.3 | 1036.3 | 1072.1 | 911.3 | 681.5 | 442.5 |
| SNK | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| FLT | 729.4 | 774.7 | 695.9 | 435.2 | 438.6 | 385 | 370.4 | 281.8 | 277.5 |
| ODF | 1794.4 | 1778.6 | 1706.3 | 1125.4 | 990.6 | 968.7 | 879.4 | 641.2 | 577.7 |
| SDF | 140.3 | 136.1 | 124.1 | 59.6 | 56 | 53 | 48.5 | 31.6 | 36.7 |
| YTN | 1721.3 | 2369.2 | 1957.1 | 1329.6 | 1927.4 | 1711.6 | 1584.6 | 1202.1 | 1096.2 |
| BTN | 13.7 | 36.1 | 34.5 | 16.9 | 29.5 | 38.5 | 66.3 | 43.5 | 16 |
| LTN | 105.3 | 232.1 | 54.1 | 193.3 | 207.6 | 506.2 | 81.1 | 110 | 144 |
| OTN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SWD | 510.5 | 447.8 | 467.8 | 347 | 413.2 | 375.7 | 402.4 | 345.8 | 267 |
| WMR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| BMR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| BIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| AMB | 317.6 | 354.2 | 415.7 | 332.9 | 357.2 | 451.1 | 442.7 | 337.4 | 286.9 |
| JCK | 1176.7 | 891.7 | 835 | 878.9 | 775.8 | 967.6 | 1013 | 956.1 | 736.8 |
| KMK | 1374.7 | 1363 | 1135.5 | 1281.9 | 1218.1 | 1098.2 | 1273.6 | 1113.9 | 1515.1 |
| SMK | 303.9 | 552.1 | 610.3 | 701.8 | 533.8 | 839.6 | 617.4 | 819.5 | 820.3 |
| SAR | 371.5 | 312.4 | 621.7 | 626.6 | 653.4 | 725.3 | 964 | 458 | 1023.2 |
| LPL | 1162.7 | 947.4 | 840.7 | 1018.6 | 1024.6 | 982.8 | 894.7 | 539.2 | 623.9 |
| DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| MEN | 495684.2 | 694272.6 | 591487.6 | 528569.9 | 585341.5 | 518362.8 | 464162.4 | 369914.8 | 408881.9 |
| PIN | 128.9 | 124.3 | 116.2 | 64.4 | 61.1 | 62.2 | 60.4 | 46.3 | 52.5 |
| MPL | 274.7 | 280.5 | 274.5 | 192.9 | 175 | 190.4 | 199.7 | 131.1 | 180.6 |
| SPL | 2543.2 | 2559.8 | 2453.7 | 2675.8 | 2919.6 | 2311.9 | 2728.5 | 2012.7 | 1327.2 |
| TIP | 521.8 | 436.8 | 351.8 | 329 | 250.9 | 597.4 | 368.9 | 242.2 | 364.8 |
| BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LGS | 1924.3 | 691.7 | 658 | 775.9 | 1053.2 | 1230.2 | 1059.9 | 907.2 | 1238.4 |
| FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 10.2 | 0.4 | 23.4 | 0 | 22.1 | 1.3 | 6.6 | 0.6 | 8.6 |
| BSH | 59092.6 | 60651.8 | 71592.5 | 65725.2 | 55472.7 | 62187.3 | 54625.3 | 43559.1 | 64925.7 |
| WSH | 39036.9 | 39187.3 | 49640.8 | 37874.8 | 38050.8 | 43369.7 | 51317.1 | 46179.7 | 60594.6 |
| PSH | 12436.6 | 5912.8 | 5417.9 | 7000.7 | 7760.6 | 6844.1 | 7015.1 | 6508.3 | 4379 |
| OSH | 9517.6 | 4902.7 | 4391.8 | 6020.2 | 4776.8 | 4178.2 | 3086.1 | 1938.6 | 1189.7 |
| TUR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| BCR | 30715.2 | 31432.3 | 31283.1 | 24722.9 | 29953.2 | 29088.6 | 27484.8 | 22718.3 | 30612.2 |
| SCR | 3171.8 | 2579.9 | 3109.9 | 3031.1 | 2918.7 | 2409.1 | 2708.8 | 2058.9 | 2180.5 |
| LOB | 2530 | 3270.3 | 2557.2 | 1479.4 | 1914 | 1774.6 | 2099.2 | 1394.3 | 1983.5 |
| SPG | 280.2 | 285.1 | 268.1 | 235.8 | 234.2 | 187.3 | 202.1 | 185.5 | 140.2 |
| CMB | 1.2 | 0.7 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| OYS | 9349.9 | 10946.5 | 11699.9 | 11622 | 10939.2 | 12291.5 | 11365.6 | 9158.7 | 8925.2 |
| BIV | 1293.1 | 1267.7 | 250.6 | 230.9 | 218.3 | 257.6 | 121.1 | 97.5 | 43.6 |
| SQU | 108 | 58.5 | 57.6 | 85 | 55.5 | 55.1 | 49.1 | 34.4 | 45.8 |

Table A.3: Continued

| Group | 2007 | 2008 | 2009 | 2010 | 2011 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| GAG | 621.3 | 678.6 | 384.6 | 264.9 | 161.4 |
| RGR | 1670.5 | 2141.6 | 1990.8 | 1582.6 | 2512 |
| SCM | 147.4 | 149.4 | 135.9 | 84.1 | 69.3 |
| SSR | 251 | 166.2 | 136.8 | 126.3 | 245.2 |
| DSR | 673.7 | 650.7 | 696.5 | 422.2 | 620.9 |
| RSN | 1360 | 1074.2 | 1135.3 | 1478.3 | 1605.7 |
| VSN | 1081.3 | 1273.8 | 1722.2 | 956.9 | 1391.3 |
| LUT | 654.7 | 831.1 | 1072.7 | 879.6 | 1043.4 |
| BIO | 1.2 | 0.7 | 0.8 | 0.6 | 0 |
| LRF | 113.3 | 99.3 | 99.1 | 71.5 | 143.6 |
| SRF | 561.9 | 538.5 | 482.2 | 344.5 | 641.4 |
| BDR | 1907.3 | 1838.2 | 2254.9 | 2079.4 | 2402.6 |
| RDR | 14.1 | 15.6 | 17.1 | 18.7 | 18.2 |
| SEA | 175.6 | 149 | 146.3 | 129.8 | 225.8 |
| SCI | 446.2 | 357.9 | 320.3 | 295 | 597 |
| LDY | 547.3 | 664.5 | 389.2 | 660.5 | 415.3 |
| MUL | 4052.1 | 4799 | 5126.8 | 4064 | 6455.8 |
| POM | 156.3 | 147.3 | 125.5 | 39.4 | 33.2 |
| SHP | 631 | 664.3 | 690.6 | 611.3 | 562.2 |
| SNK | 0 | 0 | 0 | 0 | 0 |
| FLT | 347.8 | 303.9 | 307 | 236.9 | 508.5 |
| ODF | 897.1 | 801.3 | 884.1 | 680.7 | 1416.8 |
| SDF | 64.7 | 49.7 | 43.3 | 37.2 | 91.4 |
| YTN | 1348.8 | 731.3 | 1114.4 | 302 | 658.1 |
| BTN | 32.9 | 25.2 | 17.4 | 20.5 | 3.1 |
| LTN | 127.7 | 34 | 119.6 | 266.2 | 26.7 |
| OTN | 0 | 0 | 0 | 0 | 0 |
| SWD | 337.9 | 301.2 | 398.4 | 174.1 | 320.4 |
| WMR | 0 | 0 | 0 | 0 | 0 |
| BMR | 0 | 0 | 0 | 0 | 0 |
| BIL | 0 | 0 | 0 | 0 | 0 |
| AMB | 280.6 | 228.7 | 287.1 | 452.4 | 386.4 |
| JCK | 834.9 | 647.4 | 599.3 | 633.2 | 681 |
| KMK | 694.6 | 1017.5 | 1306.6 | 1042.8 | 1208.2 |
| SMK | 500.6 | 610.9 | 890.5 | 615.7 | 660.9 |
| SAR | 3.8 | 986 | 628 | 909.3 | 5.4 |
| LPL | 731 | 768.3 | 1081.5 | 271.9 | 1007.6 |
| DWF | 0 | 0 | 0 | 0 | 0 |
| MEN | 456034.1 | 420734.8 | 528882.8 | 438650 | 623408.5 |
| PIN | 88.3 | 61.2 | 53.8 | 134.2 | 102.9 |
| MPL | 213.7 | 185.9 | 155.3 | 158.3 | 165.5 |
| SPL | 1395.2 | 1867.6 | 2009.5 | 1496.9 | 1565.7 |
| TIP | 382.2 | 117.3 | 121.4 | 175.4 | 228.7 |
| BEN | 0 | 0 | 0 | 0 | 0 |
| LGS | 477.9 | 585.6 | 650.9 | 652.9 | 471 |
| FIL | 0 | 0 | 0 | 0 | 0 |
| SMS | 0 | 0 | 0 | 0 | 0 |
| RAY | 5.5 | 15.4 | 2.8 | 1 | 3.2 |
| BSH | 52838.1 | 36466.2 | 56968.4 | 33909.1 | 54333.2 |
| WSH | 46157.7 | 44995 | 53315.2 | 42053.1 | 41705.1 |
| PSH | 2449.4 | 3286.9 | 3132.5 | 4539.3 | 3845.3 |
| OSH | 1171.1 | 894.8 | 581.4 | 1006.9 | 1746.2 |
| TUR | 0 | 0 | 0 | 0 | 0 |
| BCR | 26416.7 | 22346.4 | 27795.8 | 18694.7 | 25464 |
| SCR | 2686.7 | 2777.6 | 2420.4 | 2318.8 | 2512.9 |
| LOB | 1558.4 | 1355.4 | 1792.4 | 2398.3 | 2438.3 |
| SPG | 200.8 | 184.3 | 91.5 | 100.9 | 46.7 |
| CMB | 0 | 0 | 0 | 0 | 0 |
| OYS | 10262.4 | 9369.8 | 10358 | 7199 | 8439.7 |
| BIV | 59.4 | 66.5 | 68 | 70.7 | 76.3 |
| SQU | 23 | 33.2 | 30.4 | 39 | 60.4 |

Table A.4: United States Historical Recreational Landings by Atlantis Functional Group (tonnes)

| Group | 1980 | 1981 | 1982 | 1983 | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GAG | 992.2 | 829.9 | 1459.2 | 2893.6 | 884.7 | 2980.5 | 1631.8 | 1110.3 | 1699.8 | 1049.8 |
| RGR | 446.1 | 446.1 | 739.6 | 1577.6 | 3218 | 1533 | 1088.8 | 664.4 | 1123.1 | 1252.5 |
| SCM | 34.1 | 34.1 | 40.6 | 54.9 | 5.2 | 6.9 | 52.4 | 10.1 | 19.4 | 21.5 |
| SSR | 1236.6 | 1236.6 | 1347.4 | 4753.9 | 1678.5 | 2507.4 | 4026.8 | 2311.6 | 2976.2 | 1846.4 |
| DSR | 21090.8 | 21090.8 | 22237.9 | 12566.1 | 12845.7 | 87930.8 | 22805.7 | 10596.5 | 9651.9 | 45938.7 |
| RSN | 3669.7 | 3669.7 | 3255.4 | 5683.7 | 2095.1 | 2069.7 | 1816.5 | 1455.6 | 1757.4 | 1537.4 |
| VSN | 52.2 | 52.2 | 6.1 | 34.3 | 44.9 | 128.4 | 482.3 | 500.2 | 665.1 | 416.1 |
| LUT | 138125.2 | 138125.2 | 1535.9 | 64608 | 123888.6 | 7214.3 | 29803.7 | 54095.4 | 46023.9 | 27297.3 |
| BIO | 19.4 | 19.4 | 5.6 | 139.7 | 16.8 | 8 | 14 | 15.5 | 4.7 | 61.5 |
| LRF | 941.7 | 941.7 | 561.3 | 1262.1 | 2526.9 | 233.8 | 249.4 | 2004.8 | 277.2 | 106.7 |
| SRF | 2175.8 | 2175.8 | 5854.4 | 1239.6 | 860.1 | 660.8 | 2089.8 | 2728.3 | 1002.6 | 695.3 |
| BDR | 685.7 | 685.7 | 1349.4 | 1589.8 | 800.5 | 912.8 | 1350.7 | 1799.6 | 1098.9 | 936.3 |
| RDR | 2248.2 | 2248.2 | 3428.2 | 3703.3 | 3577.6 | 3435.9 | 2853.5 | 2675.4 | 1820.1 | 3050.1 |
| SEA | 8504.4 | 8504.4 | 10298.5 | 12081.6 | 11544.1 | 10475.4 | 12391.3 | 1665.1 | 8085.3 | 6172.4 |
| SCI | 1327.5 | 1327.5 | 1303.7 | 1112.4 | 1010 | 1045.2 | 1442 | 120.4 | 583.6 | 249.1 |
| LDY | 528.9 | 528.9 | 216.2 | 128.5 | 206.8 | 234 | 172.2 | 130.1 | 152 | 65.3 |
| MUL | 979.3 | 979.3 | 1064.9 | 6272.4 | 10711.2 | 11478.4 | 6321.3 | 2674.9 | 3190 | 1072 |
| POM | 11.4 | 11.4 | 64.6 | 345.6 | 98.6 | 29.1 | 60.9 | 62.4 | 30.4 | 50.4 |
| SHP | 875.8 | 875.8 | 1104.8 | 2032.6 | 1499.7 | 1575.1 | 1164.5 | 985.8 | 2181.2 | 2346.3 |
| SNK | 31 | 31 | 23.6 | 35 | 0 | 16.8 | 7.2 | 18.8 | 19.8 | 8.8 |
| FLT | 312.9 | 312.9 | 3096.9 | 4475.3 | 774.4 | 913.4 | 897.4 | 448.8 | 557.8 | 318.7 |
| ODF | 40392.9 | 40392.9 | 232027.2 | 57408.7 | 60719.3 | 287598.9 | 245347.4 | 214242.1 | 313932.5 | 74275 |
| SDF | 0.9 | 0.9 | 9.5 | 2.5 | 2.2 | 0.6 | 0.9 | 0.5 | 6.5 | 5.5 |
| YTN | 0 | 0 | 71.3 | 0 | 109.6 | 0 | 115.4 | 13.7 | 48.6 | 20.2 |
| BTN | 0 | 0 | 0 | 4.1 | 9.5 | 0.4 | 2.8 | 6.2 | 0.7 | 0 |
| LTN | 293.7 | 293.7 | 419.7 | 292.4 | 190.4 | 167 | 743.4 | 610 | 568.7 | 308 |
| OTN | 6 | 6 | 5 | 0.6 | 0.5 | 0 | 1.7 | 2.7 | 0 | 0 |
| SWD | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| WMR | 87.6 | 81.3 | 30.5 | 12.2 | 16.1 | 6.1 | 5.7 | 16.9 | 3.7 | 0.7 |
| BMR | 43.1 | 66.4 | 3269.1 | 3754 | 453.2 | 11427.4 | 1203.9 | 799.3 | 4995.3 | 18.1 |
| BIL | 464.9 | 464.9 | 762.6 | 13 | 682.8 | 15 | 268.8 | 207.7 | 71.7 | 12.5 |
| AMB | 261 | 283.7 | 2094.5 | 1220.8 | 592.8 | 1055.3 | 2634 | 2106.8 | 1024 | 1588.8 |
| JCK | 1828.7 | 1828.7 | 8004.8 | 30669.1 | 13265.5 | 7785.2 | 9947.4 | 8595.2 | 7307.7 | 23103.9 |
| KMK | 1695.1 | 4536.1 | 5949 | 2360 | 2521.8 | 1309.9 | 1452.2 | 3493.6 | 2704.1 | 2237 |
| SMK | 65.7 | 65.7 | 65.7 | 3717.3 | 1504.7 | 1619.5 | 11585.1 | 143.4 | 24.3 | 1759.7 |
| SAR | 3.9 | 3.9 | 2 | 7.9 | 10.7 | 22.1 | 0 | 1.2 | 0 | 0.4 |
| LPL | 21673.1 | 21673.1 | 12376.3 | 15977.2 | 8819.5 | 12481.2 | 25097.2 | 15537.4 | 10401.7 | 13981.1 |
| DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| MEN | 210 | 38 | 54 | 24 | 5 | 449 | 258 | 209 | 488 | 440 |
| PIN | 368.7 | 368.7 | 423.7 | 372.1 | 622.6 | 217.8 | 363.2 | 188.5 | 375.1 | 309.7 |
| MPL | 98 | 98 | 117.6 | 88.6 | 4.1 | 9.4 | 160.9 | 65.4 | 11.9 | 48 |
| SPL | 279.3 | 279.3 | 170.5 | 364.4 | 254.9 | 54.5 | 50.7 | 239.1 | 103.5 | 85.4 |
| TIP | 45.8 | 45.8 | 162.9 | 29.1 | 60.3 | 250 | 500 | 184.8 | 397.2 | 322.1 |
| BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LGS | 9068.6 | 9068.6 | 7832.1 | 10199.8 | 12170.8 | 11404.7 | 16225.9 | 7752.1 | 15590.6 | 8700.2 |
| FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 25.1 | 25.1 | 109.7 | 77 | 379.8 | 466.7 | 146.4 | 77.9 | 199.6 | 170.2 |
| LOB | 640.2 | 798.9 | 692.8 | 717.2 | 535.7 | 582.3 | 564.5 | 528.1 | 627.5 | 841.3 |

Table A.4: Continued

| Group | 1990 | 1991 | 1992 | 1993 | 1994 | 1995 | 1996 | 1997 | 1998 | 1999 | 2000 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GAG | 571.5 | 1246.6 | 1018.7 | 1264.6 | 907.1 | 1224.8 | 1067.5 | 1167.1 | 1596.3 | 1688.2 | 2255.5 |
| RGR | 511.5 | 805.2 | 1205.1 | 948.5 | 820.2 | 844.8 | 405.4 | 255.1 | 291.7 | 522.9 | 956 |
| SCM | 2.3 | 8.6 | 14.8 | 17.5 | 31.5 | 2.2 | 5.5 | 30.6 | 36.9 | 46.7 | 12.3 |
| SSR | 353.6 | 236 | 233.9 | 225.6 | 243.6 | 214.5 | 210.1 | 269.5 | 177.2 | 108.9 | 105 |
| DSR | 15256 | 3815 | 1195.6 | 3289.8 | 3629.9 | 3518.9 | 732.6 | 570.3 | 1096 | 1018.4 | 373.3 |
| RSN | 964 | 1511.7 | 2217.9 | 3002.3 | 2263.6 | 1936.4 | 1674.6 | 2252.8 | 1769.8 | 1434.9 | 1428.4 |
| VSN | 570 | 627.4 | 657 | 581.2 | 477.9 | 594 | 283.2 | 300.6 | 152 | 204.5 | 161.8 |
| LUT | 31589.9 | 37691.2 | 39300.8 | 40448.9 | 19691 | 24318.1 | 15058.2 | 10916.5 | 9706.1 | 14222.9 | 3585.5 |
| BIO | 19.2 | 2.8 | 0 | 3.1 | 36.3 | 9 | 2 | 0 | 0 | 0 | 0 |
| LRF | 742.2 | 157.7 | 516.5 | 576.6 | 644.8 | 432 | 267.4 | 267.4 | 252.3 | 267.2 | 156 |
| SRF | 645.4 | 820.2 | 688.3 | 762.2 | 795 | 768.4 | 982.1 | 1156.8 | 497 | 535.1 | 411 |
| BDR | 463.1 | 619.9 | 773.2 | 775.4 | 664.3 | 713.5 | 651.7 | 915.7 | 1052.7 | 658.5 | 1610.9 |
| RDR | 2201.9 | 2724.9 | 4004.6 | 4736.5 | 4161.4 | 6144.5 | 6016.4 | 6132.4 | 4492.1 | 4984.2 | 7260.7 |
| SEA | 3781.7 | 6523.2 | 4760.3 | 4591.2 | 5466.8 | 5558.2 | 5332.6 | 5209.1 | 3636 | 5318.4 | 4501.9 |
| SCI | 262.6 | 475.9 | 318.5 | 194.8 | 282.7 | 264.5 | 310.5 | 253.1 | 322.9 | 358.3 | 329 |
| LDY | 58.3 | 39.2 | 105.2 | 30.7 | 84 | 47.5 | 62.6 | 40.1 | 93.4 | 46.3 | 124.3 |
| MUL | 388.8 | 1860.5 | 1067.3 | 1501.1 | 908.9 | 660.2 | 1023.2 | 699.9 | 642.6 | 741.1 | 1102.2 |
| POM | 4.1 | 564.4 | 82.3 | 18.5 | 48.6 | 65.3 | 54.7 | 36.2 | 383 | 72.5 | 47.4 |
| SHP | 1216.3 | 1538.3 | 2346.5 | 2201.8 | 1378 | 2405.3 | 1726.1 | 1959.6 | 1790.2 | 1815.2 | 1698.2 |
| SNK | 0.6 | 7 | 12.8 | 14.8 | 6.1 | 10.9 | 5.8 | 48.1 | 13.7 | 29.8 | 10 |
| FLT | 579.9 | 676.5 | 371 | 348.8 | 292.9 | 282 | 222.9 | 251.6 | 233.2 | 327.7 | 189.6 |
| ODF | 88709.9 | 82406.3 | 38978 | 51599.6 | 61849.9 | 76264.3 | 66112 | 60576.2 | 40935.4 | 31167.5 | 29843.4 |
| SDF | 0 | 0.6 | 0.7 | 2.1 | 0.6 | 1 | 0.8 | 2.7 | 1.7 | 0 | 4.3 |
| YTN | 0 | 39.2 | 76.6 | 312.5 | 30.9 | 0 | 2.9 | 34.8 | 57.1 | 115.1 | 112.8 |
| BTN | 0 | 1.9 | 0 | 0 | 15.1 | 0 | 0 | 0 | 0 | 0 | 0.6 |
| LTN | 655.1 | 1106.1 | 679.6 | 412.1 | 609.7 | 369.4 | 359.6 | 282.4 | 313.3 | 311.4 | 259.1 |
| OTN | 0.7 | 0 | 0 | 0 | 0 | 0 | 12.7 | 2.9 | 0 | 0 | 0.6 |
| SWD | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| WMR | 1.1 | 1 | 1.1 | 0.6 | 1 | 0.8 | 0.6 | 0.8 | 0.2 | 0.1 | 0 |
| BMR | 16.5 | 196.2 | 16.4 | 9.4 | 17.1 | 18.6 | 10.7 | 13.5 | 5.7 | 10.3 | 6.2 |
| BIL | 75.7 | 144.6 | 34.6 | 122.6 | 56.9 | 81.9 | 122.4 | 5.7 | 45 | 33.6 | 3.7 |
| AMB | 429.8 | 1345.7 | 1132.2 | 1370 | 732.5 | 394.2 | 583.8 | 538.4 | 295.1 | 384.7 | 470.6 |
| JCK | 6003.4 | 9699.6 | 4751 | 69465.3 | 17795.4 | 6885.5 | 3614.3 | 5165.3 | 8563.5 | 7714.4 | 19771.1 |
| KMK | 3162.3 | 4872.4 | 3236.6 | 4144.6 | 4428.8 | 4059.7 | 4750.9 | 4461.1 | 3782.1 | 2971.1 | 3498.6 |
| SMK | 2140.8 | 2587.2 | 3060.5 | 1931.6 | 1702.4 | 1677.1 | 1265.6 | 1218 | 1247.8 | 1854.7 | 1781.7 |
| SAR | 26 | 2.2 | 0.5 | 0.2 | 0 | 28.7 | 0.9 | 0.3 | 3.4 | 2.6 | 0.9 |
| LPL | 8613.7 | 12145.9 | 13422.1 | 17121.7 | 11245.4 | 18779.2 | 13452.6 | 20680 | 23391.2 | 17655.7 | 17692.6 |
| DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| MEN | 135 | 51 | 138 | 170 | 189 | 56 | 82 | 20 | 47 | 51 | 207 |
| PIN | 257.5 | 364.8 | 402.4 | 446.7 | 475.3 | 547.3 | 397 | 604.4 | 788.1 | 487.7 | 852 |
| MPL | 57.7 | 3.5 | 14 | 27.1 | 28.8 | 15.9 | 26.5 | 229 | 44.8 | 12.3 | 19.2 |
| SPL | 59.2 | 366.7 | 136.6 | 158.9 | 168.7 | 387.3 | 147.1 | 175.6 | 146.7 | 112.1 | 90.4 |
| TIP | 300.1 | 302.1 | 274.9 | 128.6 | 88.6 | 154.6 | 170.1 | 236.9 | 173.9 | 132 | 287.4 |
| BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LGS | 6216.1 | 3521.7 | 1953.5 | 2903 | 2462.7 | 2890 | 5336.8 | 3130.8 | 2062.8 | 1366.1 | 1599.6 |
| FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 113.2 | 43.8 | 43.6 | 22.6 | 36.3 | 33.4 | 30.4 | 4.8 | 0.5 | 7.9 | 6.8 |
| LOB | 827.3 | 720.7 | 963.4 | 613.4 | 854.2 | 830.6 | 845.3 | 847.3 | 1022.5 | 568.4 | 1088.8 |

Table A.4: Continued

| Group | 2001 | 2002 | 2003 | 2004 | 2005 | 2006 | 2007 | 2008 | 2009 | 2010 | 2011 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GAG | 1828.6 | 2011.9 | 1711.5 | 2228.7 | 1618.1 | 1087.3 | 1001.3 | 1369.5 | 579.8 | 604.3 | 262.8 |
| RGR | 602.3 | 730.8 | 578.7 | 1377.6 | 664.5 | 420 | 435.3 | 390.5 | 444.7 | 338.5 | 290.2 |
| SCM | 26.4 | 26.1 | 30.6 | 55.6 | 34.9 | 66.3 | 32.2 | 49.6 | 40.3 | 22.6 | 25.2 |
| SSR | 59.5 | 88.4 | 107.9 | 218.3 | 123.5 | 108.3 | 138.8 | 118.3 | 91.4 | 10.5 | 20.9 |
| DSR | 635.7 | 608.4 | 512.5 | 895.1 | 1098.3 | 621.9 | 390.5 | 451.9 | 568.3 | 348.6 | 817.3 |
| RSN | 1537.3 | 1982.2 | 1819.9 | 1866.2 | 1489.7 | 1735.3 | 1932.8 | 1249.2 | 1666.9 | 799.2 | 1626.6 |
| VSN | 303.7 | 248.3 | 288.9 | 396.4 | 123.2 | 161.5 | 176.8 | 161.2 | 167.6 | 114.3 | 302.8 |
| LUT | 3317.8 | 9390.9 | 16721.9 | 8973.8 | 3028.6 | 25879.3 | 34000.6 | 30384.9 | 12403.3 | 9761.6 | 5762.6 |
| BIO | 2.4 | 0 | 1.5 | 2.9 | 0 | 0 | 0.6 | 0 | 0 | 0 | 0 |
| LRF | 183.4 | 240.1 | 314.7 | 412.5 | 278.3 | 234.6 | 274.5 | 274.3 | 205.2 | 177.1 | 178.3 |
| SRF | 595.4 | 460.7 | 643.8 | 795.6 | 606.5 | 488.5 | 585.1 | 842.2 | 434.7 | 2709.7 | 592.7 |
| BDR | 1224.8 | 1238.9 | 1403 | 1341.9 | 1106.9 | 1146 | 1285.1 | 1613.7 | 1357.1 | 1310.5 | 1313.5 |
| RDR | 6771.9 | 5907.1 | 6706.5 | 7321.6 | 5640.4 | 6080.8 | 6801.2 | 7327.1 | 6141.1 | 6870.8 | 7945.6 |
| SEA | 3744.6 | 3379.4 | 2848.5 | 3254.7 | 2951.2 | 3508 | 3234 | 3250.7 | 2887.2 | 1764.6 | 3216.5 |
| SCI | 625.4 | 231.7 | 271.8 | 208.1 | 153.2 | 193.3 | 122.8 | 174.3 | 102.1 | 127.6 | 129.4 |
| LDY | 130.9 | 77.4 | 223.1 | 299.3 | 156.2 | 265.7 | 145.2 | 358.1 | 206.9 | 151.7 | 150.7 |
| MUL | 917.9 | 415.4 | 442.7 | 531.3 | 732 | 1172.4 | 500.2 | 750.1 | 192.8 | 521 | 787.9 |
| POM | 93 | 75.9 | 43.5 | 142 | 34.5 | 244.1 | 61.4 | 156.7 | 37.9 | 36.9 | 11.1 |
| SHP | 2168.9 | 1900.6 | 2440 | 3147.7 | 2531.8 | 1489 | 1700.7 | 2106.8 | 1847.7 | 1606.1 | 3289.8 |
| SNK | 16.9 | 10.4 | 8 | 35.4 | 4.9 | 3.8 | 3.9 | 1.7 | 3.1 | 0 | 0 |
| FLT | 262.7 | 197.7 | 189.8 | 198.6 | 133.7 | 104.3 | 146.6 | 118.2 | 133 | 122.4 | 184.6 |
| ODF | 38494.5 | 34102.4 | 40969 | 67956.9 | 44269.9 | 48908.3 | 79342.7 | 79270.3 | 48483.9 | 70510.6 | 139676.5 |
| SDF | 0.1 | 2.2 | 0 | 0 | 0.1 | 0.2 | 0.4 | 0.1 | 0 | 1.6 | 0 |
| YTN | 350.3 | 141.9 | 455.6 | 267 | 288.1 | 337 | 204.7 | 444.4 | 121.1 | 18.4 | 417.4 |
| BTN | 0 | 0 | 0 | 0 | 3.4 | 0 | 0 | 0 | 0 | 0 | 0 |
| LTN | 268.9 | 398.7 | 269.2 | 484.7 | 158.6 | 292.7 | 265.5 | 200.4 | 235.7 | 192.1 | 198.6 |
| OTN | 0 | 0 | 0 | 0 | 0 | 0 | 4.3 | 0 | 0 | 0 | 0 |
| SWD | 0 | 0 | 0 | 0 | 33.4 | 0 | 0 | 47.1 | 308.1 | 24.3 | 7.1 |
| WMR | 0.1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.1 | 0 |
| BMR | 6.8 | 6.4 | 60.6 | 3211.9 | 6.5 | 8.9 | 4.5 | 3.5 | 3.4 | 2.2 | 4.3 |
| BIL | 0 | 0 | 2 | 0 | 0 | 8.5 | 3.4 | 0 | 0 | 0 | 0 |
| AMB | 571.1 | 928.8 | 1206.6 | 1080.2 | 655.3 | 640.5 | 489.2 | 589.8 | 723.2 | 675 | 430.2 |
| JCK | 13856 | 8727.9 | 7732 | 8953.6 | 4109.1 | 8870.1 | 20958.2 | 13329 | 15333.2 | 7257.1 | 2957.1 |
| KMK | 3442.4 | 3573.4 | 2996.3 | 2883.3 | 2537.1 | 4937.2 | 1884.8 | 881.6 | 1789.2 | 869.1 | 855.8 |
| SMK | 3233 | 2110.6 | 1909.6 | 3230.7 | 1678.3 | 2656.6 | 1818.6 | 2614.1 | 1651.6 | 1795.1 | 1944 |
| SAR | 4 | 16.8 | 13.6 | 8.5 | 3.5 | 28.5 | 3.6 | 3.8 | 0.7 | 0.3 | 0.4 |
| LPL | 11270.6 | 9787.2 | 9947.1 | 14615 | 9106.1 | 19110.4 | 16336.7 | 17906.4 | 18674.7 | 8642.5 | 17294 |
| DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| MEN | 48 | 108 | 118 | 64 | 48 | 55 | 30 | 28 | 61 | 44 | 78.7 |
| PIN | 708.7 | 761.1 | 789.4 | 1253.2 | 587.2 | 395.9 | 632.7 | 921 | 364 | 920.6 | 680.1 |
| MPL | 11.4 | 8.2 | 16.9 | 61.8 | 0.1 | 21.5 | 29.8 | 36.6 | 18.6 | 0.5 | 0.5 |
| SPL | 456 | 266.2 | 364.8 | 568 | 260.5 | 1234.2 | 559.8 | 735.5 | 1805.4 | 476.6 | 166.1 |
| TIP | 163.2 | 134.1 | 89.7 | 125.6 | 134.2 | 73.1 | 102.9 | 29.8 | 42.6 | 57.3 | 75 |
| BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LGS | 2325.8 | 1092.4 | 950.7 | 1461.6 | 908.4 | 1105.8 | 825.9 | 442.7 | 694.4 | 3318.4 | 716 |
| FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 5.6 | 8.8 | 8.1 | 1.5 | 18.6 | 0.5 | 0.3 | 13.4 | 0.3 | 1.9 | 9.5 |
| LOB | 870.7 | 540.1 | 607.7 | 572.1 | 572.1 | 572.1 | 572.1 | 572.1 | 572.1 | 572.1 | 572.1 |

Table A.5: Mexican Historical Commercial Landings by Atlantis Functional Group (tonnes)

| Group | 1980 | 1981 | 1982 | 1983 | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 | 1990 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AMB | 1242.8 | 1906.2 | 1590.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| BCR | 6282.7 | 9142.2 | 10864 | 6841.4 | 6784.1 | 6304.8 | 4891.7 | 5569.2 | 12042.1 | 7053.6 | 8360.9 |
| BFS | 1240.2 | 1905.8 | 1624.4 | 701.5 | 940.1 | 753.8 | 332 | 587.1 | 1145.6 | 1861.2 | 2339.2 |
| BIL | 1219.2 | 1874.4 | 1590.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| BIV | 3176 | 3756.8 | 3784 | 2477.4 | 3148.2 | 2292.7 | 1943.6 | 1635 | 2200 | 2225 | 2832.7 |
| BMR | 1219.2 | 1874.4 | 1590.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| BSH | 22946.2 | 25402.5 | 19493.1 | 18763.3 | 20866 | 20304.3 | 19104.9 | 18732.5 | 37423.7 | 20640.3 | 20478.5 |
| BTN | 1219.2 | 1874.4 | 1591.7 | 662 | 893.8 | 710 | 301.1 | 547.7 | 1068.2 | 1812.9 | 2288.2 |
| CMB | 3006.4 | 4316.6 | 4927 | 2144.4 | 4367.5 | 2714.5 | 1452.8 | 1879.1 | 4550.5 | 4551.6 | 4573.3 |
| DSR | 1619.3 | 2283.2 | 1977.8 | 962.6 | 1225.3 | 1097.8 | 679.3 | 1042.2 | 2233 | 1860.4 | 2882.9 |
| FLT | 1263.7 | 1963.3 | 1634 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| JCK | 2662.5 | 3829.4 | 4439 | 3598.7 | 4711.3 | 2610.2 | 1968.1 | 2375.3 | 4322.8 | 3190 | 3325.7 |
| KMK | 3186.4 | 4415.5 | 4957.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| LGS | 4310.2 | 8466.3 | 8746.8 | 9013.9 | 10056.2 | 8787.6 | 7958.1 | 8008.5 | 17597.7 | 8322.6 | 10943.8 |
| LOB | 1436.7 | 2073.3 | 1904.7 | 889.8 | 1166.6 | 904.1 | 514 | 884.5 | 2311 | 2226.5 | 2481 |
| LOG | 2.5 | 3.8 | 14.8 | 147.6 | 101.3 | 89.7 | 0.2 | 0.4 | 0 | 0 | 0 |
| LPL | 4431.1 | 6160 | 7341.3 | 3315.8 | 3808.7 | 3358.6 | 3090.1 | 3475.7 | 5186 | 4823 | 6348.6 |
| LRF | 1339.9 | 2416.9 | 1884.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| LTN | 1291.3 | 2057.7 | 1763.2 | 655.7 | 1220.5 | 799 | 294.3 | 547.3 | 1062.4 | 1909.5 | 2470.2 |
| LUT | 2590.4 | 7111.4 | 3963.6 | 693.3 | 916.9 | 738.6 | 360.2 | 589.6 | 1150.5 | 1846.7 | 2324.9 |
| MPL | 12575.3 | 23615.1 | 32905.5 | 8365.9 | 7870.1 | 7051.8 | 5727.5 | 7096.3 | 15792.3 | 5598.3 | 4303.3 |
| MUL | 7930.6 | 6968.3 | 7424.5 | 4695.4 | 6100.2 | 4472.6 | 8948.9 | 9675.4 | 19754.9 | 9703.2 | 11425.5 |
| ODF | 4878.8 | 7587.9 | 7733 | 695.7 | 3369.6 | 3246.1 | 2725.6 | 4192.4 | 8397.9 | 6081.8 | 7052.2 |
| OSH | 1702 | 2397.3 | 1988.3 | 1058.1 | 1327 | 1139.8 | 712.3 | 951.4 | 1870.4 | 2227.8 | 2689.5 |
| OYS | 47763.2 | 36080.7 | 29167 | 31051.9 | 38086.6 | 36430 | 35177.4 | 42572.7 | 95458.6 | 1809.3 | 2285.2 |
| PIN | 1472.8 | 2187.9 | 1906.6 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| POM | 1465 | 2045 | 1764.6 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| PSH | 2184.8 | 2920.1 | 2386.1 | 1460.5 | 1771.1 | 1575.3 | 1130.4 | 1355.6 | 2678.4 | 2646.2 | 3093.8 |
| RAY | 1355.5 | 1947.6 | 2077.1 | 701.5 | 940.1 | 753.8 | 332 | 587.1 | 1145.6 | 1861.2 | 2339.2 |
| RDR | 1903.7 | 2527.6 | 2225.9 | 1624.6 | 1850.9 | 1507.1 | 1386.6 | 1432.3 | 2873.1 | 2510.5 | 3078 |
| RGR | 8420.4 | 9233.2 | 8563.9 | 6178.3 | 7044.9 | 7788.7 | 7223.7 | 9455 | 22132.9 | 2729.5 | 13045.4 |
| RSN | 2361 | 3207 | 3441 | 3073.9 | 4082.9 | 3763.7 | 3977.1 | 4839 | 7881.3 | 3773.5 | 7067 |
| SAR | 1219.2 | 1986.9 | 1693.8 | 3282.1 | 6184.6 | 3575.4 | 1903.7 | 1348.4 | 2096.6 | 3684.4 | 2912 |
| SCI | 1860.6 | 2596.5 | 2372.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| SCR | 1333.2 | 2038.1 | 1799.2 | 795 | 1015.9 | 830.3 | 397.8 | 660.4 | 1309.6 | 1927.4 | 2422 |
| SDF | 1219.2 | 1874.4 | 1590.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| SEA | 2808.1 | 3775.9 | 3732.7 | 1624.6 | 1850.9 | 1507.1 | 1386.6 | 1432.3 | 2873.1 | 2510.5 | 3078 |
| SHP | 1472.8 | 2187.9 | 1906.6 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| SMK | 3487.6 | 4655.9 | 5062.7 | 3302.5 | 3457.6 | 3254.7 | 3080.7 | 3475.3 | 5168.8 | 4703.7 | 6155.2 |
| SMS | 1261.3 | 1937.2 | 1658.4 | 747.3 | 997.1 | 803.4 | 369.8 | 626.8 | 1228.9 | 1913.2 | 2393.2 |
| SNK | 3767.5 | 5234.6 | 6331.4 | 4499.5 | 3876.7 | 4030.8 | 3244.5 | 3474.8 | 6983.6 | 3709.3 | 3782.6 |
| SPL | 2763.2 | 3567.2 | 4500.2 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| SQU | 7104.1 | 8133.1 | 7449 | 8406.7 | 6115.6 | 6453.9 | 8752.9 | 7905.5 | 14791.7 | 2849.8 | 16124.3 |
| SRF | 1368 | 2209.2 | 2174.2 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| SSR | 4150.3 | 4421.5 | 3955.5 | 962.6 | 1225.3 | 1097.8 | 679.3 | 1042.2 | 2233 | 1860.4 | 2882.9 |
| SWD | 1219.2 | 1874.4 | 1590.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| TIP | 2482 | 3758.8 | 3629.4 | 3403.4 | 4307.2 | 3680.5 | 2557.6 | 2932.5 | 6058.1 | 4926.6 | 5527.6 |
| TUR | 22.9 | 34.3 | 133.1 | 1328.4 | 912 | 807.5 | 1.6 | 3.3 | 0 | 0 | 0 |
| VSN | 1395.2 | 2308.1 | 2083.9 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| WMR | 1219.2 | 1874.4 | 1590.4 | 655.7 | 883 | 704.2 | 294.3 | 547.3 | 1062.4 | 1809.3 | 2285.2 |
| WSH | 2184.8 | 2920.1 | 2386.1 | 1460.5 | 1771.1 | 1575.3 | 1130.4 | 1355.6 | 2678.4 | 2646.2 | 3093.8 |
| YTN | 1219.2 | 1874.4 | 1701.8 | 1195.9 | 1819.2 | 1201 | 876 | 576.5 | 1562.8 | 2121.4 | 2547.7 |

Table A.5: Continued

| Group | 1991 | 1992 | 1993 | 1994 | 1995 | 1996 | 1997 | 1998 | 1999 | 2000 | 2001 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AMB | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 1356.5 | 1216.9 | 957.3 | 899.4 | 789.8 |
| BCR | 9591.7 | 36327.5 | 11940.7 | 11888.7 | 11531.8 | 14014.8 | 14126.5 | 12667.3 | 12079.7 | 8473.9 | 7252 |
| BFS | 2468.1 | 10688.9 | 2319.9 | 2353.9 | 2134.9 | 1886.6 | 1390.4 | 1252.6 | 986.3 | 927.2 | 818.6 |
| BIL | 2422.2 | 10456.8 | 2264.8 | 2309 | 2092.5 | 1840.3 | 1358.1 | 1217.9 | 958.8 | 902.6 | 789.8 |
| BIV | 4119.3 | 19048.5 | 2344.8 | 2821.1 | 2782.8 | 2444.7 | 1852.9 | 1870.1 | 1578.5 | 2008.9 | 1851.2 |
| BMR | 2422.2 | 10456.8 | 2264.8 | 2308.9 | 2092.5 | 1840.3 | 1363.3 | 1222.1 | 961.1 | 913.7 | 789.8 |
| BSH | 23350.7 | 101445.4 | 19778.6 | 19285 | 19879.2 | 18265.1 | 18457.3 | 19869.9 | 16827 | 17915.4 | 18178.9 |
| BTN | 2427.3 | 11716.9 | 2272.7 | 2317.4 | 2104.1 | 1848.6 | 1367.7 | 1228.7 | 977.1 | 914.4 | 802.5 |
| CMB | 4675.9 | 22640.4 | 6165.3 | 6000.5 | 6071.5 | 3860.4 | 5607 | 3990.3 | 7025.4 | 7828.5 | 8099 |
| DSR | 3060.6 | 12342.6 | 2871.4 | 2883.8 | 2682 | 2313.9 | 1830.2 | 1701.9 | 1432.1 | 1454.3 | 1228.1 |
| FLT | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 1537.9 | 1411.1 | 1134.2 | 1001 | 948.6 |
| JCK | 4233 | 17082.8 | 4533.6 | 4719 | 4886.6 | 7352.3 | 10468.2 | 10470.2 | 5253.8 | 7688.8 | 7423.2 |
| KMK | 2422.2 | 10456.8 | 2262.9 | 2305 | 3493.6 | 3891.4 | 3714.7 | 3248.6 | 3191.2 | 2926.5 | 3085 |
| LGS | 8679.2 | 51992.8 | 10135.6 | 9925.1 | 9579 | 9920.7 | 6991.1 | 6387.5 | 6037 | 5652.5 | 5349.9 |
| LOB | 2896.8 | 13241.1 | 2801.7 | 2606.9 | 2511.3 | 2183.9 | 1823.1 | 1476.3 | 1208.2 | 1260.1 | 1184.7 |
| LOG | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LPL | 7029.9 | 36502.5 | 7283.3 | 6504.1 | 7484.8 | 9491.1 | 7997.1 | 7602 | 7402.6 | 6334.1 | 6273.8 |
| LRF | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 6399.5 | 6977.5 | 7056.3 | 7489.2 | 5403.8 |
| LTN | 2956.6 | 11539.5 | 2707 | 2691.4 | 2609.1 | 2431.8 | 2280.9 | 2210.3 | 1304.5 | 1678.2 | 1473.4 |
| LUT | 2482 | 10713.6 | 2332.1 | 2351.1 | 2137.3 | 1882.3 | 2067.4 | 3433.5 | 2902.1 | 2504 | 2865.7 |
| MPL | 3431 | 77238.4 | 6048.6 | 6414.4 | 5179.2 | 5632.6 | 4429.4 | 3737.6 | 3105 | 2715.1 | 2669.1 |
| MUL | 9233.6 | 37457.8 | 12004.8 | 12441.8 | 14258.6 | 13716.6 | 14189.7 | 12389.4 | 12137.6 | 13141.5 | 11231.4 |
| ODF | 7920.3 | 30116.2 | 8182.1 | 8689.9 | 8210.9 | 7911.2 | 8924.6 | 9212.9 | 7679.7 | 7683 | 7734.3 |
| OSH | 2887.3 | 12478.7 | 2652.2 | 2682.3 | 2487.7 | 2205.3 | 1736.5 | 1631.4 | 1310 | 1277.6 | 1176.2 |
| OYS | 2422.2 | 65833.1 | 2262.9 | 3097.8 | 3353.5 | 1840.3 | 5650.2 | 1216.9 | 979.1 | 899.4 | 789.8 |
| PIN | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 1356.5 | 1216.9 | 957.3 | 899.4 | 789.8 |
| POM | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 1934.3 | 1713.2 | 1432.7 | 1265 | 1078.3 |
| PSH | 3352.4 | 14500.7 | 3041.4 | 3059.6 | 2883 | 2570.3 | 2116.5 | 2045.9 | 1662.6 | 1655.7 | 1562.7 |
| RAY | 2468.1 | 10688.9 | 2319.9 | 2353.9 | 2134.9 | 1886.6 | 6444.3 | 6977 | 4697.6 | 3511.8 | 3189.1 |
| RDR | 2931.2 | 13695 | 3057.6 | 2885.1 | 2723.4 | 2756.5 | 2185.6 | 2215.7 | 1943 | 1973.1 | 1595.4 |
| RGR | 13913.2 | 44402.2 | 13215 | 12723.1 | 12213.4 | 9875.1 | 9884.4 | 9946.6 | 9503.6 | 10887.3 | 8679.3 |
| RSN | 7523.2 | 33762.9 | 8871.6 | 6778.9 | 6400.2 | 5998 | 5241 | 4355.9 | 4152.5 | 3435.3 | 3326.8 |
| SAR | 2913 | 241379.8 | 3165.6 | 4778 | 2400.9 | 2357.4 | 2270.9 | 4905.5 | 2201.1 | 1388.4 | 1128.2 |
| SCI | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 2453.2 | 2857.1 | 2384.3 | 1449.2 | 1228.9 |
| SCR | 2583.7 | 11039.3 | 2480.9 | 2520.8 | 2305 | 2114.4 | 1644 | 1474.8 | 1207.8 | 1070 | 935.3 |
| SDF | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 5792.6 | 4802.1 | 4247.7 | 3285.3 | 3460.6 |
| SEA | 2931.2 | 13695 | 3057.6 | 2885.1 | 6006.3 | 5595.1 | 5418.2 | 5473.4 | 5242.6 | 4903.1 | 3892.9 |
| SHP | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 1356.5 | 1216.9 | 957.3 | 899.4 | 789.8 |
| SMK | 6489.1 | 25003.9 | 6817.9 | 6037.6 | 5552.5 | 6836.2 | 4696.7 | 4555.8 | 4750.2 | 3486.7 | 3196.6 |
| SMS | 2513.9 | 10921 | 2376.9 | 2402.8 | 2177.4 | 1932.9 | 1424.3 | 1288.3 | 1015.4 | 955 | 847.4 |
| SNK | 4323.3 | 15784.7 | 4087.7 | 4328.1 | 4193.5 | 4205.9 | 3676.8 | 3286.9 | 3286.1 | 3037.8 | 3578.5 |
| SPL | 2422.2 | 13911.8 | 2262.9 | 2319.5 | 2092.5 | 1840.3 | 2175.6 | 2815.4 | 2420.6 | 1953.6 | 1993.7 |
| SQU | 16202.4 | 55342.5 | 16441.3 | 17405.1 | 19015.1 | 27494.6 | 17379.2 | 16001.3 | 18262.8 | 21255.7 | 19388.9 |
| SRF | 2422.2 | 13911.8 | 2262.9 | 2319.5 | 2092.5 | 1840.3 | 1366.4 | 1265 | 1019.9 | 940.2 | 815.2 |
| SSR | 3060.6 | 12342.6 | 2871.4 | 2883.8 | 2654.7 | 2286.7 | 2122.3 | 1980.4 | 1728.7 | 1572.2 | 1413.2 |
| SWD | 2422.2 | 10456.8 | 2266.9 | 2310.1 | 2092.5 | 1840.3 | 1363 | 1222.8 | 964 | 911.5 | 789.8 |
| TIP | 5172.6 | 24383 | 5680.1 | 5239.3 | 4639.3 | 4617.9 | 3393.3 | 3358.8 | 2699.6 | 2567.7 | 2517.5 |
| TUR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| VSN | 2422.2 | 10456.8 | 2262.9 | 2305 | 2092.5 | 1840.3 | 3282.4 | 2580.4 | 2457.8 | 2308.3 | 2743 |
| WMR | 2422.2 | 10456.8 | 2264.1 | 2307.1 | 2092.5 | 1840.3 | 1359.6 | 1217.7 | 958.9 | 907.8 | 789.8 |
| WSH | 3352.4 | 14500.7 | 3041.4 | 3059.6 | 2883 | 2570.3 | 2116.5 | 2045.9 | 1662.6 | 1655.7 | 1562.7 |
| YTN | 2860.7 | 119129.7 | 3100.1 | 3376.4 | 3091.4 | 2559.9 | 2325.4 | 2232.7 | 2668.8 | 2189.3 | 1880.7 |

Table A.5: Continued

| Group | 2002 | 2003 | 2004 | 2005 | 2006 | 2007 | 2008 | 2009 | 2010 | 2011 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AMB | 636.8 | 719.5 | 704.6 | 701.7 | 585.2 | 557.5 | 469.5 | 411.1 | 545.2 | 374.6 |
| BCR | 7783.3 | 10443.3 | 11117.1 | 10740.1 | 10710.6 | 10504.8 | 9972.3 | 8151.7 | 11912.3 | 9660.8 |
| BFS | 664.1 | 744.9 | 732.7 | 734.7 | 606.6 | 576.6 | 488.4 | 432.9 | 571.5 | 391.3 |
| BIL | 636.8 | 721.5 | 706 | 702.9 | 586.3 | 559.4 | 471.2 | 412.2 | 547 | 375.3 |
| BIV | 1617.5 | 1714.1 | 1702.7 | 2239.5 | 2767 | 2503.1 | 2231.7 | 1756.6 | 1781.3 | 1829.4 |
| BMR | 636.8 | 738.4 | 726.7 | 717.8 | 601.9 | 576.6 | 490.2 | 425.5 | 570.5 | 392.9 |
| BSH | 15626.8 | 19686 | 15392 | 22584.3 | 16008.2 | 17272.9 | 15870.8 | 16863.4 | 20449.7 | 17503.4 |
| BTN | 651.1 | 736.6 | 719.9 | 716 | 598.2 | 569.3 | 482.4 | 426.9 | 559.4 | 392 |
| CMB | 5927 | 5725.8 | 7182.4 | 6901.5 | 6689.5 | 6565.2 | 3332.1 | 3678.1 | 7348.4 | 5041 |
| DSR | 1122.1 | 1095.2 | 1072 | 1198.9 | 1006.8 | 1128.4 | 1018.2 | 978.1 | 1097.8 | 760.5 |
| FLT | 875.4 | 891.9 | 887 | 879.8 | 789.7 | 701.6 | 579.7 | 455 | 648.8 | 465.7 |
| JCK | 7452 | 9149.1 | 8832.1 | 9227.9 | 9081.2 | 8871.9 | 8654.3 | 10438.8 | 13143 | 11021.9 |
| KMK | 3213.6 | 3099.9 | 3196.2 | 2910.1 | 2803.8 | 2543.9 | 2880 | 2426.5 | 2755 | 2208.2 |
| LGS | 4342 | 4042.8 | 3958.8 | 4529.9 | 4007.7 | 3606.7 | 2880.7 | 2738.3 | 3657.1 | 2676.8 |
| LOB | 1184.5 | 1103.2 | 1048.2 | 1070.9 | 813.5 | 854.9 | 686.6 | 604.5 | 935.6 | 621.7 |
| LOG | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| LPL | 6739 | 7196.1 | 7230.6 | 6925.2 | 5974.6 | 5882.2 | 6878.1 | 6004.9 | 7236.1 | 5375.6 |
| LRF | 4149.9 | 5208.2 | 6002.6 | 3087.9 | 3012.3 | 3786 | 4540.2 | 2681.5 | 2448 | 2449.2 |
| LTN | 1272 | 1429.4 | 1405.4 | 1323.4 | 1305 | 1088.6 | 1000.9 | 1042.9 | 1360.3 | 1069.8 |
| LUT | 2399.4 | 2705.4 | 2356.9 | 2661.8 | 2151.1 | 2452.8 | 2736.4 | 2058.4 | 2400.4 | 2232.5 |
| MPL | 1838.7 | 2274.5 | 2585.6 | 2355.1 | 2686.5 | 2236.1 | 1860.6 | 1319.7 | 1721.9 | 1761.5 |
| MUL | 9263.6 | 8749.4 | 8387.3 | 8299.4 | 6888.3 | 7647.8 | 6976.5 | 7848.9 | 8096.1 | 7063.1 |
| ODF | 6991.5 | 7361 | 6670.3 | 6469.3 | 7486.4 | 5417.4 | 5151 | 5403.5 | 5893.2 | 3894.9 |
| OSH | 969.9 | 1141 | 1031 | 1188 | 927.9 | 929 | 811.8 | 776.7 | 987.6 | 755.3 |
| OYS | 636.8 | 893.7 | 788.1 | 3604.3 | 2632.5 | 2121.5 | 1683.9 | 2307.9 | 2411.2 | 1903.6 |
| PIN | 636.8 | 719.5 | 704.6 | 701.7 | 585.2 | 557.5 | 469.5 | 411.1 | 545.2 | 374.6 |
| POM | 1166.3 | 1262.9 | 1081.1 | 1223.3 | 884.3 | 938.9 | 841.6 | 887.6 | 1028.7 | 749.7 |
| PSH | 1303 | 1562.5 | 1357.4 | 1674.2 | 1270.6 | 1300.4 | 1154 | 1142.3 | 1429.9 | 1135.9 |
| RAY | 2841.1 | 3069.1 | 3262.8 | 3711.3 | 3632.7 | 3558.5 | 2914.2 | 2898.9 | 3382.9 | 3665.1 |
| RDR | 1680.4 | 2122.9 | 2039.6 | 1950.5 | 1499 | 1234.5 | 1076.9 | 1055.2 | 1359.4 | 1030.5 |
| RGR | 9371.2 | 7481.5 | 7317.2 | 9652.5 | 8175.5 | 10834.3 | 10345.5 | 10617 | 9369.6 | 7319.5 |
| RSN | 3055.6 | 3220.2 | 3419.5 | 3938.6 | 3228.1 | 3471.7 | 3094.6 | 3351.9 | 4198.2 | 3145.2 |
| SAR | 793.7 | 945.4 | 848.9 | 914.3 | 734.1 | 669 | 511 | 479.4 | 596.2 | 411.2 |
| SCI | 1208.2 | 1114.1 | 1266.2 | 1161.7 | 962.8 | 756.6 | 570 | 519.9 | 680.1 | 480.4 |
| SCR | 797.7 | 938.5 | 939.1 | 927.7 | 813.2 | 781.5 | 683.5 | 585.4 | 801.2 | 583.7 |
| SDF | 3764.3 | 3664.2 | 3446.1 | 3546.5 | 2790.2 | 2285.9 | 2217.4 | 2495.2 | 3401.2 | 2371.6 |
| SEA | 4450.8 | 5158.9 | 5620.4 | 4717.1 | 4086 | 2646.8 | 3036.1 | 2954.1 | 3646.8 | 3012.5 |
| SHP | 636.8 | 719.5 | 704.6 | 701.7 | 585.2 | 557.5 | 469.5 | 411.1 | 545.2 | 374.6 |
| SMK | 3411.4 | 3961.8 | 3876.6 | 3922.4 | 2866.1 | 3268.6 | 3903.1 | 3330.4 | 4185.9 | 2815.7 |
| SMS | 691.4 | 770.3 | 760.7 | 767.7 | 628 | 595.7 | 507.3 | 454.7 | 597.7 | 407.9 |
| SNK | 4156.2 | 5109.3 | 4309.4 | 4794.7 | 4570.2 | 4712.8 | 4408.3 | 4528.1 | 4946.3 | 3531 |
| SPL | 1597.9 | 2215.4 | 2155.2 | 2751.4 | 2132.5 | 1334.3 | 1487.7 | 601.3 | 1409.3 | 1151.4 |
| SQU | 15171 | 14914.2 | 22549.6 | 10556.9 | 26657.6 | 19226.2 | 11681.6 | 24259.2 | 22463.5 | 26057.5 |
| SRF | 643.7 | 757.6 | 729.1 | 712.8 | 597.2 | 571.7 | 472.6 | 414.4 | 1074.5 | 1061.9 |
| SSR | 1294.3 | 1273 | 1322.4 | 1440 | 1231.5 | 1239.1 | 1070.8 | 1020.2 | 1088 | 800 |
| SWD | 636.8 | 728 | 719.1 | 713.1 | 595.1 | 570.8 | 481.9 | 416.8 | 565.3 | 385.8 |
| TIP | 2275.2 | 2244.1 | 2387.6 | 2681.7 | 1869 | 1702.4 | 1604.3 | 1719.2 | 2120 | 1374.3 |
| TUR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| VSN | 2166.9 | 2143.8 | 2049.1 | 2445.6 | 2233.5 | 2481.7 | 1441.3 | 1377.8 | 1521.2 | 971.6 |
| WMR | 636.8 | 724.2 | 712.5 | 707.3 | 589.6 | 560.3 | 473.8 | 415.7 | 556.2 | 387.2 |
| WSH | 1303 | 1562.5 | 1357.4 | 1674.2 | 1270.6 | 1300.4 | 1154 | 1142.3 | 1429.9 | 1135.9 |
| YTN | 1869.9 | 2197.6 | 2021.9 | 1940.8 | 1709.4 | 1578.3 | 1576.6 | 1772.8 | 1763.2 | 1873.4 |

Table A.6: Cuban Historical Commercial Landings by Atlantis Functional Group (tonnes)

| Group | 1980 | 1981 | 1982 | 1983 | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 | 1990 | 1991 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| BCR | 0 | 80.5 | 104.4 | 132.1 | 147.5 | 153.6 | 174.8 | 132.1 | 216.2 | 227 | 0 | 139.9 |
| BFT | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1810.3 | 0 |
| BIO | 1983 | 1356.9 | 1715.6 | 2130.5 | 2234.8 | 2327 | 2419.4 | 2339.5 | 2342.7 | 2297.8 | 645 | 1691.9 |
| BIV | 0 | 263.1 | 682.3 | 525.8 | 583.4 | 653.9 | 683.4 | 660.1 | 649.4 | 698.5 | 33.5 | 475.3 |
| BMR | 102.1 | 155.3 | 159.4 | 105.3 | 44.3 | 54.9 | 34.8 | 4 | 43.6 | 18.4 | 30.6 | 38 |
| CMB | 0 | 0 | 28.4 | 281.9 | 263.5 | 250.9 | 93 | 78.2 | 80.8 | 100.4 | 96 | 30.3 |
| JCK | 30 | 13 | 28 | 23 | 38 | 44 | 42 | 56 | 24 | 24 | 3.3 | 45 |
| KMK | 5.7 | 3.3 | 0.1 | 0.3 | 1.3 | 0.7 | 0.5 | 0.1 | 0.3 | 0.3 | 197 | 1.3 |
| LGS | 369 | 702 | 561 | 445 | 422 | 335 | 319 | 470 | 524 | 471 | 1242.2 | 122 |
| LOB | 1551.3 | 1575.3 | 1704.6 | 1596.8 | 1848.7 | 1985.3 | 1728.8 | 2059 | 1899.2 | 1744.3 | 77 | 1496.8 |
| LOG | 200.3 | 155.9 | 162.8 | 170.9 | 173.4 | 201.6 | 193.4 | 149.6 | 115.8 | 88.9 | 606.3 | 52.6 |
| LPL | 723.4 | 556.3 | 628.6 | 624 | 499.8 | 498.7 | 433.9 | 392.1 | 558.3 | 593.3 | 1810.3 | 501.3 |
| LRF | 1983 | 1356.9 | 1715.6 | 2130.5 | 2234.8 | 2327 | 2419.4 | 2339.5 | 2342.7 | 2297.8 | 22.1 | 1832.6 |
| LTN | 0 | 0 | 11.6 | 2.1 | 5.3 | 5.6 | 8.4 | 19.3 | 18.6 | 29.1 | 574.7 | 22.1 |
| LUT | 825.5 | 883.8 | 900.1 | 844.2 | 735.6 | 693.4 | 701.7 | 790.1 | 715.1 | 753.5 | 8 | 777.3 |
| MPL | 0 | 0 | 0 | 0 | 5 | 8 | 10 | 15 | 17 | 22 | 46.9 | 8 |
| MUL | 214.9 | 184.8 | 216.7 | 204.1 | 249.6 | 111.3 | 132.7 | 136.9 | 68.6 | 78.1 | 44.1 | 90.7 |
| OBL | 41.7 | 46.9 | 63.4 | 9.8 | 59.2 | 45.5 | 17.5 | 59.9 | 27.3 | 19.3 | 373.9 | 29.1 |
| ODF | 189.3 | 233.5 | 318.9 | 290.6 | 308.3 | 317 | 339.6 | 328 | 372.9 | 396.5 | 794.1 | 361.6 |
| OYS | 332.8 | 514.8 | 526.6 | 446.5 | 629.2 | 516.2 | 529.5 | 330.3 | 479 | 854.6 | 0 | 757.2 |
| RAY | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 407.4 | 1.1 |
| RGR | 1025.2 | 824.6 | 734.3 | 578.2 | 724.5 | 635.3 | 669.9 | 726.6 | 766.2 | 903 | 431.2 | 320.3 |
| RSN | 373.5 | 341.6 | 367.2 | 433.7 | 382.2 | 505.1 | 429.1 | 594.3 | 491.8 | 483 | 0.4 | 52.2 |
| SCR | 119.3 | 12.1 | 28.5 | 26.3 | 25.8 | 23.2 | 17.8 | 15.2 | 12.6 | 5.1 | 3.3 | 0.4 |
| SMK | 5.7 | 3.3 | 0.1 | 0.3 | 1.3 | 0.7 | 0.5 | 0.1 | 0.3 | 0.3 | 1061.6 | 1.3 |
| SPL | 829.9 | 813.8 | 833 | 963.6 | 799.4 | 979.7 | 1058.1 | 795.2 | 1001.7 | 994 | 1810.3 | 1151.2 |
| SRF | 1983 | 1356.9 | 1715.6 | 2130.5 | 2234.8 | 2327 | 2419.4 | 2339.5 | 2342.7 | 2297.8 | 3725.9 | 1691.9 |
| SSR | 4038.1 | 2886.2 | 3702.8 | 4504.1 | 4646.2 | 4911.2 | 5090.4 | 4897 | 4843.3 | 4786.3 | 20 | 3478.9 |
| SWD | 134.9 | 138.7 | 53.6 | 44.7 | 28.8 | 33.1 | 48.3 | 57.1 | 47.4 | 23.6 | 399.4 | 16.2 |
| TUR | 350.6 | 416.3 | 414.4 | 407.5 | 338.7 | 541.5 | 421.3 | 373.1 | 324.3 | 343.7 | 2.1 | 249.2 |
| WMR | 74.2 | 40.6 | 15.8 | 39.2 | 53.6 | 75.6 | 67.2 | 21.7 | 8.4 | 7.7 | 18.6 | 3.5 |
| YTN | 241.2 | 699 | 526.1 | 277.6 | 888.3 | 667.1 | 728.4 | 371.7 | 34.3 | 31.9 | 0 | 6.3 |

Table A.6: Continued

| Group | 1992 | 1993 | 1994 | 1995 | 1996 | 1997 | 1998 | 1999 | 2000 | 2001 | 2002 | 2003 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| BCR | 166.2 | 74.9 | 122.6 | 108.6 | 131 | 102.8 | 155.5 | 82.6 | 85.3 | 77.1 | 59.9 | 113.9 |
| BFT | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 25.9 | 3.9 |
| BIO | 1639.4 | 1139.7 | 1492.6 | 1286.9 | 969.2 | 1209.5 | 962.5 | 1426.4 | 1587.3 | 773.9 | 351.5 | 723.9 |
| BIV | 475.3 | 499.6 | 573.1 | 703 | 686.3 | 638 | 899.6 | 753.9 | 956.1 | 233.6 | 138.4 | 107 |
| BMR | 40.4 | 20.6 | 11.7 | 25.4 | 12.9 | 15.9 | 3.6 | 11.4 | 16.5 | 16.8 | 10.2 | 0.9 |
| CMB | 19.9 | 33.6 | 22.5 | 28.8 | 313.3 | 460.9 | 181.2 | 305.9 | 429.2 | 432.5 | 367.9 | 137.3 |
| JCK | 36 | 19 | 61 | 36.9 | 37.4 | 25.1 | 53.5 | 17.9 | 274.1 | 229 | 19.3 | 17.7 |
| KMK | 0.2 | 1.3 | 0.1 | 1.1 | 1.2 | 0.9 | 0.5 | 0.5 | 0.4 | 0.4 | 0.5 | 0.4 |
| LGS | 142 | 74 | 88 | 120 | 123.9 | 167.8 | 270.1 | 250.3 | 199.1 | 210.7 | 206.1 | 107.7 |
| LOB | 1412.4 | 1266 | 1460.6 | 1426.3 | 1423.2 | 1413.6 | 1488.2 | 1567 | 1225.1 | 1115.6 | 1201.9 | 792.2 |
| LOG | 40.1 | 30.1 | 14.4 | 6.9 | 6.3 | 4.4 | 3.1 | 3.1 | 5.6 | 4.4 | 5 | 5 |
| LPL | 514.2 | 167.3 | 360 | 289.4 | 351.3 | 462.7 | 397.7 | 482.5 | 369.9 | 310.4 | 320.5 | 364.6 |
| LRF | 1720.6 | 1205.2 | 1564 | 1356.2 | 1010.5 | 1243.4 | 975.5 | 1430.9 | 1593.3 | 779.8 | 355.4 | 725.3 |
| LTN | 11.6 | 4.6 | 5.3 | 9.5 | 8.1 | 6 | 3.2 | 1.1 | 0.7 | 0.4 | 1.1 | 1.8 |
| LUT | 494.9 | 384 | 475.4 | 506 | 767.7 | 590.4 | 543.8 | 413.9 | 464.9 | 518 | 570.3 | 458.9 |
| MPL | 8 | 8 | 8 | 9.2 | 7.1 | 5.6 | 5 | 4.2 | 4.5 | 4.6 | 3.6 | 3.3 |
| MUL | 50.4 | 55.3 | 46.9 | 37.8 | 32.6 | 55.7 | 42.7 | 37.8 | 41.7 | 39.9 | 43.8 | 27 |
| OBL | 24.5 | 14.7 | 16.1 | 11.6 | 13 | 14 | 9.8 | 68.6 | 72.8 | 23.8 | 11.2 | 6.3 |
| ODF | 295.4 | 232.7 | 318.8 | 358.5 | 306.6 | 264.5 | 214.4 | 209.3 | 219.1 | 199.9 | 196.5 | 265.5 |
| OYS | 564.2 | 440.6 | 607.7 | 695.6 | 696.7 | 742.1 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 0 | 4.2 | 0 | 1.8 | 334.3 | 475.3 | 466.9 | 473.2 | 417.6 | 391.7 | 447.3 | 465.5 |
| RGR | 150.2 | 119.4 | 104.3 | 73.9 | 94.2 | 68.3 | 24.2 | 38.5 | 59.9 | 31.9 | 30.1 | 36.1 |
| RSN | 211.4 | 166.6 | 218.1 | 283.2 | 350.7 | 318.2 | 282.5 | 267.1 | 241.2 | 270.6 | 309.8 | 345.1 |
| SCR | 1 | 0.4 | 0.3 | 0.7 | 4.1 | 1.3 | 5.1 | 7.6 | 7.9 | 11.1 | 7 | 7.3 |
| SMK | 0.2 | 1.3 | 0.1 | 1.1 | 1.2 | 0.9 | 0.5 | 0.5 | 0.4 | 0.4 | 0.5 | 0.4 |
| SPL | 792.4 | 640.2 | 873.6 | 1067.5 | 1073.8 | 996.5 | 677.3 | 882.7 | 1079.4 | 1206.1 | 806.8 | 860.7 |
| SRF | 1639.4 | 1139.7 | 1492.6 | 1286.9 | 969.2 | 1209.5 | 962.5 | 1426.4 | 1587.3 | 773.9 | 351.5 | 723.9 |
| SSR | 3399.4 | 2357.1 | 3027.2 | 2623.8 | 2015 | 2469.5 | 1970.4 | 2882.5 | 3211.1 | 1566.4 | 734.9 | 1476.1 |
| SWD | 7.5 | 2.4 | 3 | 4.5 | 2.1 | 3 | 3 | 1.5 | 3.3 | 0.9 | 3 | 0.9 |
| TUR | 289.8 | 154.6 | 102 | 49.5 | 35.7 | 23.2 | 22.5 | 12.5 | 17.5 | 10 | 14.4 | 10 |
| WMR | 3.5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| YTN | 3.9 | 0.4 | 0 | 0 | 0 | 0 | 0 | 11.9 | 99.4 | 54.6 | 22.8 | 24.9 |

Table A.6: Continued

| Group | 2004 | 2005 | 2006 | 2007 | 2008 | 2009 | 2010 | 2011 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| BCR | 62.8 | 32.9 | 51 | 48.2 | 49.6 | 44.7 | 50.4 | 58.3 |
| BFT | 6.3 | 9.5 | 6.7 | 0 | 0 | 0 | 0 | 0 |
| BIO | 576.8 | 576.5 | 580.9 | 618 | 502.2 | 569.2 | 280.8 | 293.7 |
| BIV | 163.5 | 148.7 | 138 | 142.1 | 35.1 | 109.2 | 112.2 | 69 |
| BMR | 1.2 | 2.1 | 2.1 | 0.6 | 1.2 | 1.8 | 0.9 | 1.8 |
| CMB | 203.7 | 237.6 | 120.3 | 211.8 | 148 | 197.1 | 186.7 | 181.6 |
| JCK | 7.9 | 7.3 | 5.1 | 7.7 | 6.5 | 10.8 | 4.4 | 6.3 |
| KMK | 0.2 | 0.2 | 0.3 | 0.3 | 0.4 | 0.4 | 0.3 | 0.4 |
| LGS | 50.3 | 48.4 | 78.1 | 78.7 | 79.2 | 77.6 | 54.4 | 53.9 |
| LOB | 1137.3 | 891.5 | 669.7 | 761 | 902.1 | 647.2 | 712.5 | 769.1 |
| LOG | 3.8 | 4.4 | 1.9 | 1.9 | 0 | 0 | 0 | 0 |
| LPL | 373.5 | 369.5 | 287 | 244.5 | 241.9 | 305.1 | 209.8 | 265.2 |
| LRF | 577.9 | 577.9 | 581.2 | 618.4 | 502.2 | 569.2 | 280.8 | 293.7 |
| LTN | 2.8 | 2.8 | 1.1 | 1.4 | 2.5 | 1.4 | 4.9 | 8.1 |
| LUT | 251.7 | 159.4 | 192.3 | 212.2 | 233.1 | 227.7 | 223.5 | 335.2 |
| MPL | 3 | 3.2 | 2.9 | 3 | 3.6 | 4.5 | 5.5 | 5 |
| MUL | 16.5 | 14 | 20 | 18.2 | 21 | 37.8 | 87.2 | 85.4 |
| OBL | 17.5 | 25.2 | 16.5 | 19.6 | 6.7 | 6.7 | 9.1 | 20.7 |
| ODF | 247.6 | 196.1 | 189.6 | 183 | 137 | 150.6 | 155.8 | 184.9 |
| OYS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RAY | 497.4 | 536.6 | 534.5 | 648.9 | 674.1 | 709.5 | 559.7 | 614.6 |
| RGR | 19.6 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RSN | 219.5 | 248.2 | 223.7 | 269.5 | 265.7 | 322.4 | 296.8 | 263.2 |
| SCR | 19.3 | 16.6 | 11.2 | 13.9 | 13.1 | 0 | 0 | 0 |
| SMK | 0.2 | 0.2 | 0.3 | 0.3 | 0.4 | 0.4 | 0.3 | 0.4 |
| SPL | 871.9 | 655.2 | 944.7 | 781.6 | 876.8 | 984.9 | 912.5 | 881.3 |
| SRF | 576.8 | 576.5 | 580.9 | 618 | 502.2 | 569.2 | 280.8 | 293.7 |
| SSR | 1177.3 | 1163.2 | 1176.2 | 1245.3 | 1012.9 | 1149.4 | 567.7 | 595.2 |
| SWD | 0.9 | 0.9 | 0.6 | 0.3 | 0 | 0 | 0 | 0.3 |
| TUR | 5 | 8.1 | 6.9 | 1.3 | 0 | 0 | 0 | 0 |
| WMR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| YTN | 6.7 | 6.7 | 6.7 | 12.6 | 0.7 | 0.7 | 2.8 | 0.7 |
|  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  | 0.8 |  |  |

Table A.7: Seasonal Distribution of Atlantis Functional Group Harvested by United States Commercial Fleets

| Group | Winter (Jan.-Mar.) | Spring (Apr.-Jun.) | Summer (Jul.-Sep.) | Fall (Oct. - Dec.) |
| :---: | :---: | :---: | :---: | :---: |
| GAG | 0.2837 | 0.2954 | 0.1858 | 0.2351 |
| RGR | 0.2042 | 0.2799 | 0.2833 | 0.2325 |
| SCM | 0.2396 | 0.29 | 0.2452 | 0.2252 |
| SSR | 0.2538 | 0.3272 | 0.1991 | 0.22 |
| DSR | 0.3054 | 0.3329 | 0.1742 | 0.1875 |
| RSN | 0.4027 | 0.2362 | 0.1471 | 0.214 |
| VSN | 0.1853 | 0.3173 | 0.2728 | 0.2246 |
| LUT | 0.1997 | 0.3384 | 0.2637 | 0.1982 |
| BIO | 0.5082 | 0.088 | 0.0887 | 0.315 |
| LRF | 0.192 | 0.3119 | 0.3232 | 0.173 |
| SRF | 0.2735 | 0.3429 | 0.1905 | 0.1931 |
| BDR | 0.3265 | 0.2044 | 0.2224 | 0.2468 |
| RDR | 0.3433 | 0.0972 | 0.0937 | 0.4658 |
| SEA | 0.3625 | 0.1934 | 0.1654 | 0.2787 |
| SCI | 0.1247 | 0.3797 | 0.3344 | 0.1611 |
| LDY | 0.2833 | 0.2301 | 0.1699 | 0.3167 |
| MUL | 0.1496 | 0.0993 | 0.1345 | 0.6166 |
| POM | 0.2145 | 0.1855 | 0.3386 | 0.2614 |
| SHP | 0.5021 | 0.1957 | 0.0761 | 0.2261 |
| SNK | 0.25 | 0.25 | 0.25 | 0.25 |
| FLT | 0.0793 | 0.2154 | 0.2564 | 0.4489 |
| ODF | 0.1954 | 0.3063 | 0.2756 | 0.2227 |
| SDF | 0.1309 | 0.4605 | 0.2993 | 0.1093 |
| YTN | 0.2107 | 0.2809 | 0.306 | 0.2024 |
| BTN | 0.3786 | 0.5336 | 0.0562 | 0.0317 |
| LTN | 0.0867 | 0.3047 | 0.4285 | 0.1801 |
| OTN | 0.25 | 0.25 | 0.25 | 0.25 |
| SWD | 0.3654 | 0.2216 | 0.1881 | 0.2249 |
| WMR | 0.0515 | 0.2434 | 0.5943 | 0.1108 |
| BMR | 0.0662 | 0.2946 | 0.5315 | 0.1077 |
| BIL | 0.2209 | 0.2683 | 0.3274 | 0.1834 |
| AMB | 0.3093 | 0.2938 | 0.2472 | 0.1497 |
| JCK | 0.1049 | 0.4795 | 0.2655 | 0.1501 |
| KMK | 0.4039 | 0.031 | 0.3902 | 0.1749 |
| SMK | 0.4086 | 0.2946 | 0.1166 | 0.1802 |
| SAR | 0.0042 | 0.5258 | 0.3699 | 0.1 |
| LPL | 0.1249 | 0.3599 | 0.4028 | 0.1124 |
| DWF | 0.25 | 0.25 | 0.25 | 0.25 |
| MEN | 0.0002 | 0.3833 | 0.5244 | 0.092 |
| PIN | 0.2709 | 0.2936 | 0.2526 | 0.1829 |
| MPL | 0.2789 | 0.4624 | 0.1125 | 0.1463 |
| SPL | 0.3551 | 0.301 | 0.1431 | 0.2008 |
| TIP | 0.3479 | 0.1551 | 0.3919 | 0.1051 |
| BEN | 0.25 | 0.25 | 0.25 | 0.25 |
| LGS | 0.3661 | 0.1752 | 0.3579 | 0.1008 |
| FIL | 0.25 | 0.25 | 0.25 | 0.25 |
| SMS | 0.25 | 0.25 | 0.25 | 0.25 |
| RAY | 0.3947 | 0.3293 | 0.2436 | 0.0324 |
| BSH | 0.0414 | 0.4218 | 0.3999 | 0.1369 |
| WSH | 0.0701 | 0.141 | 0.3328 | 0.4561 |
| PSH | 0.3107 | 0.3364 | 0.1165 | 0.2364 |
| OSH | 0.2156 | 0.1519 | 0.1686 | 0.4639 |
| DBR | 0.25 | 0.25 | 0.25 | 0.25 |
| SBR | 0.25 | 0.25 | 0.25 | 0.25 |
| MAN | 0.25 | 0.25 | 0.25 | 0.25 |
| MYS | 0.25 | 0.25 | 0.25 | 0.25 |
| DOL | 0.25 | 0.25 | 0.25 | 0.25 |
| DDO | 0.25 | 0.25 | 0.25 | 0.25 |
| LOG | 0.25 | 0.25 | 0.25 | 0.25 |
| KMP | 0.25 | 0.25 | 0.25 | 0.25 |
| TUR | 0.25 | 0.25 | 0.25 | 0.25 |
| BCR | 0.1369 | 0.2992 | 0.3252 | 0.2387 |
| SCR | 0.3436 | 0.1124 | 0.003 | 0.541 |
| LOB | 0.1283 | 0.009 | 0.4946 | 0.368 |
| COR | 0.25 | 0.25 | 0.25 | 0.25 |
| CCA | 0.25 | 0.25 | 0.25 | 0.25 |
| OCT | 0.25 | 0.25 | 0.25 | 0.25 |
| SPG | 0.2086 | 0.3385 | 0.2719 | 0.1811 |
| CMB | 0.25 | 0.25 | 0.25 | 0.25 |
| INF | 0.25 | 0.25 | 0.25 | 0.25 |
| ECH | 0.25 | 0.25 | 0.25 | 0.25 |
| OYS | 0.2872 | 0.2405 | 0.1926 | 0.2798 |
| BIV | 0.2593 | 0.2247 | 0.3167 | 0.1993 |
| SES | 0.25 | 0.25 | 0.25 | 0.25 |
| EPI | 0.25 | 0.25 | 0.25 | 0.25 |
| GRS | 0.25 | 0.25 | 0.25 | 0.25 |
| ALG | 0.25 | 0.25 | 0.25 | 0.25 |
| MPB | 0.25 | 0.25 | 0.25 | 0.25 |
| LPP | 0.25 | 0.25 | 0.25 | 0.25 |
| SPP | 0.25 | 0.25 | 0.25 | 0.25 |
| DIN | 0.25 | 0.25 | 0.25 | 0.25 |
| PRO | 0.25 | 0.25 | 0.25 | 0.25 |
| JEL | 0.25 | 0.25 | 0.25 | 0.25 |
| SQU | 0.1666 | 0.2854 | 0.3481 | 0.1998 |
| LZP | 0.25 | 0.25 | 0.25 | 0.25 |
| SZP | 0.25 | 0.25 | 0.25 | 0.25 |
| PB | 0.25 | 0.25 | 0.25 | 0.25 |
| BB | 0.25 | 0.25 | 0.25 | 0.25 |
| DC | 0.25 | 0.25 | 0.25 | 0.25 |
| DL | 0.25 | 0.25 | 0.25 | 0.25 |
| DR | 0.25 | 0.25 | 0.25 | 0.25 |

Table A.8: Seasonal Distribution of Atlantis Functional Group Harvested by United States Recreational Fleets

| Group | Winter (Jan.-Mar.) | Spring (Apr.-Jun.) | Summer (Jul.-Sep.) | Fall (Oct. - Dec.) |
| :---: | :---: | :---: | :---: | :---: |
| GAG | 0.1878 | 0.2831 | 0.2292 | 0.2999 |
| RGR | 0.129 | 0.2683 | 0.3897 | 0.2131 |
| SCM | 0.0815 | 0.3257 | 0.3474 | 0.2454 |
| SSR | 0.2875 | 0.3417 | 0.1894 | 0.1814 |
| DSR | 0.3065 | 0.2758 | 0.1726 | 0.2452 |
| RSN | 0.0891 | 0.3531 | 0.3759 | 0.1819 |
| VSN | 0.0978 | 0.3585 | 0.3726 | 0.1711 |
| LUT | 0.3385 | 0.2286 | 0.2152 | 0.2177 |
| BIO | 0.2721 | 0.2862 | 0.2537 | 0.1881 |
| LRF | 0.1094 | 0.2608 | 0.3634 | 0.2664 |
| SRF | 0.105 | 0.2384 | 0.3436 | 0.3131 |
| BDR | 0.2306 | 0.2302 | 0.2246 | 0.3147 |
| RDR | 0.1071 | 0.2139 | 0.3966 | 0.2824 |
| SEA | 0.1292 | 0.2537 | 0.3788 | 0.2383 |
| SCI | 0.0546 | 0.3032 | 0.4085 | 0.2337 |
| LDY | 0.0927 | 0.3571 | 0.3625 | 0.1876 |
| MUL | 0.1629 | 0.2763 | 0.2735 | 0.2873 |
| POM | 0.1447 | 0.2374 | 0.3026 | 0.3152 |
| SHP | 0.4475 | 0.2627 | 0.0827 | 0.2071 |
| SNK | 0.1127 | 0.298 | 0.2614 | 0.3279 |
| FLT | 0.0826 | 0.3033 | 0.394 | 0.2201 |
| ODF | 0.3448 | 0.3462 | 0.1342 | 0.1748 |
| SDF | 0.0872 | 0.303 | 0.3761 | 0.2337 |
| YTN | 0.1579 | 0.2273 | 0.4593 | 0.1555 |
| BTN | 0.1111 | 0.4444 | 0.2222 | 0.2222 |
| LTN | 0.1081 | 0.3017 | 0.4262 | 0.164 |
| OTN | 0.23 | 0.27 | 0.35 | 0.15 |
| SWD | 0.2 | 0 | 0.7 | 0.1 |
| WMR | 0 | 0.25 | 0.6664 | 0.0836 |
| BMR | 0 | 0.5264 | 0.35 | 0.1236 |
| BIL | 0.2457 | 0.2038 | 0.2114 | 0.3392 |
| AMB | 0.1577 | 0.3881 | 0.2762 | 0.1779 |
| JCK | 0.1938 | 0.2674 | 0.2741 | 0.2647 |
| KMK | 0.1477 | 0.2845 | 0.3661 | 0.2017 |
| SMK | 0.1108 | 0.328 | 0.3567 | 0.2044 |
| SAR | 0.0711 | 0.3268 | 0.3104 | 0.2917 |
| LPL | 0.2553 | 0.3537 | 0.2119 | 0.1791 |
| DWF | 0.0385 | 0.2645 | 0.5021 | 0.1949 |
| MEN | 0.0193 | 0.3437 | 0.4506 | 0.1864 |
| PIN | 0.0916 | 0.2999 | 0.3846 | 0.2239 |
| MPL | 0.0602 | 0.4461 | 0.2833 | 0.2105 |
| SPL | 0.213 | 0.2319 | 0.2276 | 0.3275 |
| TIP | 0.0405 | 0.3909 | 0.4526 | 0.1159 |
| BEN | 0.5 | 0.5 | 0 | 0 |
| LGS | 0.1308 | 0.2192 | 0.4067 | 0.2433 |
| FIL | 0.25 | 0.25 | 0.25 | 0.25 |
| SMS | 0.875 | 0.125 | 0 | 0 |
| RAY | 0.1577 | 0.2999 | 0.2839 | 0.2585 |
| BSH | 0.25 | 0.25 | 0.25 | 0.25 |
| WSH | 0.25 | 0.25 | 0.25 | 0.25 |
| PSH | 0.25 | 0.25 | 0.25 | 0.25 |
| OSH | 0.25 | 0.25 | 0.25 | 0.25 |
| DBR | 0.25 | 0.25 | 0.25 | 0.25 |
| SBR | 0.25 | 0.25 | 0.25 | 0.25 |
| MAN | 0.25 | 0.25 | 0.25 | 0.25 |
| MYS | 0.25 | 0.25 | 0.25 | 0.25 |
| DOL | 0.25 | 0.25 | 0.25 | 0.25 |
| DDO | 0.25 | 0.25 | 0.25 | 0.25 |
| LOG | 0.25 | 0.25 | 0.25 | 0.25 |
| KMP | 0.25 | 0.25 | 0.25 | 0.25 |
| TUR | 0.25 | 0.25 | 0.25 | 0.25 |
| BCR | 0.25 | 0.25 | 0.25 | 0.25 |
| SCR | 0.25 | 0.25 | 0.25 | 0.25 |
| LOB | 0.1283 | 0.009 | 0.4946 | 0.368 |
| COR | 0.25 | 0.25 | 0.25 | 0.25 |
| CCA | 0.25 | 0.25 | 0.25 | 0.25 |
| OCT | 0.25 | 0.25 | 0.25 | 0.25 |
| SPG | 0.25 | 0.25 | 0.25 | 0.25 |
| CMB | 0.25 | 0.25 | 0.25 | 0.25 |
| INF | 0.25 | 0.25 | 0.25 | 0.25 |
| ECH | 0.25 | 0.25 | 0.25 | 0.25 |
| OYS | 0.25 | 0.25 | 0.25 | 0.25 |
| BIV | 0.25 | 0.25 | 0.25 | 0.25 |
| SES | 0.25 | 0.25 | 0.25 | 0.25 |
| EPI | 0.25 | 0.25 | 0.25 | 0.25 |
| GRS | 0.25 | 0.25 | 0.25 | 0.25 |
| ALG | 0.25 | 0.25 | 0.25 | 0.25 |
| MPB | 0.25 | 0.25 | 0.25 | 0.25 |
| LPP | 0.25 | 0.25 | 0.25 | 0.25 |
| SPP | 0.25 | 0.25 | 0.25 | 0.25 |
| DIN | 0.25 | 0.25 | 0.25 | 0.25 |
| PRO | 0.25 | 0.25 | 0.25 | 0.25 |
| JEL | 0.25 | 0.25 | 0.25 | 0.25 |
| SQU | 0.25 | 0.25 | 0.25 | 0.25 |
| LZP | 0.25 | 0.25 | 0.25 | 0.25 |
| SZP | 0.25 | 0.25 | 0.25 | 0.25 |
| PB | 0.25 | 0.25 | 0.25 | 0.25 |
| BB | 0.25 | 0.25 | 0.25 | 0.25 |
| DC | 0.25 | 0.25 | 0.25 | 0.25 |
| DL | 0.25 | 0.25 | 0.25 | 0.25 |
| DR | 0.25 | 0.25 | 0.25 | 0.25 |

Table A.9: Seasonal Distribution of Atlantis Functional Group Harvested by Mexican Commercial Fleets

| Group | Winter (Jan.-Mar.) | Spring (Apr.-Jun.) | Summer (Jul.-Sep.) | Fall (Oct. - Dec.) |
| :---: | :---: | :---: | :---: | :---: |
| GAG | 0.25 | 0.25 | 0.25 | 0.25 |
| RGR | 0.2416 | 0.3179 | 0.2554 | 0.1852 |
| SCM | 0.25 | 0.25 | 0.25 | 0.25 |
| SSR | 0.2503 | 0.304 | 0.2499 | 0.1958 |
| DSR | 0.2367 | 0.3119 | 0.2687 | 0.1827 |
| RSN | 0.2943 | 0.2426 | 0.2109 | 0.2522 |
| VSN | 0.2292 | 0.2505 | 0.2831 | 0.2373 |
| LUT | 0.3093 | 0.241 | 0.2438 | 0.2059 |
| BIO | 0.25 | 0.25 | 0.25 | 0.25 |
| LRF | 0.2603 | 0.2477 | 0.2665 | 0.2255 |
| SRF | 0.8907 | 0.0782 | 0.0292 | 0.0019 |
| BDR | 0.25 | 0.25 | 0.25 | 0.25 |
| RDR | 0.396 | 0.2285 | 0.1357 | 0.2398 |
| SEA | 0.3624 | 0.2137 | 0.1756 | 0.2484 |
| SCI | 0.3231 | 0.2239 | 0.215 | 0.2379 |
| LDY | 0.25 | 0.25 | 0.25 | 0.25 |
| MUL | 0.2665 | 0.1746 | 0.2319 | 0.3269 |
| POM | 0.3561 | 0.2697 | 0.1863 | 0.188 |
| SHP | 0.25 | 0.25 | 0.25 | 0.25 |
| SNK | 0.2645 | 0.258 | 0.2431 | 0.2344 |
| FLT | 0.2384 | 0.1983 | 0.1427 | 0.4206 |
| ODF | 0.2306 | 0.2558 | 0.2648 | 0.2488 |
| SDF | 0.3932 | 0.2937 | 0.1585 | 0.1546 |
| YTN | 0.1904 | 0.275 | 0.2991 | 0.2356 |
| BTN | 0.7164 | 0.2027 | 0.0032 | 0.0777 |
| LTN | 0.2825 | 0.2657 | 0.2301 | 0.2217 |
| OTN | 0.1207 | 0.1706 | 0.3159 | 0.3929 |
| SWD | 0.2286 | 0.2019 | 0.2351 | 0.3344 |
| WMR | 0.2353 | 0.1759 | 0.2716 | 0.3172 |
| BMR | 0.1843 | 0.2385 | 0.3258 | 0.2514 |
| BIL | 0.0331 | 0.6081 | 0.2894 | 0.0694 |
| AMB | 0.25 | 0.25 | 0.25 | 0.25 |
| JCK | 0.2794 | 0.2916 | 0.2069 | 0.222 |
| KMK | 0.3421 | 0.2728 | 0.2133 | 0.1718 |
| SMK | 0.3753 | 0.1318 | 0.1321 | 0.3608 |
| SAR | 0.2559 | 0.2359 | 0.3233 | 0.1849 |
| LPL | 0.3519 | 0.1992 | 0.1737 | 0.2752 |
| DWF | 0.25 | 0.25 | 0.25 | 0.25 |
| MEN | 0.25 | 0.25 | 0.25 | 0.25 |
| PIN | 0.25 | 0.25 | 0.25 | 0.25 |
| MPL | 0.2948 | 0.2832 | 0.2194 | 0.2026 |
| SPL | 0.685 | 0.0956 | 0.042 | 0.1775 |
| TIP | 0.2752 | 0.2823 | 0.1807 | 0.2618 |
| BEN | 0.2752 | 0.2823 | 0.1807 | 0.2618 |
| LGS | 0.2924 | 0.2735 | 0.2054 | 0.2287 |
| FIL | 0.25 | 0.25 | 0.25 | 0.25 |
| SMS | 0.2752 | 0.2823 | 0.1807 | 0.2618 |
| RAY | 0.2855 | 0.2503 | 0.2382 | 0.226 |
| BSH | 0.1893 | 0.2428 | 0.264 | 0.304 |
| WSH | 0.1893 | 0.2428 | 0.264 | 0.304 |
| PSH | 0.1893 | 0.2428 | 0.264 | 0.304 |
| OSH | 0.1893 | 0.2428 | 0.264 | 0.304 |
| DBR | 0.25 | 0.25 | 0.25 | 0.25 |
| SBR | 0.25 | 0.25 | 0.25 | 0.25 |
| MAN | 0.25 | 0.25 | 0.25 | 0.25 |
| MYS | 0.25 | 0.25 | 0.25 | 0.25 |
| DOL | 0.25 | 0.25 | 0.25 | 0.25 |
| DDO | 0.25 | 0.25 | 0.25 | 0.25 |
| LOG | 0.25 | 0.25 | 0.25 | 0.25 |
| KMP | 0.25 | 0.25 | 0.25 | 0.25 |
| TUR | 0.25 | 0.25 | 0.25 | 0.25 |
| BCR | 0.2568 | 0.2567 | 0.237 | 0.2496 |
| SCR | 0.2568 | 0.2567 | 0.237 | 0.2496 |
| LOB | 0.1958 | 0.2816 | 0.3932 | 0.1294 |
| COR | 0.25 | 0.25 | 0.25 | 0.25 |
| CCA | 0.25 | 0.25 | 0.25 | 0.25 |
| OCT | 0.25 | 0.25 | 0.25 | 0.25 |
| SPG | 0.25 | 0.25 | 0.25 | 0.25 |
| CMB | 0.09 | 0.5917 | 0.302 | 0.0163 |
| INF | 0.25 | 0.25 | 0.25 | 0.25 |
| ECH | 0.25 | 0.25 | 0.25 | 0.25 |
| OYS | 0.1985 | 0.2176 | 0.2813 | 0.3026 |
| BIV | 0.2522 | 0.238 | 0.2324 | 0.2774 |
| SES | 0.25 | 0.25 | 0.25 | 0.25 |
| EPI | 0.25 | 0.25 | 0.25 | 0.25 |
| GRS | 0.25 | 0.25 | 0.25 | 0.25 |
| ALG | 0.25 | 0.25 | 0.25 | 0.25 |
| MPB | 0.25 | 0.25 | 0.25 | 0.25 |
| LPP | 0.25 | 0.25 | 0.25 | 0.25 |
| SPP | 0.25 | 0.25 | 0.25 | 0.25 |
| DIN | 0.25 | 0.25 | 0.25 | 0.25 |
| PRO | 0.25 | 0.25 | 0.25 | 0.25 |
| JEL | 0.25 | 0.25 | 0.25 | 0.25 |
| SQU | 0.0281 | 0.0264 | 0.3757 | 0.5698 |
| LZP | 0.25 | 0.25 | 0.25 | 0.25 |
| SZP | 0.25 | 0.25 | 0.25 | 0.25 |
| PB | 0.25 | 0.25 | 0.25 | 0.25 |
| BB | 0.25 | 0.25 | 0.25 | 0.25 |
| DC | 0.25 | 0.25 | 0.25 | 0.25 |
| DL | 0.25 | 0.25 | 0.25 | 0.25 |
| DR | 0.25 | 0.25 | 0.25 | 0.25 |

Table A.10: Proportion of Functional Group Landings Across U.S. Commercial Atlantis Fleets.

| Group | GillnetEst | TwlShpEst | OytEst | PotCrbEst | TwlShpShf | PotCrbShf | PotLbtShf | HLReefShf | LLReefShf |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GAG |  |  |  |  |  |  |  | 0.672 | 0.244 |
| RGR |  |  |  |  |  | 0 |  |  | 0.991 |
| SCM |  |  |  |  |  |  |  | 0.48 | 0.507 |
| SSR |  |  |  |  |  | 0.01 | 0.001 | 0.525 | 0.281 |
| DSR |  |  |  | 0.013 |  | 0.373 |  | 0.121 | 0.493 |
| RSN |  |  |  |  |  |  |  | 0.961 | 0.033 |
| VSN |  |  |  |  |  |  |  | 0.997 | 0.002 |
| LUT | 0 |  |  |  |  | 0.003 | 0.001 | 0.919 | 0.065 |
| BIO |  |  |  |  |  |  |  |  |  |
| LRF | 0.002 | 0.015 |  |  | 0.004 | 0.012 | 0.007 | 0.326 | 0.042 |
| SRF | 0.001 |  |  |  |  |  |  | 0.282 | 0.692 |
| BDR | 0.025 | 0.019 |  | 0.015 | 0.066 |  |  | 0.059 | 0.791 |
| RDR |  |  |  |  |  |  |  | 1 |  |
| SEA | 0.095 | 0.003 |  |  | 0.091 |  |  | 0.783 |  |
| SCI | 0.124 | 0.068 |  | 0.001 | 0.284 |  |  | 0.371 |  |
| LDY | 0.688 |  |  |  |  |  |  | 0.042 |  |
| MUL | 0.177 |  |  | 0 | 0 |  |  | 0 |  |
| POM | 0.234 |  |  |  |  |  |  | 0.554 |  |
| SHP | 0.078 | 0.134 |  | 0.002 | 0.176 |  |  | 0.405 | 0.037 |
| SNK |  |  |  |  |  |  |  |  |  |
| FLT | 0.203 | 0.254 |  | 0.017 | 0.069 | 0.016 |  | 0.041 | 0.039 |
| ODF | 0.005 |  |  | 0.007 |  | 0.241 | 0.001 | 0.711 | 0.031 |
| SDF |  | 1 |  |  |  |  |  |  |  |
| YTN |  |  |  |  |  |  |  | 0.012 |  |
| BTN |  |  |  |  |  |  |  |  |  |
| LTN | 0.087 |  |  |  |  |  |  | 0.913 |  |
| SWD |  |  |  |  |  |  |  | 0.035 |  |
| AMB |  |  |  |  |  |  |  | 0.961 | 0.014 |
| JCK | 0.048 |  |  | 0 |  | 0.012 |  | 0.645 | 0.015 |
| KMK |  |  |  |  |  |  |  | 0.792 |  |
| SMK | 0.81 | 0 |  |  |  |  |  | 0.101 |  |
| SAR |  |  |  |  |  |  |  | 0.088 |  |
| LPL | 0.19 | 0.036 |  |  |  |  |  | 0.398 | 0.015 |
| DWF |  |  |  |  |  |  |  |  |  |
| MEN | 0 |  |  |  |  |  |  | 0 |  |
| PIN |  |  |  |  |  | 0.588 |  | 0.27 |  |
| MPL |  |  |  | 0.001 |  |  |  | 0.311 |  |
| SPL | 0 |  |  |  |  |  |  |  |  |
| TIP |  |  |  |  |  |  |  | 0.919 |  |
| LGS | 0.34 | 0.001 |  |  | 0 |  |  | 0.258 | 0.037 |
| RAY | 1 |  |  |  |  |  |  |  |  |
| BSH |  | 0.193 |  |  | 0.807 |  |  |  |  |
| WSH |  | 0.298 |  |  | 0.701 |  |  |  |  |
| PSH |  | 0.002 |  |  | 0.998 |  |  |  |  |
| OSH |  | 0.311 |  |  | 0.622 | 0 |  |  |  |
| BCR |  | 0.002 |  | 0.997 | 0 | 0 |  | 0 | 0 |
| SCR |  |  |  | 0.984 |  |  | 0.016 |  |  |
| LOB |  |  |  | 0 | 0 |  | 0.966 |  |  |
| SPG |  |  |  |  |  |  |  |  |  |
| OYS |  |  | 0.967 |  |  |  |  |  |  |
| BIV | 0.003 | 0.003 | 0.002 | 0.001 | 0.937 | 0.037 | 0.004 | 0 | 0 |
| SQU |  | 0.266 |  | 0.115 | 0.429 | 0.14 |  | 0.049 |  |

Table A.10: Continued

| Group | SeineMenShf | LLShkShf | LLPelgc | RoyalRed | OtherUS |
| :---: | :---: | :---: | :---: | :---: | :---: |
| GAG |  |  |  |  | 0.084 |
| RGR |  |  |  |  | 0.009 |
| SCM |  |  |  |  | 0.013 |
| SSR |  |  |  |  | 0.183 |
| DSR |  |  |  |  | 0 |
| RSN |  |  |  |  | 0.007 |
| VSN |  |  |  |  | 0.001 |
| LUT |  |  |  |  | 0.013 |
| BIO |  |  |  |  |  |
| LRF |  |  |  |  | 0.591 |
| SRF |  |  | 0.003 |  | 0.021 |
| BDR |  |  | 0.02 |  | 0.004 |
| RDR 0.028 |  |  |  |  |  |
| SEA |  |  |  |  | 0.028 |
| SCI |  |  |  |  | 0.151 |
| LDY |  |  |  |  | 0.27 |
| MUL | 0.006 |  |  |  | 0.816 |
| POM |  |  |  |  | 0.212 |
| SHP | 0 |  |  |  | 0.168 |
| SNK 0.018 |  |  |  |  |  |
| FLT |  |  | 0.018 |  | 0.343 |
| ODF |  |  |  |  | 0.005 |
| SDF |  |  |  |  |  |
| YTN |  |  | 0.988 |  |  |
| BTN |  |  | 1 |  |  |
| LTN |  |  |  |  |  |
| SWD |  |  | 0.965 |  |  |
| AMB |  |  |  |  | 0.025 |
| JCK |  |  |  |  | 0.28 |
| KMK |  |  | 0.208 |  |  |
| SMK | 0.002 |  | 0.061 |  | 0.027 |
| SAR |  |  |  |  | 0.912 |
| LPL | 0.016 |  | 0.338 |  | 0.008 |
| DWF |  |  |  |  |  |
| MEN | 1 |  |  |  | 0 |
| PIN |  |  |  |  | 0.142 |
| MPL |  |  | 0.003 |  | 0.685 |
| SPL |  |  |  |  | 1 |
| TIP |  | 0.081 |  |  |  |
| LGS | 0 | 0.29 | 0.072 |  | 0.002 |
| RAY |  |  |  |  |  |
| BSH |  |  |  |  | 0 |
| WSH |  |  |  |  | 0 |
| PSH |  |  |  |  |  |
| OSH |  |  |  | 0.067 | 0 |
| BCR |  |  |  |  |  |
| SCR |  |  |  |  |  |
| LOB |  |  |  |  | 0.033 |
| SPG |  |  |  |  | 1 |
| OYS |  |  |  |  | 0.033 |
| BIV |  |  | 0 |  | 0.013 |
| SQU |  |  |  |  | 0.001 |

Table A.11: Distribution of Mexican Commercial Landings Amongst Atlantis Fleets.

| Group | TwlShpMX | LLReefMX | LLShkMX | GillnetMackMX | OctpsMX | MixedMX |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| RGR |  | 0.500 |  |  |  | 0.500 |
| SSR |  | 0.500 |  |  |  | 0.500 |
| DSR |  | 1.000 |  |  |  |  |
| RSN |  | 0.500 |  |  |  | 0.500 |
| VSN |  | 0.500 |  |  |  | 0.500 |
| LUT |  | 0.500 |  |  |  | 0.500 |
| LRF |  |  |  |  |  | 1.000 |
| SRF |  |  |  |  |  | 1.000 |
| RDR |  |  |  |  |  | 1.000 |
| SEA |  |  |  |  |  | 1.000 |
| SCI |  |  |  |  |  | 1.000 |
| LDY |  |  |  |  |  | 1.000 |
| MUL |  |  |  | 0.500 |  | 0.500 |
| POM |  |  |  |  |  | 1.000 |
| SHP | 0.333 |  |  | 0.333 |  | 0.330 |
| SNK |  |  |  |  |  | 1.000 |
| FLT | 0.333 |  |  | 0.333 |  | 0.333 |
| ODF |  |  |  |  |  | 1.000 |
| SDF |  |  |  |  |  | 1.000 |
| YTN |  |  | 0.500 |  |  | 0.500 |
| BTN |  |  | 0.500 |  |  | 0.500 |
| LTN |  |  | 0.500 |  |  | 0.500 |
| OTN |  |  | 0.500 |  |  | 0.500 |
| SWD |  |  | 0.500 |  |  | 0.500 |
| WMR |  |  | 0.500 |  |  | 0.500 |
| BMR |  |  | 0.500 |  |  | 0.500 |
| BIL |  |  | 0.500 |  |  | 0.500 |
| AMB |  |  | 0.500 |  |  | 0.500 |
| JCK |  |  | 0.500 |  |  | 0.500 |
| KMK |  |  | 0.500 | 0.500 |  |  |
| SMK |  |  | 0.500 | 0.500 |  |  |
| SAR |  |  |  |  |  | 1.000 |
| LPL |  |  |  | 1.000 |  |  |
| PIN |  |  |  |  |  | 1.000 |
| MPL |  |  |  |  |  | 1.000 |
| SPL |  |  |  |  |  | 1.000 |
| TIP |  |  | 1.000 |  |  |  |
| BEN | 0.333 |  | 0.333 |  |  | 0.333 |
| LGS |  |  | 0.500 |  |  | 0.500 |
| FIL |  |  |  |  |  | 1.000 |
| SMS | 0.250 |  | 0.250 | 0.250 |  | 0.250 |
| RAY | 0.333 |  | 0.333 |  |  | 0.333 |
| BSH | 1.000 |  |  |  |  |  |
| WSH | 0.500 |  |  |  |  | 0.500 |
| PSH | 1.000 |  |  |  |  |  |
| OSH | 0.500 |  |  |  |  | 0.500 |
| BCR |  |  |  |  |  | 1.000 |
| SCR |  |  |  |  |  | 1.000 |
| LOB |  |  |  |  |  | 1.000 |
| OCT |  |  |  |  | 1.000 |  |
| CMB | 0.500 |  |  |  |  | 0.500 |
| OYS |  |  |  |  |  | 1.000 |
| BIV |  |  |  |  |  | 1.000 |
| SQU | 0.500 |  |  |  |  | 0.500 |

## A. 6 Additional Results for Section 2.3.2



Figure A.17: Species Composition Time Series of United States Commercial Landings by Overfished Status. Legend shows only the eight most common species.


Figure A.18: Species Composition Time Series of United States Commercial Landings by Overfished Status. Legend shows only the eight most common species.


Figure A.19: Species Composition Time Series of United States Species-Specific Total Commercial Data (excluding menhaden). Data used to create these plots were also used to calculate U.S. commercial landings seasonal indicators. Species compositions are shown for winter, Jan. - Mar. (a), spring, Apr. - Jun. (b), summer, Jul. - Sep. (c), and fall, Oct. - Dec. (d). Legend shows only the seven most common species.


Figure A.20: Species Composition Time Series of Seasonal United States Species-Specific Recreational Landings. Data used to create these plots were also used to calculate U.S. recreational landings seasonal indicators. Species compositions are shown for wave 1, Jan. - Feb. (a), wave 2, Mar. - Apr. (b), wave 3, May. - Jun. (c), wave 4, Jul. - Aug. (d), wave 5, Sep. - Oct. (e), and wave 6, Nov. - Dec. (f). Legend shows only the seven most common species.


Figure A.21: Species Composition Time Series of United States Species-Specific Commercial Landings by State. Legend shows only the seven most common species.


Figure A.22: Species Composition Time Series of United States Species-Specific Recreational Landings by State. Legend shows only the seven most common species.

## A.6.1 Landings Mean Trophic Level Sensitivity Analysis

For commercial and recreational data, the trends in landings mean trophic level from both species-specific data and functional group-specific data are within the trends computed for the trophic level sensitivity analysis (Figure A.23a, A.23b). The sensitivity analysis suggests that the computation of landings mean trophic level is sensitive to trophic level. Thus, the aggregation into functional groups, and the averaging of species-specific trophic levels, could be causing the slight difference between functional group-specific and species-specific landings mean trophic level trends. The trends computed for the trophic level sensitivity analysis are wider for recreational data than commercial data because the trophic level standard errors considered here were only for some fish groups, and commercial landings have considerable amounts of invertebrate landings.

The computation of landings mean trophic level appears to be particularly sensitive to trophic level, thus the aggregation of data into functional groups, and the averaging of species-specific trophic levels, is likely having some impact on the value computed for landings mean trophic level. However, the difference observed here is small enough to be considered negligible.

The results from the sensitivity analysis indicating the computation of landings mean trophic level to be particularly sensitive to trophic level is not surprising considering that an organism's trophic level is not a constant value. Since an organism's trophic level is governed by the prey consumed, the trophic level of any one species can vary spatially, temporally, and as the organism ages (e.g., Jennings et al., 1997; Hussey et al., 2011; Yurkowski et al., 2016). A full sensitivity analysis on landings
mean trophic level would be more informative for EBFM concerning the indicators robustness at various levels of species aggregation. Such an analysis should consider trophic level standard errors from laboratory studies, but if this information is not found in the literature than estimating trophic level standard errors may be necessary (e.g., Pauly and Christensen, 1995).


Figure A.23: Panels a and b show trends from the annual summaries of species-specific data (solid line), functional group-specific data (dashed line), and functional group-specific data with trophic level of multi-species functional groups being the average plus/minus the standard error across the multi-species functional groups (dotted lines).

## APPENDIX B

## Detailed Methodology and Additional Results for Chapter 3

## B. 1 Catch Data for Model Fitting

The Southeast Fisheries Science Center (SEFSC) conducts an annual bottom longline survey within the Gulf of Mexico and western Atlantic Ocean. Commercialtype longline gear is utilized so analyses based of this dataset can be related back to the commercial longline fisheries within the Gulf of Mexico. The mainline is suspended by two radar reflector high-flyers attached at each end, and held stationary with 5 kg weights located at the start, middle, and end of the mainline. The mainline consists of 100 gangions; each one is approximately 3.6 m in length and 18.3 m apart from one another. Gangions consist of 3 mm diameter monofilament line with \#15 circle hook. Each hook is baited with atlantic mackerel (Scomber scombrus). Longline gear soaks for one hour after the set is complete. During the haul in organisms retained are processed immediately. Processing includes, but is not limited to, identifying species and recording length, weight, and sex. While smaller organisms (e.g. Rhizoprionodon terraenovae) are handled by hand, larger organisms (e.g. Galeocerdo cuvier) are held in a landing sling to facilitate processing. Organisms of concern (e.g., Sphyrna lewini)
are also tagged. This survey operates along the continental shelf in depths between 9-366 meters, so Gulf-wide catch of pelagic species can not be represented by this dataset alone.

The Pelagic Observer Program, also managed by the SEFSC, provides longline catch and effort data within the Gulf of Mexico between 200 m depth and the exclusive economic zone (EEZ). Since 1992, observers have monitored the mobile U.S. pelagic longline fleet operating in the western Atlantic Ocean. Pelagic longline gear consists of a mainline suspended mid-depth by a series of high-flyers. Longline sets can extend from 10 to 40 miles, fishing 200-1000 baited hooks spaced approximately 100 meters apart. Highly migratory species Xiphias gladius, Thunnus albacares, or Thunnus obsess are often the primary target. Harvesting methodology, area, and season changes based on the targeted species. For instance, pelagic longliners targeting X. gladius set hooks during the night while those targeting tunas set hooks during the day. Information retained by observers includes species, date, time, and location.

## B. 2 Supplementing Environmental Variables into Catch Datasets

Sea surface temperature is one of the few environmental variables monitored in both longline catch datasets. It is measured at multiple points during the setting and hauling of longline gear, so the mean of the reported measurements was used as final estimate of sea surface temperature for each catch record. Approximately $25 \%$ of the bottom longline records and $5 \%$ of the pelagic longline records were missing estimates of sea surface temperature. To retain these records for statistical analysis, estimates of sea surface temperature were generated by using Interpolate PO.DAAC MODIS

L3 SST at Points tool from the Marine Geospatial Ecology Tools (MGET) toolbox in ArcGIS, which extracts estimates of sea surface temperature from the Moderate Resolution Imaging Spectroradiometer (MODIS) dataset. The tool was set to draw daytime sea surface temperature values from the aqua satellite using a linear interpolation method. Different combinations of spatial resolution ( 4 km grid or 9 km grid) and temporal resolution (daily, 8 day, monthly) were used because a single combination could not extract estimates of sea surface temperature for all of the necessary catch records. This is likely due to the patchiness of MODIS data, due to cloud cover. Combinations with fine scale spatial and temporal resolutions were attempted first (4km, daily), followed by combinations with reduced temporal resolution. Then, the combination with reduced spatial resolution and fine scale temporal resolution (9km, daily). Again, this was followed by combinations with reduced temporal resolution. If a catch record was still missing an estimate of sea surface temperature, then a value was drawn from another catch record in the corresponding longline catch dataset. The chosen catch record was one with a similar harvest date and location as the record missing sea surface temperature.

Altimetry data was collected from the Archiving, Validation and Interpretation of Satellite Oceanographic (AVISO) dataset. AVISO gathers raw data from various satellite sources which are individually processed into estimates of altimetry then merged as described by Ducet et al. (2000). The resulting altimetry data are provided in 7 -day increments, implying that measurements may have been taken up to 3 days before or after the date displayed in the dataset. Data are mapped to an equal angle grid 0.25 degrees latitude by 0.25 degrees longitude. Estimates of altimetry for catch records in the longline catch datasets were calculated as the mean of the four AVISO
data points nearest to the catch location and corresponded to the date the catch occurred.

The routine for calculating an estimate of the minimum distance between a catch event and a frontal edge is dependent on work by Cayula and Cornillon (1992). Cayula and Cornillon presented an algorithm for detecting fronts in satellite-derived sea surface temperature fields. After the initial processing of the data input, there are three different stages focusing on the detection and removal of clouds (since they can cause erroneous edge detection). Next, the algorithm detects and verifies an edge (i.e., front) at the window level using a histogram analysis and cohesion algorithm, respectively. Then, the front is detected and verified at a local level using contour following. Lastly, the fronts are extracted. The MGET toolbox contains a tool that uses the algorithm presented by Cayula and Cornillon (1992) for the detection and extraction of fronts, call the Cayula-Cornillon Fronts in ArcGIS Raster MGET tool. Preliminary work executing this tool with sea surface temperature fields produced patchy and fragmented front profiles (Figure B.1a). Sea surface temperature fields collected from the Gulf of Mexico can have weak gradients (Legeckis, 1978) which make it difficult for the Cayula and Cornillon algorithm to detect a front. The dynamics of the Gulf of Mexico are mostly driven by the physical oceanography, which relates directly to altimetry. Executing the Cayula-Cornillon Fronts in ArcGIS Raster MGET tool with altimetry profiles produces smoother front profiles (Figure B.1b). Altimetry needed to be magnified to be on a similar scale as sea surface temperature, and a factor of 1000 produced the cleanest images.

## B. 3 Other Important Model Descriptors

Preliminary investigations selecting model descriptors identified variables capable of improving the fit of some models. Table B. 1 highlights these results. Variables were used individually to fit generalized additive models with the bottom longline survey data. Some of these variables were not used as model descriptors primarily due to the lack of data available to make Gulf-wide data fishnets for model predictions.

## B.3.1 Beam Transmission

Preliminary results showed beam transmission to be a statistically important descriptor for shark-based functional groups. Beam transmission (\%), a measurement of the penetration of light through the water column, may influence the local density of an organism (e.g., fish residing in murky waters to evade predators), or it may relate to catchability (e.g., fish are more/less likely to strike a baited hook in waters with more/less light). However, beam transmission depends on many dynamic environmental processes (e.g., cloud cover, sediment, pollution, etc), thus developing seasonal, Gulf-wide estimates of beam transmission for model predictions would be inappropriate. However, CPUE modeling over a smaller temporal range should consider beam transmission as a descriptor.

## B.3.2 Density

Sea bottom density was a statistically important descriptor for shark-based functional groups, however a dataset containing estimates of sea bottom density across the Gulf could not be found. Considering the amount of deviance explained, it could prove to be useful to include this variable into future modeling efforts. Especially if
at a smaller spatial and temporal scale allowing for the possibility of collecting data for model predictions.

## B.3.3 Latitude and Longitude

The incorporation of latitude and longitude as model descriptors helped fitted models explain more deviance. Latitude and longitude were not considered as model descriptors because preliminary predictions showed an obvious bias in the regions outside of the spatial range of the catch datasets (i.e., catch rates showed a unnaturally strong north-south gradient). If predictions efforts were restricted to the northern Gulf of Mexico than latitude and longitude should be considered as potential model descriptors to improve model fits. Geospatial models could also be considered.

## B.3.4 Daytime

Daytime, a binomial factor indicating day or night, did not explain much model deviance when fitting bottom longline survey data. However, daytime would likely impact models fit with pelagic longline observer data since hooks are set differently during the day versus night due to the change in target organism. To see how daytime may influence model predictions, models presented in the chapter were re-fitted with daytime as a descriptor (spline basis dimensions were set to three and not adjusted, thus the prediction profiles without daytime in the fitted model were similar to the night time predictions). These fitted models were than predicted across the same fishnet grids, once with daytime identified as day and again with daytime identified as night (day and night measurements of other model descriptors are not available). The coastal models (fit with bottom longline data) did not show any major differences
between day and night prediction profiles, however some functional group with pelagic models had noticeable differences between day and night prediction profiles (examples in Figure B.3). The discussion in the chapter on large sharks is not significantly impacted by not including daytime into models, but some functional groups should have daytime incorporated into future modeling efforts (e.g., swordfish, and white marlin).

## B. 4 Selecting the Error Structure of Models fitting Zero-Truncated Data

Although the longline datasets record catch as counts (which is commonly modeled with the poisson distribution) catch rates (CPUE) in this study are continuous, which are commonly modeled with the log-normal and gamma distributions. First, we wanted to assess the validity for assuming a log-normal or gamma error structure (Maunder and Punt, 2004). Dong and Restrepo (1996) and Punt et al. (2000) discussed evaluating assumed error structure by comparing catch rate average and variance. A relationship where the variance in catch rate is proportional to the square of the average catch rate suggests the log-normal or gamma distribution. The functional group-specific average catch rate and variance in catch rate for each unique date are shown in Figure B.2, and functional group-specific catch data support the use of either the log-normal or gamma distribution for the CPUE data error structure.

Myers and Pepin (1990) found lognormal-based estimators to be very sensitive to violations of model assumptions, which can lead to biases as well as reduced efficiency, and they encouraged assuming a log-normal distribution only in situations when repeated samples from the same population consistently showed the distribution to be
log-normal. If a random variable follows a log-normal distribution, than the natural $\log$ of the random variable will follow a normal distribution. Log-transforming functional group-specific catch rates for both longline datasets (Figure B.4) fails to normalize catch rates for a majority of functional groups. Thus, the data are not lognormal. Other studies have achieved comparable, if not improved, model fits using the gamma distribution rather than the lognormal (Punt et al., 2000; Dick, 2004; Ortiz and Arocha, 2004).

The statistical software $R$ has three link functions associated to the gamma distribution:identity, inverse, and log. Functional group-specific catch profiles (for both the bottom longline survey and the pelagic longline observer data) were used to fit GAMs for each of the three link functions. The generic setup of the statistical model was as follows:

$$
\begin{equation*}
\eta_{Z}=s(S S H, k=3)+s(W D, k=3)+f(y)+s(S S T, k=3)+s(M D F, k=3) \tag{B.1}
\end{equation*}
$$

where $\eta_{Z}$ is the abundance index, $S S H$ is altimetry [ m$], W D$ is bottom depth [ m ], $y$ is year, $S S T$ is sea surface temperature $\left[{ }^{\circ} \mathrm{C}\right], M D F$ is minimum distance from a front $[\mathrm{m}], s()$ indicates a smooth function, $k$ indicates a smoother's basis dimension, and $f()$ indicates a factor. All basis dimensions were set to three, the default value. Table B. 2 displays the GCV and deviance explained for all of the model fits. Often, the best fit corresponds to models that used the inverse link function. This makes sense since the inverse link corresponds to the zero inflated nature of the cate rate data. Thus, this study assumes a gamma error structure with inverse link function for GAMs fitted with zero-truncated data.

## B. 5 Verification of Routine Setting Smoother Basis Dimension

Fit statistics for models fitted with training datasets for cross validation were recorded before and after to application of the basis dimension setting routine (Table B.3). Basis dimensions for smoothers in binomial data models were often unchanged after the application of the basis dimension setting routine, thus fit statistics didn't improve. However, basis dimensions for smoothers in zero-truncated data models were often increased thus improving model fits.

## B. 6 Forward Model Selection

During the forward model selection process for determining the order of model descriptors, a correlation analysis was conducted to determine if the environmental variable selected to be a model descriptor was highly correlated with selected model descriptors. A variable highly correlated with model descriptors (i.e., producing a correlation coefficient greater than 0.80 ) was not used as a model descriptor. Correlation coefficients for environmental variables considered for models fit with pelagic longline observer data (i.e., year, season, sea surface temperature, altimetry, and minimum distance from a front) are summarized in Table B.4. None of the variables were highly correlated, thus all of the considered variables were used as model descriptors for both parts of the delta framework.

Models fit with bottom longline survey data had a variety of environmental variables applicable as model descriptors, making overparameterization a concern, so the forward model selection process stopped (i.e., model descriptors were no longer ad-
ded) once the deviance explained from model fits were not improved by more than $5 \%$. The forward model selection process is presented for large pelagic fish (Table B.5), large sharks (Table B.6), skates and rays (Table B.7), and blacktip sharks (Table B.8). These tables also display correlation coefficients between environmental variables and model descriptors. If a model gains the most improvement from an environmental variable that is highly correlated to current model descriptors, than the variable providing the next best fit is considered for incorporation as a model descriptor instead. For example, consider the forward model selection results for the large pelagic fish zero-truncated data model (Table B.5). During the third iteration it is apparent that adding sea surface oxygen saturation to the model $d o+s s h$ produced the most improved model ( $57.6 \%$ to $64.9 \%$ deviance explained). Although sea surface oxygen saturation and altimetry are not highly correlated ( 0.138 correlation coefficient), the previous iteration revealed that sea surface dissolved oxygen and sea surface oxygen saturation are highly correlated ( 0.814 correlation coefficient). Thus, year was added as a model descriptor since it provided the second-best improvement to the model's fit ( $62.3 \%$ deviance explained) and is not correlated with any of the the current model descriptors ( 0.034 correlation coefficient with sea surface dissolved oxygen and 0.468 correlation coefficient with altimetry).

## B. 7 Cross Validation Results

Cross validation results for all fitted delta models are displayed in Figure B.5.

## B. 8 Developing Seasonal Fishnet Grids for Model Predictions

The development of seasonal, spatial biomass distribution profiles for each functional group was dependent on using the fitted GAMs to predict across grids of data representing hypothetical values of model descriptors. These grids were developed in ArcGIS. First, a $0.1^{\circ}$ latitude by $0.1^{\circ}$ longitude grid of geographic coordinates spanning the entire Gulf of Mexico was created using the Fishnet tool. Four versions of the grid were generated, one for each season. Next, coordinates within the grids were assigned estimates of all model descriptors.

## B.8.1 Environmental Data

To assign estimates of model descriptors to fishnets, environmental point data files needed to be converted to rasters. Bathymetry data, a polyline file, was first converted to a point file using the Feature to Point tool. Also, NCEI data (Table 3.1) was clipped to only contain points within the marine environment (i.e., removed the points representing Lake Okeechobee). Lastly, AVISO point data was split into seasonal datasets using the Select tool. Interpolation of point files to raster files was accomplished using the Kriging tool (all attributes remained set to default values). Rasters representing Gulf-wide seasonal averages were created for bathymetry (Figure B.6; assumed to not change seasonally), altimetry (Figure B.7), sea surface temperature (Figure B.8), sea bottom temperature (Figure B.9), sea bottom oxygen saturation (Figure B.10), sea surface dissolved oxygen (Figure B.11), sea bottom dissolved oxygen (Figure B.12), sea surface salinity (Figure B.13), and sea bottom salinity (Figure B.14).

Assigning values of environmental descriptors to seasonal Gulf-wide fishnet grids was done with the Extract Values to Points tool (set to bilinear interpolation). First, points in the Gulf-wide fishnet were assigned estimates of bathymetry. This allows the division of the Gulf-wide fishnet along the 250 m isobath - creating coastal and pelagic seasonal fishnets. Seasonal, environmental data were then assigned to appropriate fishnets.

## B.8.2 Minimum Distance From a Front

Fishnet grids were assigned estimates of minimum distance from a front using the routine described in Figure 3.2. The seasonal front polyline files created by seasonal AVISO averages (the point files) for calculating minimum distance from a front are shown in Figure B. 15.

## B. 9 Gulf-wide Abundance Distribution Profiles

Seasonal, spatial abundance distribution profiles spanning the entire Gulf of Mexico were developed for each functional group based on predicted abundance indices generated by the fitted statistical models. First, grids describing hypothetical seasonal conditions in the Gulf of Mexico were developed (see B.8). Fitted models were used to predict across fishnet grids to create seasonal, spatial distribution profiles.

Seasonal, Gulf-wide biomass distribution profiles were developed to improve the spatial representation of pelagic functional groups in the Gulf of Mexico Atlantis model. These profiles were developed for each functional group by 1) averaging the spatial catch rate profiles across overlapping polygons in the Gulf of Mexico Atlantis map, 2) extrapolating average catch rates for the remaining Gulf of Mexico polygon
map, then 3) calculating the proportion of catch rates in each polygon. Statistical software R was used to calculate the median catch rate for polygons overlapping the seasonal, spatial profiles of catch rates for each functional group. Median was used to prevent extreme catch rate predictions from having too much of an influence on the computed average. Two different methodologies were executed for extrapolating average catch rates across the remaining Atlantis polygon map.

Functional groups large pelagic fish, skates and rays, and large sharks were caught in both the survey and observer longline datasets, thus have GAMs for predicting across both pelagic and coastal areas in the Gulf of Mexico. Model predictions for large sharks are shown in the main paper, and the model predictions for large pelagic fish and skates and rays can be seen in Figure B. 16 and Figure B.18, respectively. Due to the difference in bottom longline and pelagic longline functional group catchability, the coastal and pelagic profiles must be converted to a common scale before they can be combined. Statistical models were developed to standardize the two longline datasets. Data used to fit these models contained both bottom and pelagic longline catch, but only data from similar spatial and temporal ranges where the two datasets overlapped. ArcGIS was used to create this data subset by selecting all the longline catch events that occurred in off the coast of Louisiana, an area where the two longline datasets intersect (Figure 3.1). Species from the large pelagic fish functional group were not retained in both of the longline datasets throughout the data subset, so another method had to be taken for extrapolating the average catch rates across the remaining Atlantis polygon map. The setup of the statistical model solving for the abundance index $\left(\eta_{Z}\right)$ was similar to Equation 3.1. The numerical descriptors include bottom depth $[\mathrm{m}]$, sea surface temperature $\left[{ }^{\circ} \mathrm{C}\right]$, altimetry $[\mathrm{m}[$,
and minimum distance from a front [m]. The categorical descriptors include year (2005-2010), season (1-4), and longline type (bottom or pelagic). Forward model selection was conducted to determine the best model for $\eta_{Z}$, for both functional groups (Table B.9). A standardization factor was calculated for average catch rates in pelagic polygons by dividing the median fitted CPUE of pelagic data by the median fitted CPUE of survey data. For large sharks the pelagic catch rate standardization factor is 0.522 , and for skates and rays it is 0.061 . Average catch rates for pelagic polygons were standardized by dividing the value by the standardization factor, thus allowing the pelagic and coastal profiles to be merged. The calculated standardization factors were used for all of the functional group's seasonal profiles.

For the remaining functional groups general assumptions were made for extrapolating estimates of the average catch rate to the remaining polygons. Online sources Fishbase (Froese and Pauly, 2016) and GulfBase (Moretzsohn et al., 2016), specifically the the Biodiversity of the Gulf of Mexico Database (Moretzsohn et al., 2011), were used to help gather information concerning species spatial distribution.

- BIL (other billfish) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.20. Catch in this group is primarily sailfish (Istiophorus albicans, but some identify I. platypterus as a world-wide species). Sailfish are known to be oceanic, spending much of the time oceanic environments (Riede, 2004). Kerstetter et al. (2010) used satellite tag data to discuss vertical and die distributions, but did not address Gulf-wide horizontal movements. To keep assumption general, polygons with depths defined as 10 , 20 , or 50 m were assigned a average catch rate of zero, and polygons with depths
defined as 200 m were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons. This was done for each seasonal profile.

- BMR (blue marlin) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.21. Comparing the results presented by Kraus et al. (2011) to polygons defined for the Gulf of Mexico Atlantis model, polygons with depths defined as 10,20 , or 50 m were assigned a average catch rate of zero, and polygons with depths defined as 200 m were assigned the smallest medium catch rate calculated when averaging across polygons. This was done for each seasonal profile.

- BTN (bluefin tuna) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.22. Considering results presented by Teo et al. (2007) polygons with depths defined as 10 or 20 m were assigned a average catch rate of zero, and polygons with depths defined as 50 or 200 m were assigned the smallest medium catch rate calculated when averaging across polygons. This was done for each seasonal profile. Bluefin tuna catch during the months of season 3 (Jul. - Sep.) are not present in the longline observer dataset, thus a catch rate profile could not be computed for this season. According to work by Block et al. (2005) bluefin tuna are found within Gulf of Mexico waters during this time. To develop a season 3 stock distribution profile for the Gulf of Mexico Atlantis model, the distribution of bluefin tuna were evenly distributed across polygons overlapping adult bluefin tuna hotspots reported by Block et al..

- DWF (deep water fish) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.23. Catch in this group consist of various species, many of the Cubiceps genus. Many of the DWF species are not coastal but rather oceanic (e.g., Cubiceps capensis (Riede, 2004)) and/or bathypelagic (e.g., Cubiceps pauciradiatus (Cervigón, 1994), Trachipterus arcticus (Muus et al., 1999)). Many of these species are concentrated around the slope Moretzsohn et al. (2011). All coastal polygons were assigned the smallest medium catch rate calculated when averaging across polygons (this was done for each seasonal profile). This aims to satisfy both observations: i) distributing DWF along the continental shelf, and ii) putting a slight emphasis on the slope, since GAMs estimates small catch rates for deep, oceanic polygons.

- FIL (filter feeding sharks) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.24. Catch in this group consist of mantas (of the family Mobulidae). Many of these species are epipelagic, centering around the slope Moretzsohn et al. (2011). All coastal polygons were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons (this was done for each seasonal profile). This encourages a strong emphasis on the slope since the GAM predicts small catch rates for deep, oceanic polygons.

- LPL (large pelagic fish) - extrapolate across coastal polygons

LPL species are retained in the bottom longline survey but after i) studying these catch records and ii) considering the coastal GAM predictions, the coastal model for LPL was not considered. The catch rate spatial profiles indicate a
strong seasonal flux, which we know is not part of the life histories of species in the LPL functional group. This flux is likely influenced by the fact that the survey operates primarily during summer and fall months. This signal is easier for the LPL model to pick up since there are very few instances of LPL species being caught in the bottom longline survey (barely enough to fit a statistical model). LPL catch records mostly consist of Remora sp. and Sphyraena barracuda, neither of which are benthic species. Thus, LPL catch records are likely incidental catch occurring when hooks are being set/hauled (i.e., traversing through the water column). Although it is clear that LPL species are within Gulf of Mexico coastal waters, the bottom longline survey is an inappropriate dataset for extrapolating information regarding the coastal distribution. Species identified in both longline catch datasets are known to use coastal waters (e.g., Remora remora (Fricke et al., 2011), Sphyraena barracuda (de Sylva, 1990), Pomatomus saltatrix (Claro, 1994)). All coastal polygons were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons. This was done for each seasonal profile.

- MPL (medium pelagic fish) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.25. Catch in this group consist of various species (e.g., Brama sp. and Megalops atlanticus). Many of these species have been found in coastal waters (e.g., Brama brama (May and Maxwell, 1986), Megalops atlanticus (Whitehead and Vergara, 1978)). For each seasonal profile, when calculating the median catch rate within polygons the median of those
catch rates was also calculated. The median catch rate, for each season, were assigned to all coastal polygons as the average catch rate.

- SMK (spanish mackerel) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.26. Two species dominate this catch: Lepidocybium flavobrunneum and Ruvettus pretiosus. Both species are benthopelagic and oceanic (Riede, 2004), as well as occupying the slope (Nakamura and Parin, 1993; Nakamura, 1995). Thus, polygons with depths defined as 10,20 , or 50 m were assigned a average catch rate of zero, and polygons with depths defined as 200 m were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons. This was done for each seasonal profile. Using the medium of catch rates will encourage a slight concentration of the functional group around the slope.

- SMS (small sharks) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.27. Some SMS species use the entire coast (Isistius brasiliensis (Kiraly et al., 2003)), and some are usually mesoand/or bathypelagic (e.g. Zameus squamulosus (Kiraly et al., 2003), Somniosus microcephalus (Muus et al., 1999; Moretzsohn et al., 2011), Squalus acanthias (Cox and Francis, 1997; Compagno, 2002)). Thus, polygons with depths defined as 10,20 , or 50 m were assigned an average catch rate of zero, and polygons with depths defined as 200 m were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons. This was done for each seasonal profile.

- SWD (swordfish) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.28. There is limited information discussing the horizontal distribution of swordfish, especially within the Gulf of Mexico. Dewar et al. (2011) presented data collected from swordfish tagged with PSAT tags, but the relatively low sample size and short deployment durations limit the utility for examining migratory patterns or stock structure. Nakamura (1985) generalizes that the stock is mostly pelagic with some instances of being in coastal habitats. Polygons with depths defined as 10 , 20, or 50 m were assigned a average catch rate of zero, and polygons with depths defined as 200 $m$ were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons. This was done for each seasonal profile.

- TIP (blacktip sharks) - extrapolate across pelagic polygons

Model predictions can be seen in Figure B.29. Blacktip sharks are primarily coastal, rarely moving through deep, oceanic waters (Compagno, 1984). All pelagic polygons were assigned an average catch rate equal to zero.

- TUR (other turtles) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.30. The TUR species identified in the catch records are known to be coastal (e.g., Chelonia mydas, Dermochelys coriacea, Eretmochelys imbricata (Moretzsohn et al., 2011)). All coastal polygons were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons. This was done for each seasonal profile.

- WMR (white marlin) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.31. White marlin generally prefer water deeper than 100 m (ICCAT, 2012). Polygons with depths defined as 10 , 20 , or 50 m were assigned a average catch rate of zero, and polygons with depths defined as 200 m were assigned the smallest medium catch rate calculated when averaging across polygons. This was done for each seasonal profile.

- YTN (yellowfin tuna) - extrapolate across coastal polygons

Model predictions can be seen in Figure B.32. Yellowfin tuna are known to be oceanic, spending much of the time oceanic environments (Riede, 2004). Polygons with depths defined as 10,20 , or 50 m were assigned a average catch rate of zero, and polygons with depths defined as 200 m were assigned an average catch rate equal to the medium of all the catch rates calculated when averaging across polygons. This was done for each seasonal profile.

## B. 10 Spatial Predications from Updated Pelagic Models: Large Sharks

Forward selection of pelagic models was originally done with training datasets for cross validation. This should have been done with the entire dataset (data for forecasting fitting). Forward selection of pelagic models was done with data for forecasting fitting, and these models were used to compute predictions. While for many functional groups predictions were the similar as those from pelagic models selected from training data, predictions for large sharks were quite different (Figure B.33). This is because the pelagic delta model selected with data for forecasting fitting
(Eqn. B.2) selected bottom depth as the variable providing the most improvement to AIC, so bottom depth was the first descriptor to have the basis dimension adjusted. This resulted in a smoothing spline for bottom depth to be more 'wiggly' than the smoothing spline reported in the chapter (Figure B.33). Predictions for large sharks presented in the chapter seem to make more ecological sense than predictions for large sharks from the model selected with data for forecasting fitting, and thus were selected to be used to parameterize the Gulf of Mexico Atlantis model. This is likely due to the smoothing spline for bottom depth being excessively 'wiggly' from the model selected with data for forecasting fitting. This highlights the limitations of using an automated routine for basis dimension setting.

$$
\begin{align*}
& g\left(\eta_{B}\right)=s(B D, 9)+f(y r)+f(s n)+s(M D F, 3)+s(S S H, 3)  \tag{B.2}\\
& g\left(\eta_{Z}\right)=s(B D, 21)+f(y r)+f(s n)+s(S S T, 9)+s(S S H, 25)+s(M D F, 18)
\end{align*}
$$

Table B.1: The deviance explained when bottom longline data was fit individually with the environmental and temporal variables considered to be model descriptors. Variables include year, daytime, latitude (LAT.DEG.N), longitude (LON.DEG.W), sea bottom depth (WATER.DEPTH), sea surface temperature (SST), sea surface height (SSH), minimum distance from a front (NEAR.DIST), beam transmission (XMISS), dissolved oxygen (OXY.MG), oxygen saturation (OXSAT), density, salinity, and some of these variables measured from the sea bottom (indicated by CTD.BT).

|  | Bernoulli Models |  |  |  | Gamma models |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | LGS | TIP | RAY | LPL | LGS | TIP | RAY | LPL |
| YEAR | 0.0268 | 0.0076 | 0.0162 | 0.0044 | 0.0408 | 0.1094 | 0.0628 | 0.1857 |
| DAYTIME | $3.13 \mathrm{E}-07$ | 0.0018 | 0.0028 | 0.0005 | 0.0002 | 0.0181 | $4.14 \mathrm{E}-05$ | 0.0134 |
| LAT.DEG.N | 0.0140 | 0.0138 | 0.0217 | 0.0714 | 0.0249 | 0.1422 | 0.0762 | 0.1592 |
| LON.DEG.W | 0.0099 | 0.0420 | 0.0779 | 0.0752 | 0.1532 | 0.2369 | 0.1930 | 0.0594 |
| WATER.DEPTH | 0.4486 | 0.3026 | 0.1032 | 0.0780 | 0.1224 | 0.1710 | 0.1659 | 0.0710 |
| SST | 0.0071 | 0.0332 | 0.0068 | 0.0441 | 0.0425 | 0.0742 | 0.1202 | 0.0420 |
| SSH | $5.62 \mathrm{E}-08$ | 0.0274 | 0.0213 | 0.0179 | 0.1253 | 0.1111 | 0.0184 | 0.1036 |
| NEAR.DIST | 0.0120 | 0.0441 | 0.0249 | 0.0344 | 0.0012 | 0.0392 | 0.0306 | $4.82 \mathrm{E}-07$ |
| XMISS | 0.0118 | 0.1820 | 0.0017 | 0.0326 | 0.0093 | 0.0636 | 0.0226 | $1.91 \mathrm{E}-06$ |
| OXY.MG | 0.0031 | 0.0156 | 0.0042 | 0.0592 | $5.91 \mathrm{E}-08$ | 0.0470 | $8.76 \mathrm{E}-07$ | 0.3034 |
| OXSAT | 0.0034 | 0.0123 | 0.0086 | 0.0471 | 0.0002 | 0.0881 | $2.82 \mathrm{E}-07$ | 0.1401 |
| DENSITY | $3.74 \mathrm{E}-07$ | 0.0995 | $8.84 \mathrm{E}-08$ | 0.0409 | 0.0276 | 0.2174 | 0.0267 | 0.0573 |
| SALINITY | 0.0047 | 0.1009 | $7.03 \mathrm{E}-07$ | 0.0097 | 0.0373 | 0.2418 | 0.0386 | 0.0556 |
| CTD.BT.Temp | 0.4260 | 0.2431 | 0.1104 | 0.0529 | 0.1609 | 0.1714 | 0.2169 | $2.23 \mathrm{E}-06$ |
| CTD.BT.XMISS | 0.0048 | 0.1612 | 0.0071 | 0.0593 | 0.2034 | 0.3179 | 0.2226 | 0.0759 |
| CTD.BT.OXY.MG | 0.0807 | 0.0334 | 0.0349 | 0.0725 | 0.0488 | 0.0683 | 0.0704 | 0.0547 |
| CTD.BT.OXSAT | 0.1778 | 0.0578 | 0.0579 | 0.0894 | 0.0335 | 0.0592 | 0.0926 | 0.1108 |
| CTD.BT.DENSITY | 0.4683 | 0.2546 | 0.0864 | 0.0247 | 0.0704 | 0.3057 | 0.1429 | $4.57 \mathrm{E}-06$ |
| CTD.BT.SAL | 0.0126 | 0.0716 | 0.0223 | 0.0046 | $1.47 \mathrm{E}-07$ | 0.3142 | 0.1141 | 0.0251 |

Table B.2: Comparing Statistical Fits of Gamma Link Functions. The three gamma link functions (identity, inverse, and $\log$ ) are being compared with the fit statistics generalized cross validation (GCV) and deviance explained. Bolded values indicated the best fit. The results for starred functional groups are derived from bottom longline survey data, and all other results are derived from pelagic longline observer data.

| Functional Group | Fit Statistic | Identity | Inverse | Log |
| :---: | :---: | :---: | :---: | :---: |
| BIL (other billfish) | GCV | 0.428 | 0.394 | 0.406 |
|  | Deviance Explained | 0.244 | 0.304 | 0.284 |
| BMR (blue marlin) | GCV | 0.337 | 0.327 | 0.334 |
|  | Deviance Explained | 0.188 | 0.215 | 0.194 |
| BTN (bluefin tuna) | GCV | 0.357 | 0.359 | 0.358 |
|  | Deviance Explained | 0.076 | 0.072 | 0.074 |
| DWF (deep water fish) | GCV | 0.068 | 0.062 | 0.065 |
|  | Deviance Explained | 0.386 | 0.436 | 0.412 |
| FIL (filter feeding sharks) | GCV | 0.161 | 0.147 | 0.156 |
|  | Deviance Explained | 0.213 | 0.296 | 0.243 |
| LGS (large sharks) | GCV | 0.790 | 0.626 | 0.594 |
|  | Deviance Explained | 0.536 | 0.633 | 0.652 |
| LGS (large sharks)* | GCV | 0.998 | 0.916 | 0.968 |
|  | Deviance Explained | 0.190 | 0.259 | 0.216 |
| LPL (large pelagic fish) | GCV | 0.993 | 0.966 | 0.966 |
|  | Deviance Explained | 0.212 | 0.234 | 0.234 |
| LPL (large pelagic fish)* | GCV | 0.206 | 0.189 | 0.200 |
|  | Deviance Explained | 0.270 | 0.366 | 0.309 |
| MPL (medium pelagic fish) | Deviance Explained | 0.252 0.048 | 0.251 0.056 | 0.252 0.052 |
| RAY (skates and rays) | GCV | 0.499 | 0.485 | 0.490 |
|  | Deviance Explained | 0.139 | 0.164 | 0.154 |
| RAY (skates and rays)* | GCV | 0.823 | 0.692 | 0.751 |
|  | Deviance Explained | 0.290 | 0.412 | 0.353 |
| SMK (spanish mackerel) | Deviance Explained | 0.661 0.064 | 0.068 | 0.067 |
| SMS (small sharks) | GCV | 0.474 | 0.398 | 0.424 |
|  | Deviance Explained | 0.405 | 0.503 | 0.469 |
| SWD (swordfish) | GCV | 0.738 | 0.662 | 0.665 |
|  | Deviance Explained | 0.349 | 0.417 | 0.413 |
| TIP (blacktip sharks)* | GCV | 0.696 | 0.685 | 0.683 |
|  | Deviance Explained | 0.159 | 0.169 | 0.174 |
| TUR (other turtles) | GCV | 0.208 | 0.138 | 0.169 |
|  | Deviance Explained | 0.511 | 0.673 | 0.601 |
| WMR (white marlin) | GCV | 0.303 | 0.293 | 0.299 |
| YTN (yellowfin tuna) | Deviance Explained | 0.187 0.669 | 0.218 | 0.202 |
|  | Deviance Explained | 0.104 | 0.112 | 0.108 |

Table B.3: Evaluating Basis Dimension Estimation Routine. This table displays the deviance explained for functional group-specific GAMs before and after the execution of the basis dimension estimation routine. Results for starred functional groups are derived from bottom longline survey data, and all other results are derived from pelagic longline observer data. All results come from models fit with training datasets developed for the cross validation.

| Functional <br> Group | Binomial Data Model |  | Zero-Truncated Data Model |  |
| :--- | :---: | :---: | :---: | :---: |
| Pre-Routine | Post-Routine | Pre-Routine | Post-Routine |  |
| other billfish | 0.305 | 0.305 | 0.277 | 0.352 |
| blue marlin | 0.207 | 0.207 | 0.192 | 0.245 |
| bluefin tuna | 0.431 | 0.431 | 0.088 | 0.088 |
| deep water fish | 0.117 | 0.117 | 0.671 | 0.671 |
| filter feeding sharks | 0.088 | 0.088 | 0.446 | 0.446 |
| large sharks | 0.197 | 0.197 | 0.648 | 0.702 |
| large sharks $^{\star}$ | 0.474 | 0.603 | 0.302 | 0.382 |
| large pelagic fish $_{\text {large pelagic fish }} \times 2$ | 0.150 | 0.150 | 0.227 | 0.321 |
| medium pelagic fish | 0.474 | 0.485 | 0.445 | 0.467 |
| skates and rays | 0.101 | 0.101 | 0.179 | 0.179 |
| skates and rays ${ }^{\star}$ | 0.221 | 0.221 | 0.359 | 0.399 |
| spanish mackerel | 0.165 | 0.264 | 0.294 | 0.582 |
| small sharks | 0.105 | 0.136 | 0.152 | 0.221 |
| swordfish | 0.225 | 0.225 | 0.520 | 0.604 |
| blacktip sharks | 0.238 | 0.261 | 0.413 | 0.479 |
| other turtles | 0.227 | 0.676 | 0.628 | 0.628 |
| white marlin | 0.085 | 0.085 | 0.724 | 0.724 |
| yellowfin tuna | 0.212 | 0.228 | 0.216 | 0.216 |

Table B.4: Summary of Correlations Among Environmental Variables for Models Fit with Pelagic Longline Observer Data. Presented are the correlation coefficients for both components of the Delta framework: models fit with binomial data and models fit with zerotruncated data. Model descriptors include year, season, sea surface temperature (SST), altimetry (SSH), bottom depth (BD), and minimum distance from a front (MDF). Functional group-specific binomial data models produce the same correlation matrix since models are fitted with the entire longline set data, but zero-truncated data models produce different correlation matrices since fitted datasets only include functional group-specific catch events. The correlation matrices are summarized here with the table below displaying the average, minimum, and maximum.

| Binomial Data Models |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Descriptors | year | season | SST | SSH | BD | MDF |
| year | - | - | - | - | - | - |
| season | -0.14 | - | - | - | - | - |
| SST | -0.16 | 0.51 | - | - | - | - |
| SSH | -0.10 | 0.29 | 0.37 | - | - | - |
| BD | 0.08 | 0.03 | 0.12 | -0.08 | - | - |
| MDF | 0.20 | 0.03 | 0.00 | -0.09 | -0.02 | - |
| Zero-Truncated Data Models |  |  |  |  |  |  |
| Descriptors | year | season | SST | SSH | BD | MDF |
| year | - | - | - | - | - | - |
| season | -0.13(-0.25-0.06) | - | - | - | - | - |
| SST | -0.14(-0.23--0.02) | 0.44(0.2-0.53) | - | - | - | - |
| SSH | -0.06(-0.16-0.14) | $0.27(0.08-0.44)$ | $0.35(0.09-0.56)$ | , | - | - |
| BD | 0.08(0-0.18) | 0.01(-0.15-0.08) | 0.1(-0.17-0.29) | $-0.07(-0.23-0.16)$ | - ${ }^{-}$ | - |
| MDF | $0.18(-0.15-0.26)$ | -0.01(-0.1-0.05) | -0.02(-0.16-0.12) | -0.1(-0.35-0.11) | 0.01(-0.12-0.17) | - |

Table B.5: Forward model selection results for GAMs fitted with large pelagic fish bottom longline survey data. Model descriptors included year, bottom depth, sea surface temperature (sst), sea bottom temperature (sbt), altimetry (ssh), minimum distance from a front (mdf), sea surface dissolved oxygen (do), sea bottom dissolved oxygen (sbdo), sea surface oxygen saturation (oxsat), sea bottom oxygen saturation (sboxsat), sea surface salinity (salinity), and sea bottom salinity (sbsalinity). Forward model selection ceased when descriptors failed to improve the deviance explained (d) more than $5 \%$. The displayed correlation coefficient $(c)$ is calculated based on the last two model descriptors. Bold-faced indicates selected models.

| Binomial Data Model | d | c | Zero-Truncated Data Model | d | c |
| :---: | :---: | :---: | :---: | :---: | :---: |
| year | 0.004 | - | year | 0.200 | - |
| depth | 0.086 | - | depth | 0.140 | - |
| sst | 0.032 | - | sst | 0.059 | - |
| ssh | 0.021 | - | ssh | 0.180 | - |
| mdf | 0.027 | - | mdf | $2.7 \mathrm{E}-7$ | - |
| do | 0.042 | - | do | 0.332 | - |
| oxsat | 0.032 | - | oxsat | 0.164 | - |
| salinity | 0.011 | - | salinity | 0.052 | - |
| sbt | 0.070 | - | sbt | 0.040 | - |
| sbdo | 0.080 | - | sbdo | 0.063 | - |
| sboxsat | 0.106 | - | sboxsat | 0.158 | - |
| sbsalinity | $1.4 \mathrm{E}-6$ |  | sbsalinity | 0.058 | - |
| sboxsat + year | 0.112 | -0.187 | do + year | 0.466 | 0.034 |
| sboxsat + depth | 0.124 | -0.500 | do + depth | 0.348 | -0.168 |
| sboxsat + sst | 0.144 | -0.263 | do + sst | 0.332 | -0.306 |
| sboxsat + ssh | 0.114 | -0.301 | do + ssh | 0.576 | 0.262 |
| sboxsat + mdf | 0.122 | 0.454 | do +mdf | 0.332 | -0.083 |
| sboxsat + do | 0.131 | 0.433 | do + oxsat | 0.379 | 0.814 |
| sboxsat + oxsat | 0.121 | 0.433 | do + salinity | 0.453 | -0.254 |
| sboxsat + salinity | 0.122 | 0.192 | do + sbt | 0.332 | 0.305 |
| sboxsat + sbt | 0.131 | 0.529 | do + sbdo | 0.351 | -0.135 |
| sboxsat + sbdo | 0.148 | 0.965 | do + sboxsat | 0.391 | 0.008 |
| sboxsat + sbsalinity | 0.106 | -0.037 | do + sbsalinity | 0.332 | -0.362 |
| sboxsat + sst + year | 0.147 | 0.133 | $\mathrm{do}+\mathrm{ssh}+$ year | 0.623 | 0.468 |
| sboxsat + sst + depth | 0.144 | -0.033 | do $+\mathrm{ssh}+$ depth | 0.576 | -0.132 |
| sboxsat + sst + ssh | 0.144 | -0.019 | do $+\mathrm{ssh}+\mathrm{sst}$ | 0.576 | -0.116 |
| sboxsat + sst + mdf | 0.154 | -0.226 | do $+\mathrm{ssh}+\mathrm{mdf}$ | 0.576 | -0.096 |
| sboxsat + sst + do | 0.172 | -0.511 | do + ssh + oxsat | 0.649 | 0.138 |
| sboxsat + sst + oxsat | 0.157 | -0.404 | do + ssh + salinity | 0.576 | 0.200 |
| sboxsat + sst + salinity | 0.146 | 0.300 | do $+\mathrm{ssh}+\mathrm{sbt}$ | 0.576 | 0.242 |
| sboxsat + sst + sbt | 0.156 | 0.048 | do + ssh + sbdo | 0.597 | 0.297 |
| sboxsat + sst + sbdo | 0.159 | -0.328 | do + ssh + sboxsat | 0.617 | 0.326 |
| sboxsat + sst + sbsalinity | 0.144 | 0.216 | do + ssh + sbsalinity | 0.576 | 0.017 |
| sboxsat + sst + do + year | 0.177 | -0.443 | do $+\mathrm{ssh}+$ year + depth | 0.626 | -0.389 |
| sboxsat + sst + do + depth | 0.189 | 0.031 | do $+\mathrm{ssh}+$ year +sst | 0.623 | -0.351 |
| sboxsat + sst + do + ssh | 0.174 | -0.194 | do $+\mathrm{ssh}+$ year +mdf | 0.623 | -0.075 |
| sboxsat + sst + do +mdf | 0.183 | 0.361 | do + ssh + year + salinity | 0.628 | 0.070 |
| sboxsat + sst + do + oxsat | 0.180 | 0.990 | do + ssh + year + sbt | 0.623 | 0.397 |
| sboxsat + sst + do + salinity | 0.177 | 0.094 | do + ssh + year + sbdo | 0.640 | 0.416 |
| sboxsat + sst + do + sbt | 0.182 | -0.056 | do $+\mathrm{ssh}+$ year + sboxsat | 0.659 | 0.462 |
| sboxsat + sst + do + sbdo | 0.175 | 0.533 | do + ssh + year + sbsalinity | 0.623 | -0.089 |
| sboxsat + sst + do + sbsalinity | 0.172 | -0.105 |  |  |  |
| sboxsat + sst + do + depth + year | 0.177 | -0.021 |  |  |  |
| sboxsat + sst + do + depth + ssh | 0.189 | 0.111 |  |  |  |
| sboxsat + sst + do + depth + mdf | 0.202 | -0.210 |  |  |  |
| sboxsat + sst + do + depth + oxsat | 0.197 | 0.033 |  |  |  |
| sboxsat + sst + do + depth + salinity | 0.177 | 0.071 |  |  |  |
| sboxsat + sst + do + depth + sbt | 0.189 | -0.878 |  |  |  |
| sboxsat + sst + do + depth + sbdo | 0.197 | -0.314 |  |  |  |
| sboxsat + sst + do + depth + sbsalinity | 0.203 | 0.049 |  |  |  |

Table B.6: Forward model selection results for GAMs fitted with large sharks bottom longline survey data. Model descriptors considered include year, bottom depth, sea surface temperature (sst), sea bottom temperature (sbt), altimetry (ssh), minimum distance from a front (mdf), sea surface dissolved oxygen (do), sea bottom dissolved oxygen (sbdo), sea surface oxygen saturation (oxsat), sea bottom oxygen saturation (sboxsat), sea surface salinity (salinity), and sea bottom salinity (sbsalinity). Forward model selection ceased when descriptors failed to improve the deviance explained ( $d$ ) more than $5 \%$. The displayed correlation coefficient (c) is calculated based on the last two model descriptors. Bold-faced indicates selected models.

| Binomial Data Model | d | c | Zero-Truncated Data Model | d | c |
| :---: | :---: | :---: | :---: | :---: | :---: |
| year | 0.008 | - | year | 0.037 | - |
| depth | 0.505 | - | depth | 0.144 | - |
| sst | 0.007 | - | sst | 0.070 | - |
| ssh | 0.017 | - | ssh | 0.146 | - |
| mdf | 0.031 | - | mdf | 0.004 | - |
| do | 0.010 | - | do | 0.002 | - |
| oxsat | 0.008 | - | oxsat | 0.004 | - |
| salinity | 0.006 | - | salinity | 0.037 | - |
| sbt | 0.454 | - | sbt | 0.168 | - |
| sbdo | 0.201 | - | sbdo | 0.050 | - |
| sboxsat | 0.198 | - | sboxsat | 0.036 | - |
| sbsalinity | 0.232 | - | sbsalinity | $1.9 \mathrm{E}-7$ | - |
|  |  |  | sbt + year | 0.205 | 0.030 |
|  |  |  | sbt + depth | 0.190 | -0.837 |
|  |  |  | $s \mathrm{sbt}+\mathrm{sst}$ | 0.223 | 0.030 |
|  |  |  | $\mathbf{s b t}+\mathbf{s s h}$ | 0.291 | -0.050 |
|  |  |  | $\mathrm{sbt}+\mathrm{mdf}$ | 0.176 | 0.193 |
|  |  |  | sbt + do | 0.183 | -0.016 |
|  |  |  | sbt + oxsat | 0.183 | -0.016 |
|  |  |  | sbt + salinity | 0.208 | -0.078 |
|  |  |  | sbt + sbdo | 0.244 | 0.235 |
|  |  |  | sbt + sboxsat | 0.237 | 0.427 |
|  |  |  | sbt + sbsalinity | 0.168 | -0.382 |
|  |  |  | $\mathbf{s b t}+\mathrm{ssh}+$ year | 0.321 | 0.288 |
|  |  |  | $s \mathrm{sb}+\mathrm{ssh}+$ depth | 0.328 | 0.115 |
|  |  |  | sbt + ssh + sst | 0.310 | -0.044 |
|  |  |  | $\mathrm{sbt}+\mathrm{ssh}+\mathrm{mdf}$ | 0.299 | -0.241 |
|  |  |  | $\mathrm{sbt}+\mathrm{ssh}+$ do | 0.297 | -0.147 |
|  |  |  | sbt $+\mathrm{ssh}+$ oxsat | 0.297 | -0.177 |
|  |  |  | sbt $+\mathrm{ssh}+$ salinity | 0.297 | -0.338 |
|  |  |  | sbt $+\mathrm{ssh}+$ sbdo | 0.314 | -0.389 |
|  |  |  | sbt + ssh + sboxsat | 0.312 | -0.377 |
|  |  |  | sbt + ssh + sbsalinity | 0.293 | -0.164 |

Table B.7: Forward model selection results for GAMs fitted with skates and rays bottom longline survey data. Model descriptors considered include year, bottom depth, sea surface temperature (sst), sea bottom temperature (sbt), altimetry (ssh), minimum distance from a front (mdf), sea surface dissolved oxygen (do), sea bottom dissolved oxygen (sbdo), sea surface oxygen saturation (oxsat), sea bottom oxygen saturation (sboxsat), sea surface salinity (salinity), and sea bottom salinity (sbsalinity). Forward model selection ceased when descriptors failed to improve the deviance explained ( $d$ ) more than $5 \%$. The displayed correlation coefficient (c) is calculated based on the last two model descriptors. Bold-faced indicates selected models.

| Binomial Data Model $\begin{array}{r}\text { year } \\ \text { depth } \\ \text { sst }\end{array}$ | d | c | Zero-Truncated Data Model | d | c |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | 0.006 | - | year | 0.025 | - |
|  | 0.127 | - | depth | 0.155 | - |
|  | $1.5 \mathrm{E}-7$ | - | sst | 0.101 | - |
| ssh | 0.015 | - | ssh | 3.E-6 | - |
| mdf | 0.027 | - | mdf | 0.015 | - |
| do | 0.003 | - | do | 0.003 | - |
| oxsat | 0.005 | - | oxsat | $1.2 \mathrm{E}-6$ | - |
| salinity | $2.5 \mathrm{E}-7$ | - | salinity | 0.034 | - |
| sbt | 0.126 | - | sbt | 0.210 | - |
| sbdo | 0.090 | - | sbdo | 0.056 | - |
| sboxsat | 0.110 | - | sboxsat | 0.098 | - |
| sbsalinity | 0.022 | - | sbsalinity | 0.092 | - |
| depth + year | 0.131 | 0.243 | sbt + year | 0.250 | -0.024 |
| depth + sst | 0.134 | 0.244 | sbt + depth | 0.231 | -0.830 |
| depth + ssh | 0.132 | 0.239 | $\mathbf{s b t}+\mathbf{s s t}$ | 0.328 | 0.062 |
| depth + mdf | 0.132 | -0.259 | sbt +ssh | 0.210 | 0.043 |
| depth + do | 0.129 | 0.031 | $s b t+m d f$ | 0.210 | 0.089 |
| depth + oxsat | 0.132 | 0.032 | sbt + do | 0.229 | -0.009 |
| depth + salinity | 0.137 | 0.071 | sbt + oxsat | 0.216 | -0.004 |
| depth + sbt | 0.142 | -0.877 | sbt + salinity | 0.325 | -0.045 |
| depth + sbdo | 0.146 | -0.314 | sbt + sbdo | 0.237 | 0.277 |
| depth + sboxsat | 0.150 | -0.500 | sbt + sboxsat | 0.221 | 0.508 |
| depth + sbsalinity | 0.127 | 0.049 | sbt + sbsalinity | 0.210 | -0.021 |
|  |  |  | $s b t+s s t+y e a r$ | 0.342 | 0.339 |
|  |  |  | sbt + sst + depth | 0.337 | -0.011 |
|  |  |  | sbt $+\mathrm{sst}+\mathrm{ssh}$ | 0.328 | 0.245 |
|  |  |  | $s \mathrm{sbt}+\mathrm{sst}+\mathrm{mdf}$ | 0.328 | -0.136 |
|  |  |  | sbt + sst + do | 0.336 | -0.468 |
|  |  |  | sbt + sst + oxsat | 0.331 | -0.369 |
|  |  |  | $\mathbf{s b t}+\mathbf{s s t}+$ salinity | 0.418 | 0.144 |
|  |  |  | sbt + sst + sbdo | 0.329 | -0.302 |
|  |  |  | sbt + sst + sboxsat | 0.329 | -0.252 |
|  |  |  | sbt + sst + sbsalinity | 0.328 | 0.241 |

Table B.8: Forward model selection results for GAMs fitted with blacktip sharks bottom longline survey data. Model descriptors considered include year, bottom depth, sea surface temperature (sst), sea bottom temperature (sbt), altimetry (ssh), minimum distance from a front (mdf), sea surface dissolved oxygen (do), sea bottom dissolved oxygen (sbdo), sea surface oxygen saturation (oxsat), sea bottom oxygen saturation (sboxsat), sea surface salinity (salinity), and sea bottom salinity (sbsalinity). Forward model selection ceased when descriptors failed to improve the deviance explained ( $d$ ) more than $5 \%$. The displayed correlation coefficient (c) is calculated based on the last two model descriptors. Bold-faced indicates selected models.

| Binomial Data Model | d | c | Zero-Truncated Data Model | d | c |
| :---: | :---: | :---: | :---: | :---: | :---: |
| year | 0.007 | - | year | 0.106 | - |
| depth | 0.307 | - | depth | 0.181 | - |
| sst | 0.033 | - | sst | 0.141 | - |
| ssh | 0.027 | - | ssh | 0.130 | - |
| mdf | 0.048 | - | mdf | 0.027 | - |
| do | 0.017 | - | do | 0.088 | - |
| oxsat | 0.010 | - | oxsat | 0.100 | - |
| salinity | 0.115 | - | salinity | 0.246 | - |
| sbt | 0.245 | - | sbt | 0.211 | - |
| sbdo | 0.023 | - | sbdo | 0.172 | - |
| sboxsat | 0.034 | - | sboxsat | 0.164 | - |
| sbsalinity | 0.066 | - | sbsalinity | 0.345 | - |
| depth + year | 0.315 | 0.243 | sbsalinity + year | 0.380 | 0.075 |
| depth + sst | 0.320 | 0.244 | sbsalinity + depth | 0.388 | 0.381 |
| depth + ssh | 0.357 | 0.239 | sbsalinity + sst | 0.382 | 0.191 |
| depth + mdf | 0.319 | -0.259 | sbsalinity + ssh | 0.351 | -0.287 |
| depth + do | 0.325 | 0.031 | sbsalinity +mdf | 0.356 | -0.123 |
| depth + oxsat | 0.323 | 0.032 | sbsalinity + do | 0.367 | -0.050 |
| depth + salinity | 0.364 | 0.071 | sbsalinity + oxsat | 0.370 | 0.013 |
| depth + sbt | 0.313 | -0.877 | sbsalinity + salinity | 0.365 | 0.584 |
| depth + sbdo | 0.344 | -0.314 | sbsalinity + sbt | 0.394 | -0.225 |
| depth + sboxsat | 0.345 | -0.500 | sbsalinity + sbdo | 0.403 | 0.059 |
| depth + sbsalinity | 0.324 | 0.049 | sbsalinity + sboxsat | 0.398 | 0.062 |
| depth + salinity + year | 0.375 | 0.029 | sbsalinity + sbdo + year | 0.422 | -0.457 |
| depth + salinity + sst | 0.378 | 0.298 | sbsalinity + sbdo + depth | 0.475 | -0.060 |
| depth + salinity + ssh | 0.395 | -0.179 | sbsalinity + sbdo + sst | 0.410 | -0.424 |
| depth + salinity + mdf | 0.380 | 0.056 | sbsalinity + sbdo + ssh | 0.403 | -0.528 |
| depth + salinity + do | 0.373 | 0.094 | sbsalinity + sbdo +mdf | 0.403 | 0.506 |
| depth + salinity + oxsat | 0.372 | 0.164 | sbsalinity + sbdo + do | 0.406 | 0.594 |
| depth + salinity + sbt | 0.367 | -0.100 | sbsalinity + sbdo + oxsat | 0.407 | 0.615 |
| depth + salinity + sbdo | 0.384 | 0.237 | sbsalinity + sbdo + salinity | 0.406 | 0.397 |
| depth + salinity + sboxsat | 0.387 | 0.192 | sbsalinity + sbdo + sbt | 0.470 | -0.043 |
| depth + salinity + sbsalinity | 0.373 | 0.407 | sbsalinity + sbdo + sboxsat | 0.458 | 0.992 |
| depth + salinity + ssh + year | 0.407 | 0.374 |  |  |  |
| $\text { depth }+ \text { salinity }+ \text { ssh }+ \text { sst }$ | 0.410 | -0.018 |  |  |  |
| depth + salinity + ssh + mdf | 0.409 | -0.254 |  |  |  |
| depth + salinity + ssh + do | 0.405 | -0.194 |  |  |  |
| depth + salinity + ssh + oxsat | 0.404 | -0.220 |  |  |  |
| depth + salinity + ssh + sbt | 0.400 | -0.029 |  |  |  |
| depth + salinity + ssh + sbdo | 0.413 | -0.329 |  |  |  |
| depth + salinity + ssh + sboxsat | 0.415 | -0.301 |  |  |  |
| depth + salinity + ssh + sbsalinity | 0.406 | -0.121 |  |  |  |

Table B.9: Forward Model Selection results for GAMs solving for catch-per-unit-effort to be used to standardize coastal and pelagic predictions of large sharks (left) and skates and rays (right). Model descriptors considered include year, season, longline type (title), bottom depth, sea surface temperature (sst), altimetry (ssh), and minimum distance from a front (mdf). Descriptors were selected based on model deviance explained (d). Forward model selection ceased when descriptors provided no improvement to model fits. Bold-faced indicates selected models.

| large sharks | $d$ | skates and rays | $d$ |
| :---: | :---: | :---: | :---: |
| year | 0.199 | year | 0.187 |
| season | 0.25 | season | 0.303 |
| title | 0.212 | title | 0.235 |
| depth | 0.377 | depth | 0.192 |
| sst | 0.167 | sst | 0.221 |
| ssh | 0.252 | ssh | 0.011 |
| mdf | $3.23 \mathrm{E}-7$ | mdf | 0.209 |
| depth + year | 0.518 | season + year | 0.451 |
| depth + season | 0.46 | $\text { season }+ \text { title }$ | 0.31 |
| depth + title | 0.451 | season + depth | 0.447 |
| $\text { depth }+ \text { sst }$ | 0.444 | season + sst | 0.349 |
| $\text { depth }+ \text { ssh }$ | 0.456 | season + ssh | 0.303 |
| $\text { depth }+ \text { mdf }$ | 0.377 | season + mdf | 0.472 |
| depth + year + season | 0.57 | season + mdf + year | 0.602 |
| depth + year + title | 0.55 | season + mdf + title | 0.475 |
| $\text { depth }+ \text { year }+ \text { sst }$ | 0.542 | season + mdf + depth | 0.534 |
| depth + year + ssh | 0.553 | season + mdf + sst | 0.472 |
| depth + year + mdf | 0.537 | season + mdf + ssh | 0.482 |
| depth + year + season + title | 0.57 | season + mdf + year + title | 0.611 |
| $\text { depth }+ \text { year }+ \text { season }+ \text { sst }$ | 0.57 | season + mdf + year + depth | $0.603$ |
| depth + year + season + ssh | 0.57 | season + mdf + year + sst | 0.602 |
| depth + year + season + mdf | $0.588$ | season + mdf + year +ssh | 0.602 |
| depth + year + season + mdf + title | 0.589 |  |  |
| depth + year + season + mdf + sst | 0.597 |  |  |
| depth + year + season + mdf + ssh | 0.588 |  |  |
| depth + year + season + mdf + sst + title | 0.598 |  |  |
| depth + year + season + mdf + sst + ssh | 0.597 |  |  |



Figure B.1: Fronts, indicated by blue lines, produced by the Cayula-Cornillon Fronts in ArcGIS Raster MGET tool in ArcGIS. Panel (a) displays the results produced when 2006 sea surface temperature data from January, February, and March (season 1) are processed. Panel (b) displays the results produced with 2006 altimetry data from season 1, scaled by 1000, are processed.


Figure B.2: The variance in catch rate is plotted against the mean catch rate. Panels (a) (d) are derived from bottom longline survey data and panels (e) - (s) are derived from pelagic longline observer data. For all images the black line displays the square of the average catch rate curve.


Figure B.3: The large sharks seasonal predictions for day ( $\mathrm{a}-\mathrm{d}$ ) are similar to the predictions for night (e-h). The swordfish seasonal predictions for day (i-l) are slightly different than the corresponding predictions for night ( $\mathrm{m}-\mathrm{p}$ ). Columns correspond to seasons.


Figure B.4: The distribution plots of normal-scale catch rates (black lines; corresponding to the primary axis) and log-transformed catch rates (grey lines; corresponding to the secondary axis). Panels (a) - (d) are derived from bottom longline survey data while panels (e) - (s) are derived from pelagic longline observer data.


Figure B.5: Observed catch rates against predicted catch rates for coastal models (a-d), and pelagic models ( $e-s$ ). Solid lines indicate the fitted linear regression and the dashed line indicates the 1:1 line. The plot title states the intercept, slope, and adjusted r-square value.


Figure B.6: Bathymetry [m] estimates across the Gulf of Mexico. The raster was developed in ArcGIS by processing bathymetry data (Table 3.1) with the Kriging tool (default settings).


Figure B.7: Seasonal average estimates of sea surface height [m] across the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea surface height data (Table 3.1) with the Kriging tool (default settings).


Figure B.8: Seasonal average estimates of sea surface temperature $\left[\mathrm{C}^{\circ}\right]$ across the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea surface temperature data (Table 3.1) with the Kriging tool (default settings).


Figure B.9: Seasonal average estimates of sea bottom temperature [ $\mathrm{C}^{\circ}$ ] across the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea bottom temperature data (Table 3.1) with the Kriging tool (default settings).


Figure B.10: Seasonal average estimates of sea bottom oxygen saturation [\%] across the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea bottom oxygen saturation data (Table 3.1) with the Kriging tool (default settings).


Figure B.11: Seasonal average estimates of sea surface dissolved oxygen [ $\mathrm{ml} / \mathrm{l}]$ across the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea surface dissolved oxygen data (Table 3.1) with the Kriging tool (default settings).


Figure B.12: Seasonal average estimates of the sea bottom dissolved oxygen [ml/l] across the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea bottom dissolved oxygen data (Table 3.1) with the Kriging tool (default settings).


Figure B.13: Seasonal average estimates of sea surface salinity [pss] the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea surface salinity data (Table 3.1) with the Kriging tool (default settings).


Figure B.14: Seasonal average estimates of sea bottom salinity [pss] the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Rasters were created in ArcGIS by processing seasonal sea bottom salinity data (Table 3.1) with the Kriging tool (default settings).


Figure B.15: Seasonal estimates of the average front locations within the Gulf of Mexico during the winter (a), spring (b), summer (c), and fall (d). Front polyline files were created by processing AVISO point data, separated by season, by the routine described in Figure 3.2 for calculating minimum distance from a front.


Figure B.16: Catch rates of large pelagic fish predicted by GAMs fitted with bottom longline survey data ( $a-d$ ), and GAMs fitted with pelagic longline observer data ( $e-h$ ). Columns correspond to season.


Figure B.17: Standard error of seasonal predictions from the large pelagic fish coastal logistic model (a-d), coastal Gamma model (e-h), pelagic logistic model (i-l), and pelagic Gamma model ( $\mathrm{m}-\mathrm{p}$ ). Columns correspond to season.


Figure B.18: Catch rates of skates and rays predicted by GAMs fitted with bottom longline survey data ( $\mathrm{a}-\mathrm{d}$ ), and GAMs fitted with pelagic longline observer data ( $\mathrm{e}-\mathrm{h}$ ). Columns correspond to season.


Figure B.19: Standard error of seasonal predictions from the skates and rays coastal logistic model ( $\mathrm{a}-\mathrm{d}$ ), coastal Gamma model (e-h), pelagic logistic model (i-l), and pelagic Gamma model ( $\mathrm{m}-\mathrm{p}$ ). Columns correspond to season.


Figure B.20: Catch rates of other billfish predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.21: Catch rates of blue marlin predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i - l). Columns correspond to season.


Figure B.22: Catch rates of bluefin tuna predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{c}$ ), standard error of seasonal predictions from the logistic model (d f), and standard error of seasonal predictions from the Gamma model (g - i). Columns correspond to season ( 1,2 , and 4 ).


Figure B.23: Catch rates of deep water fish predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i - l). Columns correspond to season.


Figure B.24: Catch rates of filter feeding sharks predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model ( $\mathrm{e}-\mathrm{h}$ ), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.25: Catch rates of medium pelagic fish predicted by GAMs fitted with pelagic longline observer data ( $\mathrm{a}-\mathrm{d}$ ), standard error of seasonal predictions from the logistic model (e - h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.26: Catch rates of spanish mackerel predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.27: Catch rates of small sharks predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.28: Catch rates of swordfish predicted by GAMs fitted with pelagic longline observer data ( $a-d$ ), standard error of seasonal predictions from the logistic model (e - h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.29: Catch rates of blacktip sharks predicted by GAMs fitted with bottom longline survey data ( $\mathrm{a}-\mathrm{d}$ ), standard error of seasonal predictions from the logistic model ( $\mathrm{e}-\mathrm{h}$ ), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.30: Catch rates of other turtles predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.31: Catch rates of white marlin predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.32: Catch rates of yellowfin tuna predicted by GAMs fitted with pelagic longline observer data $(\mathrm{a}-\mathrm{d})$, standard error of seasonal predictions from the logistic model (e h), and standard error of seasonal predictions from the Gamma model (i-l). Columns correspond to season.


Figure B.33: Panels a - d show the pelagic predictions of large sharks as presented in the chapter, from the model selected with training data for cross validation. Panels e - h show the pelagic predictions of large sharks from the model selected with data for forecast fitting. Panel i shows the smoothing spline of bottom depth for the large sharks Gamma pelagic model presented in the chapter, from the model selected with training data for cross validation. Panel j shows the smoothing spline of bottom depth for the large sharks Gamma pelagic model selected with data for forecast fitting.

## APPENDIX C

## Additional Methodology and Results for Chapter 4

## C. 1 Exploratory Calibration

## C.1. 1 Bluefin Tuna

Re-parameterization of the bluefin tuna functional group was explored in the attempt to improve diagnostics. Specifically, diet parameters, and migration parameters were adjusted. However, no investigated re-parameterizations meaningfully improved bluefin tuna diagnostics. For example, I attempted to adjust the migration parameters to i) reflect time frames described by Block et al. (2005) and Teo et al. (2007), and ii) have a longer transition period into/out of the modeling domain. Adjustments are summarized in Table C.1. These adjustments caused a sudden collapse in the bluefin tuna stock (Figure C.1), which could not be mitigated before the start of this study. Thus, the study was conducted with the original bluefin tuna parameterization of diet and migration.

## C. 2 Additional Methods

## C.2.1 Biology Input File

## C.2.1.1 Vertical Distribution of Little Tunny

Vectors describing the vertical distribution identify the proportion of functional groups in each depth layer. The first position of the vector identifies the sea floor. The last position of the vector identifies the sea surface. Starting from the last position in the vector and moving forward identifies descending depths. The original nighttime vertical distribution for little tunny (juveniles) put all corresponding biomass in the layer closest to the bottom:

VERTnight_LTN1 610.00 .00 .00 .00 .0
I adjusted it to match the nighttime distribution of juveniles of other tuna groups, putting organisms near the sea surface:

VERTnight_LTN1 60.00 .00 .00 .00 .10 .9

## C.2.1.2 Density and Nitrogen Diagnostics

The version of GoMAM I received had poor diagnostics for both the blue marlin (BMR) and white marlin (WMR) functional groups (Figure C.2). First, both functional groups were quickly collapsing (Figure C.2a,b), Second, adults were losing residual nitrogen (similar to starving), especially WMR (Figure C.2e,f). Lastly, adults were losing structural nitrogen (similar to shrinking), especially WMR (Figure C.2g,h). Since these two functional groups were of focus species for the study it was imperative to improve their diagnostics. Edits were made to the biological input file
following suggestions provided by the Atlantis modeling community (http://atlantis.cmar.csiro.au/).
C.2.1.2.1 White Marlin: Diagnostic plots suggested that WMR could be starving and information from the Atlantis wiki suggested, based on the diagnostics, to alter the predator-prey relationship parameter (pprey). This included minor adjustments to slightly expand their prey groups (Llopiz and Cowen, 2008) and allow increased consumption. Table C. 2 shows the original matrix and the one used for this study. Updated diagnostic plots are shown in Figure C.3.
C.2.1.2.2 Blue Marlin: Diagnostic plots suggested that BMR could be starving and information from the Atlantis wiki suggested, based on the diagnostics, to alter the predator-prey relationship parameter (pprey). Many attempts were made adjusting pprey parameters with no success in improving diagnostics. Based on information from the Atlantis wiki, the next attempt at improving BMR diagnostics involved adjusting Beverton Holt alpha parameters (BHalpha). The BHalpha for BMR was 3700, but other billfish functional groups had much higher settings: WMR had 150000 and BIL had 78000 (Ainsworth et al., 2015). BHalpha_BMR was iteratively adjusted until density of younger individuals improved. The value used for this study is 18500. Diagnostics for size at age specifically residual nitrogen (RN) were not stabilized. Information from the Atlantis wiki suggested to alter initial conditions for residual nitrogen (KWRR). The original value was 607435.91533 (Ainsworth et al., 2015), which was iteratively adjusted until diagnostics had improved stabilization. The value used is 400000.0 . Although BMR are still not stable, I stopped calibrating
the model here since BMR were no longer collapsing and size at age improved. The updated diagnostic plots are shown in Figure C. 3

## C.2.2 Harvest Input File

The version of GoMAM I used simulated harvest using a matrix describing constant, daily fishing mortalities, which was developed using historical landings data described in Chapter 2 of this dissertation (Perryman et al., 2015). However, 2010 landings (Table C.3). This occured because first, simulated fishing mortality rates were computed under the assumption that functional groups are within the modeling domain the entire 365 days (i.e., no migration). Thus, migrating groups are not being fished as hard as they likely are being fished in reality. Second, simulated fishing mortality rates were computed under the assumption that fleets fished the entire modeling domain (i.e., no regional spatial restrictions). Thus, fleets are not harvesting regional areas hard enough since they are fishing all across the spatial domain. Values in the fishing mortality matrix were updated for this study.

First, 2010 bycatch data (National Marine Fisheries Service, 2013) was collected and included into the 2010 landings data reported in Chapter 2 of this dissertation. Then, values for the fishing mortality matrix were iteratively adjusted until 2010 simulated catches were similar to 2010 landings data. The resulting matrix of fishing mortalities is presented in Table C.4.

## C.2.3 Additions to Fishery Closure Forcing Files

Forcing files simulating fishery spatial closures were updated to include Spring Closure, the seasonal pelagic longline spatial closure off the Louisiana coast, as well
as the Deepwater Horizon (DWH) emergency fishery closures. The DWH emergency closure drastically changed in spatial coverage in short periods of time. To provide a detailed representation of the DWH closure, alterations were treated as short-lived, individual fishery closures. Shapefiles describing all of the spatial boundaries of the DWH emergency closure were provided by National Centers for Environmental Information (2015). The Intersect tool in ArcGIS was used to compute the proportions of the the GoMAM polygons overlapping spatial closures.

Table C.1: Example of alterations made to bluefin tuna migration parameters, with rep (a,n) indicating a vector of $n$ elements of $a$.

| Parameter | jBTN_Migrate_Time |
| :--- | ---: |
| Original | 135 |
| Attempted | 180 |
| Parameter | BTN_Migrate_Time |
| Original | 364 |
| Attempted | 182 |
| Parameter | jBTN_Migrate_Return |
| Original | 340 |
| Attempted | 32 |
| Parameter | BTN_Migrate_Return |
| Original | 0 |
| Attempted |  |
| Parameter | jBTN_Migrate_Period |
| Original | 60 |
| Attempted |  |
| Parameter | BTN_Migrate_Period |
| Original | 1 |
| Attempted | 90 |
| Parameter | MigIOBox_BTNad 66 |
| Original | rep $(0,66)$ |
| Attempted | rep $(1,66)$ |



Figure C.1: Diagnostic plots commonly referred to when calibrating an Atlantis model include biomass, density, residual nitrogen (i.e., meat, fat), and structural nitrogen (i.e., bone). This figure bluefin tuna diagnostic plots from the original parameterization (a-d), and from the attempted calibration described in Section C.1.1 (e-h). Images with a solid, black line indicate trends for the entire population, while multi-colored lines indicate trends for each of the 10 cohorts (colors are a gradient, with red indicating cohort 1 and blue indicating cohort 10).


Figure C.2: Diagnostic plots commonly referred to when calibrating an Atlantis model include biomass, density, residual nitrogen (i.e., meat, fat), and structural nitrogen (i.e., bone). This figure displays these diagnostic plots for white marlin and blue marlin functional groups for the original version of the Gulf of Mexico Atlantis Model provided for this study. Images with a solid, black line indicate trends for the entire population, while multi-colored lines indicate trends for each of the 10 cohorts (colors are a gradient, with red indicating cohort 1 and blue indicating cohort 10).
Table C.2: White Marline Predator-Prey Relationship Parameters.

| pPREY1WMR1 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | GAG | RGR | SCM | SSR | DSR | RSN | VSN | LUT | BIO | LRF | SRF | BDR | RDR | SEA | SCI | LDY |
| Original | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $5.81 \mathrm{E}-04$ | 0 |
| Used | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $5.81 \mathrm{E}-04$ | 0 |
|  | MUL | POM | SHP | SNK | FLT | ODF | SDF | YTN | BTN | LTN | OTN | SWD | WMR | BMR | BIL | AMB |
| Original | 0 | 0 | 0.28 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0.28 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
|  | JCK | KMK | SMK | SAR | LPL | DWF | MEN | PIN | MPL | SPL | TIP | BEN | LGS | FIL | SMS | RAY |
| Original | 0 | 0 | 0 | 0 | $7.50 \mathrm{E}-03$ | 0.156 | $5.40 \mathrm{E}-02$ | 0 | 0.4 | $5.14 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0 | 0 | $7.50 \mathrm{E}-03$ | 0.156 | $5.40 \mathrm{E}-02$ | 0 | 0.4 | $5.14 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | 0 |
|  | BSH | WSH | PSH | OSH | DBR | SBR | MAN | MYS | DOL | DDO | LOG | KMP | TUR | BCR | SCR | LOB |
| Original | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
|  | COR | CCA | OCT | SPG | CMB | INF | ECH | OYS | BIV | SES | EPI | GRS | ALG | MPB | LPP | SPP |
| Original | 0 | 0 | 0 | $2.68 \mathrm{E}-07$ | 0 | $1.34 \mathrm{E}-02$ | 0 | 0 | $2.96 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0 | $2.68 \mathrm{E}-07$ | 0 | $1.34 \mathrm{E}-02$ | 0 | 0 | $2.96 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
|  | DIN | PRO | JEL | SQU | LZP | SZP | PB | BB | DC | DL | DR | Dcsed | Dlsed | Drsed |  |  |
| Original | 0 | 0 | 0 | $7.23 \mathrm{E}-02$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |  |  |
| Used | 0 | 0 | 0 | 7.23E-02 | 0 | 0.1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |  |  |
| $p$ PREY2WMR1 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  | GAG | RGR | SCM | SSR | DSR | RSN | VSN | LUT | BIO | LRF | SRF | BDR | RDR | SEA | SCI | LDY |
| Original | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.184 | 0 |
| Used | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.184 | 0 |
|  | MUL | POM | SHP | SNK | FLT | ODF | SDF | YTN | BTN | LTN | OTN | SWD | WMR | BMR | BIL | AMB |
| Original | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $3.15 \mathrm{E}-03$ | $8.18 \mathrm{E}-05$ | $6.20 \mathrm{E}-05$ | $1.20 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $3.15 \mathrm{E}-03$ | $8.18 \mathrm{E}-05$ | $6.20 \mathrm{E}-05$ | $1.20 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 |
|  | JCK | KMK | SMK | SAR | LPL | DWF | MEN | PIN | MPL | SPL | TIP | BEN | LGS | FIL | SMS | RAY |
| Original | 0 | 0 | 0 | $5.09 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | $8.78 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0 | $5.09 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | $8.78 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 | 0 |
|  | BSH | WSH | PSH | OSH | DBR | SBR | MAN | MYS | DOL | DDO | LOG | KMP | TUR | BCR | SCR | LOB |
| Original | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
|  | COR | CCA | OCT | SPG | CMB | INF | ECH | OYS | BIV | SES | EPI | GRS | ALG | MPB | LPP | SPP |
| Original | 0 | 0 | 0 | $2.68 \mathrm{E}-07$ | 0 | $1.34 \mathrm{E}-02$ | 0 | 0 | $2.96 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Used | 0 | 0 | 0 | $2.68 \mathrm{E}-07$ | 0 | $1.34 \mathrm{E}-02$ | 0 | 0 | $2.96 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
|  | DIN | PRO | JEL | SQU | LZP | SZP | PB | BB | DC | DL | DR | Dcsed | Dlsed | Drsed |  |  |
| Original | 0 | 0 | 0 | $7.23 \mathrm{E}-02$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |  |  |
| Used | 0 | 0 | 0 | 7.23E-02 | 0 | 0.1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |  |  |



Figure C.3: Diagnostic plots commonly referred to when calibrating an Atlantis model include biomass, density, residual nitrogen (i.e., meat, fat), and structural nitrogen (i.e., bone). This figure displays these diagnostic plots for white marlin and blue marlin functional groups for the edited version of the Gulf of Mexico Atlantis Model provided for this study. Images with a solid, black line indicate trends for the entire population, while multi-colored lines indicate trends for each of the 10 cohorts (colors are a gradient, with red indicating cohort 1 and blue indicating cohort 10).

Table C.3: Comparing Simulated Catch to Landings Data, 2010-2011. This data does not include bycatch data.

| Functional Group | Catch [Data] |  | Catch [Model] |  |
| :---: | :---: | :---: | :---: | :---: |
|  | 2010 | 2011 | 2010 | 2011 |
| GAG | 869.2541834 | 424.2282683 | 2717.601963 | 135.328075 |
| RGR | 11290.65301 | 10121.77972 | 10355.35638 | 855.230868 |
| SCM | 106.6226963 | 94.49047107 | 430.10957 | 60.083291 |
| SSR | 1792.514163 | 1661.260256 | 11435.88161 | 7621.159696 |
| DSR | 1868.696426 | 2198.741464 | 7934.120677 | 2910.945576 |
| RSN | 7834.709603 | 7485.57075 | 28111.75431 | 1718.446258 |
| VSN | 2592.430015 | 2665.645146 | 11271.09206 | 960.105507 |
| LUT | 13720.23186 | 9735.783046 | 82790.55164 | 34117.61985 |
| BIO | 281.4140106 | 293.7197801 | 178.205865 | 233.031601 |
| LRF | 2977.391752 | 3064.836201 | 9059.557854 | 3023.253913 |
| SRF | 4409.45074 | 2589.645167 | 23461.92162 | 14362.26674 |
| BDR | 3389.852953 | 3716.062303 | 2410.786693 | 823.796106 |
| RDR | 8248.862435 | 8994.278344 | 6978.791807 | 765.494759 |
| SEA | 5541.172513 | 6454.721288 | 14497.18305 | 10304.0511 |
| SCI | 1102.736522 | 1206.850755 | 23742.39079 | 31925.21771 |
| LDY | 812.1860073 | 566.0487624 | 13743.51592 | 8662.761084 |
| MUL | 12768.28809 | 14392.26876 | 29209.08988 | 18324.90213 |
| POM | 1104.983016 | 793.9470391 | 11626.27994 | 8623.964201 |
| SHP | 2762.648313 | 4226.622449 | 43213.71453 | 14866.45148 |
| SNK | 4946.304571 | 3530.981242 | 24134.13372 | 18626.22111 |
| FLT | 1008.066502 | 1158.880923 | 22315.96577 | 22307.55464 |
| ODF | 77240.27515 | 145173.1183 | 56759.784 | 23882.56042 |
| SDF | 3439.940771 | 2463.022424 | 11064.00351 | 29419.49274 |
| YTN | 2086.957513 | 2950.324009 | 3219.066524 | 1085.615481 |
| BTN | 578.271376 | 393.0733991 | 521.231866 | 27.01504 |
| LTN | 1823.500762 | 1303.117893 | 11301.83941 | 4189.151629 |
| OTN | 0 | 0 | 0 | 0 |
| SWD | 763.6312954 | 713.6085877 | 35572.25075 | 1784.454045 |
| WMR | 556.2137959 | 387.2352518 | 2613.00091 | 702.026704 |
| BMR | 573.5365847 | 398.9554236 | 5902.384574 | 610.744249 |
| BIL | 556.0498356 | 395.9402602 | 2110.671917 | 290.050668 |
| AMB | 1672.649993 | 1191.170086 | 1315.285888 | 107.521184 |
| JCK | 21037.69288 | 14666.34909 | 5635.362462 | 2791.627347 |
| KMK | 4667.319921 | 4272.613195 | 16345.66585 | 906.667556 |
| SMK | 6597.037188 | 5420.938599 | 8862.800034 | 2253.458296 |
| SAR | 1505.724651 | 416.941936 | 42940.34878 | 84499.3485 |
| LPL | 16361.29752 | 23943.53414 | 22701.33933 | 24977.71248 |
| DWF | 0 | 0 | 0 | 0 |
| MEN | 438694.0209 | 623487.1235 | 337037.1668 | 253451.2215 |
| PIN | 1600.016487 | 1157.607798 | 25572.70535 | 17976.11752 |
| MPL | 1886.170133 | 1932.425816 | 18682.85154 | 14242.08553 |
| SPL | 4295.263388 | 3764.48995 | 109872.9597 | 129306.907 |
| TIP | 2352.602061 | 1677.982504 | 3747.665649 | 4264.491933 |
| BEN | 571.476492 | 391.2722242 | 8287.027566 | 9297.838878 |
| LGS | 7682.870588 | 3917.657732 | 55627.02485 | 40864.83637 |
| FIL | 0 | 0 | 0 | 0 |
| SMS | 597.7221394 | 407.9341713 | 12291.96878 | 9877.290105 |
| RAY | 3945.50903 | 4292.319974 | 42096.43535 | 32712.62274 |
| BSH | 54358.80095 | 71836.5773 | 208778.7873 | 38726.54188 |
| WSH | 43483.0138 | 42841.01804 | 91288.34858 | 41194.00024 |
| PSH | 5969.19728 | 4981.224216 | 11606.95687 | 4051.569203 |
| OSH | 1994.476757 | 2501.451361 | 98439.45989 | 16744.92404 |
| DBR | 0 | 0 | 0 | 0 |
| SBR | 0 | 0 | 0 | 0 |
| MAN | 0 | 0 | 0 | 0 |
| MYS | 0 | 0 | 0 | 0 |
| DOL | 0 | 0 | 0 | 0 |
| DDO | 0 | 0 | 0 | 0 |
| LOG | 0 | 0 | 0 | 0 |
| KMP | 0 | 0 | 0 | 0 |
| TUR | 0 | 0 | 0 | 0 |
| BCR | 30657.36079 | 35183.043 | 230943.2357 | 318.445866 |
| SCR | 3120.00048 | 3096.57996 | 150953.5078 | 12841.43481 |
| LOB | 4618.402816 | 4401.223353 | 74373.41795 | 1655.435612 |
| SPG | 100.8752321 | 46.65601953 |  |  |
| CMB | 7535.127658 | 5222.56307 | 233252.7871 | 48234.29739 |
| OYS | 9610.124849 | 10343.27108 | 221967.8541 | 204935.8659 |
| BIV | 1964.155394 | 1974.751522 | 129823.2984 | 130481.7972 |
| SES | 0 | 0 | 0 | 0 |
| SQU | 22502.49823 | 26117.92085 | 2753883.016 | 264619.8827 |

Table C.4: Updated Fishing Mortality Matrix for Gulf of Mexico Atlantis Model.

|  | SprtEst | GillnetEst | TwlShpEst | OytEst | PotCrbEst | SprtShf | TwlShpShf | PotCrbShf |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| mFC_GAG | 0 | 0 | 0 | 0 | 0 | $9.58 \mathrm{E}-04$ | 0 | 0 |
| mFC_RGR | 0 | 0 | 0 | 0 | 0 | $1.28 \mathrm{E}-02$ | 0 | $1.53 \mathrm{E}-05$ |
| mFC_SCM | 0 | 0 | 0 | 0 | 0 | $2.69 \mathrm{E}-04$ | 0 | 0 |
| mFC_SSR | 0 | 0 | 0 | 0 | 0 | $1.23 \mathrm{E}-05$ | 0 | $1.44 \mathrm{E}-06$ |
| mFC_DSR | 0 | 0 | 0 | 0 | $6.86 \mathrm{E}-06$ | $6.96 \mathrm{E}-04$ | 0 | $1.97 \mathrm{E}-04$ |
| mFC_RSN | 0 | 0 | 0 | 0 | 0 | $3.92 \mathrm{E}-04$ | $5.12 \mathrm{E}-05$ | 0 |
| mFC_VSN | 0 | 0 | 0 | 0 | 0 | $1.94 \mathrm{E}-04$ | $2.61 \mathrm{E}-05$ | 0 |
| mFC_LUT | 0 | $1.01 \mathrm{E}-09$ | 0 | 0 | 0 | $5.54 \mathrm{E}-04$ | $1.68 \mathrm{E}-05$ | $2.55 \mathrm{E}-07$ |
| mFC _BIO | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LRF | 0 | $3.34 \mathrm{E}-07$ | $2.30 \mathrm{E}-06$ | 0 | 0 | $1.88 \mathrm{E}-04$ | $6.00 \mathrm{E}-07$ | $1.81 \mathrm{E}-06$ |
| mFC_SRF | 0 | $3.87 \mathrm{E}-07$ | 0 | 0 | 0 | $3.04 \mathrm{E}-04$ | $3.07 \mathrm{E}-06$ | 0 |
| mFC_BDR | 1.009228641 | $4.67 \mathrm{E}-02$ | $3.48 \mathrm{E}-02$ | 0 | $2.68 \mathrm{E}-02$ | 0 | 0.306036876 | 0 |
| mFC_RDR | 2.125756414 | 0 | 0.183536107 | 0 | 0 | 0 | 0 | 0 |
| mFC_SEA | $3.30 \mathrm{E}-03$ | $2.21 \mathrm{E}-05$ | $1.30 \mathrm{E}-02$ | 0 | 0 | 0 | $2.11 \mathrm{E}-05$ | 0 |
| mFC_SCI | $2.43 \mathrm{E}-05$ | $1.39 \mathrm{E}-05$ | $1.57 \mathrm{E}-03$ | 0 | 7.63E-08 | 0 | $1.59 \mathrm{E}-03$ | 0 |
| mFC_LDY | $6.40 \mathrm{E}-05$ | $1.21 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MUL | $2.03 \mathrm{E}-04$ | $2.95 \mathrm{E}-04$ | 0 | 0 | $9.55 \mathrm{E}-08$ | 0 | $3.19 \mathrm{E}-07$ | 0 |
| mFC_POM | $6.03 \mathrm{E}-06$ | $4.24 \mathrm{E}-06$ | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SHP | $2.63 \mathrm{E}-04$ | $3.51 \mathrm{E}-06$ | $6.00 \mathrm{E}-06$ | 0 | $1.08 \mathrm{E}-07$ | 0 | $7.89 \mathrm{E}-06$ | 0 |
| mFC_SNK | $5.27 \mathrm{E}-09$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_FLT | 0 | $1.82 \mathrm{E}-05$ | $8.48 \mathrm{E}-05$ | 0 | $1.51 \mathrm{E}-06$ | $3.26 \mathrm{E}-05$ | $6.15 \mathrm{E}-06$ | $1.41 \mathrm{E}-06$ |
| mFC_ODF | 0 | $1.24 \mathrm{E}-04$ | $6.81 \mathrm{E}-02$ | 0 | $1.80 \mathrm{E}-04$ | 2.688466653 | $6.81 \mathrm{E}-02$ | $6.57 \mathrm{E}-03$ |
| mFC_SDF | 0 | 0 | $1.09 \mathrm{E}-04$ | 0 | 0 | $2.81 \mathrm{E}-08$ | 0 | 0 |
| mFC_YTN | 0 | 0 | 0 | 0 | 0 | $1.42 \mathrm{E}-03$ | 0 | 0 |
| mFC_BTN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LTN | 0 | $1.79 \mathrm{E}-06$ | 0 | 0 | 0 | $1.53 \mathrm{E}-04$ | 0 | 0 |
| mFC_OTN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SWD | 0 | 0 | 0 | 0 | 0 | $2.96 \mathrm{E}-06$ | 0 | 0 |
| mFC_WMR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_BMR | 0 | 0 | 0 | 0 | 0 | $9.63 \mathrm{E}-06$ | 0 | 0 |
| mFC_BIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC _AMB | 0 | 0 | 0 | 0 | 0 | 0.991878998 | 0 | 0 |
| mFC_JCK | 0 | $1.93 \mathrm{E}-02$ | 0 | 0 | $9.10 \mathrm{E}-05$ | 1.757864834 | 0 | $4.74 \mathrm{E}-03$ |
| mFC_KMK | 0 | 0 | 0 | 0 | 0 | $4.70 \mathrm{E}-04$ | $8.78 \mathrm{E}-06$ | 0 |
| mFC_SMK | 0 | $1.36 \mathrm{E}-03$ | $6.29 \mathrm{E}-04$ | 0 | 0 | $4.95 \mathrm{E}-03$ | 0 | 0 |
| mFC_SAR | 0 | 0 | 0 | 0 | 0 | $1.17 \mathrm{E}-07$ | 0 | 0 |
| mFC_LPL | 0 | $5.61 \mathrm{E}-05$ | $1.06 \mathrm{E}-05$ | 0 | 0 | $5.08 \mathrm{E}-03$ | 0 | 0 |
| mFC_DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MEN | 0 | $2.26 \mathrm{E}-05$ | 0 | 0 | 0 | $1.27 \mathrm{E}-05$ | 0 | 0 |
| mFC_PIN | 0 | 0 | 0 | 0 | 0 | $1.52 \mathrm{E}-04$ | 0 | $1.35 \mathrm{E}-05$ |
| mFC_MPL | 0 | 0 | 0 | 0 | $5.51 \mathrm{E}-08$ | $1.71 \mathrm{E}-07$ | 0 | 0 |
| mFC_SPL | 0 | $1.43 \mathrm{E}-08$ | 0 | 0 | 0 | $5.01 \mathrm{E}-06$ | 0 | 0 |
| mFC_TIP | 0 | 0 | 0 | 0 | 0 | $6.27 \mathrm{E}-04$ | 0 | 0 |
| mFC_BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LGS | 0 | $1.66 \mathrm{E}-05$ | $2.39 \mathrm{E}-05$ | 0 | 0 | $7.41 \mathrm{E}-05$ | $2.39 \mathrm{E}-05$ | 0 |
| mFC_FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_RAY | $1.18 \mathrm{E}-06$ | $3.89 \mathrm{E}-07$ | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_BSH | 0 | 0 | $1.18 \mathrm{E}-04$ | 0 | 0 | 0 | $4.92 \mathrm{E}-04$ | 0 |
| mFC_WSH | 0 | 0 | $1.11 \mathrm{E}-03$ | 0 | 0 | 0 | $2.62 \mathrm{E}-03$ | 0 |
| mFC_PSH | 0 | 0 | $4.09 \mathrm{E}-06$ | 0 | 0 | 0 | $2.44 \mathrm{E}-03$ | 0 |
| mFC_OSH | 0 | 0 | $2.85 \mathrm{E}-05$ | 0 | 0 | 0 | $5.71 \mathrm{E}-05$ | $1.23 \mathrm{E}-09$ |
| mFC_DBR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SBR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MAN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MYS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DOL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DDO | 0 | 0 | 0 | 0 | 0 | 0 |  | 0 |
| mFC_LOG | 0 | 0 | 0 | 0 | 0 | 0 | $4.30 \mathrm{E}-05$ | 0 |
| mFC_KMP | 0 | 0 | 0 | 0 | 0 | 0 | $4.13 \mathrm{E}-06$ | 0 |
| mFC_TUR | 0 | 0 | 0 | 0 | 0 | 0 | $5.98 \mathrm{E}-07$ | 0 |
| mFC_BCR | 0 | 0 | $9.91 \mathrm{E}-07$ | 0 | $4.79 \mathrm{E}-04$ | 0 | $1.00 \mathrm{E}-07$ | $1.48 \mathrm{E}-07$ |
| mFC_SCR | 0 | 0 | 0 | 0 | $6.01 \mathrm{E}-05$ | 0 | 0 | 0 |
| mFC_LOB | 0 | 0 | 0 | 0 | $6.57 \mathrm{E}-08$ | $5.67 \mathrm{E}-05$ | 7.16E-09 | 0 |
| mFC_COR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_CCA | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_OCT | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SPG | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_CMB | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_INF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_ECH | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_OYS | 0 | 0 | 0 | $2.12 \mathrm{E}-06$ | 0 | 0 | 0 | 0 |
| mFC_BIV | 0 | $2.31 \mathrm{E}-10$ | $2.00 \mathrm{E}-10$ | 1.62E-10 | $5.51 \mathrm{E}-11$ | 0 | $7.20 \mathrm{E}-08$ | $2.84 \mathrm{E}-09$ |
| mFC_SES | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_EPI | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_GRS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_ALG | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MPB | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LPP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SPP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DIN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_PRO | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_JEL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SQU | 0 | 0 | $7.80 \mathrm{E}-09$ | 0 | $3.36 \mathrm{E}-09$ | 0 | $1.26 \mathrm{E}-08$ | $4.10 \mathrm{E}-09$ |
| mFC_LZP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SZP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC _PB | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC _BB | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DC | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |

Table C.4: (Continued)

|  | PotLbtShf | HLReefShf | LLReefShf | SeineMenShf | LLShkShf | LLPelgc | RoyalRed | OtherUS |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| mFC_GAG | 0 | 3.95E-04 | $1.44 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | $4.93 \mathrm{E}-05$ |
| mFC_RGR | 0 | 0 | 0.109742601 | 0 | 0 | 0 | 0 | $1.00 \mathrm{E}-03$ |
| mFC_SCM | 0 | $3.55 \mathrm{E}-04$ | $3.74 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | $9.94 \mathrm{E}-06$ |
| mFC -SSR | $1.56 \mathrm{E}-07$ | $7.56 \mathrm{E}-05$ | $4.04 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | $2.63 \mathrm{E}-05$ |
| mFC_DSR | 0 | $6.43 \mathrm{E}-05$ | $2.61 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | $5.41 \mathrm{E}-08$ |
| mFC_RSN | 0 | $3.72 \mathrm{E}-04$ | $1.26 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | $2.57 \mathrm{E}-06$ |
| mFC_VSN | 0 | $8.87 \mathrm{E}-04$ | $1.55 \mathrm{E}-06$ | 0 | 0 | 0 | 0 | $6.96 \mathrm{E}-07$ |
| mFC_LUT | $1.03 \mathrm{E}-07$ | $9.22 \mathrm{E}-05$ | $6.54 \mathrm{E}-06$ | 0 | 0 | 0 | 0 | $1.26 \mathrm{E}-06$ |
| mFC_BIO | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LRF | $1.10 \mathrm{E}-06$ | $4.94 \mathrm{E}-05$ | $6.37 \mathrm{E}-06$ | 0 | 0 | 0 | 0 | $8.93 \mathrm{E}-05$ |
| mFC_SRF | 0 | $9.27 \mathrm{E}-05$ | $2.28 \mathrm{E}-04$ | 0 | 0 | $1.13 \mathrm{E}-06$ | 0 | $6.98 \mathrm{E}-06$ |
| mFC_BDR | 0 | 0.109658285 | 1.459713061 | 0 | 0 | $3.76 \mathrm{E}-02$ | 0 | $8.17 \mathrm{E}-03$ |
| mFC_RDR | 0 | $4.88 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SEA | 0 | $1.81 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 | $6.46 \mathrm{E}-06$ |
| mFC_SCI | 0 | $4.16 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | 0 | $1.69 \mathrm{E}-05$ |
| mFC_LDY | 0 | $7.46 \mathrm{E}-06$ | 0 | 0 | 0 | 0 | 0 | $4.75 \mathrm{E}-05$ |
| mFC_MUL | 0 | $4.98 \mathrm{E}-07$ | 0 | $9.37 \mathrm{E}-06$ | 0 | 0 | 0 | $1.36 \mathrm{E}-03$ |
| mFC_POM | 0 | $1.00 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | 0 | $3.84 \mathrm{E}-06$ |
| mFC_SHP | 0 | $1.82 \mathrm{E}-05$ | $1.67 \mathrm{E}-06$ | $1.10 \mathrm{E}-08$ | 0 | 0 | 0 | $7.52 \mathrm{E}-06$ |
| mFC_SNK | 0 | $0$ | $0$ | 0 | 0 | 0 | 0 | $0$ |
| $\mathrm{mFC} \text { _FLT }$ | 0 | $3.67 \mathrm{E}-06$ | $3.54 \mathrm{E}-06$ | 0 | 0 | $1.58 \mathrm{E}-06$ | 0 | $3.08 \mathrm{E}-05$ |
| $\mathrm{mFC} O D F$ | $1.79 \mathrm{E}-05$ | 0.01939481 | $8.53 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | $1.30 \mathrm{E}-04$ |
| mFC.SDF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_YTN | 0 | $2.75 \mathrm{E}-05$ | 0 | 0 | 0 | $2.27 \mathrm{E}-03$ | 0 | 0 |
| mFC_BTN | 0 | 0 | 0 | 0 | 0 | $9.36 \mathrm{E}-03$ | 0 | 0 |
| mFC_LTN | 0 | $1.88 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_OTN | 0 | 0 | 0 | 0 | 0 | $8.50 \mathrm{E}-05$ | 0 | 0 |
| mFC_SWD | 0 | $4.68 \mathrm{E}-06$ | 0 | 0 | 0 | $1.83 \mathrm{E}-04$ | 0 | 0 |
| mFC_WMR | 0 | 0 | 0 | 0 | 0 | $3.53 \mathrm{E}-05$ | 0 | 0 |
| mFC_BMR | 0 | 0 | 0 | 0 | 0 | $4.74 \mathrm{E}-05$ | 0 | 0 |
| mFC_BIL | 0 | 0 | 0 | 0 | 0 | $1.82 \mathrm{E}-05$ | 0 | 0 |
| mFC_AMB | 0 | 0.85650951 | $1.26 \mathrm{E}-02$ | 0 | 0 | 0 | 0 | $2.19 \mathrm{E}-02$ |
| mFC_JCK | 0 | 0.261193261 | $6.24 \mathrm{E}-03$ | 0 | 0 | 0 | 0 | 0.11331893 |
| mFC_KMK | 0 | $5.26 \mathrm{E}-04$ | 0 | 0 | 0 | $1.38 \mathrm{E}-04$ | 0 |  |
| mFC_SMK | 0 | $1.69 \mathrm{E}-04$ | 0 | $3.16 \mathrm{E}-06$ | 0 | $1.02 \mathrm{E}-04$ | 0 | $4.50 \mathrm{E}-05$ |
| mFC_SAR | 0 | $1.42 \mathrm{E}-07$ | 0 | 0 | 0 | 0 | 0 | $1.47 \mathrm{E}-06$ |
| mFC_LPL | 0 | $1.18 \mathrm{E}-04$ | $4.50 \mathrm{E}-06$ | $4.61 \mathrm{E}-06$ | 0 | $9.99 \mathrm{E}-05$ | 0 | $2.24 \mathrm{E}-06$ |
| mFC_DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MEN | 0 | $1.22 \mathrm{E}-08$ | 0 | 0.10065426 | 0 | 0 | 0 | $1.93 \mathrm{E}-05$ |
| mFC_PIN | 0 | $6.20 \mathrm{E}-06$ | 0 | 0 | 0 | 0 | 0 | $3.25 \mathrm{E}-06$ |
| mFC_MPL | 0 | $1.97 \mathrm{E}-05$ | 0 | 0 | 0 | $1.75 \mathrm{E}-07$ | 0 | $4.33 \mathrm{E}-05$ |
| mFC_SPL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $4.72 \mathrm{E}-05$ |
| mFC_TIP | 0 | $1.76 \mathrm{E}-03$ | 0 | 0 | $1.55 \mathrm{E}-04$ | 0 | 0 | 0 |
| mFC_BEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LGS | 0 | $1.26 \mathrm{E}-05$ | $1.81 \mathrm{E}-06$ | $1.50 \mathrm{E}-08$ | $1.42 \mathrm{E}-05$ | $1.31 \mathrm{E}-05$ | 0 | $7.73 \mathrm{E}-08$ |
| mFC_FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SMS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_RAY | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_BSH | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $2.06 \mathrm{E}-07$ |
| mFC_WSH | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $1.12 \mathrm{E}-06$ |
| $\mathrm{mFC} \text { _PSH }$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $0$ |
| mFC_OSH | 0 | 0 | 0 | 0 | 0 | 0 | $6.17 \mathrm{E}-06$ | $3.85 \mathrm{E}-09$ |
| mFC_DBR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SBR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MAN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MYS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DOL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DDO | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LOG | 0 | 0 | 0 | 0 | 0 | $7.93 \mathrm{E}-06$ | 0 | 0 |
| mFC_KMP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_TUR | 0 | 0 | 0 | 0 | 0 | $9.89 \mathrm{E}-06$ | 0 | 0 |
| mFC-BCR | 0 | $2.25 \mathrm{E}-09$ | $1.21 \mathrm{E}-07$ | 0 | 0 | 0 | 0 | 0 |
| mFC_SCR | $9.71 \mathrm{E}-07$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LOB | $2.33 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 | 0 | $8.07 \mathrm{E}-06$ |
| $\mathrm{mFC} C O R$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| $\mathrm{mFC} \text { _CCA }$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC -OCT | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SPG | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_CMB | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_INF | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_ECH | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_OYS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $7.16 \mathrm{E}-08$ |
| mFC_BIV | $3.25 \mathrm{E}-10$ | $1.89 \mathrm{E}-11$ | $3.09 \mathrm{E}-11$ | 0 | 0 | $2.26 \mathrm{E}-11$ | 0 | $9.91 \mathrm{E}-10$ |
| mFC-SES | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_EPI | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_GRS | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_ALG | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MPB | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LPP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC.SPP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DIN | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| $\mathrm{mFC} \text { _PRO }$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_JEL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SQU | 0 | $1.43 \mathrm{E}-09$ | 0 | 0 | 0 | 0 | 0 | $3.66 \mathrm{E}-11$ |
| mFC_LZP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC -SZP | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC -PB | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC - $\mathrm{BB}^{\text {a }}$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DC | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DL | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DR | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |

Table C.4: (Continued)

|  | TwlShpMX | LLReefMX | LLShkMX | GillnetMackMX | OctpsMX | MixedMX | MixedCB |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| mFC_GAG | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_RGR | 0 | 0.161361716 | 0 | 0 | 0 | 0.161361716 | 0 |
| mFC_SCM | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SSR | 0 | $2.35 \mathrm{E}-04$ | 0 | 0 | 0 | $2.35 \mathrm{E}-04$ | $3.49 \mathrm{E}-04$ |
| mFC_DSR | 0 | $6.48 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 |
| mFC_RSN | 0 | $4.81 \mathrm{E}-04$ | 0 | 0 | 0 | $4.81 \mathrm{E}-04$ | $6.35 \mathrm{E}-05$ |
| mFC_VSN | 0 | $3.11 \mathrm{E}-04$ | 0 | 0 | 0 | $3.11 \mathrm{E}-04$ | 0 |
| mFC_LUT | 0 | $1.25 \mathrm{E}-04$ | 0 | 0 | 0 | $1.25 \mathrm{E}-04$ | $3.23 \mathrm{E}-05$ |
| mFC_BIO | 0 | 0 | 0 | 0 | 0 | 0 | $1.32 \mathrm{E}-02$ |
| mFC_LRF | 0 | 0 | 0 | 0 | 0 | $2.58 \mathrm{E}-03$ | $3.09 \mathrm{E}-04$ |
| mFC_SRF | 0 | 0 | 0 | 0 | 0 | $5.44 \mathrm{E}-04$ | $1.51 \mathrm{E}-04$ |
| mFC_BDR | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_RDR | 0 | 0 | 0 | 0 | 0 | 0.275696236 | 0 |
| mFC_SEA | 0 | 0 | 0 | 0 | 0 | $3.09 \mathrm{E}-03$ | 0 |
| mFC _SCI | 0 | 0 | 0 | 0 | 0 | $9.02 \mathrm{E}-05$ | 0 |
| mFC_LDY | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MUL | 0 | 0 | 0 | $9.08 \mathrm{E}-04$ | 0 | $9.08 \mathrm{E}-04$ | $2.20 \mathrm{E}-05$ |
| mFC_POM | 0 | 0 | 0 | 0 | 0 | $4.09 \mathrm{E}-04$ | 0 |
| mFC_SHP | $9.97 \mathrm{E}-06$ | 0 | 0 | $9.97 \mathrm{E}-06$ | 0 | $9.97 \mathrm{E}-06$ | 0 |
| mFC_SNK | 0 | 0 | 0 | 0 | 0 | $2.04 \mathrm{E}-03$ | 0 |
| mFC_FLT | $2.74 \mathrm{E}-05$ | 0 | 0 | $2.74 \mathrm{E}-05$ | 0 | $2.74 \mathrm{E}-05$ | 0 |
| mFC_ODF | 0 | 0 | 0 | 0 | 0 | $7.50 \mathrm{E}-02$ | $3.56 \mathrm{E}-03$ |
| mFC_SDF | 0 | 0 | 0 | 0 | 0 | $2.84 \mathrm{E}-03$ | 0 |
| mFC_YTN | 0 | 0 | $3.19 \mathrm{E}-03$ | 0 | 0 | $3.19 \mathrm{E}-03$ | $2.39 \mathrm{E}-06$ |
| mFC_BTN | 0 | 0 | $1.74 \mathrm{E}-02$ | 0 | 0 | $1.74 \mathrm{E}-02$ | 0 |
| mFC_LTN | 0 | 0 | $4.11 \mathrm{E}-04$ | 0 | 0 | $4.11 \mathrm{E}-04$ | $6.19 \mathrm{E}-06$ |
| mFC_OTN | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SWD | 0 | 0 | $8.07 \mathrm{E}-05$ | 0 | 0 | $8.07 \mathrm{E}-05$ | $1.25 \mathrm{E}-07$ |
| mFC_WMR | 0 | 0 | $6.64 \mathrm{E}-04$ | 0 | 0 | $6.64 \mathrm{E}-04$ | 0 |
| mFC CMMR | 0 | 0 | $4.45 \mathrm{E}-04$ | 0 | 0 | $4.45 \mathrm{E}-04$ | $4.06 \mathrm{E}-06$ |
| mFC_BIL | 0 | 0 | $1.16 \mathrm{E}-03$ | 0 | 0 | $1.16 \mathrm{E}-03$ | $1.28 \mathrm{E}-04$ |
| mFC_AMB | 0 | 0 | 0.431905909 | 0 | 0 | 0.431905909 | 0 |
| mFC_JCK | 0 | 0 | 3.276053863 | 0 | 0 | 3.276053863 | $3.76 \mathrm{E}-03$ |
| mFC_KMK | 0 | 0 | $6.07 \mathrm{E}-04$ | $6.07 \mathrm{E}-04$ | 0 | 0 | $2.10 \mathrm{E}-07$ |
| mFC_SMK | 0 | 0 | $3.58 \mathrm{E}-03$ | $3.58 \mathrm{E}-03$ | 0 | 0 | $9.73 \mathrm{E}-07$ |
| mFC_SAR | 0 | 0 | 0 | 0 | 0 | $1.24 \mathrm{E}-04$ | 0 |
| mFC_LPL | 0 | 0 | 0 | $1.58 \mathrm{E}-03$ | 0 | 0 | $7.78 \mathrm{E}-05$ |
| mFC_DWF | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MEN | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_PIN | 0 | 0 | 0 | 0 | 0 | $8.35 \mathrm{E}-05$ | 0 |
| mFC_MPL | 0 | 0 | 0 | 0 | 0 | $6.72 \mathrm{E}-04$ | $1.91 \mathrm{E}-06$ |
| mFC_SPL | 0 | 0 | 0 | 0 | 0 | $3.47 \mathrm{E}-05$ | $2.66 \mathrm{E}-05$ |
| mFC_TIP | 0 | 0 | $1.15 \mathrm{E}-02$ | 0 | 0 | 0 | 0 |
| mFC_BEN | $2.73 \mathrm{E}-06$ | 0 | $2.73 \mathrm{E}-06$ | 0 | 0 | $2.73 \mathrm{E}-06$ | 0 |
| mFC_LGS | 0 | 0 | $1.39 \mathrm{E}-04$ | 0 | 0 | $1.39 \mathrm{E}-04$ | $5.58 \mathrm{E}-06$ |
| mFC_FIL | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SMS | $1.28 \mathrm{E}-05$ | 0 | $1.28 \mathrm{E}-05$ | $1.28 \mathrm{E}-05$ | 0 | $1.28 \mathrm{E}-05$ |  |
| mFC_RAY | $1.51 \mathrm{E}-04$ | 0 | $1.51 \mathrm{E}-04$ | 0 | 0 | $1.51 \mathrm{E}-04$ | $7.60 \mathrm{E}-05$ |
| mFC_BSH | $1.96 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_WSH | $5.08 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | $5.08 \mathrm{E}-05$ | 0 |
| mFC_PSH | $7.22 \mathrm{E}-04$ | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_OSH | $1.98 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | $1.98 \mathrm{E}-05$ | 0 |
| mFC_DBR | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SBR | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MAN | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_MYS | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DOL | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DDO | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LOG | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_KMP | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_TUR | 0 | 0 | 0 | 0 | 0 | 0 |  |
| mFC -BCR | 0 | 0 | 0 | 0 | 0 | $1.82 \mathrm{E}-04$ | $1.10 \mathrm{E}-06$ |
| mFC_SCR | 0 | 0 | 0 | 0 | 0 | $1.42 \mathrm{E}-05$ | 0 |
| mFC_LOB | 0 | 0 | 0 | 0 | 0 | $6.16 \mathrm{E}-05$ | 7.62E-05 |
| mFC -COR | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_CCA | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_OCT | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SPG | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_CMB | $3.00 \mathrm{E}-05$ | 0 | 0 | 0 | 0 | $3.00 \mathrm{E}-05$ | $2.16 \mathrm{E}-06$ |
| mFC_INF | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_ECH | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_OYS | 0 | 0 | 0 | 0 | 0 | $4.94 \mathrm{E}-07$ | 0 |
| mFC_BIV | 0 | 0 | 0 | 0 | 0 | $1.84 \mathrm{E}-06$ | $6.95 \mathrm{E}-08$ |
| mFC_SES | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_EPI | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_GRS | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC _ALG | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC _MPB | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_LPP | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SPP | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DIN | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_PRO | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_JEL | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SQU | $1.26 \mathrm{E}-07$ | 0 | 0 | 0 | $1.24 \mathrm{E}-05$ | $1.26 \mathrm{E}-07$ | 0 |
| mFC_LZP | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_SZP | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC _PB | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC - $\mathrm{BB}^{\text {a }}$ | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC -DC | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DL | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| mFC_DR | 0 | 0 | 0 | 0 | 0 | 0 | 0 |

## C. 3 Additional Results

Functional group-specific performance measures were also computed for functional groups not identified in the main text, as well as functional group assemblages not identified in the main text. This allowed some investigation into indirect impacts from pelagic longline fisheries. There was a lot of information, and all of it could not be discussed in the main text. Some examples of additional biomass and catch metrics are presented in Table C.5, and addition results for average weight and proportion mature metrics are presented in Table C.6.
Table C.5: Additional biomass and catch performance metrics. Metrics are relative to the status quo. Metrics for functional group assemblages (i.e., elasmobranchs and reef fish) are the sum across all associated functional groups. Functional group-specific metrics are shown for the functional groups: large sharks (LGS), blacktip sharks (TIP), king mackerel (KMK), and amberjack

| Scenarios | Biomass |  |  |  |  |  | Catch |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Elasmobranchs | LGS | TIP | KMK | AMB | Reef fish | Elasmobranchs | LGS | TIP | KMK | AMB | Reef fish |
| Status Quo | 1 | 1 |  | 1 | 1 | 1 | - 1 | ${ }^{1}$ | 1 | 1 | 1 | 1 |
| All longlining $\mathrm{F} * 0$ | 1.001 | 1.010 | 1.003 | 1.044 | 0.996 | 1.041 | 0.927 | 0.858 | 0.953 | 0.965 | 0.994 | 0.945 |
| All longlining $\mathrm{F} * 0.5$ | 1.001 | 1.005 | 1.002 | 1.021 | 0.998 | 1.019 | 0.964 | 0.930 | 0.977 | 0.984 | 0.997 | 0.977 |
| All longlining F * 2 | 0.999 | 0.990 | 0.997 | 0.963 | 1.004 | 0.966 | 1.071 | 1.137 | 1.045 | 1.027 | 1.005 | 1.033 |
| Pelagic longlining F * 0 | 1.001 | 1.004 | 1.000 | 1.043 | 1.000 | 1.000 | 0.982 | 0.939 | 1.000 | 0.964 | 1.000 | 1.000 |
| Pelagic longlining $\mathrm{F} * 0.5$ | 1.000 | 1.002 | 1.000 | 1.020 | 1.000 | 1.000 | 0.991 | 0.970 | 1.000 | 0.983 | 1.000 | 1.000 |
| Pelagic longlining F * 2 | 0.999 | 0.996 | 1.000 | 0.963 | 1.000 | 1.000 | 1.018 | 1.060 | 1.000 | 1.028 | 1.000 | 1.000 |
| No DeSoto Canyon | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.002 | 1.005 | 1.000 | 1.001 | 1.000 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 |
| No PLL Spatial Closures | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.002 | 1.005 | 1.000 | 1.001 | 1.000 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.000 | 1.000 | 1.006 | 1.000 | 1.000 | 0.999 | 0.998 | 1.000 | 0.991 | 1.000 | 1.000 |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All longlining F * 0 | 1.001 | 1.010 | 1.003 | 1.027 | 0.996 | 1.051 | 0.924 | 0.877 | 0.952 | 0.995 | 0.994 | 0.939 |
| All longlining $\mathrm{F} * 0.5$ | 1.001 | 1.005 | 1.001 | 1.013 | 0.998 | 1.024 | 0.962 | 0.939 | 0.976 | 0.998 | 0.997 | 0.973 |
| All longlining F * 2 | 0.999 | 0.990 | 0.997 | 0.977 | 1.004 | 0.958 | 1.073 | 1.119 | 1.046 | 1.003 | 1.005 | 1.038 |
| Pelagic longlining F * 0 | 1.001 | 1.004 | 1.000 | 1.026 | 1.000 | 1.000 | 0.985 | 0.960 | 1.000 | 0.993 | 1.000 | 1.000 |
| Pelagic longlining $\mathrm{F} * 0.5$ | 1.000 | 1.002 | 1.000 | 1.013 | 1.000 | 1.000 | 0.992 | 0.980 | 1.000 | 0.997 | 1.000 | 1.000 |
| Pelagic longlining F * 2 | 0.999 | 0.996 | 1.000 | 0.978 | 1.000 | 1.000 | 1.015 | 1.040 | 1.000 | 1.004 | 1.000 | 1.000 |
| No DeSoto Canyon | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.011 | 1.028 | 1.000 | 1.032 | 1.000 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 |
| No PLL Spatial Closures | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.011 | 1.028 | 1.000 | 1.032 | 1.000 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.000 | 1.000 | 1.004 | 1.000 | 1.000 | 0.999 | 0.998 | 1.000 | 0.997 | 1.000 | 1.000 |
| Status Quo | 1 | 1 | 1 | 1 |  | 1 | 1 | 1 | NA | 1 | 1 | 1 |
| All longlining F * 0 | 1.001 | 1.010 | 1.000 | 1.038 | 0.996 | 1.143 | 0.892 | 0.868 | NA | 0.980 | 0.992 | 0.959 |
| All longlining $\mathrm{F} * 0.5$ | 1.000 | 1.005 | 1.000 | 1.018 | 0.998 | 1.068 | 0.947 | 0.935 | NA | 0.991 | 0.996 | 0.986 |
| All longlining F * 2 | 0.999 | 0.991 | 1.000 | 0.967 | 1.004 | 0.885 | 1.104 | 1.128 | NA | 1.015 | 1.008 | 1.000 |
| Pelagic longlining F * 0 | 1.000 | 1.004 | 1.000 | 1.037 | 1.000 | 0.999 | 0.959 | 0.950 | NA | 0.979 | 1.000 | 0.999 |
| Pelagic longlining $\mathrm{F} * 0.5$ | 1.000 | 1.002 | 1.000 | 1.018 | 1.000 | 1.000 | 0.980 | 0.975 | NA | 0.990 | 1.000 | 0.999 |
| Pelagic longlining F $* 2$ | 1.000 | 0.996 | 1.000 | 0.968 | 1.000 | 1.001 | 1.040 | 1.049 | NA | 1.016 | 1.000 | 1.001 |
| No DeSoto Canyon | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.013 | 1.016 | NA | 1.007 | 1.000 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.001 | 1.001 | NA | 1.006 | 1.000 | 1.000 |
| No PLL Spatial Closures | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.014 | 1.017 | NA | 1.014 | 1.000 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.000 | 1.000 | 1.005 | 1.000 | 1.000 | 0.999 | 0.998 | NA | 1.000 | 1.000 | 1.000 |

Table C.6: Additional average individual weight, and proportion mature performance metrics. Metrics are relative to the status quo. Metrics for functional group assemblages (i.e., elasmobranchs and reef fish) are the sum across all associated functional groups. Functional group-specific metrics are shown for the functional groups: large sharks (LGS), blacktip sharks (TIP), king mackerel (KMK), and amberjack (AMB).

|  | Average Individual Weight [Biomass] |  |  |  |  |  | Proportion Mature [Biomass] |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Scenarios | Elasmobranchs | LGS | TIP | KMK | AMB | Reef fish | Elasmobranchs | LGS | TIP | KMK | AMB | Reef fish |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 7 | 1 | 1 | 1 | 1 | 1 | 1 |
| All longlining $\mathrm{F} * 0$ | 1.001 | 1.000 | 1.002 | 1.027 | 0.999 | 1.027 | 0.999 | 1.002 | 1.002 | 1.017 | 1.000 | 1.007 |
| All longlining F * 0.5 | 1.000 | 1.000 | 1.001 | 1.013 | 1.000 | 1.013 | 1.000 | 1.001 | 1.001 | 1.009 | 1.000 | 1.003 |
| All longlining F * 2 | 0.999 | 1.000 | 0.998 | 0.977 | 1.001 | 0.978 | 1.001 | 0.998 | 0.998 | 0.984 | 1.000 | 0.994 |
| Pelagic longlining F * 0 | 1.000 | 1.000 | 1.000 | 1.026 | 1.000 | 1.000 | 1.000 | 1.001 | 1.000 | 1.017 | 1.000 | 1.000 |
| Pelagic longlining F * 0.5 | 1.000 | 1.000 | 1.000 | 1.012 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.008 | 1.000 | 1.000 |
| Pelagic longlining F $* 2$ | 1.000 | 1.000 | 1.000 | 0.978 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 0.984 | 1.000 | 1.000 |
| No DeSoto Canyon | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 |
| No PLL Spatial Closures | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.000 | 1.000 | 1.004 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.003 | 1.000 | 1.000 |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| All longlining F * 0 | 1.001 | 1.000 | 1.002 | 1.018 | 0.999 | 1.029 | 0.999 | 1.002 | 1.002 | 1.017 | 1.000 | 1.005 |
| All longlining $\mathrm{F} * 0.5$ | 1.000 | 1.000 | 1.001 | 1.009 | 1.000 | 1.014 | 1.000 | 1.001 | 1.001 | 1.008 | 1.000 | 1.003 |
| All longlining F * 2 | 0.999 | 1.000 | 0.998 | 0.985 | 1.001 | 0.976 | 1.001 | 0.998 | 0.998 | 0.985 | 1.000 | 0.995 |
| Pelagic longlining F * 0 | 1.000 | 1.000 | 1.000 | 1.017 | 1.000 | 1.000 | 1.000 | 1.001 | 1.000 | 1.016 | 1.000 | 1.000 |
| Pelagic longlining F * 0.5 | 1.000 | 1.000 | 1.000 | 1.008 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.008 | 1.000 | 1.000 |
| Pelagic longlining F * 2 | 1.000 | 1.000 | 1.000 | 0.985 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 0.985 | 1.000 | 1.000 |
| No DeSoto Canyon | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 |
| No PLL Spatial Closures | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.000 | 1.000 | 1.002 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.002 | 1.000 | 1.000 |
| Status Quo | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | - | 1 |  | 1 |
| All longlining F * 0 | 1.000 | 1.000 | 1.000 | 1.024 | 0.999 | 1.115 | 1.000 | 1.000 | 1.000 | 1.017 | 1.000 | 1.023 |
| All longlining $\mathrm{F} * 0.5$ | 1.000 | 1.000 | 1.000 | 1.011 | 1.000 | 1.055 | 1.000 | 1.000 | 1.000 | 1.008 | 1.000 | 1.011 |
| All longlining $\mathrm{F} * 2$ | 1.000 | 1.000 | 1.000 | 0.980 | 1.001 | 0.904 | 1.000 | 1.000 | 1.000 | 0.984 | 1.000 | 0.977 |
| Pelagic longlining $\mathrm{F} * 0$ | 1.000 | 1.000 | 1.000 | 1.023 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.017 | 1.000 | 1.000 |
| Pelagic longlining $\mathrm{F} * 0.5$ | 1.000 | 1.000 | 1.000 | 1.011 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.008 | 1.000 | 1.000 |
| Pelagic longlining F * 2 | 1.000 | 1.000 | 1.000 | 0.980 | 1.000 | 1.001 | 1.000 | 1.000 | 1.000 | 0.984 | 1.000 | 1.000 |
| No DeSoto Canyon | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 |
| No Spring Closure | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 |
| No PLL Spatial Closures | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 0.999 | 1.000 | 1.000 |
| Seasonal PLL Closure | 1.000 | 1.000 | 1.000 | 1.003 | 1.000 | 1.000 | 1.000 | 1.000 | 1.000 | 1.003 | 1.000 | 1.000 |

## Bibliography

Abascal, F., J. Mejuto, M. Quintans, B. García-Cortés, and A. Ramos-Cartelle. 2015. Tracking of the broadbill swordfish, Xiphias gladius, in the central and eastern North Atlantic. Fisheries Research, 162:20-28.

Adams, C. M., E. Hernandez, and J. C. Cato. 2004. The economic significance of the Gulf of Mexico related to population, income, employment, minerals, fisheries and shipping. Ocean $\mathcal{E}$ Coastal Management, 47(11):565-580.

Adams, C. M., E. Hernandez, and J. Lee. 2009. An economic overview of selected industries dependent upon the Gulf of Mexico. In D. L. Felder and D. K. Camp, eds., Gulf of Mexico: Origin, Waters, and Biota: Volume 2, Ocean and Coastal Economy, pp. 28-48. Texas A\&M University Press.

Agardy, T. S. 1997. Marine protected areas and ocean conservation. Academic Press.
Aguilar, C., G. González-Sansón, R. Hueter, E. Rojas, Y. Cabrera, A. Briones, R. Borroto, A. Hernández, and P. Baker. 2014. Captura de tiburones en la región noroccidental de Cuba. Latin American Journal of Aquatic Research, 42(3):477-487.

Ainsworth, C., H. Morzaria-Luna, I. C. Kaplan, P. S. Levin, E. A. Fulton, R. CudneyBueno, P. Turk-Boyer, J. Torre, G. D. Danemann, and T. Pfister. 2012. Effective ecosystem-based management must encourage regulatory compliance: A Gulf of California case study. Marine Policy, 36(6):1275-1283.

Ainsworth, C. H. and T. J. Pitcher. 2006. Modifying Kempton's species diversity index for use with ecosystem simulation models. Ecological Indicators, 6(3):623630.

Ainsworth, C. H., M. J. Schirripa, and H. Morzaria-Luna, eds. 2015. An Atlantis Ecosystem Model for the Gulf of Mexico Supporting Integrated Ecosystem Assessment. NOAA Technical Memorandum NMFS-SEFSC-676, 149p.

Allison, G. W., J. Lubchenco, and M. H. Carr. 1998. Marine reserves are necessary but not sufficient for marine conservation. Ecological Applications, 8(sp1).

Almany, G., S. Connolly, D. Heath, J. Hogan, G. Jones, L. McCook, M. Mills, R. Pressey, and D. Williamson. 2009. Connectivity, biodiversity conservation and the design of marine reserve networks for coral reefs. Coral Reefs, 28(2):339-351.

Amorim, S., M. N. Santos, R. Coelho, and J. Fernandez-Carvalho. 2015. Effects of 17/0 circle hooks and bait on fish catches in a southern Atlantic swordfish longline fishery. Aquatic Conservation: Marine and Freshwater Ecosystems, 25(4):518-533.

Arreguín-Sánchez, F. and E. Arcos-Huitrón. 2011. La pesca en México: estado de la explotación y uso de los ecosistemas. Hidrobiológica, 21(3):431-462.

Arreguín-Sánchez, F., M. Zetina-Rejon, and M. Ramírez-Rodríguez. 2008. Exploring ecosystem-based harvesting strategies to recover the collapsed pink shrimp (Farfantepenaeus duorarum) fishery in the southern Gulf of Mexico. Ecological Modelling, 214(2):83-94.

Ault, J. S., J. A. Bohnsack, S. G. Smith, and J. Luo. 2005. Towards sustainable multispecies fisheries in the Florida, USA, coral reef ecosystem. Bulletin of Marine Science, 76(2):595-622.

Austin, M. 2002. Spatial prediction of species distribution: an interface between ecological theory and statistical modelling. Ecological Modelling, 157(2):101-118.

Austin, M. 2007. Species distribution models and ecological theory: a critical assessment and some possible new approaches. Ecological Modelling, 200(1):1-19.

Babcock, R. C., S. Kelly, N. T. Shears, J. W. Walker, and T. J. Willis. 1999. Changes in community structure in temperate marine reserves. Marine Ecology Progress Series, 189:125-134.

Badan, A., J. Candela, J. Sheinbaum, and J. Ochoa. 2005. Upper-layer circulation in the approaches to Yucatan Channel. In W. Sturges and A. Lugo-Fernandez, eds., Circulation in the Gulf of Mexico: Observations and Models, pp. 57-69. American Geophysical Union.

Bakun, A. 1996. Patterns in the ocean: ocean processes and marine population dynamics. California Sea Grant College System, National Oceanic and Atmospheric Administration.

Bakun, A. and K. Broad. 2003. Environmental 'loopholes' and fish population dynamics: comparative pattern recognition with focus on El Nino effects in the Pacific. Fisheries Oceanography, 12(4-5):458-473.

Bakun, A. and S. J. Weeks. 2006. Adverse feedback sequences in exploited marine systems: are deliberate interruptive actions warranted? Fish and Fisheries, $7(4): 316-333$.

Balmford, A., P. Gravestock, N. Hockley, C. J. McClean, and C. M. Roberts. 2004. The worldwide costs of marine protected areas. Proceedings of the National Academy of Sciences of the United States of America, 101(26):9694-9697.

Barker, M. J. and V. Schluessel. 2005. Managing global shark fisheries: suggestions for prioritizing management strategies. Aquatic Conservation: Marine and Freshwater Ecosystems, 15(4):325-347.

Baum, J. K. and W. Blanchard. 2010. Inferring shark population trends from generalized linear mixed models of pelagic longline catch and effort data. Fisheries Research, 102(3):229-239.

Baum, J. K., D. Kehler, and R. A. Myers. 2005. Robust estimates of decline for pelagic shark populations in the northwest Atlantic and Gulf of Mexico. Fisheries, 30(10):27-30.

Baum, J. K., J. J. Meeuwig, and A. C. Vincent. 2003a. Bycatch of lined seahorses (Hippocampus erectus) in a Gulf of Mexico shrimp trawl fishery. Fishery Bulletin, 101(4):721-731.

Baum, J. K. and R. A. Myers. 2004. Shifting baselines and the decline of pelagic sharks in the Gulf of Mexico. Ecology Letters, 7(2):135-145.

Baum, J. K., R. A. Myers, D. G. Kehler, B. Worm, S. J. Harley, and P. A. Doherty. 2003b. Collapse and conservation of shark populations in the northwest Atlantic. Science, 299(5605):389-392.

Baum, J. K. and B. Worm. 2009. Cascading top-down effects of changing oceanic predator abundances. Journal of Animal Ecology, 78(4):699-714.

Beck, M. W., R. D. Brumbaugh, L. Airoldi, A. Carranza, L. D. Coen, C. Crawford, O. Defeo, G. J. Edgar, B. Hancock, M. C. Kay, et al. 2011. Oyster reefs at risk and recommendations for conservation, restoration, and management. Bioscience, $61(2): 107-116$.

Beddington, J. R. and J. G. Cooke. 1984. Estimating the response of population to exploitation from catch and effort data. In S. A. Levin and T. G. Hallam, eds., Mathematical Ecology, pp. 247-261. Springer.

Beerkircher, L. R., E. Cortes, and M. Shivji. 2002. Characteristics of shark bycatch observed on pelagic longlines off the southeastern United States, 1992-2000. Marine Fisheries Review, 64(4):40-49.

Beerkircher, L. R., D. W. Lee, C. J. Brown, and D. L. Abercrombie. 2004. SEFSC Pelagic Observer Program Data Summary for 1992-2002. NOAA Technical Memorandum NMFS-SEFSC-522, 25p.

Belcher, C. and C. A. Jennings. 2011. Identification and evaluation of shark bycatch in Georgia's commercial shrimp trawl fishery with implications for management. Fisheries Management and Ecology, 18(2):104-112.

Benoit-Bird, K. J., B. C. Battaile, S. A. Heppell, B. Hoover, D. Irons, N. Jones, K. J. Kuletz, C. A. Nordstrom, R. Paredes, R. M. Suryan, et al. 2013. Prey patch patterns predict habitat use by top marine predators with diverse foraging strategies. PLoS One, 8(1): e53348.

Bethea, D. M., M. J. Ajemian, J. K. Carlson, E. R. Hoffmayer, J. L. Imhoff, R. D. Grubbs, C. T. Peterson, and G. H. Burgess. 2015. Distribution and community structure of coastal sharks in the northeastern Gulf of Mexico. Environmental Biology of Fishes, 98(5):1233-1254.

Bethea, D. M., L. Hollensead, J. K. Carlson, C. L. Conrath, M. J. Ajemian, E. R. Hoffmayer, and G. W. Peterson. 2006. Shark Nursery Grounds and Essential Fish Habitat Studies. GULFSPAN Gulf of Mexico-FY07. An Internal Report to NOAA's Highly Migratory Species Office. Sustainable Fisheries Division.

Beverton, R. J. H. and S. J. Holt. 1957. On the dynamics of exploited fish populations. Fishery Investigations Series, 2(19). Ministry of Agriculture, Fisheries, and Food, London, UK.

Bianchi, G., H. Gislason, K. Graham, L. Hill, X. Jin, K. Koranteng, S. ManickchandHeileman, I. Paya, K. Sainsbury, F. Sanchez, et al. 2000. Impact of fishing on size composition and diversity of demersal fish communities. ICES Journal of Marine Science, 57(3):558-571.

Bigelow, K. A., C. H. Boggs, and X. He. 1999. Environmental effects on swordfish and blue shark catch rates in the US North Pacific longline fishery. Fisheries Oceanography, 8(3):178-198.

Bjørndal, T. and A. Brasão. 2006. The East Atlantic bluefin tuna fisheries: stock collapse or recovery? Marine Resource Economics, 21(2):193-210.

Blanchard, J. L., M. Coll, V. M. Trenkel, R. Vergnon, D. Yemane, D. Jouffre, J. S. Link, and Y.-J. Shin. 2010. Trend analysis of indicators: a comparison of recent changes in the status of marine ecosystems around the world. ICES Journal of Marine Science, 67(4):732-744.

Block, B. A., H. Dewar, S. B. Blackwell, T. D. Williams, E. D. Prince, C. J. Farwell, A. Boustany, S. L. Teo, A. Seitz, A. Walli, et al. 2001. Migratory movements, depth preferences, and thermal biology of Atlantic bluefin tuna. Science, 293(5533):13101314.

Block, B. A., S. L. Teo, A. Walli, A. Boustany, M. J. Stokesbury, C. J. Farwell, K. C. Weng, H. Dewar, and T. D. Williams. 2005. Electronic tagging and population structure of Atlantic bluefin tuna. Nature, 434(7037):1121-1127.

Boersma, P. D. and J. K. Parrish. 1999. Limiting abuse: marine protected areas, a limited solution. Ecological Economics, 31(2):287-304.

Bojorquez, L. F. 1998. Bycatch utilization in Mexico. In I. Clucas and F. Teutscher, eds., Report and Proceedings of the FAO/DFID Expert Consultation on Bycatch Utilization in Tropical Fisheries. Beijing, 21-28 September 1998. Rome, FAO.

Bonfil, R. 1997. Status of shark resources in the southern Gulf of Mexico and caribbean: implications for management. Fisheries Research, 29(2):101-117.

Bornatowski, H., A. F. Navia, R. R. Braga, V. Abilhoa, and M. F. M. Corrêa. 2014. Ecological importance of sharks and rays in a structural foodweb analysis in southern Brazil. ICES Journal of Marine Science, 71(7):1586-1592.

Branch, T. A., R. Watson, E. A. Fulton, S. Jennings, C. R. McGilliard, G. T. Pablico, D. Ricard, and S. R. Tracey. 2010. The trophic fingerprint of marine fisheries. Nature, 468(7322):431-435.

Branstetter, S. 1981. Biological notes on the sharks of the north central Gulf of Mexico. Contributions in Marine Science, 24:13-34.

Branstetter, S. 1987. Age and growth estimates for blacktip, Carcharhinus limbatus, and spinner, C. brevipinna, sharks from the northwestern Gulf of Mexico. Copeia, 4:964-974.

Breidt, F. J., H.-L. Lai, J. D. Opsomer, and D. A. Van Voorhees. 2010. A Report of the MRIP Sampling and Estimation Project: Improved Estimation Methods for the Access Point Angler Intercept Survey Component of the Marine Recreational Fishery Statistics Survey. NOAA Fisheries Contract Report, Silver Spring, MD.

Brill, R. W. and M. E. Lutcavage. 2001. Understanding environmental influences on movements and depth distributions of tunas and billfishes can significantly improve population assessments. American Fisheries Society Symposium, 25:179-198.

Britten, G. L., M. Dowd, C. Minto, F. Ferretti, F. Boero, and H. K. Lotze. 2014. Predator decline leads to decreased stability in a coastal fish community. Ecology Letters, 17(12):1518-1525.

Brodziak, J. and J. Link. 2002. Ecosystem-based fishery management: what is it and how can we do it? Bulletin of Marine Science, 70(2):589-611.

Brooke, S. G. 2012. Federal Fisheries Observer Programs in the United States: Over 40 Years of Independent Data Collection. NOAA Marine Fisheries Review, 38p.

Brose, U., R. J. Williams, and N. D. Martinez. 2006. Allometric scaling enhances stability in complex food webs. Ecology Letters, 9(11):1228-1236.

Burgess, G. H., L. R. Beerkircher, G. M. Cailliet, J. K. Carlson, E. Cortés, K. J. Goldman, R. D. Grubbs, J. A. Musick, M. K. Musyl, and C. A. Simpfendorfer. 2005. Is the collapse of shark populations in the Northwest Atlantic Ocean and Gulf of Mexico real? Fisheries, 30(10):19-26.

Butterworth, D. S., N. Bentley, J. A. De Oliveira, G. P. Donovan, L. T. Kell, A. M. Parma, A. E. Punt, K. J. Sainsbury, A. D. Smith, and T. K. Stokes. 2010. Purported flaws in management strategy evaluation: basic problems or misinterpretations? ICES Journal of Marine Science, 67(3):567-574.

Caddy, J. 1993. Toward a comparative evaluation of human impacts on fishery ecosystems of enclosed and semi-enclosed seas. Reviews in Fisheries Science, 1(1):57-95.

Caddy, J. 2000. Marine catchment basin effects versus impacts of fisheries on semienclosed seas. ICES Journal of Marine Science, 57(3):628-640.

Caddy, J. and A. Bakun. 1994. A tentative classification of coastal marine ecosystems based on dominant processes of nutrient supply. Ocean $\mathcal{B}$ Coastal Management, 23(3):201-211.

Caddy, J., F. Carocci, and S. Coppola. 1998a. Have peak fishery production levels been passed in continental shelf area? Some perspectives arising from historical trends in production per shelf area. Journal of Northwest Atlantic Fishery Science, 23:191-220.

Caddy, J. F., J. Csirke, S. M. Garcia, and R. J. R. Grainger. 1998b. How pervasive is "fishing down marine food webs"? Science, 282(5393):1383-1383.

Caddy, J. F. and L. Garibaldi. 2000. Apparent changes in the trophic composition of world marine harvests: the perspective from the FAO capture database. Ocean ${ }^{6}$ Coastal Management, 43(8):615-655.

Campbell, M., A. Pollack, T. Henwood, J. Provaznik, and M. Cook. 2012. Summary Report of the Red Snapper (Lutjanus campechanus) Catch During the 2011 Expanded Annual Stock Assessment (EASA). SEDAR31-DW17. SEDAR, North Charleston, SC. 27 pp.

Cardinale, B. J., J. E. Duffy, A. Gonzalez, D. U. Hooper, C. Perrings, P. Venail, A. Narwani, G. M. Mace, D. Tilman, D. A. Wardle, et al. 2012. Biodiversity loss and its impact on humanity. Nature, 486(7401):59-67.

Carlson, J. and S. Gulak. 2013. Standardized Catch Rates of Atlantic Sharpnose Sharks (Rhizoprionodon terraenovae) in the U.S. Gulf of Mexico From the Shark Bottom Longline Observer Program, 1994-2011. SEDAR 34-WP-01. SEDAR, North Charleston, SC. 22 pp.

Carlson, J., L. Hale, A. Morgan, and G. Burgess. 2010a. Standardized Catch Rates of Sandbar, Dusky and Blacknose Sharks from the Shark Fishery Bottom Longline Observer Program, 1994-2009. SEDAR21-DW-02. SEDAR, North Charleston, SC. 28 pp.

Carlson, J., L. Hale, A. Morgan, and G. Burgess. 2012. Standardized Catch Rates of Blacktip Sharks (Carcharhinus limbatus) in the U.S. Gulf of Mexico from the Shark Bottom Longline Observer Program, 1994-2010. SEDAR 29-WP-02. SEDAR, North Charleston, SC. 16 pp.

Carlson, J. and J. Osborne. 2013. Standardized Catch Rates of Bonnetheads From the Everglades National Park Creel Survey. SEDAR34-WP-02. SEDAR, North Charleston, SC. 11 pp.

Carlson, J. K. and G. R. Parsons. 2001. The effects of hypoxia on three sympatric shark species: physiological and behavioral responses. Environmental Biology of Fishes, 61(4):427-433.

Carlson, J. K., M. M. Ribera, C. L. Conrath, M. R. Heupel, and G. H. Burgess. 2010b. Habitat use and movement patterns of bull sharks Carcharhinus leucas determined using pop-up satellite archival tags. Journal of Fish Biology, 77(3):661-675.

Carollo, C. and D. J. Reed. 2010. Ecosystem-based management institutional design: balance between federal, state, and local governments within the Gulf of Mexico alliance. Marine Policy, 34(1):178-181.

Carruthers, E. H., J. D. Neilson, and S. C. Smith. 2011. Overlooked bycatch mitigation opportunities in pelagic longline fisheries: soak time and temperature effects on swordfish (Xiphias gladius) and blue shark (Prionace glauca) catch. Fisheries Research, 108(1):112-120.

Carruthers, T. R., L. T. Kell, D. D. Butterworth, M. N. Maunder, H. F. Geromont, C. Walters, M. K. McAllister, R. Hillary, P. Levontin, T. Kitakado, et al. 2016. Performance review of simple management procedures. ICES Journal of Marine Science, 73(2):464-482.

Carruthers, T. R., A. E. Punt, C. J. Walters, A. MacCall, M. K. McAllister, E. J. Dick, and J. Cope. 2014. Evaluating methods for setting catch limits in data-limited fisheries. Fisheries Research, 153:48-68.

Casini, M., J. Hjelm, J.-C. Molinero, J. Lövgren, M. Cardinale, V. Bartolino, A. Belgrano, and G. Kornilovs. 2009. Trophic cascades promote threshold-like shifts in pelagic marine ecosystems. Proceedings of the National Academy of Sciences, 106(1):197-202.

Castillo-Géniz, J. L., J. F. Márquez-Farias, M. R. De La Cruz, E. Cortés, and A. C. Del Prado. 1998. The Mexican artisanal shark fishery in the Gulf of Mexico: towards a regulated fishery. Marine and Freshwater Research, 49(7):611-620.

Cayula, J.-F. and P. Cornillon. 1992. Edge detection algorithm for SST images. Journal of Atmospheric and Oceanic Technology, 9(1):67-80.

Ceccarelli, D. M., A. J. Frisch, N. A. Graham, A. M. Ayling, and M. Beger. 2014. Habitat partitioning and vulnerability of sharks in the Great Barrier Reef marine park. Reviews in Fish Biology and Fisheries, 24(1):169-197.

Cervigón, F. 1994. Los Peces Marinos de Venezuela, 2ème édition. Fundación Científica Los Roques, Caracas. Vol. III. 296 pp.

Christensen, V., S. Guenette, J. J. Heymans, C. J. Walters, R. Watson, D. Zeller, and D. Pauly. 2003. Hundred-year decline of north Atlantic predatory fishes. Fish and Fisheries, 4(1):1-24.

Ciannelli, L., P. Fauchald, K.-S. Chan, V. N. Agostini, and G. E. Dingsør. 2008. Spatial fisheries ecology: recent progress and future prospects. Journal of Marine Systems, 71(3):223-236.

Claro, R. 1994. Características generales de la ictiofauna. In R. Claro, ed., Ecología de los peces marinos de Cuba. Instituto de Oceanología Academia de Ciencias de Cuba and Centro de Investigaciones de Quintana Roo.

Claro, R., Y. S. de Mitcheson, K. C. Lindeman, and A. R. García-Cagide. 2009. Historical analysis of Cuban commercial fishing effort and the effects of management interventions on important reef fishes from 1960-2005. Fisheries Research, 99(1):716.

Claro, R., K. C. Lindeman, and L. R. Parenti, eds. 2001. Ecology of the Marine Fishes of Cuba. Smithsonian Institution.

Coelho, R., M. N. Santos, and S. Amorim. 2012. Effects of hook and bait on targeted and bycatch fishes in an equatorial Atlantic pelagic longline fishery. Bulletin of Marine Science, 88(3):449-467.

Coleman, F. C., P. B. Baker, and C. C. Koenig. 2004a. A review of Gulf of Mexico marine protected areas: successes, failures, and lessons learned. Fisheries, $29(2): 10-21$.

Coleman, F. C., W. F. Figueira, J. S. Ueland, and L. B. Crowder. 2004b. The impact of United States recreational fisheries on marine fish populations. Science, 305(5692):1958-1960.

Coleman, F. C., C. C. Koenig, and L. A. Collins. 1996. Reproductive styles of shallow-water groupers (pisces: Serranidae) in the eastern Gulf of Mexico and the consequences of fishing spawning aggregations. Environmental Biology of Fishes, $47(2): 129-141$.

Coleman, F. C. and S. L. Williams. 2002. Overexploiting marine ecosystem engineers: potential consequences for biodiversity. Trends in Ecology $8 \mathcal{E}$ Evolution, 17(1):4044.

Collette, B., K. Carpenter, B. Polidoro, M. Juan-Jordá, A. Boustany, D. Die, C. Elfes, W. Fox, J. Graves, L. Harrison, et al. 2011. High value and long lifedouble jeopardy for tunas and billfishes. Science, 333(6040):291-292.

Compagno, L. J. V. 1984. Sharks of the world: an annotated and illustrated catalogue of shark species known to date. Food and Agriculture Organization Species Catalogue, Vol. 4, Part 2. Carcharhiniformes. FAO Fisheries Synopsis, 125:251-655.

Compagno, L. J. V. 2002. Sharks. In K. E. Carpenter, ed., The Living Marine Resources of the Western Central Atlantic. Vol. 1: Introduction, mollusks, crustaceans, hagfishes, sharks, batoid fishes and chimaeras, pp. 358-505. FAO Species Identification Guide for Fishery Purposes and American Society of Ichthyologists and Herpetologists Special Publication No. 5. FAO, Rome.

Convention on Biological Diversity. 2004. The 2020 biodiversity target: a framework for implementation. In Decisions from the Seventh Meeting of the Conference of the Parties of the Convention on Biological Diversity, Annex I, Decision VII/30, p. 351. Kuala Lumpur, Malaysia.

Cooke, J. 1999. Improvement of fishery-management advice through simulation testing of harvest algorithms. ICES Journal of Marine Science, 56(6):797-810.

Cooke, J. and J. Beddington. 1984. The relationship between catch rates and abundance in fisheries. Mathematical Medicine and Biology, 1(4):391-405.

Cortés, E. 2002a. Catches and catch rates of pelagic sharks from the northwestern Atlantic, Gulf of Mexico, and Caribbean. ICCAT Collective Volume of Scientific Papers, 54(4):1164-1181.

Cortés, E. 2002b. Stock Assessment of Small Coastal Sharks in the US Atlantic and Gulf of Mexico. SFD-01/02-152. NMFS Sustainable Fisheries Division. 133 pp.

Cox, G. J. and M. Francis. 1997. Sharks and Rays of New Zealand. Canterbury University Press.

Cramer, J. 1996. Recent trends in the catch of undersized swordfish by the U.S. pelagic longline fishery. Marine Fisheries Review, 58(3):24-32.

Dale, V. H. and S. C. Beyeler. 2001. Challenges in the development and use of ecological indicators. Ecological Indicators, 1(1):3-10.

Darimont, C. T., S. M. Carlson, M. T. Kinnison, P. C. Paquet, T. E. Reimchen, and C. C. Wilmers. 2009. Human predators outpace other agents of trait change in the wild. Proceedings of the National Academy of Sciences, 106(3):952-954.

Day, J. W. and A. Yáñez-Arancibia, eds. 2013. Gulf of Mexico: Origin, Waters, and Biota: Volume 4, Ecosystem-Based Management. Texas A\&M University Press.
de Leiva Moreno, J., V. Agostini, J. Caddy, and F. Carocci. 2000. Is the pelagicdemersal ratio from fishery landings a useful proxy for nutrient availability? A preliminary data exploration for the semi-enclosed seas around Europe. ICES Journal of Marine Science, 57(4):1091-1102.
de Mutsert, K., J. H. Cowan, T. E. Essington, and R. Hilborn. 2008. Reanalyses of Gulf of Mexico fisheries data: landings can be misleading in assessments of fisheries and fisheries ecosystems. Proceedings of the National Academy of Sciences, 105(7):2740-2744.
de Silva, J. A., R. E. Condrey, K. Anglin, and J. Rester. 1996. Bycatch in the United States Gulf of Mexico menhaden fishery. In Symp. on the Consequences and Management of Fisheries Bycatch. Dearborn Michigan, USA.
de Silva, J. A., R. E. Condrey, and B. A. Thompson. 2001. Profile of shark bycatch in the US Gulf of Mexico menhaden fishery. North American Journal of Fisheries Management, 21(1):111-124.
de Sylva, D. P. 1990. Sphyraenidae. In J. C. Quero, J. C. Hureau, C. Karrer, A. Post, and L. Saldanha, eds., Check-list of the fishes of the eastern tropical Atlantic (CLOFETA), vol. 2. JNICT, Lisbon; SEI, Paris; and UNESCO, Paris.
de Sylva, D. P. and P. R. Breder. 1997. Reproduction, gonad histology, and spawning cycles of north Atlantic billfishes (Istiophoridae). Bulletin of Marine Science, 60(3):668-697.

Dewar, H., E. D. Prince, M. K. Musyl, R. W. Brill, C. Sepulveda, J. Luo, D. Foley, E. S. Orbesen, M. L. Domeier, N. NASBY-LUCAS, et al. 2011. Movements and behaviors of swordfish in the Atlantic and Pacific Oceans examined using pop-up satellite archival tags. Fisheries Oceanography, 20(3):219-241.

Diamond, S. L., L. G. Cowell, and L. B. Crowder. 2000. Population effects of shrimp trawl bycatch on Atlantic croaker. Canadian Journal of Fisheries and Aquatic Sciences, 57(10):2010-2021.

Dicenta, A., C. Piccinetti, et al. 1980. Comparison between the estimated reproductive stocks of bluefin tuna ( $T$. thynnus) of the Gulf of Mexico and western Mediterranean. ICCAT Collected Volume of Scientific Papers, 9(2):442-448.

Dick, E. 2004. Beyond lognormal versus gamma: discrimination among error distributions for generalized linear models. Fisheries Research, 70(2):351-366.

Dickey-Collas, M. 2014. Why the complex nature of integrated ecosystem assessments requires a flexible and adaptive approach. ICES Journal of Marine Science, 71(5):1174-1182.

Die, D. J. 2006. Are Atlantic marlins overfished or endangered? Some reasons why we may not be able to tell. Bulletin of Marine Science, 79(3):529-543.

Dong, Q. and V. R. Restrepo. 1996. Notes on the poisson error assumption made to estimate relative abundance of west Atlantic bluefin tuna. ICCAT Collective Volume of Scientific Papers, 45:158-161.

Drexler, M. and C. H. Ainsworth. 2013. Generalized additive models used to predict species abundance in the Gulf of Mexico: an ecosystem modeling tool. PLoS One, 8(5):e64458.

Drymon, J. M., L. Carassou, S. P. Powers, M. Grace, J. Dindo, and B. Dzwonkowski. 2013. Multiscale analysis of factors that affect the distribution of sharks throughout the northern Gulf of Mexico. Fishery Bulletin, 111(4):370-380.

Drymon, J. M., S. P. Powers, J. Dindo, B. Dzwonkowski, and T. A. Henwood. 2010. Distributions of sharks across a continental shelf in the northern Gulf of Mexico. Marine and Coastal Fisheries, 2(1):440-450.

Ducet, N., P.-Y. Le Traon, and G. Reverdin. 2000. Global high-resolution mapping of ocean circulation from topex/poseidon and ers-1 and-2. Journal of Geophysical Research: Oceans (1978-2012), 105(C8):19477-19498.

Duffy, J. E. 2002. Biodiversity and ecosystem function: the consumer connection. Oikos, 99(2):201-219.

Duffy, J. E. 2003. Biodiversity loss, trophic skew and ecosystem functioning. Ecology Letters, 6(8):680-687.

Dulvy, N. K., J. K. Baum, S. Clarke, L. J. Compagno, E. Cortes, A. Domingo, S. Fordham, S. Fowler, M. P. Francis, C. Gibson, et al. 2008. You can swim but you can't hide: the global status and conservation of oceanic pelagic sharks and rays. Aquatic Conservation: Marine and Freshwater Ecosystems, 18(5):459-482.

Dunn, D. C., A. M. Boustany, and P. N. Halpin. 2011. Spatio-temporal management of fisheries to reduce by-catch and increase fishing selectivity. Fish and Fisheries, 12(1):110-119.

Ecosystem Principles Advisory Panel. 1999. Ecosystem-based fishery management. US National Oceanic and Atmospheric Administration. National Marine Fisheries Service, Silver Spring, MD. 54p.

Elith, J. and J. R. Leathwick. 2009. Species distribution models: ecological explanation and prediction across space and time. Annual Review of Ecology, Evolution, and Systematics, 40(1):677.

Essington, T. E., A. H. Beaudreau, and J. Wiedenmann. 2006. Fishing through marine food webs. Proceedings of the National Academy of Sciences of the United States of America, 103(9):3171-3175.

Etnoyer, P. and J. Warrenchuk. 2007. A catshark nursery in a deep gorgonian field in the Mississippi Canyon, Gulf of Mexico. Bulletin of Marine Science, 81(3):553.

Fahmi and Dharmadi. 2015. Pelagic shark fisheries of Indonesia's eastern Indian Ocean fisheries management region. African Journal of Marine Science, 37(2):259265.

Falterman, B. and J. E. Graves. 2002. A preliminary comparison of the relative mortality and hooking efficiency of circle and straight shank ("J") hooks used in the pelagic longline industry. In American Fisheries Society Symposium, pp. 80-87. American Fisheries Society.

FAO. 2003. Fisheries and Aquaculture Department. Organisation des Nations Unies pour l'alimentation et l'agriculture. Food and Agriculture Organization of the United Nations. World Wide Web electronic publication. http://www.fao.org/ fi/oldsite/FCP/en/mex/profile.htm.

FAO. 2013a. Fisheries and Aquaculture Department. Statistics-Introduction. Food and Agriculture Organization of the United Nations. World Wide Web electronic publication. http://www.fao.org/fishery/statistics/en. Accessed: 19 May 2013.

FAO. 2013b. Report of the Fourth FAO Expert Advisory Panel for the Assessment of Proposals to Amend Appendices I and II of CITES Concerning Commerciallyexploited Aquatic Species. FAO Fisheries and Aquaculture Report No. R1032. Rome, FAO. 161 pp.

Field, I. C., M. G. Meekan, R. C. Buckworth, and C. J. Bradshaw. 2009. Susceptibility of sharks, rays and chimaeras to global extinction. Advances in Marine Biology, 56:275-363.

Finkbeiner, E. M., B. P. Wallace, J. E. Moore, R. L. Lewison, L. B. Crowder, and A. J. Read. 2011. Cumulative estimates of sea turtle bycatch and mortality in USA fisheries between 1990 and 2007. Biological Conservation, 144(11):2719-2727.

Fitzhugh, G. R., E. T. Lang, and H. Lyon. 2012. Expanded Annual Stock Assessment Survey 2011: Red Snapper Reproduction. SEDAR31-DW07. SEDAR, North Charleston, SC. 33 pp .

Foley, M. M., M. H. Armsby, E. E. Prahler, M. R. Caldwell, A. L. Erickson, J. N. Kittinger, L. B. Crowder, and P. S. Levin. 2013. Improving ocean management through the use of ecological principles and integrated ecosystem assessments. Bioscience, 63(8):619-631.

Foster, D. G., S. P. Epperly, A. K. Shah, and J. W. Watson. 2012. Evaluation of hook and bait type on the catch rates in the western north Atlantic Ocean pelagic longline fishery. Bulletin of Marine Science, 88(3):529-545.

Fox, D. S. and R. M. Starr. 1996. Comparison of commercial fishery and research catch data. Canadian Journal of Fisheries and Aquatic Sciences, 53(12):2681-2694.

Frick, A. 2011. Fishery Closures - Seasonal/Area/Quota Closures In Gulf of Mexico Data Atlas. Stennis Space Center (MS): National Centers for Environmental Information. https://www.ncddc.noaa.gov/website/DataAtlas/atlas.htm. Accessed: 1 April 2016.

Fricke, R., M. Kulbicki, and L. Wantiez. 2011. Checklist of the fishes of New Caledonia, and their distribution in the Southwest Pacific Ocean (Pisces). Stuttg Beitr Natkd Ser A (Biol), 4:341-463.

Froese, R. 2004. Keep it simple: three indicators to deal with overfishing. Fish and Fisheries, 5(1):86-91.

Froese, R. and D. Pauly, eds. 2016. FishBase. World Wide Web electronic publication. www.fishbase.org, version (10/2016). Accessed: 12 October, 2016.

Fromentin, J.-M. and J. E. Powers. 2005. Atlantic bluefin tuna: population dynamics, ecology, fisheries and management. Fish and Fisheries, 6(4):281-306.

Fulton, E. A. 2002. The effects of model structure and complexity on the behaviour and performance of marine ecosystem models. Ph.D. thesis, University of Tasmania, Australia.

Fulton, E. A. 2010. Approaches to end-to-end ecosystem models. Journal of Marine Systems, 81(1):171-183.

Fulton, E. A., M. Fuller, A. Smith, and A. Punt. 2004a. Ecological indicators of the ecosystem effects of fishing: final report. Australian Fisheries Management Authority Report, 99:116.

Fulton, E. A., J. S. Link, I. C. Kaplan, M. Savina-Rolland, P. Johnson, C. Ainsworth, P. Horne, R. Gorton, R. J. Gamble, A. D. Smith, et al. 2011. Lessons in modelling and management of marine ecosystems: the atlantis experience. Fish and Fisheries, 12(2):171-188.

Fulton, E. A., J. S. Parslow, A. D. Smith, and C. R. Johnson. 2004b. Biogeochemical marine ecosystem models ii: the effect of physiological detail on model performance. Ecological Modelling, 173(4):371-406.

Fulton, E. A., A. D. Smith, and C. R. Johnson. 2004c. Biogeochemical marine ecosystem models i: IGBEM - a model of marine bay ecosystems. Ecological Modelling, 174(3):267-307.

Fulton, E. A., A. D. Smith, and A. E. Punt. 2005. Which ecological indicators can robustly detect effects of fishing? ICES Journal of Marine Science, 62(3):540-551.

Fulton, E. A., A. D. M. Smith, and D. C. Smith. 2007. Alternative Management Strategies for Southeast Australian Commonwealth Fisheries: Stage 2: Quantitative Management Strategy Evaluation. Commonwealth Scientific and Industrial Research Organization (CSIRO).

Gaines, S. D., C. White, M. H. Carr, and S. R. Palumbi. 2010. Designing marine reserve networks for both conservation and fisheries management. Proceedings of the National Academy of Sciences, 107(43):18286-18293.

Gallaway, B. J. and J. G. Cole. 1999. Reduction of juvenile red snapper bycatch in the US Gulf of Mexico shrimp trawl fishery. North American Journal of Fisheries Management, 19(2):342-355.

Game, E. T., H. S. Grantham, A. J. Hobday, R. L. Pressey, A. T. Lombard, L. E. Beckley, K. Gjerde, R. Bustamante, H. P. Possingham, and A. J. Richardson. 2009. Pelagic protected areas: the missing dimension in ocean conservation. Trends in Ecology $\mathcal{E}^{2}$ Evolution, 24(7):360-369.

Game, E. T., H. S. Grantham, A. J. Hobday, R. L. Pressey, A. T. Lombard, L. E. Beckley, K. Gjerde, R. Bustamante, H. P. Possingham, and A. J. Richardson. 2010. Pelagic MPAs: The devil you know. Trends in Ecology 83 Evolution, 25(2):63-64.

García, V. B., L. O. Lucifora, and R. A. Myers. 2008. The importance of habitat and life history to extinction risk in sharks, skates, rays and chimaeras. Proceedings of the Royal Society of London B: Biological Sciences, 275(1630):83-89.

Gascuel, D., G. Merino, R. Döring, J. N. Druon, L. Goti, S. Guenette, C. Macher, K. Soma, M. Travers-Trolet, and S. Mackinson. 2012. Towards the implementation of an integrated ecosystem fleet-based management of European fisheries. Marine Policy, 36(5):1022-1032.

Giattina, J. D. and D. T. Altsman. 1999. Gulf of Mexico program: partnership with a purpose. In H. Kumpf, K. Steidinger, and K. Sherman, eds., The Gulf of Mexico Large Marine Ecosystem: Assessment, Sustainability, and Management, pp. 3-13. Blackwell Science.

Godø, O. R., A. Samuelsen, G. J. Macaulay, R. Patel, S. S. Hjøllo, J. Horne, S. Kaartvedt, and J. A. Johannessen. 2012. Mesoscale eddies are oases for higher trophic marine life. PLoS One, 7(1):e30161. doi:10.1371/journal.pone. 0030161.

Goodyear, C. P. 1998. An analysis of the possible utility of time-area closures to minimize billfish bycatch by US pelagic longlines. ICCAT Collective Volume of Scientific Papers, 48(1):263-268.

Goodyear, C. P., J. Luo, E. D. Prince, J. P. Hoolihan, D. Snodgrass, E. S. Orbesen, and J. E. Serafy. 2008. Vertical habitat use of Atlantic blue marlin Makaira nigricans: interaction with pelagic longline gear. Marine Ecology Progress Series, 365:233-245.

Grace, M. and T. Henwood. 1997. Assessment of the distribution and abundance of coastal sharks in the US Gulf of Mexico and eastern seaboard, 1995 and 1996. Marine Fisheries Review, 59(4):23-32.

Grantham, H. S., E. T. Game, A. T. Lombard, A. J. Hobday, A. J. Richardson, L. E. Beckley, R. L. Pressey, J. A. Huggett, J. C. Coetzee, C. D. Van der Lingen, et al. 2011. Accommodating dynamic oceanographic processes and pelagic biodiversity in marine conservation planning. PLoS One, 6(2):e16552. doi:10.1371/journal.pone. 0016552 .

Grantham, H. S., S. L. Petersen, and H. P. Possingham. 2008. Reducing bycatch in the South African pelagic longline fishery: the utility of different approaches to fisheries closures. Endangered Species Research, 5(2-3):291-299.

Gray, J. S. 1997. Marine biodiversity: patterns, threats and conservation needs. Biodiversity $\mathcal{G}$ Conservation, 6(1):153-175.

Greiner, M., D. Pfeiffer, and R. Smith. 2000. Principles and practical application of the receiver-operating characteristic analysis for diagnostic tests. Preventive Veterinary Medicine, 45(1):23-41.

Grémillet, D., S. Lewis, L. Drapeau, C. D. van Der Lingen, J. A. Huggett, J. C. Coetzee, H. M. Verheye, F. Daunt, S. Wanless, and P. G. Ryan. 2008. Spatial match-mismatch in the Benguela upwelling zone: should we expect chlorophyll and sea-surface temperature to predict marine predator distributions? Journal of Applied Ecology, 45(2):610-621.

Griffiths, S. and G. Fay. 2015. Integrating recreational fisheries data into stock assessment: implications for model performance and subsequent harvest strategies. Fisheries Management and Ecology, 22(3):197-212.

Grüss, A. 2014. Modelling the impacts of marine protected areas for mobile exploited fish populations and their fisheries: what we recently learnt and where we should be going. Aquatic Living Resources, 27(3-4):107-133.

Grüss, A., E. A. Babcock, S. R. Sagarese, M. Drexler, D. D. Chagaris, C. H. Ainsworth, B. Penta, and T. T. Sutton. 2016a. Improving the spatial allocation of functional group biomasses in spatially-explicit ecosystem models: insights from three Gulf of Mexico models. Bulletin of Marine Science, 92(4):000-000.

Grüss, A., M. Drexler, and C. H. Ainsworth. 2014. Using delta generalized additive models to produce distribution maps for spatially explicit ecosystem models. Fisheries Research, 159:11-24.

Grüss, A., W. J. Harford, M. J. Schirripa, L. Velez, S. R. Sagarese, Y.-J. Shin, and P. Verley. 2016b. Management strategy evaluation using the individual-based, multispecies modeling approach OSMOSE. Ecological Modelling, 340:86-105.

Guidetti, P. 2007. Potential of marine reserves to cause community-wide changes beyond their boundaries. Conservation Biology, 21(2):540-545.

Guisan, A., T. C. Edwards, and T. Hastie. 2002. Generalized linear and generalized additive models in studies of species distributions: setting the scene. Ecological Modelling, 157(2):89-100.

Guisan, A. and N. E. Zimmermann. 2000. Predictive habitat distribution models in ecology. Ecological Modelling, 135(2):147-186.

Guitart, D. J. 1975. Las pesquíres pelágico-oceánicas de corto radio de acción en la región noroccidental de cuba. Academia de Ciencia de Cuba, Instituto de Oceanología, La Habana, pp. 1-26.

Gulf of Mexico Fishery Management Council. 1989. Amendment 1 to the Fishery Management Plan for the Reef Fish Fishery of the Gulf of Mexico. Gulf of Mexico Fisheries Management Council, Tampa, FL.

Gulf of Mexico Fishery Management Council. 2005. Gulf of Mexico Fishery Management Council Report to the National Research Council on the Recreational Fishing Survey Methods. National Oceanic and Atmospheric Administration.

Gulf of Mexico Fishery Management Council. 2016. Commercial Fishing Regulations for Gulf of Mexico Federal waters. Gulf of Mexico Fisheries Management Council, Tampa, FL. URL http://www.gulfcouncil.org.

Gulland, J. A. 1956. On the Fishing Effort in English Demersal Fisheries. Her Majesty's Stationery Office.

Hale, L. F. and J. K. Carlson. 2007. Characterization of the Shark Bottom Longline Fishery, 2005-2006. NOAA Technical Memorandum NMFS-SEFSC-554, 28p.

Hamilton, P., T. J. Berger, J. J. Singer, E. Waddell, J. H. Churchill, R. R. Leben, T. N. Lee, and W. Sturges. 2000a. DeSoto Canyon eddy intrusion study: Final report volume i: Executive summary. OCS Study MMS 2000-079, U.S. Department of Interior, Minerals Management Service, Gulf of Mexico OCS Region, New Orleans, LA. 47 pp .

Hamilton, P., T. J. Berger, J. J. Singer, E. Waddell, J. H. Churchill, R. R. Leben, T. N. Lee, and W. Sturges. 2000b. DeSoto Canyon eddy intrusion study: Final report volume ii: Technical report. OCS Study MMS 2000-080, U.S. Department of Interior, Minerals Management Service, Gulf of Mexico OCS Region, New Orleans, LA. 293 pp.

Hammerschlag, N., J. Luo, D. J. Irschick, and J. S. Ault. 2012. A comparison of spatial and movement patterns between sympatric predators: bull sharks (Carcharhinus leucas) and Atlantic tarpon (Megalops atlanticus). PLoS One, 7(9):e45958. URL http://dx.doi.org/10.1371/journal.pone. 0045958.

Hanley, J. A. and B. J. McNeil. 1982. The meaning and use of the area under a receiver operating characteristic (ROC) curve. Radiology, 143(1):29-36.

Hannan, K. M., I. Driggers, B. William, D. S. Hanisko, L. M. Jones, and A. B. Canning. 2012. Distribution of the nurse shark, ginglymostoma cirratum, in the northern Gulf of Mexico. Bulletin of Marine Science, 88(1):73-80.

Harley, S. J., R. A. Myers, and A. Dunn. 2001. Is catch-per-unit-effort proportional to abundance? Canadian Journal of Fisheries and Aquatic Sciences, 58(9):17601772.

Harrington, J. M., R. A. Myers, and A. A. Rosenberg. 2005. Wasted fishery resources: discarded by-catch in the USA. Fish and Fisheries, 6(4):350-361.

Hastie, T. J. and R. J. Tibshirani. 1986. Generalized additive models. Statistical Science, pp. 297-310.

Hastie, T. J. and R. J. Tibshirani. 1990. Generalized Additive Models, vol. 43. CRC Press.

Hayes, D. B., J. K. Brodziak, and J. B. O'Gorman. 1995. Efficiency and bias of estimators and sampling designs for determining length-weight relationships of fish. Canadian Journal of Fisheries and Aquatic Sciences, 52(1):84-92.

Heithaus, M. R., D. Burkholder, R. E. Hueter, L. I. Heithaus, H. L. Pratt, Jr, and J. C. Carrier. 2007a. Spatial and temporal variation in shark communities of the lower florida keys and evidence for historical population declines. Canadian Journal of Fisheries and Aquatic Sciences, 64(10):1302-1313.

Heithaus, M. R., B. K. Delius, A. J. Wirsing, and M. M. Dunphy-Daly. 2009. Physical factors influencing the distribution of a top predator in a subtropical oligotrophic estuary. Limnology and Oceanography, 54(2):472-482.

Heithaus, M. R., A. Frid, A. J. Wirsing, and B. Worm. 2008. Predicting ecological consequences of marine top predator declines. Trends in Ecology 8 Evolution, 23(4):202-210.

Heithaus, M. R., A. J. Wirsing, L. M. Dill, and L. I. Heithaus. 2007b. Long-term movements of tiger sharks satellite-tagged in Shark Bay, Western Australia. Marine Biology, 151(4):1455-1461.

Henwood, T., W. Ingram, and M. Grace. 2006. Shark/snapper/grouper longline surveys. SEDAR7-DW-08. SEDAR, North Charleston, SC. 22 pp.

Heupel, M. R. and C. A. Simpfendorfer. 2008. Movement and distribution of young bull sharks Carcharhinus leucas in a variable estuarine environment. Aquatic Biology, 1:277-289.

Highly Migratory Species Division. 2000. Final Supplemental Environmental Impact Statement, Regulatory Amendment 1 to the Atlantic Tunas, Swordfish, and Sharks, Fishery Management Plan. National Oceanic and Atmospheric Administration, National Marine Fisheries Service, Office of Sustainable Fisheries, Highly Migratory Species Management Division, Public Document, Silver Spring Maryland.

Highly Migratory Species Division. 2008. Final amendment 2 to the consolidated Atlantic highly migratory species fishery management plan. National Oceanic and Atmospheric Administration, National Marine Fisheries Service, Office of Sustainable Fisheries, Highly Migratory Species Management Division, Public Document, Silver Spring Maryland.

Highly Migratory Species Division. 2014. Final amendment 7 to the 2006 consolidated Atlantic highly migratory species fishery management plan. National Oceanic and Atmospheric Administration, National Marine Fisheries Service, Office of Sustainable Fisheries, Highly Migratory Species Management Division, Public Document, Silver Spring Maryland.

Hilborn, R. 2011. Future directions in ecosystem based fisheries management: a personal perspective. Fisheries Research, 108(2):235-239.

Hilborn, R., C. J. Walters, et al. 1992. Quantitative fisheries stock assessment: choice, dynamics and uncertainty. Reviews in Fish Biology and Fisheries, 2(2):177-178.

Hoffmayer, E. R., J. S. Franks, W. B. Driggers III, J. A. McKinney, J. M. Hendon, and J. M. Quattro. 2014. Habitat, movements and environmental preferences of dusky sharks, Carcharhinus obscurus, in the northern Gulf of Mexico. Marine Biology, 161(4):911-924.

Hoffmayer, E. R. and G. R. Parsons. 2003. Food habits of three shark species from the Mississippi Sound in the northern Gulf of Mexico. Southeastern Naturalist, 2(2):271-280.

Honma, M. 1973. Overall fishing intensity and catch by length class of yellowfin tuna in Japanese Atlantic longline fishery, 1956-1972. ICCAT Collective Volume of Scientific Papers, 4:20-25.

Hueter, R. E. and J. P. Tyminski. 2007. Species-specific distribution and habitat characteristics of shark nurseries in Gulf of Mexico waters off peninsular Florida and Texas. In American Fisheries Society Symposium, vol. 50, pp. 193-223. American Fisheries Society.

Humphries, N. E., N. Queiroz, J. R. M. Dyer, N. G. Pade, M. K. Musyl, K. M. Schaefer, D. W. Fuller, J. M. Brunnschweiler, T. K. Doyle, J. D. Houghton, et al. 2010. Environmental context explains lévy and brownian movement patterns of marine predators. Nature, 465(7301):1066-1069.

Hussey, N. E., S. F. Dudley, I. D. McCarthy, G. Cliff, and A. T. Fisk. 2011. Stable isotope profiles of large marine predators: viable indicators of trophic position, diet, and movement in sharks? Canadian Journal of Fisheries and Aquatic Sciences, 68(12):2029-2045.

Hutchings, J. A. 2000. Collapse and recovery of marine fishes. Nature, 406(6798):882885.

Hyrenbach, K. D., K. A. Forney, and P. K. Dayton. 2000. Marine protected areas and ocean basin management. Aquatic Conservation: Marine and Freshwater Ecosystems, 10(6):437-458.

ICCAT. 2007. Supplemental Recommendation by ICCAT Concerning Sharks. International Commission for the Conservation of Atlantic Tunas.

ICCAT. 2009. Recommendation by ICCAT on the Conservation of Thresher Sharks Caught in Association with Fisheries in the ICCAT Convention Area. International Commission for the Conservation of Atlantic Tunas.

ICCAT. 2010. Recommendation by ICCAT on Hammerhead Sharks (Family Sphyrnidae) Caught in Association with Fisheries Managed by ICCAT. International Commission for the Conservation of Atlantic Tunas.

ICCAT. 2011. Report of the 2011 Blue Marlin Stock Assessment and White Marlin Data Preparatory Meeting. International Commission for the Conservation of Atlantic Tunas.

ICCAT. 2012. Report of the 2012 white marlin stock assessment meeting. International Commission for the Conservation of Atlantic Tunas.

ICCAT. 2014a. Recommendation by ICCAT Amending the Supplemental Recommendation by ICCAT Concerning the Western Atlantic Bluefin Tuna Rebuilding Program. International Commission for the Conservation of Atlantic Tunas.

ICCAT. 2014b. Report of the 2014 Atlantic Bluefin Tuna Stock Assessment Session. International Commission for the Conservation of Atlantic Tunas, Madrid, Spain.

ICCAT. 2016a. Report of the 2014 Atlantic Bluefin Tuna Stock Assessment Session. International Commission for the Conservation of Atlantic Tunas, Madrid, Spain.

ICCAT. 2016b. Report of the 2016 ICCAT Yellowfin Tuna Stock Assessment Meeting. International Commission for the Conservation of Atlantic Tunas, San Sebastian, Spain.

ICCAT. 2016c. Report of the 2016 Sailfish Stock Assessment. International Commission for the Conservation of Atlantic Tunas, Miami, USA.

Ingram, W., T. Henwood, M. Grace, L. Jones, W. Driggers, and K. Mitchell. 2005. Catch rates, distribution and size composition of large coastal sharks collected during NOAA Fisheries Bottom Longline Surveys from the US Gulf of Mexico and US Atlantic Ocean. SEDAR11-LCS05/06-DW-27. SEDAR, North Charleston, SC. 62 pp.

Ingram Jr., G. W. 2008. Data Summary of King Mackerel (Scomberomorus cavalla) Collected During Small Pelagic Trawl Surveys in the US Gulf of Mexico, 1988-1996 and 2002-2007. SEDAR16-DW20. SEDAR, North Charleston, SC. 16 pp.

ITIS. 2012. Integrated Taxonomic Information System. World Wide Web electronic publication. http://www.itis.gov/. Accessed: 22 January 2012.

Jameson, S. C., M. H. Tupper, and J. M. Ridley. 2002. The three screen doors: can marine protected areas be effective? Marine Pollution Bulletin, 44(11):1177-1183.

Januchowski-Hartley, F. A., N. A. Graham, J. E. Cinner, and G. R. Russ. 2013. Spillover of fish naivete from marine reserves. Ecology Letters, 16(2):191-197.

Jennings, S., O. Reñones, B. Morales-Nin, N. Polunin, J. Moranta, and J. Coll. 1997. Spatial variation in the 15 n and 13 c stable isotope composition of plants, invertebrates and fishes on mediterranean reefs: implications for the study of trophic pathways. Marine Ecology Progress Series, 146:109-116.

Jones, G. P., M. Srinivasan, and G. R. Almany. 2007. Population connectivity and conservation of marine biodiversity. Oceanography, 20(3):100-110.

Juan-Jordá, M. J., I. Mosqueira, A. B. Cooper, J. Freire, and N. K. Dulvy. 2011. Global population trajectories of tunas and their relatives. Proceedings of the National Academy of Sciences, 108(51):20650-20655.

Kaplan, D. M., E. Chassot, A. Gruss, and A. Fonteneau. 2010. Pelagic MPAs: the devil is in the details. Trends in Ecology © Evolution, 25(2):62-63.

Kaplan, I. C., P. J. Horne, and P. S. Levin. 2012. Screening California Current fishery management scenarios using the Atlantis end-to-end ecosystem model. Progress in Oceanography, 102:5-18.

Kaplan, I. C. and J. Leonard. 2012. From krill to convenience stores: forecasting the economic and ecological effects of fisheries management on the US west coast. Marine Policy, 36(5):947-954.

Karnauskas, M., M. J. Schirripa, C. R. Kelble, G. S. Cook, and J. K. Craig, eds. 2013. Ecosystem status report for the Gulf of Mexico. NOAA Technical Memorandum NMFS-SEFSC-653, 52p.

Kauermann, G. and J. D. Opsomer. 2011. Data-driven selection of the spline dimension in penalized spline regression. Biometrika, 98(1):225-230.

Kellner, J. B., I. Tetreault, S. D. Gaines, and R. M. Nisbet. 2007. Fishing the line near marine reserves in single and multispecies fisheries. Ecological Applications, 17(4):1039-1054.

Kelly, S., D. Scott, and A. MacDiarmid. 2002. The value of a spillover fishery for spiny lobsters around a marine reserve in northern New Zealand. Coastal Management, $\mathbf{3 0}(2)$ :153-166.

Kempton, R. A. and L. R. Taylor. 1976. Models and statistics for species diversity. Nature, 262:818-820.

Kerstetter, D., S. Bayse, and J. Graves. 2010. Sailfish (Istiophorus platypterus) habitat utilization in the southern Gulf of Mexico and Florida Straits with implications on vulnerability to shallow-set pelagic longline gear. ICCAT Collective Volume of Scientific Papers, 65:1701-12.

Kerstetter, D. W. and J. E. Graves. 2006. Effects of circle versus J-style hooks on target and non-target species in a pelagic longline fishery. Fisheries Research, 80(2):239-250.

Kim, Y.-J. and C. Gu. 2004. Smoothing spline gaussian regression: more scalable computation via efficient approximation. Journal of the Royal Statistical Society: Series B (Statistical Methodology), 66(2):337-356.

Kiraly, S. J., J. A. Moore, and P. H. Jasinski. 2003. Deepwater and other sharks of the US Atlantic Ocean exclusive economic zone. Marine Fisheries Review, 65(4):1-20.

Kleisner, K. M. 2008. A Spatio-Temporal Analysis of Dolphinfish; Coryphaena hippurus, Abundance in the Western Atlantic: Implications for Stock Assessment of a Data-Limited Pelagic Resource. Ph.D. thesis, University of Miami, United States.

Kleisner, K. M., J. F. Walter III, S. L. Diamond, and D. J. Die. 2010. Modeling the spatial autocorrelation of pelagic fish abundance. Marine Ecology Progress Series, 411:203-213.

Kraus, R. T., R. J. D. Wells, and J. R. Rooker. 2011. Horizontal movements of Atlantic blue marlin (Makaira nigricans) in the Gulf of Mexico. Marine Biology, 158(3):699-713.

Landry Jr, A. M. and D. Costa. 1999. Status of sea turtle stocks in the Gulf of Mexico with emphasis on the kemp's ridley. In H. Kumpf, K. Steidinger, and K. Sherman, eds., The Gulf of Mexico Large Marine Ecosystem: Assessment, Sustainability, and Management, pp. 248-268. Blackwell Science.

Lang, K. L., C. B. Grimes, and R. F. Shaw. 1994. Variations in the age and growth of yellowfin tuna larvae, Thunnus albacares, collected about the Mississippi River plume. Environmental Biology of Fishes, 39(3):259-270.

Lawson, G. L., M. R. Castleton, and B. A. Block. 2009. Movements and diving behavior of Atlantic bluefin tuna Thunnus thynnus in relation to water column structure in the northwestern Atlantic. Marine Ecology Progress Series, 400:245.

Leben, R. R. 2005. Altimeter-derived Loop Current metrics. In W. Sturges and A. Lugo-Fernandez, eds., Circulation in the Gulf of Mexico: Observations and Models, pp. 181-201. American Geophysical Union.

Legeckis, R. 1978. A survey of worldwide sea surface temperature fronts detected by environmental satellites. Journal of Geophysical Research: Oceans (1978-2012), 83(C9):4501-4522.

Lester, S. E., B. S. Halpern, K. Grorud-Colvert, J. Lubchenco, B. I. Ruttenberg, S. D. Gaines, S. Airamé, and R. R. Warner. 2009. Biological effects within no-take marine reserves: a global synthesis. Marine Ecology Progress Series, 384(2):33-46.

Levin, P. S., M. J. Fogarty, G. C. Matlock, and E. M. 2008. Integrated Ecosystem Assessment. NOAA Technical Memorandum NMFS-NWFSC-92, 21p.

Levin, P. S., M. J. Fogarty, S. A. Murawski, and D. Fluharty. 2009. Integrated ecosystem assessments: developing the scientific basis for ecosystem-based management of the ocean. PLoS Biology, 7(1):e1000014. doi:10.1371/journal.pbio. 1000014.

Li, Y. and D. Ruppert. 2008. On the asymptotics of penalized splines. Biometrika, 95(2):415-436.

Link, J. 2010. Ecosystem-based fisheries management: confronting tradeoffs. Cambridge University Press.

Link, J. S. 2002. What does ecosystem-based fisheries management mean? Fisheries, 27(4):18-21.

Link, J. S., E. A. Fulton, and R. J. Gamble. 2010a. The northeast US application of ATLANTIS: a full system model exploring marine ecosystem dynamics in a living marine resource management context. Progress in Oceanography, 87(1):214-234.

Link, J. S., D. Yemane, L. J. Shannon, M. Coll, Y.-J. Shin, L. Hill, and M. de Fatima Borges. 2010b. Relating marine ecosystem indicators to fishing and environmental drivers: an elucidation of contrasting responses. ICES Journal of Marine Science, 67(4):787-795.

Llopiz, J. K. and R. K. Cowen. 2008. Precocious, selective and successful feeding of larval billfishes in the oceanic Straits of Florida. Marine Ecology Progress Series, 358:231-244.

Lowe, C. G., B. M. Wetherbee, G. L. Crow, and A. L. Tester. 1996. Ontogenetic dietary shifts and feeding behavior of the tiger shark, Galeocerdo cuvier, in Hawaiian waters. Environmental Biology of Fishes, 47(2):203-211.

Luo, J., J. S. Ault, L. K. Shay, J. P. Hoolihan, E. D. Prince, C. A. Brown, and J. R. Rooker. 2015. Ocean heat content reveals secrets of fish migrations. PLoS One, 10(10):e0141101. doi:10.1371/journal.pone.0141101.

MacKenzie, B. R., H. Mosegaard, and A. A. Rosenberg. 2009. Impending collapse of bluefin tuna in the northeast Atlantic and Mediterranean. Conservation Letters, 2(1):26-35.

Malakoff, D. 1998. Death by suffocation in the Gulf of Mexico. Science, 281(5374):190-192.

Mandelman, J. W., P. W. Cooper, T. B. Werner, and K. M. Lagueux. 2008. Shark bycatch and depredation in the US Atlantic pelagic longline fishery. Reviews in Fish Biology and Fisheries, 18(4):427-442.

Marasco, R. J., D. Goodman, C. B. Grimes, P. W. Lawson, A. E. Punt, and T. J. Quinn II. 2007. Ecosystem-based fisheries management: some practical suggestions. Canadian Journal of Fisheries and Aquatic Sciences, 64(6):928-939.

Martell, S. J., T. E. Essington, B. Lessard, J. F. Kitchell, C. J. Walters, and C. H. Boggs. 2005. Interactions of productivity, predation risk, and fishing effort in the efficacy of marine protected areas for the central Pacific. Canadian Journal of Fisheries and Aquatic Sciences, 62(6):1320-1336.

Masi, M., C. Ainsworth, and D. Jones. 2017. Using a Gulf of Mexico Atlantis model to evaluate ecological indicators for sensitivity to fishing mortality and robustness to observation error. Ecological Indicators, 74:516-525.

Mateu, J. 1997. Methods of assessing and achieving normality applied to environmental data. Environmental Management, 21(5):767-777.

Mather, F. J., J. M. Mason Jr, and A. Jones, eds. 1995. Historical document: Life history and fisheries of Atlantic bluefin tuna. NOAA Technical Memorandum NMFS-SEFSC-370, 165p.

Matsunaga, H. 2009. Tag and release of pelagic shark species by the observers on the Japanese tuna longline vessels in the Atlantic Ocean. ICCAT Collective Volume of Scientific Papers, 64(5):1690-1692.

Maunder, M. N. and A. E. Punt. 2004. Standardizing catch and effort data: a review of recent approaches. Fisheries Research, 70(2):141-159.

May, J. L. and J. G. H. Maxwell. 1986. Field guide to trawl fish from the temperate waters of Australia. CSIRO, Division of Fisheries Research.

McClanahan, T. R. and S. Mangi. 2000. Spillover of exploitable fishes from a marine park and its effect on the adjacent fishery. Ecological Applications, 10(6):1792-1805.

McKinnell, S. and M. P. Seki. 1998. Shark bycatch in the Japanese high seas squid driftnet fishery in the North Pacific Ocean. Fisheries Research, 39(2):127-138.

Medved, R. J. and J. A. Marshall. 1981. Feeding behavior and biology of young sandbar sharks, Carcharhinus plumbeus (pisces, carcharhinidae), in Chincoteague Bay, Virginia. Fishery Bulletin, 79(3):441-447.

Metcalfe, J. and P. Butler. 1984. Changes in activity and ventilation in response to hypoxia in unrestrained, unoperated dogfish (Scyliorhinus canicula L.). Journal of Experimental Biology, 108(1):411-418.

Miller, P. I., K. L. Scales, S. N. Ingram, E. J. Southall, and D. W. Sims. 2015. Basking sharks and oceanographic fronts: quantifying associations in the northeast Atlantic. Functional Ecology, 29(8):1099-1109. URL http://dx.doi.org/ 10.1111/1365-2435.12423.

Monte-Luna, P., D. Lluch-Belda, E. Serviere-Zaragoza, R. Carmona, H. ReyesBonilla, D. Aurioles-Gamboa, J. L. Castro-Aguirre, S. A. Próo, O. Trujillo-Millán, and B. W. Brook. 2007. Marine extinctions revisited. Fish and Fisheries, 8(2):107122.

Montiel, B. H. 1988. Contributión al conocimiento de los elasmobranquios de la zona costera de Tuxpan, Veracruz, México. Bachelor's thesis, Universidad Veracruzana, Facultad de Biología, Tuxpan, Veracruz, México.

Morato, T., S. D. Hoyle, V. Allain, and S. J. Nicol. 2010. Seamounts are hotspots of pelagic biodiversity in the open ocean. Proceedings of the National Academy of Sciences, 107(21):9707-9711.

Moretzsohn, F., J. Brenner, P. Michaud, J. Tunnell, and T. Shirley. 2011. Biodiversity of the Gulf of Mexico Database (BioGoMx). Version 1.0. Harte Research Institute for Gulf of Mexico Studies, Texas A\&M University-Corpus Christi, Corpus Christi, Texas.

Moretzsohn, F., J. A. Sánchez Chávez, and J. Tunnell, Jr. 2016. GulfBase: Resource Database for Gulf of Mexico Research. World Wide Web electronic publication. http://www.gulfbase.org. Accessed: 12 September, 2016.

Morgan, A., P. W. Cooper, T. Curtis, and G. H. Burgess. 2009. Overview of the US east coast bottom longline shark fishery, 1994-2003. Marine Fisheries Review, 71(1):23-38.

Morrissey, J. F. and S. H. Gruber. 1993. Habitat selection by juvenile lemon sharks, Negaprion brevirostris. Environmental Biology of Fishes, 38(4):311-319.

Morzaria-Luna, H. N., C. H. Ainsworth, I. C. Kaplan, P. S. Levin, and E. A. Fulton. 2013. Indirect effects of conservation policies on the coupled humannatural ecosystem of the upper Gulf of California. PLoS One, 8(5):e64085. doi:10.1371/journal.pone.0064085.

Muhling, B. A., J. T. Lamkin, and M. A. Roffer. 2010. Predicting the occurrence of Atlantic bluefin tuna (Thunnus thynnus) larvae in the northern Gulf of Mexico: building a classification model from archival data. Fisheries Oceanography, 19(6):526-539.

Muhling, B. A., S.-K. Lee, J. T. Lamkin, and Y. Liu. 2011. Predicting the effects of climate change on bluefin tuna (Thunnus thynnus) spawning habitat in the Gulf of Mexico. ICES Journal of Marine Science, 68(6):1051-1062.

Muhling, B. A., Y. Liu, S. Lee, J. T. Lamkin, E. Malca, J. Llopiz, G. W. J. Ingram, J. M. Quattro, J. F. Walter, D. K, M. A. Roffer, and F. Muller-Karger. 2014. Past, ongoing and future research on climate change impacts on tuna and billfishes in the western atlantic. ICCAT Collected Volume of Scientific Papers, 71(4):1716-1727.

Mullin, K. D. and H. L. J. 1999. Marine mammals of the northern Gulf of Mexico. In H. Kumpf, K. Steidinger, and K. Sherman, eds., The Gulf of Mexico Large Marine Ecosystem: Assessment, Sustainability, and Management, pp. 269 - 277. Blackwell Science.

Musick, J., S. Berkeley, G. Cailliet, M. Camhi, G. Huntsman, M. Nammack, and M. Warren Jr. 2000a. Protection of marine fish stocks at risk of extinction. AFS policy statement. Fisheries, 25(3):6-8.

Musick, J., G. Burgess, G. Cailliet, M. Camhi, and S. Fordham. 2000b. Management of sharks and their relatives (Elasmobranchii). Fisheries, 25(3):9-13.

Muus, B. J., J. G. Nielsen, P. Dahlstrøm, and B. O. Nyström. 1999. Sea fish. Scandinavian Fishing Year Book.

Myers, R. A., J. K. Baum, T. D. Shepherd, S. P. Powers, and C. H. Peterson. 2007. Cascading effects of the loss of apex predatory sharks from a coastal ocean. Science, 315(5820):1846-1850.

Myers, R. A. and P. Pepin. 1990. The robustness of lognormal-based estimators of abundance. Biometrics, 46(4):1185-1192.

Myers, R. A. and B. Worm. 2003. Rapid worldwide depletion of predatory fish communities. Nature, 423(6937):280-283.

Myers, R. A. and B. Worm. 2005. Extinction, survival or recovery of large predatory fishes. Philosophical Transactions of the Royal Society B: Biological Sciences, 360(1453):13-20.

Nakamura, I. 1985. FAO fisheries catalogue vol. 5. billfishes of the world: An annotated and illustrated catalogue of marlins, sailfishes, spearfishes and swordfishes known to date. Food and Agriculture Organization, 125(5):65.

Nakamura, I. 1995. Gempylidae. escolares. In W. Fischer, F. Krupp, W. Schneider, C. Sommer, K. E. Carpenter, and V. Niem, eds., Guia FAO para Identificatión de Especies para lo Fines de la Pesca. Pacifico Centro-Oriental, vol. 3, pp. 1106-1113. FAO, Rome.

Nakamura, I. and N. V. Parin. 1993. FAO species catalogue. vol. 15: snake mackerels and cutlassfishes of the world (families Gempylidae and Trichiuridae). an annotated and illustrated catalogue of the snake mackerels, snoeks, escolars, gemfishes, sackfishes, domine, oilfish, cutlassfishes, scabbardfishes, hairtails and frostfishes known to date. Food and Agriculture Organization, 125(15):136.

National Centers for Environmental Information. 2015. Fisheries Closures: Deepwater Horizion Support. National Oceanic and Atmospheric Administration. URL https: //www.nodc.noaa.gov/deepwaterhorizon/fisheries \_closures.html.

National Marine Fisheries Service. 1998. Annual Report to Congress on the Status of U.S. Fisheries 1997. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 29 p.

National Marine Fisheries Service. 1999. Annual Report to Congress on the Status of U.S. Fisheries 1998. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 94 p.

National Marine Fisheries Service. 2000. Annual Report to Congress on the Status of U.S. Fisheries 1999. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 104 p.

National Marine Fisheries Service. 2001. Annual Report to Congress on the Status of U.S. Fisheries 2000. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 122 p.

National Marine Fisheries Service. 2002. Annual Report to Congress on the Status of U.S. Fisheries 2001. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 142 p.

National Marine Fisheries Service. 2003. Annual Report to Congress on the Status of U.S. Fisheries 2002. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 156 p.

National Marine Fisheries Service. 2004. Annual Report to Congress on the Status of U.S. Fisheries 2003. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 24 p.

National Marine Fisheries Service. 2005. Annual Report to Congress on the Status of U.S. Fisheries 2004. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 20 p.

National Marine Fisheries Service. 2006. Annual Report to Congress on the Status of U.S. Fisheries 2005. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 20 p.

National Marine Fisheries Service. 2007. Annual Report to Congress on the Status of U.S. Fisheries 2006. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 28 p.

National Marine Fisheries Service. 2008. Annual Report to Congress on the Status of U.S. Fisheries 2007. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 23 p.

National Marine Fisheries Service. 2009. Annual Report to Congress on the Status of U.S. Fisheries 2008. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 23 p .

National Marine Fisheries Service. 2010. Annual Report to Congress on the Status of U.S. Fisheries 2009. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 20 p.

National Marine Fisheries Service. 2011. Annual Report to Congress on the Status of U.S. Fisheries 2010. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 21 p.

National Marine Fisheries Service. 2012. Annual Report to Congress on the Status of U.S. Fisheries 2011. U.S. Dept.Commerce, NOAA, Natl. Mar. Fish. Serv., Silver Spring, MD, 20 p.

National Marine Fisheries Service. 2013. U. S. National Bycatch Report First Edition Update 1. Benaka, L. R., and C. Rilling, and E. E. Seney, and H. Winarsoo, eds. U. S. Dep. Commer. 56 pp.

National Oceanic and Atmospheric Administration. 2012a. Fisheries Office of Science and Technology. Commercial Fisheries Statistics. World Wide Web electronic publication. http://www.st.nmfs.noaa.gov/commercial-fisheries/. Accessed: 16 May 2012.

National Oceanic and Atmospheric Administration. 2012b. Fisheries Office of Science and Technology. Recreational Fisheries Statistics Queries. World Wide Web electronic publication. http://www.st.nmfs.noaa.gov/recreational-fisheries/ data-and-documentation/queries/index. Accessed: 16 May 2012.

National Oceanic and Atmospheric Administration. 2014a. Fisheries Office of Science and Technology. Marine Recreational Information Program. World Wide Web electronic publication. http://www.st.nmfs.noaa.gov/recreational-fisheries/ MRIP/index.

National Oceanic and Atmospheric Administration. 2014b. Recreational Billfish Survey. World Wide Web electronic publication. http://www.sefsc.noaa.gov/rbs/.

National Oceanic and Atmospheric Administration. 2016a. Atlantic HMS Fishery Management Plans and Amendments. URL http://www.nmfs.noaa.gov/sfa/ hms/documents/fmp/index.html.

National Oceanic and Atmospheric Administration. 2016b. Fisheries Office of Science and Technology. Commercial Fisheries Statistics. Landings Background Information. World Wide Web electronic publication. https://www.st.nmfs.noaa.gov/ commercial-fisheries/commercial-landings/landings-background/index. Accessed: 21 October 2016.

National Oceanic and Atmospheric Administration. 2016c. NOAA Fisheries Reminds Pelagic Longline Fishermen that Spring Gulf of Mexico Gear Restricted Areas are in Effect from April 1 through May 31. World Wide Web electronic publication. http://www.nmfs.noaa.gov/sfa/hms/news/news_list/2016/3/a7_ spring_gom_gra_reminder_032316.html. Accessed: 1 April 2016.

Norse, E. A. 1993. Global marine biological diversity: a strategy for building conservation into decision making. Island Press.

Norton, C., I. N. Sarkar, and P. Leary. 2013. Universal Biological Indexer and Organizer (uBio). The Marine Biological Laboratory, Woods Hole Oceanographic Institution. World Wide Web electronic publication. http://www.ubio.org/. Accessed: 22 January 2013.

Nugent, I. and L. Cantral. 2005. Charting a course toward ecosystem-based management in the Gulf of Mexico. Duke Environmental Law $\mathcal{B}$ Policy Forum, 16:267-292.

Oey, L.-Y., T. Ezer, and H.-C. Lee. 2005. Loop Current, rings and related circulation in the Gulf of Mexico: a review of numerical models and future challenges. In W. Sturges and A. Lugo-Fernandez, eds., Circulation in the Gulf of Mexico: Observations and Models, pp. 31-56. American Geophysical Union.

O’Hop, J. and B. Sauls. 2012. Index of Abundance for Pre-Fishery Recruit Red Snapper from Florida Headboat Observer Data. SEDAR31-DW-09. SEDAR, North Charleston, SC. 17 pp.

Olson, D. B., G. L. Hitchcock, A. J. Mariano, C. J. Ashjian, G. Peng, R. W. Nero, and G. P. Podesta. 1994. Life on the edge: marine life and fronts. Oceanography, 7(2):52-60.

Ortega, L. A., M. R. Heupel, P. Van Beynen, and P. J. Motta. 2009. Movement patterns and water quality preferences of juvenile bull sharks (Carcharhinus leucas) in a Florida estuary. Environmental Biology of Fishes, 84(4):361-373.

Ortiz, M. and F. Arocha. 2004. Alternative error distribution models for standardization of catch rates of non-target species from a pelagic longline fishery: billfish species in the Venezuelan tuna longline fishery. Fisheries Research, 70(2):275-297.

Oxenford, H. A. and W. Hunte. 1999. Feeding habits of the dolphinfish (Coryphaena hippurus) in the eastern Caribbean. Scientia Marina, 63(3-4):317-325.

Pacheco, J., D. W. Kerstetter, F. Hazin, H. Hazin, R. Segundo, J. Graves, F. Carvalho, and P. Travassos. 2011. A comparison of circle hook and J hook performance in a western equatorial Atlantic Ocean pelagic longline fishery. Fisheries Research, 107(1):39-45.

Paine, R. T. 1980. Food webs: linkage, interaction strength and community infrastructure. Journal of Animal Ecology, 49(3):666-685.

Palomares, M. L. D. and D. Pauly, eds. 2016. SeaLifeBase. World Wide Web electronic publication. www.sealifebase.org, version (10/2016). Accessed: 12 October, 2016.

Parsons, G. R., E. R. Hoffmayer, and C. Taylor. 2005. Seasonal changes in the distribution and relative abundance of the Atlantic sharpnose shark Rhizoprionodon terraenovae in the north central Gulf of Mexico. Copeia, 2005(4):914-920.

Parsons, T. 1992. The removal of marine predators by fisheries and the impact of trophic structure. Marine Pollution Bulletin, 25(1):51-53.

Pauly, D. 2007. The Sea Around Us Project: documenting and communicating global fisheries impacts on marine ecosystems. AMBIO: a Journal of the Human Environment, 36(4):290-295.

Pauly, D. and V. Christensen. 1995. Primary production required to sustain global fisheries. Nature, 374(6519):255-257.

Pauly, D., V. Christensen, J. Dalsgaard, R. Froese, and F. Torres. 1998. Fishing down marine food webs. Science, 279(5352):860-863.

Pauly, D. and M.-L. Palomares. 2005. Fishing down marine food web: it is far more pervasive than we thought. Bulletin of Marine Science, 76(2):197-212.

Pauly, D. and R. Watson. 2005. Background and interpretation of the marine trophic indexas a measure of biodiversity. Philosophical Transactions of the Royal Society of London B: Biological Sciences, 360(1454):415-423.

Pauly, D., R. Watson, and J. Alder. 2005. Global trends in world fisheries: impacts on marine ecosystems and food security. Philosophical Transactions of the Royal Society B: Biological Sciences, 360(1453):5-12.

Pecquerie, L., L. Drapeau, P. Fréon, J. Coetzee, R. Leslie, and M. Griffiths. 2004. Distribution patterns of key fish species of the southern Benguela ecosystem: an approach combining fishery-dependent and fishery-independent data. African Journal of Marine Science, 26(1):115-139.

Peel, E., R. Nelson, and C. P. Goodyear. 2003. Managing Atlantic marlin as bycatch under ICCAT. The fork in the road: recovery or collapse. Marine and Freshwater Research, 54(4):575-584.

Pérez-Jiménez, J. C. and I. Mendez-Loeza. 2015. The small-scale shark fisheries in the southern Gulf of Mexico: understanding their heterogeneity to improve their management. Fisheries Research, 172:96-104.

Perry, H. M. and S. J. VanderKooy, eds. 2015. The Blue Crab Fishery of the Gulf of Mexico: A Regional Management Plan - 2015 Revision. Publication No. 243, Blue Crab Technical Task Force, Gulf States Marine Fisheries Commission, Ocean Springs, Mississippi.

Perryman, H. A., D. Die, and E. Babcock. 2015. An Atlantis Ecosystem Model for the Gulf of Mexico Supporting Integrated Ecosystem Assessment, chap. Catch Reconstruction, pp. 37-43. NOAA Technical Memorandum NMFS-SEFSC-676, 149p.

Petersen, S., M. Honig, P. Ryan, L. Underhill, and L. J. Compagno. 2009. Pelagic shark bycatch in the tuna-and swordfish-directed longline fishery off southern Africa. African Journal of Marine Science, 31(2):215-225.

Piet, G., H. Van Overzee, and M. Pastoors. 2010. The necessity for response indicators in fisheries management. ICES Journal of Marine Science, 67(3):559-566.

Pikitch, E. K., C. Santora, E. A. Babcock, A. Bakun, R. Bonfil, D. O. Conover, P. Dayton, P. Doukakis, D. Fluharty, B. Heneman, E. D. Houde, J. Link, P. A. Livingston, M. Mangel, M. K. McAllister, J. Pope, and K. J. Sainsbury. 2004. Ecosystem-based fishery management. Science, 305(5682):346-347.

Piñeiro, G., S. Perelman, J. P. Guerschman, and J. M. Paruelo. 2008. How to evaluate models: observed vs. predicted or predicted vs. observed? Ecological Modelling, 216(3):316-322.

Piovano, S., S. Clò, and C. Giacoma. 2010. Reducing longline bycatch: the larger the hook, the fewer the stingrays. Biological Conservation, 143(1):261-264.

Plagányi, É. E. 2007. Models for an ecosystem approach to fisheries. Tech. Rep. Technical Paper 477, FAO, Fisheries and Aquaculture, Rome, Italy.

Podestá, G. P., J. A. Browder, and J. J. Hoey. 1993. Exploring the association between swordfish catch rates and thermal fronts on US longline grounds in the western North Atlantic. Continental Shelf Research, 13(2):253-277.

Pollack, A. G. and G. W. Ingram Jr. 2014. Smoothhound Abundance Indices from NFMS Small Pelagics Surveys in the Northern Gulf of Mexico. SEDAR39-DW-08. SEDAR, North Charleston, SC. 14 pp.

Pomeroy, R., L. Garces, M. Pido, and G. Silvestre. 2010. Ecosystem-based fisheries management in small-scale tropical marine fisheries: emerging models of governance arrangements in the Philippines. Marine Policy, 34(2):298-308.

Powers, J. E. 2010. Fisheries: measuring biodiversity in marine ecosystems. Nature, 468(7322):385-386.

Powers, J. E. and M. H. Monk. 2010. Current and future use of indicators for ecosystem based fisheries management. Marine Policy, 34(3):723-727.

Priede, I. G. and P. I. Miller. 2009. A basking shark (Cetorhinus maximus) tracked by satellite together with simultaneous remote sensing ii: new analysis reveals orientation to a thermal front. Fisheries Research, 95(2):370-372.

Prince, E. D., A. R. Bertolino, and A. M. Lopez. 1989. A comparison of fishing success and average weights of blue marlin and white marlin landed by the recreational fishery in the western Atlantic Ocean, Gulf of Mexico, and Caribbean Sea, 1972-1986. In R. H. Stroud, ed., Planning the future of billfishes, research and management in the 90's and beyond. Part 1. Fishery and stock synopses, data needs, and management, pp. 159-178. National Coalition for Marine Conservation, Inc.

Prince, E. D. and C. P. Goodyear. 2006. Hypoxia-based habitat compression of tropical pelagic fishes. Fisheries Oceanography, 15(6):451-464.

Punt, A. E., D. S. Butterworth, C. L. de Moor, J. A. A. De Oliveira, and M. Haddon. 2016. Management strategy evaluation: best practices. Fish and Fisheries, 17(2):303-334. URL http://dx.doi.org/10.1111/faf. 12104.

Punt, A. E., T. I. Walker, B. L. Taylor, and F. Pribac. 2000. Standardization of catch and effort data in a spatially-structured shark fishery. Fisheries Research, 45(2):129-145.

Pya, N. and S. N. Wood. 2016. A note on basis dimension selection in generalized additive modelling. Nazarbayev University and University of Bristol.

Queiroz, N., N. E. Humphries, G. Mucientes, N. Hammerschlag, F. P. Lima, K. L. Scales, P. I. Miller, L. L. Sousa, R. Seabra, and D. W. Sims. 2016. Ocean-wide tracking of pelagic sharks reveals extent of overlap with longline fishing hotspots. Proceedings of the National Academy of Sciences, 113(6):1582-1587.

Queiroz, N., N. E. Humphries, L. R. Noble, A. M. Santos, and D. W. Sims. 2012. Spatial dynamics and expanded vertical niche of blue sharks in oceanographic fronts reveal habitat targets for conservation. PLoS One, 7(2):e32374. doi:10.1371/journal.pone. 0032374 .

R Core Team. 2014. R: A Language and Environment for Statistical Computing. R Foundation for Statistical Computing, Vienna, Austria. URL http: //www.R-project.org/.

Rabalais, N. N., R. E. Turner, Q. Dortch, D. Justic, V. J. Bierman Jr, and W. J. Wiseman Jr. 2002a. Nutrient-enhanced productivity in the northern Gulf of Mexico: past, present and future. In E. Orive, M. Elliott, and V. N. de Jonge, eds., Nutrients and Eutrophication in Estuaries and Coastal Waters, pp. 39-63. Springer.

Rabalais, N. N., R. E. Turner, and W. J. Wiseman Jr. 2002b. Gulf of Mexico hypoxia, aka "the dead zone". Annual Review of Ecology and Systematics, 33:235-263.

Rice, P. H., J. E. Serafy, D. Snodgrass, and E. D. Prince. 2012. Performance of nonoffset and 10 offset 18/0 circle hooks in the United States pelagic longline fishery. Bulletin of Marine Science, 88(3):571-587.

Richards, W. J. 1990. Results of a review of the US bluefin tuna larval assessment with a brief response. ICCAT Collected Volume of Scientific Papers, 32(2):240-247.

Riede, K. 2004. Global register of migratory species: from global to regional scales: final report of the R $\dot{B}$ D-Projekt 80805081 . Federal Agency for Nature Conservation, Bonn, Germany.

Roberts, C. M. 1997. Ecological advice for the global fisher crisis. Trends in Ecology E Evolution, 12(1):35-38.

Robson, D. 1966. Estimation of the relative fishing power of individual ships. ICNAF Research Bulletin, 3:5-14.

Rochet, M.-J. and V. M. Trenkel. 2003. Which community indicators can measure the impact of fishing? A review and proposals. Canadian Journal of Fisheries and Aquatic Sciences, 60(1):86-99.

Rogan, E. and M. Mackey. 2007. Megafauna bycatch in drift nets for albacore tuna (Thunnus alalunga) in the NE Atlantic. Fisheries Research, 86(1):6-14.

Rooker, J. R., J. R. Simms, R. D. Wells, S. A. Holt, G. J. Holt, J. E. Graves, and N. B. Furey. 2012. Distribution and habitat associations of billfish and swordfish larvae across mesoscale features in the Gulf of Mexico. PLoS One, 7(4):e34180. doi:10.1371/journal.pone. 0034180 .

Ruppert, D. 2002. Selecting the number of knots for penalized splines. Journal of Computational and Graphical Statistics, 11(4):735-757.

Russ, G. R., A. J. Cheal, A. M. Dolman, M. J. Emslie, R. D. Evans, I. Miller, H. Sweatman, and D. H. Williamson. 2008. Rapid increase in fish numbers follows creation of world's largest marine reserve network. Current Biology, 18(12):R514R515.

Safina, C. and D. H. Klinger. 2008. Collapse of bluefin tuna in the Western Atlantic. Conservation Biology, 22(2):243-246.

SAGARPA. 1980. Anuario Estadístico Pesquero 1980. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1981. Anuario Estadístico Pesquero 1981. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1982. Anuario Estadístico Pesquero 1982. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1983. Anuario Estadístico Pesquero 1983. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1984. Anuario Estadístico Pesquero 1984. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1985. Anuario Estadístico Pesquero 1985. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1986. Anuario Estadístico Pesquero 1986. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1987. Anuario Estadístico Pesquero 198\%. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1988. Anuario Estadístico Pesquero 1988. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1989. Anuario Estadístico Pesquero 1989. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1990. Anuario Estadístico Pesquero 1990. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1991. Anuario Estadístico Pesquero 1991. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1992. Anuario Estadístico Pesquero 1992. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1993. Anuario Estadístico Pesquero 1993. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1994. Anuario Estadístico Pesquero 1994. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1995. Anuario Estadístico Pesquero 1995. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1996. Anuario Estadístico Pesquero 1996. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1997. Anuario Estadístico Pesquero 1997. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1998. Anuario Estadístico Pesquero 1998. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 1999. Anuario Estadístico Pesquero 1999. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2000. Anuario Estadístico Pesquero 2000. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2001. Anuario Estadístico de Acuacultura y Pesca 2001. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2002. Anuario Estadístico de Acuacultura y Pesca 2002. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2003. Anuario Estadístico de Acuacultura y Pesca 2003. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2004. Anuario Estadístico de Acuacultura y Pesca 2004. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2005. Anuario Estadístico de Acuacultura y Pesca 2005. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2006. Anuario Estadístico de Acuacultura y Pesca 2006. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2007. Anuario Estadístico de Acuacultura y Pesca 2007. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2008. Anuario Estadístico de Acuacultura y Pesca 2008. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2009. Anuario Estadístico de Acuacultura y Pesca 2009. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2010. Anuario Estadístico de Acuacultura y Pesca 2010. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2011. Anuario Estadístico de Acuacultura y Pesca 2011. Comisión Nacional de Acuaultura y Pesca (CONAPESCA), Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA), México, D.F.

SAGARPA. 2016. Comisión Nacional de Acuacultura y Pesca (CONAPESCA). World Wide Web electronic publication. http://www.sagarpa.gob.mx/quienesomos/ datosabiertos/conapesca/Paginas/default.aspx. Accessed: 16 May 2012.

Sainsbury, K. 1998. Living marine resource assessment for the 21st century: what will be needed and how will it be provided. In F. Funk, T. J. Quinn II, J. Heifetz, J. N. Ianelli, J. E. Powers, J. F. Schweigert, P. J. Sullivan, and C.-I. Zhang, eds., Fishery Stock Assessment Models, pp. 1-40. Alaska Sea Grant College Program Report No. AK-SG-98-01. University of Alaska, Fairbanks, Alaska, USA.

Sainsbury, K. J., A. E. Punt, and A. D. Smith. 2000. Design of operational management strategies for achieving fishery ecosystem objectives. ICES Journal of Marine Science, 57(3):731-741.

Salas, S., R. Chuenpagdee, A. Charles, and J. C. Seijo. 2011. Coastal Fisheries of Latin America and the Caribbean. Technical Paper 544. FAO, Fisheries and Aquaculture. Rome, Italy.

Samhouri, J. F., A. J. Haupt, P. S. Levin, J. S. Link, and R. Shuford. 2014. Lessons learned from developing integrated ecosystem assessments to inform marine ecosystem-based management in the USA. ICES Journal of Marine Science, 71(5):1205-1215. URL http://dx.doi.org/10.1093/icesjms/fst141.

Saul, S., J. Walter, D. Die, D. Naar, and B. Donahue. 2013. Modeling the spatial distribution of commercially important reef fishes on the West Florida Shelf. Fisheries Research, 143:12-20.

Schick, R. and M. Lutcavage. 2009. Inclusion of prey data improves prediction of bluefin tuna (Thunnus thynnus) distribution. Fisheries Oceanography, 18(1):7781.

Schirripa, M. J., B. Allee, S. Cross, C. Kelble, and A. R. Parsons. 2013. Progress towards an integrated ecosystem assessment for the Gulf of Mexico. ICCAT Collective Volume of Scientific Papers, 69(4):1867-1875.

Schmitz, W. J. 2005. Cyclones and westward propagation in the shedding of anticyclonic rings from the Loop Current. In W. Sturges and A. Lugo-Fernandez, eds., Circulation in the Gulf of Mexico: Observations and Models, pp. 241-261. American Geophysical Union.

Seber, G. A. F. 1982. The estimation of animal abundance. Griffin London.

SEDAR. 2005. Stock Assessment of SEDAR 8: Caribbean Spiny Lobster. Southeast Data, Assessment, and Review, Key West, FL.

SEDAR. 2007. Stock Assessment of SEDAR 13: small coastal shark complex, Atlantic sharpnose, blacknose, bonnethead, and finetooth shark. Southeast Data, Assessment, and Review, North Charleston, SC.

Serafy, J. E., G. A. Diaz, E. D. Prince, E. S. Orbesen, and C. M. Legault. 2004. Atlantic blue marlin, Makaira nigricans, and white marlin, Tetrapterus albidus, bycatch of the Japanese pelagic longline fishery, 1960-2000. Marine Fisheries Review, 66(2):9-20.

Sethi, S. A., T. A. Branch, and R. Watson. 2010. Global fishery development patterns are driven by profit but not trophic level. Proceedings of the National Academy of Sciences, 107(27):12163-12167.

Shackell, N. L., K. T. Frank, J. A. Fisher, B. Petrie, and W. C. Leggett. 2010. Decline in top predator body size and changing climate alter trophic structure in an oceanic ecosystem. Proceedings of the Royal Society of London B: Biological Sciences, 277(1686):1353-1360.

Shepherd, T. D. and R. A. Myers. 2005. Direct and indirect fishery effects on small coastal elasmobranchs in the northern Gulf of Mexico. Ecology Letters, 8(10):10951104.

Shin, Y.-J., M.-J. Rochet, S. Jennings, J. G. Field, and H. Gislason. 2005. Using size-based indicators to evaluate the ecosystem effects of fishing. ICES Journal of Marine Science, 62(3):384-396.

Shin, Y.-J. and L. J. Shannon. 2010. Using indicators for evaluating, comparing, and communicating the ecological status of exploited marine ecosystems. 1. The IndiSeas project. ICES Journal of Marine Science, 67(4):686-691.

Shin, Y.-J., L. J. Shannon, A. Bundy, M. Coll, K. Aydin, N. Bez, J. L. Blanchard, M. de Fatima Borges, I. Diallo, E. Diaz, et al. 2010. Using indicators for evaluating, comparing, and communicating the ecological status of exploited marine ecosystems. 2. Setting the scene. ICES Journal of Marine Science, 67(4):692-716.

Sims, D. W. and V. A. Quayle. 1998. Selective foraging behaviour of basking sharks on zooplankton in a small-scale front. Nature, 393(6684):460-464.

Sims, D. W., E. J. Southall, V. A. Quayle, and A. M. Fox. 2000. Annual social behaviour of basking sharks associated with coastal front areas. Proceedings of the Royal Society of London B: Biological Sciences, 267(1455):1897-1904.

Smale, M. J. 2008. Pelagic shark fisheries in the Indian Ocean. In M. D. Camhi, E. K. Pikitch, and E. A. Babcock, eds., Sharks of the Open Ocean: Biology, Fisheries and Conservation, pp. 247-259. Blackwell Publishing.

Smith, A. D. M. 1994. Management strategy evaluation: the light on the hill. In D. A. Hancock, ed., Population dynamics for fisheries management: Australian Society for Fish Biology Workshop proceedings, pp. 249-253.

Southeast Data, Assessment, and Review. 2016. SEDAR 49 Gulf of Mexico Datalimited Species. Southeast Data, Assessment, and Review (SEDAR), North Charleston, SC.

Stachowicz, J. J., J. F. Bruno, and J. E. Duffy. 2007. Understanding the effects of marine biodiversity on communities and ecosystems. Annual Review of Ecology, Evolution, and Systematics, 38:739-766.

Stanley, D. R. and C. A. Wilson. 2004. Effect of hypoxia on the distribution of fishes associated with a petroleum platform off coastal Louisiana. North American Journal of Fisheries Management, 24(2):662-671.

Stevens, J. D., R. Bonfil, N. K. Dulvy, and P. A. Walker. 2000. The effects of fishing on sharks, rays, and chimaeras (chondrichthyans), and the implications for marine ecosystems. ICES Journal of Marine Science, 57(3):476-494.

Su, N.-J., C.-L. Su, A. E. Punt, and S.-Z. Yeh. 2008. Environmental and spatial effects on the distribution of blue marlin (Makaira nigricans) as inferred from data for longline fisheries in the Pacific Ocean. Fisheries Oceanography, 17(6):432-445.

Swain, D. P., A. F. Sinclair, and J. M. Hanson. 2007. Evolutionary response to sizeselective mortality in an exploited fish population. Proceedings of the Royal Society of London B: Biological Sciences, 274(1613):1015-1022.

Swets, J. A. 1988. Measuring the accuracy of diagnostic systems. Science, 240(4857):1285-1293.

Takahashi, M., H. Okamura, K. Yokawa, and M. Okazaki. 2003. Swimming behaviour and migration of a swordfish recorded by an archival tag. Marine and Freshwater Research, 54(4):527-534.

Teo, S. L., A. M. Boustany, and B. A. Block. 2007. Oceanographic preferences of Atlantic bluefin tuna, Thunnus thynnus, on their Gulf of Mexico breeding grounds. Marine Biology, 152(5):1105-1119.

Terborgh, J., L. Lopez, P. Nunez, M. Rao, G. Shahabuddin, G. Orihuela, M. Riveros, R. Ascanio, G. H. Adler, T. D. Lambert, et al. 2001. Ecological meltdown in predator-free forest fragments. Science, 294(5548):1923-1926.

Ubeda, A. J., C. Simpfendorfer, and M. Heupel. 2009. Movements of bonnetheads, Sphyrna tiburo, as a response to salinity change in a Florida estuary. Environmental Biology of Fishes, 84(3):293-303.
U.S. Congress. 1996. Magnuson-Stevens Fishery Conservation and Management Act, Public Law 94-265. Washington, D.C.: U.S. Congress.

Uychiaoco, A. J., H. O. Arceo, S. J. Green, T. Margarita, P. A. Gaite, and P. M. Aliño. 2005. Monitoring and evaluation of reef protected areas by local fishers in the Philippines: tightening the adaptive management cycle. Biodiversity $\mathcal{B}$ Conservation, 14(11):2775-2794.

VanderKooy, S. J., ed. 2013. GDAR01 Gulf of Mexico blue crab stock assessment report. Blue Crab Technical Task Force, Gulf States Marine Fisheries Commission, Publication No. 215, 313 pp.

Venables, W. N. and C. M. Dichmont. 2004. GLMs, GAMs and GLMMs: an overview of theory for applications in fisheries research. Fisheries Research, 70(2):319-337.

Vukovich, F. M. and G. A. Maul. 1985. Cyclonic eddies in the eastern Gulf of Mexico. Journal of Physical Oceanography, 15(1):105-117.

Wall, C. C., F. E. Muller-Karger, and M. A. Roffer. 2009. Linkages between environmental conditions and recreational king mackerel (Scomberomorus cavalla) catch off west-central Florida. Fisheries Oceanography, 18(3):185-199.

Ward, P., R. A. Myers, and W. Blanchard. 2004. Fish lost at sea: the effect of soak time on pelagic longline catches. Fishery Bulletin, 102(1):179-195.

Watson, J. W., S. P. Epperly, A. K. Shah, and D. G. Foster. 2005. Fishing methods to reduce sea turtle mortality associated with pelagic longlines. Canadian Journal of Fisheries and Aquatic Sciences, 62(5):965-981.

Weidner, D. M., G. E. Laya, and J. A. Serano, eds. 2001. World swordfish fisheries: an analysis of swordfish fisheries, market trends and trade patterns, past-presentfuture. NOAA Technical Memorandum NMFS-F/SPO-53.

Weng, K., M. Stokesbury, A. Boustany, A. Seitz, S. Teo, S. Miller, and B. Block. 2009. Habitat and behaviour of yellowfin tuna Thunnus albacares in the Gulf of Mexico determined using pop-up satellite archival tags. Journal of Fish Biology, 74(7):1434-1449.

Whitehead, P. J. P. and R. R. Vergara. 1978. Megalopidae. In W. Fischer, ed., FAO species identification sheets for fishery purposes. Western Central Atlantic (Fishing Area 31), vol. 3. FAO, Rome.

Whitfield, P. E., J. A. Hare, A. W. David, S. L. Harter, R. C. Munoz, and C. M. Addison. 2007. Abundance estimates of the Indo-Pacific lionfish Pterois volitans miles complex in the western North Atlantic. Biological Invasions, 9(1):53-64.

Wise, G., J. M. Mulvey, and G. M. Renshaw. 1998. Hypoxia tolerance in the epaulette shark (Hemiscyllium ocellatum). Journal of Experimental Zoology, 281(1):1-5.

Wiseman, W. J., W. Sturges, and H. Kumpf. 1999. Physical oceanography of the Gulf of Mexico: processes that regulate its biology. In H. Kumpf, K. Steidinger, and K. Sherman, eds., The Gulf of Mexico Large Marine Ecosystem: Assessment, Sustainability, and Management, pp. 77-92. Blackwell Science.

Wood, S. 2006a. Generalized additive models: an introduction with $R$. CRC press.
Wood, S. and M. S. Wood. 2015. Package mgcv. R package version, pp. 1-7.
Wood, S. N. 2006b. Generalized additive models: an introduction with R. CRC press.
Wood, S. N. 2011. Fast stable restricted maximum likelihood and marginal likelihood estimation of semiparametric generalized linear models. Journal of the Royal Statistical Society: Series B (Statistical Methodology), 73(1):3-36.

Wood, S. N. and N. H. Augustin. 2002. GAMs with integrated model selection using penalized regression splines and applications to environmental modelling. Ecological Modelling, $\mathbf{1 5 7}(2): 157-177$.

Woodward, G., B. Ebenman, M. Emmerson, J. M. Montoya, J. M. Olesen, A. Valido, and P. H. Warren. 2005. Body size in ecological networks. Trends in Ecology $\xi^{6}$ Evolution, 20(7):402-409.

Worm, B., E. B. Barbier, N. Beaumont, J. E. Duffy, C. Folke, B. S. Halpern, J. B. Jackson, H. K. Lotze, F. Micheli, S. R. Palumbi, et al. 2006. Impacts of biodiversity loss on ocean ecosystem services. Science, $\mathbf{3 1 4 ( 5 8 0 0 ) : 7 8 7 - 7 9 0 .}$

Worm, B., B. Davis, L. Kettemer, C. A. Ward-Paige, D. Chapman, M. R. Heithaus, S. T. Kessel, and S. H. Gruber. 2013. Global catches, exploitation rates, and rebuilding options for sharks. Marine Policy, 40:194-204.

Worm, B., H. K. Lotze, and R. A. Myers. 2003. Predator diversity hotspots in the blue ocean. Proceedings of the National Academy of Sciences, 100(17):9884-9888.

Yáñez-Arancibia, A. and J. W. Day. 2004. The Gulf of Mexico: towards an integration of coastal management with large marine ecosystem management. Ocean \& Coastal Management, 47(11):537-563.

Yáñez-Arancibia, A., J. W. Day, and E. Reyes. 2013. Understanding the coastal ecosystem-based management approach in the Gulf of Mexico. Journal of Coastal Research, 63(sp1):244-262.

Yoskowitz, D. W. 2009. The productive value of the Gulf of Mexico. In D. L. Felder and D. K. Camp, eds., Gulf of Mexico: Origin, Waters, and Biota: Volume 2, Ocean and Coastal Economy, pp. 21-27. Texas A\&M University Press.

Yuan, D. 2002. A numerical study of barotropicly forced intrusion in DeSoto Canyon. Journal of Geophysical Research: Oceans, 107(C2):1-15. URL http://dx.doi. org/10.1029/2001JC000793.

Yurkowski, D. J., S. H. Ferguson, C. A. Semeniuk, T. M. Brown, D. C. Muir, and A. T. Fisk. 2016. Spatial and temporal variation of an ice-adapted predators feeding ecology in a changing arctic marine ecosystem. Oecologia, 180(3):631-644.

Zeller, D., M. Darcy, S. Booth, M. Lowe, and S. Martell. 2008. What about recreational catch? Potential impact on stock assessment for Hawaii's bottomfish fisheries. Fisheries Research, 91(1):88-97.

Zhang, H., S. A. Ludsin, D. M. Mason, A. T. Adamack, S. B. Brandt, X. Zhang, D. G. Kimmel, M. R. Roman, and W. C. Boicourt. 2009. Hypoxia-driven changes in the behavior and spatial distribution of pelagic fish and mesozooplankton in the northern Gulf of Mexico. Journal of Experimental Marine Biology and Ecology, 381:S80-S91.


[^0]:    ${ }^{\dagger}$ Averages for both sea surface and sea bottom
    ${ }^{\ddagger}$ See Appendix B for details

[^1]:    Functional group is not considered pelagic-based fish

    * Functional group does not have enough data to fit a statistical model

