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<A>Title:

Exploring trends in abundance of Young-of-the-Year, and Age-1 Atlantic Croaker
(*Micropogonias undulatus*), Black Drum (*Pogonias cromis*), Spot (*Leiostomus xanthurus*) and
Weakfish (*Cynoscion regalis*) in relation to salinity, temperature and large-scale climatic signals
in a Mid-Atlantic estuary

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48

49 <A>Abstract

50 Atlantic Croaker (*Micropogonias undulatus*), Black Drum (*Pogonias cromis*), Spot (*Leiostomus*
51 *xanthurus*) and Weakfish (*Cynoscion regalis*) have shown species-specific varying trends in
52 abundance, despite general declines in commercial landings throughout the Delaware River
53 Estuary. Identifying how environmental factors and climatic processes affect fishes at multiple
54 life stages is needed to enhance the precision of regulatory actions for managed species. Species'
55 area and age specific indices were compared with depth, salinity, temperature, the Atlantic
56 Multidecadal Oscillation (AMO) and the North Atlantic Oscillation (NAO) to explore potential

57 relationships, as well as temporal and spatial parameters. Our results demonstrate that station (15
58 indices), salinity (14 indices), the AMO and NAO (13 indices each), depth and temperature (12
59 indices each), served as important components for models of abundance of seventeen species,
60 age, and area specific combination from 1991-2016. Significant time series trends were detected
61 in six of the modelled indices of abundance, including Bay-wide indices of Age-1 Weakfish
62 (decline), and YOY Atlantic Croaker (increase); as well as YOY Atlantic Croaker (increase) and
63 Weakfish (decline) in Delaware, and Age-1 Spot (decline) and Weakfish (increase) in Delaware.
64 Our results demonstrate how multiple fixed station surveys can be combined to quantitatively
65 assess environmental and climatic effects correlated with species-age-area specific levels of
66 abundance, suggesting climatic signals are affecting smaller scale environmental variables that in
67 turn affect relative abundance.

68

69 <A>Introduction

70 Understanding how environmental phenomena affect fisheries is a significant challenge
71 that requires immediate attention to identify how these processes impact managed species.
72 Recognizing the effects of environmental processes on fish stocks at multiple life stages can
73 provide new information to enhance precision of regulatory action and management efforts.
74 Previous studies have suggested that environmental and ecological factors need to be accounted
75 for to better understand the behavior of fish stocks and more accurately assess population levels
76 (Garcia and Cochrane 2005; Lehodey et al. 2006; Methot and Richard 2015). Environmental
77 processes, such as El Niño and La Niña in the Pacific Ocean; the North Atlantic Oscillation
78 (NAO) and the Atlantic Multidecadal Oscillation (AMO) in the Atlantic Ocean, vary over large
79 spatiotemporal scales, causing varying responses amongst fish growth rates (Fiedler et al. 1986;
80 Ottersen et al. 2001; Jonsson and Jonsson 2004; Izzo and Zydlewski 2017), predation (Stenseth
81 et al. 2002; Nye et al. 2009; Yasumiishi et al. 2016), recruitment and other population dynamics
82 (Hare and Able 2007; Nye et al. 2009; Large et al. 2013; Edwards et al. 2013; Harris et al. 2014;
83 Nye et al. 2014; Buchheister et al. 2016). Climatic processes have been linked to changing
84 physiochemical properties in estuaries (Irby et al. 2018) which can have confounding effects on
85 the fish species that rely on certain conditions seasonally within estuaries to survive (Barletta et
86 al. 2005). Understanding how components of multiple large-scale environmental phenomena

87 impact marine nekton is critical to address trends in annual variability of recruitment and
88 survival. In the northwest Atlantic Ocean, two large-scale atmospheric processes, the NAO and
89 AMO, are likely influencing the abundance of the Delaware River Estuary fish community in
90 unknown ways.

91 The NAO is observable in the Northern Hemisphere throughout the year with over one
92 third of sea-level-pressure variance occurring during winter months (December-February) over
93 the North-Atlantic (Hurrell and Deser 2010). The NAO is characteristically defined as an annual
94 index and a winter index due to the significant variances in sea-level-pressure observed in winter
95 months (Jing et al. 2019). The NAO is defined more specifically as a change in pressure
96 differences between the subtropical atmospheric high-pressure zone over the Azores and the
97 atmospheric low-pressure zone over Iceland (Otterson et al. 2001; Durkee et al. 2008). However,
98 NAO indices have been shown to influence many atmospheric and oceanographic conditions
99 such as sea surface temperature (SST), storms and precipitation, cloud cover, hydrographic
100 characteristics, mixed-layered depths, and circulation patterns (Barnston and Livezey 1987;
101 Ostermeier and Wallace 2003; Durkee et al. 2008; Hurrell and Deser 2010). Unlike El Niño/La
102 Niña and the AMO, the NAO has no defined change in periodicity between its warm and cold
103 phases (Otterson et al. 2001; Hurrell and Deser 2010). Warm (positive) phases of the NAO index
104 are defined as an intense Icelandic low and a strong Azores high-pressure gradient, with the
105 pressure difference delivering powerful winter storms crossing the Atlantic Ocean in a northern
106 direction, conversely cold (negative) phases have a weak pressure grade and produce weaker
107 storms that tend to move west to east (Hurrell 1995; Otterson et al. 2001). The NAO influences
108 atmospheric pressures causing variations in precipitation patterns, wind events, storm events and
109 climate fluctuations (Durkee et al. 2008).

110 The AMO is characterized as varying SST anomalies that encompass the North Atlantic
111 and is considered the dominant pattern of SST variability within the region (Schlesinger and
112 Ramankutty 1994; Dong et al. 2006; Knudsen et al. 2011; Alexander et al. 2014; Harris et al.
113 2014; Nye et al. 2014). Unlike the NAO, the AMO has oscillating phases that tend to switch
114 every 65-70 years based on approximately 130 years of observed and reconstructed SST data
115 (Nye et al. 2014). The AMO is defined as having warm and cool phases, with warm phases
116 described as having above average ocean temperatures and a shifting of the intertropical
117 convergence zone from the south to the north, where precipitation expands spatially (Enfield et

118 al. 2001; Knudsen et al. 2011; Nogueira et al. 2013; Nye et al. 2014). Warm phases of the AMO
119 typically cause the mixed layer depth to be much shallower and are linked to variations in
120 oceanic pressure gradients, wind speed and direction across the North Atlantic (Nye et al. 2014).
121 Conversely, the cool phase of the AMO is described as causing opposite anomalies as seen with
122 the warm phase respectively (Dijkstra et al. 2006; Alexander et al. 2014).

123 Successful recruitment of young-of-the-year (YOY) fishes has been linked to
124 environmental processes (e.g. Boehlert and Mundy 1988; Lehodey et al. 2006; Cury et al. 2008).
125 Previous studies have shown that nekton size (Hale & Targett 2018), temperature (Witting et al.
126 1999; Lankford and Targett 2001; Hare and Able 2007; Carassou et al. 2011; Yasumiishi et al.
127 2016), salinity (Lankford and Targett 1994; Able et al. 2009), freshwater input (Reist et al. 2006;
128 Carassou et al. 2011), flow (Dunning et al. 2009) and wind speed and direction (Schieler et al.
129 2014; Nye et al. 2014) can significantly affect transport, growth and survival of multiple life
130 stages of marine nekton. The NAO and the AMO have been linked to trends in fisheries
131 production, distribution, and abundance (Lehodey et al. 2006). The AMO has been correlated
132 with spatial distribution and spawning success for fish species in the Mid- and North Atlantic
133 Ocean including Striped Bass (*Morone saxatilis*; O'Connor et al. 2012) and American Shad
134 (*Alosa sapidissima*; O'Connor et al. 2012); Atlantic Mackerel (*Scomber scombrus*; Overholtz et
135 al. 2011); Atlantic Croaker (*Micropogonias undulatus*; Hare and Able 2007); Atlantic Salmon
136 (*Salmo salar*; Izzo and Zydlewski 2017). Lankford and Targett (2001) found that severe cold
137 winter temperatures were correlated to weak year-class strength of age-0 Atlantic Croaker in
138 estuaries along the Mid-Atlantic Bight. In that study, temperature was found to be a growth-
139 limiting factor in juvenile Atlantic Croaker within the Delaware Bay due to feeding temperature
140 preferences and acute thermal stress leading to significantly increased mortality at temperatures
141 below 5° C. Other environmental factors, including wind patterns, have been found to have a
142 positive correlation with larval fish ingress into estuaries within the Mid-Atlantic Bight. Schieler
143 et al. (2014) found that wind direction and speed were correlated with the ingress of larval fish
144 including Atlantic Menhaden (*Brevoortia tyrannus*), Summer Flounder (*Paralichthys dentatus*)
145 and Atlantic Croaker in Delaware Bay.

146 The mechanistic link between ocean climatic processes and year-class strength has some
147 regional stock assessment teams beginning to examine effects that atmospheric and
148 oceanographic conditions have on species-specific stock assessments (e.g., Sablefish

149 (*Anoplopoma fimbria*; Schirripa et al. 2009) and Weakfish (*Cynoscion regalis*; Jiao et al. 2012).
150 Responses of Atlantic Croaker and Weakfish to changing oscillations of the NAO have been
151 observed along the continental shelf between the North Carolina-Virginia border and Cape
152 Canaveral, Florida (Roberts et al. 2019). The findings by Roberts et al. (2019), suggest that the
153 predictability of species distributions potentially shift with phases of the NAO. Along the Eastern
154 US continental shelf, Atlantic Mackerel have been found to occupy different areas annually in
155 response to large scale climatic shifts associated with the phases of the AMO but were
156 historically located in relatively narrow bands of optimal habitat within the Mid-Atlantic region
157 during winter (Overholtz et al. 2011). However, Overholtz et al. (2011) found that overwintering
158 Atlantic Mackerel shifted their distribution much further North and East as they followed their
159 optimal habitat to higher latitudes in response to shifts in environmental conditions over a period
160 of decades. In the Hudson River Estuary, YOY Striped Bass and American Shad were both
161 found to exhibit strong relationships to the AMO; freshwater flow, water temperature, and the
162 AMO explained 46% of the total variance in the species stage specific abundance patterns within
163 the Hudson River Estuary (O'Connor et al. 2012). Juvenile American Shad were negatively
164 correlated with the AMO in the Hudson River Estuary, while Striped Bass were positively
165 correlated with the AMO (O'Connor et al. 2012). A recent study developed indices of Atlantic
166 Menhaden recruitment using independent surveys from the southern region of the Northeast
167 Atlantic and correlated the indices to the AMO (Buchheister et al. 2016). Even though different
168 correlations to the AMO were discovered in Chesapeake Bay and Southern New England
169 regions, results showed that the AMO had been one of the greatest predictors of Atlantic
170 Menhaden recruitment patterns throughout the entire Atlantic coast (Buchheister et al. 2016).

171 The Delaware River Estuary is a large coastal plain estuary, which borders the states of
172 Delaware, New Jersey, and Pennsylvania in the Mid-Atlantic Bight. The estuary is both an
173 important spawning area for many diadromous and estuarine species, seasonally occupied by a
174 host of species, serving as a nursery ground that contributes habitat services to commercial and
175 recreational fisheries along the eastern United States. Common migratory species inhabiting the
176 Delaware River Estuary include Striped Bass, American Shad, Hickory Shad (*Alosa mediocris*),
177 Alewife (*Alosa pseudoharengus*), and Blueback Herring (*Alosa aestivalis*), and several species
178 of sciaenid including Atlantic Croaker, Black Drum (*Pogonias cromis*), Weakfish, and Spot
179 (*Leiostomus xanthurus*).

180 Identifying and monitoring environmental influences on successful reproduction, growth,
181 distribution, and recruitment is essential to understand how variable environmental processes
182 may affect populations in the future, especially within dynamic environments, including coastal
183 estuaries. To examine the effect the AMO, the NAO, salinity and water temperature have on
184 managed fisheries in the Mid-Atlantic region, we have compared trends in these variables to
185 multiple indices of abundance for Atlantic Croaker, Black Drum, Spot and Weakfish within the
186 Delaware River Estuary. Annual recreational and commercial landings of Atlantic Croaker,
187 Black Drum, Weakfish and Spot have all generally decreased from 1991-2016 despite significant
188 management measures to restrict or reduce harvest. Management plans have suggested
189 monitoring environmental preferences of Atlantic Croaker, Black Drum, Spot and Weakfish to
190 understand effects on stock size, life histories, and spatial distribution (ASMFC 2017). In order
191 to better understand fluctuations in recruitment we need to better account for the effects of large-
192 scale climatic trends, such as the AMO and NAO, as well as more direct environmental factors
193 including salinity and temperature. Our first objective was to standardize indices of catch among
194 commonly shared variables to generate models of relative abundance, while incorporating
195 significant abiotic parameters associated with the AMO and NAO. Next, we sought to combine
196 multiple indices of abundance to generate Bay-wide composite indices of relative abundance.
197 Finally, we examined significant trends in abundance to determine if state specific, and Bay-
198 wide indices of species, age and area specific abundance significantly varied through time from
199 1991-2016.

200

201 <A>Methods:

202 *Study Area:*

203 The Delaware River Estuary is the second largest estuary on the Atlantic Coast of the
204 USA, and it stretches 213 km from Trenton, NJ to Cape May, NJ on the north and Cape
205 Henlopen, DE on the south (Schieler et al. 2014). The widest part of the estuary extends 45 km
206 just inside the estuary entrance, which is 18 km wide (Janzen and Wong, 2002). The Delaware
207 Bay has a mean depth of 8 m (Aristizábal and Chant 2015) with a shipping channel dredged to
208 maintain 14 m MLLW (Mean Lower Low Water) extending up the estuary to the Delaware
209 River. The estuary is weakly stratified with estuarine circulation variably dominated by wind and

210 buoyancy driven flows impacting species and stage/size specific patterns of early juvenile fish
211 ingress (Epifanio and Garvine 2001; Hale and Targett 2018).

212 *Surveys:*

213 Two data sets of fishery independent samples from the Delaware River Estuary were
214 used in this analysis; one conducted by Delaware's Division of Fish and Wildlife (DFW; Figure
215 1) and one conducted by New Jersey's Bureau of Marine Fisheries (NJMF). Since 1980, DFW
216 has used a 5.2-m semi-balloon style trawl, equipped with a 1.3-cm knotless stretch-mesh liner to
217 retain juvenile fishes to estimate relative abundance (Greco 2019). Sampling is conducted
218 monthly in Delaware Bay at 33 fixed sites and in Delaware River at 6 fixed sites, from April
219 through October. Tow durations for both surveys are standardized at 10 minutes for both
220 surveys. Catches that yield large numbers are randomly subsampled in DFW's trawl survey for
221 length and age data. In the DFW 5.2-m trawl a randomly selected 30 individual subsample for
222 each species is measured to the nearest half centimeter for fork length (FL), and the remaining
223 individuals are enumerated. Similarly, NJMF has conducted a trawl survey within New Jersey
224 state waters of Delaware Bay since 1991, using a 4.9-m trawl net equipped with a 1.3-cm
225 knotless stretch-mesh liner (Hassall 2019) contributing to indices of relative fish abundance
226 within the estuary. NJMF collects data monthly at 11 fixed stations within the bay, from April
227 through October. NJMF tow durations are standardized at 10 minutes against tide, using optimal
228 weather windows to conduct sampling similar to the DFW survey. NJMF randomly subsamples
229 50 individual fish for length, which was reported as both FL and total length (TL) for Atlantic
230 Croaker, Black Drum, Spot, and Weakfish. All data were temporally subset to provide
231 overlapping information from 1991-2016 for the purposes of this project. The total number of
232 each species caught per tow was multiplied by the proportion of catch by age based on the
233 observed size distribution for that tow to provide an estimate of the number of YOY and Age-1
234 individuals captured per haul. Fish less than 13 mm were removed from the models to account
235 for gear saturation. Age at length for YOY and Age-1 Black Drum, Atlantic Croaker, Spot and
236 Weakfish were derived from juvenile recruitment indices. Year classes tend to have lengths
237 centered around frequently occurring size classes after hatching (Michels and Greco 1995;
238 Bonzek et al. 1995). Further the YOY recruits are available to the gear seasonally and spatially in
239 the estuary and only certain months are considered in the length ranges and indices per species.
240 YOY length cutoffs for Atlantic Croaker were designated at 100 mm and under in September

241 and October; Black Drum YOY cutoffs were 300 mm and under in August, September, and
242 October; Spot YOY cutoffs were 170 mm and under in July and 200 mm and under in August,
243 September, and October; Weakfish YOY cutoffs were 110 mm and under in June, 150 mm and
244 under in July, 200 mm and under in August, 235 mm and under in September and 250 mm and
245 under in October. Age-1 length frequencies were developed using literature reviews of YOY to
246 Age-1 cutoffs and Age-1 to Age-2 cutoffs (Atlantic Croaker; Liao et al. 2019, Michels and Greco
247 1995, Bonzek et al. 1995, ASMFC 2017b); Black Drum; Liao et al. 2019, Michels and Greco
248 1995, Bonzek et al. 1995); Spot; Liao et al. 2019, Michels and Greco 1995, Bonzek et al. 1995);
249 Weakfish; Liao et al. 2019, Michels and Greco 1995, Bonzek et al. 1995, ASMFC 2006,
250 Lowerre-Barbieri 1995). Age-1 length cutoffs for Atlantic Croaker were determined to be 201 –
251 260 mm; Black Drum Age-1 fish were considered at 330 – 455 mm; Spot Age-1 fish were
252 considered at 201 – 224 mm; Weakfish Age-1 fish were considered at 254 – 275 mm.

253 *Statistical Analyses:*

254 Environmental data including surface water temperature (°C) and salinity (ppt) collected
255 at each net set/haul, as well as the monthly values of two climatic indices including the AMO
256 (<https://psl.noaa.gov/data/timeseries/AMO/>), and NAO
257 (www.cpc.ncep.noaa.gov/products/precip/CWlink/pna/nao_index.html) were used to explore
258 how these parameters influenced the species, age and area specific indices of catch. Both the
259 DFW 5.2 m and the NJMF 4.9 m surveys were standardized using a generalized additive model
260 framework where tow specific catch was modelled as a function of

$$261 \text{ Catch} \sim \text{factor(Year)} + \text{factor(Month)} + \text{s(Station)} + \text{s(Temp.)} + \text{s(Salinity)} + \text{s(Depth)} + \\ 262 \text{s(AMO)} + \text{s(NAO)}$$

263 using a negative binomial error distribution, Poisson error distribution and zero-inflated Poisson
264 error distribution for each index through time where Year and Month are fitted as fixed effect
265 factors, while Station was treated as a random effect, Temperature, Salinity, Depth, monthly
266 values for the AMO and NAO are fitted as smooth terms with thin plate regression splines as the
267 basis and k , is the number of basis functions to use for each smooth term before any
268 identifiability constraints are applied to estimate an index using the “gam” function in the
269 “mgcv” package in R 3.5.2 (R 2008) for each survey, species and age combination (Drexler and
270 Ainsworth 2013). Indices of abundance were each fit using this model formulation to standardize
271 according to year, and fixed station, as well as to account for variation in depth and month

272 between surveys while measuring if environmental and climatic variables affected catch. The
273 estimation method applied was restricted maximum likelihood (REML). Similarly, Bay-wide,
274 composite indices of abundance were modelled using the same approach, with the incorporation
275 of an additional factor to account for variability associated with the two state surveys with tow
276 specific catch was modelled as a function of

$$\text{Catch} \sim \text{factor}(\text{Year}) + \text{factor}(\text{State}) + \text{factor}(\text{Month}) + \text{s}(\text{Station}) + \text{s}(\text{Temp.}) + \\ \text{s}(\text{Salinity}) + \text{s}(\text{Depth}) + \text{s}(\text{AMO}) + \text{s}(\text{NAO}) + \text{s}(\text{Year} * \text{State})$$

279 using a negative binomial error distribution (NBGM), Poisson error distribution (PGAM) and
280 zero-inflated Poisson error distribution (ZIPGAM) for each index. For all of the candidate
281 GAMs, the best model (initial error distribution (NBGM, PGAM, ZIPGAM) and final model
282 selection) was determined with an information-theoretic approach using the Akaike information
283 criterion or AIC score (Burnham and Anderson 2002; Bucheister et al. 2016). After determining
284 the optimal error distribution among three similar, global models, the “dredge” function in the
285 “MuMIn” package of R 3.5.2 (R 2008) was used to automatically test and find the most
286 parsimonious model formulation of potential covariates for each species, area (DE, NJ), and age
287 (YOY, Age-1) specific model, as well as for Bay-wide composite indices (CI) which included an
288 interaction between Year and State. The “predict” function in the “car” package of R 3.5.2 (R
289 2008) was used to generate estimates of annual mean catch using the tow specific standardized
290 models of catch. A likelihood ratio test was used to compare models with a null model for each
291 index to determine goodness of fit. Standardized indices of abundance for each species (Atlantic
292 Croaker, Black Drum, Spot and Weakfish), age (YOY, Age-1) and survey (Composite Indices,
293 DFW 5.2 m, NJMF 4.9 m) were then fit with Autoregressive Integrated Moving Average
294 (ARIMA) models using an iterative approach of 1,000 bootstrapped runs to estimate a median
295 ARIMA fit for each index using the “surveyfit” and “surveyref” functions in the “fishmethods”
296 package of R 3.5.2 (R 2008). ARIMA models are commonly used to examine trends in a
297 population relative to a given reference point, such as survey quartiles or terminal years in
298 managed species (e.g. Atlantic Sturgeon, ASMFC 2017a) providing a mechanism to assess the
299 relative abundance of each species-area-age combination through time. The terminal year value
300 generated from the median ARIMA was then compared to both the first quartile and the survey
301 start year to estimate a probability of the terminal year being greater than 25% of the time series
302 and the survey start year using a statistical level of confidence of $\beta = 0.80$ with a Holm-adjusted

303 probability of rejecting the null hypothesis regarding normality of model residuals (Box and
304 Jenkins 1976; Helser and Hayes 1995; ASMFC 2017a). Then, we used a Mann-Kendall trend
305 test to analyze monotonic trends in the median ARIMA indices using the “mk.test” function in
306 the “trend” package of R 3.5.2 (R 2008), after correcting for family-wise error rates in the
307 statistical *p*-values using the Holm method. Finally, length frequencies of each species were
308 developed to compare the relative size distributions of Atlantic Croaker, Black Drum, Spot, and
309 Weakfish caught between the two trawl surveys from 1991-2016 (Ogle 2014; Emmanuel 2017).

310

311 <A>Results

312 Trends in the AMO index are evident between positive and negative phases over a time
313 series of reconstructed and observed sea surface temperature (SST) data at 65-70-year intervals
314 (Knudsen et al. 2011). Within our available 26-year time series, shifts from cool to warm years
315 are evident in both the NAO and the AMO. From 1991-1994, the AMO index remained in a cool
316 phase and has remained in a warm phase from 1997 through 2016 (Figure 2). Conversely, the
317 NAO had demonstrated greater variability between years than the AMO. Slightly more than half
318 (53.8 %) of the years from 1991-2016 were represented by a warm year for the NAO index with
319 reversals experienced at a frequency of one to four year intervals (Figure 2).

320 All species and age combinations were not equally represented between the two locations
321 when compared to the total number of individual fish collected for each species, age and area.
322 Higher numbers of all YOY were observed in Delaware. Between 79 – 92 % of all the YOY
323 observed for the four species were collected in Delaware compared to New Jersey. However, a
324 higher number of Age-1 Weakfish was observed in New Jersey. When examining the total
325 number of fishes caught with all covariate data present, 122,077 individuals were collected in
326 Delaware, whereas 25,202 individuals were collected in New Jersey across all species and ages
327 combined (Table 1). More than 61 % of the total number of individuals collected in Delaware
328 were comprised of YOY Weakfish. Similarly, more than 58 % of the total number of individuals
329 collected in New Jersey were attributed to YOY Weakfish. Age-1 Atlantic Croaker and Spot
330 were both collected in Delaware, at two orders of magnitude less when compared to YOY for
331 both of those species. Conversely, no Age-1 Atlantic Croaker or Spot were documented in New
332 Jersey from 1991-2016 (Table 1). However, Age-1 Weakfish were present at both locations.

333 Of the seventeen possible combinations where we had enough data to generate base
334 models of catch that included all model covariates, the negative binomial error distribution was
335 the lowest scoring AIC value of the global models in thirteen of the indices (Table 1). ZIPGAMs
336 were chosen as the final model in all three indices of YOY Black Drum indices. And a PGAM
337 was selected for in the Age-1 Atlantic Croaker index in Delaware. All final model configurations
338 were highly significant when compared to null models without covariates (p -value < 0.001). The
339 deviance explained by the final models for all species, area, and age combinations ranged from
340 23.1 – 89.2 % depending upon the model, with all dispersion values less than 2.4 for all indices,
341 suggesting reasonable levels of variance were present for all final models (Table 1).

342 Many of the environmental and climatic variable smoothing parameters were found to be
343 significant in the final models. However, the year and state interaction term included in three of
344 the five composite indices which combined data from Delaware and New Jersey was not
345 significant in any model (Table 2). Station was found to act as a significant smoothing parameter
346 in all the fifteen models in which that parameter was selected for inclusion in the final model.
347 The AMO, NAO, salinity and surface temperature collected at each tow were significant
348 smoothing parameters in twelve of the seventeen final models. While depth was found to be
349 significant in ten of the seventeen final models. Residuals were not normally distributed for six
350 of the seventeen ARIMA model fits. However, for the remaining eleven ARIMA models,
351 residuals were normally distributed (Table 3). All the normally distributed median ARIMA
352 model fits, but the Bay-wide, composite index of Age-1 Weakfish had a higher probability ($P >$
353 51 %) of being greater than the first quartile for each time series, while four of the indices had a
354 higher probability ($P > 51$ %) of being less in 2016 when compared to the survey start year of
355 1991 (Table 3).

356 Peaks in YOY Atlantic Croaker abundance were observed in New Jersey from 1996-2011
357 over the course of the time series with peaks in YOY Atlantic Croaker observed in Delaware
358 from 2001-2005 and again in 2010 and 2014 (Figure 3). The Bay-wide composite index of YOY
359 Atlantic Croaker had seven peaks in abundance throughout the course of the time series.
360 However, peaks in abundance of Age-1 Atlantic Croaker are present only at the beginning of the
361 time series in Delaware from 1991-1994 (Figure 3). GAM indices of Atlantic Croaker abundance
362 explained between 38.5 – 60.1% of the deviance for the two surveys (Table 1). The smooth term
363 for salinity was included in all three final YOY models, and significant in both the Delaware and

364 composite indices. Similarly, the AMO and the NAO were significant smoothing parameters
365 included in all three final YOY models. While a significant smooth term for depth was included
366 in the final YOY NJMF 4.9 m survey only. Station was also identified as significant smoothing
367 parameters in the YOY DFW 5.2 m and composite indices. All the smoothing parameters
368 included in the YOY DFW 5.2 m index were included in the Age-1 DFW 5.2 m index, in
369 addition to depth (Table 2). Median fit ARIMAs for the YOY DFW 5.2 m and composite index
370 for Atlantic Croaker (Figure 3) demonstrated an increasing trend from 1991-2016 with normally
371 distributed error terms and a high probability of an increase in abundance relative to the first
372 quartile for each survey ($P \geq 99\%$; Table 3). Median fit ARIMA models of both the YOY NJMF
373 4.9 m and Age-1 DFW 5.2 m indices were not normally distributed. Atlantic Croaker ranged in
374 size from 15–175 mm FL in the DFW-5.2 m survey and 13–173 mm FL in the NJMF 4.9 m
375 survey (Figure 7) throughout the time series.

376 For Black Drum, YOY were the only encountered age group in the two surveys with
377 matching covariate data. Peaks in the DFW 5.2 m survey were observed from 1993 to 2013.
378 Whereas a single large peak in abundance was observed in the NJMF 4.9 m survey in 2007
379 (Figure 4). GAMs of Black Drum abundance explained between 42.1–89.2% of the deviance
380 for each of the three surveys (Table 1). Depth was the only significant smoothing variable
381 included in all three surveys. Depth and the NAO were the only smoothing parameters included
382 in the YOY, NJMF 4.9 m GAM. The AMO, depth, NAO, salinity, station, temperature were
383 significant smoothing parameters in the YOY models for the DFW 5.2 m survey, and the Bay-
384 wide composite index (Table 2). Additionally, the interaction term between Year and State was
385 also included in the final model of the YOY composite index. The ARIMA of the YOY DFW
386 5.2 m Black Drum index had a high probability ($P \geq 76\%$, Table 3; Figures 4), that the index
387 value in 2016 was both greater than the first quartile, as well as the survey start year. However,
388 the residuals from the YOY NJMF 4.9 m survey and the composite index were not normally
389 distributed (Table 3) suggesting that the data representing the population were not normally
390 distributed and results inferred from those analyses should be ignored or used with caution.
391 Further, no significant trend in YOY Black Drum abundance was identified in the DFW 5.2 m
392 index (Table 3). Black Drum ranged in size from 75–435 mm FL in the DFW-5.2 m survey and
393 27–320 mm FL in the NJMF 4.9 m survey (Figure 7).

394 Spot like Atlantic Croaker, and Black Drum had a much higher likelihood of being
395 observed as YOY, when compared to the total number of Age-1 individuals (Table 1). Peaks in
396 YOY Spot abundance were asynchronous throughout time between the two states (Figure 5).
397 GAMs of abundance explained between 38.7 – 63.2 % of the deviance for each of the four
398 models (Table 1). Station as a random effect was the only significant smoothing parameter
399 included in the YOY NJMF 4.9 m survey. While all the smoothing parameters were both
400 significant and included in the YOY DFW-5.2 m survey, except for salinity which was dropped
401 from the final model. Additionally, all the smoothing variables of the final YOY composite index
402 were included in the final model formulation. However, the AMO and the interaction between
403 year and state, while included in the final model were not statistically significant smoothing
404 parameters in the YOY composite index. Depth, the NAO, salinity, and station were all
405 significant smoothing parameters in the Age-1 DFW-5.2 m survey index (Table 2). In 2016, the
406 median fit ARIMA of YOY Spot from the DFW 5.2 m and composite index had a 91 %
407 probability of being higher than the first quartile for the time series and lower than the value
408 observed in 1991. However, Age-1 Spot from the DFW 5.2 m index had a 77 % probability of
409 being greater than the first quartile (Table 3). No significant trends in abundance were detected
410 for any median ARIMA fit YOY Spot index. However, Age-1 Spot from the DFW-5.2 m survey
411 were found to have declined since 1991. Spot ranged in size from 15-250 mm FL in the DFW-
412 5.2 m survey and 14-224 mm FL in the NJMF 4.9 m survey (Figure 7).

413 Similar to all three of the other species explored, YOY weakfish were more frequently
414 encountered by both surveys when compared to the proportion of the total number observed of
415 Age-1 Weakfish within each survey (Table 1). YOY Weakfish had a period of relatively higher
416 abundance from 1999-2012 in the NJMF 4.9 m index; whereas YOY Weakfish were generally
417 higher in relative abundance in the DFW 5.2 m survey from 1991-2000 with peaks after that
418 period in 2007, 2011 and 2015 (Figure 6). Relative abundance was higher for the Age-1 DFW
419 5.2 m survey from 1992-2000, when compared to the rest of time series. Conversely, the Age-1
420 Weakfish index from the NJMF 4.9 m had more peaks toward the end of the time series from
421 2000-2015 (Figure 6). The Bay-wide, composite index of Weakfish appeared to be more like the
422 DFW 5.2 m index with a cluster of higher values observed from 1991-1997, however all of the
423 Age-1 annual means were less than 1. GAMs of Weakfish abundance explained between 23.1 –
424 60.5 % of the deviance in the data for the six indices (Table 1). The AMO, station as a random

425 effect and temperature, were all significant smoothing parameters included in the final models
426 for all YOY Weakfish indices. Depth was included in all three final models, as well, but not
427 significant in the YOY DFW 5.2 m Weakfish index. While the NAO and salinity were
428 significant smoothing variables for the DFW 5.2 m and the composite indices, but not included
429 in the final YOY NJMF 4.9 m index. All smoothing variables present in the final model
430 formulation were significant in the Bay-wide GAM of Age-1 Weakfish. Additionally, the
431 interaction between year and state was included in the final YOY composite index (Table 2).
432 Salinity and station as a random effect were all significant smoothing parameters in all three of
433 the Age-1 Weakfish indices. The AMO was a significant smoothing parameter in both the DFW
434 5.2 m and composite index, but not included in the final Age-1 NJMF 4.9 m index. Temperature
435 was a significant smoothing parameter in the Age-1 NJMF 4.9 m and composite index, but not
436 included in the final model of the Age-1 DFW 5.2 m index. Finally, the NAO was a significant
437 smoothing parameter in the Age-1 NJMF 4.9 m index, but not present in either of the final
438 models for the other two Age-1 indices (Table 2). The 2016 terminal time series year estimate
439 from the median fit ARIMA had a high likelihood ($P \geq 75\%$) of being greater than the first
440 quartile for each YOY Weakfish time series. However, the likelihood of the terminal year being
441 greater than the survey start year in 1991 was much lower in most of the YOY and Age-1
442 Weakfish indices. Also, the Bay-wide Age-1 Weakfish composite index had a higher likelihood
443 of being less than the first quartile ($P = 57\%$; Table 3). No significant trends were detected in the
444 YOY Weakfish for either the Bay-wide composite index or the NJMF 4.9 m index. However, the
445 YOY Weakfish DFW 5.2 m GAM was found to have declined through time. Residuals from the
446 Age-1, NJMF 4.9 m survey were not normally distributed suggesting that the data representing
447 the population are not normally distributed and results inferred from those analyses should be
448 ignored or used with caution for that index. However, the median fit ARIMA Age-1 Weakfish
449 index for the DFW 5.2 m survey was increasing, while the composite index was found to be
450 declining through time (Table 3). Weakfish showed a range in lengths between 13-250 mm FL in
451 the DFW 5.2 m survey for YOY and Age-1, and 13-250 mm FL in the NJMF 4.9 survey for
452 YOY and Age-1 (Figure 7).

453

454 <A>Discussion

455 Sciaenid species showed varying patterns in abundance through time within the Delaware
456 River Estuary from 1991-2016 demonstrating significant trends and associations with both
457 environmental covariates and larger, regional climatic indices. Each of the four species
458 experienced shifts in abundance at different time intervals within the modelled time series,
459 suggesting that underlying factors are likely affecting species-age-area specific trends in
460 abundance within the estuary. Of the environmental processes affecting abundance, temperature
461 and salinity were both consistently found to smooth GAM fits for multiple species-age-area
462 specific models of abundance for these species. Similarly, we consistently observed relationships
463 between climatic indices including the AMO and the NAO, and modelled abundance,
464 demonstrating that long term climate has a correlation between smaller scale processes like water
465 temperature, that affect the species examined in this study at multiple ages. Of the eight potential
466 composite indices (YOY & Age 1; Atlantic Croaker, Black Drum, Spot, Weakfish) that might
467 allow us to examine how environmental drivers may affect relative abundance for Sciaenids on
468 an estuary wide scale, only five of the indices had enough data to generate GAMs of abundance,
469 and only two of the indices had normally distributed, measurable trends in abundance including
470 the YOY Atlantic Croaker and Age-1 Weakfish composite indices. Of those two, YOY Atlantic
471 Croaker have increased, while Age-1 Weakfish have decreased from 1991-2016. Therefore, we
472 have demonstrated a method to aggregate data across spatial areas, standardize catch, evaluate
473 trends within an area through time and make comparisons relative to time series reference points
474 that may serve as a management guideline, and make broader assessments across a larger spatial
475 region to examine how abundance varies through time.

476 We have considered the limitations of our data and analyses in the interpretation of these
477 results. Our study used catch data across two historical fixed-station time series of unequal
478 survey sites, limiting the catch variability and inducing potential spatial bias within the sampling
479 routines. However, due to heavy maritime shipping traffic in the estuary and spatial designations
480 by each state limiting the bounds at which each survey can operate, combining fixed station data
481 became a viable option for analyzing a time series of relative abundance in the Delaware River
482 Estuary. Further, daily weather patterns, gear type, trawl speed and boat captain may have all
483 affected the concentration of individuals within a given species for each survey (Misund et al.
484 1999). Additionally, there was a higher number of stations sampled within the Delaware survey
485 when compared to the New Jersey survey, which likely skewed our results, particularly in the

486 construction of Bay-wide composite indices. Our attempt to build a composite index may reflect
487 this disparity between the number of sampling stations, as the composite models often behaved
488 similar to those from the DFW surveys. We used the most optimal GAM we could identify for
489 each individual survey and species-age-area-specific index of relative abundance so that we
490 could standardize annual catches of each species without assuming linear relationships among
491 covariates and build individual models that accounted for covariates including station and
492 interactions between year and state for composite indices. GAMs apply a series of smoothing
493 functions in an iterative approach to fit linear or nonlinear relationships between individual
494 predictors and dependent variables simultaneously (Hastie and Tibshirani 1986). As an example,
495 GAMs were found to better estimate catch when compared to GLMs because of the nonlinearity
496 associated with covariates and catch in the Alaskan Sablefish (*Anoplopoma fimbria*) longline
497 fishery (Mateo and Hanselman 2014). Overlap in ages at length was also considered within our
498 analysis, especially within the Age-1 to Age-2 cutoffs for Weakfish where large variations of
499 length at age have been observed within these age groups (Lowerre-Barbieri et al. 1995).
500 However, we believe that the estimates of catch we derived are likely a conservative
501 approximation for the true proportion at age being captured.

502 Combining each of the two surveys into a single composite index through time allowed
503 for a Bay-wide model of species-specific relative abundance by combining multiple surveys
504 throughout the estuary standardized by a similar suite of potential covariates. Abundance data
505 vary annually, reflecting population level fluctuations, survey sampling variability, and variable
506 catchability (Pennington 1986; ASMFC 2017a). We utilized ARIMAs to filter measurement
507 error from process variability to identify trends in relative abundance (Box and Jenkins 1976;
508 Helser and Hayes 1995; ASMFC 2017a) for individual surveys, and Bay-wide composite
509 indices, and generate comparisons with those filtered data with environmental variables. Finally,
510 the AMO and NAO vary at multi-decadal frequencies, so attempting to compare trends in
511 relative abundance with 26 years of data may fail to adequately capture the effect these climatic
512 processes have on Sciaenid abundance through time. However, we were fortunate enough to
513 have data that spanned positive and negative phases of both the AMO and NAO, allowing for an
514 exploration of how the AMO and NAO affect relative abundance. Despite the limitations of our
515 data and the assumptions associated with our methods, we have managed to estimate relative
516 abundance for four species at two ages, using standardized indices of abundance, generate

517 statistically significant Bay-wide trends in relative abundance for four species, and make
518 comparisons of a subset of those trends to large scale, climatic drivers and environmental
519 variables including temperature and salinity.

520 The abundance of Atlantic Croaker was related to the AMO, depth, the NAO, salinity,
521 station as a random effect, temperature and an interaction between year and state within the
522 Delaware River Estuary. These results suggest climatic, and environmental processes are
523 correlated to fluctuations in abundance at both ages. Further, our results demonstrate that the two
524 fixed station surveys asynchronously capture peaks in abundance, suggesting that Atlantic
525 Croaker may be utilizing different areas within the bay through time based on active habitat
526 selection, physical processes affecting their ingress into the estuary or a combination of
527 biophysical transport processes (Hale & Targett 2018). Trends in species abundance based on
528 statistics for the Mann-Kendall trend tests of median fitted ARIMAs for each index demonstrate
529 an increasing trend in abundance of YOY Atlantic Croaker in Delaware Bay. Additionally, we
530 found that multiple environmental variables and climatic phenomena were correlated to
531 modelled abundance through time including temperature, salinity, depth, station, the AMO and
532 the NAO depending upon the survey.

533 The connection between environmental variables that are controlled by large scale
534 climatic signals and abundance in Atlantic Croaker has been previously observed. Data from
535 Lankford and Targett (2001) suggested that Atlantic Croaker have an optimal temperature
536 window at age-0 in Delaware Bay, with substantial mortality when exposed to prolonged water
537 temperatures below 5°C. Similarly, work conducted in the Gulf of Mexico suggested that
538 Atlantic Croaker distribution can be influenced by abiotic factors, such as hypoxia and water
539 temperature, causing spatial disturbances within the species' preferred niche and dispersing
540 Atlantic Croaker to cooler offshore waters where growth was limited due to reduced growth
541 energy (Craig and Crowder 2005). Witting et al. (1999) found that in the Little Egg Harbor, NJ,
542 temperature affected the annual phenology and duration of seasonal larval fish assemblages, with
543 Atlantic Croaker being most abundant as part of the fall assemblage. Similarly, our findings
544 suggest that the AMO, which is directly related to SST (Edwards et al. 2013; Large et al. 2013),
545 affects YOY Atlantic Croaker abundance within the Delaware River Estuary (Lankford and
546 Targett 2001; Miller et al. 2003). Previous work, including a study by Hare and Able (2007)
547 correlated 'outbursts' of recruitment and year class strength of juvenile Atlantic Croaker with

548 above average minimum winter temperatures and the wintertime NAO index across several Mid-
549 Atlantic States, demonstrating that warm winters provide an increase in optimal Atlantic Croaker
550 habitat in Mid-Atlantic estuaries. Our findings, much like previous studies, show an increase in
551 Atlantic Croaker abundance during warm temperatures within Delaware Bay. Our findings with
552 the AMO and NAO index suggest that physical mechanisms controlling SST, wind driven flows
553 and precipitation (Enfield et al. 2001) are correlated to juvenile abundance, based on reported
554 size at maturity and landings (ASMFC 2010; ASMFC 2017b) in the estuary. Similarly, larval
555 Atlantic Croaker were previously suggested to be reliant upon physical forcing mechanisms at
556 early life history stages to promote ingress into Delaware Bay (Hale and Targett 2018)
557 suggesting that environmental processes are acting throughout life history on this species in a
558 compounding manner with behavioral modification to enhance or depress survival.

559 Comparable to Atlantic Croaker, we found that multiple climatic drivers and
560 environmental variables significantly affected models of abundance of YOY Black Drum in
561 Delaware, and at an estuarine scale. Other researchers have similarly identified links between
562 climatic signals and Black Drum. Zimmerman (2016) found a strong positive correlation with the
563 AMO and Black Drum landings, with significant lags occurring at 9 years prior to a given year,
564 suggesting that increased catches were linked with positive AMO phases nearly a decade prior.
565 Although the study by Zimmerman (2016) dealt with adult Black Drum primarily, the findings
566 do show that the species has some likely relationship with the AMO. Despite the relatively brief
567 time series we have available (26 years) compared to the 65-year time series examined by
568 Zimmerman (2016) coupled with the longevity of the species (maximum age = 67 years,
569 ASMFC 2015) we still identified the AMO as a significant smoothing parameter of YOY Black
570 Drum in the composite index and Delaware. However, the AMO was not correlated to the YOY
571 Black Drum index in New Jersey suggesting that some difference exists between the two data
572 series. Further, Black Drum were the lowest encountered of the four species explored. Juvenile
573 Black Drum use of salt marshes and tidal creeks as nurse habitat (Odell et al. 2017), as
574 compared to more open habitat accessible to the trawl surveys may be a cause of the low
575 interaction rates observed in our study, due to the lack of sampling in these areas by the two
576 surveys. Further, the two surveys may have differences in how they trawl, that affect catch rates,
577 which we failed to account for in our models of abundance. Differences such as area sampled,
578 depth, number of stations, equipment, and operational variations between the two surveys likely

579 impact catch rates. However, we attempted to control for these parameters by including station
580 and an interaction between year and state where applicable. Similar to the results of our study,
581 Thomas and Smith (1973) examined Black Drum YOY and suggested that factors such as
582 bottom type, current, and temperature are more important for suitable nursery habitat than
583 salinity alone in tidal creeks in the upper Delaware Bay. Further, Black Drum chorusing activity
584 associated with spawning has been correlated with water temperatures offshore of North
585 Carolina and Georgia suggesting that water temperature may affect spawning behavior (Rice et
586 al. 2016) and the subsequent presence of YOY. Black Drum were significantly correlated to
587 dissolved oxygen levels during associated spawning runs in Louisiana, where supersaturated
588 environments were found to have large aggregations of spawning adults (Saucier & Baltz 1993).
589 Additionally, Saucier and Baltz (1993) suggested that the highly oxygenated environments may
590 be sought after because of the size of the Black Drum eggs and the need for higher
591 concentrations of dissolved oxygen to assist in diffusion through the egg envelope. During
592 different phases of the AMO there may be an influx or decrease of freshwater (Enfield et al.
593 2001; Nye et al. 2014), and variability within SST, which can influence the amount of dissolved
594 oxygen within a given area, changing the physiochemical properties of the water (Irby et al.
595 2018), and potentially causing Black Drum to search for suitable habitat for spawning or juvenile
596 periods of residency.

597 A Bay-wide, composite index of YOY Spot included smoothing parameters for all of the
598 environmental variables and climatic phenomena tested, similar to the Bay-wide composite index
599 of Black Drum. In fact, all smoothing variables, except for the AMO and the interaction between
600 year and state were significantly related to the modelled abundance of Spot in Delaware Bay and
601 all but, salinity were correlated between YOY Spot collected in Delaware. Conversely, YOY
602 Spot collected in the New Jersey survey were best explained by salinity and station as a random
603 effect without the inclusion of other environmental or climatic variables. However, the total
604 number of YOY Spot was an order of magnitude lower between Delaware and New Jersey, again
605 suggesting that there is variability in the collection of a species by area, like what was found with
606 Black Drum. However, it is again worth nothing, that a greater number of stations were sampled
607 in Delaware when compared to the New Jersey survey. Our findings demonstrate that YOY Spot
608 abundance in Delaware Bay is significantly correlated to temperature, salinity, depth, and the

609 NAO. Like the results of our study, Spot in Chesapeake Bay were positively associated with
610 increased temperature and salinity (Schaffler et al. 2013; Love and May 2007).

611 Models of Weakfish abundance were significantly related to environmental and climatic
612 processes depending upon the age of the fish and the survey area. Bay-wide indices of abundance
613 demonstrate associations among temperature, salinity, station, an interaction between year and
614 state, and the AMO for YOY and Age-1 ages and the NAO for YOY. The AMO was a
615 significant smoothing variable in five of the six final models of Weakfish abundance, suggesting
616 that this climatic phenomenon is correlated to Weakfish abundance. Previous studies have found
617 similar associations between climatic indices and Weakfish abundance, as well as Weakfish
618 mortality. Both the AMO and the NAO have been previously correlated to Weakfish natural
619 mortality for ages 1-6 using data from Connecticut to North Carolina (Jiao et al. 2011).
620 However, the AMO (annual average) was linearly related to natural mortality (Jiao et al. 2011).
621 The last negative phase of the AMO, which occurred from the late 1950s into the 1980s, was
622 linked with higher total catches of Weakfish, most notably in the 1980s. However, the AMO
623 moved into a warm, positive phase in the early 1990s, remaining positive through recent time
624 with associated record low total catch rates (ASMFC 2009). A similar trend was observed
625 through historical catch records in the Delaware Bay from 1880-1933, where catches of
626 Weakfish were high into 1929 followed by decline through 1933 (Nesbit 1954). Higher catches
627 of Weakfish coincide with the negative phase of the AMO during this time period within
628 Delaware Bay, with lower catches correlated with the warm AMO phase starting in the early
629 1930s (Nesbit 1954).

630 In addition to large scale climatic drivers, smaller scale, environmental variables have
631 been found to affect Weakfish concentration as well. Weakfish studied in Great Bay, NJ, utilized
632 areas of higher salinities and water temperatures during summer suggesting those environmental
633 factors may influence their habitat selection (Turnure et al. 2015). Temperature may have an
634 influence on male Weakfish disturbance call duration during spawning which may help female
635 Weakfish find a suitable mate and discriminate males based on size (Connaughton et al. 2000).
636 Based on our results, we believe that juvenile Weakfish abundance may be significantly
637 influenced by the AMO through the effect of the AMO on water temperature. Feeding behavior
638 of juvenile Weakfish has been correlated to temperature, salinity and prey interaction rates
639 within Delaware Bay (Greca and Targett 1996), with results demonstrating these

640 physiochemical attributes can significantly affect growth rates and induce a stress response if
641 high temperatures and lower salinities are encountered (Lankford and Targett 1994). Hare et al.
642 (2016) suggested that Atlantic Croaker, Spot and Weakfish all had the same moderate levels of
643 climatic vulnerability, but the relative degree of certainty according to the standard error of the
644 index was much less for Weakfish compared to the other two species. The relative significance
645 of the environment could be tied to both natural factors, including trends in early life history and
646 natural mortality, as well as confounding anthropogenic factors, including a depleted population
647 status and subsequent interactions between those natural (e.g. mortality, competition) and
648 anthropogenic factors (destruction of environment and environmental conditions).

649 Beyond the findings for these species in Delaware Bay, the results of previous analyses
650 suggest that fish species are affected by large-scale environmental processes across broad
651 geographic areas, with species being affected at different life stages, and altering attributes
652 associated with trophic ecology. Environmental processes have been correlated with changes in
653 predation patterns (Yasumiishi et al. 2016), growth of marine nekton (Charnov and Gillooly
654 2004; Yasumiishi et al. 2016; Izzo and Zydlewski 2017) and success of year classes among
655 nekton in freshwater (Reist et al. 2006) and marine ecosystems (Lankford and Targett 2001;
656 Brander and Mohn 2004; Hare and Able 2007). In the Pacific, greater oceanic recruitment and
657 growth rates of juvenile Sockeye Salmon (*Oncorhynchus nerka*) have been positively correlated
658 with SST before entering oceanic waters along the eastern Bearing Sea during warming events,
659 allowing Sockeye Salmon to reach adult maturity more rapidly, aiding in survival (Yasumiishi et
660 al. 2016). Further, survival of nekton is greatly affected by feeding and feeding encounter rates
661 of different prey species (Chao and Musick 1977) with annual variability generated by physical
662 environmental conditions including water column stability (Carassou et al. 2011). In addition,
663 environmental variables and climatic processes have been found to generate variance in the
664 patterns of spatial distribution among marine species due to habitat preferences and potentially
665 overlapping concentrations of prey (Overholtz et al. 2011; Sagarese et al. 2011).

666 The AMO and NAO have been correlated to recruitment and life stages of many species
667 in the Atlantic Ocean. Specifically, Buchheister et al. (2016) found that an Atlantic Menhaden
668 recruitment index in Delaware showed a significant positive relationship when correlated to a
669 lagged AMO. Our results suggest the AMO's influence on the Delaware River Estuary
670 ecosystem has the potential to alter individual year classes, and subsequently affect population

671 abundance which in turn affects the potential productivity of a fishery of an individual species,
672 generating ecological implications for other species and fisheries. Similarly, the NAO also has
673 been linked to trends in the production, distribution, and abundance of fishes. The NAO has been
674 found to broadly affect anadromous (e.g. Striped Bass, O'Connor et al. 2012) and catadromous
675 (e.g. American Eel (*Anguilla rostrata*), Friedland et al. 2007) species, as well as impact multiple
676 life history stages of other species including the larval stages of Blueback Herring, American
677 Eel, Weakfish, Spot, Atlantic Croaker, and Black Drum (Love et al. 2009) and adult stages of
678 Alewives, Blueback Herring, Atlantic Herring (*Clupea harengus*), and Atlantic Mackerel
679 (Turner et al. 2017). Similarly, Hare and Able (2007) found that increased catches of adult
680 Atlantic Croaker in the 1950s, 1970s, and 1990s were significantly correlated with the warm
681 phase of the NAO index showing the effect of the NAO on Atlantic Croaker recruitment. North
682 Atlantic Cod (*Gadus morhua*) recruitment was positively correlated to the NAO in three cod
683 stocks in the North Sea, Baltic Sea, and Irish Sea, but had a negative correlation in Iceland
684 (Brander and Mohn 2004).

685 Understanding how environmental variables affect the relative abundance of managed
686 fishes is important to improve the characterization of population variability and assess stock
687 status. We successfully documented species-age-area specific relationships that suggest
688 temperature, salinity, and climatic drivers that impact those variables including the NAO and the
689 AMO can have measurable effects on the abundance of four Sciaenid species at multiple points
690 through time at variable spatial scales. Bay-wide composite indices of abundance provide insight
691 into how environmental variables impact abundance in juvenile fishes across a broader area.
692 Nonlinear relationships exist between catch, temperature and salinity for YOY Atlantic Croaker,
693 Black Drum, Spot, Weakfish, and Age-1 Weakfish in Delaware Bay (Supplemental Figures)
694 which generate variable points in time and space that can alter juvenile survival and growth in an
695 estuary. These age and species-specific responses to environmental and climatic variables occur
696 within an optimum band to produce elevated levels of abundance within a range of observed
697 values for a suite of covariates. We suggest that long term climatic conditions are generating
698 smaller scale environmental fluctuations that present unique physiochemical water conditions
699 that vary in spatial and temporal scale, generating species, age and area specific responses in the
700 abundance of marine fishes observed in Mid-Atlantic estuaries. Further, we have demonstrated a
701 need to explore relationships between environmental factors and fishes, so that stock assessments

702 and management action can incorporate or at least consider incorporating climatic variability and
703 respond accordingly to changes in environmental conditions, thus more accurately portraying
704 natural variability of trends in abundance.

705

706 <A>Figures

707 Table 1. Species, area, age, n (total number of fish observed with matching covariate data), delta
708 AIC scores for similar base models to determine optimal error structure, the final model selection
709 with deviance explained and dispersion from 1991-2016.

710 Table 2. Associated p-values for the smoothing parameters of AMO, depth (m), NAO, salinity
711 (ppt), station, temperature (°C), and an interaction between Year * State for each species, area,
712 and age specific final model with significant variables highlighted in yellow, unused covariates
713 from the final model formulation were left blank.

714 Table 3. Summary statistics for median ARIMA results and the Mann-Kendall trend tests of
715 median fitted ARIMAs for each index by species, area and age. W = Shapiro-Wilk statistic for
716 normality, adj. p-value= Holm-adjusted probability of rejecting the null hypothesis regarding
717 normality of model residuals, (θ) = moving average parameter, SE = standard error of theta, σ_c^2 =
718 variance of index, $P(2016 < 25^{\text{th}} \text{ pctl})$ = the probability that the terminal year index value
719 observed in 2016 was less than the first quartile for the time series, $P(2016 < 1991)$ = the
720 probability that the terminal year index value observed in 2016 was less than the survey start
721 year in 1991, S = Kendall Score, σ^2 = variance of Kendall Score, τ = Kendall's tau statistic, adj.
722 p-value = the Holm-adjusted probability of the Mann-Kendall time series trend being significant,
723 Trend= trend result (increasing, decreasing or n.s.= not significant). ARIMA fits with non-
724 normally distributed residuals are highlighted in yellow.

725 Figure 1. A map of Delaware Bay and Delaware River. Trawl station coordinates are marked
726 with symbols and each trawl survey is denoted by color within the estuary.

727 Figure 2. The annual AMO, annual NAO indices by year (1991-2016) with the standard error of
728 the mean for each index.

729 Figure 3. Standardized (GAM) indices of abundance for Atlantic Croaker by age, survey (DE-
730 DFW 5.2 m; NJ-NJMF 4.9 m; CI-composite index) and year from 1991-2016, with the error bars
731 representing the 95 % Confidence Interval for each annual mean in the upper four panels. While
732 the lower four panels represent the corresponding median fitted ARIMA of standardized indices

733 of abundance by species, year and survey for the standardized index immediately above. In the
734 lower plots, the dots represent the $\ln(\text{index} + 0.01)$, and the gray line represents the ARIMA
735 index, while the red line represents the first quartile for the time series for each ARIMA.

736 Figure 4. Standardized (GAM) indices of abundance for YOY Black Drum by survey (DE-DFW
737 5.2 m; NJ-NJMF 4.9 m; CI-composite index) and year from 1991-2016, with the error bars
738 representing the 95 % Confidence Interval for each annual mean in the upper three panels. While
739 the lower three panels represent the corresponding median fitted ARIMA of standardized indices
740 of abundance by species, year and survey for the standardized index immediately above. In the
741 lower plots, the dots represent the $\ln(\text{index} + 0.01)$, and the gray line represents the ARIMA
742 index, while the red line represents the first quartile for the time series for each ARIMA.

743 Figure 5. Standardized (GAM) indices of abundance for Spot by age, survey (DE-DFW 5.2 m;
744 NJ-NJMF 4.9 m; CI-composite index) and year from 1991-2016, with the error bars representing
745 the 95 % Confidence Interval for each annual mean in the upper four panels. While the lower
746 four panels represent the corresponding median fitted ARIMA of standardized indices of
747 abundance by species, year and survey for the standardized index immediately above. In the
748 lower plots, the dots represent the $\ln(\text{index} + 0.01)$, and the gray line represents the ARIMA
749 index, while the red line represents the first quartile for the time series for each ARIMA.

750 Figure 6. Standardized (GAM) indices of abundance for Weakfish by age, survey (DE-DFW 5.2
751 m; NJ-NJMF 4.9 m; CI-composite index) and year from 1991-2016, with the error bars
752 representing the 95 % Confidence Interval for each annual mean in the upper six panels. While
753 the lower six panels represent the corresponding median fitted ARIMA of standardized indices
754 of abundance by species, year and survey for the standardized index immediately above. In the
755 lower plots, the dots represent the $\ln(\text{index} + 0.01)$, and the gray line represents the ARIMA
756 index, while the red line represents the first quartile for the time series for each ARIMA.

757 Figure 7. A. Length frequency of Atlantic Croaker by survey and across all years from 1991-
758 2016. DFW - Mean: 44 mm, Median: 25 mm. NJMF - Mean: 31 mm, Median: 23 mm.

759 B. Length frequency of Black Drum by survey and across all years from 1991-2016. DFW -
760 Mean: 171 mm, Median: 170 mm. NJMF - Mean: 166 mm, Median: 165 mm.

761 C. Length frequency of Spot by survey and across all years from 1991-2016. DFW - Mean: 137
762 mm, Median: 140 mm. NJMF - Mean: 129 mm, Median: 130 mm.

763 D. Length frequency of Weakfish by survey and across all years from 1991-2016. DFW - Mean:
764 96 mm, Median: 80 mm. NJMF - Mean: 70 mm, Median: 57 mm.

765 Supplemental Figures:

766 Supplemental Figure 1. A plot of the NBGM, YOY Atlantic Croaker composite index (CI)
767 component smooth functions including AMO, NAO, salinity (ppt), station as a random effect and
768 temperature (°C) with two standard errors above and below the estimate of the smooth expressed
769 as dashed lines.

770 Supplemental Figure 2. A plot of the ZIPGM, YOY Black Drum composite index (CI)
771 component smooth functions including AMO, depth (m), NAO, salinity (ppt), station as a
772 random effect, temperature (°C) and the interaction between Year and State (labelled as Year)
773 with two standard errors above and below the estimate of the smooth expressed as dashed lines.

774 Supplemental Figure 3. A plot of the NBGM, YOY Spot composite index (CI) component
775 smooth functions including the AMO, depth (m), NAO, salinity (ppt), station as a random effect,
776 temperature (°C) and the interaction between Year and State (labelled as Year) with two standard
777 errors above and below the estimate of the smooth expressed as dashed lines.

778 Supplemental Figure 4. A plot of the NBGM, YOY Weakfish composite index (CI) component
779 smooth functions including the AMO, depth (m), NAO, salinity (ppt), station as a random effect,
780 temperature (°C) and the interaction between Year and State (labelled as Year) with two standard
781 errors above and below the estimate of the smooth expressed as dashed lines.

782 Supplemental Figure 5. A plot of the NBGM, Age-1 Weakfish composite index (CI)
783 component smooth functions including the AMO, depth (m), salinity (ppt), station as a random
784 effect and temperature (°C) with two standard errors above and below the estimate of the smooth
785 expressed as dashed lines.

786

787 <A>References

788 Able K.W., M.M. Jones, D.A. Fox. 2009. Large Nektonic Fishes in Marsh Creek Habitats in the
789 Delaware Bay Estuary. *Northeastern Naturalist* 16(1):27-44.

790

791 Alexander M.A., K. Halimeda Kilbourne, J.A. Nye. 2014. Climate variability during warm and
792 cold phases of the Atlantic Multidecadal Oscillation (AMO) 1871-2008. *Journal of Marine*
793 *Systems* 133:14-26.

794
795 Anderson, M.J. 2001. A new method for non-parametric multivariate analysis of variance.
796 *Austral Ecology* 26: 32-46.
797
798 Aristizábal M. F., and R.J. Chant. 2015. An observational study of salt fluxes in Delaware Bay.
799 *Journal of Geophysical Research: Oceans* 120:2751–2768.
800
801 ASMFC. 2006. Weakfish Benchmark Stock Assessment: Weakfish Stock Assessment Report,
802 Terms of Reference and Advisory Report, and Technical Committee Supplemental Material.
803 Washington (DC): ASMFC.
804
805 ASMFC. 2009. Weakfish Benchmark Stock Assessment: Weakfish Stock Assessment Report,
806 Terms of Reference and Advisory Report, and Technical Committee Supplemental Material.
807 Washington (DC): ASMFC.
808
809 ASMFC. 2010. Atlantic Croaker Benchmark Stock Assessment: Atlantic Croaker Stock
810 Assessment Report for Peer Review. Washington (DC): ASMFC.
811
812 ASMFC. 2015. Black Drum Benchmark Stock Assessment: Black Drum Stock Assessment
813 Report, Terms of Reference and Advisory Report, and Technical Committee Supplemental
814 Material. Washington (DC): ASMFC.
815
816 ASMFC. 2016. Weakfish Benchmark Stock Assessment: Weakfish Stock Assessment Report,
817 Terms of Reference and Advisory Report, and Technical Committee Supplemental Material.
818 Washington (DC): ASMFC.
819
820 ASMFC. 2017a. Atlantic Sturgeon Benchmark Stock Assessment: Atlantic Sturgeon Stock
821 Assessment Report, Terms of Reference and Advisory Report, and Technical Committee
822 Supplemental Material. Washington (DC): ASMFC.
823

824 ASMFC. 2017b. Review of the Atlantic States Marine Fisheries Commission Fishery
825 Management Plan for Atlantic Croaker (*Micropogonias undulatus*). Washington (DC): ASMFC.
826

827 ASMFC. 2017c. Review of the Atlantic States Marine Fisheries Commission Fishery
828 Management Plan for Spot (*Leiostomus xanthurus*). Washington (DC): ASMFC.
829

830 ASMFC. 2017d. Review of the Atlantic States Marine Fisheries Commission Fishery
831 Management Plan for Weakfish (*Cynoscion regalis*). Washington (DC): ASMFC.
832

833 Barletta M., A. Barletta-Bergan, U. Saint-Paul, and G. Hubold. 2005. The role of salinity in
834 structuring the fish assemblages in a tropical estuary. The Fisheries Society of the British Isles,
835 Journal of Fish Biology, 66, 45–72.
836

837 Barnston, A. G., and R. E. Livezey. 1987: Classification, seasonality and persistence of low-
838 frequency atmospheric circulation patterns. Monthly Weather Review 115:1083-1126.
839

840 Boehlert, G.W., and B.C. Mundy. 1988. Roles of Behavioral and Physical Factors in Larval and
841 Juvenile Fish Recruitment to Estuarine Nursery Areas. American Fisheries Society Symposium
842 3:51-67.
843

844 Bonzek C., P.J. Greer, and H. Austin. 1995. VIMS Juvenile Fish Trawl Survey, Juvenile Indices
845 1979-1994. Virginia Institute of Marine Science.
846

847 Box, G. E. P. and G. M. Jenkins. 1976. Time series analysis: forecasting and control, revised Ed.
848 Holden-Day Oakland, CA 375 pp.
849

850 Brander K., and R. Mohn. 2004. Effect of the North Atlantic Oscillation on recruitment of
851 Atlantic cod (*Gadus morhua*). Canadian Journal of Fisheries and Aquatic Sciences 61:1558-
852 1564.
853

854 Buchheister A., T.J. Miller, E.D. Houde, D.H. Secor, and R.J. Latour. 2016. Spatial and temporal
855 dynamics of Atlantic Menhaden (*Brevoortia tyrannus*) recruitment in the Northwest Atlantic
856 Ocean. *ICES Journal of Marine Science* 73(4):1147-1159.

857

858 Burnham K. P., and D.R. Anderson. 2002. *Model Selection and Multimodel Inference: A*
859 *Practical Information-Theoretic Approach*. Springer, New York, NY. 488 pp.

860

861 Carassou, L., B. Dzwonkowski, F.J. Hernandez, S.P. Powers, K. Park, W.M. Graham and J.
862 Mareska. 2011. Environmental Influences on Juvenile Fish Abundances in a River-Dominated
863 Coastal System, *Marine and Coastal Fisheries* 3(1):411-427.

864

865 Chao, L.N., and J.A. Musick. 1977. Life history, feeding habits, and functional morphology of
866 juvenile sciaenid fishes in the York River Estuary, Virginia. *Fishery Bulletin* 75:657-702.

867

868 Charnov E.L., and J.F. Gillooly. 2004. Size and Temperature in the Evolution of Fish Life
869 Histories. *Integrative and Comparative Biology* 44(6):494-497.

870

871 Connaughton M.A., M.H. Taylor, M.L. Fine. 2000. Effects of fish size and temperature on
872 Weakfish disturbance calls: Implications for the mechanism of sound generation. *The Journal of*
873 *Experimental Biology* 203:1503-1512.

874

875 Craig K.J., and L.B. Crowder. 2005. Hypoxia-induced habitat shifts and energetic consequences
876 in Atlantic Croaker and brown shrimp on the Gulf of Mexico shelf. *Marine Ecology Progress*
877 *Series* 294:79-94.

878

879 Curry P.H., Y. Shin, B. Planque, J.M. Durant, J. Fromentin, S. Kramer-Schadt, N.C. Stenseth, M.
880 Travers, V. Grimm. 2008. Ecosystem oceanography for global change in fisheries. *Trends in*
881 *Ecology & Evolution* 23(6):338-346.

882

883 Dijkstra H.A., L. Te Raa, M. Schmeits, J. Gerrits. 2006. On the physics of the Atlantic
884 Multidecadal Oscillation. *Ocean Dynamics* 56:36-50.

885
886 Dong B., R.T. Sutton, and A.A. Scaife. 2006. Multidecadal modulation of El Niño-Southern
887 Oscillation (ENSO) variance by Atlantic Ocean sea surface temperatures. *Geophysical Research*
888 *Letters* 33:L08705 1-4.
889
890 Drexler M., and C.H. Ainsworth. 2013. Generalized Additive Models Used to Predict Species
891 Abundance in the Gulf of Mexico: An Ecosystem Modeling Tool. *PLoS ONE* 8(5): e64458.
892
893 Dunning D.J., Q.E. Ross, K.A. McKown, J.B. Socrates. 2009. Effect of Striped Bass Larvae
894 Transported from the Hudson River on Juvenile Abundance in Western Long Island Sound.
895 *Marine and Coastal Fisheries: Dynamics, Management, and Ecosystem Science* 1:343-353.
896
897 Durkee J.D., J.D. Frye, C.M. Fuhrmann, M.C. Lacke, H.G. Jeong, and T.L. Mote. 2008. Effects
898 of the North Atlantic Oscillation on precipitation-type frequency and distribution in the eastern
899 United States. *Theoretical and Applied Climatology* 94:51-65.
900
901 Edwards M., G. Beaugrand, P. Helaouët, J. Alheit, S. Coombs. 2013. Marine Ecosystem
902 Response to the Atlantic Multidecadal Oscillation. *PLoS ONE* 8(2):e57212.
903
904 Emmanuel J. 2017. Histogram of Length Frequency Distribution Using ggplot2.
905 [https://jethroemmanuel.netlify.app/2017/12/16/histogram-length-frequency-distribution-using-](https://jethroemmanuel.netlify.app/2017/12/16/histogram-length-frequency-distribution-using-ggplot2/)
906 [ggplot2/](https://jethroemmanuel.netlify.app/2017/12/16/histogram-length-frequency-distribution-using-ggplot2/)
907
908 Enfield D.B., A.M. Mestas-Nuñez, P.J. Trimble. 2001. The Atlantic multidecadal oscillation and
909 its relation to rainfall and river flows in the continental U.S. *Geophysical Research Letters*
910 28(10):2077-2080.
911
912 Epifanio, C. E., and R. W. Garvine. 2001. Larval transport on the Atlantic continental shelf of
913 North America: A review. *Estuarine, Coastal and Shelf Science* 52:51–77.
914

915 Fiedler P.C., R.D. Methot, R.P. Hewitt. 1986. Effects of California El Niño 1982-1984 on the
916 northern anchovy. *Journal of Marine Research* 44:317-338.
917

918 Garcia S.M., and K.L. Cochrane. 2005. Ecosystem approach to fisheries: a review of
919 implementation guidelines. *ICES Journal of Marine Science* 62(3):311-318.
920

921 Grecoy P.A., and T.A. Targett. 1996. Spatial Patterns in Condition and Feeding of Juvenile
922 Weakfish in Delaware Bay. *Transactions of the American Fisheries Society* 125:5, 803-808.
923

924 Greco M.J., 2019. Coastal finfish assessment survey. Federal Aid in Fisheries Restoration
925 Project F-42-R-28. Annual Report. Delaware Division of Fish and Wildlife, Dover.
926

927 Hale E.A., T.E. Targett. 2018. Vertical distribution of larval Atlantic menhaden (*Brevoortia*
928 *tyrannus*) and Atlantic Croaker (*Micropogonias undulatus*): Implications for vertical migratory
929 behaviour and transport. *Fisheries Oceanography* 00:1–10.
930

931 Hare J.A., and J.J. Govoni. 2005. Comparison of average larval fish vertical distributions among
932 species exhibiting different transport pathways on the southeast United States continental shelf.
933 *Fisheries Bulletin* 103: 728-736.
934

935 Hare J.A., W.E. Morrison, M.W. Nelson, M.M. Stachura, E.J. Teeters, R.B. Griffis, M.A.
936 Alexander, J.D. Scott, L. Alade, R.J. Bell, A.S. Chute, K. Curtis, T.H. Curtis, D. Kircheis, J.F.
937 Kocik, S.M. Lucey, C.T. McCandless, L.M. Milke, D. Richardson, E. Robillard, H.J. Walsh, C.
938 McManus, K.E. Marancik, and C.A. Griswold. 2016. A Vulnerability Assessment of Fish and
939 Invertebrates to Climate Change on the Northeast U.S. Continental Shelf. *PLoS ONE* 11(2):1-30.
940

941 Hare J.A., and Able K.W. 2007. Mechanistic links between climate and fisheries along the east
942 coast of the United States: explaining population outbursts of Atlantic Croaker (*Micropogonias*
943 *undulatus*). *Fisheries Oceanography* 16(1):31-45.
944

945 Harris V., M. Edwards, and S.C. Olhede. 2014. Multidecadal Atlantic climate variability and its
946 impact on marine pelagic communities. *Journal of Marine Systems* 133:55-69.
947

948 Hassall, A. 2019. Determination of relative abundance of selected finfish species in Delaware
949 Bay, New Jersey. Federal aid to Sport Fish Restoration Act. Project F-15-R-57. Annual Report.
950 New Jersey Department of Environmental Protection Division of Fish and Wildlife Marine
951 Fisheries Administration Bureau of Marine Fisheries.
952

953 Hastie T., and R. Tibshirani. 1986. Generalized Additive Models. *Statistical Science* 1:3, 297
954 318.
955

956 Helser T.E., and D. B. Hayes. 1995. Providing quantitative management advice from stock
957 abundance indices based on research surveys. *Fishery Bulletin* 93:290-298.
958

959 Helser T.E., Punt A. E. and R. D. Methot. 2004. A generalized linear mixed model analysis of a
960 multi-vessel fishery resource survey. *Fisheries Research* 70, 251–264.
961

962 Hill G., M. Fine, and J. Musick. 1987. Ontogeny of the Sexually Dimorphic Sonic Muscle in
963 Three Sciaenid Species. *Copeia* 1987(3), 708-713.
964

965 Hurrell J.W. 1995. Decadal Trends in the North Atlantic Oscillation: Regional Temperatures and
966 Precipitation. *Science* 269:676-679.
967

968 Hurrell J.W., and C. Deser. 2010. North Atlantic climate variability: The role of the North
969 Atlantic Oscillation. *Journal of Marine Systems* 79:231-244.
970

971 Irby I.D., M.A.M. Friedrichs, F. Da, K.E. Hinson. 2018. The competing impacts of climate
972 change and nutrient reductions on dissolved oxygen in Chesapeake Bay. *Biogeosciences* 15:
973 2649-2668.
974

975 Izzo L.K., and J. Zydlewski. 2017. Retrospective Analysis of Seasonal Ocean Growth Rates of
976 Two Sea Winter Atlantic Salmon in Eastern Maine Using Historic Scales. *Marine and Coastal*
977 *Fisheries* 9(1):357-372.

978

979 Janzen C.D., and K.C. Wong. 2002. Wind-forced dynamics at the estuary-shelf interface of a
980 large coastal plain estuary. *Journal of Geophysical Research* 107(C10):3138.

981

982 Jiao Y., E.P. Smith, R. O'Reilly and D.J. Orth. 2012. Modelling non-stationary natural mortality
983 in catch-at-age models. *ICES Journal of Marine Science* 69(1):105-118.

984

985 Jing Y.J., Y.C. Li, Y.F. Xu and G.Z. Fan. 2019. Influences of the NAO on the North Atlantic
986 CO₂ Fluxes in Winter and Summer on the Interannual Scale. *Advances in Atmospheric Sciences*
987 36(11): 1288-1298.

988

989 Johnson J.C. 2013. Quantification of life history parameters in South Carolina spot (*Leiostomus*
990 *xanthurus*). Thesis. Print. ProQuest LLC Dissertations Publishing 1536459.

991

992 Jonsson N., and B. Jonsson. 2004. Size and age of maturity of Atlantic salmon correlate with the
993 North Atlantic Oscillation Index (NAOI). *Journal of Fish Biology* 64:241-247.

994

995 Knudsen M.F., M. Seidenkrantz, B.H. Jacobsen, and A. Kuijpers. 2011. Tracking the Atlantic
996 Multidecadal Oscillation through the last 8,000 years, *Nature Communications*. 2(178):1-8.

997

998 Lankford T.E., and T.E. Targett. 1994. Suitability of estuarine nursery zones for juvenile
999 weakfish (*Cynoscion regalis*): effects of temperature and salinity on feeding, growth and
1000 survival. *Marine Biology* 119:611-620.

1001

1002 Lankford T.E., and T.E. Targett. 2001. Low-Temperature Tolerance of Age-0 Atlantic Croakers:
1003 Recruitment Implications for U.S. Mid-Atlantic Estuaries. *Transactions of the American*
1004 *Fisheries Society* 130(2):236-249.

1005

1006 Large S.I., G. Fay, K.D. Friedland, and J.S. Link. 2013. Defining trends and thresholds in
1007 responses of ecological indicators to fishing and environmental pressures. *ICES Journal of*
1008 *Marine Science* 70:755-767.

1009

1010 Larkin N.K., and D.E. Harrison. 2005. On the definition of El Niño and associated seasonal
1011 average U.S. weather anomalies. *Geophysical Research Letters* 32(L13705):1-4.

1012

1013 Lehodey P., J. Alheit, M. Barange, T. Baumgartner, G. Beaugrand, K. Drinkwater, J.-M.
1014 Fromentin, S.R. Hare, G. Ottersen, R.I. Perry, C. Roy, C.D. Van der Lingen, and F. Werner.
1015 2006. Climate Variability, Fish, and Fisheries. *Journal of Climate* 19(20):5009-5030.

1016

1017 Liao H., C. Jones, and J.L. Gilmore. 2019. 2018 Final Report Virginia and Chesapeake Bay
1018 Finfish Ageing and Population Analysis. Center for Quantitative Fisheries Ecology Old
1019 Dominion University. Funded by contract No. F-126-R-16 from the Virginia Saltwater
1020 Recreational Development Fund through the Virginia Marine Resources Commission.

1021

1022 Love J.W., and E.B. May. 2007. Relationships Between Fish Assemblage Structure and Selected
1023 Environmental Factors in Maryland's Coastal Bays. *Northeastern Naturalist* 14(2):251-268.

1024

1025 Love J.W., P. Chigbu, and E.B. May. 2009. Environmental Variability Affects Distributions of
1026 Coastal Fish Species (Maryland). *Northeastern Naturalist* 16(2):255-268.

1027

1028 Lowerre-Barbieri S.K., M.E. Chittenden, and L.R. Barbieri. 1995. Age And Growth Of
1029 Weakfish, *Cynoscion Regalis*, In The Chesapeake Bay-Region With A Discussion Of Historical
1030 Changes In Maximum Size. *Fishery Bulletin*, 93(4), 643-656.

1031

1032 Mateo I., and D. H. Hanselman. 2014. A comparison of statistical methods to standardize catch-
1033 per-unit-effort of the Alaska longline sablefish fishery. U.S. Dep. Commer., NOAA Tech.
1034 Memo. NMFS-AFSC-269, 71 p.

1035

1036 Methot Jr., and D. Richard (editor). 2015. Prioritizing fish stock assessments. U.S. Department
1037 of Commerce. NOAA Tech. Memo NMFS-F/SPO152, 31 p.
1038

1039 Michels S.F., M.J. Greco. 1995. Delaware's Juvenile Recruitment Indices 1978-1994. Federal
1040 Aid in Fisheries Restoration Project F-42-R-28. Annual Report. Delaware Division of Fish and
1041 Wildlife, Dover.
1042

1043 Miller M.J., D.M. Nemerson, K.W. Able. 2003. Seasonal distribution, abundance, and growth of
1044 young-of-the-year Atlantic croaker (*Micropogonias undulatus*) in Delaware Bay and adjacent
1045 marshes. Fishery Bulletin 101(1):100-115.
1046

1047 Misund O.A., N. Luyeye, J. Coetzee, and D. Boyer. 1999. Trawl sampling of small pelagic fish
1048 off Angola: effects of avoidance, towing speed, tow duration, and time of day. ICES Journal of
1049 Marine Science 56:275-283.
1050

1051 Nesbit R.A. 1954. Weakfish migration in relation to its conservation. United States Department
1052 of the Interior/ Fish and Wildlife Service, Special Scientific Report: Fisheries No. 115,
1053 Washington DC
1054

1055 Nogueira R.C., B.D. Keim, D.P. Brown, and K.D. Robbins. 2013. Variability of rainfall from
1056 tropical cyclones in the eastern USA and its association to the AMO and ENSO. Theoretical and
1057 Applied Climatology 112:273-283.
1058

1059 Nye J.A., J.S. Link, J.A. Hare, and W.J. Overholtz. 2009. Changing spatial distribution of fish
1060 stocks in relation to climate and population size on the Northeast United States continental shelf.
1061 Marine Ecological Progress Series 393:111-129.
1062

1063 Nye J.A., M.R. Baker, R. Bell, A. Kenny, K.H. Kilbourne, K.D. Friedland, E. Martino, M.M.
1064 Stachura, K.S. Van Houtan, and R. Wood. 2014. Ecosystem effects of the Atlantic Multidecadal
1065 Oscillation. Journal of Marine Systems 133:103-116.
1066

1067 Nye, J.A., Targett, T.E., and T.E. Helser. 2008. Reproductive characteristics of weakfish in
1068 Delaware Bay: implications for management. *North American Journal of Fisheries Management*
1069 27: 1-11.

1070

1071 O'Connor M.P., F. Juanes, and K. McGarigal. 2012. Findings on American Shad and Striped
1072 Bass in the Hudson River Estuary: A Fish Community Study of the Long-Term Effects of Local
1073 Hydrology and Regional Climate Change. *Marine and Coastal Fisheries: Dynamics,*
1074 *Management, and Ecosystem Science* 4:327-336.

1075

1076 Odell J., D. H. Adams, B. Boutin, W. Collier II, A. Deary, L. N. Havel, J. A. Johnson Jr., S. R.
1077 Midway, J. Murray, K. Smith, K. M. Wilke, and M. W. Yuen. 2017. Atlantic Sciaenid Habitats:
1078 A Review of Utilization, Threats, and Recommendations for Conservation, Management, and
1079 Research. Atlantic States Marine Fisheries Commission Habitat Management Series No. 14,
1080 Arlington, VA.

1081

1082 Ogle D. 2014. fishR Vignette – Size Structure Analysis.
1083 <http://derekogle.com/fishR/examples/oldFishRVignettes/SizeStructure.pdf>

1084

1085 Ostermeier G.M., and J.M. Wallace. 2003. Trends in the North Atlantic Oscillation-Northern
1086 Hemisphere Annular Mode during the Twentieth Century. *Journal of Climate Notes and*
1087 *Correspondence* 16:336-341.

1088

1089 Otterson G., B. Planque, A. Belgrano, E. Post, P.C. Reid, and N.C. Stenseth. 2001. Ecological
1090 effects of the North Atlantic Oscillation. *Oecologia* 128:1-14.

1091

1092 Overholtz W.J., J.A. Hare, and C.M. Keith. 2011. Impacts of Interannual Environmental Forcing
1093 and Climate Change on the Distribution of Atlantic Mackerel on the U.S. Northeast Continental
1094 Shelf. *Marine and Coastal Fisheries* 3(1):219-232.

1095

1096 Pearson, J. C. 1929. Natural history and conservation of redfish and other commercial sciaenids
1097 on the Texas coast. *Bulletin of the Bureau of Fisheries* 44:129-214.

1098
1099 Pennington, M. 1986. Some statistical techniques for estimating abundance indices from trawl
1100 surveys. *Fishery Bulletin* 84(3): 519-525.
1101
1102 R Development Core Team (2008). R: A language and environment for statistical computing. R
1103 Foundation for Statistical Computing, Vienna, Austria. ISBN 3-900051-07-0, URL
1104 <http://www.R-project.org>.
1105
1106 Reist J.D., F.J. Wrona, T.D. Prowse, M. Power, B.J. Dempson, R.J. Beamish, J.R. King, T.J.
1107 Carmichael, and C.D. Sawatzky. 2006. General Effects of Climate Change on Arctic Fishes and
1108 Fish Populations. *AMBIO: A Journal of the Human Environment* 35(7):370-380.
1109
1110 Rice A.N., J.L. Morano, K.B. Hodge, and C.A. Muirhead. 2016. Spatial and temporal patterns of
1111 toadfish and Black Drum chorusing activity in the South Atlantic Bight. *Environmental Biology*
1112 *of Fishes* 99:705-716.
1113
1114 Roberts S.M., A.M. Boustany, P.N. Halpin, and R.R. Rykaczewski. 2019. Cyclical climate
1115 oscillation alters species statistical relationships with local habitat. *Marine Ecology Progress*
1116 *Series* 614: 159–171.
1117
1118 Sagarese S.R., R.M. Cerrato, and M.G. Frisk. 2011. Diet Composition and Feeding Habits of
1119 Common Fishes in Long Island Bays, New York. *Northeastern Naturalist* 18(3):291-314.
1120
1121 Saucier M.H., and D.M. Baltz. 1993. Spawning site selection by spotted seatrout, *Cynoscion*
1122 *nebulosus*, and Black Drum, *Pogonias cromis*, in Louisiana. *Environmental Biology of Fishes*
1123 36:257-272.
1124
1125 Schaffler J.J., J.V. Montfrans, C.M. Jones, and R.J. Orth. 2013. Fish Species Distribution in
1126 Seagrass Habitats of Chesapeake Bay are Structured by Abiotic and Biotic Factors. *Marine and*
1127 *Coastal Fisheries: Dynamics, Management, and Ecosystem Science* 5:114-124.
1128

1129 Schieler B.M., E.A. Hale, and T.E. Targett. 2014. Daily variation in ingress of fall-spawned
1130 larval fishes into Delaware Bay in relation to alongshore and along-estuary wind components.
1131 Estuarine, Coastal and Shelf Science 151:141-147.
1132

1133 Schirripa M.J., C.P. Goodyear, and R.M. Methot. 2009. Testing different methods of
1134 incorporating climate data into the assessment of US West Coast sablefish. ICES Journal of
1135 Marine Science 66:1605-1613.
1136

1137 Schlesinger M.E., and N. Ramankutty. 1994. An oscillation in the global climate system of
1138 period 65-70 years. Nature 367:723-726.
1139

1140 Stenseth N.C., A. Mysterud, G. Ottersen, J.W. Hurrell, K. Chan, and M. Lima. 2002. Ecological
1141 Effects of Climate Fluctuations. Science 297:1292-1296.
1142

1143 Sweka, J. 2015. Proposed Method for Estimating Adult Horseshoe Crab Abundance for Adaptive
1144 Resource Management in the Delaware Bay. Report to the Delaware Bay Ecosystem Technical
1145 Committee
1146

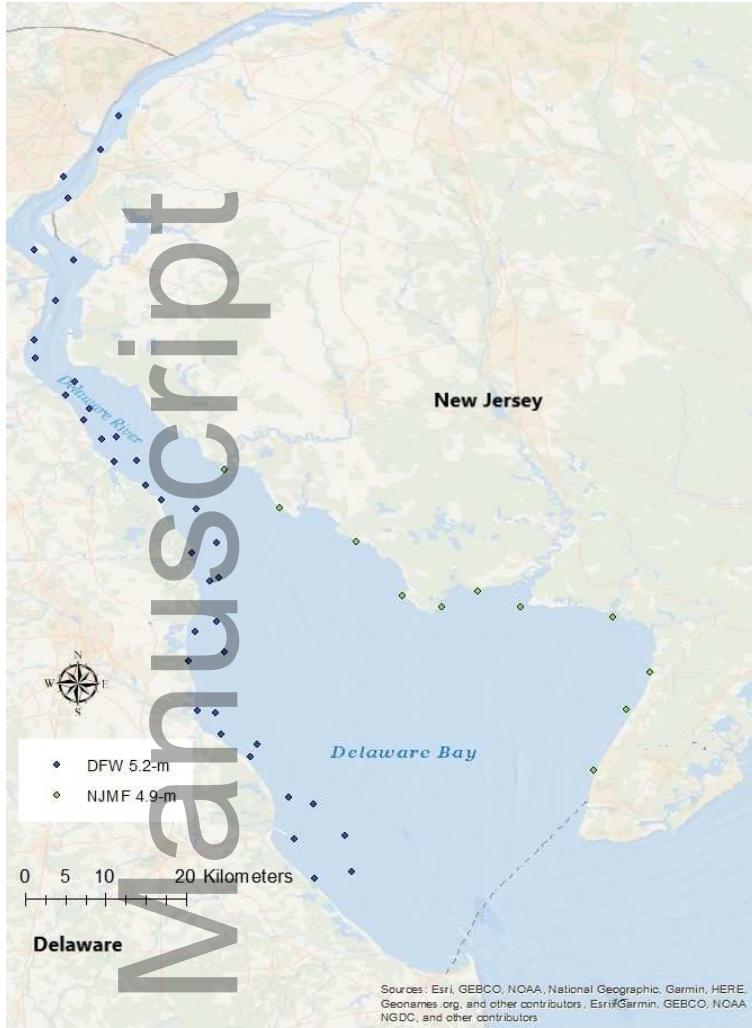
1147 Thorson J. T., and C. Minto. 2015. Mixed effects: a unifying framework for statistical modelling
1148 in fisheries biology. ICES Journal of Marine Science 72:1245–1256.
1149

1150 Thomas D.L., and B.A. Smith. 1973. Studies of Young of the Black Drum, *Pogonias cromis*, in
1151 Low Salinity Waters of the Delaware Estuary. Chesapeake Science 14(2):124-130.
1152

1153 Turner S.M., J.A. Hare, D.E. Richardson, and J.P. Manderson. 2017. Trends and Potential
1154 Drivers of Distribution Overlap of River Herring and Commercially Exploited Pelagic Marine
1155 Fishes on the Northeast U.S. Continental Shelf. Marine and Coastal Fisheries 9(1):13-22.
1156

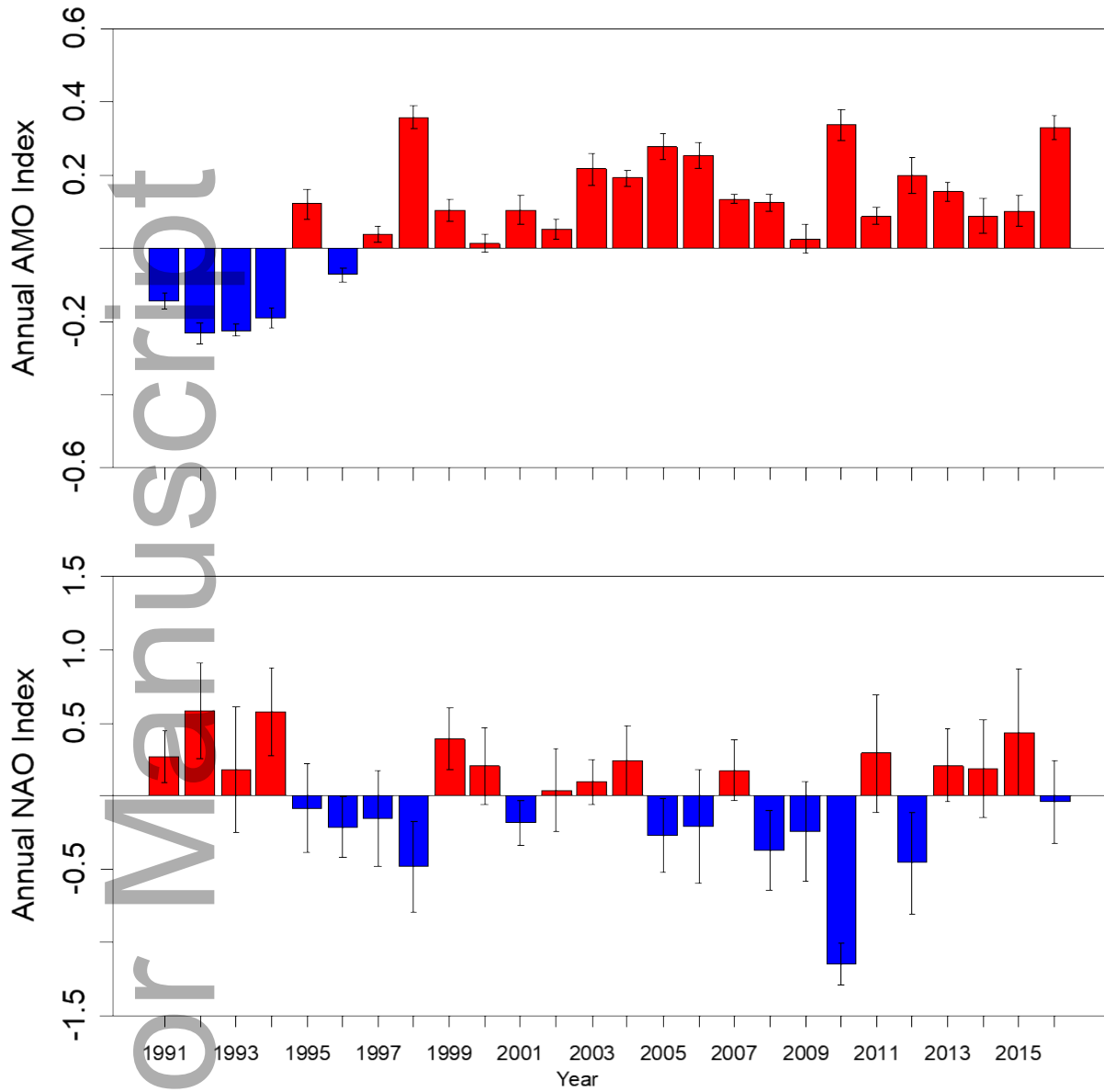
1157 Turnure J.T., T.M. Grothues, and K.W. Able. 2015. Seasonal residency of adult Weakfish
1158 (*Cynoscion regalis*) in a small temperate estuary based on acoustic telemetry: a local perspective
1159 of a coast wide phenomenon. Environmental Biology of Fishes 98:1207-1221.

1160
1161 Venables, W. M., and C. M. Dichmont. 2004. GLMS, GAMS, and GLMMs: an overview of
1162 theory for applications in fisheries research. *Fisheries Research* 70:319–337.
1163
1164 Witting D.A., K.W. Able, and M.P. Fahay. 1999. Larval fishes of a Middle Atlantic Bight
1165 estuary: assemblage structure and temporal stability. *Canadian Journal of Fisheries and Aquatic*
1166 *Sciences* 56:222-230.
1167
1168 Yasumiishi E.M., E.V. Farley, G.T. Ruggerone, B.A. Agler, and L.I. Wilson. 2016. Trends and
1169 Factors Influencing the Length, Compensatory Growth, and Size-Selective Mortality of Juvenile
1170 Bristol Bay, Alaska, Sockeye Salmon at Sea. *Marine and Coastal Fisheries: Dynamics,*
1171 *Management, and Ecosystem Science* 8:315-333.
1172
1173 Zimmerman J.M. 2016. Age, Growth, and Reproduction of Black Drum (*Pogonias cromis*) in
1174 Delaware Bay, USA. Thesis, Print.
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1177 Figure 1.



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1179 Figure 2.



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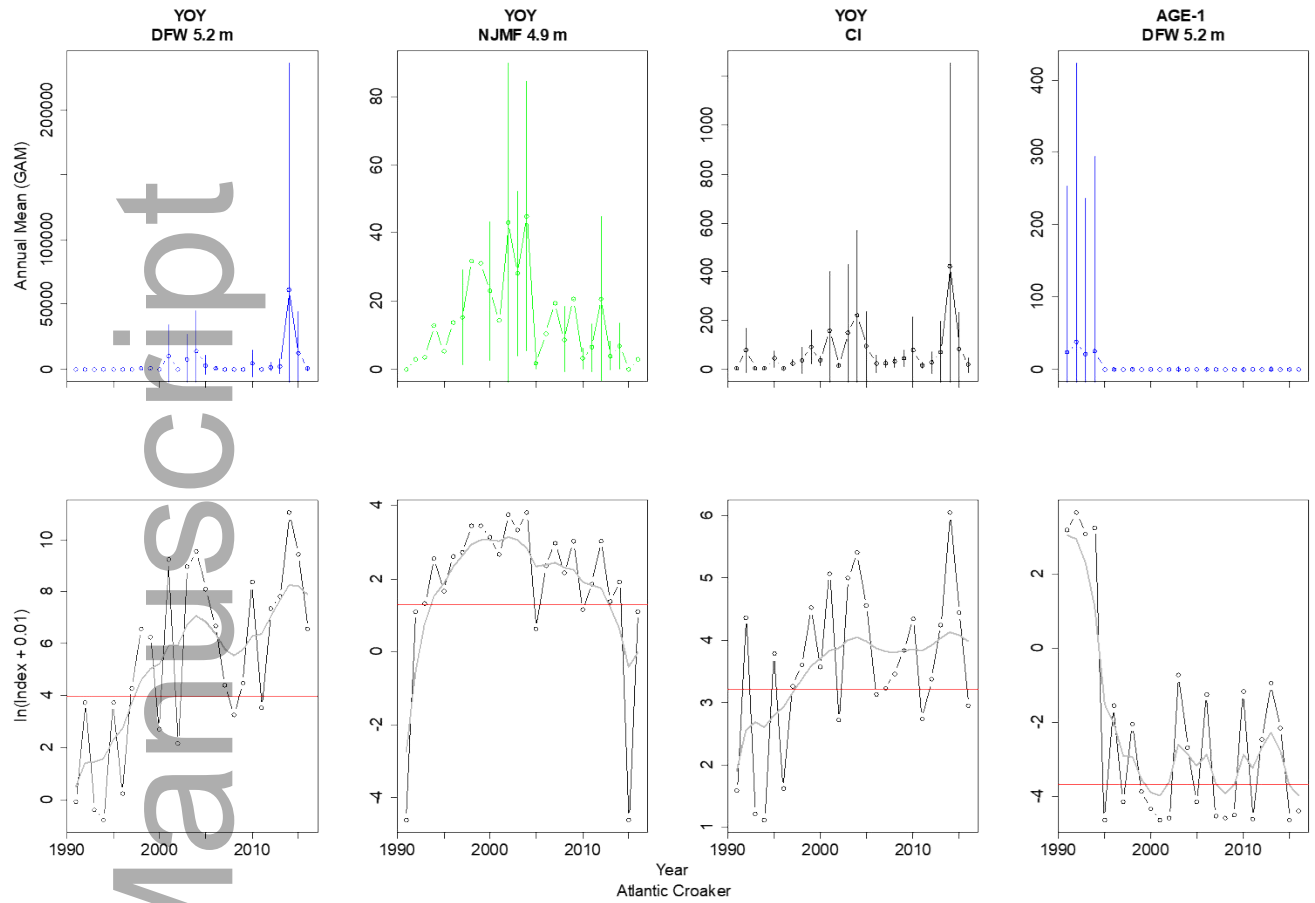
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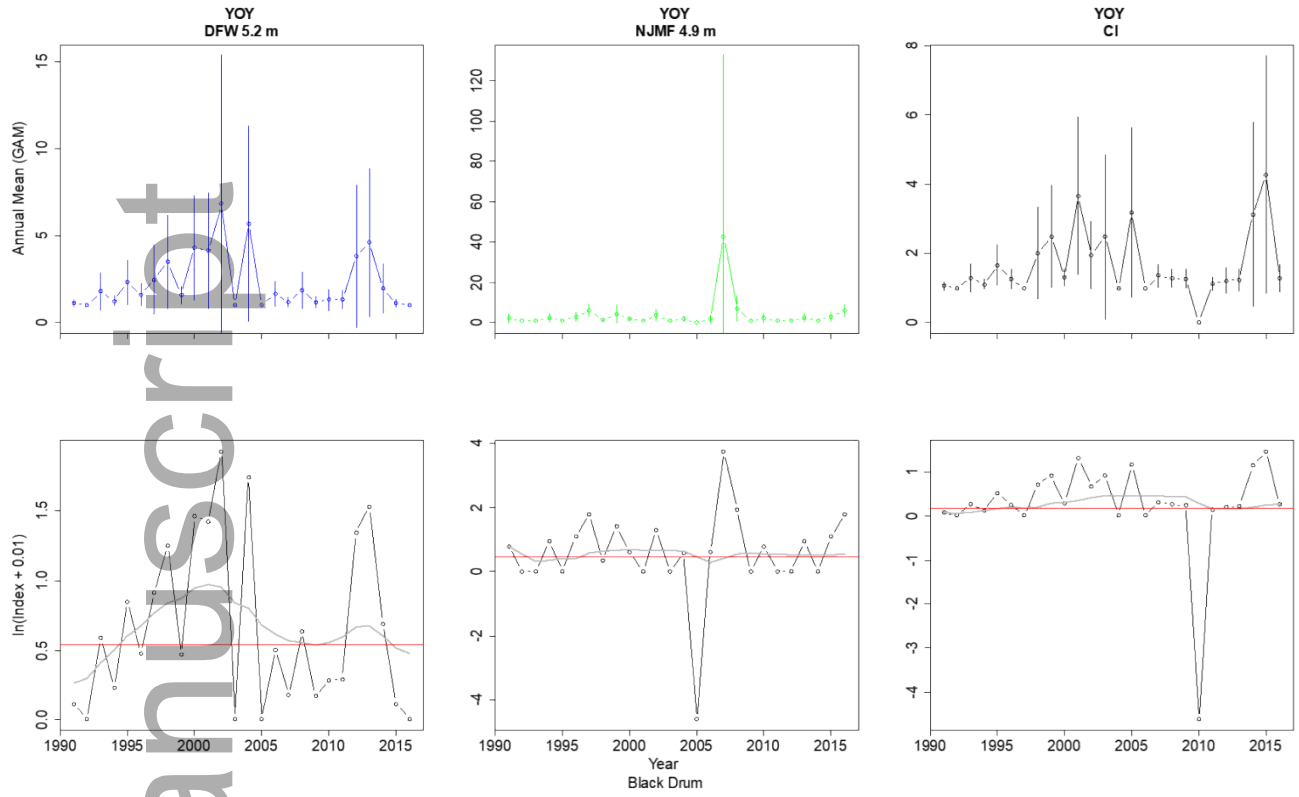
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Figure 3.



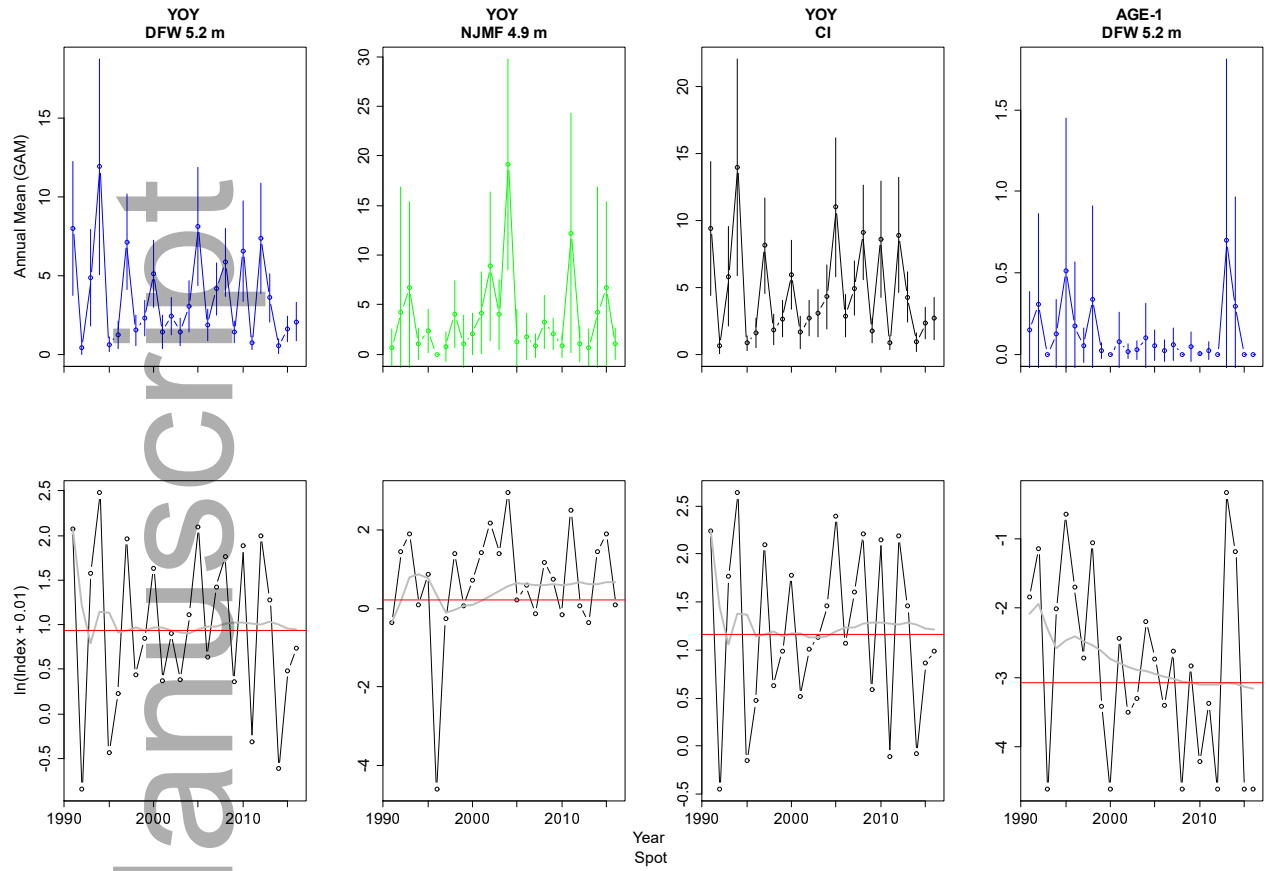
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Figure 4.



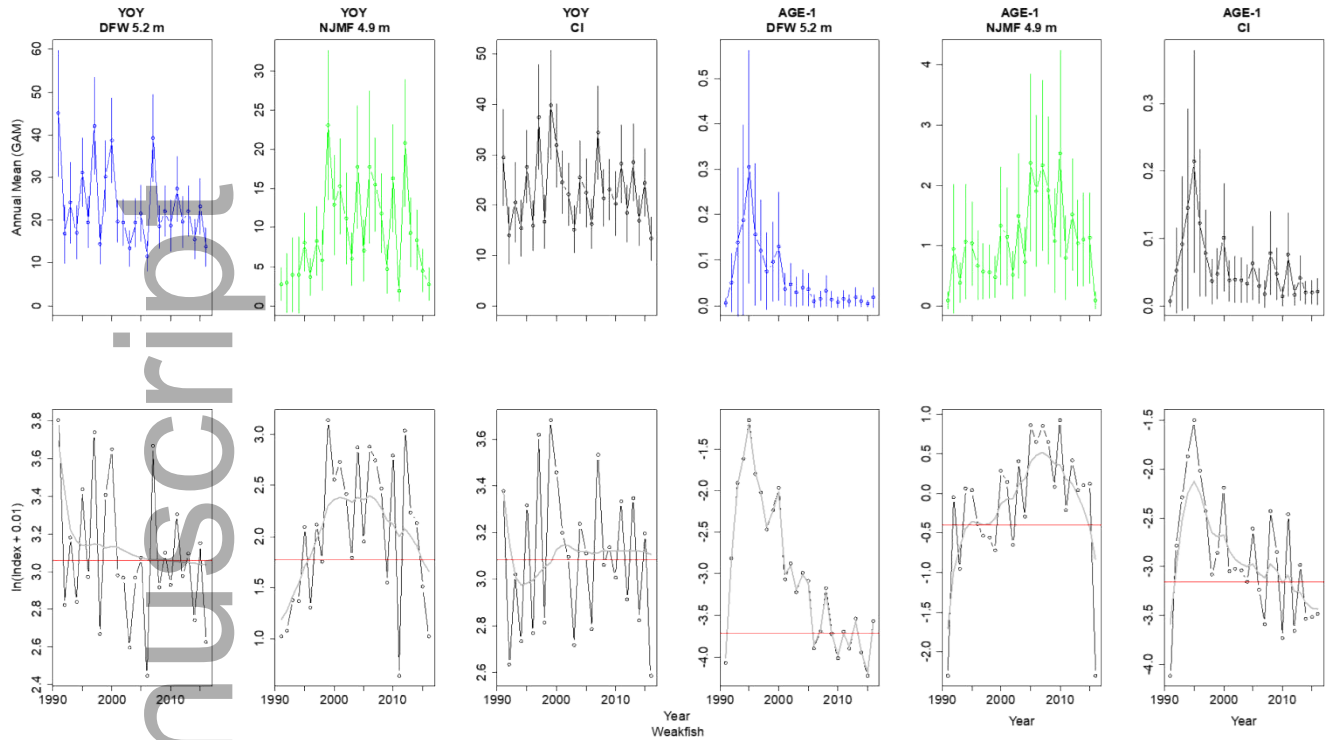
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Figure 5.



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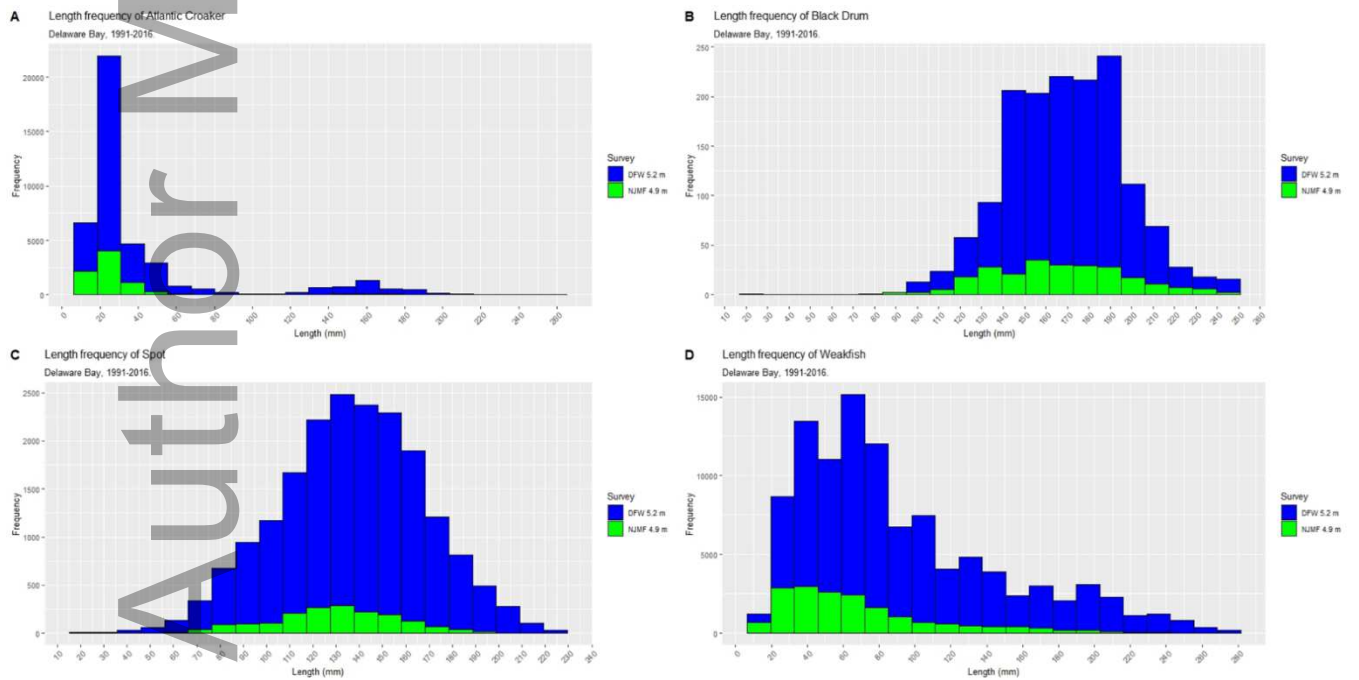
Figure 6.



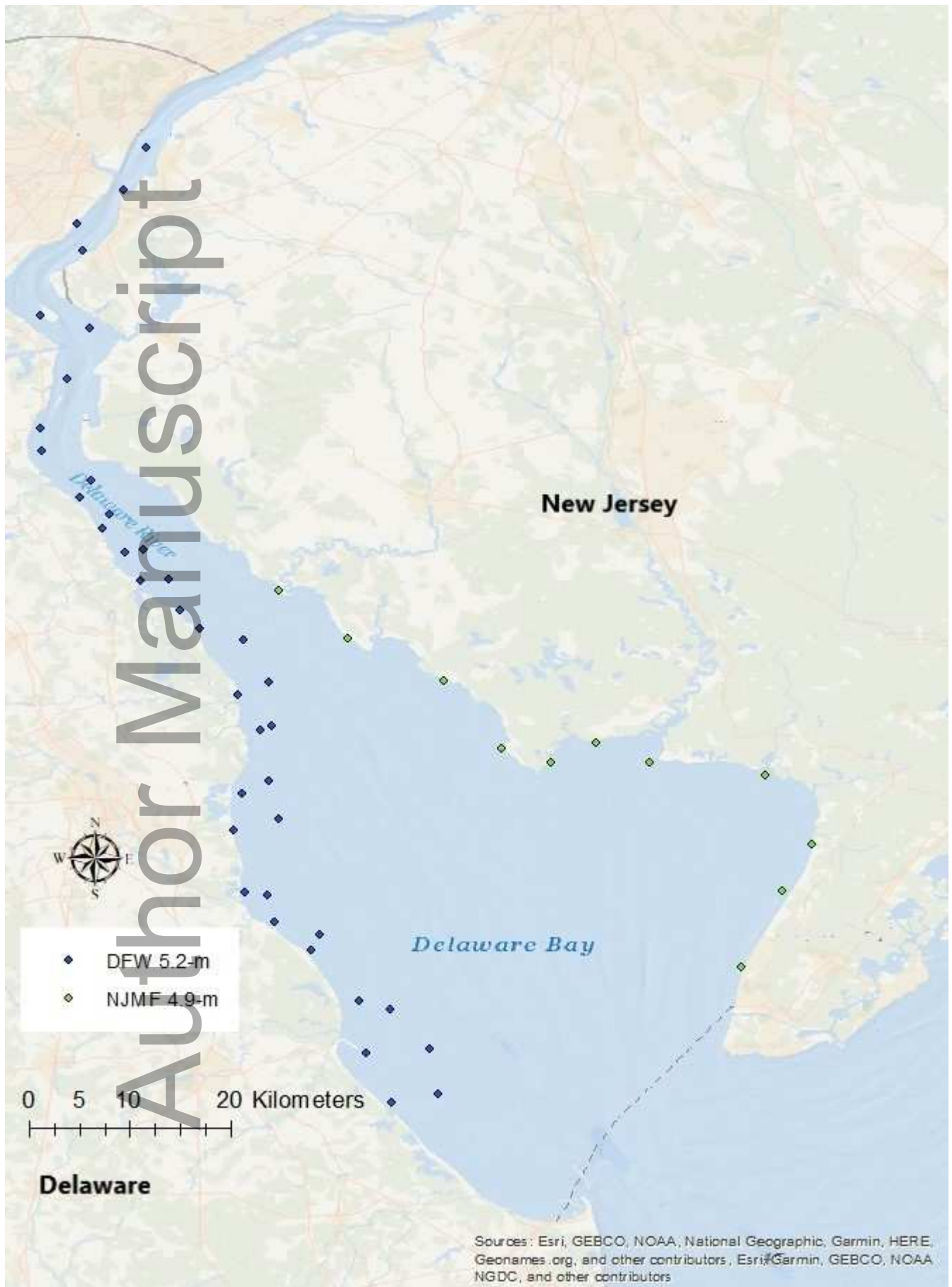
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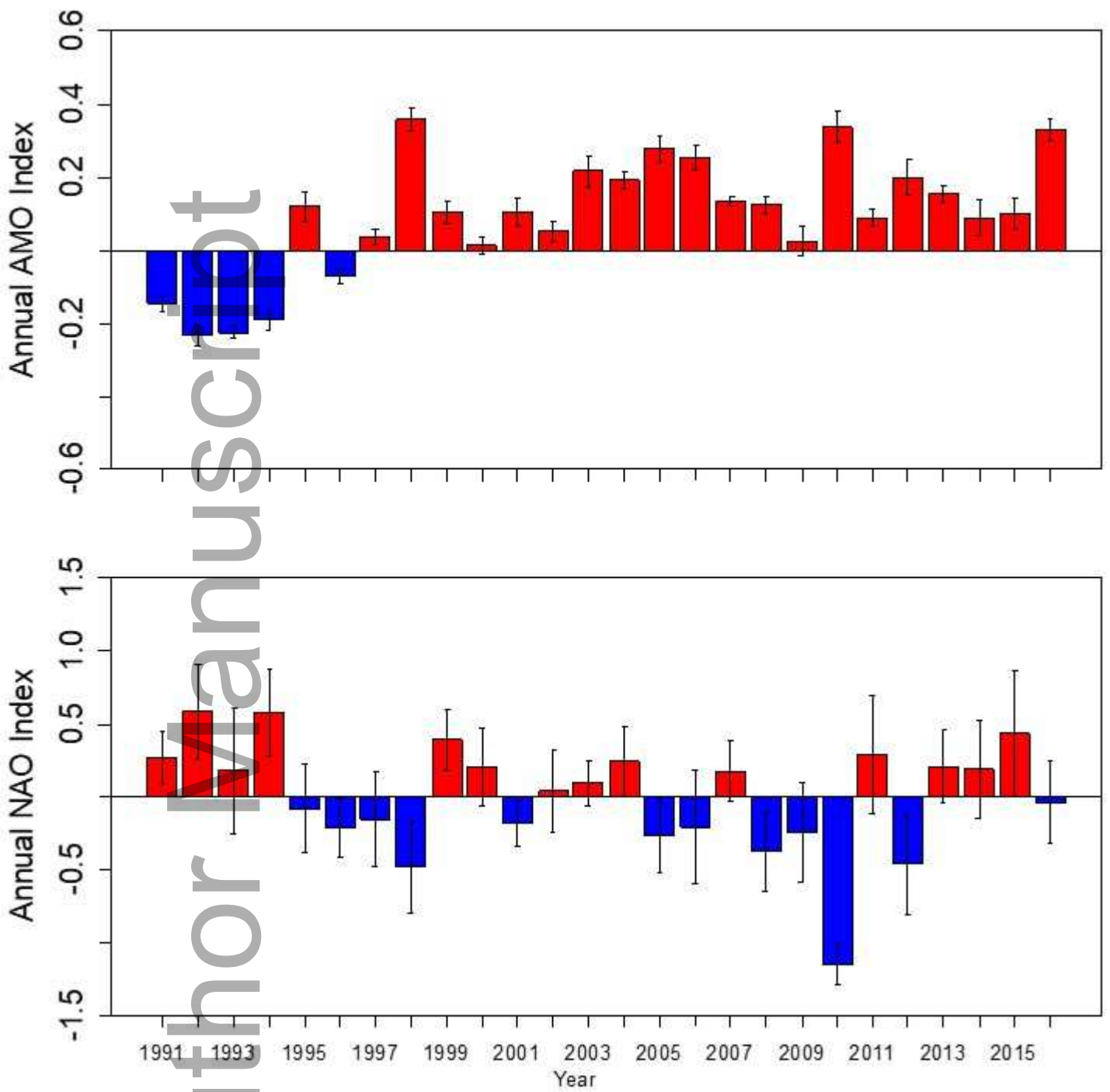
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1240 Figure 7.

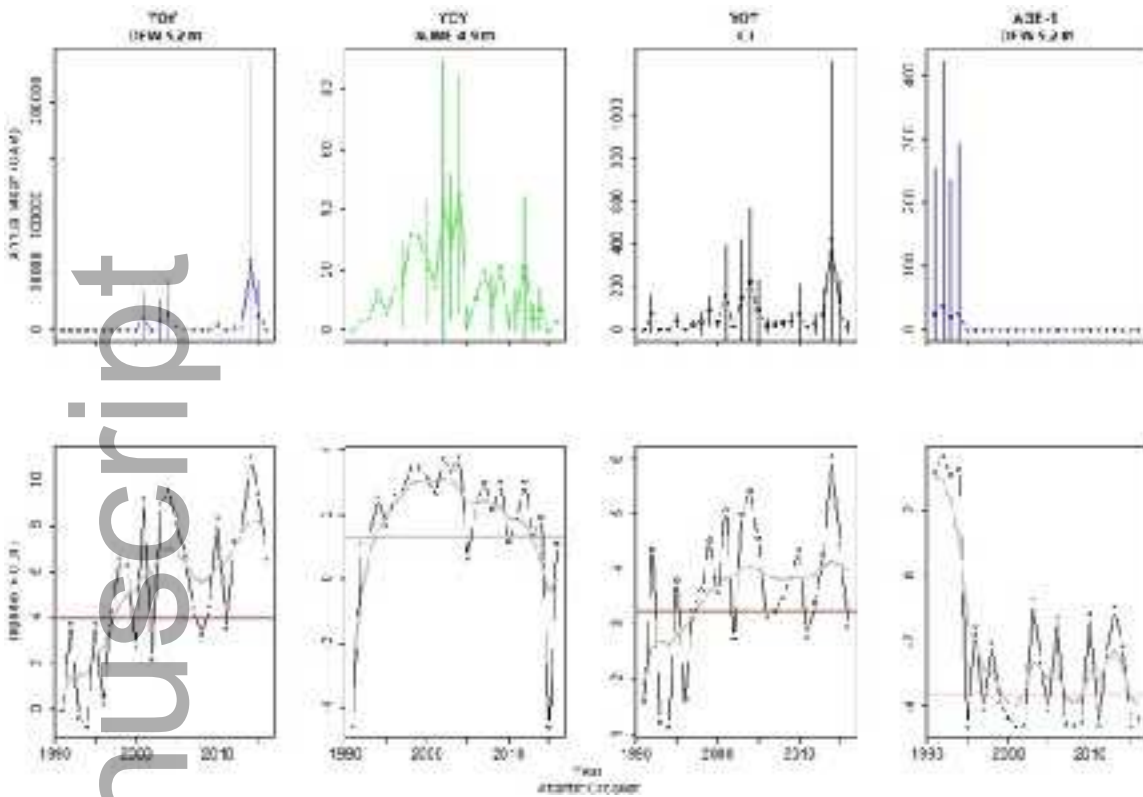


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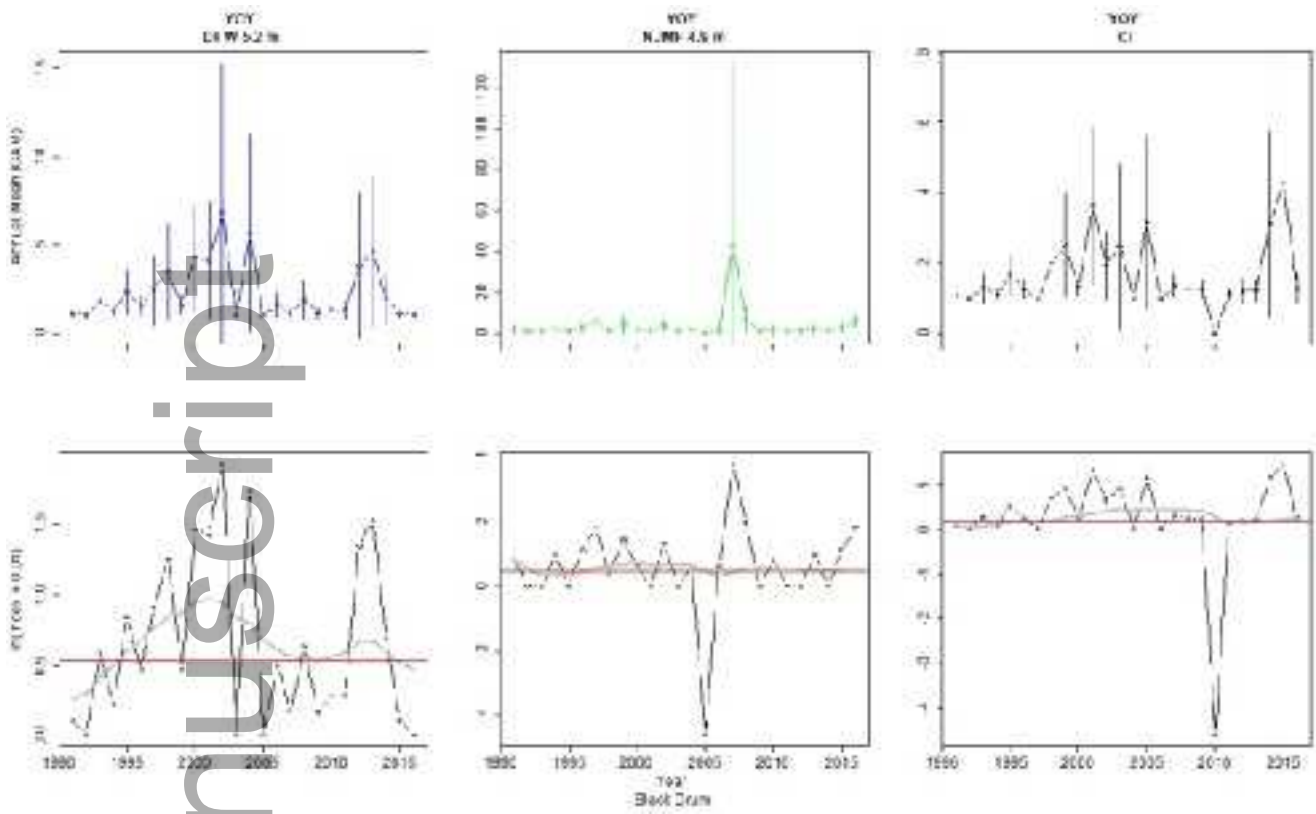




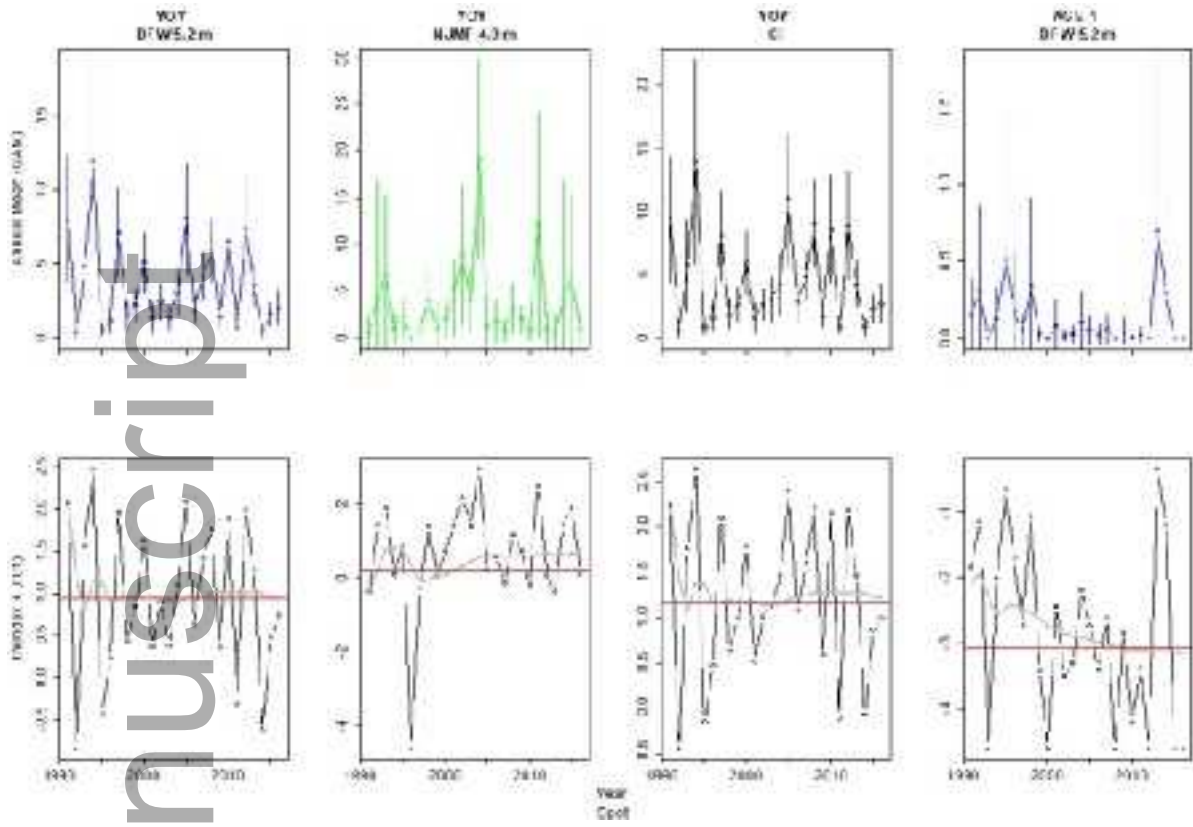
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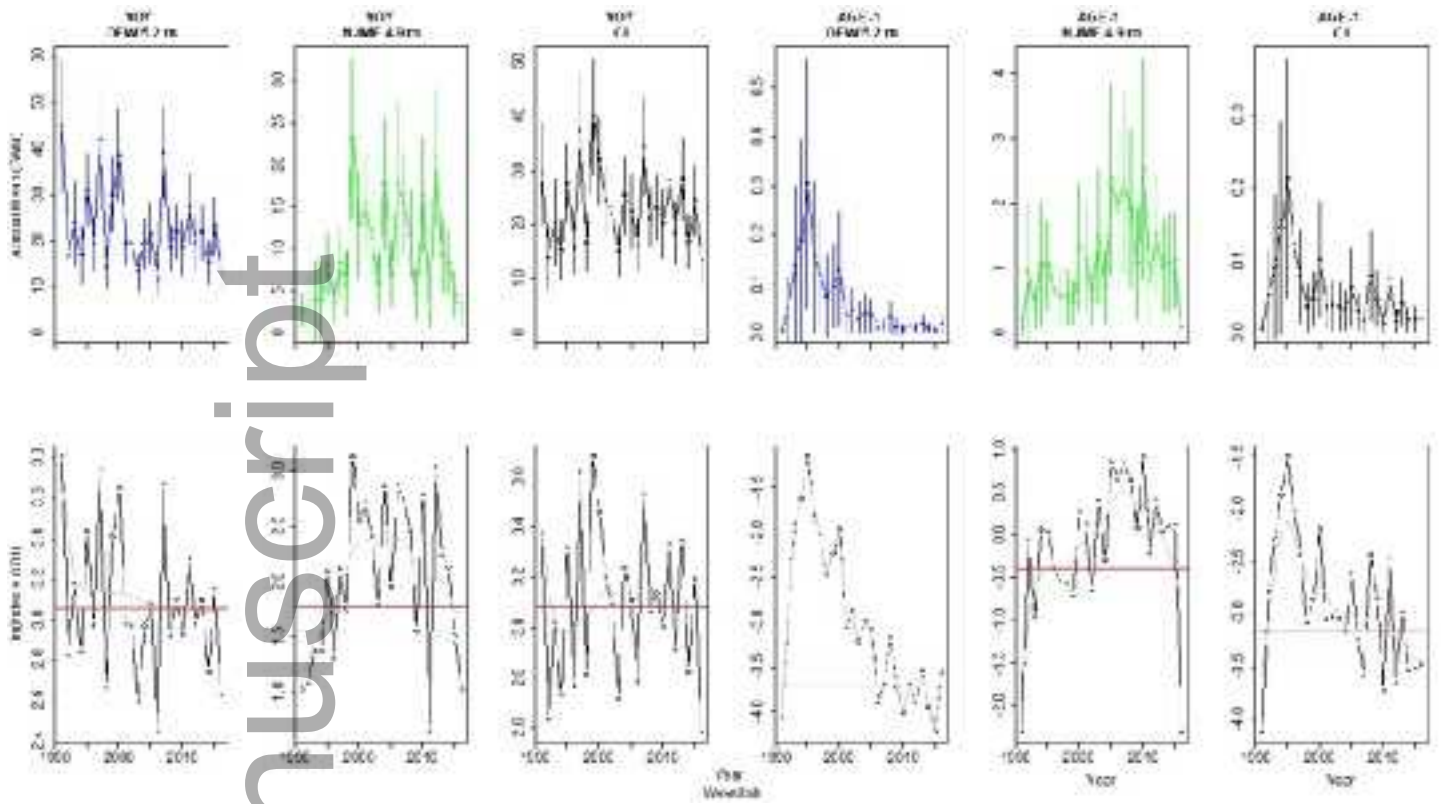
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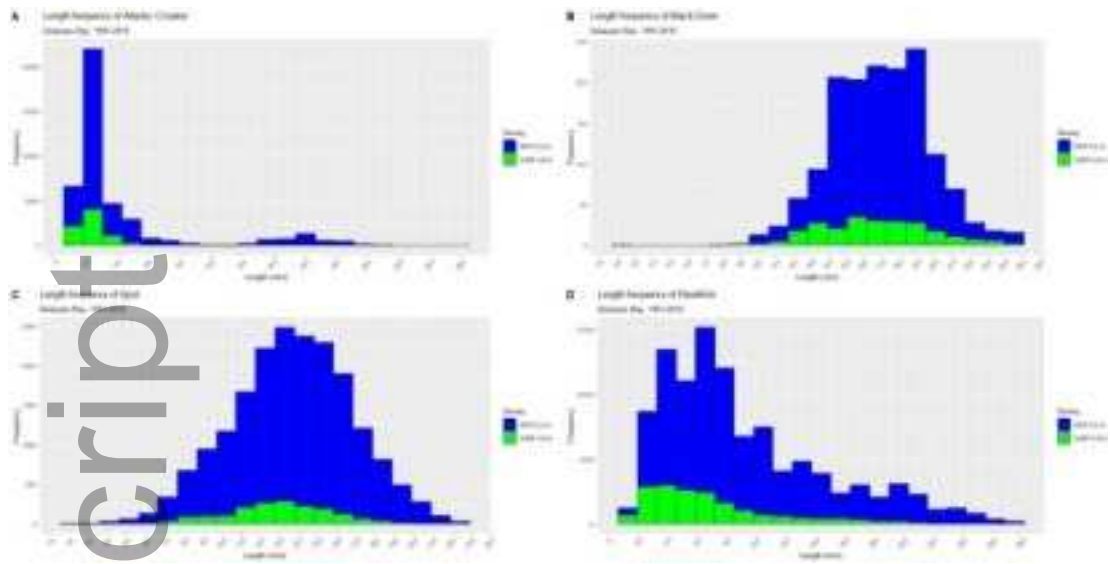
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