

Indices of abundance in the Gulf of Mexico reef fish complex: A comparative approach using spatial data from vessel monitoring systems

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Abstract

The Gulf of Mexico reef fish complex is socioeconomically important and is exploited by a vertical line fishery capable of high resolution spatial targeting. Indices of abundance derived from fishery dependent catch-per-unit-effort (CPUE) data are an important input to the assessment of these stocks. Traditionally, these indices have been derived from standardized logbook data, aggregated at a coarse spatial scale, and are limited to generating predictions for observed spatiotemporal strata. Understanding how CPUE is spatially distributed, however, can help identify range contractions and avoid hyperstability or hyperdepletion, both of which can mask the true population dynamics. Vessel monitoring systems (VMS) can provide complete, high-resolution distributions of CPUE used to create abundance indices. Here we compare two methods — spatial averaging of VMS-derived catch and effort data and the result of generalized linear models applied to logbook data for generating indices, to evaluate the use of VMS-derived abundance indices in assessments of reef fish stocks. This work suggests that in fisheries where targeting occurs at very fine spatial scales, abundance indices derived from high-resolution, spatiotemporally complete data may more accurately reflect the underlying dynamics of the stock.

Keywords: CPUE standardization; vessel monitoring systems; reef fish; abundance index; simulation; Gulf of Mexico

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37 1. Introduction

38 Abundance indices are an important input for stock assessments. Fisheries-dependent data,
39 such as catch-per-unit-effort (CPUE), are a common source of information for estimating trends
40 in abundance, as they typically represent a more spatiotemporally complete and cost effective
41 sample than fisheries-independent data (Ward, 2005).

42 Despite the availability of fishery dependent data, they may not be reliable as catch rates may
43 not adequately track abundance. Nominal CPUE are widely regarded as disproportionate to
44 abundance (Beverton and Holt, 1957; Harley et al., 2001) due to hyperstability - abundance
45 declining faster than CPUE, or hyperdepletion - CPUE declining faster than abundance (Hilborn,
46 and Walters 1992). These sources of non-linearity between CPUE and abundance can be
47 introduced through gear effects (saturation and handling time; Deriso and Parma, 1987)),
48 changes in fishing power (Bishop et al., 2004; Ye and Dennis, 2009), and interference between
49 vessels (Gillis and Peterman, 1998). In addition, discrepancies between the spatial distributions
50 of species abundance and fishing effort can exacerbate the issue if fishers are not representatively
51 sampling the underlying abundance distributions (Clark and Mangel, 1979; Paloheimo and
52 Dickie, 1964; Rose and Kulka, 1999; Rose and Leggett, 1991; Swain and Sinclair, 1994).

53 Bias in the relationship between CPUE and inferred abundance due to spatial distributions
54 are typically addressed using one of two approaches: standardization or spatial imputation. Catch
55 rates can be standardized using generalized linear models (GLMs) (Maunder and Punt, 2004;
56 Nelder and Wedderburn, 1972) to separate the abundance trend from other factors. If spatial
57 nominal CPUE data are available, they can be used to infer abundance trends provided they are
58 spatially and/or temporally imputed to account for unfished areas and changes in the
59 distributions of fishing effort (Walters, 2003). Abundance indices generated from spatially
60 imputed nominal CPUE data that randomly sample the entire underlying distribution have been
61 shown to track abundance accurately (Yu et al., 2013). However, for both of these approaches,
62 the level of data aggregation is important to consider. Bias in the inferred abundance can occur if
63 the level of data aggregation is too coarse such that fishing effort is no longer randomly sampling
64 abundance within spatiotemporal strata (Campbell, 2004; Carruthers et al., 2010). Spatially
65 averaging data on a fine spatial scale is more likely to represent the underlying abundance
66 distribution of non-transient species (Carruthers et al., 2011).

67 Vessel monitoring systems (VMS) have transformed the analysis of fisheries-dependent
68 spatial information. The high-resolution vessel location data provided by VMS have given
69 fisheries scientists and managers a better understanding of the spatial distribution of effort (Lee
70 et al., 2010; Mills et al., 2007), fisher behavior (Davie and Lordan, 2011; Vermard et al., 2010),
71 and the abundance distributions of targeted stocks (Bertrand et al., 2008; Vinther and Eero,
72 2013). Linking self-reported logbook catch records to VMS data has allowed for the creation of
73 species-specific distributions of CPUE in European trawl fisheries for groundfish (Gerritsen and
74 Lordan, 2011; Witt and Godley, 2007) and the vertical line fishery targeting reef fish in the Gulf
75 of Mexico (Ducharme-Barth and Ahrens, 2017).

76 The vertical line fishery in the Gulf of Mexico is a valuable commercial fishery (NMFS
77 2015, 2016) that targets a diverse complex comprised primarily of snappers, e.g. *Lutjanus spp*,
78 and groupers, e.g. *Epinephelus spp* (Scott-Denton et al., 2011). The four most commercially
79 encountered species (red snapper *Lutjanus campechanus*, vermilion snapper *Rhomboplites*
80 *aurorubens*, red grouper *Epinephelus morio*, and gag grouper *Mycteroperca microlepis*) can be
81 characterized by an association with easily identifiable hard bottom structure (Grimes, 1978;
82 Grimes and Huntsman, 1980; Lindberg et al., 2006; Moran, 1988) and high site fidelity

83 (Coleman et al., 2010, 2011). The vertical line gear (multiple baited lines dropped vertically from
84 a stationary or slowly drifting vessel) fished in multiple short sets (~20 minutes) allows for high
85 resolution spatial targeting of the hard bottom structure and the targeted fish stocks (Pollack et
86 al., 2013; SAFMC, 2009; Scott-Denton et al., 2011). This combination of targeting behavior and
87 species characteristics predisposes the fishery to the risk of hyperstability, particularly in the
88 absence of spatial information on where catches occur.

89 Given the unique set of coinciding circumstances between vertical line fisheries and reef fish
90 behavior, it is worthwhile to evaluate if developing abundance indices from higher resolution
91 catch and effort data from VMS gives a more accurate approximation of the underlying
92 abundance trends. Ideally, one would be able to work with data at a spatial resolution where
93 sampling is representative of the underlying abundance (Walters, 2003). However, the fishing
94 behavior of the vertical line fleet makes it unlikely that data aggregated at all but the finest scales
95 (e.g. reef or artificial structure) meet this criterion. The current practice for generating abundance
96 indices in this fishery is through the standardization of commercial logbook catch records
97 aggregated to a coarse statistical grid, at best a 1 degree spatial grid, using a two-step delta-GLM
98 (Lo et al., 1992; Stefansson, 1996). A delta-GLM is the product of two GLMs: a logistic model
99 that describes the presence-absence of positive catches and an additional model (with normally
100 distributed error structure in this case) that describes the magnitude of $\log(\text{CPUE})$ for catches
101 greater than 0. This paper evaluates two methods of creating abundance indices as applied in a
102 vertical line fishery for reef fish, and more generally in fisheries able to achieve a high level of
103 spatial targeting of non-transient species.

104 We conducted analyses to compare abundance indices derived from the same input catch
105 data using two methods: the delta-GLM standardization (status-quo) and spatial averaging of
106 VMS derived CPUE distributions. The first analysis evaluated the agreement between indices
107 generated from the two methods utilizing as input commercial logbook catch records from a suite
108 of reef fish stocks that make up a large proportion of the catch by the vertical line fleet in the
109 Gulf of Mexico. Agreement was assessed in two ways: (i) by calculating the correlation between
110 the indices from the two methods, and (ii) by calculating the change in abundance inferred by
111 each method. Instances of poor agreement between the two methods provided motivation for
112 determining which method more accurately tracked abundance.

113 A simulation analysis was used to assess how well each method captured the true population
114 abundance trend under different effort and abundance scenarios. Corresponding catch and VMS
115 records were simulated and passed as input to the two methods to create abundance indices. The
116 deviations of the indices from the true trend were calculated to determine which method was
117 more accurate under the various scenarios. A principal component analysis (PCA) identified
118 characteristics of scenarios where there were large disparities in the accuracy of the two
119 methods. Previous simulation studies investigated the effects of spatial aggregation, changing
120 distribution of effort, and imputing unfished spatiotemporal strata on indices for pelagic fisheries
121 standardized with GLMs (Campbell, 2004, 2015; Carruthers et al., 2010, 2011; Lynch et al.,
122 2012). Other have studies investigated how geostatistical averaging of VMS-informed catch rates
123 compared to a fisheries-independent measure of abundance in a scallop fishery (Walter et al.
124 2014a, b). This work represents the first direct comparison of abundance indices derived from
125 delta-GLM standardization and spatial averaging of VMS derived CPUE distributions.

126 **2. Material and Methods**

127 This study aimed to address the potential fine-scale spatial targeting problem in conventional
128 CPUE standardization by evaluating the use of VMS data for estimating population trends.

129 Multiple analyses, conducted in R 3.3.2 (R Core Team, 2016), were used to compare the delta-
130 GLM and VMS methods. An overview of the fishery and the species included in the study can be
131 found in section 2.1 and a description of the two data sources informing each method can be
132 found in section 2.2. The first step was to use the same fisheries data to estimate abundance
133 indices using the two methods for every study species. Detail on how abundance indices were
134 constructed for each method can be found in section 2.3. The next step was to assess the
135 agreement in species abundance indices estimated using the two methods. This was done using a
136 non-parametric approach described in section 2.4. Calculating the agreement between indices
137 constructed using the same catch data, but with different methodologies allowed us to identify if
138 there were noticeable differences between the abundance indices created.

139 A simulation study was used to evaluate which method was more accurate in estimating
140 abundance under a suite of scenarios governing how effort and abundance were distributed
141 spatially. The base simulation described in section 2.5.1 was designed to simulate fine scale
142 targeting in a multi-species fishery on a 1/12th degree spatial grid. Section 2.5.2 describes how
143 the base simulation was modified for each scenario. In each scenario, abundance indices for each
144 species were calculated using the two methods along with the deviation from the true simulated
145 population trend (described in section 2.5.3). This allowed us to identify how sensitive the
146 accuracy of each method was with respect to changes in broad patterns of effort and abundance.
147 A multivariate analysis (described in section 2.5.4.) was used to identify the effort and
148 abundance characteristics of species-scenario combinations where the two methods predicted
149 diverging abundance trends.

150 The base simulation made the simplifying assumption that sampling by the fishery did not
151 affect abundance, as this feedback was not necessary in the direct comparison of the ability of
152 the two methods to handle fine-scale spatial data. However, making this assumption ignored the
153 potential effects of in-year sequential depletion occurring at scales smaller than the spatial grid
154 used in the simulation. Hyperstability could occur in fisheries targeting small aggregations or
155 reefs within a cell if vessels move from reef to reef fishing down each in turn. A modification to
156 the base simulation (described in Section 2.5.5) was used to explore how sequential depletion at
157 the cell level affected the estimated abundance indices' ability to capture the true abundance
158 trend.

159 *2.1. Study Frame*

160 The study frame for this project was the vertical line reef fish fishery within the Gulf of
161 Mexico EEZ (Fig. 1) during 2007–2013. Vertical line fishing consists of dropping multiple
162 baited hooks on a single line or multiple lines deployed vertically from a stationary or slowly
163 drifting vessel. These lines are predominantly retrieved using mechanical means such as electric
164 or hydraulic reels though they may also be retrieved by hand. Fishing occurs in distinct
165 spatiotemporal sets defined as the period that hooks are being fished from a vessel at that
166 location. Multiple drops of the gear can occur during each fishing set. A change in location or
167 prolonged period with hooks out of the water represents a change to a new fishing set. Species
168 were included in the analysis if they were within the top 25 of catch by weight over the study
169 period (Table 1). Two pelagic species in the top 25 were excluded as they were likely targeted
170 using non-vertical line gear.

171 *2.2. Data*

172 This study used two data sets: VMS-derived spatial CPUE and commercial logbook self-
173 reported catch records (CLB). VMS use was required for all vessels holding a commercial Gulf

174 of Mexico Reef Fish Permit starting in 2007,. Vessel positions are reported every 60 minutes at a
175 resolution of ~0.1 meters. Reported positions were excluded from the analysis if they occurred
176 outside of the study frame, were assumed not to represent fishing activity (<5 km from land), or
177 corresponded to non-vertical line gear. The resulting data set contained 2,769,857 VMS entries
178 spanning the study period (except for July and August 2010; these data were unavailable).

179 To determine whether the vessel positions corresponded to fishing activity, VMS points were
180 classified as fishing or not fishing using a two-step random forest classification algorithm
181 (Ducharme-Barth and Ahrens, 2017). A unit of effort in the fishery was defined to be a VMS
182 point classified as fishing. Spatial distributions were generated at monthly intervals using the
183 GPS information associated with each VMS entry. Effort points were aggregated on a 1/12th
184 degree spatial grid (roughly 10 km x 10 km). The species-specific catch in pounds for each trip
185 in the CLB was uniformly distributed to all effort points associated with that trip. Spatial CPUE
186 by species was defined in each grid cell as the total catch weight across all trips divided by the
187 number of effort points across all trips.

188 A Monte Carlo simulation method was used to propagate classification uncertainty into the
189 spatial distributions by generating 201 CPUE values for each grid cell. The method applied a
190 two-step process that combined variability in the predicted state (fishing or not-fishing) for each
191 VMS entry due to the random forest model and uncertainty in the predicted state accounting for
192 the classification accuracy of the model. Thus, each of the 201 values represent an alternative
193 fishing scenario that can be used to create an individual abundance index. The number of values,
194 201, generated for each cell was selected because variability across scenarios had stabilized
195 when including more than 100 scenarios, and using greater numbers of scenarios became
196 computationally challenging. Ducharme-Barth and Ahrens (2017) provide further detail of the
197 VMS classification process and Monte Carlo simulation methods.

198 The second data source was the CLB records that corresponded to the VMS points. Within
199 the study period, the CLB contained 31,643 unique vertical line fishing trips targeting reef fish.
200 Trips were retained in the analysis if they indicated that a vertical line gear (hand line, hand gear,
201 or hydraulic/electric reel) was used on that trip. A small percentage of retained trips (2%)
202 indicated that multiple gears were used. Logbook variables considered for CPUE standardization
203 included year, month, area fished, days away, number of crew, season, and region. Season was
204 determined from month (1 – Jan, Feb, March; 2 – April, May, June; 3 – July, August, September;
205 4 – October, November, December). The region (Fig. 1) was assigned based on the reported area
206 or statistical zone. Species CPUE by trip was defined as catch in pounds per hook-hours fished.
207 Hook-hours fished is the product of number of lines fished, hooks fished per line, and total hours
208 fished.

209 2.3. Abundance Indices

210 2.3.1. VMS

211 Annual abundance indices were created from VMS-derived spatial CPUE distributions for
212 each of the 201 fishing scenarios using a combination of temporal imputation and spatial
213 averaging (Walters, 2003). Within a fishing scenario, 82 monthly spatial CPUE distributions
214 were computed to span the time series of seven years (minus two missing months). Cells were
215 identified for temporal imputation if they were empty in a month but fished in another month.
216 Empty cells were filled with the average value of that cell from the two previous months. If a cell
217 was empty to begin the study period, but was fished in a later month, all months leading up to the
218 first month fished were filled with the value of the first month fished. Following imputation, cell
219 values were averaged within month to generate a monthly abundance index. For the two missing

220 months of data (July and August 2010), CPUE was imputed as the average of the two adjoining
221 months. The monthly abundance indices were summed within year to create the annual
222 abundance indices. Repeating this process across fishing scenarios resulted in 201 annual
223 abundance indices. This allowed for the calculation of uncertainty as the 95% inter-quantile
224 range around the median for each year in the abundance index. The resulting indices were
225 rescaled to Z scores e.g. mean of zero and standard deviation of one.

226 2.3.2. Delta-GLM (status-quo)

227 The most practical comparison would be between the VMS-derived abundance index and a
228 corresponding commercial vertical line index used in the SouthEast Data, Assessment, and
229 Review (SEDAR) process. The SEDAR process provides assessments for stocks in the southeast
230 United States, including the Gulf of Mexico. Unfortunately, there was not a complete set of
231 indices from the SEDAR process spanning the study period for all species. Additionally,
232 variables used to standardize CPUE tended to vary slightly among different species (Bryan,
233 2013; Bryan and McCarthy, 2015; McCarthy, 2011; Saul, 2013; Smith et al., 2015; Smith and
234 Goethel, 2015). For this study, species-specific indices derived from CLB data were created
235 using a common framework that best approximated the various approaches used in the SEDAR
236 process.

237 Abundance indices were created from CLB records corresponding to trips that likely
238 encountered the target species. These trips were identified using a logistic regression model of
239 multi-species presence-absence data taken from the CLB records (Stephens and MacCall, 2004).
240 Then a delta-GLM (Lo et al., 1992; Stefansson, 1996) was used to standardize the $\log(\text{CPUE})$ of
241 the target species. Explanatory variables were selected for inclusion separately in each of the two
242 delta-GLM sub-models according to Akaike information criterion (AIC), with the candidate
243 variables being year, temporal strata (season or month), region, days away, and crew. All
244 variables were categorical, and days away and crew number were binned (1,2,3,4,5,6,7,8,9,10+
245 and 1,2,3,4,5+ respectively). At minimum, the two sub-models had to contain a year effect, a
246 temporal effect (season or month), and a region effect. Only one temporal effect could be
247 considered in a sub-model at a time. All effects in the model were assumed to be fixed.
248 Interactions between spatial and temporal strata were not considered as there were incomplete
249 observations of strata combinations for some of the species considered. Imputing the catch rate
250 of missing strata was not considered since this technique is not commonly used in the SEDAR
251 process. To ensure that bias did not enter the delta-GLM parameter estimates due to the uneven
252 distribution of observations across spatiotemporal strata in the models (Campbell, 2004), the
253 observations were reweighted such that each spatiotemporal strata received equal weight in the
254 models (Campbell, 2015).

255 The predictions for both sub-models across a table of all possible spatiotemporal strata
256 (Walters, 2003) were multiplied together and back transformed from log space to give a single
257 expected CPUE in each strata. For models where days away and crew were selected, the modal
258 observation for that variable was used in all predictions across spatiotemporal strata (Campbell,
259 2015). Predictions within year were averaged across temporal strata (season or month) and a
260 weighted average across regions was used to generate the annual abundance index (Campbell
261 2015). When averaging across regions, the assigned weights were proportional to the areas of the
262 regions. The standard error for the annual abundance index was constructed from the
263 uncertainties associated with the two sub-models according to the method described in Campbell
264 (2015). Lastly, the indices were rescaled to Z scores.

265 2.4. Abundance index agreement

266 One of the purposes of this study was to assess the agreement between the indices generated
 267 from the two methods, VMS (V) and delta-GLM (C). We assess agreement using two methods: a
 268 standard metric of agreement, correlation, and a metric relevant to fishery managers that
 269 measures whether the two indices imply the same overall change in abundance.

270 Given the autocorrelation in time series data, a conventional calculation of correlation and
 271 significance would not be appropriate. To account for the auto-correlated nature of the data as
 272 well as the uncertainty in each index, we used a non-parametric modification of surrogate data
 273 testing to test if the temporal structure of the indices resulted in a meaningful correlation between
 274 the two methods. Surrogate data testing is a proof by contradiction technique used in time series
 275 analysis to detect non-linearity (Schreiber and Schmitz, 2000; Theiler et al., 1992). Surrogate
 276 data testing works by calculating a given metric for the original time series and comparing it to a
 277 distribution of metrics calculated from many surrogate data sets generated by some null model. If
 278 the metric from the original time series falls outside of the distribution of metrics from the
 279 surrogate data, then the original time series is different from the null model. In our case, because
 280 there is uncertainty around each time series, we compared two distributions to each other rather
 281 than a point estimate to a distribution. This modification is outlined in Fig. 2. For each pair of
 282 indices, V and C (Fig. 2 A), two new indices, v and c , were created (Fig. 2 B):

$$\left(\begin{array}{c} \mu \\ \mu \end{array} \right) \quad (1)$$

$$\left(\begin{array}{c} \mu \\ \mu \end{array} \right) \quad (2)$$

283 where $\mu_{V,t}$ and $\mu_{C,t}$ correspond to the means of V and C at time t , and $\sigma_{V,t}$ and $\sigma_{C,t}$ correspond to
 284 the standard errors of V and C at time t . The new indices, v and c , account for the uncertainty
 285 associated with the abundance indices while maintaining the temporal structure of those indices.
 286 A Pearson's point-wise correlation can then be calculated between each pair of the indices v , c .
 287 Two surrogate indices, v' and c' , can be formed by taking v and c and randomly rearranging their
 288 order (Fig. 2 C). A correlation is then calculated between each pair of the indices v' and c' .
 289 Repeating the process of creating indices v , c , v' , and c' (Fig. 2 B, C) 10,000 times resulted in a
 290 distribution of correlations where the temporal structure was preserved and a surrogate
 291 distribution of correlations where the temporal structure was rearranged (Fig. 2 D). The mode of
 292 the distribution where temporal structure was preserved gives the correlation between the two
 293 indices. Values closer to 1 show a positive correlation between indices and values closer to -1
 294 show a negative correlation between indices.

295 The overlapping coefficient (OVL) is a commonly used metric for assessing the similarity
 296 between two distributions (Inman and Bradley, 1989; Rom and Hwang, 1996) and non-
 297 parametric estimates of OVL are robust to strong assumptions on the shape and variance of the
 298 distributions (Clemons and Bradley, 2000; Stine and Heyse, 2001). An OVL of 0 indicates the
 299 two distributions are completely dissimilar and an OVL of 1 indicates the two distributions are
 300 identical. The OVL, referred to as OVL_{Corr} , between the two distributions indicates the similarity
 301 of the correlations between the two indices accounting for the temporal autocorrelation and error
 302 associated with each index. In the current case, low OVL values indicate that the distributions of
 303 correlation with and without temporal structure are highly dissimilar and that the temporal
 304 structure resulted in a meaningful correlation. High OVL values indicate that a random temporal
 305 structure was just as likely to achieve the same level of correlation between indices.

306 We used the same 10,000 simulated indices, v and c , to assess if both indices inferred the
 307 same change in stock abundance. Inferred change in stock abundance for each index was

308 calculated as the difference between the mean of the first two years of the index and the mean of
 309 the last two years of the index. Each index was already scaled relative to its mean and standard
 310 deviation so this allowed for comparisons of the change in inferred stock abundance between
 311 indices. For each of the 10 000 indices the inferred change in stock abundance was calculated.
 312 This resulted in two distributions, one for the change in stock abundance inferred by the VMS
 313 method and the other for the change inferred by the delta-GLM method. The OVL, referred to as
 314 OVL_{Change} , between these two distributions was calculated. Low values of OVL indicated that the
 315 two distributions were dissimilar and that the two methods, VMS and delta-GLM, inferred
 316 different changes in stock abundance.

317 2.5. Simulation

318 2.5.1. Base simulation

319 To further evaluate the two methods, we designed a spatial simulation test to replicate the
 320 spatiotemporal dynamics of the underlying species abundance distributions and the vertical line
 321 fishery. A simulated fishing fleet was distributed across a multi-species fishery comprised of 15
 322 species. Fishing and species abundance patterns were simulated at an annual scale for 7 years
 323 and across a $1/12^{\text{th}}$ degree spatial grid.

324 The spatial distributions of abundance were simulated to be representative of reef fish species
 325 encountered in the Gulf of Mexico (Table 2). For each species, the base abundance distributions
 326 were smoothed versions of average annual distributions of spatial CPUE from the VMS data.
 327 Each base abundance distribution was rescaled to sum to 3×10^6 so that each species started
 328 the simulation with the same abundance. An annual abundance trend was applied to each species
 329 using a first order random walk:

(3)

330 where $a_{i,s,t}$ is the abundance of species s in cell i in year t , and ϵ is a normally distributed error
 331 term applied to each cell (ϵ is defined in more detail in Section 2.5.2.). Summing abundance
 332 across cells within years gave the true abundance trend for each species.

333 Fishing trips were simulated to be representative of the characteristics observed in the CLB
 334 and VMS datasets. The total number of trips, $TotalTrips_t$, in any given year t of the simulation
 335 was a random draw from the following distribution.

$$(\mu \quad 740) \quad (4)$$

336 Three variables defined each fishing trip f where μ_f : the number of VMS
 337 points or locations fished on a trip, l_f ; the trip length in days, $DAYS_f$; and the number of
 338 crew, c_f . The parameters used to define the distribution for these variables were estimated
 339 from the VMS and CLB data sets.

$$l(\mu \quad 06) \quad (5)$$

$$on(\lambda \quad 8) \quad (6)$$

$$ormal(\mu \quad 4) \quad (7)$$

340 In any trip, if 0 was drawn for any of these variables it was replaced with 1. Additionally,
 341 μ_f was rounded to the nearest integer value. The spatial distribution of effort was simulated
 342 by selecting an initial fishing location for each fishing trip, and then allowing additional
 343 movements to other cells for the remaining locations in VMS_f . The initial location or cell for a

344 fishing trip was allocated in accordance with a simple gravity model such that near shore cells
 345 with high expected revenues had the greatest chance of being selected.:

$$\sum p \quad (8)$$

$$(d \quad) \quad (9)$$

346 where $l_{f,t}$ was the initial cell for fishing trip f in year t , $d_{i,t}$ was the relative distance from shore
 347 in cell i , $r_{i,t}$ was the relative expected revenue in cell i in year t , and $v_{s,t}$ was the value of species
 348 s in year t . The annual value of species was taken as the average annual price per pound reported
 349 in the NOAA Annual Commercial Landing Statistics (NOAA, 2017). Movement to adjacent
 350 cells within a fishing trip was simulated according to a Queen's Case random walk with a 60%
 351 chance of staying in the same cell at each move. Out of bound cells were either on land or had a
 352 depth beyond 600m as either of these represent unlikely fishing locations for vertical line gear.

353 Each simulated fishing trip recorded the grid cells fished, the region corresponding to the
 354 initial fishing location, and the total catch of each species. The catch at each location, $C_{f,i,t}$, was
 355 a function of abundance, $A_{s,i,t}$, and vessel catchability, $Q_{f,i}$. The vessel catchability was defined
 356 by $Q_{f,i} = \frac{1}{1 + \exp(-\beta d_{i,t})}$ and a spatiotemporally correlated normally distributed random error, $\phi_{i,t}$.

$$(10)$$

$$l(\mu \quad 54) \quad (0.8) \quad (11)$$

357 The parameters used to define $Q_{f,i}$ and $\phi_{i,t}$ were selected so that the simulation produced
 358 realistic catch rates, representative of what the CLB data showed, given the scale of abundance.
 359 Additionally, we assumed that vessels with greater numbers of crew would be able to achieve
 360 higher catch rates because of reduced handling times. The species-specific catch was zero-
 361 inflated to account for occasions where no catches were made at that cell despite fishing effort.
 362 The error term $\phi_{i,t}$ was constructed as a first order random walk of Gaussian random fields
 363 (GRF) using the RandomFields package in R (Schlather et al., 2015):

$$GRF(\mu = 0, \sigma = 0.025, scale = 5) \quad (12)$$

364 2.5.2. Scenarios

365 The simulation applied a full factorial design considering three factors, each with two levels,
 366 resulting in eight scenarios (Table 3). To quantify variability, each scenario was simulated 100
 367 times. The factors considered were species abundance pattern, how effort was distributed, and
 368 changes in spatial targeting. For the first factor, species included in the simulated fishery could
 369 have one of two abundance patterns, global or local. In the global case, the ϵ in Eq. (3) was the
 370 same for each cell. In the local case, ϵ was different for each cell and defined as a first order
 371 random walk of GRFs in the same way as $\phi_{i,t}$ but with $\sigma = 0.25$. This approach simulated a
 372 scenario where there were localized patterns in abundance due to regional patterns in
 373 oceanographic conditions. For the second factor of the simulation, effort was distributed in one
 374 of two ways. In the first case, there were no restrictions on the initial fishing location ($l_{f,t}$). The
 375 second case allocated $l_{f,t}$ to the four main spatial regions in proportion to the observed regional
 376 effort distribution from the fishery. This represented a scenario where vessels were unwilling to
 377 travel very far from their home port. The third factor controlled changes in spatial targeting by
 378 manipulating $r_{i,t}$ in the gravity model. The first case did not force a change in spatial targeting,

379 and the values of $r_{i,t}$ were held constant across years. The second case forced a spatial targeting
 380 change midway through the simulation, by manipulating revenues ($p_{s,t}$) for two of the 15
 381 species. The baseline values for species 7 and 9 were \$3.34/pound and \$2.65/pound,
 382 respectively. However, in this second case, in years 1-3 the value for species 9 was set to
 383 $\$10^7$ /pound and in years 5-7 the value for species 7 was changed to $\$10^7$ /pound. This had the
 384 effect of concentrating effort in the SEGOM region over the first three years of the simulation,
 385 opening the distribution of effort up in the fourth year, and then driving effort to the WGOM
 386 region in the final three years of the simulation. This case demonstrates an instance where the
 387 fishery dramatically changed its spatial targeting behavior due to changes in species desirability
 388 driven by regulatory or socioeconomic factors.

389 2.5.3. Abundance indices

390 Species-specific abundance indices were calculated for each simulation using the methods
 391 described in Section 2.3, albeit with slight changes accounting for simplifying assumptions made
 392 in the simulation. In the VMS method, spatial distributions of species CPUE were constructed at
 393 an annual scale by uniformly allocating $C_{f,s}$ across all cells visited by a specific trip. Temporal
 394 imputation followed the method in Section 2.3.1, but at an annual time step instead of a monthly
 395 time step. Species abundance indices were created by taking the average of each imputed annual
 396 CPUE distribution.

397 The simulation testing approach provided an opportunity to test the effects of spatial strata
 398 size and the inclusion of spatial interactions in the status-quo standardization procedure. Four
 399 delta-GLM formulations were used in each simulation to estimate abundance indices: large strata
 400 and no interactions (delta-GLM I), large strata with interactions (delta-GLM II), small strata and
 401 no interactions (delta-GLM III), and small strata with interactions (delta-GLM IV). The large
 402 strata correspond to the four main regions in the Gulf of Mexico (Fig. 1), and the small strata to
 403 the 10 subdivided regions (Fig. 1). Formulations with interactions allowed for sub-models that
 404 include year and region interactions to be included in the selection of the best model. Each of
 405 these formulations modified the same base delta-GLM. The base delta-GLM standardized
 406 $\log(\text{CPUE})$ as a function of year, region, days away, and crew. CPUE from a given trip was
 407 defined at the set level as $C_{f,s} / \text{VMS}_f$.

408 Species abundance indices were created following Section 2.3.2. Trips from the simulated
 409 logbook that were likely to have targeted a given species were identified using the method of
 410 (Stephens and MacCall, 2004). CPUE from these trips were standardized using the delta-GLMs
 411 described in the previous paragraph. Inclusion of interaction terms in the construction of species
 412 abundance indices followed the suggestions made in Campbell (2015).

413 The ability of each method to capture the true trend was assessed in each simulation and for
 414 each species by calculating the root-mean-square deviation (RMSD) between the estimated
 415 abundance index and the true index. The RMSD between two indices is defined as:

$$\sqrt{\frac{\sum_{t=1}^n (e_s - t_s)^2}{n}} \quad (13)$$

416 All indices, both true and estimated, were scaled relative to their means and standard
 417 deviations, making values of RMSD comparable across species and scenarios.

418 2.5.4. Multivariate analysis

419 We used a principal component analysis (PCA) to identify the characteristics of scenarios of
 420 particular concern where the two methods estimated diverging trends in abundance. PCA is a
 421 multivariate technique that clusters observations in ordination space (McGarigal et al., 2000),
 422 and gives meaning to where observations are positioned relative to each other based on the
 423 principal component axes and the included variables. Principal component axes are orthogonal
 424 compositions of the included variables, with each axis explaining some proportion of the total
 425 variability in the observations. When plotted, observations and variables with positive values for
 426 a given principal component indicate positive correlation with that axis, and conversely negative
 427 values for an axis indicate negative correlation. Nine variables (all scaled relative to their mean
 428 and standard deviation) characterizing the simulations (Table 4) were used in the PCA. The first
 429 two principal components, respectively explaining 37.91% and 15.39% of the total variability,
 430 were retained for this analysis.

431 2.5.5. Sequential depletion simulation

432 We made three modifications to scenario 6 (Table 3) of the base simulation to explore the
 433 potential effects of in-year sequential depletion on the method's ability to estimate the true
 434 abundance trend. We chose the effort and abundance patterns of scenario 6 as our baseline since
 435 it provided a realistic approximation of the fishery without drastic changes in spatial targeting.
 436 The three modifications were 1) within cell abundance ($a_{i,s,t}$) was distributed across reefs, 2)
 437 within cell effort was distributed across reefs, and 3) catches were subtracted from abundance at
 438 that reef within year. The number of fishable reefs in cell i was defined as a random draw from a
 439 Poisson distribution.

$$on(\lambda = 7) \quad (14)$$

440 If the value 0 was drawn for any cell, it was replaced with 1. In the base simulation, fished
 441 cells were visited approximately 4-5 times a year. We simulated the number of reefs per cell with
 442 $\lambda = 7$ to ensure the likelihood of sequential depletion occurring at the cell level. Cell abundance
 443 at the start of a year was randomly allocated across reefs associated with that cell. Effort
 444 characteristics and cells fished within each trip were simulated in the same way as in the base
 445 simulation. For each cell fished on a trip, a reef within that cell was then randomly selected using
 446 a multinomial distribution. The probability of selecting a particular reef within a cell was equal
 447 to the proportion of total cell abundance at that reef. Catch was then defined at the reef level
 448 according to equations 10 and 11, and then subtracted from the available abundance at that reef
 449 in that year. If the catch value generated by equation 11 was greater than the available abundance
 450 at that particular reef, the catch was set equal to the available abundance. Abundance indices
 451 were then calculated in the same way as described in Section 2.5.3 for the VMS and delta-GLM I
 452 methods. When calculating the RMSD, the true abundance was taken as the mean of the starting
 453 and ending abundances for each year.

454 3. Results

455 Using the same catch records, abundance indices (Fig. 3) were estimated for each species
 456 listed in Table 2 using both the VMS and delta-GLM methods. Those that showed the strongest
 457 degree of positive correlation and lowest OVL_{Corr} (Table 5) included two species that were
 458 subject to high levels of directed targeting across a wide expanse of available fishing grounds,
 459 red snapper and gag grouper, as well as two species that are caught in association with them,
 460 gray triggerfish and black grouper, respectively. In general, most species showed some level of

461 positive correlation, with both approaches revealing similar trends, though values of OVL_{Corr}
462 were notably large. The greater the combined uncertainty between the two approaches, the
463 higher the OVL_{Corr} in the relationship even if the mean trajectories appeared to correlate visually,
464 e.g., yellowtail snapper, hogfish, and mutton snapper. In these three cases, the delta-GLM I
465 indices all showed greater uncertainty than the VMS. All three of these species have relatively
466 restricted spatial distributions of catch in an area of the Gulf of Mexico (SEGOM) that is subject
467 to lower levels of fishing effort relative to the other regions. A delta-GLM approach attempting
468 to standardize abundance at the Gulf-wide scale, like that currently used, could estimate higher
469 levels of uncertainty due to fewer observations in spatial strata outside of the geographic core of
470 the species catch distributions.

471 Two species were of particular concern, red porgy and mangrove snapper, as the two
472 methods appeared to estimate inverse trends. This was corroborated by looking at the overall
473 change in stock abundance inferred by each method for these two species as the OVL_{Change} was
474 zero for both. For both of these species the VMS method indicated an overall increase in stock
475 abundance and the delta-GLM indicated an overall decrease. Additionally, there were 10 other
476 species where the OVL_{Change} indicated meaningful differences ($OVL_{\text{Change}} < 0.05$) and/or inferred
477 different patterns of stock abundance. Clearly, these conflicting results were driven by
478 differences in how the data were standardized or how spatial information was handled. However,
479 without knowing the true trend, it was impossible to determine which method provided more
480 accurate estimation. This issue demonstrated the need for our simulation study.

481 The simulation generated 100 sets of catch and effort data across eight scenarios. Using each
482 simulated data set, five abundance indices (VMS and delta-GLMs I-IV) were created for each
483 species within each scenario. Of the 15 species included in the simulation, the results for five of
484 them are shown as representative of the diversity of patterns exhibited across all species. Species
485 1-2 and 7-8 were characterized by broad spatial distributions, while species 9 had a very
486 restricted spatial distribution. Additionally, species 7 and 9 were used in the target switching
487 scenarios with effort switching on or off (respectively).

488 A clear pattern emerged in the simulated abundance indices (Fig. 4). The VMS indices (blue)
489 were consistently able to track the true abundance (black) for each species, across scenarios. Of
490 the three factors manipulated to create the scenarios, abundance pattern and spatial targeting
491 shifts both negatively affected performance of the simulated delta-GLM I indices (red). As
492 expected, changing the abundance pattern from global (scenarios 1-4) to local (scenarios 5-8)
493 had a negative effect on the delta-GLM I performance, since that particular formulation was
494 unable to account for asymmetrical changes in abundance at scales smaller than the considered
495 strata. Introducing a shift in spatial targeting had a subtler effect on the delta-GLM I indices.
496 These indices appeared to be biased high for species in time periods when they were directly or
497 indirectly targeted with greater effort. This effect is most clearly shown in scenarios 3 and 4
498 across all species. Effort targeting increased in the first three years for species 9, in year 4 for
499 species 1 and 2, and in the last 3 years for species 7 and 8. Manipulating the effort distribution
500 by restricting it within certain regions did not appear to alter the ability of either method to
501 distinguish the true trend.

502 Accounting for additional delta-GLM formulations offered improvement but did not change
503 the overall pattern that VMS indices more closely approximated the true trend (Fig. 5). As
504 expected, the formulations using the smaller spatial strata provided an improvement in the delta-
505 GLM indices. Allowing for models with spatial-temporal interaction terms to be included in the
506 model selection process had mixed results. In most cases, including interactions resulted in a best

507 model that either improved or did not meaningfully change the fit, even if inclusion of
508 interaction terms were unwarranted (global abundance scenarios). However, there were cases
509 where the unwarranted inclusion of interaction terms resulted in a diminished ability to estimate
510 the true trend. In scenarios (Fig. 4, scenarios 3-4 for species 9) where the species occupied a
511 restricted spatial range, a spatial shift in targeting occurred, and small spatial strata were used in
512 the delta-GLM; the AIC indicated a mis-specified model as the best performer, which resulted in
513 poor estimation of the true trend.

514 In addition to evaluation of methods, the simulation study was also able to replicate the
515 prediction of inverse trends first observed in the actual data (Fig. 4, Species 8). A multivariate
516 visualization (Fig. 6) showed the particular abundance and effort characteristics associated with
517 this observation. An abundance decline and range contraction occurred simultaneously with a
518 shift in spatial targeting. This resulted in a case where simulated fishing effort became
519 increasingly able to target “hot spots” of abundance even as the stock decreased in range and
520 total abundance. The increased correlation between effort and abundance shown by the
521 increasing trend in Lee’s L supported this. This dynamic was likely what proved problematic in
522 the delta-GLM approaches, as effort was sampling non-randomly within the spatial strata
523 considered, and thus introducing upward-biased catches into the analysis.

524 Accounting for in-year sequential depletion did not appear to make a meaningful difference
525 in the method’s ability to estimate the true population trend. In-year decreases in abundance
526 averaged -49.85 % (std. dev. = 5.94) across all 15 species and 100 sets of data. Comparing the
527 RMSD of the two methods (VMS and delta-GLM I) from scenario 6 to those from the depletion
528 scenario (Fig. 7) did not indicate deteriorations in either method, nor any change in their relative
529 performances.

530 **4. Discussion**

531 This paper shows that in fisheries where non-transient species are easily targeted at fine
532 spatial scales, spatial averaging of high resolution CPUE data provides a robust estimate of
533 abundance trends. Even in simulated cases where there were pronounced shifts in both the spatial
534 distributions of effort and abundance, the VMS indices could more closely track the true
535 abundance pattern relative to the status-quo delta-GLM method. This may allow VMS indices to
536 serve as a bridge across significant perturbing events that may alter the spatial targeting pattern
537 of the fishery provided catchability has remained relatively constant during the transition.
538 Additionally, the pairing of high-resolution spatial data with catch rate information can also lead
539 to the creation of region-specific indices of abundance, which can be used as input in spatial
540 stock assessments (Booth, 2000) and be an important layer (Babcock et al., 2005; St Martin and
541 Hall-Arber, 2008) in the marine spatial planning process (Gilliland and Laffoley, 2008).

542 Inferences on species trends targeted in the vertical line fishery for reef fish in the Gulf of
543 Mexico may be limited due to the unquantified impacts of changing management practices. The
544 emergence of inverse trends in both the actual and simulated data indicates that a spatial shift
545 may have occurred at either the species or fleet level and that the VMS index may more
546 accurately reflect abundance. However, either method would be susceptible to bias if the
547 implementation of an individual fishing quota system (IFQ) on the grouper-tilefish sector of the
548 fishery in 2010 (GMFMC, 2008) resulted in a sudden shift in catchability due to quota
549 consolidation among more efficient vessels (Yandle and Dewees, 2008) or increased rates of
550 discarding so that landings data became uncorrelated with abundance (Turner, 1997). This issue
551 could partially be addressed by crafting abundance indices from a reference fleet of vessels, with
552 assumed constant efficiency, which fished before and after the implementation of the IFQ

553 system. Improving knowledge of discarding behavior through mandatory reporting or increased
554 observer coverage could also explain changes in catchability. In addition to the potential IFQ
555 influences on catchability, the multi-species nature of the reef fish fishery in the Gulf of Mexico
556 could also affect catchability as a result of substructure within the fleet. For example, there exist
557 several sub-fleets within the fishery, including those targeting shallow-water grouper, red
558 snapper, and deep-water species (Scott-Denton et al., 2011). Though all targeted species are
559 susceptible to capture by vertical line gear, subtle differences in gear configuration among sub-
560 fleets could result in differential species-specific catchabilities. If differences in catchabilities are
561 large and sub-fleet distribution is non-random, spatial biases in catch rate could be introduced. A
562 good understanding of vessel membership among sub-fleets would be critical to addressing this
563 potential source of bias as abundance indices could be derived from the spatial CPUE
564 distribution corresponding to each sub-fleet and then averaged together.

565 Though not explicitly accounted for, the VMS indices were robust to the simulated sources
566 of variability in catchability in the form of trip-level uncertainty and regional trends. This is
567 likely a function of how the nominal spatial CPUE distributions used for the creation of those
568 indices were defined. In defining spatial CPUE across all trips at the grid cell level, individual
569 trip or vessel effects were averaged out provided there were a large number of unique samples
570 within that cell. A limited number of trips in a given cell could reintroduce a bias in catch rates
571 due to trip or vessel effects. Imputing values for cells with limited numbers of trips using
572 regression could diminish this source of bias in the spatial averaging process used to create the
573 abundance indices.

574 Targeting species at spatial scales finer than what is modeled has the potential to introduce
575 hyperstability due to sequential-depletion. The simulation used to explore the effects of
576 sequential depletion was not exhaustive and it is possible that hyperstability occurred at the grid
577 cell level, but was masked due to the variability in abundance across cells and/or across years.
578 Future work is needed to further examine the issue of sequential depletion and how aggregation
579 scale affects our ability to observe fine scale processes. The high-resolution nature of VMS data
580 makes it uniquely positioned to address this issue as it allows for aggregation at the same spatial
581 scale that targeting is occurring.

582 Abundance indices derived from using the delta-GLM method were shown to be just as
583 effective provided that the model was correctly specified to match the scale and dynamics of the
584 underlying population. Improperly specifying the delta-GLM through the inclusion of
585 unwarranted interaction terms or the use of inappropriately sized spatial strata led to decreased
586 predictive ability. Earlier studies showed that AIC may select an overly complex model as best
587 from a pool of candidate models (Carruthers et al., 2010; Kadane and Lazar, 2004). This result
588 arose in the simulation in some cases as interaction models were incorrectly selected when there
589 was in fact only a global trend in abundance. In a worst case scenario, specifying a model with
590 inappropriately large strata resulted in an inverse trend being predicted by the delta-GLM.
591 Further simulation of that scenario with smaller strata did improve the mean RMSD, though it
592 still did not achieve the accuracy of the VMS-derived approach.

593 In scenarios where the two methods appeared to be equally effective in tracking the true
594 abundance trend, determined by their overlapping RMSD distributions, there still existed visual
595 differences in predicted trend. Particularly in scenarios where a spatial shift in targeting
596 occurred, slight anomalies were introduced in species trends using the delta-GLM method. This
597 difference between the two approaches could be meaningful in a stock assessment, particularly if
598 it causes the abundance trend to conflict with other data sources. Issues with conflicting data are

generally dealt with by either dropping the offending data source or reweighting it in the model (Maunder and Piner, 2017). Given the importance placed on maintaining a fit to the abundance trend during the data weighting process (Francis 2011, 2017), changing the data weighting to better fit the anomalous time series could have a large impact on the assessment output (Maunder et al., 2017; Punt, 2017).

One of the advantages of the VMS approach is comparative simplicity. The only major decision required is specifying the imputation rule for filling in unfished areas. Though not an overly complicated model structure, a delta-GLM requires a relatively large amount of expert knowledge of the fishery to correctly specify the sub-models. Some of the decisions required include choice of variables used for standardizing CPUE, the number and size of spatial strata, whether to include interaction terms, imputation method for unfished strata combinations, model selection criteria, model error structure, and model effects structure. Additionally, a precursor to the application of a delta-GLM model is to identify trips targeting the focal species using a method such as that of Stephens and MacCall (2004). Currently, there is no general guidance regarding how changing the selected trips affects the estimated abundance index or associated uncertainty. Averaging across a spatial catch rate distribution comprising all available catch records avoids this potential added source of uncertainty.

An extension of the delta-GLM, the spatiotemporal delta-generalized linear mixed model (delta-GLMM) is growing in popularity, though it is limited to regions where commercial logbooks include high resolution spatial data at the individual fishing set or tow level (Thorson and Barnett, 2017; Thorson et al., 2015). These models have shown the ability to accurately track abundance trends in multi-species fisheries where vessel targeting behaviors occur at multiple spatial scales (Thorson et al., 2016), provided the estimation model is correctly specified. Until the data requirements for this approach are met through observer coverage or electronic logbooks, creating indices from VMS-derived spatial CPUE data appears to be a suitable stepping stone from more commonly used delta-GLM approaches. Alternatively, the VMS-derived spatial CPUE could be used as input for the spatiotemporal delta-GLMM models.

This analysis demonstrates the utility of using high resolution CPUE distributions derived from VMS data to generate indices of abundance. The VMS method is comparatively simpler than delta-GLMs, and robust to changes in species and effort distributions. This approach shows much potential to incorporate high resolution spatial information about the fishery, and ultimately to improve stock assessments of non-transient species such as reef fishes in the Gulf of Mexico.

5. Acknowledgements

This research was funded through the NOAA Fisheries - Sea Grant Population and Ecosystem Dynamics Fellowship (grant # NA15OAR4170182) awarded to N.D.D. and through the NOAA Fisheries - RTR program (grant # NA11NMF4550121). The authors thank A. Punt, T. Carruthers as well as an anonymous reviewer for their thoughtful commentary which improved the quality of this manuscript. The authors declare no conflict of interest in this research.

Figure Captions

Figure 1: The Gulf of Mexico EEZ with the spatial regions considered in the analysis. The colored areas denote the four main regions: western Gulf (WGOM), northern Gulf (NGOM), northeastern Gulf (NEGOM), and southeastern Gulf (SEGOM). The lines indicate the 10 subdivided regions for the smaller spatial strata considered.

645

646 Figure 2: Diagram explaining how to calculate the correlation between two indices (A). The
 647 uncertainty of the initial indices is shown by the shaded regions. For each index, a new index is
 648 created by resampling from the uncertainty of the initial index (B). A correlation is calculated
 649 between the two new indices and is shown in green. For each new index in B, an additional index
 650 is formed by rearranging the order (C). A correlation is calculated between the two rearranged
 651 indices and is shown in orange. The process shown in panels B and C is repeated 10 000 times
 652 resulting in the two distributions of correlations (D). The time series correlation of the two initial
 653 indices is given by the mode of the distribution of correlations with order preserved (green). The
 654 overlapping coefficient (OVL_{Corr}) is given by the overlap of the two distributions.

655

656 Figure 3: Indices of abundance with associated uncertainty constructed using the two
 657 methods. The blue corresponds to the VMS index with the median estimate and the 95% inter-
 658 quantile range shown. Red corresponds to the delta-GLM index with the mean and the 95%
 659 confidence intervals shown.

660

661 Figure 4: Simulated abundance indices for five selected species, where each line represents a
 662 different prediction. The black line is the true abundance. Blue corresponds to the VMS-derived
 663 index and red corresponds to the estimate from a delta-GLM I index. The scenario number is
 664 denoted in the top right corner of each panel.

665

666 Figure 5: Violin plots showing the RMSD between predicted and true abundance for five
 667 selected species. The black line inside each violin signifies the 95% inter-quantile range, the
 668 black bar the 50% inter-quantile range, and the white dot the median RMSD. Moving from left to
 669 right within each panel the violins correspond to each method: VMS, delta-GLM I, delta-GLM
 670 II, delta-GLM III, and delta-GLM IV. The scenario number is denoted in the top right corner of
 671 each panel.

672

673 Figure 6: Principal components biplot for six of the nine variables used in the analysis. The
 674 lines represent the different variables, and the colored dots represent each species-method-
 675 scenario combination. For the three trend variables, blue is decreasing, red is increasing, and
 676 yellow is stationary. For the abundance pattern blue signifies global trends and red signifies local
 677 trends. For the targeting pattern blue indicates no switch in spatial targeting and red indicates a
 678 switch in spatial targeting. For the remaining variable, blue shows a low RMSD and red shows a
 679 high RMSD. The large colored dots highlight scenarios 7 and 8 for species 8.

680

681 Figure 7: Violin plots showing the RMSD from Scenario 6 for two methods: VMS (blue) and
 682 delta-GLM I (red). The pair on the left are without simulated sequential depletion, and the pair
 683 on the right (shaded region) are with simulated sequential depletion.

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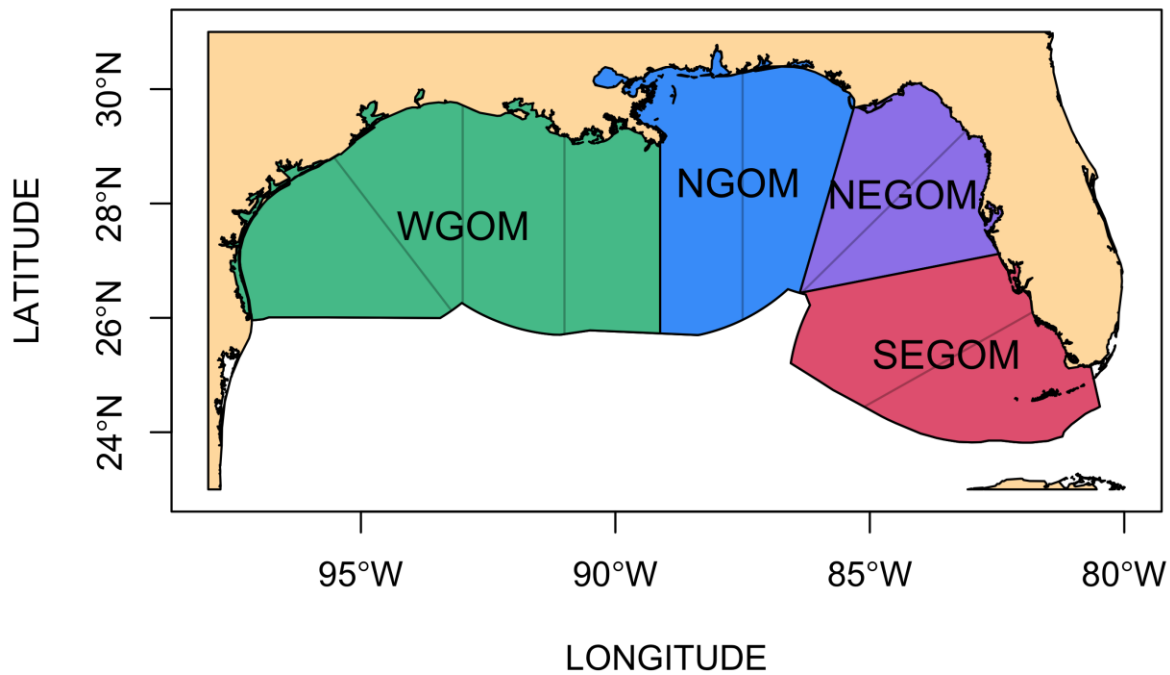
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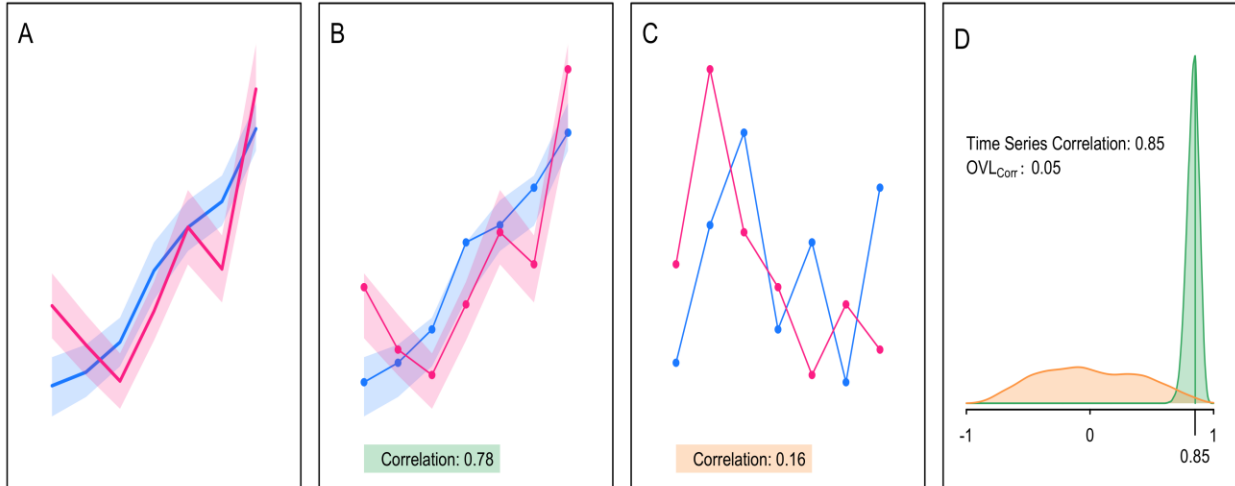
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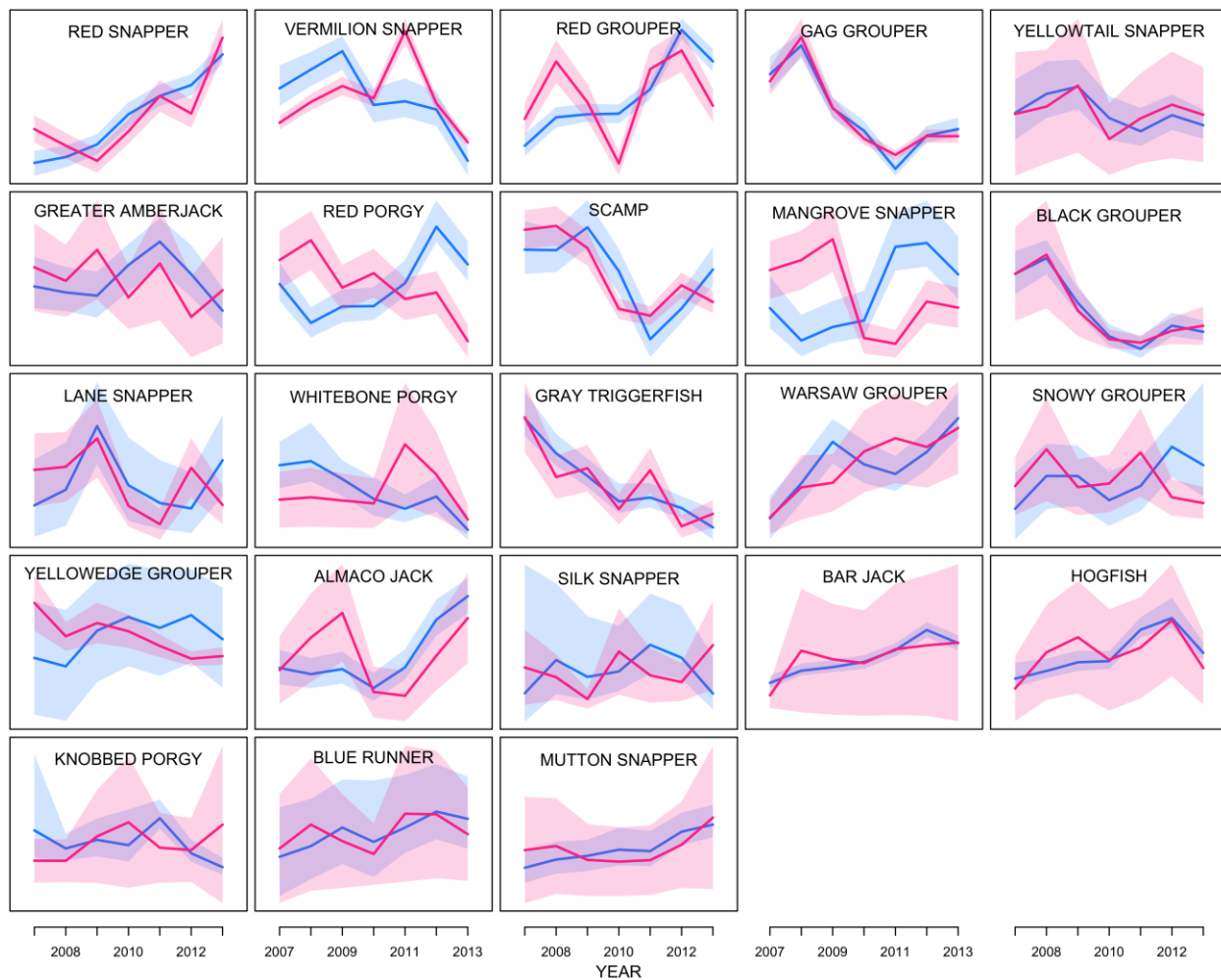
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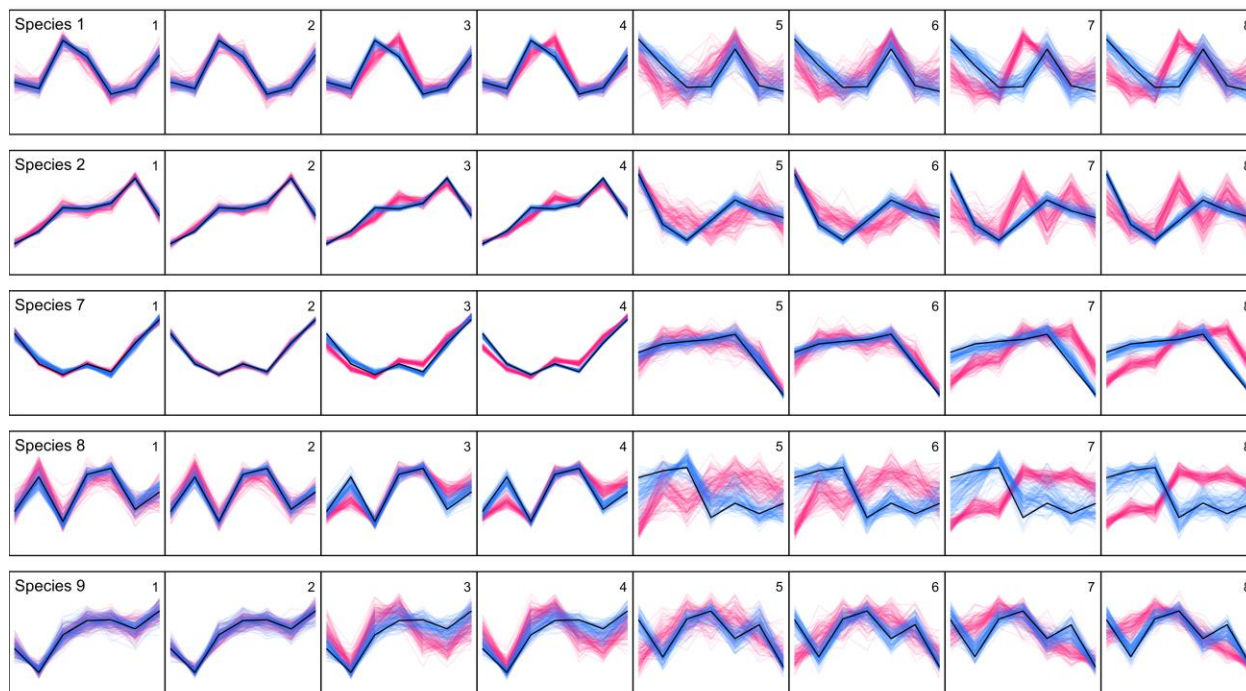
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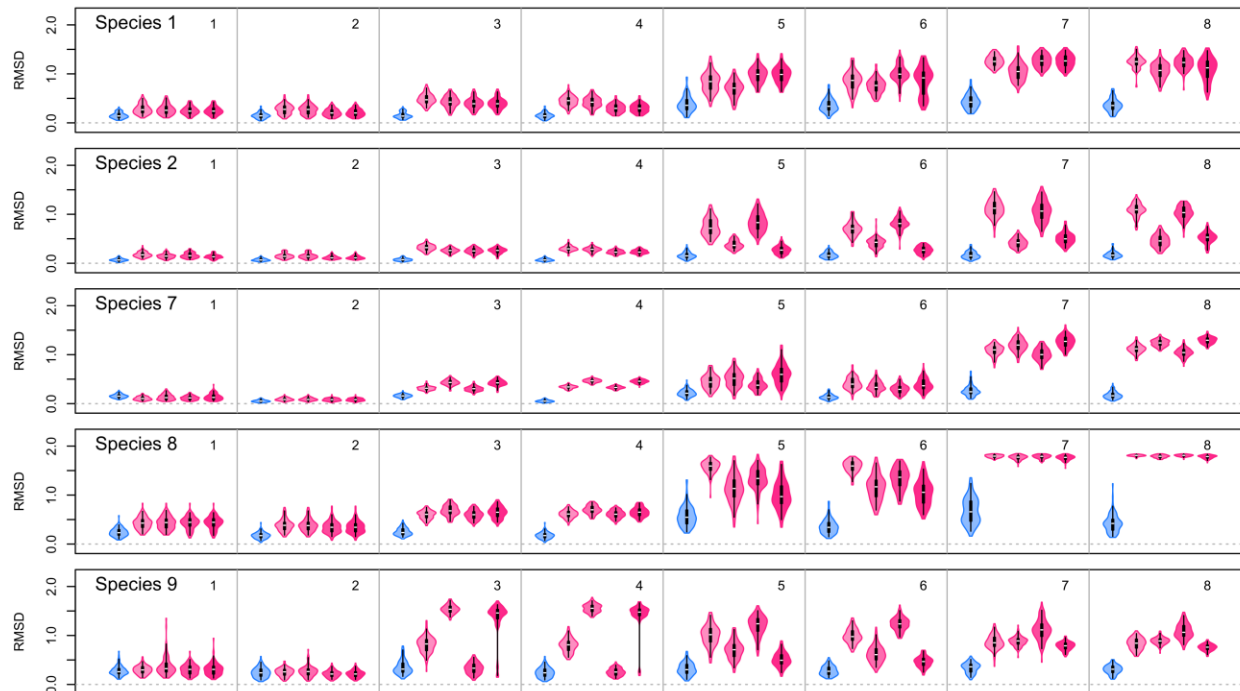
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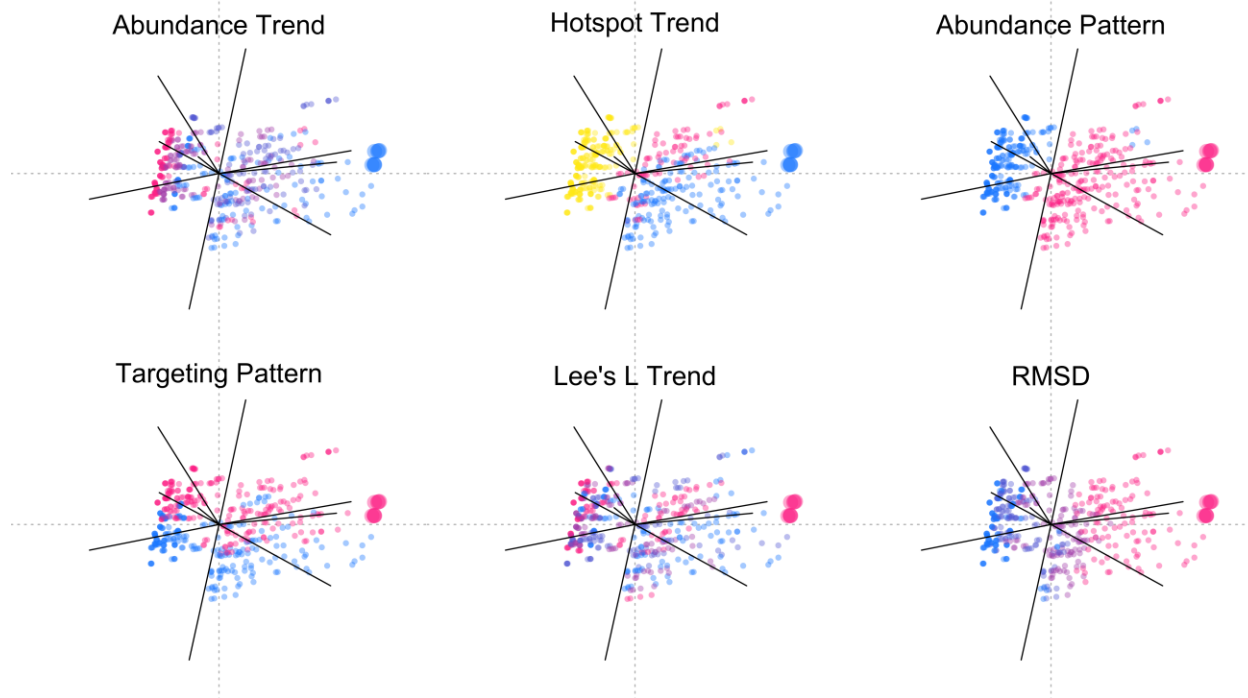
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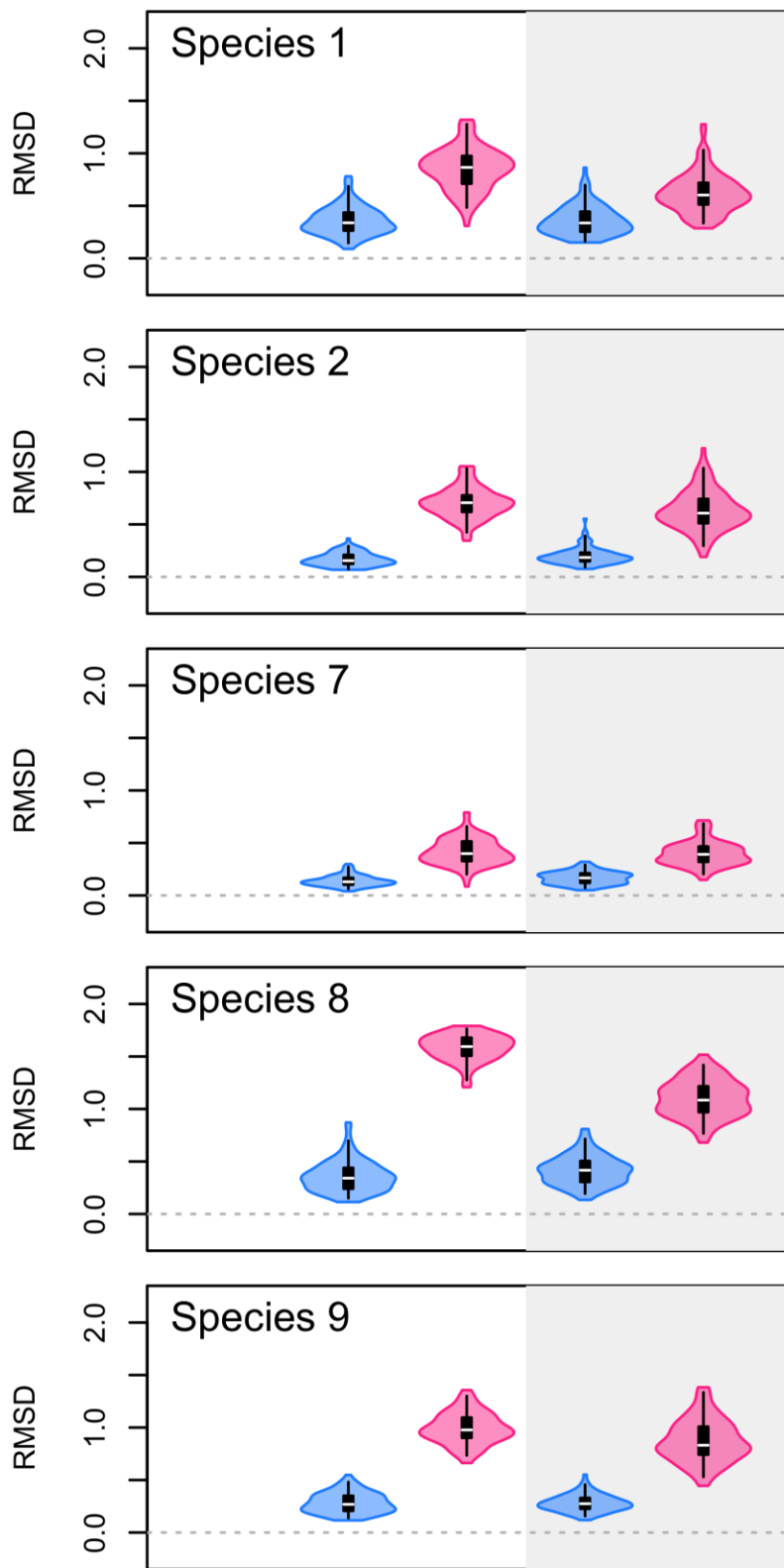
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959 *Table 1* Species occurring in the top 25 of catch by the vertical line fleet. The * indicates species
 960 in the Gulf of Mexico Reef Fish Management Plan, and the # indicates species removed from the
 961 analysis.

Scientific Name	Common Name
Balistes caprisus	Gray Triggerfish *
Calamus leucosteus	Whitebone Porgy
Calamus nodosus	Knobbed Porgy
Caranx crysos	Blue Runner
Caranx ruber	Bar Jack
Epinephelus flavolimbatus	Yellowedge Grouper *
Epinephelus morio	Red Grouper *
Epinephelus nigritus	Warsaw Grouper *
Epinephelus niveatus	Snowy Grouper *
Lachnolaimus maximus	Hogfish *
Lutjanus analis	Mutton Snapper *
Lutjanus campechanus	Red Snapper *
Lutjanus griseus	Mangrove Snapper *
Lutjanus synagris	Lane Snapper *
Lutjanus vivanus	Silk Snapper *
Mycteroperca bonaci	Black Grouper *
Mycteroperca microlepis	Gag Grouper *
Mycteroperca phenax	Scamp *
Ocyurus chrysurus	Yellowtail Snapper*
Pagrus pagrus	Red Porgy
Rachycentron canadum	Cobia #
Rhomboplites aurorubens	Vermilion Snapper *
Scomberomorus cavalla	King Mackerel #
Seriola dumerili	Greater Amberjack *
Seriola rivoliana	Almaco Jack *

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964 *Table 2* The 15 species used to inform the simulation and their approximate geographic
 965 distribution denoted as proportion of abundance in each region.

Number	Species	WGOM	NGOM	NEGOM	SEGOM
1	Red Grouper	0.00	0.03	0.52	0.45
2	Gag Grouper	0.07	0.12	0.60	0.21
3	Black Grouper	0.20	0.07	0.31	0.43
4	Warsaw Grouper	0.85	0.06	0.05	0.04
5	Snowy Grouper	0.26	0.21	0.18	0.35
6	Yellowedge Grouper	0.58	0.10	0.11	0.21
7	Red Snapper	0.83	0.14	0.02	0.01
8	Vermilion Snapper	0.65	0.29	0.05	0.01
9	Yellowtail Snapper	0.02	0.00	0.00	0.98
10	Mangrove Snapper	0.35	0.12	0.26	0.27
11	Mutton Snapper	0.00	0.01	0.00	0.97
12	Red Porgy	0.15	0.48	0.28	0.09
13	Gray Triggerfish	0.54	0.27	0.10	0.09
14	Whitebone Porgy	0.06	0.75	0.17	0.03
15	Hogfish	0.00	0.03	0.82	0.15

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967 *Table 3* Description of each scenario used in the simulation.

Scenario	Abundance Pattern	Effort Distribution	Targeting Shift
1	Global	Restricted	No
2	Global	Unrestricted	No
3	Global	Restricted	Yes
4	Global	Unrestricted	Yes
5	Local	Restricted	No
6	Local	Unrestricted	No
7	Local	Restricted	Yes
8	Local	Unrestricted	Yes

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971 *Table 4* Description of variables used in PCA

Variable	Description
1 - Species abundance trend	The slope of the true trend in abundance.
2 - Species hotspot trend	The slope of the trend in the annual number of cells that had abundance values greater than or equal to 2/3 of the maximum abundance value.
3 - Correlation with true trend	The time series correlation between the estimated trend and the true trend as described in section 2.4.
4 - Pseudo-significance of correlation to the true trend	The pseudo-significance of correlation to the true trend as described in section 2.4.
5 - Average annual Lee's L correlation between the distributions of effort and species abundance	Lee's L is a bivariate measure of the spatial correlation between two distributions (Lee 2001). Lee's L is bounded between -1 and 1 with values greater than 0 indicating a positive correlation.
6 - Trend in Lee's L correlation between the distributions of effort and species abundance	The slope of the trend in the annual Lee's L correlation between the distributions of effort and species abundance.
7 - Abundance pattern	A binary variable indicating either a global or local abundance pattern.
8 - Targeting shift	A binary variable indicating the presence or absence of a spatial targeting shift.
9 - Root-mean-square deviation	The RMSD between the estimated and true trend calculated as shown in Eq. 13.

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976 *Table 5* The metrics of agreement, mean correlation and mean inferred change in stock
 977 abundance, and their respective overlapping coefficients (OVLs) between the two estimated
 978 indices of abundance for each species arranged in order (highest to lowest) of proportion of fleet-
 979 wide catch.

Common Name	Correlation	OVL _{Corr}	Change _{VMS}	Change _{GLM}	OVL _{Change}
Red Snapper	0.85	0.05	2.23	1.53	0.04
Vermilion Snapper	0.41	0.34	-1.64	-0.30	0.00
Red Grouper	0.51	0.27	2.19	0.31	0.00
Gag Grouper	0.97	0.01	-1.75	-1.86	0.72
Yellowtail Snapper	0.44	0.73	-1.09	0.04	0.53
Greater Amberjack	-0.03	0.92	-0.13	-1.30	0.43
Red Porgy	-0.55	0.33	1.80	-2.08	0.00
Scamp	0.69	0.17	-1.02	-1.72	0.17
Mangrove Snapper	-0.57	0.28	1.67	-1.00	0.00
Black Grouper	0.95	0.09	-1.85	-1.87	0.60
Lane Snapper	0.33	0.67	0.45	-0.60	0.38
Whitebone Porgy	-0.14	0.69	-2.04	0.06	0.06
Gray Triggerfish	0.86	0.09	-2.22	-1.96	0.63
Warsaw Grouper	0.67	0.32	2.05	2.02	0.78
Snowy Grouper	-0.25	0.75	1.80	-1.55	0.05
Yellowedge Grouper	-0.28	0.70	1.84	-1.93	0.03
Almaco Jack	0.45	0.50	1.90	0.50	0.10
Silk Snapper	-0.32	0.76	-0.17	0.47	0.70
Bar Jack	0.55	0.59	2.23	1.62	0.23
Hogfish	0.58	0.51	1.78	1.18	0.40
Knobbed Porgy	-0.42	0.81	-1.38	1.50	0.20
Blue Runner	0.24	0.81	2.25	0.77	0.62
Mutton Snapper	0.57	0.68	2.30	1.11	0.40

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