MixFishSim: highly resolved spatiotemporal simulations for exploring mixed fishery dynamics

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Abstract

Most fisheries exploit a variety of spatially and temporally heterogeneous fish populations using species-unselective gear that can result in unintended, unwanted catch of low quota or protected species. Reducing these unwanted catches is crucial for biological and economic sustainability of 'mixed fisheries' and implementation of an ecosystem approach to fishing.

If fisheries are to avoid unwanted catch a good understanding of spatiotemporal fishery dynamics is required. However, traditional scientific advice is limited by a lack of highly resolved knowledge of population distributions, population movement and how fishers interact with different fish populations. This reflects the fact that data on fish location at high temporal and spatial resolutions is expensive and difficult to collect. Proxies inferred from either scientific surveys or commercial catch data are often used to model distributions, usually with sparse data at limited spatial and temporal resolution.

To understand how data resolution impacts inference on mixed fisheries interactions, we develop a highly resolved spatiotemporal simulation model incorporating: i) delay-difference population dynamics, ii) population movement

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using Gaussian Random Fields to simulate patchy, heterogeneously distributed

populations, and iii) fishery dynamics for multiple fleet characteristics based on

species targeting under an explore-exploit strategy via a mix of correlated ran-

dom walk movement (for exploration) and learned behaviour (for exploitation)

phases of the fisheries.

We simulate 50 years of fishing and use the results from the fisheries catch

to draw inference on the underlying population structures. We compare this

inference to a simulated fixed-site sampling design commonly used for fisheries

monitoring purposes and the true underlying population structures. We i) use

the results to establish the potential and limitations of fishery-dependent data

in providing a robust picture of spatiotemporal distributions; and ii) simulate an

area closure based on areas defined from the different data sources at a range

of temporal and spatial resolutions and assess their effectiveness on reducing

catches of a fish population.

We conclude from our simulations that commercial data, while containing

bias, provides a useful tool for managing catches in mixed fisheries if applied at

the correct spatiotemporal scale.

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1. Introduction

Fishers exploit a variety of fish populations that are heterogeneously dis-

tributed in space and time, with varying knowledge of species distributions and

using species non-selective fishing gear. In doing so fisheries catch an assemblage

of species and may discard over-quota catch when managed by single species

quotas and fishers exhaust one or more quota. This may lead to overexploita-

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tion of fish populations (Ulrich et al., 2011; Batsleer et al., 2015). Discarding of fish in excess of quota limits the ability to maintain fishing mortality within sustainable limits (Alverson et al., 1994; Crowder et al., 1998; Rijnsdorp et al., 2007) and the ability to manage for the biological and economic sustainability of fisheries. As such, there is increasing interest in technical solutions such as gear and spatial closures as measures to reduce unwanted catch (Kennelly and Broadhurst, 2002; Catchpole and Revill, 2008; Bellido et al., 2011; Cosgrove et al., 2019).

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Changes to spatial fishing patterns have been proposed as a method to reduce discards (Holmes et al., 2011; Little et al., 2014; Dunn et al., 2014). Implementation of avoidance measures is, however, restricted by lack of knowledge of fish and fishery spatiotemporal dynamics and understanding of the scale at which processes become important for management. Understanding the correct scale for spatial measures is crucial for implementation at a resolution that ensures effective management (Dunn et al., 2016) while minimising economic impact. For example, a scale that promotes species avoidance for vulnerable or low quota species while allowing continuance of sustainable fisheries for available quota species.

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Identifying an appropriate scale has been a challenge in the past that has
led to ineffectual measures with unintended consequences such as limited impact
towards the management objective or increased benthic impact on previously
unexploited areas (e.g. the cod closure in the North Sea (Rijnsdorp et al., 2001;
Dinmore et al., 2003)). More refined spatial information has since become available through the combination of logbook and Vessel Monitoring System (VMS)
data (Lee et al., 2010; Bastardie et al., 2010; Gerritsen et al., 2012; Mateo et al.,
2016) and more real-time spatial management has been possible (e.g. Holmes
et al., 2011). Such information is, however, derived from an inherently biased
sampling programme, targeted fishing, where fishers establish favoured fishing
grounds through an explore-exploit strategy (Bailey et al., 2018) where they

search for areas with high catches and then use experience to return to areas
where they've experienced high catch in the past.

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- We ask two fundamental questions regarding spatiotemporal inference derived from observational data:
- 1. How does sampling-derived data reflects the underlying population structures?
- 2. How does data aggregation and source impact on spatial fisheries management measures?
- To answer these questions we i) develop a simulation model where population dynamics are highly-resolved in space and time. Being known directly rather 48 than inferred from sampling or commercial catch, we can use the population model to validate how inference from fisheries-dependent and fisheries indepen-50 dent sampling relates to the real population structure in a way we could not with real data. We ii) compare, at different spatial and temporal aggregations, the 'real population' distributions to samples from fisheries-dependent and fish-53 eries independent catches to test if these are a true reflection of the relative density of the populations. We then iii) simulate a fishery closure to protect a species based on different spatial and temporal data aggregations. We use these evaluations to draw inference on the utility of commercial data in supporting management decisions. 58

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[We find..]

2. Materials and Methods

A simulation model that is modular and discrete-event based was developed.

This approach enables efficient computation by allowing for sub-modules implemented on time-scales appropriate to capture the characteristic of the different processes (Figure 1). The following sub-modules were included to capture the

full system: 1) Population dynamics, 2) Recruitment dynamics, 3) Population
 movement, 4) fishery dynamics.

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Population dynamics (fishing and natural mortality, growth) operate on a daily time-step, while population movement occurs on a weekly time-step. Recruitment takes place periodically each year for a set time duration specified for each population, while the fishing module operates on a tow-by-tow basis (i.e. multiple events a day).

In the model system population movement is driven by random (diffusive) and directed (advective) processes and we incorporate characterisation of a number of different fishing fleet dynamics exploiting four fish populations with different spatial and population demographics. The following describes the implementation of each of the sub-modules.



Figure 1: Schematic overview of the simulation model. Blue boxes indicate fleet dynamics processes, the green boxes population dynamics processes while the white boxes are the time steps at which processes occur; t = tow, tmax is the total number of tows; (Rec), (Pop Movement), (Pop Dynamics) logic gates for recruitment periods, population movement and population dynamics for each of the populations, (Past Knowledge) a switch whether to use a random (exploratory) or past knowledge (exploitation) fishing strategy.

2.1. Population dynamics

The basic population level processes are simulated using a modified two-stage Deriso-Schnute delay difference model (Deriso, 1980; Schnute, 1985; Dichmont et al., 2003) occurring at a daily time-step. A daily time-step was chosen to discretise continuous population processes on a biologically relevant and computationally tractable timescale. Under the population dynamics module population biomass growth and depletion for pre-recruits and recruited fish are modelled separately as a function of previous recruited biomass, intrinsic population growth and recruitment functionally linked to the adult population size. Biomass for each cell c is incremented each day d as follows (the full parameter list is detailed in Table 1):

$$B_{c,d+1} = (1+\rho)B_{c,d} \cdot e^{-Z_{c,d}} - \rho \cdot e^{-Z_{c,d}} \times (B_{c,d-1} \cdot e^{-Z_{c,d-1}} + Wt_{R-1} \cdot \alpha_{d-1} \cdot R_{\tilde{y}(c,y,d-1)}) + Wt_{R} \cdot \alpha_{d} \cdot R_{\tilde{y}(c,y,d)}$$
(1)

where ρ is Brody's coefficient, shown to be equal to e^{-K} when K is the growth rate from a von Bertalanffy logistic growth model (Schnute, 1985). Wt_{R-1} is the average weight of fish prior to recruitment, while Wt_R is the average recruited weight. α_d represents the proportion of fish recruited during that day for the year, while $R_{c,\tilde{y}}$ is the annual recruits in cell c for year y.

Mortality $Z_{c,d}$ can be decomposed to natural mortality, $M_{c,d}$, and fishing mortality, $F_{c,d}$, where both $M_{c,d}$ and $F_{c,d}$ are instantaneous rates with $M_{c,d}$

mortality, $F_{c,d}$, where both $M_{c,d}$ and $F_{c,d}$ are instantaneous rates with $M_{c,d}$ fixed and $F_{c,d}$ calculated by solving the Baranov catch equation (Hilborn and

Walters, 1992) for $F_{c,d}$:

$$C_{c,d} = \frac{F_{c,d}}{F_{c,d} + M_{c,d}} * (1 - e^{-(F_{c,d} + M_{c,d})}) * B_{c,d}$$
 (2)

where $C_{c,d}$ is the summed catch from the fishing model across all fleets and vessels in cell c for the population during the day d, and $B_{c,d}$ the daily biomass

for the population in the cell. Here, catch and fishing mortality are the sum of those across all fleets and vessels, where $F_{fl,v,c,d,p} = E_{fl,v,c,d} \cdot Q_{fl,p} \cdot B_{c,d,p}$ with fl, v and p the fleet, vessel and population respectively and E and Q fishing effort and catchability.

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2.2. Recruitment dynamics

Recruitment is modelled through a function relating the adult biomass to recruits at time of recruitment. In *MixFishSim*, it can be modelled either either as a stochastic Beverton-Holt stock-recruit form (Beverton and Holt, 1957):

$$\bar{R}_{c,d} = \frac{(\alpha * S_{c,d})}{(\beta + S_{c,d})}$$

$$R_{c,d} \sim \log N[(\log(\bar{R}_{c,d}), \sigma^2)]$$
(3)

Where α is the maximum recruitment rate, β the spawning stock biomass (SSB) required to produce half the maximum stock size, S current stock size and σ^2 the variability in the recruitment due to stochastic processes, or a stochastic Ricker form (Ricker, 1954):

$$\bar{R}_{c,d} = B_{c,d} * e^{(\alpha - \beta * B_{c,d})}$$

$$R_{c,d} \sim \log N[(\log(\bar{R}_{c,d}), \log(\sigma^2))]$$
(4)

where α is the maximum productivity per spawner and β the density dependent reduction in productivity as the SSB increases. In our example application the Beverton-Holt form of stock recruit relationship was used for all populations though either functional form can be chosen.

119 2.3. Population movement dynamics

To simulate fish population distribution in space and time a Gaussian spatial process was employed to model habitat suitability for each of the populations on a 2d grid.

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We first defined a Gaussian random field process, $\{S(c):c\in\mathbb{R}^2\}$, where for any set of cells c_1,\ldots,c_n , the joint distribution of $S=\{S(c1),\ldots S(c_n)\}$

is multivariate Gaussian with a *Matérn* covariance structure, where the correlation strength weakens with distance. This enables us to model the spatial autocorrelation observed in animal populations where density is more similar in nearby locations (Tobler, 1970; F. Dormann et al., 2007) and we change the parameters to implement different spatial structures for the populations.

The habitat for each of the populations was generated with the *RFSimulate* function of the *RandomFields* R package (Schlater et al., 2015), which simulates a Gaussian Random Field process given a user defined error model and correlation structure. We define a stationary habitat field and combine with a temporally dynamic thermal tolerance field to imitate two key drivers of population dynamics. Each population was initialised at a single location, and subsequently moveed according to a probabilistic distribution based on habitat suitability (represented by the normalised values from the GRFs), temperature and distance from current cell:

$$Pr(J|I) = \frac{e^{-\lambda * d_{IJ}} \cdot (Hab_{J,p}^2 \cdot Tol_{J,p,wk})}{\sum\limits_{c=1}^{C} e^{-\lambda * d} \cdot (Hab_{c,p}^2 \cdot Tol_{c,p,wk})}$$
(5)

Where d_{IJ} is the euclidean distance between cell I and cell J, λ is a given rate of decay, $Hab_{J,p}^2$ is the squared index of habitat suitability for cell J and population p, with $Tol_{J,p,wk}$ the temperature tolerance for cell J by population p in week wk (see below).

During pre-defined weeks of the year the habitat quality is modified with user-defined spawning habitat locations, resulting in each population having concentrated areas where spawning takes place. In the simulations the populations move towards these cells in the weeks prior to spawning, resulting in directional movement towards the spawning grounds.

An advection-diffusion process controls population movement, with a timevarying temperature covariate used to change the interaction between time and suitable habitat on a weekly time-step. Each population p was assigned a thermal tolerance with mean, μ_p and variance, σ_p^2 so that each cell and population temperature suitability is defined that:

$$Tol_{c,p,wk} = \frac{1}{\sqrt{(2\pi \cdot \sigma_p^2)}} \cdot \exp\left(-\frac{(T_{c,wk} - \mu_p)^2}{2 \cdot \sigma_p^2}\right)$$
 (6)

Where $Tol_{c,p,wk}$ is the tolerance of population p for cell c in week wk, $T_{c,wk}$ is
the temperature in the cell given the week and μ_p and σ_p^2 the mean and standard
deviation of the population temperature tolerance.

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The final process results in a population structure and movement pattern unique to each species, with population movement occurring on a weekly basis.

The decision to model population movement on a weekly timescale was to reflect that fish tend to aggregate in species specific locations which have been observed to last around one to two weeks (Poos and Rijnsdorp, 2007b). Therefore this process approximated the demographic shifts in fish populations throughout a year with seasonal spawning patterns (e.g. Figure S5).

2.4. Fleet dynamics

The fleet dynamics can be broadly categorised into three components; fleet 169 targeting - which determined the fleet catch efficiency and preference towards a particular species; trip-level decisions, which determined the initial location to 171 be fished at the beginning of a trip; and within-trip decisions, determining move-172 ment from one fishing spot to another within a trip. Together, these elements 173 implement an explore-exploit type strategy for individual vessels to maximise their catch from an unknown resource distribution Bailey et al. (2018). The 175 decision to use an individual based model for fishing vessels was taken because 176 fishers are heterogeneous in their location choice behaviour due to different 177 objectives, risk preference and targeting preference (Van Putten et al., 2012). 178 Therefore in the simulations fleet dynamics are the productive of individual experiences rather than pre-defined group dynamics.

2.4.1. Fleet targeting

Each fleet of n vessels was characterised by both a general efficiency, Q_{fl} , and a population specific efficiency, $Q_{fl,p}$. Thus, the product of these parameters $[Q_{fl} \cdot Q_{fl,p}]$ affects the overall catch rates for the fleet and the preferential targeting of one population over another. This, in combination with the parameter choice for the step-function defined below (as well as some randomness from the exploratory fishing process) determined the preference of fishing locations for the fleet. All species prices were kept the same across fleets and seasons.

2.4.2. Trip-level decisions

Several studies (e.g. Hutton et al., 2004; Tidd et al., 2012; Girardin et al., 190 2015) have confirmed past activity and past catch rates are strong predictors of 191 fishing location choice. For this reason, the fleet dynamics sub-model included a 192 learning component, where a vessel's initial fishing location in a trip was based 193 on selecting from previously successful fishing locations. This was achieved by calculating an expected revenue based on the catches from locations fished in 195 the preceding trip as well as the same month periods in previous years and the 196 travel costs from the port to the fishing grounds, and choosing randomly from 197 the top 75 % of fishing events as defined by the expected profit, which has a 198 seasonal component.

200 2.4.3. Within-trip decisions

Fishing locations within a trip are initially determined by a modified ran-201 dom walk process. As the simulation progresses the within-trip decision become 202 gradually more influenced by experience gained from past fishing locations (as 203 per the initial trip-level location choice), moving location choice towards areas 204 of higher perceived profit. A random walk was chosen for the exploratory fishing 205 process as it is the simplest assumption commonly used in ecology to describe 206 207 optimal animal search strategy for exploiting homogeneously distributed prey about which there is uncertain knowledge (Viswanathan et al., 1999). In a random walk, movement is a stochastic process through a series of steps. These 209

steps have a length, and a direction that can either be equal in length or take
some other functional form. The direction of the random walk was also correlated (known as 'persistence') providing some overall directional movement
(Codling et al., 2008).

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We use a Lévy flight which is a particular form of random walk charac-215 terised by a heavy-tailed distribution of step-length. The Lévy flight has re-216 ceived a lot of attention in ecological theory in recent years as having shown to 217 have very similar characteristics as those observed by animals in nature, and 218 being a near optimum searching strategy for predators pursuing patchily dis-219 tributed prey (Viswanathan et al., 1999; Bartumeus et al., 2005; Sims et al., 220 2008). Bertrand et al. (2007) showed that Peruvian anchovy fishermen have a 221 stochastic search pattern similar to that observed with a lévy flight. However, it remains a subject of debate (e.g. see Edwards et al., 2011; Reynolds, 2015), 223 with the contention that search patterns may be more simply characterised as 224 random walks (Sakiyama and Gunji, 2013) with specific patterns related to the 225 characteristics of the prey field (Sims et al., 2012). 226

For our implementation of a random walk directional change is based on a negatively correlated circular distribution where a favourable fishing ground is likely to be "fished back over" by the vessel returning in the direction it came from. The step length (i.e. the distance travelled from the current to the next fishing location) is determined by recent fishing success, measured as the summed value of fish caught (revenue, Rev),

$$Rev = \sum_{p=1}^{P} L_p \cdot Pr_p \tag{7}$$

where L_p is landings of a population p, and Pr_p price of a population. Here, when fishing is successful vessels remain in a similar location and continue to exploit the local fishing grounds. When unsuccessful, they move some distance away from the current fishing location. The movement distance retains some degree of stochasticity, which can be controlled separately, but is determined by the relationship:

$$StepL = e^{log(\beta_1) + log(\beta_2) - (log(\frac{\beta_1}{\beta_3}))} * Rev$$
 (8)

Where β_1 , β_2 and β_3 are parameters determining the shape of the step function in its relation to revenue, so that, a step from (x1,y1) to (x2, y2) is defined by:

$$(x2, y2) = x1 + StepL \cdot \cos(\frac{\pi \cdot Br}{180}),$$

$$y1 + StepL \cdot \sin(\frac{\pi \cdot Br}{180})$$

$$with \quad Br_{t-1} < 180, Br_t = 180 + \sim vm[(0, 360), k]$$

$$Br_{t-1} > 180, Br_t = 180 - \sim vm[(0, 360), k]$$

$$(9)$$

where k the concentration parameter from the von Mises distribution which we correlate with the revenue so that $k = (Rev + 1/RefRev) * max_k$, where max_k is the maximum concentration value, k, and RefRev is parametrised as for β_3 in the step length function. A realised example of the step length and turning angle relationships to revenue can be seen at Figure S15.

2.4.4. Local population depletion

Where several fishing vessels exploit the same fish population competition 248 is known to play an important role in local distribution of fishing effort (Gillis 249 and Peterman, 1998). If several vessels are fishing on the same patch of fish, local depletion and interference competition will affect fishing location choice of the fleet as a whole (Rijnsdorp, 2000; Poos and Rijnsdorp, 2007a). In order 252 to account for this behaviour, the fishing sub-model operates spatially on a 253 daily time-step so that for future days the biomass available to the fishery is 254 reduced in the areas fished. The cumulative effect is to make heavily fished 255 areas less attractive as a future fishing location choice as reduced catch rates will be experienced. 257

2.5. Fisheries independent survey

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A fisheries-independent survey is simulated where fishing on a regular grid begins each year at the same time for a given number of stations (a fixed station survey design). Catches of the populations at each station are recorded but not removed from the population. This provides a fishery independent snapshot of the populations at a regular spatial intervals each year, similar to scientific surveys undertaken by fisheries research agencies.

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2.6. Software

The simulation framework is implemented in the statistical software package R (R Core Team, 2017) and available as an R package from the authors github site (www.github.com/pdolder/MixFishSim).

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3. Parameterisation

3.1. Population models

We parametrised the simulation model for four populations with different 273 demographics; growth rates, natural mortality and recruitment functions (Table 274 4). Habitat preference (Figure S1) and temperature tolerances (Figures S3, 275 S4) were unique to each population resulting in differently weekly distribution 276 patterns (Figures S5-S7). In addition, each of the populations has two defined spawning areas which result in the populations moving towards these areas in 278 pre-defined weeks (Figure S2) with population-specific movement rates (Table 279 4). The individual habitat preferences and thermal tolerances result in different 280 spatial habitat use for each population (Figure S9) and consequently different 281 seasonal exploitation patterns (Fishing mortality in Figure S10).

283 3.2. Fleet parametrisation

The fleets were parametrised to reflect five different characteristic fisheries with unique exploitation dynamics (Table 5). By setting different catchability parameters $(Q_{fl,p})$ we create different targeting preferences between the fleets and hence spatial dynamics. The stochasticity in the random walk process

ensures that within a fleet different vessels have slightly different spatial distributions based on individual experience. The step function was parametrised 289 dynamically within the simulations as the maximum revenue obtainable was not known beforehand. This was implemented so that vessels take smaller steps 291 when fishing at a location that yields landings value in the top 90th percentile 292 of the value experienced in that year so far (as defined per fleet in Table 5). 293

With increasing probability throughout the simulation, fishing locations were 295 chosen based on experience of profitable catches built up in the same month from 296 previous years and from the previous trip. 'Profitable' in this context was de-297 fined as the locations where the top 70 % of expected profit would be found 298 given previous trips revenue and cost of movement to the new fishing location. This probability was based on a logistic sigmoid function with a lower asymptote of 0 and upper asymptote of 0.95, and a growth rate which ensures the upper 301 asymptote (where decisions are mainly based on past knowledge) is reached ap-302 proximately halfway through the simulation. 303

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3.3. Survey settings 305

The survey simulation was set up with a fixed gridded station design with 306 100 stations fished each year, starting on day 92 and ending on day 112 (5 stations per day) with same catchability parameters for all populations ($Q_p =$ 1). This approximates a real world survey design with limited seasonal and 309 spatial coverage. 310

3.4. Example research question 311

To illustrate the capabilities of MixFishSim, we investigate the influence of 312 the temporal and spatial resolution of different data sources on the reduction in 313 catches of a population given spatial closures. To do so, we set up a simulation to run for 50 years based on a 100×100 square grid (undetermined units), with five fleets of 20 vessels each and four fish populations. Fishing takes place four times a day per vessel and five days a week, while population movement is every week.

We allow the simulation to run unrestricted for 30 years, then implement spatial closed areas for the last 20 years of the simulation based on data (either derived from the commercial catches, fisheries-independent survey or the 'real population') used at different spatial and temporal scales.

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The following steps are undertaken to determine closures:

- 1. Extract data source
- 2. Aggregate according to desired spatial and temporal resolution
- 3. Interpolate across entire area at desired resolution using simple kriging using the *interp* function from the R package akima (Akima, 2006).
- 4. Close area covering top 5 % of catch rates
- In total 28 closure scenarios were run which represent combinations of:
- data types: commercial logbook data, survey data and 'real population',
- temporal resolutions: weekly, monthly and yearly closures,
- spatial resolutions: 1 x 1 grid, 5 x 5 grid, 10 x 10 grid and 20 x 20 grid,
- closure basis: highest 5 % of catch rates for the protected species
- Survey closures were on an annual basis only, as this was the most temporally resolved survey data available.

7 4. Results

8 4.1. Simulation dynamics

It can be seen from a single vessels movements during a trip that the vessel exploits four different fishing grounds, three of them multiple times (Figure S11), while across several trips fishing grounds that are further apart are fished (Figure

S12). These different locations relate to areas where the highest revenue were experienced, as shown by Figure S13, where several trips for the vessel overlaid on the revenue field, i.e.

$$\sum_{c=1}^{c} \sum_{s=1}^{s} B_{s,c} \cdot Q_{s,c}$$

Vessels from the same fleet (and therefore targeting preference) exploit similar but slightly different fishing grounds depending on their own personal experience during the explore phase of the fishery (Figure S14), which is the result of the randomness in the correlated random walk step function, with distance moved during the exploitation phase and the direction stochastically related to the revenue experienced on the fishing ground (Figure S15).

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- Jas 4.2. How does sampling-derived data reflect the underlying population structure?
 In order to answer this question we compare different spatial and temporal aggregations of the 'real population' distributions to:
 - a) fisheries-independent data: the inferred population from a fixed-site sampling survey design as commonly used for fisheries monitoring purposes;
 - b) **fisheries-dependent data:** the inferred population from our fleet model which includes fishery-induced sampling dynamics.

Figure 2 shows the aggregated catch composition from each of the data 353 sources over a ten-year period (to average seasonal patterns) at different spa-354 tial resolutions. The finer spatial grid for the real population (top left) and commercial data (top middle) show visually similar patterns, though there are 356 large unsampled areas in the commercial data from a lack of fishing activity 357 (particularly in the lower left part of the sampling domain). The survey data at 358 this spatial resolution displays very sparse information about the spatial distri-359 butions of the populations. The slightly aggregated data on a 5 x 5 grid shows similar patterns and, while losing some of the spatial detail, there remains good 36 consistency between the 'real population' and the commercial data. Survey data 362

starts to pick out some of the similar patterns as the other data sources, but lacks spatiotemporal coverage. The spatial catch information on a 10×10 and 20×20 grid lose a significant amount of information about the spatial resolutions for all data sources, and some differences between the survey, commercial and 'real population' data emerge.

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Figure 3 shows the consequences of different temporal aggregations of the data over a ten-year period, with weekly (top), monthly (middle) and yearly 370 (bottom) catch compositions from across an aggregated 20 x 20 area. By com-371 parison to the 'real population', the monthly aggregation captures the major 372 patterns seen in the weekly data, albeit missing more subtle differences. The 373 yearly data results in a constant catch pattern due to the aggregation process 374 (sometimes known as an aggregation bias). The commercial data on a weekly basis shows some of the same patterns as the 'real population', though the first 376 species (in red) is less well represented and some weeks are missing catches 377 from the area. The monthly data shows some consistency between the 'real 378 population' and commercial data for species 2 - 4, though species 1 remains 379 under-represented. On an annual basis, interestingly the commercial data un-380 der represents the first species (in red) while the survey over represents species 381 1. This is likely due to the biases in commercial sampling, with the fisheries 382 not targeting the areas where species 1 are present, and the biases in the survey 383 sampling from over representation of the spatial distribution.

385 4.3. How does data aggregation and source impact on spatial fisheries manage-386 ment measures?

We implemented a spatial closure using the different data sources and spatial and temporal aggregations as outlined in the protocol in Section 3.4. We used this to assess the efficacy of a closure in reducing fishing mortality on species 3, given availability of data and its use at different resolutions in order to evaluate the trade-offs in data sources.

The trend in fishing mortality for each species show that in most cases the

fishery closure was successful in reducing fishing mortality on the species of interest (species 3; Figure 4), though interestingly the largest reductions in fishing mortality happened immediately after the closures, following which the fisheries "adapted" to the closures and fishing mortality increased again somewhat. The exception to the success was the closures implemented based on the coarsest spa-tial (20 x 20) and temporal resolution (yearly) which were ineffective with all data sources. As expected, closures based on the "known" population distribu-tion were most effective, with differing degrees of success using the commercial data. Fishing mortality rates on the other species changed in different propor-tions, depending on whether the displaced fishing effort moved to areas where the populations were found in greater or lesser density.

A regression tree (using the R package REEMtree (Sela and Simonoff, 2012)) highlights that the factor most contributing to differences in fishing mortality before and after the closure was the population (72 % showing that the closures were effective for population 3), followed by data resolution (21 %), data type (7 %) with the least important factor the timescale (< 1 %). In general the finer the spatial resolution of the data used the greater reduction in fishing mortality for population 3 after the closures (Figure 5). The notable outliers are the commercial data at the coarsest spatial resolution (20×20) at a yearly and weekly timescale, where closures were nearly as effective as the fine-scale resolution. In this case the closures were sufficiently large to protect a core are of the habitat for the population, but this was achieved in a fairly crude manner by closing a large area - including area where the species was not found (Figure S17) which may have consequences in terms of restricting the fishery in a much larger area than necessary.

5. Discussion

Our study evaluates the importance of data scaling and considers poten-421 tial bias introduced through data aggregation when using fisheries data to infer 422 spatiotemporal dynamics of fish populations. Understanding how fishers ex-423 ploit multiple heterogeneously distributed fish populations with different catch limits or conservation status requires detailed understanding of the overlap of 425 resources; this is difficult to achieve using conventional modelling approaches 426 due to species targeting in fisheries resulting in preferential sampling (Martínez-427 Minaya et al., 2018). Often data are aggregated or extrapolated which requires 428 assumptions about the spatial and temporal scale of processes. Our study explores the assumptions behind such aggregation and preferential sampling to 430 identify potential impacts on management advice. With modern management 431 approaches increasingly employing more nuanced spatiotemporal approaches in 432 order to maximise productivity while taking account of both the biological and 433 human processes operating on different time-frames (Dunn et al., 2016), understanding assumptions behind the data used - increasingly a combination of 435 logbook and positional information from vessel monitoring systems - is vital to 436 ensure measures are effective. 437

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5.1. Simulation dynamics

We employ a simulation approach to model each of the population and fishery dynamics in a hypothetical 'mixed fishery', allowing us to i) evaluate the consequences of different aggregation assumptions on our understanding of the spatiotemporal distribution of the underlying fish populations, and ii) evaluate the effectiveness of a spatial closure given those assumptions.

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Our approach is unique in that it captures fine scale population and fishery dynamics and their interaction in a way not usually possible with real data and thus not usually considered in fisheries simulations. While other simulation frameworks seek to model individual vessel dynamics based on inferred dynamics from VMS and logbook records (Bastardie et al., 2010), or as a system to
identify measures to meet particular management goals (Bailey et al., 2018), our
framework allows users to explore the assumptions in modelling observational
data and evaluate the underlying dynamics of such approaches at a fine spatial
and temporal scale. This offers the advantage that larger scale fishery patterns
are emergent properties of the system rather and results can be compared to
those obtained under a statistical modelling framework.

457

Typically, simulation models which treat fish as individuals are focussed on exploring the inter- and intra- specific interactions among fish populations (e.g. Shin et al. (2004))

COMPARE AGAINST OTHER MODELS: Osmose, Atlantis,... do they use rationale actors and economic profit making?

463 5.2. How does sampling-derived data reflect the underlying population structure?

Our results demonstrate the importance of data scale and resolution when using observational data to support management measures. In doing so it highlights depends on scale of process: pop movement etc... Important to consider how fishers interact / adapt to changes with the resource and mgmt.

468

5.3. How does data aggregation and source impact on spatial fisheries management measures?

It seems clear that spatial disaggregation is more important than the temporal disaggregation of the commercial data... WHY

Closure scenarios demonstrate potential tor reduce F - not as high as with real pop, but good. Make link to other studies – read up on these.

475

```
5.4. Future application for framework
       The what next:
477
478
       Real world spatiotemp closures rarely been able to consider these issues / de-
479
    signed with these issues fully in mind - NS cod closures, plaice and trevose box...
481
        Use of commercial data increasing - likely to become more important in
482
    future. Also collaborative approach with industry, e.g. hotspot mapping, spa-
483
    tiotemp advice...
485
        Other potential uses of the model
486
487
        Survey design - random, fixed, stratified - all possible to explore
488
        Assumptions in spatiotemporal models: GAMs, GLMMs, geostatistical mod-
490
    els
491
       Natural mortality estimation
492
        Commercial index standardization methods - different spatial scales, indi-
    vidual vessels, data aggregation – all these can be explored.
495
        Sampling scheme design
496
497
       Testing fleet dynamics models at an aggregated level
       Bigger picture stuff:: LO, increasing desire for more nuanced spatiotemp
500
    mgmt... Wider applicability: birds, wildlife??
501
    5.5. Model assumptions and caveats
    6. Conclusions
        Study shows ....
504
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505

```
This is important because ....

This is important because ....

How we might apply this in future ....
```

510 Abbreviations

Detail any unusual ones used.

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518 Appendices

Table 1: Description of	of variables fo	r nonulation	dynamics sub-modu	le-

Variable	Meaning	Units					
Population dynamics							
Delay-difference model							
$B_{c,d}$	Biomass in cell c and day d	kg					
$Z_{c,d}$	Total mortality in cell c for day d	-					
$R_{c,\tilde{y}}$	Annualy recruited fish in cell	yr^{-1}					
ho	Brody's growth coefficient	yr^{-1}					
Wt_R	Weight of a fully recruited fish	kg					
Wt_{R-1}	Weight of a pre-recruit fish	kg					
α_d	Proportion of annually recruited fish recruited during	-					
	$\mathrm{day}\ d$						
Baranov ce	atch equation						
$C_{c,d}$	Catch from cell c for day d	kg					
$F_{c,d}$	Instantaneous rate of fishing mortality in cell \boldsymbol{c} on	-					
	$\mathrm{day}\ d$						
$M_{c,d}$	Instantaneous rate of natural mortality in cell \boldsymbol{c} on	-					
	$\operatorname{day} d$						
$B_{c,d}$	Biomass in cell c on day d	kg					
	Recruitment dynamics						
$\tilde{R}_{c,d}$	is the recruitment in cell c for day d	d^{-1}					
$B_{c,d}$	is the Biomass in cell c for day d	d^{-1}					
α	the maximum recruitment rate	kg					
β	the biomass required to produce half the maximum	kg					
	rate of recruitment						

Tal	Table 2: Description of variables for population movement sub-module					
Variable	Meaning	Units				
	Population movement dynamics					
Habitat me	odel					
a	b	С				
Thermal to	plerance					
$T_{c,wk}$	Temperature for cell in week	$^{\circ}\mathrm{C}$				
μ_p	Mean of the thermal tolerance for population	$^{\circ}\mathrm{C}$				
σ_p^2	Standard deviation of thermal tolerance for the pop-	$^{\circ}\mathrm{C}$				
	ulation					
Population	movement model					
λ	decay rate for population movement	-				
$Hab_{c,p}^2$	Square of habitat suitability for cell \boldsymbol{c} and population	-				
	p					
$Tol_{c,p,wk}$	Thermal tolerance for population p in cell c at week	-				
	wk					
d_{IJ}	euclidean distance between cell I and cell J	-				

Table 3: Description of variables for fleet dynamics sub-module

Variable	Meaning	Units					
	Short-term fleet dynamics						
Rev	Revenue from fishing tow	€					
L_p	Landings of population p	kg					
Pr_p	Average price of population p	$\in \ \mathrm{kg}^{-1}$					
StepL	Step length for vessel	euclidean					
		distance					
Br	Bearing	degrees					
k	Concentration parameter for Von mises distribution	-					
eta_1	shape parameter for step function	-					
eta_2	shape parameter for step function	-					
β_3	shape parameter for step function	-					

Table 4: Population dynamics and movement parameter setting

Parameter	Pop 1	Pop 2	Pop 3	Pop 4	
Habitat quality					
Matérn ν	1/0.015	1/0.05	1/0.01	1/0.005	
Matérn κ	1	2	1	1	
Anisotropy	1.5, 3, -3, 4	1,2,-1,2	2.5,1,-1,2	0.1,2,-1,0.2	
Spawning areas (bound	40,50,40,50;	50,60,30,40;	30,34,10,20;	50,55,80,85;	
box)	80,90,60,70	80,90,90,90	$60,\!70,\!20,\!30$	30,40,30,40	
Spawning multiplier	10	10	10	10	
Movement λ	0.1	0.1	0.1	0.1	
Population dynamics					
Starting Biomass	1e5	2e5	1e5	1e4	
Beverton-Holt Recruit 'a'	6	27	18	0.3	
Beverton-Holt Recruit 'b'	4	4	11	0.5	
Beverton-Holt Recruit σ^2	0.7	0.6	0.7	0.6	
Recruit week	13-16	12-16	14-16	16-20	
Spawn week	16-18	16-19	16-18	18-20	
K	0.3	0.3	0.3	0.3	
wt	1	1	1	1	
wt_{d-1}	0.1	0.1	0.1	0.1	
M (annual)	0.2	0.1	0.1 0.2		
Movement dynamics					
μ	12	15	17	14	
σ^2	8	9	7	10	

Table	5: Fleet dyn	amics para	meter setti	ng	
Parameter	Fleet	Fleet	Fleet	Fleet	Fleet
	1	2	3	4	5
Targeting preferences	pop	pop	-	pop 4	pop
	2/4	1/3			2/3
Price Pop1	100	100	100	100	100
Price Pop2	200	200	200	200	200
Price Pop3	350	350	350	350	350
Price Pop4	600	600	600	600	600
Q Pop1	0.01	0.02	0.02	0.01	0.01
Q Pop2	0.02	0.01	0.02	0.01	0.03
Q Pop3	0.01	0.02	0.02	0.01	0.02
Q Pop4	0.02	0.01	0.02	0.05	0.01
Exploitation dynamics					
step function β_1	1	2	1	2	3
step function β_2	10	15	8	12	7
step function β_3	Q90	Q90	Q85	Q90	Q80
step function rate	20	30	25	35	20
Past Knowledge	${ m T}$	${ m T}$	${f T}$	${ m T}$	T
Past Year & Month	${ m T}$	${ m T}$	T	T	T
Past Trip	${ m T}$	Τ	Τ	T	T
Threshold	0.7	0.7	0.7	0.7	0.7
Fuel Cost	3	2	5	2	1

Table 6: Fishing mortality effects of the closure scenarios. Results show the fishing mortality before the closure (f_before) and after the closure (f_after) and the percentage change in f (f_change). The results are ordered by most effective scenario first, least effective last.)

scenario	metric	pop	f_before	f_after	f_change	timescale	basis	data_type	resolution
9	F	spp_3	1.08	0.29	-73.47	weekly	high_pop	real_pop	1.00
10	F	${\rm spp_3}$	1.08	0.29	-72.94	monthly	$high_pop$	$real_pop$	1.00
11	F	${\rm spp_3}$	1.08	0.35	-68.04	yearly	$high_pop$	$real_pop$	1.00
45	F	${\rm spp_3}$	1.08	0.58	-46.70	yearly	$high_pop$	commercial	20.00
1	F	spp_3	1.08	0.58	-46.21	weekly	$high_pop$	commercial	1.00
23	F	${\rm spp_3}$	1.08	0.59	-45.27	weekly	high_pop	real_pop	5.00
2	F	${\rm spp} _3$	1.08	0.59	-45.06	monthly	$high_pop$	commercial	1.00
7	F	spp_3	1.08	0.60	-44.48	yearly	$high_pop$	survey	1.00
24	F	${\rm spp_3}$	1.08	0.61	-43.20	monthly	$high_pop$	$real_pop$	5.00
3	F	${\rm spp_3}$	1.08	0.64	-40.82	yearly	$high_pop$	commercial	1.00
25	F	${\rm spp_3}$	1.08	0.65	-39.94	yearly	$high_pop$	$real_pop$	5.00
17	F	${\rm spp_3}$	1.08	0.67	-38.11	yearly	$high_pop$	commercial	5.00
15	F	${\rm spp_3}$	1.08	0.71	-34.38	weekly	$high_pop$	commercial	5.00
43	F	${\rm spp_3}$	1.08	0.71	-34.31	weekly	$high_pop$	commercial	20.00
16	F	spp_3	1.08	0.73	-32.58	monthly	$high_pop$	commercial	5.00
51	F	${\rm spp_3}$	1.08	0.78	-27.92	weekly	$high_pop$	$real_pop$	20.00
37	F	${\rm spp_3}$	1.08	0.78	-27.76	weekly	$high_pop$	$real_pop$	10.00
39	F	${\rm spp_3}$	1.08	0.79	-26.98	yearly	$high_pop$	$real_pop$	10.00
38	F	${\rm spp_3}$	1.08	0.81	-25.47	monthly	$high_pop$	$real_pop$	10.00
21	F	${\rm spp_3}$	1.08	0.81	-25.21	yearly	$high_pop$	survey	5.00
35	F	${\rm spp_3}$	1.08	0.81	-25.05	yearly	$high_pop$	survey	10.00
44	F	spp_3	1.08	0.87	-19.91	monthly	$high_pop$	commercial	20.00
52	F	${\rm spp_3}$	1.08	0.88	-18.39	monthly	$high_pop$	$real_pop$	20.00
30	F	${\rm spp_3}$	1.08	0.96	-11.06	monthly	$high_pop$	commercial	10.00
29	F	spp_3	1.08	0.98	-9.80	weekly	$high_pop$	commercial	10.00
31	\mathbf{F}	spp_3	1.08	1.03	-4.36	yearly	$high_pop$	commercial	10.00
53	\mathbf{F}	${\rm spp} _3$	1.08	1.06	-1.64	yearly	$high_pop$	$real_pop$	20.00
49	F	spp_3	1.08	1.07	-1.01	yearly	$high_pop$	survey	20.00



Figure 2: Data aggregation at different spatial resolutions over a ten year period

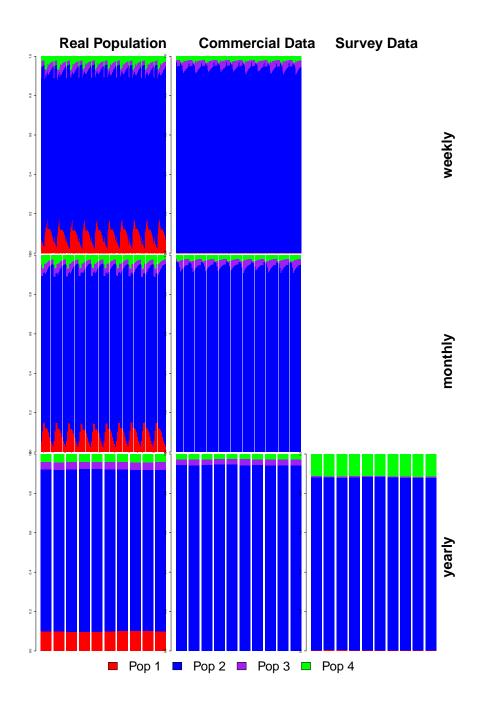


Figure 3: Data aggregation at different temporal resolutions over a ten-year period

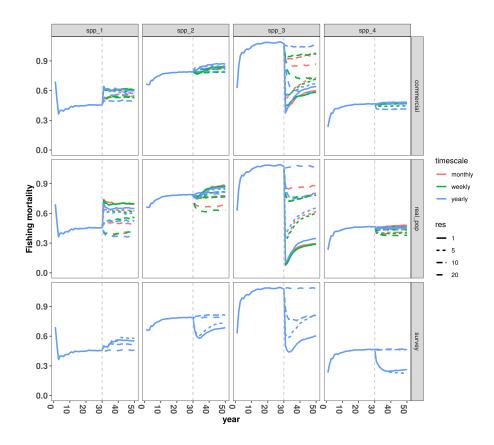


Figure 4: Comparison of closure scenarios effect on fishing mortality trends. Line colour denotes the timescale, while linestyle denotes the spatial resolution.

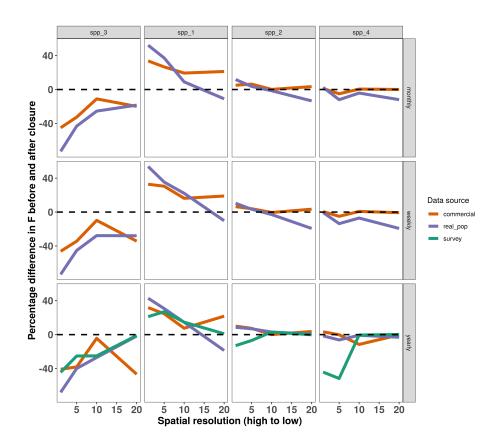


Figure 5: Comparison of closure scenario effectiveness based on different spatial and temporal resolutions.

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