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Claire Kermorvant, Nathalie Caill-Milly, Damien Sous, Iosu Paradinas, Muriel Lissardy, et al.. Detecting the effects of inter-annual and seasonal changes in environmental factors on the striped red mullet population in the Bay of Biscay. Journal of Sea Research (JSR), 2021, 169, pp.102008. 10.1016/j.seares.2021.102008 . hal-03602279

HAL Id: hal-03602279 https://hal.science/hal-03602279

Submitted on 13 Feb 2023 $\,$

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Detecting the effects of inter-annual and seasonal changes in environmental factors on the striped red mullet population in the Bay of Biscay

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Preprint submitted to Fisheries Research

¹ Abstract:

Climate fluctuations affect a wide range of environmental factors that may impact fish populations with 2 various response mechanisms. For a major part of world fisheries stocks, only abundance time-series are 3 available. Relations between abundance of these species and ocean variables, including long term trends 4 and patterns of seasonal change, remain sparsely documented. The main aim of our work is to determine 5 to what extent effects of inter-annual and seasonal changes in environmental factors can be identified and 6 quantified on a selected time-series of abundance data-set: the striped red mullet population within the Bay 7 of Biscay. 8 Landings per unit of effort (LPUE) for striped red mullet data are compared with environmental 8 covariates extracted from the Copernicus web database. Our strategy is to decompose the patterns of change 9 in fish populations into long-term and seasonal components and to analyse the possible dependency of both 10 components on environmental factors. The overall tendencies are a decrease of the long-term component 11 and a modification of the seasonality, with additional spatial variability. The observed tendencies and 12 seasonal changes are discussed against the spatio-temporal patterns of change in environmental factors. 13 High spatial heterogeneity within the Bay of Biscay was found, both for striped red mullet long-term trends 14 and seasonal changes. Most inter-annual and seasonal changes in striped red mullet abundance can be related 15 to environmental factors. In addition to the well-known annual fluctuation, most coastal sites revealed a 16 bi-annual seasonality possibly related to the recruitment process. 17

18 Keywords:

Bay of Biscay; Climate change; Environmental modelling; Fisheries indicators; *Mullus surmuletus*; Time series decomposition.

21 1. Introduction

The fluctuations of ocean variables are a primary driver for fish populations (García-Barón et al., 2020). 22 This environmental control has taken on particular importance in the context of global climate change, which 23 affects numerous oceanic factors that may impact marine fauna (Brander, 2010). Due to the complexity of the 24 mechanisms involved and the interactions between space- and time -varying driving factors, understanding 25 the effect of climate change on these populations may prove to be tricky (Rijnsdorp et al., 2009). The effects 26 of climate change on environmental factors, such as modifications of the water cycle, are expected to differ 27 among geographic areas (IPCC, 2007), both at global and regional scales. For instance, river runoff and 28 precipitation are expected to increase/decrease in northern/southern Europe, respectively (Frei et al., 2006). 29 Likewise, ocean water properties are exposed to global change-induced modifications, such as the significant 30 increase in salinity, which was observed during the three last decades in the northern part of the Atlantic 31 ocean (Stott et al., 2008). 32

Several effects of environmental changes due to global warming on fish populations have already been 33 reported (Both et al., 2006; Free et al., 2019). The review of Rijnsdorp et al. (2009) highlights that these 34 changes may result in four response mechanisms. Firstly, changes in environmental parameters can induce a 35 physiological response (Cohen et al., 2018). Secondly, modifications in environment may induce a behavioural 36 response, such as moving into new suitable areas. The third type of response may be through population 37 dynamics (i.e. changes in the balance between rates of mortality, growth, and reproduction in combination 38 with dispersal) which could result in the establishment of new populations in new areas, in the abandonment 39 of traditional sites or in a regime shift, with significantly higher or lower abundance than in other historical 40 periods. Finally, a fourth response to environmental changes may be changes at ecosystem level (i.e. in 41 productivity and/or trophic interactions). It should be emphasized that, even if this is expected to take on 42 worrying proportions in the context of global change and increased fishing pressure, the characterization of 43 the response of fish populations to the fluctuations of ocean variables is already a major challenge for ocean 44 ecology scientists. 45

From an ecological perspective, the study of the relationships between environmental factors and popula-46 tions is generally carried out using three main categories of models (Melo-Merino et al., 2020): mechanistics 47 models (Kearney and Porter, 2009), process-oriented models (Peterson et al., 2015) and correlative models 48 (Huntley et al., 1995; Araujo and Guisan, 2006; Franklin et al., 2009; Box, 2012). A direct application of 49 such modelling strategies in a marine ecosystem remains very challenging. A major limitation is that most 50 environmental data layers are not know in the entire three-dimensional ocean (Assis et al., 2016). Another 51 difficulty arises from the lack of knowledge on dynamics and behaviour for most species (Bentlage et al., 52 2013). For fisheries stock assessment, the International Council for Exploration of the Sea (ICES) divide 53

fish stocks into 6 categories. Categories 2 to 6 belong to data-limited categories (Lart, 2019). The majority
 of the world's fisheries, by number, are classified in these categories (Costello et al., 2012; Dowling et al.,
 2019). A stock can be considered as data-limited for various reasons, including a lack or a limited amount

⁵⁷ of reliable data, incomplete surveys and/or poor sampling leading to uncertainties about the biology of the ⁵⁸ species (Le Quesne et al., 2013). For ICES categories 5 and 6, abundance is only indirectly assess through

selected proxies, and ecology and behaviours are almost unknown. In such a context of data limitation which
 inhibits comprehensive understanding of fish behaviour, the alternative approach followed here is first to use

⁶¹ a non-causal statistical approach, i.e. to identify and quantify dependencies between the level of interest and ⁶² the available environmental covariates without any *a priori* knowledge of the driving mechanisms, and then ⁶³ to infer *a posteriori* causal relationships by combining statistical results and expert knowledge.

Apart from the data quality issue, the extraction of entangled trends and patterns in time-series data 64 remains a major challenge for data processing science. A widely used approach is to split the time series 65 into several components, possibly selected from an *a priori* knowledge of the driving factors (Hyndman and 66 Athanasopoulos, 2018). A number of decomposition methods are available in literature in relation to the 67 very variable nature of the data-sets, including for instance intermittency, shortness, instationarity, non-68 linear response to driving factors (Huang et al., 1998). Standard data processing books, see e.g. Brockwell 69 et al. (1991), or more recently Shumway and Stoffer (2017), review and detail a large panel of decomposition 70 methods, the latter with examples of application under R software. Under global change effect, the data from 71

the Bay of Biscay, situated at mid-latitude, may be expected to respond to climate factors affected by both 72

global long term trend and annual seasonal fluctuations. A helpful time-series decomposition can be therefore 73

be based on the extraction of long term trend and seasonality from the time-series, with an additional residual 74

(Ferguson et al., 2008). Such a time-series decomposition into trend, seasonal and residual components is 75 still rarely used in fish ecology (Broekhuizen and McKenzie, 1995; Beare and McKenzie, 1999). Among the 76

rather rare attempts, Plaza et al. (2018) used time-series decomposition as a method of data filtering, to 77

identify contrasts between data features and technical fisheries literature, and Hsieh et al. (2009) analysed 78

links between larval anchovy data and environmental variables. More generally, time-series decomposition 79 methods, while widely used in data processing, are not familiar to ecologists (Hsieh et al., 2009).

80

The present study is, to our knowledge, the first assessment of the effect of changes in environmental 81 factors on data-limited fish populations using the time-series decomposition method. The target species 82 is the striped red mullet, Mullus surmuletus Linnaeus, 1758. The striped red mullet is distributed in the 83 Mediterranean Sea and along the Eastern Atlantic coasts, from the English Channel to the northern part 84 85 of West Africa (Labropoulou et al., 1997). At a large scale (Atlantic ocean), northerly migration of striped red mullet during a phase with demonstrated temperature increases due to climate change has already 86 been demonstrated (McCarty et al., 2001; Hulme, 2002; Beare et al., 2005). Striped red mullet abundance 87 increases in the English Channel (Vaz et al., 2004) and the North Sea. Eastern English Channel and the 88 Bay of Biscay are two main areas of catches of striped red mullet in the Atlantic Ocean (Mahe et al., 2014). 89 Individuals in the Bay of Biscay belong to the sub-population individualized in Western Europe by ICES. 90 It extends from the Celtic sea to the waters bordering the Iberian Peninsula with a significant presence in 91 the Bay of Biscay. It is a predominantly benchic and gregarious species living in small groups of up to a 92 dozen individuals. It can also be solitary or live just in pairs. Young fishes live in lower salinity coastal areas 93 while adults have a more offshore distribution (Oskarsson et al., 2019). Pajuelo et al. (1997) find young 94 striped red mullet between 10 and 60 m deep in Canaries Islands while (Reñones et al., 1995) report values 95 between 30 and 90m in Majorcan waters. The striped red mullet has an inter-depth migration because 96 of its reproduction (Kousteni et al., 2019). The recruitment occurs in shallow habitat (García-Rubies and 97 Macpherson, 1995), spawns in deep habitats and continue to disperse into deep water after reproduction 98 (Machias and Labropoulou, 2002). On average, striped red mullet abundance is highest in surface waters 99 down to 100 m depth (easily observable from 0 to 30 m) (Mahé et al., 2005), but large specimens have been 100 observed below 300 m depth. In the north Atlantic, mature striped red mullet depths is between 100 and 200 101 m on average (Leaute et al., 2018). Several factor may intervene in the abundance of the striped red mullet. 102 Benthic habitat composition (Ajemian et al., 2016) or substrate type (Lombarte et al., 2000; Mahé et al., 103 2005) are important. As for all other species, other factors can affect striped red mullet and may have an 104 important impact on its abundance. In particular, population may be affected by trophic web parameters, 105 such availability of prevs and abundance of predators, or factors related to natural mortality such as diseases 106

and parasites. Recent study reports life-history traits of the striped red mullet in Mediterranean (Kousteni 107 et al., 2019). Longevity of striped red mullet is estimated at 11.75 years and the length at 50% of maturity 108 is 153.3 mm for females and 139.2 mm for males. 109

Following the ICES classification categories (ICES, 2012), Bay of Biscay sub-population is currently in 110 data-limited stocks (DLS) category 5: it is a stock for which only landing data are available, as described 111 above. From a fish population perspective, the Bay of Biscay remains a limited area with a total surface 112 aire of approximately 223 000 km². No substantial scientific literature is available to qualify the striped red 113 mullet stock status within the Bay of Biscay and its relationship with ocean variables, whether in the past. 114 at present or in the global change context. 115

The aim of the present paper is to determine whether modifications of oceanic environmental factors, such 116 as changes in long term or seasonal trends, have an impact on the striped red mullet population within the 117 Bay of Biscay. The analysis will be based on selected professional LPUE (Landing Per Unit Effort) as a proxy 118 of striped red mullet stock abundance in the Bay of Biscay, as suggested by Caill-Milly et al. (2019). Time-119 series decomposition was first applied on striped red mullet abundance to achieve a rough understanding of 120 the population dynamics through space and time. Secondly, the analysis was dedicated to the identification 121 and quantification of possible links between trends and seasonal fluctuations of both striped red mullet 122

¹²³ abundance and environmental covariates. The results and methodology were then discussed, keeping in ¹²⁴ mind the challenge of the limited time span of the available time-series on for striped red mullet index of ¹²⁵ abundance.

126 2. Methodology

127 2.1. Database

128 2.1.1. Fishery data

The fishery database used for the present paper (Kermorvant et al., 2020) was provided by the SACROIS 129 (SACROIS) algorithm developed by Ifremer. SACROIS is a cross validation tool for fisheries statistics, 130 in response to Article 145 of the EU regulation (EC Reg. 404/2011). SACROIS algorithm cross-checks 131 information from fishing forms, fishing fleet registers, logbooks, sales notes, VMS data and the scientific 132 census of fishing activity calendars (Mateo et al., 2016). We have selected this database because we considered 133 that it is the most complete available database regarding production and fishing effort data for the French fleet 134 operating in the Bay of Biscay. The SACROIS data are spatially aggregated by ICES rectangles assembled 135 monthly. ICES rectangles extend about 0.5 and 1 degree in longitude and latitude, respectively, with an 136 area of approximately 30 nautical squared miles. The present study was based on a set of six coastal ICES 137 rectangles selected because they were sufficiently documented to allow time and time-frequency analysis 138 (see Figure 1A). These selected rectangles do not integrate the full ecological niche of striped red mullet 139 in the Bay of Biscay, but were each inside the niche and contain sufficient information to be considered 140 as representative of the Bay of Biscay striped red mullet population ICES (2019). LPUE was provided 141 for a given gear, gear mesh, day and ICES statistical rectangle. Specific vessels, i.e. small trawl vessels 142 (7.9–15.8 m) with a gear mesh of 70–79 mm, were selected in the SACROIS database following the criteria 143 of Caill-Milly et al. (2019) for striped red mullet. The time series of the selected mean trawlers LPUE was 144 available from 2005 to 2018. LPUE is not a direct estimator of abundance. However, applying, (i), a filtering 145 on vessel and gear characteristics and, (ii), a selection of a period without significant changes in regulation 146 [see Caill-Milly et al. (2019)], we considered that the selected daily LPUE was a relevant proxy for striped 147 red mullet abundance. In fisheries-dependent data, bias can be introduced to abundance estimates when 148 sampling was not randomized (Potts and Rose, 2018). Striped red mullet is of high commercial interest 149 but is not specifically targeted by trawlers in the Bay of Biscay (it is a part of set of sought-after demersal 150 species). Therefore, even if the sampling scheme was not randomized, it was not directed. Futhermore. 151 the International Council For the Exploration of the Sea (ICES) identified striped red mullet LPUE as a 152 potential abundance index for this stock (Oskarsson et al., 2019; ICES, 2020). Note that, as the present 153 analysis requires continuous time-resolved database, other available sources of data at lower sampling rates 154 (such the EVHOE campaign in the Bay of Biscay and Celtic Sea) were not considered. 155

156 2.1.2. Environmental data

Environmental data was extracted from the Copernicus portal (https://www.copernicus.eu). Values for 157 surface temperature (in degree Celcius), surface salinity (in grams per liter), phytoplankton concentration 158 (in $mmolC/m^3$), chlorophyll concentration (in mg/m^3 , and chemicals (Silicate (Si), Phosphate (PO_4^{3-}), 159 Ammonium (NH_4^+) , Nitrate (NO_{3-}) and Dissolved Oxygen $(mmol/m^3)$ were selected from 2005 to 2018 160 within the Bay of Biscay. For surface temperature and surface salinity, main statistical values (table 1) were 161 computed for each environmental variable by ICES rectangle. For phytoplankton concentration, chlorophyll 162 concentration, and chemicals, a first statistical study showed a high correlation. A principal component 163 analysis (PCA - available in appendix) was therefore used to reduce the number of variables. Based on the 164 eigenvalues of the three first axes, three new variables were produced, namely PC1, PC2 and PC3. PC1 165 is driven by Si, PO_4^{3-} and NH_4^+ and explain 57.2% of variability, PC2 by phytoplankton, net primary 166 production and NO_{3-} and explain 18.6% of variability. PC3 by Dissolved Oxygen and explain 10.2% of 167 variability. 168

169 Tab.1 here

170 2.2. Data processing

171 2.2.1. LPUE time-series decomposition

Our strategy to analyse the spatio-temporal dynamics of striped red mullet abundance was to assume an additive decomposition of the LPUE in terms of a Long Term Trend (LTT), a Seasonal Trend (ST) and a residual R. *LTT* aimed to describe the interannual changes in LPUE while ST was dedicated to characterizing the amplitude of the seasonal fluctuations. LPUE time-series exhibited a very high variance and needed to be log transformed to be modelled at a given time t (in months from January 2005 to December 2018) and

location r, Y_{rt} being here below the value of $log(LPUE_{rt})$, the decomposition was written:

$$Y_{rt} = LTT_{rt} + ST_{rt} + R_{rt}.$$
(1)

178 2.2.2. Long Term Trends

For each rectangle r, a moving median $(MM(Y_{rt}))$ over a 48-month window was applied to remove intraannual or seasonal effects and then investigate appropriate model for estimating the LTT. We proposed to investigate the Long Term Trends applying a smoothing function to time:

$$MM(Y_{rt}) = \alpha_r + f_r(t) + \epsilon_{rt}, \tag{2}$$

where ϵ_{rt} follows a normal distribution $\mathcal{N}(0, \sigma^2)$. The smooth function of each rectangle $f_r(\cdot)$ is defined using spline bases [see Wood (2017)]. In practice we used generalized additive model (GAM) through the mgcv R package provided by Wood (2017). Note that an intercept α_r by rectangle was required due to some optimization constraints used in mgcv (see Chapter 5 and 7 of Wood (2017)).

The fitted model (2) provided a smooth estimation of the time-dependent inter-annual trend for each rectangle: $\widehat{LTT_{rt}} = \widehat{\alpha}_r + \widehat{f_r(t)}$.

188 2.2.3. Seasonal Trends

The seasonal component was obtained by removing the estimated trends from the initial log(LPUE)data:

$$ST_{rt} = Y_{rt} - \widehat{LTT}_{rt}.$$

¹⁹¹ The subsequent analysis is twofold:

• The first bulk Seasonal Trend model in each rectangle was obtained from dynamic harmonic regression Young et al. (1999):

$$ST_{rt} = \sum_{j=1}^{K_r} \{\alpha_{0,r} cos(\omega_j t) + \alpha_{1,r} sin(\omega_j t)\} + \epsilon_{rt}$$
(3)

where $\omega_j = \frac{2\pi}{m}j$ are the harmonic frequencies associated with the seasonality in the series, *m* being the annual number of months, i.e. 12. $\epsilon_{rt} \sim \mathcal{N}(0, \sigma_r^2)$. Depending on the selected rectangle, a range of harmonics $(2 < K_r < 6)$ were tested to provide the best model based on AIC comparisons (Sakamoto et al., 1986). This first bulk Seasonal Trend model in each rectangle provided an overview of the LPUE seasonality.

• Further insight on LPUE spatio-temporal dynamics was provided by a time-frequency analysis of the seasonal component. A discrete Fourier analysis (Bloomfield, 2004) was performed on the seasonal component for each ICES rectangle to provide the spectral density (i) over the complete time-series and (ii) over a moving 48-month window for time-resolved analysis. From the latter analysis, the amplitude of seasonal fluctuations associated with annual (12 months) and biannual (6 months) periods were extracted, named $ST_{LPUE,rt}^{12}$ and $ST_{LPUE,rt}^{6}$, respectively:

$$ST_{LPUE,rt}^{n} = \sqrt{\int_{f_1}^{f_2} SD(u)du}$$

$$\tag{4}$$

where $SD(\cdot)$ is the spectral density (Stoica et al., 2005), n = 6 or 12 for biannual and annual fluctuations, respectively and f_1 and f_2 are the frequency boundaries taken at 5 and 8, and 9 and 16 for the biannual and annual fluctuations.

208 2.3. Dependency on environmental variables

209 2.3.1. Long Term Trends

Models for Long Term Trends were defined for environmental covariates time-series similarly as done for LPUE (see Eq. 2). Links between estimated long term trends of LPUE (noted \widehat{LTT}_{LPUE}) and various environmental covariates were first tested using an uni-variate linear model. Five relevant environmental variables were then selected for the multivariate linear model: the fitted trend of median water salinity \widehat{LTT}_{MS} , the fitted trend of median surface temperature \widehat{LTT}_{MT} , and the fitted trends of PC1, PC2 and PC3 variables \widehat{LTT}_{PC1} , \widehat{LTT}_{PC2} and \widehat{LTT}_{PC3} , respectively. For each rectangle, we modeled:

$$\widehat{LTT}_{LPUE,t} = \beta_0 + \beta_1 \widehat{LTT}_{MT,t} + \beta_2 \widehat{LTT}_{MS,t} + \beta_3 \widehat{LTT}_{PC1,t} + \beta_4 \widehat{LTT}_{PC2,t} + \beta_5 \widehat{LTT}_{PC3,t} + \epsilon_t, \quad (5)$$

where $\epsilon_t \sim \mathcal{N}(0, \sigma^2)$.

217 2.3.2. Seasonality

The seasonal component of environmental covariates was processed by time-frequency analysis similarly to the LPUE data. For each variable and each rectangle, the annual and biannual amplitudes of seasonal fluctuations were obtained from the time-resolved spectral density over a 48-month moving window.

A linear model was built by rectangle to test the relationship between the annual and biannual amplitudes of seasonal fluctuations of LPUE and environmental covariates data:

$$ST_{LPUE,rt}^{n} = \gamma_{0} + \gamma_{1}ST_{MT,t}^{n} + \gamma_{2}ST_{MS,t}^{n} + \gamma_{3}ST_{PC1,t}^{n} + \gamma_{4}ST_{PC2,t}^{n} + \gamma_{5}ST_{PC3,t}^{n} + \epsilon_{t}.$$
(6)

where n = 6 and 12 for biannual and annual fluctuations and ϵ_t follows $\mathcal{N}(0, \sigma^2)$.

224 2.3.3. Models performance

Some parameters were selected for each linear model created for LTT and ST. Firstly, estimates of the 225 coefficient of the linear regression and associated p-value were recorded. Estimates is the mean change in 226 the LTT or ST for one unit of change in the covariate. A low p-value (e.g. <0.05) means changes in the 227 LTT or ST value are related to changes in the covariate. Secondly, the global R^2 and relative importance 228 per covariate were calculated. The model global R^2 indicates the proportion of variance of LTT or ST 229 explained by the created model. Relative importance of covariate enables us to determine the share of 230 variability accountable for each environmental covariate. In other words, relative importance of covariate is 231 the proportionate contribution each covariates makes to global R^2 . Relative importance of covariates was 232 calculated with the relimpo package of R software (Grömping et al., 2006). The used metric is "lmg", which 233 provides a decomposition of the model explained variance (Grömping et al., 2006). 234

235 3. Results

236 3.1. LPUE time-series decomposition



Figure 1: Time-series decomposition in striped red mullet abundance. A: Map of studied rectangles. B, C and D: Long Term Trend (LTT) and associated GAM error, Seasonal Trend (ST) and Residual (R) of log(LPUE) by ICES statistical rectangle, respectively.

Figure 1A presents the locations of the ICES rectangles in this study. Data were recorded from 01/01/2005237 to 12/31/2018. Due to moving average processing, the two first and the two last years of data were deleted. 238 Data are so presented from 01/01/2007 to 12/31/2016. The first observation in Figure 1B was an overall 239 decreasing of \widehat{LTT} of log(LPUE) during time, with an average value of approximately 20 % between 2006 240 and 2017. To go further we used a linear model by rectangle to assess the ability of LTT to explain initial 241 log(LPUE). LTT variable is significant (p-value < 0.01) in almost all linear models to explain log(LPUE) 242 data, but not for rectangles R23R7 and R22E7. A finer analysis of Figure 1B revealed a significant spatial 243 variability in time evolution of striped red mullet abundance with three main periods: 2006-2010, 2010-2014 244 and 2014-2017. First, during the 2006-2010 period, R23E7, R23E6 and R21E7 showed weakly fluctuating 245 values, R20E8 and R21R8, situated at the southeast of the study area, showed a significant increase and 246

R22E7 showed a clear decreasing trend. The second and third periods, respectively 2010-2014 and 20142017, showed more spatial consistency. The former period was characterized by a global decrease for all
zones while the latter period showed overall more stable values.

Seasonal patterns computed using Equation 3 also highlighted spatial variability, see Figure 1C. Sig-250 nificant intra-annual fluctuations were observed for most ICES rectangles. A striking observation was the 251 time and space variability of these seasonal fluctuations. Such variability in the frequency domain calls for 252 more detailed spectral analysis which will be presented in Section 3.3. The only zone clearly dominated by 253 uni-modal distribution is R23E6, i.e. the most north-westerly rectangle studied. Other rectangles showed 254 additional periodic bi-modal or higher frequencies fluctuations. The time-change was also spatially variable. 255 with a quite steady seasonal shape for R20E8, R21E8, R23E6 and R23E7 rectangles and more visible change 256 for R21E7 and R22E7. For this latter area, the seasonal signal nearly drops to zero between 2009 and 2011 257 before regaining values similar to other zones. 258

Figure 1D shows decomposition residuals. For all sites, the residuals did not exhibit any discernible pattern. This confirmed that the selected additive decomposition had extracted most of the relevant information in the striped red mullet abundance time-series.

262 3.2. Long term effects

Figure 2 depicted the comparison between \widehat{LTT} s for striped red mullet abundance and selected environ-263 mental covariates. Results were first presented in qualitative terms, while a more quantitative analysis is 264 provided hereafter by the linear model. \widehat{LTT} for log(LPUE) was described and compared in the previous 265 section (see Figure 1B). \widehat{LTT} for environmental covariates generally shown a strong spatial variability. The 266 surface salinity (MS) was lower for coastal rectangles R23E7, R21E8 and R20E8 due to the continental 267 freshwater inputs, in particular by the Garonne and Charente rivers. For most rectangles, the overall trend 268 was a slight decrease of salinity affected by a rise during the 2010-2012 period. The surface temperature 269 (MT) shown the expected meridional variation (higher temperature in the south) and an overall increase for 270 all rectangles. For the northern areas (R23E6, R23E7 and R22E7), the temperature tended to vary in the 271 opposite way to the salinity: low temperatures were associated with high salinity during the 2009-2012 period 272 while rising temperatures were associated with lowering salinity after 2012. This trend was less marked for 273 the southern rectangles. The PCA-reduced environmental variables were generally well connected in space. 274 \widehat{LTT} for PC1, PC2 and PC3 were maximal in coastal regions R21E8, R20E8 and R22E7 and decreased off-275 shore in R21E7 and R23E6. The only exception was the R23E7 coastal rectangle for which PC1 was maximal 276 while PC2 and PC3 was minimal. Changes over time for PC1 were generally very weak. Stronger dynamics 277 were observed for PC2, for which low values were associated with low temperatures and high salinity events 278 and, inversely, to rise after 2012 when salinity/temperature are generally decreasing/increasing. For PC3. 279 the signal was nearly stable for R20E8 and R21E7 and mainly decreasing for R21E8, R22E7 and R23E6, 280 without any straightforward relation with other environmental covariates. For R23E7, PC3 followed PC2 281 trends, with a strong link with surface temperature. 282



Figure 2: Long Term Trends (LTT) for striped red mullet abundance and environmental covariates for all studied ICES rectangles. LPUE is landing per unit of effort, expressed in monthly mean of daily kg by rectangle, MS is salinity in grams per liter and MT is temperature in degrees Celsius.

The linear model presented in Equation (5) was used to assess quantitatively the relationships between 283 \widehat{LTT} for striped red mullet abundance and environmental variables. The results were summarized in Table 284 2. The first observation was that all R^2 values were above 0.8 (excepting R22E7 which shows $R^2 = 0.65$), 285 meaning that trends in environmental variables were able to explain at least 80% of trends in striped red 286 mullet abundance. Note that these R2 values were not directly related to the raw data and they had to 287 be interpreted on the explained variance of the extracted long term trends $\hat{L}L\hat{T}$ of the LPUE obtained 288 from 2 by the long term trends of environmental variables. Our main goal was to investigate and highlight 289 the importance of variables to explain striped red mullet LPUE. Another observation was that striped red 290 mullet abundances were predominantly positively affected by salinity for five out of six rectangles. For the 291 coastal rectangles R20E8, R21E08 and R23E7, striped red mullet abundance could be linked to continental 292 freshwater inputs. The role of temperature was mainly significant in the two northern zones R23E6 and 293 R23E7, with striped red mullet abundance favored by higher temperature. PC1 generally played a weak 294 role except in the most northwestern zone R23E6/R23E7. PC2 was negatively correlated to abundance 295 in all rectangles except the most southeastern ones R21E8 and R20E8. PC3 displayed stronger positive 296 correlations only for the northern zones R23E6, R23E7 and R22E7. 297

²⁹⁸ Tab. 2 here

299 3.3. Seasonality

Figure 1 revealed the spatial and temporal variability of the LPUE seasonal component using a bulk 300 harmonic approach. Further insight was provided by the spectral density computed, for each rectangle, over 301 the complete time series, see Figure 3. The spatial discrimination was straightforward: coastal rectangles 302 R20E8, R21E8 and R23E7 were the only ones to show both biannual (6-month period) and annual (12-303 month period) fluctuations. The offshore rectangles R23E6 and R21E7 shown a single annual seasonality, 304 although less marked for R21E7 than for the northwestern R23E6. The remaining rectangle R22E7 had a 305 noisy spectral density. This lack of consistency for the spectrum was related to the fluctuations of seasonality, 306 both in amplitude and period, during the acquisition period revealed in Figure 1. 307



Figure 3: Spectral density of log(LPUE) for each ICES rectangle over the complete time series.

Based on the time-resolved spectral density over a 48-month moving window, the amplitudes of annual and biannual seasonal fluctuations (Eq. 4) were computed in each rectangle for log(LPUE) and environmental covariates. An illustration of the results for rectangle R23E7 is presented in Figure 4. The spectral density computed over the complete time-series (Figure 4A) revealed that the biannual (6-month) peak observed in log(LPUE) can not be connected to any similar seasonal component in the selected set of environmental covariates while the annual (12-month) peak in log(LPUE) surely had a link with the annual peak observed in all selected environmental covariates.

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The time-resolved spectral analysis of biannual and annual fluctuations over a 48-month moving window was shown in Figures 4B and C, respectively. Biannual seasonality of abundance was variable but the level observed between the start and the end of the studied period did not show any trend and was qualitatively uncorrelated to environmental covariates. Annual seasonality for log(LPUE) shown slightly greater change over time, with minimal values reached between 2009 and 2013. Annual seasonality for environmental covariates also revealed more consistent patterns over time.



Figure 4: Time-frequency analysis for rectangle R23E7. A: spectral density for log(LPUE) and environmental covariates over the full time-series. B and C: time-resolved magnitude of seasonal fluctuations over a 48-month moving window for biannual and annual components, respectively, for log(LPUE) and environmental covariates.

The multivariate linear model presented in Equation 6 was used to evaluate comprehensively the relationship between abundance and covariate seasonality. Complete results were shown in Table 3.

Figure 3 previously demonstrated that the annual seasonality of striped red mullet abundance was well devel-324 oped for rectangles R21E8, R23E6 and R23E7 and to a lesser extent for rectangles R20E8 and R21E7. Table 325 3 further indicates that the dependency of abundance seasonality on environmental covariates seasonality 326 was not systematic and not spatially uniform. PC1 annual seasonality appeared to affect abundance annual 327 seasonality in rectangles R20E8 and R23E6, with negative and positive correlations, respectively. R20E8 328 was furthermore connected to the annual fluctuations of salinity and temperature: abundance seasonality 329 was favored by strong temperature and weak salinity fluctuations. R21E7 and R21E8 were connected to 330 salinity and temperature annual fluctuations, respectively, while R23E7 annual seasonality did not show any 331 significant relation with environmental parameters. 332

Biannual seasonality of log(LPUE) was well established in the three coastal rectangles R20E8, R21E8 and R23E7 (Fig.3). For the latter rectangle, the biannual seasonality was not related to environmental covariates seasonality, similarly as for annual seasonality. PC1 biannual seasonality was observed to affect both R20E8 and R21E8, while salinity biannual seasonality only affected R20E8.

337 Tab. 3 here

338 4. Discussion

The aims of the present study were (i) to assess long-term and seasonal changes in striped red mullet 339 abundance in the Bay of Biscay, and (ii) to identify the possible relationships between changes in abundance 340 and environmental factors. This study was carried out on the basis of an indicator of abundance, which 341 was the sole information available for striped red mullet natural fish populations within the Bay of Biscay. 342 As no appropriate methodology to tackle this problematic with these data already existed in literature, we 343 had to develop a new one. Thus, the main aim of this study was not to present a new methodology, but 344 to better understand the effects of the environment on striped red mullet abundance. Here, we will discuss 345 results and compare them with other studies, but some conclusion will remains hypothtical due to a very 346 poor documentation in literature on the ecology of striped red mullet. 347

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Long term trends were inferred from the abundance data using a moving average with a smoothing function. The resulting trends were clearly non-linear, thus ruling the use of linear approaches (see Hess (2001); Lentka and Smulko (2019) for trends detection). Seasonal components were processed using timefrequency analysis with moving-window spectral computations, which highlighted the strong fluctuations of the abundance seasonality both in space and time. Other tools for time-frequency analysis could have been used here such as the wavelet analysis (Huang et al., 1998). It is important to note that time-frequency analysis is strongly constrained by the need to have continuous and long enough time-series.

The selected environmental variables also shown non-linear long term trends. These observations result from 356 the complex interactions between a large number of processes affected by fluctuations over wide ranges of 357 space and time scales which were outside of the scope of the present study. However, from a data-processing 358 perspective, it is again worth noting to note that, in our study zone, coupled ocean-atmosphere systems such 359 as the North-Atlantic Oscillation (NAO) (Hurrell, 1995) or other fluctuating large scale climate patterns 360 (Castelle et al., 2017) provided additional variability to the global change track. For instance, the overall 361 warming of the ocean surface observed from the present analysis of Copernicus database, well documented 362 in the literature (Peck and Pinnegar, 2019; deCastro et al., 2009; Koutsikopoulos et al., 1998; Planque et al., 363 2003), combines with inter- and intra-annual oscillations and spatial variability. Similarly, salinity and PCA-364 reduced environmental variables showed strong time and space variability. Such results could be related 365 with climate-driven patterns (Roessig et al., 2004) of atmosphere properties, precipitation, land runoff, river 366 discharge, etc. This reinforces the relevance of the present approach to determine trends and relationships 367 between fish populations and environmental factors. 368

the study of long term trends in striped red mullet abundance shown a high spatial heterogeneity within 369 the Bay of Biscay even on a relatively small scale. The dominant trend was a decrease in striped red 370 mullet abundance. This decrease varied among the studied ICES spatial rectangles. But, for almost all 371 rectangles, we could link 80% of this trend variance to environmental covariates trends. The dependency on 372 373 environmental covariates was also spatially variable. In coastal rectangles, long term trends of abundance can had a slight positive link with surface salinity trends. While for more off-shore rectangles, striped red 374 mullet abundance increased along with salinity. Striped red mullet salinity preference was detected to be 375 between 35.1 and 35.6 grams per liter (Desbrosses, 1935). These values were for adults fishes at depths 376 between 100 and 200 meters. One possible explanation for positive link in trends between striped red mullet 377 abundance and salinity in coastal rectangles is that the striped red mullet is at the lower limit of its salinity 378 ecological niche in coastal rectangles (see table 1). When salinity decreases in coastal zones, due to inland 379 freshwater inputs, striped red mullet abundance decreases. More precise information about the striped red 380 mullet ecological niche is needed to consolidate this hypothesis of a positive link between striped red mullet 381 abundance trends and salinity in coastal zones. 382

In the southern Bay of Biscay, striped red mullet have already been detected as species with an affinity 383 for warm temperatures (García-Rodríguez et al., 2011). In our study area, in the northern Bay of Biscay, 384 mean sea surface temperature was around 14.5 degrees Celsius for the colder rectangles and 15 degrees 385 Celsius for the warmer rectangles. We did not detect links between temperature trends and striped red 386 mullet abundance except for the two coldest rectangles. In these two northern rectangles, abundance trends 387 decreased when temperature trends decreased and conversely, which tends to confirm the observation of 388 García-Rodríguez et al. (2011). For a similar species *M. barbatus* in Aegean Sea, abundance has been proved 389 to be higher in warm bottom water $(19^{\circ}C)$ than in colder waters (Maravelias et al., 2007). In addition, this 390 species seems to avoid the cold bottom waters ($< 16^{\circ}$ C) (Maravelias et al., 2007). An hypothesis to explain 301 the freshness of the two northern rectangles is they are in a river freshwater inputs and up-welling zone. 392 This hypothesis is favoured by the study of Puillat et al. (2006) demonstrating some events of up-welling-393 favourable winds exactly in the zone were we found the lowest temperature, and a positive link with striped 394 red mullet abundance. In these rectangles, red-mullet is at the edge of its favoured life condition regarding 395 temperature. When water temperature decreases, striped red mullet falls outside its life preferendum and 396 abundance decreases. 397

³⁹⁸ Up-welling and river discharges also induce chemical transports or dissolution from sediments. In this study,

³⁹⁹ we have attempted to find a relationship between striped red mullet abundance and chemical concentrations

in the sea water. As strong correlations were initially observed between different chemical species, we chose 400 to perform a dimension reduction with PCA. But, interpretation of linear model estimates of PCA reduced 401 dimensions can be delicate. Only the two southern rectangles do not shown a link with PCA reduced 402 dimensions. For the four other rectangles, trends in striped red mullet abundance were linked to chemicals 403 and primary production. The link was positive with PC3. As PC3 had a positive contribution of O_2 404 concentration, we can say striped red mullet abundances were positively linked to O_2 concentration. This 405 result is consistent with studies highlighting an increase of capture probability of an other mullet species 406 along with dissolved oxygen rate (Maes et al., 2007). In the four northern rectangles, trends in striped 407 red mullet abundance were also positively linked with PC1. PC1 is built from a negative contribution of 408 Si, PO_4^{3-} and NH_4^+ , meaning that striped red mullet abundance was negatively linked to Si, PO_4^{3-} and NH_4^+ concentrations. The effect of chemicals, phytoplankton and net primary production rates on striped 409 410 red mullet abundance were never investigated. Further investigations are required to confirm the present 411 original observations. 412

The seasonal study revealed high variability of striped red mullet abundance seasonality within the 413 Bay of Biscay. In the most off-shore rectangle, striped red mullet abundance shown well defined annual 414 oscillations while in the most shoreward ICES rectangles, additional biannual oscillations were observed. 415 In the two remaining ICES rectangles, no clear periodic pattern was observed. Multiple peaks per year 416 when studying seasonality in fisheries stock abundance were already reported in the case of multiple species 417 studies (Hsieh et al., 2009), but such a variability for a single species was not reported before, to the best 418 of our knowledge. One peak in seasonality of annual abundance occured at intermediate depth. Following 419 the literature on striped red mullet ecology, spawning commonly occurs in spring and early summer (May 420 and June) (Desbrosses, 1935; N'Da and Déniel, 1993; Mahé et al., 2005; Leaute et al., 2018). Even if 421 massive movement of spawners during reproduction has not been reported for striped red mullet, our best 422 hypothesis regarding the annual seasonal peak in striped red mullet abundance may reflect movements during 423 reproduction process. But, to our knowledge, it has to date never been statistically proven. We also modeled 424 this one peak annual seasonal fluctuation of striped red mullet abundance along with similar environmental 425 annual seasonal fluctuations. This dependency was not systematic and not spatially uniform. Temperature 426 was the most relevant parameter linked with seasonal fluctuation in our study, but it was only significant 427 in three rectangles. This result supports our hypothesis of the annual spawning peak. Higher sea water 428 temperature has been reported during striped red mullet spawning in Israel and Canary Islands (Suquet and 429 Person-Le Ruyet, 2001). For Brittany, striped red mullet spawning is observed in relation with increasing 430 photo-period and temperature (Suquet and Person-Le Ruyet, 2001). One hypothesis is that the second peak 431 may be explained by the recruitment process corresponding to the arrival in the catches of the individuals 432 born the year before. This hypothesis is supported by the fact that biannual abundance seasonality occurs in 433 shallow coastal rectangles and that multiple recruitment peaks have also been observed for different species 434 (see, for example Abella et al. (1995) or Hatfield (1996)). We tried to model the biannual peak of seasonality 435 in striped red mullet abundance with the corresponding peak in environmental covariates but we were not 436 able to link the three coastal rectangles to the same set of environmental variables. One rectangle can be 437 linked to low fluctuation of PC1, one to low fluctuation of PC3 and the third to all variables excepting 438 PC3. The two rectangles where no seasonality was detected are in deep zones. Seasonality of striped red 439 mullet abundance was clearly depth driven. Further studies will be needed to better reveal the link between 440 seasonality in striped red mullet abundance and environmental variables. 441

A critical issue for a deeper understanding of the dependency of fish abundance on the oceanic variable is 442 the proper definition of space and time scales. The focus was here on the Bay of Biscay over the 2005-2018 443 period. The Copernicus database used to define environmental covariates revealed a mean increasing trend of 444 surface temperature over the considered period, following a century-long tendency (Garcia-Soto and Pingree, 445 2012) for the Bay of Biscay. This trend may appear contradictory with the large scale decrease of North 446 Atlantic sea surface temperature noted for instance by Piecuch et al. (2017) over the 2005-2015 period, but 447 is indeed an indicator of the heterogeneity of the ocean dynamics. This underlines that any generalization of 448 local findings should be made with particular caution and vice versa. From an ecological point of view, the use 449 of ICES rectangles to study the impact of the environment on the red mullet population in the Bay of Biscay 450

raises questions. ICES rectangles correspond to the finest spatial mesh common to the mandatory declarative 451 data of fishermen and not homogeneous ecological units. For example, several habitats may be represented 452 in the same rectangle or conversely, a single habitat may occupy several rectangles. The development of an 453 habitat-delimited analysis will certainly be a fruitful but ambitious and long-term challenge. The present 454 study was strictly bounded by the source data, i.e. by the fishermen's logbooks and fishing forms filled in 455 by ICES rectangle. Moreover there is a crucial need to simultaneously dynamically refine and extend in 456 space and time the mapping of both marine environmental parameters and fish abundance. Developing a 457 model for such a small portion of the ocean is useful to be able to understand the effect of environmental 458 changes on local fisheries. At this scale, most of the fisheries are local small industries. Understanding 459 the link between environment and resources preserves local socio-economic activities, and it is of very high 460 importance. While costly, such an approach appears now to be a prerequisite to decipher the response of 461 marine fauna to the fluctuations of ocean variables. This is already true for the relationships existing at the 462 present time (García-Barón et al., 2020; Véron et al., 2020) and it will become even more necessary to move 463 forward towards a resource-preserving marine spatial planning strategy in the context of global change. 464

The approach we developed to tackle our applied research question can be applied in a wide range of 465 fish population studies, in order to monitor the effect of climate change on fish population distribution 466 at various scales and tohelp fishermen anticipate changes. The analysis was carried out from abundance 467 time-series, which is the sole information available for a number of natural fish populations. It had also 468 been performed considering the main environmental factors available over the study area, mainly focused in 469 the surface layers. When available, additional factors can be easily implemented and tested such as more 470 complete three-dimensional information (Assis et al., 2016). For the study of striped red mullet, knowledge 471 on benthic habitat composition (Ajemian et al., 2016) or on substrate type (Lombarte et al., 2000; Mahé 472 et al., 2005) can improve the study. Substrate type layer is not yet available for the entire Bay of Biscay. 473 One perspective will be to include new layers in this study, once one is available. Finally, a comprehensive 474 understanding of the time-resolved population dynamics should encompass each driving factor, including 475 trophic web parameters and additional mortality causes. 476

477 5. Acknowledgments

Funding was provided by the Energy Environment Solutions (E2S-UPPA) consortium and the BIGCEES
project from E2S-UPPA ("Big model and Big data in Computational Ecology and Environmental Sciences"),
the Queensland Department of Environment and Science (DES) and the ARC Centre of Excellence for
Mathematical and Statistical Frontiers (ACEMS). The data that support the findings of this study is openly

⁴⁸² available in SEANOE (Kermorvant et al., 2020).

483 We do not declare any conflict of interest.

6. Tables

		R20E8	R21E7	R21E8	R22E7	R23E6	R23E7
Salinity	Min	28.51	32.36	29.10	31.80	32.27	26.76
	Max	34.65	35.72	34.57	35.53	35.67	34.58
	Med	33.17	34.85	33.12	34.13	34.80	32.10
	\mathbf{Sd}	1.27	0.67	0.88	0.66	0.65	1.52
Temperature	Min	7.32	8.56	7.09	7.76	8.57	7.32
	Max	22.16	22.54	21.22	21.64	21.52	20.89
	\mathbf{Med}	15.26	15.25	14.85	14.84	14.44	14.47
	\mathbf{Sd}	4.23	3.77	4.16	3.79	3.34	3.80

Table 1: Main statistical values of two environmental variables (temperature and salinity) recorded over the 2005–2018 period in the selected ICES rectangles. Temperature is expressed in degrees Celcius, salinity in grams of salt per liter of water

Table 2: Results of the linear model to assess the relationship between long term trends of stripped red mullet abundance and environmental variables. Models estimates (β), P-values, R^2 and relative importance (Imp) of variables obtained with linear models on each rectangle. Results with P-values < 0.005 are in bold. Standard deviations of models estimates are available in appendix.

	Type	Intercept	\widehat{LTT}_{MS}	\widehat{LTT}_{MT}	\widehat{LTT}_{PC1}	\widehat{LTT}_{PC2}	\widehat{LTT}_{PC3}	\mathbb{R}^2
R20E8	β	-6.906	0.433	-0.35	0.438	-0.879	0.373	
	P_value	0.358	$<\!0.001$	0.219	0.141	0.337	0.611	
	Imp.		0.231	0.246	0.102	0.214	0.069	0.862
R21E7	β	-7.495	0.102	0.205	-0.873	-1.739	1.743	•
	P_value	0.036	0.142	0.1	$<\!0.001$	$<\!0.001$	$<\!0.001$	
	Imp.		0.147	0.308	0.083	0.295	0.069	0.902
R21E8	β	-15.951	0.524	0.071	0.358	-0.731	-0.31	
	P_value	0.004	$<\!0.001$	0.772	0.096	0.097	0.67	
	Imp.		0.366	0.163	0.051	0.162	0.087	0.828
R22E7	β	31.395	-0.959	0.177	-0.842	-3.379	2.824	
	P_value	0.011	$<\!0.001$	0.618	$<\!0.001$	$<\!0.001$	0.002	
	Imp.		0.091	0.153	0.077	0.19	0.136	0.647
R23E6	β	-56.003	0.732	1.855	-1.067	-5.673	6.577	
	P_value	$<\!0.001$	0.001	$<\!0.001$	0.003	$<\!0.001$	$<\!0.001$	
	Imp.		0.04	0.214	0.22	0.164	0.254	0.892
R23E7	β	-23.493	0.434	0.632	0.574	-0.922	2.213	
	P_value	$<\!0.001$	$<\!0.001$	$<\!0.001$	$<\!0.001$	$<\!0.001$	$<\!0.001$	
	Imp.		0.215	0.14	0.207	0.16	0.15	0.8725

Table 3: Results of the linear model to assess the relationship between seasonal trends of stripped red mullet abundance and environmental variables. Model estimates(β), P-values, R^2 and relative importance (Imp) of environmental covariates seasonality in abundance seasonality obtained with lm models on each rectangles. ".12" and ".6" notations correspond to the annual and biannual fluctuations, respectively. Results with p-values < 0.005 are in bold. Standard deviations of models estimates are available in appendix.

	Type	Intercept	\hat{ST}_{MS}	\hat{ST}_{MT}	\hat{ST}_{PC1}	\hat{ST}_{PC2}	\hat{ST}_{PC3}	R^2
R20E8.6	β	0.171	0.063	0.077	-0.365	0.298	0.069	
	P_value	$<\!0.001$	0.441	0.031	$<\!0.001$	0.008	0.598	
	Imp.	•	0.009	0.119	0.146	0.059	0.051	0.383
R20E8.12	β	-0.167	-0.28	0.278	-0.721	-0.207	0.462	•
	P_value	0.087	$<\!0.001$	$<\!0.001$	$<\!0.001$	0.456	0.003	
	Imp.		0.152	0.28	0.272	0.27	0.008	0.982
R21E7.6	β	0.141	-0.271	-0.045	-0.182	0.2	-0.056	
	P_value	$<\!0.001$	0.002	0.006	$<\!0.001$	0.007	0.286	
	Imp.		0.167	0.088	0.244	0.075	0.095	0.668
R21E7.12	β	0.232	0.225	-0.141	0.073	0.64	-0.212	
	P_value	$<\!0.001$	$<\!0.001$	0.177	0.571	0.161	0.404	
	Imp.		0.289	0.04	0.034	0.057	0.046	0.466
R21E8.6	β	0.343	-0.765	-0.193	-0.906	0.629	0.024	
	P_value	$<\!0.001$	$<\!0.001$	$<\!0.001$	$<\!0.001$	$<\!0.001$	0.895	
	Imp.		0.261	0.089	0.138	0.069	0.012	0.569
R21E8.12	β	0.227	0.16	-0.25	0.123	-0.018	0.693	
	P_value	0.011	0.014	$<\!0.001$	0.004	0.877	$<\!0.001$	
	Imp.	•	0.144	0.305	0.1	0.266	0.095	0.910
R22E7.6	β	0.152	-0.128	-0.006	0.087	0.092	-0.24	
	P_value	$<\!0.001$	0.231	0.82	0.07	0.062	0.002	
	Imp.		0.015	0.044	0.021	0.025	0.123	0.229
R22E7.12	β	0.648	0.478	0.37	0.157	-0.973	-1.629	
	P_value	$<\!0.001$	0.252	0.003	0.253	$<\!0.001$	$<\!0.001$	
	Imp.		0.102	0.068	0.027	0.066	0.256	0.520
R23E6.6	β	0.103	-0.026	-0.08	0.26	-0.236	0.24	•
	P_value	0.012	0.897	0.128	0.211	0.253	0.089	•
	Imp.		0.008	0.023	0.054	0.012	0.078	0.175
R23E6.12	β	0.403	-0.131	-0.248	1.124	-0.588	-0.092	•
	P_value	0.003	0.647	0.026	$<\!0.001$	$<\!0.001$	0.456	
	Imp.		0.032	0.243	0.409	0.036	0.241	0.961
R23E7.6	β	0.183	-0.024	0.068	-0.016	-0.076	0.148	•
	P_value	$<\!0.001$	0.307	0.001	0.505	0.004	0.002	
	Imp.	•	0.02	0.072	0.007	0.097	0.077	0.273
R23E7.12	β	-0.341	0.025	0.241	-0.228	0.192	-0.106	•
	P_value	0.002	0.532	0.029	0.005	0.13	0.347	
	Imp.		0.097	0.074	0.131	0.057	0.084	0.442

7. Figure legends

Figure 1: Time-series decomposition in striped red mullet abundance. A: Map of studied rectangles. B, C and D: Long Term Trend (\widehat{LTT}) , Seasonal Trend (ST) and Residual (R) of log(LPUE) by ICES statistical rectangle, respectively.

Figure 2: Long Term Trends (\widehat{LTT}) for striped red mullet abundance and environmental covariates for all studied ICES rectangle. LPUE is landing per unit of effort, expressed in monthly mean of daily kg by rectangle MS is salinity in gram per liter and MT is temperature in degree Celsius, PC1, PC2 and PC3 are the

three first axes of PCA of phytoplankton concentration, chlorophyll concentration, and chemicals variables. Figure 3: Spectral density of log(LPUE) for each ICES rectangle over the complete time series.

Figure 4: Time-frequency analysis for rectangle R23E7. A: spectral density for log(LPUE) and environmental covariates over the full time-series. B and C: time-resolved magnitude of seasonal fluctuations over a 48-month moving window for biannual and annual components, respectively, for log(LPUE) and environmental covariates. MS is salinity in gram per liter and MT is temperature in degree Celsius, PC1, PC2 and PC3 are the three first axes of PCA of phytoplankton concentration, chlorophyll concentration, and chemicals variables.

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