

~~Spatial and temporal representation~~ Temporal Representation of marine fish occurrences available online ~~Marine Fish Occurrences Available Online~~

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Abstract

Despite the 243,000 ~~species of~~ marine species described by 2022, our
21 knowledge about the oceanic biodiversity is still incomplete. This knowl-
edge gap carries potentially adverse and far-reaching consequences for the
24 preservation of marine ecosystems, particularly in the context of the ongo-
ing human-induced alterations to our biosphere and the rapid progression of
climate change and global environmental shifts.

However, recently, a large number of online repositories ~~cataloging~~,
27 ~~storing and distributing biodiversity information~~ ~~hosting taxonomic information~~
~~have emerged, which catalogue, store and distribute biodiversity information,~~
~~including taxonomic~~ and species occurrence data ~~have emerged recently~~. Fish-
30 Base, the Global Biodiversity Information Facility (GBIF) and the Ocean
Biodiversity Information System (OBIS) are part of these publicly available
repositories representing a variety of sources that have exploded in number.
33 However, despite the incredible accumulation of biodiversity records, not all

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the information is ~~really actually~~ useful, nor does it represent any new knowledge regarding global species richness patterns.

3 In this study, we assessed the spatial and temporal representativeness of marine fish records (order Actinopterygii) found in the GBIF and OBIS global repositories. ~~We have developed a~~ The methodological framework that

6 we developed relies on a series of non-parametric estimators for computing species richness from incidence data. This methodology employs hexagonal grids as sampling units that overlay marine bioregions across the globe.

9 Using standard ecological and spatial analysis tools, we identify regions that are adequately represented in terms of available records and therefore have more reliable data, as well as regions with few records that do not rep-

12 resent current species richness. We overlap these results with the location of marine protected areas and fishing exploitation zones to understand the anthropogenic effect on marine ichthyofauna. We additionally evaluate hy-

15 potheses regarding the taxonomic, geographic, and temporal distribution of information biases to deepen our current understanding of public records of species occurrences worldwide.

18 Considering that more than 40 years of information was analyzed, the results showed that, on a global scale, the primary data on marine fish available on GBIF and OBIS platforms are still far from being representative

21 and complete. Only 1.14% of the records were useful for our analyses. In addition, we found that the information seems to be biased towards coastal areas, regions close to developed countries, and areas where there is a large

24 fishing activity. Finally, the best represented species and families are those with a small body size, which use shallow habitats and ~~have commercial are~~ are usually recognized as having commercial or cultural value.

27 *Keywords:* Ecoinformatics, Ecological Information Biases, Marine Fish, Spatial and Temporal Representativeness, Species Richness

1. Introduction

30 Currently, the more than 243,000 species included in the World Register of Marine Species database ([WORMS, 2022](#)) suggests that only 11% to 78% of all marine species have been discovered, revealing a striking picture
33 of vastly incomplete knowledge that may have serious implications for marine conservation ([Luypaert et al., 2020](#)). Moreover, ongoing climate change

represents one of the greatest threats to biodiversity (Malhi et al., 2020; Turner et al., 2020) and has already been documented to modify the distribution of marine species (Lenoir et al., 2020). Some of the ~~described effects~~
3 ~~effects described~~ includes the invasion of non-native species leading to massive species turnover that may ~~lead to result in~~ the local extinction of large
6 ~~proportions share~~ of species (Cheung et al., 2009).

It is crucial to recognize that species richness, ~~while being a diversity metric among many,~~ is, in itself, an aggregate variable ~~subsuming the overall variety of life quantifying the end result of the splitting and lumping of the tree of life as a product of evolutionary processes~~ (Marquet et al., 2004). Consequently, numerous endeavors have been directed towards the development of more comprehensive diversity indices, giving rise to significant scientific literature, aimed at describing ecological heterogeneity (Tuomisto, 2011; Moreno and Rodríguez, 2011; Daly et al., 2018). However, within 9 this literature, there appears to be a shifting focus towards examining the ramifications of biodiversity loss. This shift involves the adoption of new terminology designed to provide pragmatic concepts, such as “species inventory”, “taxonomic inventory”, or “inventory completeness”, which are intended to convey more precise messages to policymakers, summarizing the richness of biodiversity (Pereira et al., 2013; Butchart et al., 2010). Nevertheless, while the scientific community engages in debates over the use of 12 biodiversity terminology, it is important to note that species richness continues to offer a concise and easily manageable description of variability across 15 various other parameters characterizing the biota in both spatial and temporal dimensions (Appeltans et al., 2012). Species richness remains an essential feature for comprehending how diversity evolves in response to natural and 18 anthropogenic influences within biomes, regions, and ecosystems (Troia and 21 McManamay, 2017; Magurran and McGill, 2011).

Likewise, biodiversity can also be assessed through life history traits, 24 which are modulated by both evolutionary factors and ~~the variation in habitats~~

~~and ecosystems~~ habitat ecosystem variations (Neigel, 1997; Hutchings and Baum, 2005). We now know that biodiversity is more likely an expression of the heterogeneity of such life history traits. Alò et al. 2021, for example, ~~shows~~ show that while some of the fish diversity is certainly due to environmental processes, a large fraction of such richness variance is also determined by evolved life history traits related, for example, to migratory habits. Therefore, evaluating how life history traits impact richness metrics should deepen our understanding of fish diversity patterns.

While still short of having a robust and standardized biodiversity infrastructure (Heberling et al., 2021), ~~a~~ there is great diversity of online repositories with taxonomic information and species occurrences dataexist. Among the most important databases hosting marine information are FishBase, a platform that hosts information on the taxonomy of fish, their ecology, trophic information, habitat, and history of uses dating back to more than 250 years (Froese and Pauly, 2000); and the Global Biodiversity Information Facility (GBIF), a platform that stores and allows for the free access to species occurrence records from around the world. GBIF is currently one of the repositories hosting the largest amount of such data in the world (Telenius, 2011; GBIF: The Global Biodiversity Information Facility , 2021); and finally Ocean Biodiversity Information System (OBIS), which houses data on the occurrence and abundance of species from exclusively marine environments (OBIS: Ocean Biodiversity Information System, 2021). Records entered in these repositories are often used for research related to biodiversity assessment, taxonomic reviews, red listing of threatened species, species distribution, and generation of ecological niche models, among others (Yesson et al., 2007). GBIF currently offers more than 1.62 billion occurrence records and OBIS more than 63 million, which increase considerably each year (GBIF: The Global Biodiversity Information Facility , 2021; OBIS: Ocean Biodiversity Information System, 2021).

The records of both platforms come from a wide variety of sources col-

lected following different methodologies at different temporal and spatial scales~~introducing~~, which introduces a great variety of biases (Beck et al., 2014; Zizka et al., 2020). Among these, three main types of biases have been described: (i) taxonomic, this occurs when some species and/or families are better sampled than other rarer species (Chandler et al., 2017); (ii) geographic, when data input is unevenly distributed across geographic regions and may prove to obscure ~~inter-region~~interregional comparisons (Yang et al., 2013; Yesson et al., 2007); and (iii) temporal, which may be prevalent when comparing different time periods as data coverage is unevenly distributed over time (Chandler et al., 2017; Yang et al., 2013). While these biases introduce some uncertainty regarding reliability of species richness descriptions obtained from online platforms (Beck et al., 2014; García-Roselló et al., 2015), they have largely been used to provide an extensive overview of macroecological patterns of distribution not available otherwise (Mora et al., 2008; Troia and McManamay, 2017).

Still, identifying how sampling ~~effort~~is efforts are distributed across space and time is a ~~necessary required~~ step to interpret biodiversity patterns and reduce biases~~as understanding the distribution of our biota is essential to design~~as understanding our biota distribution is critical for well-designed protection efforts. This may be achieved through different weighting schemes for records in areas with sufficient sampling that provide a more reliable contribution compared to underrepresented regions (Phillips et al., 2009; Hortal et al., 2008; Yang et al., 2013).

We here assessed the spatial and temporal representativeness of marine fish records available in the global GBIF and OBIS repositories at the ~~level of marine bioregions~~marine bioregions' level in order to pinpoint the location of records that best quantify ~~the diversity of marine fishes~~marine fish diversity. The result is a spatial representativeness analysis that we then overlay on marine conservation areas (UNEP-WCMC and IUCN, 2022) and fisheries exploitation areas (FAO, 2014) to learn whether marine conservation efforts,

as well as large fisheries, are located in areas of high species richness or ~~in areas of~~areas which insufficient data coverage.

Finally, we also analyzed the potential effect ~~that some attributes could have of some attributes~~ on the incidence of more records in global database repositories. Specifically, we evaluated three research questions related to how body size, habitat depth, and commercial use ~~relates relate~~ to the representation of marine fish occurrences. We ask whether: (i) a better representation in ~~the~~ online platforms may be due to the ~~over sampling oversampling~~ of larger fish, ~~caused by its easy identification; that resulting from an easier identification;~~ (ii) shallow areas provide easy access to sampling; and (iii) economic and commercial ~~interest have elicit interests have elicited~~ a larger representation of culturally relevant species ~~among in~~ online biodiversity repositories.

2. Methods

2.1. Species data

We use all ~~recorded occurrences from the order Actinopterygii hosted in the recorded occurrences of the Actinopterygii order hosted in the~~ GBIF and OBIS repositories (GBIF.org, 2021; OBIS.org, 2021). Following Alò et al. (2021), ~~evolutionarily evolutionary~~ older taxa, such as Cephalaspidomorphi, were excluded from this analysis. ~~The libraries Libraries rgbif and robis of~~ the statistical package R were used for data extraction (Chamberlain, 2017; Provoost and Bosch, 2020; R Core Team, 2018). ~~Both repositories have collaborated since 2001, sharing data on the co-occurrence of marine life~~ (OBIS.org, 2021). Nevertheless, recent investigations have shown significant disparities in data contributions, revealing remarkable low shares of shared data (Chollett and Robertson, 2020; Moudry and Devillers, 2020). Noteworthy distinctions exist between the two platforms, encompassing diverse data sources and methodologies, along with substantial variations in temporal and spatial scales associated with data collection (Zizka et al., 2020). Due to

these disparities, scholars recommend a thorough examination and refinement of these data repositories (Bonnet-Lebrun et al., 2023). To enhance the quality and reliability of the information, a comprehensive series of filters has been systematically applied to our analysis. To minimize errors associated with the public usage of GBIF and OBIS repositories, we curated the dataset following Zizka et al. (2020) and filtered the dataset by the columns labeled “scientific name”, “family”, “year”, “Longitudelongitude” and “Latitudelatitude”. We retained all taxonomic information down to the species level. Any record and removed records with NA values was removed in these columns. We also removed any duplicated record all duplicate records with identical latitude and longitude data, as well as any record records collected before 1980 (see Alò et al., 2021; García-Roselló et al., 2015). Each record was further assigned to a marine bioregions bioregion following Costello et al. (2017). Spatial data manipulation and plotting was performed with the aid of the following libraries: *sf*, *dplyr*, and *cartography* (Giraud and Lambert, 2016; Pebesma, 2018; Wickham et al., 2021). We finally labeled and removed any-all exotic species record using the *distribution()* function provided by the *rfishbase* library (Boettiger et al., 2012; Froese and Pauly, 2021). To limit our analysis to species occurring within their native range, each record was checked against the classification of FAO fisheries area classification for consistency (FAO, 2014). A summary of the number of records is provided in Appendix A.

2.2. Data Analysis by Bioregion

Once the database was cleaned, a subset of the data was created data subset was extracted for each of the 30 bioregions. For each bioregion, a count of records, species and families was made, and families were counted, and the Shannon diversity index was calculated using the *vegan* library in R (Oksanen et al., 2020).

2.2.1. Spatial Representativeness Analysis

To assess the spatial representativeness of the data, we evaluated the spatial representativeness of the data's spatial representativeness. Bioregions were gridded into hexagonal cells of equal surface area to maximize the fit to bioregions' areas using a cylindrical equal area projection (i.e. EPSG Code:54034). We approximated a $1^\circ \times 1^\circ$ hexagonal lattice by computing cells of 10^4 square-kilometers, resulting in a total of 57,067 cells. In the appendix, we evaluated two additional spatial resolutions: 5° and 10° , respectively, using a gridcell of $\approx 5^\circ$ and $\approx 10^\circ$ lattice with a total of 3,029 and 953 cells, respectively, using a 2.5×10^5 and 10^7 square-kilometers in order to analyze square-kilometer gridcell to assess different biodiversity macropatterns (Titensor et al., 2010).

The expected species richness (S_{exp}) was computed as the average mean between three non-parametric richness estimators so that $S_{exp} = \frac{1}{3} \sum_i^3 S_i$, where S_i is Chao2 (S_{chao}), Bootstrap ($S_{bootstrap}$) and Jackknife 1 ($S_{jackknife1}$) (see Magurran and McGill, 2011, for individual definition of indices). Such averaging seeks (see Magurran and McGill, 2011, for individual index definitions). The purpose of this averaging is to minimize biases and potential errors of under- or overestimation by using a single richness estimator following the work of (Mora et al., 2008; Troia and McManamay, 2017) by Mora et al. (2008) and Troia and McManamay (2017).

We then produced a species representativeness index (SRI) by comparing the observed richness (S_{obs}) per cell to S_{exp} (Troia and McManamay, 2017), $SRI_i = \frac{S_{obs}}{S_{exp}}$. This index indicates the degree of representativeness of records is an undersampling index that points to the records' representativeness to quantify the actual species richness in each cell (i). Its value ranges from 0 to 1, where 0 represents an unsampled cell and 1 represents a fully sampled one cell.

Since the Species Richness Index (SRI) serves as an indicator of how databases depict is used as a metric to assess the databases accuracy in

depicting actual species richness, it is reasonable to categorize cells arbitrarily. Consequently, we propose a systematic categorization of cells into three classes: "Few," "Sufficient," and "Adequately representative" of estimated species richness. We establish these classifications, "low," "medium" and "high", based on the frequency distribution of SRI (as depicted, as illustrated in Fig. A.1). Some cells contain only one species record and are labeled. Cells with only one record are identified as having insufficient records (IR) to estimate for estimating S_{est} . Certain cells may exhibit limited knowledge with SRI falling within Those with an SRI in the range (0, 0.60), while others may demonstrate a sufficient level of species diversity knowledge for a comprehensive representation if SRI falls within are categorized as low, while those falling within the interval (0.60, 0.85). Additionally, some cells will possess an adequate representativeness level if SRI falls within may be characterized as having a medium level of representativeness. Furthermore, cells within the range (0.85, 1.00). Cells with one or no records are treated as distinct classes, as are cells with a single record, in order to identify those cells with insufficient records for SRI estimation. Maps displaying are identified as high, meaning an adequate representation of species diversity. Fig. A.2 show maps illustrating the raw values for observed species richness (S_{obs}), expected species richness (S_{exp}), and SRI can be found in Fig. A.2 values.

2.2.2. Temporal Representativeness Analysis

We constructed species accumulation curves, employing using years as the units of sampling sampling unit, to examine the temporal distribution of data records within each bioregion. To assess the adequacy of the sample sample's adequacy, we focused on data from the last four years of data (2016-2020), representing the final 10% of each accumulation curve. We employed used a linear fit following the rescaling of the SRI to facilitate statistically comparable slope measurements. Slopes approaching zero suggest bioregions that have been adequately sampled, whereas slopes deviating from zero indicate insufficient sampling efforts over time.

2.2.3. Gap Analysis

We overlaid the spatial representativeness map (§2.2.1) with ~~shapefiles~~
3 ~~of~~ Marine Protected Areas (~~MPA~~)~~MPAs~~ ~~shapefiles~~ (UNEP-WCMC and
IUCN, 2022) and fishing exploitation areas reported by (FAO, 2014). The
superposition of these layers allowed us to calculate the extent of protection
6 offered by ~~MPA~~~~MPAs~~ for each bioregions on a cell basis, and the extent
of cells in designated fishing zones. ~~This exercise allows to jointly assess the~~
~~relationship between~~~~Based on this exercise, the relationship among~~ two op-
9 posing human impacts and current uncertainties about marine fish diversity
~~can be assessed.~~

2.2.4. Bias Assessment

12 ~~The evaluation of potential biases generated by~~ Potential biases resulting
from body size, habitat depth, and cultural value of species (§2.1) ~~was~~
assessed from the fishbase database were assessed using Fishbase repository
15 information (Froese and Pauly, 2021). We ~~generated~~ developed a frequency
distribution plot for ~~the reported length of each species~~ each species' length
~~reported~~ in the database, employing ~~intervals of equal~~ 30 bins bin intervals.
18 Habitat depth ~~were~~~~was~~ determined according to the classification of oceanic
layers used in Costello et al. 2010 – (i.e. epipelagic = 0 - 200 m, mesopelagic
= 200 - 1,000 m, and bathypelagic = 1,000 - 4,000 m). A pie chart is used to
21 show how cultural values are represented in the database.

All data and scripts are available (see Appendix A).

3. Results

24 3.1. Records by Bioregions

Approximately 1.14% of the total ~~published occurrences in reported occurrences~~
of the order Actinopterygii were retained in our analysis. That is, from the
27 71,670,596 ~~downloaded records off~~ records downloaded from the GBIF and
OBIS repositories, 820,004 were considered useful (see Appendix A). This

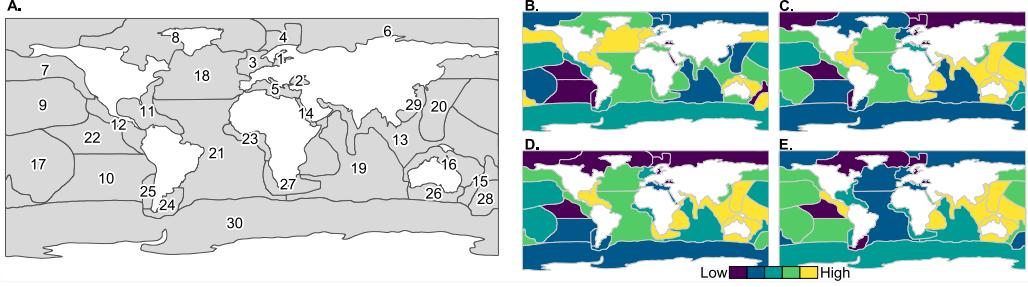


Fig. 1: Marine bioregions and spatial diversity distribution used in this study. **A.** The 30 marine bioregions from Costello et al. (2017) used in this study. Number are identification labels in Table 1. **B.** Records by bioregion; **C.** Overall species richness across bioregions; **D.** Family richness; and **E.** Shannon diversity index. Note that values in **C-E** have been normalized standardized for display illustration purposes. See Table 1 for actual values and a detailed map of observed and expected richness in Fig. A.2.

subset consisted of 10,371 species in 361 families. The most represented families in our dataset are Scombridae, Pleuronectidae, and Gadidae with 103,762, 57,018, and 52,079 records, respectively. The species with the largest representation frequency are *Hippoglossoides platessoides*, *Mola mola*, and *Coryphaena hippurus* with 30,885, 21,042 and 21,089 records, respectively.

The analysis at bioregion bioregions' level (Table 1) shows a large variability. The count of records varies Record counts vary across three orders of magnitudes, that is i.e., from 2.68×10^5 records in the Caribbean Sea and the Gulf of Mexico (11), down to 1.02×10^2 in the Black Sea (2). The bioregion bioregions with the largest species richness and diversity index is the are the Indo-Pacific Seas and the Indian Ocean (13), with 2.95×10^3 recorded species and a Shannon index of 6.93, followed by the Coral Sea bioregion (16), with 2.93×10^3 species and a Shannon index of 6.75. Likewise, the Coral Sea also presents the largest number of families. It is interesting to note that, while being Notably the Southern Ocean (30) is the largest bioregion (i.e. in km^2), the Southern Ocean show the fewest in square kilometers, it has the smallest number of records and the lowest number of species and families across all bioregions. The Black Sea (2) and Norwegian the Norwegian Sea (4) are the bioregions with bioregions have the lowest number of record records and

Table 1: Area (1,000 km²) and counts-of records, species richness, family richness, and Shannon diversity counts for each bioregion. The largest values for highest value in each column is highlighted.

ID	Bioregion	Area	Records	Species	Families	Shannon
1	Inner Baltic Sea	415	8,902	72	30	2.46
2	Black Sea	537	102	37	22	3.21
3	NE Atlantic	2,053	87,377	310	104	3.90
4	Norwegian Sea	1,132	3,046	93	35	2.16
5	Mediterranean	2,859	12,532	372	101	3.39
6	Arctic Seas	10,276	2,506	114	23	3.90
7	North Pacific	12,974	78,070	839	156	4.50
8	North American Boreal	8,001	9,709	162	48	2.99
9	Mid-Tropical N Pacific Ocean	32,685	9,310	615	127	4.59
10	South-East Pacific	21,952	386	190	89	4.97
11	The Caribbean and the Gulf of Mexico	8,427	268,066	1,703	209	4.49
12	Gulf of California	6,184	7,639	885	148	5.93
13	Indo-Pacific Seas and Indian Ocean	37,090	16,967	2,947	215	6.93
14	Gulfs of Aqaba, Aden, Suez, Red Sea	830	926	352	72	5.51
15	Tasman Sea	3,592	1,003	380	120	5.36
16	Coral Sea	7,658	40,107	2,929	249	6.75
17	Mid South Tropical Pacific	23,418	6,083	811	123	5.18
18	Offshore and NW North Atlantic	16,012	130,994	897	190	3.46
19	Offshore Indian Ocean	31,076	1,263	337	116	4.06
20	Offshore W Pacific	10,291	6,363	1,839	232	6.81
21	Offshore S Atlantic	41,435	11,960	990	188	3.79
22	Offshore Mid-E Pacific	13,815	687	79	37	3.04
23	Gulf of Guinea	3,325	6,816	384	138	3.95
24	Argentina	2,665	8,701	115	52	2.83
25	Chile	1,739	250	100	54	4.36
26	Southern Australia	3,824	15,643	1,011	201	5.75
27	Southern Africa	4,371	19,954	1,142	210	4.16
28	New Zealand	6,293	53,879	558	154	3.66
29	North West Pacific	2,457	1,767	869	182	6.46
30	Southern Ocean	62,161	8,996	294	57	3.98

Shannon index value, respectively. Fig. 1 illustrates the location of the 30 marine bioregions and their respective richness and diversity values.

3 3.2. Geographic Analysis

Fig.2 shows the cell classification according to SRI (§2.2.1). As expected, no bioregion is completely sampled at the $\approx 1^\circ$ resolution. In fact, at this resolution, large empty regions with no records are observed. The bioregions with the largest area classified as Adequate/high representativeness are the Northeast Atlantic (3) (37.53%), the Caribbean and the Gulf of Mexico (11) (29.26%), and the Inland Baltic Sea (1) (24.37%). It should be noted that such cells are mostly correspond from mostly correspond to coastal areas in the northern hemisphere. On the other hand, the bioregions that present a greater surface without records correspond to the largest surface area without records are the Southeast Pacific (10) (96.3%), the Arctic Sea (6) (94.9%), and the Southern Ocean (30) (93.7%). While the bioregions with the larger surface with sufficient largest surface area and medium representativeness of records are the Gulf of Guinea (23) (32%), the Norwegian Sea (4) (22.3%), and the Gulf of California (12) (21.6%). Additional results for $5^\circ \times \approx 5^\circ$ and $\approx 10^\circ \times 10^\circ$ spatial resolution grids are available shown in Appendix C.

3.3. Temporal Analysis

Bioregions show similar trends of data accumulation across the four decades analyzed here (Fig. 3). While a significant increase is apparent in the time period between 2005 and 2010, such increase is not significant for 14 out of the 30 bioregions. The Caribbean and the Gulf of Mexico (11) is the bioregion with the largest increases in data contribution to the dataset, while the Black Sea (2) is the bioregion with the lowest rate of data contribution shows the lowest data contribution rate in the 40 years span between 1980 and 2020. (See Appendix D for further analysis).

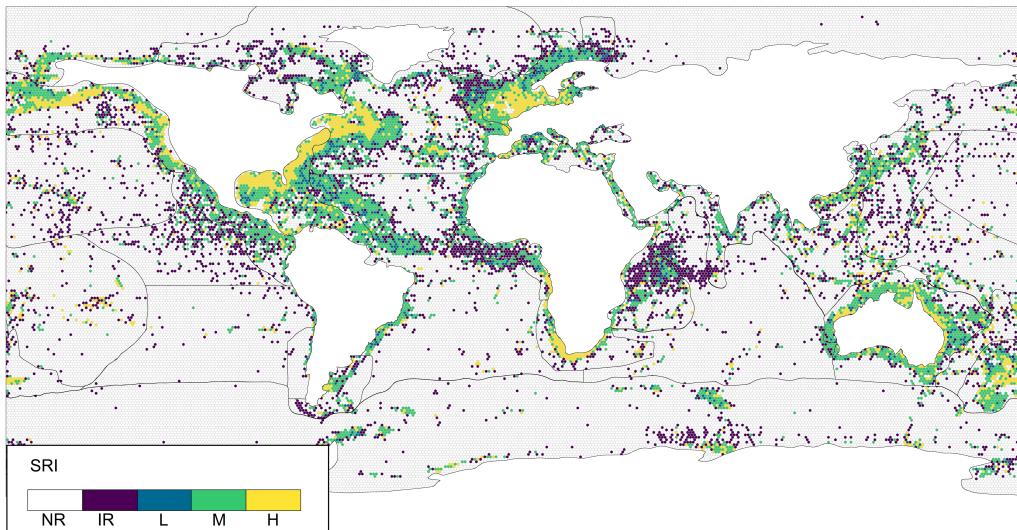


Fig. 2: Spatial Representativeness Index (SRI) in $\sim 1^\circ$ hexagonal lattice. **IR** shows cells with insufficient records to evaluate S_{est} . **A H** are cells with an adequate high representativeness of species richness, i.e. $SRI > 0.85$. **S M** are cells considered as having a sufficient medium representativeness, i.e. $SRI \in (0.60, 0.85)$. **F L** cells are cells with few low representativeness of species records and are thus not considered to be representative of actual species richness, i.e. $SRI \in (0, 0.6)$. **NR** are cells with no records ($SRI = NA = \tilde{NA}$). Raw values for SRI, S_{obs} and S_{est} are shown in the appendix (Fig. A.2).

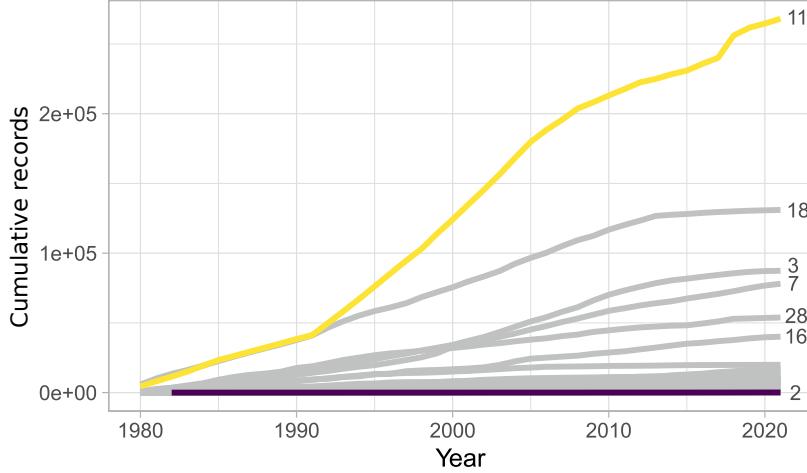


Fig. 3: Records of accumulation rate for each bioregion across the four decades analyzed. The blue-yellow line is the accumulation of fish records in the Caribbean and the Gulf of Mexico bioregion (11) and while the red-purple line shows is the accumulation rate in the Black Sea (2). Numbers as The numbers at the end of each timeseries time series correspond to the bioregion ID in Table 1.

We categorized classified the slopes of the final 10% of each accumulation curves curve in Fig. 4. Fourteen bioregions show a slope less than 1. The 3 Mediterranean Sea (5) stands out with the lowest slope value (0.47), while the Black Sea (2) is the bioregion with the steepest final slope (3.13).

3.4. Gap analysis Analysis and fishing exploitation areas Fishing Exploitation Areas

The bioregions with the largest area covered by protected areas are the Coral Sea (16), the northeast Northeast Atlantic (3) and New Zealand (28) covering 9 a. covering 37.3%, 17.4%, and 16% of their respective surface areas. Regarding the sampling level of these bioregions these bioregions' sampling level, the Offshore Indian Ocean (19), the Gulf of Aqaba, Aden, Suez, Red Sea (14) 12 ; and Coral Sea (16) are the bioregions with the highest percentages of cell sampled as Adequate inside of their share of cells with high representativeness, hence well sampled within protected areas (83%, 63.8%, and 59.8% respec-

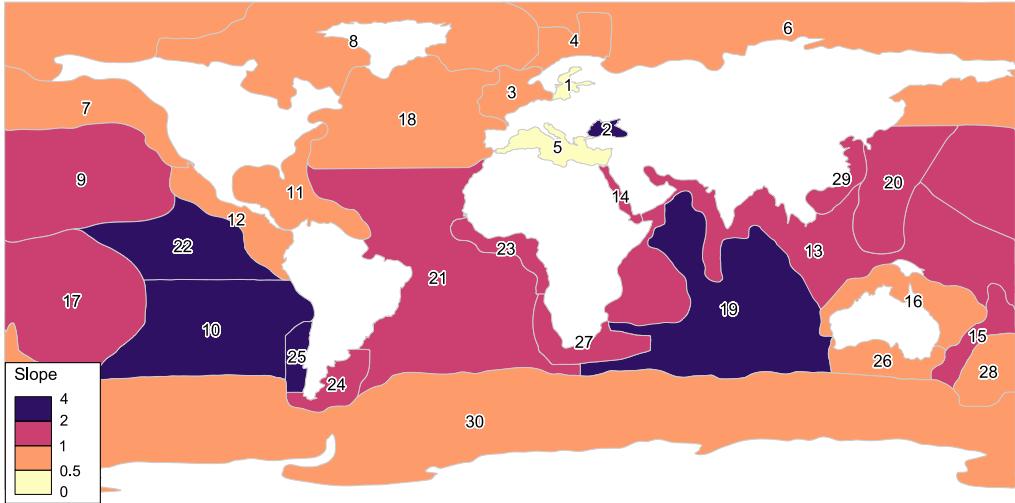


Fig. 4: Graphical representation—Illustration of the slope values of the species accumulation curve for each bioregion. The slope corresponds to the final 10% of the species accumulation curve. See §2.2.2 for details regarding the analysis.

tively). While In turn, the Arctic Seas (6), North American boreal—the North American Boreal (8) and Mid-South—Mid-South Tropical Pacific are 3 the bioregions with protected areas with the highest percentage of cell with no showing the highest share of cells without records (86.2%, 83.8%, and 81.2%). respectively). (See Appendix E).

6 The FAO areas with the largest area categorized as surface area classified as Adequate/high representativeness correspond to the northwest—Northwest Atlantic (22.1 %), Northeastern part of the Pacific Ocean (14.6 %), and 9 Western part of the Atlantic Ocean (12.6 %) (Table 3). These FAO areas correspond to regions of in the Pacific Ocean (North Pacific, North West Pacific, Mid-tropical—Mid-Tropical N Pacific Ocean and Indo-Pacific seas—Seas 12 and Indian Ocean, as well as the Gulf of California and Caribbean and the Caribbean and the Gulf of Mexico). Largest FAO areas with NR cells correspond to the Antarctic part of the Pacific Ocean, the Antarctic part of 15 the Atlantic Ocean and the Southeastern part of the Atlantic Ocean in the Southern Ocean, Offshore S Atlantic, and Southern Africa.

Table 2: Results of overlapping ~~Marine Protected Areas~~MPAs and SRI grid. ID is the identification number given to each bioregion (see Table 1 for ~~bioregion~~bioregions’ names). Area corresponds to the ~~percentage share~~ of surface area covered by ~~marine protected areas~~MPAs. ~~NR~~NR is the ~~percentage share~~ of cells with *No Records*; ~~IR~~IR is the ~~percentage share~~ of cells with *Insufficient records*Insufficient Records; ~~FL~~FL is the ~~percentage share~~ of classified cells with *Few low number of* records; ~~SM~~SM, the ~~percentage share~~ of classified cells with *Sufficient medium number of* records, and ~~AH~~AH, the ~~percentage share~~ of classified cells with *Adequate high number of* records. The highest values for each column ~~is~~are highlighted.

ID	Area <u>km²</u>	NR	IR	FL <ins>FL</ins> %	SM <ins>SM</ins>	AH <ins>AH</ins>
1	0.03	2.38	4.30	5.22	49.08	39.01
2	12.89	26.71	27.61	10.49	35.19	0.00
3	9.74	3.23	1.35	0.94	40.86	53.62
4	0.15	11.16	19.63	5.18	56.69	7.34
5	0.09	5.71	6.77	11.20	47.95	28.37
6	5.01	86.16	5.97	0.02	4.65	3.20
7	0.00	26.48	4.24	1.26	23.77	44.25
8	1.23	83.82	7.77	0.62	6.70	1.11
9	0.69	69.58	15.77	0.00	6.40	8.25
10	17.36	73.51	24.58	0.00	0.80	1.11
11	0.28	20.13	6.25	3.44	29.98	40.21
12	0.83	0.33	1.23	8.85	61.35	28.25
13	0.45	50.52	11.94	0.88	25.17	11.50
14	0.25	8.87	0.00	1.59	25.65	63.88
15	4.06	57.18	15.27	0.00	7.29	20.26
16	16.00	3.10	0.49	1.10	35.52	59.79
17	0.20	81.19	11.43	0.00	3.07	4.31
18	4.91	35.87	16.63	0.77	25.64	21.09
19	2.78	11.88	0.74	0.00	4.34	83.04
20	2.56	34.06	9.68	6.85	35.31	14.10
21	3.79	51.06	19.35	0.38	21.35	7.86
22	0.16	41.98	27.54	0.00	23.22	7.26
23	13.83	9.92	4.92	7.98	61.33	15.85
24	0.21	40.86	20.64	0.24	23.97	14.29
25	0.07	65.27	0.50	0.05	1.29	32.89
26	0.28	30.04	12.83	6.89	38.62	11.63
27	1.45	16.78	5.75	0.15	18.71	58.62
28	0.00	45.36	2.25	0.00	13.71	38.67
29	37.29	20.49	17.05	2.68	42.50	17.29
30	1.66	64.05	7.12	4.89	19.86	4.08

Table 3: Results of overlapping FAO fishery exploitation areas and SRI grid. The surface area corresponding to each bioregion, and the percentage share of surface area of each classification. Area is Areas are in thousands of square km²; NR is the percentage share of cells with No Records; IR is the percentage share of cell cells with Insufficient Record; F-L is the percentage share of classified cells with Few low number of records; S-M is the percentage share of classified cells with Sufficient medium number of records, and A-H is the percentage share of classified cells with Adequate a high number of records. The highest values for each column are highlighted.

FAO Area Name	Area km ²	NR	IR	L %	M	H
Arctic Sea	4,086	93.22	3.13	0.29	2.61	0.75
Northwestern part of the Atlantic Ocean	874	31.19	11.66	5.69	29.37	22.08
Northeastern part of the Atlantic Ocean	3,223	66.29	12.54	2.55	13.63	4.99
Western part of the Atlantic Ocean	1,285	30.84	13.09	7.91	35.60	12.55
Eastern Central part of the Atlantic Ocean	1,208	52.61	24.09	3.44	18.37	1.19
Mediterranean Sea and the Black Sea	309	46.39	15.43	5.24	24.77	8.17
Southwestern part of the Atlantic Ocean	1,731	82.49	5.85	1.69	8.55	1.42
Southeastern part of the Atlantic Ocean	1,765	89.92	4.19	0.15	2.13	3.61
Antarctic part of the Atlantic Ocean	2,310	93.31	2.80	0.20	2.93	0.76
Western part of the Indian Ocean	2,621	72.45	16.11	1.03	8.51	1.89
Eastern part of the Indian Ocean	3,029	85.40	4.69	0.82	7.39	1.70
Antarctic and South of the Indian Ocean	1,977,29	85.71	7.76	0.56	4.33	1.64
Northwestern part of the Pacific Ocean	2,259	73.55	12.40	0.94	10.32	2.79
Northeastern part of the Pacific Ocean	968	55.13	12.65	1.34	16.26	14.62
Western Central part of the Pacific Ocean	2,963	70.45	12.56	0.43	11.58	4.98
Eastern Central part of the Pacific Ocean	4,141	79.36	11.30	0.31	6.94	2.09
Southwestern part of the Pacific Ocean	3,097	85.04	4.42	0.97	6.40	3.17
Southeastern part of the Pacific Ocean	2,997	91.16	6.00	0.10	2.30	0.44
Antarctic part of the Pacific Ocean	2,361	93.47	4.57	0.21	1.42	0.33

3.5. Evaluation of Biases

We evaluated biases for body size, habitat depth, and cultural value for 10,371 marine fish species identified in our database (§3.1).

3.5.1. Body size

The range 10-40 cm range is the most frequently occurring size length, responding to the interval between the 1st and 3rd quartile (Fig. 5A). Three species stand out with the highest numbers of records, *Scomber scombrus*, *Lagodon rhomboides* and *Mallotus villosus* with 20,995, 19,563 and 13,609 records respectively. These species are distributed mainly in the Northeast

Atlantic (3) and Offshore and Northwest North Atlantic (18) bioregions. While In turn, the families that accumulate the greatest number of records 3 correspond to are Sparidae , Scombridae and Labridae with 24,837, 21,719, and 21,035 records, respectively. These families are mainly distributed in the Caribbean Sea and the Gulf of Mexico and (11), and in the Northeast 6 Atlantic (3).

3.5.2. Habitat ~~depth~~Depth

The depth range most commonly observed among records is eentered 9 around about 50 meters and decreases as depth increases, particularly from the epipelagic to the mesopelagic zone zones, as illustrated in Fig.5B. Among the species with the highest number of recorded occurrences, *Mola mola*, 12 *Coryphaena hippurus*, and *Lagodon*-*L. rhomboides* stand out, with 21,089, 21,042, and 19,563 occurrences in the databases, respectively. These species are distributed mainly around the Caribbean Sea and the Gulf of Mexico (11) 15 bioregions, as well as the following bioregions: Offshore and NW North Atlantic (18) and the South Atlantic Coast (21). The families that accumulate a greater larger number of records correspond to Scombridae, Gadidae, Spari- 18 dae with 63,572, 38,876, and 30,041 records, respectively. These are mostly distributed in the northern hemisphere. That ; that is, the Caribbean and the Gulf of Mexico (11), Offshore and NW of the North Atlantic (18) and 21 part of the South Atlantic Ocean Coast (21) bioregions.

3.5.3. Cultural ~~value~~Value

Finally, when analyzing the most frequent cultural value represented 24 across our dataset (Fig. 6), “Commercial” use of the species emerges as the most important with a 73.4% among records, followed by the category “No interest” (5.03%), and “Subsistence fishing” (3.08%).

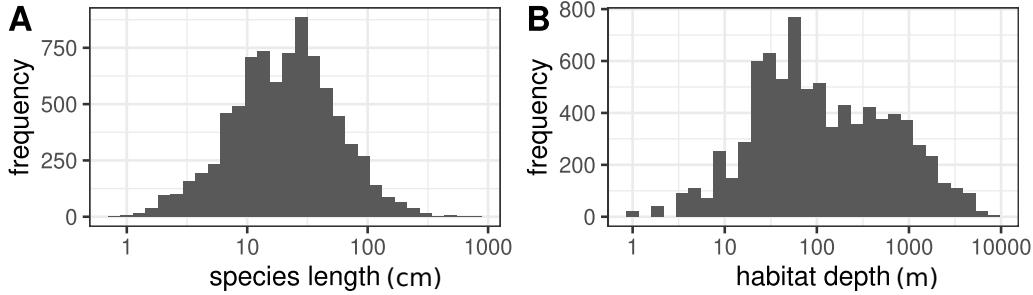


Fig. 5: Distribution of marine fish records in GBIF and OBIS ~~categorized~~ classified by body length and habitat depth. **A**, Relationship between record number and species length (\log_{10}); and **B**, Relationship between record number and habitat depth (\log_{10}) .

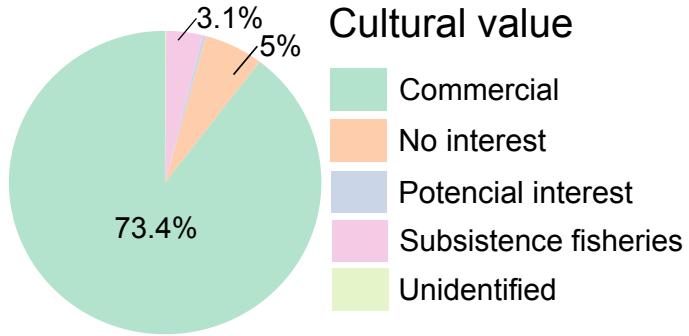


Fig. 6: Frequency of marine fish representation in GBIF and OBIS repositories according to ~~importance of~~ cultural usevalue.

4. Discussion

Our work provides a methodological framework based on a set of non-parametric estimators to quantify the potential number of species from incidence data (Chao et al., 2009). We ~~employed~~ used hexagons due to their suitability as a tessellation that conforms more effectively to the shape of a spheroid compared to square grid cells. We also placed special emphasis on cleaning the occurrence data in their taxonomy (Jin and Yang, 2020), and any potential input errors associated with large and massive datasets (Zizka et al., 2020). ~~This led us to focus on only~~ Hence, we only focused on evaluating marine species in the order Actinopterygii (Alò et al., 2021).

Publicly accessible occurrence records are growing rapidly, partly due to the significant advances significant progresses in ecoinformatics (Lenoir et al., 2020; Oliver et al., 2021). These databases harbor a growing variety of sources, including museum specimens, field observations, acoustic and visual sensors, and citizen science efforts (Amano et al., 2016). However, despite the incredible accumulation of biodiversity records, not all the data is really useful, nor does it represent new insights into the distribution of species (Bayraktarov et al., 2019; Zizka et al., 2020). That is why a systematic evaluation of the integrity and coverage of this information is required (Troia and McManamay, 2017).

There is an extensive bibliography that evaluates the record quality available for different taxonomic groups. Some examples are: legumes on a global scale (Yesson et al., 2007), lepidoptera from Great Britain, and woody plants in Panama (Chao et al., 2009), global marine biodiversity (Tittensor et al., 2010), vascular plants in China (Yang et al., 2013), marine fish on a global scale (Mora et al., 2008; García-Roselló et al., 2015), freshwater fish in the USA (Troia and McManamay, 2017; Pelayo-Villamil et al., 2018), and terrestrial mammals on a global scale (Oliver et al., 2021), among many others. Assuming that that not all data available in these repositories are is useful for biodiversity analyses, several efforts have proposed parametric and non-parametric estimators for data cleaning and species richness analysis, including ModestR (García-Roselló et al., 2013), KnowBr (Lobo et al., 2018), and RWizard (Guisande and Lobo, 2019) among these.

Striving for simplicity, we employ the ratio of observed to expected species richness (SRI) as a means to indicate the spatial distribution of undersampled regions. While acknowledging the potential for misrepresentation, particularly in cases of extremely low observed richness, we mitigate this concern by confining our analysis to locations with more than one observed species record. This approach offers a straightforward method for identifying areas that warrant additional sampling.

We evaluated two additional grid sizes (i.e. 2.5×10^4 and 10^7 km^2), and like other studies, our results show that the coarser the resolution used, the greater the overestimation is, in terms of area. That is, the richness index will indicate that a large area is, indeed, well sampled when in reality, the occurrence records could in fact be localized in a very small area. On the contrary, the finer the scale of analysis, the more localized and deficient the sampling is (Tittensor et al., 2010; García-Roselló et al., 2015; Meyer et al., 2015; Troia et al., 2010; García-Roselló et al., 2015; Meyer et al., 2015; Troia and McManamay, 2015).

Considering that more than 40 years of data were analysed, our results demonstrated that on a global scale, the primary marine fish data available on the GBIF and OBIS platforms are still far from being representative and complete. Compared with other studies evaluating the same taxonomic group (Mora et al., 2008; García-Roselló et al., 2015), although we obtained similar macroecological patterns, only 1.14% of the records extracted from both repositories were useful for our analyses. The large percentage of the A large share of occurrences presented input errors or did not have the necessary data to generate a reliable analysis lacked the data required to develop reliable analyses (Yesson et al., 2007; García-Roselló et al., 2014).

We also found evidence of strong information biases in the records explored. On the one hand, when analyzing the families and species with the greatest representation, they coincide with groups of fish of commercial interest, demonstrating match groups of commercial interest fish, pointing to the existence of taxonomic bias of the data data taxonomic biases (Melo-Merino et al., 2020). This is the case of the families Scombridae, Pleuronectidae and Gadidae, which include species of nutritional importance nutritionally-relevant species such as tuna, cod, haddock, among others (Cohen et al., 1990). The same is true for the species with the largest number of records, *H. platessoides* (Pleuronectidae), *C. hippurus* (Coryphaenidae), and *M. mola* (Molidae); while the first two are species exploited by the

fishing industry, ~~with the exception of~~ sunfish (*M. mola*) ~~which~~ has a wide distribution and is mostly associated with scientific and recreational ~~interest~~
3 ~~interests~~ (Pope et al., 2010).

The unequal contribution of data at the spatial level is another factor that must be considered ~~to work when dealing~~ with data available ~~on in~~
6 ecoinformatic platforms. ~~There is a clear preference for~~ We show a clear
~~geographic bias in the sampling of~~ certain regions and/or ecosystems~~as a~~
9 ~~result of geographical bias~~. The literature indicates that the ~~highest data~~
contribution rates correspond to largest data contributions come from developed countries (Yesson et al., 2007; Chandler et al., 2017), and those coastal regions with ~~better high~~ road connectivity (Chandler et al., 2017;
12 Melo-Merino et al., 2020). This ~~information uncertainty~~ is also particularly prevalent in under-sampled marine habitats, such as the deep sea (Webb et al., 2010). Our results ~~coincide with what is~~ match what has been described in the literature, regardless of the ~~size of the grid that was used to generate the analysis, the~~ grid size used for the analysis. The bioregions that include the ~~Atlantic Northeast Atlantic~~ (3), the Caribbean and the Gulf of Mexico (11), and the ~~Baltic Sea are the regions with the highest number of area sampled as Adequate~~ associated mainly with coastal areas ~~Inland Baltic Sea~~ (1) are regions classified with a high representativeness. However, the number of cells with insufficient ~~data to generate a records to generate an unbiased diversity analysis, is also worrisome. For instance~~ is also of concern.
15 ~~For example~~, our results show that these cells are distributed in more internal areas of the bioregions, zones where sampling is likely to be more difficult.
18 While
21 ~~While, on the other hand,~~ the bioregions that include the South and Southeast Pacific (including the southern coast of South America), the Southern Ocean, and the Arctic ~~Seas Sea~~ are the regions ~~with the least spatial representativeness of records, the proportion where the share~~ of cells without records (*NR*) exceeds 90%. The ~~absencee lack~~ of data samples over this
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extensive area renders any endeavor to depict species richness and distribution highly unreliable ~~within these bioregions~~ (as noted by Yang et al., 2013; 3 Troia and McManamay, 2017). These marine regions encompassing both the water column and the seabed beyond ~~the territorial jurisdiction of countries constitute national jurisdictions make up~~ nearly half of the Earth's surface 6 and sustain ~~a~~ substantial abundance and diversity of life, as highlighted by (Visalli et al., 2020). Nonetheless, when scrutinizing the occurrence data for marine ichthyofauna, these regions remain the least sampled areas.

9 Finally, the ~~time bias of the data~~ ~~data's time bias~~ is also present in our study. Differences in species identification and sampling methodologies over the decades have resulted in ~~the production of~~ databases of variable quality. 12 However, the current era is characterized by more accurate data thanks to improvements in individual capture and identification tools (Costello et al., 2015; Jin and Yang, 2020). For these reasons, our approach considers occurrence records since 1980, ~~however, ;~~ the coverage of occurrence data, ~~however,~~ is uneven over time when comparing ~~between~~ marine bioregions. Despite evaluating more than ~~assesing~~ four decades of data, ~~still sampling efforts are~~ 15 ~~still insufficient in~~ 46% of marine bioregions ~~have insufficient sampling efforts~~. Not surprisingly, the Caribbean and ~~the~~ Gulf of Mexico (11) ~~bioregion is the region is the bioregion~~ with the largest ~~input of data~~, demonstrating once again that ~~data input, once again showing that the~~ geographic sampling bias has strong effects ~~impacts~~ on spatial predictions of species richness (Yang 21 et al., 2013). Future sampling efforts should focus on bioregions at low or equatorial latitudes, areas where ~~biogeographic studies show that~~ marine biodiversity is concentrated according to biogeographic studies (Costello et al., 24 2017).

27 All the biases that we have described, added to ~~the inherent problems in data capture, foster and deepen various typical data capture issues, promote and deepen several~~ information gaps that affect ~~thwart~~ the effective spatio-temporal ~~quantification of biodiversity~~ biodiversity quantification (Magur- 30

ran and McGill, 2011). In this study, we have overlapped our estimates of species richness with the global marine protected areas declared up to MPAs declared up until the beginning of the year 2022 (UNEP-WCMC and IUCN, 2022), and the areas of fishing exploitation reported by the fishing exploitation areas reported by FAO (FAO, 2014).

This exercise demonstrates the importance of public databases that can faithfully reflect the taxonomic and biogeographical knowledge available for each region of the world (Pelayo-Villamil et al., 2018). Our results indicate that (Pelayo-Villamil et al., 2018). According to our results, the North West Pacific bioregion (19) has the largest area covered by marine protected areas MPAs. However, its percentage of adequately sampled cells share of cells with high representativeness is low compared to other bioregions. This latter result is of certain concern as this bioregion is considered a conservation hotspot among other bioregions such as the Coral Sea (16), a bioregion region with a relatively large percentage of adequately share of highly sampled cells (Ramírez et al., 2017). However, we found a low proportion share of well-sampled cells in both regions, demonstrating pointing to the existence of important information gaps, at least for fish of the order Actinopterygii. We emphasize the need to correct these information gaps so that conservation efforts that seek the implementation of new marine protection areas can have reliable datasets as not to underestimate the biodiversity of species can rely on dependable data, including the design and implementation of new MPAs (Sala et al., 2021).

In the same way Along these lines, by overlapping the bioregions with the fishing exploitation zones, we determined that the North Pacific (7), the North West Pacific (29), Mid-tropical the Mid-Tropical N Pacific Ocean (9) and, and the Indo-Pacific seas Seas and Indian Ocean (13) bioregions, as well the Gulf of California (21) and Caribbean and, and the Caribbean and the Gulf of Mexico (11), are the regions with the highest representation of the data data representation and where fishing activity is concentrated.

According to (Kroodsma et al., 2018), the area corresponding to the central Atlantic and Northeast Pacific present little intense fishing ~~effort~~efforts, while
3 the regions associated with the Northeast Atlantic, the Northeast Atlantic (Europe) regions, and the Northwest Pacific are known to have ~~a~~ huge fishing development~~and that is~~, where fishing efforts are concentrated worldwide.
6 The ~~southeastern~~Southeastern Atlantic Ocean (FAO area 47 and 88), part of the Pacific Ocean (FAO area 88) and Antarctica (FAO area 48 and 88) are the regions with the highest ~~percentage~~share of cells without records
9 ($NR = >93\%$). When compared with the findings ~~of~~(Kroodsma et al., 2018) by Kroodsma et al. 2018, these areas ~~agree with~~match the “holes” without fishing effort data, which is explained by the geographical remoteness and the
12 lack of technological development ~~necessary for the required for~~ fisheries to extend to new domains (Visalli et al., 2020). This ~~limits both the exploitation issue restricts both the extraction~~ of marine resources ~~and the collection of~~
15 ~~data as well as data collection.~~

The research questions addressed in this study were essential ~~for comprehending the prevailing trends in data collection and laying to understand the prevailing~~
18 ~~data collection trends and to lay~~ the groundwork for potential corrective measures ~~to than can~~ mitigate the described biases. Our initial inquiry regarding fish body size does not imply a straightforward association between
21 larger records and larger body lengths. Instead, we observe a distinct hump-shaped distribution in ~~the frequency distribution~~frequency distributions, akin to well-documented macroecological patterns observed in various taxa (Smith
24 et al., 2014; Allen et al., 2006). It is worth noting that mid-sized fish species account for the highest number of records. Among these, species such as *S. scombrus* (Scombridae), *L. rhomboides* (Sparidae), and *M. villosus* (Osmeridae) stand out for their numerous records, ~~and~~; they are predominantly distributed in well-sampled regions such as the Mediterranean Sea ~~-(5)~~, the Caribbean and the Gulf of Mexico ~~and the Caribbean~~(11), and the Atlantic
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28
29
30 Ocean ~~(e.g. bioregion 3)~~. Furthermore, the inverse relationship between fish

size and abundance, and consequently, the frequency of human ~~utilization~~use, whether for scientific research or commercial purposes, is a well-established
3 concept (Pauly and Palomares, 2005).

This variation in sampling ~~effort~~efforts results in a noticeable overrepresentation of these species, exacerbating the existing ~~taxonomic bias~~taxonomic bias. Conversely, the correlation between the number of records and habitat depth indicates that the pelagic zone ~~exhibits a significant concentration of data~~shows a significant data concentration, which appears to align with areas more readily accessible for data collection (Melo-Merino et al., 2020). It has been pointed out that ~~the concentration of species~~species concentration decreases as the ~~depth of the ocean increases~~, ocean increases its depth; however, it is precisely these areas that have been ~~the~~ least sampled and where there is ~~the greatest probability~~a larger chance of discovering new species (Costello et al., 2017). This demonstrates the need to concentrate efforts on the deeper regions of the water column (mesopelagic, bathyal, and abyssal) for a more equitable representation of marine ecosystems. Finally, a straightforward examination of cultural value ~~among~~within marine records unmistakably reveals that ~~species of marine fish~~marine fish species with more favorable or ~~economically advantageous utility to~~economic advantages for humans tend to have stronger ~~representation within the analyzed databases~~representations within the databases discussed. This observation is likely connected to the significant role of the fishing industry as one of the primary sources of information contributing to platforms such as OBIS, as previously discussed OBIS (Zhang and Grassle, 2002).

Today, marine ecosystems and their biodiversity face the ~~great challenges of climate change and the impact~~major climate change challenge as well as the impacts of human activity, especially ~~those on~~species considered key food resources for survival (Hollowed et al., 2013; Ramírez et al., 2017; O'Hara et al., 2021). It is ~~necessary to focus and strengthen~~important to focus on and further the study of ~~those areas with very~~areas with few or no records,

since the descriptions of the geographic ranges of the species describing the species geographic ranges and their temporal dynamics are fundamental measures is a key measure for the evaluation of the real state of biodiversity (Lenoir et al., 2020; Oliver et al., 2021). Having actual biodiversity state (Lenoir et al., 2020; Oliver et al., 2021) Counting on more reliable data will allow for the implementation of effective conservation actions to be implemented.

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18

Appendix A. The database

Table A.1 below shows the data loss for each criterion that we have used
³ to clean our database. We downloaded 71,670,596 records from GBIF and OBIS. Only 820,004 records were useful for our analyses.

Database state	Number of records
Original records from GBIF and OBIS	71,670,596
Data curation (following Zizka et al. (2020))	5,380,439
Taxonomically filtered data	5,007,322
Deletion of data outside the native range	820,004

Table A.1: Criteria for filtering occurrence data from GBIF and OBIS using bioregions.

Files of the 10,371 marine fish species and their attributes (body size,
⁶ habitat depth, and cultural value) from FishBase may be found in the GitHub project page of this manuscript: http://github.com/vapizarro/stp_fishes

Appendix B. Species Representativeness Analysis (SRI)

⁹ For each cell (i), the SRI is the simple ratio between the observed number of species S_{obs} and the expected number of species (S_{exp}): $SRI_i = S_{obs}/S_{exp}$. Maps for the smaller resolution analyzed ($\sim 1^\circ \times 1^\circ \sim 1^\circ$) are in Fig. A.2.

12 Appendix C. Grids resolutions

For spatial representation analysis we evaluated two additional spatial resolutions ($5^\circ \times 5^\circ \sim 5^\circ = 3,021$ cells, and $10^\circ \times 10^\circ \sim 10^\circ = 958$ cells).
¹⁵ Table C.2 contains the results of this analysis for these grids. We have also mapped these results (see Figure A.3), to understand how the effect of spatial resolution on the evaluation of biodiversity macropatterns. Finally, we also
¹⁸ plot the frequency of cells for each SRI category for the three grid sizes (R1= $1^\circ \times 1^\circ \sim 1^\circ$; R5= $5^\circ \times 5^\circ \sim 5^\circ$; R10= $10^\circ \times 10^\circ \sim 10^\circ$) to understand how the data is distributed in our analyses (see Figure A.4)

ID	R1 ($\sim 1^\circ$)					R5 ($\sim 5^\circ$)					R10 ($\sim 10^\circ$)				
	NR	IR	L	M	H	NR	IR	L	M	H	NR	IR	L	M	H
1	18.49	15.13	5.04	36.97	24.37	0.00	16.67	0.00	33.33	50.00	16.67	16.67	0.00	33.33	33.33
2	68.75	19.79	1.04	10.42	0.00	10.00	40.00	10.00	30.00	10.00	40.00	0.00	0.00	40.00	20.00
3	15.74	6.54	3.39	36.80	37.53	3.57	3.37	0.00	17.86	75.00	0.00	0.00	0.00	10.00	90.00
4	46.35	22.34	7.93	22.13	1.25	28.13	9.38	6.25	40.63	15.63	30.77	15.38	0.00	23.08	30.77
5	42.39	14.75	4.92	27.87	10.07	14.29	3.57	0.00	32.14	50.00	16.67	8.33	0.00	8.33	66.67
6	94.96	2.21	0.13	1.87	0.83	82.13	5.64	0.31	6.58	5.33	62.65	13.25	1.20	10.84	12.05
7	63.24	11.24	0.87	14.46	10.19	17.09	9.40	3.42	29.06	41.03	7.69	7.69	2.56	35.90	46.15
8	79.52	11.27	0.89	7.17	1.15	43.93	11.56	4.05	32.37	8.09	32.69	11.54	3.85	40.38	11.54
9	88.74	8.71	0.00	1.57	0.99	28.74	22.99	2.87	38.51	6.90	7.69	9.62	21.15	38.08	13.46
10	96.31	2.41	0.04	0.88	0.36	70.87	15.75	0.79	7.09	5.51	51.28	15.38	2.56	20.51	10.26
11	23.82	8.42	5.65	32.85	29.26	8.62	0.00	0.00	18.97	72.41	0.00	10.53	0.00	5.26	84.21
12	35.59	21.61	2.45	35.59	4.76	14.29	4.76	2.38	47.62	30.95	5.88	11.76	0.00	17.65	64.71
13	67.52	15.80	1.01	12.00	3.67	13.76	12.84	7.34	44.95	21.10	9.46	6.76	2.70	44.59	36.49
14	45.83	10.83	2.50	30.83	10.00	46.15	0.00	0.00	7.69	46.15	25.00	0.00	0.00	0.00	75.00
15	74.52	13.06	0.00	7.07	5.35	20.00	6.67	6.67	40.00	26.67	37.50	12.50	0.00	37.50	12.50
16	36.68	10.95	3.84	34.65	13.88	5.77	7.69	3.85	28.85	53.85	10.53	0.00	0.00	21.05	68.42
17	91.36	4.90	0.00	1.57	2.17	47.93	19.01	0.00	20.66	12.40	25.00	8.33	0.00	36.11	30.56
18	48.29	16.27	3.78	22.06	9.61	6.50	7.32	7.32	43.09	35.77	10.26	5.13	0.00	28.21	56.41
19	90.40	6.93	0.06	2.27	0.35	53.45	18.39	3.45	17.82	6.90	31.48	12.96	1.85	35.19	18.52
20	63.61	17.35	1.43	13.56	4.04	8.20	8.20	9.84	44.26	29.51	15.00	5.00	0.00	45.00	35.00
21	74.78	9.63	2.84	11.48	1.27	34.68	13.51	3.15	28.38	20.27	21.21	9.09	0.00	27.27	42.42
22	76.12	18.00	0.00	5.10	0.78	33.33	6.17	16.05	43.21	1.23	9.09	4.55	9.09	59.09	18.18
23	34.65	32.02	2.89	24.41	6.04	25.93	0.00	14.81	51.85	7.41	0.00	18.18	0.00	36.36	45.45
24	63.07	17.89	1.38	13.53	4.13	25.93	7.41	3.70	51.85	11.11	20.00	10.00	0.00	50.00	20.00
25	88.02	4.96	0.83	5.37	0.83	52.63	10.53	0.00	21.05	15.79	42.86	0.00	0.00	28.57	28.57
26	60.93	7.04	2.41	22.41	7.22	27.27	12.12	0.00	30.30	30.30	16.67	0.00	0.00	33.33	50.00
27	66.84	10.35	0.70	8.07	14.04	27.78	16.67	0.00	22.22	33.33	7.69	7.69	0.00	23.08	61.54
28	59.84	9.17	2.13	17.67	11.19	30.19	7.55	1.89	30.19	30.19	30.00	10.00	0.00	25.00	35.00
29	41.96	19.87	2.84	26.81	8.52	15.00	10.00	5.00	40.00	30.00	12.50	12.50	0.00	37.50	37.50
30	93.74	3.49	0.20	1.97	0.59	69.45	11.02	1.00	10.52	8.01	42.29	16.57	2.86	22.29	16.00

Table C.2: Surface area as a percentage share of each bioregion (ID) for every SRI category for each of the three grid sizes (R1= $1^\circ \times 1^\circ \sim 1^\circ$; R5= $5^\circ \times 5^\circ \sim 5^\circ$; R10= $10^\circ \times 10^\circ \sim 10^\circ$). Values show the surface area as a percentage share of each bioregion for every SRI category (see §2.2.1). ID is the identification number given to each bioregion (Table 1). **A-H** are cells with an adequate-a high representativeness of species richness (i.e. SRI > 0.85). **S** **M** are cells considered as having a sufficient-medium representativeness (i.e. SRI ∈ (0.60, 0.85)). **F-L** cells are cells with few-a low number of records and are thus not considered to be representative of actual species richness (i.e. SRI ∈ (0, 0.6)). **NR-NR** as cells with no records-no records (SRI=NA/NA), and **IR-IR** as cell with insufficient records-insufficient records to apply SRI.

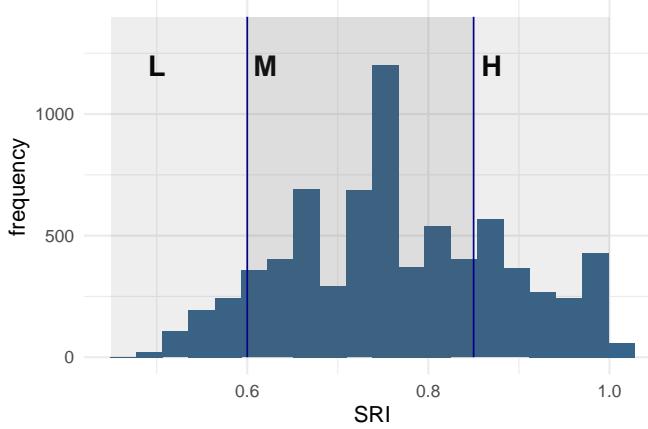


Fig. A.1: Classification of SRI values based on its frequency distribution. This histogram displays the frequency distribution of SRI (Species Richness Index) values and the corresponding class selection thresholds. Cells are categorized as follows: SRI < 0.6 are classified as “Few low” representativeness (L), “SRI falling in the range (0.6, 0.85) as “Sufficient medium” representativeness (M), and SRI > 0.85 as “Adequate high” representativeness (H).

Appendix D. Bioregions slopes

We evaluated the slopes of the last 10% of the accumulation curves of
³ each bioregion in our temporal representation analysis. Table D.3 shows the result for each bioregion.

Appendix E. GAP Analysis

⁶ We plotted the percentage of surface with marine protected areas of share of surface areas with MPAs in each bioregion (Fig A.5), and the percentage share of cells of each FAO Area area for each category of SRI value (Fig A.6).

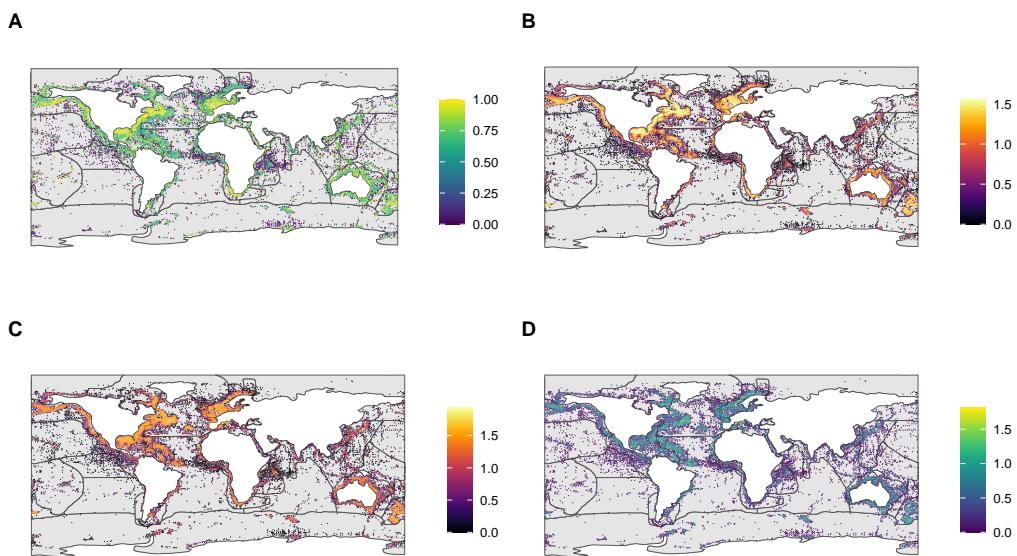


Fig. A.2: SRI and Species richness S depicted from GBIF and OBIS databases. **A.** Species representativeness index; **B.** Observed species richness (S_{obs}); **C.** Expected species richness (S_{exp}); **D.** Difference between raw estimated and observed richness. The difference has been \log_{10} transformed after subtraction.

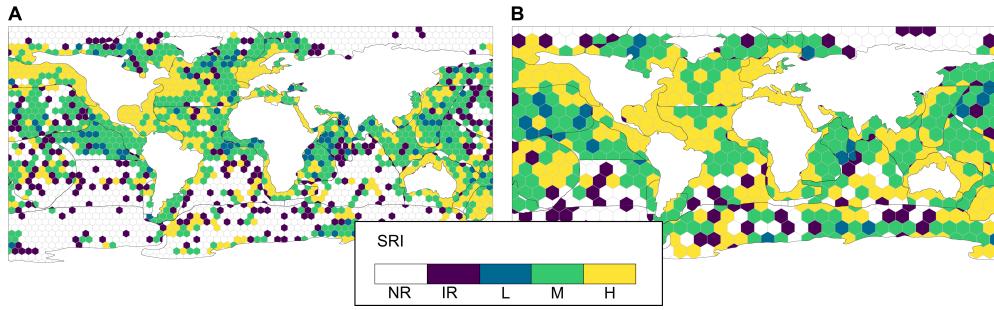


Fig. A.3: Spatial representativeness index (SRI) mapping of cells of size: A= $5^\circ \times 5^\circ \sim 5^\circ$; B= $10^\circ \times 10^\circ \sim 10^\circ$. The categorization of the cells corresponds to the level reached by the SRI, where SRI > 0.85: Amount of data Adequate is high for the representation of species richness (“A”H); SRI=0.60-0.85: Amount of data can be considered Sufficient of medium representativeness (“S”M); SRI=0-0.60: Amount of records Few is low (“F”L); and SRI = NA: cells with no records (“NR”NR). **IR** are cells with insufficient records to evaluate species diversity representativeness.

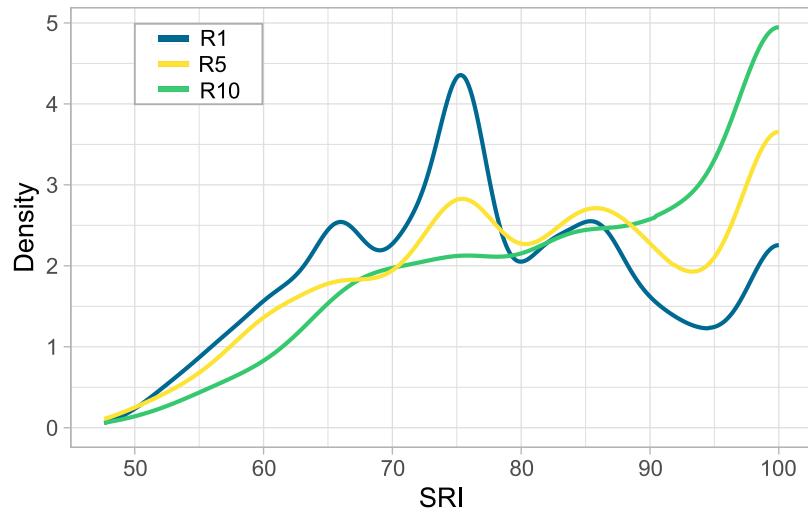


Fig. A.4: Density probability distribution of SRI in three grids of different sizes: R1= $1^\circ \times 1^\circ \sim 1^\circ$ (blue line); R5= $5^\circ \times 5^\circ \sim 5^\circ$ (red line); and R10= $10^\circ \times 10^\circ \sim 10^\circ$ (yellow line).

Bioregion	Slope
1	0.35
2	1.16
3	1.79
4	0.91
5	1.76
6	0.65
7	4.44
8	1.37
9	6.18
10	4.87
11	10.37
12	7.57
13	32.86
14	4.90
15	6.78
16	21.62
17	10.10
18	6.59
19	12.44
20	23.21
21	11.70
22	1.85
23	4.42
24	3.49
25	2.12
26	7.74
27	12.29
28	4.82
29	14.08
30	2.74

Table D.3: Final slope (10%) of the accumulation curves for each bioregion

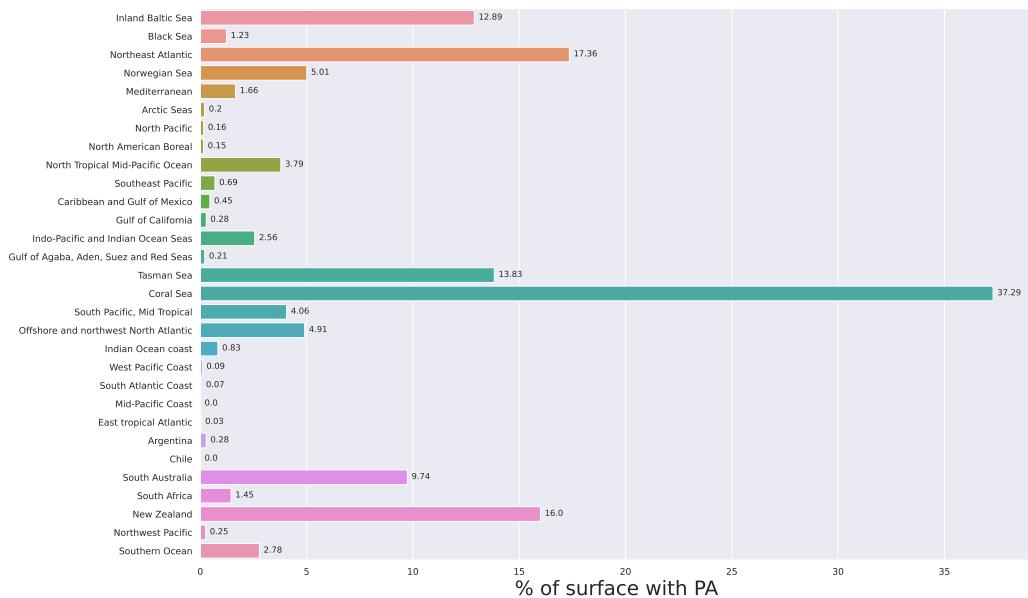


Fig. A.5: Percentage Share of surface area with marine protected areas—MPAs by bioregions.

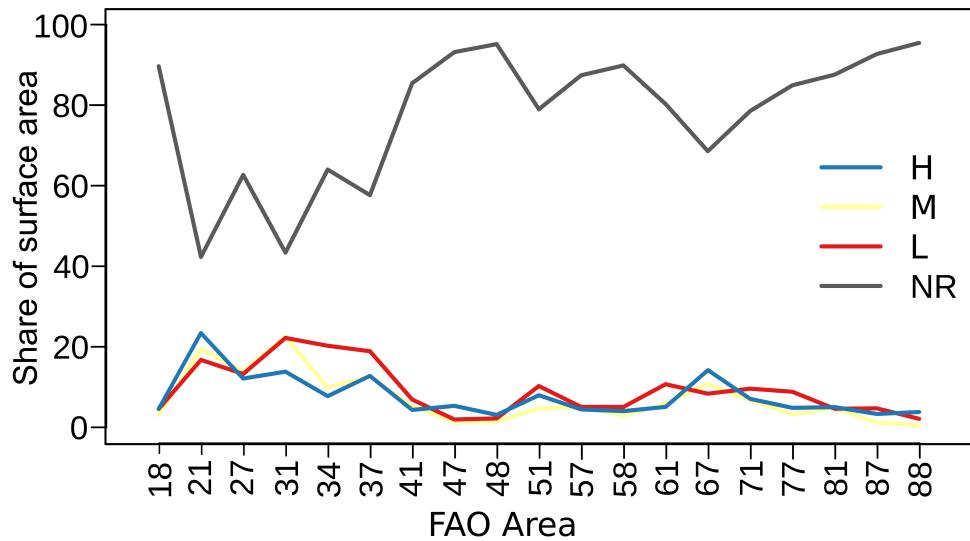


Fig. A.6: Percentage Share of cells of each FAO Area-area for each category of SRI value category. Amount of data Adequate for the representation $SRI = >0.85$: High representativeness of observed species richness ("A" H); $SRI=0.60-0.85$: Amount Medium representativeness of data can be considered Sufficient observed species richness ("S" M); $SRI=0-0.60$: Amount Low representativeness of records Few observed species richness ("F" L); and $SRI= \text{NA}$: cells with no records ("NR" NR).