ARTICLE 2

# Predicting species distributions in the open oceans with convolutional neural networks.

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#### **Abstract**

As biodiversity plummets due to anthropogenic disturbances, the conservation of oceanic species is made harder by limited knowledge of their distributions and migrations. Indeed, tracking species distributions in the open ocean is particularly challenging due to scarce observations, and the complex and variable nature of the ocean system. In this study, we propose a new method that leverages deep learning, specifically convolutional neural networks (CNNs), to capture spatial features of environmental variables. This novelty eliminates the need to predefine these features before modelling and creates opportunities to discover unexpected correlations. Our aim is to present the results of the first trial of this method in the open oceans, discuss limitations, and provide feedback for future improvements or adjustments.

In this case study, we considered 38 taxa which include pelagic fishes, elasmobranchs, marine mammals, as well as marine turtles and birds. We trained a model to make probability predictions from the environmental conditions at any specific point in space and time, using species occurrence data from the Global Biodiversity Information Facility (GBIF) and environmental data from various sources. These variables included sea surface temperature, chlorophyll concentration, salinity, and fifteen others.

During the testing phase, the model was applied to environmental data at locations where species occurrences were recorded. The model accurately predicted the observed taxon as the most likely taxon in 69% of cases and included the observed taxon among the top three most likely predictions in 89% of cases. These findings show the adequacy of deep learning for species distribution modelling in the open ocean and demonstrate the relevance of CNNs for prospective modelling of the impacts of future ocean conditions on oceanic species.

Additionally, this black box model was then analysed with explicability tools to understand which variables had an influence on the model's predictions. While variable importance was species-dependent, we identified finite-size Lyapunov exponents (FSLEs), sea surface temperature, pH, bathymetry and salinity as the most influential variables, in that order. These insights can prove valuable for future species-specific movement ecology studies.

Keywords: deep learning; megafauna; open oceans; pelagic species; species distribution models

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1. Introduction

1.1 Background

The open ocean is a vast and complex ecosystem that covers over 70% of the Earth's surface, yet it remains one of the least understood and studied ecosystems on our planet [1]. It plays a critical role in regulating the Earth's climate, nutrient cycles, and biogeochemical cycles (including carbon sequestration), making it a vital component of all life on Earth [2].

However, the ocean is facing a range of human-induced threats, including over-fishing, pollution, and climate change [3, 4, 5]. These threats can have serious consequences for marine biodiversity, and therefore negatively impact the livelihoods of millions of people who depend on the oceans for their food or income [6].

To focus on solving the most pressing challenges, it is essential to understand how marine life is distributed within open oceans. Species distribution models can provide valuable insights into where different species are likely to be found, what environmental factors are driving their distribution, and the influence of temporal variations [7]. By developing accurate and reliable models, we can identify areas that are most at risk and develop effective conservation strategies to protect these ecosystems.

Furthermore, changes in the Earth's climate are already affecting ocean conditions, namely warming waters, ocean acidification, and sea level rise, among others [8]. This makes it even more urgent to understand the link between environmental variables and species distributions, to be able to predict how marine biodiversity may respond to these changes. This information is critical for informing decision-making and management efforts to ensure the long-term sustainability of marine ecosystems and the services they provide to society.

Therefore, studying species distribution in open oceans is essential for advancing our understanding of these complex ecosystems and for developing effective conservation and management strategies to protect them.

## 1.2 Existing methods for predicting species distributions

There is a wide variety of Species Distribution Models (SDMs) discussed in literature [9]. This is generally done through modelling a species-specific *environmental niche*: the area where environmental conditions are favourable to the species [10]. Predictors are chosen empirically to best fit the observed species' occurrences with specific environmental conditions.

Usually, SDMs use environmental data at the exact location where the prediction is computed, which is insufficient to represent the full nature of the environmental seascape around animals. Even averaging data over a buffer area does not fully represent the environmental conditions, as it cannot convey bathymetry features such as seamounts or trenches. The same applies to other variables, which spatial structure may be more important than punctual or average values: algal blooms, temperature fronts, eddies, etc. Yet these spatial structures are essential to understanding species distributions [11, 12].

A solution to this shortcoming is to include the values of these environmental data in the neighbourhood of species occurrences, but the number of variables then becomes much larger than the number of observations. This is unfit for statistical models and requires a feature extraction step to summarize input data as fewer significant variables. This work may be made manually, which enables the model to take advantage of scientists' expert knowledge. This is how some spatial features are added into SDMs [13], but it limits the performance of the model to the scope of existing knowledge and prevents the discovery of previously unknown influential factors.

Furthermore, SDMs rarely take into account the high temporal variability of environmental data [14], which seriously hinders the prediction of highly mobile species distributions.

This calls for new methods to fully take into account the complex spatial structures of environmental seascapes and their influence on species distributions.

# 1.3 Potential benefits of using deep learning for modelling marine species distribution

Convolutional neural networks (CNNs) were invented for image recognition, so they have embedded feature extractors that are designed to detect multiple levels of details using convolution layers [15]. With image classification, one can identify the following levels, from most precise to coarser:

- 1. Values of specific pixels
- 2. Value of a small group of pixels: textures, edges
- 3. Association of several groups of pixels: shapes, geometric features
- 4. Association of several shapes: objects, animals, plants
- 5. Average and extreme values on the whole image: brightness/tint

This is especially useful with environmental data raster layers (from satellite observations or models) as it enables the model to detect the same various levels of details on environmental variables. Here are some examples of the same levels of detail, applied to environmental variables:

- 1. Values at the point of occurrence
- 2. Homogeneity of the variable in the neighbourhood of occurrence: temperature fronts, slopes
- 3. Geographic features: bays, underwater canyons, river plumes
- 4. Complex shapes: current structures, cyclones
- 5. Average and extreme values over the buffer zones

The use of CNNs to model species distributions was successfully developed for terrestrial plants [16]. The CNN architecture proved especially useful to capture spatial features, as well as to transfer knowledge from better-known species to lesser-known species [17]. While these studies were mostly based on satellite imagery (Sentinel-2), we cannot rely on this information in the open oceans. Another significant difference is the high temporal variability of the oceanic seascape. This is why we present an adaptation of their work that relies only on environmental data.

# 1.4 Objectives of the study

Through the present study, we explore and report the possibilities that deep-learning-based SDMs offer in the open oceans. We first give a detailed overview of the data that was used to build our model. Then we show the results that we obtained, including performance metrics and distribution maps. Finally, we show an example of how this method may be used to simulate a change in future ocean conditions and how probabilities predictions respond. We also point out the limits that we have found with our methodology choices and suggest ways to improve the results' quality in the future.

2. Methods

The main step of our process is to build a model to link environmental data to species presence. To achieve this, we used freely available occurrence data from the Global Biodiversity Information Facility (GBIF) [18], and downloaded environmental data in a buffer around each of their locations, at the date of their occurrence.

After training, this provided us with a model which takes environmental data as input and outputs a vector of observation probabilities (one for each taxon). It is important to note that as training data

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is presence-only, we cannot predict abundance or any absolute measure of presence. The predictions are observation probabilities, relative to the 38 taxa that are the subject of this study. The full process is summarized in Figure 1 and is explained in detail in this section.

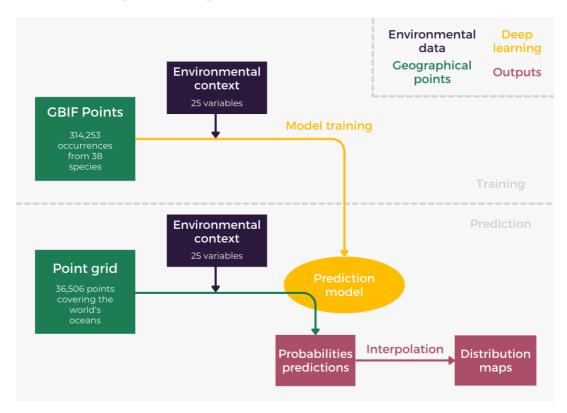


Figure 1. Summary view of the analysis process: model training in the top half and predictions in the bottom half.

## 2.1 Description of the occurrence data

Thirty-eight marine species or genera were selected for the proof of concept that is described in the present article. They include large pelagics, elasmobranchs, turtles, sea mammals, and two species of marine birds (*cf.* table 1). These taxa may be replaced or complemented with others in the future.

Presence data for these taxa were downloaded from the GBIF. This database contains species occurrences that are free to access and download, which is essential to reproducibility. Some flawed data is unavoidably present in the database, but small errors in the geographical coordinates are not a problem as the oceanographic landscape that we consider has limited precision due to environmental data resolution. We removed points located on the continents, and no other filtering was conducted on geographical precision. Furthermore, convolutional neural networks are known to be robust against occasional labelling mistakes [19].

Digital object identifiers (DOIs) for the download of each species are available in table 1. When there were more than 10,000 occurrences of a taxon, a random sample of 10,000 occurrences was selected. In all cases, GBIF identifiers of occurrences that were actually used are available in the training data set CSV files (*id* column).

This added up to 314,253 occurrences for all taxa, depicted in figure 2.

**Table 1.** Species that were included in the study, coloured by taxonomic class. The last column is the digital object identifier (DOI) of downloaded archives.

English name	Taxonomic name	DOI
Yellowfin tuna	Thunnus albacares	10.15468/dl.gr2wbb
Longfin tuna	Thunnus alalunga	10.15468/dl.aqjv3y
Atlantic bluefin tuna	Thunnus thynnus	10.15468/dl.nnyeyb
Southern bluefin tuna	Thunnus maccoyii	10.15468/dl.tw97qj
Bigeye tuna	Thunnus obesus	10.15468/dl.c96qpp
Skipjack tuna	Katsuwonus pelamis	10.15468/dl.6y2zzm
Frigate Tuna	Auxis thazard	10.15468/dl.kfm6kq
Sailfish	Istiophorus	10.15468/dl.f48dug
Black marlin	Istiompax indica	10.15468/dl.b5acky
Blue marlin	Makaira	10.15468/dl.sygtaw
Swordfish	Xiphias gladius	10.15468/dl.hazqd2
Dolphinfish	Coryphaena	10.15468/dl.q67bqt
Humphead wrasse	Cheilinus undulatus	10.15468/dl.9g76hq
Oceanic Whitetip	Carcharhinus longimanus	10.15468/dl.b5ws4q
Whitetip	Carcharhinus albimarginatus	10.15468/dl.vpc772
Silk shark	Carcharhinus falciformis	10.15468/dl.vg4rwh
Sandbar shark	Carcharhinus plumbeus	10.15468/dl.7fczpa
Grey reef shark	Carcharhinus amblyrhynchos	10.15468/dl.ccqyws
Mako shark	Isurus oxyrinchus	10.15468/dl.h5akxk
Blue shark	Prionace glauca	10.15468/dl.zqkssk
Devil ray	Mobula mobular	10.15468/dl.p4e2sx
Reef manta	Mobula alfredi	10.15468/dl.bkjkgu
Eagle ray	Myliobatis	10.15468/dl.3u3v7k
Humpback whale	Megaptera novaeangliae	10.15468/dl.yzg4n3
Fin whale	Balaenoptera physalus	10.15468/dl.r9kaq8
Blue whale	Balaenoptera musculus	10.15468/dl.28f7xd
Bottlenose	Tursiops	10.15468/dl.bec9p4
Spinner dolphin	Stenella longirostris	10.15468/dl.xz5eds
Common dolphin	Delphinus delphis	10.15468/dl.u5be7v
Sperm whale	Physeter macrocephalus	10.15468/dl.7pf4ue
Harbour porpoise	Phocoena phocoena	10.15468/dl.afr2fn
Southern right whale	Eubalaena australis	10.15468/dl.e3hdkj
Green turtle	Chelonia mydas	10.15468/dl.6gs9rp
Loggerhead	Caretta caretta	10.15468/dl.dmb6ds
Hawksbill turtle	Eretmochelys imbricata	10.15468/dl.e6w44w
Emperor penguin	Aptenodytes forsteri	10.15468/dl.s5unhs
Wedge-tailed shearwater	Puffinus pacificus	10.15468/dl.vyztue
Acropora coral	Acropora	10.15468/dl.vg752f

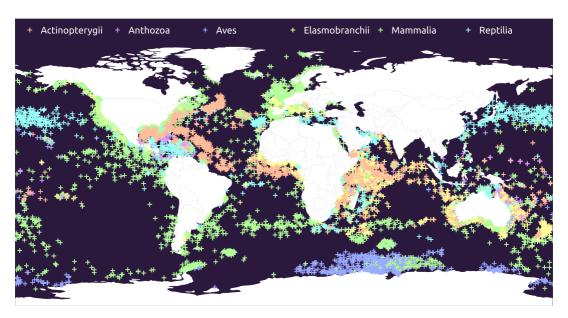


Figure 2. Random sample (10%) of the training data set, coloured by taxonomic class.

## 2.2 Description of the environmental data used as inputs

Eighteen environmental variables were considered, some from satellite observations and others from models. Three of them contain two components: surface wind, geostrophic current and finite-size Lyapunov exponents (FSLEs). Temperature and chlorophyll values were also included 15 and 5 days before the occurrences, as it was previously demonstrated that marine animals may have a delayed response to some variables, especially temperature [20]. Finally, four geographical variables were added (cf. section 2.3.2). This amounts to a total of 29 layers of data, shown in table 2.

# 2.3 Data preparation

#### 2.3.1 Enrichment

Environmental data were downloaded in a buffer around the occurrences using the GeoEnrich python package, which was developed for this purpose and made available to other researchers for a wide range of uses in the GitHub IRDG2OI/geoenrich repository [21]. A spatial buffer of 115km was used, to include at least one data point from the least precise data (2° resolution). This is consistent with values of daily potential movement for fast animals that may travel up to 120 km per day [22, 23].

These data arrays with various resolutions (minimum  $1 \times 1$  for wave height, maximum  $241 \times 241$  for bathymetry) also have various horizontal dimensions due to the longitude contraction closer to the poles. They were all interpolated (up-scaled or down-scaled depending on the initial resolution) to fit the same  $32 \times 32$  grid centred around the occurrence.

#### 2.3.2 Ocean basin and hemisphere

An initial goal of the study was to produce a geography-agnostic model, which means that two points with the same oceanic conditions, wherever they are, should yield the same predictions. But this is ecologically wrong for one main reason: there are natural barriers that animals cannot cross, namely continents and for some species, the warm waters around the equator. Therefore we added four binary variables: three for the main oceanic basins and one for the hemisphere.

The world's oceans were split into three main basins: the Atlantic, Indian and Pacific oceans. Very few of the occurrences were located in the Arctic Ocean: they were assigned the closest of these ocean basins. Occurrences from the Southern Ocean were more numerous and are not separated from these three oceans by any physical barrier, so they could be assigned to the closest one.

It is important to note that the Ocean basin and the hemisphere are the only geographical information provided to the model. This is by design to avoid learning the observation bias that is present in the training data.

Table 2. 29 layers used as input data

Variable	Source	Source type
Bathymetry	GEBCO [24]	Observations
Salinity	Copernicus [25]	Observations
Wave Height	Copernicus [26]	Observations
Surface wind (u)	CCMP [27]	Observations
Surface wind (v)	CCMP [27]	Observations
Oxygen	Copernicus [28, 29]	Models
рН	Copernicus [28, 29]	Models
FSLEs (strength)	Aviso [30]	Observations
FSLEs (orientation)	Aviso [30]	Observations
Geostrophic Current (u)	Copernicus [31, 32]	Observations
Geostrophic Current (v)	Copernicus [31, 32]	Observations
Eddy kinetic energy	Calculated	
Chlorophyll	OCCI [33]	Observations
Sea surface temperature	MUR [34]	Observations
Mixed layer thickness	Copernicus [25]	Observations
Diatoms	Copernicus [35]	Observations
Dinophytes	Copernicus [35]	Observations
Haptophytes	Copernicus [35]	Observations
Green algae	Copernicus [35]	Observations
Prochlorophytes	Copernicus [35]	Observations
Prokaryotes	Copernicus [35]	Observations
Chlorophyll (D-15)	OCCI [33]	Observations
Sea surface temperature (D-15)	MUR [34]	Observations
Sea surface temperature (D-5)	MUR [34]	Observations
Chlorophyll (D-5)	OCCI [33]	Observations
Atlantic Ocean	Calculated	
Indian Ocean	Calculated	
Pacific Ocean	Calculated	
North hemisphere	Calculated	

## 2.3.3 Feature scaling

All data were scaled to the [0,1] interval, and saved into a data cube (32 × 32 geographical pixels ×29 layers). Outliers (highest and lowest 1% of the values of the training data set) were clipped and the scaling factors were saved to be reapplied to any subsequent input data.

There are some missing data because of natural phenomena such as clouds, or because the occurrences were out of the data set time range. In that case, we used the median value of the variable over the tile. If data was missing over the whole tile, we used the median value over the whole data set instead.

Figure 3 shows an example of all the data that are included in the data cube used for training, with the feature scaling reversed in order to show the real values. The figure does not show the four binary geographical variables.

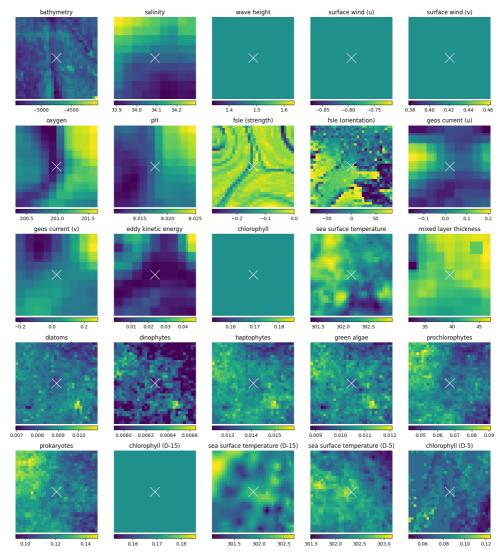


Figure 3. Environmental variables around the point of coordinates -14.389°S, 78.918°E on March 20th, 2021.

# 2.4 Training the model

The modelling technique that we describe in this study was developed for plant species distributions [17]. We used the *Malpolon* framework [36] after some adaptations to our use case. It was built on top of PyTorch [37] and PyTorch Lightning [38] frameworks.

The *Malpolon* framework implements a convolutional network with the *resnet50* feature extractor [15]. We adapted it to use 29 inputs channels, and 38 numerical outputs converted to relative probabilities by a Softmax function. It was trained in two sessions by minimizing the Binary Cross Entropy loss: one with a .1 learning rate, and another one with a .01 learning rate to fine-tune the weights.

The target probabilities for training were set using one-hot encoding, *ie.* a one for the observed species and zeroes for all others. This follows the principle of assumed negatives [39]: we assume that only the observed species is present at the point of observation. This equates to considering the pseudo-absences of the other species. This is obviously wrong, but as we work with a limited number of species in an extensive area and period of time, chances are slim that the model receives contradictory information.

# 2.5 Evaluation metrics and performance assessment

Training data were randomly split into three sets: training (60%), validation (20%) and test (20%). The validation set was used to assess and improve performance during the training phase, while the test set was used after training to compute the final performance of the model, on data that it had never seen before.

The accuracy of the final version of the model was 69%, which means that in 69% of cases, the most likely taxon according to the model was the same as the one that was actually observed. See table 3 for more complete accuracy results.

Table 3. Probability that the observed taxon is among the Top N predictions of the model, for 11 values of N

Top N	Probability
1	69.15%
2	83.19%
3	89.13%
4	92.83%
5	95.13%
6	96.63%
7	97.48%
8	97.99%
9	98.43%
10	98.75%
38	100.00%

A confusion matrix was computed on the test data set and is shown in Figure 4. It shows that some taxa were easily identified by the model (the top two being *Aptenodytes forsteri* and *Mobula alfredi*). Others were harder to predict, the worst two being *Istiompax indica* and *Carcharhinus longimanus*.

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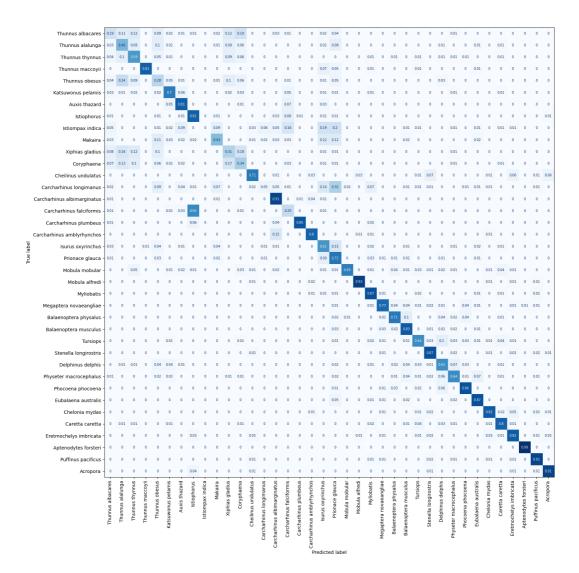


Figure 4. The confusion matrix shows the top predictions of the model on the test data set, for each observed taxon.

# 2.6 From probability predictions to distribution maps

After training the model, we used it on new data to generate distribution maps. As the environmental data download phase can be quite slow, we had to choose to focus either on time extent or spatial extent, but not both at the same time. This is why we chose to compute two different outputs:

- Global species distribution maps at four dates in 2021.
- Regional species distribution maps for the Southwestern Indian Ocean at 53 dates in 2021.

It is important to note that the model may be used at any date in any area; the only limitation is the availability of environmental data, and the time required to download them.

Two grids covering both areas were generated. They comprised 36,506 points for the global oceans and 3,001 for the Southwestern Indian Ocean. Environmental data were downloaded for each of these points and run through the model, which led to 38 relative probability predictions for each of

these points, at each requested date. For each taxon, these probabilities were interpolated over the whole area to generate rasterized distribution maps.

It is worth noting that since we are working with relative probabilities (because of the softmax layer), the absolute values have little purpose. Therefore no scale is provided for all distribution maps: they should be interpreted relatively to one another, across species, time or space.

#### 2.7 Influence of variables

To study the influence of variables, a new model was trained after removing chlorophyll and sea surface temperature at D-5 and D-15, as well as Eddy Kinetic Energy. Indeed, these layers are highly correlated with chlorophyll and sea surface temperature on the day of occurrence, and geostrophic current respectively.

It is worth noting that this model has almost the same accuracy (69.08%) as the previously described one (69.15%), which shows that the 5 variables that were removed have very little influence on the classification.

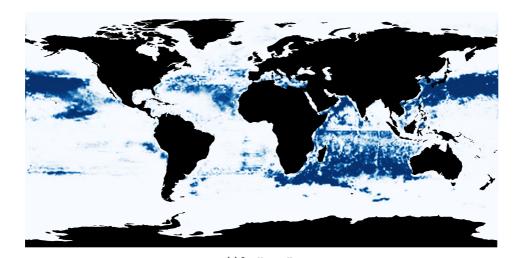
Afterwards, the most determining variables were calculated using the integrated gradients method for a random sample (N=1,000) of the points on the world grid [40], using the Captum python package [41]. They were then aggregated over the geographical area (sum of absolute values) and these values were grouped by taxon (the top prediction).

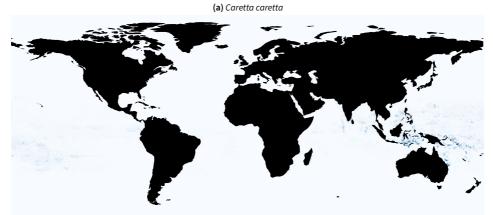
3. Results

# 3.1 Presentation of the species distributions maps

## 3.1.1 Global oceans

Distribution maps were calculated on four dates, all in 2021, corresponding to both solstices and both equinoxes, for the thirty-eight taxa. Figure 5 shows these maps for three species on the spring equinox. All 152 distribution maps are available online [42].





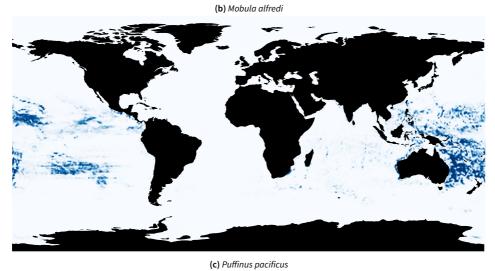
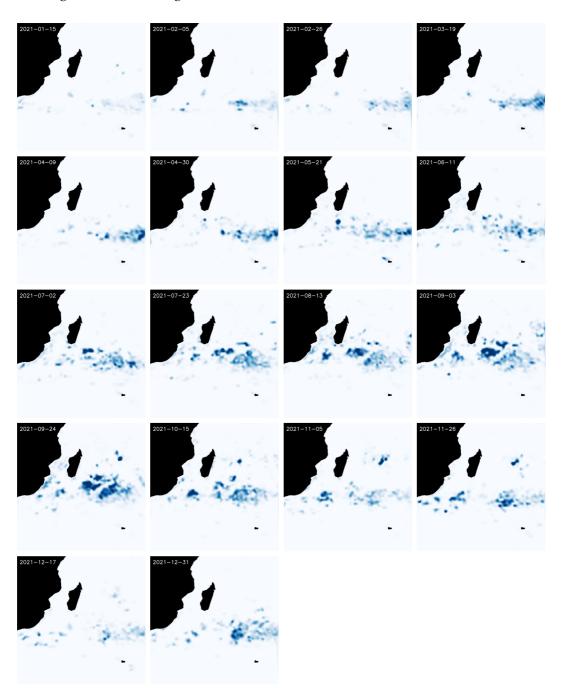


Figure 5. Examples of distribution maps on March 20th, 2021.

# 3.1.2 Southwestern Indian Ocean

In this case, distribution maps were calculated each week of 2021, for the 38 taxa. To make visualization easier, they were exported as animated GIFs that are available online [43]. An example for *Prionace glauca* is shown in Figure 6.



**Figure 6.** Distribution maps for *Prionace glauca* every three weeks of 2021. The contrast was increased to improve visibility: colours from this figure should not be compared to other figures.

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# 3.2 Comparison of predicted distribution maps to established maps

Validation of the distribution maps is challenging because existing distribution maps rarely take into account temporal variability, except sometimes broad seasonal variations. Yet the maps that we produce are highly dependent on time, see Figure 6 for instance.

We compared some of our distribution maps to established ones, to check for inaccuracies. See Figure 7 for a few examples.

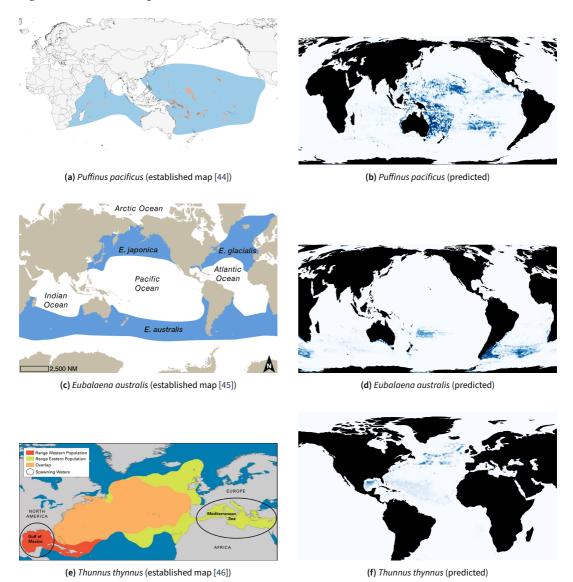


Figure 7. Comparison between established distribution maps and deep-learning generated maps

**Puffinus pacificus** The prediction map 7a is coherent with the established one 7b, although there is a significant difference between the Indian and Pacific oceans. Since established maps are usually binary (presence/absence), the difference we see between the two oceans cannot be (in)validated. It

is possible that the Pacific Ocean is more suitable to this species than the Indian Ocean, or there may be an under-representation of occurrences in our data set.

**Eubalaena australis** The predicted distribution 7d fits within the known geographical range of *Eubalaena australis* 7c, and there is a strong disparity of the prediction density within this area. Again, no assumption can be made on the plausibility of the predictions, as this heterogeneity may be caused by temporal variation, or it may not fit reality. Another possibility is that the established geographical range is not fully used by the species.

**Thunnus thynnus** The predicted range for *Thunnus thynnus* 7f is within the established range 7e, but it does not include all of it. Specifically, the Mediterranean Sea and the Bay of Biscay are excluded, even though there is a major population living in these areas [47]. After checking our input data, this shortcoming can be explained by the under-representation of this population in the occurrences used for training. This will be discussed further in section 4.2.2.

# 3.3 Analysis of determining variables

Over the predictions for the 2021 Global use case, the most influential variables were FSLEs (strength), sea surface temperature, pH, bathymetry, salinity, and FSLEs (orientation), in this order. See table 4 for a full accounting of variable influence.

Figure 8 shows the median integrated gradient for each taxon. Note that *Istiompax indica* is not present in this chart as it was never predicted to be the most likely taxon.

#### 3.4 Effect of a 2°C increase in sea surface temperature

Predictions were computed after adding 2°C to sea surface temperature, leaving all other variables unchanged. In the context of climate change, this is a tentative projection but it is theoretical, as there are significant and complex correlations between future changes in various environmental variables. Our model does not provide abundance results, but relative probabilities indicate which taxa benefit or suffer from this change.

There is also a change in presence areas, illustrated in Figure 9. For *Caretta caretta* and, to a smaller extent, *Eubalaena australis*, there is an expected poleward shift: away from the equator and towards cooler water, as projected previously [48]. The evolution of the *Katsuwonus pelamis* range is harder to interpret: this species may not be that sensitive to warmer water because it already lives in the equatorial band, so the change may be caused by the interaction between temperature and other variables. In reality, a temperature rise would cause a decrease in oxygen levels (not modelled here), which might drive the *Katsuwonus pelamis* population away from the equator [49].

Evolution maps were calculated by subtracting the  $+2^{\circ}$ C projection from the corresponding 2021 predictions, for each geographical point. This process creates new raster layers with a -1 to 1 range reflecting the evolution of predicted probabilities.

**Table 4.** Statistics of the influence of variables on a sample of 1,000 predictions ( $\times$ 1,000, sorted by median).

Variable	mean	std	min	25%	50%	75%	max
FSLEs (strength)	1.08	0.8	0.12	0.53	0.8	1.36	3.87
SST	0.77	0.71	0	0.24	0.56	1.1	3.66
рН	0.64	0.62	0	0.25	0.41	0.8	3.35
Bathymetry	0.67	0.76	0	0.17	0.39	0.85	5.25
Salinity	0.6	0.53	0	0.24	0.38	0.75	2.83
FSLEs (orientation)	0.53	0.42	0.04	0.24	0.37	0.64	2.62
Geos. current (u)	0.4	0.34	0	0.18	0.28	0.49	2.15
Geos. current (v)	0.39	0.31	0.05	0.18	0.28	0.49	1.56
Surface wind (v)	0.37	0.28	0.04	0.17	0.26	0.45	1.36
Oxygen	0.42	0.45	0	0.06	0.24	0.66	3.04
Surface wind (u)	0.33	0.26	0.03	0.15	0.23	0.4	1.21
Pacific Ocean	0.41	0.56	0	0	0.23	0.63	3.11
Wave height	0.18	0.14	0.02	0.08	0.13	0.22	0.7
Mixed layer thickness	0.26	0.37	0	0.05	0.12	0.34	2.83
Prochlorophytes	0.07	0.16	0	0.02	0.03	0.08	2.9
Green algae	0.08	0.15	0	0.01	0.03	0.08	2.23
Haptophytes	0.07	0.14	0	0.01	0.03	0.07	2.27
Prokaryotes	0.03	0.05	0	0.01	0.02	0.03	1.15
Chlorophyll	0.03	0.04	0	0.01	0.01	0.03	0.53
Dinophytes	0.04	0.08	0	0	0.01	0.03	0.83
Diatoms	0.03	0.07	0	0	0.01	0.03	0.72
Atlantic Ocean	0.19	0.43	0	0	0	0.16	2.6
Indian Ocean	0.25	0.6	0	0	0	0	3.28
North hemisphere	0.42	0.7	0	0	0	0.56	3.66

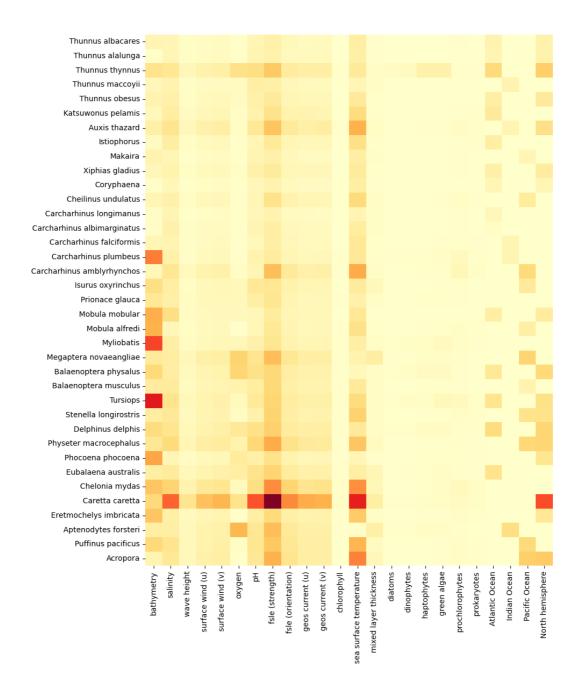
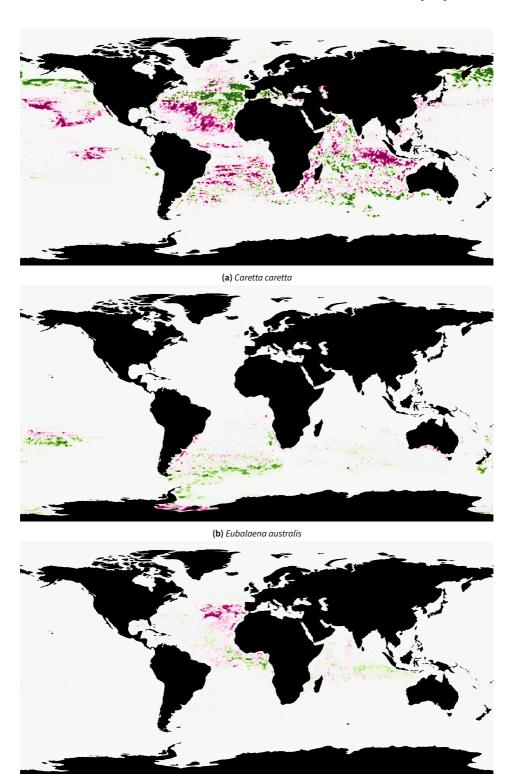


Figure 8. Variables that had the most influence on the determination of each taxon presence



(c) Katsuwonus pelamis

**Figure 9.** Examples of evolution maps after adding +2°C. Green areas are more suitable with 2°C more, while pink areas are less suitable. Maps for all 38 species on all 4 dates are available online. [50]

4. Discussion

# 4.1 Ecological interpretation of the results, implications for offshore species distributions

The variables that were identified are coherent with past research. Specifically, FSLEs were identified as a particularly important movement predictor for top marine predators [51]. Sea surface temperature was also expected to be an important predictor, as it is the most frequently used descriptor in marine SDMs [14] and was identified as the most relevant factor in an SDM review [52].

This study also demonstrates a high sensitivity to temporal variations in environmental conditions, as shown in figure 6. This highlights the need for distribution models of fast-moving species to consider these variations, instead of relying only on averaged values..

There are some surprises in influential variables: bathymetry was not a good predictor of *Acropora* coral distribution, which is contradictory with their need for light. A possible explanation is that the model may have used other variables as a proxy for low depths.

# 4.2 Benefits and limitations of using deep learning for SDMs in the open ocean

This method holds promise in helping researchers uncover new correlations between the oceanic conditions and species distributions: implicit feature extraction allows the use of more numerous and more complex features. But there is a drawback: determining features have to be studied afterwards, and it is not straightforward. In this study, we showed the variables that had the most influence on average. This needs to be complemented by a deeper study of the nature of the determining features.

We noted three main limitations of our method, namely performance metrics, biases in the input data, and some undetected patterns.

4.2.1 Accuracy

The accuracy of the model could still be improved, depending on the ecological feasibility. Indeed, if some species are frequently seen together, there is no way for the model to discriminate between the two. In that case, this uncertainty will show as a .5 mistake rate even though it is the correct result. A way to improve the final accuracy score would be to group species by traits. But this would remove the possibility of studying differences between similar species.

For example, the confusion matrix in figure 4 shows that *Xiphias gladius* and *Coryphaena* are often predicted instead of each other. This may be the result of these two taxa having similar habitats, and the low resulting score does not necessarily mean that the predictions are wrong.

Accuracy is not an ideal metric for this use case, but the scarcity of training data is very limiting in that aspect. This is why we provided a Top-N score in table 3 for a more complete performance assessment.

4.2.2 Observer bias

Most observation data in the open ocean come from fishing vessels, which target certain species. This causes observations to mostly include target species or frequently associated species. Furthermore, fishing boats tend to target some areas based on outputs of fishing guidance models so it creates an artificial correlation between the parameters used in these models and the presence of animals.

The strength of deep learning in this context is that it makes no assumption when there is no data: it replicates the results from similar well-known areas. This partly compensates for sampling effort heterogeneity. But this only works when there is a homogeneous population, unlike *Thunnus thynnus* which has two separate stocks (West and East Atlantic). This explains why the model failed

to extrapolate from West Atlantic data and to predict high probabilities in the Mediterranean Sea and the Bay of Biscay.

Finally, some data come from scientific tracking of individual animals, so these individuals may be over-represented in our data and reflect their preferences rather than the general tendency of their species. The large amount of occurrences that we used help tackle this bias.

These biases would be better tackled with more available data, which is a serious issue in the open oceans. Little data is produced relative to the size of the oceans, and a large part of this data is not shared publicly. More data is key to better models and more trustworthy distribution maps.

# 4.2.3 Undetected patterns

Detection of seasonal migrations is incomplete. For instance, we should see the *Megaptera novae-angliae* distribution spreading north during the southern winter. The model also did not catch the *Thunnus thynnus* seasonal spawning in summer in the Mediterranean Sea.

# 4.3 Suggestions to further improve the modelling methods

#### 4.3.1 Occurrence data

As a first experiment, occurrence data were selected randomly for this study. Even though the aim should not be to have a perfect fit between observation data and model predictions, sourcing training data from many different providers rather than randomly would help reduce observer bias.

#### 4.3.2 Environmental data

It has been suggested to include 3D environmental data, as most variables vary with depth and occurrence data are not limited to the surface [53]. Such data could easily be included in the input data with no change to our method. Additional data may be beneficial, in particular the distance to the nearest coastline or level of anthropisation.

## 4.3.3 Other use cases

The present method could be used at different scales, in particular in coastal areas. This would require a significant change in the input variables, as the resolution of globally available environmental data is a limiting factor. They could be replaced by satellite or drone images, as well as locally available (more precise) environmental data.

5. Conclusion

# 5.1 Main findings and their significance

The present method provides a way to estimate species distribution at all dates and all areas, provided environmental data is available. While this makes it difficult to judge accuracy (there is often no reference data), it provides a baseline that can be calculated for any species (that have enough existing observations). Researchers working on terrestrial plants have also shown that such models may be used to infer species distribution for rare species, by extrapolating results from co-occurring species [17].

# 5.2 Implications for management and conservation of offshore species

We hope this method will be developed further and used on other endangered species, together with existing methods and field observation. The technique that we presented would be especially useful in the hands of scientists who are experts in the life cycle of specific species. It would help them

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increase scientific knowledge of their distributions, which is essential for decision-makers to target areas of interest for conservation.

# 5.3 Recommendations for future research and potential applications

While the accuracy of our distribution maps is difficult to assess, there is exceptional room for improvement and further research. All the blocks in Figure 1 can be modified, either to adapt the process to a different use case or to try to improve the quality of the results. Here are some examples of potential changes:

- To study other species, the initial choice of species can be changed, for example, to focus on sedentary species or a specific area.
- To improve accuracy, the occurrence data may be selected in other ways that are not random.
- To investigate the influence of other variables, they may be added to the variable set.
- To study the long-term effect of environmental conditions, some variables may be included with a longer time lag such as months or years.

The results we presented in this article are a small part of what can be achieved with this model. Many other scientific questions can be investigated both with the model we provide (already trained) or with other models trained with the same method.

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# Open data statement

Code

The code that was used to prepare the data, train the model, and export the outputs is available on GitHub in the IRDG2OI/deep-sdm-oceans repository and on Zenodo [54].

Input data

The input data include the CSV file describing the geographical points, the standardized numpy arrays of corresponding environmental data, and the standardization factors. They are available on Zenodo [55] for each use case:

- Training data (includes train+validation+test)
- · Prediction data for the world at 4 dates
- Prediction data for the Western Indian Ocean at 53 dates

Modelling

We provide the model checkpoint and configuration file [56], so researchers can make predictions with the presently described model.

We also provide the code that was used for training so researchers can adapt it to their needs and retrain a new model [54]. It consists of Python files based on a custom version of Malpolon.	38
Results	38
The distribution maps were uploaded to Zenodo for easy visualisation, in three repositories:	38
<ul> <li>Global predictions as PNGs and GeoTIFFs [42]</li> <li>Western Indian Ocean predictions as GIFs and GeoTIFFs [43]</li> </ul>	38
<ul> <li>+2°C hypothesis as PNGs and GeoTIFFs [50]</li> </ul>	38

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