Identification of candidate pelagic marine protected areas through a seabird seasonal-, multispecific- and extinction risk-based approach

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Abstract. With increasing pressure on the oceans from environmental change, there has been a 1 2 global call for improved protection of marine ecosystems through the implementation of Marine 3 Protected Areas (MPAs). Here, we used Species Distribution Modelling (SDM) of tracking data 4 from 14 seabird species to identify key marine areas in the southwest Atlantic Ocean, valuing areas based on seabird species occurrence, seasonality and extinction risk. We also compared 5 overlaps between the outputs generated by the SDM and layers representing important human 6 7 threats (fishing intensity, ship density, plastic and oil pollution, ocean acidification), and 8 calculated loss in conservation value by using fishing and ship density as cost layers. The key 9 marine areas were located on the southern Patagonian Shelf, overlapping extensively with areas 10 of high fishing activity, and did not change seasonally, while seasonal areas were located off south and southeast Brazil and overlapped with areas of high plastic pollution and ocean 11 acidification. Non-seasonal key areas were located off northeast Brazil on an area of high 12 13 biodiversity, and with relatively low human impacts. We found support for the use of seasonal areas depending on the seabird assemblage used, because there was a loss in conservation value 14 15 for the seasonal compared to the non-seasonal approach when using 'cost' layers. Our approach, 16 accounting for seasonal changes in seabird assemblages and their risk of extinction, identified 17 additional candidate areas for incorporation in the network of pelagic MPAs.

18 Keywords: Ecologically and Biologically Significant Areas; Important Bird Areas; seabird
19 tracking; species distribution modelling; threats; Zonation

20 Introduction

21 Oceans are facing rapid and profound changes in their characteristics and structure (Halpern et al., 2008; Rockström et al., 2009; Valdés et al., 2009). Ocean changes are so widespread that 22 only <4% of the global ocean area is considered to be experiencing very low human impacts 23 24 (Halpern et al., 2015). Hence, there is a need for a major international effort to protect and 25 maintain the functioning of marine ecosystems, through the implementation of Marine Protected Areas (MPAs) (Rockström et al., 2009; Kachelriess et al., 2014). Despite a global agreement to 26 designate MPAs covering 10% of the global ocean (Secretariat of the Convention on Biological 27 28 Diversity, 2006), only 2.98% is currently protected, and <1% of the global ocean comprise 'no-29 take' zones, where no harvesting of marine resources is allowed (Marine Conservation Institute 30 2015). Furthermore, few established MPAs effectively and integrally protect their targeted 31 species, so their value is likely to be even lower than expected (Jameson, Tupper & Ridley, 32 2002; Mora & Sale, 2011; Edgar et al., 2014).

To truly protect the diversity of marine systems, the design of MPAs must account for 33 34 its location (Jameson et al., 2002; Sundblad, Bergström & Sandström, 2011), connectivity with 35 other areas (Sundblad et al., 2011) number and proportion of protected oceanographic features (e.g. upwellings, seamounts; Sundblad et al., 2011; Dunn et al., 2014), species occurring there 36 37 and their seasonality (Hyrenbach, Forney & Dayton, 2000; Mann & Lazier, 2006), climatic 38 variability (Jameson et al., 2002), isolation from human impacts (Rife et al., 2013; Edgar et al., 39 2014), human activities (Charles & Wilson, 2009; Edgar et al., 2014; Mazor et al., 2014) and, 40 socio-economic use of the area (Charles & Wilson, 2009). Most MPAs are designed to improve 41 fisheries management, i.e. to manage commercial fish stocks in a more sustainable way (e.g. 42 (Pitchford, Codling & Psarra, 2007; Horta e Costa et al., 2013ab; Stevenson, Tissot & Walsh, 43 2013). Few studies have accounted for other environmental factors in the design of MPAs. For 44 instance, ship movements (Dalton, 2004; Halpern et al., 2008), exposure to plastics (Lebreton, 45 Greer & Borrero, 2012; van Sebille, England & Froyland, 2012; Wilcox, Van Sebille & Hardesty, 2015), oil pollution (Wiese & Robertson, 2004; Moreno et al., 2013; Kark et al., 46

2015), and ocean acidification (Harvey, Gwynn-Jones & Moore, 2013) present substantial
impacts to marine biodiversity, and should be considered in MPA design (Kelleher, 1999;
Pomeroy *et al.*, 2005).

50 Wide-ranging, pelagic top predators have been widely proposed as useful tools for 51 pointing out important areas which can add conservation value to MPA networks (Hooker & 52 Gerber, 2004; Sergio et al., 2008; Ronconi et al., 2012). Top-predators integrate factors 53 affecting seasonal variation in abundance and distribution of their lower trophic-level prey over 54 broad areas (Montevecchi et al., 2012; Tancell et al., 2012; Michael, Jahncke & Hyrenbach, 55 2014). Seabirds, in this context, are relatively easy to sample at their breeding colonies and to 56 track them at sea by using electronic devices, providing information on foraging movements at various spatio-temporal scales (Phillips et al., 2006a; Delord et al., 2014; Thiebot et al., 2014). 57 58 Several important marine areas have been identified based on the distribution and occurrence of 59 seabird species and proposed to integrate the current network of MPAs. The marine Important 60 Bird Areas (mIBAs) represent one of the most recent wide-ranging international efforts to 61 identify relevant areas for marine conservation (Lascelles et al., 2012, 2016).

62 Here, we applied spatial distribution modelling to tracking data from 14 pelagic seabird 63 species in the southwest Atlantic Ocean (Food and Agriculture Organization, FAO, Fishing 64 Area 41). The use of tracking data is deliberately used to identify important areas in offshore 65 waters since many important "coastal" areas have been identified for MPAs or MIBAs through 66 other approaches and investigations of coastal seabirds. We identified key areas based solely on 67 the climatic and oceanographic variables irrespective of the distribution of threats, a posteriori 68 we quantified sensitivity of species to threats and quantified the costs to human activities within 69 those areas that are important for the birds, by imposing restrictions of activities within the areas 70 and calculating changes in conservation value. Such approach is justified by evidences that the 71 distribution of threats matches oceanographic conditions targeted by marine animals (Karpouzi, 72 Watson & Pauly, 2007; Hatch et al., 2008; Titmus & Hyrenbach, 2011; Krüger et al., 2016a). 73 Our objectives were: (1) identify important areas for pelagic seabirds throughout the year; (2) 74 quantify the sensitivity of the identified important areas for seabird assemblages to 75 environmental pressures; (3) compare our spatially-explicit results to the existing networks of 76 proposed or established protected areas identified by international organizations, and (4) carry 77 out a cost analysis evaluation of the implementation of seasonal vs non-seasonal protected areas. 78 FAO Fishing Area 41 encompasses a wide range of latitudes from tropical to Antarctic waters, 79 including a large array of habitats and oceanographic conditions, the largest continental shelf in 80 the Southern Hemisphere and a highly productive convergence zone that support a biodiversity-81 rich ecosystem, and major fisheries (Vasconcellos & Csirke, 2011). Despite several proposals to 82 identify key areas for conservation in pelagic ecosystems, either in the form of Ecologically and 83 Biologically Significant Areas (EBSAs) (Dunn et al., 2014) or mIBAs (Lascelles et al., 2012), 84 the designated MPAs inside FAO41 are mostly restricted to coastal regions (Marine Conservation Institute 2015, http://www.mpatlas.org/explore/). Thus, such network of MPAs 85 does not protect marine pelagic species and systems. In this sense, our analysis prioritizes 86 habitats used by pelagic seabird communities weighted by their seasonality and extinction risk 87 88 (which may protect other pelagic species) as pelagic habitats are underrepresented in the MPAs 89 network within FAO41. The key areas identified with this study may be seen as supporting the 90 already proposed MPAs as add-ons, which indentifies pelagic and seasonal areas that were not 91 considered by previous methods.

92 Materials and methods

93 Tracking data and bird assemblages

We used geolocator (Global Location Sensor or GLS) tracking data from 14 seabird species,
occurring at the southwest Atlantic Ocean during part or all of the annual cycle (Table 1).
Deployment and other details are provided elsewhere (Grémillet *et al.*, 2000; Croxall *et al.*,
2005; Phillips *et al.*, 2006a; González-Solís, Croxall & Afanasyev, 2008; González-Solís *et al.*,
2009; Mackley *et al.*, 2010; Quillfeldt *et al.*, 2013; Ramírez *et al.*, 2013; Reid *et al.*, 2013;
Missagia *et al.*, 2015, Krüger *et al.*, 2016a,b). The data were modelled in three different periods:
summer (Oct.-Mar.), winter (Apr.-Sep.) and year-round (Jan.-Dec.). For summer and winter

101 periods, we removed bird locations from months when the number of points was below10% of 102 the maximum number for that species in any month in order to remove the interference of 103 migratory movements from the seasonal data, but all available locations were included in the 104 year-round models. Each species was classified as one of two groups, based on its core 105 distribution relative to the mean position of the northern boundary of the Subtropical Front 106 (Burls & Reason, 2006): Southern species (i.e. species with a distribution largely south of 35°S 107 within the area) and Northern species (i.e. species mainly distributed north of 35°S within the 108 area). This categorisation was based on previous evidence for segregation of seabird 109 communities resulting mostly from latitudinal gradients in sea surface temperature (Péron et al., 110 2010; Krüger & Petry, 2011; Navarro et al., 2015; Quillfeldt et al., 2015).

111 Environmental variables

112 Our Species Distribution Models (SDMs) used 10-year average data (from summer, winter and year-round) for the following oceanographic variables: chlorophyll-a concentration (CHL), 113 114 CHL anomaly, CHL gradients, sea surface temperature (SST), SST anomaly, SST gradients, 115 Sea Surface Height (SSH), SSH anomaly, wind speed, water depth, and minimum distance to 116 coast (DCOA) (Supporting Information Fig. S1). All variables, except DCOA, anomalies and gradients, were downloaded as monthly mean composite raster images from NOAA 117 118 CoastWatch Browser (http://coastwatch.pfeg.noaa.gov/). SST anomalies and CHL anomalies 119 for each month were calculated as the difference between the average value for a given month 120 and year, and the average for that month over a 10-year period in that grid cell. Gradients were 121 generated by calculating the standard deviation of each cell in relation to the adjacent cell values 122 (Sidhu et al., 2012; Li et al., 2015). All raster files were processed in ArcMap 10.2.

123 Species Distribution Models

124 Tracking data was combined with environmental predictors to generate SDMs from presence-

125 only data using the Maximum Entropy (MaxEnt) software (Oppel et al., 2012; Quillfeldt et al.,

126 2013) on a 1°x1°spatial grid (e.g. Pinet *et al.*, 2011; Ramírez *et al.*, 2013; Missagia *et al.*, 2015),

127 as the mean error of geolocation tags is usually lower than 1° (Wilson et al., 1992; Phillips et

128 al., 2004; Nielsen & Sibert, 2007). Geolocators are useful to detect and model core areas for 129 animals' distribution (Quillfeldt et al., 2013; Krüger et al., 2016a, 2016b) and compared to more 130 accurate methods like GPS or Argos, geolocators tend to slightly inflate home range of animals 131 (Phillips et al., 2004). To compensate this potential inflation, we were conservative in the next 132 steps of the analysis by only assuming areas of high importance values (see next section) to propose the key areas. Furthermore, geolocators are able to collect data for a longer period of 133 134 time that is highly limited by battery size in GPS and PTTs, and this may also represent a barrier 135 to sample distribution data on smaller seabirds like gadfly petrels and prions which can not 136 carry much weight. The tracking data were divided into training and test data by randomly 137 setting aside 10% of the tracking dataset for spatial evaluation of the models (Araújo & Guisan, 138 2006; Austin, 2007). The first step of the modelling consisted of principal component analyses 139 (PCA) to eliminate any potential effect of multicollinearity. This is a recommended procedure 140 as our goal was to model species distributions, rather than to verify relationships between 141 species occurrence and the environment (Merow, Smith & Silander, 2013). We extracted the 142 scores for components with eigenvalues above 1. With those scores, we ran MaxEnt on the 143 presence-only positions, 50 times, with a bootstrap procedure to obtain a prediction of the 144 average distribution (Edrén et al., 2010). We assessed the accuracy of models using the area 145 (AUC) under the Receiver Operating Characteristic (ROC) curve. The AUC estimates the 146 likelihood that a randomly selected presence point is located in a raster cell with a higher 147 probability value for species occurrence than a randomly generated point (Phillips, Anderson & 148 Schapire, 2006).

149 Calculating area importance values

The outputs from MaxEnt were used to calculate area importance values for each bird
assemblage using the Zonation software (Moilanen *et al.*, 2005; Moilanen & Wintle, 2006;
Leathwick *et al.*, 2008). Zonation is assumed as one of the best programs to set out conservation
priorities when efforts focus on ecological communities and habitat connectivity (i.e.,
Blumentrath, 2011; Delavenne *et al.*, 2012). Zonation generates a raster with pixels representing

155 the importance value, taking into account the probability of occurrence of each species given its 156 weighting (see below) and response to habitat. The final output is scaled from 0-1, representing 157 the least (0) to most important (1) areas. Each species was weighted by its IUCN conservation 158 status as follows: Least Concern (LC) = 1; Near Threatened (NT) = 2; Vulnerable (VU) = 3; 159 Endangered (EN) = 4 and Critically endangered (CR) = 5. Boundary Quality Penalty Curves (BQPCs; Moilanen & Wintle, 2007), which are measures of species responses to habitat 160 161 connection, were constructed empirically based on the IUCN conservation status. We generated responses with changing slopes for CR = -0.01, EN = -0.008, VU = -0.004, NT = -0.002 and LC162 = 0. Thus, the proportion of area occupied by each species decreases with increasing habitat 163 164 disconnection (except for LC) and the intensity of the decrease (slope) is proportional to the 165 level of threat.

166 We evaluated the sensitivity of the weighting and habitat connectivity response (more 167 details in Supporting Information SI). Changing the values for these two variables had no 168 substantial effect on the final output, except for a complete null model considering all species as 169 equivalent, the output from which was unrealistic and did not match the species distributions 170 (Supporting Information Fig. S2, S3, S4). Probability of occurrence in a given pixel diminishes 171 when habitat characteristics differ from those within surrounding cells, using a home range-172 based grid size (Leathwick et al., 2008). Uncertainty analysis (Moilanen & Wintle, 2006) was disabled in the computation (Leathwick et al., 2008). The importance value was calculated for 173 174 each assemblage during the three specified periods (summer, winter and year-round). We were 175 conservative to propose key areas due geolocators errors (see previous section) and selected 176 areas that fell within 1% (90% of importance value) of the distribution threshold (Moilanen et 177 al., 2005), areas of 0.5% (95%) and 0.1 % (99%) threshold (Oppel et al., 2012), in a nested 178 design – these areas are within the 1% threshold. The locations of the 0.1 % candidate MPAs were compared with those of existing MPAs (Marine Conservation Institute 2015), EBSAs 179 180 (Dunn et al., 2014) and mIBAs (Birdlife International 2015).

181 Environmental threats

We examined spatial risk from five environmental threats: ocean acidification, oil pollution, 182 183 floating plastics, shipping and fishing intensity (Supporting Information SII, Fig.S5), which are 184 environmental stressors largely recognized as stressors to which marine animals, particularly seabirds, are exposed and vulnerable, by means of spatial overlap and evidence of impact: 185 186 acidification (Grémillet & Boulinier, 2009), oil pollution (Camphuysen & Heubeck, 2001; Wiese & Robertson, 2004; Moreno et al., 2013), plastic (Titmus & Hyrenbach, 2011; Jiménez et 187 188 al., 2015; Wilcox et al., 2015), shipping (Hatch et al., 2008), fisheries (Anderson et al., 2011; Croxall et al., 2012). These variables were standardized to have a mean of 0 and an SD of 1 189 190 (Zuur, Ieno & Smith, 2007).

191 Cost analysis

192 We evaluated costs for the proposed key areas, using the layers from Fishing Vessels Density 193 and overall Ship Density (Supporting Information SII and Fig. S5) as cost layers (Dalton, 2004; 194 Leathwick et al., 2008) in the Zonation software. We are using those layers as 'costs' because 195 both are the most manageable factors from the most widespread human threats affecting marine 196 fauna worldwide. Impact of fisheries on marine fauna are well described in literature, like food 197 depletion through overfishing and direct mortality through bycatch (Becker & Beissinger, 2006; Cury et al., 2011) but the vessel traffic is a potential source of oil (Halpern et al., 2008, 2015; 198 199 Hatch et al., 2008) and noise (Morton & Symonds, 2002; Weilgart, 2007; Codarin et al., 2009) 200 pollution, and birds may collide against vessels during the night (Black, 2005; Glass & Ryan, 201 2013).

Zonation uses cost layers to reduce the value (importance to conservation based on the previously specified parameters) of a given cell, thus cost is interpreted as a variable that reduce the conservation value of a given area. The "cost" variables were used here as a cost/limitation imposed to human activities, in the perspective that a no-cost represent no changes in the current activities, and a cost represent restrictions to activities. As our valuing of areas considers no influence of the human activities, sole the seabirds distribution regarding environmental variables, this represents the hypothetical value of the area under no constraint by human

209 activities. Then it is possible to measure differences on the conservation value of an area when 210 comparing different cost scenarios with the previous area valued under no influence of human 211 activities. We used two cost constraint scenarios to evaluate how the intensity of human activities may change the conservation value of candidate MPAs (i.e. key marine areas) if 212 213 shipping and fishing were not regulated (no-cost, scenario 1) or partially regulated (cost, scenario 2). Costs constraints were calculated to each time-frame (Summer, Winter and Year-214 215 round). The resulting values for each cost model were then subtracted from the respective area 216 importance value within the 99, 95 and 90 threshold values (the proposed areas based on seabird distribution only), to evaluate how much each of the human activities change the conservation 217 218 values within the proposed key areas.

219 Statistical Analysis

220 We used the pixel value of the rasters as point information, and the centroid of longitude and 221 latitude (N=1697) as the geographic position of each grid cell. To evaluate the degree of correspondence between the distribution probabilities for the two seabird communities (MaxEnt 222 223 outputs) and the distributions of the environmental threats, we used Canonical Correspondence 224 Analysis (CCA) using 'cca' function from the 'vegan' package (Oksanen et al., 2013) within the R environment (R Core Team 2015). Species probabilities were arcsine transformed. Species 225 matrix entered the analysis as dependent matrix, and the threats entered the analysis as the 226 227 independent matrix, with geographical coordinates used as covariables to control for spatial 228 auto-correlation (cca[Species Matrix ~ Threat Matrix + Latitude * Longitude]). Each time-frame 229 was analyzed in separate. After running the analysis, we quantified the Euclidian distance of the 230 species from the threats in the CCA bi-dimensional space to compare shifts in group exposition 231 to threats.

We compared the change of value within the proposed key areas between cost models, time-frames and assemblages through a Generalized Linear Mixed Model in *'lme4'* package within R-environment (Bates *et al.*, 2015). As threshold values for proposed key areas (90%,

235 95% and 99% of importance values) are in a nested design, we entered the threshold as a236 random factor in the model.

237 **Results**

238 Species Distribution Models

Many of the environmental variables were highly correlated (Supporting Information Table S1). 239 240 The power of the models using PC1 and PC2 to predict species distribution was high (AUC=0.79 \pm 0.06), although the models were less accurate for species with large latitudinal 241 ranges, such as Cory's Shearwater (Calonectris borealis), Cape Verde Shearwater (Calonectris 242 243 edwardsii), Great Shearwater (Ardenna gravis) and Wandering Albatross (Diomedea exulans). 244 It was clear that dividing the data into two seasons resulted in a slightly more accurate prediction of distributions than using data from the entire year (AUC summer = 0.80 ± 0.06 ; 245 246 AUC winter = 0.83 ± 0.05 ; AUC year-round = 0.76 ± 0.06), probably because species occupy 247 only part of the annual distribution in any one season (Fig. S6, S7, S8).

248 Assemblages and Environmental Change Factors

Canonical Correspondence Analysis yielded slight different results for each time frame, for Summer the constrained analysis captured 74.1% of data variability (axis1= 83.0%, axis2= 12.3%), 70.7% for Winter (axis1= 81.9%, axis2 = 15.9%) and 77.2% for all Year (axis1 = 81.9%, axis2 = 14.9%). Probability of occurrence of species (species matrices) was significantly matched by the distribution of threats for Summer ($F_{8,1638}$ =585.49, P=0.001), Winter ($F_{8,1638}$ = 494.48, P=0.001) and all Year ($F_{8,1638}$ = 691.34, P=0.001).

There was a clear segregation between assemblages within the bi-dimensional space of the CCA which showed that Northern species were more exposed with plastic pollution and acidification, while Southern species were associated with Fisheries and Ship Traffic (Fig. 1a,b,c). However, such trend was not obvious in Winter (Fig.1b) as the two Northern species (TA and TP) were highly pelagic and occurred in areas of low cumulative threat occurrence (see Supporting Information Fig. S5, Fig. S7). In Winter two sub-groups of Southern seabirds were differentially exposed to Acidification and Plastic Pollution (AP, GHA, WA) or to Fisheries and
Ship Traffic (BBA, NGP, SGP). Two species were displaced from threat influence in two cases
(MS and WCP) which seem to be caused by their wide latitudinal distribution. This was
reflected in the bi-dimensional distance of each group from threats, where Northern seabirds
increased their distance from threats in Winter comparing to Summer and all Year, but Southern
seabirds decreased their distance from threats in Winter comparing to Summer and all Year,
with exception of WCP (Fig. 1d-i).

268 Proposed key marine areas

269 The areas of highest value for Southern species did not vary seasonally, and were off the 270 southern tip of South America and near the Falkland Islands (Fig.2). In contrast, results for 271 Northern species were highly seasonal: a) during summer, the highest value areas were off the 272 central coast of Brazil, whereas in the winter, these extended to oceanic waters in the mid-south Atlantic, b) considering the whole year, the highest value areas were coastal and pelagic waters 273 off northern Brazil (Fig.2). The zonation value increased with the species occurrence probability 274 (Fig. 3a,c,e) and with increased number of species (Fig. 3b,d,f). It means the high valued areas 275 276 are a good representation for the occurrence of species.

277 The candidate protected areas based on the Southern assemblage were concentrated in 278 the same area during summer, winter and year-round (Fig.4). The three temperate areas were 279 located south of Isla de los Estados and northwest of the Yaghan Basin. The candidate MPAs 280 for Southern seabirds overlapped in the north with the current Isla de los Estados MPA 281 (Argentinean Ecological and Provincial Reserve) and IBA (proposed to protect BBA and Sooty 282 Shearwater Ardenna grisea) and in the south with Southwest 33 IBA (proposed for GHA), but most areas of high value for Southern seabirds lacked any protection under current regimes 283 284 (Fig.4). On the other hand, the candidate protected areas for Northern seabirds were completely different among time-frames. The candidate summer area on North overlapped with part of the 285 286 proposed EBSA off southern Brazil (proposed due to occurrence of a strong upwelling) and the 287 IBA on Arquipélago dos Alcatrazes (proposed based on the occurrence of Magnificent 288 Frigatebirds Fregata magnificens and the Brown Boobies Sula leucogaster). Only a small 289 fraction of a designated MPA overlapped the key area identified for Northern seabirds in the 290 summer: the Litoral Centro Environmental Protection Area. During winter, the key area for 291 Northern seabirds was in pelagic waters, over the gyre northeast of the Rio Grande Rise, where 292 there is a large elevation of the seabed. Currently no EBSA or IBA has been proposed in this 293 region. Finally, the year-round key area for Northern seabirds was on the shelf slope and pelagic 294 waters off northern Brazil, reaching as far as an oceanic ridge, where there is a strong upwelling 295 from the Amazon River (Fig.4). Roughly, one-third of this area overlapped with the 296 Amazonian-Orinoco EBSA, proposed due to the enhanced marine productivity occurring within 297 this area (Fig.4). The overlap of our proposed key areas with existing MPAs was minimal (less 298 than 1% Table 2). The percentage of mIBAs and EBSAs overlapped by our key areas was also 299 minimal, however overlap of Northern areas with EBSAs was substantially greater than 300 Southern areas, while Southern Areas were more overlapped by proposed mIBAs than Northern 301 areas (Table 2).

302 Cost analysis

The use of cost constraints for Southern seabirds resulted in few spatial differences of values when using Fishing Density as a cost layer, but higher conservation values in pelagic areas shifted north when using Ship Density as a cost layer (Supporting Information Fig. S9). On the other hand, using Fishing Density as a cost layer for Northern seabirds displaced the zones of high values to spread south, and to spread to pelagic areas when Ship Density was the cost layer (Supporting Information Fig. S10).

The differences of both cost models were significant among Time-Frames ($\chi^2_{26,6222}$ = 4143.9, P<0.001). The higher changes in values for Southern occurred when we used Fishing Density as a cost layer during summer, for both No Cost and Cost models, for winter when using Ship Density No Cost model and for year when using Ship Density Cost model (Fig.5, Supporting Information Table S2). On the other hand, the change in values for Northern were higher for year-round for both No Cost and Cost models and for both Fishing and Ship Density cost layer, despite using Ship Density there was also a high change of value during summer forboth cost models (Fig.5, Supporting Information Table S2).

317 Discussion

318 In our study, we considered a modelling approach that distinguishes between ecological groups 319 and accounts for seasonality in distribution of pelagic seabirds, with a view to identify a more 320 representative network of key sites that would be more effective year-round MPAS. This 321 approach located areas where currently there are few designated or proposed MPAs, suggesting 322 that a community integrated approach can be an effective way to propose new pelagic MPAs 323 (Yorio, 2009; Ronconi et al., 2012; Thaxter et al., 2012) in addition to proposals which use one 324 or few attributes of one area, such as the presence of seamounts, reefs, or the occurrence of one focal species (Arcos et al., 2012; Grecian et al., 2012; Dunn et al., 2014). 325

326 High Valued Areas, Threats and Benefits

327 We placed the most important area for Southern seabird species within the southwest Atlantic at the southern Argentina shelf slope, and around the Falkland Islands. This area has long been 328 329 recognized as important for marine predators and conservation of biodiversity (Croxall & 330 Wood, 2002). This is reflected in the large number of proposed BirdLife International mIBAs 331 (BirdLife International 2015), and several designated coastal MPAs, including the large reserves 332 of Namuncurá and Isla de losEstados (Argentine National Parks; Marine Conservation Institute 333 2015). These waters are used by several and abundant seabird species during the breeding and non-breeding seasons, including birds from Patagonia (Yorio et al., 1999; Yorio, 2009), 334 Falkland Islands (Grémillet et al., 2000; White et al., 2002), South Georgia (Croxall & Wood, 335 2002; Croxall et al., 2005; Phillips et al., 2006a; Navarro et al., 2015), Tristan da Cunha and 336 337 Gough (Ronconi, Ryan & Ropert-Coudert, 2010; Reid et al., 2013), and New Zealand (Nicholls 338 et al., 2002). Marine mammals also rely on this area during part of the year, including Fin 339 (Balaenoptera physalus), Sei (B. borealis), Minke (B. acutorostrata), Sperm (Physester 340 macrocephalus), Southern Bottlenose (Hyperoodon planifrons) and Long-finned Pilot 341 (Globicephala melas) whales, Hourglass (Lagenorhynchus cruciger), Peale's (L. australis) and

Commerson's (Cephalorhynchus commersonii) dolphins, Fur Seals (Arctocephalus spp.) and 342 343 Southern Elephant Seals (Mirounga leonina) (White et al., 2002). The attractiveness of this area 344 for marine top predators results from the extensive, year-round upwelling that extends from sub-345 Antarctic to temperate waters, which supports a rich food web, including diverse communities 346 of abundant squid and fishes (Acha et al., 2004; Miloslavich et al., 2011). As a consequence of 347 such rich communities, both fishing vessels and seabirds target those areas (Grémillet et al., 348 2000, 2008; Wakefield, Phillips & Belchier, 2012). In fact we found a higher "exposition" of seabird to fisheries, which is well-known for the study area (Xavier et al., 2004; Bugoni et al., 349 2008; Jiménez et al., 2010, Krüger et al., 2016a), and elsewhere (Anderson et al., 2011; 350 Lewison et al., 2014). In the other hand, the vessel traffic is a potential source of threat for 351 352 marine fauna, by oil (Halpern et al., 2008, 2015; Hatch et al., 2008) and noise (Morton & 353 Symonds, 2002; Weilgart, 2007; Codarin et al., 2009) pollution.

354 The different time-frame approaches resulted in completely different value outputs for 355 Northern seabird species, which reflects the seasonality of species present in the area throughout 356 the year. This is mostly a consequence of migratory species such as Deserta's Petrel, Trindade 357 Petrel and Cory's Shearwater. It also reflects the different types of habitats targeted by these 358 species. The highly valued area during summer supports a highly diverse coastal ecosystem, 359 including corals reefs (Roberts et al., 2002) and mangroves (Polidoro et al., 2010), and a 360 relatively narrow shelf with several seamounts (Miloslavich et al., 2011), supporting several 361 taxa, particularly sharks, cetaceans and euphausiids (Tittensor et al., 2010), and is intensively used by the species in this study, as evidenced by other authors (Dias et al., 2011; Catry et al., 362 363 2013; Dias, Granadeiro & Catry, 2013; Ramírez et al., 2013; Missagia et al., 2015; Krüger et al., 2016a). During winter, the most important areas matched with an area of low biodiversity 364 (Tittensor et al., 2010) and low productivity near the South Atlantic tropical gyre. Short-term 365 366 decreases in chlorophyll concentration within gyres due to climate shifts are a potential issue of concern, with implications for management (Gregg, Casey & McClain, 2005; Polovina, Howell 367 & Abecassis, 2008; Irwin & Oliver, 2009). This area also overlaps with the non-breeding 368

369 distribution of the Critically Endangered Tristan Albatross (Reid et al., 2013), and the at-sea 370 distribution during the breeding season and immediate post-breeding period of the Vulnerable 371 Trindade Petrel (Krüger et al., 2016b). The important areas year-round matched the upwelling 372 where the Amazon and Orinoco river plumes reach seamounts and islands, such as Fernando de 373 Noronha and São Pedro and São Paulo (Kitchingman et al., 2008; Miloslavich et al., 2011). It is 374 important to emphasize that the high valued (99%) year-round areas reflected more the suitable 375 habitat for the species than the bulk of species distribution (i.e. Gonzáles-Solís et al., 2009, Ramírez et al., 2013, 2015), despite the lower values (0.90) embraced those areas presented by 376 377 those authors. However the 99% valued area is recognized as biologically important for the 378 intense upwelling, which supports a biodiverse ecosystem (Tittensor et al., 2010; Miloslavich et 379 al., 2011; Selig et al., 2014), so highliting the value of this area for conservation, which is used 380 less frequently by the seabird species in this study

381 Many of our study species ingest large amounts of plastic debris (Petry & Fonseca, 2002; Jiménez et al., 2015), and the sensitivity of the Northern species to Plastic pollution 382 383 evidences that. The drifting model used in this study (Van Sebille et al., 2012) assumes that 384 plastic particles are carried by currents and accumulate in zones of lower current speed, mainly 385 gyres (Van Sebille, 2015). Titmus & David Hyrenbach (2011) found that some seabird species 386 target those same areas of plastic concentration, because they tend to be characterised by a high 387 occurrence of squid and flying fish (Titmus & Hyrenbach, 2011; Wilcox et al., 2015). Our 388 results provide circumstantial supporting evidence, but highlight the need for more studies to evaluate the overlap of seabird distribution with floating plastics, and the associated risk. 389 390 Similarly, direct evaluations of the effects and implications of spatial variation in rates of ocean 391 acidification for seabirds are lacking. Acidification is linked to enhanced levels of CO₂ in the atmosphere released by human activities (Cao & Caldeira, 2008). Severe decreases in pH may 392 393 affect lower trophic levels in food webs (Cao & Caldeira, 2008; Pörtner & Peck, 2010; Hale et al., 2011), with consequent impacts on top predators (Grémillet & Boulinier, 2009). 394

The assemblage approach to identify candidate MPAs revealed important areas missed in previous efforts to define both EBSAs (Dunn *et al.*, 2014) and BirdLife International mIBAs (Lascelles *et al.*, 2012; Ronconi *et al.*, 2012), although in other respects there was some overlap. As the currently designated MPAs are all coastal (Marine Conservation Institute 2015), our analyses of tracking data from pelagic seabirds showed almost no overlap with areas that are presently protected by national legislation.

401 **Potential Costs**

402 We found an assemblage and time-frame differential change in costs that suggests that the use 403 of seasonal protected areas could mean less impact on human activities to achieve high 404 conservation value as proposed by Hyrenbach, Forney & Dayton (2000). For instance, by 405 changing the fishing density by half, the changes for values within the key Southern areas did 406 not change substantially in relation to the no-cost model. On the other hand, for Northern areas, 407 the changes in value were relatively higher when applying cost layers, and for all the cost models the non-seasonal approach always resulted in a higher change in values. That's probably 408 409 a result of the higher seasonality for the species occurrence in the northern areas, whereas the 410 occurrence of species in the south off Patagonia and subantarctic waters is more constant 411 throughout the year. Leathwick et al. (2008) also showed that changes in conservation value of an area compared between models with and without use of cost layers can vary from 5% to 412 10%, but in our case the mean differences could reach as high as 50% for Northern seabirds. 413

414 Conclusions

By assigning seabirds to different assemblages, we were able to detect additional candidate areas for protection not recognised in previous marine spatial planning initiatives for FAO Region 41 in the southwest Atlantic Ocean. Hence, our results help guiding conservation decisions at both national and international levels, in terms of potential new MPAs and implementation of those already designated or proposed as EBSAs or mIBAs. Our analysis support the network of areas proposed as mIBAs in sub-Antarctic waters; the ultimate goal of BirdLife International is that these areas receive statutory protection (BirdLife International

422 2015). We also call attention for the need to delineate protected areas in pelagic tropical waters
423 both within the Brazilian EEZ, and in international waters. Those unprotected regions are
424 important seasonally or year-round, and are subject to a wide array of Human-related threats.

While in most cases the establishment of MPAs is aimed at improving management of fishing activities (Pitchford *et al.*, 2007; Stevenson *et al.*, 2013) several other factors may threaten species and ecosystems. Human activities, such as fishing, ship traffic and oiling, may be controllable to a large extent if there is a means of enforcing compliance with effective management regimes, but other threats associated with dynamic features of natural systems, such as ocean acidification and plastic pollution, present considerable challenges in the design and implementation of MPA networks (Conroy *et al.*, 2011).

The application of our approach to other oceans should reveal new areas to be incorporated in conservation networks. The benefits elsewhere would be highly dependent on the degree of seasonality in predator distributions, but it seems likely there are parallel situations where areas are only used for part of the year by migratory species. We envisage that this technique can be used on finer temporal (e.g. incubating versus chick rearing periods) and spatial (e.g. by the use of GPS or PTT devices) scales.

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Table 1 Seabird species included in the study. Smoothing cell size corresponds to the number of grid cells used for distribution smoothing in Zonation software for the analysis of habitat connectivity, and reflects the minimum area of habitat required for a species to occur. Cell sizes differ according to body size, following the approach of Leathwick et al (2008). Each species was assigned to one of two assemblages based on its core distribution relative to the mean position of the northern boundary of the Subtropical Front (Burls and Reason 2006): Temperate (TE) and tropical (TR) species. IUCN Status: LC=Least Concern, NT=Near Threatened, VU=Vulnerable, EN=Endangered, CR=Critically Endangered. Colony of the tracked animals is presented. Number of tracked individuals (N), tracked years and source of the tracking data.

CommonName	Species	Abbreviatio n	Size Range (cm)	Assemblag e	IUCN Status	Colony	N	Years	Source
Tristan Albatross	Diomedea dabbenena	ТА	110	Northern	CR	Gough Island	34	2004, 2005, 2006	Reid et al (2013)
Wandering Albatross	Diomedea exulans	WA	120-135	Southern	VU	Bird Island (South Georgia)	18	2003	Mackley et al (2010)
Black-browed Albatross	Thalassarche melanophris	BBA	80-96	Southern	NT	Bird Island (South Georgia), New Island (Falklands)	57	1996, 1997, 2002, 2003	Grémillet et al (2000), Mackley et al (2010)
Grey-headed Albatross	Thalassarche chrysostoma	GHA	70-85	Southern	EN	Bird Island (South Georgia)	35	2003, 2006	Croxallet al (2005)
Northern Giant Petrel	Macronectes halli	NGP	80-95	Southern	LC	Bird Island (South Georgia)	25	1999, 2001	González- Solís et al (2008)
Southern Giant Petrel	Macronectes giganteus	SGP	85-100	Southern	LC	Bird Island (South Georgia), Elephant Island (South Shetlands)	40	1999, 2001, 2011	González- Solís et al (2008), Krüger et al (2016)
White-chinned Petrel	Procellaria aequinoctialis	WCP	51-58	Southern	VU	South Georgia	11	2003, 2004	Phillips et al (2006)
Antarctic Prion	Pachyptila desolata	AP	25-27	Southern	LC	South Georgia	10	2009, 2010	Quillfeldt et al (2013)
Deserta's Petrel	Pterodroma deserta	DP	35	Northern	VU	Bugio Island	24	2007, 2008, 2009,	Ramírez et al (2013)

Trindade Petrel	Pterodroma arminjoniana	ТР	37-40	Northern	VU	Trindade Island	4	2013,201 4	Krüger et al (2016.)
Cory's Shearwater	Calonectris borealis	CS	46	Northern	LC	Berlenga, Azores and Canary Islands	41	2002, 2003, 2004, 2011, 2012	González- Solís et al (2009), Missagia et al (2016)
Great Shearwater	Ardenna gravis	GS	46-51	Southern	LC	Gough Island	37	2008- 2012	González- Solís
Manx Shearwater	Puffinus puffinus	MS	30-35	Southern	LC	Heimeaey, Iceland	10	2006,200 7	González- Solíset al (2009)
Cape Verde Shearwater	Calonectris edwardsii	CVS	34	Northern	NT	Cape Verde Isalnds	26	2006,200 7	González- Solíset al (2009)

Assemblage		IBA confirmed	IBA proposed	EBSA	MPAs
South	Overlap with	0.058	1.195	0.000	0.000
North		0.000	0.382	3.559	0.002
South	Overlapped by	0.053	8.397	0.000	0.026
North		0.000	1.139	15.345	0.037

Table 2 Percentage of proposed key areas overlapping and overlapped by the IBAs, EBSAs and current MPAs.

List of Figures

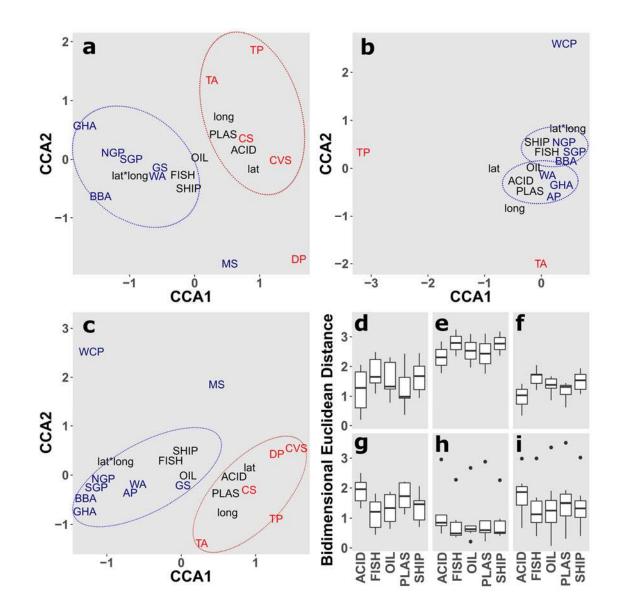
Figure 1 Canonical Correspondence Analysis bi-plots presenting the position of Northern species (red labels), Southern species (blue labels) and environmental – threats and coordinates – (black labels) for Summer (a), Winter (b) and all Year (c). Boxplots presenting the distribution frequency of the bi-dimensional distance of species from threats for Northern species during Summer (d), Winter (e) and all Year (f), and Southern species during Summer (g), Winter (h) and all Year (i). Top outliers in 'h' and 'i' are White-chinned Petrel values.

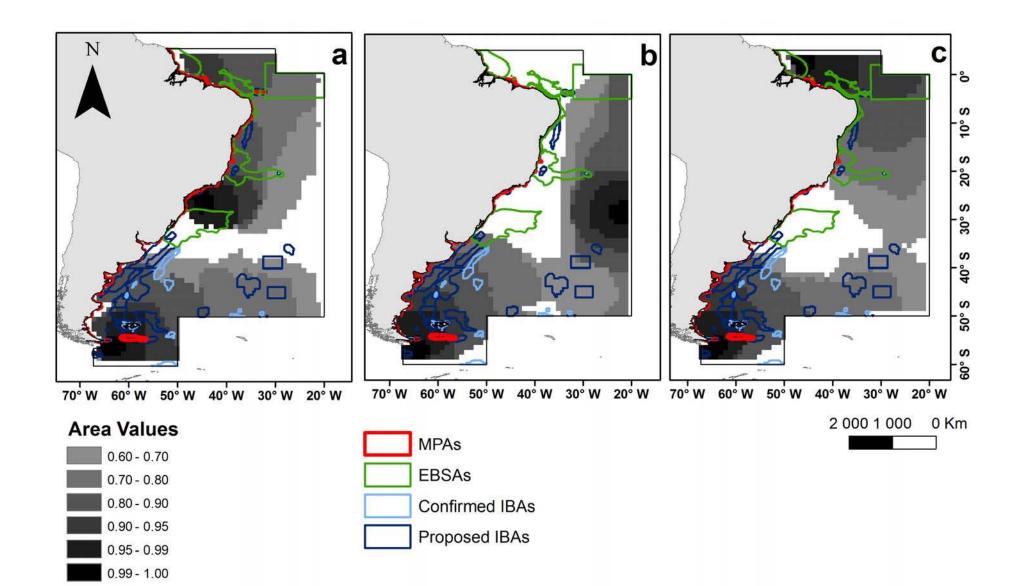
Figure 2 Zonation outputs in proportion to value based on the probability of occurrence of species given their weight and response to fragmentation for Summer (a), Winter (b) and yearround (c). Also shown the Official Marine Protected Areas (MPAs, red lines), Ecologically or Biologically Significant Areas (EBSAs; green lines; https://www.cbd.int/ebsa/) and confirmed (light blue lines) or proposed (dark blue lines) BirdLife marine IBAs (http://maps.birdlife.org/marineIBAs/default.html).

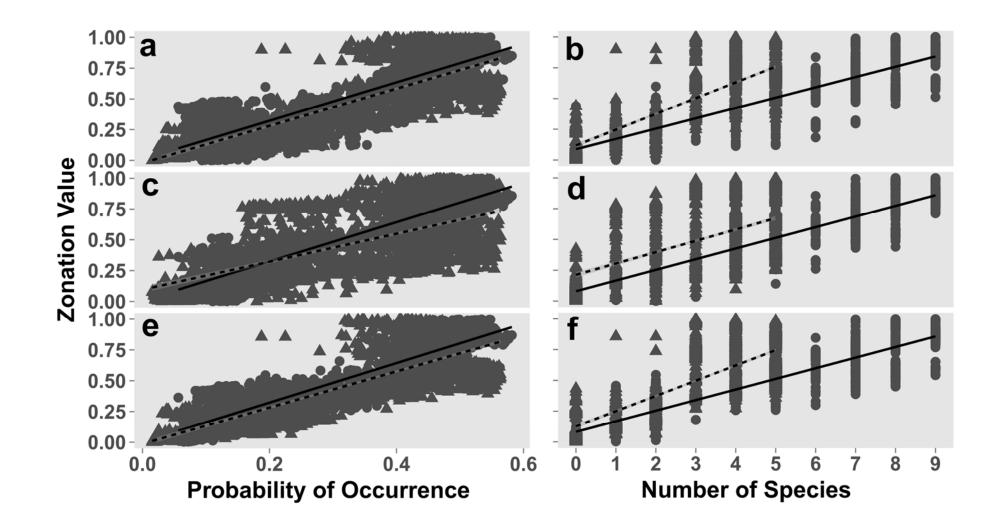
Figure 3 The zonation calculated area value in relation to the Species Occurrence Probability (a,c,e) and number of species occurrence (b,d,f). Linear trend for Southern (solid line) and Northern (dashed line) seabird assemblages.

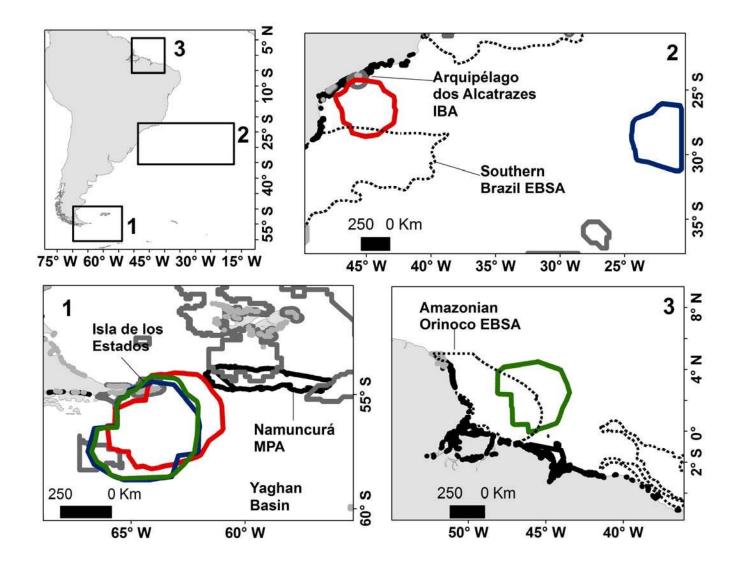
Figure 4 Key marine areas for 99% value threshold overlapped with Confirmed (light grey) or Proposed (dark grey) IBAs (http://maps.birdlife.org), Ecologically or Biologically Significant Marine Areas (EBSAs; dashed line; https://www.cbd.int/ebsa/) and designated MPAs (black line; http://www.mpatlas.org). Southern (1), Northern (2 and 3) seabirds during summer (red line), winter (blue line) and all year (green line).

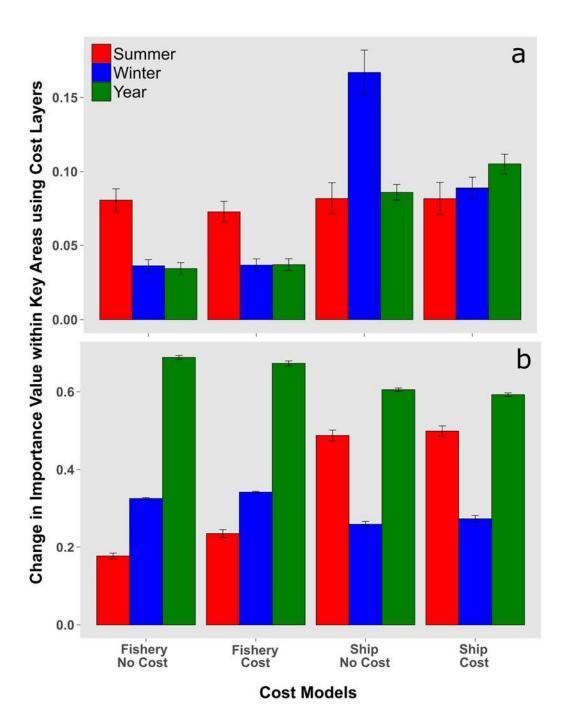
Figure 5 Change in values (proportion of reduction) within the proposed marine areas (thresholds 90%, 95% and 99%) when using Fishing and Ship Densities as cost layers to calculate importance values in Zonation. Southern (a) and Northern (b) seabirds.











Electronic Supplementary Material for Animal Conservation research article:

Identification of candidate pelagic marine protected areas through a seabird seasonal-, multispecific- and extinction risk-based approach

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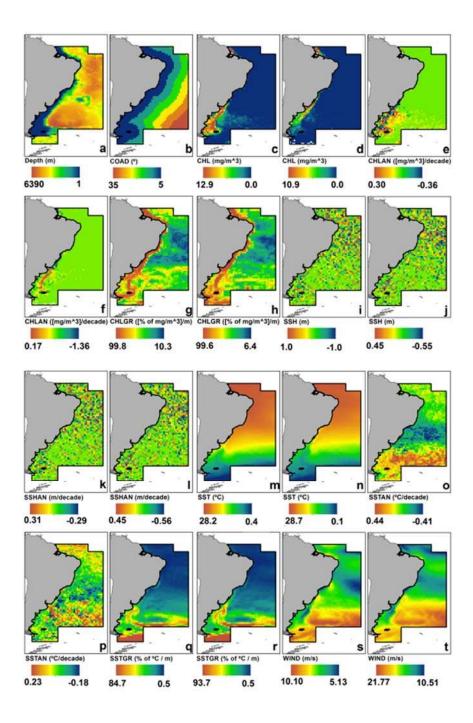


Figure S1 Environmental variables used for the Species Distribution Modelling: (a) depth in meters; (b) Distance to coast (COAD) in degrees; chlorophyll-*a* concentration (CHL) in mg/m³, during (c) summer and (d) winter; chlorophyll-a concentration anomaly CHLAN in decade variability of mg m⁻³, during (e) summer and (f) winter; CHL gradient (CHLGR) in percentage of mgm⁻³ variation per meter during (g) summer and (h) winter; sea surface height SSH in meters during (i) summer and (j) winter; sea surface height anomaly (SSHAN) in variability of meters per decade during (k) summer and (l) winter; sea surface temperature SST in °C during (m) summer and (n) winter; sea surface temperature anomaly (SSTAN) in variability of °C per decade during (o) summer and (p) winter; sea surface temperature gradients in percentage of °C change per meter during (q) summer and (r) winter; wind speed in meters per second during (s) summer and (t) winter. Year variables were the average of both summer and winter.

SI Sensitivity analysis of the Zonation outputs when using different weighting values

The aim of this section is to provide an analysis of the method we used to weight species distribution by the IUCN conservation status, and the habitat connectivity response (in Zonation software called as *Boundary-Quality Penalty Curves BQPCs*). We used the full weight method that is in the main text to give value to the year-round distribution of all species.

Full model: Each species was weighted by its IUCN conservation status as follows, least concern (LC) = 1; near threatened (NT) = 2; vulnerable (VU) = 3; endangered (EN) = 4 and critically endangered (CR) = 5. Boundary Quality Penalty Curves, which are measures of species responses to habitat connection, were constructed empirically based on the IUCN conservation status. We generated linear responses with changing slopes for CR = -0.01, EN = -0.008, VU = -0.004, NT = -0.002 LC = 0.0.

We also generated three other different weight methods: a null model where the weight given for all species is the same (1) and the response to habitat connection (β^h) is constant; a model where LC weights 1 with β^h constant and the other levels (NT, VU, EN, CR) weights 2 with β^h = -0.002 (weight model #1), a model where LC weights 1 with β^h constant, NT weights 2 with β^h = -0.002, and other levels (VU, EN, CR) weight 3 with β^h = -0.004 (weight model #2). Then we tested via simple regression how these three methods are similar to the full model and tested how the probability of species occurrence is related to those weighting outputs through cubic regression.

We found that a null model with equal weight and no β^h variation means that the software will give more value to pixels isolated from the edges of the study area, without being realistic nor reflecting any oceanographic process intrinsic to the species distribution, but probably embracing the edges of most species distribution (Fig. S5). It means that high values in this case were concentrated in the middle of the study area (Fig. S5) and were completely opposed to the output we proposed (F_{1,1652} = 3.29, β = -0.14, P=0.07). On the other hand, one single differentiation on the weights in two (F_{1,1652} = 31.18, β = 0.18, P<0.001) and three (F_{1,1652} = 1556.6, β = 0.8, P<0.001) groups was enough to approximate the values to the full model output (Fig. S6).The null model weakly reflected the overall distribution of species (R²=0.04, F_{1,1652} = 24.55, P = 0.09), while the other weight models #1 (R² = 0.53, F_{1,1652} = 613.6, P<0.001), #2 (R² = 0.63, F_{1,1652} = 952.2, P<0.001) and the full model (R² = 0.69, F_{1,1652} = 1238.0, P<0.001) did significantly explained the species distribution (Fig. S7).

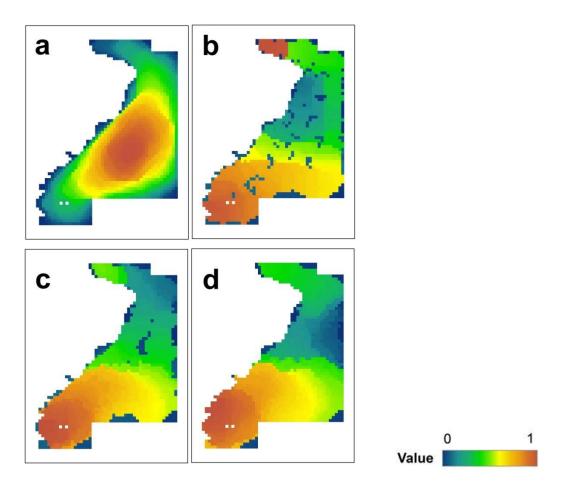


Figure S2 Zonation output showing the how much the values attributed to each pixel change by the weighting method used in building the (a) null model, (b) weight model #1, (c) weight model #2 and (d) the full model.

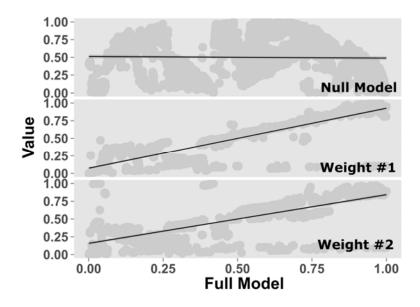


Figure S3 Regression between the full model and the new weight models. Lines are linear trends \pm SE.

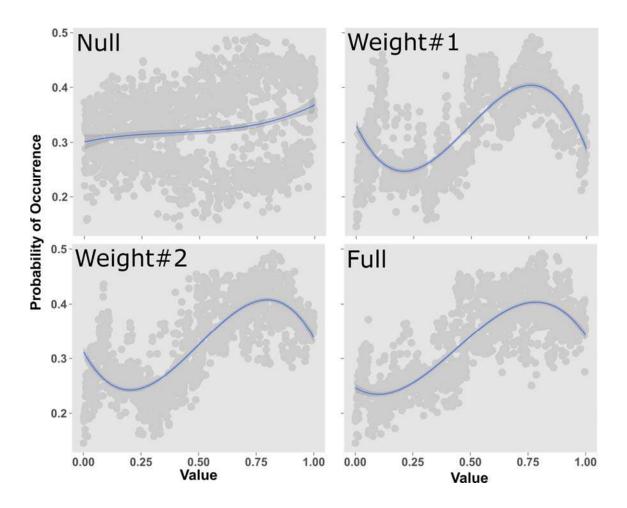


Figure S4 Relation between the overall species probability of occurrence and zonation value for each of the four models. Lines are cubic trend \pm SE.

SII Mehods for calculating environmental threat variables.

A total of five threat variable were used: ocean surface acidification, oil pollution risk, plastic accumulation, ship density and fishing intensity (Fig. S8). All environmental variables were standardized to have a mean of 0 and an SD of 1 due to differing ranges of variables. Such standardization was executed by dividing each absolute value by the respective maximal value of the environmental variable (Zuur, Ieno & Smith, 2007).

Ocean surface acidification was measured in terms of normalized changes in aragonite saturation state between pre-industrial and modern times, which is a compound that enhances when Ph decreases (Halpern *et al.*, 2008). The acidification image was generated by Halpern *et al.*(2008), and we are using it as it is in the publication.

For risk of oil pollution we used the areas of oil accumulation proposed by Halpern *et al.*(2008). A Nearest Natural Neighbor Interpolation was used to calculate a risk based on the distance from these pollution areas, being the cells presented by Halpern *et al.*(2008) with value 1.0, and decreasing according to the distance.

Plastic accumulation was estimated using a dynamic particle flotation model based on movement of buoys (van Sebille, England & Froyland, 2012; van Sebille *et al.*, 2015; van Sebille, 2014). The model takes in account the coastal population density as a proxy for source of pollution, and predicts the probability of the movements of those particles due to ocean currents for each two month periods along a $1^{\circ}x1^{\circ}$ spatial grid. As there are different outputs if we use different periods of the year as the start of the modeling, we run models for all the two month periods along ten years. Each 2 month period output images were used to calculate an average image. This means that for cells with higher values there is a greater probability of plastic accumulation accounting for all possible scenarios. Thus the final images accounted also for the movement of the particles instead of only the final destination of the particles at the end of the 10-year period. We also considered a model with a constant particle release and a single release in time, to make the model more realistic, and calculated a mean of both.

Ship density was computed from the ship density shapefiles made available by the PASTA-MARE project (LuxSpace, 2010) which takes in account three month Satellite Automatic Identification System S-AIS movement of 62000 vessels on a global scale. The technical report (LuxSpace, 2010) shows a similarity between their results and two longer term data bank of the Voluntary Observing Ship from the World Meteorological Organization WMO-VOS (Halpern *et al.*, 2008) which uses one year data of 3374 commercial and research vessels, and Automated Mutual-assistance Vessel Rescue system AMVER that uses 3809 commercial vessels. We believe that the PASTA-MARE project is more accurate in terms of characterizing the ship density per area in relative terms because it uses all the available data for

any type and flag of vessels, while the more long-term data for VOS and AMVER are probably biased for the more representative types of vessels. We correlated the Halpern *et al.* (2008) with the PASTA-MARE results using a spatial correlation analysis on Spatial Analysis in Macroecology SAM software (Rangel, Diniz-Filho & Bini, 2010), and found a significant adjustment between both (R=0.61, F=28.8, P<0.001) meaning that the short term data from PASTA-MARE is a fair approximation for a year-long data bank. Fishing intensity was measured as the fishing vessel density from the PASTA-MARE data (LuxSpace, 2010).

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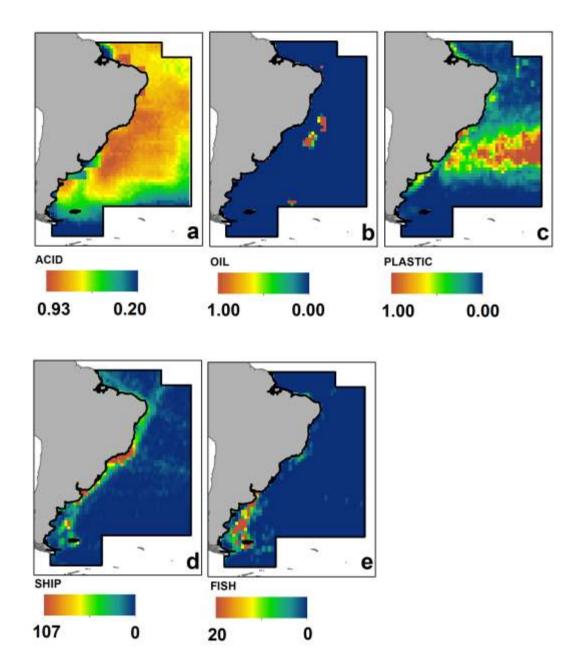


Figure S5 Environmental change variables: (a) ocean surface acidification (ACID) measured in terms of normalized changes in aragonite saturation state between pre-industrial and modern times (see Halpern et al 2008); (b) Risk of oil pollution (OIL) measured as a distance probability given the known points of oil spills or oil platforms from Halpern et al (2008) as value 1; (c) 10-year mean percentage of plastic pollution (PLASTIC) based on the particle drift model (Van Sebille et al 2012 2015); (d) ship density (SHIP) as the average abundance of vessels per grid cell (LuxSpace 2010); (e) fishing intensity (FISH) is a measure of density of fishing vessels.

Table S1 Loadings from the PCA. The values represent the proportion of each variable that is explained by each of the PCA axis, and the signal (- or +) indicates the relation of the variables with the axes. Variables with higher correlation on the same axis are not independent from each other, meaning that they vary together. Table presents results from the axis whose eigenvalues were above 1.0. Higher values for each variable by season are presented in bold. Chlorophyll (CHL), CHL anomaly (CHLAN), CHL gradient (CHLGR), coast distance (COAD), Depth (DEPTH), Sea Surface Height (SSH), SSH anomaly (SSHAN), Sea Surface Temperature (SST), SST anomaly (SSTAN), SST gradient (SSTGR), wind speed (WIND).

Variables	Summer				Winter				Year			
	AX1	AX2	AX3	AX4	AX1	AX2	AX3	AX4	AX1	AX2	AX3	AX4
CHL	0.12	-0.64	0.10	-0.17	-0.03	0.60	0.00	0.57	0.01	-0.65	-0.06	-0.48
CHLAN	0.11	0.05	0.01	-0.98	0.09	-0.21	0.04	-0.77	0.12	0.22	0.01	0.85
CHLGR	0.43	-0.70	0.07	-0.01	0.57	0.67	0.06	-0.07	0.46	-0.74	-0.03	0.15
COAD	0.24	0.72	-0.03	0.02	0.25	-0.73	-0.10	0.34	0.31	0.70	0.03	-0.35
DEPTH	-0.13	0.87	-0.07	-0.08	-0.17	-0.83	-0.08	0.11	-0.08	0.86	0.04	-0.11
SSH	-0.03	0.09	0.99	0.01	-0.01	-0.09	0.99	0.05	-0.01	0.06	-0.95	0.00
SSHAN	-0.04	0.10	0.99	0.01	-0.01	-0.10	0.99	0.05	-0.01	0.07	-0.95	0.03
SST	-0.94	-0.03	-0.01	-0.08	-0.94	0.00	-0.02	0.05	-0.93	0.06	0.00	-0.02
SSTAN	0.75	0.16	0.01	-0.04	0.26	-0.16	-0.06	0.42	0.72	0.12	-0.03	-0.11
SSTGR	0.84	-0.03	0.00	0.09	0.85	0.08	0.01	-0.10	0.83	-0.12	0.01	0.09
WIND	0.83	0.23	0.00	0.00	0.78	-0.41	-0.03	0.10	0.82	0.29	0.01	-0.10

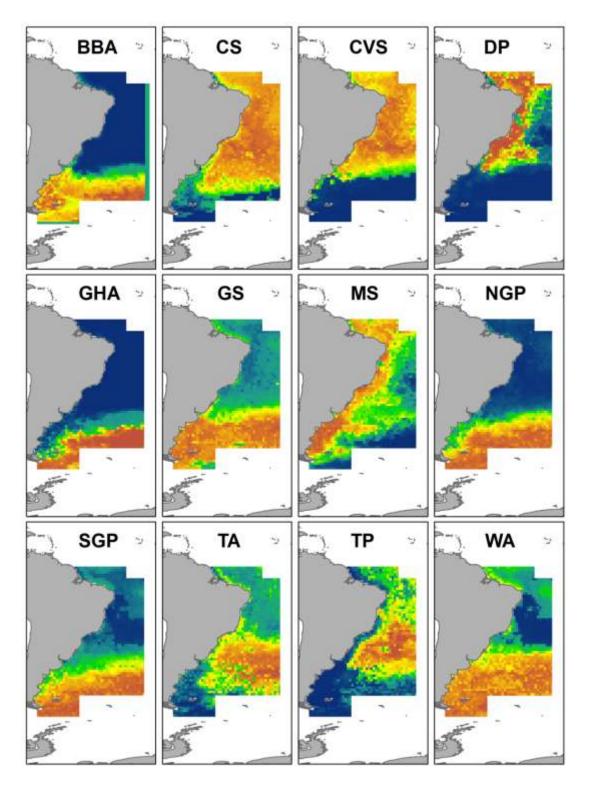


Figure S6 Summer probability of occurrence of Black-browed Albatross (BBA), Cory's Shearwater (CS), Cape Verde Shearwater (CVS), Deserta's Petrel (DP), Grey-headed Albatross (GHA), Great Shearwater (GS), Manx Shearwater (MS), Northern Giant Petrel (NGP), Southern Giant Petrel (SGP), Tristan Albatross (TA), Trindade Petrel (TP), Wandering Albatross (WA), as estimated by MaxEnt models. Probability varies from 0 (dark blue) to 1 (dark red).

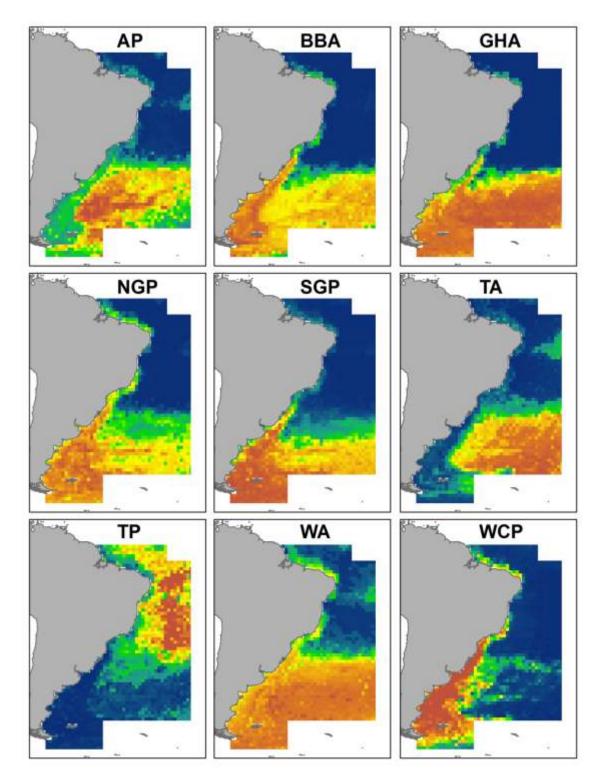


Figure S7 Winter probability of occurrence of Antarctic Prion (AP), Black-browed Albatross (BBA), Grey-headed Albatross (GHA), Northern Giant Petrel (NGP), Southern Giant Petrel (SGP), Tristan Albatross (TA), Trindade Petrel (TP), Wandering Albatross (WA), Whitechinned Petrel (WCP), as estimated by MaxEnt models.Probability varies from 0 (dark blue) to 1 (dark red).

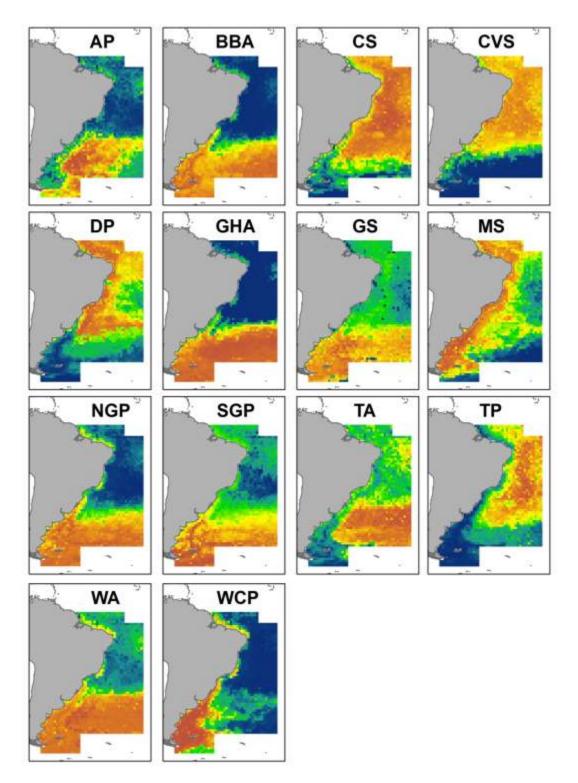


Figure S8 Year-round probability of occurrence of Antarctic Prion (AP), Black-browed Albatross (BBA), Cory's Shearwater (CS), Cape Verde Shearwater (CVS), Deserta's Petrel (DP), Grey-headed Albatross (GHA), Great Shearwater (GS), Manx Shearwater (MS), Northern Giant Petrel (NGP), Southern Giant Petrel (SGP), Tristan Albatross (TA), Trindade Petrel (TP), Wandering Albatross (WA), White-chinned Petrel (WCP), as estimated by MaxEnt models. Probability varies from 0 (dark blue) to 1 (dark red).

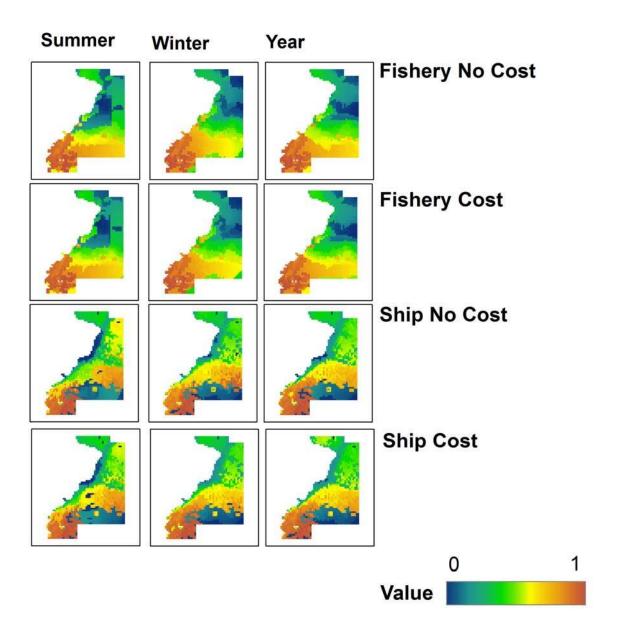


Figure S9 Zonation value for the different cost models, for southern seabirds.

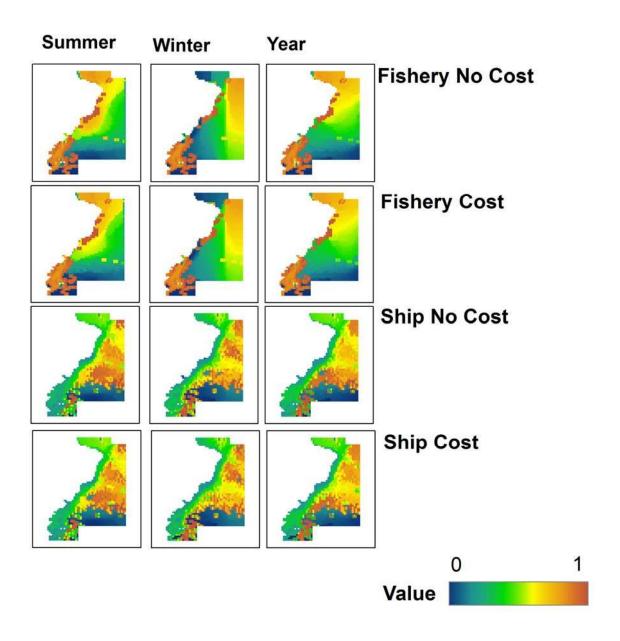


Figure S10 Zonation value for the different cost models models, for northern seabirds.

Models	β	SE	t
(Intercept)	0.0913	0.0142	6.44
Assemblage:Northern	0.0966	0.0113	8.58
Time-Frame:Winter	-0.0441	0.0113	-3.92
Time-Frame:Year	-0.0460	0.0113	-4.09
Cost-Model:Fishery Cost	-0.0078	0.0113	-0.7
Cost-Model:Ship No Cost	0.0015	0.0113	0.13
Cost-Model:Ship Cost	0.0014	0.0113	0.13
Assemblage:Northern* Time-Frame:Winter	0.1936	0.0162	11.93
Assemblage:Northern:* Time-Frame:Year	0.5585	0.0159	35.06
Assemblage:Northern* Cost-Model:Fishery Cost	0.0667	0.0159	4.19
Assemblage:Northern* Cost-Model:Ship No Cost	0.3094	0.0159	19.42
Assemblage:Northern* Cost-Model:Ship Cost	0.3206	0.0159	20.13
Time-Frame:Winter* Cost-Model:Fishery Cost	0.0083	0.0159	0.52
Time-Frame:Year* Cost-Model:Fishery Cost	0.0105	0.0159	0.66
Time-Frame:Winter* Cost-Model:Ship No Cost	0.1292	0.0159	8.11
Time-Frame:Year* Cost-Model:Ship No Cost	0.0502	0.0159	3.15
Time-Frame:Winter* Cost-Model:Ship Cost	0.0513	0.0159	3.22
Time-Frame:Year* Cost-Model:Ship Cost	0.0694	0.0159	4.36
Assemblage:Northern* Time-Frame:Winter* Cost-Model:Fishery Cost	-0.0509	0.0229	-2.22
Assemblage:Northern* Time-Frame:Year* Cost-Model:Fishery Cost	-0.0847	0.0225	-3.76
Assemblage:Northern* Time-Frame:Winter* Cost-Model:Ship No Cost	-0.5062	0.0229	-22.07
Assemblage:Northern* Time-Frame:Year* Cost-Model:Ship No Cost	-0.4450	0.0225	-19.76
Assemblage:Northern* Time-Frame:Winter* Cost-Model:Ship Cost	-0.4255	0.0229	-18.55
Assemblage:Northern* Time-Frame:Year* Cost-Model:Ship Cost	-0.4885	0.0225	-21.69

Table S2 Generalized Linear Mixed Model results comparing change in values for the proposed key marine areas between seabird assemblages (Southern and Northern) and time-frames (summer, winter, year) when using different cost models (no cost and cost for Fishing Density and Ship Density).