



HAL
open science

Meeting Paris agreement objectives will temper seabird winter distribution shifts in the North Atlantic Ocean

M. Clairbaux, W. Cheung, P. Mathewson, W. Porter, N. Courbin, Jérôme Fort, H. Strøm, B. Moe, P. Fauchald, S. Descamps, et al.

► To cite this version:

M. Clairbaux, W. Cheung, P. Mathewson, W. Porter, N. Courbin, et al.. Meeting Paris agreement objectives will temper seabird winter distribution shifts in the North Atlantic Ocean. *Global Change Biology*, 2021, 27, pp.1457-1469. 10.1111/gcb.15497 . hal-03088572

HAL Id: hal-03088572

<https://hal.science/hal-03088572>

Submitted on 28 Oct 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

1 **Meeting Paris agreement objectives will temper seabird winter distribution**
2 **shifts in the North Atlantic Ocean**

3 M. Clairbaux^{1*}, W. Cheung², P. Mathewson³, W. Porter³, N.Courbin¹, J. Fort⁴, H. Strøm⁵, B.
4 Moe⁶, P. Fauchald⁶, S. Descamps⁵, H. Helgason⁵, V.S. Bråthen⁶, B. Merkel⁵, T. Anker-
5 Nilssen⁶, I.S. Bringsvor⁷, O. Chastel⁸, S. Christensen-Dalsgaard⁶, J. Danielsen⁹, F. Daunt¹⁰, N.
6 Dehnhard⁶, K.E. Erikstad^{6, 11}, A. Ezhov¹², M. Gavrilov^{13, 14}, Y. Krasnov¹², M. Langset⁶, S.H.
7 Lorentsen⁶, M. Newell¹⁰, B. Olsen⁹, T.K. Reiertsen⁶, G. Systad⁶, T.L. Thórarinsson¹⁵, M.
8 Baran¹⁶, T. Diamond¹⁶, A.L. Fayet¹⁷, M.G Fitzsimmons¹⁸, M. Frederiksen¹⁹, H.G. Gilchrist²⁰,
9 T. Guilford¹⁷, N.P. Huffeldt²¹, M. Jessopp²², K.L. Johansen¹⁹, A.L. Kouwenberg²³, J.F
10 Linnebjerg¹⁹, L. McFarlane Tranquilla²³, M. Mallory²⁴, F.R. Merkel¹⁹, W. Montevecchi²⁵, A.
11 Mosbech¹⁹, A. Petersen²⁶, D. Grémillet^{8, 27}

12 ¹ CEFE, Univ Montpellier, CNRS, EPHE, IRD, Univ Paul Valéry Montpellier 3, Montpellier,
13 France

14 ² Changing Ocean Research Unit, Institute for the Oceans and Fisheries, The University of
15 British Columbia, Vancouver, B.C., V6T 1Z4, Canada

16 ³ Department of Integrative Biology, University of Wisconsin, Madison, WI, USA

17 ⁴ Littoral, Environnement et Sociétés (LIENSs), UMR7266 CNRS - La Rochelle Université, 2
18 rue Olympe de Gouges, 17000 La Rochelle, France

19 ⁵ Norwegian Polar Institute, Fram Centre, P.O. Box 6606 Langnes, 9296 Tromsø, Norway

20 ⁶ Norwegian Institute for Nature Research - NINA, PO Box 5685 Torgarden, NO-7485
21 Trondheim, Norway

22 ⁷ Norwegian Ornithological Society, Sandgata 30 B, NO-7012 Trondheim, Norway

23 ⁸ Centre d'Etudes Biologiques de Chizé (CEBC), UMR 7372 CNRS-Univ. La Rochelle,
24 France

25 ⁹ Faroe Marine Research Institute, P.O. Box 3051, Nóatún 1, FO-110 Tórshavn, Faroe Islands

26 ¹⁰ UK Centre for Ecology & Hydrology, Bush Estate, Penicuik EH26 0QB, UK

27 ¹¹ Centre for Biodiversity Dynamics, Department of Biology, Norwegian University of
28 Science and Technology, 7491 Trondheim, Norway

29 ¹² Murmansk Marine Biological Institute, 17 Vladimirskaia street, 183010 Murmansk, Russia

30 ¹³ Association Maritime Heritage, Saint Petersburg, Russia

31 ¹⁴ National Park Russian Arctic, 57 Sovetskikh Kosmonavtove Avenue, Archangelsk, Russia

32 ¹⁵ Northeast Iceland Nature Research Centre, Hafnarstétt 3, 640 Húsavík, Iceland

33 ¹⁶ Atlantic Laboratory for Avian Research, University of New Brunswick, P.O. Box 4400,
34 Fredericton NB, Canada E3B 5A3

35 ¹⁷ Department of Zoology, University of Oxford, South Parks Road, Oxford OX1 3PS, UK

36 ¹⁸ Wildlife Research Division, Environment and Climate Change Canada, 6 Bruce St., Mount
37 Pearl, NL, Canada A1N 4T3

38 ¹⁹ Aarhus University, Dept. of Bioscience, Frederiksborgvej 399, 4000 Roskilde, Denmark

39 ²⁰ National Wildlife Research Centre, Environment and Climate Change Canada, Ottawa

40 ²¹ Greenland Institute of Natural Resources, Kivioq 2, 3900 Nuuk, Greenland & Aarhus
41 University, Dept. of Bioscience, Frederiksborgvej 399, 4000 Roskilde, Denmark

42 ²² School of Biological, Earth and Environmental Sciences, University College Cork,
43 Distillery Field, North Mall, Cork, Ireland

44 ²³ Bird Studies Canada, Sackville, New Brunswick, Canada

45 ²⁴ Biology, Acadia University, 15 University Avenue, Wolfville, Nova Scotia, Canada, B4P
46 2R6

47 ²⁵ Psychology and Biology Departments, Memorial University of Newfoundland, St. John's,
48 NL A1C 5S7, Canada

49 ²⁶ Brautarland 2, 108 Reykjavik, Iceland

50 ²⁷ Percy FitzPatrick Institute, DST/NRF Centre of Excellence, University of Cape Town,
51 Rondebosch, South Africa

52

53 * Corresponding author: clairbauxm@gmail.com

54

55

56

57

58

59

60

61

62

63

64

65

66

67

68 **Abstract**

69 We explored the implications of reaching the Paris Agreement Objective of limiting
70 global warming to $<2^{\circ}\text{C}$ for the future winter distribution of the North Atlantic seabird
71 community. We predicted and quantified current and future winter habitats of five North
72 Atlantic Ocean seabird species (*Alle alle*, *Fratercula arctica*, *Uria aalge*, *Uria lomvia* and
73 *Rissa tridactyla*) using tracking data for ~1500 individuals through resource selection
74 functions based on mechanistic modeling of seabird energy requirements, and a dynamic
75 bioclimate envelope model of seabird prey. Future winter distributions were predicted to shift
76 with climate change, especially when global warming exceed 2°C under a “no mitigation”
77 scenario, modifying seabird wintering hotspots in the North Atlantic Ocean. Our findings
78 suggest that meeting Paris agreement objectives, will limit changes in seabird selected habitat
79 location and size in the North Atlantic Ocean during the 21st century. We thereby provide key
80 information for the design of adaptive marine protected areas in a changing ocean.

81 **Keywords:** DBEM, Energy requirement, Mechanistic habitat selection, NicheMapper TM,
82 Paris Agreement, RCP scenarios, Seabird migration, Seabird distributions

83

84

85

86

87

88

89

90 The main target of the 2015 Paris Agreement under the United Nations Framework
91 Convention on Climate Change (UNFCCC) is to limit temperature increase to less than 2°C
92 above preindustrial levels at the global scale (United Nations Framework Convention on
93 Climate Change (UNFCCC), 2015). Recent studies have demonstrated the potential benefits
94 of this target for marine ecosystems (Sumaila et al., 2019; Trisos et al., 2020) but its relevance
95 and our capacity to reach the Paris Agreement Objectives (PAO) are still being debated
96 (Gasser, Guivarch, Tachiiri, Jones, & Ciais, 2015). Notably, there are few estimates of these
97 benefits for marine top predators (but see Jenouvrier et al., 2020) despite their key roles in
98 marine ecosystem functioning and their support to cultural ecosystem services
99 (Hammerschlag et al., 2019). It is therefore essential to examine the possible effects of
100 climate change on these species and the interest of avoiding such climate impacts.

101 Seabirds are among the most threatened of all bird groups (Dias et al., 2019), with
102 demonstrated sensitivity to direct (through physiological modifications or change in extreme
103 events exposure) and indirect (through trophic mechanisms or modifications of their critical
104 habitats) climate change impacts (Sydeman, Poloczanska, Reed, & Thompson, 2015).
105 Seabirds are also ecological sentinels of marine ecosystems across their life cycles (Durant et
106 al., 2009; Lescroël et al., 2016), and the subject of long-term monitoring studies throughout
107 the world (Paleczny, Hammill, Karpouzi, & Pauly, 2015). Yet, because of technological
108 limitations and practical difficulties, most of these studies dealing with climate change
109 impacts on seabirds focus on population processes (Descamps et al., 2017) or on their
110 responses during the breeding season (Frederiksen, Anker-Nilssen, Beaugrand, & Wanless,
111 2013), rather than climate change impacts on their at-sea distributions during the non-
112 breeding period. This is critical since conditions encountered during the non-breeding period
113 strongly shape seabird population dynamics (Alves et al., 2013). Knowledge of climate effects
114 on seabird wintering distributions is therefore essential for global marine spatial planning and

115 conservation schemes. Community-wide analyses allowing direct comparison between
116 species and providing a more complete picture of climate change impacts are then particularly
117 valuable.

118 In this context, we explored the implications of succeeding or failing to reach the Paris
119 Agreement Objective of limiting global warming to $<2^{\circ}\text{C}$ on the North Atlantic Ocean seabird
120 community distribution, during the non-breeding period (October-February). We used Global
121 Location Sensors (GLS) to track the inter-breeding movements of five species across the
122 North Atlantic Ocean and adjacent seas (see the map of the studied area in Supplemental
123 materials I) and then modeled seabird energy requirement and seabird prey fields for each
124 location. Mechanistic Resource Selection Functions (RSF) allowed us to determine how the
125 balance between prey availability and energy requirements explained seabird winter habitat
126 selection at the time of our investigations. Based on those mechanistic RSFs, we predicted
127 current seabird community distributions during winter and determined the location and the
128 size of selected habitats for each species before projecting them to the time periods 2045-2055
129 and 2090-2100. To this aim, we used two greenhouse gas concentration pathways: the
130 Representative Concentration Pathway RCP2.6 and RCP8.5, which correspond to low and
131 high radiative forcing, respectively. While RCP8.5 is often qualified as a “no mitigation”
132 scenario, RCP2.6 assumes strong mitigation policies under which global warming is projected
133 to be on average $<2^{\circ}\text{C}$ relative to preindustrial levels (Meinshausen et al., 2006; Vuuren et al.,
134 2011). Overall, comparing current and future distributions and size of selected habitats, we
135 tested the hypothesis that limiting greenhouse gas emissions will reduce community-level
136 seabird winter habitat change in the North Atlantic Ocean.

137

138

139 **Methods**

140 *Winter geolocation of the North Atlantic Ocean seabird community*

141 We focused on the five species which represent >75% of the total number of seabirds
142 breeding in the North Atlantic Ocean and adjacent seas (Barrett et al., 2006): little auk (*Alle*
143 *alle*), Atlantic puffin (*Fratercula arctica*), Brünnich's guillemot (*Uria lomvia*), common
144 guillemot (*Uria aalge*) and black-legged kittiwake (*Rissa tridactyla*). Despite the major focus
145 on the Alcidae seabird family and the Charadriiformes order, this sample included species
146 with a wide range of prey species and foraging behaviors (del Hoyo, Elliott, & Sargatal,
147 1996). Global Location Sensors (GLS) recording year-round locations were deployed and
148 retrieved on 1532 individuals from 39 breeding colonies across the North Atlantic Ocean
149 during summer fieldwork (see Supplemental materials XII for details). Each GLS tag
150 recorded light levels, which were used to calculate two locations per day with an accuracy of
151 +/- 200 km (Lisovski et al., 2012; Phillips, Silk, Croxall, Afanasyev, & Briggs, 2004).
152 Different models of GLS tags were used, and the raw light data were analyzed with
153 corresponding software (see Supplemental materials XII for detail); some of these data were
154 previously analyzed in other studies (Amélineau et al., 2018; Fayet et al., 2017; Fort et al.,
155 2013; Fort et al., 2013; Frederiksen et al., 2016, 2011; Linnebjerg et al., 2013; McFarlane
156 Tranquilla et al., 2015; Merkel, 2019; Montevecchi et al., 2012; Tranquilla et al., 2013). We
157 kept locations acquired during the winter period (October-February) and erroneous locations
158 obtained during the two weeks around either side of the equinoxes (Lisovski et al., 2012)
159 and/or those falling on land or outside of the study area (100°W–100°E, 30°N–90°N) were
160 removed.

161

162

163 *Modeling current and future energyscapes*

164 We used the mechanistic model Niche Mapper TM (Porter & Mitchell, 2006) to model the
165 current and future energy expenditures for each species across the North Atlantic Ocean basin
166 for each wintering month (October-February). This model contains two sub-modules: a
167 microclimate model, which provides environmental data for the near surroundings of the
168 animal at each location, and an animal module, which uses outputs from the microclimate
169 model, together with animal morphological, behavioral and physiological characteristics.
170 These inputs are used to solve heat balance equations between the animal's body and its
171 environment, and to find the metabolic rate needed for the animal to maintain its body
172 temperature at the particular time with the considered behavior.

173 Environmental data (sea surface temperature, air temperature, wind speed, cloud cover and
174 relative humidity) used to parameterize the microclimate model were selected from the
175 outputs of the Earth system model HadGEM2-ES (Collins et al., 2011; Martin et al., 2011).
176 This climatic model includes different components (troposphere, land surface and hydrology,
177 aerosols, ocean, sea ice, terrestrial carbon cycle and ocean biogeochemistry) which are
178 coupled on a 24h time-step with an atmospheric resolution of 1.875°x1.25° and an oceanic
179 resolution of 1° (increasing to 1/3° at the equator) (Collins et al., 2011). We used HadGEM2-
180 ES outputs because empirical data were not available for all of our study areas, precluding the
181 calculation of global energyscapes (spatialized energy requirements). Also, HadGEM2-ES has
182 been shown to perform well when simulating recent and past mechanisms such as deep
183 convection in North Atlantic Ocean (Heuzé, 2017), North-West Atlantic Ocean physical
184 process (Lavoie, Lambert, ben Mustapha, & Baaren, 2013), Arctic sea ice melt (Wang &
185 Overland, 2012) and North-East Atlantic Ocean climatology (Perez, Menendez, Mendez, &
186 Losada, 2014; Zappa, Shaffrey, & Odges, 2013). Relative humidity was calculated following

187 Nadeau & Puiggali (1995) and Nayar et al. (2016). Daily data were averaged to obtain
188 monthly means and were interpolated in a 1° grid.

189 As Niche Mapper™ has already been parameterized to model energy expenditure in little
190 auks, Brünnich's guillemots and common guillemots (Fort et al., 2013; Fort, Porter, &
191 Grémillet, 2009), we re-used most Niche Mapper™ input values from these previous studies.
192 Missing values and values required to parameterize Niche Mapper™ for black-legged
193 kittiwakes and Atlantic puffins, were sourced from the literature (see Supplemental materials
194 XIII) and supplemented with dedicated measurements. Notably, feather reflectivity was
195 measured with a double beam spectrophotometer (CARY 5000 UV-VIS-NIR, Agilent, USA)
196 with a deuterium-tungsten-mercury light source. We used an integrative sphere to measure
197 spectral and diffuse reflectance with a 1 nm resolution across all wavelengths between 300
198 and 2500 nm. This range covers approximately 98% of the solar spectrum that reaches the
199 Earth's surface. Reflectance spectra relative to a Spectralon white standard were then
200 computed with the Cary WinUV software. For each species, measurements were made on one
201 ventral and dorsal patch for six individuals. We then calculated the reflectivity of each sample
202 following the same method as in Medina et al (Medina et al., 2018). For each species the
203 results were averaged across patches.

204 Other morphological properties such as leg length, as well as head and body diameters, were
205 measured on bird carcasses of five Atlantic puffins and four black-legged kittiwakes. Since
206 Niche Mapper has relatively low sensitivity with respect to the variables concerned by such
207 measurements (Amélineau et al., 2018; Fort et al., 2009), potential biases induced by using
208 such measurements on carcasses instead of live organisms had limited impacts on our results.

209 All input data are available in Supplemental materials XIII. Main drivers of the modeled
210 energy requirements were identified through sensitivity analysis in previous studies
211 (Amélineau et al., 2018; Fort et al., 2009).

212 *Modeling current and future seabird prey abundance*

213 We identified the main winter prey of each seabird species using bibliographic information
214 (see Supplemental materials XIV for details). Little auks and black-legged kittiwakes are
215 mainly zooplanktivorous during winter. Other studied species are assumed to lower their
216 trophic level outside the breeding season but could still prey on Nereid worms and on various
217 benthopelagic/pelagic fish such as Atlantic herring (*Clupea harengus*), capelin (*Mallotus*
218 *villosus*), Atlantic cod (*Gadus morhua*) or lesser sand-eel (*Ammodytes marinus*).

219 We modeled the relative abundance of fish species to 1950 across the North Atlantic Ocean,
220 using a Dynamic Bioclimate Envelope Model (DBEM) (Cheung et al., 2016, 2009). The
221 DBEM simulates changes in abundance of marine fishes since 1950 according to
222 oceanographic conditions. For each fish species considered, habitat suitability was modeled
223 according to temperature, salinity, dissolved oxygen concentration, bathymetry and sea-ice
224 extent. These environmental preferences were then linked to a population model which takes
225 into account the carrying capacity of the habitat, growth, mortality, and larval and adult
226 dispersion. DBEM outputs are annual averages but, considering the very large spatial scale
227 and resolution of our study, we assumed that relative abundance patterns were maintained at
228 the scale of months and that the winter period changes can be reflected in the annual
229 projections.

230 Fish characteristics required as inputs to the DBEM model (see Cheung et al., 2016) for
231 details) were obtained from the Sea Around Us catch database (<http://www.seaaroundus.org>)
232 and from FishBase (<https://www.fishbase.se/search.php>). Environmental data needed for
233 current and future modeling (see Supplemental materials XV for details) were obtained from
234 the Earth system model HadGEM2-ES (Collins et al., 2008; Martin et al., 2011). We used
235 HadGEM2-ES because empirical environmental data were not available at the right spatial
236 and temporal scales. Equally, since empirical information on current and monthly spatial and

237 temporal abundances of each zooplankton species taken by seabirds was not available, we
238 used the monthly total concentration of zooplankton provided by the ocean biogeochemical
239 model, Diat-HadOCC, included within the Earth system model HadGEM2-ES to model the
240 exchange of carbon dioxide between the atmosphere and the oceanic biosphere (Totterdell,
241 2019). Zooplankton concentration is one of the 13 biogeochemical state variables represented
242 by Diat-HadOCC, and is modeled as advected by ocean currents and mixed by physical
243 processes. Zooplankton biomass is enhanced by grazing of miscellaneous phytoplankton,
244 diatoms and detritus, and diminished by respiration and by density-dependent predation
245 through higher trophic levels.

246 Finally, due to missing species-specific physiological data, the relative abundance of several
247 fish species could not be calculated with DBEM (see Supplemental materials XIV for the list).
248 According to FishBase, those fish are zooplanktivorous and are mainly benthopelagic. We
249 therefore used bathymetry (using General Bathymetric Chart of the Oceans (GEBCO), 30 arc-
250 second interval grid) and zooplankton concentration as a proxy of their availability to diving
251 seabirds. All seabird prey data were interpolated in a 1° grid.

252 *Habitat selection of different seabird species*

253 To analyze seabird habitat use and predict current and future winter distributions, we used
254 Resource Selection Functions (RSF) (Boyce & McDonald, 1999; Manly, McDonald, Thomas,
255 McDonald, & Erickson, 2002). RSF compare environmental variables at locations used by the
256 animal with those variables at a set of locations available to the animal (Manly, McDonald,
257 Thomas, McDonald, & Erickson, 2007). In our case, RSF compared for each species the
258 energy expenditure and prey availability at GLS locations with those expected at a set of
259 random points generated in our study area (100°W–100°E, 30°N–90°N). The random points
260 were created using the same temporal distribution as the GLS data. RSF were fitted using
261 generalized linear mixed models with a binomial error distribution and a logit link. To take

262 into account individual heterogeneity and variation between colonies, we used a weighted
263 logistic regression with random intercepts (individual and colony) and random slopes, with
264 weight=1 for locations used and weight=1000 for available locations (Muff, Signer, &
265 Fieberg, 2019). To avoid power issues that arise with random intercept, individuals with
266 fewer than 100 locations (threshold obtained after a kernel stability analysis conducted with
267 the BRB-MKDE software (Benhamou, 2011), March 2018 version) for the entire winter
268 period were not included in the model.

269 For each species, we built a RSF model with month, energy expenditure and relative
270 abundance of each prey as covariates. When computational issues or collinearity problems
271 arose, we reduced model complexity by selecting the relative abundance of the prey most
272 often eaten by the seabird species considered. We tested for the trade-off between main prey
273 abundance and energy requirement by adding an interaction term between the two
274 predictors. Moreover, when considered as seabird prey, zooplankton concentration was linked
275 to sea surface temperature through an interaction term to allow selection of zooplankton
276 species associated with cold or warm water. Finally, when bathymetry was considered as an
277 availability proxy for some benthopelagic prey, we linked it to zooplankton concentration
278 through an interaction term. We allowed non-linear effects for each covariate via a spline
279 function. RSF models are detailed in Supplemental materials II.

280 Variables used in our final RSF analysis were scaled and were not correlated as tested with a
281 Pearson pairwise correlation test and a variance inflation factor analysis ($VIF < 3$ in all
282 models). We evaluated each model using block cross-validation (Roberts et al., 2017) with
283 80% of the data to fit the RSF and 20% to test it. The corresponding output (rated “k-fold rs”
284 in Supplemental materials II) is based on the Spearman’s rank correlation (Boyce, Vernier,
285 Nielsen, & Schmiegelow, 2002) and varies between 0 (low predictive performance) and 1

286 (good predictive performance). RSF analyses were performed in R (version 3.5) with the
287 glmmTMB package (Brooks et al., 2017).

288 For each species, we mapped RSF results after splitting them in 10 quantile bins (Morris,
289 Proffitt, & Blackburn, 2016). More than 60% of the GLS locations occurred in the 4 last bins,
290 and we therefore defined selected habitat as pixels with a binned RSF score ≥ 7 . To identify
291 wintering hotspots, we then summed, for each pixel of the map, the number of species for
292 which the pixel considered was classified as a selected habitat. Finally, the area occupied by
293 selected habitats were calculated.

294 *Current and future predictions*

295 Current conditions were modeled with outputs from the HadGEM2-ES algorithm, according
296 to the historical extended simulation (1950-2018). To assess the benefit of reaching the PAO
297 and limiting temperature increase to the 2°C target at the global scale, environmental
298 variables and models (see above) were considered under Intergovernmental Panel on Climate
299 Change (IPCC)'s RCP2.6 and RCP8.5 scenarios at the medium term (2045-2055) and the
300 longer term (2090-2100). The RCP2.6 scenario is based on a low radiative forcing and
301 assumes that strong mitigation policies will limit global greenhouse gas emissions and reach
302 the goal of negative emissions in the second half of the 21st century. Stabilization at this low
303 level of emissions will limit global mean temperature increase to 2°C above preindustrial
304 levels with high probability (Meinshausen et al., 2006). According to the HadGEM2-ES
305 model, the global mean temperature increase will reach this 2°C threshold just before mid-
306 century before slowly decreasing (Caesar et al., 2013). In contrast, the RCP8.5 scenario
307 follows a high radiative forcing trajectory with a continuous increase in global greenhouse gas
308 emissions across the 21st century. According to the HadGEM2-ES model, global mean
309 temperature will increase $>2.8^{\circ}\text{C}$ above preindustrial levels in 2050, and $>6^{\circ}\text{C}$ in at the end of
310 the 21st century (Caesar et al., 2013).

311 **Results**

312 Predictors of seabird distributions and current community wintering hotspots

313 All Resource Selection Functions (RSF) had high predictive power of species
314 distributions (see “k-fold rs” in Supplemental materials II). Mechanistic RSF showed that
315 seabirds selected winter habitat by balancing two major constraints: prey availability and their
316 own energy requirements (see Figure 1 and Supplemental materials III). This trade-off was,
317 nonetheless, best predicted by seabird energy requirements for all studied species (see
318 Supplemental materials III).

319 **Figure 1:** The trade-off between zooplankton concentration and bird energy requirements predicts
320 little auk habitat selection. An optimum was reached at a scaled energy requirement of -0.6 and a
321 scaled zooplankton concentration of 1.3. See Supplemental materials III for other species.

322 Species-specific selected habitat was defined using a threshold approach on the developed
323 RSF’s (see Methods for details). These habitats were in turn combined to identify multi-
324 species wintering hotspots (sensu (Fort, Beaugrand, Grémillet, & Phillips, 2012; Grecian et
325 al., 2016)) by counting for each pixel of the map, the number of species for which the pixel
326 considered was classified as a selected habitat. Given seabird energy requirements and prey
327 availability (see Supplemental III and IV), birds were predicted to use areas off southwestern
328 Iceland and along the Norwegian coast between October and November, while the Gulf of
329 Maine was predicted to be a major wintering hotspot for the selected seabird community
330 between December and February (Figure 2 and Supplemental materials V). At the beginning
331 of the winter period, the southern coast of Greenland and the Barents Sea were predicted to be
332 important areas for the seabird community. Yet, between December and February, the
333 HadGEM2-ES climatic model showed scarce zooplankton resources to seabirds at those high-
334 latitudes (see Supplemental materials IV), and the areas were then predicted to be used only
335 by the two guillemot species (Figure 2 and Supplemental materials V). Further, the area off
336 Newfoundland and Nova Scotia was predicted to be a wintering hotspot between October and

337 January, but the composition of the seabird community varied with the month considered:
338 little auks and black-legged kittiwakes were joined by Atlantic puffins between October and
339 December, and by Brünnich's guillemots between November and January (Figure 2 and
340 Supplemental materials V). Finally, the Mid-Atlantic Ocean and the North Sea were predicted
341 to be selected by black-legged kittiwakes, little auks and Atlantic puffins between November
342 and February, but those species were also predicted to aggregate in the eastern Celtic Sea in
343 January and in the Bay of Biscay in February (Figure 2 and Supplemental materials V).

344 **Figure 2:** Count of species for which the pixel considered was a selected habitat (RSF binned score
345 ≥ 7), in October (a), November (b), December (c), January (d) and February (e). Graticules are
346 displayed at 15° intervals and the map is projected as North Pole Lambert Azimuthal Equal Area.

347 Climate change impacts on seabird distribution and selected habitat size

348 Modeled spatial trends of future selected habitats for seabird were generally similar under
349 RCP2.6 and RCP8.5 scenarios. However, the magnitude of projected impacts on seabird
350 biogeography was generally more pronounced under the higher CO₂ emission scenario
351 (RCP8.5) and towards the end of the 21st century (Figure 3 and 4, Supplemental materials X
352 and XI). Moreover, our projections showed that climate change is predicted to modify prey
353 availability and seabird energy requirements (see Supplemental materials VI and VII),
354 inducing loss and gain of selected habitats and resulting in a general northward shift of
355 wintering areas (see Figure 3 and Supplemental materials VIII and IX).

356 **Figure 3:** Selected habitat (in purple) of Brünnich's guillemots predicted for January in 2006-2015
357 (a) and in 2090-2100 according to the RCP2.6 scenario (b) and the RCP8.5 scenario (c). Maps for
358 other months, species and time periods (2045-2055) are provided in Supplemental materials VIII and
359 IX. Graticules are displayed at 15° intervals and the map is projected as North Pole Lambert
360 Azimuthal Equal Area.

361 **Figure 4:** Count of species for which the pixel considered is predicted to be a selected habitat in
362 January 2006-2015 (a) and in January 2090-2100 according to the RCP2.6 scenario (b) and the
363 RCP8.5 scenario (c). Maps for other months and time periods (2045-2055) are provided in
364 Supplemental materials X and XI. Graticules are displayed at 15° intervals and the map is projected as
365 North Pole Lambert Azimuthal Equal Area.

366

367 In October, all five seabird species were predicted to be present in the Labrador Sea and south
368 of Greenland. The Barents Sea was predicted to remain an important seabird wintering site in
369 October, but its attractiveness for black-legged kittiwakes, Brünnich's guillemots and little
370 auks was predicted to diminish as global warming intensifies. Further, in November, seabirds
371 were projected to shift towards areas off southern Greenland and in the Labrador Sea, where
372 conditions were predicted to improve in terms of energy requirements and prey availability
373 (see Supplemental materials VI and VII). Thereby, black-legged kittiwakes and little auks
374 were expected to progressively abandon the Norwegian coast. At the seabird community
375 level, marine areas east off Newfoundland and the Gulf of St Lawrence were predicted to
376 become increasingly important as wintering hotspots (see Figure 4 and Supplemental
377 materials X and XI). All considered species were predicted to aggregate in these areas in
378 December, January and February, where the birds will probably benefit from lower energy
379 requirements relative to present (see Supplemental materials VI and VII). During these
380 months black-legged kittiwakes, little auks and Atlantic puffins were predicted to continue to
381 winter in the mid-Atlantic Ocean, but the corresponding hotspots were predicted to shift
382 northward/westward. The North Sea was predicted to become a preferred habitat in January,
383 except under the RCP8.5 scenario at the end of the 21st century. For this time period and
384 scenario, seabird energy requirements were predicted to remain unchanged in the North Sea,
385 but prey availability was predicted to diminish (see Supplemental materials VII), leading to a
386 decrease in overall habitat quality.

387 The projected biogeographic shifts will significantly impact seabird winter habitat sizes (see
388 Figure 5). Selected habitats size was predicted to increase for common guillemots (for
389 example, by 21% and 47% by 2045-2055 and up to 38% and 102% by 2090-2100, in January
390 according to RCP2.6 and RCP8.5 scenarios, respectively). Between 2045 and 2055, selected
391 habitat of Brünnich's guillemots was predicted to increase slightly (up to 32% in October

392 according to the RCP2.6 scenario) but the persisting temperature increase will probably limit
393 this expansion and even lead to a reduction of selected habitat size for the RCP8.5 scenario.
394 Change in selected habitat size was predicted to be limited for the Atlantic puffin (+/- 9% or
395 less regardless of the time period and climatic scenario). For black-legged kittiwakes,
396 predicted selected winter habitat will initially show little change in size (+/- 9% or less)
397 between 2045 and 2055, except in October, during which the range will expand by ca. 20%
398 according to both scenarios. Later in the 21st century, our results nonetheless suggest a strong
399 reduction (up to 63% in October but around 22% for the rest of the winter period) in kittiwake
400 winter habitat for the RCP8.5 scenarios. Finally, for little auks, winter habitat range was
401 generally predicted to decline as global warming increases, except in October between 2045
402 and 2055 when it will increase slightly relative to today.

403 **Figure 5:** Variation relative to present (in %) of the size of predicted selected habitat for (a) little
404 auks (b) black-legged kittiwakes (c) Atlantic puffins (d) common guillemots (e) Brünnich's
405 guillemots. The y-axis scales are different between panels.

406 **Discussion**

407 To our knowledge, our study is the first to investigate climate change impacts on the
408 winter distribution of a seabird community at an ocean basin scale with a mechanistic
409 approach. Our work predicts that global warming will likely induce substantial changes in
410 seabird prey field distributions and spatialized energy requirements (energyscapes, sensu
411 Amélineau et al., 2018). These two predictors of seabird habitat selection are predicted to
412 strongly shape the location and size of seabird selected habitats in the North Atlantic Ocean
413 during winter. Thereby, global warming will result in northward distribution shifts of varying
414 magnitude, depending on the seabird species and considered time period. More specifically,
415 areas such as Newfoundland, the Gulf of St Lawrence, and southern Greenland will become
416 increasingly attractive as seabird wintering hotspots because of diminishing seabird energy
417 requirements. Crucially, our broad-scale analyses confirms that meeting Paris Agreement

418 Objectives will limit habitat range shifts of the North Atlantic Ocean seabird community in
419 21st century in comparison with shifts induced by a “no-mitigation” scenarios.

420 Despite these advances, we recommend caution in interpreting our results. First, since direct
421 measurements of seabird prey availability and energy requirements are lacking at the spatio-
422 temporal scales needed for a community-wide, ocean-basin-scale study, we used a variety of
423 modeling tools to gain this information. Those entail potential approximations and biases
424 which could be difficult to assess. For example, as most commonly-used biogeochemical
425 models (Arhonditsis & Brett, 2004), the Diat-HadOCC biogeochemical model included
426 within HadGEM2-ES has not been evaluated on its ability to model zooplankton
427 concentration. However, considering its ability to model chlorophyll concentration, this model
428 seems to respect geographical distribution and seasonal cycle of the carbon cycle in the North
429 Atlantic Ocean (Totterdell, 2019). Further, due to some lack of information about seabird
430 winter diets, we extrapolated available dietary information, using those for birds from a
431 subsample of all colonies. In the same line, the availability of seabird prey was modeled
432 without taking into account the impacts of fisheries. By altering the difference in relative prey
433 abundance between locations used by seabirds and random points, fisheries could diminish
434 the role of prey availability in modelled seabird habitat selection. Moreover, the availability of
435 seabird prey was modeled on an annual basis and future works should add seasonal resolution
436 to the DBEM analysis. However, considering the very large spatial scale and resolution of our
437 study, we assumed that relative prey abundance patterns were maintained at the scale of
438 months, and that changes occurring during the winter period were adequately reflected by
439 annual projections. Further, we limited the complexity of RSF models, by only considering
440 seabird energy requirements and relative prey abundances, omitting the roles of abiotic factors
441 such as sea ice concentration or day length as well as biotic factors like competition, predation
442 or costs of transport (linked with migratory distance from the breeding ground). Notably, day

443 length may have an incidence on foraging times, although seabird have shown surprising
444 abilities to cope with the polar night (Ostaszewska, Balazy, Berge, & Johnsen, 2017).
445 Moreover, by modeling future distributions, we assume that the statistical link between
446 seabird distributions, their energyscapes and prey fields will hold across time (Grémillet &
447 Charmantier, 2010). Finally, our approach is based on model stacking, but allows a better
448 understanding and representation of ecological processes, translating interactions of
449 organismal functional traits with their environment into key fitness components throughout
450 the inclusion of mechanistic models into correlative Habitat Suitability Models. RSF analyses
451 yielded high predictive power and accuracy, providing support to this approach. In this
452 context, limitations of studies based on DBEM and NicheMapperTM have been previously
453 described and positively rated (Cheung et al., 2009; Fort, Porter, & Grémillet, 2011),
454 suggesting that our work can indeed provide valuable predictions.

455 *A mechanistic framework to predict wildlife distribution hotspots in a changing world*

456 On the basis of mechanistic and biodynamic model outputs, Resource Selection Functions
457 allowed us to predict seabird wintering habitats. We thereby elaborated a general
458 methodological framework which may be applied to the spatial ecology of any animal
459 tracking environmental gradients and resources. With respect to the North Atlantic Ocean
460 seabird community, we confirmed the great importance of areas such as the mid-Atlantic
461 Ocean (Afonso et al., 2020; Barrett et al., 2006; Bennison, Jessopp, Bennison, & Jessopp,
462 2015), the Barents Sea (Gabrielsen, 2009), the North Sea (Harris, Daunt, Newell, Phillips, &
463 Wanless, 2010), the Icelandic coast (Garðarsson, 1999) and southwest Greenland (Boertmann,
464 Lyngs, Merkel, & Mosbech, 2004), for wintering seabirds. We highlighted the importance of
465 seabird energy requirements as predictor of winter habitat selection (Figure 1). Prey
466 availability contributed to shape seabird distributions at finer spatial scales (Amélineau et al.,
467 2018), but we modeled that habitat choice generally resulted from a trade-off between energy

468 requirements and prey availability, and was predicted to be explained by the former (see
469 Figure 1 and Supplemental materials III). This result is in agreement with recent
470 macroecological studies (Somveille, Rodrigues, & Manica, 2018), and the magnitude of the
471 energy requirement/prey availability trade-off may explain marked inter-species differences in
472 seabird winter habitat dynamics under the influence of climate change. Yet, for all species
473 considered, global warming was predicted to induce a significant decrease in seabird winter
474 energy requirements following similar geographical features (see Supplemental materials VI
475 and VII). Therefore, observed inter-species differences appeared to be related to variation in
476 prey availability. For zooplanktivorous species, the decrease in energy requirements at high
477 latitude was not predicted to be accompanied by a zooplankton concentration increase, thus
478 limiting the northward expansion of their wintering habitat. However, as the northward shift
479 of zooplankton was sufficient to decrease the relative probability of selection at the southern
480 margin of their range, little auks and black-legged kittiwakes were predicted to experience an
481 overall decrease in their selected habitat size. Conversely, for both species of guillemots, the
482 decrease in energy requirements in the high Arctic was predicted to be accompanied by an
483 increase in the abundance of cod and capelin, thus greatly increasing the probability of habitat
484 selection in large areas such as the Russian Arctic. Further, decreases in the abundance of cod
485 and capelin in the southern part of guillemot ranges seemed to remain within a range of
486 favorable values. Those may not impact the relative probability of selection there, resulting in
487 an overall gain of selected habitat. Further work will nonetheless be necessary to fully
488 understand population consequences of these changes (Carneiro et al., 2020). Finally, the
489 greater amplitude of latitudinal shifts in the abundance of fish species, relative to zooplankton
490 concentration observed in our study, may be explained by the different temporal scales
491 (annual versus monthly) considered, and/or by the impacts of other factors beyond
492 temperature (such as day length or sea ice concentration) at high latitudes. Those factors may

493 have a greater impact on the northward expansion of zooplankton than on fish latitudinal
494 shifts.

495 *Seabirds tracking climate change*

496 Latitudinal shifts in response to climate change have been predicted and observed for many
497 marine organisms, including top predators (Cristofari et al., 2018), small pelagic fish (Perry,
498 Low, Ellis, & Reynolds, 2005) and zooplankton (Beaugrand et al., 2019). The High Arctic is
499 already colonized by many organisms which currently occur at lower latitudes (Fossheim et
500 al., 2015), and a series of endemic species might become extinct. Even though birds may
501 adjust their migratory behavior in response to climate change (Visser, Perdeck, van Balen, &
502 Both, 2009), seabirds generally seem to not shift their reproductive timing according to
503 warmer ocean temperatures (Keogan et al., 2018), and remain extremely faithful to their
504 breeding sites (Newton, 2010). Therefore, we speculate that the North Atlantic Ocean seabird
505 community might be under pressure from climate change consequences through two main
506 mechanisms. During the breeding season, site fidelity tends to expose seabird breeders to
507 spatio-temporal mismatches with their food base and lead them into ecological traps (Durant,
508 Hjernmann, Ottersen, & Stenseth, 2007; Grémillet et al., 2008). During the non-breeding
509 season, even though there might be some fidelity to wintering sites, we might predict that
510 seabirds will be more likely to track environmental conditions and shift spatio-temporally,
511 following a trade-off between their energy requirements and prey availability. As a
512 consequence, there is a strong potential for shifts in distances and/or directions between
513 seabird breeding and wintering sites, which will reshape migratory corridors and locations of
514 stopover sites (Newton, 2010; Clairbaux et al., 2019). Assessing and quantifying these
515 changes in response to global warming for each colony will provide a better understanding of
516 their magnitude, and of consequences for individual populations. Moreover, shifts in marine
517 ecosystem composition and structure caused by global warming likely require seabirds to

518 change their diets and behaviours (Amélineau et al., 2019). The ability of seabirds to cope
519 with such shifts, and the mechanisms (plasticity and/or microevolution) that underpin
520 potential responses, are difficult to assess without long-term studies based on individual
521 monitoring (Teplitsky & Charmantier, 2019). As long-lived organisms, we nonetheless expect
522 that seabirds are more likely to respond through plasticity, rather than microevolution, when
523 facing the consequences of climate change (Sauve, Divoky, & Friesen, 2019). Therefore,
524 reaching Paris Agreement Objectives will decrease the risk that distribution discrepancies
525 between seabirds and their prey arise and will also make it more likely that environmental
526 shifts remain within the limits of seabird plastic responses (Descamps et al., 2017).

527 *Conservation implications*

528 Global warming is one of the four most serious threats to seabirds, along with invasive
529 species, bycatch, and overfishing of prey species (Dias et al., 2019; Grémillet et al., 2018). All
530 of these challenges, which also compromise many other components of marine biodiversity,
531 have to be urgently addressed. Restoring marine life may be feasible by 2050 if major
532 pressures, such as global warming, are mitigated (Duarte et al., 2020) and, as our analysis
533 shows, reaching PAO will be an effective way to mitigate climate change impacts on the
534 North Atlantic Ocean seabird community. Our analyses provide new information and methods
535 to identify seabird wintering hotspots in the North Atlantic Ocean, and allow modeling future
536 spatio-temporal dynamics and community alteration under the threat of climate change. This
537 is key to defining and managing marine protected areas (MPAs), which have emerged as
538 powerful means to protect entire ecosystems from overfishing and effects of bycatch for
539 which seabirds function as ecological indicators and umbrella species (Pichegru, Grémillet,
540 Crawford, & Ryan, 2010). In practice, a first essential step towards defining MPAs using
541 seabird at sea-movement data is to map marine Important Bird and Biodiversity Areas
542 (Donald et al., 2019) (IBA). Several IBAs/MPAs have already been identified in the North

543 Atlantic Ocean (see, <http://datazone.birdlife.org/site/mapsearch> and
544 <http://www.mpatlas.org/map/mpas/> for respective maps), most of them in coastal areas, with
545 the exception of the MPAs recently established within the OSPAR Convention (Charlie
546 Gibbs, north of the Azores and Milne seamount complex areas). In this context, our analysis
547 features an important contribution to the upcoming marine IBA/MPA network, as we
548 predicted currently unprotected community-level seabird wintering hotspots off
549 Newfoundland, southern Greenland, Iceland and in the Barents Sea. Furthermore, our work
550 emphasizes that marine spatial planning involving IBAs/MPAs will have to be adaptive, with
551 protected areas adjusted according to species range shifts under the impact of global changes.
552 MPA benefits will be negatively impacted by climate change (Bruno et al., 2018; Trisos et al.,
553 2020), and thus reaching Paris Agreement Objectives will help to maintain their effectiveness
554 in the future. Our modeling work is a clear example of how community-level winter
555 distributions of marine top predators are likely to shift in the North Atlantic Ocean, and we
556 provide a complete methodological framework allowing the forecasting of such range shifts
557 according to different climate change scenarios. Overall, we emphasize the effectiveness of
558 Paris Agreement Objectives for the spatial management of seabird community under the
559 impact of climate change, and of dynamic MPAs defined using track-based data on at-sea
560 movements of seabirds and other marine top predators (Péron, Authier, & Grémillet, 2018).

561 **Acknowledgements**

562 We warmly thank the many fieldworkers involved in this study. Data collection in
563 Norway, including Svalbard and Jan Mayen, was supported by the Norwegian seabird
564 program SEAPOP (www.seapop.no). Thanks to the SEATRACK-program for making
565 tracking data available for the analyzes (<http://seatrack.seapop.no/map/>). Fieldwork on little
566 auks in Northwest Greenland was funded by the Bureau of Minerals and Petroleum,
567 Greenland Government. The New Brunswick Wildlife Trust Fund provided funding for GLS

568 tags on Machias Seal Island. We also thank the University of New Brunswick, the Federal
569 Government of Canada and the Government of Greenland and the Danish Ministry of
570 Environment and Food. Thanks to the Polar Continental Shelf Program of Natural Resources
571 Canada (Grant 213-08), core funding from Environment and Climate Change Canada. Thanks
572 to the French Polar Institute Paul Emile Victor for funding the ADACLIM program (IPEV
573 program 388 to D.Grémillet and J.Fort) and the ORNITHO-ENDOCRINO program (IPEV
574 program 330 to O.Chastel). We thank Simon Benhamou for his help with BRB-MKDE, Doris
575 Gomez, Christine Andraud and Aurélie Tournié for their valuable help regarding feather
576 reflectivity measurements and Katherine Brownlie for carcass measurements. Many thanks to
577 Benjamin Robira for his advice. Finally, thanks to Aude Caizergues for the loan of her
578 powerful computers.

579 **Author Contributions**

580 Conceived and designed the project: D.G., J.F, M.C. Software developers and data providers:
581 W.P, P.M, J.F, H.S, B.M, P.F, S.D, H.H, V.S.B, B.M, T.A-N, I.S.B, O.C, S.C-D, J.D, F.D, N.D,
582 K.E.E, A.E, M.G, Y.K, M.L, S.H.L, M.N, B.O, T.K.R, G.S, T.L.T, M.B, T.D, A.L.F, M.G.F, M.F,
583 H.G.G, T.G, N.P.H, M.J, K.L.J, A.L.K, J.F.L, L.MT, M.M, F.R.M, W.M, A.M, A.P, D.G. Analyzed
584 the data: M.C, W.C, P.M, N.C. Wrote the paper: M.C., D.G. All authors reviewed the
585 manuscript.

586 **Competing Interests statement**

587 No competing interests.

588 **Data availability statement**

589 The datasets analyzed during this study are available from the corresponding author on
590 reasonable request.

591

592

593 **Code availability statement**

594 The different R codes used during this study are available from the corresponding author on
595 reasonable request.

596 **References**

- 597 Afonso, P., Fontes, J., Giacomello, E., Magalhães, M. C., Martins, H. R., Morato, T., ...
598 Vandepierre, F. (2020). The Azores : A Mid-Atlantic Hotspot for Marine Megafauna
599 Research and Conservation. *Frontiers in Marine Science*, 6(826), 1–9.
600 <https://doi.org/10.3389/fmars.2019.00826>
- 601 Alves, J. A., Gunnarsson, T. G., Hayhow, D. B., Appleton, G. F., Potts, P. M., Sutherland, W.
602 J., & Gill, J. A. (2013). Costs, benefits, and fitness consequences of different migratory
603 strategies. *Ecology*, 94(1), 11–17. <https://doi.org/10.1890/12-0737.1>
- 604 Amélineau, F., Fort, J., Mathewson, P. D., Speirs, D. C., Courbin, N., Perret, S., ... Grémillet,
605 D. (2018). Energyscapes and prey fields shape a North Atlantic seabird wintering hotspot
606 under climate change. *Royal Society Open Science*, 5:171883.
607 <https://doi.org/10.1098/rsos.171883>
- 608 Amélineau, Françoise, Grémillet, D., Harding, A. M. A., Walkusz, W., Choquet, R., & Fort, J.
609 (2019). Arctic climate change and pollution impact little auk foraging and fitness across
610 a decade. *Scientific Reports*, 9(1), 1014. <https://doi.org/10.1038/s41598-018-38042-z>
- 611 Arhonditsis, G. B., & Brett, M. T. (2004). Evaluation of the current state of mechanistic
612 aquatic biogeochemical modeling. *Marine Ecology Progress Series*, 271, 13–26.
613 <https://doi.org/10.1021/es061030q>
- 614 Barrett, R. T., Chapdelaine, G., Anker-Nilssen, T., Mosbech, A., Montevecchi, W. A., Reid, J.
615 B., & Veit, R. R. (2006). Seabird numbers and prey consumption in the North Atlantic.
616 *ICES Journal of Marine Science*, 63(6), 1145–1158.
617 <https://doi.org/10.1016/j.icesjms.2006.04.004>
- 618 Beaugrand, G., Conversi, A., Atkinson, A., Cloern, J., Chiba, S., Fonda-Umani, S., ...
619 Edwards, M. (2019). Prediction of unprecedented biological shifts in the global ocean.
620 *Nature Climate Change*, 9(3), 237–243. <https://doi.org/10.1038/s41558-019-0420-1>
- 621 Benhamou, S. (2011). Dynamic approach to space and habitat use based on biased random
622 bridges. *PLoS ONE*, 6(1). <https://doi.org/10.1371/journal.pone.0014592>
- 623 Bennison, A., Jessopp, M., Bennison, A., & Jessopp, M. (2015). At-sea surveys confirm a
624 North Atlantic biodiversity hotspot At-sea surveys confirm a North Atlantic biodiversity
625 hotspot. *Bird Study*, 0(0), 1–5. <https://doi.org/10.1080/00063657.2015.1011601>
- 626 Boertmann, D., Lyngs, P., Merkel, F. R., & Mosbech, A. (2004). The significance of
627 Southwest Greenland as winter quarters for seabirds. *Bird Conservation International*,
628 14(2004), 87–112. <https://doi.org/10.1017/S0959270904000127>
- 629 Boyce, M. ., & McDonald, L. L. (1999). Resource selection functions. *TREE*, 14(7), 268–272.
630 [https://doi.org/10.1016/S0169-5347\(99\)01593-1](https://doi.org/10.1016/S0169-5347(99)01593-1)

631

632 Boyce, M. S., Vernier, P. R., Nielsen, S. E., & Schmiegelow, F. K. A. (2002). Evaluating
633 resource selection functions. *Ecological Modelling*, *157*, 281–300.
634 [https://doi.org/10.1016/S0304-3800\(02\)00200-4](https://doi.org/10.1016/S0304-3800(02)00200-4)

635 Brooks, M. E., Kristensen, K., van Benthem, K. J., Magnusson, A., Berg, C. W., Nielsen, A.,
636 ... Bolker, B. M. (2017). glmmTMB balances speed and flexibility among packages for
637 zero-inflated generalized linear mixed modeling. *R Journal*, *9*(2), 378–400.
638 <https://doi.org/10.32614/rj-2017-066>

639 Bruno, J. F., Bates, A. E., Cacciapaglia, C., Pike, E. P., Amstrup, S. C., Van Hooedonk, R., ...
640 Aronson, R. B. (2018). Climate change threatens the world's marine protected areas.
641 *Nature Climate Change*, *8*(6), 499–503. <https://doi.org/10.1038/s41558-018-0149-2>

642 Caesar, J., Palin, E., Liddicoat, S., Lowe, J., Burke, E., Pardaens, A., ... Kahana, R. (2013).
643 Response of the HadGEM2 Earth System Model to future greenhouse gas emissions
644 Pathways to the year 2300. *J.Climate*, *26*(10), 3275–3284.
645 <https://doi.org/https://doi.org/10.1175/JCLI-D-12-00577.1>

646 Carneiro, A. P. B., Pearmain, E. J., Opper, S., Clay, T. A., Phillips, R. A., Bonnet-Lebrun, A.-
647 S., ... Dias, M. P. (2020). A framework for mapping the distribution of seabirds by
648 integrating tracking, demography and phenology. *Journal of Applied Ecology*, *57*(3),
649 514–525. <https://doi.org/10.1111/1365-2664.13568>

650 Cheung, W. W. L., Jones, M. C., Reygondeau, G., Stock, C. A., Lam, V. W. Y., & Frölicher,
651 T. L. (2016). Structural uncertainty in projecting global fisheries catches under climate
652 change. *Ecological Modelling*, *325*, 57–66.
653 <https://doi.org/10.1016/j.ecolmodel.2015.12.018>

654 Cheung, W. W. L., Lam, V. W. Y., Sarmiento, J. L., Kearney, K., Watson, R., & Pauly, D.
655 (2009). Projecting global marine biodiversity impacts under climate change scenarios.
656 *Fish and Fisheries*, *10*(3), 235–251. <https://doi.org/10.1111/j.1467-2979.2008.00315.x>

657 Clairbaux, M., Fort, J., Mathewson, P., Porter, W., Strøm, H., & Grémillet, D. (2019).
658 Climate change could overturn bird migration : Transarctic flights and high-latitude
659 residency in a sea ice free Arctic. *Scientific Reports*, *9*(17767), 1–13.
660 <https://doi.org/10.1038/s41598-019-54228-5>

661 Collins, W. J., Bellouin, N., Doutriaux-Boucher, M., Gedney, N., Halloran, P., Hinton, T., ...
662 Woodward, S. (2011). Development and evaluation of an Earth-System model –
663 HadGEM2. *Geosci. Model Dev.*, *4*(4), 1051–1075. [https://doi.org/10.5194/gmd-4-1051-](https://doi.org/10.5194/gmd-4-1051-2011)
664 2011

665 Collins, W. J., Bellouin, N., Doutriaux-Boucher, M., Gedney, N., Hinton, T., Jones, C. D., ...
666 Woodward, S. (2008). *Evaluation of HadGEM2 model*.

667 Cristofari, R., Liu, X., Bonadonna, F., Cherel, Y., Pistorius, P., Le Maho, Y., ... Trucchi, E.
668 (2018). Climate-driven range shifts of the king penguin in a fragmented ecosystem.
669 *Nature Climate Change*, *8*(3), 245–251. <https://doi.org/10.1038/s41558-018-0084-2>

670 del Hoyo, J., Elliott, H., & Sargatal, J. (1996). *Handbook of the Birds of the World- Volume 3-*
671 *Hoatzin to Auks*. Lynx Edicions.

672

673

- 674 Descamps, S., Anker-Nilssen, T., Barrett, R. T., Irons, D. B., Merkel, F., Robertson, G. J., ...
675 Zelenskaya, L. (2017). Circumpolar dynamics of a marine top-predator track ocean
676 warming rates. *Global Change Biology*, 23(9), 3770–3780.
677 <https://doi.org/10.1111/gcb.13715>
- 678 Dias, M. P., Martin, R., Pearmain, E. J., Burfield, I. J., Small, C., Phillips, R. A., ... Croxall,
679 J. P. (2019). Threats to seabirds: A global assessment. *Biological Conservation*, 237,
680 525–537. <https://doi.org/https://doi.org/10.1016/j.biocon.2019.06.033>
- 681 Donald, P. F., Fishpool, L. D. C., Ajagbe, A., Bennun, L. A., Bunting, G., Burfield, I. J., ...
682 Wege, D. C. (2019). Important Bird and Biodiversity Areas (IBAs): the development and
683 characteristics of a global inventory of key sites for biodiversity. *Bird Conservation*
684 *International*, 29(2), 177–198. <https://doi.org/10.1017/S0959270918000102>
- 685 Duarte, C. M., Agusti, S., Barbier, E., Britten, G. L., Castilla, J. C., Gattuso, J.-P., ... Worm,
686 B. (2020). Rebuilding marine life. *Nature*, 580(7801), 39–51.
687 <https://doi.org/10.1038/s41586-020-2146-7>
- 688 Durant, J. M., Hjernmann, D., Frederiksen, M., Charrassin, J. B., Le Maho, Y., Sabarros, P. S.,
689 ... Stenseth, N. C. (2009). Pros and cons of using seabirds as ecological indicators.
690 *Climate Research*, 39(2), 115–129. <https://doi.org/10.3354/cr00798>
- 691 Durant, Joël M., Hjernmann, D., Ottersen, G., & Stenseth, N. C. (2007). Climate and the match
692 or mismatch between predator requirements and resource availability. *Climate Research*,
693 33(3), 271–283. <https://doi.org/10.3354/cr033271>
- 694 Fayet, A. L., Freeman, R., Anker-Nilssen, T., Diamond, A., Erikstad, K. E., Fifield, D., ...
695 Guilford, T. (2017). Ocean-wide drivers of migration strategies and their influence on
696 population breeding performance in a declining seabird. *Current Biology*, 27, 3871–
697 3878. <https://doi.org/10.1016/j.cub.2017.11.009>
- 698 Fort, J., Beaugrand, G., Grémillet, D., & Phillips, R. A. (2012). Biologging, remotely-sensed
699 oceanography and the continuous plankton recorder reveal the environmental
700 determinants of a seabird wintering hotspot. *PLoS ONE*, 7(7): e41194.).
701 <https://doi.org/10.1371/journal.pone.0041194>
- 702 Fort, J., Moe, B., Strøm, H., Grémillet, D., Welcker, J., Schultner, J., ... Mosbech, A. (2013).
703 Multicolony tracking reveals potential threats to little auks wintering in the North
704 Atlantic from marine pollution and shrinking sea ice cover. *Diversity and Distributions*,
705 *Wiley*, 19(10), 1322–1332. <https://doi.org/10.1111/ddi.12105>
- 706 Fort, J., Porter, W. P., & Grémillet, D. (2009). Thermodynamic modelling predicts energetic
707 bottleneck for seabirds wintering in the northwest Atlantic. *Journal of Experimental*
708 *Biology*, 212, 2483–2490. <https://doi.org/10.1242/jeb.032300>
- 709 Fort, J., Porter, W. P., & Grémillet, D. (2011). Energetic modelling: A comparison of the
710 different approaches used in seabirds. *Comparative Biochemistry and Physiology - A*
711 *Molecular and Integrative Physiology*, 158(3), 358–365.
712 <https://doi.org/10.1016/j.cbpa.2010.05.004>
- 713 Fort, J., Steen, H., Strøm, H., Tremblay, Y., Grønningsæter, E., Pettex, E., ... Grémillet, D.
714 (2013). Energetic consequences of contrasting winter migratory strategies in a sympatric
715 Arctic seabird duet. *Journal of Avian Biology*, 44(3), 255–262.
716 <https://doi.org/10.1111/j.1600-048X.2012.00128.x>

- 717 Fossheim, M., Primicerio, R., Johannesen, E., Ingvaldsen, R. B., Aschan, M. M., & Dolgov,
718 A. V. (2015). Recent warming leads to a rapid borealization of fish communities in the
719 Arctic. *Nature Climate Change*, 5(7), 673–677. <https://doi.org/10.1038/nclimate2647>
- 720 Frederiksen, M., Anker-Nilssen, T., Beaugrand, G., & Wanless, S. (2013). Climate, copepods
721 and seabirds in the boreal Northeast Atlantic - current state and future outlook. *Global*
722 *Change Biology*, 19(2), 364–372. <https://doi.org/10.1111/gcb.12072>
- 723 Frederiksen, M., Descamps, S., Erikstad, K. E., Gaston, A. J., Gilchrist, H. G., Grémillet, D.,
724 ... Thórarinnsson, T. L. (2016). Migration and wintering of a declining seabird, the thick-
725 billed murre *Uria lomvia*, on an ocean basin scale: Conservation implications. *Biological*
726 *Conservation*, 200, 26–35. <https://doi.org/10.1016/j.biocon.2016.05.011>
- 727 Frederiksen, M., Moe, B., Daunt, F., Phillips, R. A., Barrett, R. T., Bogdanova, M. I., ...
728 Anker-Nilssen, T. (2011). Multicolony tracking reveals the winter distribution of a
729 pelagic seabird on an ocean basin scale. *Diversity and Distributions*, 1–13.
730 <https://doi.org/10.1111/j.1472-4642.2011.00864.x>
- 731 Gabrielsen, G. W. (2009). Seabirds in the Barents Sea. In *Ecosystem Barents Sea* (Tapir
732 Acad, p. 587).
- 733 Garðarsson, A. (1999). The density of seabirds west of Iceland. *Rit Fiskideildar*, 16, 155–169.
- 734 Gasser, T., Guivarch, C., Tachiiri, K., Jones, C. D., & Ciais, P. (2015). Negative emissions
735 physically needed to keep global warming below 2 ° C. *Nature Communications*,
736 6(7958). <https://doi.org/10.1038/ncomms8958>
- 737 Grecian, W. J., Witt, M. J., Attrill, M. J., Bearhop, S., Becker, P. H., Egevang, C., ... Votier,
738 S. C. (2016). Seabird diversity hotspot linked to ocean productivity in the Canary
739 Current Large Marine Ecosystem. *Biology Letter*, 12.
- 740 Grémillet, D., & Charmantier, A. (2010). Shifts in phenotypic plasticity constrain the value of
741 seabirds as ecological indicators of marine ecosystems. *Ecological Applications*, 20(6),
742 1498–1503. <https://doi.org/10.1890/09-1586.1>
- 743 Grémillet, D., Lewis, S., Drapeau, L., Van Der Lingen, C. D., Huggett, J. A., Coetzee, J. C.,
744 ... Ryan, P. G. (2008). Spatial match-mismatch in the Benguela upwelling zone: Should
745 we expect chlorophyll and sea-surface temperature to predict marine predator
746 distributions? *Journal of Applied Ecology*, 45(2), 610–621.
747 <https://doi.org/10.1111/j.1365-2664.2007.01447.x>
- 748 Grémillet, D., Ponchon, A., Paleczny, M., Palomares, M.-L. D., Karpouzi, V., & Pauly, D.
749 (2018). Persisting Worldwide Seabird-Fishery Competition Despite Seabird Community
750 Decline. *Current Biology*, 28, 1–5. <https://doi.org/10.1016/j.cub.2018.10.051>
- 751 Hammerschlag, N., Schmitz, O. J., Flecker, A. S., Lafferty, K. D., Sih, A., Atwood, T. B., ...
752 Cooke, S. J. (2019). Ecosystem Function and Services of Aquatic Predators in the
753 Anthropocene. *Trends in Ecology and Evolution*, 34(4), 369–383.
754 <https://doi.org/10.1016/j.tree.2019.01.005>
- 755 Harris, M. P., Daunt, F., Newell, M., Phillips, R. A., & Wanless, S. (2010). Wintering areas of
756 adult Atlantic puffins *Fratercula arctica* from a North Sea colony as revealed by
757 geolocation technology. *Marine Biology*, 157(4), 827–836.
758 <https://doi.org/10.1007/s00227-009-1365-0>
- 759

- 760 Heuzé, C. (2017). North Atlantic deep water formation and AMOC in CMIP5 models. *Ocean*
761 *Science*, 13, 609–622.
- 762 Jenouvrier, S., Holland, M., Iles, D., Labrousse, S., Landrum, L., Garnier, J., ... Barbraud, C.
763 (2020). The Paris Agreement objectives will likely halt future declines of emperor
764 penguins. *Global Change Biology*, 26(3), 1170–1184. <https://doi.org/10.1111/gcb.14864>
- 765 Keogan, K., Daunt, F., Wanless, S., Phillips, R. A., Walling, C. A., Agnew, P., ... Lewis, S.
766 (2018). Global phenological insensitivity to shifting ocean temperatures among seabirds.
767 *Nature Climate Change*, 8(4), 313–318. <https://doi.org/10.1038/s41558-018-0115-z>
- 768 Lavoie, D., Lambert, N., ben Mustapha, S., & Baaren, A. Van Der. (2013). *Projections of*
769 *Future Physical and Biogeochemical Conditions in the Northwest Atlantic from CMIP5*
770 *Global Climate Models*.
- 771 Lescroël, A., Mathevet, R., Péron, C., Authier, M., Provost, P., Takahashi, A., & Grémillet,
772 D. (2016). Seeing the ocean through the eyes of seabirds: A new path for marine
773 conservation? *Marine Policy*, 68, 212–220. <https://doi.org/10.1016/j.marpol.2016.02.015>
- 774 Linnebjerg, J. F., Fort, J., Guilford, T., Reuleaux, A., Mosbech, A., & Frederiksen, M. (2013).
775 Sympatric Breeding Auks Shift between Dietary and Spatial Resource Partitioning
776 across the Annual Cycle. *PLoS ONE*, 8(8), 1–10.
777 <https://doi.org/10.1371/journal.pone.0072987>
- 778 Lisovski, S., Hewson, C. M., Klaassen, R. H. G., Korner-Nievergelt, F., Kristensen, M. W., &
779 Hahn, S. (2012). Geolocation by light: Accuracy and precision affected by
780 environmental factors. *Methods in Ecology and Evolution*, 3(3), 603–612.
781 <https://doi.org/10.1111/j.2041-210X.2012.00185.x>
- 782 Manly, B.F.J, McDonald, L. ., Thomas, D. ., McDonald, T. ., & Erickson, W. . (2002).
783 *Resource Selection by Animals: Statistical Design and Analysis for Field Studies*.
784 London: Kluwer Academic Publisher.
- 785 Manly, Bryan F. J., McDonald, L. L., Thomas, D. L., McDonald, T. L., & Erickson, W. P.
786 (2007). *Resource Selection by Animals Statistical Design and Analysis for Field Studies*
787 *Second Edition*. Kluwer Academic Publisher. <https://doi.org/10.1007/0-306-48151-0>
- 788 Martin, T. H. D. T. G. M., Bellouin, N., Collins, W. J., Culverwell, I. D., Halloran, P. R.,
789 Hardiman, S. C., ... Wiltshire, A. (2011). The HadGEM2 family of Met Office Unified
790 Model climate configurations. *Geosci. Model Dev.*, 4(3), 723–757.
791 <https://doi.org/10.5194/gmd-4-723-2011>
- 792 McFarlane Tranquilla, L., Montevecchi, W. A., Hedd, A., Regular, P. M., Robertson, G. J.,
793 Fifield, D. A., & Devillers, R. (2015). Ecological segregation among Thick-billed
794 Murres (*Uria lomvia*) and Common Murres (*Uria aalge*) in the Northwest Atlantic
795 persists through the nonbreeding season. *Canadian Journal of Zoology*, 93, 447–460.
796 <https://doi.org/10.1139/cjz-2014-0315>
- 797 Medina, I., Newton, E., Kearney, M. R., Mulder, R. A., Porter, W. P., & Stuart-Fox, D.
798 (2018). Reflection of near-infrared light confers thermal protection in birds. *Nature*
799 *Communications*, 9:3610, 1–7. <https://doi.org/10.1038/s41467-018-05898-8>
- 800 Meinshausen, M., Hare, B., Wigley, T. M. L., Van Vuuren, D., Den Elzen, M. G. J., & Swart,
801 R. (2006). Multi-gas emissions pathways to meet climate targets. *Climatic Change*,
802 75(1–2), 151–194. <https://doi.org/10.1007/s10584-005-9013-2>

- 803 Montevecchi, W. A., Hedd, A., McFarlane Tranquilla, L., Fifield, D. A., Burke, C. M.,
804 Regular, P. M., ... Phillips, R. A. (2012). Tracking seabirds to identify ecologically
805 important and high risk marine areas in the western North Atlantic. *Biological*
806 *Conservation*, 156, 62–71. <https://doi.org/10.1016/j.biocon.2011.12.001>
- 807 Morris, L. R., Proffitt, K. M., & Blackburn, J. K. (2016). Mapping Resource Selection
808 Functions in Wildlife Studies: Concerns and Recommendations. *Appl Geogr*, 76, 173–
809 183. <https://doi.org/10.1016/j.apgeog.2016.09.025>. Mapping
- 810 Muff, S., Signer, J., & Fieberg, J. (2019). Accounting for individual - specific variation in
811 habitat - selection studies : Efficient estimation of mixed - effects models using Bayesian
812 or frequentist computation. *Journal of Animal Ecology*, 1–13.
813 <https://doi.org/10.1111/1365-2656.13087>
- 814 Nadeau, J.-P., & Puiggali, J.-R. (1995). Sechage: des processus physiques aux procedes
815 industriels. Paris (France) Technique et Documentation - Lavoisier.
- 816 Nayar, K. G., Sharqawy, M. H., Banchik, L. D., & V, J. H. L. (2016). Thermophysical
817 properties of seawater : A review and new correlations that include pressure dependence.
818 *Desalination*, 390, 1–24. <https://doi.org/10.1016/j.desal.2016.02.024>
- 819 Newton, I. (2010). *Bird Migration* (Collins Ne).
- 820 Ostaszewska, K., Balazy, P., Berge, J., & Johnsen, G. (2017). Seabirds During Arctic Polar
821 Night : Underwater Observations from Svalbard Archipelago , Norway Seabirds During
822 Arctic Polar Night : Underwater Observations from. *Waterbirds*, 40(3), 302–308.
823 <https://doi.org/10.1675/063.040.0301>
- 824 Paleczny, M., Hammill, E., Karpouzi, V., & Pauly, D. (2015). Population trend of the world’s
825 monitored seabirds, 1950-2010. *PLoS ONE*, 10(6), 1–11.
826 <https://doi.org/10.1371/journal.pone.0129342>
- 827 Perez, J., Menendez, M., Mendez, F. J., & Losada, I. J. (2014). Evaluating the performance of
828 CMIP3 and CMIP5 global climate models over the north-east Atlantic region. *Clim.*
829 *Dyn.*, 43, 2663–2680. <https://doi.org/10.1007/s00382-014-2078-8>
- 830 Péron, C., Authier, M., & Grémillet, D. (2018). Testing the transferability of track-based
831 habitat models for sound marine spatial planning. *Diversity and Distributions*, 1–16.
832 <https://doi.org/10.1111/ddi.12832>
- 833 Perry, A. L., Low, P. J., Ellis, J. R., & Reynolds, J. D. (2005). Climate Change and
834 Distribution Shifts in Marine Fishes. *Science*, 308(5730), 1912 LP – 1915.
835 <https://doi.org/10.1126/science.1111322>
- 836 Phillips, R. A., Silk, J. R. D., Croxall, J. P., Afanasyev, V., & Briggs, D. R. (2004). Accuracy
837 of geolocation estimates for flying seabirds. *Marine Ecology Progress Series*, 266, 265–
838 272. <https://doi.org/10.3354/meps266265>
- 839 Pichegru, L., Grémillet, D., Crawford, R. J. M., & Ryan, P. G. (2010). Marine no-take zone
840 rapidly benefits endangered penguin. *Biology Letters*, 6(4), 498–501.
841 <https://doi.org/10.1098/rsbl.2009.0913>

842

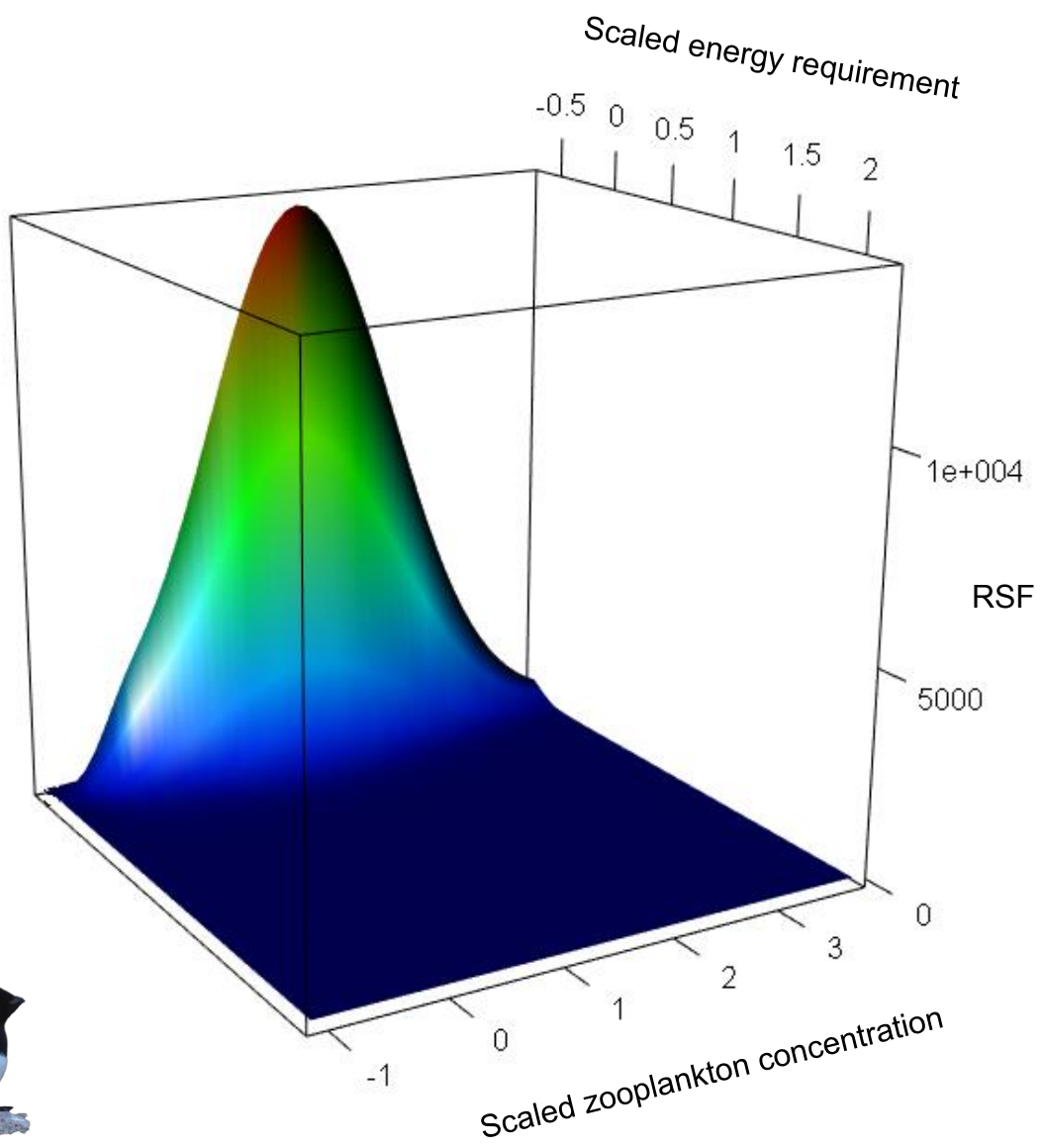
843

- 844 Porter, W. P., & Mitchell, J. W. (2006). *Method and system for calculating the spatial-*
845 *temporal effects of climate and other environmental conditions on animals.* In:
846 *http://www.patentstorm.us/patents/7155377-fulltext.html (ed. U.P Office).Wisconsin*
847 *Alumni Research Foundation, USA.*
- 848 Roberts, D. R., Bahn, V., Ciuti, S., Boyce, M. S., Elith, J., Guillera-arroita, G., ... Dormann,
849 C. F. (2017). Cross-validation strategies for data with temporal , spatial , hierarchical , or
850 phylogenetic structure. *Ecography*, *40*, 913–929. <https://doi.org/10.1111/ecog.02881>
- 851 Sauve, D., Divoky, G., & Friesen, V. L. (2019). Phenotypic plasticity or evolutionary change?
852 An examination of the phenological response of an arctic seabird to climate change.
853 *Functional Ecology*, *33*(11), 2180–2190. <https://doi.org/10.1111/1365-2435.13406>
- 854 Somveille, M., Rodrigues, A. S. L., & Manica, A. (2018). Energy efficiency drives the global
855 seasonal distribution of birds. *Nature Ecology & Evolution*, *2*(6), 962–969.
856 <https://doi.org/10.1038/s41559-018-0556-9>
- 857 Sumaila, U. R., Tai, T. C., Lam, V. W. Y., Cheung, W. W. L., Bailey, M., Cisneros-
858 montemayor, A. M., ... Gulati, S. S. (2019). Benefits of the Paris Agreement to ocean
859 life , economies , and people. *Science Advances*, *5*(:eaau3855), 1–9.
- 860 Sydeman, W. ., Poloczanska, E., Reed, T. ., & Thompson, S. A. (2015). Climate change and
861 marine vertebrates. *Science*, *350*(6262), 171–193. Retrieved from
862 <http://dx.doi.org/10.1126/science.aac9874>
- 863 Teplitsky, C., & Charmantier, A. (2019). Evolutionary consequences of climate change in
864 birds. In P. O. Dunn & A. P. Moller (Eds.), *Effects of Climate Change on Birds-Second*
865 *Edition* (Oxford Uni, pp. 134–146).
866 <https://doi.org/10.1093/oso/9780198824268.003.0011>
- 867 Totterdell, I. J. (2019). Description and evaluation of the Diat-HadOCC model v1 . 0 : the
868 ocean biogeochemical component of HadGEM2-ES. *Geoscientific Model Development*,
869 4497–4549.
- 870 Tranquilla, L. A. M., Montevecchi, W. A., Hedd, A., Fifield, D. A., Burke, C. M., Smith, P.
871 A., ... Phillips, R. A. (2013). Multiple-colony winter habitat use by murrens *Uria* spp . in
872 the Northwest Atlantic Ocean : implications for marine risk assessment. *Marine Ecology*
873 *Progress Series*, *472*, 287–303. <https://doi.org/10.3354/meps10053>
- 874 Trisos, C. H., Merow, C., & Pigot, A. L. (2020). The projected timing of abrupt ecological
875 disruption from climate change. *Nature*, (January 2019), 1–6.
876 <https://doi.org/10.1038/s41586-020-2189-9>
- 877 United Nations Framework Convention on Climate Change (UNFCCC). (2015). *The Paris*
878 *Agreement. Technical Report United Nations Framework Convention on Climate*
879 *Change.*
- 880 Visser, M. E., Perdeck, A. C., van Balen, J. H., & Both, C. (2009). Climate change leads to
881 decreasing bird migration distances. *Global Change Biology*, *15*(8), 1859–1865.
882 <https://doi.org/10.1111/j.1365-2486.2009.01865.x>
- 883 Vuuren, D. P. Van, Stehfest, E., den Elzen, M. G. J., Kram, T., Vliet, J. Van, Deetman, S., ...
884 van Ruijven, B. (2011). RCP2 . 6 : exploring the possibility to keep global mean
885 temperature increase below 2 ° C. *Climatic Change*, *109*, 95–116.
886 <https://doi.org/10.1007/s10584-011-0152-3>

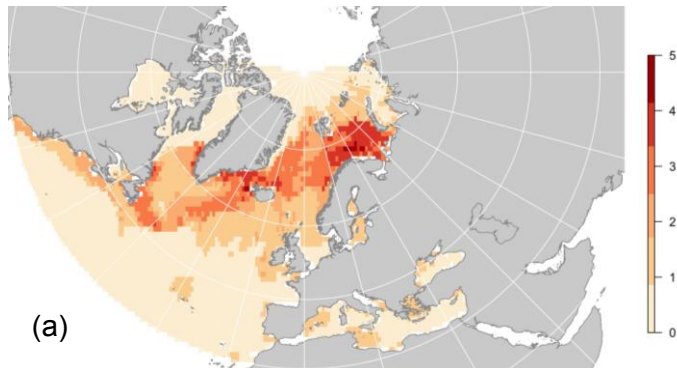
887 Wang, M., & Overland, J. E. (2012). A sea ice free summer Arctic within 30 years: An update
888 from CMIP5 models. *Geophysical Research Letters*, 39, 1–6.
889 <https://doi.org/10.1029/2012GL052868>

890 Zappa, G., Shaffrey, L. C., & Odges, K. I. (2013). The Ability of CMIP5 Models to Simulate
891 North Atlantic Extratropical Cyclones. *American Meteorological Society*, 26.
892 <https://doi.org/10.1175/JCLI-D-12-00501.1>

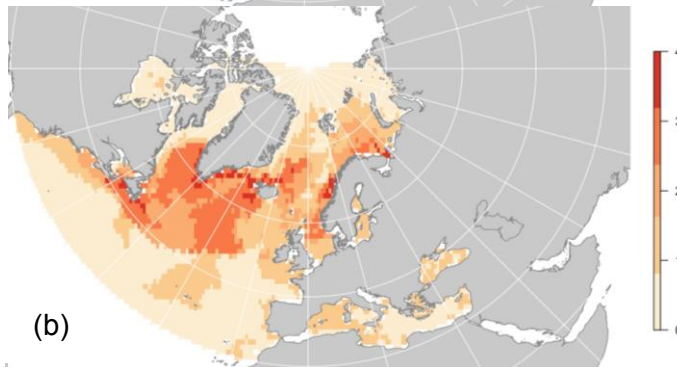
893



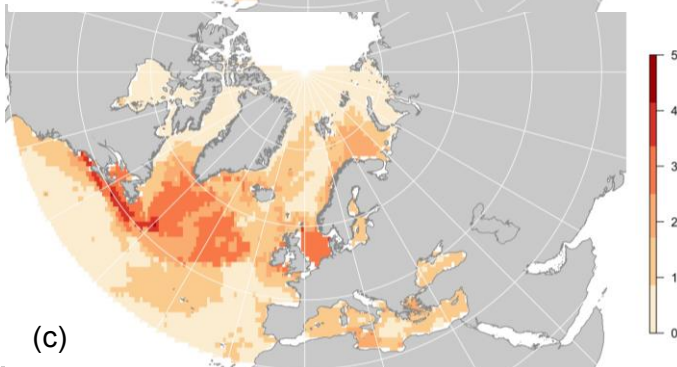
OCTOBER



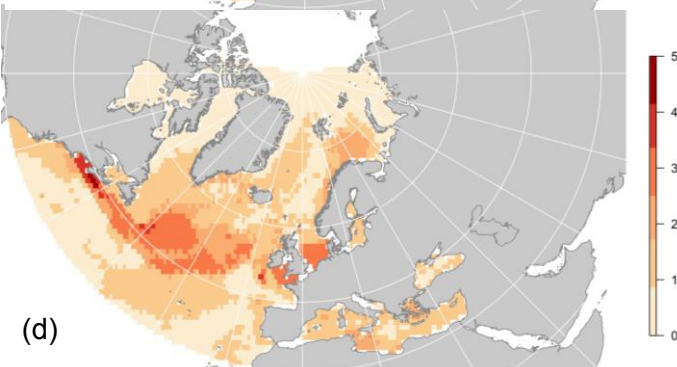
NOVEMBER



DECEMBER



JANUARY



FEBRUARY

