

1 **Biogeography of cephalopods in the Southern Ocean**
2 **using habitat suitability prediction models**

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19

20 Abstract

21 Our understanding of how environmental change in the Southern Ocean will affect marine
22 diversity, habitats, and distribution remain limited. The habitats and distributions of Southern
23 Ocean cephalopods are generally poorly understood, and yet such knowledge is necessary for
24 research and conservation management purposes, as well as for assessing the potential
25 impacts of environmental change. We used net-catch data to develop habitat suitability
26 models for 15 of the most common cephalopods in the Southern Ocean. Using modelled
27 habitat suitability, we assessed favourable areas for each species and examined the
28 relationships between species distribution and environmental parameters. The results
29 compared favourably with the known ecology of these species and with spatial patterns from
30 diet studies of squid predators. The individual habitat suitability models were overlaid to
31 generate a “hotspot” index of species richness, which showed higher numbers of squid
32 species associated with various fronts of the Antarctic circumpolar current. Finally, we
33 reviewed the overall distribution of these species and their importance in the diet of Southern
34 Ocean predators. There is a need for further studies to explore the potential impacts of future
35 climate change on Southern Ocean squid.

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40 Keywords: Biogeography, Southern Ocean, Cephalopods, Habitat suitability models

41

42 **Introduction**

43 Habitat suitability models can contribute significantly to our understanding of species
44 niche requirements and can predict the potential distributions of species (Hirzel et al., 2006).
45 Certain regions of the Antarctic are among the most rapidly warming areas on Earth (Turner
46 et al., 2009). However, our understanding of how these changes affect marine diversity,
47 habitats, and distribution remain limited, particularly regarding pelagic taxa in the Southern
48 Ocean (Xavier et al., 2006; Griffiths, 2010).

49 In the Southern Ocean, defined here as the region south of the Subtropical Front, all
50 known squid are oceanic pelagic species with high levels of endemism (Collins and
51 Rodhouse, 2006). As most pelagic cephalopods have a short life span, rapid and labile
52 growth, and semelparous maturation patterns (Boyle and Rodhouse, 2005), it seems likely
53 that they will respond relatively rapidly to environmental change. Antarctic squid are also a
54 poorly studied group despite considered to be commercially exploitable in the future (Xavier
55 et al., 2007). For the Southern Ocean, it has been suggested that predicted temperature
56 increases, and/or changes in sea ice extent, are unlikely to have major effects on squid other
57 than changes in distribution near the limits of their range (Rodhouse, 2013). However, the
58 likely consequences of ecosystem change on the distribution of squid fauna in the Southern
59 Ocean are not well understood (Constable et al., 2014; Kennicutt II et al., 2014; Xavier et al.,
60 2014).

61 The objective of this study was to estimate the spatial distribution of suitable habitats
62 of a number of common squid species from the Southern Ocean. We review our predicted
63 distributions against previously published distribution estimates (Xavier et al., 1999;
64 Rodhouse et al., 2014), the known distribution of the sampling effort (Griffiths, 2010), and
65 the presence of the studied species in the diet of key top predators in different areas of the

66 Southern Ocean.

67

68 **Materials and Methods**

69 Occurrence data were taken from the SCAR Biogeographic Atlas of the Southern
70 Ocean (De Broyer et al., 2014). This compilation was based upon Xavier et al. (1999), with
71 additional data drawn from the Ocean Biogeographic Information System (OBIS, 2013),
72 biodiversity.aq, the Australian Antarctic Data Centre, and the National Institute of Water and
73 Atmospheric Research (NIWA, 2014). Duplicate records (identified by exact matches in
74 species name and position) were removed. Figure 1 shows the study region and the names of
75 features mentioned in the text. Figure 2 shows the complete set of occurrence records used.

76 The available species occurrence records were in presence-only form, and so the
77 habitat suitability modelling was conducted using the Maxent software package (v3.3.3k)
78 (Phillips et al., 2006). Maxent does not provide a direct estimate of the probability of
79 presence of the species across its range, but rather an index of habitat suitability (effectively,
80 utilized habitat relative to the background environmental conditions). This index is
81 nonlinearly related to the probability of presence (Phillips et al., 2009). Maxent allows for
82 nonlinear model terms by formulating a series of features from the predictor variables. Due to
83 relatively limited sample sizes, we constrained the complexity of most models by considering
84 only linear, quadratic, and product features. A multiplier of 3.0 was used on automatic
85 regularization parameters to discourage overfitting (Radosavljevic and Anderson, 2014);
86 otherwise, default Maxent settings were used. A 10-fold cross-validation procedure was used
87 to assess model performance (using the area under the receiver-operating curve) and variable
88 permutation importance, with values averaged over the 10 fitted models. The final predicted

89 distribution for each species was based on a single model fitted using all data. The squid
90 presence records come from a mixture of sources: some dedicated marine science surveys
91 with a designed sampling strategy, but also other sources such as fishing vessels. The
92 presence records are therefore biased, in that they were not drawn at random from across the
93 range of each species. To reduce the effects of this bias on the fitted models, the background
94 points were sampled from the locations of all squid records, rather than randomly sampled
95 from across the region of interest (Phillips et al., 2009). 1000 background points were used
96 for each model.

97 Species distribution and habitat suitability modelling in the Southern Ocean relies on
98 predictor variables drawn from remote-sensing and model sources in order to obtain synoptic
99 coverage at suitable spatial and temporal resolution. Predictor variables (**Error! Reference**
100 **source not found.**)(Smith and Sandwell, 1997; Spreen et al., 2008; Feldman and McClain,
101 2010; Garcia et al., 2010; Rio et al., 2011; Trübenbach et al., 2013) were chosen from a
102 collection of Southern Ocean layers (Raymond, 2012). These variables were selected as
103 indicators of ecosystem structure and processes including water mass properties, sea ice
104 dynamics, and productivity (see biological relevance, **Error! Reference source not found.**).
105 We used a combination of predictive performance and expert opinion, including
106 interpretation of the fitted responses, to select appropriate variables for each model. The
107 selection process was also used to avoid including multiple, highly-correlated predictor
108 variables within any one model.

109 Records of squid as prey items were extracted from the Southern Ocean dietary
110 database (Raymond et al., 2011). These data were not used as part of the model fitting
111 process, which was based entirely on the net catch data, but rather as informal evaluation of
112 the predicted habitat distributions from the models. This comparison between the spatial

113 distribution of the diet records and the spatial pattern of predicted habitat was not done in a
114 formal manner, because the geographic location of a diet record indicates where that diet
115 sample was obtained (usually a breeding colony); the prey item in question may not be local
116 to that colony. Some predators (e.g. sperm whales, wandering albatrosses) retain squid beaks
117 in their stomachs for long periods of time (Clarke, 1980; Xavier et al., 2005) and/or have long
118 foraging ranges (e.g. albatrosses) (Phillips et al., 2008), so they may have consumed that prey
119 item a considerable distance from the colony. Further, the absence of a prey item from a
120 predator's diet may be due to factors such as availability (e.g. deep prey beyond the diving
121 reach of air-breathing predators) or prey preference, rather than disjunct spatial distributions
122 (Xavier et al., 2013).

123 We identified cephalopod hotspots using an index of species richness derived from the
124 individual species habitat models. We then converted the predicted habitat suitability for each
125 species to a binary presence/absence layer by applying a threshold, such that habitat
126 suitability values above the threshold were converted to presences. The threshold used for
127 each species was the average of the thresholds (for each of the 10 training models) chosen to
128 maximize the test area under the receiver-operating curve (Phillips et al., 2006). The binary
129 layers were summed to give the number of species estimated to be present in each pixel
130 (Ballard et al., 2012). The results of this study are available from the Australian Antarctic
131 Data Centre.

132

133 **Results**

134 The results of the modelling, including the predictor variables used in each
135 model, are summarised in Table 2. For each species we provide two maps: one showing the

136 catch data used to fit the model as well as the diet data available for that squid species as
137 prey, and the second figure for each species showing the predicted habitat suitability map.

138

139 Family Bathyteuthidae

140 *Bathyteuthis abyssicola* (Figure 3a,b). The predicted habitat suitability suggests a circumpolar
141 distribution (i.e. occurring in all three sectors of the Southern Ocean). The most favourable
142 habitat was predicted to lie between the Southern Antarctic Circumpolar Current Front
143 (SACCF) and the Sub-Antarctic Front (SAF), with more moderate values of habitat suitability
144 extending from roughly 45°S up to the Antarctic shelf.

145

146 Family Brachioteuthidae

147 *Slosarczykovia circumantarctica* (Figure 4a,b). The predicted habitat suitability suggests a
148 circumpolar distribution, with meridional limits between approximately the SACCF and the
149 SAF. Zonally, the most favourable habitat was predicted in the Scotia Sea in the Atlantic
150 sector, particularly around the Antarctic Peninsula and South Georgia, in the Indian sector
151 (with higher values at Kerguelen shelf and eastern waters) and south of the Tasman Sea
152 between Tasmania and New Zealand. While net catches of this species were sparse outside of
153 the southwest Atlantic sector, diet records were present in the Indian and Pacific sectors,
154 broadly matching the predicted habitat distribution.

155

156 Family Cranchiidae

157 *Galiteuthis glacialis* (Figure 5a,b). The predicted habitat suitability clearly indicates a
158 circumpolar distribution, bounded to the south by the Antarctic continent and to the north by
159 the SAF.

160

161 *Mesonychoteuthis hamiltoni* (Figure 6a,b). The predicted habitat suitability suggests a
162 circumpolar distribution extending relatively close to the Antarctic continent but not into
163 shallow areas such as the continental shelf or the Kerguelen Plateau. To the north, suitable
164 habitat appears to be delimited by the SAF. The highest values of habitat suitability extended
165 from the Weddell Sea in the Atlantic sector to 60 °E (west of the Kerguelen archipelago), and
166 between 180 °E and 120 °W in the Ross/Amundsen seas region.

167

168 Family Gonatidae

169 *Gonatus antarcticus* (Figure 7a,b). The predicted habitat suitability suggests a circumpolar
170 distribution, with patches of highly suitable habitat over the south part of the Patagonian shelf
171 (around the Falkland Islands and Cape Horn), in the Scotia Sea and to the east in the Atlantic
172 sector, in the Indian sector (northern Kerguelen Plateau and Prydz Bay) and in the Pacific
173 sector (the Ross Sea, and eastwards along the continental shelf to the Antarctic Peninsula).
174 Similarly to *S. circumantarctica*, catch records were almost exclusively restricted to the
175 southwest Atlantic sector, whereas diet records were circumpolar, as was the predicted habitat
176 distribution.

177

178 Family Histioteuthidae

179 *Histioteuthis atlantica* (Figure 8a,b). The predicted habitat suitability indicated a circumpolar
180 distribution north of approximately 60 °S (50 °S in the Atlantic sector), away from the coldest
181 waters of the Southern Ocean. The predicted distribution of *H. atlantica* was restricted to
182 more northerly regions than that of the closely-related *H. eltaninae*.

183

184 *Histioteuthis eltaninae* (Figure 9a,b). The model predictions indicated that suitable habitat is
185 widespread across the Southern Ocean, excluding shallow areas such as continental shelves
186 and undersea banks and ridges.

187

188 Family Loliginidae

189 *Doryteuthis gahi* (Figure 10a,b). Predicted habitat for this species was limited to continental
190 shelves, particularly the Patagonian shelf (agreeing well with all net capture and predator diet
191 locations), and in South Chilean waters (in the Pacific). Areas of suitable habitat, albeit more
192 restricted in extent, were also predicted around South Georgia, the Kerguelen Islands, and
193 New Zealand.

194

195 Family Neoteuthidae

196 *Alluroteuthis antarcticus* (Figure 11a,b). Predicted habitat was circumpolar, bounded
197 approximately by the SACCF to the north, and by the Antarctic continental shelf to the south.

198

199 Family Ommastrephidae

200 *Martialia hyadesi* (Figure 12a,b). The model predicted spatially-patchy areas of suitable
201 habitat, generally downstream of land masses. The principal areas of predicted habitat were
202 around the South American shelf, in the north Scotia Sea close to South Georgia, in the
203 Indian sector (Prince Edward, Crozet and Kerguelen shelf archipelagos and to the east of the
204 latter islands) and south and southwest of New Zealand.

205

206 *Todarodes filippovae* (Figure 13a,b). The predicted habitat suitability was clearly
207 circumpolar, bounded to the south by the SAF, away from the coldest waters of the Southern
208 Ocean. While catches were largely confined to the eastern Indian and Pacific sectors
209 (Tasmania through to South America), a small number of diet and catch records from the
210 western and central Indian sectors (approximately 30–80 °E) provided some corroboration of
211 the circumpolar habitat prediction.

212

213 Family Onychoteuthidae

214 *Kondakovia longimana* (Figure 14a,b). The predicted habitat suitability was circumpolar,
215 consistent with the catch and diet records, but spatially patchy. Areas of most suitable habitat
216 were found in the Scotia Sea, particularly around South Georgia and the South Sandwich
217 islands, in the Indian sector (Kerguelen waters and further south) and south of the Tasman
218 Sea around 60 °S. Patches of suitable habitat were also predicted for parts of the Antarctic
219 continent shelf (e.g. the western Antarctic Peninsula, Prydz Bay, and the Dumont d'Urville
220 Sea).

221

222 *Moroteuthis ingens* (Figure 15a,b). Predicted areas of suitable habitat were patchy, generally
223 restricted to regions around 50 °S or further north. Oceanic waters were generally predicted to
224 be unsuitable habitat, compared to areas above and around continental or island shelves,
225 particularly around the Falkland Islands, Crozet, the Kerguelen Plateau in the Indian sector,
226 south of Tasmania, and on the New Zealand shelf in the Pacific sector. North of about 60 °S,
227 the shelf distribution of *M. ingens* is complementary to that of *H. atlantica*.

228

229 *Moroteuthis robsoni* (Figure 16a,b). The predicted habitat suitability was broadly circumpolar
230 with a southern boundary at approximately the SAF, away from the coldest waters of the
231 Southern Ocean, and with an affinity for mid-depth regions (e.g. shelf slopes).

232

233

234 Family Psychroteuthidae

235 *Psychroteuthis glacialis* (Figure 17a,b). Suitable habitat was predicted to be circumpolar,
236 extending northwards from the Antarctic continent to approximately the APF. Areas of
237 highest habitat suitability were found in patches in the southern Scotia Sea, Weddell, and
238 Ross seas, and in coastal waters around the Antarctic continent.

239

240 **“Hotspot” regions in the distribution of cephalopods from the Southern Ocean**

241 The individual species habitat suitability predictions were combined to
242 produce an index of species richness (Figure 18). The highest predicted values (8 or more

243 species) occurred in a largely-circumpolar band, approximately from the Polar Front south to
244 the northernmost extent of sea ice. Areas of nine or more species were found in the southwest
245 Atlantic sector. The lowest values occurred over the shelf around the Antarctic continent, and
246 to the north of the sub-Antarctic front (but note that these latter waters are home to other
247 species of squid not considered in this study).

248

249 **Discussion**

250 **Biases and uncertainties**

251 To our knowledge, this is the first study to develop habitat suitability predictions for
252 these common cephalopod species of the Southern Ocean. The modelling component of this
253 study presented a number of challenges. Data systems such as OBIS currently provide the
254 most comprehensive occurrence data for biogeography but are aggregated from a variety of
255 diverse sources with differences in aspects such as survey design and sampling techniques.
256 One possible approach is to use the aggregated dataset merely as an index of the available
257 data and follow each component dataset back to its original, detailed source. However, this is
258 rarely practical for large-scale studies. Although we have attempted to account for the spatial
259 distribution of survey efforts in the modelling procedure, these results should still be treated
260 with caution, particularly for species with small sample sizes or where one particular area
261 dominates the occurrence record.

262 The predictor variables used were drawn from satellite and similar sources. The
263 information from such variables rarely provides direct characterization of the primary
264 processes affecting the species distribution. For example, there are no direct estimates of
265 squid prey distributions. Instead, these variables typically provide proxy information such as

266 water mass properties or primary productivity. The spatial and temporal scales of this
267 information often do not match the scales experienced by the animals. Furthermore, predictor
268 variables in the Southern Ocean are typically highly correlated because of the strong
269 latitudinal and seasonal gradient that affects oceanic and atmospheric conditions. Because of
270 these factors, it is rarely obvious which particular predictor variable is the most appropriate
271 proxy to use in a given model. Predictive performance offers some guidance, but should not
272 be relied upon exclusively (Raymond et al., 2014). Squid are also notorious for their net
273 avoidance ability, and scientific nets typically catch only juvenile individuals (Collins and
274 Rodhouse, 2006).

275 In order to help assess the influence of these issues on the results, we used *Doryteuthis*
276 *gahi* as a validation species, because it is well known to be coastally distributed (up to 350 m
277 depth) in areas of the Patagonian shelf and eastern Pacific Ocean from southern Peru to
278 Southern Chile (Arkhipkin et al., 2013). The predicted habitat suitability for this species
279 broadly matched the expected pattern. Small areas of suitable habitat were predicted in a few
280 locations where this species is unlikely to be present (e.g. around New Zealand). This
281 highlights the fact that the outputs from these models are predictions of suitable habitat and
282 do not take into account other processes that govern species distributions such as dispersion,
283 competition, and trophic dependencies. Indeed, combining food web models with species
284 distribution models to predict spatial variation in community composition remains an active
285 area of research in biodiversity modelling (Pellissier et al., 2013; Constable et al., 2014).

286 Generally, each predicted habitat distribution matched the picture provided by the
287 combination of occurrence and predator diet records. For *Kondakovia longimana*, the
288 predicted habitat was much more circumpolar in nature than the observed catch records, but
289 that circumpolar pattern was consistent with predator diet observations. Similarly, suitable

290 habitat for *Gonatus antarcticus* was predicted to include areas close to the Antarctic
291 continent, well away from observed net catches. However, emperor penguins have been
292 recorded to feed on this species there (Cherel and Kooyman, 1998). Some minor
293 discrepancies were also noted. The sea ice zone was not predicted to be suitable habitat for
294 *Moroteuthis ingens*, apparently contradicting an emperor penguin diet record from Auster
295 colony, near Mawson station (Robertson et al., 1994). However, this species has never been
296 recorded in any other emperor diet studies (Xavier and Cherel, 2009) and so this record may
297 have been a misidentified *M. knipovitchi*, an Antarctic species of the same family with
298 broadly similar beak characteristics (Xavier and Cherel, 2009). Finally, *S. circumantarctica*
299 has occasionally been caught by nets in warm waters near New Zealand (around 45 °S) but
300 this was not predicted to be a suitable habitat (Figure 4b).

301 The models showed that ocean areas with generally higher levels of habitat suitability
302 exist around 50 °S in the Atlantic and Indian sectors and 60 °S in the Pacific sector, where the
303 majority of the fronts are distributed (i.e. Polar Front, sub-Antarctic Front and subtropical
304 Fronts; see Xavier et al. 1999), reinforcing that these areas regions are broadly ecologically
305 relevant, including for cephalopods.

306

307 **Physical and biogeochemical ocean mechanisms influencing the distribution of squid in** 308 **the Southern Ocean**

309 The Southern Ocean is characterized by high surface macro-nutrient concentrations
310 and relatively low iron concentrations, and so iron input typically leads to increased
311 productivity. Several low-latitude species (e.g. *Doryteuthis gahi*, *Martialia hyadesi*) were
312 associated with regions where atmospheric iron deposition is strongly correlated with

313 increased productivity (as measured by satellite-based chlorophyll, e.g. east of Patagonia, on
314 the Falkland plateau) (Erickson et al., 2003). Away from land in the open Southern Ocean,
315 areas of elevated productivity tend to be driven by upwelling of nutrients, often caused by the
316 interaction of the Antarctic circumpolar current flow with large-scale bathymetric features,
317 such as mid-ocean ridges (Sokolov and Rintoul, 2007). Some such areas are known to be
318 foraging grounds for predators such as seabirds (Raymond et al., 2010). Thus water depth can
319 potentially influence cephalopod distribution, even in deep, mid-ocean areas well away from
320 shelves.

321 Broadly speaking, cold, nutrient-rich waters upwell south of the Polar Front and
322 subduction (i.e. downwelling) occurs north of the Polar Front (Sarmiento et al., 2004).
323 Several species in this study (e.g. *Todarodes filippovae*, *Galiteuthis glacialis*) featured a
324 strong contrast across the Subantarctic Front. Upwelling in the Weddell and Ross gyres may
325 also play a role in forming suitable habitat for some species, such as *Psychroteuthis glacialis*
326 (Figure 17a,b). Three species (*Mesonychoteuthis hamiltoni*, *Bathyteuthis abyssicola* and
327 *Slosarczykovia circumantarctica*) were found to have a potential affinity for areas with low
328 oxygen minima, suggesting that these species may use this ecological niche close to or within
329 the oxygen minimum zone. Although low oxygen levels are known to greatly limit the
330 abundance, vertical distribution, and ecology (e.g. predation, food competition) of numerous
331 marine animals, some species of squid (e.g. jumbo squid *Dosidicus gigas*) are known to thrive
332 in such harsh environments (Trübenbach et al., 2013).

333 This study indicates that large-scale physical and biogeochemical properties can
334 influence the suitability of a given region, often in remarkably different ways for different
335 cephalopod species. The 15 species modelled here can be discussed in terms of three broad
336 spatial groupings: those distributed in cold waters close to the Antarctic continent, those in

337 relatively warm waters to the north, and those with less constrained distribution (i.e.
338 extending into both warm and cold waters).

339

340 **Habitat suitability of “cold” water cephalopod species**

341 The species that clearly have suitable habitat close to the Antarctic continent were
342 *Alluroteuthis antarcticus*, *Galiteuthis glacialis*, *Mesonychoteuthis hamiltoni*, *Psychroteuthis*
343 *glacialis*. These species have been recognized as typical Antarctic water species with a
344 suggested circumpolar distribution (Xavier et al., 1999; Rodhouse et al., 2014), consistent
345 with our results. *A. antarcticus* is occasionally caught in nets in the Atlantic and Indian
346 sectors of the Southern Ocean (Rodhouse, 1989; Lu and Williams, 1994) but also in the diet
347 of albatrosses in the Pacific sector (Xavier et al., 2014). *G. glacialis* is one of the most
348 abundant (i.e. most commonly caught in midwater research nets) and widely distributed squid
349 species in the colder waters of the Southern Ocean. *M. hamiltoni* is arguably the largest squid
350 species in the world, growing to ten metres or more in length (Collins and Rodhouse, 2006).
351 Its habitat is typically in circumpolar colder waters (see results; Xavier et al. 1999). It is
352 occasionally caught by longline fisheries (as a by-catch), and is found in top predator diets
353 (Xavier and Cherel, 2009). Finally, *P. glacialis* is considered to be abundant with a
354 circumpolar distribution in high Antarctic areas (Filippova and Pakhomov, 1994; Xavier et
355 al., 1999). This is supported by our habitat suitability predictions, suggesting that this species
356 may be abundant close to the continent as previously thought, but also in oceanic waters (see
357 results). Evidence of *P. glacialis* living near the bottom at the shelf break area (300–1000 m)
358 (Lu and Williams, 1994; Collins et al., 2004), applies particularly for the Scotia Sea region in
359 our habitat suitability predictions. *P. glacialis*, like *A. antarcticus*, *G. glacialis* and *M.*

360 *hamiltoni*, is also found in the diets of a wide range of top predators, including albatrosses,
361 penguins, seals, whales and toothfish species (Xavier and Cherel, 2009).

362

363 **Habitat suitability of “mixed” (i.e. cold and warm) water cephalopod species**

364 A group of species were predicted to be broadly distributed, from close to the
365 Antarctic continent to warmer waters north of 60 °S: *Bathyteuthis abyssicola*, *Slosarczykovia*
366 *circumantarctica*, *Histioteuthis eltaninae*, *Kondakovia longimana* and *Gonatus antarcticus*.

367 *B. abyssicola* occurs in all three sectors of the Southern Ocean (Roper, 1969), but
368 occurs very rarely in the diet of top predators (Xavier and Cherel, 2009), probably because it
369 lives at great depths (Roper, 1969). *S. circumantarctica* is considered to be the most abundant
370 squid in the upper layers of pelagic waters in the Southern Ocean, generally deeper than 400
371 m by day and migrating towards the surface by night (Collins and Rodhouse, 2006). The
372 suitable habitat of *S. circumantarctica* in this study predicted to be circumpolar but not close
373 to the Antarctic continent (only at the Antarctic Peninsula islands; see results) occurring
374 regularly in research nets in the Scotia Sea (Rodhouse and Piatkowski, 1995; Rodhouse et al.,
375 1996; Collins et al., 2004). *S. circumantactica* is the most important squid species (by
376 frequency of occurrence and by number) in the diet of Antarctic fur seals breeding at South
377 Georgia in most years (British Antarctic Survey, unpubl. data). *H. eltaninae* is distributed the
378 furthest south of the species of the family Histioteuthidae, with a circumpolar distribution
379 (Rodhouse and Piatkowski, 1995; Rodhouse et al., 1996; Xavier et al., 1999; Collins and
380 Rodhouse, 2006), occurring in small numbers in research nets (Rodhouse and Piatkowski,
381 1995; Xavier and Cherel, 2009), but never close to the Antarctic continent (matching model
382 predictions here; see results). Although it has been suggested that *H. eltaninae* is more

383 abundant in proximity to land and oceanic ridges (Roper et al., 1984), this is not expressed in
384 our predictions (but note that the habitat suitability model for this species used only a single
385 predictor variable (depth), and so should be treated with caution). *H. eltaninae* occurs in a
386 wide range of predator diets (Cherel and Klages, 1998; Xavier and Cherel, 2009). *K.*
387 *longimana* has a circumpolar distribution (Xavier, 1997; Xavier et al., 1999), matched by our
388 study results, ranging from close to the Antarctic continent coasts to north of 60 °S (Cherel
389 and Weimerskirch, 1999). This species also reaches large sizes (Rodhouse et al., 2014), but
390 not as large as *M. hamiltoni*. Although rare in research nets (Collins et al., 2004), *K.*
391 *longimana* is one of the most important species (by number and by mass) in numerous
392 predators in the Southern Ocean, including wandering and grey-headed albatrosses (Clarke,
393 1980; Croxall and Prince, 1996; Xavier et al., 2003b; Cherel et al., 2004; Xavier and Cherel,
394 2009). *G. antarcticus* has a circumpolar distribution reaching as far south as the Antarctic
395 continent (Xavier et al., 1999), a finding mirrored by our results. This species occurs
396 occasionally in nets (Rodhouse et al., 1996; Collins and Rodhouse, 2006) but is more
397 commonly found in the diet of seabirds and seals (Croxall and Prince, 1996; Cherel and
398 Klages, 1998; Xavier et al., 2002; Cherel et al., 2004; Xavier and Cherel, 2009).

399

400 **Habitat suitability of “warm” water cephalopod species**

401 Five species were predicted to be distributed in the warmer waters of the Southern
402 Ocean: *Histioteuthis atlantica*, *Martialia hyadesi*, *Todarodes filippovae*, *Moroteuthis ingens*
403 and *Moroteuthis robsoni*. *H. atlantica* has a circumpolar distribution (Xavier et al., 1999;
404 Rodhouse et al., 2014), and is more northerly distributed than *H. eltaninae*, as reflected by
405 our predicted habitat suitability for both histioteuthid species. *H. atlantica* is known to occur
406 in oceanic waters (Roper et al., 1984), as suggested by our predictions (see results) but it has

407 also been caught in shallow waters (Voss et al., 1998). *H. atlantica* is important in the diet of
408 albatrosses, sharks and many other predators that forage in the warmer waters of the Southern
409 Ocean (Xavier and Cherel, 2009). *M. hyadesi* is found further south than *H. atlantica*, but
410 never close to the Antarctic continent (Rodhouse, 1998a; Xavier et al., 1999). Our results are
411 consistent with this, indicating that the sea ice zone comprises unsuitable habitat for this
412 species. *M. hyadesi* is the squid species that has attracted most attention with regard to future
413 commercial exploitation (Rodhouse, 1997), and is present in the diet of a wide range of top
414 predators (Xavier and Cherel, 2009), being particularly important in the diet of grey-headed
415 albatrosses in some years (Xavier et al., 2003a). *T. filippovae* has a similar circumpolar
416 distribution to *H. atlantica*, and extending further north than the region modelled here (to 35
417 °S) (Pethybridge et al., 2013). It is common around seamounts and slope waters, up to a 1000
418 m depth (Roeleveld, 1998; Xavier et al., 1999). While generally found in relatively warm
419 waters, *T. filippovae* is periodically caught further south (Rodhouse, 1998b). *Todarodes* spp.
420 are present in the diet of toothed whales, wandering albatrosses, seals, sharks and fish (Smale,
421 1996; Xavier and Cherel, 2009). *M. ingens* is mostly associated with shelves (Cherel and
422 Duhamel, 2003) but is also found in bathyal waters (Rodhouse et al., 2014). Given these
423 depth differences, more sampling and genetics work must be carried out to verify if it is truly
424 a single species and not a group of similar species. *M. ingens* is common in the diet of
425 penguins, albatrosses, petrels, whales, seals and the southern opah (Clarke, 1980; Green and
426 Burton, 1993; Cherel et al., 1996; Croxall and Prince, 1996; Cherel and Klages, 1998; Xavier
427 and Cherel, 2009). *M. robsoni* may exhibit a circumpolar distribution (Rodhouse, 1990;
428 Rodhouse et al., 2014), extending as far south as the Scotia Sea but nevertheless still a warm
429 water cephalopod species. Like *M. ingens*, most specimens of *M. robsoni* have been caught in
430 shelf/near shelf waters with a small number specimens being caught in oceanic waters, and so
431 more sampling and genetics work must be carried out to verify the nature of this species. *M.*

432 *robsoni* is an oceanic species (Roper et al., 1984) that occurs occasionally in the diets of
433 Southern Ocean predators that forage sufficiently far north (Imber, 1992; Cherel et al., 2004;
434 Xavier and Cherel, 2009).

435 The “hotspots” in the distribution of cephalopods in the Southern Ocean are related to
436 oceanic waters, across various fronts. This is consistent with the tendency of top predators to
437 target oceanic fronts, potentially to catch squid (Rodhouse et al., 1996; Xavier et al., 2004).
438 Further research should concentrate on these areas to improve our understanding of the
439 abundance and population dynamics of Southern Ocean cephalopods (Xavier et al., 2015).
440 Several species (e.g. *K. longimana*, *G. antarcticus*, *M. hyadesi*, *M. knipovitchi*) have
441 commercial potential in the future (Xavier et al., 2007), although the biology and ecology of
442 some species (particularly *M. knipovitchi*) remain poorly known (Collins et al., 2004; Collins
443 and Rodhouse, 2006). There is also a need for studies to explore the potential impacts of
444 future climate change on Southern Ocean squid.

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456 **References**

- 457 Arkhipkin AI, Hatfield EMC, Rodhouse PGK. 2013. *Doryteuthis gahi*, Patagonian long-
 458 finned squid. Rosa R, Pierce G, O'Dor RK editors. Advances in squid biology, ecology and
 459 fisheries. Part I - Myopsid squids: New York, Nova Science Publishers, p123-157.
- 460 Ballard G, Jongsomjit D, Veloz SD, Ainley DG. 2012. Coexistence of mesopredators in an
 461 intact polar ocean ecosystem: The basis for defining a Ross Sea marine protected area.
 462 Biological Conservation 156: 72–82.
- 463 Boyle P, Rodhouse PG. 2005. Cephalopods ecology and fisheries: Blackwell Science, Oxford.
 464 452p.
- 465 Chérel Y, Duhamel G. 2003. Diet of the squid *Moroteuthis ingens* (Teuthoidea:
 466 Onychoteuthidae) in the upper slope waters of the Kerguelen Islands. Marine Ecology
 467 Progress Series 250: 197-203.
- 468 Chérel Y, Duhamel G, Gasco N. 2004. Cephalopod fauna of subantarctic islands: new
 469 information from predators. Marine Ecology Progress Series 266: 143-156.
- 470 Chérel Y, Klages N. 1998. A review of the food of albatrosses. Robertson G, Gales R editors.
 471 Albatross Biology and Conservation. Chipping Norton, Australia: Surrey Beatty & Sons,
 472 p113-136.
- 473 Chérel Y, Kooyman GL. 1998. Food of emperor penguins (*Aptenodytes forsteri*) in the
 474 western Ross Sea, Antarctica. Marine Biology 130: 335-344.
- 475 Chérel Y, Ridoux V, Rodhouse PG. 1996. Fish and squid in the diet of king penguin chicks
 476 *Aptenodytes patagonicus* during winter at sub-Antarctic Crozet Islands. Marine Biology 126:
 477 559-570.
- 478 Chérel Y, Weimerskirch H. 1999. Spawning cycle of onychoteuthid squids in the southern
 479 Indian Ocean: new information from seabird predators. Marine Ecology Progress Series 188:
 480 93-104.
- 481 Clarke M. 1980. Cephalopoda in the diet of sperm whales of the southern hemisphere and
 482 their bearing on sperm whale biology. Discovery Reports, p1-324.
- 483 Collins MA, Allcock AL, Belchier M. 2004. Cephalopods of the South Georgia slope. Journal
 484 of the Marine Biological Association of the United Kingdom 84: 415-419.
- 485 Collins MA, Rodhouse PGK. 2006. Southern ocean cephalopods. Advances in Marine
 486 Biology 50: 191-265.
- 487 Constable AJ, Melbourne-Thomas J, Corney SP, Arrigo K, Barbraud C, Barnes D, Bindoff N,
 488 Boyd P, Brandt A, Costa DP, Davidson A, Ducklow H, Emmerson L, Fukuchi M, Gutt J,
 489 Hindell MA, Hofmann EE, Hosie G, Iida T, Jacob S, Johnston NM, Kawaguchi S, Koubbi P,
 490 Lea M-A, Makhado A, Massom R, Meiners K, Meredith M, Murphy E, Nicol S, Richerson K,
 491 Riddle MJ, Rintoul SR, Walker Smith Jr. W, Southwell C, Stark JS, Sumner M, Swadling K,
 492 Takahashi K, Trathan PN, Welsford D, Weimerskirch H, Westwood K, Wienecke B, Wolf-
 493 Gladrow D, Wright S, Xavier JC, Ziegler P. 2014. Change in Southern Ocean ecosystems I:
 494 How changes in physical habitats directly affect marine biota. Global Change Biology 20:
 495 3004-3025.
- 496 Croxall JP, Prince PA. 1996. Cephalopods as prey: seabirds. Philosophical Transactions of
 497 the Royal Society, Series B 351: 1023-1043.
- 498 De Broyer C, Koubbi P, Griffiths HJ, Raymond B, Udekem d'Acoz Cd, Van de Putte AP,
 499 Danis B, David B, Grant S, Gutt J, Held C, Hosie G, Huettmann F, Post A, Ropert-Coudert

500 Y. 2014. Biogeographic Atlas of the Southern Ocean: Scientific Committee on Antarctic
501 Research, Cambridge, UK. 400p.

502 Erickson DJI, Hernandez JL, Ginoux P, Gregg WW, McClain C, Christian J. 2003.
503 Atmospheric iron delivery and surface ocean biological activity in the Southern Ocean and
504 Patagonian region. *Geophys. Res. Lett.* 30: 1609.

505 Feldman GC, McClain CR. 2010. Ocean Color Web & MODIS Aqua Reprocessing. Kuring
506 N, Bailey SW editors: NASA Goddard Space Flight Center

507 Filippova JA, Pakhomov EA. 1994. Young squid in the plankton of Prydz Bay, Antarctica.
508 *Antarctic Science* 6: 171-173.

509 Garcia HE, Locarnini RA, Boyer TP, Antonov JI. 2010. World Ocean Atlas 2009, Volume 3:
510 Dissolved Oxygen, Apparent Oxygen Utilization, and Oxygen Saturation: Washington, DC:
511 U.S. Government Printing Office. 1-344p.

512 Green K, Burton HR. 1993. Comparison of the stomach contents of southern elephant seals,
513 *Mirounga leonina*, at Macquarie and Heard Islands. *Marine Mammal Science* 9: 10-22.

514 Griffiths HJ. 2010. Antarctic Marine Biodiversity: What Do We Know About the Distribution
515 of Life in the Southern Ocean? *PLoS ONE* 5: e11683.

516 Hirzel AH, Le Lay G, Helfer V, Randin C, Guisan A. 2006. Evaluating the ability of habitat
517 suitability models to predict species presences. *Ecological Modelling* 199: 142-152.

518 Imber MJ. 1992. Cephalopods eaten by wandering albatrosses *Diomedea exulans* L. breeding
519 at six circumpolar localities. *Journal of the Royal Society of New Zealand* 22: 243-263.

520 Kennicutt II MC, Chown SL, Cassano JJ, Liggett D, Massom R, Peck LS, Rintoul SR, Storey
521 J, W. V., Vaughn DG, Wilson TJ, Sutherland WJ, Allison I, Ayton J, Badhe R, Baeseman J,
522 Barrett PJ, Bell RE, Bertler N, Bo S, Brandt A, Bromwich D, Cary C, Clark MS, Convey P,
523 Costa ES, Cowan D, DeConto R, Dunbar R, Elfring C, Escutia C, Francis J, Fricker HA,
524 Fukuchi M, Gilbert N, Gutt J, Havermans C, Hik D, Hosie G, Jones C, Kim Y, Le Maho Y,
525 Lee S, Leppe M, Leichenkova G, Li X, Lipenkov V, Lochte K, López-Martínez J, Lüdecke C,
526 Lyons WB, Marensi S, Miller H, Morozova P, Naish T, Nayak S, Ravindra R, Retamales J,
527 Ricci CA, Rogan-Finnemore M, Ropert-Coudert Y, Samah AA, Sanson L, Scambos T,
528 Schloss I, Shiraishia K, Siegert MA, Simões J, Sparrow MD, Storey B, Wall DH, Walsh JC,
529 Wilson G, Winter J-G, Xavier JC, Yang H. 2014. Six priorities for Antarctic Science. *Nature*
530 512: 23-25.

531 Lu CC, Williams R. 1994. Contribution to the biology of squid in the Prydz Bay region,
532 Antarctica. *Antarctic Science* 6: 223-229.

533 NIWA. 2014. New Zealand fish and squid distributions from research bottom trawls.
534 Southwestern Pacific OBIS, National Institute of Water and Atmospheric Research (NIWA),
535 Wellington, New Zealand, 486781 records, Online
536 <http://nzobisipt.elasticbeanstalk.com/resource.do?r=obisprovider> released on May 8,
537 2014. – In.

538 OBIS. 2013. Data from the Ocean Biogeographic Information System: Web.
539 <http://www.iobis.org> (consulted on 2013/09/01).

540 Pellissier L, Rohr RP, Ndiribe C, Pradervand J-N, Salamin N, Guisan A, Wisz M. 2013.
541 Combining food web and species distribution models for improved community projections.
542 *Ecology and Evolution* 3: 4572-4583.

543 Pethybridge HR, Nichols PD, Virtue P, Jackson GD. 2013. The foraging ecology of an
544 oceanic squid, *Todarodes filippovae*: The use of signature lipid profiling to monitor
545 ecosystem change. *Deep Sea Research Part II: Topical Studies in Oceanography* 95: 119-128.

546 Phillips RA, Croxall JP, Silk JRD, Briggs DR. 2008. Foraging ecology of albatrosses and
547 petrels from South Georgia: two decades of insights from tracking technologies. *Aquatic*
548 *Conservation: Marine and Freshwater Ecosystems* 17: S6-S21.

549 Phillips SJ, Anderson RP, Schapire RE. 2006. Maximum entropy modeling of species
550 geographic distributions. *Ecological Modelling* 190: 231–259.

551 Phillips SJ, Dudík M, Elith J, Graham CH, Lehmann A, Leathwick J, Ferrier S. 2009. Sample
552 selection bias and presence-only distribution models: implications for background and
553 pseudo-absence data. *Ecological Applications* 19: 181–197.

554 Radosavljevic A, Anderson RP. 2014. Making better MAXENT models of species
555 distributions: complexity, overfitting and evaluation. *Journal of Biogeography* 41: 629–643.

556 Raymond B. 2012. Polar Environmental Data Layers. Australian Antarctic Data Centre -
557 CAASM Metadata
558 (https://data.aad.gov.au/aadc/metadata/metadata_redirect.cfm?md=/AMD/AU/Polar_Environmental_Data).
559

560 Raymond B, Lea M-A, Patterson T, Andrews-Goff V, Sharples R, Charrassin J-B, Cottin M,
561 Emmerson L, Gales N, Gales R, Goldsworthy SD, Harcourt R, Kato A, Kirkwood R, Lawton
562 K, Ropert-Coudert Y, Southwell C, van den Hoff J, Wienecke B, Woehler EJ, Wotherspoon
563 S, Hindell MA. 2014. Important marine habitat off east Antarctica revealed by two decades of
564 multi-species predator tracking. *Ecography*.

565 Raymond B, Marshall M, Nevitt GA, Gillies CL, van den Hoff J, Stark JS, Losekoot M,
566 Woehler EJ, Constable AJ. 2011. A Southern Ocean dietary database. *Ecology* 92: 1188.

567 Raymond B, Shaffer SA, Sokolov S, Woehler EJ, Costa DP, Einoder L, Hindell M, Hosie G,
568 Pinkerton M, Sagar PM. 2010. Shearwater foraging in the Southern Ocean: the roles of prey
569 availability and winds. *PLoS ONE* 5: e10960.

570 Rio MH, Guinehut S, Larnicol G. 2011. New CNES-CLS09 global mean dynamic
571 topography computed from the combination of GRACE data, altimetry, and in situ
572 measurements. *Journal of Geophysical Research: Oceans* 116: C07018.

573 Robertson G, Williams R, Green K, Robertson L. 1994. Diet composition of emperor penguin
574 chicks *Aptenodytes forsteri* at two Mawson Coast colonies, Antarctica. *Ibis* 136: 19-31.

575 Rodhouse PG. 1989. Pelagic cephalopods caught by nets during the Antarctic research cruises
576 of the 'Polarstern' and 'Walther Herwig', 1985 - 1987. *Archiv für Fischerei Wissenschaft* 39:
577 111-121.

578 Rodhouse PG. 1990. Cephalopod fauna of the Scotia Sea at South Georgia: potential for
579 commercial exploitation and possible consequences. Kerry KR, Hempel G editors. *Antarctic
580 Ecosystems: Ecological Change and Conservation*. Berlin: Springer-Verlag, p289-298.

581 Rodhouse PG. 1997. Precautionary measures for a new *Martialia hyadesi* (Cephalopoda,
582 Ommastrephidae) fishery in the Scotia Sea: an ecological approach. *CCAMLR Science* 4:
583 125-140.

584 Rodhouse PG. 1998a. Large and meso-scale distribution of the ommastrephid squid *Martialia
585 hyadesi* in the Southern Ocean: a synthesis of information relevant to fishery forecasting and
586 management. *Korean Journal of Polar Research* 8: 145-154.

587 Rodhouse PG. 1998b. *Todarodes filippovae* in the Southern Ocean: an appraisal for
588 exploitation and management. *Large pelagic squids*. Tokyo: Japan Marine Fishery Resources
589 Research Center, p207-215.

590 Rodhouse PG. 2013. Role of squid in the Southern Ocean pelagic ecosystem and the possible
591 consequences of climate change. *Deep Sea Research Part II: Topical Studies in
592 Oceanography*. 95: 129-138.

593 Rodhouse PG, Piatkowski U. 1995. Fine-scale distribution of juvenile cephalopods in the
594 Scotia Sea and adaptive allometry of the brachial crown. *Marine Biology* 124: 111-117.

595 Rodhouse PG, Prince PA, Trathan PN, Hatfield EMC, Watkins JL, Bone DG, Murphy EJ,
596 White MG. 1996. Cephalopods and mesoscale oceanography at the Antarctic Polar Front:

597 satellite tracked predators locate pelagic trophic interactions. *Marine Ecology Progress Series*
598 136: 37-50.

599 Rodhouse PG, Xavier JC, Griffiths H. 2014. Southern Ocean squid. De Broyer C, Koubbi P,
600 Griffiths H, Danis B, David B, Grant S, Gutt J, Held C, Hosie G, Huettmann F, Post A,
601 Raymond B, Ropert-Coudert Y, van de Putte A editors. *The CAML / SCAR-MarBIN*
602 *Biogeographic Atlas of the Southern Ocean*. Cambridge: Scientific Committee on Antarctic
603 Research, p284-289.

604 Roeleveld MAC. 1998. The status and importance of cephalopod systematics in Southern
605 Africa. *S. Afr. J. Mar. Sci.* 20: 1–16.

606 Roper C, Sweeney M, Nauen C. 1984. *FAO species catalogue, Cephalopods of the World. An*
607 *annotated illustrated catalogue of species of interest to fisheries: FAO Fish Synop.* 1-227p.

608 Roper CFE. 1969. Systematics and zoogeography of the worldwide bathypelagic squid
609 *Bathyteuthis* (Cephalopoda: Oegopsida). *U.S. National Museum Bulletin* 291: 210pp.

610 Sarmiento JL, Gruber N, Brzezinski MA, Dunne JP. 2004. High-latitude controls of
611 thermocline nutrients and low latitude biological productivity. *Nature* 427: 56-60.

612 Smale MJ. 1996. Cephalopods as prey. IV. Fishes. *Phil.Trans.R.Soc.Lond.B* 351: 1067-1081.

613 Smith WHF, Sandwell DT. 1997. Global seafloor topography from satellite altimetry and ship
614 depth soundings. *Science* 277: 1957–1962.

615 Sokolov S, Rintoul R. 2007. On the relationship between fronts of the Antarctic Circumpolar
616 Current and surface chlorophyll concentrations in the Southern Ocean. *Journal of*
617 *Geophysical Research* 112: C07030.

618 Spreen G, Kaleschke L, Heygster G. 2008. Sea ice remote sensing using AMSR-E 89 GHz
619 channels. *Journal of Geophysical Research* 113: C02S03.

620 Trübenbach K, Teixeira T, Diniz M, Rosa R. 2013. Hypoxia tolerance and antioxidant
621 defense system of juvenile jumbo squids in oxygen minimum zones. *Deep Sea Research Part*
622 *II: Topical Studies in Oceanography* 95: 209-217.

623 Turner J, Bindshadler R, Convey P, di Prisco G, Fahrbach E, Gutt J, Hodgson D, Mayewski
624 P, Summerhayes C. 2009. *Antarctic climate change and the environment: Scientific*
625 *Committee for Antarctic Research*, Cambridge, UK. 256 ppp.

626 Voss NA, Nesis KN, Rodhouse PG. 1998. Systematics, biology and biogeography of the
627 family Histioteuthidae (Oegopsida). Voss NA, Veechione, M., Toll, R. O., & Sweeney, M. J.
628 editor. *Systematics and Biogeography of Cephalopods: Smithsonian contribution to Zoology*,
629 p293-373.

630 Xavier JC. 1997. *Distribution of cephalopods in Antarctic, sub-Antarctic and subtropical*
631 *waters. Faculdade das Ciencias do Mar e do Ambiente. Faro: Universidade do Algarve,*
632 *Portugal, p 87.*

633 Xavier JC, Allcock L, Cherel Y, Lipinski MR, Gomes-Pereira JN, Pierce G, Rodhouse PGK,
634 Rosa R, Shea L, Strugnell J, Vidal E, Villanueva R, Ziegler A. 2015. Future challenges in
635 cephalopod research. *Journal of the Marine Biological Association of the UK* 95: 999-1015.

636 Xavier JC, Cherel Y. 2009. *Cephalopod beak guide for the Southern Ocean: British Antarctic*
637 *Survey.* 129p.

638 Xavier JC, Cherel Y, Roberts J, Piatkowski U. 2013. How do cephalopods become available
639 to seabirds: can fish gut contents from tuna fishing vessels be a major food source of deep-
640 dwelling cephalopods? *ICES J. of Marine Science* 70: 46-49.

641 Xavier JC, Croxall JP, Cresswell KA. 2005. Boluses: an effective method to assess the
642 proportions of cephalopods in the diet of albatrosses. *Auk* 122: 1182-1190.

643 Xavier JC, Croxall JP, Reid K. 2003a. Inter-annual variation in the diet of two albatross
644 species breeding at South Georgia: implications for breeding performance. *Ibis* 145: 593-610.

645 Xavier JC, Croxall JP, Trathan PN, Rodhouse PG. 2003b. Inter-annual variation in the
646 cephalopod component of the diet of wandering albatrosses *Diomedea exulans* breeding at
647 Bird Island, South Georgia. *Marine Biology* 142: 611-622.

648 Xavier JC, Geraint GA, Croxall JP. 2006. Determining large scale distribution of pelagic
649 cephalopods, fish and crustaceans in the South Atlantic from wandering albatross (*Diomedea*
650 *exulans*) foraging data. *Ecography* 29: 260-272.

651 Xavier JC, Rodhouse PG, Purves MG, Daw TM, Arata J, Pilling GM. 2002. Distribution of
652 cephalopods recorded in the diet of Patagonian toothfish (*Dissostichus eleginoides*) around
653 South Georgia. *Polar Biology* 25: 323-330.

654 Xavier JC, Rodhouse PG, Trathan PN, Wood AG. 1999. A Geographical Information System
655 (GIS) atlas of cephalopod distribution in the Southern Ocean. *Antarctic Science* 11: 61-62.

656 Xavier JC, Trathan PN, Croxall JP, Wood AG, Podestá GP, Rodhouse PG. 2004. Foraging
657 ecology and interactions with fisheries of wandering albatrosses at South Georgia. *Fisheries*
658 *Oceanography* 13: 324-344.

659 Xavier JC, Walker K, Elliot G, Cherel Y, Thompson D. 2014. Cephalopod fauna of South
660 Pacific waters: new information from breeding New Zealand wandering albatrosses. *Marine*
661 *Ecology Progress Series* 513: 131-142.

662 Xavier JC, Wood AG, Rodhouse PG, Croxall JP. 2007. Interannual variations in cephalopod
663 consumption by albatrosses at South Georgia: implications for future commercial exploitation
664 of cephalopods. *Marine and Freshwater Research* 58: 1136-1143.

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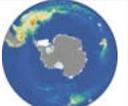
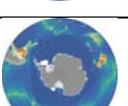
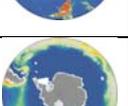
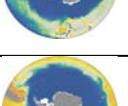
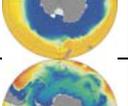
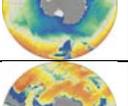
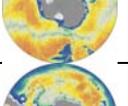
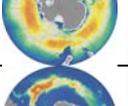
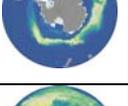
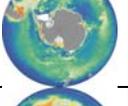
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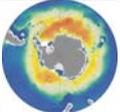
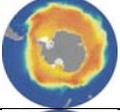
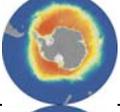
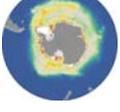
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Table 1. Predictor variables used in the species distribution habitat suitability modeling.

Variable	Description	Source and references	Biological relevance
Sea surface temperature (SST)	Sea surface temperature summer climatology, calculated over the 2002/03 to 2012/13 austral summer seasons	MODIS Aqua (Feldman and McClain 2010)	General water mass properties including positions of fronts, which can represent areas of different prey and of prey-aggregation
Sea ice cover	The average proportion of the year for which sea ice is present. Concentration data from 1-Jan-2003 to 31-Dec-2010 was used. The fraction of time each pixel was covered by sea ice of at least 85% concentration was calculated	AMSR-E satellite estimates of daily sea ice concentration (Spreen et al. 2008)	Indicator of sea ice cover, including polynyas, which affects ecosystem structure and prey availability
Depth	Measured and estimated seafloor topography from satellite altimetry and ship depth soundings	Smith and Sandwell (1997) V15.1	Water mass properties
Sea surface height (SSH)	Mean dynamic topography (sea surface height relative to geoid)	CNES-CLS09 Mean Dynamic Topography v1.1 (Rio et al. 2011)	Water mass properties
Chlorophyll- <i>a</i> (Chl- <i>a</i>)	Near-surface chlorophyll- <i>a</i> summer climatology, calculated over the 2002/03 to 2012/13 austral summer seasons	MODIS Aqua (Feldman and McClain 2010)	Productivity, ecosystem structure
Oxygen minimum	Minimum dissolved oxygen value in the top 1000m of the water column	World Ocean Atlas 2009 annual climatology (Garcia et al. 2010)	Water mass properties, potential habitat niche for cephalopods (Trübenbach et al. 2013)

Table 2. Habitat suitability modeling results summary (N occ.: Number of occurrences, AUC: area under the receiver-operating characteristic, OW: open water, PF: polar front, SAF: Sub-Antarctic Front, SIZ: sea ice zone. For variable names see Table 1). The thumbnail maps are reproductions of figures 3b-17b, and are included here to allow a convenient comparison of the broad spatial patterns in the modelling results.

Species	N occ.	Train /test AUC	Variables used in model (permutation importance, %)		Typical habitat from model predictions					
					SST (°C)	Depth (m)	Ice	O ₂ min. (ml/l)	Chl- <i>a</i> (mg m ⁻³)	Water mass
<i>Doryteuthis gahi</i>	149	0.99/0.99	Chl (60.5) Depth (34.3) Ice (5.1)		≥9	<400	OW		>0.75	
<i>Martialia hyadesi</i>	260	0.94/0.94	Chl (47.2) SST (43.8) Ice (5.8) Depth (3.2)		3–15		OW		>0.3	PF to SAF
<i>Moroteuthis ingens</i>	3808	0.61/0.61	Depth (65.5) SST (27.4) SSH (4.8) Ice (2.3)		3–16	300–1500	OW		>0.2	
<i>Moroteuthis robsoni</i>	342	0.75/0.75	SSH (41.5) Chl (32.5) O ₂ min (16.4) Depth (9.7)		7–18	>1000	OW		0.15–0.55	SAF and north
<i>Todarodes filippovae</i>	1173	0.62/0.61	SSH (84.9) Depth (15.1)		≥10		OW			SAF and north
<i>Histioteuthis atlantica</i>	106	0.89/0.89	Depth (85.4) SST (14.6)		≥5	>3000	OW		<0.6	
<i>Histioteuthis eltaninae</i>	110	0.90/0.90	Depth (100)			>3000	OW–70% cover		<0.65	
<i>Bathyteuthis abyssicola</i>	548	0.91/0.91	Depth (53.8) SST (37.2) O ₂ min (9.0)		≤12	>2500	OW–70% cover	<4.8	<0.7	
<i>Slosarczykovia circumantarctica</i>	1304	0.96/0.96	SSH (74.4) Ice (19.6) Chl (4.2) SST (1.8)		0–7	>1500	OW–20% cover	4–4.5	0.15–0.65	SAF to parts of SIZ
<i>Gonatus antarcticus</i>	120	0.83/0.82	SST (66.2) Chl (33.8)		< 12				>0.4	
<i>Kondakovia longimana</i>	100	0.95/0.94	SSH (82.8) Chl (12.1) Ice (5.2)		<6	>500	OW–70% cover		>0.15	PF and south

<i>Mesonychoteuthis hamiltoni</i>	234	0.93/ 0.92	SSH (57.5) Depth (37) O ₂ min (5.4)		≤11	>2200		<4.75		SAF and south
<i>Galiteuthis glacialis</i>	1449	0.88/ 0.88	SST (79.5) SSH (20.5)		≤6	>500	SIZ- OW			PF and south
<i>Alluroteuthis antarcticus</i>	124	0.93/ 0.93	SST (98.6) Depth (1.4)		≤3	>500	SIZ and OW			South of PF
<i>Psychroteuthis glacialis</i>	316	0.94/ 0.93	SST (66.6) SSH (24.0) Chl (7.4) Ice (2.0)		≤3	>500	SIZ and OW			South of SAF

