Sexual segregation in a highly pagophilic and sexually dimorphic marine predator

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Abstract : Sexual segregation is common in many species and has been attributed to intra-specific competition, sex-specific differences in foraging efficiency or in activity budgets and habitat choice. However, very few studies have simultaneously quantified sex-specific foraging strategies, at sea distribution, habitat use, and trophic ecology. Moreover, these studies come from low latitude areas reflecting a lack of evidence for polar species. We investigated sexual segregation in snow petrels *Pagodroma nivea* and combined movement, foraging trip efficiency, stable isotope and oceanographic data to test whether sexual segregation results from sex-specific habitat use. Breeding birds foraging in the Dumont d'Urville sea, Antarctica, were tracked during incubation. Some similarities between males and females foraging characteristics did not support the sexual segregation hypothesis. Indeed, space-use sharing and utilization distribution, $\delta^{13}C$ values and foraging trip performances (trip duration, length, speed and directions, mass gain, proportion mass gain) were similar between males and females.. However, there was support for sexual segregation in foraging characteristics linked to foraging habitats. Females foraged less than males in areas with higher sea ice concentration (SIC >70%) and had lower $\delta^{15}N$ values in plasma, blood cells and feathers. Foraging efficiency (proportionate daily mass gain while foraging), was greater for females than for males, and was greater for larger females with deeper bills. Females were more efficient than males during short (<2 days) foraging trips, and for females, but not for males, mass gain, proportion mass gain and body condition at return from a foraging trip were positively correlated to SIC of the foraging areas. Together, these results suggest an absence of sexual segregation at large spatial scales in snow petrels during incubation, but strongly support habitat segregation between high (>70%) more profitable SIC (males) and low SIC areas (females), probably driven by intra-specific competition. Therefore, male and female snow petrels segregate at small spatial scales mainly determined by habitat (SIC) characteristics.

Keywords : bio-logging, competition, foraging, isotopic niche, *Pagodroma nivea*, sea ice concentration, snow petrel

1 Introduction

Sexual segregation occurs in a many living animals including invertebrates (Hochkirch et al. 2007, Romey & Wallace 2007) and vertebrates (Ruckstuhl & Neuhaus 2005, Wearmouth & Sims 2008) but also in plants (Harder et al. 2000). Investigating sexual segregation is of particular relevance from a fundamental point of view to understand how and why the sexes differentially distribute themselves and the consequences on population processes and dynamics. It is also relevant from a management and conservation point of view since sex specific distribution influences overlap with spatial distribution of human activities and/or contamination gradient (Carravieri et al. 2014). Two main concepts have been proposed to describe sexual segregation: social segregation, where males and females tend to form single-sex groups within the same or homogeneous habitat; and habitat segregation, where males and females use different habitats within a home range and with habitats differing in their amount or quality of forage distributed heterogeneously or patchily (Conradt 2005, Ruckstuhl 2007). Both social segregation and habitat segregation can or cannot lead to spatial or temporal segregation, which have been be considered as auxiliary concepts (Conradt 2005, Ruckstuhl 2007). However, the distinction between habitat segregation and spatial segregation is scale-dependent: spatial segregation is a mechanism to avoid competing for the same habitat by choosing different locations, while habitat segregation is a mechanism to avoid competing for resources at the same location. These concepts can also be understood in the framework of equalizing and stabilizing mechanisms applied to movement ecology (Chesson 2000, Jeltsch et al. 2013).

Several hypotheses have been proposed to explain social and habitat segregations (Conradt 2005, Ruckstuhl 2007, Wearmouth & Sims 2008). In solitary animals, social segregation is unlikely to occur since by definition a single animal is not social (Conradt 1998, Neuhaus & Ruckstuhl 2004), except perhaps in rare cases (Martin & Da Silva 2004). Four main hypotheses explain habitat segregation in solitary species (Ruckstuhl 2007, Wearmouth & Sims 2008). The forage-selection hypothesis, which incorporates the scramble competition hypothesis, suggests sex differences in nutritional requirements linked to sex-specific differences in body size (Gross 1998). The larger sex individuals select habitats where intake rates are high whereas the smaller sex individuals are constrained to sites where they can obtain a high-quality food (Beier 1987, Barboza & Bowyer 2000). Alternatively, one sex may forage more efficiently, thus outcompeting and excluding the other (scramble-competition hypothesis or intersexual competition hypothesis) (Clutton□Brock et al. 1987). The activity-budget hypothesis, initially developed for group-living species, was extended to solitary species and to species with unequal reproductive investment (Wearmouth & Sims 2008). This hypothesis proposes that sex differences in activity budgets will increase with divergence in the body size of the sexes. Therefore, the sex-specific energy requirements will result in sex-specific habitat used due to allometric relationships between body size and metabolic rate. Finally, the predation-risk hypothesis proposes sexual differences in risk of predation and in reproductive strategies (Main et al. 1996), and the thermal niche-fecundity hypothesis assumes that sex differences occur in the temperature at which fecundity is maximized (Sims 2005).

Sexual segregation has been widely studied among terrestrial animals, particularly mammals, but only relatively recently in marine organisms (Wearmouth & Sims 2008). Yet, despite an ongoing interest in sexual segregation in marine animals such as seabirds and marine mammals (Lewis et al. 2002, Elliott et al. 2010, Phillips et al. 2011, Mancini et al.

2013, Baylis et al. 2016, Kernaléguen et al. 2016), the underlying causes and the mechanisms driving habitat segregation remain poorly understood. In addition, very few studies focused on between-sex differences in habitat segregation in relation to dynamic oceanographic features (Pinet et al. 2012, Cleasby et al. 2015, Paiva et al. 2017), thereby limiting our ability to distinguish between the concurrent sexual segregation hypotheses. Moreover, these studies come from temperate or tropical areas reflecting a lack of evidence for polar species. However, foraging strategies may differ between polar, temperate and tropical oceanographic environments, at least in seabirds (Baduini & Hyrenbach 2003, Weimerskirch 2007). Furthermore, for practical, technical and ethical reasons most studies that have investigated sexual segregation on marine animals have focused on large species (Phillips et al. 2011), complicating the possibility to discriminate between the various hypotheses proposed to explain sexual segregation.

In this study we aimed to quantify sexual differences in the foraging strategies, at sea distribution, habitat use, and trophic ecology of a sexually dimorphic polar seabird, the snow petrel, *Pagodroma nivea*, during the incubation period. Snow petrels are endothermic animals, therefore excluding the thermal niche-fecundity hypothesis as an explanatory hypothesis. Since predation on this species is occasional and no sex-specific predation is known to occur (Barbraud 1999), the predation-risk hypothesis can be discounted. Therefore, both the forage-selection hypothesis and the activity budget hypothesis can be highlighted as possible mechanisms for segregation in this species. There is considerable overlap between the forage-selection hypothesis and the activity-budget hypothesis, complicating our ability to make clear predictions to distinguish between the two, and to estimate the relative support of each hypothesis (Wearmouth & Sims 2008). Nevertheless, using GPS tracking data, isotopic data and environmental data we addressed the following main questions: (1) do female snow petrels differ from males in their foraging tactics, distribution and habitat use?; (2) how are

126 body reserves regulated during incubation in the two sexes?; and (3) do sex-specific
127 morphological characteristics influence foraging efficiency? Based on results from
128 comparative studies suggesting that dimorphic seabird species from polar/temperate regions
129 are more prone to show trophic or spatial segregation than dimorphic species from the tropics
130 (Mancini et al. 2013), and on a relationship between sexual segregation in diet and sexual size
131 dimorphism in seabirds (Phillips et al. 2011), we predicted sexual segregation in diet and/or
132 spatial segregation in the snow petrel, which is one of the most sexually dimorphic seabird
133 species (Croxall 1982, Fairbairn & Shine 1993).
134

135 # 2 Material and methods

136 ## 2.1 Study species

137 The snow petrel is endemic to Antarctica and the Southern Ocean, with a circumpolar
138 breeding distribution (Croxall et al. 1995). It is a specialist forager and ship-based
139 observations indicate that this is the most pagophilic species amongst flying seabirds,
140 occurring only where there is some degree of sea ice cover (Griffiths 1983, Ainley et al. 1984,
141 1986), generally within the marginal ice zone and areas of heavy ice concentrations (Ainley et
142 al. 1992, 1993). Snow petrels forage by flying rapidly along the edges of ice floes, ice shelves
143 and icebergs in search of its prey (Ainley et al. 1984). The species feeds primarily on fish,
144 including the myctophid *Electrona antarctica* in oceanic waters and the pelagic notothenid
145 *Pleuragramma antarctica* (Antarctic silverfish) in neritic waters; they prey also upon
146 swarming crustaceans, the Antarctic (*Euphausia superba*) and ice (*E. crystallorophias*) krill,
147 and the hyperiid amphipod *Themisto gaudichaudii* (Ainley et al. 1984, 1991, Ridoux &
148 Offredo 1989, Van Franeker & Williams 1992, Ferretti et al. 2001). At Pointe Géologie
149 (Adélie Land), undetermined fish dominated the chick diet in 1982 (Ridoux & Offredo 1989)
150 and fish items identified in 1994 were all Antarctic silverfish (authors'unpublished data). Prey

151 are caught by dipping and surface-seizing (Harper et al. 1985) generally on the wing but also
152 by ambush feeding (Ainley et al. 1984).

153 Snow petrels breed in crevices and under boulders. Adult birds arrive at the colonies in
154 late October to copulate before departing at sea for a two to three week pre-laying exodus, and
155 females lay a single egg in early December (Mougin 1968, Isenmann 1970). Incubation lasts
156 ≈44 days on average during which males and females alternately incubate their egg until
157 hatching (Brown 1966, Barbraud et al. 1999). After hatching the chick is guarded by parents
158 alternating short spells until it attains homeothermy. Then the chick is left unattended and
159 regularly fed by both parents until fledging, which occurs on average ≈47 days after hatching.
160 Adults leave the colony during the first two weeks of March before dispersing at sea where
161 they remain in the sea ice zone during the non-breeding period (Delord et al. 2016).
162

163 **2.2 Fieldwork**

164 Fieldwork was carried out at Ile des Pétrels (66°40'S, 140°01'E), Pointe Géologie
165 archipelago, Adélie Land, East Antarctica, between 7 December 2015 and 17 January 2016.
166 This corresponds to the incubation period. On average 550 pairs of snow petrels breed on Ile
167 des Pétrels in dense colonies or in loosely aggregated nests (CEBC-CNRS unpublished data).
168 By daily visits at 36 nests, we studied laying dates and the duration of the foraging trips and
169 incubation shifts of 36 males and 36 females until hatching. Incubating birds were identified
170 using their metal ring number. Sixty five snow petrels (n = 36 females and n = 29 males) were
171 tracked with GPS loggers (nanoFix-Geo; PathTrack Limited, UK) during the incubation
172 period. We tracked only one foraging trip per bird to minimize disturbance and to ensure
173 independence between trips. The devices weighed 2.2 g, which represented between 0.5% and
174 0.8% of the birds' mass, thus well below the 3% threshold advised by Phillips et al.(2003).
175 Birds were manually captured at the nest and weighted (± 5 g) in a bag with a Pesola spring

176 balance before being equipped with a GPS. The birds were initially sexed by vocalization
177 when approached on the nest and handled (male calls have a lower pitch and a lower rhythm
178 than those of females (Guillotin & Jouventin 1980, Barbraud et al. 2000). GPS units were
179 deployed on birds about to leave for a foraging trip (i.e. when both partners were at the nest)
180 and were attached to the two central tail feathers using Tesa® tape. The GPS recorded
181 locations at 15, 30, 40 or 60 min intervals. Several intervals (15 min, n = 15; 30 min, n = 4; 40
182 min, n = 43; 60 min, n = 3) were tested to estimate the minimum interval frequency that
183 allowed the GPS battery to last for a complete foraging trip. Birds were recaptured on the day
184 they returned to the nest following their foraging trip, weighed, measured (wing length ± 1
185 mm with a ruler, tarsus length, bill length, and bill depth ± 0.1 mm with calipers) and the
186 loggers were recovered. All birds were recaptured but three birds lost their GPS during the
187 foraging trip. Data from all other GPS (n = 62) were retrieved successfully.
188
189 **2.3 Tissue sampling, molecular sexing and stable isotopes**
190 Adults equipped with GPS and 24 additional individuals (11 females and 13 males) were
191 sampled during incubation for stable isotope and molecular sexing analyses. A blood sample
192 from the alar vein was taken immediately after capture of the bird upon return from a foraging
193 trip using a 1-mL heparinized syringe and a 25-gauge needle and maintained at 4°C until
194 being processed. Collected blood volumes ranged from 0.50 to 0.80 mL. Blood samples were
195 separated into plasma and blood cells by centrifugation at 12,000 rpm for 5 min, within 2-3
196 hours of sampling and stored frozen at -20°C until analyses at the laboratory. For each
197 individual, 6 whole body feathers were pulled out from the upper chest and stored dry in
198 sealed individual plastic bags for stable isotope analysis.

From a subsample of blood cells, the sex was determined by polymerase chain reaction amplification of part of two highly conserved genes present on the sex chromosomes as detailed in Weimerskirch et al. (2005).

Stable carbon ($\delta^{13}C$) and nitrogen ($\delta^{15}N$) isotope ratios in the blood cells, plasma and body feathers of snow petrels were determined to investigate the trophic choices of each sex and consistency of their foraging niche over time. The isotopic method was validated in the Southern Ocean for several seabird species: $\delta^{15}N$ values mainly define the trophic position, with values increasing with trophic level (Cherel et al. 2010), and $\delta^{13}C$ values indicate the latitude of the foraging habitat (Cherel & Hobson 2007, Jaeger et al. 2010). Plasma has a half-life of about 3 days (Hobson & Clark 1993), a shorter period than the average trip duration during incubation (≈7 days, Barbraud et al. 1999), and represents prey ingestion and trophic ecology during the last trip before sampling (Cherel et al. 2005a). Blood cells have a half-life of about 30 days (Hobson & Clark 1993) and represent dietary information integrated over a few months. Feathers contain dietary information at the time they were grown, because keratin is inert after synthesis (Hobson & Clark 1992, 1993, Bearhop et al. 2002). In snow petrels body moult is a gradual process extending over at least 4 months in summer and autumn. It begins during incubation, but most body feathers grow in the weeks following completion of breeding, i.e. from February to April (Maher 1962, Beck 1969). Therefore, isotopic values of body feathers contain information about diet near the end of the previous breeding season and the beginning of the previous non-breeding season.

Feathers (one single feather per bird) were cleaned to remove surface contaminants using a 2:1 chloroform:methanol solution followed by two methanol rinses. They were then oven dried for 48 h at 50°C and cut into small pieces using stainless steel scissors. Blood cells and plasma samples were freeze-dried and powdered. Since avian plasma, unlike blood cells, contains a high and variable lipid content that affect its $\delta^{13}C$ values, lipids were removed from

224 plasma samples using chloroform/methanol (Cherel et al. 2005a, Cherel et al. 2005b). Then,
225 tissue sub-samples were weighed with a microbalance (aliquots mass: ≈ 0.3 mg dw), packed
226 in tin containers, and nitrogen and carbon isotope ratios were subsequently determined at the
227 laboratory LIENSs by a continuous flow mass spectrometer (Thermo Scientific Delta V
228 Advantage) coupled to an elemental analyser (Thermo Scientific Flash EA 1112). Results are
229 presented in the usual δ notation relative to Vienna PeeDee Belemnite and atmospheric N_2 for
230 $\delta^{13}C$ and $\delta^{15}N$, respectively. Replicate measurements of internal laboratory standards
231 (acetanilide and peptone) indicate measurement errors <0.15 ‰ for both $\delta^{13}C$ and $\delta^{15}N$
232 values.
233
234 **2.4 Foraging analysis and spatial usage**
235 Spatial and statistical analyses were performed using R 3.2.1 using the "stats" package (R
236 Development Core Team 2015) and "*adehabitatLT*" package (Calenge 2006, Calenge et al.
237 2009). From the GPS recorded data, foraging trips were reconstructed and data were
238 rediscretized to have one location each 40 min. Some of the trips were largely incomplete
239 (return journey not initiated; n = 15 corresponding to the 15 min intervals) because of battery
240 limitations and were removed from the analysis. For each complete (n = 40) and incomplete
241 (return journey initiated; n = 7) foraging trip, we computed the following foraging indices:
242 maximum distance to the colony (*Dmax*, km), average movement speed (*MS*, km h^{-1}) and
243 daily distance covered (*Dday*, km d^{-1}). For each complete trip, we calculated the additional
244 following metrics: total distance travelled (*Dtotal*, km) and trip duration (*T*, h). Spatial
245 distribution of snow petrels was investigated by producing utilization distributions (UDs 25%,
246 50%, 75% and 95%; Worton 1989) for each individual, using kernel analysis with a cell size
247 of $0.1° \times 0.1°$ and a smoothing parameter (h) that was estimated using the ad hoc method href.
248 Grid cell size was based on the mean accuracy of the devices (≈ 10 m), the mean maximum

speed of flying snow petrels (see Results) and on the time interval between two GPS locations (40 min). To investigate whether space use differed between sexes, we calculated observed overlaps in each UD representing the high core (25%), core (50%), middle (75%) and general (90%) use areas using utilization distribution overlap index (UDOI), which is the most appropriate measure of quantifying similarity among UD estimates (Fieberg & Kochanny 2005). The extent of overlap between male and female home ranges was estimated using Bhattacharyya's affinity (BA), which ranges from 0 (no overlap) to 1 (complete overlap). Using these metrics we performed a randomization procedure to test the null hypothesis that there was no difference in the spatial distribution of males and females at the population level (Breed et al. 2006). The sex of each bird was randomly assigned using the observed sex ratio in our data set and the overlap metric between males and females was calculated for 25%, 50%, 75% and 95% kernels. We performed 1000 randomizations of our dataset from which the probability of accepting the null hypothesis was calculated as the proportion of random overlaps that were smaller than the observed overlap. Since we were testing only if the observed overlap was smaller than random overlap, we considered this as a one-tailed test. Second, we tested the null hypothesis that there was no difference in the extent of overlap in spatial distribution of males and females at the individual level.

For each foraging trip we also calculated the following metrics from the phenotypic data: the body mass change (Δm, in g) between departure and arrival of a foraging trip, the daily mass gain (Mday, in $g.day^{-1}$) calculated as the ratio between Δm and the trip duration, the proportion mass gain calculated as the ratio between Δm and mass at departure for a foraging trip, and the proportion daily mass gain calculated as the ratio between Daym and mass at departure for a foraging trip. A body condition index before departure and after return from a foraging trip was also calculated. To estimate the body condition we used the body measurements to calculate the scale mass index (SMI) as recommended by Peig and Green

(2009, 2010). The SMI adjusts the mass of all individuals to that expected if they had the same body size. We used the score of the first axis of a principal component analysis (PC1) combining wing, bill, tarsus lengths and bill depth to characterize body size. PC1 accounts for 70.9% of the total variance and all measurements are highly correlated with PC1 (Pearson's r > 0.80; P < 0.001). The SMI was calculated for each individual *i* according to the formula:

$$SMI_i = M_i * \left(\frac{L_0}{L_i}\right)^b$$

where M_i and L_i are, respectively, the body mass and the PC1 score of the individual *i*, L_0, is the value of PC1 for the whole studied population and *b* the slope estimate of the RMA (Reduced Major Axis) regression of log-transformed body mass on log-transformed PC1.

2.5 Foraging habitat covariates

To investigate the foraging habitats used by males and females, the tracking locations were categorized as occurring during commuting (outward and inward) or foraging (middle) stage of foraging trips, as commonly used for central-place foragers. Among Procellariformes the distinction between these stages varies greatly between species and breeding stages (Weimerskirch et al. 1997, Phillips et al. 2009). Moreover, at the individual level defining objectively the transition between such behaviors may prove to be difficult (Phillips et al. 2009, Wakefield et al. 2009). To avoid this pitfall, we applied the method used by Wakefield et al. (2009) and Phillips et al. (2009) to determine the stage of the trips at which the transitions occurred at the population level. For each location within a foraging trip the ratio d_{col}/D_{max} was calculated, where d_{col} is the distance from the colony and D_{max} is the maximum distance from the colony reached during that trip. The ratio t/T was also calculated, where *t* is the time elapsed since the beginning of the trip and *T* is the total trip time. Then, the total variance in d_{col}/D_{max} for all locations occurring before t/T was plotted against t/T. The point of inflexion of this curve was determined as well as the value of t/T at this point. Tracking

298 locations recorded before this point were classified as those corresponding to the outward trip.
299 Similarly, the total variance in d_{col}/D_{max} occurring after t/T was plotted against t/T and the t/T
300 value from which a monotonic decrease of the variance began was recorded. Tracking
301 locations recorded after this point were classified as those corresponding to the return trip, and
302 locations between both points were considered as foraging locations.
303 Previous studies have shown that the snow petrel is a sea ice obligate species and remains
304 highly associated with sea ice year round (Griffiths 1983, Ainley et al. 1984, 1986, 1992,
305 1993, Delord et al. 2016). We therefore used sea ice concentration (SIC) to describe the
306 foraging habitat of snow petrels. Although sea surface temperature is commonly used to
307 describe foraging habitats in seabirds, there are very few sea surface temperature observations
308 in regions covered by sea ice, especially in the Southern Ocean (Rayner et al. 2003).
309 Therefore this covariate could not be used. We used passive-microwave estimates of daily sea
310 ice concentration from the Special Sensor Microwave Imager (SSMI/I) brightness
311 temperatures (12.5 × 12.5 km resolution) from the Institut Français de Recherche pour
312 l'Exploitation de la Mer (Ifremer, ftp://ftp.ifremer.fr/ifremer/cersat/products/gridded/psi-
313 concentration/data/antarctic). We also used bathymetry data (ocean depth at one-minute
314 horizontal spatial resolution) obtained from NOAA's ETOPO
315 (https://sos.noaa.gov/datasets/etopo1-topography-and-bathymetry/) as an additional habitat
316 variable. Daily sea ice concentration and depth values were extracted for each foraging
317 location (therefore excluding the commuting part of the trips at sea) on each track using
318 bilinear interpolation from the native ice and depth grids using "*raster*" package in R
319 (Hijmans 2018). Since snow petrels are highly associated with the sea ice region (as defined
320 by the region within >15% sea ice concentration isocline, Cavalieri et al. 1991), the SIC data
321 were filtered to retain SIC values >15%.
322

323 **2.6 Statistical analysis**

324 Isotopic niche of the two sexes was established using the metric SIBER (Stable Isotope
325 Bayesian Ellipses), which is based on a Bayesian framework that confers a robust comparison
326 to be made among data sets concerning different sample sizes (Jackson et al. 2011). The area
327 of the standard ellipse (SEA_C, an ellipse having a 40% probability of containing a
328 subsequently sampled datum) was used to compare female and male isotopic values and their
329 overlap in relation to the total niche width (i.e. both sexes combined), and a Bayesian estimate
330 of the standard ellipse and its area (SEA_B) was used to test whether females' isotopic niche is
331 narrower than males' isotopic niche (Jackson et al. 2011). The *standard.ellipse* and
332 *convexhull* functions were used to calculate these metrics from SIBER implemented in the
333 package '*SIAR'* (Parnell et al. 2010) under R.

334 Consistency in foraging niche was estimated following Votier et al. (2010) and Ceia et al.
335 (2012), by regressing stable isotope ratios in plasma on those of blood cells to obtain an index
336 of consistency in carbon source (habitat) and trophic level. Since $\delta^{13}C$ has a trophic
337 component, we used the studentized residuals of the relationship with $\delta^{15}N$ in the same tissue
338 (male plasma: $F_{1,25}$=1.438 , P=0.242, r=0.233; male blood cells: : $F_{1,36}$=0.838, P=0.366,
339 r=0.151; female plasma: $F_{1,33}$=1.470 , P=0.234, r=0.206; female blood cells: $F_{1,45}$=6.507,
340 P=0.014, r=0.355) to determine the degree of short-term repeatability in $\delta^{13}C$ independently
341 of trophic effects. Longer-term foraging consistency was estimated by regressing stable
342 isotope values of blood cells (actual breeding period) with those of feathers (most likely the
343 end of the previous breeding period and subsequent fall at sea). We also used the residuals to
344 correct the trophic component associate with $\delta^{13}C$ by regressing these values upon $\delta^{15}N$
345 signatures in feathers (male: $F_{1,36}$=1.945, P=0.172, r=0.226; female: $F_{1,45}$=5.863, P=0.020,
346 r=0.340).

347 Foraging probability was modelled using a binomial generalized additive mixed model
348 (GAMM) in the '*gamm4'* package in R (Wood et al. 2017). This allowed for the possibility of
349 nonlinear responses to environmental covariates, which we expected. The response variable
350 was the tracking location, which was coded as 1 for a foraging location and as 0 for a
351 commuting location, and explanatory variables were sea ice concentration and bathymetry.
352 Because interactions between the variable sex and environmental covariates would be difficult
353 to interpret in complex nonlinear models, separate models were developed for male and
354 female birds. Models included sea ice concentration and bathymetry as fixed factors, and bird
355 identity as a random term to account for pseudoreplication issues. The smoothing parameter
356 was chosen automatically using generalized cross-validation. To model spatial auto-
357 correlation an isotropic thin plate spline was included, set up as a two dimensional smoother
358 based on both x and y coordinates (Cleasby et al. 2015). To ascertain whether collinearity
359 between covariates may have occurred we examined the correlations between environmental
360 variables using a Spearman correlation coefficient since covariates were not normally
361 distributed. We assumed that a correlation of greater than r_s 0.4 was problematic, but the
362 correlation was below this threshold ($r_s = 0.14$).
363 Foraging intensity was modelled using GAM in the '*gam'* package in R (Hastie &
364 Tibshirani 1990). Foraging intensity was defined based on the frequency distribution of the
365 tracking locations classified as foraging only. The environmental covariates were divided into
366 K classes. Then, within each class the number of foraging locations was extracted and the
367 count was used as the response variable. A GAM with a quasi-Poisson distribution was then
368 fitted to the data. Separate models were developed for male and female birds. Models
369 included sea ice concentration and bathymetry as fixed factors. The smoothing parameter was
370 chosen automatically using generalized cross-validation. For SIC we used K=1% SIC classes
371 and for bathymetry we used K=50 m classes.

372 Differences between sexes in body measurements, sea ice characteristics used, and
373 foraging trip metrics were tested using Student's t-tests and differences in stable isotope date
374 using Student's t-tests and Wilcoxon rank tests. Since we performed a large number of tests
375 when comparing male and female body size measurements, isotopic values and foraging trip
376 characteristics, we used the Benjamini-Hochberg procedure (Benjamini & Hochberg 1995) to
377 control for false discovery rate. We chose a false discovery rate (q*) of 0.10 when applying
378 the Benjamini-Hochberg procedure. This choice was motivated by the fact that this study was
379 conducted on a single year and was exploratory. In such cases setting a FDR to an extremely
380 low value results in decreasing the statistical power for detecting genuine effects and several
381 authors recommend setting FDR to a relatively large value (Yoccoz 1991, Field et al. 2004,
382 Roback and Askins 2005).
383

384 # 3 Results

385 Male snow petrels were larger than females, particularly for bill length and bill depth, and
386 were 10% heavier than females (Table 1). Bill length, bill depth and body mass were the most
387 sexually dimorphic phenotypic traits.
388

389 ## 3.1 Spatial distribution of males and females and habitat differences

390 Males and females foraged in offshore waters to the east and to the west of the colony in
391 equal proportions (χ^2=0.03, p=0.86, Figure 1). Space-use sharing was similar between males
392 and females as the UDOI was not significantly lower than the null expectation for 25%, 50%,
393 75% or 95% UDs (Table 2). The 95% UDOI was > 1, indicating a higher than normal overlap
394 between male and female UDs relative to uniform space use, i.e. male and female UDs were
395 non-uniformly distributed and had a high degree of overlap. By contrast, the 25% UDOI was
396 relatively close to 0 indicating less overlap between male and female UDs relative to uniform

space use. Males and females UDs were also similar whatever the UDs considered since BA were not significantly lower than the null expectation for 25%, 50%, 75% or 95% UDs (Table 2).

In average males foraged in areas with higher SIC than females (Table 3). Fitted models on foraging probability contained sex-specific smoothers for bathymetry and SIC (Table 4). For females, the GAMM model explained 10% of the deviance of foraging probability. All smoothers for SIC and bathymetry were significant (Table 4). Foraging probability increased sharply with increasing SIC up to 30% and more smoothly for high SIC (Figure 2). Foraging probability showed a first peak at depth of ≈600 m and a second and high peak at depth of ≈1600 m. Foraging probability sharply increased at depths >2500 m but sample size was small and there was high uncertainty. Both the random intercept for bird identity and the spatial smoother were significant.

For males, the model explained 4.6% of the deviance of foraging probability. All smoothers for SIC and bathymetry, the random intercept for bird identity and the spatial smoother were significant (Table 4). Male foraging probability varied non-linearly with SIC and bathymetry. It increased smoothly with increasing SIC, and was higher when SIC was higher than ≈90% (Figure 2). Foraging probability also increased with bathymetry up to ≈600 m and remained relatively stable until ≈2000 m from which it increased.

Female foraging intensity was non-linearly related to SIC and bathymetry (Table 5). Foraging intensity increased with SIC up to a maximum for SIC ≈40% and then decreased for higher SIC (Figure 3). Lowest foraging intensity was observed for SIC >80%. Foraging intensity showed a rather bimodal distribution as a function of bathymetry. It was maximal in waters ≈400 m deep, then decreased to reach a minimum at ≈1400 m, and increased again for water depths between ≈2000-2700 m. Male foraging intensity was non-linearly related to SIC and bathymetry (Table 5). It showed a bimodal distribution as a function of SIC, with a

maximum for SIC ≈36% and a second peak for SIC ≈85% (Figure 3). As for females, male foraging intensity was bimodal as a function of bathymetry. It was maximal in waters ≈400 m deep, then decreased to reach a minimum at ≈1500 m, and increased up to a second peak in waters ≈2400 m deep.

3.2 Stable isotope ratios

Male plasma, blood cells and feathers had significantly 0.6-0.8‰ higher $\delta^{15}N$ values than those of females (Table 6). There was no difference in $\delta^{13}C$ values between males and females, except for plasma for which males had higher values. Males and females had similar SEA_B for all tissues (Figure 4). Overlap between SEA_B areas for males and females was 0.462, 0.586 and 0.599 for blood cells, plasma and feathers, respectively.

Strong significant positive relationships were found in $\delta^{15}N$ between blood cells and plasma (males: $F_{1,25}$=18.846, P<0.001, r=0.656; females: $F_{1,33}$=31.679, P<0.001, r=0.700; Figure 5), but not between feathers and blood cells (males: $F_{1,36}$=0.036, P=0.850, r=0.032; females: $F_{1,45}$=0.062, P=0.805, r=0.037). No significant positive relationship was found in residual $\delta^{13}C$ between blood cells, plasma and feathers (all p's>0.243).

There was no significant relationship between isotopic values and body measurements or body condition (all p's>0.08).

3.3 Foraging trip performance and foraging efficiency

Foraging trip duration, length, speed and directions (Table 7), as well as mass gain and proportion mass gain (Table 8) did not differ between males and females. Foraging efficiency, measured as the proportionate daily mass gain while foraging, was significantly greater for females than for males (Table 8), and was greater for larger females with deeper bills (PC1: $F_{1,20}$=5.279, P=0.033, r=0.457; bill depth: $F_{1,20}$=8.630, P=0.008, r=0.549). In females, but not

in males, foraging efficiency decreased with the duration of the foraging trip (Figure 6). Females were more efficient than males during short (<2 days) foraging trips, but for trips longer than 2 days, foraging efficiency was similar in males and in females (daily mass gain: t_{39}=0.397, P = 0.693; proportion daily mass gain : t_{39}=0.862, P = 0.394).

3.4 Regulation of the foraging trips

To investigate how birds regulate foraging trips according to the depletion of their body reserves, we correlated the body condition at departure with the duration of the foraging trips and the mass gain metrics while foraging. Foraging trip duration was not correlated to body condition at departure (Pearson correlation coefficient: p = 0.417 for females, p = 0.576 for males), but mass gain and proportionate mass gain were negatively related to body condition at departure for both sexes (all P's<0.005; Figure 6). In addition, in males, but not in females, daily mass gain and proportionate daily mass gain were negatively correlated to body condition at departure (males: all P's<0.010; females: all P's>0.429). Male (but not female) body condition at return from a foraging trip was positively correlated to the time spent at sea (Pearson correlation coefficient: p = 0.05).

3.5 Factors affecting mass gain at sea

For females, but not for males, mass gain, proportion mass gain and body condition at return from a foraging trip were positively correlated to mean and maximum sea ice concentration of the foraging trip locations (females: all P<0.049; males: all P>0.232; Figure 7). For males and females, there was no relationship between bathymetry and mass gain, proportion mass gain, and body condition at return (all P>0.100).

4 Discussion

472 This study provides clear evidence of sexual segregation and foraging tactics in snow petrels.
473 In accordance with our prediction, we found evidence for sexual segregation in diet, with
474 males feeding on average on higher trophic level prey when compared to females, but no
475 evidence for spatial segregation as indicated by spatial data and $\delta^{13}C$ isotopic data. Males and
476 females differed in their usage of sea ice, providing evidence for sex-specific habitat
477 segregation.
478

479 **4.1 Differences in habitat use**

480 During incubation males and females foraged predominantly in pack-ice areas over the deep
481 Antarctic continental shelf and adjacent continental margin (500-900 m, due to the isostatic
482 effect of the ice sheet), and to a lesser extent in oceanic waters. These results are consistent
483 with previous observational work at sea showing that high densities of breeding snow petrels
484 in the Ross Sea were found within the pack ice along the continental slope (Ainley et al.
485 1984). The low tissue $\delta^{13}C$ values of snow petrels is a consistent characteristic of consumers
486 foraging in high-Antarctic waters (Cherel 2008, Cherel et al. 2011). Blood cell, plasma and
487 feather $\delta^{13}C$ values were similar in males and females, which indicates that both sexes foraged
488 offshore in pelagic waters without an obvious neritic-oceanic $\delta^{13}C$ gradient in high-Antarctica
489 (Cherel et al. 2011). Blood cell and plasma $\delta^{13}C$ values of birds from Adélie Land were
490 similar to values obtained from snow petrel muscle tissue in the Weddell Sea (Rau et al. 1992)
491 between 64°S and 66°S, but were slightly lower than those measured in whole blood of birds
492 from Hop Island (Hodum & Hobson 2000). However, the shape of the relationships between
493 foraging intensity and SIC suggested that males and females used different sea ice habitats.
494 Female foraging intensity was highest for SIC between ≈20% and ≈40%, and then decreased
495 non-linearly for higher SIC, with a sharp decrease for SIC higher than ≈70%. By contrast
496 male foraging intensity remained high for high SIC. Therefore, although foraging intensity

497 decreased with increasing SIC for both sexes, males foraged more intensively in high sea ice
498 concentration areas (> 70%) than females. Males and females made greater use of pack-ice
499 areas over the continental shelf and continental margin than of oceanic pack-ice areas, but
500 males were more likely to forage and foraged more intensively on the continental margin (-
501 550 to -950 m) than females.

502 Few studies have simultaneously quantified between-sex differences in habitat use and
503 foraging behavior in marine species in relation to dynamic oceanographic features such as sea
504 ice. In the northern gannet (*Morus bassanus*), sexual segregation was driven largely by spatial
505 and habitat segregation with males, smaller than females, mainly foraging in coastal mixed
506 waters where net primary production was high, and females mainly foraging in offshore
507 stratified waters (Cleasby et al. 2015). Similarly, the sex-specific habitat use reported in the
508 monomorphic Barau's petrel (*Pterodroma baraui)* during the prelaying exodus (males used
509 more frequently marine areas with high productivity) can be partly explained by spatial
510 segregation between sexes (Pinet et al. 2012). During the incubation and chick rearing period,
511 they did not found evidence for habitat segregation and foraging areas largely overlapped. In
512 the Adélie penguin (*Pygoscelis adeliae*) at Pointe Géologie, females foraged more intensively
513 in areas of higher sea ice concentration than males during the guard stage, and there was
514 spatial segregation between sexes with females foraging further from the colony than males
515 (Widmann et al. 2015). Using a multiyear comprehensive dataset, Paiva et al. (2017) found
516 that sexual segregation in foraging areas and foraging habitats of Cory's shearwaters
517 (*Calonectris borealis*) varied between years, with greater sexual (habitat and spatial)
518 segregation during years when sea surface temperatures were higher and chlorophyll *a*
519 concentrations were lower, presumably corresponding to lower food availability. In favorable
520 years no spatial segregation was observed and habitat segregation was low. The hatching
521 success of snow petrels during the 2015/2016 breeding season was 46.9%, i.e. lower than the

522 long-term average of 63.3% (Chastel et al. 1993), suggesting that environmental conditions
523 were relatively poor. However, we did not observed spatial segregation between sexes but
524 foraging habitat use differed, with males foraging more frequently in high sea ice
525 concentration areas than females. Such a pattern was found in the wandering albatross
526 (*Diomedea exulans*) at South Georgia in which, despite no clear sexual segregation at large
527 scales, sex-specific microhabitat selection was found during the chick-rearing period,
528 resulting in sexual segregation in core foraging areas (Pereira et al. 2018). Multiple years of
529 tracking are needed to shed light into the effects of environmental stochasticity (sea ice
530 variability) on habitat segregation and spatial segregation.

531 As opposed to other highly sexually size-dimorphic seabirds (wandering albatross:
532 Weimerskirch et al. 1993, giant petrels *Macronectes spp.*: Gonzáles-Solís et al. 2000, boobies
533 *Sula spp.*: Weimerskirch et al. 2009, frigatebirds *Fregata spp.*: Hennicke et al. 2015) snow
534 petrels did not show spatial segregation in their foraging habitat during incubation. Spatial
535 segregation in snow petrels may occur during other periods of the year such as during the
536 chick-rearing period when which food requirements are particularly high for provisioning the
537 chick. Alternatively, this lack of spatial segregation may be constrained by the specific
538 foraging habitat requirements of snow petrels. These seabirds forage exclusively in a sea ice
539 environment, which is limited during the breeding season around breeding colonies and may
540 thus constraint males and females to spatially overlap at a broad spatial scale.

541

542 **4.2 Influence of sex on diet and foraging tactics**

543 The snow petrel diet is relatively well known during the chick-rearing period and isotopic
544 data together with prey biometric data suggest that snow petrels mainly feed on postlarvae
545 and juvenile Antarctic silverfish (*Pleuragramma antarcticum*) (Ridoux & Offredo 1989,
546 Hodum & Hobson 2000, Pinkerton et al. 2013). Although, snow petrel diet during incubation

547 remains poorly known, $\delta^{15}N$ values obtained in our study are similar or slightly higher than
548 those found in other studies during the chick rearing period (Hodum & Hobson 2000, Delord
549 et al. 2016), suggesting a similar diet. Nevertheless, and despite large overlap in their core
550 isotopic niches as indicated by the standard ellipse areas, female snow petrels had lower $\delta^{15}N$
551 values than males for all tissues sampled, which suggests they were feeding on lower trophic
552 level prey than males. Similar results were found by Tartu et al. (2014) for blood cells during
553 the pre-laying period. We speculate that there might be at least two reasons for this. First,
554 compared to males, females may feed more frequently on other prey than Antarctic silverfish,
555 such as crustaceans which are situated at a lower trophic level than Antarctic silverfish.
556 Indeed, diet studies indicate that snow petrels also feed on crustaceans such as *Euphausia*
557 *superba*, *E. crystallorophias*, *Themisto gaudichaudii*, and other amphipods (Ainley et al.
558 1984, Ridoux & Offredo 1989) which have lower $\delta^{15}N$ values than Antarctic silverfish
559 (Pinkerton et al. 2013). Second, females may feed on Antarctic silverfish in similar
560 proportions than males but on smaller sized individuals (i.e. younger). It is known that $\delta^{15}N$
561 values increase with body length (and age) in Antarctic silverfish from ≈7-8‰ in larvae (10-
562 20 mm standard length) to ≈10-11‰ in juvenile and adult fish (Giraldo et al. 2011, Pinkerton
563 et al. 2013). It is currently unknown whether sea ice concentration and characteristics
564 differentially affect the spatial distribution of Antarctic silverfish age-classes. However, it is
565 likely that females fed more on crustaceans than on young silverfish since crustaceans have
566 much lower $\delta^{15}N$ values than young silverfish (Cherel 2008). Thus, our results suggest that
567 males ate more silverfish in areas with higher sea ice concentration.

568 The strong positive relationship between plasma $\delta^{15}N$ and blood $\delta^{15}N$ indicates short term
569 (over weeks) consistency in trophic level between successive foraging trips during incubation.
570 Values of $\delta^{15}N$ in plasma and feathers did not differ in both sexes (Appendix 1), but blood
571 $\delta^{15}N$ were smaller than feather and plasma $\delta^{15}N$ in both sexes, suggesting that males and

572 females fed on lower trophic level prey prior to incubation than during the breeding season.
573 Short and long term consistency in foraging water masses was also low as indicated by the
574 lack of relationship between plasma and blood $\delta^{13}C$, and between feather and blood $\delta^{13}C$,
575 respectively. Indeed tracking data indicated that birds foraged on the continental shelf,
576 continental margin, and to a lesser extent in oceanic waters. Values of $\delta^{13}C$ in feathers were
577 higher than those in blood and plasma for both sexes (Appendix 1), suggesting that during the
578 latter part of the breeding season and the beginning of the non-breeding season snow petrels
579 foraged in more oceanic waters (snow petrels start molting during the chick rearing period
580 and until early May (Beck 1969, 1970, Delord et al. 2016). This period coincides with the sea
581 ice growth and its northward extension.

582 The negative relationship between mass gain (and proportion daily mass gain) during a
583 foraging trip and body condition at departure for a foraging trip (i.e. at the end of fasting
584 while incubating the egg), indicated that males and females were able to regulate their body
585 reserves as found in other Procellariiform species (Chaurand & Weimerskirch 1994,
586 Gonzáles-Solís et al. 2000). Although both sexes regulated body condition, this ability seemed
587 greater for females than for males. Indeed, body condition at departure for a foraging trip was
588 lower in females than in males, but similar for both sexes at return from a foraging trip despite
589 similar trip durations. This is further supported by the fact that females had higher daily mass
590 gains and proportion daily mass gains than males. However, this greater ability in females
591 may be partly explained by the fact that females undertook short foraging trips during which
592 mass gain was particularly high (Figure 1). Although some males also made short foraging
593 trips, mass gain was still lower than female mass gain during these trips. Therefore, these
594 results suggest that female foraging efficiency was similar in males and females, except
595 during short (<2 days) foraging trips during which females appeared more efficient. We
596 suspect that some females undertook short foraging trips during their incubation shift in order

597 to restore their body condition to avoid abandoning the egg while their partner was foraging at
598 sea. This could result from the lower fasting capacities of females compared to males due to
599 their smaller body size (Barbraud & Chastel 1999).

600 Interestingly, the ability of females (but not of males) to restore their body condition
601 during a foraging trip was affected by sea ice concentration. Indeed, female body condition at
602 return from a foraging trip was positively related to sea ice concentration in the foraging area,
603 contrary to males. This suggests that areas with heavy sea ice concentration were more
604 profitable. This is further supported by the positive relationship between male (but not
605 female) body condition at return from a foraging trip and time spent at sea, and given that
606 males foraged more frequently in high sea ice concentration areas. Thus, foraging on highly
607 nutritional preys such as silverfish in high sea ice concentration areas might be more efficient
608 to restore body condition that feeding in more open water areas.

609 Body condition at the start of a foraging trip was not related to the time spent at sea,
610 suggesting that the time spent at sea was not only dependent on the restoration of body
611 condition. Although only a few birds returned to undertake the next incubation shift after
612 losing mass (n = 3, 6.3%) or without gaining mass (n = 3, 6.3%), this suggests that mass gain
613 alone does not explain the decision to return to the colony. Perhaps birds took into account the
614 increased probability of partners deserting the egg with the increasing duration of the foraging
615 trip (Tveraa et al. 1997).

616 Thus, incubating female snow petrels seemed more efficient at restoring their body
617 condition during a foraging trip despite similar trip duration, length or speed, while foraging
618 areas were identical to those of males at a broad spatial scale. However, this higher efficiency
619 mainly concerned short (<2 days) foraging trips. In addition, our results show that females
620 foraging in high sea ice concentration areas foraged more efficiently (this relationship holds
621 when excluding foraging trip <2 days), and female fed on lower trophic level preys that

622 males. Together, these results suggest that areas with high sea ice concentration may be more
623 profitable for resource acquisition, perhaps due to higher abundance, availability or quality of
624 prey such as the Antarctic silverfish.
625
626 **4.3 Factors underlying sexual segregation**
627 Sex differences in foraging behavior could result from the influence of sexual size
628 dimorphism on foraging efficiency and intra-specific competition (forage-selection hypothesis
629 and scrambled competition hypothesis). The positive relationship between female bill depth
630 and proportion daily mass gain suggests that foraging efficiency is size dependent in females,
631 which are smaller than males. Our results also suggest that the most favorable areas were
632 areas of high sea ice concentration (females body condition at return increased with increase
633 sea ice concentration, male body condition at return increased with foraging trip length),
634 which were used less frequently by females. Therefore, it is possible that females were
635 excluded from high sea ice concentration areas via direct competition. This could possibly
636 indicate that male and female snow petrels try to avoid competition and thus diverged in
637 habitat preference in more profitable areas, where intra-specific competition might be more
638 intense. Such a mechanism was also proposed to explain sex-specific differences in broad
639 scale foraging areas in highly sexually size dimorphic species (wandering albatross:
640 Weimerskirch et al. 1993, Shaffer et al. 2001; giant petrels: Gonzáles-Solís et al. 2000), but
641 also in foraging habitat at a microhabitat scale (Pereira et al. 2018). A major assumption of
642 the intersexual competition hypothesis is that prey capture should be a function of bill size
643 (Selander 1966, Shine 1989). Although we do not have the data in hand to test this prediction
644 explicitly, we note that $\delta^{15}N$ values suggested that females consumed lower sized prey than
645 males (crustaceans vs fish). Females with thicker bills were also more efficient during their

646 foraging trip, suggesting they were feeding on more profitable prey, and bill size was among
647 the most sexually dimorphic phenotypic trait in this species.
648 Sex-specific niche divergence and habitat segregation can also arise from a difference
649 between sexes in parental roles and investment (the activity budget hypothesis, Clarke et al.
650 1998, Thaxter et al. 2009, Weimerskirch et al. 2009, Pinet et al. 2012). Although males
651 undertake a greater investment in chick provisioning though higher feeding frequencies
652 (Barbraud et al. 1999), there is little differentiation in the reproductive role of male and
653 female snow petrels during incubation. Males make slightly shorter foraging trips than
654 females during incubation (Isenmann 1970, Barbraud et al. 1999), but in average the total
655 time spent foraging during the incubation period is very similar for both sexes (males: average
656 19.8 days, females: average 21.0 days, Barbraud 1999), indicating that the roles of male and
657 female snow petrels do not appear to differ substantially during incubation. Therefore, it
658 seems unlikely that such limited constraints related to reproductive role specialization could
659 explain why female snow petrels foraged less intensively in high sea ice concentration areas;
660 this hypothesis can probably, therefore, be discounted. Sex-specificity in flight performance
661 may also be responsible for sexual segregation (Shaffer et al. 2001, Phillips et al. 2004).
662 Indeed, sexual dimorphism in wing area and wing loading in several albatross species may
663 partially explain large-scale sexual segregation in foraging areas in these species: sex-specific
664 foraging locations were likely influenced by activity budgets since smaller birds are more
665 efficient flyers. Therefore, other aspects of the morphology not measured here, such as wing
666 loading and agility, may be important. Female snow petrels appear to have a lower aspect
667 ratio and lower wing loading than males (Spear & Ainley 1998), suggesting they might be
668 less flight efficient but more maneuverable than males. However, since there was no spatial
669 segregation between sexes at large-spatial scales, environmental conditions potentially
670 affecting flight efficiency (wind speed) were identical for males and females. Thus, these

aspects are also unlikely to be of importance in snow petrels to explain sex-specific foraging habitat use during incubation.

Overall, our study demonstrates sex-specific foraging tactics in a highly sexually size dimorphic species during the incubation period, probably driven by intra-specific competition. Results indicate an absence of sexual segregation at a broad-spatial scale, but suggest that sexual segregation in snow petrels is mediated by habitat segregation at a microhabitat scale. Males foraged more intensively than females in high sea ice concentration areas, which seemed to be more profitable in terms of resource acquisition as results suggest that males ate more fish in these areas. Studying sex-specific foraging tactics during the entire breeding period, thus including the pre-laying exodus and the chick-rearing period, is however necessary to better understand the underlying drivers of sexual segregation in snow petrels and in marine predators in general (Pinet et al. 2012). Sexual segregation in foraging behavior may also vary between years as a function of environmental conditions (Cleasby et al. 2015, Paiva et al. 2017), highlighting the need for multi-year tracking studies.

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Table 1. Body measurements of male and female snow petrels and percentage of difference between sexes for each measurement. For t-tests homogeneity of variances we checked using a Brown and Forsythe tests (Brown and Forsythe 1974), and corrected p values are reported (uncorrected in brackets). Significant differences with a false detection rate of 0.10 are shown in bold. Δ is the difference in %. The sample size is 47 individuals.

	Sex	Mean (SD)	Range	Δ	t-test
Wing length (mm)	Male	298.3 (6.8)	287.0-311.0	2.3	**t_{45}=4.215**
	Female	290.0 (6.7)	280.0-302.0		**p<0.001** (<0.001)
Tarsus length (mm)	Male	40.2 (1.2)	38.0-42.7	4.5	**t_{45}=4.846**
	Female	38.4 (1.2)	35.9-41.0		**p<0.001** (<0.001)
Bill length (mm)	Male	24.4 (1.0)	22.1-26.6	9.0	**t_{45}=8.488**
	Female	22.2 (0.8)	20.9-23.7		**p<0.001** (<0.001)
Bill depth (mm)	Male	10.8 (0.5)	9.6-11.8	8.3	**t_{45}=6.408**
	Female	9.9 (0.4)	8.9-10.9		**p<0.001** (<0.001)
Body mass[1] (g)	Male	389.0 (30.4)	331.5-464.0	10.3	**t_{45}=6.014**
	Female	340.5 (24.0)	300.0-385.0		**p<0.001** (<0.001)
Body mass[2] (g)	Male	431.4 (34.0)	365.0-495.0	10.3	**t_{45}=4.467**
	Female	387.0 (33.9)	320.0-460.0		**p<0.001** (<0.001)

[1]before a foraging trip; [2]on return from a foraging trip.

Table 2. Estimated overlap in utilization distributions (UD) between male and female snow petrels from Ile des Pétrels, Adélie Land, East Antarctica. UDOI: Utilization distribution overlap index. BA: Bhattacharyya's affinity

UD (%)	Observed UDOI	Randomized UDOI	p	Observed BA	Randomized BA	p
25	0.062	0.064	0.417	0.127	0.123	0.452
50	0.226	0.243	0.262	0.379	0.398	0.273
75	0.502	0.545	0.227	0.634	0.662	0.212
95	1.215	1.229	0.439	0.858	0.863	0.359

Table 3. Mean, maximum and variance in sea ice concentration, mean, minimum and maximum bathymetry for foraging localities of male and female snow petrels from Ile des Pétrels, Adélie Land, East Antarctica. For t-tests homogeneity of variances we checked using a Brown and Forsythe tests (Brown and Forsythe 1974), corrected p values are reported (uncorrected in brackets). Significant differences with a false detection rate of 0.10 are shown in bold.

	Sex	Mean (SD)	Range	t-test
Mean SIC (%)	Male	54.8 (14.6)	30.4-77.9	**t_{44}=2.600**
	Female	44.0 (13.4)	16.8-72.5	**p=0.063** (0.013)
Maximum SIC (%)	Male	83.8 (16.8)	43.9-99.9	t_{44}=1.867
	Female	72.6 (23.8)	19.8-100.0	p=0.171 (0.069)
Variance in SIC	Male	375.7 (224.2)	69.5-840.2	t_{44}=1.772
	Female	264.0 (198.3)	3.9-749.9	p=0.139 (0.083)
Mean bathymetry (m)	Male	503.8 (248.7)	285-1490	t_{45}=0.911
	Female	582.4 (341.2)	259-1706	p=0.367 (0.367)
Maximum bathymetry (m)	Male	1354.1 (632.1)	809-2973	t_{45}=1.285
	Female	1617.4 (771.4)	613-3223	p=0.257 (0.205)

Table 4. Generalized Additive Mixed Model (GAMM) results for foraging probability of male and female snow petrels as a function of sea ice concentration (SIC), bathymetry (BAT) and spatial autocorrelation (s(x,y)). edf indicates the estimated degrees of freedom.

Variable	Sex	Smoother edf (p value)	Estimate (SE)	σ^2 (SE)
Intercept	Male		3.543 (1.101)	
	Female		3.087 (1.365)	
SIC	Male	**7.97 (<0.001)**		
	Female	**1.00 (0.003)**		
BAT	Male	**3.89 (<0.001)**		
	Female	**1.00 (0.044)**		
s(x,y)	Male	**22.96 (<0.001)**		
	Female	**24.52 (<0.001)**		
Random intercept for bird ID	Male			10.250 (3.201)
	Female			1.174 (0.343)

Table 5. Generalized Additive Model (GAM) results for foraging intensity of male and female snow petrels as a function of sea ice concentration (SIC) and bathymetry (BAT). edf indicates the estimated degrees of freedom.

Variable	Sex	Smoother edf (p value)	Scale	Adjusted R^2
SIC	Male	**4.99 (<0.001)**	2.50	0.223
	Female	**4.11 (<0.001)**	2.25	0.663
BAT	Male	**6.47 (<0.001)**	16.01	0.779
	Female	**6.82 (<0.001)**	9.28	0.863

Table 6. Stable isotope values in blood cells, plasma and feathers of male and female snow petrels sampled Ile des Pétrels, Adélie Land, East Antarctica. Values are mean ± SD. SEA_B are Bayesian approximation of the standard ellipse area. Values in brackets indicate n and range for $\delta^{13}C$ and $\delta^{15}N$, and 95% credible interval for SEA_B. p indicates the probability that SEA_B of males and females differ. Sample sizes are 47 individuals for blood cells and feathers, and 46 individuals for plasma. For t-tests homogeneity of variances we checked using a Brown and Forsythe tests (Brown and Forsythe 1974), corrected p values are reported (uncorrected in brackets). Significant differences with a false detection rate of 0.10 are shown in bold.

Tissue	$\delta^{13}C$(‰)			$\delta^{15}N$(‰)			SEA_B		p
	Male	Female		Male	Female		Male	Female	
Plasma	-25.46 ± 0.35 (-26.01;-24.56)	-25.66 ± 0.22 (-26.06;-25.26)	**t_{83}=2.105 p=0.057** (0.039)	12.11 ± 0.74 (10.70;13.38)	11.49 ± 0.97 (9.04;12.83)	**t_{83}=3.418 p=0.002** (0.001)	0.69 (0.45;0.99)	0.79 (0.55;1.09)	0.311
Blood cells	-25.79 ± 0.21 (-26.13;-25.35)	-25.72 ± 0.27 (-26.13;-25.19)	t_{83}=1.491 p=0.168 (0.140)	10.65 ± 0.68 (9.37;12.17)	9.96 ± 0.65 (8.36;11.46)	**t_{83}=5.568 p<0.001** (<0.001)	0.41 (0.29;0.55)	0.39 (0.29;0.52)	0.586
Feather	-23.68 ± 0.71 (-25.05;-22.06)	-23.50 ± 0.67 (-25.00;-22.49)	t_{83}=0.335 p=0.738 (0.738)	12.11 ± 1.35 (8.75;14.03)	11.34 ± 1.35 (8.61;14.10)	**t_{83}=4.289 p<0.001** (<0.001)	2.41 (1.72;3.28)	2.65 (1.94;3.49)	0.331

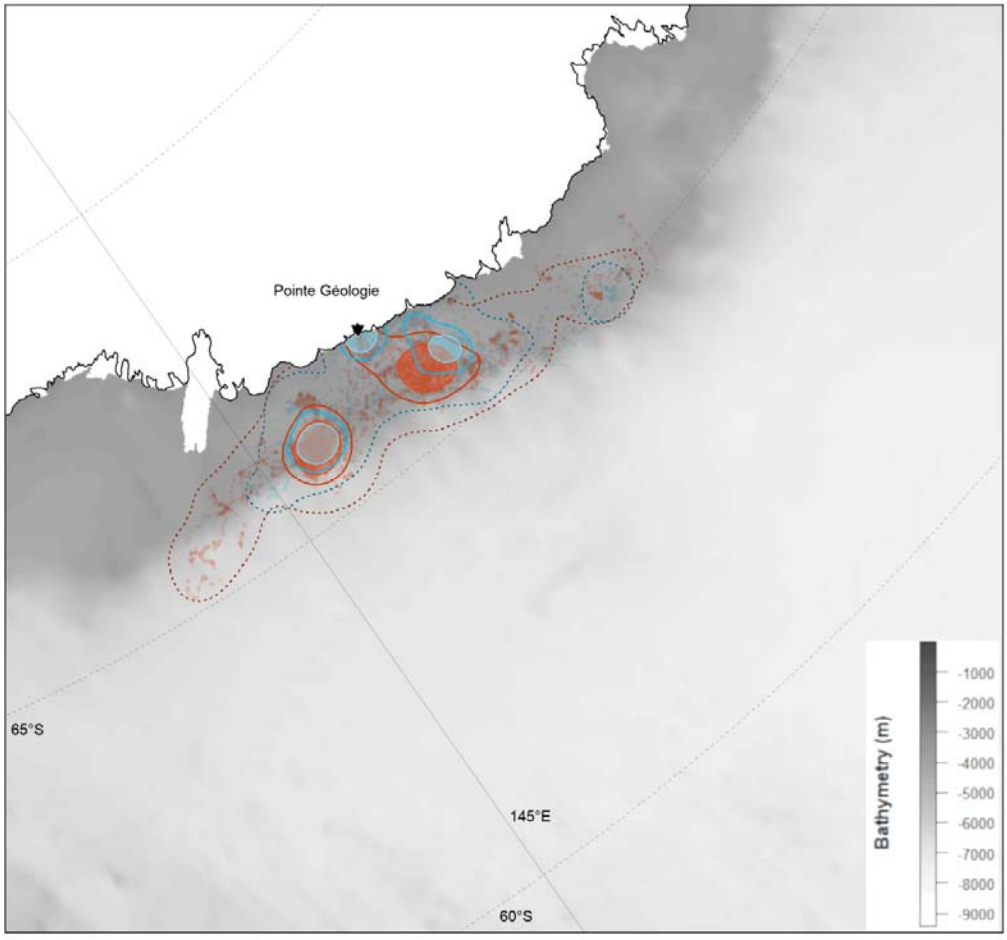
Table 7. Summary of foraging trip metrics for snow petrels from Ile des Pétrels, Adélie Land, East Antarctica. For t-tests homogeneity of variances we checked using a Brown and Forsythe tests (Brown and Forsythe 1974), corrected p values are reported (uncorrected in brackets). Significant differences with a false detection rate of 0.10 are shown in bold.

	Sex	Mean (SD)	Range	t-test
Trip duration (d)	Male	5.2 (1.8)	1.1-8.0	t_{45}=0.979
	Female	4.8 (1.9)	0.7-8.0	p=1.160 (0.333)
Trip length (km)	Male	851.6 (352.1)	302.0-1957.5	t_{45}=0.292
	Female	856.4 (389.0)	173.8-1803.8	p=0.772 (0.772)
Trip mean speed ($km \cdot h^{-1}$)	Male	2.0 (0.6)	1.2-3.5	t_{45}=0.762
	Female	2.1 (0.4)	1.7-3.2	p=0.788 (0.450)
Trip maximum speed ($km \cdot h^{-1}$)	Male	13.5 (2.6)	9.2-21.4	t_{45}=0.670
	Female	13.3 (2.5)	9.5-18.9	p=0.709 (0.506)
Trip start direction (°)	Male	161.5 (114.0)	0.8-355.2	t_{45}=1.066
	Female	204.6 (143.2)	2.6-348.6	p=2.045 (0.292)
Trip mean direction (°)	Male	190.9 (110.4)	70.1-307.9	t_{45}=0.336
	Female	209.2 (117.6)	68.8-342.7	p=0.862 (0.739)
Trip end direction (°)	Male	167.5 (114.0)	46.9-282.8	t_{45}=0.773
	Female	184.8 (68.6)	102.9-291.6	p=1.035 (0.444)

Table 8. Summary of metrics of foraging trip efficiency for snow petrels from Ile des Pétrels, Adélie Land, East Antarctica. For t-tests homogeneity of variances we checked using a Brown and Forsythe tests (Brown and Forsythe 1974), corrected p values are reported (uncorrected in brackets). Significant differences with a false detection rate of 0.10 are shown in bold.

	Sex	Mean (SD)	Range	t-test
Mass at departure (g)	Male	389.0 (30.4)	331.5-464.0	**t_{45}=6.014**
	Female	340.5 (24.0)	300.0-385.0	**p<0.001** (<0.001)
Mass at return (g)	Male	431.4 (34.0)	365.0-495.0	**t_{45}=4.467**
	Female	387.0 (33.9)	320.0-460.0	**p<0.001** (<0.001)
BCI[1] at departure	Male	377.1 (25.2)	336.2-437.7	**t_{45}=2.633**
	Female	358.1 (24.0)	321.3-414.1	**p=0.031** (0.012)
BCI at return	Male	419.5 (31.2)	372.5-479.3	t_{45}=1.636
	Female	405.0 (29.3)	361.0-458.4	p=0.145 (0.109)
Δmass (g)	Male	42.4 (38.1)	-20-110	t_{45}=0.392
	Female	46.6 (34.6)	-10-95	p=0.697 (0.697)
Daily mass gain ($g \cdot day^{-1}$)	Male	9.2 (8.2)	-4.2-28.0	t_{45}=1.707
	Female	14.6 (13.2)	-2.0-45.0	p=0.152 (0.095)
Proportion mass gain (%)	Male	0.11 (0.10)	-0.05-0.29	t_{45}=0.886
	Female	0.14 (0.11)	-0.03-0.31	p=0.435 (0.381)
Proportion daily mass gain (%)	Male	0.02 (0.02)	-0.01-0.07	**t_{45}=2.064**
	Female	0.04 (0.04)	-0.01-0.12	**p=0.089** (0.045)

[1] Body Condition Index



1133
1134 Figure 1. Foraging ranges of male (blue) and female (red) snow petrels during incubation
1135 sampled at Ile des Pétrels, Adélie Land, East Antarctica. Dots show raw location data. Kernel
1136 density based utilization distributions at 95% (dotted lines), 50% (solid lines) and 25% (filled
1137 areas). Bathymetry shown in grey and land in white. Ile des Pétrels is shown as a black
1138 triangle.
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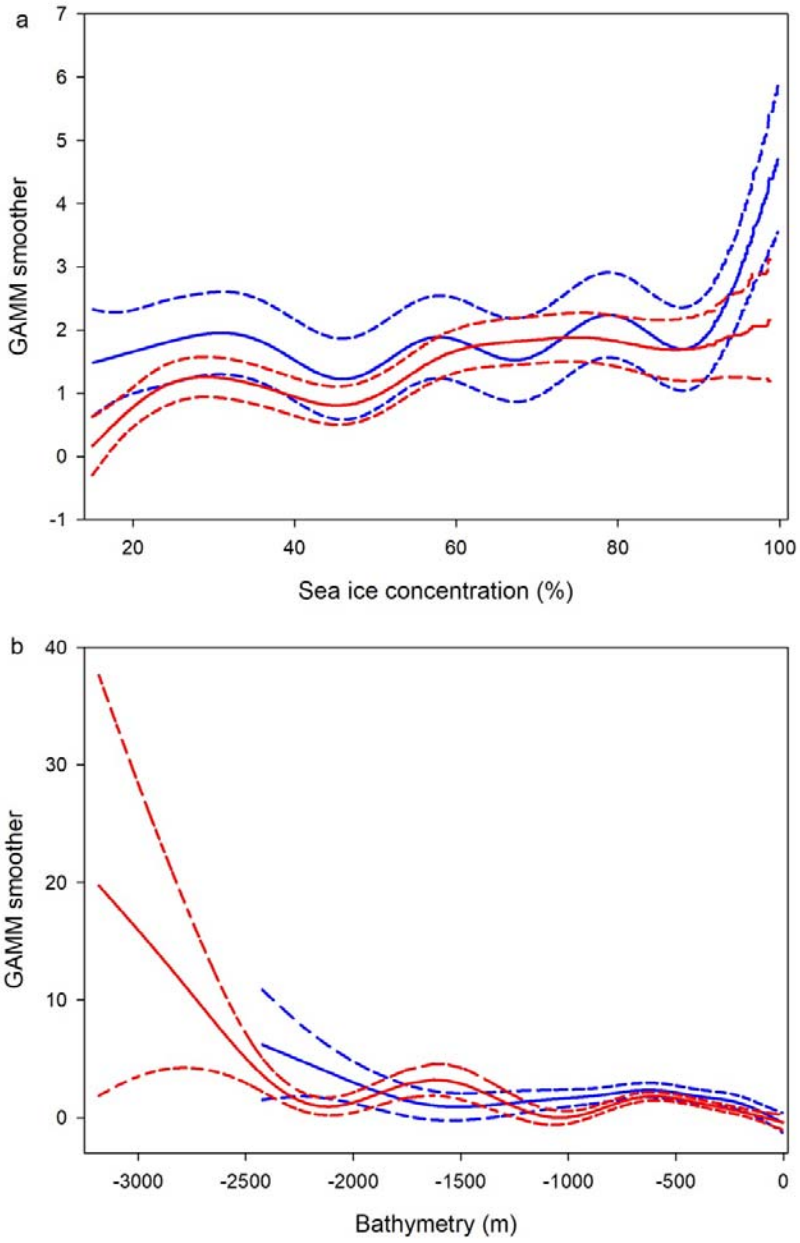

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1144 Figure 2. Foraging probability habitat selection functions for (a) sea ice concentration and (b)

1145 bathymetry). Plots show the predicted curve from the model (solid line) and 95% confidence

1146 intervals (dashed lines) for male (blue) and female (red) snow petrels sampled at Ile des

1147 Pétrels, Adélie Land, East Antarctica. GAMM: generalized additive mixed model.

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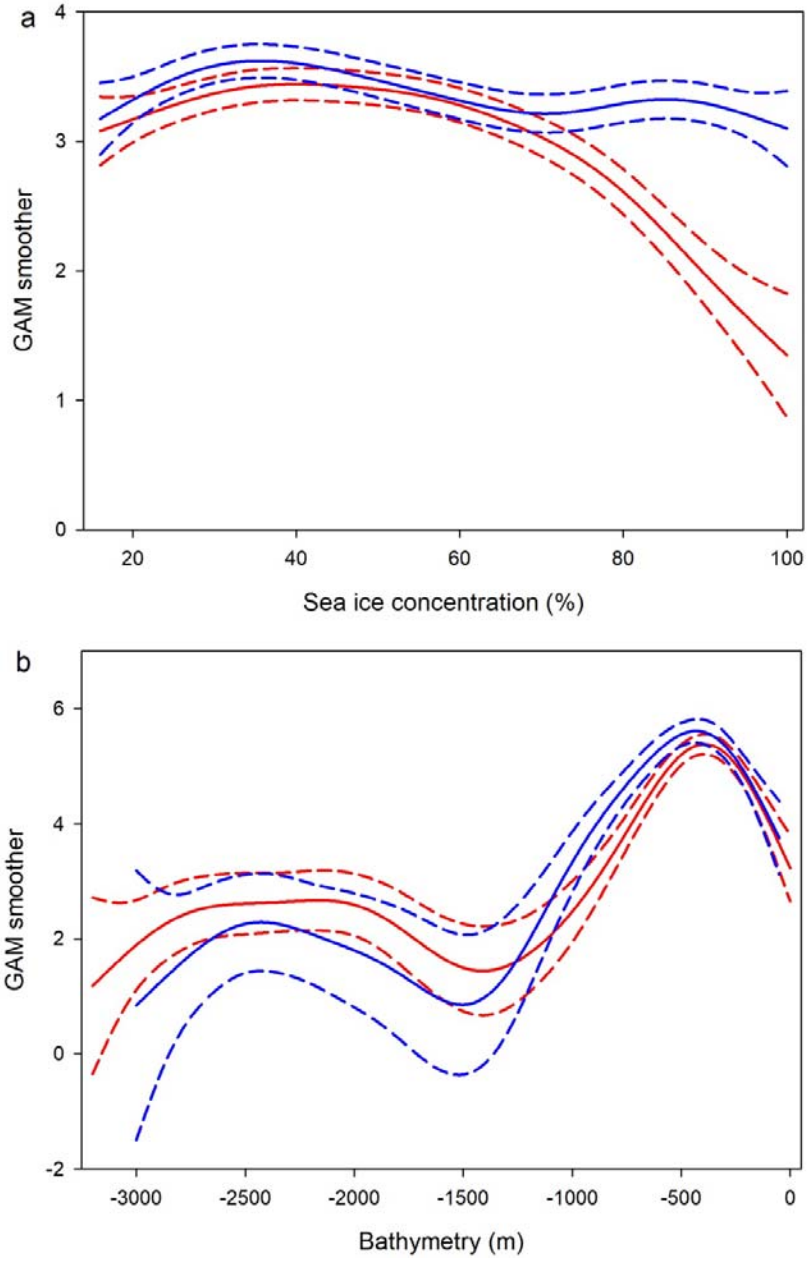
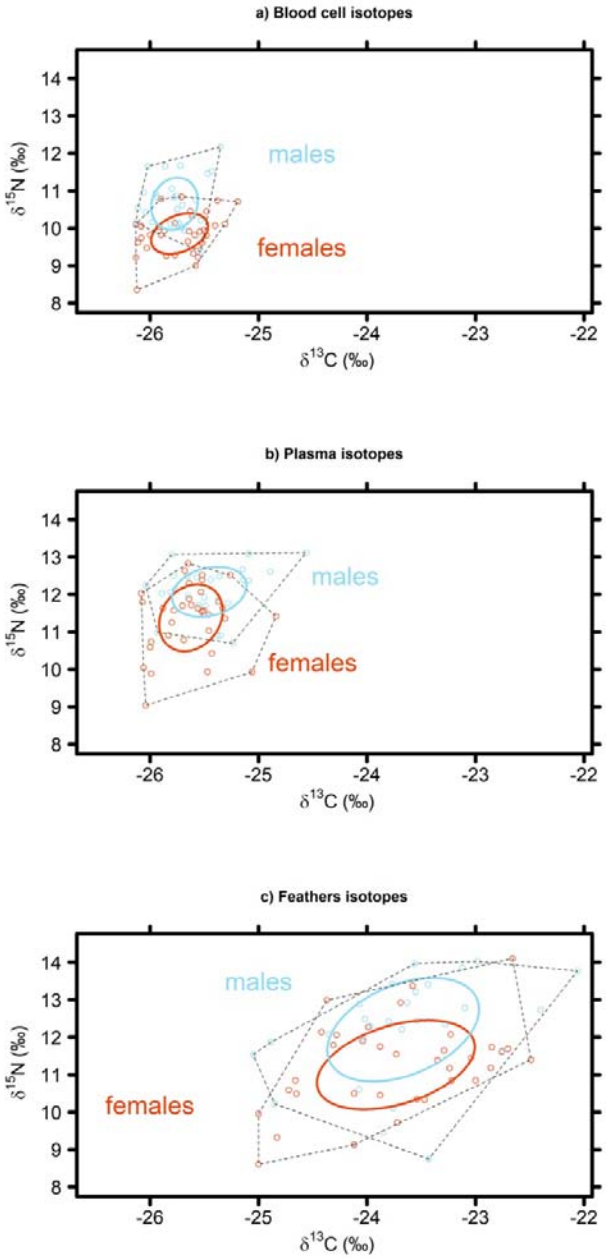


Figure 3. Foraging intensity habitat selection functions for (a) sea ice concentration and (b) bathymetry). Plots show the predicted curve from the model (solid line) and 95% confidence intervals (dashed lines) for male (blue) and female (red) snow petrels sampled at Ile des Pétrels, Adélie Land, East Antarctica. GAM: generalized additive model.



1157

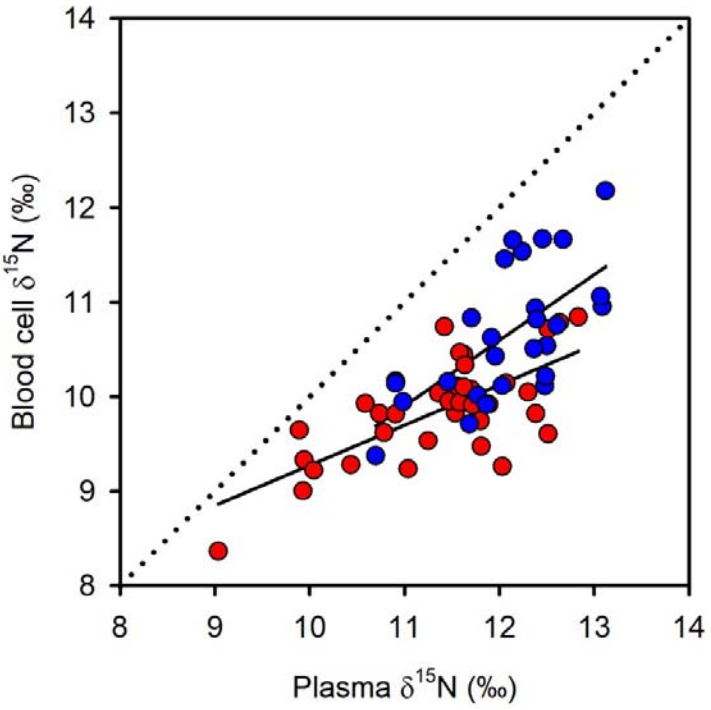
1158 Figure 4. Isotopic niche area based on stable isotope values ($\delta^{15}N$ and $\delta^{13}C$) in blood cells

1159 (top), plasma (middle) and body feathers (bottom) of male (blue) and female (red) snow

1160 petrels breeding at Ile des Pétrels, Pointe Géologie, Antarctica during the incubation period.

1161 The areas of the standard ellipses are represented by the solid lines, and the layman metric of

1162 convex hull area by black dotted lines.



1163

1164 Figure 5. Relationships between blood cells and plasma $\delta^{15}N$ values for male (n = 27, blue)

1165 and female (n = 35, red) snow petrels sampled at Ile des Pétrels, Adélie Land, East Antarctica.

1166 Males: $F_{1,22}=15.203$, $P<0.001$, $R^2 = 0.409$; females: $F_{1,20}=24.300$, $P<0.001$, $R^2 = 0.549$.

1167

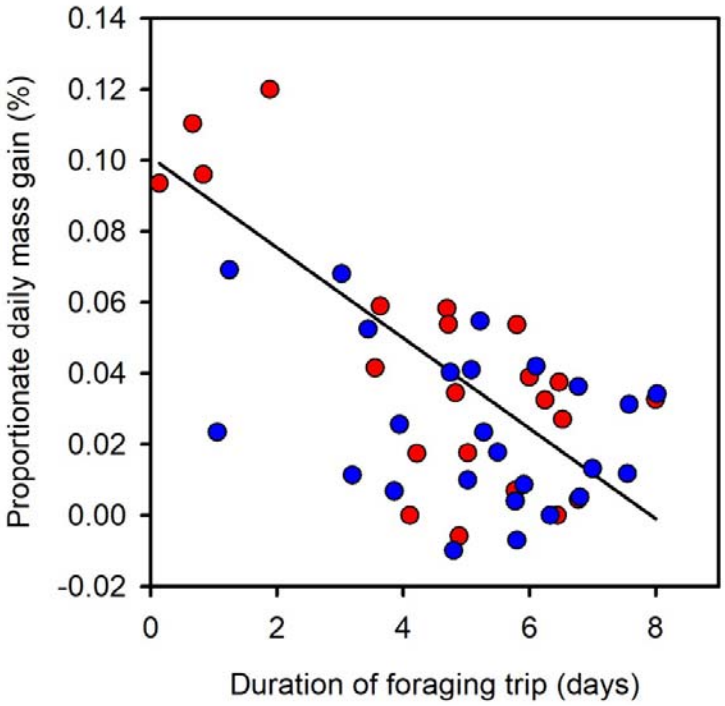
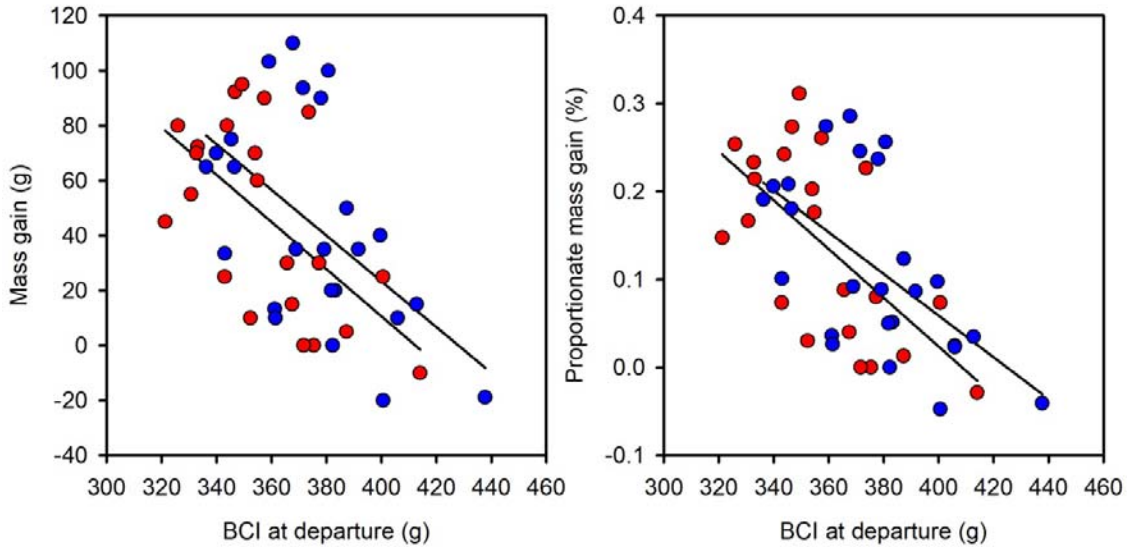


Figure 6. Foraging efficiency (proportionate daily mass gain while foraging) as a function of the total duration of the foraging trip for male (blue) and female (red and solid line) snow petrels sampled at Ile des Pétrels, Adélie Land, East Antarctica. For females: $F_{1,20}$=25.349, P<0.001, R^2 = 0.559.



1174

1175 Figure 7. Mass gain and proportionate mass gain as a function of body condition before a

1176 foraging trip for male (blue) and female (red) snow petrels sampled at Ile des Pétrels, Adélie

1177 Land, East Antarctica. Male mass gain: $F_{1,23}$=10.010, P=0.004, r^2=0.303; female mass gain:

1178 $F_{1,20}$=11.071, P=0.003, r^2=0.356; male proportionate mass gain: $F_{1,23}$=12.361, P=0.002,

1179 r^2=0.350; female proportionate mass gain: $F_{1,20}$=13.258, P=0.002, r^2=0.399.

1180

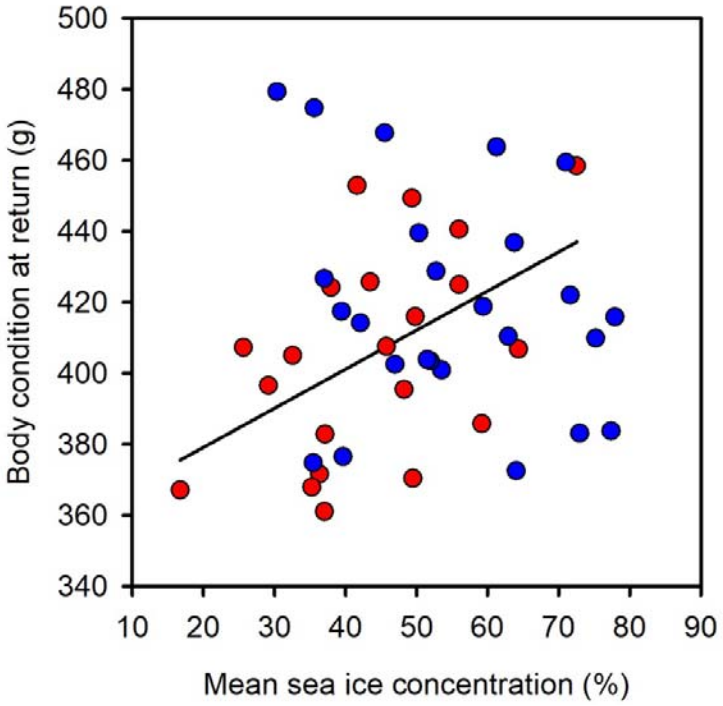


Figure 8. Body condition at return from a foraging trip as a function of the mean sea ice concentration of the foraging trip locations for male (blue) and female (red and solid line) snow petrels sampled at Ile des Pétrels, Adélie Land, East Antarctica. For females: $F_{1,19}$=6.106, P=0.023, R^2 = 0.243.

Appendix I. Testing for differences in $\delta^{15}N$ (‰) and $\delta^{13}C$ (‰) values between tissues for male and female snow petrels sampled Ile des Pétrels, Adélie Land, East Antarctica. t indicates Student's t-tests with df, Z indicates Wilcoxon rank test. ** indicates $P < 0.01$, *** indicates $P < 0.001$ after applying the Benjamini-Hochberg procedure with a false discovery rate of 0.10. Values above diagonal are for $\delta^{15}N$ (‰), values below diagonal are for $\delta^{13}C$(‰). $\delta^{15}N$ (‰) and $\delta^{13}C$ (‰) values in feathers were corrected following Cherel et al. (2014a) before comparison with blood cells. $\delta^{13}C$ (‰) values for plasma were normalized following Post et al. (2007) and Cherel et al. (2014b).

	Plasma	Blood	Feather
Male			
Plasma	-	t_{47}=7.206***	t_{47}=0.008
Blood cells	Z=2.743**	-	t_{47}=3.060**
Feather	t_{47}=17.166***	t_{47}=29.033***	-
Female			
Plasma	-	t_{42}=0.039	t_{42}=0.435
Blood cells	t_{42}=0.039	-	t_{42}=2.967**
Feather	t_{42}=23.219***	Z=4.107***	-