

35 night and close to the bottom during daylight hours. Also, the seal doubled the prey capture
36 attempts in December compared to November. This study reveals the importance of ocean physical
37 properties on seal's diving and foraging behavior, and how this changes, although small, can impact
38 on seals diet and body composition during their post-breeding trips.

39

40 **Keywords :** Southwestern Atlantic Ocean, Patagonian Shelf Slope, Elephant Seals, *Mirounga*
41 *leonina*, Malvinas Current.

42

43 **Acknowledgments**

44 This work was funded through a collaboration of the National Observation Service Mammal as
45 Ocean observer (SNO-MEMO), CNES-TOSCA and WCS. This work was supported by European
46 Organization for the Exploitation of Meteorological Satellites (EUMETSAT)/Centre National de la
47 Recherche Spatial (CNES) through project DSP/OT/07-2118 and by Agencia Nacional de
48 Promoción Científica y Tecnológica (Argentina), project PICT 2018-02433.

49

50

51 **1. Introduction**

52

53 The Argentine Continental Shelf is located in the Southwestern Atlantic (Figure 1). The shelf is
54 broad and relatively shallow, with almost 850 km in its wider zone, a surface area of about 1.2
55 million km², an extension of 2400 km from north to south, and a smooth slope with a variable
56 maximum depth ranging between 70 and 200 m [Parker et al., 1996; Violante and Cavallotto, 2011;
57 Violante et al., 2014]. East of the 200 m isobath, the shelf-break is characterized by a steep slope
58 that goes from 200 to 4000 m depth. In the upper portion of the slope, biological activity is
59 enhanced due to large chlorophyll-a (chl-a) blooms during the austral spring and summer [Saraceno
60 et al., 2005; Romero et al., 2006; Garcia et al., 2008, 2011]. The Malvinas Current (MC), which
61 detaches from the Antarctic Circumpolar Current's northern front, flows northward along the shelf-
62 break. The MC carries cold and rich nutrient waters that may play a role in the large chl-a
63 concentrations through interaction with topography and consequent upwelling [Matano and Palma,
64 2008]. The collision of the MC with the Brazil Current at ~38°S, makes the confluence region one of

65 the most energetic of the world, with eddies and meanders reaching 50°S eastward of the slope
66 [Piola et al., 1987; Chelton et al., 1990; Saraceno et al., 2009]. Thus, the shelf-break front and the
67 upper slope encompass a complex region that concentrates a diversity of top predators and other
68 species, some exploited by industrial fisheries [Martinetto et al., 2020; [Rey and Huettmann, 2020](#)].
69 The Southern elephant seal, *Mirounga leonina*, that breed and molt along the coast of the Península
70 Valdés represent one of the main top predators inhabiting these waters. These continuous and deep-
71 diving seals spend several months at sea foraging, coming back to the location from where they left
72 to breed or moult [Lewis et al., 2004]. After leaving the shore, post-breeding females cross the
73 continental shelf in 3-7 days. They forage for approximately two months in deep waters off the
74 shelf, while adult males usually forage along the outer part of the shelf break [[Campagna et al.,](#)
75 [1998, 1999](#)]. [Post-breeding females on deep waters show a dive rate of 2.2 dives/hr and a mean dive](#)
76 [duration of 22.8 minutes, where the mean dive depth range from the ocean surface to the bottom of](#)
77 [the continental shelf \(~80 m\) and beyond the shelf-break \(~ 600 m\)](#) [[Campagna et al., 1998](#)]. This
78 continuous and deep diving behavior lead to the idea of using seals as ocean samplers of physical,
79 biogeochemical and biological parameters by deploying on them small electronic devices
80 (biologgers) allowing the conduction of a broad range of studies linking seals foraging behavior to
81 the oceanographic environment [[MEOP project](#); [Boehlert et al., 2001](#); [Guinet et al., 2014](#); [Della](#)
82 [Penna et al., 2015](#)].

83
84 Taking advantage of the high frequency temperature, salinity, light, pressure and accelerometry
85 measurements obtained with the instruments carried by one seal, we investigated how ocean bio-
86 physical features changed and how its foraging behavior and success responded to those
87 environmental changes. This paper also aimed at investigating the consistency between satellite
88 measurements (temperature and chl-a) as well as CMEMS (Copernicus Marine Environment
89 Monitoring Service) model outputs with the in-situ hydrographic profiles provided by one seal.

90

91 **2. Data and Methods**

92 **2.1 In situ data**

93

94 As part of an international collaboration between the Centre National de la Recherche Scientifique
95 (CNRS), the Wildlife Conservation Society (WCS), the University of Tasmania (UTAS) and the
96 University of Buenos Aires (UBA), nine adult female seals were tracked along the Patagonian shelf
97 and outer ocean during the austral spring 2018. Following deployment procedures described in
98 Campagna et al. [1998, 1999, 2006], seals were equipped with positional Argos and/or GPS
99 systems, as well as different high frequency sampling sensors that record depth (pressure),
100 temperature, conductivity, fluorescence, irradiance (light), and acceleration (Table 1). In oceanic
101 Case 1 waters (i.e. waters whose optical properties are determined primarily phytoplankton
102 concentration and dissolved organic matter), light attenuation (i.e. the variation of the irradiance
103 signal with depth) quantifies the amount of light arriving at the ocean surface and is independent to
104 solar angle and cloud cover, thus, it can be used as a proxy of phytoplankton concentration [Bricaud
105 et al., 1998; Mobley et al., 2004; Jaud et al., 2012; Bayle et al., 2015]. The higher the phytoplankton
106 concentration the greater is light attenuation from the surface within the water column (Jaud et al.,
107 2012). Acceleration measurements associated with head movements provide an index of prey
108 capture attempts (PCA) (see Goulet et al., 2019). Satellite tags and data loggers were deployed in
109 October 2018, Península Valdés, Argentina, on breeding seal females prior to their departure to the
110 sea for their post-breeding foraging trip. At the end of their foraging trip, females return to shore.
111 After being located thanks to the satellite tag, they were recaptured to recover the loggers. For this
112 study we focus in only one seal (ID 14903) that, after leaving Península Valdés on October 25,
113 2018, spent two months foraging in a relatively small region over the slope, near the 600 m isobath,
114 and returned to the same departure location on January 10, 2019 (Figure 1). The studied seal was
115 equipped with a CTD-SRDLT tag [Boehme et al., 2009], including Argos transmitter (PPT), light
116 and accelerometry (12.5 Hz) sensors (Table 1). After recovering, the tag was calibrated following

117 the methodology from MEOP described in Roquet et al., [2001] and Siegelman et al., [2019]. The
118 accuracy of each profile location is of the order of 2 km [Boehme et al., 2009].

119
120

121

122

123

124

125

126

127

128

129

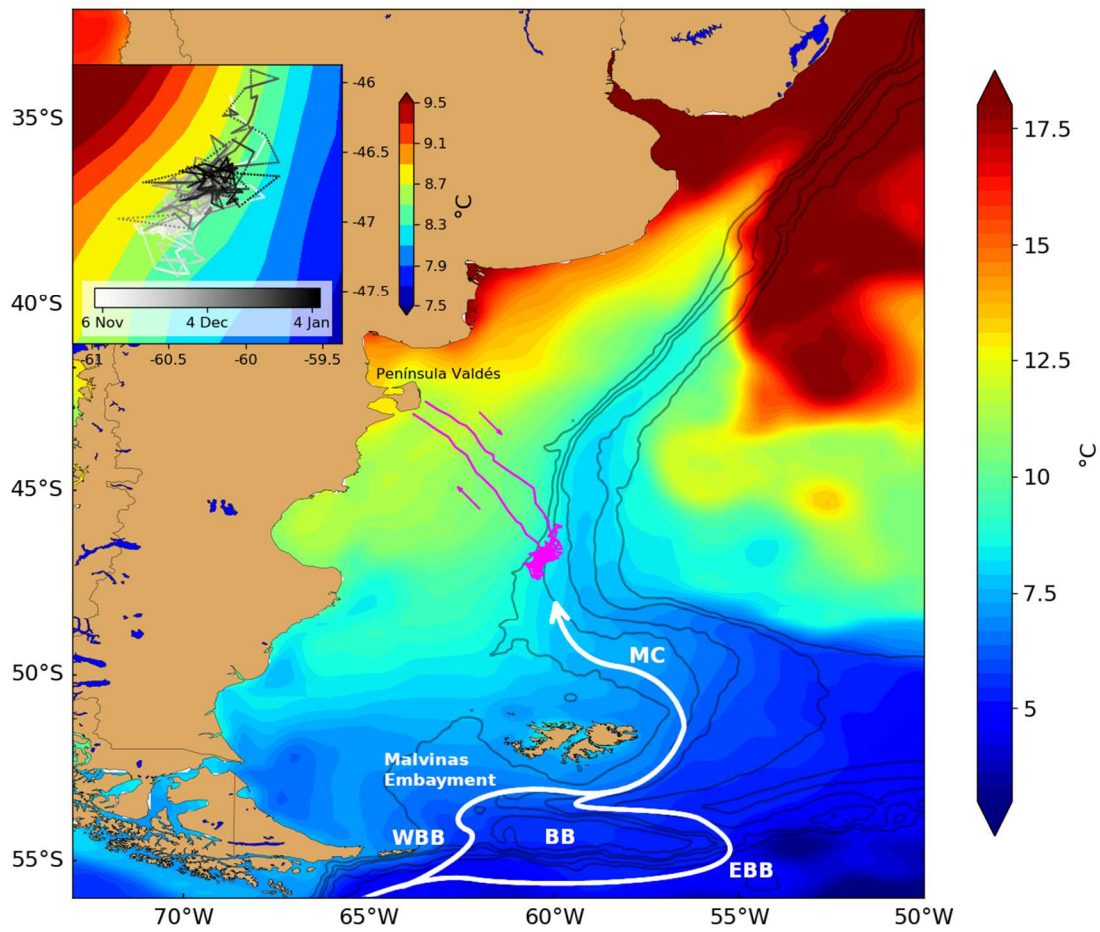
130

131

132

133

134



135 Figure 1. Tracks for the 73-day trip of one seal (magenta line) overlapping monthly-mean sea surface temperature (°C) for November
136 2018 from OSTIA (see section 2.3). Magenta arrows indicate direction of trajectory. The 200, 600, 1000, 2000 and 3000 m isobaths
137 are indicated with thin black lines. The white arrow illustrates the main path of the upper slope portion of Malvinas Current (MC)
138 through the western (WBB) and eastern (EBB) passages of the Burdwood Bank (BB). The top left inset zooms into the trajectory on
139 the slope. Note that the scale of the temperature contours is different.

140

141

142

143

144

145

146

147 Table 1. Main characteristics of the instruments deployed on the studied seal.

CTD-SRDL	Range	Accuracy	Resolution	Frequency
Temperature	-5 to 35 °C	± 0.005°C	0.001°C	2 Hz
Salinity	0 to 50	± 0.01	0.002	2 Hz
Pressure	0 to 2000 dBar	2 dBar ± 0.01% / K	0.05 dBar	2 Hz
Light	8.31×10^{-7} to 3000 $\mu\text{mol}/\text{m}^2\text{s}$	1%	0.1%	2 Hz

148

149

150 For the purpose of this study, and based on the sensor's accuracy, only dives deeper than 15 meters
 151 were considered. Bottom depth, light and the temperature profiles associated with each dive were
 152 extracted following the methodology described in Dragon et al., [2012a]. Each dive was divided
 153 into a descent, bottom and ascent phase. The bottom phase corresponds to the period between the
 154 end of the descent and the beginning of the ascent. The different dive phases were defined
 155 according to Dragon et al., [2012a] using a custom-written MATLAB code (version 7.0.1; available
 156 on request). Accelerometer data was processed following Viviant et al., [2010] and Gallon et al.,
 157 [2013] using custom-written MATLAB code (available on request) to assess PCA. While PCA
 158 provide fine scale insights of local variations of prey encounters, seal net foraging gain can be
 159 assessed by monitoring change in seal body density (i.e. the change in the relative proportion in
 160 between lipid and protein content) through its foraging trip. This is achieved by monitoring of the
 161 descent rate, when the seal is sinking passively through the water column during special
 162 resting/food processing dives qualified as 'drift dives' [Crocker et al., 1997; Biuw et al., 2003]. The
 163 descent speed during these drift phases is directly related to buoyancy, with more buoyant seals (i.e.
 164 with a higher lipid proportion) sinking at a slower rate compared with leaner ones [Crocker et al.,
 165 1997; Biuw et al., 2003; Miller et al., 2012]. However, such 'drift rate' data obtained from time-
 166 depth recorder tags can only provide a general temporal trend in body composition change and with
 167 a temporal lag of about 3 to 6 days compared to change of PCA rate [Dragon et al., 2012b; Richard

168 et al., 2014, 2016]. Each dive was attributed to either day or night periods. This was defined
169 according to the solar angle, taking into account the geographical location of the seal provided by
170 the Argos transmitter and the time at the beginning of the dive (see Guinet et al., 2014).

171
172 The animal spent most of the time in a boundary region with strong year-round zonal temperature
173 gradients (Figure 1, inset-panel), generated by the cold MC waters and the warmer continental shelf
174 waters [Saraceno et al., 2004; Piola et al., 2013]. Over the two-month period, the seal tended to dive
175 mostly in the southwestern part of the foraging area in November and in the northeastern part in
176 December (Figure 1, inset-panel). Thus, to be able to compare the monthly mean temperature and
177 salinity between November and December, the zonal movement of the seal was taken into account.
178 To achieve this, we kept only with a sub-sample of the data (~28% of the total dives in the upper
179 slope, i.e. 1146 from 4150 total profiles) after dividing the area in equal longitudinal bins and
180 selecting the same amount of profiles on each longitude bin for November and December,
181 discarding the rest. The selection of the discarded profiles was done randomly. This methodology
182 was repeated a hundred times, to compute finally a mean temperature averaged from the hundred
183 samples.

184 185 **2.2 Numerical model and reanalysis data**

186
187 The Operational Mercator global ocean analysis and forecast system (PSY4QV3R1) is based on
188 NEMO ocean model [Madec et al., 2019] and provides 3D global ocean forecasts updated daily,
189 starting in January 2016 for daily outputs and January 2007 for monthly outputs. It has a 1/12°
190 degree horizontal resolution with regular longitude/latitude equi-rectangular projection, resolved
191 over 50 vertical levels, ranging from 0 to 5500 m. Model outputs include daily and monthly mean
192 files of temperature, salinity, currents, sea level and mixed layer depth among others. Comparison
193 with satellite and Argo float data showed that the system correctly reproduced the general
194 circulation and the complex hydrographic features of the Southwestern Atlantic Ocean (Artana et

195 al., 2018b). Model data is available through Copernicus Marine Environment Monitoring Service
196 (CMEMS, <http://marine.copernicus.eu/>). To compare seal data with model, we downloaded daily
197 values of temperature and salinity for the levels corresponding to the 0-600 m depth range in seal's
198 foraging domain (46.2-47.3°S and 60.7-59.8°W). Valid seal measurements (after sensors stabilize in
199 the water column) begin at ~18 m depth, thus, model values from 18 m were interpolated to seal
200 location. Finally, a 4-hour moving average was applied to both series. For the temperature-salinity
201 diagram (Figure 4), seal profiles were selected when the distance between a given profile and the
202 nearest model grid point was less than 1 km. This procedure provided a total of 205 profiles for
203 each data set. Also, to have the same number of vertical values per profile, we selected the values
204 from the seal records that coincide with a depth level of the model (i.e. 32 levels, from the surface
205 to 550 m).

206 Cloud cover mentioned in Section 3.2 is a product of ERA5 global reanalysis. We used monthly
207 values of cloud area fraction in seal's foraging region, with a spatial resolution of 0.25°. Data is
208 available at <https://cds.climate.copernicus.eu/cdsapp#!/home>

209

210 **2.3 Satellite data**

211

212 Sea surface temperature (SST) considered here was produced by the Group for High Resolution
213 Sea Surface Temperature (GHRSSST) and is provided by the Operational Sea Surface Temperature
214 and Ice Analysis (OSTIA). The product merges in situ data, infra-red and micro-wave radiometers
215 and has a 0.05° x 0.05° horizontal resolution [Bell et al., 2000; Stark et al., 2007; Donlon et al.,
216 2012]. Near real time L4 product was downloaded from <http://marine.copernicus.eu/>. To compare
217 SST with seal temperature measurements at 18 m (Figure 3a), data was interpolated to seal location
218 and finally a 4-hour moving average applied.

219

220 Monthly mean composites and daily gridded L3 chlorophyll-a concentration MODIS-Aqua ocean
221 color data, with a spatial resolution of 4 km from the Ocean Biology Processing Group
222 (NASA/GSFC/OBPG) were downloaded from <https://oceandata.sci.gsfc.nasa.gov>.

223

224 **3. Results**

225

226 Sixty-three days out of the 73 day-long foraging trip were spent on the slope of the Patagonian
227 shelf break (Figure 1). The seal covered 4,500 kilometers and performed a total of 4,951 dives (i.e.
228 hydrographic profiles), from which 92% of them occurred over the slope. In this study we focus
229 only on the data provided by the seal while foraging over the Patagonian slope from 4 of November
230 2018 to 6 of January 2019.

231

232 **3.1 Physical environmental changes**

233

234 Temperature and salinity profiles evidenced contrasted oceanographic changes over the study
235 period between a warming upper layer (0-35 m) and cooling waters below (from 50 m to the
236 bottom; Figure 2). In the upper layer, the depth-averaged temperature increased (1.6°C) within the
237 two months due to higher net incoming heat radiation associated with the arrival of austral summer
238 (Figure 2a). This change in temperature was accompanied by a salinity increase of 0.09 in the upper
239 35 m, enhanced in the last two weeks (Figure 2b). Waters below the upper layer show a very
240 different behavior, with a mean temperature decrease in December of 0.25°C (Figure 2a) and an
241 increase in salinity of 0.04 (Figure 2b). If we minimize the zonal displacement of the seal (see
242 methods, Section 2.1), the temperature decrease and salinity increase in the subsurface layer
243 accounts for 0.15°C and 0.03 respectively.

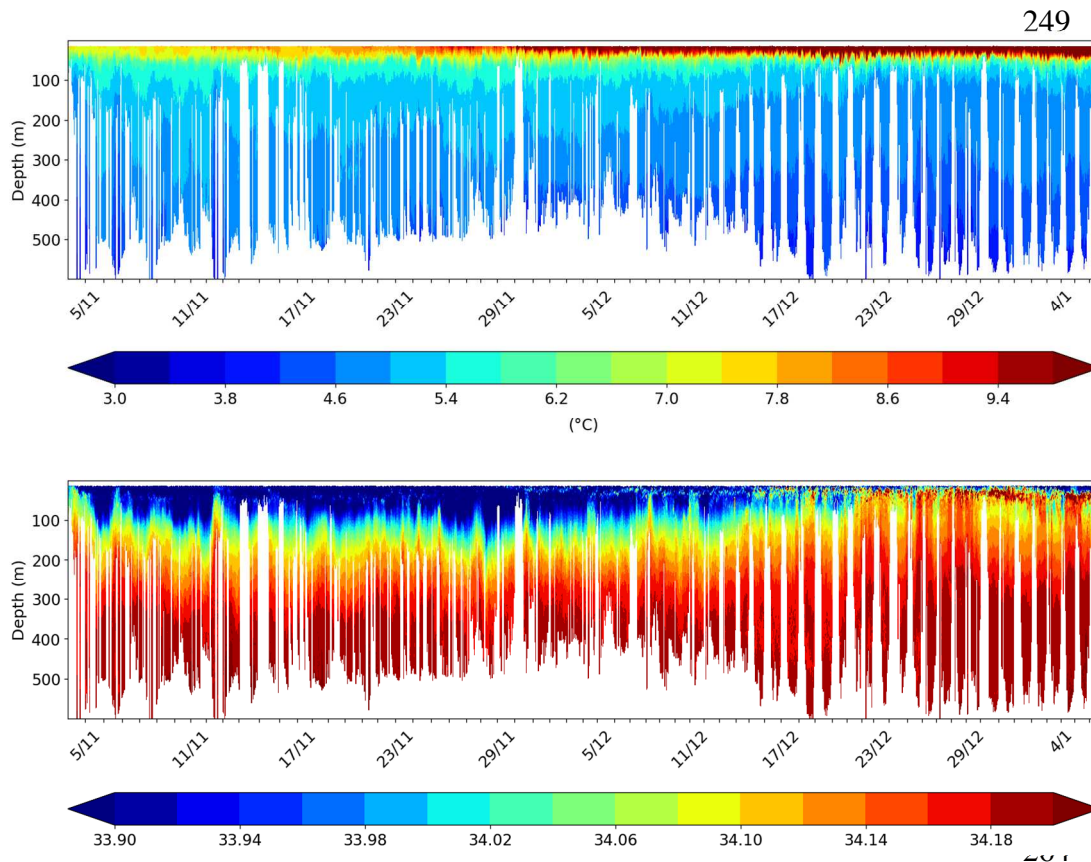
244

245

246

247

248

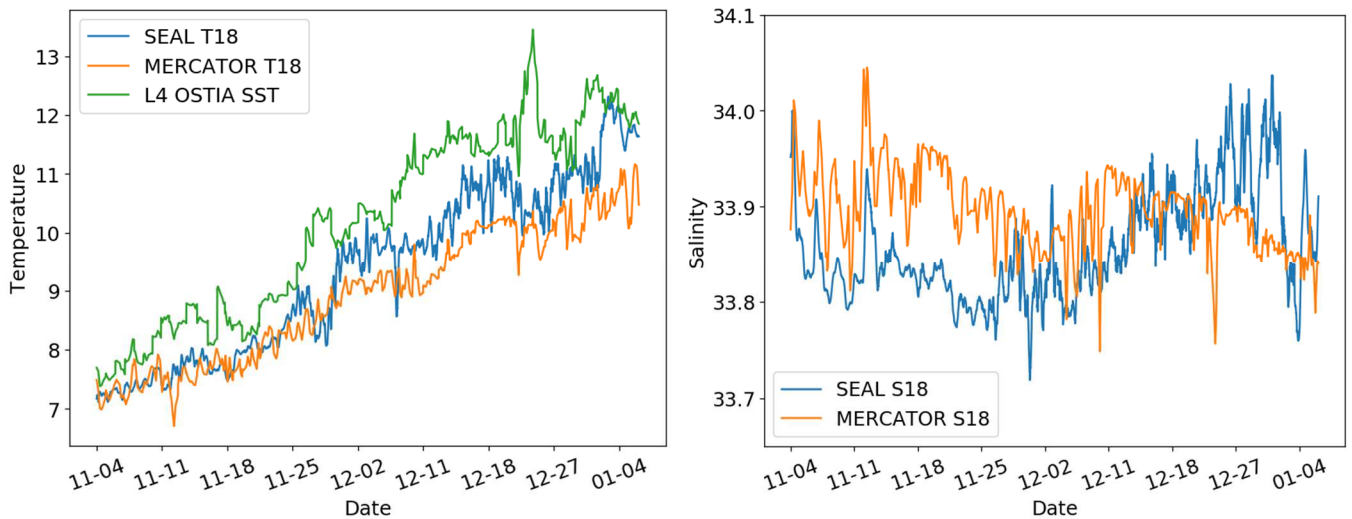


265 Figure 2. Temperature (°C, upper panel), and salinity (bottom panel) profiles measured by the seal in the Patagonian slope (top and
266 bottom panels respectively).

267

268 Near surface temperature (T18) and salinity (S18) values obtained by the seal were compared with
269 those from Mercator model and satellite observations following procedures described in Section 2
270 (Figure 3). The seal T18 signal presented visible discrepancies compared to Mercator temperature at
271 18 m and to OSTIA L4 surface values. Temperatures recorded by the seal were lower than OSTIA
272 during both months, and slightly higher than Mercator in December. Also, there were amplitude
273 differences between seal data and both series that result in a root mean square error of 0.62°C with
274 OSTIA and 1°C with Mercator. Nevertheless, they all described the same temperature increase
275 pattern during both months, and were highly correlated ($r = 0.91$ with Mercator and $r = 0.94$ with
276 OSTIA, at the 99% significance level) (Figure 3, left panel). As we were unable to compare our data

277 with satellite surface salinity measurements with spatial and temporal resolution comparable to the
 278 horizontal diving range of the seal, we only compared salinity recorded by the seal and Mercator
 279 outputs at 18 m (Figure 3, right panel). Mercator salinity showed higher values in November and
 280 early December compared to our data (root mean square error = 0.08), and a salinity decrease since
 281 December 11, the opposite of what was observed in our records, that increased during all December.
 282 Also, the two-month salinity time series were not significantly correlated; yet, if we split the series
 283 by month we found a significant correlation in November ($r = 0.5$), and no correlation in
 284 December.



285

286 Figure 3. Four hours moving average of temperature at 18 m from seal records (blue) and Mercator (orange) and at surface for
 287 OSTIA (green) (°C, left panel). Right panel shows the salinity from seal records at 18 m (blue), Mercator (orange).

288

289 Subsurface data obtained by the seal in the upper slope were compared with those from Mercator
 290 model in a temperature-salinity diagram (Figure 4) following the methodology explained in Section
 291 2.2. Following the water masses definitions described by Piola and Gordon [1989] and by
 292 Maamaatuaiahutapu et al. [1994], the temperature and salinity distribution indicated that the
 293 foraging region of the seal was composed mostly of three water masses: Subantarctic Surface
 294 Waters (temperature $> 5^{\circ}\text{C}$, salinity ~ 33.82), Subantarctic Mode Waters (temperature $\sim 4.7^{\circ}\text{C}$,
 295 salinity ~ 34.19) and Upper Antarctic Intermediate Waters (temperature $< 4^{\circ}\text{C}$, salinity ~ 34.18)

296 (Figure 4a). A closer inspection of the bottom water distribution (blue box in Figure 4a) is coherent
297 with the $\sim 0.15^{\circ}\text{C}$ December cooling observed in Figure 2, while a salinity change was
298 imperceptible (Figure 4b). Some major differences arise when comparing seal data with Mercator
299 temperature-salinity distribution. Subantarctic Surface Water for model distribution were less
300 disperse in salinity but showed a strong dispersion in the transition through Subantarctic Mode
301 Waters, that, in turn, presented a higher temperature, reaching 6°C (Figure 4c). The Antarctic
302 Intermediate Waters for model data (blue box in Figure 4c) appeared as two distinct waters masses,
303 one fresher and other saltier and slightly colder, the former being from November and the latter
304 from December (Figure 4d). The cooling observed in December may be due to intrusions of the MC
305 core [Piola et al., 2010; Matano et al., 2010; Combes and Matano, 2014] or to the increased
306 transport of Upper Antarctic Intermediate Waters from the Eastern passage of the Burdwood Bank
307 [Piola and Gordon, 1989; Matano et al., 2019].

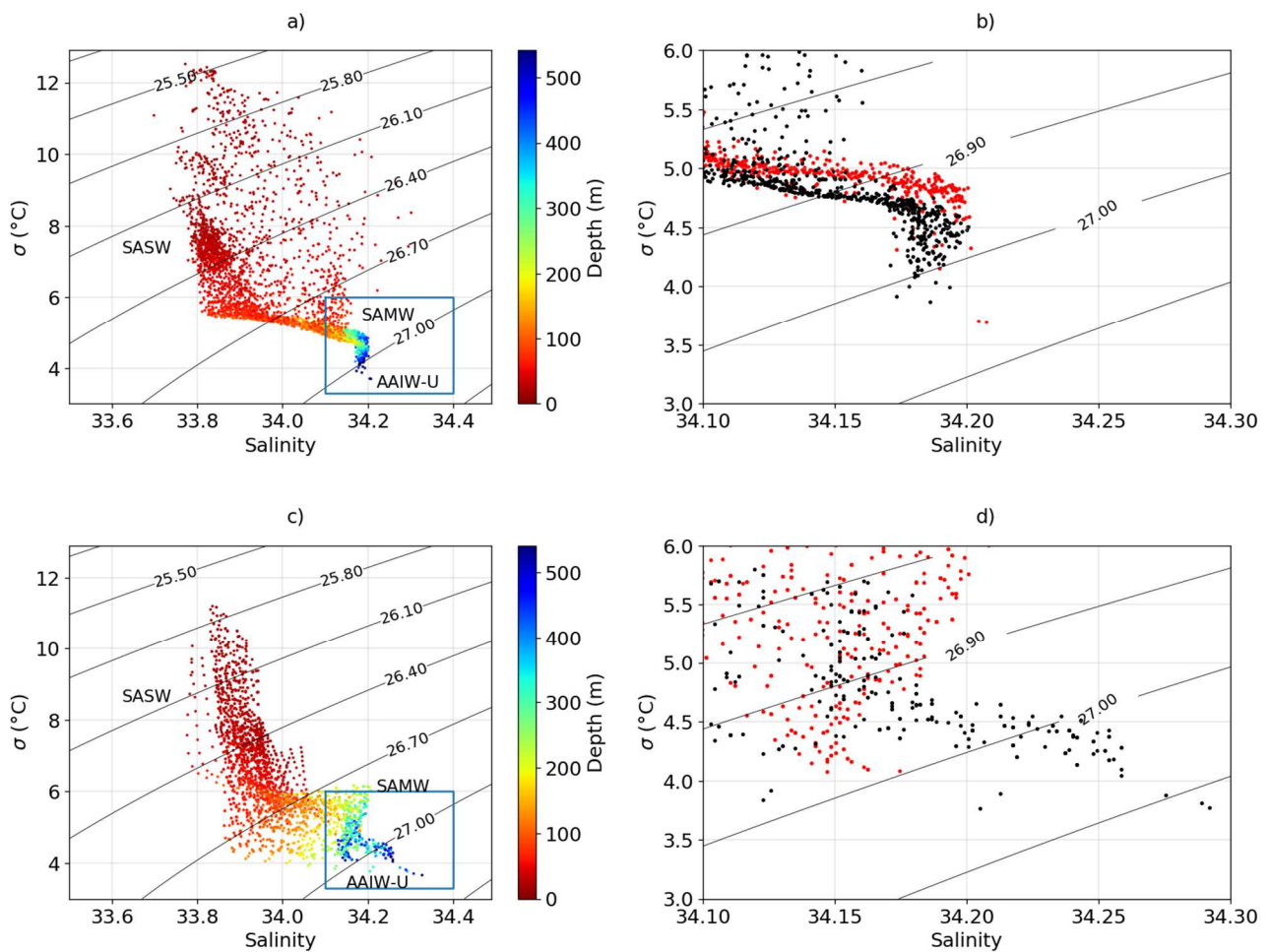
308

309

310

311

312



313
 314 Figure 4. Potential temperature – salinity diagrams from data recorded by the seal (a) and Mercator outputs (c). The colorbar
 315 indicates the depth of each dot and the gray lines the potential density isolines (sigma-theta). Deep waters (indicated with a blue box
 316 on both diagrams) are amplified in panel b) for the data recorded by the seal and panel d) for Mercator. The red dots in panels b and d
 317 correspond to November data and the black ones to December. The different water masses are indicated in panels a,c as SASW
 318 (Subantarctic Surface Waters), SAMW (Subantarctic Mode Waters) and AAIW-U (Upper Antarctic Intermediate Waters).

319

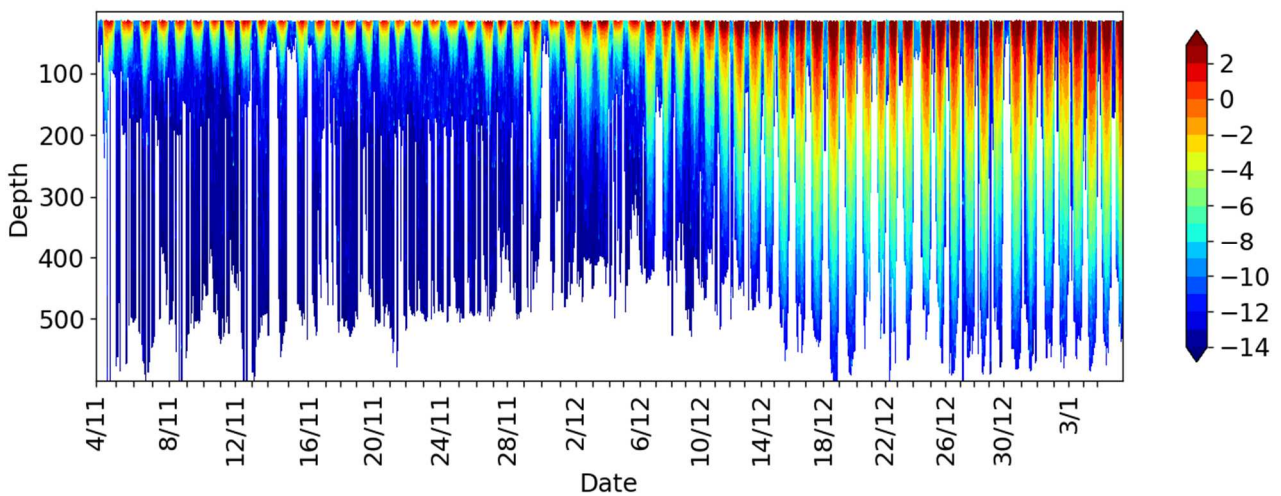
320 3.2 Change in phytoplankton concentration

321

322 Light attenuation in open ocean waters partially depends on the concentration of inorganic and
 323 organic particles suspended within the water column. In particular, phytoplankton (quantified
 324 through active measurement of chlorophyll-a fluorescence) constitutes the main source of particles
 325 in suspension within the euphotic layer, and it has been shown to be the main cause of light
 326 attenuation for Case 1 waters [Bricaud et al., 1998; Jaud et al., 2012]. In November, the irradiance
 327 in the Patagonian upper slope region, where the seal spent time, reached more than -8 ln

328 ($\mu\text{mol}/\text{m}^2/\text{s}$) only in the upper 150 m of the water column (Figure 5). However, the same amount of
329 irradiance reached 500 m depth in December, revealing a sharp decline in phytoplankton
330 concentration. Seasonal solar incidence angle and daylight duration also contributed, in a minor
331 way, to light increase in December. Cloudiness also affects the amount of light that reaches the
332 ocean surface. However, total cloud cover was higher in December compared to November (not
333 shown). Thus, the irradiance increase observed between November and December can be mostly
334 explained by the decrease in chlorophyll-a (chl-a hereafter) concentration.

335



336 Figure 5. Amount of light (irradiance) as logarithmic photosynthetic photon flux density ($\ln(\mu\text{mol}/\text{m}^2/\text{s})$) along depth (m) measured
337 by the seal in the slope region during November and December.

338

339 In-situ assessments of phytoplankton concentration were consistent with the MODIS-Aqua ocean
340 color satellite measurements over the study period for November 2018 (Figure 6a). This result
341 revealed large amounts of chl-a ($> 6 \text{ mg}/\text{m}^3$) over the region where the seal dove, and over the 200
342 m isobath, the shelf-break front (SBF). These large concentrations of chl-a generated a barrier of
343 suspended particles that inhibited light beyond the upper layers. In December 2018 (Figure 6b), the
344 chl-a concentration decreases ($< 2 \text{ mg}/\text{m}^3$), thus allowing the light to reach greater depths. The chl-a
345 concentration decrease was also observed in daily MODIS-Aqua ocean color chl-a and surface chl-a
346 concentration estimated by seal irradiance data and a linear functional model (LFM) (Figure 6c).

347 The linear functional model described in Bayle et al. 2015 is a predictive tool enabling to infer chl-a
348 concentration from irradiance along the water column through a per-profile functional approach
349 based on a learning dataset of concomitant vertical profiles. Light in the ocean has a direct
350 correlation with the vertical distribution of many marine organisms, ranging from zooplankton [Liu
351 et al., 2003] to fish [Batty et al., 1990] and marine mammals [Horning and Trillmich, 1999].
352 Elephant seals depend on suitable and abundant prey (fishes and squid) present at the fronts
353 [Campagna et al., 2007; Eder et al., 2010]. The area frequented by the studied animal, is associated
354 with the SBF. The SBF has high concentrations of chl-a and high primary production, which would
355 attract organisms of different trophic levels [Martinetto et al., 2020]. Moreover, the SBF is an area
356 of intense fishing activity [Portela et al., 2012; Alemany et al., 2014] (Figure 6a,b). The main
357 commercial species, at different times of the year, are Patagonian grenadier (*Macruronus*
358 *magellanicus*), shortfin squid (*Illex argentinus*) and Argentine hake (*Merluccius hubbsi*) [Portela et
359 al., 2012; Alemany et al., 2014]. Figure 6a,b shows that the fishing effort (in hours, Global Fishing
360 Watch, 2020) in the SBF was much larger in December 2018 (18031 hours) than in November 2018
361 (4841 hours), and that the international fishing fleet concentrated north of the seal foraging ground.
362 Figure 6 also shows that the largest fishing activity along the shelf-break did not correspond to the
363 period of time when the largest chl-a concentration was observed.

364

365

366

367

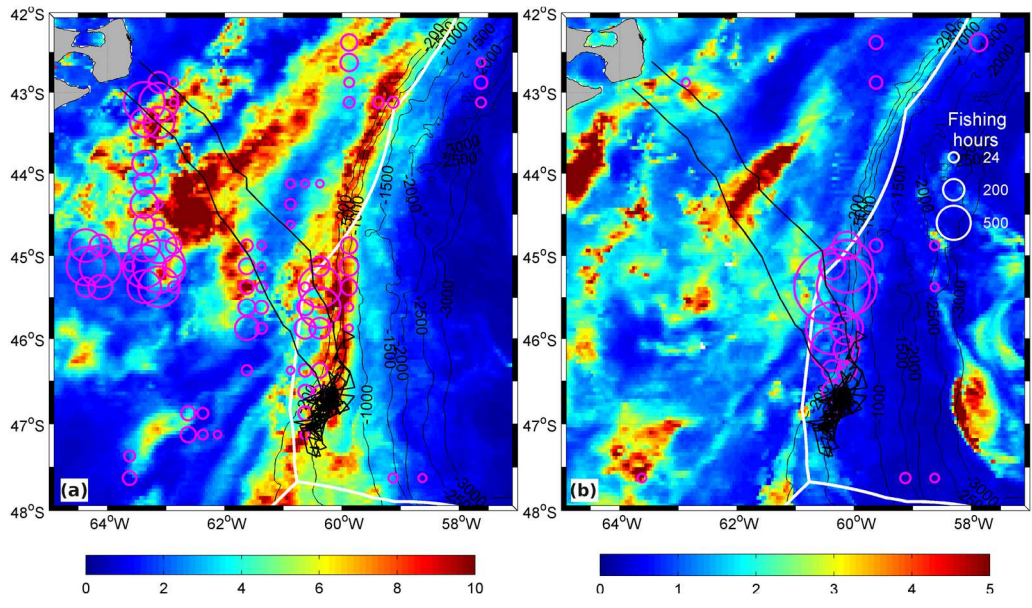
368

369

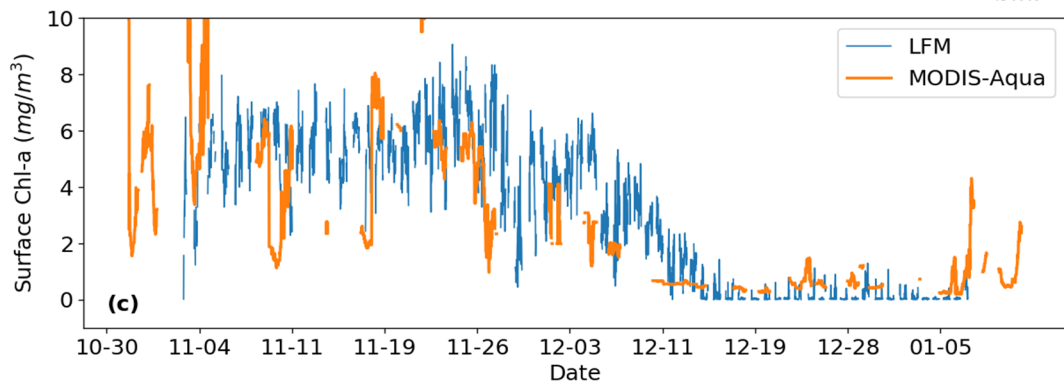
370

371

372
373
374
375
376
377
378
379
380
381
382
383
384
385
386
387



388



396
397
398
399
400
401
402
403
404

Figure 6. Modis-Aqua monthly mean chl-a concentration (mg/m^3) for November (a) and December (b) 2018. The magenta circles represent the fisheries activity for each month, the size is proportional to the fishing effort (hours). The white line represents the limit of the Argentine Exclusive Economic Zone. The black lines indicate the seal trajectory and the isobaths. Daily surface chl-a concentration (c) in mg/m^3 for satellite Modis-Aqua (orange) and estimated by the LFM method (blue).

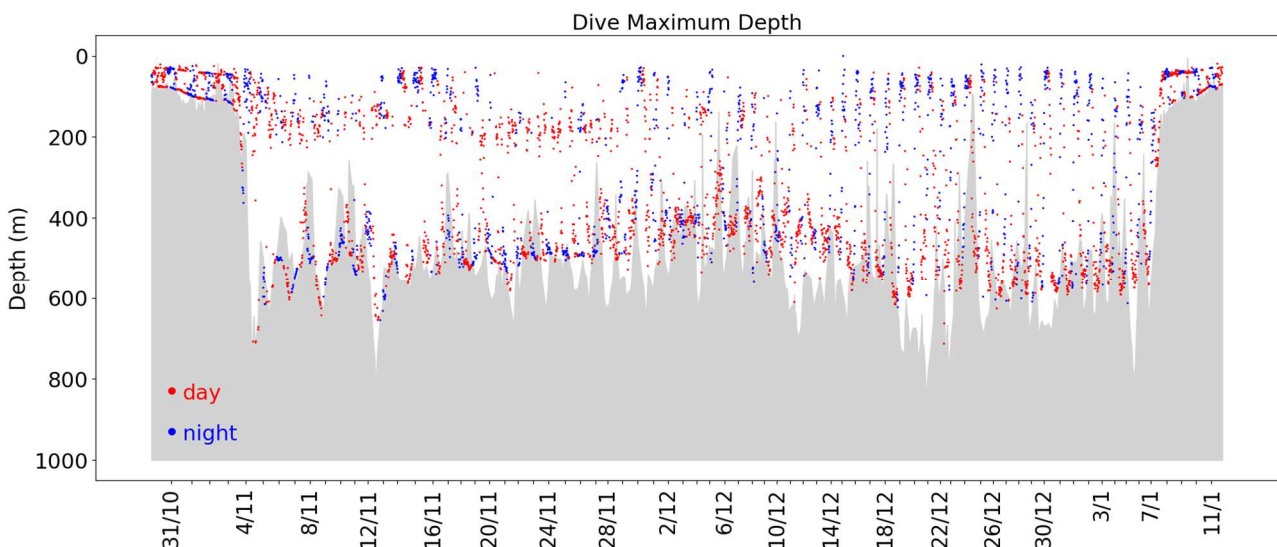
3.3 Elephant seal foraging behavior

405

406 The overall mean diving depth of the seal was 263.8 ± 192.0 m at night, with a mean dive duration
407 of 17.4 ± 4.3 minutes. During the day, the mean diving depth was 331.5 ± 186.7 m, with a mean
408 dive duration of 19.8 ± 3.9 minutes.

409 To better understand how ocean dynamics and light can affect elephant sea foraging behavior we
410 analyzed dive maximum depth (DMD) for each dive, and number of PCA performed. In November,
411 DMD were either on shallow waters (<200 m) or sweeping the ocean bottom, avoiding the middle
412 depths and with a minor difference between day and night (Figure 7). The sea floor (shading gray in
413 figures 7 and 8) is an interpolation of GEBCO 2019 bathymetry (15 arc-second resolution) to seal
414 diving locations, and DMD extending beyond the sea floor are due to possible errors on seal
415 location or bathymetric chart product. In December, the animal dove predominantly (58%) near the
416 surface at night (blue dots in Figure 7), and reached the ocean bottom (80%) during the day (red
417 dots in Figure 7). The mean DMD increased from 337.5 ± 173.7 m in November to 359.4 ± 180.4 m
418 in December and the mean dive duration increased from 17.6 ± 3.5 minutes in November (9.4 ± 3.3
419 minutes in the bottom phase) to 19.6 ± 4.5 minutes in December (11.1 ± 4.0 minutes in the bottom
420 phase).

421



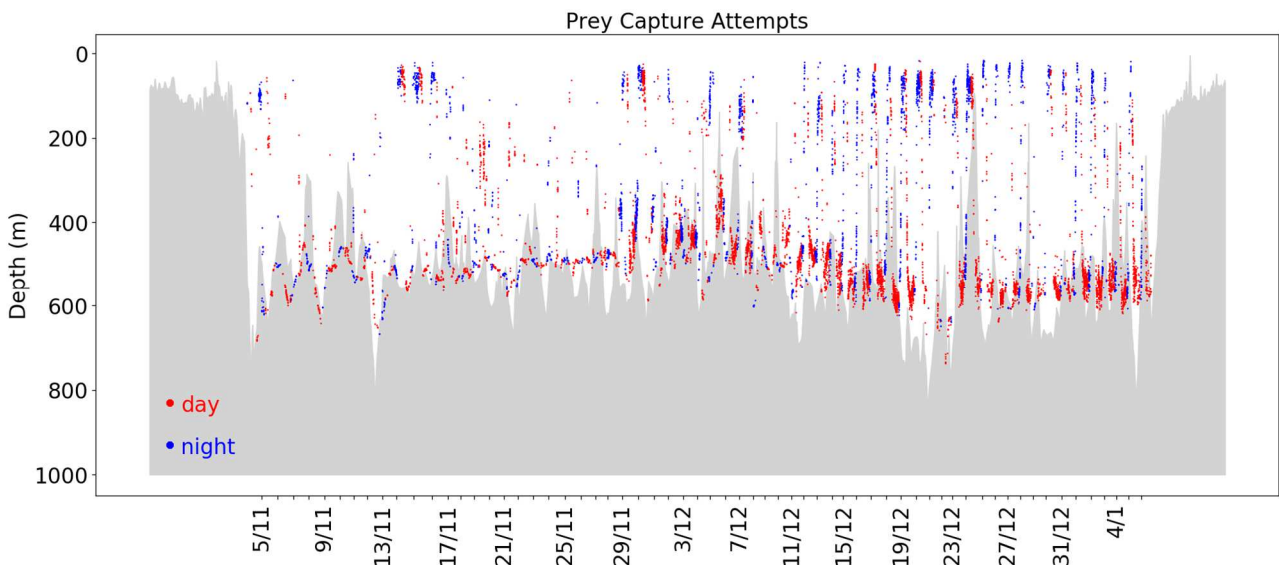
422 Figure 7. Dive maximum depth (DMD) with different colors for day (red) and night (blue). The shading gray is the bottom

423 topography from GEBCO 2019 interpolated along the elephant seal diving trip.

424

425 Compared to November, December was characterized by a significant increase in number of PCA
426 (287 versus 125 PCA per day) (Figure 8). Statistics for both DMD and PCA are summarized in
427 Table 2. In agreement with early studies [Guinet et al., 2014; Gallon et al., 2013; Schreer et al.,
428 2001], the majority of the PCA (~86%) occurred during the bottom phase of the dive. In November,
429 77% of the observed PCA were skimming the ocean bottom, both during day (red dots) and night
430 (blue dots) and with the particularity that an increase of PCA near the surface can be seen between
431 13-17 of November (Figure 8). In December, the majority of the PCA occurs in the upper (< 300 m)
432 part of the water column during night (54%) and in the lower part (> 300 m) during daylight hours
433 (89%). A strong decrease of PCA, especially near the ocean bottom was observed between 21-23 of
434 December.

435



436 Figure 8. Depth location of prey capture attempts (PCA) with different colors for day (red) and night (blue). The shading gray is the
437 bottom topography from GEBCO 2019 interpolated along the elephant seal diving trip.

438

439

440

441

442

443

444

445

446

447

448 Table 2. Total dives and percentages of DMD and PCA during November and December 2018. Upper water column (< 300 m) and
 449 lower water column (> 300 m).

	Dives		DMD		PCA	PCA	
	Total	Mean Duration (minutes)	Upper Water Column (%)	Lower Water Column (%)	Total	Upper Water Column (%)	Lower Water Column (%)
<u>November</u>	2093	17.6	40	60	3745	23	77
Day	1285	18.5	41	59	2194	20	80
Night	808	16.8	38	62	1551	29	71
<u>December</u>	2050	19.6	34	66	8617	24	76
Day	1287	21.2	20	80	5967	11	89
Night	763	18.0	58	42	2650	54	46

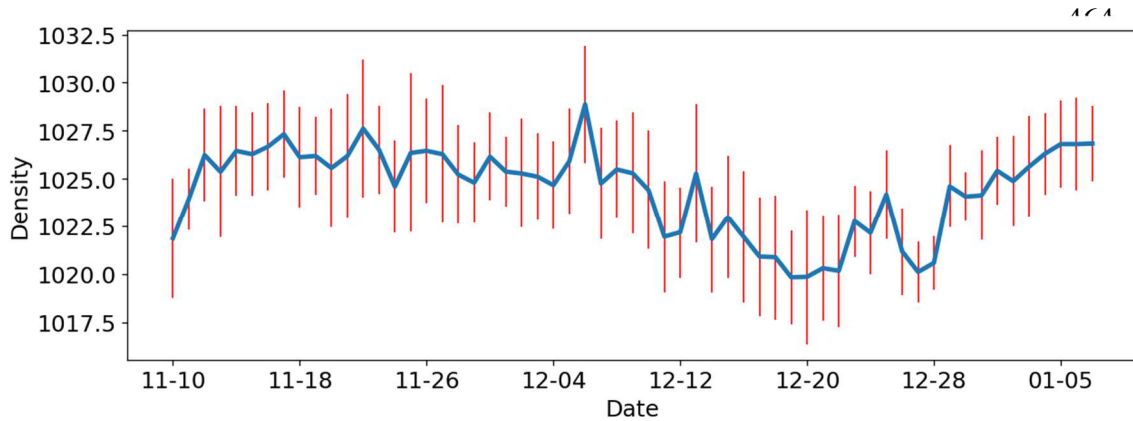
450

451

452 The monitoring of the female body density, reveals that seal density increased to remain high
 453 through the whole month of November and started to decrease (i.e. improvement of body condition
 454 due to an increase in lipid content, which is linked to a higher prey ingestion) by early December
 455 (Figure 9), which matches well with the change in environmental conditions (keeping in mind that
 456 there is a lag of 3 to 7 days between prey ingestion and body density changes [Dragon et al., 2012b;
 457 Richard et al., 2014, 2016]).

458

459
460
461
462
463



472 Figure 9. Seal daily mean density (kg/m^3) estimated along the trip (blue line) and standard deviations (red lines).

473

474 **4. Summary and Discussion**

475

476 High frequency data collected along the upper portion of the Patagonian slope during November
477 and December 2018 by one post-breeding female elephant seal equipped with multiple sensors was
478 used to investigate the interaction between ocean dynamics and seal diving and foraging behavior.
479 Measurements obtained by the seal showed a decrease in temperature (0.15°C) and a salinity
480 increase (0.03) below the mixed layer during December in the Patagonian slope, at $46\text{-}47.5^\circ\text{S}$.
481 Temperature outputs from Mercator model at 18 m and OSTIA satellite data were in agreement with
482 the increase pattern observed from seal data at 18 m. Nevertheless, our records showed higher
483 temperatures than Mercator in December, and discrepancies in amplitude oscillations. On the other
484 hand, salinity outputs from Mercator showed major differences with that obtained by the seal at 18
485 m, mostly in December, when Mercator data showed an opposite pattern. The differences between
486 seal and Mercator data may be due to a mismatch between seal location and ARGOS positioning
487 system, but also to the fact that seal's data used for the comparison was from around 18 meters

488 (where sensors stabilize and give more accurate measurements). Given that temperature and salinity
489 profiles recorded by the seal indicate a mixed layer depth of ~25 m in November and ~18 m in
490 December (not shown), measuring at this depth level may lead to uncertainties, mostly in
491 December. Also, SRLD-CTD sensors could suffer an erosion of its accuracy during the seal's trip,
492 specially for salinity measurements [Santini et al., 2018]. Light data also revealed a significant
493 increase of irradiance in December (almost reaching the ocean bottom) associated with a decrease
494 of chl-a in the upper levels revealed by our records and satellite Modis-Aqua data. Indeed, the
495 largest concentration of chl-a values in the Patagonian shelf and slope are usually observed in
496 November, during the austral spring bloom [Saraceno et al., 2005; Romero et al., 2006; Rivas et al.,
497 2006] and start decreasing in December due to nutrient consumption and ocean stratification that
498 inhibits the upwelling of nutrients to the euphotic zone.

499
500 Fishing effort in December (larger compared to November,) was concentrated north of the seal
501 foraging ground. The spatial mismatch between the fishing activity and the seal foraging ground
502 can be due to several causes, as seal trying to avoid fishing lines and nets and/or a different spatial
503 distribution or differences in the targeted fish/squids species by seal and fisheries. Indeed, the seal
504 might be eating preys that are not of commercial interest, like lantern fish (Myctophids) that are
505 very common in this area [Figueroa et al., 1998; Eder et al., 2010; McGovern et al., 2019]. Yet, we
506 do not have data to explain the spatial distribution mismatch. Also, the largest fishing activity along
507 the shelf-break did not corresponded to the period of time when the largest chl-a concentration was
508 observed. This fact can be explained considering different possibilities: (i) the most interesting
509 commercial species arrived in December with the colder and saltier water mass observed above; (ii)
510 the species that took advantage of the large phytoplankton bloom in November grow enough so in
511 December became prey of commercial species. Last but not least, the fact that both in November
512 and in December relatively large chl-a concentrations were observed along the upper portion of the

513 Patagonian slope in comparison to adjacent waters is a clear indication that a physical forcing
514 related to the bathymetry drives the large concentration of chl-a observed.

515

516 The elephant seal also showed a distinct behavior during the two-months trip. In December, the
517 PCA increased more than double with respect to November with the seal diving preferentially at
518 surface during the night and in the bottom layers during the day. The increase in PCA observed in
519 December can be attributed to a higher concentration of prey items due to the arrival of preys with
520 cold nutrient rich Antarctic Intermediate Waters. This result was also reflected in the larger fishing
521 effort detected in December. The density change of the seal along the trip, reflected quite well the
522 increment of PCA at the beginning of December. Moreover, the density increase observed around
523 the 23-26 of December matched with the observed decrease in PCA between 21-23 of December.
524 The increase of PCA in December can be seen both in the surface and bottom layers, suggesting that
525 the arrival preys migrated along the whole water column in response of light variability. Results
526 also showed that in December DMD increased 6% compared to November (66% of the dives in
527 December were below 300 m depth against 60% in November) and an increase of 2 minutes in
528 dives duration. These results are in agreement with previous studies that found a negative
529 correlation between surface chl-a concentration and both DMD and dive duration [Dragon et al.,
530 2010].

531

532 The relationship between prey location in the water column and light is known to dominate seals
533 diving behavior [Jaud et al., 2012; Guinet et al., 2014]. Several studies have shown that light level
534 precisely controls the vertical distribution of many species, in particular myctophids, which are part
535 of the seals diet [Catul et al., 2011]. Lower surface chl-a concentration means higher light
536 penetration, thus a deeper distribution of myctophids species. According to this, seal behavior in
537 December suggests that shallower dives during the night are related to preys that tend to concentrate
538 near the surface. During the day, as light increases in the water column, prey items tend to displace

539 to higher depths, and so does the seal, with predominant DMD and PCA near the seabed in
540 December.

541
542 We can expect that low [irradiance \(light\)](#) in November due to high chl-a concentration should be
543 associated with myctophids prey located in shallower waters, conditioning the seal to dive and
544 forage closer to the surface. However, November data showed that the seal dived predominantly
545 near the seabed, where most of the PCA were detected. This result suggests that a different prey
546 item, other than myctophids, can be the main diet for seals in the ocean bottom. This hypothesis is
547 in agreement with McIntyre et al., [2011] which came to the conclusion that seals can have different
548 forage strategies and dives, reaching greater depths at night compared with the day and that the
549 change can be associated with foraging of different prey resources. Furthermore, Goulet et al.,
550 [2020] suggest that female seals in Península Valdés are mainly pursuing larger or harder to catch
551 prey that do not produce bioluminescence as a predator defense and are probably more dispersed in
552 the water column.

553
554 The Patagonian slope is a very complex region, where subantarctic waters and bathymetry give
555 raise to a unique area of ecological processes and one of the greatest fisheries in the world. The
556 uniqueness of simultaneous high resolution in-situ data (physical, biogeochemical and biological)
557 collected by seals are a valuable resource to address oceanographic research and to calibrate and
558 validate ocean numerical models and satellite data. In particular, the data obtained from one seal
559 allowed us to improve our knowledge of the MC in the upper portion of the Patagonian slope, at
560 47°S, and also to link the sharp change in the water masses to the animal behavior. To deepen this
561 study, future campaigns in the region are thought to track fully equipped male elephant seals that
562 are known to dive preferentially in the Patagonian slope. Also, it is planned to incorporate
563 photographic cameras to the seals to try to unveil the prey items. Complementary data from
564 fisheries will also help elucidate the diving behavior and forage strategies in the region.

565

566 **References**

567

568 Alemany D., Acha E.M., and Iribarne O.O. (2014). Marine fronts are important fishing areas for
569 demersal species at the Argentine Sea (Southwest Atlantic Ocean). *Journal of sea research*, 87,
570 56-67

571

572 Artana C., Lellouche J.M., Sennéchaël N., and Provost C. (2018a). The open-ocean side of the
573 Malvinas Current in Argo floats and 24 years of Mercator Ocean high-resolution (1/12) physical
574 reanalysis. *Journal of Geophysical Research: Oceans*, 123, 8489–8507.
575 <https://doi.org/10.1029/2018JC014528>

576

577 Artana, C., Lellouche, J.M., Park, Y.H., Garric, G., Koenig, Z., Sennéchaël, N., et al. (2018b).
578 Fronts of the Malvinas Current System: Surface and Subsurface Expressions Revealed by
579 Satellite Altimetry, Argo Floats, and Mercator Operational Model Outputs. *Journal of*
580 *Geophysical Research: Oceans*, 123(8), 5261–5285. <https://doi.org/10.1029/2018JC013887>

581

582 Batty R., Blaxter J., Richard J. (1990). Light intensity and the feeding behaviour of herring, *Clupea*
583 *harengus*. *Marine Biology* 107: 383–388

584

585 Bayle S., Monestiez P., Guinet C., Nerini D. (2015). Moving toward finer scales in oceanography:
586 Predictive linear functional model of Chlorophyll *a* profile from light data. *Prog. Oceanogr.*
587 Volume 134, Pages 221-231, ISSN 0079-6611, doi : 10.1016/j.pocean.2015.02.001

588

589 Bell M. J., Forbes R.M. and Hines A. (2000). Assessment of the FOAM global data assimilation
590 system for real-time operational ocean forecasting. *J. of Marine Systems*, 25(2000) 1-22

591

592 Biuw, M., McConnell, B., Bradshaw, C.J.A., Burton, H. and Fedak, M. (2003). Blubber and
593 buoyancy: monitoring the body condition of free-ranging seals using simple dive characteristics.
594 J. Exp. Biol. 206, 3405–3423. (doi:10.1242/jeb.00583)

595

596 Blanke B., and Raynaud S. (1997). Kinematics of the Pacific Equatorial Undercurrent: a Eulerian
597 and Lagrangian approach from GCM results. J. Phys. Oceanogr., 27, 1038-1053

598

599 Boehlert G.W., Costa D.P., Crocker D.E., Green P., O'Brien T., Levitus S., Le Boeuf B.J. (2001).
600 Autonomous Pinniped Environmental Samplers: Using Instrumented Animals as Oceanographic
601 Data Collectors. J. Atmos. Oceanic Technol. 18 (11): 1882–1893

602

603 Boehme L., Lovell P., Biuw M., Roquet F., Nicholson J., Thorpe S.E., Meredith M.P. and Fedak M.
604 (2009). Technical note: Animal-borne CTD-satellite relay data loggers for real-time
605 oceanographic data collection. Ocean Sci., 5, 685–695

606

607 Bricaud A., Morel A., Babin M., Allali K., Claustre H. (1998) .Variations of light absorption by
608 suspended particles with chlorophyll a concentration in oceanic (case 1) waters: Analysis and
609 implications for bio-optical models. Journal of Geophysical Research 103: 31,033–031,044

610

611 Campagna C., Quintana F., Le Boeuf B.J., Blackwell S., Crocker D. (1998). Diving behaviour and
612 foraging ecology of female southern elephant seals from Patagonia. Aquatic Mammals 4, 1–11

613

614 Campagna C., Fedak M.A., McConnell B.J. (1999). Postbreeding distribution and diving behavior
615 of adult male southern elephant seals from Patagonia. Journal of Mammalogy 80, 1341–1352

616

617 Campagna, C., Piola, A.R., Rosa Marin, M., Lewis, M., & Fernández, T. (2006). Southern elephant
618 seal trajectories, fronts and eddies in the Brazil/Malvinas Confluence. *Deep-Sea Research Part I:
619 Oceanographic Research Papers*, 53(12), 1907–1924. <https://doi.org/10.1016/j.dsr.2006.08.015>

620

621 Campagna C., Piola A.R., Marin M.R., Lewis M., Zajaczkovski U., and Fernández T. (2007). Deep
622 divers in shallow seas: Southern elephant seals on the Patagonian shelf. *Deep Sea Research Part I:
623 Oceanographic Research Papers*, 54(10), 1792-1814

624

625 Catul V., Gauns M., Karuppasamy P. (2011). A review on mesopelagic fishes belonging to family
626 Myctophidae. *Reviews in Fish Biology and Fisheries* 21: 339–354

627

628 Chelton D.B., Schlax M. G., Witter D.L. and Richman J.G. (1990). GEOSAT altimeter observations
629 of the surface circulation of the Southern Ocean, *J. Geophys. Res.*, 95, 17,877–17,903

630

631 [Combes, V., and Matano, R.P. \(2014\). A two-way nested simulation of the oceanic circulation in the
632 Southwestern Atlantic, *J. Geophys. Res. Oceans*, 119, 731–756, doi:10.1002/2013JC009498](#)

633

634 Crocker, D., Le Boeuf B. and Costa, D. (1997). Drift diving in female northern elephant seals:
635 Implications for food processing. *Canadian Journal of Zoology*. 75. 27-39. 10.1139/z97-004

636

637 Della Penna, A., De Monte, S., Kestenare, E. et al., (2015). Quasi-planktonic behavior of foraging
638 top marine predators. *Sci Rep* 5, 18063. <https://doi.org/10.1038/srep18063>

639

640 Donlon C.J., Martin M., Stark J., Roberts-Jones J., Fiedler E. and Wimmer W. (2012). The
641 Operational Sea Surface Temperature and Sea Ice Analysis (OSTIA) system. Remote Sensing of
642 the Environment. doi: 10.1016/j.rse.2010.10.017 2011
643

644 Dragon A.C., Monestiez P., Bar-Hen A., Guinet C. (2010). Linking foraging behaviour to physical
645 oceanographic structures: Southern elephant seals and mesoscale eddies east of Kerguelen
646 Islands. Progress in Oceanography 87: 61–71
647

648 Dragon A.C., Bar-Hen A., Monestiez P., Guinet C. (2012a). Horizontal and vertical movements as
649 predictors of foraging success in a marine predator. Mar Ecol Prog Ser 447:243–257.
650 10.3354/meps09498
651

652 Dragon A.C., Bar-Hen A., Monestiez P., Guinet C. (2012b). Comparative analysis of methods for
653 inferring successful foraging areas from Argos and GPS tracking data. Mar Ecol Prog Ser
654 452:253-267. <https://doi.org/10.3354/meps09618>
655

656 Eder E.B., Lewis M.N., Campagna C. and Koch P.L. (2010). Evidence of demersal foraging from
657 stable isotope analysis of juvenile elephant seals from Patagonia. Marine Mammal Science, 26(2),
658 430-442
659

660 Figueroa D.E, Díaz de Astarloa J.M., Martos P. (1998). Mesopelagic fish distribution in the
661 southwest Atlantic in relation to water masses. Deep-Sea Res I Oceanogr Res Pap 45:317-332
662

663 Gallon S., Bailleul F., Charrassin J.B., Guinet C., Bost C.A., Handrich Y., et al. (2013). Identifying
664 foraging events in deep diving southern elephant seals, *Mirounga leonina*, using acceleration data
665 loggers. *Deep Sea Res Part II Top Stud Oceanogr.* 2013;88–89: 14–22
666

667 Garcia, V.T, Garcia, C.A.E., Mata, M.M., Pollery, R.C., Piola, A.R., Signorini, S.R., McClain, C.R.,
668 Iglesias-Rodríguez, M.D. (2008). Environmental factors controlling the phytoplankton blooms at
669 the Patagonia shelf-break in spring. *Deep-Sea Research Part II*, 55, 1150-1166.
670 doi:10.1016/j.dsr.2008.04.011
671

672 Garcia, C.A.E., Garcia, V.T, Dogliotti, A.I., Ferreira, A., Romero, S.I., Mannino, A., Souza, M.S.,
673 Mata, M.M. (2011). Environmental conditions and bio-optical signature of a coccolithophorid
674 bloom in the Patagonian shelf. *Journal of Geophysical Research*, 116, C03025.
675 doi:10.1029/2010JC006595
676

677 Global Fishing Watch. (2020). <http://www.globalfishingwatch.org>
678

679 Goulet, P., Guinet, C., Swift, R., Madsen, P.T., & Johnson, M. (2019). A miniature biomimetic sonar
680 and movement tag to study the biotic environment and predator-prey interactions in aquatic
681 animals. *Deep Sea Research Part I: Oceanographic Research Papers*, 148, 1-11
682

683 Goulet, P., Guinet, C., Campagna, C., Campagna, J., Tyack, P.L., & Johnson, M. (2020). Flash and
684 grab: deep-diving southern elephant seals trigger anti-predator flashes in bioluminescent prey.
685 *Journal of Experimental Biology*, 223(10)
686

687 Guerrero, R.A., Baldoni, A.G., Benavides, H.R. (1999). Oceanographic conditions at the southern
688 end of the Argentine continental slope. INIDEP DOC. CIENT., 5, pp. 7-22
689

690 Guinet C., Vacquié-Garcia J., Picard B., Bessigneul G., et al. (2014). Southern elephant seal
691 foraging success in relation to temperature and light conditions: insight into prey distribution.
692 *Mar Ecol Prog Ser* 499:285-301. <https://doi.org/10.3354/meps10660>
693

694 Horning M., Trillmich F. (1999). Lunar cycles in diel prey migrations exert a stronger effect on the
695 diving of juveniles than adult Galapagos fur seals. *Proceedings of the Royal Society of London*
696 *Series B: Biological Sciences* 266:1127–1132
697

698 Jaud T., Dragon A.C., Garcia J.V., Guinet C. (2012). Relationship between Chlorophyll a
699 Concentration, Light Attenuation and Diving Depth of the Southern Elephant Seal *Mirounga*
700 *leonina*. *PLoS ONE* 7(10): e47444. doi:10.1371/journal.pone.0047444
701

702 Lewis, M., Campagna, C., & Zavatti, J. (2004). Annual cycle and inter-annual variation in the haul-
703 out pattern of an increasing southern elephant seal colony. *Antarctic Science*, 16, 219-226
704

705 Liu S.H., Sun S., Han B.P. (2003). Diel vertical migration of zooplankton following optimal food
706 intake under predation. *Journal of plankton research* 25: 1069–1077
707

708 [Maamaatuaiahutapu, K., Garçon, V. C., Provost, C., Boulahdid, M. and Bianchi, A. A. \(1994\).](#)
709 [Spring and winter water mass composition in the Brazil-Malvinas Confluence, *Journal of*](#)
710 [Marine Research](#), 52, 397–426. <https://doi.org/10.1357/0022240943077064>
711

712 Madec G., Bourdallé-Badie R., Chanut J., Clementi E., Coward A., Ethé C., et al. (2019). NEMO
713 ocean engine (Version v4.0). Notes Du Pôle De Modélisation De L'institut Pierre-simon Laplace
714 (IPSL). Zenodo. <http://doi.org/10.5281/zenodo.3878122>
715

716 Martinetto, P., Alemany, D., Botto, F., Mastrángelo, M., Falabella, V., Acha, E.M., et al. (2020).
717 Linking the scientific knowledge on marine frontal systems with ecosystem services. *Ambio*,
718 49(2), 541–556. <https://doi.org/10.1007/s13280-019-01222-w>
719

720 Matano R.P., and Palma E.D. (2008). On the upwelling of downwelling currents. *Journal of*
721 *Physical Oceanography*, 38, 2482–2500. <https://doi.org/10.1175/2008JPO3783.1>
722

723 [Matano, R.P., Palma, E.D., Piola, A.R. \(2010\). The influence of the Brazil and Malvinas Currents on](#)
724 [the southwestern Atlantic Shelf circulation. *Ocean Science*, 6\(4\), 983–995.](#)
725 <https://doi.org/10.5194/os-6-983-2010>
726

727 Matano, R.P., Palma, E.D., & Combes, V. (2019). The Burdwood Bank circulation. *Journal of*
728 *Geophysical Research: Oceans*, 124. <https://doi.org/10.1029/2019JC015001>
729

730 McGovern, K.A., Rodríguez, D.H., Lewis, M.N., & Davis, R.W. (2019). Diving classification and
731 behavior of free-ranging female southern elephant seals based on three-dimensional movements
732 and video-recorded observations. *Marine Ecology Progress Series*, 620, 215-232
733

734 McIntyre T., Bornemann H., Plötz J., Tosh C.A., Bester M.N. (2011). Water column use and forage
735 strategies of female southern elephant seals from Marion Island. *Mar Biol* 158: 2125–2139
736

737 [MEOP \(Marine Mammals Exploring the Oceans Pole to Pole\) project. www.meop.net](http://www.meop.net)
738

739 Miller, P.J.O., Biuw, M., Watanabe, Y.Y., Thompson, D. and Fedak, M.A. (2012). Sink fast and
740 swim harder! Round-trip cost-of-transport for buoyant divers. *J. Exp.Biol.* 215, 3622-3630
741

742 [Mobley, C.D., Stramski, D., Bissett, W.P., Boss, E., \(2004\). Optical modeling of ocean waters: Is the](#)
743 [Case 1 –Case 2 classification still useful? *Oceanography* 17 \(2\), 60-67](#)
744

745 Parker G., Violante R.A. and Paterlini C.M. (1996). Fisiografía de la Plataforma Continental.
746 In : Ramos, V. & Turic, M. (eds) *Geología y Recursos Naturales de la Plataforma Continental*
747 *Argentina. Relatorio XIII Congreso Geológico Argentino. Asociación Geológica Argentina,*
748 *Buenos Aires, 1–16*
749

750 Piola A.R., Figueroa H.A., and Bianchi A.A. (1987). Some aspects of the surface circulation
751 south of 20°S revealed by First GARP Global Experiment drifters, *J. Geophys. Res.*, 92,
752 5101-5114
753

754 Piola A.R., Gordon A.L. (1989). Intermediate waters in the southwest South Atlantic. *Deep Sea*
755 *Research Part A. Oceanographic Research Papers, Volume 36, Issue 1, 1989, Pages 1-16, ISSN*
756 *0198-0149, [https://doi.org/10.1016/0198-0149\(89\)90015-0](https://doi.org/10.1016/0198-0149(89)90015-0)*
757

758 [Piola, A.R., Martínez Avellaneda, N., Guerrero, R.A., Jardón, F.P., Palma, E.D., Romero, S.I.](#)
759 [\(2010\). Malvinas-slope water intrusions on the northern Patagonia continental shelf, *Ocean Sci.*,](#)
760 [6, 345–359](#)
761

762 Piola A.R., Franco B.C., Palma E.D., Saraceno M. (2013). Multiple jets in the Malvinas Current. J.
763 Geophys. Res. Oceans, 118 (2013), pp. 2107-2117, 10.1002/jgrc.20170
764

765 Portela J., et al. (2012). Management Strategies to Limit the Impact of Bottom Trawling on VMEs
766 in the High Seas of the SW Atlantic, Marine Ecosystems, Dr. Antonio Cruzado (Ed.), ISBN: 978-
767 953-51-0176-5, InTech
768

769 Rey, A.R., Huettmann, F. (2020). [Telecoupling analysis of the Patagonian Shelf: A new approach to](#)
770 [study global sea bird-fisheries interactions to achieve sustainability. Journal for Nature](#)
771 [Conservation, 53, 125748. <https://doi.org/10.1016/j.jnc.2019.125748>](#)
772

773 Richard G., Vacquié-Garcia J., Jouma'a J., Picard B., Génin A., Arnould J.P.Y., et al. (2014).
774 Variation in body condition during the post-moult foraging trip of southern elephant seals and its
775 consequences on diving behaviour. J Exp Biol. 2014;217: 2609–2619. pmid:24803471
776

777 Richard G., Cox S.L., Picard B., Vacquié-Garcia J., Guinet C. (2016) Southern Elephant Seals
778 Replenish Their Lipid Reserves at Different Rates According to Foraging Habitat. PLoS ONE
779 11(11): e0166747. doi:10.1371/journal.pone.0166747
780

781 Rivas, A.L., Dogliotti A.I., Gagliardini D.A. (2006). Seasonal variability in satellite-measured
782 surface chlorophyll in the Patagonian Shelf; Pergamon-Elsevier Science Ltd; Continental Shelf
783 Research; 26; 6; 4-2006; 703-720
784

785 Romero S.I., Piola A.R., Charo M., and Garcia C.A. E. (2006). Chlorophyll-a variability off
786 Patagonia based on SeaWiFS data. *Journal of Geophysical Research*, 111, C05021.
787 <https://doi.org/10.1029/2005JC003244>
788

789 Roquet F., Charrassin J.-B., Marchand S., Boehme L., Fedak M., Reverdin G., and Guinet C.
790 (2011). Validating hydrographic data obtained from seal-borne satellite-relayed data loggers. *J.*
791 *Atmosph. And Ocean. Tech.*, 28:787-801. doi: 10.1175/2010JTECHO801.1
792

793 Santini M.F., Souza R.B., Wainer I., Muelbert M.M. (2018). Temporal analysis of water masses and
794 sea ice formation rate west of the Antarctic Peninsula in 2008 estimated from southern elephant
795 seals' SRDL–CTD data. *Deep Sea Research Part II: Topical Studies in Oceanography*, Vol. 149,
796 p58-69. <https://doi.org/10.1016/j.dsr2.2018.02.013>
797

798 Saraceno M., Provost C., Piola A.R., Bava J., & Gagliardini A. (2004). Brazil Malvinas Frontal
799 System as seen from 9 years of advanced very high resolution radiometer data. *Journal of*
800 *Geophysical Research C: Oceans*, 109(5). <https://doi.org/10.1029/2003JC002127>
801

802 Saraceno M., Provost C., & Piola A.R. (2005). On the relationship between satellite-retrieved
803 surface temperature fronts and chlorophyll a in the western South Atlantic. *Journal of*
804 *Geophysical Research: Oceans*, 110(11), 1–16
805

806 Saraceno M., Provost C., & Zajaczkovski U. (2009). Long-term variation in the anticyclonic ocean
807 circulation over the Zapiola Rise as observed by satellite altimetry: Evidence of possible
808 collapses. *Deep-Sea Research Part I: Oceanographic Research Papers*, 56(7)
809

810 Schreer J.F., Kovacs K.M., O’Hara Hines R. J. (2001). Comparative diving patterns of pinnipeds
811 and seabirds. *Ecol Monogr.* 2001;71: 137–162
812

813 Siegelman L., Roquet F., Mensah V., Rivière P., Pauthenet E., Picard B., and Guinet C. (2019).
814 Correction and accuracy of high- and low-resolution CTD data from animal-borne instruments.
815 *Journal of Atmospheric and Oceanic Technology.* doi: 10.1175/JTECH-D-18-0170.1
816

817 Stark J.D., Donlon C.J., Martin M.J. and McCulloch M. E. (2007). OSTIA: An operational, high
818 resolution, real time, global sea surface temperature analysis system., *Oceans 07 IEEE Aberdeen,*
819 *conference proceedings. Marine challenges: coastline to deep sea. Aberdeen, Scotland. IEEE.*
820

821 Violante R.A. and Cavallotto J.L. (2011). The record of the LGM sea-level position at the Argentina
822 Continental Shelf: an evidence of the complex processes involved in the post-glacial relative sea
823 level rise. In : *Abstracts of the 5th IGCP 526 Conference ‘Continental Shelves: Risks, Resources*
824 *and Record of the Past’, Victoria (BC), Canada. UNESCO-IGCP, Paris, 128–129*
825

826 Violante R., et al. (2014). Chapter 6 The Argentine continental shelf: morphology, sediments,
827 processes and evolution since the Last Glacial Maximum. 10.1144/M41.6.
828

829 Viviant M., Trites A.W., Rosen D.A. S., Monestiez P., Guinet C. (2010). Prey capture attempts can
830 be detected in Steller sea lions and other marine predators using accelerometers. *Polar Biol* 33:
831 713–719
832
833
834