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Deleterious effects of mercury contamination on immunocompetence, liver function and egg volume in an antarctic seabird

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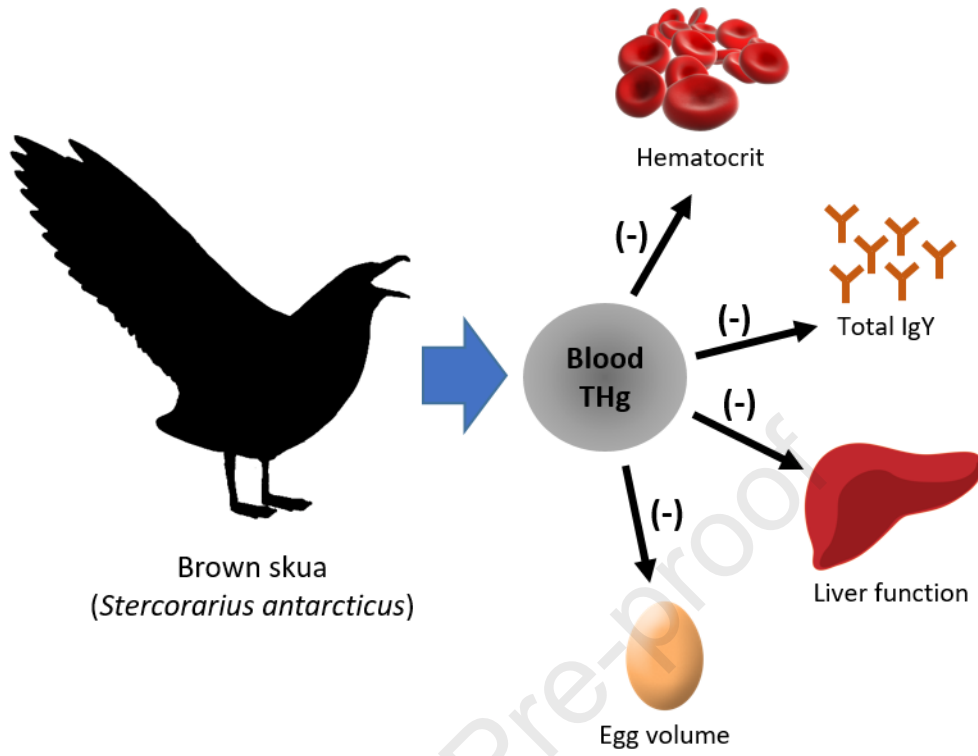
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CReditT authorship contribution statement

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1 **Deleterious effects of mercury contamination on immunocompetence, liver**
2 **function and egg volume in an Antarctic seabird**

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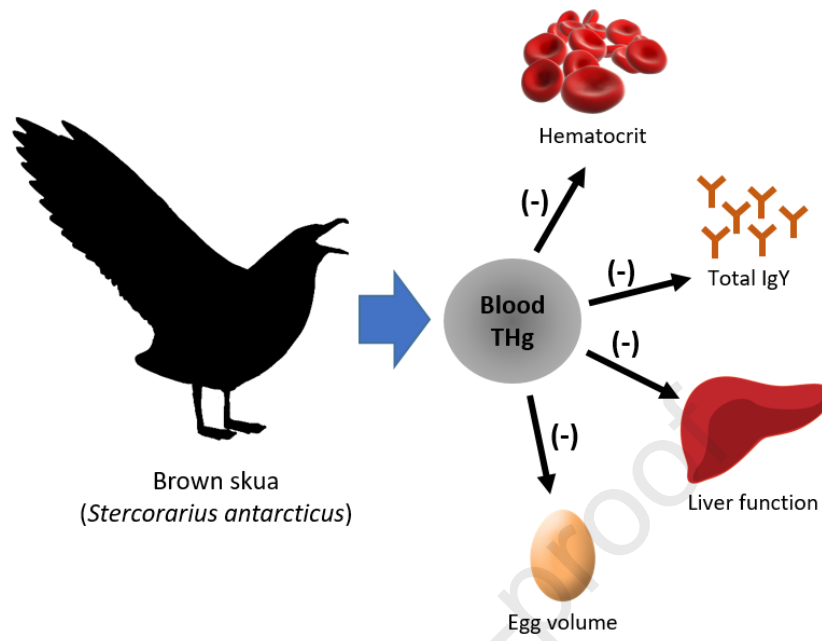
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GRAPHICAL ABSTRACT



25

HIGHLIGHTS

- 26 - Blood THg concentrations were measured in brown skuas on the Antarctic Peninsula.
- 27 - Higher blood THg concentrations had deleterious effects on physiology.
- 28 - Higher blood THg concentrations disrupted immune and liver function.
- 29 - Higher blood THg concentrations were associated with lower egg volume.

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ABSTRACT

31 Mercury (Hg) is a globally important pollutant that can negatively impact metabolic,
32 endocrine and immune systems of marine biota. Seabirds are long-lived marine top
33 predators and hence are at risk of bioaccumulating high Hg concentrations from their
34 prey. Here, we measured blood total mercury (THg) concentrations and relationships with
35 physiology and breeding parameters of breeding brown skuas (*Stercorarius antarcticus*)
36 ($n = 49$ individuals) at Esperanza/Hope Bay, Antarctic Peninsula. Mean blood THg
37 concentrations were similar in males and females despite the differences in body size and
38 breeding roles, but differed between study years. Immune markers (hematocrit,
39 Immunoglobulin Y [IgY] and albumin) were negatively correlated with blood THg
40 concentrations, which likely indicates a disruptive effect of Hg on immunity. Alanine
41 aminotransferase (GPT) activity, reflecting liver dysfunction, was positively associated
42 with blood THg. Additionally, triacylglycerol and albumin differed between our study
43 years, but did not correlate with Hg levels, and so were more likely to reflect changes in
44 diet and nutritional status rather than Hg contamination. Egg volume correlated
45 negatively with blood THg concentrations. Our study provides new insights into the
46 sublethal effects of Hg contamination on immunity, liver function and breeding
47 parameters in seabirds. In this Antarctic species, exposure to sublethal Hg concentrations
48 reflects the short-term risks which could make individuals more susceptible to
49 environmental stressors, including ongoing climatic changes.

50

51 **Keywords:** pollution; mercury; seabirds; physiology; reproduction; Antarctica

52

1. INTRODUCTION

53 Mercury (Hg) is a global environmental pollutant that can have deleterious consequences
54 for humans and wildlife (Corsolini 2009; Tan et al., 2009; Driscoll et al., 2013). Although
55 Hg occurs naturally, its availability has increased in marine and terrestrial environments
56 in different organic and inorganic forms (MeHg, inorganic Hg, etc) (Aronson et al., 2011).
57 The gaseous, elemental form of Hg (Hg^0) can spread through atmospheric transport from
58 emission sources to distant regions (Calle et al., 2015). Once deposited in marine
59 environments, the inorganic form of Hg (Hg^{II}) is methylated by microorganisms to the
60 more toxic form, methyl-Hg (MeHg, $[\text{CH}_3\text{Hg}]^+$). MeHg bioaccumulates within the tissues
61 of marine organisms and biomagnifies through marine food webs (Bargagli 2008;
62 Driscoll et al., 2013; Ibañez et al., 2022a; Matias et al., 2022). Seabirds, which are
63 typically long-lived and feed at high trophic positions, are potentially at risk of
64 accumulating high Hg levels in their tissues from dietary exposure (Bearhop et al., 2000;
65 Bargagli 2008; Tavares et al., 2013; McKenzie et al., 2021; Mills et al., 2020; 2022;
66 Ibañez et al., 2022a).

67 Hg in Antarctic ecosystems derives from natural and human sources, and the Antarctic
68 continent constitutes a sink for Hg, which condensates in colder regions after evaporation
69 and long-range transportation from lower latitudes (Angot et al., 2016). Artisanal and
70 small-scale gold mining, which releases large quantities of Hg into the environment, is
71 mostly concentrated in the Southern Hemisphere (Keane et al., 2023). There are also local
72 sources of Hg, including volcanic activity and the release of Hg stored in sea ice which
73 increase its bioavailability for microbial methylation (Cossa 2013; de Ferro et al., 2014;
74 Gionfriddo et al., 2016). Human activity that is associated with nearby research stations
75 on the Antarctic Peninsula increases during spring and summer, which results in the

76 release of pollutants (heavy metals and organic compounds) from waste-disposal sites,
77 construction materials and compounds used for treating effluent (Acero et al., 1999).

78 Hg is a neurological, immune and endocrine disruptor (Tartu et al., 2013; Whitney and
79 Cristol 2017), and, ultimately, can Hg contamination can have short- or long-term fitness
80 consequences for seabirds (Bustnes et al., 2007; Roos et al., 2012; Dietz et al., 2019;
81 Chételat et al., 2020; Mills et al., 2020; Goutte et al., 2014a, 2014b). Hg contamination
82 may also negatively impact body condition (Tan et al., 2009; Ackerman et al., 2016;
83 Chételat et al., 2020), although, in the wild, body condition indices are unreliable
84 indicators of Hg sublethal effects (Carravieri et al., 2022). Although many studies have
85 examined inter- and intraspecific variation in Hg contamination of seabirds in the
86 Southern Ocean (Carravieri et al., 2014a; 2016; 2017; Mills et al., 2020; 2022; Quillfledt
87 et al., 2023), few have tested for relationships with fitness parameters. Indeed, these have
88 only been investigated for the wandering albatross (*Diomedea exulans*) (Tavares et al.,
89 2013; Carravieri et al., 2014b; 2014c; Goutte et al., 2014a; Bustamante et al., 2016),
90 Antarctic petrel (*Thalassoica antarctica*) (Carravieri et al., 2018; 2021), grey-headed
91 albatross (*Thalassarche chrysostoma*) (Mills et al., 2020), and brown skua (*Stercorarius*
92 *antarcticus*) (Ibañez et al., 2022a) and south polar skua (*S. maccormicki*) (Goutte et al.,
93 2014b).

94 Brown skuas breed on the Antarctic continent and sub-Antarctic islands and are
95 opportunistic predators that feed on a wide diversity of prey in both terrestrial and marine
96 environments (Reinhardt et al., 2000; Phillips et al., 2004; Ritz et al., 2008; Carneiro et
97 al., 2015; Graña Grilli and Montalti 2015; Borghello et al., 2019; Ibañez et al., 2022b).
98 As the brown skua is a migratory seabird, birds are exposed to pollutants not just during
99 the breeding season but also in the nonbreeding season when they visit regions with

100 higher anthropogenic pressure (Albert et al., 2022). At Bahía Esperanza/Hope Bay–
101 located on the Antarctic Peninsula–brown skuas breed close to large colonies of Adélie
102 penguins (*Pygoscelis adeliae*) and gentoo penguins (*Pygoscelis papua*). They feed mainly
103 on penguins and to a lesser extent on marine prey (e.g., fishes and invertebrates)
104 (Borghello et al., 2019; Ibañez et al., 2022b). Macro-plastics have been found recently in
105 skua diet samples collected during the breeding season (Ibañez et al., 2020), which may
106 represent a possible route for chemical pollutants such as Hg (Hamilton et al., 2023).
107 Overall, there has been little research concerning the physiological effects of Hg
108 contamination on Antarctic seabirds (Carravieri et al., 2021; Goutte et al., 2014), and
109 more studies are needed to understand the potential sublethal effects. In the present study,
110 we focus on the sublethal effects of Hg contamination of brown skuas. Total Hg (the sum
111 of inorganic and organic Hg) concentration was measured in red blood cells of brown
112 skuas at Esperanza/Hope Bay. Our aims were to: (i) compare blood THg concentrations
113 between two breeding seasons and sexes; and (ii) relate blood THg concentrations to
114 markers of energy metabolism, immunocompetence and liver function, and to egg
115 volume.

116

117 **2. MATERIALS AND METHODS**

118 *2.1. Fieldwork and sample collection*

119 Fieldwork for this study was conducted at Bahía Esperanza/Hope Bay, Antarctic
120 Peninsula (63°24'S, 57°01'W) (Fig. 1), from November to January during the 2018/19
121 and 2019/20 breeding seasons. Brown skuas were sampled during the early incubation
122 stage (5–10 days after clutch completion). Blood samples (~2 ml) were obtained from the
123 brachial vein using a 25-G needles ($n = 49$, $n = 24$ and $n = 25$ in 2018/19 and 2019/20,
124 respectively). Sampled birds included both individuals from 16 nests and one adult from

125 17 nests. Once in the laboratory (within 2–6 hours after extraction) serum and red blood
126 cells were separated by centrifugation (2000 rpm for 10 min), placed in sterile plastic
127 eppendorf tubes and were stored frozen (-20°C) prior to laboratory analyses.

128 Eggs were measured (length and breadth) using digital calipers and volumes (mm^3)
129 calculated as $0.00048 \times \text{length (mm)} \times \text{breadth (mm)}^2$ (Phillips et al., 2004). We calculated
130 the total clutch volume as the average of the volume of both eggs. The sex of birds was
131 initially assigned morphologically based on body size, and later confirmed by DNA
132 analysis (Fridolfsson and Ellegren 1999; Phillips et al., 2002).

133

134

2.2. Total Hg analysis

135 Studies have demonstrated that Hg in blood is associated predominantly with the cellular
136 fraction (i.e., red blood cells) rather than plasma (Bond and Robertson 2015; Renedo et
137 al., 2018). THg in seabird RBCs is mostly ($>90\%$) MeHg (Renedo et al., 2018; Albert et
138 al., 2019). Prior to analysis, red blood cells were freeze-dried and homogenized. Blood
139 THg concentrations were measured using an Advanced Mercury Analyser
140 spectrophotometer (Altec AMA 254) (LIENSs, France). For each sample, a minimum of
141 two aliquots (range: 1.02–1.86 mg dry weight [dw]) were analyzed, and the means and
142 relative standard deviation (RSD) among measurements were calculated (all samples
143 RSD $<10\%$). THg concentrations are presented in $\mu\text{g g}^{-1}$ dw. Accuracy was tested using
144 certified reference material (CRM; dogfish liver DOLT-5, NRC, Canada; certified Hg
145 concentration: $0.44 \pm 0.18 \mu\text{g g}^{-1}$ dw) every 10 samples. Recovery of the CRM was 97.8
146 $\pm 1.7\%$. Blanks were analyzed at the beginning of each set of samples. The limit of
147 quantification of the AMA was 0.1 ng and the detection limit of the method was $0.005 \mu\text{g}$
148 g^{-1} dw.

149

150

2.3. Hematological determinations

151 Serum concentrations of five energy metabolism markers (including total proteins, uric
152 acid, triacylglycerol, cholesterol and glucose), albumin, aspartate aminotransferase
153 (GOT) and alanine aminotransferase (GPT) enzymes were measured in each bird using
154 colorimetric commercial kits (Wiener Lab). All assays were conducted using an
155 automatic analyzer (Ibañez et al., 2015).

156 Serum circulating levels of Immunoglobulin Y (IgY) were determined by direct ELISA
157 using peroxidase conjugated anti-chicken IgY antibodies (Sigma, St Louis, MO, USA,
158 A-9046) (Martínez et al., 2003; Ibañez et al., 2018). For this, 96-well microtiter plates
159 (Nunc PolySorp; Nunc, Roskilde, Denmark) were coated during 1 h at 37 °C with serum
160 samples diluted (1/30,000) in 0.1 M carbonate-bicarbonate buffer (pH = 9.6). Then the
161 plates were washed three times with PBS- 0.05% Tween 20 and incubated with 1% non-
162 fat milk (Nestlé coffee-mate) in PBS-Tween-20 during 1 h at 37 °C to block the free
163 binding sites. After new washing, the wells were incubated with peroxidase- conjugated
164 anti-chicken IgY. Finally, the wells were washed and ABTS [2,2-azino-di (3-
165 ethylbenzthiazoline sulfonate)] was added as substrate. After incubating for 30 min (at
166 room temperature) color development was stopped with oxalic acid 2% and then read as
167 optical density (OD) at 405 nm.

168 To determine the hematocrit value a heparinized capillary was filled in the laboratory
169 with 100 µl of blood from the Eppendorf tube that contained heparinized blood obtained
170 in the field. The capillary was then centrifuged at 5000 rpm for 15 min (Ibañez et al.,
171 2015), and a digital caliper used to measure the length (mm) of the red blood cell fraction

172 and the total blood volume. Hematocrit values are presented as a percentage of total
173 volume.

174

175 *2.4. Statistical analysis*

176 Data were analysed using R (R Core Team 2015). Blood THg concentrations,
177 physiological parameters and egg volumes were checked for normality and homogeneity
178 of variances using Shapiro-Wilk and Levene's tests, respectively. Differences in blood
179 THg between sexes and seasons were tested using general linear mixed models (GLMMs;
180 Gaussian distribution and identity link function), with the individual identity and the
181 breeding pair included as random effects to account for partners potentially being more
182 similar than non-partners in blood THg concentrations. Relationships between
183 physiological markers (total proteins, uric acid, triacylglycerol, cholesterol, hematocrit,
184 albumin, IgY, GOT and GPT), egg volume and blood THg concentrations were also
185 tested using GLMMs, with sex (except in the case of egg volume) and season included as
186 covariates, and individual identity included as a random effect. All GLMMs were fitted
187 using the "nlme" package in R (Pinheiro et al. 2017).

188

189 **3. RESULTS**

190 *3.1. Sex and annual variation in Hg contamination*

191 Detectable blood THg concentrations were found in all samples from brown skuas at
192 Bahía Esperanza/Hope Bay in 2018/19 (mean \pm SD, $0.73 \pm 0.22 \mu\text{g g}^{-1}$ dw) and 2019/20
193 ($0.91 \pm 0.44 \mu\text{g g}^{-1}$ dw) (Table 1). There were no significant differences in blood THg
194 concentrations (log-transformed) between sexes (est = 0.23, p = 0.48) or seasons (est =
195 0.19, p = 0.56) (Table 1).

196

197 3.2. *Physiological markers and egg volume*

198 Physiological parameters, in particular immune and hepatic functions, did not differ
199 between sexes (all $p > 0.05$), but were negatively associated with blood THg
200 concentrations. Significant negative relationships were found between albumin (est = -
201 0.31, $p < 0.0001$), hematocrit (%) (est = -9.27, $p < 0.0001$) and IgY (est = -0.06, $p < 0.05$)
202 and blood THg concentrations (Fig. 2; Table 2). A significant positive relationship was
203 found between GPT activity and blood THg concentrations (est = 13.10, $p < 0.01$) (Fig.
204 2; Table 2). Mean albumin levels differed significantly between seasons (est = 0.31, $p <$
205 0.0001). Triacylglycerol levels also varied annually (est = 55.89, $p < 0.001$) and were
206 unrelated to blood THg concentrations (est = -11.04, $p = 0.55$) (Table 2). Uric acid, total
207 proteins, cholesterol, glucose and GOT show no relationship with sex, season or blood
208 THg (Table 2). Egg volume (mm^3) showed a significant negative relationship with blood
209 THg concentrations (est = -16.40, $p < 0.05$) (Fig. 3), and did not differ significantly
210 between seasons (est = 4.25, $p = 0.19$).

211

212

4. DISCUSSION

213 In our study, blood THg concentrations of brown skuas ranged from 0.41 to 2.33 $\mu\text{g g}^{-1}$
214 dw (equivalent to 0.10 to 0.58 $\mu\text{g g}^{-1}$ wet weight [ww], assuming a 79% moisture content;
215 Eagles-Smith et al., 2008; Ackerman et al., 2016). These concentrations are comparable
216 to those found in adults of other Antarctic seabirds, including snow petrels (*Pagodroma*
217 *nivea*) (Tartu et al., 2015), and Antarctic petrels (*Thalassoica antarctica*) (Carravieri et
218 al., 2021). However, they were lower than those reported in skuas at lower latitude
219 breeding colonies in the southwest Atlantic Ocean sector of the Southern Ocean (Mills et
220 al., 2022), but were comparable to those associated with reduced breeding success in
221 south polar skuas at Adélie Land (Goutte et al., 2014b). Despite the low values of blood

222 THg measured here, we found significant negative effects on immunity and hepatic
223 enzymes, and on breeding parameters.

224

225 4.1. *Impacts of Hg contamination on immunocompetence*

226 Hg is often associated with immunosuppressive effects at sublethal levels, but mostly in
227 captive studies (Fallacara et al., 2011; Kenow et al., 2007; Lewis et al., 2013a). Despite
228 the relatively low concentrations (initially suggesting a low risk of MeHg toxicity;
229 Ackerman et al., 2016), IgY and hematocrit were negatively related to blood THg
230 concentrations in our study (Fig. 2). These results indicate a negative impact of Hg
231 contamination on the immune status of brown skuas. Our results agree with previous
232 studies on captive zebra finches (*Taeniopygia guttata*) that have shown negative effects
233 of Hg exposure on B-cell proliferation (Lewis 2012; Lewis et al., 2013). Also, impaired
234 macrophage phagocytosis was related to high Hg levels in black-footed albatrosses
235 (*Phoebastria nigripes*) in the North Pacific (Finkelstein et al., 2007). Hematocrit values
236 decreased with blood THg concentrations, which agrees with the hemolytic and anemia-
237 inducing effects of Hg (Zolla et al., 1997). Erythrocytes are an important target of Hg and
238 the majority of Hg in blood is found in the cellular fraction (~90%) (Bond and Robertson
239 2015). *In vitro* studies have demonstrated that the exposure of erythrocytes to low
240 concentrations of Hg induce structural changes in the external surface of the membrane.
241 These changes are mediated by the translocation of phosphatidylserine to the external
242 surface of the erythrocyte cell membrane as a signal that may prompt cellular apoptosis
243 (Eisele et al., 2006; Lim et al., 2010). Another possible explanation behind the decrease
244 in hematocrit may be an association with lowered renal function, also linked to hemolytic
245 processes (Rivarob et al., 1983; Chitra et al., 2013).

246 Serum IgY and albumin are useful indicators of health in birds (Lumeij 1987; Ibañez et
247 al., 2018). Fitness traits are mediated by hormones including luteinizing hormone, which
248 is a pituitary hormone involved in the onset of breeding (Dawson et al., 2001);
249 corticosterone, an adrenal hormone in the stress response (Wingfield and Sapolsky 2003);
250 and prolactin, a pituitary hormone involved in the expression of parental care (Angelier
251 and Chastel 2009). Trace metal pollution may have different effects on hormones of the
252 hypothalamic–pituitary–adrenal (HPA) axis, such as corticosterone (Tan et al., 2009;
253 Tartu et al., 2013). Serum IgY and albumin concentrations were negatively related to
254 blood THg concentrations of brown skuas in our study (Fig. 2). One possible explanation
255 for this may be increasing immunosuppressive effects and catabolic activity associated
256 with endocrine disruption of the HPA axis (Coutinho and Chapman 2011), as Hg
257 accumulates in the pituitary gland and thyroid in vertebrates (Colborn et al., 1993; Tan et
258 al., 2009; Meyer et al., 2014; Tartu et al., 2013). In this scenario, the levels of Hg found
259 in brown skuas at Esperanza/Hope Bay may induce protein catabolism and
260 immunosuppression (IgY and albumin decline) because corticosterone release is
261 disrupted (Ibañez et al., 2018). Another possible explanation is impaired protein synthesis
262 in the liver. However, these hypotheses would need to be confirmed in future studies on
263 skuas, particularly as a Hg-induced deterioration in immunocompetence is likely to affect
264 disease risk and, ultimately, population dynamics.

265

266 *4.2. Impacts of Hg contamination on liver function*

267 GPT was positively related to blood THg concentrations of brown skuas. This suggests
268 that Hg contamination impacted the liver function of brown skuas. The liver is the major
269 organ involved in the biotransformation, metabolism, protein synthesis and detoxification
270 processes, which also reflect its susceptibility to pollutants, leading to tissue damage.

271 GOT and GPT enzymes are considered to be biochemical markers of impaired liver
272 function. Liver-cell damage, such as degeneration and necrosis, may increase GOT and
273 GPT levels (Gowda et al., 2009; Mari et al., 2010; Ibañez et al., 2015; Yang et al., 2015;
274 Choi et al., 2017). The biological mechanism of association between Hg exposure and
275 liver dysfunction is mainly explained by oxidative stress, cell death, and impaired
276 metabolism (Malhi et al., 2010). For instance, in male rodents exposed to Hg, levels of
277 GOT, GPT, and gamma glutamyl-transferase activities were elevated, and tissue damage
278 or necrotic changes observed in most livers (Waddam 2009). Also, histological analyses
279 described degenerative changes and lysed areas in liver parenchyma in Hg-exposed
280 zebrafish (Maricella et al., 2016). Hg exposure in zebrafish induced deregulation of
281 oxidative stress, intrinsic apoptotic pathways, and resulting hepatotoxicity through cell
282 death, mitochondrial dysfunction, endocrine disruption, and metabolic disorders (Ung et
283 al., 2010; Maricella et al., 2016). In the spectacled caiman (*Caiman crocodilus*), negative
284 associations between Hg and alkaline phosphatase activity (a liver cytoplasmic enzyme
285 involved in the hepatocytic functions) were observed (Lemaire et al., 2018). Therefore,
286 our results suggest that variation in GPT activity in response to elevated blood Hg levels,
287 may indicate hepatotoxicity even at the low levels of contamination found in Antarctica.

288

289 4.3. Annual variations in Hg contamination and association with physiology

290 In a previous study of brown skuas at Hope Bay, we found annual variation in blood THg
291 concentrations was related to trophic ecology, but had no impact on adult body condition
292 (Ibañez et al., 2022a). Blood THg concentrations were higher in 2019/20 than 2018/19
293 (by $\sim 0.2 \mu\text{g g}^{-1} \text{ dw}$) (Ibañez et al., 2022a). This was potentially due to changes in diet
294 composition, and the consumption of more contaminated prey (Braune et al., 2014). Also,
295 annual fluctuations in environmental conditions may influence Hg transport, MeHg

296 production, and bioavailability to marine predators and their prey (Cossa et al., 2011;
297 Driscoll et al., 2013; Renedo et al., 2020). In the current study, mean albumin and
298 triacylglycerol levels differed between years, but were not linked directly to blood Hg
299 contamination. Albumin and triacylglycerol are markers of nutritional status (Ibañez et
300 al., 2018). In this scenario, annual dietary differences may affect nutritional status. This
301 may in turn moderate the effects of Hg on metabolism, with potentially limited food
302 resources (thus lower Hg levels) associated with lower albumins and triacylglycerols.

303

304 4.4. Impacts of Hg contamination on egg volume

305 Hg contamination of seabirds may induce changes on body condition that ultimately
306 affect breeding fitness and survival (Evers et al., 2008; Labocha and Hayes 2012;
307 Ackerman et al., 2016). Effects of Hg on reproduction can be reflected at different levels
308 including egg neglect (Tartu et al., 2015), lower breeding success (Tartu et al., 2016),
309 reproductive failure (Mills et al., 2020), or population dynamics (Goutte et al., 2014b,
310 Goutte et al., 2015). However, the effects of Hg contamination during the non-breeding
311 season on subsequent breeding success are poorly known. In little auks (*Alle alle*), Fort
312 et al. (2014) suggested a carry-over effect, in that individuals with the highest Hg
313 concentrations laid smaller eggs. In the present study, egg volume was negatively
314 associated with female blood THg concentrations (Fig. 3). Although not significant, THg
315 concentrations were slightly higher in males than females in both seasons (Table 1; Mills
316 et al., 2022; Ibañez et al., 2022a), possibly because egg production provides a route
317 through which females are able to eliminate Hg (Robinson et al. 2012; Ackerman et al.
318 2020). An explanation for the association of smaller eggs laid with higher blood THg
319 concentrations of brown skuas relates to the trophic ecology and Hg contamination during
320 female pre-laying exodus or differences in prey consumption prior to sampling.

321 Therefore, our results support a previous study which also suggested carry-over effects
322 of Hg on the reproduction of great skuas (*Stercorarius skua*) with a specific influence of
323 female winter distribution and Hg contamination on egg volume (Albert et al., 2022).
324 However, this would need to be confirmed in future studies by measuring THg in feathers
325 grown during the nonbreeding season to infer the degree of Hg contamination since the
326 last moult (Fort et al., 2014).

327 The threshold of Hg toxicity in seabirds appears to be related to the latitude of the study
328 site. Toxicity appears to differ in Antarctic compared with lower latitudes such as
329 subantarctic or subtropical environments (Goutte et al., 2014b; Carravieri et al., 2021).
330 Blood THg concentrations of brown skuas in our study would initially suggest a low risk
331 of MeHg toxicity (Ackerman et al., 2016). These sublethal THg concentrations had
332 deleterious effects on physiology and egg volume, but there was no relationship with
333 breeding success (Ibañez et al., 2022a). Goutte et al. (2014b) reported short-term effects
334 of Hg on breeding success in brown skuas and south polar skuas, though concentrations
335 were higher than in brown skuas at our study site. Selenium (Se)-Hg interactions are often
336 observed in the blood and internal tissues of marine predators, and Se has a protective
337 effect against Hg toxicity when Se is in molar excess (Carravieri et al., 2017; 2020;
338 Manceau et al., 2021). Our results may also be explained by the presence of Se, as was
339 reported for brown skua chicks from the Southern Ocean (Carravieri et al., 2017), south
340 polar skuas (Goutte et al., 2014b) and the spectacled caiman (Lemaire et al., 2018). If so,
341 higher concentrations of Se at Antarctic latitudes may reduce the negative effects of Hg
342 on physiology and reproduction; however, this hypothesis requires further investigation.

343

5. CONCLUSIONS

344 Our study demonstrated negative effects of Hg contamination on physiology and breeding
345 parameters in brown skuas on the Antarctic Peninsula. Despite low blood THg
346 concentrations, which are below or similar to those of other Antarctic seabirds, the
347 association between Hg and the physiological and breeding parameters are of concern.
348 The detrimental impact on egg volume highlights the importance of investigating the
349 relationships between blood THg and the hormones that play a role in stress responses
350 and reproductive decisions, as well as, Hg concentration in chicks during development.
351 This is particularly as skuas and other predators in the Antarctic may become more
352 susceptible to pollutants and other environmental stressors, given the evidence for rapid,
353 ongoing climatic changes in the region.

354

355 **CRediT authorship contribution statement**

356 **Andrés E Ibañez:** Conceptualization, Methodology, Investigation, Resources, Formal
357 Analysis, Visualization, Writing – Original Draft. **William F Mills:** Methodology,
358 Investigation, Resources, Writing – Review & Editing. **Paco Bustamante:** Investigation,
359 Writing – Review & Editing. **Lara M Morales:** Investigation, Writing – Review &
360 Editing. **Diego S Torres:** Investigation, Writing – Review & Editing. **Beatriz D´Astek:**
361 Investigation, Resources. **Rocío Mariano-Jelicich:** Investigation, Writing – Review &
362 Editing. **Richard A Phillips:** Conceptualization, Methodology, Investigation, Resources,
363 Supervision, Writing – Review & Editing. **Diego Montalti:** Investigation, Resources,
364 Supervision, Writing – Review & Editing

365

366 **Conflict of interest**

367 The authors declare that they have no known competing financial interests or personal
368 relationships that could have appeared to influence the work reported in this paper.

369

370 Ethical approval

371 All applicable international, national, and institutional guidelines for sampling, care and
372 experimental use of animals for the study were followed as established by the Article III,
373 Annex II of the Madrid Protocol, Law 24.216 (Taking, Harmful Intrusion and
374 Introduction of Species) within the framework of the projects evaluated and approved by
375 the Environment Office of the IAA and Dirección Nacional del Antártico (DNA) (permits
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380

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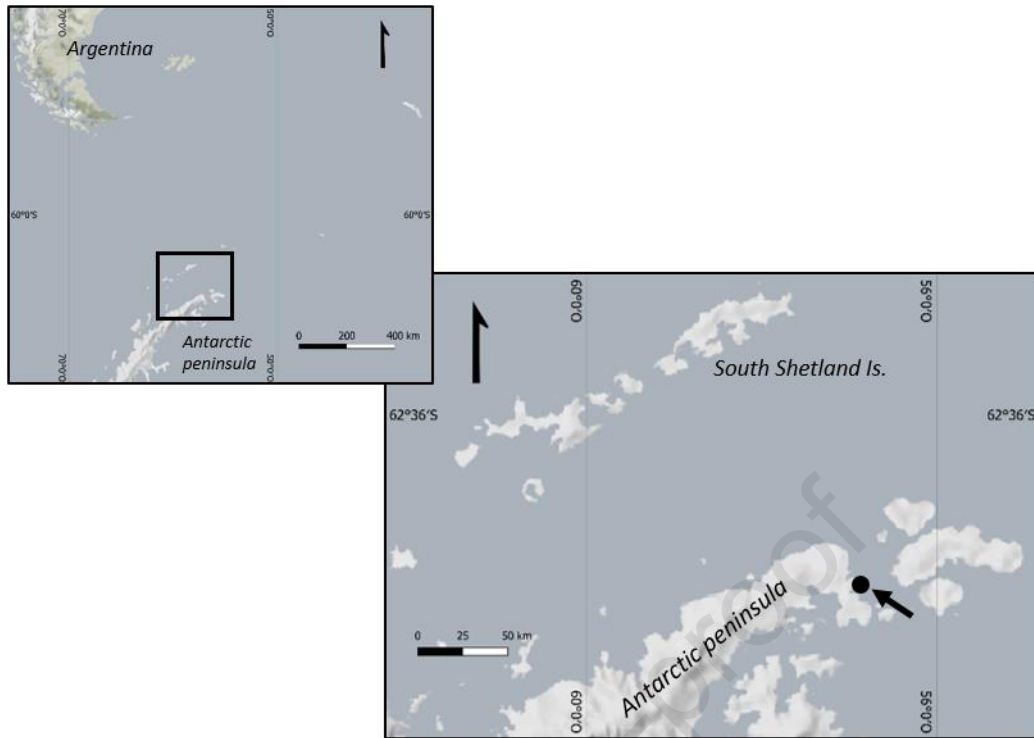


Figure 1. Location of the study site (black circle), Esperanza/Hope Bay, Antarctic Peninsula ($63^{\circ}24' S$, $57^{\circ}01' W$).

Table 1. Total mercury (THg) concentrations ($\mu\text{g g}^{-1}$ dw) in red blood cells of adult male and female brown skuas (*Stercorarius antarcticus*) at Esperanza/Hope Bay, Antarctic Peninsula ($63^{\circ}24'S$, $57^{\circ}01'W$), in the 2018/19 and 2019/20 breeding seasons. Data are means \pm SDs.

Year	<i>N</i>	Sex	THg ($\mu\text{g g}^{-1}$ dw)	Range ($\mu\text{g g}^{-1}$ dw)
2018-2019	11	F	0.66 ± 0.25	0.41 - 1.21
	13	M	0.79 ± 0.18	0.48 - 1.06
	24	Both	0.73 ± 0.22	0.41 - 1.21
2019-2020	13	F	0.82 ± 0.39	0.49 - 1.94
	12	M	1.02 ± 0.50	0.44 - 2.33
	25	Both	0.91 ± 0.44	0.44 - 2.33

Table 2. Serum hematological markers of adult brown skuas (*Stercorarius antarcticus*) at Esperanza/Hope Bay, Antarctic Peninsula (63°24'S, 57°01'W), in the 2018/19 and 2019/20 breeding seasons. Data are means \pm SDs. Uric acid (mg/dl), triacylglycerol (mg/dl), total proteins (d/dl), cholesterol (mg/dl), glucose (mg/dl), albumin (g/dl), GOT (U/l) and GPT (U/l). Serum hematological markers that showed an association with season are indicated with ^a ($p < 0.05$).

Parameter	Season 2018-2019		Season 2019-2020	
	Mean \pm SD	Min-Max	Mean \pm SD	Min-Max
Uric acid	8.126 \pm 2.776	3.87 - 14.37	6.234 \pm 2.069	2.250 - 9.56
Triacylglycerols	63.41 \pm 18.40	31 - 99	118.1 \pm 49.76	53 - 286
Total proteins	2.809 \pm 0.392	2.1 - 3.7	2.790 \pm 0.497	1.9 - 3.5
Cholesterol	245 \pm 58.06	103.6 - 354.9	244.8 \pm 65.63	113.3 - 351.5
Glucose	340.6 \pm 43.41	269 - 420	330.6 \pm 49.95	170 - 379
Albumin	1.020 \pm 0.135	0.73 - 1.260	1.282 \pm 0.233	0.87 - 1.63
GOT	75.95 \pm 27.20	25 - 160	105.4 \pm 20.79	69 - 153
GPT	27.70 \pm 12.84	9.1 - 63	27.27 \pm 7.192	15 - 39
Hematocrit (%)	45.60 \pm 4.967	35.6 - 53.5	41.33 \pm 6.204	30.10 - 30.90
IgY (OD 405nm)	0.215 \pm 0.069	0.11 - 0.38	0.237 \pm 0.075	0.104 - 0.430

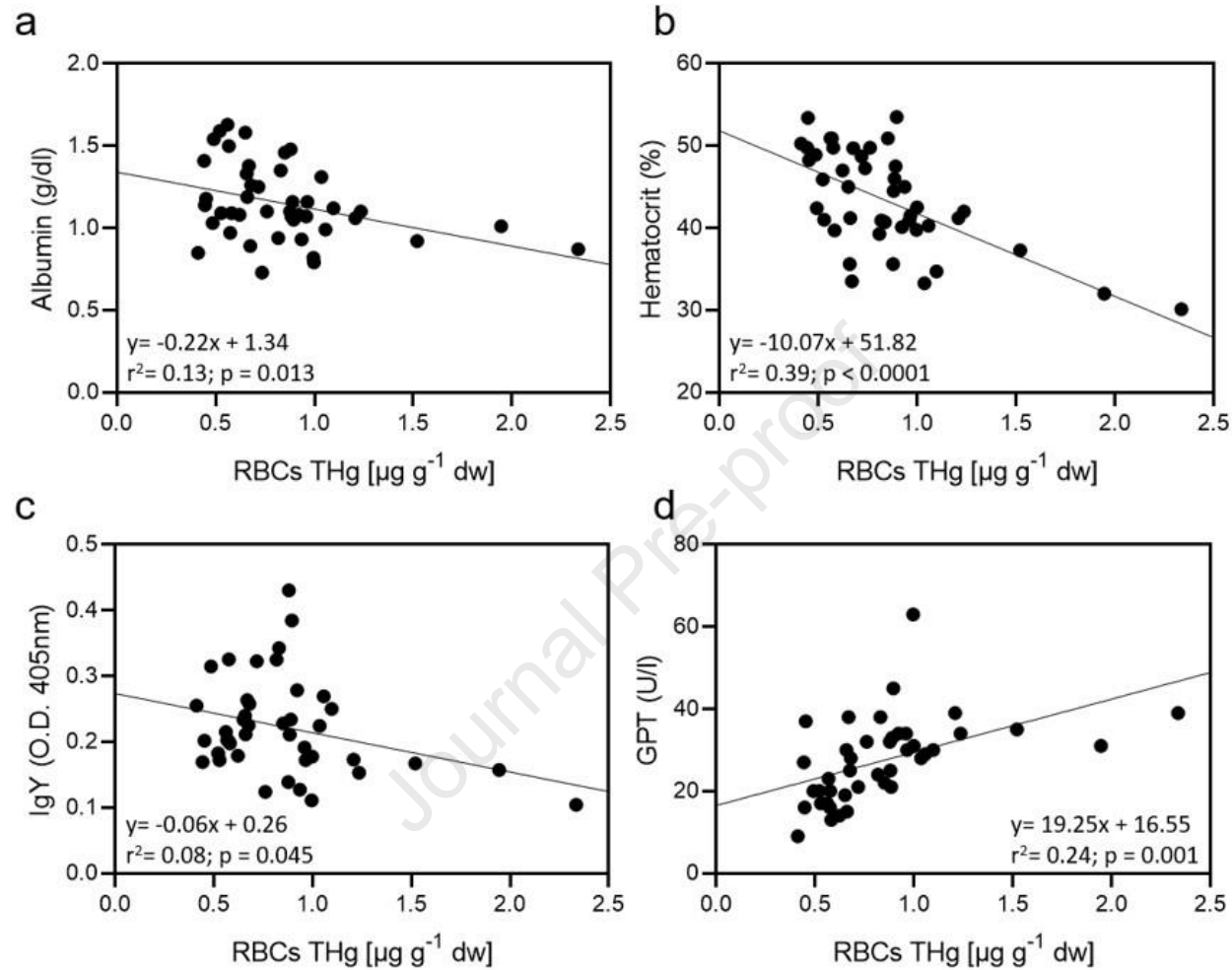


Figure 2. Relationships between red blood cells THg concentrations ($\mu\text{g g}^{-1} \text{ dw}$) and **(a)** albumin (g/dl), **(b)** hematocrit (%), **(c)** IgY (OD 405nm) and **(d)** GPT (U/l) in the blood of adult brown skuas (*Stercorarius antarcticus*) at Esperanza/Hope Bay, Antarctic Peninsula ($63^{\circ}24'S$, $57^{\circ}01'W$) in the 2018/19 and 2019/20 breeding seasons.

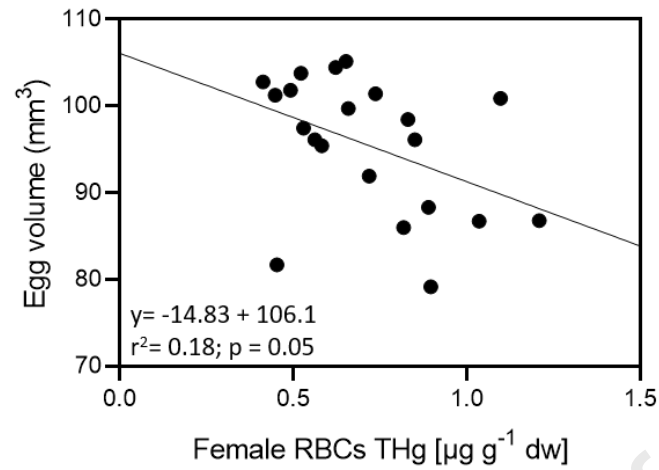


Figure 3. Relationships between red blood cells THg concentrations ($\mu\text{g g}^{-1} \text{dw}$) and egg volume (mm^3) of female brown skuas (*Stercorarius antarcticus*) at Esperanza/Hope Bay, Antarctic Peninsula ($63^{\circ}24'S$, $57^{\circ}01'W$) in the 2018/19 and 2019/20 breeding seasons.

HIGHLIGHTS

- Blood THg concentrations were measured in brown skuas on the Antarctic Peninsula.
- Higher blood THg concentrations had deleterious effects on physiology.
- Higher blood THg concentrations disrupted immune and liver function.
- Higher blood THg concentrations were associated with lower egg volume.

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Declaration of interests

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

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