

2 **Mid-Holocene skua remains from King George Island, Antarctica**

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7 **Abstract** Seven avian remains from Mid-Holocene strata
8 of the southeastern coast of Potter Peninsula (King George
9 Island: Isla 25 de Mayo, South Shetland Islands, Antarc-
10 tica) are reported. They were described and assigned to the
11 brown skua *Stercorarius antarcticus lonnbergi* (Aves,
12 Laridae), a living species currently breeding in the area.
13 The presence of penguins in the same sequence is in
14 agreement with the current dynamics of the coastal eco-
15 systems of Antarctica. Nowadays, penguins and skuas
16 frequent these same environments.

17
18 **Keywords** Skuas · Laridae · Mid-Holocene ·
19 South Shetland Islands · Antarctica

20
21 **Kurzfassung** Aus dem mittleren Holozän der südöstli-
22 chen Küste der Potter-Halbinsel (King George Island: Isla
23 25 de Mayo, South Shetland Islands, Antarktis) werden
24 sieben Rippen beschrieben, die dem Braunen Skua

Stercorarius antarcticus lonnbergi (Aves, Laridae) zuge- 25
ordnet werden. Diese Art kommt auch heute noch in der 26
Region vor. Das gemeinsame Vorkommen von Pinguinen 27
in der selben Fundschicht steht im Einklang mit der der- 28
zeitigen Dynamik der Ökosysteme der antarktischen Küs- 29
ten. Heute leben auch noch Pinguine und Raubmöwen in 30
derselben Umgebungen. 31
32

Schlüsselwörter Raubmöwen · Laridae · Mid Holozän · 33
Süd-Shetland-Inseln · Antarktis 34

Introduction 35

36 Skuas are marine birds related to gulls, waders, auks, and 36
37 skimmers (Charadriiformes, Laridae). They have been 37
38 classically divided into two genera; the smaller species 38
39 were assigned to *Stercorarius* Brisson 1760, while the 39
40 larger species were placed in *Catharacta* Brünnich 1764. 40
41 Proposals based on genetic studies have included all spe- 41
42 cies in the genus *Stercorarius* (Cohen et al. 1997; 42
43 Andersson 1973, 1999). The latter systematic arrangement 43
44 is followed here. 44

45 The fossil record of skuas (Aves, Laridae) is scant; only 45
46 isolated remains have been found (mainly in Europe). 46
47 A tarsometatarsus of *Stercorarius pomarinus* was found in 47
48 the Upper Pleistocene of Spain (Sánchez Marco 2006). 48
49 Remains of this species have also been reported in the 49
50 Upper Pleistocene of the northern coast of Italy (Cassoli 50
51 1980) and the Middle Pleistocene of France (a tarsometat- 51
52 tarsus of fossil subspecies *Stercorarius pomarinus philippi*, 52
53 Mourer-Chauviré 1975). 53

54 Besides the above-mentioned records, at least three 54
55 different species of jaegers and one of skua were recog- 55
56 nized in Lee Creek Mine (North Caroline, USA), assigned 56

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57 probably to Yorktown Formation (Early Pliocene; Olson
58 and Rasmussen 2001). The first one was identified as
59 *Stercorarius aff. pomarinus*, and it is represented by two
60 fragmentary humeri, an ulna, and a coracoid very similar to
61 those of living species. Also, an isolated coracoid from the
62 Pliocene of Florida (USA) was assigned to an undescribed
63 species of *Stercorarius*, slightly larger than *S. pomarinus*
64 (Emslie 1995). The second species was identified as
65 *Stercorarius aff. parasiticus* (two incomplete humeri and a
66 carpometacarpus). The last species of jaeger is *Stercorarius*
67 *aff. longicaudus*, represented by two humeral epiphyses
68 and a carpometacarpus. The remains assigned to *Cathar-*
69 *racta* are a fragmentary ulna, two carpometacarpi, and
70 two tarsometatarsi, the first mention for the Northern
71 Hemisphere (Olson and Rasmussen 2001).

72 The only extinct species is *Stercorarius shufeldti* How-
73 ard 1946, slightly smaller than *Catharacta*, from the
74 Middle Pleistocene of Oregon, USA (Howard 1946). An
75 ulna, a carpometacarpus, and three tarsometatarsi probably
76 assignable to *C. skua hamiltoni* were described from
77 Amsterdam Island (Indian Ocean). Although no radiocar-
78 bon datings were made, Worthy and Jouventin (1999)
79 estimated the age at between a few hundred and a few
80 thousand years old. In addition, *Stercorarius skua* was
81 found in levels of the Holocene Nerja cave (Tyrberg and
82 Hernández 1995).

83 Others fossils of *Stercorarius* correspond to the Miocene
84 and Pliocene of North Carolina, USA and the middle
85 Holocene from the South Shetland Islands, Antarctica
86 (Olson and Rasmussen 2001; del Valle et al. 2002).
87 Finally, some Holocene remains from New Zealand (North
88 and South Islands, Chatham Island) assigned to *Catharacta*
89 *skua lonnbergi* (= *Stercorarius antarcticus lonnbergi*) were
90 mentioned by Fordyce (1982) and compiled by Turbott
91 (1990) and Millener (1991, 1999) and bibliography cited
92 therein.

93 In this context, the goal of the present contribution is the
94 study of new Holocene remains assigned to skuas found in
95 the South Shetland Islands, Antarctica, including their
96 systematic assignment and the descriptions of these bones.
97 Additionally, some other comments concerning the geo-
98 logical and geographical context are made.

99 Materials and methods

100 Almost 3 m thick of fossiliferous sediments were exca-
101 vated and bones were manually collected. Penguin bones
102 were ¹⁴C-dated at Weizmann Institute of Science, Rehovot
103 (Israel). Recovered skua remains were measured and
104 described following the terminology of Baumel (1993).
105 Because the species are quite homogeneous concerning
106 their qualitative characters, measurements have often been

used in systematics (Worthy and Jouventin 1999; Acosta
Hospitaleche et al. 2009). We determined the remains
according to Remsen et al. (2009).

The study area

The Pingfo I locality (62°15'26.483"S, 58°37'08.530"W,
17.3 m a.s.l.) is located on the southeastern coast of Potter
Peninsula, King George Island (Isla 25 de Mayo), South
Shetland Islands, Antarctica (Fig. 1). It is within the
Potter Peninsula Antarctic Specially Protected Area
(ASPA 132).

The exposed depositional sequence shows a monoto-
nous succession of sedimentary facies with high degree
of conformity. It includes skuas, penguins, marine
mammals, and seaweed, distributed in seven beds
between 14.7 and 16.7 m a.s.l. Combined with the
taphonomy of the fossil remains, it allows us to establish
that bed 1 bearing skuas is older in geological terms than
beds 3 and 5 from where radiocarbon-dated penguins
were exhumed (Fig. 2). Reservoir correction ages are ca.
4540 and 4450 years BP for beds 3 and 5, respectively
(see del Valle et al. 2002).

Bird remains recovered from these beds are complete,
non-reworked, and free from erosion. They were probably
accumulated in a marine beach environment without sig-
nificant transport (del Valle et al. 2002). The taphonomical
analysis of the remains suggests that bodies from a nearby
breeding colony were deposited during mid-Holocene
times (Montalti et al. 2009).

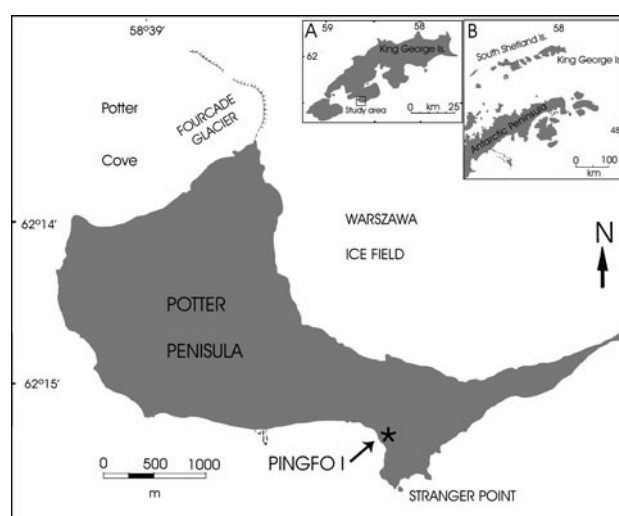


Fig. 1 Location map showing the Pingfo I locality, southeastern coast of Potter Peninsula (King George Island, South Shetland Islands, Antarctica)

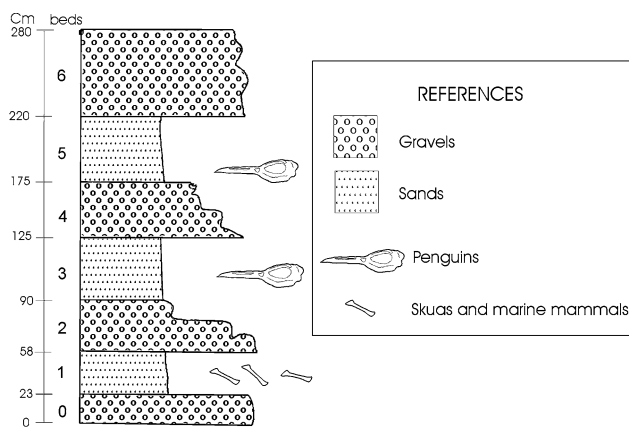


Fig. 2 Sedimentary sequence exposed at Pingfo I locality, Potter Peninsula (King George Island, South Shetlands Islands, Antarctica)

Systematic paleontology

- 135 Family Laridae Rafinesque, 1815
- 136 Genus *Stercorarius* Brisson, 1760
- 137 Species *Stercorarius antarcticus* Lesson, 1831
- 138 Subspecies *Stercorarius antarcticus lonnbergi* Mathews,
- 139 1912

Assigned materials

- 142 Femur (IAA 797), humerus (IAA 798), incomplete tibio-
- 143 tarsus (IAA 800), tarsometatarsus (IAA 799), carpometa-
- 144 carpus (IAA 801), mandible (IAA 802), pelvis (IAA 803).

Description

146 The mandible (Fig. 3a, b) is complete, and both rami
 147 mandibulae are articulated. A lateral deformation was
 148 caused during taphonomic processes (Fig. 3a, b). The
 149 processus mandibulae medialis is robust, and the processi
 150 mandibulae lateralis are small. The fenestra mandibularis
 151 cranialis is open, while the fenestra mandibularis caudalis
 152 is completely obliterated.

153 The humerus (Fig. 3c, d) has a subtriangular fossa
 154 pneumotricipitalis, and it is surrounded by a conspicuous
 155 tuberculum ventrale. Both crus dorsale fossae and crus
 156 ventrale fossae are well developed. The tuberculum dorsale
 157 is elongated and small. The crista deltopectoralis is widely
 158 expanded and its sharpened distal end projects cranially.

159 The processus supracondylaris dorsalis projects crani-
 160 ally and proximally. The epicondylus ventralis is rounded
 161 and slightly marked.

162 The sulcus humerotricipitalis is wide and connects to the
 163 fossa olecrani. The impression coracobrachialis is wide and

shallow, and connects to the narrow and deep sulcus liga- 164
 165 mentaris transversus. The fossa m. brachialis is very
 166 expanded.

The carpometacarpus (Fig. 3e, f) has a trochlea carpalis 167
 168 with a sharp end, and a broad notch joining it to the pro-
 169 cessus extensorius. The os metacarpale alulare is hook-
 170 shaped. The triangular sulcus interosseus is broad and
 171 connected with the sulcus tendinosus that reaches half of
 172 the os metacarpale majus. The fossa infratrochlearis is
 173 rounded.

The pelvis, os coxae, and synsacrum (Fig. 4a, b) are 174
 175 articulated in one piece. Fifteen vertebrae compose the
 176 synsacrum, whose processi transversi are joined so that the
 177 foramina intertransversaria open between them. The crista
 178 spinosi synsacri is uniformly developed along the vertebral
 179 column axis. The ala preacetabularis is notably longer than
 180 the ala postacetabularis. The foramen acetabuli is rounded
 181 and much smaller than the foramen ilioischadicum, which
 182 is anteroposteriorly elongated. They are both divided by a
 183 prominent antitrochanter. The sulcus antitrochantericus is
 184 narrower at its posterior end.

The femur (Fig. 5a, b) has an extended crista trochan- 185
 186 teris and a very well-developed impression iliotrochante-
 187 ricae. The sulcus intercondylaris is symmetrical with
 188 respect to both distal condyles. The epicondylus medialis
 189 and lateralis are weak. The trochanter femoris is small, and
 190 the fossa trochanteris is minimally developed. The crista
 191 supracondylaris medialis is weak and limits a fossa popli-
 192 tea of irregular perimeter.

The tibiotarsus (Fig. 5c, d) belongs to a subadult spec- 193
 194 imen, as the proximal end is not fully developed.

The sulcus extensorius is broad, and its lateral and medial 195
 196 limits are represented by poorly defined crests. The pons
 197 supratendinosus is broad and runs obliquely to the axis of the
 198 diaphysis. The epicondylus medialis is not conspicuous, but
 199 more developed than the epicondylus lateralis.

The tarsometatarsus (Fig. 5e, f) has a sturdy and rounded 200
 201 eminentia intercotylaris aligned with the trochlea meta-
 202 tarsi III. The cotyla medialis is displaced medially towards
 203 the trochlea metatarsi II, while the cotyla lateralis is aligned
 204 with the trochlea metatarsi IV. The fossa infracotylaris
 205 dorsalis is deep, and the tuberositas musculi tibialis cranialis
 206 is elongated. Both foramina vascularia proximalia open in
 207 both faces, caudal and cranial. The elongated and large
 208 foramen vasculare distale opens into the fossa supra-
 209 trochlearis plantaris. The sulcus extensorius extends
 210 becoming shallower and narrower from the foramina vas-
 211 cularia proximalia to the foramen vasculare distale. The
 212 crista medialis hypotarsi is very well developed.

Measurements. Femur: total length 66 mm, proximal 213
 214 width 14.6 mm, distal width 19 mm. Humerus: total length
 215 141 mm, lateromedial anterior width 28 mm, distal width
 216 20 mm. Tibiotarsus: total length 18.5 mm, distal width

Author Proof

Fig. 3 Mandible and elements of the wing of *Stercorarius antarcticus lonnbergi*: **a** mandible IAA 802 (*dorsal view*), **b** mandible IAA 802 (*ventral view*), **c** carpometacarpus IAA 801 (*anterior view*), **d** carpometacarpus IAA 801 (*posterior view*), **e** humerus IAA 798 (*anterior view*), **f** humerus IAA 798 (*posterior view*). Scale bar: 10 mm

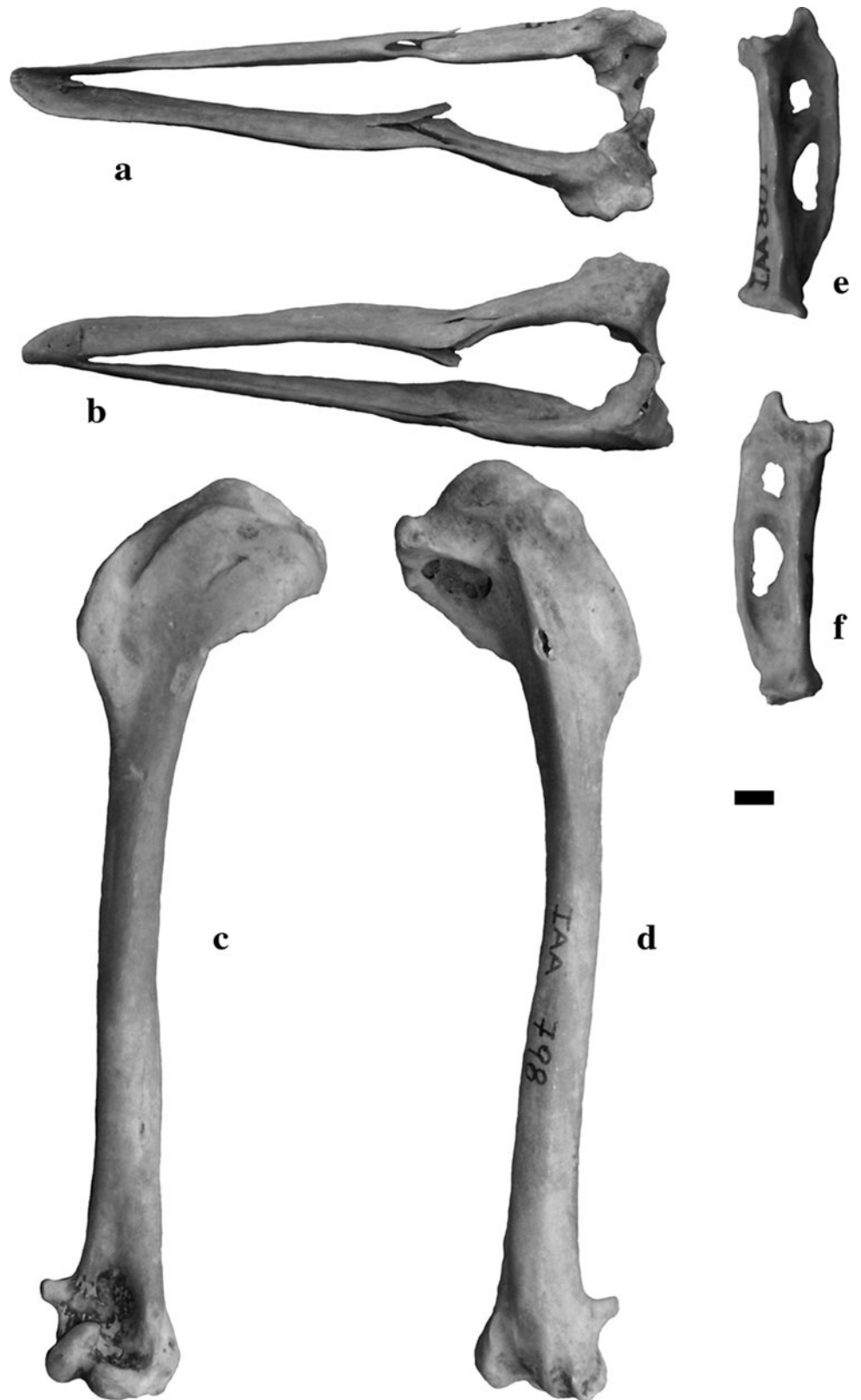
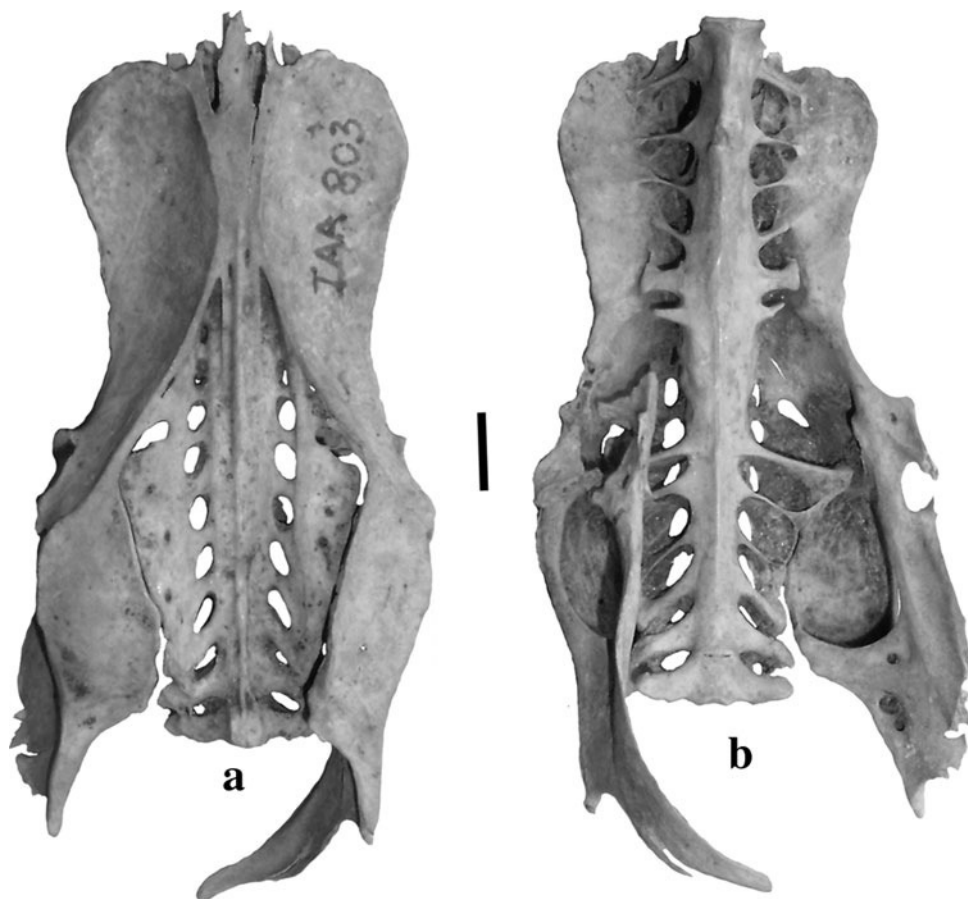


Fig. 4 Pelvis of *Stercorarius antarcticus lonnbergi*: **a** Os coxae and synsacrum IAA 803 in dorsal view, **b** in ventral view. Scale bar: 10 mm



217 12.7 mm. Tarsometatarsus: total length 73 mm, proximal
218 width 14 mm, distal width 13.5 mm.

219 **Paleoenvironmental context**

220 Because the sea level reached its present height between
221 5,000 and 6,000 ¹⁴C-calibrated year BP (Lambeck and
222 Chappel 2001), most of the observed Holocene uplift in the
223 South Shetland Islands is due to glacio-isostasy (Pallàs
224 et al. 1997). This suggests that the sedimentary sequence
225 may allow quantification of the glacio-isostatic uplift of the
226 coast since deposition (del Valle et al. 2002).

227 Additionally, the values of age versus altitude a.s.l. fall
228 between the predicted relative sea-level curve, according to
229 the ICE-3G glacial rebound model (Tushingham and
230 Peltier 1992), and the expected relative sea-level curves
231 corresponding to Maxwell Bay (Bahía Guardia Nacional)
232 area (King George Island: Isla 25 de Mayo) (Pallàs et al.
233 1997; del Valle et al. 2002; Fig. 3).

234 The recorded birds probably inhabited King George
235 Island before the “climate optimum” of the Antarctic
236 Peninsula (between 4,000 and 3,000 years BP; see, e.g.,
237 Ingólfsson et al. 1998). Moreover, the coastal occupation

238 period, reported by del Valle et al. (2002), might match the
239 cooling period in Antarctica, from 8,000 to 4,000 years BP
240 (Lorius et al. 1979; Cias et al. 1992).

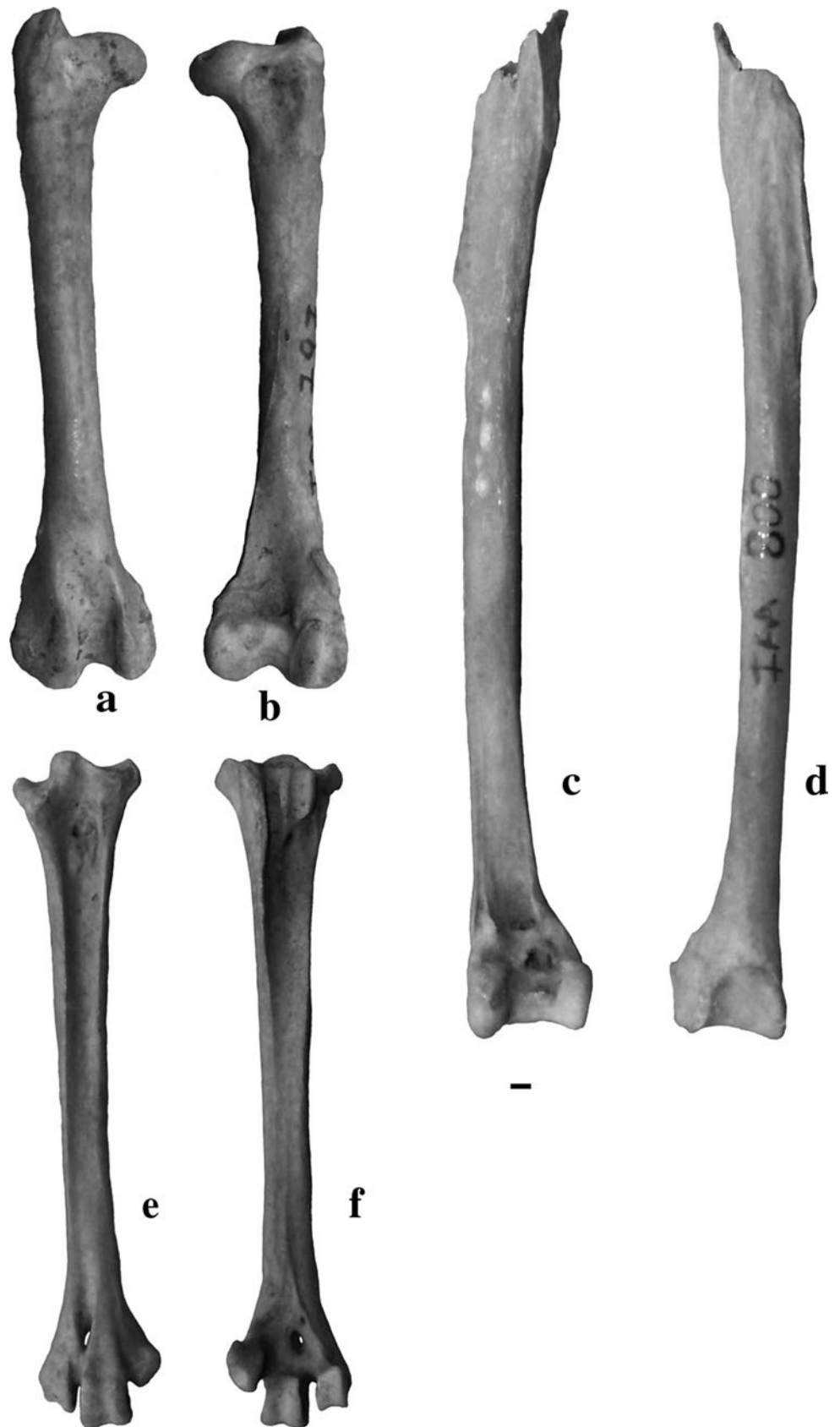
241 The fossiliferous sediments were assigned to a deposi-
242 tional environment, which is in agreement with the sea-
243 sonally open marine conditions proposed for the Antarctic
244 Peninsula between 8,000 and 4,000 years BP (e.g.,
245 Ingólfsson et al. 1998). The sedimentary, paleontological,
246 and chronological data reported herein are in agreement
247 with the quaternary paleoenvironmental chronology known
248 for the region (del Valle et al. 2002).

249 **Results and conclusions**

250 The brown skua is the largest and heaviest of all the
251 southern skuas, and its identification is partially based on
252 size measurements. However, osteological features repre-
253 sent a legitimate and necessary tool for taxonomic deter-
254 minations (Acosta Hospitaleche et al. 2009). The presence
255 of the following characters allows its assignment to species
256 level as well as the distinction of this species from *Sterc-*
257 *orarius maccormicki*, the most similar in morphological
258 terms. The processi mandibulae medialis of the mandible

Author Proof

Fig. 5 Elements of the leg of *Stercorarius antarcticus lonnbergi*: **a** femur IAA 797 (anterior view), **b** femur IAA 797 (posterior view), **c** tibiotarsus IAA 800 (anterior view), **d** tibiotarsus IAA 800 (posterior view), **e** tarsometatarsus IAA 799 (anterior view), **f** tarsometatarsus IAA 799 (posterior view). Scale bar: 10 mm



259 are robust. In the humerus, the tuberculum ventrale is
260 clearly defined, the crus dorsale and ventrale fossae are
261 well developed, and the fossa m. brachialis is extensive. In
262 the tarsometatarsus, the trochlea carpalis has an acute
263 proximal end, the os metacarpale alulare is hook-shaped,
264 the sulcus interosseus is wide, the fossa infratrochlearis is
265 proximodistal, and the end of the processus pisiformis is
266 sharper. In the femur, the crista trochanteris is well
267 extended, and the sulcus intercondylaris is shallow and
268 symmetric with respect to the condyles. In the tibiotarsus
269 the pons supratendinosus is broad. In the tarsometatarsus
270 the eminentia intercotylaris is rounded and sturdy, the
271 tuberositas musculi tibialis cranialis is notable, the edges of
272 the trochlea metatarsi III are weak, and the sulcus
273 extensorius is deep.

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279 References

280 Acosta Hospitaleche, C., D. Montalti, and L. Marti. 2009. Skeletal
281 morphoanatomy of the Brown Skua *Stercorarius antarcticus*
282 *lonnbergi* and the South Polar Skua *Stercorarius maccormicki*.
283 *Polar Biology* 32: 759–774.
284 Andersson, M. 1973. Behaviour of the Pomarine Skua, *Stercorarius*
285 *pomarinus* Temm with comparative remarks on Stercorariinae.
286 *Ornis Scandinavica* 4: 1–16.
287 Andersson, M. 1999. Hybridization and skua phylogeny. *Proceeding*
288 *of Royal Society of London B* 266: 1579–1585.
289 Baumel, J., and L.M. Witmer. 1993. Osteologia. In *Handbook of*
290 *avian anatomy: Nomina Anatomica Avium*, ed. Baumel, J., King,
291 A., Breazile, J.E., Evans, H., and J.C. Vanden Bergue, 45–132.
292 Cambridge, Massachusetts.
293 Cassoli, P.F. 1980. L'avifauna del Pleistocene superiore delle Arene
294 Candide (Liguria). *Memoria dell'Istituto Italiano di Paleontolo-*
295 *gia Umana* 3: 155–234.
296 Cias, P., J.R. Petit, J. Jouzel, C. Lorius, N.I. Barkov, V. Lipenkov, and
297 V. Nicolaiev. 1992. Evidence for an early Holocene climatic
298 optimum in the Antarctic deep ice-core record. *Climate*
299 *Dynamics* 6: 169–177.
300 Cohen, B.L., A.J. Baker, K. Blechschmidt, D.L. Dittmann, R.W.
301 Furness, J.A. Gerwin, A.J. Helbig, J. de Korte, H.D. Marshall,
302 R.L. Palma, H.-U. Peter, R. Ramli, J. Siebold, M.S. Willcox,
303 R.H. Wilson, and R.M. Zink. 1997. Enigmatic phylogeny of
304 skuas (Aves: Stercorariidae). *Proceeding Royal Society London*
305 *B* 264: 181–190.
306 del Valle, R.A., D. Montalti, and M. Inbar. 2002. Mid-Holocene
307 macrofossil-bearing raised marine beaches at Potter Peninsula,
308 King George Island, South Shetland Islands. *Antarctic Sciences*
309 14: 263–269.
310 Emslie, S.D. 1995. An Early Irvingtonian Avifauna from Leisey Shell
311 Pit, Florida. *Bulletin of the Florida Museum of Natural History*
312 37: 299–344.
313 Fordyce, E. 1982. The Australasian marine vertebrate record and its
314 climatic and geographic implications. In *The fossil vertebrate*

record of Australasia, ed. P.V. Rich, and E.M. Thompson, 595–
627. Australia: Monash University Offset. 315
Howard, H. 1946. A review of the Pleistocene birds of Fossil
lake, Oregon. *Carnegie Institute of Washington Publication* 551:
141–195. 316
Ingólfsson, O., C. Hjort, P.A. Berkman, S. Björck, E. Colhoun, I.D.
Goodwin, B. Hall, K. Hirakawa, M. Melles, P. Möller, and M.L.
Prentice. 1998. Antarctic glacial history since the last glacial
maximum: An overview of the record on land. *Antarctic*
Sciences 10: 326–344. 317
Lambeck, K., and J. Chappel. 2001. Sea level change through the last
glacial cycle. *Science* 292: 679–686. 318
Lorius, C., L. Merlivat, J. Jouzel, and M. Pouchet. 1979. A 30 000-yr
climatic record from Antarctic ice. *Nature* 280: 642–648. 319
Millener, P.R. 1991. The Quaternary avifauna of New Zealand. In
Vertebrate palaeontology of Australasia, ed. P. Vickers-Rich,
J.M. Monaghan, R.F. Baird, and T.H. Rich, 1317–1344.
Melbourne: Pioneer Design Studio in cooperation with the
Monash University Publications Committee. 320
Millener, P.R. 1999. The history of the Chatham Islands' bird fauna
of the last 7000 years - a chronicle of change and extinction. In
Avian paleontology at the close of the 20th Century, ed. S.L.
Olson, 85–109. *Proceedings of the 4th International meeting of*
the Society of Avian Paleontology and Evolution, Washington. 321
Montalti, D., Acosta Hospitaleche, C., and R. del Valle. (in press
2009). New Holocene penguin assemblages at South Shetland
Islands, Antarctica. *Neues Jahrbuch für Geologie und Palaon-*
tologie-Abhandlungen. 322
Mourer-Chauviré, C. 1975. Les oiseaux du Pléistocène moyen et
supérieur de France. *Documents des Laboratoires de Géologie*
de la Faculté des Sciences de Lyon 64: 1–624. 323
Olson, S., and P.C. Rasmussen. 2001. Miocene and Pliocene birds
from the Lee Creek Mine, North Carolina. In *Geology and*
paleontology of the Lee Creek Mine, North Carolina, eds. Ray,
C.E., and D.J. Bohaska, Smithsonian Contributions to Paleobi-
ology 90: 233–365. 324
Pallás, R., T.S. James, F. Sàbat, J.M. Vilaplana, and D.R. Grant. 1997.
Holocene uplift in the South Shetland Islands: Evaluation of
tectonics and glacio-isostasy. In *The Antarctic region: Geolog-*
ical evolution and processes, ed. C. Ricci, 861–868. Siena: Terra
Antarctic. 325
Remsen, J.V., Jr., Jaramillo, A., Nores, M.A., Robbins, M.B.,
Schulenberg, T.S., Stiles, F.G., da Silva, C., Stotz, D.F. and
K.J. Zimmer. 2009. A classification of the bird species of South
America. American Ornithologists' Union. [http://www.museum.](http://www.museum.lsu.edu/~Remsen/SACCBaseline.html)
[lsu.edu/~Remsen/SACCBaseline.html](http://www.museum.lsu.edu/~Remsen/SACCBaseline.html). Accessed 9 November
2009. 326
Sánchez Marco, A. 2006. Aves fósiles del Auriñaciense (Pleistoceno
superior) de El Castillo (Cantabria, España). *Zona Arqueologica*
7: 114–121. 327
Turbott, E.G. 1990. *Checklist of the birds of New Zealand and the*
Ross Dependency, Antarctica, 3rd ed. Auckland: Ornithological
Society of New Zealand & Random Century New Zealand Ltd. 328
Tushingham, A.M., and W.R. Peltier. 1992. Validation on the ICE-3G
model of Würm-Wisconsin deglaciation using a global data base
of relative sea level histories. *Journal of Geophysics Research*
97: 3285–3304. 329
Tyrberg, T., and F. Hernández. 1995. First fossil record of the Great
skua. *Bulletin of the British Ornithological Club* 115: 167–168. 330
Worthy, T.H., and P. Jouventin. 1999. The fossil avifauna of
Amsterdam Island, Indian Ocean. *Smithsonian Contributions to*
Paleobiology 89: 39–65. 331
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