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1 A new fossil rorqual aff. *Balaenoptera bertae* specimen from the Shinazawa
2 Formation (late Pliocene to early Pleistocene), Yamagata, Japan

3

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16

17 **Abstract.** More than 23 extinct species and 10 extant species of the
18 Balaenopteridae are known. Our knowledge of the family Balaenopteridae is
19 increasing quickly, however, few fossil records support a circum-north Pacific
20 distribution of balaenopterid genera and species. Because of limited preservations,
21 most rorqual fossils reported from the western North Pacific can only be
22 identified to the family level. A skull from the Shinazawa Formation (late
23 Pliocene to early Pleistocene) in Yamagata, Japan, is identified as aff.
24 *Balaenoptera bertae* by possessing two diagnostic features of the species: large
25 occipital condyles, and a posteriorly elongate postglenoid process. Combination

26of four more features also support that the specimen is a closely related to *B.*
27*bertae*. The specimen is probably a slightly older individual than the holotype of
28*B. bertae*, based on the estimated bizygomatic width and slightly longer posterior
29process of the tympanoperiotic. The first and only report of *B. bertae* was from
30the Pliocene Purisima Formation in California, USA. The specimen from Japan is
31incompletely preserved, but implies that the occurrence of *B. bertae* in the
32western North Pacific for the first time, as many living balaenopterids are
33distributed across the North Pacific, such as *Balaenoptera musculus*, *B. physalus*,
34*B. borealis*, *B. acutorostrata*, and *Megaptera novaeangliae*.

35

36Key Words: Balaenopteridae, Mysticeti, Cetacea, Neogene, Quaternary, Pacific

37

38

Introduction

39Our knowledge of the fossil record of the rorqual whales, family Balaenopteridae,
40has been improving recently, such establishing new balaenopterids from Belgium,
41the Netherlands, Italy, California, Peru, Chile and Japan, and taxonomic revision
42of European problematic fossils (Deméré *et al.*, 2005; Bisconti, 2007, 2010;
43Bosselaers and Post, 2010; Boessenecker, 2013a; Bisconti and Bosselaers, 2016;
44Marx and Kohno, 2016; Bisconti *et al.*, 2019, 2020a; Bisconti and Bosselaers,
452020). More than 23 extinct species and 10 extant species are known (Rosel *et al.*,
462021). Among fossil balaenopterids, few records suggest a circum North Pacific
47distribution because there are only a few records of extinct balaenopterid species
48from the western side of the North Pacific.

49 The Miocene through Pleistocene fossil record of balaenopterid whales in Japan
50has improved recently (Oishi *et al.*, 1985; Hasegawa *et al.*, 2002; Kohno *et al.*,

512007; Kimura *et al.*, 2015; Tanaka and Taruno, 2019). Some records with species
52level identification are as follows; *Megaptera novaeangliae* from the middle
53Pleistocene (about 0.12 to 0.15 Ma) at Chiba (Nagasawa and Mitani, 2004),
54*Burtinopsis tatsunokuchiensis* from the early Pliocene at Iwate (Hatai *et al.*,
551963), and *Miobalaenoptera numataensis* from the late Miocene at Hokkaido
56(Tanaka and Watanabe, 2019). These limited fossil records from the western
57North Pacific are insufficient to document the past distribution of balaenopterids
58across the Pacific. However, fossils of other whale families such the pygmy right
59whale from Okinawa and the right whale from Taiwan provide their past global
60distribution (Tsai *et al.*, 2017; Tsai and Chang, 2019).

61 Here, we describe a balaenopterid cranium from the late Pliocene to early
62Pleistocene found at Yamagata, Japan, which was preliminary reported by
63Nagasawa *et al.* (2003). The specimen is incompletely preserved but tentatively
64identified to species level. This record implies that there was a widely distributed
65extinct balaenopterid species across the Pacific.

66

67

Materials and method

68Morphological terminology follows Mead and Fordyce (2009). The specimen
69was measured using calipers in mm. Distances are either horizontal or vertical,
70unless identified as point to point (Table 1). Comparison was made with all
71named balaenopterid fossils with the skull (Table 2) by combination of
72examination on literatures and through our own examination of specimens.

73

74 *Institutional Abbreviation.*— YPM, Yamagata Prefectural Museum, Japan.

75

76 **Systematic paleontology**

77 Order Cetacea Brisson, 1762

78 Neoceti Fordyce & de Muizon, 2001

79 Suborder Mysticeti Gray, 1864

80 Family Balaenopteridae Gray, 1864

Genus *Balaenoptera* La Cépède, 1804

81 *Balaenoptera bertae* Boessenecker, 2013a

82 aff. *Balaenoptera bertae*

83 Figures 2 to 4 and Table 1

84

85 *Diagnosis.*—YPM 11852 preserves the supraoccipital, right exoccipital,
86 basioccipital, right squamosal, right alisphenoid, and parietals. YPM 11852 can
87 be identified as a member of the Balaenopteridae by having combination of the
88 temporal fossa invisible in dorsal view, which is overhanged by the temporal and
89 nuchal crests, widely expanded posterior portion and anteriorly converging
90 supraoccipital in dorsal view, weak protrusion of the posterolateral corner of the
91 exoccipital in ventral view, and posteriorly well projected postglenoid process in
92 ventral view (Bisconti and Bosselaers, 2016).

93 YPM 11852 is identified as aff. *Balaenoptera bertae* by having two diagnostic
94 features of the species stated in 2013 such having large occipital condyles, and a
95 posteriorly elongate postglenoid process (Boessenecker, 2013a). The ventral part
96 of the postglenoid process of YPM 11852 is broken, but preserved part shows
97 slender process with a strong excavation at the base of the postglenoid process,
98 which is the same condition as the holotype of *B. bertae*. In addition, there are
99 four shared features (see discussion) such a large fossa for the sternocephalicus,

100dorsally wider and ventrally narrower foramen magnum, occipital condyle
101borders the lateral and ventral sides, but not dorsolateral sides of the foramen
102magnum, and straight nuchal crest at the level of the subtemporal crest in dorsal
103view. Comparison with the holotype of *B. bertae*, YPM 11852 has more slender
104zygomatic process and straight lateral border of the supraoccipital. As Tsai and
105Fordyce (2014) mentioned, different heterochronic processes characterizing
106different mysticete clades. These differences might be affected by ontogenetic
107variation as these can be seen among *Balaenoptera acutorostrata* in Figure 2 of
108Nakamura and Kato (2014) (see more in ontogeny section).

109 *Locality.*—YPM 11852 was found ventral up in the Shinazawa Formation at the
110Shinazawa sandstone quarry, in Shonai Town (former Tachikawa Town),
111Yamagata Prefecture, Japan (Figures 1 and 2): Latitude 38°41'15.87"N, longitude
112140° 0'14.97"E on 10 October 2000 by a student of Yamagata University on a
113field trip (Nagasawa *et al.*, 2003). The specimen was excavated by R. Abe and S.
114Oba on 4 November 2000.

115 *Horizon.*—Several fossiliferous stratigraphic units are exposed in the vicinity of
116the YPM 11852 locality near Shinazawa, including the middle Miocene Aosawa
117and Kusanagi Formations, the upper Pliocene Shinazawa Formation, and terrace
118deposits (Honda and Applied Geological Society of Yamagata, 2016).

119 The Shinazawa Formation is exposed in a small geographic area where YPM
12011852 was discovered (Zinbo, 1965). This unit at the YPM 11852 locality is
121reported that the sequence of fossil locality is about 90 m in thickness and is
122overlain by unnamed unconsolidated Pleistocene sediments (Nagasawa *et al.*,
1232003). The Shinazawa Formation of the fossil locality can be divided into three
124parts. The lower part is about 40 m thick, which is composed mainly of massive

125 fine sandstone with burrows and bioturbation, and intercalated thin silt and fine
126 gravel layers. The middle part, about 20 m thick, is formed by mainly massive
127 fine sandstone with burrows and bioturbation, shell fragments, carbonaceous
128 materials, and shell bed. Grain size in the middle part coarsens upwards. The
129 upper part, about 30 m thick, consists of cross-laminated coarse sandstone at the
130 bottom, alternated coarse sandstone and fine gravel at the middle, and fine
131 sandstone with a lignite layer at the top.

132 YPM 11852 was collected from the lower part. The fossil horizon is
133 approximately 10 m below the 20 cm thick shell marker bed. Other vertebrate
134 remains have been reported, such as a skull (still unprepared), mandible, rib and
135 an atlas of cetacean specimens belonging to different individuals from different
136 horizon of the lower and middle parts, and two sirenian ribs from the upper part
137 of the unit at the Shinazawa quarry (Nagasawa *et al.*, 2003). Foraminifera
138 *Elphidium* from the lower part of the formation also indicates that the
139 environment was shallow marine (Ozawa *et al.*, 1986). The area was shallowing
140 upward sequence.

141 The Shinazawa Formation has not been directly dated, but thought to be the late
142 Pliocene (Honda and Applied Geological Society of Yamagata, 2016), based on
143 fossils such as a Pliocene plant *Comptonia kidoi* (Suzuki, 1961) and various
144 mollusks belonging to the Omma-Manganzian fauna (Otuka, 1939), such as
145 *Yabepecten tokunagai*, *Mizuhopecten yessoensis*, *Turritella saishuensis* and
146 others (Sato, 2000). The Omma-Mangazian fauna is composed of mainly cold
147 water species and existed from the Pliocene to the early Pleistocene age in the
148 northeastern Sea of Japan area (Amano, 2007). A previous geological map
149 (Ozawa *et al.*, 1986) did not use the term Shinazawa Formation, and recognized

150the fossil locality area as the Kan-nonji Formation (lower Pleistocene) in the
151northern Shonai area based on similar lithology. At present, it is not possible to
152correlate the two formations because lack of dating for the Shinazawa Formation.
153Based on the mollusc fauna, we estimate the age of YPM 11852 as late Pliocene,
154and also include possibility of the early Pleistocene.

155

156

General description

157 *Ontogeny*.—YPM 11852 can be identified as an immature individual. It has
158visible parietal/squamosal and alisphenoid/pterygoid sutures; and does not
159preserve other cranial sutures. The holotype of *Balaenoptera bertae* identified as
160an immature individual also has a visible parietal/squamosal suture
161(Boessenecker, 2013a). The posterior process of tympanoperiotic grows during
162ontogeny (Bisconti, 2001). YPM 11852 shows slightly longer posterior process
163of the tympanoperiotic (about 105.0 mm long, based on the fossa of the skull for
164the posterior process) than that of *B. bertae* (about 80.0 mm long). In addition,
165estimated bizygomatic width of YPM 11852 from the right side is 719 mm+. It is
166wider than that of the holotype of *B. bertae*: 614 mm (Boessenecker, 2013a).
167Thus, YPM 11852 possibly was slightly ontogenetically older than the holotype
168of *B. bertae*.

169 *Supraoccipital, exoccipital and basioccipital*.—The supraoccipital is triangular in
170dorsal view (Figure 3D) with an almost straight nuchal crest. The posterolateral
171part of the nuchal crest is expanded laterally. The occipital tuberosities are
172formed as a horizontal ridges and depressions ventral to the ridges. The foramen
173magnum is teardrop shaped with a bluntly triangular ventral margin. The
174occipital condyles are wide and project posteroventrally beyond the surface of

175the occipital shield. The intercondyloid notch is shallow. Ventrally, the
176exoccipital has a weakly depressed paroccipital concavity (Figure 4). Medial to
177the paroccipital concavity, there is an anteroposteriorly running shallow jugular
178notch medially. The lateral end of the exoccipital stops medial to the postglenoid
179process and forms a strong posteroexternal angle.

180 *Squamosal.*—The zygomatic process projects anterolaterally and is bounded
181medially by a shallow and vertical squamosal crease. A shallow squamosal cleft
182is developed, running horizontally from the posterior end of the alisphenoid to
183the dorsal broken base of the zygomatic process. A small elliptical opening filled
184by matrix (24.5 mm long, 13.5 mm high) located anterior to the squamosal cleft,
185ventral to the parietal/squamosal suture on the temporal fossa; this might be a
186junction of the parietal, squamosal and alisphenoid. The postglenoid process is
187slender (preserved length is 88.0 mm) and projects posterolaterally. The lateral
188surface of the zygomatic process is weakly excavated, and the anterior tip of the
189zygomatic process orients anterolateral. Posterior to the postglenoid process,
190there is a shallow transverse groove of the external auditory meatus. In ventral
191view, the external auditory meatus is narrow anteroposteriorly (about 14 mm
192long) but wider medially (about 27 mm long). On the lateral surface of the
193squamosal, dorsal to the external auditory meatus, there is a shallow depression
194of the fossa for the sternocephalicus. On the ventral surface and just lateral to a
195broken anterior process of the periotic, only the base of the falciform process is
196preserved.

197 *Pterygoid.*—The right pterygoid (Figure 3E) is exposed laterally within the
198temporal fossa, located anterior to the squamosal, and contacts the alisphenoid.

199 *Parietal.*—The parietal shows a weakly excavated surface and forms the

200temporal wall and a dorsoventrally thin nuchal crest. The parietal/squamosal
201suture is visible.

202 *Alisphenoid*.— Based on relative position with other structures, the presumed
203alisphenoid has a small exposure (17.9 mm long, 6.0 mm high) on the
204anterolateral surface of the temporal fossa, anteroventral to the squamosal and
205dorsal to the pterygoid. The alisphenoid contacts with the squamosal cleft (Figure
2063E). Its anterior part was possibly damaged as mentioned above.

207 *Periotic*.—The incomplete right periotic preserves only the dorsal part of the
208anterior and posterior processes and showing fresh broken surface (Figure 4).
209The lateral end of the posterior process of the tympanoperiotic is damaged, and
210the posterior process was possibly exposed on the lateral surface of the skull. The
211anterior process was at least 25.5 mm long. The preserved pars cochlearis is 48.5
212mm long anteroposteriorly, 43.5 mm wide mediolaterally. The ventral surface of
213the pars cochlearis is broken away, and the section of a presumed internal
214acoustic meatus is exposed, and its preserved anteroposterior length is 13.5 mm.
215The remnants of a long posterior process projects posterolaterally from the
216damaged pars cochlearis. Though incomplete, the posterior process appears
217laterally narrowing (maximum length of the part is 15.5 mm). The posterior
218process was probably about 105.0 mm long (preserved maximum length is 86.0
219mm) when complete.

220

221

Discussion

222**Comparison with fossil balaenopterids and linkage in the North Pacific of**
223**the Balaenopteridae**

224 As mentioned above, YPM 11852 is identified as aff. *Balaenoptera bertae*,

225because it has two diagnostic features of the species, such as large occipital
226condyles occupied one third of the bizygomatic width, and a posteriorly elongate
227postglenoid process (Boessenecker, 2013a). YPM 11852 and *B. bertae* also have
228four shared features such as having 1) a large fossa for the sternocephalicus on
229the lateral surface of the squamosal, 2) a dorsally wider and ventrally narrower
230foramen magnum, and 3) a occipital condyle borders the lateral and ventral sides,
231but not dorsolateral sides of the foramen magnum, and 4) a straight nuchal crest
232at the level of the subtemporal crest. YPM 11852 is closely related to *B. bertae*
233by the two diagnostic features and four shared features.

234 Having elongate and posteroventrally projecting postglenoid processes can be
235seen also in *Marzanoptera tersillae*, which formed a clade with *Balaenoptera*
236*bertae* by phylogenetic analysis of Bisconti *et al.* (2020b). In comparison with
237presumed closely related *Marzanoptera tersillae*, YPM 11852 and *Balaenoptera*
238have slenderer postglenoid process, larger occipital condyles, and also dorsally
239wide and ventrally narrow foramen magnum. On the other hands, *M. tersillae*
240shows an elliptical foramen magnum.

241 Of note, Bisconti *et al.* (2020b) established the genus *Marzanoptera* with
242combination of some diagnostic features on the skull and periotic, such a
243remarkably short premaxilla, strongly narrowed ascending process of the maxilla
244and so on. *Balaenoptera bertae* was transferred to the genus as *Marzanoptera*
245*bertae*. However, the premaxilla, maxilla and body of the periotic were not
246preserved on the holotype of *Balaenoptera bertae*, which is currently only one
247specimen of the species (Boessenecker, 2013a). Thus, we reserve to use the
248genus name *Marzanoptera* for *Balaenoptera bertae* here.

249 In comparison with all named balaenopterids preserving skulls (Table 2),

250 "*Balaenoptera*" *siberi* also possesses a long postglenoid process (longer than in
251 *Norrisanima miocaena*, *Parabalaenoptera baulinensis*, *Nehalaennia devossi*,
252 *Incakujira anillodefuego*, *Archaebalaenoptera castriarquati*, *Plesiobalaenoptera*
253 *hubachi*), but the postglenoid process of "*B.*" *siberi* is shorter than in *B. bertae*
254 and YPM 11852. *Archaebalaenoptera liesselensis* also shows larger occipital
255 condyles, occupying almost one half of the exoccipital width compare with other
256 balaenopterids ("*Balaenoptera*" *siberi*, *Norrisanima miocaena*, *Protolorqualus*
257 *wilfriedneesi*, *Fragilicetus velponi*, *Parabalaenoptera baulinensis*, *Nehalaennia*
258 *devossi*, *Incakujira anillodefuego*, "*Balaenoptera*" *ryani*, *Marzanoptera tersillae*,
259 *Protolorqualus cuvieri*, *Plesiobalaenoptera hubachi*).

260 A large fossa for the sternocephalicus can be seen on not only YPM 11852 and
261 *B. bertae*, but also on *Norrisanima miocaena*, *Protolorqualus wilfriedneesi* and
262 *Diunatans luctoretemergo*. *Parabalaenoptera baulinensis* and *Marzanoptera*
263 *tersillae* have a much smaller single fossa for the sternocephalicus.
264 *Archaebalaenoptera liesselensis*, *Fragilicetus velponi*, *Incakujira anillodefuego*
265 and *Plesiobalaenoptera hubachi* possess two fossae for the sternocephalicus. The
266 foramen magnum is rounded to dorsoventrally long elliptical in most
267 balaenopterids (*Norrisanima miocaena*, *Protolorqualus wilfriedneesi*,
268 *Parabalaenoptera baulinensis*, *Nehalaennia devossi*, *Marzanoptera tersillae* and
269 *Plesiobalaenoptera hubachi*), but only YPM 11852, *B. bertae* and *Incakujira*
270 *anillodefuego* have dorsally wider and ventrally narrower foramen magnum.
271 YPM 11852, *B. bertae* and most of other balaenopterids ("*Balaenoptera*" *siberi*,
272 *Norrisanima miocaena*, *Protolorqualus wilfriedneesi*, *Fragilicetus velponi*,
273 *Nehalaennia devossi*, *Incakujira anillodefuego*, and "*Balaenoptera*" *ryani*)
274 possess occipital condyles that are positioned ventrolaterally, and do not extend

275to the dorsolateral margin of the foramen magnum. Only *Parabalaenoptera*
276*baulinensis*, *Marzanoptera tersillae* and *Plesiobalaenoptera hubachi* have
277occipital condyles that completely encircle the foramen magnum. Among
278balaenopterids, the lateral margin of the exoccipital at the level of the
279subtemporal crest in dorsal view is posteriorly concave (*Norrisanima miocaena*,
280*Archaeobalaenoptera liesselensis*, *Diunatans luctoretmergo*, *Incakujira*
281*anillodefuego*, *Marzanoptera tersillae*, *Archaeobalaenoptera castriarquati* and
282*Plesiobalaenoptera hubachi*), posteriorly convex (*Fragilicetus velponi*,
283*Nehalaennia devossi*, *Protororqualus cuvieri*), or straight (YPM 11852, *B. bertae*,
284"*Balaenoptera*" *siberi*, *Protororqualus wilfriedneesi* and *Parabalaenoptera*
285*baulinensis*). Combination of the diagnostic features and other features'
286conditions support that YPM 11852 is the most closely related to *B. bertae*.
287 Presumed closely related specimens, YPM 11852 and the holotype of
288*Balaenoptera bertae* show that the extinct balaenopterid was distributed both the
289eastern and western North Pacific during the late Pliocene to possibly into the
290early Pleistocene. Currently, many balaenopterids have circum North Pacific
291distributions, including *Balaenoptera musculus*, *B. physalus*, *B. borealis*, *B.*
292*acutorostrata*, and *Megaptera novaeangliae* (Jefferson *et al.*, 2008). There are
293some other marine mammal species and/or genera known to have a circum-North
294Pacific distribution during the Pliocene to early Pleistocene; *Eschrichtius*
295spp.(Tsai and Boessenecker, 2015; Kimura *et al.*, 2018), *Balaenoptera physalus*
296(Tsai and Boessenecker, 2017), *Herpetocetus* (Oishi *et al.*, 1985; Boessenecker,
2972013a, 2013b) *Callorhinus gilmorei* and *Thalassoleon* (Repenning and Tedford,
2981977; Kohno, 1992; Kohno and Yanagisawa, 1997), and *Hydrodamalis*
299(Domning and Furusawa, 1995). This new record/YPM 11852 indicates that the

300 extinct rorqual species *Balaenoptera bertae* can be added to this list of marine
301 mammals with a circum-North Pacific distribution during the pre-glacial
302 Pliocene-Pleistocene interval.

303

304

Conclusion

305 We report a skull YPM 11852 from the Shinazawa Formation (late Pliocene to
306 early Pleistocene) in Yamagata, Japan, as aff. *Balaenoptera bertae*. The
307 specimen shows two diagnostic features of the species such as having large
308 occipital condyles, and a posteriorly elongate postglenoid process. Combination
309 of the four more shared features also support this. YPM 11852 is probably
310 slightly ontogenetically older individual than the holotype of *B. bertae*, based on
311 the estimated bizygomatic width and slightly longer posterior process of the
312 tympanoperiotic. YPM 11852 suggests that fossil balaenopterids had a circum
313 North Pacific distribution, like several modern balaenopterid species. YPM
314 11852 is incompletely preserved, however nonetheless demonstrates the
315 occurrence of *Balaenoptera bertae* and/or closely related taxa in the western
316 North Pacific for the first time.

317

318

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333 **Figure 1.** A-B, Maps showing the locality of YPM 11852. The base map of A is
334 from a topographic map published by the Geospatial Information Authority of
335 Japan. C, Stratigraphic sections of the locality modified from Nagasawa *et al.*
336 (2003).

337 **Figure 2.** Field photo of YPM 11852, aff. *Balaenoptera bertae*.

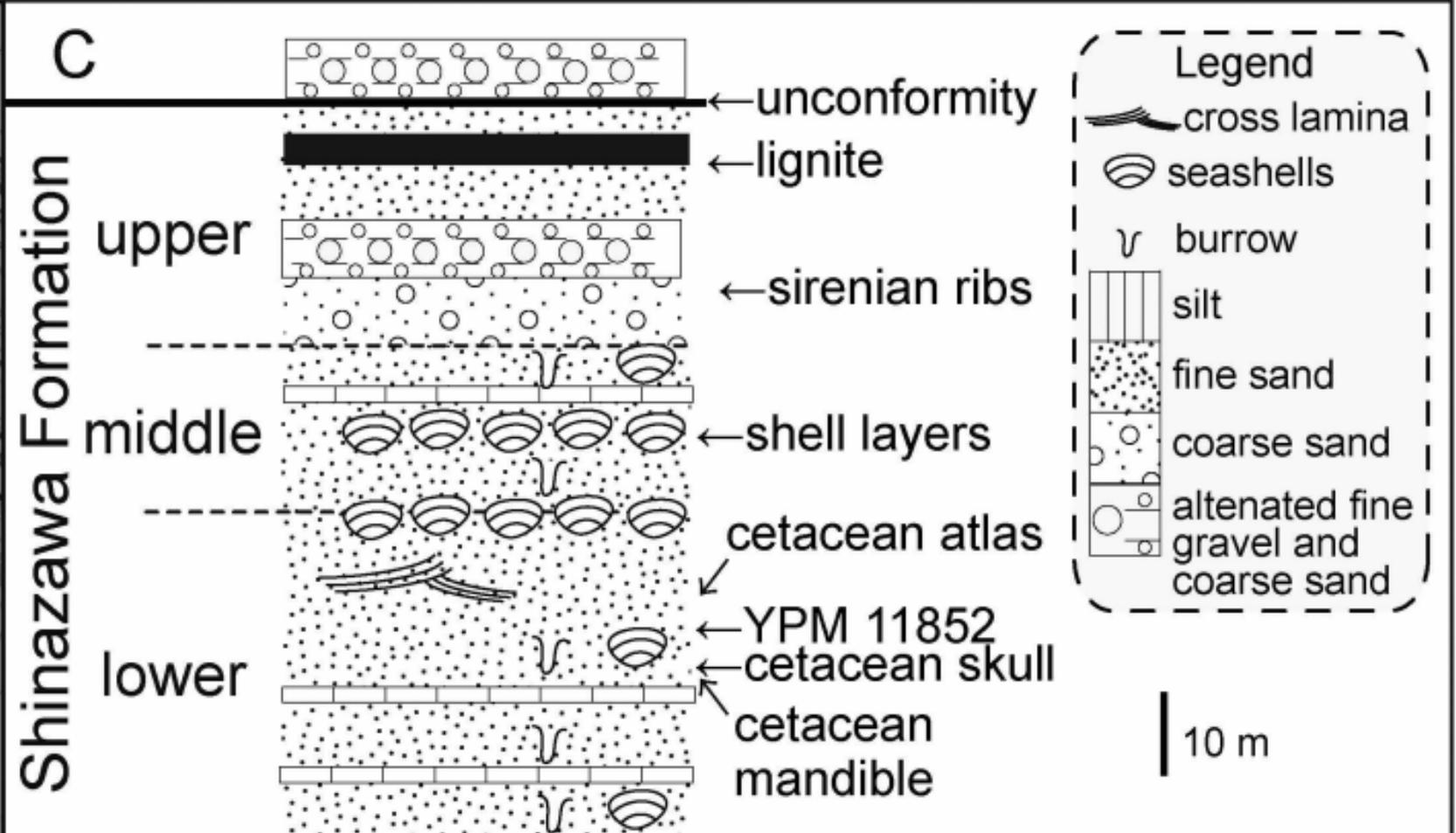
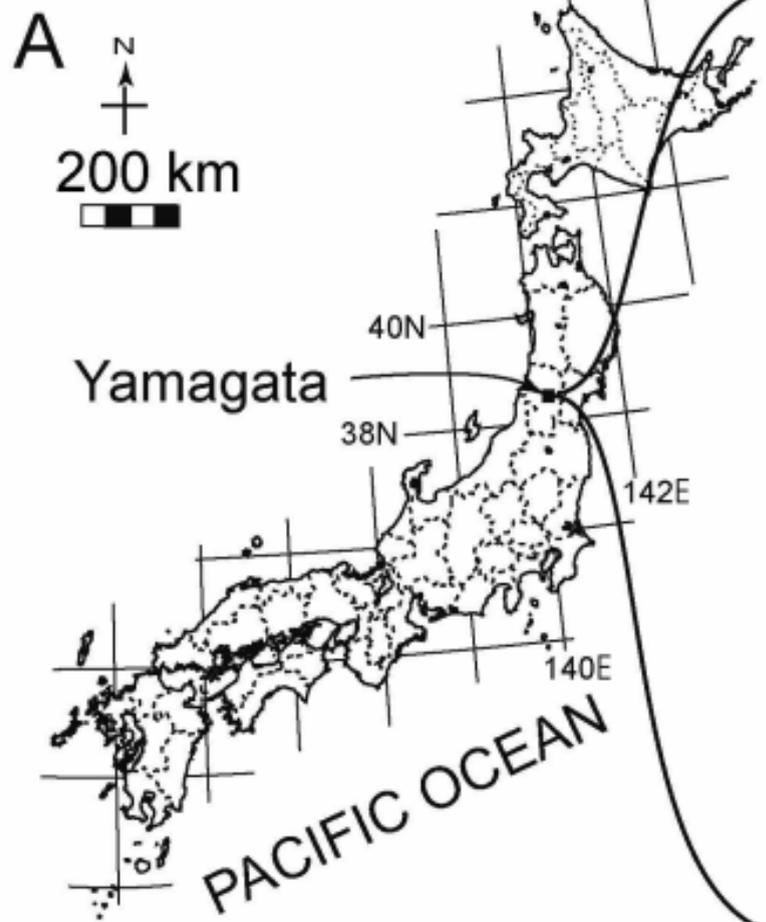
338 **Figure 3.** A-E, Skull of YPM 11852, aff. *Balaenoptera bertae*. A, ventral view.
339 B, right lateral view. C, left lateral view. D, dorsal view. E, anterior view. F,
340 posterior view.

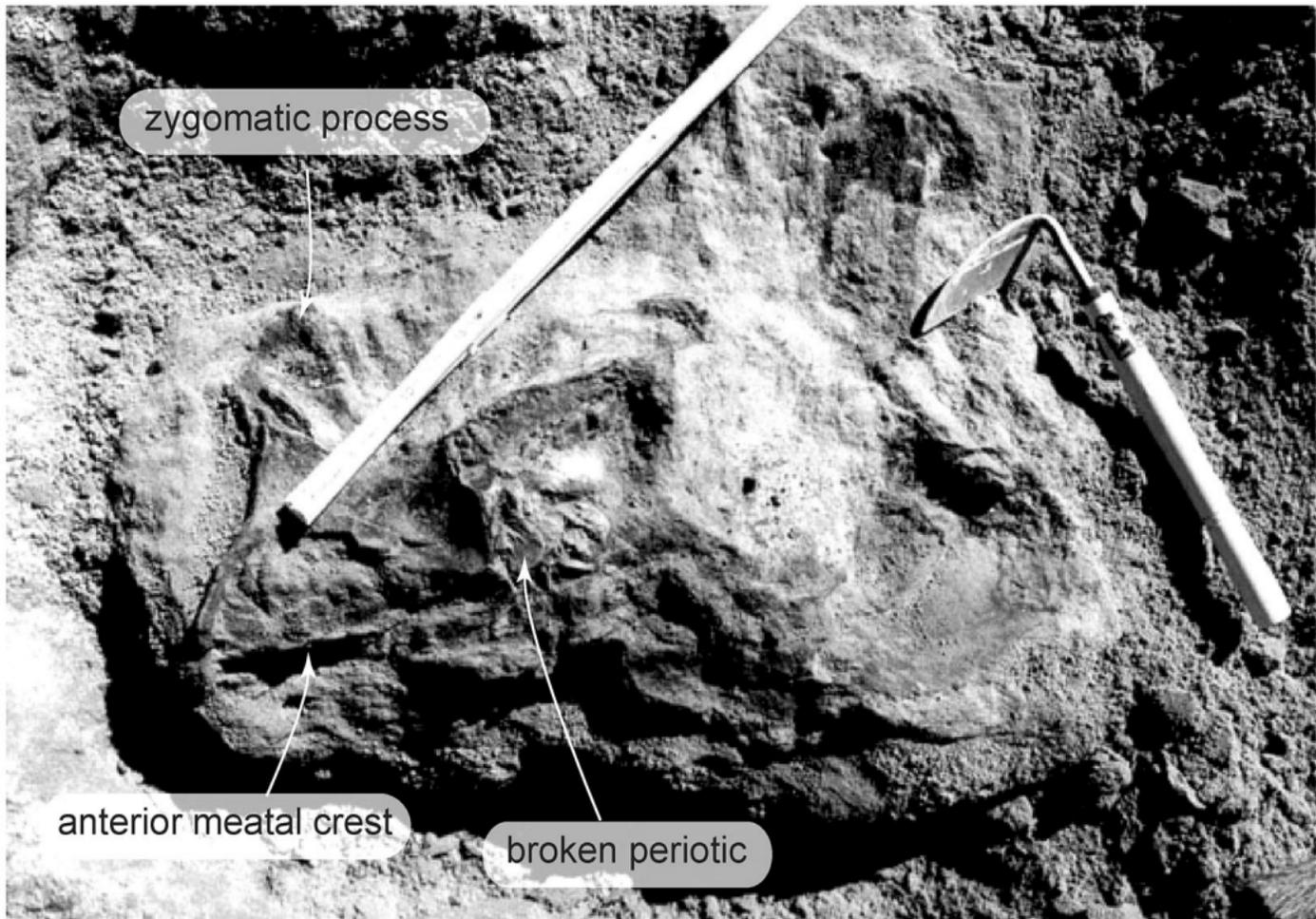
341 **Figure 4.** Squamosal and broken periotic of YPM 11852, aff. *Balaenoptera*
342 *bertae* in ventral view.

343 **Table 1.** Measurements of skull, aff. *Balaenoptera bertae* (YPM 11852) in mm.
344+ are maximum preserved measurements of broken parts. * means only right side.

345 **Table 2.** Compared named fossil balaenopterids. * means directly observed
346 specimens.

347

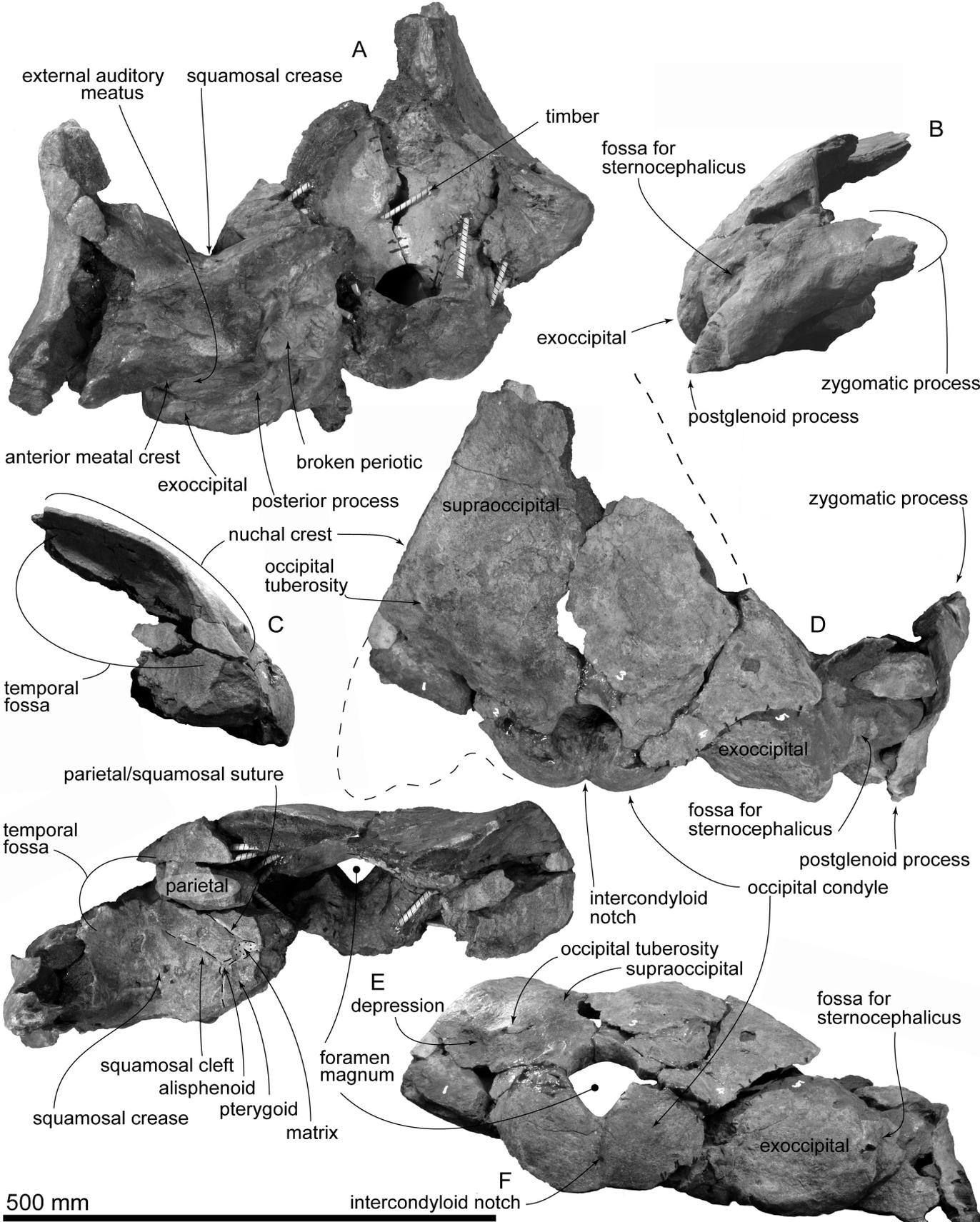


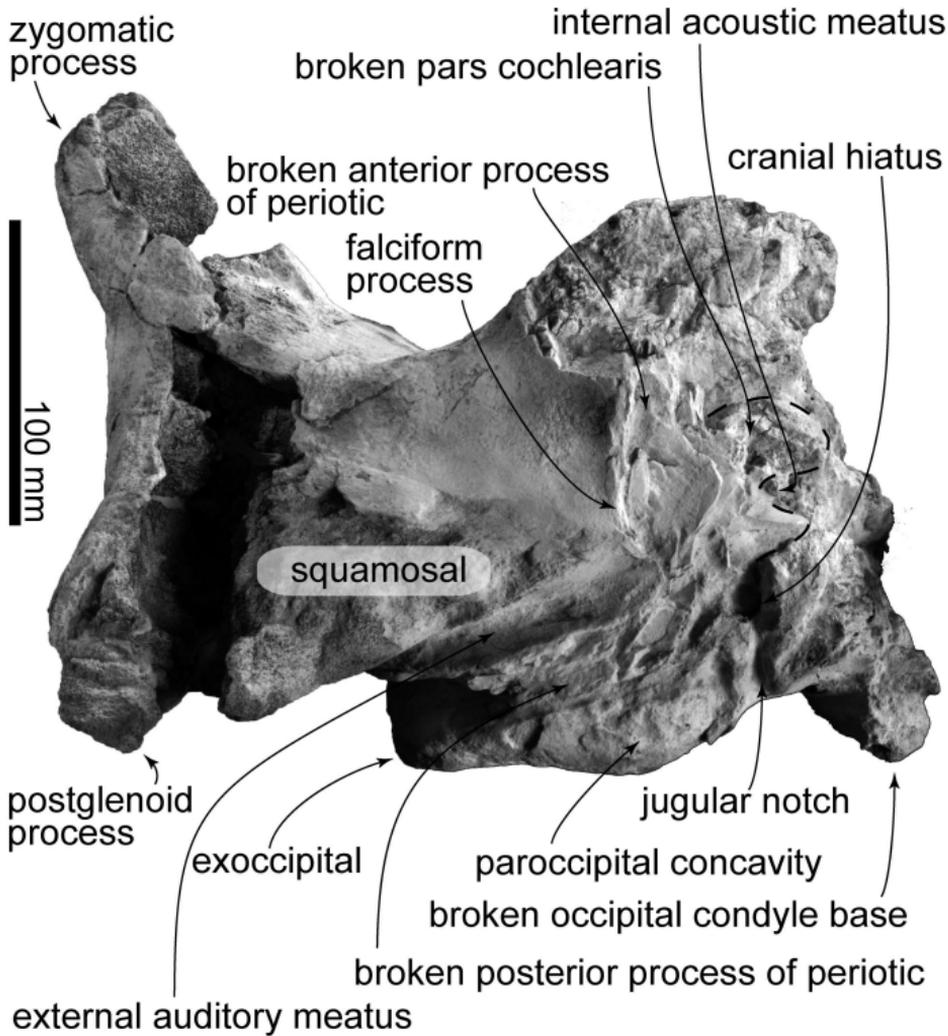


zygomatic process

anterior meatal crest

broken periotic





Squamosal length	230 +
Height between postglenoid process to dorsal border of zygomatic process	110 +
Median line to lateral border of exoccipital	240 *
Median line to lateral border of zygomatic process	355 *
Maximum width of occipital condyles	180
Foramen magnum height	65
Foramen magnum width	58
Preserved maximum length, from ventral border of occipital condyles to anterior most of supraoccipital	395

Accepted manuscript

Scientific name	Age	Localities	Reference
YPM 11852, aff. <i>Balaenoptera bertae</i>	Late Pliocene to Early Pleistocene	Yamagata, Japan	This study *
<i>Archaeobalaenoptera liesselensis</i>	Late Pleistocene	Noord Brabant, The Netherlands	Bisconti et al, 2020a
<i>Balaenoptera bertae</i>	Late Pliocene	Half Moon Bay, California	Boessenecker, 2013a *
<i>Marzanoptera tersillae</i>	Pliocene	Serra Domenico locality, Italy	Bisconti et al, 2020b
<i>Protororqualus wilfriedneesi</i>	Early Pliocene	Wommelgem, Belgium	Bisconti and Bosselaers, 2020
<i>Archaeobalaenoptera castriarquati</i>	Early Pliocene	Rio Carbonari, Italy	Bisconti, 2007; Bisconti et al, 2020a
<i>Plesiobalaenoptera hubachi</i>	Early Pliocene	Bahia de Guayacan, Chile	Dathe, 1983; Bisconti et al, 2020b
<i>Fragilicetus velponi</i>	Early Pliocene	Antwerp, Belgium	Bisconti and Bosselaers, 2016
<i>Diunatans luctoretemergo</i>	Early Pliocene	Antwerp, Belgium	Bosselaers and Post, 2010
<i>Miobalaenoptera numataensis</i>	Late Miocene	Hokaido, Japan	Tanaka and Watanabe, 2019 *
<i>Parabalaenoptera baulinensis</i>	Late Miocene	Balinas Point, California	Zeigler et al, 1997 *
<i>Nehalaennia devossi</i>	Late Miocene	Westerschelde, the Netherlands	Bisconti et al, 2019
<i>Incakujira anillodefuego</i>	Late Miocene	Aguada de Lomas, Peru	Marx and Kohno, 2016 *