

Endocranial anatomy and life habits of the Early Triassic archosauriform *Proterosuchus fergusi*

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3 1 **Endocranial anatomy and life habits of the Early Triassic archosauriform**
4
5 2 *Proterosuchus fergusi*
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12 5 EMILY E. BROWN^{1*}, RICHARD J. BUTLER¹, MARTÍN D. EZCURRA^{1,2}, BHART-
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40 17 **Abstract:** Proterosuchids are an important group of carnivorous basal archosauriforms
41
42 18 characterised by a bizarre and enigmatic downturned premaxilla that overhangs the lower jaw.
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44 19 They are particularly significant because they radiated in the immediate aftermath of the
45
46 20 Permian–Triassic mass extinction, and represent one of the best known ‘disaster taxa’
47
48 21 following that event. While traditionally considered semi-aquatic, recent histological studies
49
50 22 and geological data have suggested that they more likely inhabited terrestrial environments.
51
52 23 By utilising computed tomographic (CT) data, we virtually reconstruct the brain endocast and
53
54 24 endosseous labyrinths of two adult specimens of *Proterosuchus fergusi* from the earliest
55
56 25 Triassic of South Africa, in an attempt to understand its life habits within the context of basal
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3 26 archosauriform evolution. Endocasts reveal that the brain cavity is tubular in shape and the
4
5 27 endosseous labyrinths are highly pyramidal. The angle of the lateral semi-circular canal
6
7 28 suggests that *P. fergusi* naturally held its head upwards $\sim 17^\circ$, while the length of the cochlear
8
9 29 duct suggests its auditory abilities were specialised towards low-frequency sounds.
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11
12 30 Furthermore, beam theory analysis suggests that the rostrum of *P. fergusi* is highly resistant to
13
14 31 both bending and torsion when compared to modern crocodylians, although this resistance is
15
16 32 neither enhanced or reduced by the overhanging premaxilla. Comparative anatomical analyses
17
18 33 suggest *P. fergusi* was likely a semi-aquatic, generalist apex predator capable of surviving the
19
20 34 harsh environmental perturbations of the Early Triassic.
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27 36 KEYWORDS: Archosauriformes, endocranial anatomy, brain cavity, semi-circular canals,
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29 37 digital reconstruction, Early Triassic
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35
36 40 The end-Permian mass extinction is thought to have resulted in the loss of up to $\sim 96\%$ of life
37
38 41 on Earth, including $\sim 70\%$ of terrestrial vertebrates (Raup 1979; Jablonski 1995; Benton &
39
40 42 Twitchett 2003; Chen & Benton 2012). Despite the desolate greenhouse conditions of the
41
42 43 Early Triassic, the opening up of new ecosystems and niches allowed for an adaptive
43
44 44 radiation of surviving clades (Chen & Benton 2012; Viglietti *et al.* 2013; Foth *et al.* 2016;
45
46 45 Ezcurra & Butler 2018). Originating in the middle–late Permian, the diapsid clade
47
48 46 Archosauriformes successfully gained a foothold in the Early Triassic world, and gave rise to
49
50 47 groups that would go on to dominate in the Mesozoic (e.g. crocodylomorphs, dinosaurs,
51
52 48 pterosaurs) as well as continuing to make up a considerable portion of the modern fauna (e.g.
53
54 49 birds, crocodylians) (Ezcurra *et al.* 2014; Pinheiro *et al.* 2016; Ezcurra & Butler 2018).
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3 50 The basal archosauriform group Proterosuchidae, characterised by an enigmatic
4
5 51 downturned premaxilla that overhangs the anterior margin of the lower jaw, is often referred
6
7 52 to as a ‘disaster’ clade that thrived for a geologically short interval following the end-Permian
8
9 53 extinction (e.g. Ezcurra 2016; Button *et al.* 2017). The clade attained a near cosmopolitan
10
11 54 distribution, with specimens found in China, India, and South Africa (Ezcurra *et al.* 2013;
12
13 55 Ezcurra 2016). The type species of this clade, *Proterosuchus fergusi* Broom 1903, is known
14
15 56 from the *Lystrosaurus* Assemblage Zone (AZ) of the Karoo Supergroup of South Africa.
16
17 57 Although well sampled, with eleven highly informative specimens, many aspects of the
18
19 58 species’ life habits and functional morphology are still unknown (Ezcurra & Butler 2015a).
20
21 59 Traditionally, *P. fergusi* was considered semi-aquatic due to its superficial similarity to
22
23 60 modern crocodylians and the then-presumed wet climate of the Early Triassic Karoo Basin
24
25 61 (Broili & Schröder 1934; Tatarinov 1961; Reig 1970). Further evidence for an aquatic affinity
26
27 62 included the predominately horizontal orientation of vertebral zygapophyses, allowing for
28
29 63 lateral flexure of the body, which crocodiles use to propel themselves through the water
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31 64 (Cruickshank 1972).
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38 65 However, proterosuchid ecology has since been disputed, with several authors
39
40 66 suggesting a more terrestrial life habit (Cruickshank 1972; Welman 1998; Botha-Brink &
41
42 67 Smith 2011). Cruickshank (1972) and Welman (1998) noted the lack of dorsally positioned
43
44 68 external nares, which are an adaptation typical of aquatic and semi-aquatic diapsids such as
45
46 69 plesiosaurs, mosasaurs, phytosaurs and crocodylians (Sereno 1991; Nesbitt *et al.* 2009).
47
48 70 Cruickshank (1972) also noted the presence of well-ossified limbs, carpus and tarsus, and the
49
50 71 vertical orientation of the occipital elements, and suggested *P. fergusi* was largely terrestrial.
51
52 72 Furthermore, Botha-Brink & Smith (2011) combined sedimentological data and an
53
54 73 osteohistological analysis of several *P. fergusi* limb bones, which provided no evidence of
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56 74 osteological specialisation to aquatic life (such as pachyosteosclerosis or osteoporosis), to
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3 75 suggest a terrestrial mode of life for this species. However, palaeohistological analyses are not
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5 76 always accurately indicative of ecology, with, for example, a similar study finding no aquatic
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7 77 osteological specialisations in the marine teleosaurids, *Steneosaurus* and *Teleosaurus* (Hua &
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9 78 De Buffrenil 1996).

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13 79 *Proterosuchus fergusi* is both the earliest new tetrapod following the onset of the
14
15 80 Triassic and the best-sampled basal archosauriform species from the Karoo Basin (Smith &
16
17 81 Botha 2005; Botha-Brink & Smith 2011; Smith *et al.* 2012). Therefore, consensus on the life
18
19 82 habits of this species is crucial to understanding the faunal recovery following the end-
20
21 83 Permian mass extinction. While the neuroanatomy and inner ear morphology of archosaurs
22
23 84 have been extensively studied, non-archosaurian archosauromorphs have been widely
24
25 85 neglected, in part due to their comparatively poor fossil record. To date, the brain endocast of
26
27 86 only one basal archosauriform has been described in the literature, the proterochampsian
28
29 87 *Tropidosuchus romeri* (Trotteyn & Paulina-Carabajal 2016). Several basal archosauriform
30
31 88 endosseous labyrinths have been virtually reconstructed (Sobral *et al.* 2016a; Trotteyn &
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33 89 Paulina-Carabajal 2016); however, none are complete.

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39 90 This study presents the first in-depth assessment of the endocranial anatomy of the
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41 91 basal archosauriform *P. fergusi*. Based on data from the endocranial reconstructions, we
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43 92 further discuss the life habits and ecology of *P. fergusi* and implications for the early
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45 93 evolution of Archosauriformes.

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50 51 95 **METHODS**

52 53 54 96 *Specimens*

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57 97 The skulls of two specimens of *Proterosuchus fergusi* were used in this study, RC 846
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59 98 (Rubidge Collection, Wellwood, Graaff-Reinet, South Africa) and SNSB-BSPG 1934 VIII

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3 99 514 (Staatliche Naturwissenschaftliche Sammlungen Bayerns/Bayerische Staatssammlung für
4
5 100 Paläontologie und Geologie, Munich, Germany) (Fig. 1). Both specimens are large adults
6
7 101 from the *Lystrosaurus* AZ of the Karoo Basin of South Africa (Ezcurra & Butler 2015b). The
8
9 102 left side of SNSB-BSPG 1934 VIII 514 is nearly complete, but the right side has been subject
10
11 103 to severe deformation and loss of information (Fig. 1B-C). The specimen also previously
12
13 104 underwent substantial reconstruction in an attempt to adhere disarticulated and fractured
14
15 105 elements, especially in the premaxillary region (Broili & Schröder 1934). RC 846 is far more
16
17 106 complete but the skull is moderately transversely compressed and its posterior region has
18
19 107 been moderately crushed. The premaxilla is also largely complete and articulated in this
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21 108 specimen.
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30 110 *Scanning*

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32
33 111 RC 846 was CT scanned at the University of Texas High-Resolution X-ray CT Facility
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35 112 Archive. The braincase of RC 846 is disarticulated from the rest of the skull, allowing it to be
36
37 113 μ CT scanned separately (hereafter 'RC 846 μ CT'). SNSB-BSPG 1934 VIII 514 was CT
38
39 114 scanned at the Klinikum rechts der Isar (Munich). Datasets consist of 548 coronal slices (1024
40
41 115 x 1024 pixels, voxel size 0.211 mm) for RC 846, 457 oblique ($\sim 15^\circ$ ventroposterior deviation)
42
43 116 coronal slices (1024 x 1024 pixels, voxel size 0.117 mm) for RC 846 μ CT, and 1229 coronal
44
45 117 slices (768 x 526 pixels, voxel size 0.5 mm) for SNSB-BSPG 1934 VIII 514. Original CT
46
47 118 data for SNSB-BSPG 1934 VIII 514 are available in the Dryad Digital Repository:
48
49
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51 119 <https://datadryad.org/review?doi=doi:10.5061/dryad.XXXX>. Original CT data for RC 846 are
52
53 120 archived at Yale University and are available from B.-A.S. upon request.
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122 *Virtual Endocast Construction*

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3 123 The CT data were imported into SPIERSedit (2.20, www.spiers-software.org). Areas of
4
5 124 interest such as the endosseous labyrinths, braincase and nerves were manually segmented
6
7 125 using the Masks and Curves tools following Balanoff *et al.* (2016) (see Brown *et al.* 2019, fig.
8
9 126 S1). Interactive 3D PDFs of the endocranial reconstructions are provided as supplementary
10
11 127 information.
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18 129 *Morphometric Outline Analysis*

20
21 130 Elliptical Fourier analysis (EFA) was used in this study to quantify the morphological
22
23 131 variability of brain and inner ear endocasts among diapsid groups. EFA statistically compares
24
25 132 the co-ordinates of complex 2D shapes or outlines. EFA is very versatile as it can be used to
26
27 133 compare shapes where homologous features are hard to distinguish (Crampton 1995). As a
28
29 134 result, EFA is widely used in both the palaeontological and biological sciences to study
30
31 135 interspecific (e.g. Crampton 1995; Bonhomme *et al.* 2013; Vidal *et al.* 2014; Lautenschlager
32
33 136 2014; Lautenschlager *et al.* 2018) and intraspecific (e.g. Polihronakis 2006; Ramajo *et al.*
34
35 137 2013) anatomical variation.
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43 139 *Brain Outline Preparation.* An outline of the most complete *Proterosuchus fergusi*
44
45 140 endocranial reconstruction (RC 846) was drawn in Adobe Illustrator (CS5, www.adobe.com).
46
47 141 The brain cavities of 69 extinct and extant archosauriforms compiled from the literature were
48
49 142 also outlined in Adobe Illustrator. For consistency, the left lateral view was always used. If
50
51 143 unavailable, the right lateral view was reversed. Natural, artificial (e.g. latex) and virtual
52
53 144 endocasts were used for comparative taxa. Brain cavity outlines and taxonomic information of
54
55 145 all comparative taxa are available in Table S1. Outline co-ordinates were digitised in tpsDig2
56
57 146 (v2.31, Rohlf 2010) as 1000 x/y pairs.
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6 148 *Inner Ear Outline Preparation.* The inner ear is divided into two regions, both with different
7
8 149 functions: the semi-circular canals are part of the vestibular region, responsible for balance
9
10 150 and co-ordination; whereas the endosseous cochlear duct (ECD) is part of the auditory
11
12 151 system, responsible for hearing. To determine whether these features have evolved separately
13
14
15 152 the inner ear was outlined and analysed both with and without the ECD. Analysis without the
16
17 153 ECD also allowed the inclusion of the basal archosauriform *Euparkeria capensis*, for which
18
19 154 the ECD could not be reconstructed (Sobral *et al.* 2016a). Prior to this study, *E. capensis* was
20
21 155 the only non-archosaurian archosauriform for which three articulated semi-circular canals had
22
23 156 been virtually reconstructed.
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27 157 For this analysis, a computerised outline procedure was used. Images of the left
28
29 158 labyrinth from extinct and extant diapsid taxa (n = 94, with ECD; n = 99, without ECD) were
30
31 159 collected from the literature. These were then converted into binary images by altering the
32
33 160 threshold in Adobe Photoshop (CS5, www.adobe.com). At this stage for the analysis
34
35 161 excluding the ECD, the polygonal lasso tool was used to select and remove the ECD. Images
36
37 162 were then vectorised using the 'Live Trace' tool in Adobe Illustrator before using the expand
38
39 163 tool to make the vector editable. The 'Outline Stroke' tool was then used to create a
40
41 164 computer-generated outline. All endosseous labyrinths were outlined in lateral view. If the left
42
43 165 labyrinth was unavailable, the right labyrinth was reversed. Inner ear outlines and taxonomic
44
45 166 information of all comparative taxa are available in the supplementary information. Co-
46
47 167 ordinates for the outlines were digitised in tpsDig2 as 1000 x/y pairs for outlines with the
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49 168 ECD and 750 x/y pairs for outlines without the ECD.
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3 170 *Elliptical Fourier Analysis*. Outline co-ordinates were analysed respectively in PAST
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5 171 (v3.19, Hammer *et al.* 2001) using EFA. Outlines were smoothed ten times to eliminate pixel
6
7 172 noise, and 23 Fourier harmonics were found to describe the outlines of all sampled taxa
8
9 173 sufficiently (average Fourier power > 99%). A Procrustes superimposition was performed
10
11 174 before undergoing principal component analysis (PCA). Taxa were assigned to a phylogenetic
12
13 175 group and broad ecological group (aquatic, semi-aquatic, terrestrial) (see Table S1). To test
14
15 176 whether morphological variation was significantly different between group variables
16
17 177 (ecological and phylogenetic) the non-parametric one-way PERMANOVA test was used with
18
19 178 10,000 permutations. Due to the ambiguity surrounding the ecology of *Proterosuchus fergusi*,
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21 179 this taxon was excluded from analyses between ecology.
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33 182 *Morphometric Landmark Analysis*

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35 183 This analysis is an iteration of that used by Yi & Norell (2015), Neenan *et al.* (2017) and
36
37 184 Neenan *et al.* (2019) to compare the endosseous labyrinths of extant snakes, extinct
38
39 185 sauropterygians and *Massospondylus* individuals, respectively. However, unlike those
40
41 186 mentioned this analysis compares each of the different semi-circular canals (anterior semi-
42
43 187 circular canal, ASC; lateral semi-circular canal, LSC; posterior semi-circular canal, PSC)
44
45 188 separately.
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50 189 The semi-circular canals generated from RC 846 μ CT and RC 846 were used for this
51
52 190 analysis. Additionally, the inner ears of comparative extinct and extant diapsid taxa (n=37,
53
54 191 ASC; n=51, LSC; n=35, PSC) were sourced from the literature. The left labyrinth was
55
56 192 selected for this analysis due to its relative availability, but if unavailable the right labyrinth
57
58 193 was reversed.
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3 194 Screenshots of each semi-circular canal were taken perpendicular to the plane to
4
5 195 remove perspective skew. The external and internal surface of each canal was outlined in
6
7 196 Adobe Illustrator using the automated approach described above. The dashed line stroke tool
8
9
10 197 was used to find 21 equally-spaced points along the external surface of each canal and 11
11
12 198 along the inner surface of each canal. These dashed lines were used as a reference point for
13
14 199 accurate and consistent landmark placing. The outlines with their reference points were
15
16 200 imported into tpsDig2 and landmarks were placed in a designated order. Co-ordinates were
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18 201 exported from tpsDig2 and imported into MorphoJ (v1.06d,
19
20 202 www.flywings.org.uk/morphoj_page.htm). To mitigate the effects of varying orientation,
21
22 203 sizing, and distance between landmarks of different canals, a Procrustes superimposition was
23
24 204 performed (following Neenan *et al.* 2017). Principal component analysis was run and plotted
25
26 205 to show geometric variability. A canonical variate analysis (CVA) was used to test the
27
28 206 morphological variance of diapsid semi-circular canals between different phylogenetic and
29
30 207 ecologic groupings, respectively. Due to the ambiguity surrounding the ecology of
31
32 208 *Proterosuchus fergusi*, this taxon was excluded from analyses between ecology.
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209

210 *Head posture*

211 The orientation of the LSC ('horizontal semi-circular canal') has been widely used to infer the
212 head posture of fossil taxa (Rogers 1998; Witmer *et al.* 2003; Sampson & Witmer 2007;
213 Sereno *et al.* 2007; Witmer *et al.* 2008; Witmer & Ridgely 2009; Neenan & Scheyer 2012;
214 Benoit *et al.* 2017), however not without criticism (Hullar 2006; Taylor *et al.* 2009; Marugán-
215 Lobón *et al.* 2013). As part of the vestibular system, it is thought that many species habitually
216 hold their LSC in a horizontal orientation (Lebedkin 1924; de Beer 1947). For example, the
217 LSC orientation in the basal sauropterygian *Placodus gigas* suggests that it was down tilted

1
2
3 218 ~20° (Neenan & Scheyer 2012). Neenan & Scheyer (2012) suggested that the downturned
4
5 219 head posture was adapted for aquatic herbivory and concluded that *Pl. gigas* was fully
6
7 220 adapted for aquatic life despite being one of the most basal sauropterygians.
8
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13 222 *Orientation of the lateral semi-circular canal.* Screenshots of the isolated LSC and whole
14
15 223 skull (skull at 75% opacity) were taken in SPIERS View to show both the skull and LSC. The
16
17 224 orientation of each LSC was measured in ImageJ using the angle tool. Lateral semi-circular
18
19 225 canals reconstructed from RC 846 μ CT were excluded from this analysis because the scans
20
21 226 only contained the braincase region. Often in these analyses, the ‘horizontal’ is measured as
22
23 227 the bottom surface of the dentary (e.g. Witmer & Ridgely 2009). However, as both of the
24
25 228 mandibles in RC 846 and SNSB-BSPG 1934 VIII 514 are significantly deformed, the
26
27 229 maxillary tooth-line was used instead as the horizontal plane. The left and right LSC planes
28
29 230 were measured separately. The mean average was calculated from all four values.
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36 37 38 232 *Auditory Abilities*

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40
41 233 The mean hearing frequency and hearing range of *Proterosuchus fergusi* were estimated
42
43 234 following methods from Walsh *et al.* (2009). The endosseous cochlear duct (ECD) length is
44
45 235 considered a proxy of auditory ability. The study from Walsh *et al.* (2009) used a linear
46
47 236 regression of scaled ECD length and sensitivity data of extant avian and reptilian taxa to
48
49 237 estimate the auditory abilities of several extinct Aves. This has since been repeated with the
50
51 238 Early Jurassic marine crocodylomorph *Steneosaurus cf. gracilirostris* (Brusatte *et al.* 2016)
52
53 239 and with the basal eusuchian *Lohuecosuchus megadontos* (Serrano-Martínez *et al.* 2018). As
54
55 240 individuals can typically only hear within their own vocal range, estimations of auditory
56
57 241 acuity can be informative regarding vocality (Narins *et al.* 2004). Vocality and auditory
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1
2
3 242 ability are also considered to be indicators of sociality and cognitive ability, since larger more
4
5 243 complex groups require better communicative skills to work together efficiently (Blumstein &
6
7 244 Armitage 1997; Freeberg *et al.* 2012; Freeberg & Krams 2015; Sewall 2015). This is
8
9 245 corroborated by Walsh *et al.* (2009) who found longer ECD length to be significantly
10
11 246 correlated with animals living in larger group sizes.
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15 247
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18 248 *Regressions.* This study uses ECD length (scaled to basicranial length) and hearing sensitivity
19
20 249 data derived from Walsh *et al.* (2009). Additional data on the hearing sensitivity of extant
21
22 250 archosaurs was also collected from the literature (Corfield *et al.* 2013; Bonke *et al.* 2015)
23
24 251 giving a total of 26 comparative taxa. RC 846 produced the most complete endocast and so
25
26 252 was the only proterosuchid specimen used in the analysis. The ECD length of RC 846 was
27
28 253 measured and scaled to the basicranial length (measured from the basisphenoid-presphenoid
29
30 254 suture to the caudal-most tip of the occipital condyle, S. Walsh 2018 pers. comm.).
31
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35 255 All scaled ECD lengths were log-transformed. Log-scaled ECD length for extant taxa
36
37 256 was plotted against respective mean hearing and hearing range data. Both an ordinary least
38
39 257 squares linear regression and a phylogenetic generalized least square regression were
40
41 258 calculated for this analysis. These regression lines were used to predict mean hearing and
42
43 259 hearing range for *Proterosuchus fergusi* based on the scaled and log-transformed ECD length.
44
45 260 Analyses were conducted using R (RStudio v1.1.423, www.rstudio.com). The R packages ape
46
47 261 (5.3, Paradis *et al.* 2004), geiger (2.0.6.1, Harmon *et al.* 2007), nlme (3.1-137, Pinheiro *et al.*
48
49 262 2018) and phytools (0.6-60, Revell 2012) were used to calculate the phylogenetic generalized
50
51 263 least square (PGLS) regression with a Brownian motion model of trait evolution. The analysis
52
53 264 used a phylogenetic tree of all comparative taxa generated from <http://timetree.org/>.
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3 266 *Snout Beam Analysis*
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6 267 Beam theory was used to investigate the relative rostral resistance of *Proterosuchus fergusi* to
7
8 268 bending and torsion. Beam theory calculates the potential bending of an object based on the
9
10 269 density and distribution of material around the neutral axis (the centre of an object, where
11
12 270 there is no tension or compression) (Therrien *et al.* 2005; Cuff & Rayfield 2013). For
13
14 271 example, a hollow tube is more resistant to bending than an infilled tube because material is
15
16 272 distributed further from the neutral axis. To test the biomechanical resistance of rostra and
17
18 273 lower jaws, this technique has been previously applied to a range of fossil taxa (Therrien *et al.*
19
20 274 2005; Cuff & Rayfield 2013; Foffa *et al.* 2014).
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28 276 *Slice Preparation.* Slices were chosen along the rostrum at intervals. The first 20% of the
29
30 277 rostrum was sliced at intervals of 2% as the premaxillary area is of specific interest in
31
32 278 *Proterosuchus fergusi*. Due to the extent of the overhanging premaxilla in *P. fergusi*, a
33
34 279 straight beam may not be considered appropriate for this taxon. Therefore, for the rostral 20%
35
36 280 of the snout both a straight beam and curved beam were used in the analysis for *P. fergusi*.
37
38 281 For the curved beam analysis, the rostrum of *P. fergusi* was sliced at intervals of 4% rather
39
40 282 than 2% to prevent slices overlapping. The latter 80% of the rostrum was sliced at intervals of
41
42 283 10%. RC 846 was used for this analysis because the premaxillae of SNSB-BSPG 1934 VIII
43
44 284 514 have undergone significant reconstruction and may not represent the true original shape
45
46 285 (Fig. 1). As RC 846 was scanned along the coronal axis, the raw CT slices could be used in
47
48 286 this analysis.
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54 287 For comparative analysis, five extant taxa were chosen: four pseudosuchians, *Alligator*
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56 288 *mississippiensis* (American alligator), *Crocodylus moreletii* (Morelet's crocodile), *Crocodylus*
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58 289 *rhombofer* (Cuban crocodile) and *Tomistoma schlegelii* (false gharial); and one
59
60

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3 290 actinopterygian, *Atractosteus spatula* (alligator gar). *Atractosteus spatula* was included
4
5 291 because of the slight overhang of the premaxilla, similar to that of *P. fergusi*.
6
7

8 292 The skull of *A. spatula* was downloaded as CT data from Digimorph.org. The
9
10 293 specimen was scanned along the coronal plane so slices for this analysis could be taken
11
12 294 directly from the raw CT-scans. Skulls of *C. moreletii*, *C. rhombifer*, and *T. schlegelii* were
13
14 295 downloaded as STL files from Digimorph.org. *Alligator mississippiensis* (juvenile) was
15
16 296 downloaded from Digimorph.org as CT data sliced along the horizontal plane so the raw
17
18 297 slices could not be used in this analysis. *Alligator mississippiensis* was exported from
19
20 298 SPIERSview as an STL file. These four STL files were imported into Blender (2.79a,
21
22 299 www.blender.org) and manually sliced using the Boolean Modifier tool. A visual reference
23
24 300 was used to accurately slice at the correct positions along the rostrum. All slices were
25
26 301 rendered against the same flat plane for consistency. For the curved beam analysis RC 846
27
28 302 was imported into Avizo Lite (9, Thermo Fisher Scientific) and the rotate tool was used create
29
30 303 the required slices.
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36 304 All slices were imported into Adobe Photoshop where mandibles were manually
37
38 305 removed. Teeth can vastly increase the proportion of material in the slice, especially in
39
40 306 specimens with larger teeth (Cuff & Rayfield 2013). To standardise these effects, teeth were
41
42 307 removed and the alveoli flattened in all slices. Additionally, in *P. fergusi*, the sediment was
43
44 308 manually removed from the slices and rostrum walls were slightly corrected by
45
46 309 symmetrisation. All slices were then converted into binary images by altering the image
47
48 310 threshold.
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56 312 *Second moments calculation and analysis.* Prepared slices were imported into ImageJ (1.51j8,
57
58 313 www.imagej.nih.gov/ij). The second moments of area in the dorsoventral (I_x) and
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1
2
3 314 mediolateral (Iy) directions and the polar moment of inertia (J) were calculated using the
4
5 315 ImageJ macro, MomentMacroJ (v1.4, www.hopkinsmedicine.org/fae/mmacro.html). In an
6
7 316 additional analysis, rostra were scaled to the rostral length of *Proterosuchus fergusi* (aspect
8
9 317 ratio was maintained) to determine whether size difference influenced results (see
10
11 318 Supplementary material). All results were log transformed and plotted using R. A paired t-test
12
13 319 was computed between *P. fergusi* and comparative taxa for the entire rostrum and the first
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15 320 anterior 20% of the rostrum both at true size and scaled size and also using both a straight and
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17 321 curved beam.
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323 RESULTS

324

325 *Virtual Endocast Reconstruction*

326 *Brain Cavity.* The endocranial reconstructions of *Proterosuchus fergusi* are generally linear in
327 shape and organisation (Fig. 2), similar to that of modern-day adult crocodylians (Jirak &
328 Janacek 2017). The brain cavity of *P. fergusi* is straighter than many archosauriforms
329 including erythrosuchids (Gower & Sennikov 1996), phytosaurs (Holloway *et al.* 2013;
330 Lautenschlager & Butler 2016) and many crocodylomorphs (Witmer *et al.* 2008; Witmer &
331 Ridgely 2009; Kley *et al.* 2010), but analogous to the thalattosuchians *Pelagosaurus typus*
332 and *Steneosaurus gracilirostris* (Brusatte *et al.* 2016; Pierce *et al.* 2017).

333 Due to breaks at the frontal-parietal suture in both RC 846 and SNSB-BSPG 1934
334 VIII 514, the elongate olfactory tracts typical of many archosauriforms and non-
335 archosauriform diapsids (Witmer *et al.* 2008; Witmer & Ridgely 2009; Ezcurra 2014;
336 Lautenschlager & Butler 2016; Trotteyn & Paulina-Carabajal 2016) could not be fully
337 reconstructed in *P. fergusi*. The endocast for RC 846 μ CT lacks the anterior portion of the

1
2
3 338 olfactory region, which would lie on the ventral surface of the frontals. This region is also
4
5 339 missing in SNSB-BSPG 1934 VIII 514 due to the poor preservation of the frontals. Shallow
6
7 340 concave fossae along the ventral surface of the frontals of RC 846 allow for the partial
8
9 341 reconstruction of the dorsal surfaces of the olfactory bulbs (Fig. 2C-D). The ventral extent of
10
11 342 the bulbs cannot be determined due to the lack of constraining bones in the region. The bulbs
12
13 343 are slightly laterally expanded and rostrally taper out into two separate structures. Unlike in
14
15 344 the basal archosauromorph *Tasmaniosaurus triassicus* (Ezcurra 2014), the olfactory bulbs of
16
17 345 *P. fergusi* are exceeded in width by the cerebral hemispheres. The morphology of these bulbs
18
19 346 allows for an estimation of the size and extent of the olfactory tracts and olfactory region as a
20
21 347 whole. Compared to other archosauriforms the olfactory region is relatively short, analogous
22
23 348 to the thalattosuchian *Pelagosaurus typus* (Pierce *et al.* 2016), but considerably larger than the
24
25 349 ornithosuchid *Riojasuchus tenuisiceps* (Baczko & Desojo 2016).

30
31 350 The forebrain is bulbous and horizontal while the mid-brain is more anteroventrally
32
33 351 directed. The floccular lobes are relatively small and do not extend through the anterior semi-
34
35 352 circular canal, unlike in many archosaurs (e.g. Witmer & Ridgely 2009; Lautenschlager *et al.*
36
37 353 2012; Lautenschlager & Butler 2016). The hind brain is mediolaterally narrow, especially
38
39 354 between the endosseous labyrinths, but expands ventrally more than other parts of the
40
41 355 braincase. Ventral to the hindbrain, pituitary fossae are partially preserved in both specimens,
42
43 356 but not sufficiently to reconstruct. The main body of the brain cavity (forebrain-hindbrain) sits
44
45 357 underneath the parietal and post-parietal bones.

46
47 358 The carotid artery canal could only be reconstructed in SNSB-BSPG 1934 VIII 514.
48
49 359 The carotid canal extends ventrally from the braincase before dividing into two canals that
50
51 360 then extend away from each other laterally. This is unlike the condition in many
52
53 361 archosauriformes where the carotid artery extends out from the pituitary fossa (Witmer *et al.*
54
55 362 2008; Witmer & Ridgely 2009; Lautenschlager & Butler 2016). This suggests the pituitary is

1
2
3 363 considerably reduced in *P. fergusi* and projects posteriorly. A bony margin separating the
4
5 364 pituitary from the diencephalon is not recognisable in the CT data.
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8 365 A small ventrally positioned nerve canal reconstructed in SNSB-BSPG 1934 VII I 514
9
10 366 may be the trochlear nerve canal (iv). However, it does not sit rostrally to the trigeminal nerve
11
12 367 (v) as occurs in phytosaurs (Lautenschlager & Butler, 2016). In RC 846 and SNSB-BSPG
13
14 368 1934 VIII 514 a group of nerves were reconstructed on the lateral surface of the endocast and
15
16 369 may be the three branches of the trigeminal nerve canal (v). It seems to be in a position
17
18 370 similar to the trigeminal nerve in thalattosuchians (Brusatte *et al.* 2016; Pierce *et al.* 2016) but
19
20 371 more anteriorly positioned than in phytosaurs (Lautenschlager & Butler 2016). Similar
21
22 372 reconstructions on RC 846 μ CT were considered too dorsally positioned to be cranial nerves
23
24 373 and so were identified as venous canals.
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29 374 Anterior to the carotid artery canal on SNSB-BSPG 1934 VIII 514 is a small
30
31 375 ventrolaterally-directed nerve canal, possibly the abducens nerve (vi). This nerve is also
32
33 376 reconstructed in RC 846. In all of the reconstructions a cranial nerve sits immediately anterior
34
35 377 to the endosseous labyrinths. This nerve is similar in position to what has been interpreted as
36
37 378 the facial nerve (vii) in many extinct and extant archosauriforms (Gower & Sennikov 1996;
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39 379 Sampson & Witmer 2007; Witmer *et al.* 2008; Witmer & Ridgely 2009; Knoll *et al.* 2012;
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41 380 Lautenschlager *et al.* 2012; Lautenschlager *et al.* 2014; Lautenschlager & Butler 2016; Pierce
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43 381 *et al.* 2017). The ventral portion of the braincase of RC 846 is poorly preserved, limiting the
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45 382 reconstructions available. The endosseous labyrinths are relatively large in comparison to the
46
47 383 braincase, making up on average 7.9% the volume of the brain cavity.
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56 385 *Inner Ear.* The endosseous labyrinths of all three specimens were reconstructed completely
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58 386 (Fig. 3). The vestibular regions (top portion) of the endosseous labyrinths of *Proterosuchus*
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3 387 *fergusi* have a pyramidal shape superficially comparable to the thalattosuchians *Steneosaurus*
4
5 388 *gracilirostris* (Brusatte *et al.* 2016) and *Pelagosuchus typus* (Pierce *et al.* 2017). While in
6
7 389 lateral view the anterior semi-circular canal (ASC) looks substantially larger than the
8
9 390 posterior semi-circular canal (PSC), the canals are actually relatively equal in their
10
11 391 proportions. This morphological feature is shared among many non-archosaurian
12
13 392 archosauriformes and early pseudosuchians (Brusatte *et al.* 2016; Lautenschlager & Butler,
14
15 393 2016; Pierce *et al.* 2016; Sobral *et al.* 2016b) but is thought to be a plesiomorphic trait lost in
16
17 394 extinct and extant crocodylomorphs, which tend to have considerably larger anterior canals
18
19 395 compared to the posterior canal (Georgi & Sipla 2008; Witmer *et al.* 2008; Witmer & Ridgely
20
21 396 2009; Pierce *et al.* 2017). The ASC and LSC are elliptical in shape, whereas the PSC is
22
23 397 straighter. The PSC of SNSB-BSPG 1934 VIII 514 has a pronounced ventrally-directed kink,
24
25 398 unlike the condition in RC 846. This feature is more pronounced in the right PSC and
26
27 399 therefore likely the result of post-mortem deformation of the specimen (Fig. 3C-D).
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34 400 The endosseous cochlear duct (ECD) is significantly less ventrally extended than in
35
36 401 other archosauriform reconstructions (e.g. Lautenschlager & Butler 2016; Brusatte *et al.*
37
38 402 2016; Pierce *et al.* 2017; Leahey *et al.* 2015; Witmer *et al.* 2008). The fenestra vestibuli
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40 403 (which connects the inner and middle ears) is visible in RC 846 and possibly in SNSB-BSPG
41
42 404 1934 VIII 514 and extends posterolaterally away from the labyrinth.
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47 48 49 406 *Morphometric Outline Analysis*

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52 407 *Brain Outlines.* EFA was used to compare the brain cavity endocast of *Proterosuchus*
53
54 408 *fergusi* to that of various extant and extinct archosauriforms. As the olfactory bulbs are only
55
56 409 reconstructed in RC 846, this endocast was outlined and analysed. The first three principal
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58 410 component (PC) axes (Fig. 4 & Brown *et al.* 2019, fig. S2-3) account for 79.2% of shape
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3 411 variation. *Proterosuchus fergusi* and the proterochampsian *Tropidosuchus romeri* (the only
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5 412 other non-archosaurian archosauriform with an endocast sufficiently complete enough to be
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7 413 included in the analysis; Trotteyn & Paulina-Carabajal 2016) fall well within the non-
8
9 414 phytosaur pseudosuchian morphospace in PC1 vs. PC2 (Fig. 4). Minimum spanning trees
10
11 415 show that *P. fergusi* is placed closest to *Sebecus icaeorhinus* (early–mid Eocene sebecid
12
13 416 crocodylomorph) and *Gavialis gangeticus* (gharial) in PC1 vs. PC2 (Brown *et al.* 2019, fig.
14
15 417 S4) and PC1 vs. PC3 (Brown *et al.* 2019, fig. S5), respectively. Both PC plots show clear
16
17 418 divisions between phylogenetic groups, with groups with similar ecological niches such as
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19 419 birds and pterosaurs, and phytosaurs and crocodylomorphs, overlapping each other in
20
21 420 morphospace. PERMANOVA tests show significant differences between the endocranial
22
23 421 morphology of semi-aquatic and terrestrial taxa in the analysis (Table 1) as well as significant
24
25 422 support for the separation in morphospace of many groups within the clade Archosauriformes
26
27 423 (Table 2).
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36 425 *Inner Ear Outlines*. EFA was used to test the anatomical variations between the inner ears of
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38 426 diapsids, both with (w/) and without (w/o) the ECD. The first two PC axes account for 70.1%
39
40 427 and 72.9% of shape variation with (Fig. 5A) and without the ECD (Fig. 5B), respectively. In
41
42 428 both PC plots, RC 846 overlaps in morphospace with aquatic, semi-aquatic and terrestrial
43
44 429 taxa. RC 846 μ CT similarly lies within all three morphospaces when the ECD is excluded
45
46 430 (Fig. 5B), but lies outside all morphospace when the ECD is included (Fig. 5A). Minimum
47
48 431 spanning trees (Brown *et al.* 2019, fig. S7A & S7B) show that both RC 846 and RC 846 μ CT
49
50 432 lie closest to terrestrial squamates, with the exception of RC 846 μ CT which comes out
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52 433 closest to the sauropodomorph *Antarctosaurus wichmannianus* when the ECD is excluded
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54 434 (Brown *et al.* 2019, fig. S7B).
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3 435 Plots show substantial morphospace overlap between different ecological groupings.
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5 436 However, PERMANOVA tests (Table 3) show significant differences between the inner ear
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7 437 morphology of taxa from all ecologies when the ECD is included, and between aquatic taxa
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9 438 and both terrestrial ($p = 0.0013$) and semi-aquatic taxa ($p = 0.0282$) when excluded.
10
11 439 Phylogenetic groupings are generally not very distinguishable in either iteration, but at higher
12
13 440 taxonomic ranks ‘archosauromorphs’ and ‘outgroup taxa’ (testudines, sauropterygians and
14
15 441 squamates) show a clear divergence when the ECD is included (Brown *et al.* 2019, fig. S8A)
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17 442 compared to when the ECD is excluded (Brown *et al.* 2019, fig. S8B). PERMANOVA tests
18
19 443 (Table 4) show that when the ECD is included, basal archosauriforms (which includes solely
20
21 444 *Proterosuchus fergusi* in this iteration) are significantly different to all groups apart from
22
23 445 sauropterygians ($p = 0.2562$), testudines ($p = 0.1189$) and phytosaurs (0.0995). However, when
24
25 446 the ECD is excluded, basal archosauriforms (which includes both *P. fergusi* and *Euparkeria*
26
27 447 *capensis* in this iteration) are significantly different to fewer groups, including more derived
28
29 448 archosauriform groups: Phytosauria ($p = 0.7363$), non-phytosaurian pseudosuchians ($p =$
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31 449 0.1340) and sauropods ($p = 0.0618$).
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41 *Morphometric Landmark Analysis*

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44 452 The PC plots (Fig. 6) demonstrate the variation in the morphology of the semi-circular canals
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46 453 of diapsids based on landmarks. The three PC axes (Fig. 6A-C) account for 81.2%, 74.1% and
47
48 454 80.8% of shape variation across the ASC, LSC and PSC, respectively. CVA results (Table 5)
49
50 455 support significant differences between ecological groupings in most cases, with terrestrial
51
52 456 and aquatic taxa differing significantly across all semi-circular canals ($p = 0.0002$, ASC; $p =$
53
54 457 0.0050 , LSC; $p = 0.0051$, PSC). RC 846 is recovered in morphospace occupied by terrestrial,
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56 458 semi-aquatic and aquatic taxa in ASC and LSC, but places outside occupied morphospace in
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3 459 the PSC plot. Whereas RC 846 μ CT is displaced from all occupied morphospace in both the
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5 460 PSC and LSC plots, but overlaps with aquatic taxa in the ASC plot (Fig. 6A). All three PC
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7 461 plots seem to separate taxa by phylogenetic group relatively well; however, CVA results show
8
9 462 very few significant results between phylogenetic groupings (Table S2). Across all semi-
10
11 463 circular canals, basal archosauriforms (solely *P. fergusi*) are only significantly different to
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13 464 Aves ($p= 0.0231$, ASC; $p= 0.0208$, LSC; $p= 0.0211$, PSC).
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20 466 *Head Posture*

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23 467 Relative to the horizontal, the average orientation of the LSC plane is 17.19° (Fig. 7A),
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25 468 suggesting *Proterosuchus fergusi* had a significantly upright ‘alert’ head posture (Fig. 7B).
26
27 469 This is in contrast to *Crocodylus johnstoni* (freshwater crocodile), which has a horizontal head
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29 470 posture when the LSC plane is aligned to the horizon (Fig. 7C; Witmer *et al.* 2008).
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36 472 *Auditory Abilities*

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39 473 Both the linear and PGLS regression (Fig. 8; solid line and dashed line, respectively) support
40
41 474 significant relationships between ECD length and both mean hearing and hearing range in
42
43 475 extant taxa (Fig. 8). On the basis of these relationships, the auditory abilities of *Proterosuchus*
44
45 476 *fergusi* have been estimated. Both regression lines reveal considerably different
46
47 477 reconstructions, with the linear regression predicting a mean hearing frequency of $\sim 525\text{Hz}$
48
49 478 and a hearing range of $\sim 780\text{Hz}$ (similar to that of *Alligator mississippiensis* and the
50
51 479 rhynchocephalian *Sphenodon punctatus*), and the PGLS regression predicting a mean hearing
52
53 480 frequency of $\sim 1275\text{Hz}$ and a hearing range of $\sim 2150\text{Hz}$ (similar to that of the terrestrial
54
55 481 squamate *Ptyodactylus hasselquistii*).
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3 482
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56 483 *Snout Beam Analysis*
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89 484 Plots generally show similar trends in all taxa, with second moment and moment of inertia
10
11 485 values generally increasing steadily posteriorly along the rostra (Fig. 9). Pertaining to the
12
13 486 anterior 20% of the rostrum of *Proterosuchus fergusi*, differences between the straight and
14
15 487 curved beam (Fig. 9, solid and dashed line, respectively) analysis are relatively consistent
16
17 488 across all plots; with the straight beam showing a more posteriorly positioned peak (~15%
18
19 489 along the rostrum length) compared to the curved beam (~4% along the rostrum length).
20
21
2223 490 *Tomistoma schlegelii* (false gharial) is the only comparative taxa that is significantly different
24
25 491 to *P. fergusi* across all iterations of the analysis and all moments of resistance (Table S4).
26
2728 492 Values of I_x , which represents resistance to dorsoventral bending, (Fig. 9A-B) are highest in
29
30 493 *Proterosuchus fergusi*, even when other taxa are size-corrected. True size plots show that
31
32 494 *Atractosteus spatula* (alligator gar) is the least resistant (Fig. 9A), but when scaled *Tomistoma*
33
34 495 *schlegelii* is the least (Fig. 9B). Paired t-test results from the straight beam iteration show that
35
36 496 there is no significant difference in resistance along the whole rostrum between *P. fergusi* and
37
38 497 the crocodylids when scaled (Table S4), but it is worth noting that this is not the case when a
39
40 498 curved beam is used instead.
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4445 499 Resistance to mediolateral bending (I_y) is very similar between *Proterosuchus fergusi*
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47 500 and the crocodylids when scaled (Fig. 9C) This is also demonstrated by the paired t-test
48
49 501 results, which show no significant differences in I_y values between *P. fergusi* and the
50
51 502 crocodylids at true size (Table S4). However, when taxa are size-corrected *P. fergusi* has
52
53 503 relatively poor resistance compared to the other taxa (Fig. 9D).
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5657 504 Resistance to torsion (J) is the sum of I_x and I_y and so follows the general trends of I_x
58
59 505 and I_y . *Proterosuchus fergusi* has J values very similar to *Crocodylus rhombifer* and
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3 506 *Crocodylus moreletii* at true size (Fig. 9E; Table S4). When taxa are scaled, all species have
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5 507 very similar torsion resistance, except *T. schlegelii* which is much lower (Fig. 9F).
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11 509 **DISCUSSION**
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17 511 *Life habits of Proterosuchus*
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19
20 512 *Hearing.* The lower portion of the inner ear contains the ECD, which comprises sound-
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22 513 detecting cells that are stimulated by sound waves transmitted from the middle ear (Sobral *et*
23
24 514 *al.* 2016b). The virtual reconstruction of the ECD of *Proterosuchus fergusi* has permitted an
25
26 515 estimation of the mean hearing frequency (MHF) and hearing range (HR). In order to
27
28 516 understand the effects that phylogeny has on these interpretations, a PGLS regression was
29
30 517 also used in addition to the linear regression proposed by Walsh *et al.* (2009). The regressions
31
32 518 gave different results, with sensitivity range estimations of ~0.1–0.9 kHz and ~0.2–2.4 kHz
33
34 519 for the linear and PGLS regressions, respectively. However, despite differences, results from
35
36 520 both iterations concur that *P. fergusi* was probably more specialised to lower frequencies,
37
38 521 much like modern crocodiles (Vergne *et al.* 2009; Walsh *et al.* 2009). Following the results of
39
40 522 the linear regression, acoustic estimations for *P. fergusi* are on the lower end of sensitivity
41
42 523 values known for modern crocodylians (Walsh *et al.* 2009), as well as the extinct
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44 524 crocodyliforms such as the thalattosuchian *Steneosaurus cf. gracilirostris* (Brusatte *et al.*
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46 525 2015) and the basal euschian *Lohuecosuchus megadontos* (Serrano-Martínez *et al.* 2019), for
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48 526 which auditory acuity has also been estimated.
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55 527 Estimations of auditory ability can also be informative in understanding the vocal
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57 528 complexity of a species. Considered to have resulted from coevolution of the vocal and
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59 529 auditory systems, individuals can typically only perceive frequencies within their own vocal
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3 530 range (Narins *et al.* 2004). Vocality of a species can be used to infer behavioural aspects such
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5 531 as social complexity and ecology (Walsh *et al.* 2009; Walsh *et al.* 2014).
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8 532 Sociality and vocality are integrally linked, with research suggesting social complexity
9
10 533 has driven both vocal complexity and cognition in vertebrates (Blumstein & Armitage 1997;
11
12 534 Freeberg *et al.* 2012; Freeberg & Krams 2015; Sewall 2015). Species that form larger and
13
14 535 more complex social systems require a more complex vocal system to work together
15
16 536 effectively. Therefore, vocal range, and thus auditory range, can be indicative of group size in
17
18 537 a species. This is supported by results from Walsh *et al.* (2009) which found significant
19
20 538 correlations between increased ECD length and larger social aggregations. Following this, the
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22 539 small ECD reconstructed in *P. fergusi* could suggest that it lived in small groups, or was even
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24 540 solitary.
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29 541 In closed environments where visual communication is less effective (such as dense
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31 542 jungles), vocal and auditory complexity is highly advantageous (Garrick & Lang 1977;
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33 543 Brown & Waser 1984). In this study we find the relatively low acoustic complexity
34
35 544 interpreted for *P. fergusi* to be consistent with the proposed absence of forests and vegetation
36
37 545 following the Permo-Triassic mass extinction in the Karoo (Smith 1995; Ward *et al.* 2000;
38
39 546 Smith *et al.* 2012). However, it is worth noting that while vocalisation and hearing ability in
40
41 547 mammals is thought to be suggestive of habitat type (Brown & Waser 1984; de la Torre &
42
43 548 Snowdon 2002), within living archosaur groups the relationship remains ambiguous (Garrick
44
45 549 & Lang 1977; Nicholls & Goldizen 2006; Mason & Burns 2015).
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54 551 *Olfaction.* The recognition of olfactory cues (sense of smell) is important for interspecific
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56 552 communication (e.g. kin recognition, finding a mate, territorial markers), navigation (e.g.
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58 553 foraging, predation, finding shelter and other resources), and avoiding dangers (e.g. predator
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3 554 identification, poisonous food) (Dial & Schwenk 1996; Hemila & Reuter 2008; Krause *et al.*
4
5 555 2012; Müller *et al.* 2018). The olfactory bulbs, located anterior to the forebrain, contain
6
7 556 olfactory receptor (OR) proteins that are responsible for detecting odorants (Freitag *et al.*
8
9 557 1998). The size of the olfactory bulbs generally corresponds to the quantity of OR proteins
10
11 558 (Steiger *et al.* 2009), meaning that the enlargement of the olfactory bulbs is often indicative of
12
13 559 increased olfactory abilities. Following this, the size of the olfactory bulbs has been widely
14
15 560 used as a proxy for olfactory acuity in fossil taxa (Benton 1983; Brochu 2000; Kundrát 2007;
16
17 561 Witmer *et al.* 2008; Witmer & Ridgely 2009; Zelenitsky *et al.* 2009; Bourke *et al.* 2014; Sales
18
19 562 & Schultz 2014). The presence of olfactory fossae along the ventral surface of the frontals has
20
21 563 allowed for the reconstruction of the olfactory bulbs in *Proterosuchus fergusi* (Fig. 2). The
22
23 564 olfactory bulbs are analogous in relative size and shape to many modern crocodylians (Pritz
24
25 565 1975; Witmer *et al.* 2008; Jirak & Janacek 2017). Many crocodylians are considered to have a
26
27 566 well-developed sense of smell, capable of quickly locating carrion both in the water and on
28
29 567 land (Weldon *et al.* 1990). This suggests that proterosuchids may have had a similar olfactory
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31 568 acuity to modern crocodiles.
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38 569 A latex endocast of *Tasmaniosaurus triassicus*, a sister-taxon to Archosauriformes
39
40 570 (Ezcurra 2016) (previously considered a proterosuchid (Camp & Banks 1978)), revealed
41
42 571 olfactory bulbs that were 1.4 times wider than the maximum width of the cerebrum (Ezcurra
43
44 572 2014). In *P. fergusi* the olfactory bulbs are only ~60% the maximum width of the cerebrum,
45
46 573 suggesting significant differences in the life habits of these two Early Triassic carnivores.
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48 574 Ezcurra (2014) suggested the large olfactory bulbs in *T. triassicus* were indicative of a
49
50 575 terrestrial ecology, following a study showing larger olfactory bulbs in fully terrestrial
51
52 576 carnivorans in comparison to those with an aquatic affinity (Gittleman 1991). While the
53
54 577 relationship between olfaction and habitat is more ambiguous in reptiles than mammals
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56 578 (Marek *et al.* 2015; Müller *et al.* 2018), the striking difference in relative olfactory width in
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3 579 these two phylogenetically similar taxa suggests that *P. fergusi* may have been more adapted
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5 580 to aquatic environments than *T. triassicus*.

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11 582 *Prey Choice*. Historically proterosuchids were considered aquatically-based predators,
12
13 583 feeding on fish (Tatarinov 1961) and tetrapods, such as the dicynodont therapsid *Lystrosaurus*
14
15 584 (Reig 1970). *Proterosuchus fergusi* occurs between 5–14 metres above the Permo-Triassic
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18 585 boundary in the lower levels of the *Lystrosaurus* AZ of the Karoo Basin (Smith & Botha-
19
20 586 Brink 2014). During the deposition of these rocks there was a high abundance of potential
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22
23 587 prey items for *P. fergusi*, including dicynodont (e.g. *Lystrosaurus* spp.) and therocephalian
24
25 588 (e.g. *Moschorhinus* and *Promoschorhynchus*) synapsids (Smith & Botha-Brink 2014).
26
27 589 However, as no direct evidence of proterosuchid diet has been found (i.e. preserved gut
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29 590 contents), only indirect evidence such as tooth morphology and general anatomy can be used
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31
32 591 to infer feeding habit (Ezcurra *et al.* 2013).

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35 592 Results of the rostral beam analysis show that the more oreinirostral morphology of *P.*
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37 593 *fergusi* was highly resistant to bending forces and torsion, comparable to the typically
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39 594 platyrostral morphology in *Crocodylus rhombifer* and *Crocodylus moreletii* (Fig. 9, Table
40
41 595 S4). These results are consistent with finite element analyses from Rayfield & Milner (2008)
42
43 596 comparing oreinirostral and platyrostral morphologies. The two *Crocodylus* species included
44
45 597 in this analysis are generalists, feeding on a variety of prey items, including fish, amphibians,
46
47 598 reptiles, birds and mammals (Platt *et al.* 2006; Milián-García *et al.* 2011). Juvenile and
48
49 599 subadult crocodiles have an even more varied diet, with large quantities of invertebrates,
50
51 600 gastropods and small vertebrates (Platt *et al.* 2006). Similarly, in *P. fergusi* the more gracile
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53 601 skull and relatively longer teeth of juvenile specimens has been thought to suggest that
54
55 602 different ontogenetic stages favoured different prey items, possibly to decrease intraspecific
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3 603 competition (Ezcurra & Butler 2015b). *Proterosuchus fergusi* may have had a generalist diet
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5 604 like modern crocodylians that change their diet through ontogeny from largely invertebrates
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7 605 to large terrestrial vertebrates. *Proterosuchus fergusi* and the platyrostral crocodylians
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9 606 (*Alligator mississippiensis*, *C. rhombifer*, *C. moreletii*) show significant differences in
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11 607 bending resistance to the longirostrine false gharial (*Tomistoma schlegelii*) in the analyses
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13 608 (Fig. 9, Table S4). This difference reflects the more specialist life habit of *T. schlegelii* with a
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15 609 predominately piscivorous diet (Pierce *et al.* 2008).
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23 611 *Ecological adaptations.* Despite the presence of eleven well preserved specimens of
24
25 612 *Proterosuchus fergusi*, there is currently no consensus on their ecology, or on the ecology of
26
27 613 proterosuchids as a whole (Ezcurra *et al.* 2013). Original interpretations of a semi-aquatic
28
29 614 habit for *P. fergusi* were in part due to the supposedly wet climate and palustrine
30
31 615 environments presumed of the *Lystrosaurus* AZ. However, recent studies have shown that the
32
33 616 Karoo Basin at this time was drastically different. Sedimentological analyses show the region
34
35 617 was predominately semi-arid with vast braided rivers resulting from lack of vegetation (Smith
36
37 618 1995; Ward *et al.* 2000; Smith *et al.* 2012) and the climate fluctuated between droughts,
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39 619 sporadic flood events and extreme cold snaps (Smith 1995; Smith & Ward 2001; Viglietti *et*
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41 620 *al.* 2013; Smith & Botha-Brink 2014).
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46 621 Endocranial reconstructions of the two *P. fergusi* specimens have not allowed us to
47
48 622 make sound interpretations of its ecology based on palaeoneurology. Statistical tests of the
49
50 623 brain cavity outline morphology (Table 1) show significant differences only between
51
52 624 terrestrial and semi-aquatic taxa. This difference however is likely explained by the
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54 625 evolutionary divergence between predominately terrestrial avemetatarsalians and
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56 626 predominately semi-aquatic pseudosuchians. Similar morphometric analyses on the
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3 627 endosseous labyrinth of saurians (Fig. 5) better separate groups based on ecology (Table 3),
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5 628 but also seem to be highly constrained by phylogeny (Table 4). The landmark-based analysis
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7 629 on individual semi-circular canals (Fig. 6) appears to be more appropriate for looking at
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9 630 ecology; statistical analyses show significant differences between the majority of ecological
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11 631 groupings (Table 5) and comparatively poor separation of phylogenetic groupings (Table S2).
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13 632 Across all three canals, *P. fergusi* resides consistently close to predominately semi-aquatic
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15 633 and aquatic taxa from various phylogenetically-distant groups (particularly sauropterygians,
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17 634 squamates and pseudosuchians). While the relationships between ecology, locomotion and
18
19 635 function of a taxon and the endocranial anatomy are currently uncertain, several studies have
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21 636 recently attempted to elucidate this by looking at the morphology of the endosseous labyrinths
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23 637 (Cuthbertson *et al.* 2015; Yi & Norell 2015; Benson *et al.* 2017; Neenan *et al.* 2017). Based
24
25 638 on our results, the endosseous labyrinths seem to be most appropriate for this type of analysis,
26
27 639 particularly when individual semi-circular canals are separately analysed. If morphological
28
29 640 differences are found to be indicative of ecological variation, with endosseous labyrinths
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31 641 comparable to various amphibious taxa, it is reasonable to postulate that *P. fergusi* may have
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33 642 had the sensory and balance organs required for semi-aquatic life.
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40 643 In life, the endosseous labyrinths (semi-circular canals) are part of the vestibular
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42 644 system, responsible for balance and co-ordination. Endolymph fluid moves back and forth
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44 645 through each of the canals as the skull rotates, triggering receptors at the end of the canals that
45
46 646 send signals to the brain (Clack 2016). Unlike the auditory organs, the function of these
47
48 647 vestibular organs has largely remained constant over vertebrate evolution (Carey & Amin
49
50 648 2006) allowing extant animals to be used as analogues to inform about the balance of early
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52 649 tetrapods. For nearly a century many tetrapods have been thought to habitually hold the lateral
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54 650 semi-circular canal (LSC, or sometimes ‘horizontal semi-circular canal’) horizontally
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56 651 (Lebedkin 1924; de Beer 1947; Duijm 1951). Following this, over the last two decades the
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3 652 LSC has been widely used to infer the head posture (typically ‘alert’ or ‘resting’) of various
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5 653 fossil groups (Rogers 1998; Witmer *et al.* 2003; Sampson & Witmer 2007; Sereno *et al.* 2007;
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7 654 Witmer *et al.* 2008; Witmer & Ridgely 2009; Neenan & Scheyer 2012; Benoit *et al.* 2017).
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10 655 The average orientation of the LSC in *P. fergusi* specimens SNSB-BSPG 1934 VIII 514 and
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12 656 RC 846 reconstructed in this study was $\sim 17^\circ$ (Fig. 7A), suggesting *P. fergusi* held its head
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14 657 tilted upwards (Fig. 7B). Among the array of taxa for which head posture has been inferred in
15
16 658 this way, up tilted expressions are generally uncommon, especially to the degree found in *P.*
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18 659 *fergusi* (Marugán-Lobón *et al.* 2013). While the use of the LSC in inferring head posture has
19
20 660 been met with scepticism, due to variation of inferred head postures found within
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22 661 phylogenetically similar groups (Hullar 2006; Taylor *et al.* 2009; Marugán-Lobón *et al.*
23
24 662 2013), several studies have suggested a possible relationship between the LSC and the
25
26 663 resulting head posture linked to ecological factors (e.g. diet, feeding behaviour) and supported
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28 664 by osteological correlates (i.e. occipital position) (Sereno *et al.* 2007; Neenan & Scheyer
29
30 665 2012; Schellhorn 2018). If LSC orientation does show a behavioural and/or ecological signal,
31
32 666 the range of LSC orientations recorded in the two specimens of *P. fergusi* indicate an
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34 667 upturned head posture and could be interpreted as an adaptation to semi-aquatic life. When
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36 668 upturned $\sim 17^\circ$ the bones of the cranial roof lie almost horizontally and the vertically
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38 669 positioned external nares are about level with orbit height. When in the water, *P. fergusi*
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40 670 would be able to keep its eyes and nostrils out of the water while keeping a low profile close
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42 671 to the waterline with the rest of its body remaining fully submerged, alike to modern day
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44 672 crocodiles. As previously discussed, dorsally-positioned external nares are an adaptation to
45
46 673 aquatic and semi-aquatic life that has convergently evolved in numerous groups over
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48 674 vertebrate evolution. Perhaps the upturned head posture of *P. fergusi* represents an alternative
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50 675 secondary adaptation for a semi-aquatic lifestyle among early diapsids. This would concur
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52 676 with the upwards tilting skull of *Lystrosaurus* (Benoit *et al.* 2017), which were
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3 677 contemporaneous to proterosuchids and have likewise also been interpreted as possibly semi-
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5 678 aquatic (Retallack *et al.* 2003; Ray *et al.* 2005).
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11 680 *Functional morphology of the premaxilla*
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14 681 The overhanging premaxilla that characterises proterosuchids is enigmatic in origin and
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16 682 function. While unique in its accentuation, the snout morphology is analogous with several
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18 683 extant and extinct vertebrate groups. Within the clade Archosauriformes, ornithosuchids such
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20 684 as *Riojasuchus* and *Ornithosuchus* exhibit a superficially similar snout morphology to
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22 685 *Proterosuchus*. However, the comparatively few specimens known for ornithosuchids mean
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24 686 that limited information can be obtained about the functional morphology and evolution of
25
26 687 this trait within archosauriforms. Modern analogues allow for the study of functional and
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28 688 sexual dimorphic aspects of this bizarre snout morphology. While phylogenetically distant,
29
30 689 actinopterygians including *Oncorhynchus kisutch* (coho salmon) (Ezcurra 2017) and (to a
31
32 690 lesser extent) many lepisosteids (gar) have a downturned premaxilla that does not fully
33
34 691 occlude with the lower jaw. In *O. kisutch*, this trait is sexually selected for and found only in
35
36 692 the hooknose adult male morphotype of the species (Fleming & Gross 1994). By contrast, in
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38 693 lepisosteids, this trait is thought to be functional in predation, for manipulating prey following
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40 694 capture (Lauder & Norton 1980).
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47 695 Convergent features in fossil and living taxa have long been interpreted as inferring shared
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49 696 function, even between highly phylogenetically-distant groups (e.g. Ji *et al.* 2006; Field *et al.*
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51 697 2011; O'Brien *et al.* 2016; Vullo *et al.* 2016; Arbour & Zanno 2019). Vullo *et al.* 2016
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53 698 commented on the morphological similarities between the jaws of spinosaurid dinosaurs and
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55 699 an extant group of anguilliform fish (pike conger eels) and suggested that this inferred shared
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57 700 feeding behaviours. The lepisosteid *Atractosteus spatula* (alligator gar) was among
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3 701 comparative taxa analysed for rostral resistance because of its overhanging premaxilla. Albeit
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5 702 far less accentuated compared to *Proterosuchus fergusi*, comparing results between the two
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7 703 overhanging premaxillae may have identified signals unique to this trait. The various
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9
10 704 statistical analyses tend to show significant differences between *P. fergusi* and *A. spatula*
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12 705 (Table S4). These results may suggest that *P. fergusi* did not share a similar method of
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14 706 predation to lepisosteids.

17 707 Growing and maintaining an oversized premaxilla with up to 18 ankylotheodont
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19 708 (deeply rooted and fused to the bone) teeth that would undergo periodic replacement, made
20
21 709 the snout a costly phenotype (Ezcurra 2017). As such, without a morphological function, this
22
23 710 trait would likely been negatively selected out of the population by natural selection
24
25 711 (Andersson 1994). To date no physiological function has been suggested and mechanical
26
27 712 functions such as digging have been ruled out after macroscopic analysis on the premaxillary
28
29 713 teeth (Ezcurra 2017). The ontogenetic changes in the snout morphology (Ezcurra & Butler
30
31 714 2015b) and phenotypic costliness led Ezcurra (2017) to propose social and sexual selection as
32
33 715 a possible explanation.

36 716 Snout beam analyses suggest that the premaxilla of *P. fergusi* is considerably resistant
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38 717 to dorsoventral bending and torsion compared to the crocodylians (Fig. 9; Table S4). The
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40 718 overhanging premaxilla does not seem to enhance or reduce resistance to mediolateral
41
42 719 bending. The high resistance of the premaxilla to dorsoventral bending may be an indication
43
44 720 of a specialist predation method. If this specialist function was beneficial enough it would
45
46 721 outweigh the costliness of the premaxilla. However advanced biomechanical analyses such as
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48 722 finite element analysis would be required to test this hypothesis. It is possible that this
49
50 723 phenotype may have been positively selected for under sexual selection, with a larger
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52 724 overhanging premaxilla being a signal for increased physical strength. While presumably
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54 725 sexual dimorphism was as widespread among fossil saurians as it is today, recognising sexual

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3 726 dimorphism and even distinguishing gender is extremely difficult and problematic in the
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5 727 saurian fossil record (Bonnar *et al.* 2008; Ezcurra 2017; Mallon 2017). One aspect of sexual
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7 728 selection that could possibly be ruled out is the ‘handicap principle’ proposed by Zahavi
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9 729 (1975). The handicap principle states that survival of an individual to sexual maturity with a
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11 730 ‘handicap’ (in this instance the costly overhanging premaxilla) is evidence of overall fitness
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13 731 and therefore a suitable mate (Zahavi 1975; 1977). Resistance tests show that the premaxilla
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15 732 is not detrimental to rostral strength, therefore it is likely not a ‘handicap’ and this might not
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17 733 be a viable explanation of the phenotype.
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25 735 *Implications for the evolution of Archosauriformes*

28 736 *Evolution of the archosauriform brain.* The endocasts of *Proterosuchus fergusi* reconstructed
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30 737 in this study are important contributions to understanding the evolution of basal
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32 738 archosauriform neuroanatomy (Fig. 10). While the palaeoneurology of archosaurian clades
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34 739 (particularly Dinosauria) have been extensively studied, stem archosaurs have been widely
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36 740 overlooked. Prior to this study, the endocranial anatomy of only three specimens of non-
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38 741 archosaurian archosauriform had been virtually reconstructed (Trotteyn & Paulina-Carabajal
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40 742 2016; Fabbri *et al.* 2017) to add to the small number of informative artificial endocasts
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42 743 (Benton 1983; Gower & Sennikov 1996; Wharton 2000). Trotteyn & Paulina-Carabajal
43
44 744 (2016) reconstructed the endocranium of the proterochampsian *Tropidosuchus romeri*, while
45
46 745 Fabbri *et al.* (2017) reconstructed the endocranium of *P. fergusi* (RC 846 – although Fabbri *et*
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48 746 *al.* incorrectly identified the specimen as ‘RC 96’ which actually corresponds to a dicynodont
49
50 747 (Ezcurra 2015)) and *E. capensis*. However, the endocasts from Fabbri *et al.* (2017) are
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52 748 incompletely figured and not described in the text and are therefore largely uninformative in
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54 749 regard to the morphology of the early archosauriform brain cavity.
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3 750 Results of the morphometric analysis show that convergence in brain morphology has
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5 751 occurred multiple times in the archosauriform lineage, particularly in groups with similar life
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7 752 habits. A clear example of this is the significant overlap between bird and pterosaur brain
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9 753 morphology, both of which share a volant mode of life. These analyses also highlight
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11 754 similarities between the brain cavities of pseudosuchians and basal archosauriforms,
12
13 755 especially when compared to avemetatarsalians. In some archosauriform groups (particularly
14
15 756 non-avian theropods and avialans), endocranial morphology has been tentatively linked to
16
17 757 orbit size, locomotion, ecology and organisation of cranial elements (Bhullar *et al.* 2012;
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19 758 Kawabe *et al.* 2013; Balanoff & Bever 2017; Fabbri *et al.* 2017). Perhaps with further
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21 759 research the correlations between form and function of the endocranial anatomy will become
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23 760 better understood and explain some of the similarities between relatively phylogenetically-
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25 761 distant taxa that our study displays.
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34 763 *Evolution of the archosauriform inner ear.* The functional morphology and early evolution of
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36 764 the endosseous labyrinth is important for understanding the rise of archosaurs in the
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38 765 Mesozoic. Prior to this study no non-archosaurian archosauriform inner ear had been
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40 766 completely reconstructed (Fig. 11), leaving a significant gap in our understanding of how the
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42 767 vestibular and auditory systems have evolved in stem archosaurs.
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46 768 In many eusuchians, the ASC is significantly longer compared to the PSC (Georgi &
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48 769 Sipla 2008; Witmer *et al.* 2008; Brusatte *et al.* 2016; Pierce *et al.* 2017). However, in more
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50 770 basal archosauriforms, the ASC and PSC are much more equal in size (Kley *et al.* 2010;
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52 771 Brusatte *et al.* 2016; Lautenschlager & Butler 2016; Sobral *et al.* 2016b; Pierce *et al.* 2017).
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54 772 Pierce *et al.* (2017) proposed that more equally proportioned anterior and posterior canals are
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56 773 plesiomorphic for Archosauriformes. The relatively equal size of the ASC and PSC in the
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3 774 endosseous labyrinths reconstructed for *P. fergusi* is consistent with these interpretations. The
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5 775 relatively equal size of the ASC and PSC in phytosaurs has been suggested to be indicative of
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7 776 limited neck movement and semi-aquatic habitat (Sobral & Müller 2017). However, this
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9
10 777 seems unlikely as extant crocodiles inhabit semi-aquatic ecosystems and have relatively
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12 778 limited neck movements. Further research is required to determine the functionality of this
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15 779 derived trait in eusuchians.

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18 780 The inner ears of archosaurs are significantly different to those of other diapsid
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20 781 groups. One of the most striking differences is the comparatively long ECD possessed by both
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22 782 crocodile-line and bird-line archosaurs. This trait seems to be confined to archosaurs, with
23
24 783 most diapsid groups outside of Archosauromorpha (e.g. squamates, rhychocephalians,
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26 784 testudines and sauropterygians) having relatively short cochlear ducts. Pierce *et al.* (2017)
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28 785 suggested that within Pseudosuchia the elongated cochlear duct evolved crownward to
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31 786 phytosaurs due to the small ECD present in the phytosaurs *Ebrachosuchus* and *Parasuchus*
32
33 787 reconstructed by Lautenschlager & Butler (2016). However, Lautenschlager & Butler (2016)
34
35 788 stated that the two phytosaur specimens were dorsoventrally compressed by ~40% and so the
36
37 789 raw endocasts are not representative of true scale. When retrodeformed (Fig. 11E-F) the
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40 790 cochlear ducts are superficially and morphometrically similar to other pseudosuchians (Fig.
41
42 791 5A). Following this, the presence of the derived cochlear trait in both crocodile-line and bird-
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44 792 line archosaurs suggests the evolution of the trait occurred before Avemetatarsalia and
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46 793 Pseudosuchia diverged. However, prior to this study, the lack of labyrinth reconstructions in
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49 794 stem archosaurs meant that the origin of this trait could not be further constrained.

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53 795 Reconstructions of the endosseous labyrinth reveal *P. fergusi* to have a relatively
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55 796 small ECD compared to crown archosaurs (Fig. 2; Fig. 11B). Morphometric analyses of the
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57 797 inner ear place *P. fergusi* close to non-archosaurian diapsids when the ECD is included,
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59 798 suggesting that the small ECD is a plesiomorphic trait (Fig. 5A). Following this it could be

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2
3 799 postulated that the elongated ECD first evolved in archosauromorphs crownward of *P.*
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5 800 *fergusi*. As *P. fergusi* is the most basal member of the clade Archosauriformes (Nesbitt 2011),
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7 801 the most parsimonious inference is that a small cochlear duct is a plesiomorphic trait found in
8
9 802 basal diapsids and retained in lepidosauromorphs and basal archosauromorphs. Subsequently,
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11 803 the derived trait of an elongate duct evolved somewhere in stem archosaurs and was retained
12
13 804 in both avemetatarsalians and pseudosuchians. As a ventrally-extended ECD has been linked
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15 805 to increased auditory acuity (as previously discussed), perhaps the evolution of this derived
16
17 806 trait marked a significant increase in auditory ability, vocality, sociality and cognition, all of
18
19 807 which are thought to be integrally linked (Blumstein & Armitage 1997; Freeberg *et al.* 2012;
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21 808 Freeberg & Krams 2015; Sewall 2015). These developments may potentially have contributed
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23 809 to the radiation of archosaurs during the Triassic and Early Jurassic.
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32 811 **CONCLUSIONS**

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35 812 This study presents the first in-depth assessment of the braincase and inner ear of the Early
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37 813 Triassic basal archosauriform *Proterosuchus fergusi*. Morphological comparisons of
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39 814 endocranial reconstructions, supplemented by analyses on the skull posture, rigidity and
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41 815 auditory abilities, have revealed much about the probable life habits of *P. fergusi* and the
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43 816 evolution of the brain and inner ear through early archosauriform evolution. Endocranial
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45 817 reconstructions reveal *P. fergusi* had: (1) an endocranial cavity with low-angle brain flexures;
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47 818 (2) medium-sized olfactory bulbs; (3) pyramidal and sub-equally-sized semi-circular canals;
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49 819 (4) a small cochlear duct.
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54 820 The brain morphology of *P. fergusi* is superficially and statistically similar to modern
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56 821 crocodylians and phytosaurs, and different to birds, pterosaurs and non-avian theropods.
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58 822 Assessment of the endocranial anatomy suggests that *P. fergusi* is estimated to be specialised
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3 823 to lower frequency sounds, with a proficient sense of smell, both comparable to modern
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5 824 crocodiles. *Proterosuchus fergusi* held its head upwards $\sim 17^\circ$, perhaps representing an
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7 825 alternative adaptation to semi-aquatic life than dorsally-positioned nares and orbits. The
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9 826 overhanging premaxilla remains largely enigmatic; with beam theory analyses indicating that
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11 827 generally this unusual morphology neither strengthens or weakens the snout and therefore
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13 828 purely non-functional selection pressures for this trait may be ruled out.
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17 829 The small ECD of *P. fergusi* suggests basal archosauromorphs retained an ancestrally
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19 830 small ECD which later evolved to become elongate in more derived archosauriforms, before
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21 831 the divergence of avemetarsalians and pseudosuchians. The increase in auditory complexity
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23 832 as a result of an elongated ECD may indicate the coevolution of increased cognitive abilities,
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25 833 vocality and sociality in stem archosaurs, which could in turn have contributed to the
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27 834 radiation of archosaurs in the Triassic, forging the way for dinosaurs to dominate global
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29 835 faunas during the Mesozoic.
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44
45 841 University of Birmingham.
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58 1275 FIG 1. Specimens of *Proterosuchus fergusi* analysed in this study. A–D, SNSB-BSPG 1934
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60 1276 VIII 514; E–H, RC 846 (proposed neotype). Skulls are shown as photographs in dorsal (A, E),

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3 1277 right lateral (B, F), and left lateral (C, G) views, with outline drawings (D, H) of the most
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5 1278 complete lateral view. Outline drawing of RC 846 (H) is taken from Ezcurra & Butler
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8 1279 (2015a). All scale bars equal 50 mm.
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14 1281 FIG 2. Brain endocast reconstructions of *Proterosuchus fergusi*. (A–B) SNSB-BSPG 1934

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16 1282 VIII 514; (C–D) RC 846; (E–F) RC 846 μ CT. Reconstructions show dorsal views of

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18 1283 endocasts in situ (A, C, E), dorsal and left lateral views of isolated endocasts (B, D, F).
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21 1284 Abbreviations: ca, carotid artery canal; cb, cerebellum; ch, cerebral hemispheres; el,

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23 1285 endosseous labyrinth; mcv, middle cerebral vein; mo, medulla oblongata; ob, olfactory bulb;

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25 1286 ot, olfactory tract; iv, trochlear nerve canal; v, trigeminal nerve canal; vi, abducens nerve; vii,

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27 1287 facial nerve canal. Colour key: cranial endocast (blue); endosseous labyrinth (pink); cranial

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29 1288 nerves (yellow); arterial canals (red); smaller venous canals (dark blue). Scale bars: 50 mm

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32 1289 (A, C); 25 mm (E); 10 mm (B, D, F).
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38 1291 FIG 3. Endosseous labyrinths of *Proterosuchus fergusi* from highest to lowest CT resolution.

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40 1292 A, B, left (A) and right (B) labyrinths of RC 846 (μ CT). C, D, left (C) and right (D) labyrinths

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42 1293 of SNSB-BSPG 1934 VIII 514. E, F, left (E) and right (F) labyrinths of RC 846. Labyrinths

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44 1294 are shown in (from left to right) lateral, dorsal, medial, anterior and posterior views.
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47 1295 Abbreviations: asc, anterior semi-circular canal; cc, crus communis; ecd, endosseous cochlear

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49 1296 duct; fv, fenestra vestibuli; lsc, lateral semi-circular canal; psc, posterior semi-circular canal.
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52 1297 All scale bars equal 5 mm.
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58 1299 FIG 4. Morphospace plots of archosauriform brain outlines based on elliptic Fourier analysis

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60 1300 and after principal component analysis. Abbreviations: NAAR, non-archosaurian

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3 1301 archosauriform; NPPS, non-phytosaurian pseudosuchians; PHYT, Phytosauria; PTER,
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5 1302 Pterosauria; SAUR, Sauropodomorpha; ORTH, Ornithischia; NATH, non-avian theropods;
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7 1303 AVES, Aves. † denotes extinct taxa. For taxonomic information see Brown *et al.* (2019, Fig.
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9 1304 S9). Silhouettes used include work by S. Hartman, M. Witton, N. Tamura and T. M. Keesey
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11 1305 (see <http://phylopic.org> for full licensing information).

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18 1307 FIG 5. Morphospace plots of avian and reptile inner ear outlines based on elliptic Fourier
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20 1308 analysis and after principal component analysis, both with and without the endosseous
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22 1309 cochlear duct (ECD). (A) with ECD; (B) without ECD. Colour key: Aquatic (dark blue);
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24 1310 Semi-aquatic (light blue); Terrestrial (yellow). Abbreviations: SQUA, Squamata; SAUO,
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26 1311 Sauropterygia; TEST, Testudines; NAAR, non-archosaurian archosauriform; PHYT,
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28 1312 Phytosauria; NPPS, non-phytosaurian pseudosuchians; ORNI, Ornithischia; SAUR,
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30 1313 Sauropodomorpha; NATH, non-avian theropods; AVES, Aves. For taxonomic information
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32 1314 see Brown *et al.* (2019, Fig. S6).

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40 1316 FIG 6. Morphospace plots of avian and reptile semi-circular canal landmarks after principal
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42 1317 component analysis. PC1 versus PC2. (A) ASC; (B) LSC; (C) PSC. For taxonomic
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44 1318 information see Brown *et al.* (2019, fig. S9).

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51 1320 FIG 7. Discerning the ‘alert’ head posture in life from the orientation of the lateral semi-
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53 1321 circular canal (LSC). (A) Orientation of the LSC when head posture of *Proterosuchus fergusi*
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55 1322 is horizontal in both specimens studied. (B) Head posture of *P. fergusi* in life following mean
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57 1323 average orientation of all LSC planes studied. (C) Comparative head posture of *Crocodylus*
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59 1324 *johnstoni* following the orientation of the LSC plane (Witmer *et al.* 2008). *Proterosuchus*

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3 1325 *fergusi* and *C. johnstoni* skull figures modified from Ezcurra & Butler (2015a) and Witmer *et*
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5 1326 *al.* (2008) respectively.

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11 1328 FIG 8. Correlation plot between scaled/log transformed ECD length and variables of hearing
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13 1329 sensitivity for extant reptiles and Aves. Both a linear regression (solid line) and PGLS
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15 1330 regression (dashed line) were used to estimate the auditory capabilities of *Proterosuchus*
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17 1331 *fergusi* (indicated by the hollow circle). Sensitivity variables: (A) mean hearing frequency;
18
19 1332 (B) hearing range. Abbreviations: SQUA, Squamata; RHYN, Rhynchocephalia; TEST,
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21 1333 Testudines; CROC, Crocodylomorpha; AVES, Aves. For taxonomic information see Brown
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23 1334 *et al.* (2019, fig. S10).

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31 1336 FIG 9. Change in the resistance to dorsoventral bending (lx), mediolateral bending (ly), and
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33 1337 torsion (J) along the rostra in *Proterosuchus fergusi* and comparative extant taxa. (A) log lx
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35 1338 true size; (B) log lx scaled size; (C) log ly true size; (D) log ly scaled size; (E) log J true size;
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37 1339 (F) log J scaled size. In *Proterosuchus fergusi*, both a straight beam (solid line) and a curved
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39 1340 beam (dashed line) were used when analysing the anterior 20% of the rostrum.

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46 1342 FIG 10. Evolution of the brain cavity in Archosauriformes. Left lateral view of the brain
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48 1343 cavity of saurians redrawn from the literature. (A) snake *Dispholidus typus* (Allemand *et al.*
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50 1344 2017); (B) *Proterosuchus fergusi* (this study); (C) proterochampsian *Tropidosuchus romeri*
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52 1345 (Trotteyn & Paulina-Carabajal 2016); (D) *Parasuchus angustifrons* (retrodeformed,
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54 1346 Lautenschlager & Butler 2016); (E) *Ebrachosuchus neukami* (retrodeformed, Lautenschlager
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56 1347 & Butler 2016); (F) *Riojasuchus tenuisiceps* (Baczko & Desojo 2016); (G) *Pelagosaurus typus*
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58 1348 (Pierce *et al.* 2017); (H) *Caiman crocodilus* (Jirak & Janacek 2017); (I) *Crocodylus seamensis*

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3 1349 (Kawabe *et al.* 2009); (J) *Anhanguera santanae* (Witmer *et al.* 2003); (K) *Tyrannosaurus rex*

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5 1350 (Witmer *et al.* 2008); (L) *Bubo virginianus* (Witmer *et al.* 2008). † denotes extinct taxa.

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7 1351 Phylogeny following Nesbitt (2011) and Ezcurra (2016). Brain cavities not to scale.

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13 1353 FIG 11. Evolution of the endosseous labyrinth in Archosauriformes. Lateral view of the left

14 1354 labyrinth of archosauriforms redrawn from the literature. (A) snake *Ptyas mucosa* (reversed,

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16 1355 Yi & Norell 2015); (B) *Proterosuchus fergusi* (this study); (C) proterochampsian

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18 1356 *Tropidosuchus romeri* (reversed, Trotteyn & Paulina-Carabajal 2016); (D) *Euparkeria*

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20 1357 *capensis* (reversed, Sobral *et al.* 2016); (E) *Parasuchus angustifrons* (retrodeformed,

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22 1358 Lautenschlager & Butler 2016); (F) *Ebrachosuchus neukami* (retrodeformed, Lautenschlager

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24 1359 & Butler 2016); (G) *Simosuchus clarki* (reversed, Kley 2010); (H) *Pelagosaurus typus* (Pierce

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26 1360 *et al.* 2017); (I) *Steneosaurus cf. gracilirostris* (Brusatte *et al.* 2016); (J) *Crocodylus johnstoni*

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28 1361 (Brusatte *et al.* 2016); (K) *Alligator mississippiensis* (Brusatte *et al.* 2016); (L) *Stegosaurus*

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30 1362 *stenops* (Leahey *et al.* 2015); (M) *Camarasaurus lentus* (Witmer *et al.* 2008); (N)

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32 1363 *Tyrannosaurus rex* (Witmer *et al.* 2008); (O) *Bubo virginianus* (Witmer *et al.* 2008). †

33
34 1364 denotes extinct taxa. Abbreviations: ASC, anterior semi-circular canal; ECD, endosseous

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36 1365 cochlea duct; LSC, lateral semi-circular canal; PSC, posterior semi-circular canal. Phylogeny

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38 1366 following Nesbitt (2011), Ezcurra (2016) and Leardi *et al.* (2017). Labyrinths not to scale.

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49 1368 Table 1. Results of one-way PERMANOVA test on endocranial outlines between ecological
50 1369 groupings using all axes. *Proterosuchus* is excluded. Results in bold show statistical
51 1370 significance at $p = 0.05$.

52
53 1371 Permutation N 10000

54 1372 Total sum of squares 6.193

55 1373 Within-group sum of squares 5.098

56 1374 F 7.082

57 1375 p 0.0002

Ecological groupings	p-value (w/ ECD)
Aquatic/Semi-aquatic	0.0998
Aquatic/Terrestrial	0.4222
Semi-aquatic/Terrestrial	<0.0001

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1377 Table 2. Results of one-way PERMANOVA test on endocranial outlines between
 1378 phylogenetic groupings and using all axes. Abbreviations: NAAR, non-archosaurian
 1379 archosauriform; PHYT, Phytosauria; NPPS, non-phytosaurian pseudosuchians; PTER,
 1380 Pterosauria; ORTH, Ornithischia; SAUR, Sauropodomorpha; NATH, non-avian theropods;
 1381 AVES, Aves. Results in bold show statistical significance at $p = 0.05$.

1382 Permutation N 10000

1383 Total sum of squares 6.255

1384 Within-group sum of squares 2.825

1385 F 10.75

1386 p 1.00E-04

p-values	PHYT	NPPS	PTER	ORNI	SAUR	NATH	AVES
NAAR	0.4312	0.7721	0.104	0.0215	0.0190	0.0612	0.0064
PHYT		0.1084	0.0183	0.0043	0.0033	0.0175	0.0002
NPPS			0.0015	0.0001	0.0001	0.0002	0.0001
PTER				0.0126	0.0310	0.0136	0.0406
ORNI					0.0086	0.0263	0.0001
SAUR						0.0014	0.0001
NATH							0.0001

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1390 Table 3. Results of one-way PERMANOVA test on inner ear outlines between ecological
 1391 groupings with (left) and without (right) the endosseous cochlea duct (ECD) and using all
 1392 axes. *Proterosuchus* is excluded. Results in bold show statistical significance at $p = 0.05$.

1393 Permutation N 10000/10000

1394 Total sum of squares 7.52/ 5.685

1395 Within-group sum of squares 6.546/ 5.315

1396 F 6.772/ 3.345

1397 p 0.0001/ 0.0044

Ecological groupings	p-value (w/ ECD)	p-value (w/o ECD)
Aquatic/Semi-aquatic	0.0099	0.0282
Aquatic/Terrestrial	0.0003	0.0013
Semi-aquatic/Terrestrial	0.0099	0.2968

1398

1399 Table 4. Results of one-way PERMANOVA test on inner ear outlines between phylogenetic
 1400 groupings with (left) and without (right) the endosseous cochlea duct (ECD) and using all

1401 axes. Abbreviations: SQUA, Squamata; SAUO, Sauropterygia; TEST, Testudines; NAAR,
 1402 non-archosaurian archosauriform; PHYT, Phytosauria; NPPS, non-phytosaurian
 1403 pseudosuchians; ORNI, Ornithischia; SAUR, Sauropodomorpha; NATH, non-avian
 1404 theropods; AVES, Aves. Results in bold show statistical significance at $p = 0.05$.

1405 Permutation N 10000

1406 Total sum of squares 1.658

1407 Within-group sum of squares 0.522

1408 F 10.26

1409 p 9.999E-05

p-values (w/o ECD \ w/ ECD)	SQUA	SAUO	TEST	NAAR	PHYT	NPPS	ORNI	SAUR	NATH	AVES
SQUA		0.0002	0.0005	0.0010	0.0065	0.0001	0.0005	0.0001	0.0001	0.0001
SAUO	0.0101		0.4432	0.2562	0.1680	0.0002	0.0015	0.0001	0.0001	0.0001
TEST	0.0032	0.0307		0.1189	0.1078	0.0001	0.0014	0.0001	0.0001	0.0001
NAAR	0.0303	0.3823	0.0007		0.0995	0.0027	0.0290	0.0034	0.0014	0.0007
PHYT	0.2860	0.6812	0.0320	0.7363		0.0671	0.1380	0.0150	0.0119	0.0040
NPPS	0.0392	0.3248	0.0185	0.1340	0.6042		0.0023	0.0001	0.0001	0.0001
ORNI	0.0028	0.3494	0.0002	0.0063	0.0392	0.0072		0.2123	0.0080	0.0013
SAUR	0.0003	0.0200	0.0001	0.0618	0.1232	0.0001	0.0005		0.1188	0.0015
NATH	0.0001	0.0003	0.0001	0.0047	0.0162	0.0001	0.0001	0.0319		0.0011
AVES	0.0001	0.0001	0.0001	0.0004	0.0053	0.0001	0.0001	0.0001	0.0001	

1410

1411 Table 5. Results of canonical variate analysis on semi-circular canal landmarks using all axes.
 1412 *Proterosuchus* was excluded. Results in bold show statistical significance at $p = 0.05$.

1413 Permutation N 10000

Ecological groupings	ASC		LSC		PSC	
	Procrustes	p-value	Procrustes	p-value	Procrustes	p-value
Terrestrial/Semi-Aquatic	0.1937	0.0146	0.1093	0.2442	0.1921	0.0178
Terrestrial/Aquatic	0.2970	0.0002	0.2646	0.0050	0.2168	0.0051
Semi-Aquatic/Aquatic	0.1655	0.0075	0.1778	0.0314	0.1099	0.3103

1414

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1416

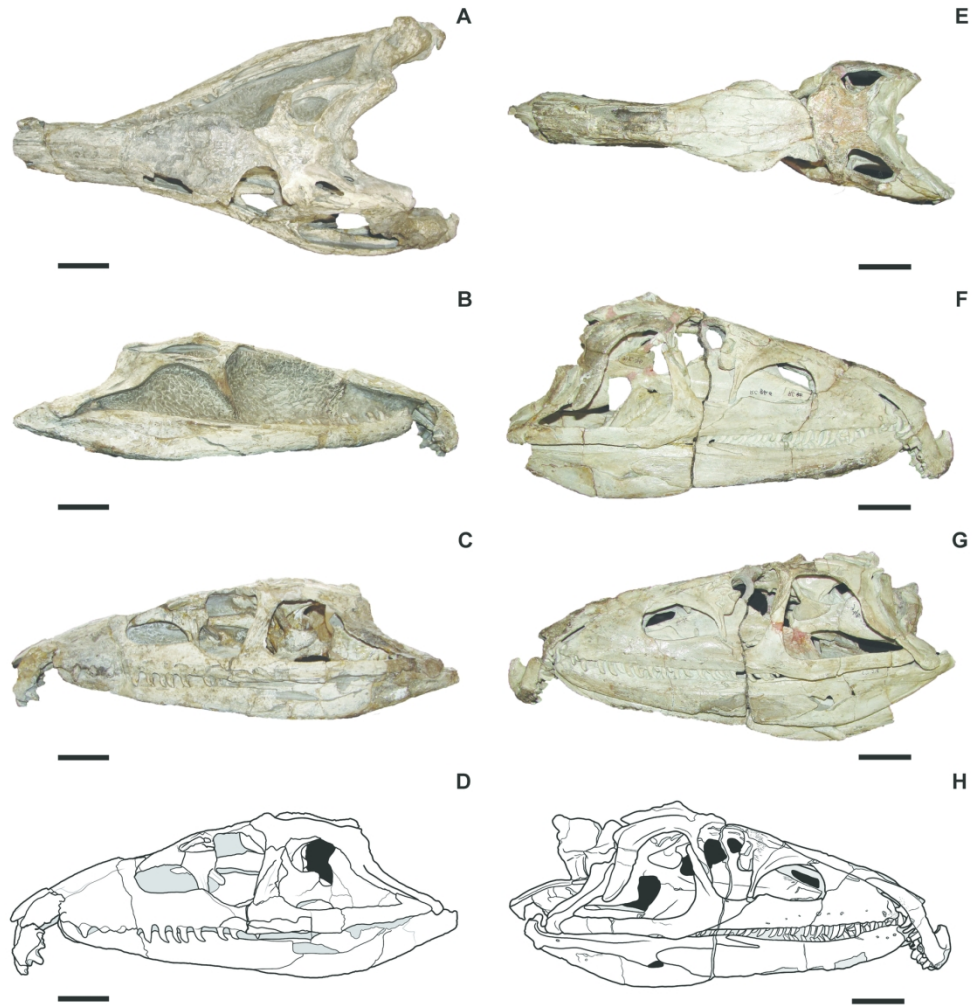


FIG 1. Specimens of *Proterosuchus fergusi* analysed in this study. A–D, SNSB-BSPG 1934 VIII 514; E–H, RC 846 (proposed neotype). Skulls are shown as photographs in dorsal (A, E), right lateral (B, F), and left lateral (C, G) views, with outline drawings (D, H) of the most complete lateral view. Outline drawing of RC 846 (H) is taken from Ezcurra & Butler (2015a). All scale bars equal 50 mm.

165x170mm (300 x 300 DPI)

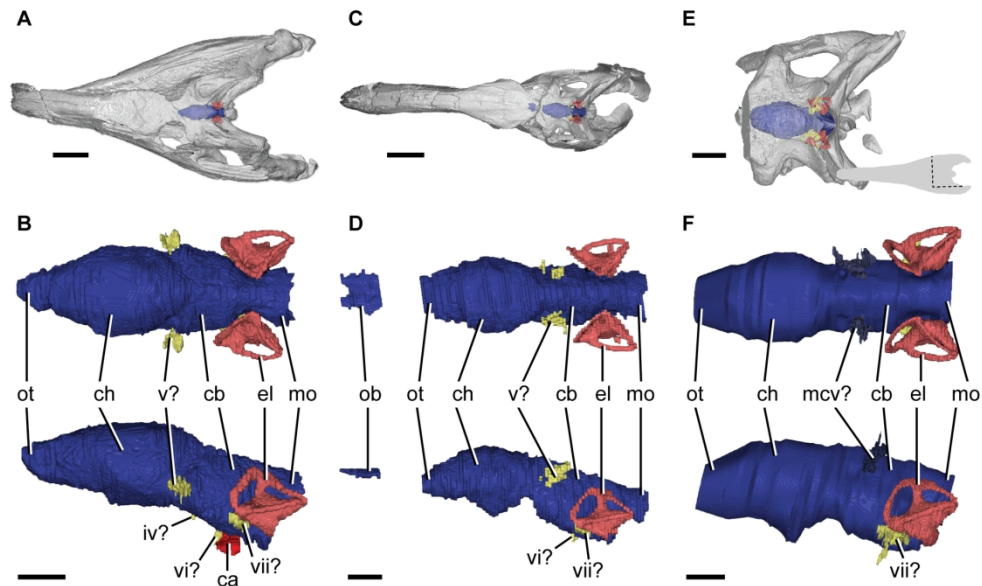


FIG 2. Brain endocast reconstructions of *Proterosuchus fergusi*. (A–B) SNSB-BSPG 1934 VIII 514; (C–D) RC 846; (E–F) RC 846 μ CT. Reconstructions show dorsal views of endocasts in situ (A, C, E), dorsal and left lateral views of isolated endocasts (B, D, F). Abbreviations: ca, carotid artery canal; cb, cerebellum; ch, cerebral hemispheres; el, endosseous labyrinth; mcv, middle cerebral vein; mo, medulla oblongata; ob, olfactory bulb; ot, olfactory tract; iv, trochlear nerve canal; v, trigeminal nerve canal; vi, abducens nerve; vii, facial nerve canal. Colour key: cranial endocast (blue); endosseous labyrinth (pink); cranial nerves (yellow); arterial canals (red); smaller venous canals (dark blue). Scale bars: 50 mm (A, C); 25 mm (E); 10 mm (B, D, F).

170x100mm (300 x 300 DPI)

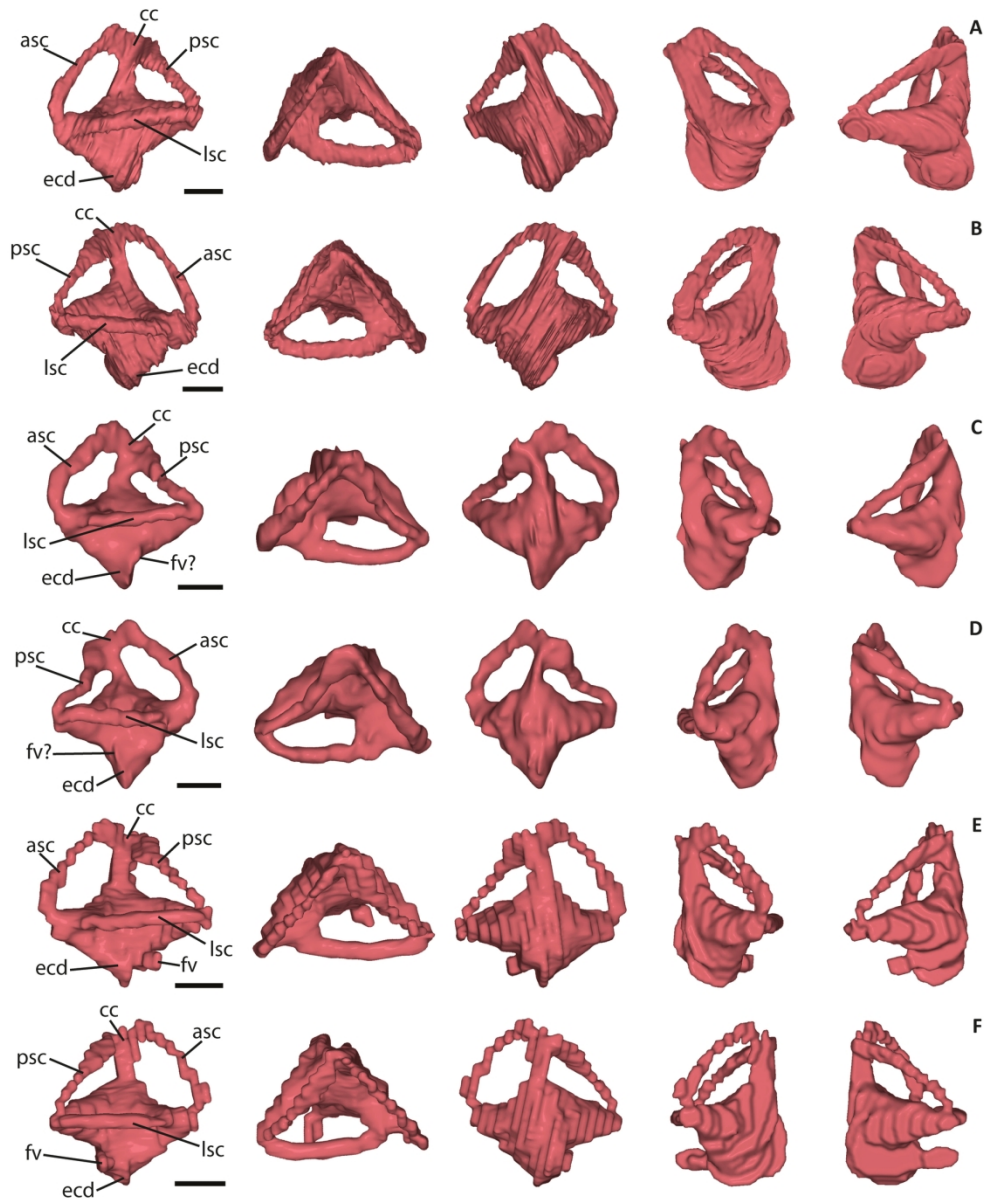


FIG 3. Endosseous labyrinths of *Proterosuchus fergusi* from highest to lowest CT resolution. A, B, left (A) and right (B) labyrinths of RC 846 (μ CT). C, D, left (C) and right (D) labyrinths of SNSB-BSPG 1934 VIII 514. E, F, left (E) and right (F) labyrinths of RC 846. Labyrinths are shown in (from left to right) lateral, dorsal, medial, anterior and posterior views. Abbreviations: asc, anterior semi-circular canal; cc, crus communis; ecd, endosseous cochlear duct; fv, fenestra vestibuli; lsc, lateral semi-circular canal; psc, posterior semi-circular canal. All scale bars equal 5 mm.

161x194mm (300 x 300 DPI)

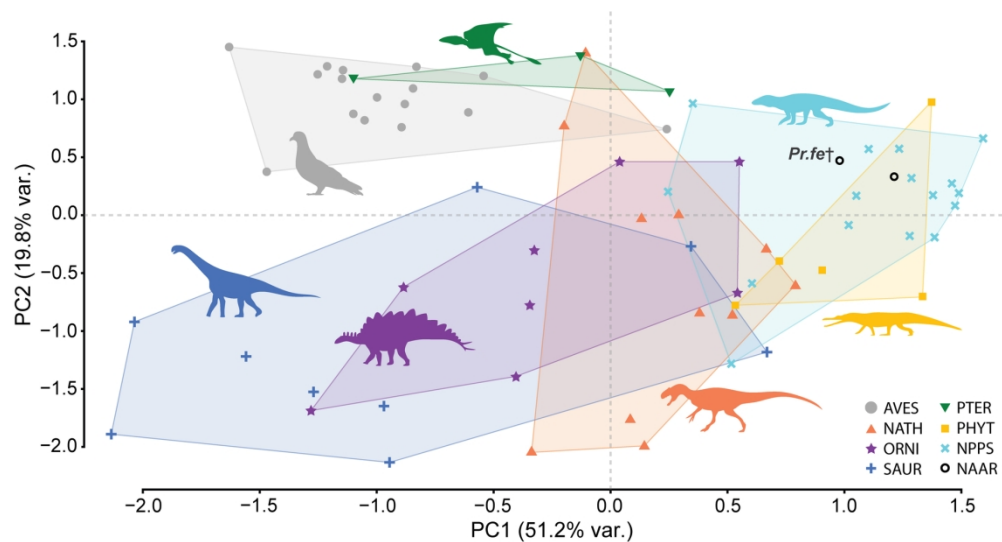


FIG 4. Morphospace plots of archosauriform brain outlines based on elliptic Fourier analysis and after principal component analysis. Abbreviations: NAAR, non-archosaurian archosauriform; NPPS, non-phytosaurian pseudosuchians; PHYT, Phytosauria; PTER, Pterosauria; SAUR, Sauropodomorpha; ORTH, Ornithischia; NATH, non-avian theropods; AVES, Aves. † denotes extinct taxa. For taxonomic information see Brown et al. (2019, Fig. S9). Silhouettes used include work by S. Hartman, M. Witton, N. Tamura and T. M. Keesey (see <http://phylopic.org> for full licensing information).

166x88mm (300 x 300 DPI)

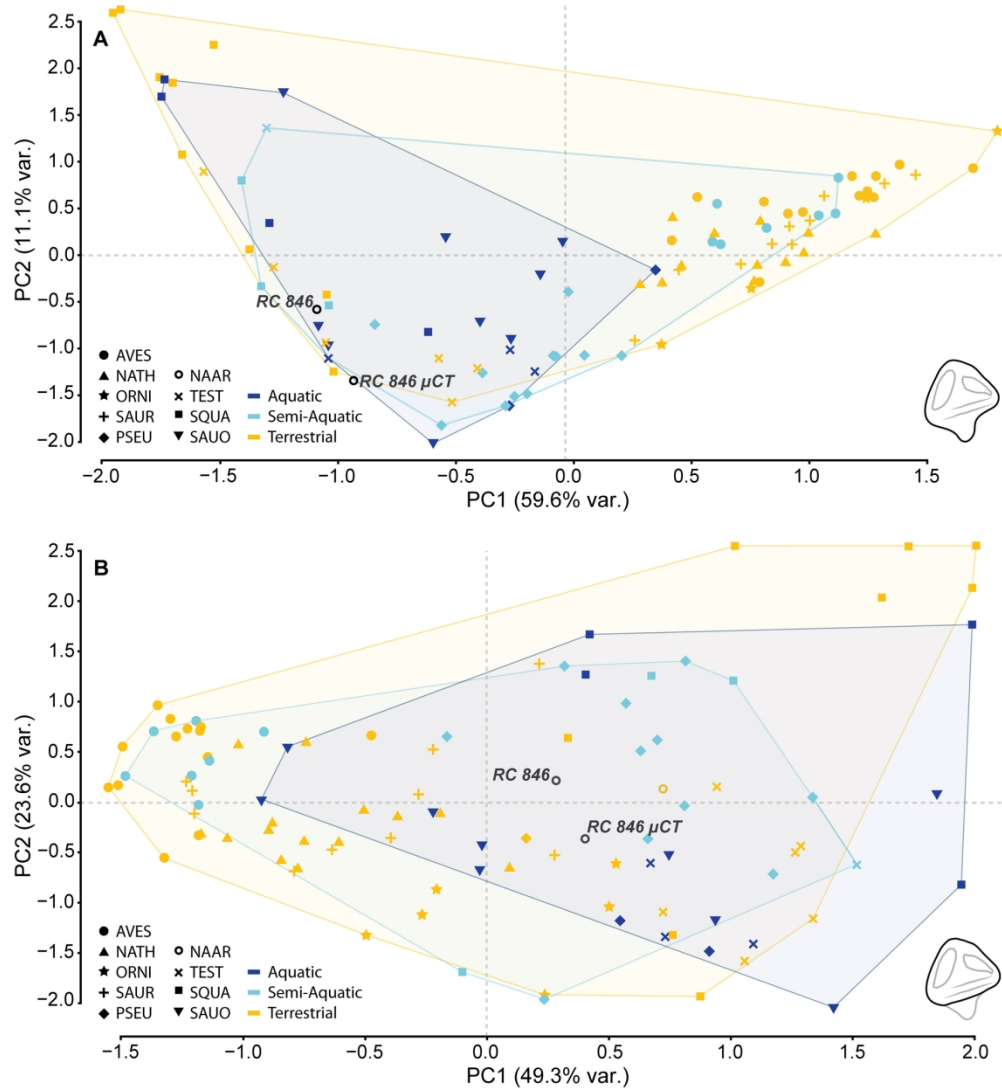


FIG 5. Morphospace plots of avian and reptile inner ear outlines based on elliptic Fourier analysis and after principal component analysis, both with and without the endosseous cochlear duct (ECD). (A) with ECD; (B) without ECD. Colour key: Aquatic (dark blue); Semi-aquatic (light blue); Terrestrial (yellow). Abbreviations: SQUA, Squamata; SAUO, Sauropterygia; TEST, Testudines; NAAR, non-archosaurian archosauriform; PHYT, Phytosauria; NPPS, non-phytosaurian pseudosuchians; ORNI, Ornithischia; SAUR, Sauropodomorpha; NATH, non-avian theropods; AVES, Aves. For taxonomic information see Brown et al. (2019, Fig. S6).

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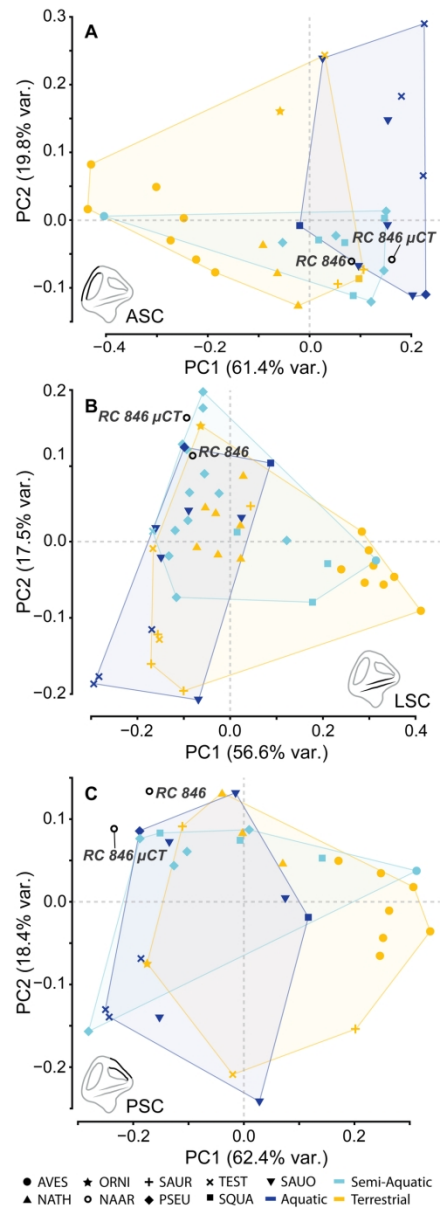


FIG 6. Morphospace plots of avian and reptile semi-circular canal landmarks after principal component analysis. PC1 versus PC2. (A) ASC; (B) LSC; (C) PSC. For taxonomic information see Brown et al. (2019, fig. S9).

80x226mm (300 x 300 DPI)

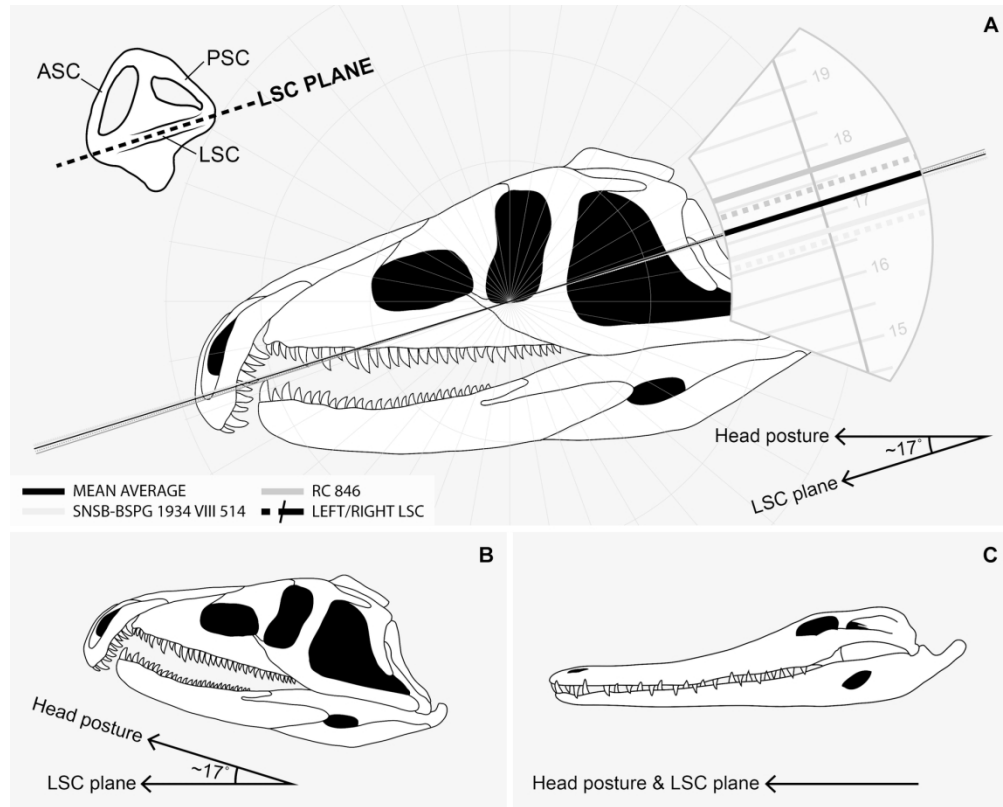


FIG 7. Discerning the 'alert' head posture in life from the orientation of the lateral semi-circular canal (LSC). (A) Orientation of the LSC when head posture of *Proterosuchus fergusi* is horizontal in both specimens studied. (B) Head posture of *P. fergusi* in life following mean average orientation of all LSC planes studied. (C) Comparative head posture of *Crocodylus johnstoni* following the orientation of the LSC plane (Witmer et al. 2008). *Proterosuchus fergusi* and *C. johnstoni* skull figures modified from Ezcurra & Butler (2015a) and Witmer et al. (2008) respectively.

165x133mm (300 x 300 DPI)

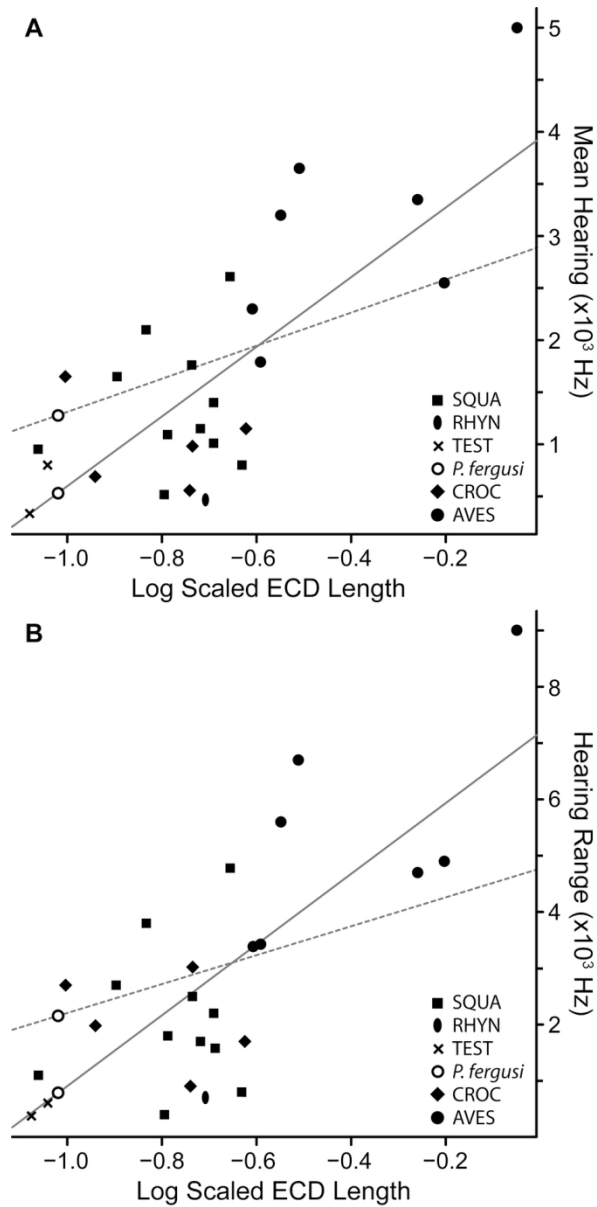


FIG 8. Correlation plot between scaled/log transformed ECD length and variables of hearing sensitivity for extant reptiles and Aves. Both a linear regression (solid line) and PGLS regression (dashed line) were used to estimate the auditory capabilities of *Proterosuchus fergusi* (indicated by the hollow circle). Sensitivity variables: (A) mean hearing frequency; (B) hearing range. Abbreviations: SQUA, Squamata; RHYN, Rhynchocephalia; TEST, Testudines; CROC, Crocodylomorpha; AVES, Aves. For taxonomic information see Brown et al. (2019, fig. S10).

81x165mm (300 x 300 DPI)

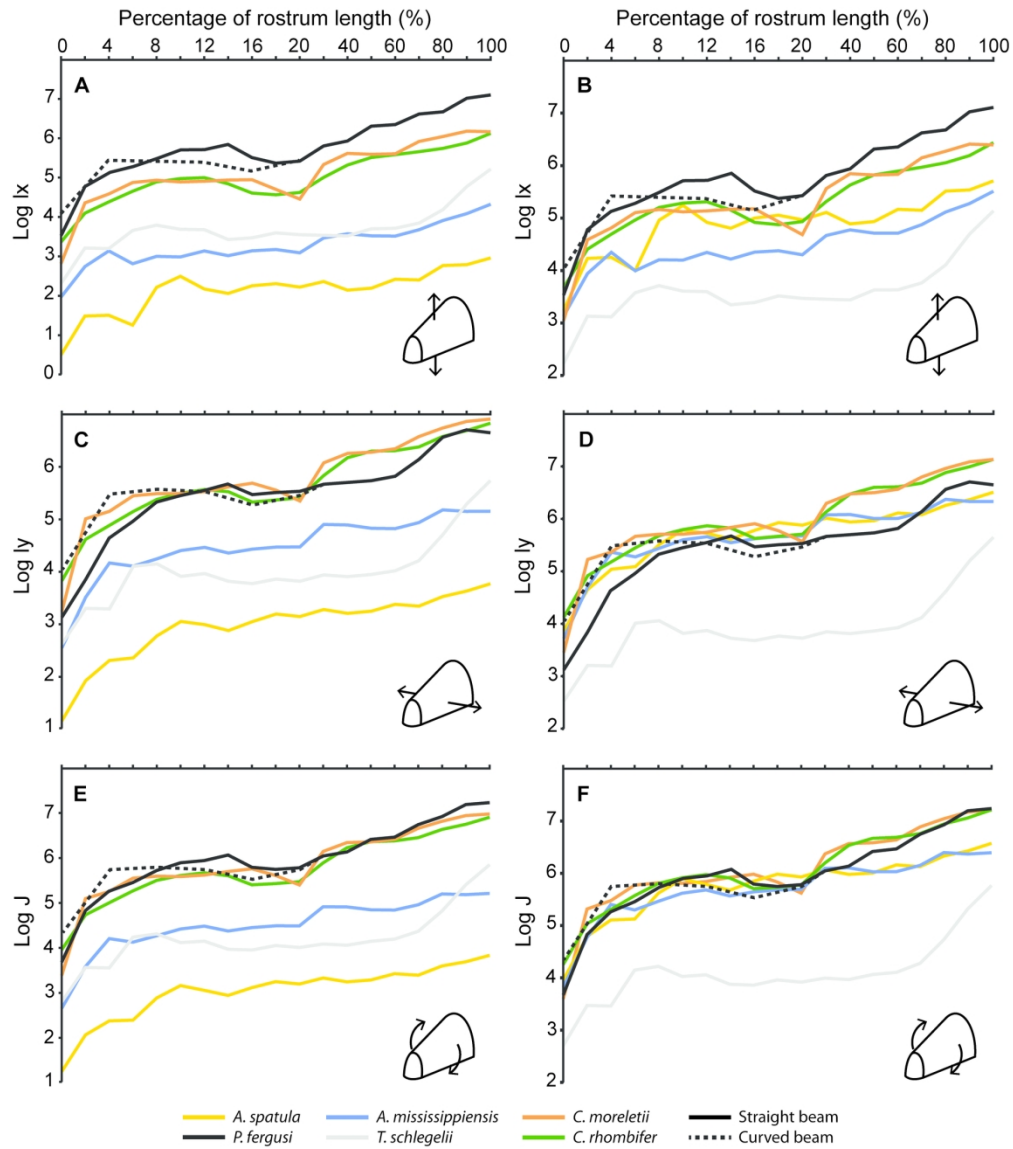


FIG 9. Change in the resistance to dorsoventral bending (I_x), mediolateral bending (I_y), and torsion (J) along the rostra in *Proterosuchus fergusi* and comparative extant taxa. (A) log I_x true size; (B) log I_x scaled size; (C) log I_y true size; (D) log I_y scaled size; (E) log J true size; (F) log J scaled size. In *Proterosuchus fergusi*, both a straight beam (solid line) and a curved beam (dashed line) were used when analysing the anterior 20% of the rostrum.

166x189mm (300 x 300 DPI)

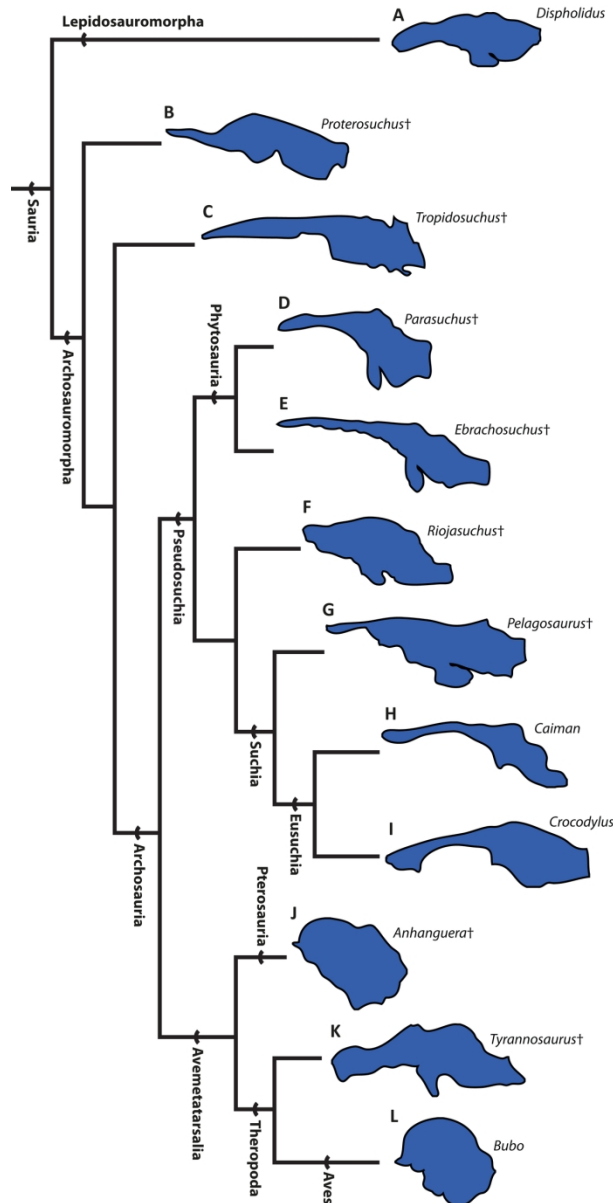


FIG 10. Evolution of the brain cavity in Archosauriformes. Left lateral view of the brain cavity of saurians redrawn from the literature. (A) snake *Dispholidus typus* (Allemand et al. 2017); (B) *Proterosuchus fergusi* (this study); (C) proterochampsian *Tropidosuchus romeri* (Trotteyn & Paulina-Carabajal 2016); (D) *Parasuchus angustifrons* (retrodeformed, Lautenschlager & Butler 2016); (E) *Ebrachosuchus neukami* (retrodeformed, Lautenschlager & Butler 2016); (F) *Riojasuchus tenuisiceps* (Baczko & Desojo 2016); (G) *Pelagosaurus typus* (Pierce et al. 2017); (H) *Caiman crocodilus* (Jirak & Janacek 2017); (I) *Crocodylus seamensis* (Kawabe et al. 2009); (J) *Anhanguera santanae* (Witmer et al. 2003); (K) *Tyrannosaurus rex* (Witmer et al. 2008); (L) *Bubo virginianus* (Witmer et al. 2008). † denotes extinct taxa. Phylogeny following Nesbitt (2011) and Ezcurra (2016). Brain cavities not to scale.

109x217mm (300 x 300 DPI)

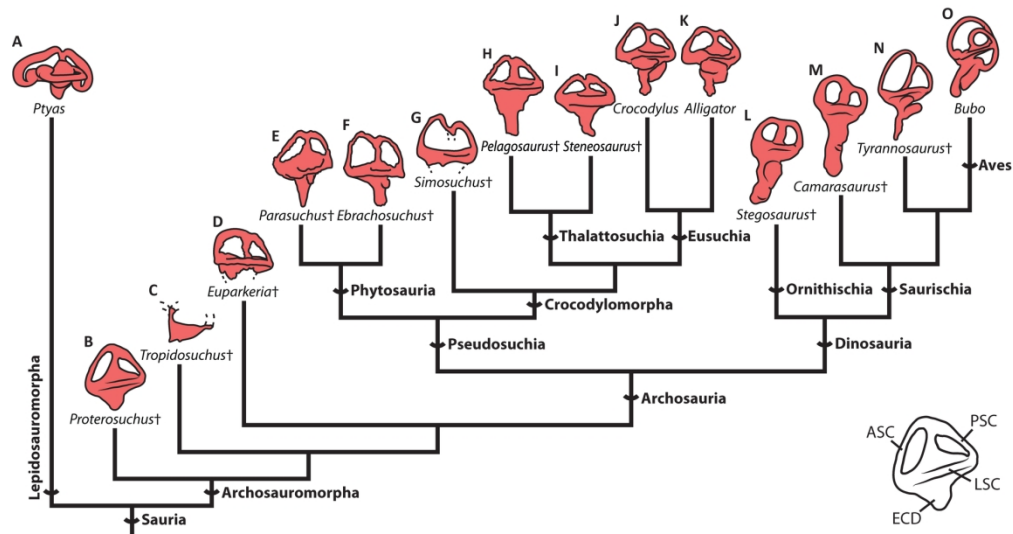


FIG 11. Evolution of the endosseous labyrinth in Archosauriformes. Lateral view of the left labyrinth of archosauriforms redrawn from the literature. (A) snake *Ptyas mucosa* (reversed, Yi & Norell 2015); (B) *Proterosuchus fergusi* (this study); (C) proterochampsian *Tropidosuchus romeri* (reversed, Trotteyn & Paulina-Carabajal 2016); (D) *Euparkeria capensis* (reversed, Sobral et al. 2016); (E) *Parasuchus angustifrons* (retrodeformed, Lautenschlager & Butler 2016); (F) *Ebrachosuchus neukami* (retrodeformed, Lautenschlager & Butler 2016); (G) *Simosuchus clarki* (reversed, Kley 2010); (H) *Pelagosaurus typus* (Pierce et al. 2017); (I) *Steneosaurus cf. gracilirostris* (Brusatte et al. 2016); (J) *Crocodylus johnstoni* (Brusatte et al. 2016); (K) *Alligator mississippiensis* (Brusatte et al. 2016); (L) *Stegosaurus stenops* (Leahey et al. 2015); (M) *Camarasaurus lentus* (Witmer et al. 2008); (N) *Tyrannosaurus rex* (Witmer et al. 2008); (O) *Bubo virginianus* (Witmer et al. 2008). † denotes extinct taxa. Abbreviations: ASC, anterior semi-circular canal; ECD, endosseous cochlea duct; LSC, lateral semi-circular canal; PSC, posterior semi-circular canal. Phylogeny following Nesbitt (2011), Ezcurra (2016) and Leardi et al. (2017). Labyrinths not to scale.

166x87mm (300 x 300 DPI)