

***Xenoposeidon* is the earliest known rebbachisaurid sauropod dinosaur (#21907)**

1

First submission

Editor guidance

Please submit by **6 Dec 2017** for the benefit of the authors (and your \$200 publishing discount).



Structure and Criteria

Please read the 'Structure and Criteria' page for general guidance.



Raw data check

Review the raw data. Download from the location [described by the author](#).



Image check

Check that figures and images have not been inappropriately manipulated.

Privacy reminder: If uploading an annotated PDF, remove identifiable information to remain anonymous.

Files

Download and review all files from the [materials page](#).

5 Figure file(s)



Structure your review

The review form is divided into 5 sections.

Please consider these when composing your review:

1. BASIC REPORTING

2. EXPERIMENTAL DESIGN

3. VALIDITY OF THE FINDINGS

4. General comments

5. Confidential notes to the editor






 You can also annotate this PDF and upload it as part of your review

When ready [submit online](#).





Editorial Criteria

Use these criteria points to structure your review. The full detailed editorial criteria is on your [guidance page](#).





BASIC REPORTING

-  Clear, unambiguous, professional English language used throughout.
-  Intro & background to show context. Literature well referenced & relevant.
-  Structure conforms to [PeerJ standards](#), discipline norm, or improved for clarity.
-  Figures are relevant, high quality, well labelled & described.
-  Raw data supplied (see [PeerJ policy](#)).

EXPERIMENTAL DESIGN

-  Original primary research within [Scope of the journal](#).
-  Research question well defined, relevant & meaningful. It is stated how the research fills an identified knowledge gap.
-  Rigorous investigation performed to a high technical & ethical standard.
-  Methods described with sufficient detail & information to replicate.

VALIDITY OF THE FINDINGS

-  Impact and novelty not assessed. Negative/inconclusive results accepted. *Meaningful* replication encouraged where rationale & benefit to literature is clearly stated.
-  Data is robust, statistically sound, & controlled.
-  Conclusions are well stated, linked to original research question & limited to supporting results.
-  Speculation is welcome, but should be identified as such.

Standout reviewing tips

3



The best reviewers use these techniques

Tip

Support criticisms with evidence from the text or from other sources

Example

Smith et al (J of Methodology, 2005, V3, pp 123) have shown that the analysis you use in Lines 241-250 is not the most appropriate for this situation. Please explain why you used this method.

Give specific suggestions on how to improve the manuscript

Your introduction needs more detail. I suggest that you improve the description at lines 57- 86 to provide more justification for your study (specifically, you should expand upon the knowledge gap being filled).

Comment on language and grammar issues

The English language should be improved to ensure that an international audience can clearly understand your text. Some examples where the language could be improved include lines 23, 77, 121, 128 – the current phrasing makes comprehension difficult.

Organize by importance of the issues, and number your points

- 1. Your most important issue*
- 2. The next most important item*
- 3. ...*
- 4. The least important points*

Please provide constructive criticism, and avoid personal opinions

I thank you for providing the raw data, however your supplemental files need more descriptive metadata identifiers to be useful to future readers. Although your results are compelling, the data analysis should be improved in the following ways: AA, BB, CC

Comment on strengths (as well as weaknesses) of the manuscript

I commend the authors for their extensive data set, compiled over many years of detailed fieldwork. In addition, the manuscript is clearly written in professional, unambiguous language. If there is a weakness, it is in the statistical analysis (as I have noted above) which should be improved upon before Acceptance.

***Xenoposeidon* is the earliest known rebbachisaurid sauropod dinosaur**

Michael Taylor Corresp. 1

¹ Department of Earth Sciences, University of Bristol

Corresponding Author: Michael Taylor
Email address: dino@miketaylor.org.uk

Xenoposeidon proneneukos is a sauropod dinosaur represented by a single partial dorsal vertebra, NHMUK R2095, which consists of the centrum and the base of a tall neural arch. Despite its fragmentary nature, it is recognisably distinct from all other sauropods, and is here diagnosed with five unique characters. One character previously considered unique is here recognised as shared with *Rebbachisaurus garasbae*: an “M”-shaped arrangement of laminae on the lateral face of the neural arch. Following the more complete *Rebbachisaurus garasbae*, these laminae are now interpreted as ACPL and lateral CPRL, which intersect anteriorly; and PCDL and CPOL, which intersect posteriorly. Similar arrangements are also seen in some other rebbachisaurid specimens (though not all, possibly due to serial variation), but never in non-rebbachisaurid sauropods. *Xenoposeidon* is therefore referred to Rebbachisauridae. Due to its elevated parapophysis, the holotype vertebra is considered a posterior dorsal despite its elongate centrum. Since *Xenoposeidon* is from the Berriasian–Valanginian (earliest Cretaceous) Ashdown Beds Formation of the Wealden Supergroup of southern England, it is the earliest known rebbachisaurid by some 10 million years. Electronic 3D models were invaluable in determining *Xenoposeidon*'s true affinities: descriptions of complex bones such as sauropod vertebrae should always provide them where possible.

1 *Xenoposeidon* is the earliest known rebbachisaurid 2 sauropod dinosaur

3 Michael P. Taylor. Department of Earth Sciences, University of Bristol, Bristol BS8 1RJ,
4 England.
5 dino@miketaylor.org.uk

6 Abstract

7 *Xenoposeidon proneneukos* is a sauropod dinosaur represented by a single partial dorsal vertebra,
8 NHMUK R2095, which consists of the centrum and the base of a tall neural arch. Despite its
9 fragmentary nature, it is recognisably distinct from all other sauropods, and is here diagnosed
10 with five unique characters. One character previously considered unique is here recognised as
11 shared with *Rebbachisaurus garasbae*: an “M”-shaped arrangement of laminae on the lateral face
12 of the neural arch. Following the more complete *Rebbachisaurus garasbae*, these laminae are
13 now interpreted as ACPL and lateral CPRL, which intersect anteriorly; and PCDL and CPOL,
14 which intersect posteriorly. Similar arrangements are also seen in some other rebbachisaurid
15 specimens (though not all, possibly due to serial variation), but never in non-rebbachisaurid
16 sauropods. *Xenoposeidon* is therefore referred to Rebbachisauridae. Due to its elevated
17 parapophysis, the holotype vertebra is considered a posterior dorsal despite its elongate centrum.
18 Since *Xenoposeidon* is from the from the Berriasian–Valanginian (earliest Cretaceous) Ashdown
19 Beds Formation of the Wealden Supergroup of southern England, it is the earliest known
20 rebbachisaurid by some 10 million years. Electronic 3D models were invaluable in determining
21 *Xenoposeidon*'s true affinities: descriptions of complex bones such as sauropod vertebrae should
22 always provide them where possible.

Table of Contents

Introduction.....	2
Anatomical Abbreviations.....	2
Institutional Abbreviations.....	3
Reinterpretation.....	3
Serial position.....	6
Revised Reconstruction.....	7
Systematic Palaeontology.....	7
Discussion.....	8
Age.....	8
Wealden Rebbachisaurids.....	9
3D models of complex bones.....	9
Acknowledgements.....	9
References.....	10
Figure Captions.....	12
Supplementary Files.....	13

Introduction

Xenoposeidon proneneukos is a neosauropod sauropod dinosaur from the Berriasian–Valanginian (earliest Cretaceous) Ashdown Beds Formation of the Wealden Supergroup of southern England. It is represented by a single partial mid-to-posterior dorsal vertebra, NHMUK R2095 (BMNH R2095 at the time of the original description by Taylor and Naish 2007). This element consists of the centrum and the base of a tall neural arch, broken off below the transverse processes and zygapophyses. Despite its fragmentary nature, it is recognisably different from all other sauropods, and Taylor and Naish (2007) diagnosed it on the basis of six characters that they considered unique among sauropods.

D’Emic (2012:651) asserted that “the absence of diagnostic features renders *Xenoposeidon* a nomen dubium”. However, his assessment was mistaken in several respects. For example, the extension of the base of the neural arch to the posterior extremity of the centrum is clearly not, as he asserted, due to damage. D’Emic claimed that dorsal vertebrae illustrated by Osborn and Mook (1921:plates LXIX and LXXII) have forward-sloping neural arches resembling those of *Xenoposeidon*: in reality, only one posterior dorsal vertebrae out of four complete dorsal columns illustrated in that monograph shows a forward slope, and it differs so much from its fellows that this can only be interpreted as the result of crushing. D’Emic further claimed that the lamina patterns observed in *Xenoposeidon* can be recognised in other sauropods, but I have been unable find morphology resembling them in the descriptions he suggests: Osborn and Mook 1921 for *Camarasaurus*, Riggs 1903 for *Brachiosaurus* (probably a typo for Riggs 1904, which also does not depict similar patterns), Carballido et al. 2011 for *Tehuelchesaurus*. A similar pattern does appear in *Rebbachisaurus*, as will be discussed below. D’Emic (2012:651) is probably correct that the “asymmetric neural canal” described by Taylor and Naish (2007:1553–1554) is a misreading of the tall centroprezygapophyseal fossae as being the anterior portion of the neural canal: as Taylor and Naish pointed out, “The vacuity is filled with matrix, so the extent of its penetration posteriorly into the neural arch cannot be assessed”. Nevertheless, the shape and size of the fossa is unique among sauropods, and it is bounded by laminae which do not seem to be medial CPRLs. In summary, *Xenoposeidon proneneukos* is a valid, diagnosable taxon, *contra* D’Emic (2012).

Taylor and Naish (2007:1554–1557) compared the *Xenoposeidon* vertebra to those of the main neosauropod groups — Diplodocoidea, Camarasauridae, Brachiosauridae and Titanosauria — and concluded that it could not be convincingly referred to any of these groups. Their phylogenetic analysis (pp. 1157–1558 and figure 6) corroborated this by recovering *Xenoposeidon* as a neosauropod in all most parsimonious trees, but in a polytomy with all other neosauropods, wholly unresolved save that the clade Flagellicaudata was preserved in all MPTs.

In light of Wilson and Allain’s (2015) redescription of *Rebbachisaurus garasbae*, and the availability of more photographs and models of rebbachisaurid material, it has now become possible to reinterpret the idiosyncratic system of laminae found in *Xenoposeidon*, and to refer it confidently to an existing family-level clade.

Anatomical Abbreviations

- aEI — average elongation index *sensu* Chure et al. 2010: length of a centrum divided by the average of the height and width of the posterior articular surface.

- ACPL — anterior centroparapophyseal lamina.
- CPOL — centropostzygapophyseal lamina.
- CPRF — centroprezygapophyseal fossa.
- CPRL — centroprezygapophyseal lamina.
- EI — elongation index *sensu* Wedel et al. 2000: length of a centrum divided by the height of the posterior articular surface.
- PCDL — posterior centrodiapophyseal lamina.
- PCPL — posterior centroparapophyseal lamina.
- POSL — postspinal lamina.
- Postzyg — postzygapophysis.
- PPDL — paradiapophyseal lamina.
- Prezyg — prezygapophysis.
- PRPL — prezygaparapophyseal lamina.
- PRSL — prespinal lamina.
- SDL — spinodiapophyseal lamina.

Institutional Abbreviations

- IWCMS — Isle of Wight County Museum Service at Dinosaur Isle, Sandown, Isle of Wight, England.
- MIWG — Museum of Isle of Wight Geology (now Dinosaur Isle Visitor Centre), Sandown, Isle of Wight, England.
- MNHN — Muséum National d'Histoire Naturelle, Paris, France.
- NHMUK — the Natural History Museum, London, England.
- NMC — Canadian Museum of Nature (previously National Museum of Canada), Ottawa, Ontario, Canada.
- “WN” — “without number”, an informal designation for specimens awaiting accession.

Reinterpretation

Taylor and Naish’s (2007) history, geography, geology and description of the *Xenoposeidon* specimen requires no revision, and should continue to be considered definitive: this paper does not supersede it, but should be read in conjunction with it.

The illustrations of the specimen in the original paper, however, were in monochrome and omitted the dorsal and ventral views. The present paper supplements these illustrations with a colour depiction from all six cardinal directions (Figure 1), and a high-resolution 3D model of the specimen (supplementary file AA).

More importantly, Taylor and Naish’s (2007) interpretation of some features of the vertebra, particularly the “M”-shaped complex of laminae on the lateral faces of the neural arch, was mistaken. Although the neural spine and dorsal part of the neural arch are missing, including the pre- and postzygapophyses and lateral processes, they wrote that “sufficient laminae remain to allow the positions of the processes to be inferred with some certainty”. But their inferences were incorrect. Taylor and Naish (2007:1553) interpreted the cross-shaped structure on the anterodorsal part of the left lateral face of the neural arch as the site of the parapophysis, despite the lack of any articular facet in that location. This influenced their interpretation of the four laminae that met at that point as the ACPL below, the PPDL above, the PRPL anteriorly and an

unnamed accessory infrapapophyseal lamina posteroventrally, which they interpreted as homologous with a PCPL (Figure 2A). Similarly, they did not attempt to identify either the long lamina running up the posterior edge of the lateral face of the neural arch (designating it only “posterior lamina”) or the lamina forming a shallow “V” with the “accessory infrapapophyseal lamina”, simply calling it an “accessory postzygapophyseal lamina” (Figure 2A)

Among the various unusual features of the *Xenoposeidon* vertebra, the “M”-shaped set of laminae is immediately apparent in lateral view (Figure 3A): a line can be traced from the anterior margin of the neural arch’s lateral face up the ACPL to the cross that was interpreted as the parapophysis, then posteroventrally down the “accessory infrapapophyseal lamina”, then posterodorsally up the “accessory postzygapophyseal lamina” and finally down the posterior margin of the neural arch’s lateral face, along the “posterior lamina”. Photographs of other specimens that were available to us at this time did not apparently manifest similar features.

But subsequent work on *Rebbachisaurus garasbae* (Wilson 2012:100, Wilson and Allain 2015) — and an associated video of the rotating vertebra (see acknowledgements) — show that *Rebbachisaurus* has a similar complex of laminae (Figure 3B), which are described by Wilson and Allain (2015:6) as the second of the eight autapomorphies that they listed for the species: “infrazygapophyseal laminae (lat. CPRL, CPOL) that intersect and pass through neighbouring costal laminae (ACPL, PCDL) to form an ‘M’ shape”.

Because the illustrated dorsal vertebra of *Rebbachisaurus* — MNHN MRS 1958 — is substantially complete, it is possible to follow the trajectories of the laminae that participate in the “M” to their apophyses, and so determine their true identities. The two vertically oriented laminae — the outer pillars of the “M” — continue up past the top of the “M”. The anterior one supports the parapophysis, and the posterior supports the diapophysis. And the two laminae that form the valley in the middle of the “M” support the prezygapophyses and postzygapophyses: in both cases, as noted by Wilson and Allain, they intersect the vertical lamina before continuing to meet their respective zygapophyses. The four laminae that make up the “M”, from anterior to posterior, are therefore the ACPL, posterior part of the lateral CPRL, anterior part of the CPOL and PCDL. Of these, the intersection between the ACPL and lateral CPRL is clearly visible in left lateral view of MNHN MRS 1958. The intersection between the CPOL and PCDL is less apparent in this view, though clear in three dimensions. Both laminae continue dorsally beyond this intersection, but their paths are somewhat changed at the point of contact, with the dorsal portion of the PCDL inclining more anteriorly, and the rod-like CPOL apparently passing through the sheet of bone formed by the PCDL to meet the postzygapophysis.

The referred *Rebbachisaurus garasbae* specimen NMC 50844 described and illustrated by Russell (1996:388–390 and figure 30) is also broadly consistent with this morphology. It is not possible to be definite about the laminar intersection based only on line drawings of the specimen from the four cardinal directions, but, as illustrated in Russell’s figure 30c, the lateral CPRL does appear to pass through the ACPL. The CPOL seems in this specimen to originate posterior to the PCDL, not intersecting with it. But this difference from the holotype dorsal may be serial variation since, as Russell notes, the relatively longer centrum of his specimen indicates a more anterior serial position than for the holotype’s dorsal vertebra; and this interpretation is corroborated by the observation that, based on lamina trajectories, the anteroposterior distance between the parapophysis and diapophysis was less in NMC 50844 than in the holotype.

In light of these *Rebbachisaurus* specimens, the mysterious laminae of *Xenoposeidon* are easily explained. It is now apparent that the cross on the side of the *Xenoposeidon* vertebra is not the site of the parapophysis, as Taylor and Naish (2007:1553) proposed, but merely the intersection

of two laminae that pass right through each other: the ACPL, running dorsolaterally, and the lateral CPRL, extending anterodorsally to the (missing) prezygapophysis (Figure 2B). Similarly, the “posterior lamina” is the PCDL, and it intersects with the CPOL, though the intersection is lost in NHMUK R2095 (Figure 2B). Both the parapophysis and diapophysis of the *Xenoposeidon* vertebrae would have been located some distance above the preserved portion, the former anterior to the latter.

It appears from Dalla Vecchia (1999:figure 47, left part) that in the holotype and only vertebra of *Histriasaurus boscarollii*, “WN-V6”, the CPOL on the right side of the vertebra intersects with the PCDL in the same way as in *Rebbachisaurus*, though it is not possible to determine whether the lateral CPRL similarly intersects the ACPL. Dorsal vertebrae of other rebbachisaurid sauropods, however, do not appear to feature the distinctive “M” and intersecting laminae of *Rebbachisaurus* and *Xenoposeidon*:

- The 3D model of a dorsal vertebra of *Nigersaurus* (Serenio et al. 2007) shows that the lateral CPRLs originate anterior to the ACPLs and the CPOLs posterior to the PCDLs, so that there is no intersection. A subtle “V” shape does appear high up on the lateral faces of the neural arch, between the ACPL and the PCDL, but it seems unrelated to the lateral CPRL and CPOL.
- Unpublished 3D models of an anterior dorsal neural arch and a more posterior dorsal vertebra of *Katapultisaurus* (pers. comm., Lucio M. Ibiricu) as illustrated in figures 3A and 5A of Ibiricu et al. (2017) show that in both vertebrae, the lateral CPRLs originate anterior to the ACPLs, and the CPOLs seem to originate posterior to the PCDLs — though damage to the posterior portion makes the latter uncertain.
- The laminae do not appear to intersect in the illustrated dorsal vertebra of *Demandasaurus* (Fernández-Baldor et al. 2011:figure 9).
- The sole known vertebra of *Nopcsaspondylus* seems to have an entirely different pattern of lamination (Mannion 2010:figure 5) with no lamina intersections like those of MNHN MRS 1958.

No determination can be made for other rebbachisaurids as they are insufficiently preserved (e.g. *Limaysaurus*, *Amazonsaurus*), or illustrated (e.g. *Cathartesaura*), or simply lack posterior dorsal vertebral material (e.g. *Rayososaurus*, *Tataouinea*, *Comahuesaurus*, *Zapalasaurus*).

However, we cannot rule out the possibility that complete and well-preserved posterior dorsal vertebrae of most or all rebbachisaurids have *Rebbachisaurus*-like intersecting laminae: even in those species for which a well-preserved vertebra lacks them, this could be due to serial variation, with these features only fully developing in the most posterior dorsals.

Xenoposeidon, then, resembles *Rebbachisaurus* in the possession of a distinctive “M” on the lateral face of the neural arch, in the intersecting lateral CPRL and ACPL, and in the elevation of the parapophysis above the level of the prezygapophysis — a complex of related features. Although at first glance they appear rather different, *Xenoposeidon* and *Rebbachisaurus*, while geometrically different, are topologically similar.

Regarding the significance of the elevated parapophysis, since no complete or nearly complete rebbachisaurid dorsal column has been described, comparisons with other, better represented sauropods are warranted. In the probable basal diplodocoid *Haplocanthosaurus*, the dorsal margin of the parapophyseal facet reaches the level of, and is coincident with, the prezygapophyseal facet around dorsal vertebra 7 or 8, but never rises any higher than this in more

posterior vertebrae (Hatcher 1903:plate I). In the more distantly related diplodocid diplodocoids *Apatosaurus* and *Diplodocus*, the parapophysis never migrates far enough dorsally to reach a position level with the prezygapophyses, even in the most posterior dorsals (Gilmore 1936:plate XXV; Hatcher 1901:plates VII, VIII).

Taylor and Naish (2007:1554) argued that *Xenoposeidon* could not at that time be convincingly referred to Rebbachisauridae because *Rebbachisaurus* differs from NHMUK R2095 in five ways: “possession of a very prominent PCPL, large and laterally diverging prezygapophyses, depressions at the base of the neural arch (Bonaparte 1999:173), lateral foramina not set within fossae, and a strongly arched ventral border to the centrum.” Of these features, the first is now recognised as occurring in *Xenoposeidon*; the second appears to be an outright error, as the prezygapophyses of *Rebbachisaurus* meet on the midline, and in any case the situation in *Xenoposeidon* is not known. “Depressions at the base of the neural arch” seems to be a mistranslation of Bonaparte’s original Spanish, “profundas depresiones en la base de la espina neural”, which refers not to the neural arch but the neural spine, and since this portion is not preserved in *Xenoposeidon*, it is not informative for our purposes. The 3D model of the *Rebbachisaurus* dorsal shows that in fact its lateral foramina are set in shallow depression, similar in quality if not in degree to those of *Xenoposeidon*. This leaves the stronger arching of the ventral border of the centrum in *Rebbachisaurus*, a feature that in isolation is not convincing. In conclusion, the weight of morphological evidence supports including *Xenoposeidon* within Rebbachisauridae. This is in accordance with the observation of Taylor and Naish (2007:1557), in whose phylogenetic analysis “various most-parsimonious trees also recover *Xenoposeidon* in many other positions, including as a ... rebbachisaurid.”

Serial position

The serial position of the *Rebbachisaurus garasbae* holotype dorsal vertebra MNHN MRS 1958 is not definitely known. However, it has been uniformly referred to as a posterior dorsal, most likely due to the very elevated position of its parapophyses and Lavocat’s (1954) initial assessment of it as “une des dernières dorsales” (one of the last dorsals) — perhaps made with knowledge of the spatial relation of bones in the quarry.

The position of the *Xenoposeidon proneneukos* holotype vertebra NHMUK R2095 is of course even more difficult to determine in light of the limited nature of the specimen, though its similarity to MNHN MRS 1958 suggests a similar position. Taylor and Naish (2007:1553) wrote that “the high position of the parapophysis on the neural arch of R2095 indicates a mid to posterior placement of the vertebra within the dorsal column, but, because the prezygapophyses must have been dorsal to it, it was probably not among the most posterior vertebrae in the sequence.” With the location of the parapophysis now interpreted as significantly higher than previously thought, and probably well above the prezygapophysis, an even more posterior position is indicated.

This posterior serial position is surprising in light of the anteroposterior length of the *Xenoposeidon* centrum. Its posterior articular surface measures 160 mm high by 170 mm wide, while the length of even the preserved portion of the centrum is 190 mm, and it must have been at least 200 mm long when complete (Taylor and Naish 2007:table 1). As noted by Taylor and Naish (2007:1554), “the length of the centrum, especially in so posterior a dorsal vertebra, argues against [a diplodocoid identity]: the posterior dorsal centra of diplodocoids typically have EI < 1.0, compared with 1.25 for R2095” — or 1.21 using the aEI of Chure et al. (2010:384). However, rebbachisaurids may be unusual among diplodocoids in this respect — perhaps

unsurprisingly, as they diverged early from the line leading to diplodocids, with their characteristically short dorsal centra, and likely retained something more similar to the ancestral neosauropod condition. Wilson and Allain (2015:8) give the centrum measurements of MNHN MRS 1958 as posterior height 231 mm, posterior width 220 mm and length 220 mm. This yields an aEI of 0.98, meaning that the *Xenoposeidon* centrum is only 24% more elongate than that of *Rebbachisaurus*. This is a significant difference, but not an outlandish one. For comparison, the centrum of the basal rebbachisaurid *Histriasaurus boscarollii* holotype “WN-V6” is relatively elongate, with its posterior articular surface measuring 150 mm high and centrum length of “more than 200 mm” (Dalla Vecchia 1998:122) yielding an EI of > 1.33. Also, the aEIs of the last four dorsal vertebrae of the *Brachiosaurus altithorax* holotype FMNH PR 25107 are 1.34, 1.27, 1.19 and 0.96 (calculated from the table of Riggs 1904:34): so aEIs of sauropod dorsals can vary, within two serial positions of the same individual, from values below that of MNHN MRS 1958 to above that of NHMUK R2095.

In conclusion, while the evidence regarding the serial position of NHMUK R2095 remains equivocal, it suggests a more posterior position than previous inferred — it can be fairly confidently described as “posterior” rather than “mid-to-posterior” — but it is unlikely to be the very last dorsal.

Revised Reconstruction

In light of the reassignment of *Xenoposeidon* to Rebbachisauridae, and the reinterpretation of its laminae, I present a new reconstruction of how the vertebra NHMUK R2095 might have looked when complete (Figure 4). As in MNHN MRS 1958, the parapophysis and diapophysis are both elevated above the zygapophyses. The lateral CPRL and ACPL meet at a point where they project outwards about the same distance from the vertebra, as is apparent from the preserved portion of the vertebra; but the CPOL is assumed to pass through a sheet-like PCDL as in *Rebbachisaurus*, because it is clear from breakage in NHMUK R2095 that the PCDL extended further from the body of the neural arch than the preserved portion indicates. The neural spine, composed as in *Rebbachisaurus* of pre- and post-spinal laminae together with the left and right SDLs, is shown fading out at the top, as there is no way to determine its height. The condyle that is the centrum’s anterior articular surface is reconstructed as only slightly convex, as in *Rebbachisaurus*.

It is instructive to compare this with the original reconstruction of the vertebrae (Taylor and Naish:figure 5). The new reconstruction has a taller neural arch, a far more elevated parapophysis, a more posteriorly located diapophysis (no longer dorsal to the parapophysis) and a shallower condyle, as that of the original reconstruction was drawn with those of brachiosaurs in mind.

Systematic Palaeontology

Dinosauria Owen, 1842
 Saurischia Seeley, 1888
 Sauropodomorpha Huene, 1932
 Sauropoda Marsh, 1878
 Neosauropoda Bonaparte, 1986
 Rebbachisauridae Sereno et al., 1999
Xenoposeidon Taylor and Naish, 2007

285 *Xenoposeidon proneneukos* Taylor and Naish, 2007

286 **Holotype.** NHMUK R2095, the Natural History Museum, London. A mid posterior dorsal
287 vertebra consisting of partial centrum and neural arch.

288 **Revised diagnosis:** Differs from all other sauropods in the following characters:

- 289 1. neural arch covers dorsal surface of centrum, with its posterior margin continuous with
290 that of the centrum;
- 291 2. neural arch slopes anteriorly 35 degrees relative to the vertical;
- 292 3. broad, flat area of featureless bone on lateral face of neural arch;
- 293 4. very large, teardrop-shaped centroprezygapophyseal fossa.
- 294 5. arched laminae form vaulted boundary of centroprezygapophyseal fossa.

295 The “arched laminae” of #5 are not the medial CPRLs, as these arise from the neural arch
296 pedicels — and the laminae arising from the pedicels cannot instead be regarded lateral CPRLs,
297 as those laminae are located on the lateral face of the neural arch, intersecting with the ACPLs.
298 Furthermore, the point where the supporting laminae meet at the top of their arch is located some
299 way posterior to the inferred location of the prezygapophyses (Figure 5).

300 Discussion

301 Age

302 As shown by the Wilson and Allain (2015:table 1), the 19 then-recognised rebbachisaurids (of
303 which 13 had been named) span the middle third of the Cretaceous. The earliest recognised taxon
304 is *Histriasaurus boscarollii* from the upper Hauterivian or lower Barremian limestones of
305 southwest Istria, Croatia. Seven taxa, of which five are named, survived at least to the
306 Cenomanian (earliest Late Cretaceous), of which two (*Katepensaurus goicoecheai* and
307 *Limaysaurus tessonei*) may be from the Turonian age.

308 As discussed by Taylor and Naish (2007:1547–1548), the precise location and horizon where
309 NHMUK R2095 was excavated was not recorded in the specimen’s original brief description,
310 which only said “the Wealden of Hastings” (Lydekker 1893:276). However, records of the
311 collection of Philip James Rufford, who collected the specimen, indicate that the most likely
312 location is Ecclesbourne Glen, a mile or two east of Hastings, East Sussex (see discussion in
313 Taylor and Naish 2007:1548). The units exposed at Ecclesbourne Glen are part of the Ashdown
314 Beds Formation, which straddles the Berriasian/Valanginian boundary; but the part of the
315 formation at that location is from the earlier Berriasian age. If this assessment is correct, then
316 *Xenoposeidon* is from the very earliest Cretaceous, giving it an age of around 140 million years
317 — about 10 million years earlier than *Histriasaurus*.

318 This early age is consonant with a basal position within Rebbachisauridae, a possibility that is
319 corroborated by *Xenoposeidon*’s camerate internal morphology compared with the camellate
320 centra of most rebbachisaurids. However, further material will be required before numerical
321 phylogenetic work can firmly establish its position within the group.

Wealden Rebbachisaurids

Although *Xenoposeidon* is the first named Rebbachisaurid from the Wealden Supergroup of southern England, other material from this unit has been referred to Rebbachisauridae. Naish and Martill (2001:plate 36, opposite page 236) illustrated some isolated sauropod teeth IWCMS.2001.201–203, and these were referred to Rebbachisauridae by Sereno and Wilson (2005:174). Mannion (2009) described a partial rebbachisaurid scapula MIWG 6544. Finally, Mannion et al. (2011) described a proximal caudal neural arch MIWG 5384, which they also interpreted as rebbachisaurid. All of these specimens are from the Barremian Wessex Formation of the Isle of Wight, so they could all belong to the same species or genus. However, since the likely Berriasian age of NHMUK R2095 makes it 10–15 Mya older than these specimens, it is unlikely that they belong to *Xenoposeidon*, but to some other as yet-unnamed rebbachisaurid. Thus it is likely that the Wealden Supergroup contains at least two rebbachisaurid sauropods.

3D models of complex bones

Electronic 3D models were invaluable in determining *Xenoposeidon*'s true affinities. Most obviously, the model of the *Xenoposeidon* vertebra itself, created by Heinrich Mallison, has functioned as an invaluable proxy for the fossil itself when I am unable to visit the NHMUK, and I have consulted it many times in writing this paper. I would also have been unable to determine to my own satisfaction whether the *Katepensaurus* dorsals feature intersecting laminae like those of *Rebbachisaurus* without the models provided by Lucio M. Ibiricu. Although no true model is available for the *Rebbachisaurus* dorsal itself or for the dorsal vertebrae of *Nigersaurus*, rotating videos were crucial in enabling me to understand their morphology. When interpreting specimens for which no such models exist, such as Russell's (1996) referred *Rebbachisaurus* specimen NMC 50844, the conclusions reached using only 2D representations — whether photographs or drawings — are much less well founded.

Techniques such as photogrammetry (see e.g. Falkingham 2012; Mallison and Wings 2014) are reducing the barriers to the creation of high-quality 3D models in full colour. Doing so is now inexpensive in both time and money. In light of our discipline's goal of making palaeontology more accessible and reproducible, then, it should become increasingly routine in the 21st Century to provide 3D models as a standard part of the description of complex bones such as sauropod vertebrae.

Acknowledgements

I thank Sandra D. Chapman (Natural History Museum, London) for access to the *Xenoposeidon* specimen, and Heinrich Mallison (Palaeo3D) who went far beyond the call of duty in building the 3D model of NHMUK R2095 and talking me through aspects of photogrammetry. I am also grateful to Jeff Wilson (University of Michigan) and Ronan Allain (Muséum National d'Histoire Naturelle, Paris) for sharing high-resolution photographs of the French *Rebbachisaurus* vertebra, and to Mathew J. Wedel (Western University of Health Sciences) and Darren Naish (University of Southampton) for helpful discussion. Lucio M. Ibiricu kindly provided access to unpublished 3D models of an anterior dorsal neural arch and a more posterior dorsal vertebra of *Katepensaurus*.

As noted in Taylor (2015), this project began when I recognised the true identity of the curious laminae on the *Xenoposeidon* vertebra while viewing a rotating video of the *Rebbachisaurus garasbae* holotype dorsal vertebra MNHN MRS 1958 on the University of Michigan Museum of

365 Paleontology's UMORF web-site (University of Michigan Online Repository of Fossils) at
 366 <https://umorf.ummp.lsa.umich.edu/wp/gallery/vertebrate-animations/>. This video was based on a
 367 3D reconstruction created from CT scans performed at the AST-RX (Accès Scientifique à la
 368 Tomographie à Rayons X) of the MNHN by F. Goussard.

369 References

- 370 Bonaparte, Jose F. 1986. Les dinosaures (Carnosaures, Allosauridés, Sauropodes, Cétiosauridés)
 371 du Jurassique moyen de Cerro Cónдор (Chubut, Argentina). *Annales de Paléontologie* **72**:325–
 372 386.
- 373 Bonaparte, Jose F. 1999. Evolucion de las vertebras presacras en Sauropodomorpha [Evolution of
 374 the presacral vertebrae in Sauropodomorpha]. *Ameghiniana* **36**(2):115–87.
- 375 Carballido, Jose L., Oliver W. M. Rauhut, Diego Pol and Leonardo Salgado. 2011. Osteology and
 376 phylogenetic relationships of *Tehuelchesaurus benitezii* (Dinosauria, Sauropoda) from the
 377 Upper Jurassic of Patagonia. *Zoological Journal of the Linnean Society* **163**:605–662.
 378 doi:10.1111/j.1096-3642.2011.00723.x
- 379 Chure, Daniel, Brooks B. Britt, John A. Whitlock and Jeffrey A. Wilson. 2010. First complete
 380 sauropod dinosaur skull from the Cretaceous of the Americas and the evolution of sauropod
 381 dentition. *Naturwissenschaften* **97**(4):379–91. doi:10.1007/s00114-010-0650-6
- 382 Dalla Vecchia, Fabio M. 1998. Remains of Sauropoda (Reptilia, Saurischia) in the Lower
 383 Cretaceous (Upper Hauterivian/Lower Barremian) Limestones of SW Istria (Croatia).
 384 *Geologia Croatica* **51**(2):105–134.
- 385 Dalla Vecchia, Fabio M. 1999. Atlas of the sauropod bones from the Upper Hauterivian-Lower
 386 Barremian of Bale/Valle (SW Istria, Croatia). *Natura Nascosta* **18**:6–41.
- 387 D'Emic, Michael D. 2012. The early evolution of titanosauriform sauropod dinosaurs. *Zoological*
 388 *Journal of the Linnean Society* **166**:624–671.
- 389 Falkingham, Peter L. 2012. Acquisition of high resolution 3D models using free, open-source,
 390 photogrammetric software. *Palaeontologia Electronica* **15**(1):1T. 15 pages. [http://palaeo-](http://palaeo-electronica.org/content/issue1-2012technical-articles/92-3d-photogrammetry)
 391 [electronica.org/content/issue1-2012technical-articles/92-3d-photogrammetry](http://palaeo-electronica.org/content/issue1-2012technical-articles/92-3d-photogrammetry)
- 392 Fernández-Baldor, Fidel Torcida, José Ignacio Canudo, Pedro Huerta, Diego Montero, Xabier
 393 Pereda Suberbiola and Leonardo Salgado. 2010. *Demandasaurus darwini*, a new
 394 rebbachisaurid sauropod from the Early Cretaceous of the Iberian Peninsula. *Acta*
 395 *Palaeontologica Polonica* **56**(3):535–552. doi:10.4202/app.2010.0003
- 396 Gilmore, Charles W. 1936. Osteology of *Apatosaurus* with special reference to specimens in the
 397 Carnegie Museum. *Memoirs of the Carnegie Museum* **11**:175–300 and plates XXI–XXXIV.
- 398 Hatcher, Jonathan B. 1901. *Diplodocus* (Marsh): its osteology, taxonomy and probable habits,
 399 with a restoration of the skeleton. *Memoirs of the Carnegie Museum* **1**:1–63 and plates I–XIII.
- 400 Hatcher, J. B. 1903. Osteology of *Haplocanthosaurus* with description of a new species, and
 401 remarks on the probable habits of the Sauropoda and the age and origin of the *Atlantosaurus*
 402 beds; additional remarks on *Diplodocus*. *Memoirs of the Carnegie Museum* **2**:1–75 and plates
 403 I–VI.

- 404 Huene, Friedrich von. 1932. Die fossile Reptile-Ordnung Saurischia, ihre Entwicklung und
405 Geschichte. *Monographien zur Geologie und Palaeontologie* (Serie 1) **4**:1–361.
- 406 Ibiricu, Lucio M., Matthew C. Lamanna, Rubén D.F. Martínez, Gabriel A. Casal, Ignacio A.
407 Cerda, Gastón Martínez and Leonardo Salgado. 2017. A novel form of postcranial skeletal
408 pneumaticity in a sauropod dinosaur: Implications for the paleobiology of Rebbachisauridae.
409 *Acta Palaeontologica Polonica* **62**(2):221–236. doi:10.4202/app.00316.2016
- 410 Lavocat, René J. M. 1954. Sur les Dinosauriens du continental intercalaire des Kem-Kem de la
411 Daoura. [On the dinosaurs of the Continental Intercalaire of the Kem Kem of the Daoura].
412 *Comptes Rendus 19th International Geological Congress* **1952**(1):65–68. English translation
413 by Matthew C. Lamanna provided by the *Polyglot Paleontologist* at
414 http://paleoglot.org/files/Lavocat_54.pdf
- 415 Lydekker, Richard. 1893. On a sauropodous dinosaurian vertebra from the Wealden of Hastings.
416 *Quarterly Journal of the Geological Society, London* **49**:276–280.
- 417 Mallison, Heinrich, and Oliver Wings. 2014. Photogrammetry in paleontology — a practical
418 guide. *Journal of Paleontological Techniques* **12**:1–31.
- 419 Mannion, Philip D. 2009. A rebbachisaurid sauropod from the Lower Cretaceous of the Isle of
420 Wight, England. *Cretaceous Research* **30**:521–526. doi:10.1016/j.cretres.2008.09.005
- 421 Mannion, Philip D. 2010. A revision of the sauropod dinosaur genus '*Bothriospondylus*' with a
422 redescription of the type material of the middle jurassic form '*B. madagascariensis*'.
423 *Palaeontology* **53**(2):277–296. doi:10.1111/j.1475-4983.2009.00919.x
- 424 Mannion, Philip D., Paul Upchurch and Stephen Hutt. 2011. New rebbachisaurid (Dinosauria:
425 Sauropoda) material from the Wessex Formation (Barremian, Early Cretaceous), Isle of Wight,
426 United Kingdom. *Cretaceous Research* **32**(6):774–780. doi:10.1016/j.cretres.2011.05.005
- 427 Marsh, Othniel C. 1878. Principal characters of American Jurassic dinosaurs, part I. *American*
428 *Journal of Science* (Series 3) **16**:411–416.
- 429 Naish, Darren, and David M. Martill. 2001. Saurischian dinosaurs I: Sauropods. pp. 185–241 in:
430 Martill, David M., and Darren Naish (eds.). *Dinosaurs of the Isle of Wight*. Palaeontological
431 Association, London.
- 432 Osborn, Henry Fairfield, and Charles C. Mook. 1921. *Camarasaurus*, *Amphicoelias* and other
433 sauropods of Cope. *Memoirs of the American Museum of Natural History*, new series
434 **3**(3):247–387, and plates LX–LXXXV.
- 435 Owen, Richard. 1842. Report on British fossil reptiles, Part II. *Reports of the British Association*
436 *for the Advancement of Science* **11**:60–204.
- 437 Riggs, Elmer S. 1903. *Brachiosaurus altithorax*, the largest known dinosaur. *American Journal*
438 *of Science* **15**(4):299–306.
- 439 Riggs, Elmer S. 1904. Structure and relationships of opisthocoelian dinosaurs. Part II, the
440 Brachiosauridae. *Field Columbian Museum, Geological Series* **2**(6):229–247, plus plates
441 LXXI–LXXXV.
- 442 Russell, Dale A. 1996. Isolated dinosaur bones from the Middle Cretaceous of the Tafila,
443 Morocco. *Bulletin du Muséum National d'Histoire Naturelle*, 4ème série – section C: Sciences
444 de la Terre, Paléontologie, Géologie, Minéralogie **18**(2–3):349–402.

- 445 Seeley, Harry G. 1888. On the classification of the fossil animals commonly named Dinosauria.
446 *Proceedings of the Royal Society of London* **43**:165–171.
- 447 Sereno, Paul C., and Jeffrey A. Wilson. 2005. Structure and evolution of a sauropod tooth battery.
448 pp. 157–177 in: Wilson, Jeffrey A., and Kristina Curry-Rogers (eds.), *The Sauropods:*
449 *Evolution and Paleobiology*. University of California Press, Berkeley.
- 450 Sereno, Paul C., Jeffrey A. Wilson, Lawrence A. Witmer, John A. Whitlock, Abdoulaye Maga,
451 Oumarou Ide, Timothy A. Rowe. 2007. *Nigersaurus taqueti* (On-line), Digital Morphology.
452 Accessed November 14, 2017 at
453 http://digimorph.org/specimens/Nigersaurus_taqueti/dorsal_vertebra/
- 454 Sereno, Paul C., Allison L. Beck, Didier. B. Dutheil, Hans C. E. Larsson, Gabrielle. H. Lyon,
455 Bourahima Moussa, Rudyard W. Sadleir, Christian A. Sidor, David J. Varricchio, Gregory P.
456 Wilson and Jeffrey A. Wilson. 1999. Cretaceous sauropods from the Sahara and the uneven
457 rate of skeletal evolution among dinosaurs. *Science* **282**:1342–1347.
- 458 Taylor, Michael P. 2015. Is *Xenoposeidon* a rebbachisaur? *Sauropod Vertebra Picture of the Week*,
459 July 14, 2015. Accessed November 15, 2017 at [https://svpow.com/2015/07/14/is-](https://svpow.com/2015/07/14/is-xenoposeidon-a-rebbachisaur/)
460 [xenoposeidon-a-rebbachisaur/](https://svpow.com/2015/07/14/is-xenoposeidon-a-rebbachisaur/)
- 461 Taylor, Michael P., and Darren Naish. 2007. An unusual new neosauropod dinosaur from the
462 Lower Cretaceous Hastings Beds Group of East Sussex, England. *Palaeontology* **50**(6):1547–
463 1564. doi: 10.1111/j.1475-4983.2007.00728.x
- 464 Wedel, Mathew J., Richard L. Cifelli and R. Kent Sanders. 2000. Osteology, paleobiology, and
465 relationships of the sauropod dinosaur *Sauroposeidon*. *Acta Palaeontologica Polonica*
466 **45**(4):343–388.
- 467 Wilson, Jeffrey A. 2012. New vertebral laminae and patterns of serial variation in vertebral
468 laminae of sauropod dinosaurs. *Contributions from the Museum of Paleontology, University of*
469 *Michigan* **32**(7):91–110. <http://hdl.handle.net/2027.42/92460>
- 470 Wilson, Jeffrey A., and Ronan Allain. 2015. Osteology of *Rebbachisaurus garasbae* Lavocat,
471 1954, a diplodocoid (Dinosauria, Sauropoda) from the early Late Cretaceous-aged Kem Kem
472 beds of southeastern Morocco. *Journal of Vertebrate Paleontology* **35**(4):e1000701.
473 doi:10.1080/02724634.2014.1000701

474 Figure Captions

475 **Figure 1.** NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, shown
476 from all six cardinal directions. Top row: **A.** dorsal view, with anterior to the left. Middle row, left
477 to right: **B.** anterior, **C.** left lateral, **D.** posterior and **E.** right lateral view. Bottom row: **F.** ventral
478 view, with anterior to the left. Scale bar = 200 mm.

479 **Figure 2.** NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left
480 lateral view, with interpretative drawing. **A.** The incorrect interpretation of the laminae from
481 Taylor and Naish (2017:figure 4A), with identifying captions greyed out since they are largely
482 incorrect. **B.** The revised interpretation of the same laminae, based on the similar arrangement in
483 *Rebbachisaurus garasbae*. Scale bar = 200 mm.

484 **Figure 3.** Centra and neural arches of posterior dorsal vertebrae from two rebbachisaurid
485 sauropods (not to scale), highlighting the distinctive “M” shape formed by laminae high on the

neural arch. **A.** NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*.
B. MNHN MRS 1958, a posterior dorsal vertebra from the holotype specimen of *Rebbachisaurus*
garasbae.

Figure 4. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left lateral view, interpreted as a rebbachisaurid. This interpretation is modelled primarily on MNHN MRS 1958, a posterior dorsal vertebra from the holotype specimen of *Rebbachisaurus garasbae*. The CPOL passes through a sheetlike PCDL, as in *Rebbachisaurus*; but the lateral CPRL forms a cross-shaped junction with the ACPL, each of these laminae equally interrupting the trajectory of the other. Abbreviations as used in the text. Scale bar = 200 mm.

Figure 5. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left anteroventrolateral view, highlighting the three sets of laminae related to the prezygapophyses. The trajectories of the medial CPRLs (which emerge from the neural arch pedicels) and the lateral CPRLs (which intersect with the APCLs) indicate the approximate position of the prezygapophyses. The additional arched laminae form the margins of the large, teardrop-shaped CPRF, but meet at a position some way below and posterior to the presumed location of the prezygapophyseal facets. Breakage of both medial CPRLs and the left ACPL and PCDL is indicated by cross-hatching. Note that, from this perspective, the lateral CPRL appears to turn a corner where it intersects with the ACPL, such that the posteroventral portion of the lateral CPRL appears contiguous with the dorsal portion of the ACPL. This is an illusion brought about by the eminence at the point of intersection. As always, this is much easier to see in three dimensions. Abbreviations as used in the text.

Supplementary Files

Supplementary file 1. Three-dimensional surface model (11 million polygons) of NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*. A 3D polygon mesh file was created by Heinrich Mallison in Agisoft Photoscan Pro version 1.3.0 (agisoft.com), from 95 high resolution digital photographs by the author. All 95 images aligned, and resulted in a dense point cloud at maximum resolution of 20,900,043 points and 44,871,128 polygons. Scaling was based on a single 10 cm scale bar created from a high quality scale bar placed in the pictures with the specimen. Available from <https://doi.org/10.6084/m9.figshare.5605612.v2>

Figure 1

NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, shown from all six cardinal directions.

Figure 1. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, shown from all six cardinal directions. Top row: **A.** dorsal view, with anterior to the left. Middle row, left to right: **B.** anterior, **C.** left lateral, **D.** posterior and **E.** right lateral view. Bottom row: **F.** ventral view, with anterior to the left. Scale bar = 200 mm.

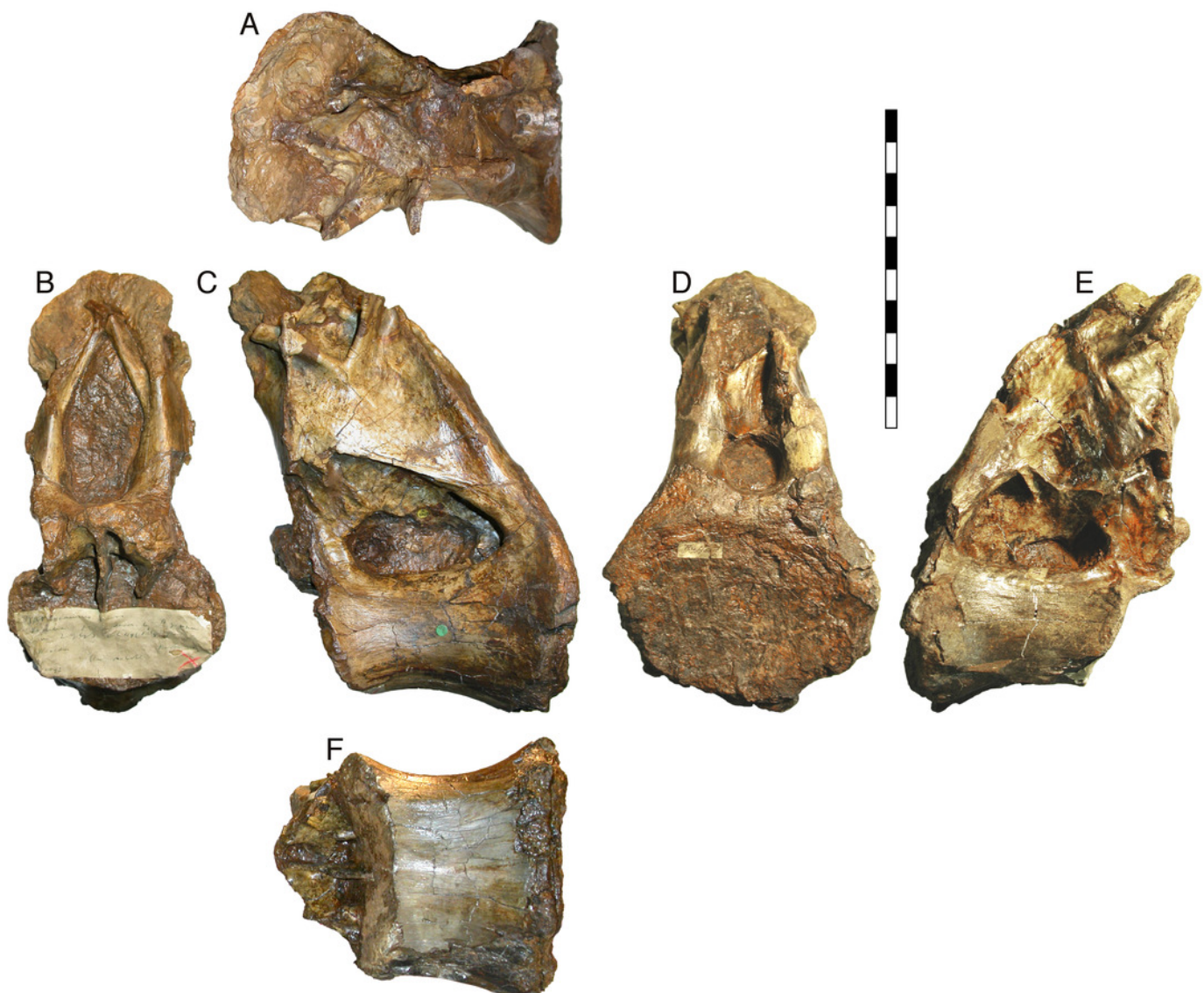


Figure 2

Figure 2. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left lateral view, with interpretative drawing.

Figure 2. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left lateral view, with interpretative drawing. **A.** The incorrect interpretation of the laminae from Taylor and Naish (2017:figure 4A), with identifying captions greyed out since they are largely incorrect. **B.** The revised interpretation of the same laminae, based on the similar arrangement in *Rebbachisaurus garasbae*. Scale bar = 200 mm.

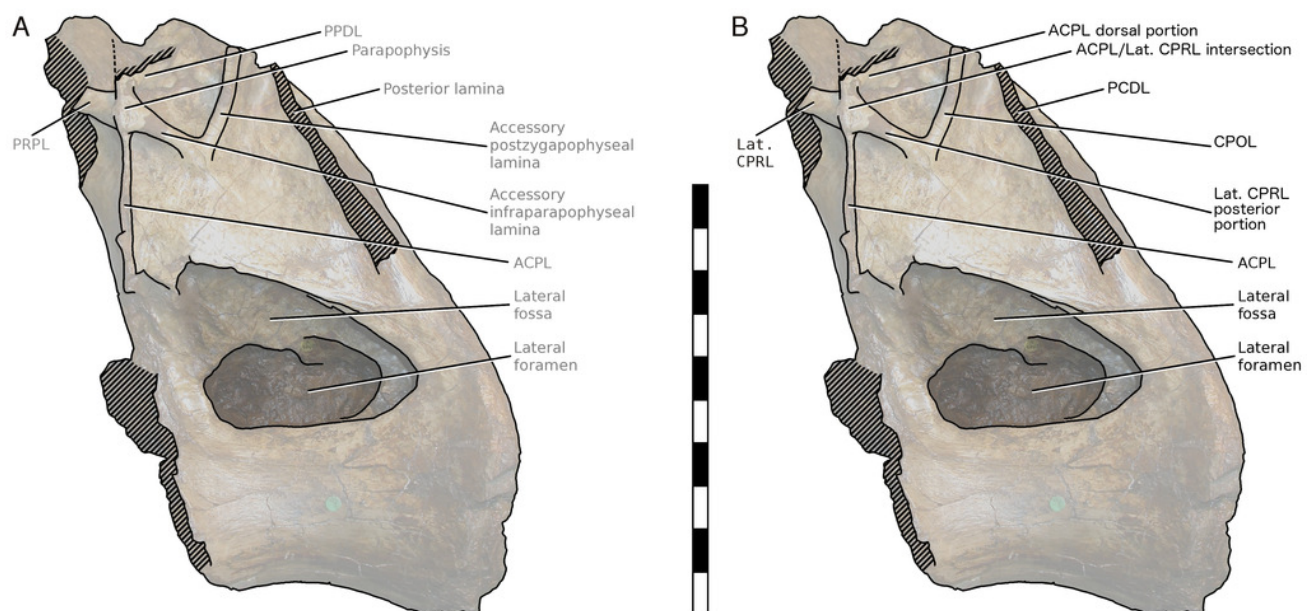


Figure 3

Figure 3. Centra and neural arches of posterior dorsal vertebrae from two rebbachisaurid sauropods (not to scale), highlighting the distinctive “M” shape formed by laminae high on the neural arch.

Figure 3. Centra and neural arches of posterior dorsal vertebrae from two rebbachisaurid sauropods (not to scale), highlighting the distinctive “M” shape formed by laminae high on the neural arch. **A.** NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*. **B.** MNHN MRS 1958, a posterior dorsal vertebra from the holotype specimen of *Rebbachisaurus garasbae*.

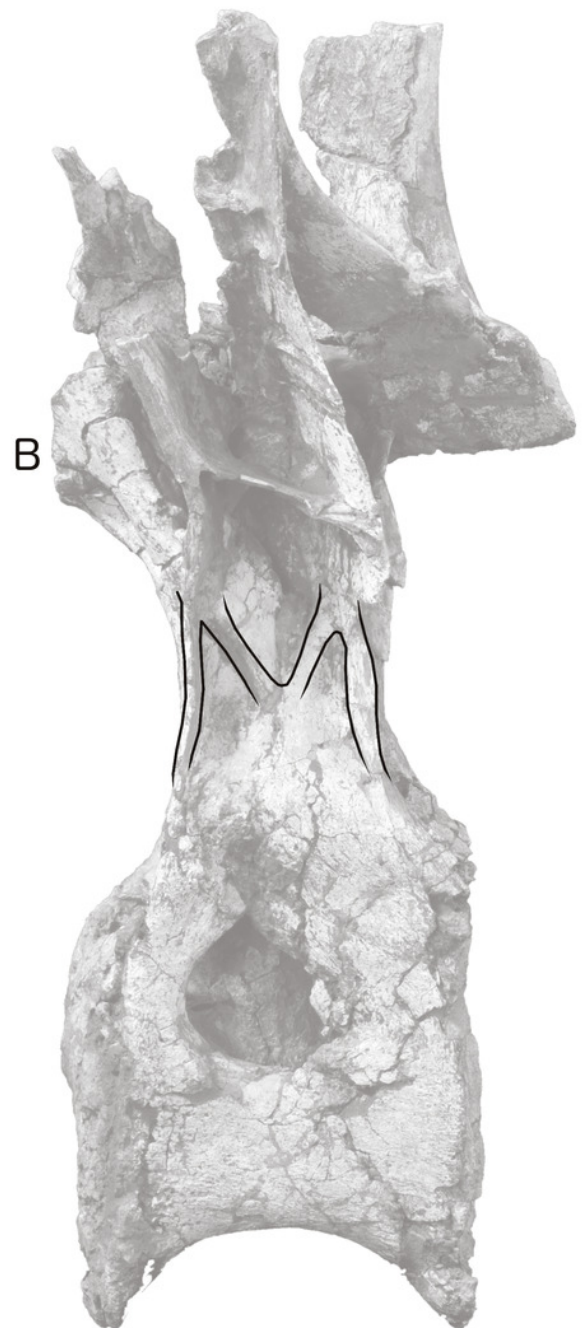


Figure 4

Figure 4. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left lateral view, interpreted as a rebbachisaurid.

Figure 4. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left lateral view, interpreted as a rebbachisaurid. This interpretation is modelled primarily on MNHN MRS 1958, a posterior dorsal vertebra from the holotype specimen of *Rebbachisaurus garasbae*. The CPOL passes through a sheetlike PCDL, as in *Rebbachisaurus*; but the lateral CPRL forms a cross-shaped junction with the ACPL, each of these laminae equally interrupting the trajectory of the other. Abbreviations as used in the text. Scale bar = 200 mm.

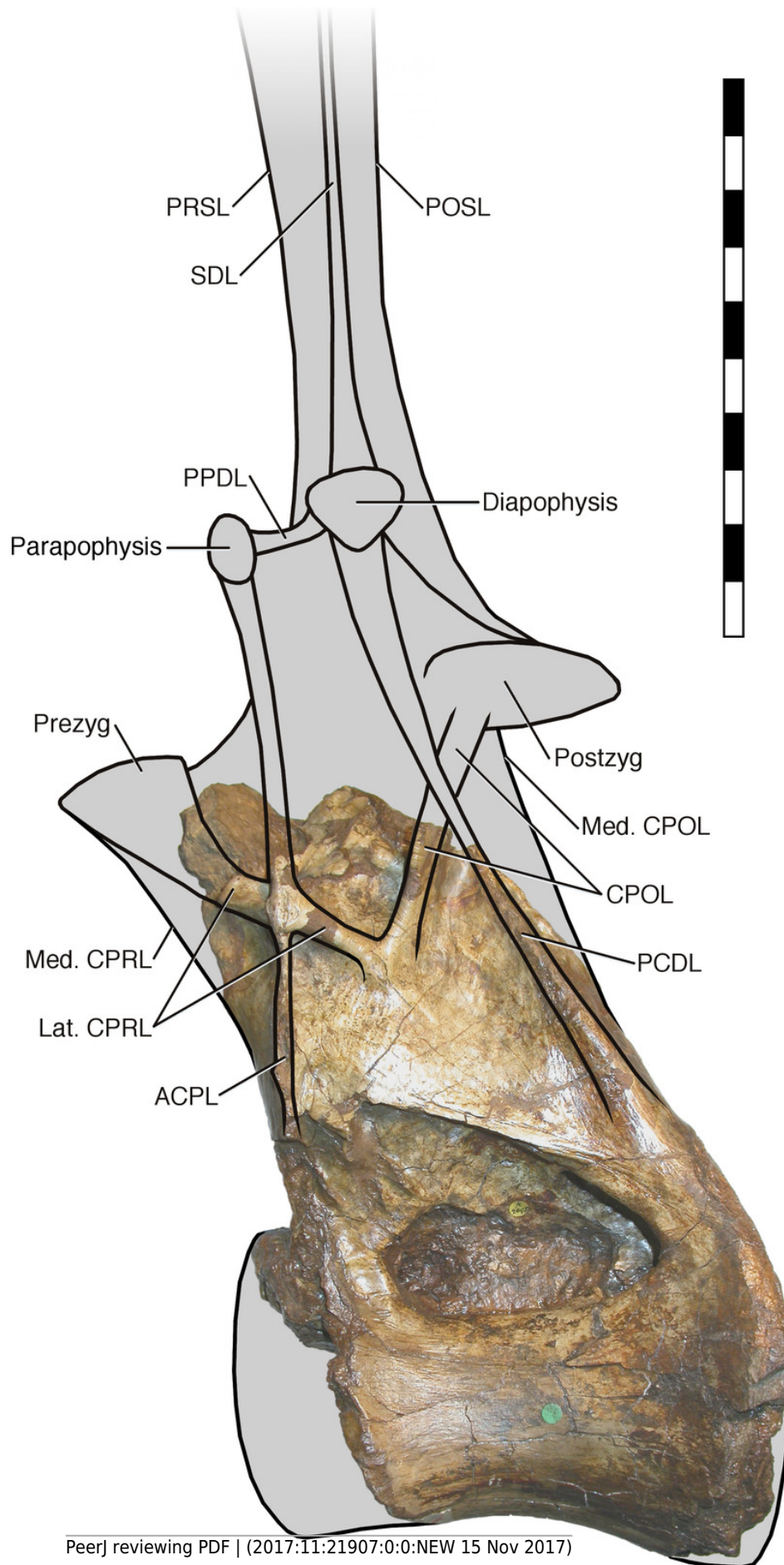


Figure 5

Figure 5. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left anteroventrolateral view, highlighting the three sets of laminae related to the prezygapophyses.

Figure 5. NHMUK R2095, the holotype and only vertebra of *Xenoposeidon proneneukos*, in left anteroventrolateral view, highlighting the three sets of laminae related to the prezygapophyses. The trajectories of the medial CPRLs (which emerge from the neural arch pedicels) and the lateral CPRLs (which intersect with the APCLs) indicate the approximate position of the prezygapophyses. The additional arched laminae form the margins of the large, teardrop-shaped CPRF, but meet at a position some way below and posterior to the presumed location of the prezygapophyseal facets. Breakage of both medial CPRLs and the left ACPL and PCDL is indicated by cross-hatching. Note that, from this perspective, the lateral CPRL appears to turn a corner where it intersects with the ACPL, such that the posteroventral portion of the lateral CPRL appears contiguous with the dorsal portion of the ACPL. This is an illusion brought about by the eminence at the point of intersection. As always, this is much easier to see in three dimensions. Abbreviations as used in the text.

