



Research



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Oldest record of Machimosaurini (Thalattosuchia, Teleosauroida): teeth and scavenging traces from the Middle Jurassic (Bajocian) of Switzerland

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The Jurassic period was a time of major diversification for Mesozoic marine reptiles, including Ichthyosauria, Plesiosauria and thalattosuchian Crocodylomorpha. The latter originated in the Early Jurassic and thrived during the Late Jurassic. Unfortunately, the Middle Jurassic, a crucial time in their evolution, has a poor fossil record. Here, we document the first evidence of macrophagous/durophagous Machimosaurini-tribe teleosauroid thalattosuchians from the late Bajocian (*ca* 169 Ma) in the form of three robust tooth crowns with conical blunt shapes and anastomosed pattern of thick enamel ridges towards the apex, associated with the skeleton of a large ichthyosaur lacking preserved tooth crowns. The tooth crowns were found on the posterior section of the lower jaw (left angular), a lacrimal and the axis neural arch of the ichthyosaur. In addition, some of the distal sections of the posterior dorsal ribs of the ichthyosaur skeleton exhibit rounded bite marks and some elongated furrows that fit in size and shape with the Machimosaurini teeth. These marks, together with the absence of healing in the rib bone are

interpreted here as the indicators of peri- to post-mortem scavenging by a Machimosaurini teleosauroid after the large ichthyosaur carcass settled on the floor of a shallow ocean.

1. Introduction

Many Mesozoic marine reptile clades have their origins in the Triassic period, but for some of the most diverse and long-lived groups, an important radiation happened during the Early Jurassic (*ca* 201–145 Ma). Ichthyosaurs and plesiosaurs, which first appeared in the Early and Late Triassic, respectively [1,2], and thalattosuchian crocodylomorphs, which originated in the early Lower Jurassic (Hettangian–Sinemurian [3]), diversified and reached a diversity peak during the Toarcian [4]. For the Middle Jurassic, however, species diversity and rock record analyses indicate a drastic decline in the richness of marine reptile species, especially for shallow marine inhabitants during the Aalenian to Bajocian (*ca* 174–168 Ma). Diversity levels recovered only in the late Middle Jurassic (Bathonian/Callovian), a phenomenon linked to sea-level change, general availability of marine fossiliferous formations, and the effect of highly productive Lagerstätten [5–7]. Thus, early Middle Jurassic reptiles constitute rare finds, with only a few valid species described globally (e.g. [8–10]).

The uniquely marine crocodylomorph clade Thalattosuchia comprises two groups, Metriorhinoidea and Teleosauroidea [11]. Following the Toarcian, their record is characterized by Aalenian, Bathonian and Oxfordian materials, with five of the six Bathonian species belonging to Machimosauridae, a speciose clade inhabiting the western part of the Tethys Ocean [12]. Within Machimosauridae, certain taxa show a distinctly robust jaw morphology and blunt tooth shape that make them different from other machimosaurids having a specialized macrophagous/durophagous diet [13–15]. These taxa constitute the Machimosaurini tribe within Machimosauridae. *Yoridiosuchus boutillieri* [16] is the oldest Machimosaurini-tribe teleosauroid, appearing in the Bathonian, and the group gets more firmly established based on the record of *Lemmysuchus obtusidens* [17] from the UK and France in the Callovian. The Machimosaurini record extends to the Kimmeridgian (Late Jurassic), except for *Machimosaurus rex*, the largest machimosaurine teleosauroid from the early Lower Cretaceous of Tunisia [12,18]. It is noteworthy that the appearance of the macrophagous/durophagous morphotype in crocodylomorphs coincides with the appearance of large macropredatory pliosaurs in the Bajocian of Germany, France and Switzerland [9,19].

The purpose here is to describe three tooth crowns from the Bajocian, which constitute the oldest record of the Machimosaurini tribe. These teeth are considered a rare find indeed, as they were recovered in association with the skeletal remains of a new Middle Jurassic ichthyosaur [10]. Finally, the bite marks on the ichthyosaur skeleton are identified as potential scavenging traces left by the teeth of the Machimosaurini.

2. Material and methods

2.1. Material

The ichthyosaur skeleton (PIMUZ A/III 5279 [10]) and the three Machimosaurini teeth (two larger tooth crowns, PIMUZ A/III 5281a and PIMUZ A/III 5281b, associated with a lacrimal and the left angular of the ichthyosaur skeleton, respectively; PIMUZ A/III 5281c, a smaller tooth crown associated with the axis neural arch) were found in the Oberegg quarry, Auenstein, Canton Aargau, Switzerland (figure 1a). They were recovered from the Lower Acuminata beds of the Hauptrogenstein Formation (*subfurcatum/niortense* zone), which are late Bajocian (figure 1b) in age [20]. The original association of the teeth and the ichthyosaur skeleton was kept intact so that the teeth were still attached to the bones by a thin layer of limestone matrix, an exception being the third tooth, which was isolated to facilitate the examination of its morphology.

2.2. Methods

The documentation of the teeth and the bite marks on the ichthyosaur ribs was performed using a Keyence vhx-7000 digital microscope. In addition, the smaller of the tooth crowns (PIMUZ A/III

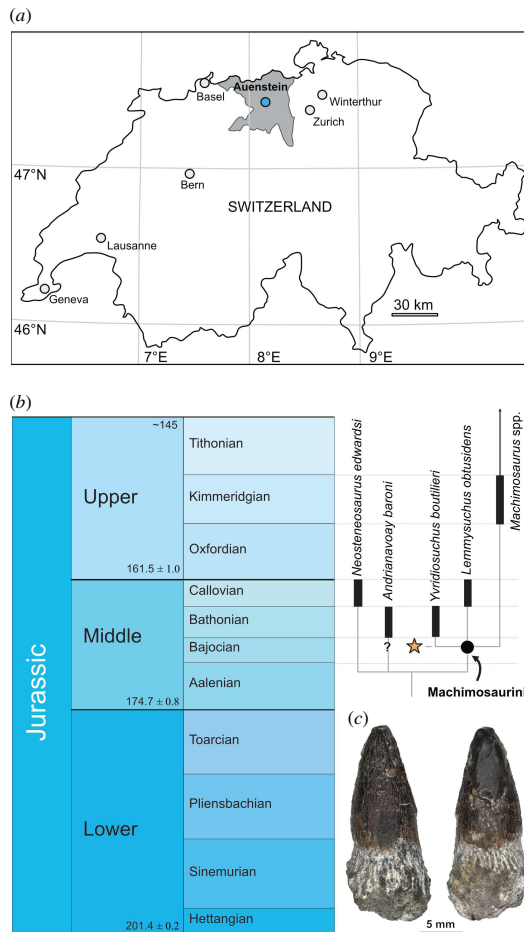


Figure 1. Finding location and phylogenetic framework of Swiss Machimosaurini tooth crowns. (a) Map of Switzerland with Canton of Aargau (in grey) and location of Auenstein indicated. (b) Time scale and abbreviated phylogeny based on Johnson *et al.* [12] with hypothesized position of new Machimosaurini indet. indicated by an orange star. (c) Isolated larger tooth crown (PIMUZ A/III 5281a).

5281c) was computed tomography (CT)-scanned in-house at the University of Zurich with a Nikon XTH 225 ST CT scanner. The CT scan was taken with a voltage of 141 kV and a current of 206 μ A, yielding a voxel size of 0.01027787 mm, with no filter used. The reconstruction of the digital stack (originally 3142 rotational projections, two frames, 250.0 ms exposure and 2000 tif slices in stack) for virtual three-dimensional reconstruction was done using MIMICS Innovation Suite 25.0 (Materialise NV, Leuven, Belgium).

2.3. Abbreviations

PIMUZ collections of the Department of Paleontology, University of Zurich, Zurich, Switzerland.

3. Results

3.1. Systematic palaeontology

- Crocodylomorpha Walker, 1968 ([21] *sensu* [22])
- Thalattosuchia Fraas, 1901 ([23] *sensu* [24])
- Teleosauroida Geoffroy Saint-Hilaire, 1831 ([25] *sensu* [24])
- Machimosauridae Johnson *et al.*, 2020 ([12] *sensu* [26])
- Machimosaurinae Johnson *et al.*, 2020 ([12] *sensu* [26])
- Tribe Machimosaurini Jouve *et al.*, 2016 ([27])
- Machimosaurini indet. (figures 1 and 2)

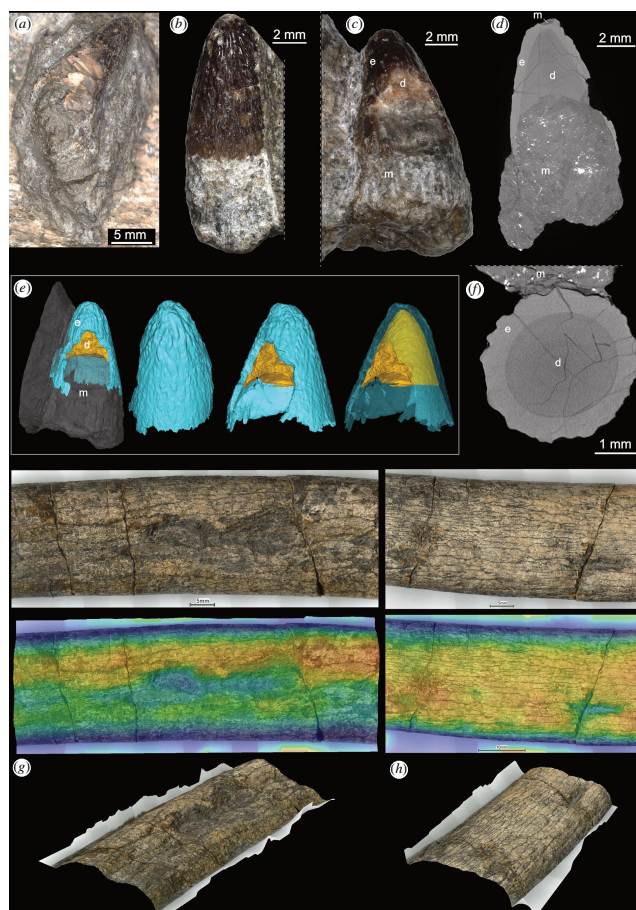


Figure 2. Photographs, CT scan and three-dimensional images of Swiss Machimosaurini tooth crowns and selected scavenging traces on the ichthyosaur ribs (PIMUZ A/III 5279). (a) Tooth crown PIMUZ A/III 5281b on left angular. (b,c) Tooth crown PIMUZ A/III 5281c on the axis neural arch. (d,f) CT scan images showing longitudinal and horizontal sections of PIMUZ A/III 5281c. (e) Three-dimensional model of PIMUZ A/III 5281c in different views, not to scale. (g,h) Normal, false colour and angled views of the ribs showing bite marks. Abbreviations: d, dentine; e, enamel; m, sediment matrix.

3.2. Anatomical description of tooth crowns

All teeth exhibit a wide conical crown shape, which are blunt and robust and show neither carinae or labiolingual (= mediolateral) compression nor identifiable true or false denticles (compared to [28,29]). An anastomosed pattern of enamel ridges extends apicobasally over the whole crown. There appears to be no curvature at the apices. All crowns also experienced some local damage to the enamel and dentine; however, whether this damage occurred during the animal's life, fossilization or preparation cannot be elucidated. The enamel thickness increases apically, although, without additional CT scanning, the boundary between the pulp cavity and dentine is difficult to discern in the larger teeth. In the two larger specimens, the dentine infilling is visible due to breaks in the enamel caps in both cases.

PIMUZ A/III 5281a (figure 1c) is a larger tooth crown, 15.53 mm long and up to 8.92 mm wide. About one-third of the apex of the crown was damaged and needed to be stabilized with a dark synthetic resin. Otherwise, the crown appears mostly complete with the apex having experienced slight erosion. The enamel ridges appear to be shallower at the base of the tooth crown to become more distinct, higher and more strongly branching towards the apex. Parallel marks in the sediment matrix proximal to the crown are identified as preparation traces left by pneumatic tools and are not part of any root tissue. PIMUZ A/III 5281b (figure 2) is very similar in proportion, being 15.41 mm high as preserved and 9.83 mm wide. The apex is complete without showing any damage at the tip or any worn facets. In PIMUZ A/III 5281c (figure 2), the crown is 9.08 mm high as preserved and 6.82 mm wide. The internal pulp cavity is restricted to about the lower half of the preserved crown, with dentine filling the upper half. The thickness of the outer enamel layer increases from the base of the crown to the apex, reaching a thickness of 0.5–0.65 mm. Apically, the enamel is locally damaged, leaving behind a shallow depression filled with sediment. The enamel ridges on the apex of the tooth appear

to have an anastomosed (= wrinkled) texture, but they are not closely situated to one another and are more knobbly and 'chaotic'/disorganized compared with other machimosaurins such as *Yoridiosuchus boutillieri*, *Lemmingsuchus obtusidens* and *Machimosaurus* spp. [12,29–31]. The apicobasal enamel ridges are not well developed but appear to extend parallel from the perceived crown base to approximately three-quarters of the entire tooth. The enamel–dentine junction is smooth rather than scalloped, such that the ridges are constituted entirely by thickening of the enamel layer.

3.3. Bone modifications on the ichthyosaur ribs

Several ribs in the posterior trunk region of the ichthyosaur skeleton PIMUZ A/III 5279 show surficial marks on the bone cortex and upper parts of the cancellous interior (figure 2). These bone modifications are localized, often in rows and sometimes appear on opposite sides of the ribs. Furthermore, these modifications can be identified as furrows, pits and deeper punctures, which led to the collapse of the rib cortex [32] but did not pierce the ribs completely. The collapse of the rib cortex also argues against chemical or parasitic bone modification (e.g. [33]). The pitted and punctured rib bone also indicates that the tooth causing these modifications was of wide and blunt shape, likely lacking carinae [32]. There is no indication of healing or calluses forming around the marks.

4. Discussion and conclusions

The three tooth crowns are identified as those belonging to Machimosaurini indeterminate due to their blunt and wide crown shape, the anastomosed ('wrinkled') pattern of enamel ridges on the tooth apex and the absence of scalloping of the enamel–dentine junction underlying the enamel ridges. These three major dental characteristics of the group [12,29] set them apart from ichthyosaur and plesiosaur/pliosaur teeth. In these other marine reptile groups, apicobasal enamel ridges remain usually separate or show few anastomosing junctions [9,34–37]; in ichthyosaurs in particular, enamel ridges are formed through scalloping of the enamel–dentine junction rather than solely through thickening of the enamel layer [38]. The thickened enamel cap (more than 0.5 mm) of the crown, underlying massive dentine patch with little pulp cavity expansion towards the apex and overall tooth robustness, can further be interpreted as indicative of a macrophagous or durophagous diet [34].

Besides being blunt and robust in shape, the new Swiss specimens show round cross-sections with no mediolateral compression and short and often branching enamel ridges along the lower half of the crown, rather than longer parallel ones. The teeth of early diverging Machimosaurini such as *Lemmingsuchus obtusidens* and *Yoridiosuchus boutillieri*, while also being large and robust and lacking mediolateral compression, are typically carinated and their crowns showing apicobasal curvature and denticulation on their mostly parallel-trending enamel ridges [28,30,31]. Round cross-sections and lack of carinae have been noted for some machimosaurine teeth as well, as seen in specimens pertaining to Machimosaurini indet. from the Corallian Group, UK [35, fig. 9], but these teeth are apicobasally curved with more ordered enamel ridges along the non-apical crown.

The lack of apparent tooth crown curvature in the new Swiss specimens could indicate that either (i) there simply was no curvature in any of these teeth or (ii) the teeth come from the posterior part of the tooth row in which teeth might be less curved compared to anterior teeth (as in *Yoridiosuchus* [30]). If the former would be the case, this together with the 'irregular' anastomosing pattern of the enamel ridges, the shorter branching enamel ridges, and the lack of carinae and denticulation of these Bajocian specimens might hint at the presence of a new, potentially early diverging taxon within Machimosaurini. Given the scarcity of the material, however, this remains speculative for now, while at the same time hinting at the hitherto untapped potential of filling the gaps in marine crocodyliform evolution, particularly in the Middle Jurassic.

The bone modifications found on the ichthyosaur ribs likely appeared peri- or post-mortem as bone remodelling and healing processes are absent. The large size of the ichthyosaur and the inferred thick layer of soft tissue overlying the ribs make scavenging a more plausible source for these traces rather than predation. Taphonomic interpretation suggests that the carcass lay exposed on the seafloor for a prolonged period prior to burial [10]. Furthermore, only posterior thoracic ribs bear the bite marks, which suggests that the scavenging machimosaurin(s) accessed the tissues from behind like many other scavengers today [39,40]. Despite the tooth crowns being found more anterior on the ichthyosaur skeleton and not in direct vicinity of the traces on the ribs, a close fit of tooth crown morphology with the shape of the bone modifications indicates that the latter are consistent with scavenging traces (e.g.

[41]) of a small- to medium-sized macrophagous Machimosaurini thalattosuchian crocodyliform that co-inhabited the marine environment in the Bajocian.

Ethics. Dealing exclusively with fossils, this work did not require specific ethical approval. Permission to excavate the fossils was obtained by the Department Bau, Verkehr und Umwelt of the Kanton Aargau, Aarau, Switzerland, and by the company Jura Cement Fabriken AG, Wildeggen, Switzerland.

Data accessibility. All the new Swiss fossils mentioned in the text are officially accessioned in the collections of the Department of Paleontology, University of Zurich, and can be studied upon request. The CT data set and the 3D surface model (in .ply format) are provided on Morphosource repository project ID: 000593818 [42].

Declaration of AI use. We have not used AI-assisted technologies in creating this article.

Authors' contributions. C.K.: conceptualization, data curation, funding acquisition, project administration, resources, validation, writing—review and editing; T.M.S.: conceptualization, data curation, formal analysis, funding acquisition, investigation, visualization, writing—original draft, writing—review and editing; M.M.J.: investigation, validation, writing—original draft; E.E.M.: investigation, validation, writing—review and editing; D.B.: investigation, validation, writing—review and editing; F.M.: investigation, validation, writing—review and editing.

All authors gave final approval for publication and agreed to be held accountable for the work performed therein.

Conflict of interest declaration. We declare we have no competing interests.

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References

- Wintrich T, Hayashi S, Houssaye A, Nakajima Y, Sander PM. 2017 A Triassic plesiosaurian skeleton and bone histology inform on evolution of a unique body plan. *Sci. Adv.* **3**, e1701144. (doi:10.1126/sciadv.1701144)
- Moon BC, Stubbs TL. 2020 Early high rates and disparity in the evolution of ichthyosaurs. *Commun. Biol.* **3**, 68. (doi:10.1038/s42003-020-0779-6)
- Benani H, Nehili A, Ouzzaoui LA, Jouve S, Boudad L, Masrour M, Jalil NE, Arrad TY. 2023 Discovery of the teleosauroid crocodylomorph from the Early Jurassic of Chaara cave, Middle Atlas of Morocco. *J. Afr. Earth Sci.* **198**, 104804. (doi:10.1016/j.jafrearsci.2022.104804)
- Wilberg EW, Turner AH, Brochu CA. 2019 Evolutionary structure and timing of major habitat shifts in Crocodylomorpha. *Sci. Rep.* **9**, 514. (doi:10.1038/s41598-018-36795-1)
- Mannion PD, Benson RBJ, Carrano MT, Tennant JP, Judd J, Butler RJ. 2015 Climate constrains the evolutionary history and biodiversity of crocodylians. *Nat. Commun.* **6**, 8438. (doi:10.1038/ncomms9438)
- Benson RBJ, Butler RJ, Lindgren J, Smith AS. 2010 Mesozoic marine tetrapod diversity: mass extinctions and temporal heterogeneity in geological megabiases affecting vertebrates. *Proc. Biol. Sci.* **277**, 829–834. (doi:10.1098/rspb.2009.1845)
- Benson RBJ, Butler RJ. 2011 Uncovering the diversification history of marine tetrapods: ecology influences the effect of geological sampling biases. In *Comparing the geological and fossil records: implications for biodiversity studies* (eds AJ McGowan, AB Smith), pp. 191–208. London, UK: Geological Society. (doi:10.1144/SP358.13)
- Druckenmiller PS, Maxwell EE. 2014 A Middle Jurassic (Bajocian) ophthalmosaurid (Reptilia: Ichthyosauria) from the Tuxedni Formation, Alaska and the early divergence of the clade. *Geol. Mag.* **151**, 41–48. (doi:10.1017/S0016756813000125)
- Sachs S, Madzia D, Thuy B, Kear BP. 2023 The rise of macropredatory pliosaurs near the Early-Middle Jurassic transition. *Sci. Rep.* **13**, 17558. (doi:10.1038/s41598-023-43015-y)
- Miedema F, Bastiaans D, Scheyer TM, Klug C, Maxwell EE. 2024 A large new Middle Jurassic ichthyosaur shows the importance of body size evolution in the origins of the Ophthalmosauridae. *BMC Ecol. Evol.* **24**, 34. (doi:10.1186/s12862-024-02208-3)
- Foffa D, Young MT, Brusatte SL, Graham MR, Steel L. 2017 A new metriorhynchid crocodylomorph from the Oxford Clay Formation (Middle Jurassic) of England, with implications for the origin and diversification of Geosaurini. *J. Syst. Palaeontol.* **16**, 1–21. (doi:10.1080/14772019.14772017.11367730)
- Johnson MM, Young MT, Brusatte SL. 2020 The phylogenetics of Teleosauroidea (Crocodylomorpha, Thalattosuchia) and implications for their ecology and evolution. *PeerJ* **8**, e9808. (doi:10.7717/peerj.9808)
- Johnson MM, Foffa D, Young MT, Brusatte SL. 2022 The ecological diversification and evolution of Teleosauroidea (Crocodylomorpha, Thalattosuchia), with insights into their mandibular biomechanics. *Ecol. Evol.* **12**, e9484. (doi:10.1002/ecs3.9484)
- Karl HV, Tichy G. 2004 The structure of fossil teeth of chelonochagous crocodiles (Diapsida: Crocodylia). *Stud. Geol. Salmant* **40**, 116–124.

15. Vignaud P. 1997 La morphologie dentaire des *Thalattosuchia* (Crocodylia, Mesosuchia). *Palaeovertebrata* **26**, 35–59.
16. Eudes-Deslongchamps E. 1867 Notes sur le squelette et la restauration du *Teleosaurus cadomensis*. *Bulletin de La Société Linnéenne de Normandie, Caen, série 2* **2**, 381–473.
17. Andrews CW. 1909 XXXVIII. — On some new Steneosaus from the Oxford Clay of Peterborough. *Ann. Mag. Nat. Hist.* **3**, 299–308. (doi:10.1080/00222930908692579)
18. Fanti F, Miyashita T, Cantelli L, Mnasri F, Dridi J, Contessi M, Cau A. 2016 The largest thalattosuchian (Crocodylomorpha) supports teleosaurid survival across the Jurassic–Cretaceous boundary. *Cretac. Res.* **61**, 263–274. (doi:10.1016/j.cretres.2015.11.011)
19. Sachs S, Klug C, Kear BP. 2019 Rare evidence of a giant pliosaurid-like plesiosaur from the Middle Jurassic (lower Bajocian) of Switzerland. *Swiss J. Palaeontol.* **138**, 337–342. (doi:10.1007/s13358-019-00200-9)
20. Gonzalez R, Wetzel A. 1996 Stratigraphy and paleogeography of the Hauptrogenstein and Klingnau Formations (middle Bajocian to late Bathonian), northern Switzerland. *Eclogae Geol. Helv.* **89**, 695–720.
21. Walker AD. 1968 *Protosuchus*, *Proterochampsia*, and the origin of phytosaurs and crocodiles. *Geol. Mag.* **105**, 1–14. (doi:10.1017/S0016756800046434)
22. Irmis RB, Nesbitt SJ, Sues HD. 2013 Early Crocodylomorpha. In *Anatomy, phylogeny and palaeobiology of early Archosaurs and their kin* (eds SJ Nesbitt, JB Desojo, RB Irmis), pp. 275–302. London, UK: Geological Society. (doi:10.1144/SP379.24)
23. Fraas E. 1901 Die Meerkrokodile (*Thalattosuchia* n. g.) eine neue Sauriergruppe der Juraformation. *Jahreshefte Des Vereins Für Vaterländische Naturkunde, Württemberg* **57**, 409–418.
24. Young MT, De Andrade MB. 2009 What is *Geosaurus*? Redescription of *Geosaurus giganteus* (*Thalattosuchia*: Metriorhynchidae) from the Upper Jurassic of Bayern, Germany. *Zool. J. Linn. Soc.* **157**, 551–585. (doi:10.1111/j.1096-3642.2009.00536.x)
25. Geoffroy Saint-Hilaire E. 1831 Recherches sur de grands sauriens trouvés à l'état fossile vers les confins maritimes de la Basse Normandie, attribués d'abord au crocodile, puis déterminés sous les noms de *Télosaurus* et *Sténo-saurus*. Paris, France: Firmin Didot frères.
26. Johnson MM, Young MT, Brignon A, Brusatte SL. 2022 Addition to 'The phylogenetics of Teleosauroida (Crocodylomorpha; Thalattosuchia) and implications for their ecology and evolution'. *Bull. Phylogenet. Nom.* **1**, 1–7. (doi:10.11646/bpn.1.1.1)
27. Jouve S, Mennecart B, Douteau J, Jalil NE. 2016 The oldest durophagous teleosauroid (Crocodylomorpha, Thalattosuchia) from the lower Bathonian of central High Atlas, Morocco. *Palaeontology* **59**, 863–876. (doi:10.1111/pala.12262)
28. Young MT, Beatty BL, Brusatte SL, Steel L. 2015 First evidence of denticulated dentition in teleosaurid crocodylomorphs. *Acta Palaeontol. Pol.* **60**, 661–671. (doi:10.4202/app.00002.02013)
29. Young MT, Steel L, Brusatte SL, Foffa D, Lepage Y. 2014 Tooth serration morphologies in the genus *Machimosaurus* (Crocodylomorpha, Thalattosuchia) from the Late Jurassic of Europe. *R. Soc. Open Sci.* **1**, 140269. (doi:10.1098/rsos.140269)
30. Johnson MM, Young MT, Brusatte SL. 2020 Re-description of two contemporaneous mesorostrine teleosauroids (Crocodylomorpha: Thalattosuchia) from the Bathonian of England and insights into the early evolution of Machimosaurini. *Zool. J. Linn. Soc.* **189**, 449–482. (doi:10.1093/zoolinnean/zlzo37)
31. Johnson MM, Young MT, Steel L, Foffa D, Smith AS, Hua S, Havlik P, Howlett EA, Dyke G. 2018 Re-description of '*Steneosaurus*' *obtusidens* Andrews, 1909, an unusual macrophagous teleosaurid crocodylomorph from the Middle Jurassic of England. *Zool. J. Linn. Soc.* **182**, 385–418. (doi:10.1093/zoolinnean/zlx035)
32. Drumheller SK, Brochu CA. 2016 Phylogenetic taphonomy: a statistical and phylogenetic approach for exploring taphonomic patterns in the fossil record using crocodylians. *PALAIOS* **31**, 463–478. (doi:10.2110/palo.2016.030)
33. Zonneveld JP, AbdelGawad MK, Miller ER. 2022 Ectoparasite borings, mesoparasite borings, and scavenging traces in early Miocene turtle and tortoise shell: Moghra Formation, Wadi Moghra, Egypt. *J. Paleontol.* **96**, 304–322. (doi:10.1017/jpa.2021.92)
34. McCurry MR, Evans AR, Fitzgerald EMG, McHenry CR, Bevirt J, Pyenson ND. 2019 The repeated evolution of dental apicobasal ridges in aquatic-feeding mammals and reptiles. *Biol. J. Linn. Soc.* **127**, 245–259. (doi:10.1093/biolinnean/blz025)
35. Foffa D, Young MT, Brusatte SL. 2018 Filling the Corallian gap: new information on Late Jurassic marine reptile faunas from England. *Acta Palaeontol. Pol.* **63**, 287–313. (doi:10.4202/app.00455.2018)
36. Foffa D, Young MT, Brusatte SL. 2015 Evidence of macrophagous teleosaurid crocodylomorphs in the Corallian Group (Oxfordian, Late Jurassic) of the UK. *PeerJ* **3**, e1497. (doi:10.7717/peerj.1497)
37. Weryński Ł, Błazejowski B. 2023 Late Jurassic teeth of plesiosauroid origin from the Owadów-Brzezinki Lagerstätte, Central Poland. *PeerJ* **11**, e15628. (doi:10.7717/peerj.15628)
38. Maxwell EE, Caldwell MW, Lamoureux DO. 2012 Tooth histology, attachment, and replacement in the Ichthyopterygia reviewed in an evolutionary context. *Palaeontol. Z.* **86**, 1–14. (doi:10.1007/s12542-011-0115-z)
39. Weldon PJ, Ferguson MWJ. 1993 Chemoreception in crocodylians: anatomy, natural history, and empirical results. *Brain Behav. Evol.* **41**, 239–245. (doi:10.1159/000113845)
40. Platt SG, Rainwater TR, Snider S, Garel A, Anderson TA, McMurtry ST. 2007 Consumption of large mammals by *Crocodylus moreletii*: field observations of necrophagy and interspecific kleptoparasitism. *Southwest. Nat.* **52**, 310–317. (doi:10.1894/0038-4909(2007)52[310:COLMBC]2.0.CO;2)
41. Meyer CA. 1991 Burial experiments with marine turtle carcasses and their paleoecological significance. *PALAIOS* **6**, 89. (doi:10.2307/3514956)
42. Morphosource. *The CT data set and the 3D surface model*. Morphosource repository. See <https://www.morphosource.org/catalog/media?f.project=000593818>