

Microraptor: A Systematic View of Its Three Species: Cranial Elements

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Abstract

The genus *Microraptor* is classified within the Dromaeosauridae. It was first found in the Early Cretaceous Jiufotang Formation in Liaoning, China. It has played a central role in the debate over the supposed evolutionary relationship between dinosaurs and birds and the supposed “feathered” dinosaurs. This genus has been considered a feathered dinosaur by most secular scientists and some creation scientists. *Microraptor* has been used to support the hypothesis that birds evolved from dinosaurs and that dinosaurs had feathers. This paper examines the anatomical skull features of the first three described *Microraptor* species (*Microraptor zhaoianus*, *Microraptor gui*, and *Microraptor hanqingi*) to understand whether they align more closely with bird or dinosaur characteristics. This analysis is performed through a literature review of the first original publications of the first three species of *Microraptor*. The original descriptions of the bone structures in the publications were compared to determine whether they follow a dinosaurian or bird-like pattern. That was done by comparing the secular literature on extinct birds and dinosaurs.

The findings indicate that the *Microraptor*'s skull exhibits numerous characteristics that are also shared with extinct birds. This study raises questions about the interpretation of *Microraptor* as a feathered dinosaur and instead supports its classification as an extinct bird. The result of this analysis can contribute to the ongoing debate about whether those anatomical structures support a dinosaurian or a bird classification.

Keywords: *Microraptor*, Dromaeosauridae, feathered dinosaurs, dinosaur-bird relationship, cranial morphology, avian characteristics, extinct birds

Introduction

Microraptor is a recognized and classified genus of Dromaeosauridae from the Early Cretaceous of the Jiufotang Formation in Liaoning, Northeastern China. This formation records many specimens of birds and supposedly feathered dinosaurs. This formation is above the Yixian Formation. Both are part of the Jehol Group, which gained fame in the 1990s with the discovery of *Sinosauropteryx*, the first supposedly feathered dinosaurs.

Microraptor has been accepted as one of the few dinosaur genera identified as having feathers by secular and some creation scientists (Wood and Garner n.d.). *Microraptor* has played a central role in the ongoing debate about feathered dinosaurs. However, many secular researchers have supported the possibility that the supposedly feathered dinosaurs are birds (Bechly 2024 a, b).

Here, I analyze the morphological features of the first three species of *Microraptor* that have been identified, described, and published. Since hundreds of specimens represent *Microraptor* (Alexander et al. 2010), it is impractical to analyze each one of the specimens individually. This analysis focuses on the cranial elements to examine whether they align with a bird or dinosaur structure based on current secular literature.

For now, the scientific consensus is that this genus has three species: *Microraptor zhaoianus* gen. et sp. nov. (Xu, Zhou, and Wang 2000), *Microraptor gui* sp. nov. (Xu et al. 2003), and *Microraptor hanqingi* (Gong et al. 2012). The synonymy (when two or more different scientific names are used for the same fossil species or genus) proposed by Turner, Makovicky, and Norell (2012) and Senter et al. (2004) for this genus is not discussed here.

Based on the anatomical analyses of the skulls of these three species, the view of *Microraptor* as a feathered dinosaur is contested. The analyzed skull structure was listed, mentioned, identified, and described in the first three original publications of *Microraptor* specimens (Gong et al. 2012; Xu et al. 2003; Xu, Zhou, and Wang 2000).

Understanding *Microraptor* as a bird can help support the idea that other specimens with similar features can be classified as birds instead of feathered dinosaurs. That is why evaluating whether the features listed, mentioned, identified, and described for this genus align more with those of birds or dinosaurs is necessary.

The Three Species of *Microraptor*—General View

Microraptor zhaoianus was discovered with a partially articulated skeleton, lacking the middle

portion of the body. One specimen (V 12330) was used for its description (Xu, Zhou, and Wang 2000).

Most of the six specimens found of *Microraptor gui* were articulated well, and the holotype is an almost complete skeleton. The holotype (IVPP V13352) and a referred specimen were used for its description (Xu et al. 2003).

M. hanqingi is a nearly complete specimen, showing a mostly articulated postcranial skeleton with a disarticulated pectoral girdle and left forearm. One specimen (LVH 0026) was used for its description (Gong et al. 2012).

Nothing is said in the publication about the quality of preservation of the skeletal material for *M. zhaoianus* and *M. gui*. Regarding *M. hanqingi*, it is mentioned that one of the sides of the skeleton exhibits, “well-preserved skeletal elements and details not evident on the original surface” (Gong et al. 2012).

The first specimens analyzed for the publication of *M. zhaoianus* were identified as mature (Xu, Zhou, and Wang 2000). There is no mention of the degree of maturity of *M. gui* and *M. hanqingi* (Gong et al. 2012; Xu et al. 2003).

Skull Bones

In the secular scientific field, information in the skull has been necessary to understanding dinosaur-bird evolution. According to Bhullar et al. (2012), “The unique bird skull houses two highly specialized systems: the sophisticated visual and neuromuscular coordination system allows flight coordination and exploitation of diverse visual landscapes, and the astonishing variations of the beak enable a wide range of avian lifestyles.” Similarly, Felice et al. (2020) highlight that some defining avian traits are localized specifically in the skull, “key ‘avian’ features being localised to the skull (e.g., edentulous beak, kinetic palate, encephalised brain).”

These morphological variations in birds’ skulls emphasized by Bhullar et al. (2012) and Felice et al. (2020), are understood in the secular worldview as results of evolutionary processes. Recognizing that secular scientists interpret anatomical structures through the lens of evolutionary ideas is essential when analyzing any morphological feature in this context. Neutrality does not exist, and the evolutionary worldview has been applied to interpreting traits to tell an evolutionary story of those traits, not accounting for shared design and functionality.

The Three Species of *Microraptor* and Their Skulls

Not many of the skull features of *Microraptor zhaoianus* are described and discussed in the original publication (Xu, Zhou, and Wang 2000) of this

species. The information mentioned is that there are similarities between specific features of the premaxilla and those of *Archaeopteryx* and *Sinornithosaurus*. Additionally, certain features of the maxilla are like those of *Archaeopteryx* and troodontids (one of the groups that belong to Maniraptora).

On *M. gui* not much was said in its original publication (Xu et al. 2003) about skull morphology. The morphological information described for the skull included an identifiable triradiate postorbital, one of the bones that contribute to the orbital structure.

The skull of *M. hanqingi* is the most fully described. The cranial elements are the skull and mandible. They were found to be closely associated. The skull varies in preservation, and many cranial bones can be identified. They are mostly disarticulated. Surface ornamentation is not present. The skull has a length of 92.5 mm, but this measurement is based on the length of the mandible. The identified cranial bones are “right and left maxilla, the remnants of both premaxillae, nasals, lacrimal, squamosal, dentaries, posterior mandibular elements including the articular, surangular, prearticular, angular, and splenial” (Gong et al. 2012).

Premaxilla

A premaxilla is mentioned in the original publications of *M. zhaoianus* as, “The premaxilla is similar to that of *Archaeopteryx* and *Sinornithosaurus* in that it has a sloping anterior margin. As in *Archaeopteryx* and troodontids, the maxilla contributes to the border of the external naris” (Xu, Zhou, and Wang 2000). And for *M. hanqingi* as, “Preservation of the skull varies, and the cranial bones that can be identified include the right and left maxilla, remnants of both premaxillae...” (Gong et al. 2012)

M. zhaoianus has a sloping anterior margin. This feature in *M. hanqingi* is indicated based on the outline of the maxilla. Xu and Wu (2001) cited the premaxilla’s sloping anterior margin as a bird-like feature. No information on this structure is noted for *M. gui* in its original publication.

Maxilla

Not much is mentioned about the maxilla of *M. zhaoianus*. There is only a comparison of its contribution to the external border of the naris as in *Archaeopteryx* and troodontids (Xu, Zhou, and Wang 2000). Nothing is said about the maxilla of *M. gui*.

M. hanqingi has a maxilla that is approximately 43.5 mm long and a height projection of 20.6 mm (Gong et al. 2012). Due to its outline, the skull exhibits a trait like that of *Archaeopteryx* and *Sinornithosaurus*. *M. hanqingi*’s maxilla is preserved with four teeth in the original position. However, there are 14 alveoli in

it. The maxillary presents teeth that are longer and more recurved compared to the dentary teeth (Gong et al. 2012).

Dentary

The dentary of *M. hanqingi* presents an arrangement of alveoli that are close together (Gong et al. 2012). It accommodates as many as 16 teeth, possibly. Therefore, the number of teeth for *M. hanqingi* is unclear. The dentary has short, broad teeth compared to the maxillary teeth. They also present a uniform size.

M. zhaoianus shows a dentary with at least 15 dentary teeth (Xu, Zhou, and Wang 2000). Its dentary teeth are packed as in troodontids. Troodontids are recognized as a bird group (Feduccia 2020). No information about the dentary of *M. gui* is mentioned.

Teeth

Xu, Zhou, and Wang (2000) write that at least 15 teeth are present, arranged like those in troodontids. There are no anterior serrations on the teeth. However, there are serrations on the posterior teeth. Basal constriction is found on the posterior teeth, and the crown is less compressed. The constriction is between the root and the crown. The authors mention those two features as bird-like and write that *M. zhaoianus* presents, “the more bird-like teeth” (Xu, Zhou, and Wang 2000).

No teeth are mentioned for *M. gui*. For *M. hanqingi*, the number of teeth cannot be certain. However, the authors write that it may be 16. Four teeth are preserved in position. There are minute serrations in one tooth (Gong et al. 2012). The authors of the original publication of *M. hanqingi* (Gong et al. 2012) mention that the serration pattern varies in the skulls known of *Microraptor*. They also write that serration is ambiguous in their specimen (Gong et al. 2012).

The presence of teeth is not problematic for *Microraptor*. Teeth have been noted as a dinosaurian feature, as modern birds do not possess teeth. However, some extinct birds have teeth like *Hesperornis* and *Ichthyornis* (Martin and Stewart 1977), *Yanornis* and *Yixianornis* (Zhou and Zhang 2001), and *Sapeornis* (Wang et al. 2017).

Regarding serration, as it is now known, no existing literature mentions serrated teeth in extinct birds. Also, heterodonty (different types of teeth) is not yet known for extinct birds.

Postorbital

No postorbital information is mentioned for *M. zhaoianus*. A triradiate postorbital was identified in *M. gui* and *M. hanqingi*. For *M. gui*, “Little can be said about the cranial morphology but a tri-radiate

postorbital is identifiable” (Xu et al. 2003). And for *M. hanqingi*, “The holotype of *M. gui* has a skull, but the anterior portions are dubious and the posterior cranial elements are too badly crushed to reconstruct the cranium although it confirms the presence of a tri-radiate postorbital in this genus. The postorbital of *M. hanqingi* is similar” (Gong et al. 2012).

This feature is not found in modern birds (Bhullar et al. 2016) but is present in many dinosaurs (Nesbitt 2011). The presence of a postorbital was also confirmed in a *Microraptor* specimen found in China in 2014 (Pei et al. 2014). However, although extant birds do not possess a postorbital, they have it as an embryo. This bone fuses with the frontal before hatching. This fusion occurs during embryonic development in birds (Smith-Paredes et al. 2018). Thus, the presence of an unfused postorbital may not be a diagnostic feature for dinosaurs. In fact, recent research shows that fossil forms of several taxa of birds and dinosaurs matured much more slowly, having had longer gestation times than their modern counterparts. For example, Yang and Sander (2018) argued that the reason modern birds no longer develop teeth is because they spend less time in development than their fossil forebears. Though, as yet speculative, longer development and slower maturation times might have led to a longer time for the postorbital to fuse.

That is, it is possible that the slower development observed in some fossil taxa allowed the postorbital bone to remain unfused for a longer duration, potentially persisting into post-hatching stages. Raising the possibility that an unfused postorbital may reflect differences in developmental timing rather than a fundamental taxonomic distinction.

Smith-Paredes et al. (2018; Supplementary Figure 9) also state regarding the postorbital:

A postorbital has been found in a few enantiornithes [sic] preserving the temporal region. In many others, the area is crushed beyond recognition, which prevents identification of the element, but was probably present as inferred by phylogenetic bracketing. However, in some members of Enantiornithes, the postorbital is reduced (see *Rapaxavis* in O'Connor and Chiappe 2011), which suggests some taxa could have lost it in parallel to Euornithes. In the lineage leading to modern birds, an independent postorbital was lost at some point between the origin of Euornithes and the origin of Ornithurae; this cannot be established more precisely because preservation of basal Euornithes does not allow recognition of the presence or absence of a postorbital.

Although an evolutionary worldview is being applied to interpret the data, the citation above demonstrates that a postorbital is found in extinct birds. Enantiornithes are extinct birds with teeth and various features, including a postorbital. Modern

birds do not exhibit many of the characteristics of Enantiornithes. The postorbital seems to be lost in the group (Euornithes) that includes the modern birds (Neornithes). Supporting these observations, Wellnhofer (2009) mentions the presence of a postorbital in an enantiornithine bird from the Lower Cretaceous in Spain. He also mentioned *Confuciusornis* from the Lower Cretaceous of China and wrote that this bird had a distinct postorbital.

O'Connor and Chiappe (2011) state that, when discussing the postorbital bone, “This element—absent among modern birds—is retained within Enantiornithes but is only definitively preserved in three specimens: LP 4450, *Pengornis* and *Shenqiornis*. In LP 4450 and *Shenqiornis*, it exhibits a typical T-shaped design, but in *Pengornis*, the caudodorsal (squamosal) ramus is either absent or broken. The postorbital of these specimens shows considerable variation in size and morphology.” A postorbital is also present in *Navaornis hestiae*, found in the Cretaceous of Brazil. This new genus and species was identified as Enantiornithes (Chiappe et al. 2024).

Additional evidence of a postorbital is present in extinct birds (fig. 1). A complete, well-preserved postorbital in *Jeholornis prima*, a bird from the Early Cretaceous of Jehol, China, exhibits a triradiate configuration (Hu et al. 2022). Also, *Sapeornis* is a bird from the Early Cretaceous in China. According to its

description, it features a Y-shaped postorbital (Zhou and Zhang 2003). Furthermore, *Yuanchuavis*, a bird from the Early Cretaceous of China, also presents a triradiate postorbital (Wang et al. 2022).

It is also important to note that the presence or absence of a postorbital process might not indicate reptilian status for the reasons listed above. Postorbital bones are not always present in reptiles and thus should not be taken as a reptilian feature for *Microraptor*. The Amphisbaenia tribe, which includes burrowing worm-like or snake-like lizards, does not present a postorbital bone (Williston and Gregory 1925). It is also excluded from the extinct reptiles *Araucoscelis* and *Hyperodapedon* (Williston and Gregory 1925). Additionally, the postorbital bar is sometimes incomplete in lizards, snakes, and therapsids (Williston and Gregory 1925).

Moreover, Rauhut (2013) discusses the presence of a triradiate postorbital configuration in a new reconstruction of the *Archaeopteryx* skull. Although the article does not mention the word “triradiate,” the skull reconstruction in Fig. 5 of the publication reveals a triradiate structure for the postorbital. He uses this new reconstruction to support a more theropod-like skull for *Archaeopteryx*. The author also cites that a bird, *Confuciusornis*, has a postorbital structure. Even though he does not mention whether it is a triradiate structure. Hu et al. (2020) cite the

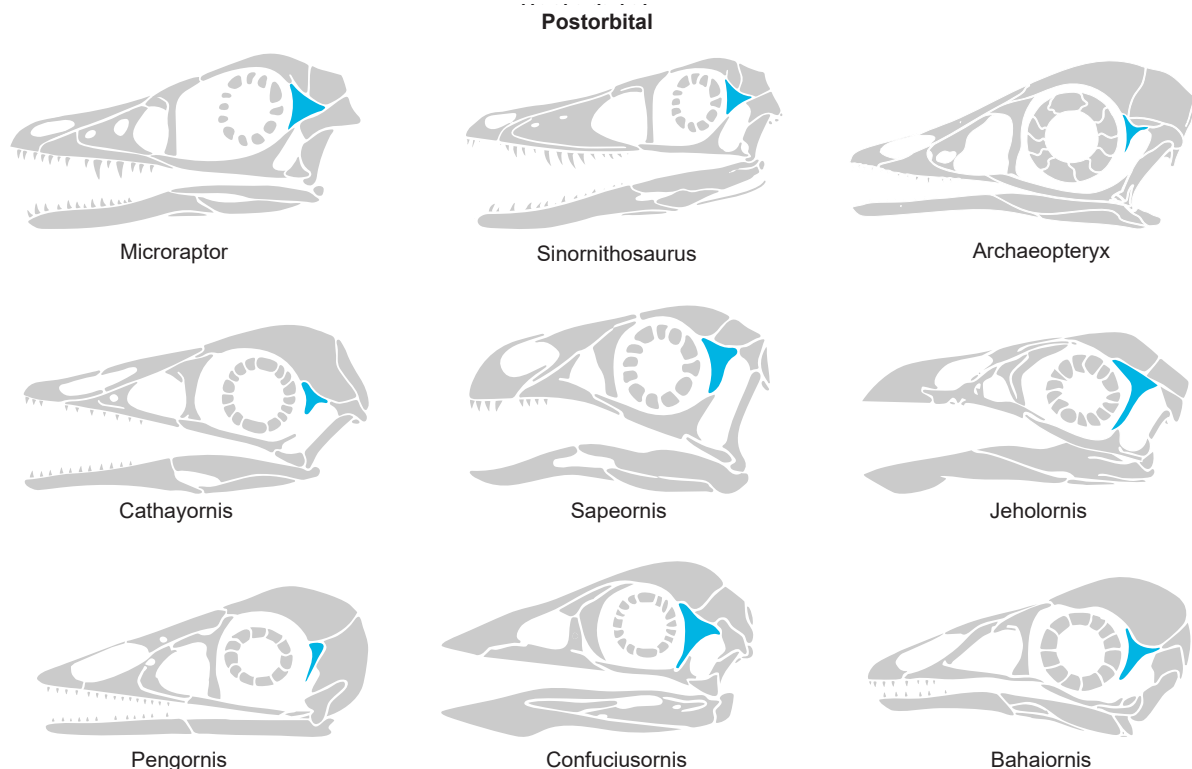


Fig. 1. *Microraptor* and *Sinornithosaurus* adapted from Hartman (n.d.) and Paul (2016) reconstructions. *Hesperornis* adapted from Everhart (2011). *Archaeopteryx* skull adapted from Wellnhofer (2009). *Cathayornis*, *Confuciusornis*, and *Sapeornis* adapted from Hartman (n.d.) reconstructions and Wang et al. (2017). *Jeholornis* adapted from Hu et al. (2023). *Pengornis* and *Bahaornis* adapted from Zelenkov (2017). Artwork by Joel Leineweber, used by permission.

well-developed postorbital in *Confuciusornis*. He also mentions that some Enantiornithes preserve some of the features found in *Archaeopteryx*. *Archaeopteryx*'s status as a bird has been disputed based less on anatomy and more on evolutionary bias, as Haynes (2022, 2023) has demonstrated.

The postorbital configuration is related to the kinesis (movement/mobility) of the skull of birds and dinosaurs (Wang and Hu 2017). Wang et al. (2022) cite Bock (1964) and Zusi (1984) and say that “avian cranial kinesis has been demonstrated to have improved feeding performance by increasing biting force, jaw closing speed, and food handling precision.”

Finally, Rahut (2013) notes that the possibility of the absence of a postorbital bar was, “one of the key arguments that led Bühler (1985) to propose a bird-like cranial kinesis in *Archaeopteryx*.” However, Hu et al. (2020) mention that “the skull of *Sapeornis* [an extinct bird] was largely akinetic, as in non-avian dinosaurs and palaeognaths (Hu et al. 2019).” Then, on the postorbital, it seems reasonable to conclude that the presence of a closed postorbital and an akinetic (immobile) skull can be features of a bird. Fig. 1 shows the list of extinct birds that present a postorbital bone.

Despite the evolutionary perspectives, extensive paleontological and anatomical data demonstrate that postorbital bones existed in multiple extinct bird and reptile taxa. Their presence or absence is subject to considerable developmental, functional, and biomechanical variability, cautioning against oversimplified interpretations of their taxonomic significance.

Lacrimal

The other feature of the cranial element is the lacrimal bone that is part of the orbit. This bone is found in one species (*M. hanqingi*). The feature of this bone for this species is described as, “The lacrimal is T-shaped and about 19.5 mm tall. The anterior ramus is 10.7 mm long; the posterior ramus is much shorter” (Gong et al. 2012).

However, extinct birds, such as those in the Enantiornithes group, exhibit the same T-shaped bone. Wellnhofer (2009) also mentions the lacrimal bone in *Archaeopteryx*, which has a T-bone shape. He explains that the mobility this bone configuration allows is in line with the prokinetic condition (independent motion of the upper beak relative to the rest of the skull) of the skull, as described by Bühler (1985). That is a condition present in most modern birds. However, Elzanowski (2002) mentioned that Wellnhofer (2009) suggested rhynchokinesis (ability to flex the upper part of the beak independently from the rest of the skull), but due to the *Archaeopteryx*'s upper jaw, its mobility might differ from the modern

cranial kinesis categories. A T-shaped bone is also identified by Chiappe et al. (2024) as a lacrimal bone in Enantiornithes, specifically *Navaornis hestiae*.

Clarke (2009), analyzing two birds (*Apatornis* and *Ichthyornis*), discusses the presence of lacrimal and states, “In Aves, the lacrimal can articulate with (or coossify to) both the nasal and frontal, but may articulate exclusively with either of these two bones (Cracraft 1968). In *Struthio camelus* and *Rhea americana*, for example, the lacrimal usually contacts only the nasal (Cracraft 1968), unlike the condition in *Ichthyornis* dispar, while in tinamous, *Apteryx*, and *Gallus gallus*, it contacts the frontal and nasal (Cracraft 1968). In Anhimidae, it also articulates exclusively with the nasal (Cracraft 1968). *Hesperornis regalis* has the lacrimal primarily contacting the nasal, although also contacting the anterolateral portion of the frontal (Bühler, Martin, and Witmer 1988).” As shown above, a T-shape is necessary to articulate or touch the two bones (nasal and frontal), rather than just articulating with the jugal in a bar shape.

O'Connor and Chiappe (2011) state on a T-shaped lacrimal in two birds: “The morphology of this bone is definitively clear in only two specimens: *Pengornis* (IVPP V15336) and LP 4450.” *Yuanchuavis* also presents a T-shaped lacrimal, as shown in Wang et al. (2022). *Sapeornis* is also a bird that presents a T-shaped lacrimal bone, as described by Hu et al. (2020). The authors also mention that the *Sapeornis*' lacrimal is like, “*Archaeopteryx* and some enantiornithines like *Pengornis*, *Parapengorn* and *Pterygornis* in overall shape.” The same happens to *Cratonavis*, a bird from the Early Cretaceous of China that presents a T-shaped lacrimal (Li et al. 2023).

A new species of fossil bird, belonging to the Anseriformes, was discovered in the Late Cretaceous system of Antarctica. *Vegavis iaii* was identified and described based on a nearly complete skull. This species also presents a T-shaped lacrimal. Little is known about this bone, except that it remains unfused. However, the figure on page 148 of the article shows a lacrimal with a T shape. (Torres et al. 2025). As shown in fig. 2, several extinct birds share the same features of the lacrimal found in the *Microraptor*. Fig. 2 shows the extinct birds that present a lacrimal bone.

Nasals

The characteristics of the nasal bones are cited in one species, *M. hanqingi*, “The nasals are 35 mm long and are thin, paired elements that widen anteriorly and posteriorly. They are widest (9.2 mm) across the anterior portion of the bone just before the concave border of the nares” (Gong et al. 2012). This aligns with a lightweight cranial structure. In contrast, in dinosaurs, nasal bones were often involved in supporting strong jaw muscles or forming crests for

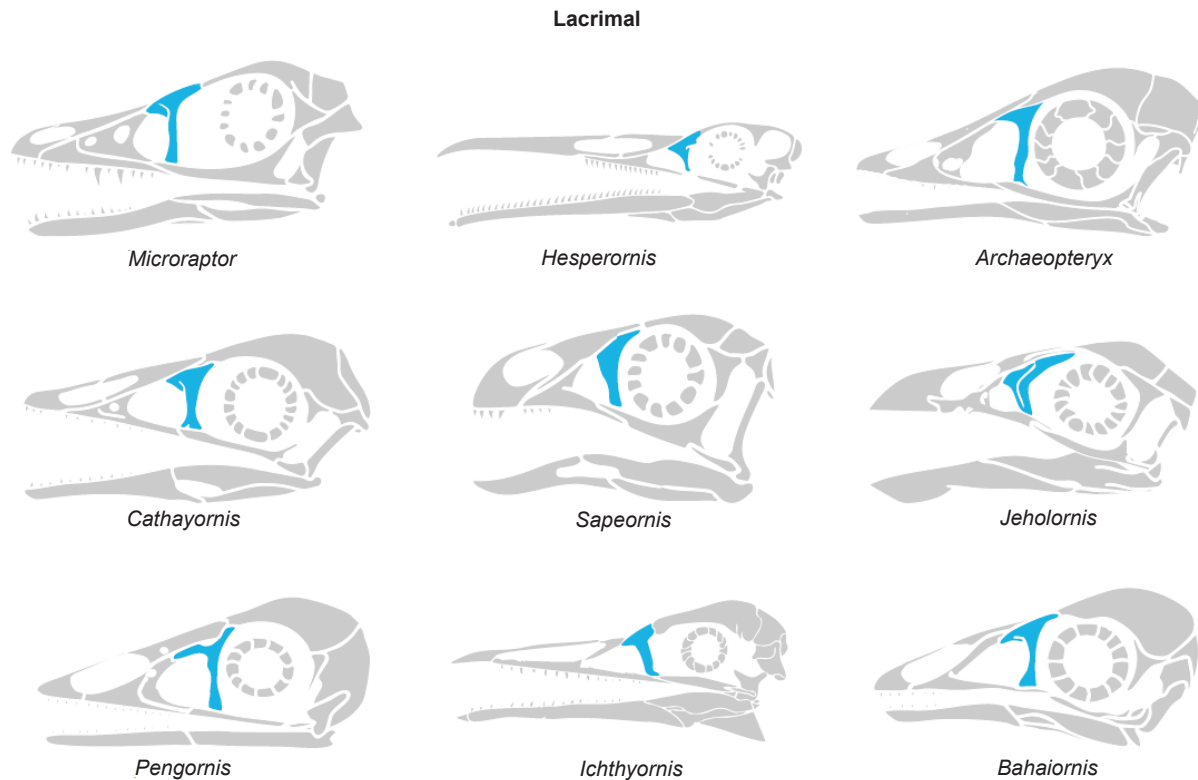


Fig. 2. *Microraptor* and *Sinornithosaurus* adapted from Hartman (n.d.) and Paul (2016) reconstructions. *Hesperornis* adapted from Everhart (2011). *Archaeopteryx* skull adapted from Wellnhofer (2009). *Cathayornis*, *Confuciusornis*, and *Sapeornis* adapted from Hartman (n.d.) reconstructions and Wang et al. (2017). *Jeholornis* adapted from Hu et. al. (2023). *Pengornis* and *Bahaionis* adapted from Zelenkov (2017). Artwork by Joel Leineweber, used by permission.

display or combat. The characteristics cited for the nasal bones of *M. hangingi* point to a bird-like feature. The nasal bones do not conform to a dinosaurian structure for this genus.

Quadrate

A quadrate (a bone that helps to connect the lower jaw to the rest of the skull) is mentioned for *M. zhaoianus* (Xu, Zhou, and Wang 2000), but it is not described in any of the specimens of those three species (*M. zhaoianus*, *M. gui*, and *M. hanqingi*). Wellnhofer (2009) mentions the single-headed feature as theropod-like and the double-headed feature as bird-like. However, a typical enantiornithine skull presents a single-headed quadrate (O'Connor and Chiappe 2011). Showing that a quadrate with that feature should not point to a reptilian-like skull.

It is also important to note that the word “theropod” has changed. When its definition was created, it only referred to dinosaurs. After 1985/1986, due to evolutionary influences, the term “theropod” began to include both dinosaurs and birds (Haynes 2022, 2023).

To summarize the anatomical information discussed in the previous sections and present a more straightforward comparison among the three described species of *Microraptor* (*M. zhaoianus*, *M. gui*, and *M. hanqingi*). Table 1 shows structured

information of their cranial features. This comparative overview outlines the elements mentioned or described in each of the first three original publications of the species and contextualizes the significance of their preserved bones based on current paleontological literature. The table highlights patterns that may challenge the prevailing interpretations regarding *Microraptor*.

Conclusion

The anatomical analysis of *Microraptor* presented in this study questions the prevailing view that it was a feathered dinosaur and instead supports its classification as an extinct bird. The assumption that *Microraptor* must be a dinosaur based on certain skull features is weakened by facts showing that these traits also exist in extinct birds and even in some modern bird embryos. All the features examined on the skull of the first three original publications of *Microraptor* species show similarities with those of known extinct birds. The presence of multiple traits in *Microraptor* that are also found in birds suggests that *Microraptor* is a bird, not a dinosaur with feathers. Although the skull features might not be considered conclusive (since the features analyzed might not be diagnostic) for the status of *Microraptor* as a bird, the results in this paper seem to weigh in

Table 1. Cranial morphological features analyzed of the three described species of *Microraptor*.

Feature	<i>M. zhaoianus</i>	<i>M. gui</i>	<i>M. hanqingi</i>	Notes / Interpretation
Premaxilla	Present; sloping anterior margin (like <i>Archaeopteryx</i> , <i>Sinornithosaurus</i>)	Not mentioned	Remnants preserved	Sloping premaxilla considered bird-like
Maxilla	Contributes to external naris; similarity to <i>Archaeopteryx</i> and troodontids	Not mentioned	Present; 43.5 mm; 14 alveoli; 4 teeth preserved	Longer and more recurved teeth in the maxilla than the dentary
Dentary	At least 15 teeth; packed like troodontids	Not mentioned	~16 alveoli; teeth short, broad, uniformly sized	Tooth shape and packing varies; posterior serrations ambiguous
Tooth Serration	No anterior serrations; posterior serrations present; basal constriction	Not mentioned	One tooth with minute serrations; pattern ambiguous	Tooth serration has not been described in extinct birds
Postorbital	Not mentioned	Triradiate postorbital	Triradiate postorbital	Present in extinct birds like <i>Jeholornis</i> , <i>Confuciusornis</i>
Lacrima	Not mentioned	Not mentioned	T-shaped; 19.5 mm tall	T-shapes also found in <i>Archaeopteryx</i> , <i>Sapeornis</i> , enantiornithines
Nasal Bones	Mentioned but not described	Not mentioned	Thin, paired, widen anteriorly and posteriorly	Gracile form aligns with avian-like cranial lightness
Quadrate	Mentioned but not described	Not mentioned	Not mentioned	Avian/dinosaurian classification is inconclusive without it

favor of it. This conclusion is particularly relevant considering the ongoing debate about the nature of “feathered dinosaurs.” Understanding *Microraptor* as an extinct bird, rather than a dinosaur with feathers, may help us recognize many other supposed feathered dinosaurs as just extinct birds.

Future research is planned to include postcranial characteristics. This study contributes to a growing body of information suggesting that our understanding of Mesozoic birds might be blurred with evolutionary influence. Moreover, many of the so-called feathered dinosaurs are, in fact, birds.

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