Shouldering the challenge of deciphering avian palate evolution

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Abstract

Wilken et al. (1) investigate the evolution of avian palatal kinesis using comparative morphology and
biomechanical modelling. While the study's topic and approach are timely, its conclusions are marred
by inadequate taxon sampling and morphological misinterpretations.

Main text

Reinterpreting the *Janavis* pterygoid (2) as a juvenile bird coracoid (1) is contradicted by clear
morphological and developmental evidence (Fig. 1). The concave, ossified 'sternal articulation' of the
would-be coracoid (1) is unossified in juvenile bird coracoids, yet resembles the quadrate articulation
of the pterygoid in several extant taxa (Fig. 1B). Similarly, *contra* (1), the basipterygoid articulation in
Janavis closely resembles that of several extant neognathous birds in its form and position (Fig. 1C).

The authors identify a supposed glenoid and acrocoracoid on the purported coracoid (1), yet in
developing bird coracoids these structures only arise after the scapular cotyle, which is absent in the
fossil. Importantly, the positions of these structures do not shift relative to one-another throughout
extant bird ontogeny, which would be necessary for the element to come to resemble a coracoid (Fig.
1B). Since ichthyornithine postcranial morphology and growth generally fall within the range of
variation of extant birds (3), there is no reason to think these would be any different in *Janavis*.
Crucially, the authors of (1) must have reduced the size of the *Janavis* scapula by ~50% to 'match' it
with the putative coracoid (see their Supplementary Fig. 14A; our Fig. 1A), yet this is unmentioned.

The above issues notwithstanding, the ichthyornithine hemipterygoid alone—with an articular condyle
like those of prokinetic neognaths —suggests a mobile intrapterygoid joint (Fig. 2A). Wilken et al.
seemingly agree ("…*Ichthyornis...* possesses a segmented, potentially propulsive pterygoid..." (1));
yet, they interpret palatal kinesis as a neognath autapomorphy by asserting that stem bird
musculature was incompatible with palate mobility. This rests upon misinterpretations of the functional
correlates of quadrate morphology. First, elongate orbital processes in *Ichthyornis* and *Hesperornis*
are declared to impose constraints on kinesis due to articulation with the neurocranium (1). However,
many neognaths exhibit equivalently long (or longer) orbital processes closely approaching the
neurocranium without precluding prokinesis (Fig. 2B; (5); https://doi.org/10.5281/zenodo.15619496)
Second, *contra* (1), the bicondylar nature of the *Ichthyornis* quadrate would not have precluded
coupled palatal kinesis, as evidenced by bicondylar Anseriformes (Fig. 2B) (6). Further morpho-
functional misconceptions underpin problematic cranial linkage models in (1) (Fig. 2C), yet the
authors' own ancestral reconstructions of cranial linkages and protractor force seemingly support
palatal mobility as plesiomorphic for all crown birds (their Figs. 3A, 4), rather than being apomorphic
for Neognathae as they argue.

Contemporary research in avian macroevolution seeks to evaluate associations between brain and
skull transformations (7-9); however, this link is weakly substantiated by (1), as several taxa with

palatal kinesis (e.g., Anseriformes) share plesiomorphically low degrees of encephalization with
akinetic palaeognaths and stem birds (10).

Conclusions

The origin of palatal kinesis was a key milestone in avian evolutionary history—careful morphological
interpretation will be crucial to clarify how this major transformation occurred and influenced avian
evolutionary history.

Figures

Fig. 1. The pterygoid of *Janavis finalidens* is not a juvenile coracoid. A) The scapula of *Janavis*
was downscaled to 50% of its real size by (1) (see their Fig. S14). When correctly scaled (Fig. 1a and
Extended Data Fig 2b-d in (2)) the pterygoid and scapula do not match (nor does the pterygoid
resemble a coracoid in the first place—see main text). **B)** If interpreted as an immature coracoid, the
pterygoid of *Janavis* differs from all known avian coracoids in: 1) glenoid development ontogenetically
preceding the scapular cotyle, procoracoid, and acrocoracoid; 2) exhibiting a 'dorsally' convex and
'laterally' flat shaft; 3) the authors' (1) 'sternal process' (presumably the sternolateral process) being
medially, rather than laterally, oriented; 4) exhibiting fully ossified concave surfaces on both omal and
sternal ends, which are instead open and hollow or formed by unfinished bone in immature bird
coracoids (the sternal end of the coracoid is only concave in adult, highly derived Apodiformes); 5)
being highly pneumatic with a simple trabecular structure, as pneumaticity only develops late in post-
hatching development in extant birds. **C)** If correctly interpreted as a right pterygoid (*contra* (2), where
it was considered a left pterygoid), all features of the element are found in fossil and extant
neognathous bird pterygoids, including: 1) a large, medially-oriented basipterygoid process facet
reaching the rostral end (present in Galliformes, *contra* (1)); 2) a laterally and often dorsally open,
cup-shaped palatine contact (present across Galloanserae); 3) a tubular shaft that is medially flat and
dorsally convex (present in Galliformes and Pelagornithidae); 4) a simple, concave quadrate articular
surface (present in Anseriformes); 4) a marked orbito-pterygoid contact process (present in
Galliformes and some Anseriformes); 5) a heavily pneumatised internal structure with simple
trabeculae (present across Galloanserae). Bold letters indicate anatomical views: caudal (C), dorsal
(D), lateral (L), medial (M), rostral (R). Cross-sectional (CS) planes are indicated with dotted lines over
each element. All scale bars equal 5 mm.

Fig. 2. Morphology of the suspensorial system in Ichthyornithes suggests prokinetic
capabilities. A) *Ichthyornis* exhibited a mobile intrapterygoid (hemipterygoid-pterygoid) articulation,
like the condition in unsegmented immature (imm.) neognaths (hemipterygoid highlighted in teal
shade). The condylar morphology of the pterygoid articular contact resembles that of prokinetic
Galliformes and Neoaves, and contrasts with the long or wide overlapping contacts of palaeognaths.
Long rostral processes of the palatine are obscured for clarity. **B)** The quadrates of *Ichthyornis* and
Hesperornis (bicondylar, with long orbital processes interpreted as contacting the braincase (1))
would not preclude palatal kinesis. Both features are present across a wide range of prokinetic
neognath taxa unsampled by (1). This includes taxa where the orbital process 1) wraps around and
closely approaches the neurocranium; 2) terminates in an expansion whose lateral surface anchors
the M. pseudotemporalis profundus; and 3) is presumably associated with depressions or 'cotylae' on
the braincase (e.g., *Ardea*; see https://doi.org/10.5281/zenodo.15619496). **C)** The ichthyornithine
linkage model proposed by (1) enforces immobility at contacts between bones (e.g., the jugal and the
lacrimal) which do not preclude prokinesis in extant birds. We present corrected linkage values
following the scheme of (1), indicated in green, and illustrate the support for these from previous
reconstructions of the skull of *Ichthyornis*. Bold letters indicate anatomical views: caudal (C), dorsal

(D), lateral (L), ventral (V). All scale bars equal 2.5 mm. Digitally segmented *Struthio camelus* palatine in A courtesy of Annabel K. Hunt.

References

1. A. T. Wilken *et al.*, Avian cranial kinesis is the result of increased encephalization during the origin of birds. *Proceedings of the National Academy of Sciences* **122**, e2411138122 (2025).

2. J. Benito, P.-C. Kuo, K. E. Widrig, J. W. M. Jagt, D. J. Field, Cretaceous ornithurine supports a neognathous crown bird ancestor. *Nature* **612**, 100-105 (2022).

3. J. Benito *et al.*, 40 new specimens of *Ichthyornis* provide unprecedented insight into the postcranial morphology of crownward stem group birds. *PeerJ* **10**, e13919 (2022).

4. C. R. Torres, M. A. Norell, J. A. Clarke, Bird neurocranial and body mass evolution across the end-Cretaceous mass extinction: The avian brain shape left other dinosaurs behind. *Science Advances* **7**, eabg7099 (2021).

5. P.-C. Kuo, G. Navalón, R. B. J. Benson, D. J. Field, Macroevolutionary drivers of morphological disparity in the avian quadrate. *Proceedings of the Royal Society B: Biological Sciences* **291**, 20232250 (2024).

6. P.-C. Kuo, R. B. J. Benson, D. J. Field, The influence of fossils in macroevolutionary analyses of 3D geometric morphometric data: A case study of galloanseran quadrates. *Journal of Morphology* **284**, e21594 (2023).

7. B.-A. S. Bhullar *et al.*, How to Make a Bird Skull: Major Transitions in the Evolution of the Avian Cranium, Paedomorphosis, and the Beak as a Surrogate Hand. *Integrative and Comparative Biology* 10.1093/icb/icw069 (2016).

8. A. M. Balanoff, G. S. Bever, T. B. Rowe, M. A. Norell, Evolutionary origins of the avian brain. *Nature* **501**, 93-96 (2013).

9. L. M. Chiappe, G. Navalon *et al.*, Cretaceous bird from Brazil informs the evolution of the avian skull and brain. *Nature* **635**, 376-381 (2024).

10. D. T. Ksepka *et al.*, Tempo and Pattern of Avian Brain Size Evolution. *Current Biology* **30**, 2026-2036.e2023 (2020).

A Scale of *Janavis finalidens* palatal & pectoral elements

Wilken et al. (2025) wrong scale
Supplementary fig. 14a

Scapula 50% of real size

Benito et al. (2022) correct scale

Left scapula

Shoulder girdle of *Ichthyornis dispar*
Specimen FHSM 18702

Scapula

Coracoid

B *Janavis finalidens* pterygoid interpreted as a juvenile left coracoid

Janavis finalidens†
NHMM RD 271
Ichthyornithes

No procoracoid process or scapular cotyla
"Glenoid"
Ossified & absent "acrocoracoid process" (ACP)
Convex "dorsal" surface (DS)
D
O
Simple trabecular structure (TS)
CS
"Glenoid"
L
Pneumatic foramen
Concave & solid "sternal end" (SE)
Flat "lateral" surface (LS)
C
"Medially"-oriented "sternal process" (SP)

Anser cygnoides
29 days old
Anseriformes

Unossified omal end
D
O
CS
Flat DS (F-DS)
Apneumatic (AP) Pneumatic in adult
No TS
L
L-SP
Convex LS (C-LS)
Unossified "hollow" SE
C

Thalassornis leuconotus
1 month old
Anseriformes

Scapular cotyla (SC)
Procoracoid process (PCP)
D
O
F-DS
Unossified acrocoracoid process (ACP)
Laterally-oriented SP (L-SP)
CS
AP
No glenoid
SC
No TS
L
C-LS
L-SP
L-SP
Unossified "hollow" SE
C

Stercorarius antarcticus
Immature indet.
Charadriiformes

Incipient ACP
PCP
SC
D
O
L-SP
Incipient glenoid
F-DS
AP Pneumatic in adult
CS
Incipient glenoid
SC
No TS
L
C-LS
SC
L-SP
Unfinished bone SE
C
L-SP

Gallinula chloropus
Immature indet.
Gruiformes

ACP
PCP
Glenoid
SC
D
O
L-SP
Glenoid
Concave DS
AP
CS
Glenoid
SC
Complex TS
L
C-LS
SC
L-SP
Narrow ossified SE
C
L-SP

Ichthyornis dispar†
KUVP 2281
Ichthyornithes

ACP
PCP
SC
D
O
L-SP
Glenoid
F-DS
CS
AP
Glenoid
SC
Complex TS
L
C-LS
L-SP
Narrow ossified SE
C
L-SP

C *Janavis finalidens* pterygoid interpreted as a right pterygoid

Janavis finalidens†
NHMM RD 271
Ichthyornithes

Basipterygoid process contact (BPC)
Pneumatic foramen (PF)
Flat medial surface (F-MS)
M
R
Laterally-open palatine contact (L-PC)
Orbito-pterygoid contact process (OPC)
Simple trabecular structure (TS)
CS
L-PC
PF
Convex dorsal surface (C-DS)
D
OPC
Concave quadrate articular surface (C-QAS)
C

Dasornis toliapica†
NHMUK PV 44096
Pelagornithidae

BPC
M
R
BPC
F-MS
L-PC
Simple TS
CS
PF
L-PC
PF
C-QAS
PF
D
C-DS
PF

Nettapterornis oxfordi†
NHMUK PVA 5922
Anseriformes

BPC
L-PC
M
R
BPC
OPC
CS
L-PC
Simple TS
L-PC
PF
C-QAS
C-DS
OPC
D
OPC
C-QAS
C

Chauna torquata
Adult
Anseriformes

BPC
M
R
L-PC
BPC
PF
L-PC
OPC
CS
Simple TS
C-QAS
OPC
C-DS
D
OPC
C-QAS
C

Megapodius nicobariensis
Adult
Galliformes

BPC
M
R
L-PC
BPC
F-MS
L-PC
OPC
Simple TS
CS
L-PC
C-QAS
OPC
D
OPC
C-QAS
C

Gallus gallus
Adult
Galliformes

BPC
M
R
L-PC
BPC
F-MS
OPC
L-PC & Palatine
Simple TS
CS
C-QAS
L-PC
OPC
C-DS
D
OPC
C-QAS
C

A Caudal palatine-hemipterygoid complex (right) & relation with kinetic capabilities

Ichthyornis dispar†
AMNH FARB 32773
Ichthyornithes
Likely prokinetic

Struthio camelus
Palaeognathae
Not prokinetic

Rhea americana
Palaeognathae
Not prokinetic

Gallus gallus
Galliformes
Prokinetic

Porphyrio martinicus Imm.
Gruiformes
Prokinetic

Porphyrio martinicus Adult
Gruiformes
Prokinetic

B Quadrate morphology (left), approach to braincase (BC) & relation with kinetic capabilities

Ichthyornis dispar†
AMNH FARB 32773 &
BHI 6421 composite
Ichthyornithes
Bicondylar
Close BC approach?
Likely prokinetic

Crypturellus tataupa
Palaeognathae
Tricondylar
No BC approach
Not prokinetic

Ptilopachus petrosus
Galliformes
Bicondylar
No BC approach
Prokinetic

Podiceps taczanowskii
Mirandornithes
Tricondylar
Close BC approach
Prokinetic

Ardea alba
Pelecaniformes
Tricondylar
Close BC approach
Prokinetic

Momotus momota
Coraciiformes
Tricondylar
Close BC approach
Prokinetic

C Wilken et al. 2025 cranial linkage system of *Ichthyornis* reconstruction is erroneous

Ichthyornis skull reconstruction modified from Torres et al. (2021). Single articulated skull (AMNH FARB 32773).

Ichthyornis skull reconstruction from Benito et al. (2022a). Composite from several specimens.