# Supporting Information 

# Evolutionary patterns of vane barb angles in birds 

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## Included:

1. Dataset and methods
2. Aves with different flight styles plotted in a wing phylomorphospace described by leading and trailing vane barb angles (Figure S1).
3. Aves with different flight styles plotted in a wing phylomorphospace described by trailing vane barb and barb angle asymmetry values. (Figure S2)
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8.Measurements used for extant birds from Feo et al. (2015) (Table S1)
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## Dataset

Values for trailing vane angle and barb angle asymmetry for $25 \%$ feather length are significantly different for the Feo et al. sample. We chose to use the $50 \%$ values in this reanalysis since these were most similar to the measurement locations available for the majority fossil feathers sampled and because Feo et al. only provided $50 \%$ measurements for non-volant extant taxa. For Mesozoic taxa Measured values were compared to published values for specimens V15336, V13156, DNHM-D3078, BMNH-Ph000881 and Berlin specimen of Archaeopteryx utilized in that analysis.

## Methods

(a) Phylogenetic signal and ancestral state reconstruction

Pagel's lambda [1] and Blomberg's K [2] were performed in R v. 3.0.1 [3] using the Phytools package (function phylosig [4]) to assess phylogenetic signal of barb angles. The influence of the phylogeny increases with lambda from 0 (no phylogenetic signal) and 1 (strong phylogenetic signal). When lambda $=0$, a star phylogeny results with all tips radiating from a basal node, describing a model where traits evolve independent of the phylogeny. When lambda = 1, trait evolution followed a Brownian motion model, where branch lengths would be proportional to divergence. Blomberg's $K$ statistic is used to test whether the observed distribution of traits exhibits more or less divergence than expected for traits evolving under Brownian motion. Values of $K$ close to 1 indicate trait similarity is proportional to divergence and a Brownian motion model of evolution fits the data. $K>1$ indicates that close relatives are more similar than expected, and $\mathrm{K}<1$ indicates more divergence between taxa than expected under a Brownian model.

Mesquite (v. 2.75 [5]) was used to map barb angles onto the reference phylogeny. Each character was traced onto the tree using the 'reconstruct ancestral state' module of Mesquite with weighted squared change parsimony [6]. Given the tree and observed character distribution, this method finds the ancestral states that minimize the number of steps of
character change.
1000 time-calibrated trees for the possible phylogenetic affinities of these 73 birds were sampled from the posterior distribution of Jetz et al [7] (http://www.birdtree.org). These trees use the Hackett et al [8] topology as a backbone. A majority rules consensus tree was built by Mesquite [5]. As taxa within passerines and rails are not fully resolved, the consensus tree was further resolved following recent phylogenetic hypotheses for passerines and rails [9, 10]. For Mesozoic taxa, we generate a fossil subtree with the timePaleoPhy function in paleotree [11] based on published fossil ages and branch [12, 13] (summarized in table S3). We grafted this time-calibrated tree of extinct taxa to the Aves tree with the bind.tip function in Phytools.
(b) Statistical analysis

All statistical analyses were conducted using the $R$ statistical computing environment v.3.4.3 ( $R$ Development Core Team, 2017). To account for phylogeny via PGLS. ANOVA on PGLS models were conducted in $R$ using the procD.pgls function in Geomorph package (residual randomisation permutation procedure) [14] and pairwise comparision was made in RRPP package [15]. The procD.pgls function performs ANOVA and regression models in a phylogenetic context under a Brownian motion model of evolution. The approach is derived from the statistical equivalency between parametric methods utilizing covariance matrices and methods based on distance matrices [14]. The randomization of residuals in a permutation procedure was used with 1000 iterations. F statistics are calculated from sums of squares based on coefficients that have appropriately accounted for phylogenetic relatedness. Effect-sizes (Z scores) are computed as standard deviates of the $F$ sampling distributions generated [14].

The function Phylogenetic generalized least-squares (pGLS) were also performed in $R$ package Caper [16] to assess the relationship between trailing vane barb angle and barb angle asymmetry. We categorized flight styles for living birds as those defined by Bruderer et al [17]. Specifically, "continuous flapping", "flapping and soaring", "flapping and gliding" and "passerine type flight". Flightless species and fossils were categoried as disticnt flight styles in this study. Measurements were log transformed to obtain a normal distribution as methods used have the assumption of normally distributed data.


Figure S1. Aves with different flight styles and Mesozoic taxa plotted in a wing phylomorphospace described by leading and trailing vane barb angles. Red, continuous flapping "CF"; blue, flapping and soaring "FS"; purple, flapping and gliding "FG"; green, passerine type flight "PT"; brown, flightless "FL"; black, fossil data.


Figure S2. Aves with different flight styles and Mesozoic taxa plotted in a wing phylomorphospace described by trailing vane barb and barb angle asymmetry values. Red, continuous flapping "CF"; blue, flapping and soaring "FS"; purple, flapping and gliding "FG"; green, passerine type flight "PT"; brown, flightless "FL"; black, fossil data.

Interestingly, we found that distinct from other water birds, loons showed very reduced barb angles and angle asymmetry ( 11.829 and 30.508 for leading and trailing vane barb angles), similar to those of grebes (Figures 1,2, S2).

As barb angles were measured from different feathers and positions, we also mapped the values of barb angle variance across modern flight species. Similar patterns were recovered in ancestral state reconstructions of angle variance values. The leading vane and angle asymmetry values show reduced variance in galloanseres and passerines. Variance for trailing vane barb angles is also small in passerines (Figures S3-S5).


Fig S3. Changes in trailing vane barb angle variance from different feathers (inner, middle and outer) and positions ( $25 \%$ and $50 \%$ from feather tip) across extant flight species.


Fig. S4 Changes in vane barb angle asymmetry variance from different (inner, middle and outer) and positions ( $25 \%$ and $50 \%$ from feather tip) across extant flight species.


Fig. S5 Changes in Leading vane barb angle variance from different (inner, middle and outer) and positions ( $25 \%$ and $50 \%$ from feather tip) across extant flight species.


Fig. S6 Ancestral state reconstruction for leading (a), trailing (b) vane barb angles and barb angle asymmetry (c) showing the general evolutionary pattern for simplified phylogeny. Diagrams show the barb geometry. Grey, leading vane barb; purple, trailing vane barb.

Table S1. Measurements used for extant birds from Feo et al. (2015)*

| Species | Ld | Tr | Diff | flight style |
| :--- | :--- | :--- | :--- | :--- |
| Coragyps_atratus | 16.008 | 37.495 | 21.488 | FS |
| Cathartes_aura | 18.222 | 46.757 | 28.535 | FS |
| Elanus_leucurus | 14.887 | 40.099 | 25.212 | FS |
| Buteo_jamaicensis | 13.223 | 37.100 | 23.878 | FS |
| Accipiter_gentilis | 8.849 | 38.133 | 29.284 | FS |
| Accipiter_striatus | 11.916 | 33.505 | 21.589 | FS |
| Anser_fabalis | 9.269 | 44.457 | 35.187 | CF |
| Anas_discors | 6.226 | 45.384 | 39.158 | CF |
| Tachyeres_brachypterus | 7.067 | 21.171 | 14.104 | FL |
| Tachyeres_leucocephalus | 5.885 | 30.917 | 25.031 | FL |
| Calypte_anna | 12.664 | 43.836 | 31.172 | CF |
| Chaetura_pelagica | 15.973 | 44.779 | 28.806 | FG |
| Chordeiles_acutipennis | 7.250 | 44.577 | 37.327 | FG |
| Jacana_jacana | 20.919 | 47.810 | 26.891 | CF |
| Numenius_americanus | 6.903 | 47.417 | 40.514 | CF |
| Larus_marinus | 11.022 | 47.423 | 36.401 | CF |
| Zenaida_asiatica | 14.690 | 50.156 | 35.466 | CF |
| Streptopelia_decaocto | 10.968 | 53.943 | 42.975 | CF |
| Columba_livia | 19.787 | 53.782 | 33.996 | CF |
| Columbina_inca | 13.582 | 40.570 | 26.989 | CF |
| Columbina_passerina | 16.558 | 39.732 | 23.175 | CF |
| Todus_mexicanus | 11.874 | 48.230 | 36.356 | PT |
| Coccyzus_americanus | 21.291 | 47.710 | 26.419 | CF |
| Falco_columbarius | 16.050 | 39.114 | 23.064 | FG |
| Callipepla_californica | 9.340 | 32.963 | 23.623 | CF |
| Alectoris_chukar | 9.855 | 37.650 | 27.794 | CF |
| Dendragapus_canadensis | 10.427 | 33.645 | 23.218 | CF |
| Numida_meleagris | 9.331 | 30.138 | 20.807 | CF |
| Gavia_immer | 11.829 | 30.508 | 18.679 | CF |
| Habroptila_wallacii | 22.393 | 45.832 | 23.440 | FL |
| Atlantisia_rogersi | 16.218 | 42.375 | 26.156 | FL |
| Fulica_gigantea | 14.321 | 31.000 | 16.679 | FL |
| Porzana_atra | 15.295 | 31.664 | 16.369 | FL |
| Aramidopsis_plateni | 24.205 | 51.094 | 26.890 | FL |
| Megacrex_inepta | 41.273 | 70.174 | 28.901 | FL |
| Gallirallus_australis | 26.919 | 49.250 | 22.331 | FL |
| Gallirallus_sylvestris | 18.205 | 44.733 | 26.528 | FL |
| Gallirallus_rovianae | 22.229 | 31.150 | 8.921 | FL |
| Rallus_limicola | 10.985 | 33.354 | 22.369 | CF |
| Tyrannus_vociferans | 12.215 | 44.285 | 32.070 | PT |
| Pipra_mentalis | 12.715 | 50.799 | 38.085 | PT |
| Aphelocoma_californica | 11.925 | 32.040 | 20.115 | CF |
| Cyanocitta_stelleri | 18.498 | 32.615 | 14.118 | CF |
| Corvus_brachyrhynchos | 17.885 | 31.449 | 13.564 | CF |
| Pica_nuttalli | 19.881 | 32.875 | 12.995 | CF |
| Hirundo_rustica | 19.154 | 37.410 | 18.256 | FG |
| Petrochelidon_pyrrhonota | 16.564 | 38.299 | 21.736 | FG |
| Progne_subis | 21.154 | 45.293 | 24.139 | FG |
|  |  |  |  |  |


| Tachycineta_bicolor | 21.071 | 44.374 | 23.303 | FG |
| :--- | :--- | :--- | :--- | :--- |
| Tachycineta_thalassina | 20.222 | 39.624 | 19.402 | FG |
| Oporornis_tolmiei | 17.854 | 33.617 | 15.762 | PT |
| Protonotaria_citrea | 18.292 | 36.756 | 18.464 | PT |
| Dendroica_coronata | 15.581 | 34.123 | 18.542 | PT |
| Seiurus_motacilla | 16.396 | 40.809 | 24.414 | PT |
| Vermivora_celata | 13.224 | 31.267 | 18.043 | PT |
| Junco_hyemalis | 14.478 | 26.672 | 12.194 | PT |
| Zonotrichia_albicollis | 14.593 | 31.064 | 16.471 | PT |
| Passerella_iliaca | 16.000 | 32.891 | 16.891 | PT |
| Pipilo_maculatus | 13.719 | 27.074 | 13.354 | PT |
| Spizella_pusilla | 14.567 | 34.880 | 20.313 | PT |
| Casmerodius_albus | 14.982 | 59.920 | 44.937 | CF |
| Pelecanus_erythrorhynchos | 21.610 | 43.411 | 21.801 | FS |
| Phalacrocorax_harrisi | 21.789 | 43.270 | 21.481 | FL |
| Sphyrapicus_varius | 12.498 | 44.609 | 32.112 | PT |
| Podiceps_taczanowskii | 16.903 | 28.605 | 11.702 | FL |
| Daption_capense | 9.175 | 49.622 | 40.447 | FS |
| Thalassarche_melanophrys | 10.181 | 37.892 | 27.711 | FS |
| Nymphicus_hollandicus | 10.707 | 38.358 | 27.652 | CF |
| Micrathene_whitneyi | 12.011 | 35.665 | 23.655 | CF |
| Athene_cunicularia | 12.207 | 44.568 | 32.361 | CF |
| Surnia_ulula | 10.595 | 40.984 | 30.389 | CF |
| Crypturellus_soui | 6.378 | 27.558 | 21.180 | CF |
| Trogon_violaceus | 8.041 | 28.254 | 20.213 | CF |
| La, leading vane angle | Traling | vane ange; Diff | $b a n$ | aifer |

Ld, leading vane angle; Tr, trailing vane angle; Diff, barb angle differences; CF, continours flapping; FS, flapping and soaring; FG, flapping and gliding; PT, passerine type flight; FL, flightless *Specimen numbers and localities can be found from Feo et al. (2015).

Table S2 Measurements used for fossil taxa

| Specimen No. | Species | Ld | Tr | Diff |
| :--- | :--- | ---: | ---: | ---: |
| STM-7-145 | Archaeornithura meemannae | 10.62 | 23.16 | 12.54 |
| V13631 | Yixianornis grabaui | 17.00 | 28.29 | 11.29 |
| V11309 | Longirostravis hani | 9.85 | 20.12 | 10.27 |
| STM-24-1 | Eopengornis martini | 7.42 | 23.59 | 16.16 |
| V15336 | Eopengornis martini | 11.13 | 19.46 | 8.33 |
| V13156 | Confuciusornis sanctus | 11.37 | 23.14 | 11.77 |
| CUGB-P1401 | Confuciusornis sanctus | 10.36 | 24.38 | 14.03 |
| AGB5488 | Confuciusornis sanctus | 9.22 | 22.75 | 13.53 |
| V13178 | Confuciusornis sanctus | 8.27 | 20.69 | 12.41 |
| V13175 | Confuciusornis sanctus | 11.87 | 23.02 | 11.15 |
| V11977 | Eoconfuciusornis zhengi | 12.50 | 17.34 | 4.84 |
| DNHM-D3078 | Sapeornis chaoyangensis | 7.84 | 13.78 | 5.93 |
| Berlin specimen | Archaeopteryx lithographica | 11.11 | 19.27 | 8.17 |
| BMNH-Ph000881 | Microraptor gui | 12.74 | 18.48 | 5.74 |
| V13352 | Microraptor gui | 12.90 | 19.08 | 6.18 |
| NGMC 97-4-A DSC2551 | Caudipteryx zoui | 18.72 | 21.66 | 2.93 |

Ld, leading vane angle; Tr, trailing vane angle; Diff, barb angle differences; STM, Shandong Tianyu Natural History Museum; V, Institute of Vertebrate Paleontology and Paleoanthropology; CUGB, China University of Geosciences (Beijing); DNHM, Dalian Natural History Museum; BMNH, Beijing Museum of Natural History; NGMC, National Geological Museum of China.

Table S3 Fossil age (Ma) data for paleotree [31].

| Species | Min_age | Max_age | Start | End | Midpoint |
| :--- | :--- | :--- | ---: | ---: | ---: |
| Yixianornis_grabaui | Aptian | Aptian | 126.3 | 113 | 119.65 |
| Longirostravis_hani | Late_Barremian | Early_Aptian | 128.55 | 119.65 | 124.1 |
| Eopengornis_martini | Hauterivian | Hauterivian | 133.9 | 130.8 | 132.35 |
| Confuciusornis_sanctus | Late_Barremian | Early_Aptian | 128.55 | 119.65 | 124.1 |
| Eoconfuciusornis_zhengi | Hauterivian | Hauterivian | 133.9 | 130.8 | 132.35 |
| Sapeornis_chaoyangenis | Aptian | Aptian | 126.3 | 113 | 119.65 |
| Archaeopteryx_lithographica | Early_Tithonian | Early_Tithonian | 152.1 | 147.7 | 149.9 |
| Microraptor_gui | Aptian | Aptian | 126.3 | 113 | 119.65 |
| Caudipteryx_zoui | Aptian | Aptian | 126.3 | 113 | 119.65 |
| Archaeornithura_meemannae | Hauterivian | Hauterivian | 133.9 | 130.8 | 132.35 |

10. Images showing the angle measurements of Mesozoic fossil taxa.


Yixianornis grabaui (V13631)


Longirostravis hani (V11309)


Confuciusornis sanctus(V13156)


Confuciusornis sanctus (V13175)


Confuciusornis sanctus (V13178a)


Confuciusornis sanctus (V13178b)


Confuciusornis sanctus (AGB5488)


Confuciusornis sanctus (CUGB-P1401)


Eoconfuciusornis zhengi (V11977)


Sapeornis chaoyangensis (DNHM-D3078)


Archaeopteryx lithographica (Berlin specimen)


Microraptor gui (V13352)


Caudipteryx zoui (NGMC 97-4-A DSC2551)

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