


Article

## Relationships of mass properties and body proportions to locomotor habit in terrestrial Archosauria

Peter J. Bishop\* , Karl T. Bates, Vivian R. Allen, Donald M. Henderson, Marcela Randau, and John R. Hutchinson 

**Abstract.**—Throughout their 250 Myr history, archosaurian reptiles have exhibited a wide array of body sizes, shapes, and locomotor habits, especially in regard to terrestriality. These features make Archosauria a useful clade with which to study the interplay between body size, shape, and locomotor behavior, and how this interplay may have influenced locomotor evolution. Here, digital volumetric models of 80 taxa are used to explore how mass properties and body proportions relate to each other and locomotor posture in archosaurs. One-way, nonparametric, multivariate analysis of variance, based on the results of principal components analysis, shows that bipedal and quadrupedal archosaurs are largely distinguished from each other on the basis of just four anatomical parameters ( $p < 0.001$ ): mass, center of mass position, and relative forelimb and hindlimb lengths. This facilitates the development of a quantitative predictive framework that can help assess gross locomotor posture in understudied or controversial taxa, such as the crocodile-line *Batrachotomus* (predicted quadruped) and *Postosuchus* (predicted biped). Compared with quadrupedal archosaurs, bipedal species tend to have relatively longer hindlimbs and a more caudally positioned whole-body center of mass, and collectively exhibit greater variance in forelimb lengths. These patterns are interpreted to reflect differing biomechanical constraints acting on the archosaurian *Bauplan* in bipedal versus quadrupedal groups, which may have shaped the evolutionary histories of their respective members.

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### Introduction

Archosauria (the “ruling reptiles”) is a diverse, long-lived (~250 Myr) saurian clade that dominated terrestrial, aquatic, and aerial ecosystems throughout the Mesozoic Era, and persists to the present day in the form of about 23 crocodylian and 10,000 bird species (Oaks 2011; Jetz et al. 2012). Throughout their history, archosaurs have displayed disparate body plans and divergent locomotor habits, including obligate bipedal and quadrupedal species, as well as volant, semi-aquatic, and marine forms (Fig. 1). The majority of archosaurs were land dwelling, and it is well known that many important innovations related to terrestrial locomotion evolved throughout

archosaur history (Bakker 1971; Charig 1972; Parrish 1986; Hutchinson and Gatesy 2000; Hutchinson 2006; Sullivan 2015). Terrestrial locomotor evolution in archosaurs is also characterized by repeated instances of morphological and functional convergence, involving members of both the ornithodiran (bird-line) and pseudosuchian (crocodile-line) lineages (Walker 1964; Parrish 1986; Sereno 1991; Carrano 2000; Nesbitt and Norell 2006; Bates and Schachner 2012; Bates et al. 2012; Kubo and Kubo 2012; Maidment and Barrett 2012; Grinham et al. 2019).

Given their long evolutionary history and diverse array of body sizes and shapes,

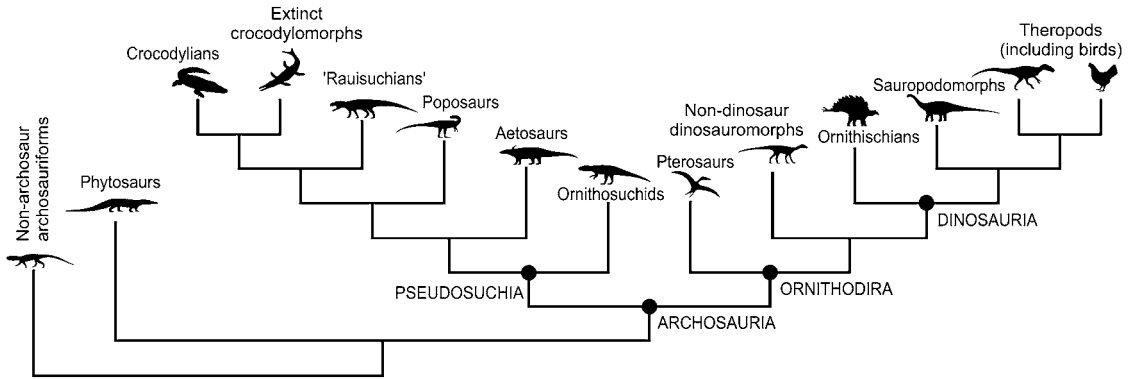


FIGURE 1. Simplified phylogeny of Archosauria based on current consensus of interrelationships (e.g., Nesbitt 2011; Ezcurra 2016), illustrating the diversity of body forms and locomotor habits across the group. Major clades are also indicated.

terrestrial archosaurs provide a good case study for investigating how size and gross body proportions may relate to each other and locomotor behavior. Yet extant archosaur species occupy a very select subset of the total range of observed archosaur postcranial morphologies, and concomitantly exhibit disparate locomotor patterns that almost certainly do not capture the full range of historical diversity in archosaur locomotor behavior (Hutchinson and Gatesy 2000; Hutchinson 2006). There is no archosaur—indeed, any animal—alive today that is of a comparable size and shape to extinct taxa like *Postosuchus* or *Iguanodon*. Although extant species may be highly useful for the development of “lower-level inferences” such as soft tissue anatomy, their utility can become increasingly uncertain for “higher-level inferences” such as locomotor function and behavior (Witmer 1995; Hutchinson 2006; Bates and Falkingham 2018). Thus, the usefulness of extant archosaur species as a basis for deriving broad inferences (grounded in homology or analogy) of locomotion for *all* extinct archosaurs is uncertain. Understanding locomotor habit in extinct archosaurs has therefore frequently relied on interpreting bony anatomy or footprints preserved in the fossil record. For instance, hoof-shaped manual unguals suggest at least some use of a quadrupedal stance (Maidment and Barrett 2014), and narrow-gauge footprints left by a single pair of feet may indicate at least intermittent parasagittal bipedalism (Thulborn 1990). The insights afforded by these lines of evidence can be

important, but they are limited by the incompleteness of the fossil record (i.e., such evidence is only preserved for some species, and even then footprints can rarely be assigned to a known species), and perhaps more critically, they often cannot be generalized across large clades of disparate body plans.

An alternative approach to investigating locomotor patterns across a clade as morphologically diverse as Archosauria is one based on phenomena that affect all species in a universal fashion. For example, the manner of stance and gait of all archosaur species are (or were) fundamentally dictated by Newtonian mechanics, insofar as the size and proportions of any terrestrial animal greatly influence its ability to support and move its body on land. Body mass is paramount, as the mass that an animal must support against gravity increases more quickly with respect to size (e.g., length) than the cross-sectional area of supportive tissues such as muscle and bone (Alexander 1985; Biewener 1990; Vogel 2003; Campione and Evans 2012). Limb size, as well as intra- and inter-limb proportions, will dictate the range of possible kinematics such as stride length and patterns of inter-limb coordination (Alexander and Jayes 1983; Gatesy and Biewener 1991; Sellers et al. 2009; Gatesy and Pollard 2011), as well as potential locomotor performance (Garland and Janis 1993; Carrano 1999). The location of the whole-body center of mass (COM), which represents the collective distribution of mass throughout the body, will also exert a strong influence on posture and movements. To maintain stability, an animal must

keep its COM over the “polygon of support” formed by its feet (however many) when standing (Alexander 2006; Winter 2009), and this principle can be extended to dynamically stable locomotion as well (Herr and Popovic 2008).

Many previous studies have estimated body mass or COM in extinct archosaurs through a variety of approaches, yet these have seldom been used in a comparative context to address broad questions concerning locomotor disparity or evolution (Henderson and Snively 2003; Allen et al. 2013; Bates et al. 2016). Here, previously published estimates of terrestrial archosaur mass properties and body proportions are synthesized with new computational models for additional taxa, spanning five orders of magnitude in mass and almost 250 Myr of evolution. The aims of the present study were twofold: first, to investigate how different aspects of body shape and size may or may not statistically relate to each other and gross locomotor habit (here taken to mean bipedal vs. quadrupedal stance) in terrestrial Archosauria; and second, to develop a generalized and quantitative predictive framework that can be used to help assess locomotor habit in uncertain or enigmatic taxa.

## Material and Methods

*Computational Modeling.*—With the exception of the models of *Rapetosaurus* (Bates et al. 2016) and *Dreadnoughtus* (Lacovara et al. 2014), it was assumed here that the models used in the core component of the present study were of adult (or near-adult) animals. Determining the ontogenetic status of many extinct archosaurs is not straightforward (Hone et al. 2016), but the models used consistently fell at or near the upper end of the known size range for each of the taxa studied and were therefore assumed to represent “adult” morphology. In terms of the anatomical parameters investigated (see “Parameters Investigated”) and broad comparative approach of the study, it was felt that this assumption was justified, although the extremely limited number of known specimens for some taxa renders it difficult to test.

Previously published three-dimensional (3D) digital volumetric models for 72 taxa were used (see Supplementary Material, Supplementary Table S1), which includes 2 non-

archosaur archosauriforms, 6 pseudosuchians (including two extant taxa), 1 non-dinosaur dinosauromorph, 15 ornithischians, 29 theropods (including five extant taxa), and 19 sauropodomorphs. These data were synthesized with new models for an additional eight taxa (one extant, seven extinct). For all the extinct taxa, and some of the extant taxa, digital volumetric models were produced using one of three established protocols (Fig. 2): a 3D slicing method that discretizes the body into a series of elliptical frusta (Henderson 1999); a convex hull-based method whereby convex hulls are fit around each major body segment (Sellers et al. 2012; Bates et al. 2016); and a spline- or hoop-based method whereby a series of polygonal hoops are fit around each body segment and then lofted together (Allen et al. 2009, 2013). All three methods have been previously demonstrated to produce accurate results for a variety of extant tetrapod taxa (for which, admittedly, the approximate body geometry is already known).

Models for three extant avian taxa (*Gallus*, *Anas*, and *Buteo*) were generated directly from computed tomographic scanning of whole carcasses (*Gallus*: Allen et al. 2013; *Anas* and *Buteo*: Macaulay et al. 2017), and the values for mass and COM location used for these models were taken from the “best guess” results; see Allen et al. (2013) and Macaulay et al. (2017) for details on density assignment. The same approach was also used to generate the new model of *Crocodylus moreletii* (machine tube voltage 120 kV, tube current 233 mA, exposure 750 ms, pixel resolution 0.977 mm, slice thickness 1.5 mm; scan segmentation in Mimics 19.0 [Materialize NV, Leuven, Belgium]), with density assignments following “best guess” values of Allen et al. (2013) and Macaulay et al. (2017). For most other models, the values for mass and COM in all other models were taken directly from the original, published results. Two exceptions to this were as follows:

1. For the theropod models of Allen et al. (2013), a “best guess” model did not exist, but rather there were end-member extremes used to delimit the plausible ranges of mass and COM in the original study. In the present study, the mass used was the average

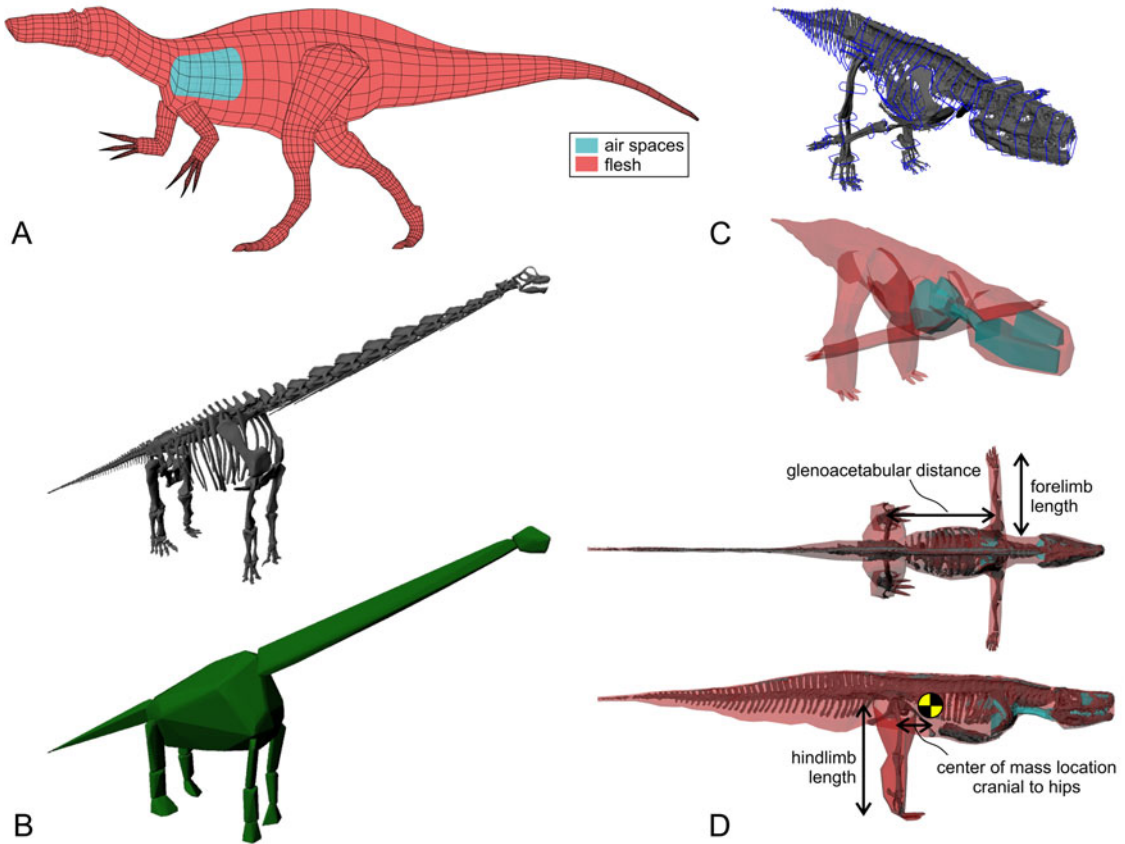


FIGURE 2. Digital modeling of extinct archosaurs. A, 3D slicing method, here with the theropod *Suchomimus*. B, Convex-hull method, here with the sauropod *Giraffatitan*. C, Spline- or hoop-based method, here with the raiuisuchian *Batrachotomus* (new analysis). Whereas 3D slicing is based on 2D illustrations in different views, the other methods are based on mounted skeletal material. D, Anatomical parameters extracted from each model used in the analyses. Glenoacetabular distance was measured parallel to the sagittal plane, and forelimb and hindlimb lengths were the sum of the lengths of the respective propodia, zeugopodia, and metapodia (or their corresponding flesh segments).

of the “maximal mass” and “minimal mass” models, and the COM location used was the average of the “maximally cranial” and “maximally caudal” models.

- For the sauropodomorph models of Bates et al. (2016), mass and COM were determined following an approach that differed slightly from the original study. Low-density (e.g., lung) volumes were ignored, and rather the “+21%” convex-hull models for all segments were used to calculate segment mass properties directly, assuming a density of  $1000 \text{ kg/m}^3$  for the tail and limbs and a density of  $850 \text{ kg/m}^3$  for the trunk, neck, and head. The purpose of this alternative protocol was to enable total consistency with new models that were

generated in this study via the convex-hull technique.

Seven new extinct taxa were added to the present study, and were generated as follows. *Tenontosaurus*, *Trilophosaurus*, *Stenaulorhynchus*, and *Protoceratops* were modeled using the convex-hull method, as described earlier for the sauropodomorphs of Bates et al. (2016), based on digital skeletal geometries previously published by Clauss et al. (2017). *Edmontonia* was also modeled using the same convex-hull method, but this was based on a digital skeletal sculpt based on fossil material (Sellers 2016). *Batrachotomus* and *Muttaborrasaurus* were modeled using the spline-based method, and were based on digital skeletons generated from

photogrammetry of mounted skeletons (*Batrachotomus*: Staatliches Museum für Naturkunde, Stuttgart; *Muttaborrasaurus*: Queensland Museum, Brisbane), using the software Agisoft Photoscan 1.04 (Agisoft LLC, Russia). The lungs of both taxa were modeled as filling the cranial portion (cranial half in the “maximally caudal” model, cranial quarter in the “maximally cranial” model) of the thoracic cavity (Allen et al. 2009, 2013); no abdominal or other air sacs were modeled owing to the paucity of evidence for postcranial skeletal pneumaticity in pseudosuchians and ornithischians (Butler et al. 2012).

*Comparison of Modeling Methods.*—Previous studies that have used digital volumetric modeling of extinct taxa have typically employed only a single modeling protocol, whereas the current study draws upon data from three. In addition to technical details concerning model generation and density assignment, there are at least three important differences between the methods involved:

1. Whereas the convex-hull and spline-based methods are based on 3D digital skeletal reconstructions, the 3D slicing method is based on two-dimensional reconstructions (lateral and dorsal views).
2. Both the convex-hull and spline-based methods create models in a standard posture, with the forelimbs held out either directly laterally or ventrally from the body, and the hindlimbs directed downward (Fig. 2C); often the vertebral column is also held in a straightened fashion for at least part of its length. In contrast, models generated using the 3D slicing method have a more natural, lifelike pose, but one that is not necessarily standardized across models.
3. Both the 3D slicing and spline-based methods involve a significant element of manual investigator input in the generation of (albeit more natural) flesh contours, whereas the convex-hull method is almost entirely automated. Nonetheless, in the spline-based method of Allen et al. (2009), the contours are based on empirically derived scaling factors, regarding how “inflated” contour cross sections are with respect to the underlying skeleton; it hence lies between the 3D slicing and convex-hull methods in terms of objectivity.

The collective differences between the three approaches raise the possibility that there may exist one or more systematic differences in estimates of mass or COM location between the modeling methods.

As the comparability of different protocols’ results has yet to be formally examined, an attempt was made here to assess method consistency. This assessment was unfortunately restricted by the strong historical bias between methodology and locomotor habit in the current dataset (chi-square test for association,  $\chi^2 = 30.5$ ,  $p < 0.001$ ; Supplementary Table S1); for instance, of the taxa with “known” locomotor habit (see “Statistical Analyses”), only one quadruped was modeled using the spline method. Moreover, there was also strong bias between methodology and clade ( $\chi^2 = 56.2$ ,  $p < 0.001$ ; Supplementary Table S1). Such bias not only limits the ability to distinguish systematic differences between modeling methods from genuine differences due to locomotor habit or clade, but it also limits the ability to delimit the true nature of any such difference between methods and thence apply appropriate corrections to the dataset (e.g., corrections may be disproportionately applied to some locomotor habits or clades). Here, estimates for mass and COM location for 27 taxa that had been modeled using multiple protocols were compiled, using both previously published results and new models generated here via 3D slicing or convex hulling (see Supplementary Material, Supplementary Table S2). Generation of convex-hull models followed the same protocol as outlined earlier. Not all 27 taxa were modeled using all three methods, but in each case at least two had been used. As different-sized skeletons were often used as the basis upon which these models were built, the attempt was made to first remove the effect of size by normalizing the estimates of mass and COM location (distance cranial to the hips) by glenoacetabular distance (GA):

$$\text{mass}^* = \frac{\sqrt[3]{\text{mass}}}{\text{GA}}, \quad (1)$$

$$\text{COM}^* = \frac{\text{COM}}{\text{GA}}. \quad (2)$$

While the normalization expressed in equation

1 assumes isometry, it is important to recognize that the normalized values were only compared vis-à-vis for (at worst) slightly different-sized models of the same taxon. They were not compared across models (taxa) of widely varying sizes or proportions, for which the assumption of isometry is clearly untenable. Given that within a taxon there would be relatively little scope for size differences to begin with (indeed, in some cases, the same underlying skeleton was used for multiple methods), the assumption of isometry being violated here was considered negligible.

For a given pair of methods, normalized values of mass and COM location were compared using the *smatr* package (v. 3.4-3; Warton et al. 2012) in R v. 3.5.2 (R Core Team 2012). A major axis (MA) regression forced through the origin was fit to the data and then tested for whether its slope was statistically different from 1.0 (significance level set at 0.05); if there was a difference, this suggested that one method tended to systematically under- or overestimate a given parameter compared with the other. The use of a zero-intercept MA regression here was simply to test the congruence between a given pair of methods, such that the MA slope gives a measure of systematic deviation. This assumed that if there was a systematic difference in one method compared with another, that difference would be proportional to the magnitude of the quantity being compared; given the manner in which the digital models were constructed, it was considered implausible that a negative-slope (or nonzero intercept) relationship could occur between two methods.

Regression indicated that some systematic differences did indeed exist, as far as can be

determined with the current sample (Table 1). The convex-hull method gave different estimates from the spline method in terms of both COM and mass, and from the 3D slicing method in terms of COM; yet there was no detectable difference between 3D slicing and spline methods. Assuming that these differences also extended to the main dataset, values for mass and COM derived from the convex-hull method in the main dataset were “normalized” with respect to the spline method (Supplementary Table S1), using the MA regression slopes as correction factors:

$$\text{mass}_{\text{spline-corrected}} = \frac{\text{mass}_{\text{convex hull}}}{0.90717^3} \quad (3)$$

$$\text{COM}_{\text{spline-corrected}} = \frac{\text{COM}_{\text{convex hull}}}{1.5434} \quad (4)$$

This restricted analysis of, and correction for, systematic differences between modeling methods is admittedly weak, and consequently in all remaining analyses statistical significance was conservatively identified with an alpha level of 0.01. Evidently, the issue deserves a more thorough and rigorous investigation in the future, which will first require the generation of many more models to reduce the aforementioned distribution biases. Until such detailed investigation is undertaken, it is prudent that future studies that focus on a single taxon should employ multiple methods (e.g., Otero et al. 2019) as a way of assessing the sensitivity of the results to the modeling method used.

*Parameters Investigated.*—For each model, four anatomical parameters were determined, in addition to GA (Fig. 2D): body mass (BM),

TABLE 1. Results for comparison between different modeling approaches for size-normalized estimates of mass and center of mass (mass\* and COM\*, respectively; see eqs. 1 and 2), using major axis regression forced through the origin. Statistically significant results are noted in boldface, indicating systematic bias between modeling approaches; daggers (†) indicate that a significant difference was removed following correction as described in the text. Approaches are listed as ordinate versus abscissa in the regressions.

| Comparison                 |       | <i>n</i> | Slope   | <i>r</i> <sup>2</sup> | <i>F</i> | <i>p</i>        |
|----------------------------|-------|----------|---------|-----------------------|----------|-----------------|
| Convex hull vs. 3D slicing | mass* | 21       | 0.878   | 0.926                 | 4.049    | 0.0579          |
|                            | COM*  | 21       | 1.49277 | 0.838                 | 16.638   | <b>0.0006</b> † |
| Convex hull vs. spline     | mass* | 13       | 0.90717 | 0.988                 | 9.063    | <b>0.0100</b>   |
|                            | COM*  | 13       | 1.5434  | 0.857                 | 13.856   | <b>0.0026</b> † |
| 3D slicing vs. spline      | mass* | 8        | 1.10112 | 0.930                 | 0.742    | 0.4176          |
|                            | COM*  | 8        | 1.12146 | 0.841                 | 0.418    | 0.5387          |

COM location cranial to the hips ( $COM_x$ ), hind-limb length (HL), and forelimb length (FL). The use of these four relatively simple parameters is justified on the basis that each has direct physical relevance to terrestrial locomotor biomechanics (see “Introduction”). It is worth reiterating here that only volumetric models were used in this study, as they give internally consistent and mechanistically based estimates of both BM and  $COM_x$ , in contrast to other methods that compute BM only, such as those based on propodial minimal circumferences (e.g., Campione et al. 2014). Such mechanistic estimates can in turn be more lucidly related to other aspects of locomotor biomechanics.

Due to the vast range in absolute size across the taxa sampled, the raw values for each parameter were corrected to account for differences in size. For BM, the base-10 logarithm was taken, which also helped reduce the sample distribution’s skewness. For the linear metrics of  $COM_x$ , HL, and FL, a linear regression of each metric against GA was computed in R and residuals were extracted; to account for suspected differences in allometric trajectories between GA and the different metrics, the base-10 logarithm of each metric was taken first before computing the regressions. This use of residuals was favored over the computation of basic ratios (e.g., HL/GA), as it reduces heteroscedasticity and nonlinearity in the resulting dataset.

In the original dataset, the archosauromorph *Trilophosaurus* had a negative  $COM_x$  position (Supplementary Table S1), caused at least in part by its long and proximally deep tail, which is incompatible with the above approach of computing logarithms. The long tail would presumably have been dragged on the ground in life for some of its length (Gregory 1945), and this would have shifted the COM of that part of the body supported by the limbs to lying cranial to the hips by some amount. As such, COM of this taxon was nominally set as 1 cm cranial to the hips. Given that *Trilophosaurus* is the most stemward taxon in the current dataset, this was judged as an acceptable modification to facilitate statistical analysis.

*Statistical Analyses.*—Phylogenetically informed statistical analyses of the size-corrected anatomical variables were conducted

in R using a fully resolved, time-calibrated phylogenetic tree of the study taxa, which follows the “traditional” hypothesis of dinosaur interrelationships (see Supplementary Fig. S1). The topology of the tree was based on Nesbitt (2011) and Ezcurra (2016) for Archosauromorpha and Pseudosuchia; Carrano and Sampson (2008), Carrano et al. (2012), Jetz et al. (2012), Allen et al. (2013), and Loewen et al. (2013) for Theropoda; Nair and Salisbury (2012), Otero and Pol (2013), Tschoop et al. (2015), and Bates et al. (2016) for Sauropodomorpha; and Butler et al. (2007, 2011), Prieto-Márquez (2010), McDonald (2012), and Farke et al. (2014) for Ornithischia. The ages used for terminal taxa and internal nodes for this tree are reported in Supplementary Tables S3 and S4, respectively. Analyses were also conducted using a second tree of markedly different topology, based on the “Ornithoscelida hypothesis” of dinosaur interrelationships (Baron et al. 2017; see Supplementary Fig. S2); the ages of terminal taxa remained identical to those of the first tree, whereas the ages used for the internal nodes are reported in Supplementary Table S5.

Phylogenetic principal components analysis (pPCA; Revell 2009) was used to explore how morphological variation related to locomotor habit, using the phytools package for R (v. 0.6; Revell 2012), where the evolutionary correlation matrix was derived assuming a Brownian model of trait evolution. To assess the influence of phylogeny, the  $K$  statistic (and associated  $p$ -value) of Blomberg et al. (2003) was also determined in phytools. In this study, the locomotor habit of each taxon was classified a priori based on current consensus (see also Thulborn 1990; Maidment and Barrett 2014; Grinham et al. 2019 and references cited therein), as either obligate biped ( $n = 33$ ), obligate quadruped ( $n = 34$ ), or “other” ( $n = 13$ ), where the last category contained taxa hypothesized to be facultatively bipedal or taxa whose habits are controversial or previously not assessed in detail. Locomotor mode is here defined as the stance adopted during straight-line, level, quasi-steady walking.

The pPCA served primarily as an aid to qualitatively assess broad patterns. The distinction between bipedal and quadrupedal

principal component (PC) scores was quantitatively tested with a one-way, nonparametric, multivariate analysis of variance (PERMANOVA) in the *vegan* package for R (v. 2.5-6; Oksanen et al. 2019). Disparity in PC scores was estimated for bipeds and quadrupeds using Procrustes distance regression in the *geomorph* package in R (v. 3.1.1; Adams et al. 2018). To investigate how the size-corrected anatomical parameters themselves related to each other and locomotor habit, three analyses were undertaken. First, a phylogenetic multivariate analysis of variance (pMANOVA) was conducted, using phylogenetic Procrustes distance regression, to test whether there was an overall difference between bipeds and quadrupeds considering all four parameters together. Second, a phylogenetic analysis of variance (pANOVA) was conducted, using the *phytools* package, to test whether there was a difference between bipeds and quadrupeds considering each parameter independently. Finally, pairwise comparisons between parameters were conducted for bipeds and quadrupeds via MA regression, using the *smatr* package. This involved correcting the values for each parameter for phylogenetic signal, using phylogenetic generalized least squares in the *caper* package in R (v. 0.5.2; Orme et al. 2015), the residuals from which were then used in the regressions. The slopes for a given pairwise MA fit were compared between bipedal and quadrupedal taxa using the *smatr* package.

**Predictive Framework.**—Linear discriminant analysis (LDA) was used to produce a predictive framework for archosaur locomotor mode based on the original (size-corrected) anatomical parameters, using the *MASS* package for R (v. 7.3-50; Venables and Ripley 2002). Instead of deriving a single predictive model, 11 variants were tested here, which differed in the combination of anatomical variables (two, three, or all four) used in the training dataset. In addition, as birds were found to occupy a distinct position in morphospace (see “Results”), two variants of the training dataset were also tested: one with all “known” taxa (i.e., 33 bipeds and 33 quadrupeds) and one excluding birds (i.e., 25 bipeds and 33 quadrupeds); as *Trilophosaurus* was found to be a morphological outlier (see “Results”), it was

excluded from both training datasets. Hence, a total of 22 different training datasets were tested to identify which resulted in predictions most consistent with the postures assigned to “known” bipeds and quadrupeds (see Supplementary Table S6). The consistency of each LDA with predefined assignments for “known” taxa was assessed using leave-one-out testing (jackknifing) of the training dataset and computing the total success rate. Subsequently, each LDA was then used to predict locomotor habit for the “other” taxa, for which locomotor habit is currently uncertain or controversial. It is worth noting that LDA will always classify a given subject into one of the available categories, regardless of how congruous it actually is with other data in the same category.

Initial analyses with additional models suggested that LDA may produce spurious results for immature specimens with morphologies (or even stance) that are qualitatively different from those of adults. As such, in addition to the original 13 “other” taxa used in the statistical analysis, two previously published models were included to explore the relevance of the LDA for assessing ontogenetic effects on locomotor mode: a juvenile *Alligator* (Bates et al. 2016) and a hatchling *Mussaurus* (Otero et al. 2019).

**Monte Carlo Simulation.**—Two of the primary variables used in this study, BM and  $COM_X$ , are estimates derived from computational volumetric models; as estimates they may therefore carry an attendant level of error from the “true” value (were it ever able to be known). In turn, error in the baseline dataset may affect the results obtained and the conclusions drawn from them. To explore the effect of error in BM and  $COM_X$  estimates, a Monte Carlo simulation was performed with 1000 replicates, wherein BM and  $COM_X$  were allowed to randomly vary up to a prescribed amount, and the pPCA and LDA were recomputed. Based on the results of Henderson (1999) and Allen et al. (2009), both BM and  $COM_X$  were allowed to vary by up to  $\pm 15\%$  of their original values.

## Results

**Patterns of Anatomical Variation.**—The results of pPCA differed only marginally between the



two phylogenies used (traditional vs. Ornithoscelida hypotheses of dinosaur interrelationships). Comparison of the PC scores between the two trees using symmetric Procrustes superimposition in the *vegan* package showed that the score sets were not significantly different (multivariate correlation coefficient = 0.999,  $p < 0.001$ ). Moreover, the relative difference between PC scores for a given taxon between the two phylogenetic hypotheses was on average 0.015% for PC 1, 0.357% for PC 2, 0.148% for PC 3, and 0.370% for PC 4; that is, the score a given taxon received on average differed by  $< 1\%$  between the two phylogenies used. Therefore, only the results derived from using the traditional phylogeny are presented herein. Despite the minimal difference in pPCA results between the two tree topologies, three of the four anatomical parameters showed a low to modest influence of phylogeny (BM:  $K = 0.57$ ,  $p = 0.001$ ; HL:  $K = 1.11$ ,  $p = 0.001$ ; FL:  $K = 0.39$ ,  $p = 0.001$ ). This indicated that the phylogenetic effect originated from at least one crownward part of the tree; because the difference between the two trees used occurred very deep within the phylogeny, the scope for phylogenetic effects in more crownward parts remained unaltered.

Each PC was loaded moderately to strongly by at least two anatomical parameters (Table 2); PC 1 accounted for 47.47% of the total data variance, PC 2 accounted for 25.61%, PC 3 accounted for 19.4%, and PC 4 accounted for 7.52% of the total variance. There was a very strong distinction between bipedal and quadrupedal taxa in PC space, both visually (Fig. 3, Supplementary Movie S1) and as indicated by PERMANOVA ( $F_{1,65} = 16.25$ ,  $p < 0.001$ ). Some of the 13 “other” taxa tended to bridge between the bipedal and quadrupedal

morphospaces in PC space, often plotting near the region of overlap between the two morphospaces, whereas others plotted either within one particular morphospace or outside both (Fig. 3, Supplementary Movie S1). Birds (except the giant, flightless *Struthio*) occupied their own distinct region of the bipedal morphospace, characterized by the lowest (most negative) PC 3 values; comparably low values were not present in any quadruped. *Trilophosaurus* was an outlier for the quadrupedal morphospace, its position in PC space—closer to bipeds than quadrupeds—being driven heavily by its caudal COM<sub>X</sub> position (Fig. 3, Supplementary Movie S1).

Procrustes disparity of the bipedal morphospace of PC scores was not significantly different from that of the quadrupeds (108.7 vs. 92.2, respectively;  $p = 0.49$ ), a result that remained unaltered when *Trilophosaurus* was excluded from consideration. Distilling this down to each PC, however, it was apparent that there were marked differences between bipeds and quadrupeds (Table 3). Quadrupeds showed more than double the variation in PC 1 than bipeds, a result that appears to have been driven in large part by ornithischians. Conversely, bipeds showed more than six times the variation in PC 3 than quadrupeds, a result that probably stemmed in large part from the high variance in theropods (including birds), given that theropods comprise the majority of bipeds in the current sample. Indeed, PC 3 was loaded most strongly by BM and COM<sub>X</sub> (Table 2), two parameters for which phylogenetically directed trends within Theropoda have been previously noted (Turner et al. 2007; Allen et al. 2013; Lee et al. 2014; Benson et al. 2018). Bipedal and quadrupedal morphospaces were well (but not completely) separated along the PC 2 axis (Fig. 3, Supplementary Movie S1), which was mostly loaded by COM<sub>X</sub> and hindlimb length (Table 2); this suggests that, broadly speaking, bipedal archosaurs have longer hindlimbs and a COM<sub>X</sub> located closer to the hips.

In terms of the anatomical parameters themselves, pANOVA indicated that bipeds and quadrupeds were statistically indistinct when each parameter was considered separately (cf. Table 4). However, more holistically, pMANOVA of the anatomical parameters showed

TABLE 2. Loadings of each anatomical parameter on the principal components resulting from phylogenetic principal components analysis (pPCA). BM, body mass; COM<sub>X</sub>, center of mass location cranial to the hips; HL, hindlimb length; FL, forelimb length.

| Parameter        | PC 1    | PC 2    | PC 3    | PC 4    |
|------------------|---------|---------|---------|---------|
| BM               | -0.6    | 0.2159  | 0.7703  | -0.007  |
| COM <sub>X</sub> | -0.5018 | 0.7722  | -0.3286 | -0.2097 |
| HL               | -0.7139 | -0.6108 | -0.1052 | -0.326  |
| FL               | -0.8817 | -0.0919 | -0.252  | 0.3881  |

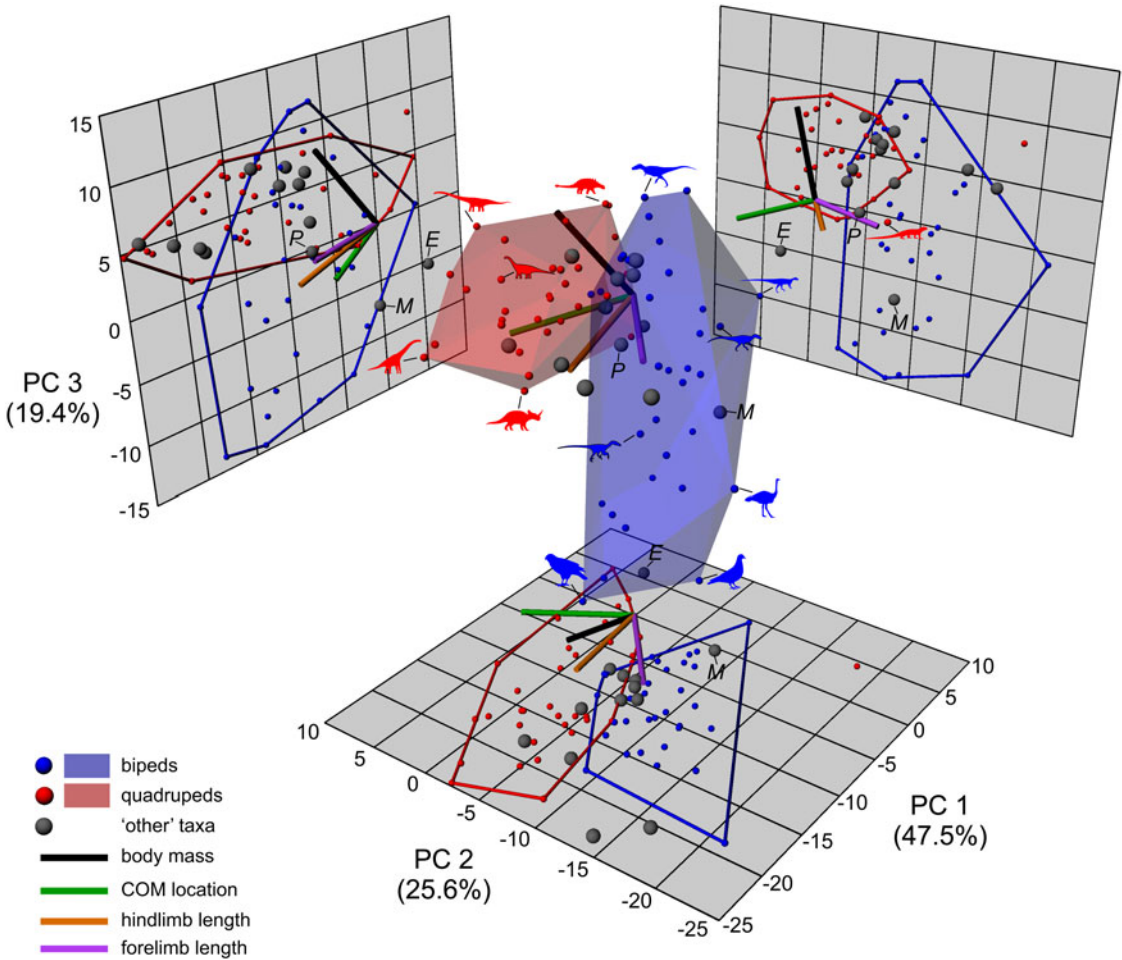


FIGURE 3. 3D plot of the first three principal component (PC) scores for each taxon, which collectively accounted for 92.5% of the variation in the dataset. The regions of space occupied by bipedal and quadrupedal taxa are delimited by convex hulls (generated using Meshlab 1.3.3; Cignoni et al. 2008), for visualization purposes only; the convex hull for quadrupeds was generated excluding the outlier *Trilophosaurus*. Also plotted are the loading vectors for each anatomical parameter, and silhouettes that illustrate body shapes at the extremes of the bipedal and quadrupedal morphospaces. The enigmatic taxa *Euparkeria* (E), *Postosuchus* (P) and *Marasuchus* (M) are also labeled. COM, center of mass.

that bipeds were distinct from quadrupeds ( $F_{1,65} = 44.74$ ,  $r^2 = 0.41$ ,  $p < 0.001$ ), and were almost twice as disparate morphologically

than quadrupeds (Procrustes variances of 2.38 and 1.22, respectively;  $p = 0.005$ ). Bipeds displayed more than double the variance in BM

TABLE 3. Means ( $\mu$ ) and variances ( $\sigma^2$ ) for each principal component (PC) score, parsed by both locomotor category and major clade.

|                 | PC 1    |            | PC 2   |            | PC 3   |            | PC 4   |            |
|-----------------|---------|------------|--------|------------|--------|------------|--------|------------|
|                 | $\mu$   | $\sigma^2$ | $\mu$  | $\sigma^2$ | $\mu$  | $\sigma^2$ | $\mu$  | $\sigma^2$ |
| Bipeds          | -8.965  | 25.205     | -8.779 | 13.496     | -0.798 | 60.747     | -5.017 | 12.648     |
| Quadrupeds      | -11.203 | 60.975     | -2.126 | 19.792     | 4.525  | 9.11       | -0.248 | 5.098      |
| Theropoda       | -9.983  | 18.083     | -8.961 | 15.077     | -0.74  | 69.085     | -5.375 | 13.097     |
| Sauropodomorpha | -15.39  | 21.642     | -3.37  | 12.117     | 6.279  | 2.701      | -0.549 | 4.763      |
| Ornithischia    | -11.02  | 56.288     | -5.234 | 19.717     | 3.203  | 10.642     | -2.153 | 4.457      |
| Pseudosuchia    | -3.887  | 15.273     | -0.96  | 6.353      | 0.864  | 3.819      | -1.666 | 5.262      |

TABLE 4. Means and variances for each anatomical parameter, parsed by both locomotor category and major clade. Note that for center of mass location cranial to the hips (COM<sub>X</sub>), hindlimb length (HL), and forelimb length (FL), these are the residuals derived from normalizing to body size (glenoacetabular distance).

|                 | log <sub>10</sub> (mass) |                | COM <sub>X</sub> |                | HL     |                | FL     |                |
|-----------------|--------------------------|----------------|------------------|----------------|--------|----------------|--------|----------------|
|                 | μ                        | σ <sup>2</sup> | μ                | σ <sup>2</sup> | μ      | σ <sup>2</sup> | μ      | σ <sup>2</sup> |
| Bipeds          | 1.573                    | 2.366          | -0.049           | 0.034          | 0.071  | 0.011          | -0.044 | 0.04           |
| Quadrupeds      | 3.177                    | 1.155          | 0.06             | 0.063          | -0.076 | 0.015          | 0.047  | 0.025          |
| Theropoda       | 1.679                    | 2.598          | -0.031           | 0.033          | 0.091  | 0.009          | -0.03  | 0.044          |
| Sauropodomorpha | 3.887                    | 0.302          | 0.062            | 0.025          | -0.009 | 0.005          | 0.118  | 0.019          |
| Ornithischia    | 2.747                    | 1.276          | -0.004           | 0.037          | -0.004 | 0.02           | 0.02   | 0.018          |
| Pseudosuchia    | 1.772                    | 0.323          | 0.067            | 0.004          | -0.15  | 0.01           | -0.114 | 0.006          |

than quadrupeds, a result again strongly driven by theropods (Table 4), whose range in absolute values of BM in the current dataset spans more than five orders of magnitude (compared with a little over three orders of magnitude across all quadrupeds).

The results for all pairwise comparisons of phylogenetically corrected anatomical values are reported in Table 5, and pairwise plots of phylogenetically reduced residuals are presented in Figure 4. In both bipeds and quadrupeds, FL was significantly and positively correlated with HL, although the relationships were statistically indistinguishable between the two groups; furthermore, it cannot be discounted that these results are not partly driven by spurious correlation with GA, as both variables were normalized with respect to this distance in deriving residuals. In addition, FL was significantly and positively correlated to COM<sub>X</sub> in bipeds; that is, a relatively more cranial COM location was tied to relatively longer forelimbs (cf. Allen et al. 2013). Unconventionally, COM<sub>X</sub> and BM were found to be positively correlated in bipeds, but the significance of this

trend was driven solely by the outlying residuals for *Heterodontosaurus* (Fig. 4).

*Linear Discriminant Analysis.*—Of the 22 variants in LDA tested, three of the four best-performing variants excluded birds from the training dataset and had success rates exceeding 98% (Supplementary Table S6). Indeed, excluding birds from the training dataset almost always resulted in an improvement in classification success rate. The single best LDA used COM<sub>X</sub> + HL + FL (“model 21”) as the combination of anatomical variables in the training dataset, with a classification success rate of 100% (Supplementary Table S6). Typically, the assignment of the training dataset taxa into bipedal or quadrupedal categories was relatively unambiguous; for instance, in model 21, more than 80% of the training taxa had posterior probabilities exceeding 0.95. These results further affirmed the strong distinction between bipeds and quadrupeds in the current sample. The most frequently misclassified training dataset taxa were the pseudosuchian *Riojasuchus* (misclassified as bipedal, 12 out of 22 times) and the sauropod *Neuquensaurus* (misclassified

TABLE 5. Results for pairwise comparison of the phylogenetically corrected values for each anatomical parameter by major axis regression. Statistically significant results are noted in boldface. \*When *Heterodontosaurus* is removed from consideration, the correlation degenerates to becoming nonsignificant (cf. Fig. 4). COM<sub>X</sub>, center of mass location cranial to the hips; BM, body mass; HL, hindlimb length; FL, forelimb length.

|                         | Bipeds  |                |               | Quadrupeds |                |               | Slope test, bipeds vs. quadrupeds |        |
|-------------------------|---------|----------------|---------------|------------|----------------|---------------|-----------------------------------|--------|
|                         | Slope   | r <sup>2</sup> | P             | Slope      | r <sup>2</sup> | P             | t                                 | P      |
| COM <sub>X</sub> vs. BM | 0.0814  | 0.2691         | <b>0.002*</b> | 0.098      | 0.0422         | 0.2441        | 0.0368                            | 0.8479 |
| HL vs. BM               | -0.0034 | 0.0015         | 0.8281        | 0.049      | 0.0937         | 0.0784        | 2.7236                            | 0.0989 |
| FL vs. BM               | 0.0414  | 0.1007         | 0.0719        | 0.0749     | 0.0908         | 0.0834        | 0.4912                            | 0.4834 |
| HL vs. COM <sub>X</sub> | 0.0113  | 0.0002         | 0.9382        | 0.013      | 0.0008         | 0.8731        | 0.0001                            | 0.9917 |
| FL vs. COM <sub>X</sub> | 0.6976  | 0.2497         | <b>0.0031</b> | 0.3259     | 0.1677         | 0.0162        | 2.315                             | 0.1281 |
| FL vs. HL               | 1.9758  | 0.3222         | <b>0.0006</b> | 1.8966     | 0.3812         | <b>0.0001</b> | 0.0139                            | 0.9061 |

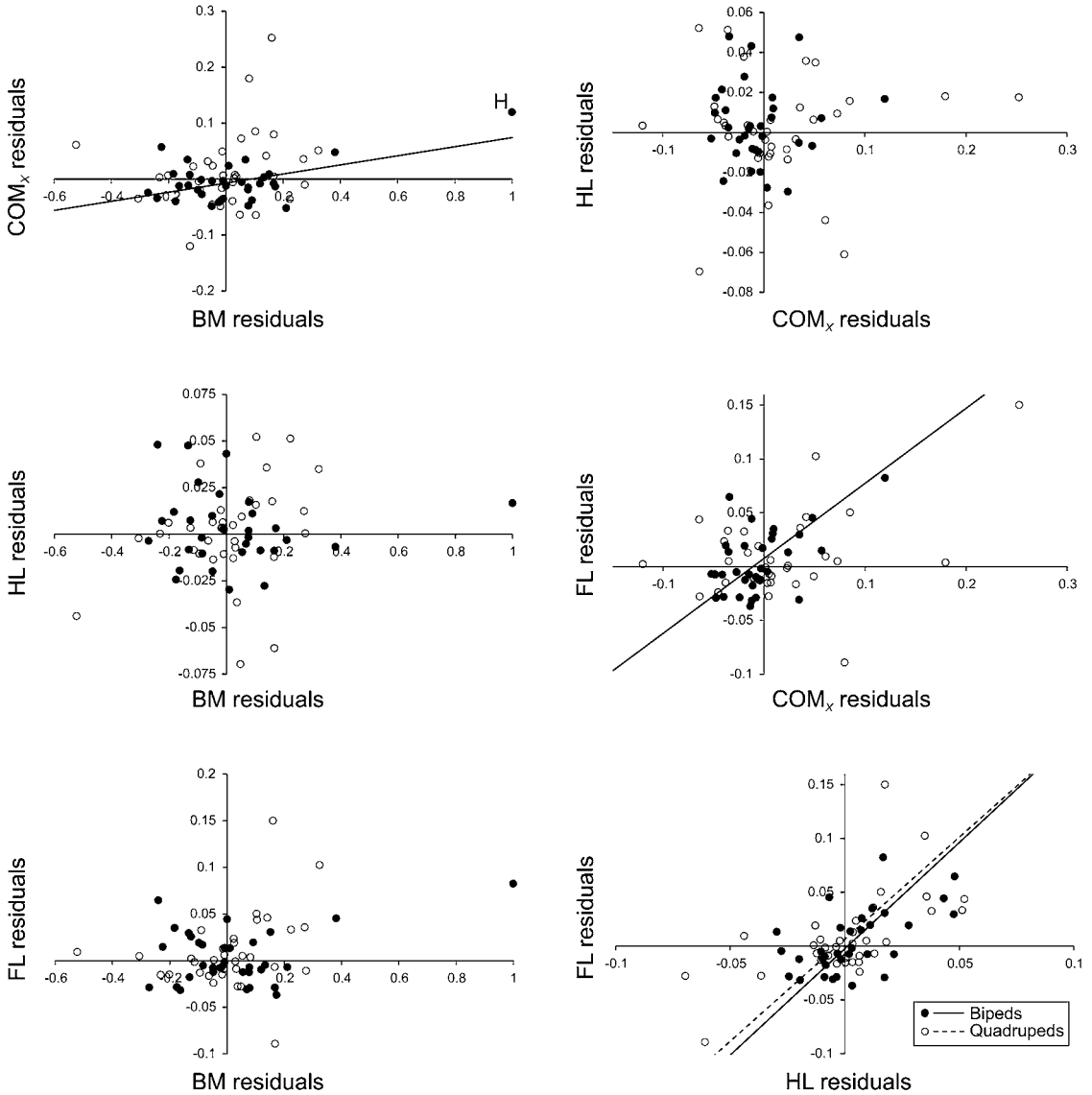


FIGURE 4. Pairwise comparisons of phylogenetically corrected anatomical parameters for both bipeds (solid circles) and quadrupeds (open circles). Where a statistically significant correlation was identified (via major axis regression), the regression line is also plotted; solid line for bipeds; dashed line for quadrupeds. Note that the significant correlation identified between mass and center of mass location cranial to the hips ( $COM_x$ ) for bipeds is driven by the residuals for *Heterodontosaurus* (indicated by “H”). BM, body mass; HL, hindlimb length; FL, forelimb length.

as bipedal, 11 out of 22 times), which had  $COM_x$  residuals among the lowest, if not the lowest, values for their respective clades (i.e., caudal  $COM_x$  location), potentially explaining their relatively frequent misclassifications. Both taxa are known from the majority of the skeleton (Otero 2010; von Baczko et al. 2019), suggesting underlying causes other than reconstruction error, such as apomorphic anatomies. For

instance, *Neuquensaurus* was quite small for a sauropod, and moreover it possessed a very broad pelvis, potentially resulting in a more caudal  $COM_x$  location.

The classifications assigned to the 15 “other” taxa for the four best-performing models are reported in Table 6. The classifications broadly corresponded to qualitative observations of how each taxon plots in PC space (Fig. 3,

TABLE 6. Classifications and posterior probabilities ( $Pr$ ) resulting from linear discriminant analysis applied to the 15 “other” taxa, for the four best-performing models. The correct classification rate on the training dataset is also given for each model.

| Taxon                        | Model 11 (98.48%) |       | Model 20 (98.28%) |       | Model 21 (100%) |       | Model 22 (98.28%) |       |
|------------------------------|-------------------|-------|-------------------|-------|-----------------|-------|-------------------|-------|
|                              | Prediction        | $Pr$  | Prediction        | $Pr$  | Prediction      | $Pr$  | Prediction        | $Pr$  |
| <i>Euparkeria</i>            | Quadruped         | 0.562 | Quadruped         | 0.821 | Quadruped       | 0.999 | Quadruped         | 0.981 |
| <i>Batrachotomus</i>         | Quadruped         | 0.884 | Quadruped         | 0.941 | Quadruped       | 0.955 | Quadruped         | 0.959 |
| <i>Postosuchus</i>           | Biped             | 0.984 | Biped             | 0.995 | Biped           | 0.847 | Biped             | 0.981 |
| <i>Alligator</i> (juvenile)  | Biped             | 0.94  | Biped             | 0.898 | Biped           | 0.525 | Biped             | 0.919 |
| <i>Marasuchus</i>            | Biped             | 0.999 | Biped             | 0.993 | Biped           | 0.584 | Biped             | 0.998 |
| <i>Muttaborrasaurus</i>      | Quadruped         | 0.987 | Quadruped         | 0.986 | Quadruped       | 0.767 | Quadruped         | 0.981 |
| <i>Tenontosaurus</i>         | Quadruped         | 0.973 | Quadruped         | 0.995 | Quadruped       | 0.834 | Quadruped         | 0.962 |
| <i>Iguanodon</i>             | Quadruped         | 0.975 | Quadruped         | 0.958 | Quadruped       | 0.979 | Quadruped         | 0.995 |
| <i>Lambeosaurus</i>          | Quadruped         | 0.823 | Quadruped         | 0.893 | Quadruped       | 0.848 | Quadruped         | 0.902 |
| <i>Gryposaurus</i>           | Biped             | 0.985 | Biped             | 0.944 | Biped           | 0.997 | Biped             | 0.999 |
| <i>Edmontosaurus</i>         | Biped             | 0.999 | Biped             | 1     | Biped           | 1     | Biped             | 1     |
| <i>Plateosaurus</i>          | Quadruped         | 0.696 | Quadruped         | 0.791 | Biped           | 0.677 | Biped             | 0.504 |
| <i>Lufengosaurus</i>         | Biped             | 0.982 | Biped             | 0.998 | Biped           | 0.999 | Biped             | 0.999 |
| <i>Mussaurus</i> (adult)     | Quadruped         | 0.583 | Quadruped         | 0.541 | Biped           | 0.829 | Biped             | 0.676 |
| <i>Mussaurus</i> (hatchling) | Biped             | 0.953 | Biped             | 0.738 | Quadruped       | 0.997 | Quadruped         | 0.637 |

Supplementary Movie S1). As an explicit “facultatively bipedal” category was not included, the posterior probabilities determined may give some tentative indication of how often bipedalism and quadrupedalism were used during normal activity. The pseudosuchians *Postosuchus* and *Batrachotomus* were strongly supported as a biped and quadruped, respectively. The enigmatic archosauriform *Euparkeria* was also classified as a quadruped, although in PC space it plotted well clear of the bipedal and quadrupedal morphospaces, and hence this result should be viewed with caution. In general concordance with a recent study (Otero et al. 2019), adult *Mussaurus* was frequently classified as a biped, whereas its hatchling was frequently classified as a quadruped (including in the best-performing model). The juvenile *Alligator* model was frequently (mis)classified as a biped (73% of LDA variants), whereas the jackknifed classifications of the adult *Crocodylus* models (quadrupedal) were frequently correct (70% of LDA variants).

**Monte Carlo Simulation.**—Simulation indicated that the results of pPCA were very robust to error in BM and  $COM_x$ . Of the 1000 replicates, not one resulted in a set of PC scores that was significantly different from (i.e., uncorrelated with) the scores obtained in the main analysis using the original estimates, as indicated by symmetric Procrustes superimposition ( $p < 0.001$ ). The first three PCs still accounted for 92% of the dataset’s variation on average

(mean  $\pm$  SD for each PC: PC 1 =  $47.1 \pm 2.2\%$ , PC 2 =  $25.9 \pm 1\%$ , PC 3 =  $19.4 \pm 2.3\%$ , PC 4 =  $7.6 \pm 0.4\%$ ). The average direction of the loading vectors across the 1000 replicates was within  $5^\circ$  of the original directions in the main analysis, and the angular standard deviation of the loading vectors’ directions in PC 1–3 space was  $17^\circ$  or less, indicating a narrow or acute “cone of variation” (cf. Fig. 3). Procrustes disparity ( $104.7 \pm 3.5$  for bipeds and  $90.9 \pm 3.1$  for quadrupeds,  $p > 0.5$ ) and PERMANOVA results ( $F_{1,65} = 15.8 \pm 0.6$ ,  $p < 0.001$ ) also did not differ significantly from the results of the main analysis.

In terms of the LDA, model 21 remained the best-performing combination of anatomical variables, with a classification success rate of  $98.53 \pm 1.33\%$ . The distribution of posterior probabilities for classification of the 15 “other” taxa were also highly consistent with the original probabilities obtained in the main analyses; Figure 5 illustrates the distributions for the model 21 combination. Nonetheless, the spread of probabilities for the juvenile *Alligator* and *Marasuchus* encompassed a considerable proportion of quadrupedal classifications, suggesting that the main analysis’ classifications for these two taxa should be viewed tentatively.

## Discussion

**Patterns of Anatomical Variation.**—By synthesizing data derived from 3D computational

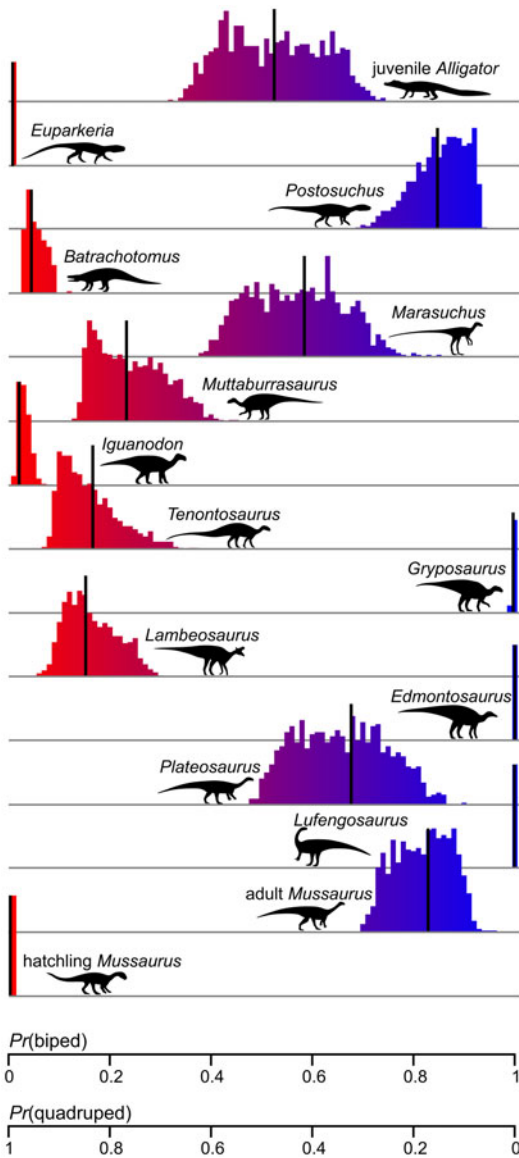


FIGURE 5. Distribution of posterior probabilities ( $Pr$ ) obtained for each of the 15 “other” taxa under model 21 linear discriminant analysis (LDA) across the 1000 replicates of the Monte Carlo simulation, binned into 100 intervals. Thick vertical lines denote probabilities obtained for each taxon in the main analysis using the original estimates for body mass (BM) and center of mass location cranial to the hips ( $COM_x$ ).

models, this study principally sought to examine how mass properties and gross body proportions relate to each other and locomotor habit—here taken to mean bipedal versus quadrupedal stance—in terrestrial archosaurs. The results of pPCA indicated that, despite concerns regarding

methodological bias in generating estimates, or potential error associated with this, bipedal and quadrupedal species were largely distinct from each other in morphospace (Fig. 3). Procrustes distance regression of all the original anatomical parameters themselves also recognized a clear distinction between the two groups. Yet few differences existed between bipeds and quadrupeds on an individual parameter-by-parameter basis (Table 4), and no PC was loaded primarily by just one anatomical parameter (Table 2). These findings highlight how a holistic, multivariate approach can facilitate more rigorous, quantitative assessment of locomotor habit in extinct archosaurs.

Following on from this, the single clearest distinction between bipedal and quadrupedal taxa revealed in the present study was bivariate (Fig. 3, Supplementary, Movie S1): overall, bipeds have relatively longer hindlimbs than quadrupeds (paralleling the finding of Kubo and Kubo 2012) and have a COM located closer to the hips. This result is hardly surprising, as biological and mechanical intuition would lead to an expectation of this pattern. From a biological perspective, longer hindlimbs reflect greater investment of biomass in them as locomotive organs, and therefore likely reflect greater reliance on the hindlimbs during daily activity. From a mechanical perspective, the COM must be kept over a “polygon of support” formed by the feet during static and dynamic activities (Alexander 2006; Herr and Popovic 2008; Winter 2009), the size of which is more restricted—and posteriorly positioned—in a biped. The size and mobility of the hindlimbs therefore constrains how far the COM is able to be located cranial to the hips in a biped (Hutchinson 2006; Gatesy et al. 2009), although postural differences can also modulate this relationship, as occurs in birds (Hutchinson and Allen 2009; Allen et al. 2013).

Quadrupedal archosaurs are (or were) probably also subject to their own form of locomotor constraint. Compared with bipeds, the current sample of quadrupeds exhibited substantially less variance in FL and a similar variance in HL (Table 4), suggesting that in order to coordinate stable, efficient progression, the dimensions of the limbs and intervening trunk in quadrupeds need to be more closely coupled

with one another (see also Thulborn 1982). The adoption of bipedality and removal of this constraint can therefore enable uncoupling of fore- and hindlimb locomotor modules (Gatesy and Dial 1996) and in turn promote greater evolutionary variability in limb size. This raises a possible evolutionary scenario within bipedal archosaurs, whereby relaxation of the “coupled proportions” constraint may have facilitated greater anatomical, functional, and perhaps ecological innovation, which in turn may have expedited the diversification of bipedal archosaurs across the Triassic/Jurassic boundary and into the later Mesozoic. For example, constraint relaxation should have been conferred by bipedality in the immediate ancestors of dinosaurs (Grinham et al. 2019). Such relaxation would then have presaged the uncoupling of forelimb and hindlimb allometries at the origin of birds, in association with the acquisition of powered flight (Dececchi and Larsson 2013), which subsequently led to phenotypic release in both limbs and increased rates of evolution in stem birds (Benson and Choiniere 2013).

*Predicting Locomotion in Extinct Archosaurs.*—

A secondary aim of this study was to develop a quantitative framework that could be used to help predict locomotor habit in extinct terrestrial archosaurs, and that bipeds and quadrupeds were largely distinct in PC space has facilitated this. Although beyond the scope of the present study, this new framework has the potential to provide new rigor to assessments of locomotor evolution within Archosauria, such as the frequency and phyletic distribution of shifts from quadrupedalism to bipedalism (facultative and obligate; Kubo and Kubo 2012; McPhee et al. 2018; Grinham et al. 2019; Chappelle et al. 2020). This would first require the generation of digital volumetric models for additional taxa, particularly those surrounding inferred transitions, which can be a nontrivial and time-intensive process.

Interestingly, it was found that excluding birds from the training dataset almost always resulted in improved classification accuracy. Birds occupied a distinct region of PC space in the current study, with strongly negative values along PC 3 reflecting their overall smaller BM (compared with both bipeds and

quadrupeds), and to a lesser extent their more cranial  $COM_x$  (compared with bipeds). It is possible that the tendency toward a more cranial  $COM_x$  makes birds more quadruped-like than most other nonavian bipedal archosaurs, impeding the ability of LDA to accurately separate out the two locomotor categories. As most birds are volant, and therefore not strictly terrestrial in their locomotor habit, the current study’s findings echo previous arguments that birds are not a good candidate as modern analogues for studying locomotion in Archosauria as a whole (e.g., Gatesy and Middleton 1997; Carrano 1998; Hutchinson and Gatesy 2000).

Many of the predictions of the best-performing LDAs in the current study concur with previous assessments, such as quadrupedality in *Batrachotomus* (Gower and Schoch 2009) and *Iguanodon* (Norman 1980), bipedality in *Lufengosaurus* (Bates et al. 2016; McPhee et al. 2018) and *Postosuchus* (Weinbaum 2013), and bipedality in adult *Mussaurus* but quadrupedality in its hatchlings (Otero et al. 2017, 2019). There are also conflicts with previous assessments, most notably the hadrosaurids *Gryposaurus* and *Edmontosaurus* being consistently classified here as bipeds. This is in contrast with the quadrupedal classification obtained for a third hadrosaurid, *Lambeosaurus*, despite the fact that all three taxa possess numerous osteological features suggestive of the use of quadrupedal stance (Dilkes 2001; Maidment and Barrett 2014), and reflected by many Late Cretaceous quadrupedal trackways attributed to hadrosaurids generally (Lockley and Wright 2001). What, then, might explain this discrepancy?

The discord surrounding hadrosaurids has been noted previously: on the basis of propodial proportions, the method of McPhee et al. (2018) was observed to have some difficulty with quadrupedal taxa that possess biped-like humeral proportions. Hadrosaurids tend to have gracile, “biped-like” forelimbs, and this may be responsible for the (potential) misclassifications obtained in the present study as well, which only used gross body proportions as a basis for classifications. In future studies, it would be worthwhile to combine the metrics used in the current study with those used by others, such as the proportions of individual

limb bones (McPhee et al. 2018) or the vertebral column (Christian and Preuschoft 1996), into a single framework. Moreover, as discrete osteological features have also frequently aided interpretations of locomotor habit (e.g., Mallison 2010; VanBuren and Bonnan 2013; Maidment and Barrett 2014), it would also be interesting to incorporate such discrete measures into a quantitative predictive framework.

One further misclassification of the LDAs was that the juvenile *Alligator* model was often misclassified as bipedal, yet there is no evidence of sustained, habitual, or facultative bipedalism in any extant Crocodylia. This result parallels a bipedal misclassification obtained for a juvenile *Caiman* by McPhee et al. (2018). It is known that extant crocodylians undergo substantial changes through ontogeny, in both cranial and postcranial proportions (Dodson 1975; Allen et al. 2009; Iijima and Kubo 2019). Ontogenetic effects may therefore distort relationships between individual bone proportions and mass,  $COM_X$  or other posturally relevant parameters (e.g., see Brassey et al. 2015). The present study did not aim to explicitly account for ontogenetic effects, and yet as noted earlier, the LDAs frequently detected an ontogenetic shift in locomotor mode in the early sauropodomorph *Mussaurus*, concordant with the interpretation of Otero et al. (2019). While the same computational models were used in the present study and that of Otero et al. (2019), they were analyzed and interpreted differently (one mechanistically, the other statistically), such that the concision of interpretations further supports a shift in locomotor mode as *Mussaurus* matured. Ontogenetic changes in locomotor mode have been proposed for other extinct terrestrial archosaurs (e.g., Heinrich et al. 1993; Dilkes 2001; Zhao et al. 2013; Słowiak et al. 2019; Chapelle et al. 2020), and it would be interesting to explore the applicability of the current study's approach to testing these interpretations in the future.

### Conclusion

By considering multiple parameters that describe whole-body mass properties and proportions, bipedal and quadrupedal archosaur

taxa were largely distinguished from each other. The results presented here should nevertheless be viewed tentatively, as it remains to be fully determined how methodological differences (in deriving estimates of BM and  $COM_X$ ) may influence the nature of the resulting comparative dataset, and hence any of the findings and interpretations made here. It is also important to recognize that the present study's dataset is biased toward two principal clades (Theropoda and Sauropodomorpha), and future sampling of additional disparate taxa (e.g., bipedal poposaurs, basal marginocephalians, and long-legged crocodylomorphs or silesaurids) may potentially reduce the degree of distinction between bipedal and quadrupedal taxa. Furthermore, the mean body mass of the present study's taxa was 3393 kg, with more than half being >200 kg, and greater sampling of small-bodied taxa could enrich insights on the relationships between body form, size, and locomotor habit. Despite these limitations, it is clear that a more holistic approach is required to fully understand locomotor behavior and evolution within archosaurs. Additionally, the findings of the present study suggest that important biomechanical constraints on the *Bauplan* of archosaurs (limb and trunk proportions) influenced the scope and extent of anatomical variation in both bipedal and quadrupedal species over their 250 Myr history.

### Acknowledgments

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P.J.B., V.R.A., and J.R.H. conceived the study design; P.J.B., K.T.B., V.R.A., D.M.H., and J.R.H. produced digital models and



contributed data; P.J.B. and M.R. conducted statistical analyses; all authors contributed to results interpretation and writing the article.

### Literature Cited

- Adams, D., M. Collyer, and A. Kaliontzopoulou. 2018. geomorph: geometric morphometric analyses of 2D/3D landmark data. <https://CRAN.R-project.org/package=geomorph>.
- Alexander, R. M. 1985. Body support, scaling, and allometry. Pp. 26–37 in M. Hildebrand, D. M. Bramble, K. F. Liem, and D. B. Wake, eds. *Functional vertebrate morphology*. Harvard University Press, Cambridge, Mass.
- Alexander, R. M. 2006. *Principles of animal locomotion*. Princeton University Press, Princeton, N.J.
- Alexander, R. M., and A. S. Jayes. 1983. A dynamic similarity hypothesis for the gaits of quadrupedal mammals. *Journal of Zoology* 201:135–152.
- Allen, V., H. Paxton, and J. R. Hutchinson. 2009. Variation in center of mass estimates for extant saurians and its importance for reconstructing inertial properties of extinct archosaurs. *Anatomical Record* 292:1442–1461.
- Allen, V., K. T. Bates, Z. Li, and J. R. Hutchinson. 2013. Linking the evolution of body shape and locomotor biomechanics in bird-line archosaurs. *Nature* 497:104–107.
- Bakker, R. T. 1971. Dinosaur physiology and the origin of mammals. *Evolution* 25:636–658.
- Baron, M. G., D. B. Norman, and P. M. Barrett. 2017. A new hypothesis of dinosaur relationships and early dinosaur evolution. *Nature* 543:501–506.
- Bates, K. T., and P. L. Falkingham. 2018. The importance of muscle architecture in biomechanical reconstructions of extinct animals: a case study using *Tyrannosaurus rex*. *Journal of Anatomy* 233:625–635.
- Bates, K. T., and E. R. Schachner. 2012. Disparity and convergence in bipedal archosaur locomotion. *Journal of the Royal Society Interface* 9:1339–1353.
- Bates, K. T., S. C. R. Maidment, M. R. Allen, and P. M. Barrett. 2012. Computational modelling of locomotor muscle moment arms in the basal dinosaur *Lesothosaurus diagnosticus*: assessing convergence between birds and basal ornithischians. *Journal of Anatomy* 220:212–232.
- Bates, K. T., P. D. Mannion, P. L. Falkingham, S. L. Brusatte, J. R. Hutchinson, A. Otero, W. I. Sellers, C. Sullivan, K. A. Stevens, and V. Allen. 2016. Temporal and phylogenetic evolution of the sauropod dinosaur body plan. *Royal Society Open Science* 3:150636.
- Benson, R. B. J., and J. N. Choiniere. 2013. Rates of dinosaur limb evolution provide evidence for exceptional radiation in Mesozoic birds. *Proceedings of the Royal Society of London B* 280:20131780.
- Benson, R. B. J., G. Hunt, M. T. Carrano, and N. E. Campione. 2018. Cope's rule and the adaptive landscape of dinosaur body size evolution. *Palaeontology* 61:13–48.
- Biewener, A. A. 1990. Biomechanics of mammalian terrestrial locomotion. *Science* 250:1097–1103.
- Blomberg, S. P., T. Garland, Jr, and A. R. Ives. 2003. Testing for phylogenetic signal in comparative data: behavioural traits are more labile. *Evolution* 57:717–745.
- Brassey, C. A., S. C. R. Maidment, and P. M. Barrett. 2015. Body mass estimates of an exceptionally complete *Stegosaurus* (Ornithischia: Thyreophora): comparing volumetric and linear bivariate mass estimation methods. *Biology Letters* 11:20140984.
- Butler, R. J., R. M. H. Smith, and D. B. Norman. 2007. A primitive ornithischian dinosaur from the Late Triassic of South Africa, and the early evolution and diversification of Ornithischia. *Proceedings of the Royal Society of London B* 274:2041–2046.
- Butler, R. J., J. Liyong, C. Jun, and P. Godefroit. 2011. The postcranial osteology and phylogenetic position of the small ornithischian dinosaur *Changchunsaurus parvus* from the Quantou Formation (Cretaceous: Aptian–Cenomanian) of Jilin Province, North-eastern China. *Palaeontology* 54:667–683.
- Butler, R. J., P. M. Barrett, and D. J. Gower. 2012. Reassessment of the evidence for postcranial skeletal pneumaticity in Triassic archosaurs, and the early evolution of the avian respiratory system. *PLOS ONE* 7:e34094.
- Campione, N. E., and D. C. Evans. 2012. A universal scaling relationship between body mass and proximal limb bone dimensions in quadrupedal terrestrial animals. *BMC Biology* 10:60.
- Campione, N. E., D. C. Evans, C. M. Brown, and M. T. Carrano. 2014. Body mass estimation in non-avian bipeds using a theoretical conversion to quadruped stylopodial proportions. *Methods in Ecology and Evolution* 5:913–923.
- Carrano, M. T. 1998. Locomotion in non-avian dinosaurs: integrating data from hindlimb kinematics, in vivo strains, and bone morphology. *Paleobiology* 24:450–469.
- Carrano, M. T. 1999. What, if anything, is a cursor? Categories versus continua for determining locomotor habit in mammals and dinosaurs. *Journal of Zoology* 247:29–42.
- Carrano, M. T. 2000. Homoplasy and the evolution of dinosaur locomotion. *Paleobiology* 26:489–512.
- Carrano, M. T., and S. D. Sampson. 2008. The Phylogeny of Ceratosauria (Dinosauria: Theropoda). *Journal of Systematic Palaeontology* 6:183–236.
- Carrano, M. T., R. B. J. Benson, and S. D. Sampson. 2012. The phylogeny of Tetanurae (Dinosauria: Theropoda). *Journal of Systematic Palaeontology* 10:211–300.
- Chapelle, K. E. J., R. B. J. Benson, J. Stiegler, A. Otero, and J. N. Choiniere. 2020. A quantitative method for inferring locomotor shifts in amniotes during ontogeny, its application to dinosaurs, and its bearing on the evolution of posture. *Palaeontology* 63:229–242.
- Charig, A. J. 1972. The evolution of the archosaur pelvis and hindlimb: and explanation in functional terms. Pp. 121–155 in K. A. Joysey and T. S. Kemp, eds. *Studies in vertebrate evolution*. Oliver & Boyd, Edinburgh.
- Christian, A., and H. Preuschoft. 1996. Deducing the body posture of extinct large vertebrates from the shape of the vertebral column. *Palaeontology* 39:801–812.
- Cignoni, P., M. Callieri, M. Corsini, M. Dellepiane, F. Ganovelli, and G. Ranzuglia. 2008. MeshLab: an open-source mesh processing tool. Pp. 129–136 in *Sixth Eurographics Italian Chapter Conference*, Salerno, Italy.
- Clauss, M., I. Nurutdinova, C. Meloro, H. C. Gunga, D. Jiang, J. Koller, B. Herkner, P. M. Sander, and O. Hellwich. 2017. Reconstruction of body cavity volume in terrestrial tetrapods. *Journal of Anatomy* 230:325–336.
- Dececchi, T. A., and H. C. E. Larsson. 2013. Body and limb size dissociation at the origin of birds: uncoupling allometric constraints across a macroevolutionary transition. *Evolution* 67:2741–2752.
- Dilkes, D. W. 2001. An ontogenetic perspective on locomotion in the Late Cretaceous dinosaur *Maiaasaura peeblesorum* (Ornithischia: Hadrosauridae). *Canadian Journal of Earth Sciences* 38:1205–1227.
- Dodson, P. 1975. Functional and ecological significance of relative growth in *Alligator*. *Journal of Zoology* 175:315–355.
- Ezcurra, M. D. 2016. The phylogenetic relationships of basal archsauromorphs, with an emphasis on the systematics of proterosaur archosauriforms. *PeerJ* 4:e1778.
- Farke, A. A., W. D. Maxwell, R. L. Cifelli, and M. J. Wedel. 2014. A ceratopsian dinosaur from the Lower Cretaceous of western North America, and the biogeography of Neoceratopsia. *PLoS ONE* 9:e112055.

- Garland, T., Jr., and C. M. Janis. 1993. Does metatarsal/femur ratio predict maximal running speed in cursorial mammals? *Journal of Zoology* 229:133–151.
- Gatesy, S. M., and A. A. Biewener. 1991. Bipedal locomotion: effects of speed, size and limb posture in birds and humans. *Journal of Zoology* 224:127–147.
- Gatesy, S. M., and K. P. Dial. 1996. Locomotor modules and the evolution of avian flight. *Evolution* 50:331–340.
- Gatesy, S. M., and K. M. Middleton. 1997. Bipedalism, flight, and the evolution of theropod locomotor diversity. *Journal of Vertebrate Paleontology* 17:308–329.
- Gatesy, S. M., and N. S. Pollard. 2011. Apples, oranges, and angles: comparative kinematic analysis of disparate limbs. *Journal of Theoretical Biology* 282:7–13.
- Gatesy, S. M., M. Bäker, and J. R. Hutchinson. 2009. Constraint-based exclusion of limb poses for reconstructing theropod dinosaur locomotion. *Journal of Vertebrate Paleontology* 29:535–544.
- Gower, D. J., and R. R. Schoch. 2009. Postcranial anatomy of the rauisuchian archosaur *Batrachotomus kupferzellensis*. *Journal of Vertebrate Paleontology* 29:103–122.
- Gregory, J. T. 1945. Osteology and relationships of *Trilophosaurus*. University of Texas Publications 4401:273–359.
- Grinham, L. R., C. S. VanBuren, and D. B. Norman. 2019. Testing for a facultative locomotor mode in the acquisition of archosaur bipedality. *Royal Society Open Science* 6:190569.
- Heinrich, R. E., C. B. Ruff, and D. B. Weishampel. 1993. Femoral ontogeny and locomotor biomechanics of *Dryosaurus lettowvorbecki* (Dinosauria, Iguanodontia). *Zoological Journal of the Linnean Society* 108:179–196.
- Henderson, D. M. 1999. Estimating the masses and centers of mass of extinct animals by 3-D mathematical slicing. *Paleobiology* 25:88–106.
- Henderson, D. M., and E. Snively. 2003. *Tyrannosaurus* en pointe: allometry minimized rotational inertia of large carnivorous dinosaurs. *Biology Letters* 271:S57–S60.
- Herr, H., and M. Popovic. 2008. Angular momentum in human walking. *Journal of Experimental Biology* 211:467–481.
- Hone, D. W. E., A. A. Farke, and M. J. Wedel. 2016. Ontogeny and the fossil record: what, if anything, is an adult dinosaur? *Biology Letters* 12:20150947.
- Hutchinson, J. R. 2006. The evolution of locomotion in archosaurs. *Comptes Rendus Palevol* 5:519–530.
- Hutchinson, J. R., and V. Allen. 2009. The evolutionary continuum of limb function from early theropods to birds. *Naturwissenschaften* 96:423–448.
- Hutchinson, J. R., and S. M. Gatesy. 2000. Adductors, abductors, and the evolution of archosaur locomotion. *Paleobiology* 26:734–751.
- Iijima, M., and T. Kubo. 2019. Allometric growth of limb and body proportions in crocodylians. *Journal of Zoology*.
- Jetz, W., G. H. Thomas, J. B. Joy, K. Hartmann, and A. O. Moores. 2012. The global diversity of birds in space and time. *Nature* 491:444–448.
- Kubo, T., and M. O. Kubo. 2012. Associated evolution of bipedality and cursoriality among Triassic archosaurs: a phylogenetically controlled evaluation. *Paleobiology* 38:474–485.
- Lacovara, K. J., M. C. Lamanna, L. M. Ibric, J. C. Poole, E. R. Schroeter, P. V. Ullman, K. K. Voegelé, Z. M. Boles, A. M. Carter, E. K. Fowler, V. M. Egerton, A. E. Moyer, C. L. Coughenour, J. P. Schein, J. D. Harris, R. D. Martínez, and F. E. Novas. 2014. A gigantic, exceptionally complete titanosaurian sauropod dinosaur from southern Patagonia, Argentina. *Scientific Reports* 4:6196.
- Lee, M. S. Y., A. Cau, D. Naish, and G. J. Dyke. 2014. Sustained miniaturization and anatomical innovation in the dinosaurian ancestors of birds. *Science* 345:562–566.
- Lockley, M. G., and J. L. Wright. 2001. Trackways of large quadrupedal ornithomorphs from the Cretaceous: a review. Pp. 428–442 in D. H. Tanke and K. Carpenter, eds. *Mesozoic vertebrate life*. Indiana University Press, Bloomington.
- Loewen, M. A., R. B. Irmis, J. J. W. Sertich, P. J. Currie, and S. D. Sampson. 2013. Tyrant dinosaur evolution tracks the rise and fall of Late Cretaceous oceans. *PLoS ONE* 8:e79420.
- Macaulay, S., J. R. Hutchinson, and K. T. Bates. 2017. A quantitative evaluation of physical and digital approaches to centre of mass estimation. *Journal of Anatomy* 231:758–775.
- Maidment, S. C. R., and P. M. Barrett. 2012. Does morphological convergence imply functional similarity? A test using the evolution of quadrupedalism in ornithischian dinosaurs. *Proceedings of the Royal Society of London B* 279:3765–3771.
- Maidment, S. C. R., and P. M. Barrett. 2014. Osteological correlates for quadrupedality in ornithischian dinosaurs. *Acta Palaeontologica Polonica* 59:53–70.
- Mallison, H. 2010. The digital *Plateosaurus* II: An assessment of the range of motion of the limbs and vertebral column and of previous reconstructions using a digital skeletal mount. *Acta Palaeontologica Polonica* 55:433–458.
- McDonald, A. T. 2012. Phylogeny of basal iguanodonts (Dinosauria: Ornithischia): an update. *PLoS ONE* 7:e36745.
- McPhee, B. W., R. B. J. Benson, J. Botha-Brink, E. M. Bordy, and J. N. Choiniere. 2018. A giant dinosaur from the earliest Jurassic of South Africa and the transition to quadrupedality in early sauropodomorphs. *Current Biology* 28:3143–3151.
- Nair, J. P., and S. W. Salisbury. 2012. New anatomical information on *Rhoetosaurus brownei* Longman, 1926, a gravisaurian sauropodomorph dinosaur from the Middle Jurassic of Queensland, Australia. *Journal of Vertebrate Paleontology* 32:369–394.
- Nesbitt, S. J. 2011. The early evolution of archosaurs: relationships and the origin of major clades. *Bulletin of the American Museum of Natural History* 352:1–292.
- Nesbitt, S. J., and M. A. Norell. 2006. Extreme convergence in the body plans of an early suchian (Archosauria) and ornithomimid dinosaurs (Theropoda). *Proceedings of the Royal Society of London B* 273:1045–1048.
- Norman, D. B. 1980. On the ornithischian dinosaur *Iguanodon bernisartensis* from the Lower Cretaceous of Bernisart (Belgique). *Mémoires Institut royal des Sciences naturelles de Belgique* 178:1–105.
- Oaks, J. R. 2011. A Time-calibrated species tree of Crocodylia reveals a recent radiation of the true crocodiles. *Evolution* 65:3285–3297.
- Oksanen, J., F. G. Blanchet, M. Friendly, R. Kindt, P. Legendre, D. McGlinn, P. R. Minchin, R. B. O'Hara, G. L. Simpson, P. Solymos, M. H. H. Stevens, E. Szoecs, and H. Wagner. 2019. *vegan*: community ecology package. <https://CRAN.R-project.org/package=vegan>.
- Orme, D., R. Freckleton, G. Thomas, T. Petzoldt, S. Fritz, N. Isaac, and W. Pearse. 2015. *caper*: comparative analyses of phylogenetics and evolution in R. <https://CRAN.R-project.org/package=caper>.
- Otero, A. 2010. The appendicular skeleton of *Neuquensaurus*, a Late Cretaceous saltasaurine sauropod from Patagonia, Argentina. *Acta Palaeontologica Polonica* 55:399–426.
- Otero, A., and D. Pol. 2013. Postcranial anatomy and phylogenetic relationships of *Mussaurus patagonicus* (Dinosauria, Sauropodomorpha). *Journal of Vertebrate Paleontology* 33:1138–1168.
- Otero, A., V. Allen, D. Pol, and J. R. Hutchinson. 2017. Forelimb muscle and joint actions in Archosauria: insights from *Crocodylus johnstoni* (Pseudosuchia) and *Mussaurus patagonicus* (Sauropodomorpha). *PeerJ* 5:e3976.
- Otero, A., A. R. Cuff, V. Allen, L. Sumner-Rooney, D. Pol, and J. R. Hutchinson. 2019. Ontogenetic changes in the body plan of the sauropodomorph dinosaur *Mussaurus patagonicus* reveal

- shifts of locomotor stance during growth. *Scientific Reports* 9:7614.
- Parrish, J. M. 1986. Locomotor adaptations in the hindlimb and pelvis of the Thecodontia. *Hunteria* 1:1–35.
- Prieto-Márquez, A. 2010. Global historical biogeography of hadrosaurid dinosaurs. *Zoological Journal of the Linnean Society* 159:503–525.
- R Core Team. 2012. R: a language and environment for statistical computing. R Foundation for Statistical Computing, Vienna.
- Revell, L. J. 2009. Size-correction and principal components for interspecific comparative studies. *Evolution* 63:3258–3268.
- Revell, L. J. 2012. phytools: an R package for phylogenetic comparative biology (and other things). *Methods in Ecology and Evolution* 3:217–223.
- Sellers, W. I. 2016. Animal Simulation Laboratory. <http://www.animalsimulation.org>, accessed 21 November 2019.
- Sellers, W. I., P. L. Manning, T. Lyson, K. A. Stevens, and L. Margetts. 2009. Virtual palaeontology: gait reconstruction of extinct vertebrates using high performance computing. *Palaeontologia Electronica* 12:11A.
- Sellers, W. I., J. Hepworth-Bell, P. L. Falkingham, K. T. Bates, C. A. Brassey, V. M. Egerton, and P. L. Manning. 2012. Minimum convex hull mass estimations of complete mounted skeletons. *Biology Letters* 8:842–845.
- Sereno, P. C. 1991. Basal archosaurs: phlogenetic relationships and functional implications. *Society of Vertebrate Paleontology Memoir* 2:1–53.
- Ślowiak, J., V. S. Tereschenko, and L. Fostowicz-Frelik. 2019. Appendicular skeleton of *Protoceratops andrewsi* (Dinosauria, Ornithischia): comparative morphology, ontogenetic changes, and the implications for non-ceratopsid ceratopsian locomotion. *PeerJ* 7:e7324.
- Sullivan, C. 2015. Evolution of hind limb posture in Triassic Archosauriforms. Pp. 107–124 in K. P. Dial, N. H. Shubin, and E. L. Brainerd, eds. *Great transformations in vertebrate evolution*. University of Chicago Press, Chicago.
- Thulborn, R. A. 1982. Speeds and gaits of dinosaurs. *Palaeogeography, Palaeoclimatology, Palaeoecology* 38:227–256.
- Thulborn, T. 1990. *Dinosaur tracks*. Chapman and Hall, London.
- Tschopp, E., O. Mateus, and R. B. J. Benson. 2015. A specimen-level phylogenetic analysis and taxonomic revision of Diplodocidae (Dinosauria, Sauropoda). *PeerJ* 3:e857.
- Turner, A. H., D. Pol, J. A. Clarke, G. M. Erickson, and M. A. Norell. 2007. A basal dromaeosaurid and size evolution preceding avian flight. *Science* 317:1378–1381.
- VanBuren, C. S., and M. F. Bonnan. 2013. Forearm posture and mobility in quadrupedal dinosaurs. *PLoS ONE* 8:74842.
- Venables, W. N., and B. D. Ripley. 2002. *Modern applied statistics with S*. Springer, New York.
- Vogel, S. 2003. *Comparative biomechanics: life's physical world*. Princeton University Press, Princeton, N.J.
- von Baczko, M. B., J. B. Desojo, and D. Ponce. 2019. Postcranial anatomy and osteoderm histology of *Riojasuchus tenuisiceps* and phylogenetic update on ornithosuchidae (Archosauria, Pseudosuchia). *Journal of Vertebrate Paleontology* 39:5.
- Walker, A. D. 1964. Triassic reptiles from the Elgin area: *Ornithosuchus* and the origin of carnosaurs. *Philosophical Transactions of the Royal Society of London B* 248:53–134.
- Warton, D. I., R. A. Duursma, D. S. Falster, and S. Taskinen. 2012. SMATR 3—an R package for estimation and inference about allometric lines. *Methods in Ecology and Evolution* 3:257–259.
- Weinbaum, J. C. 2013. Postcranial skeleton of *Postosuchus kirkpatricki* (Archosauria: Paracrocodylomorpha), from the Upper Triassic of the United States. Pp. 525–553 in S. J. Nesbitt, J. B. Desojo, and R. B. Irmis, eds. *Anatomy, phylogeny and palaeobiology of early archosaurs and their kin*. Geological Society of London, London.
- Winter, D. A. 2009. *Biomechanics and motor control of human movement*. Wiley, Hoboken, N.J.
- Witmer, L. M. 1995. The extant phylogenetic bracket and the importance of reconstructing soft tissues in fossils. Pp. 19–33 in J. J. Thomason, ed. *Functional morphology in vertebrate paleontology*. Cambridge University Press, Cambridge.
- Zhao, Q., M. J. Benton, C. Sullivan, P. M. Sander, and X. Xu. 2013. Histology and postural change during the growth of the ceratopsian dinosaur *Psittacosaurus lujiatunensis*. *Nature Communications* 4:2079.