Postcranial anatomy of the hyperodapedontine rhynchosaur Teyumbaita sulcognathus (Azevedo and Schultz, 1987) from the Late Triassic of Southern Brazil

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**INTRODUCTION**

_Teyumbaita sulcognathus_ was originally described by Azevedo and Schultz (1987) as _Scaphonyx sulcognathus_, and later transferred to a new genus by Montefeltro et al. (2010), based on a set of cranial autapomorphies that emphasized its highly modified morphology. The few postcranial elements of its holotype (UFRGS-PV-0232T) were described by Schultz (1986), but the skull anatomy was described and a new generic name was assigned to this highly autapomorphic taxon recently, when its skull anatomy was described and a new generic name was assigned to this highly autapomorphic taxon. Here, the postcranial skeleton of _Teyumbaita sulcognathus_ is for the first time fully described based on the holotype and the two more complete referred specimens. Rhynchosaur postcranial anatomy has usually been considered to be rather conservative, but _T. sulcognathus_ shows unforeseen morphological variation. Autapomorphic traits were added to the diagnosis of _T. sulcognathus_ and intraspecific variation was also identified. In addition, six new phylogenetically informative postcranial characters were recognized. Some of these represent apomorphies of clades such as Rhynchosauridae (axis with ventral keel, crest on the anteromedial surface of tibial shaft) and Hyperodapedontinae (postaxial cervical vertebrae with ventral keel, supinator process composed of a low supinator ridge and the ligament groove), revealing new support for their monophyly. The rhynchosaur diversity of the Otter Sandstone Formation (England) was also evaluated based on a phylogenetic analysis. The results suggest that the postcranial EXEMS 79/1992 is more likely related to _Fodonyx spenceri_ than to _Bentonyx sidensis_.

**POSTCRANIAL ANATOMY OF THE HYPERODAPEDONTINE RHYNCHOSAUR TEYUMBAITA SULCOGNATHUS (AZEVEDO AND SCHULTZ, 1987) FROM THE LATE TRIASSIC OF SOUTHERN BRAZIL**

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**ABSTRACT**—_Teyumbaita sulcognathus_ is a peculiar endemic Brazilian rhynchosaur that remained somewhat obscure until recently, when its skull anatomy was described and a new generic name was assigned to this highly autapomorphic taxon. Here, the postcranial skeleton of _Teyumbaita sulcognathus_ is for the first time fully described based on the holotype and the two more complete referred specimens. Rhynchosaur postcranial anatomy has usually been considered to be rather conservative, but _T. sulcognathus_ shows unforeseen morphological variation. Autapomorphic traits were added to the diagnosis of _T. sulcognathus_ and intraspecific variation was also identified. In addition, six new phylogenetically informative postcranial characters were recognized. Some of these represent apomorphies of clades such as Rhynchosauridae (axis with ventral keel, crest on the anteromedial surface of tibial shaft) and Hyperodapedontinae (postaxial cervical vertebrae with ventral keel, supinator process composed of a low supinator ridge and the ligament groove), revealing new support for their monophyly.

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**SUPPLEMENTAL DATA**—Supplemental materials are available for this article for free at www.tandfonline.com/UJVP.

**ARTICLE**

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arch; nc, neural channel; ns, neural spine; p, parapophyses; pa, popliteal area; pb, posterior boss; pc, posterior condyle; pp, postero-lateral peg; ppaxns, posterior process of axis neural spine; ptz, postzygapophysis; pz, prezygapophysis; ve, vertebra centrum; vk, ventral keel; sd, shaft depression; sg, superficial groove; sgb, supraglenoid buttress; sr, supinator ridge. tasc, trochanteric area for the insertion of M. subcoracoscapularis; tc, tibial crest; tp, transverse process.

SYSTEMATIC PALEONTOLOGY

DIAPSIDA Osborn, 1903
RHYNCHOSAURIA (Gervais, 1859) Osborn, 1903
HYPERODAPEDONTINAE Chatterjee, 1969 (nom. trans. ex Lydekker, 1885)

TEYUMBAITA Montefeltro, Langer, and Schultz, 2010
TEYUMBAITA SULCOCNATHUS (Azevedo and Schultz, 1987) (Figs. 1–18)

Diagnosis—Rhynchosaur with anterior margin of nasal concave at midline; prefrontal separated from the ascending process of the maxilla; palatal ramus of pterygoid expanded laterally within the palatines; markedly depressed dorsal surface of the exoccipital; and a single tooth lingually displaced from the main medial tooth-bearing area of the maxilla.

Appended Postcranial Diagnosis—Glenoid cavity formed mostly by the scapula; groove with elevated borders on the external surface of the entepicondyle; and transverse bar on the posterior surface of the astragalus extending from the lateral astragalar peg to the centrale articular facet (on the opposite side of the bone).

Description and Comparisons—The following description is based on the holotype (UFRGS-PV-0232T) and referred specimens (UFRGS-PV-0298T and PV-0290T) of Teyumbaita sulcognathus. The three specimens together have most of the postcranial elements preserved, although the holotype has the greater number of elements, including cervical, trunk, and caudal vertebrae, pectoral girdle, forelimbs, and hind limbs. UFRGS-PV-0298T has the axis and few trunk vertebrae, and UFRGS-PV-0290T has four trunk vertebrae and part of the left hind limb, including the astragalus. The postcranial remains were described following the osteological nomenclature employed by Romer (1966), Langer and Schultz (2000), and Whatley (2005). Previous studies suggested that rhynchosaurs had the forelimb in a partly sprawling position, but the hind limbs in a semiflexed position (Benton, 1983; 1990). Following this suggestion, the positional terms used in the forelimb description refer to a horizontally oriented humerus, whereas the hind limb elements are described vertically oriented. The postcranial remains of T. sulcognathus were compared within the context of total rhynchosaur diversity, although special attention was given to the morphology of taxa with more complete postcrania. Otischalkia eldæae Hunt and Lucas, 1991, was not used for the comparisons given that its rhynchosaurian affinity has been challenged (Long and Murry, 1995; S. Nesbitt and M. Stocker, pers. comm., March 2012). Clades mentioned in the description follow the definitions of Langer et al. (2010).

As previously described for the cranial remains of UFRGS-PV-0232T and PV-0298T (Montefeltro et al., 2010), the preserved postcranal bones are lightly built (Tables 1–4) in comparison with other hyperodapedontines (Huene, 1929, 1942; Sill, 1970; Chatterjee, 1974; Benton, 1983). As an example, the holotype femur has the ratio of the total length/shaft width of 6.9 and the same ratio in UFRGS-PV-0290T is 5.2. In contrast, UFRGS-PV-0290T is larger and more robust than the other specimens of T. sulcognathus.

Vertebral Column—The atlas-axis complex is virtually complete in the holotype, the atlantal intercentrum and neural arches are preserved separately (Fig. 1) from the remaining elements (atlantal centrum, axial centrum, and intercentrum), which are articulated with the next four neck vertebrae (Figs. 2, 3). In UFRGS-PV-0298T, only the axial neural arch is preserved but not fully prepared, preventing an accurate description.

The atlantal intercentrum is a dorsally concave and ventrally convex semicylindrical element. This morphology resembles the condition observed in Stenaulorhynchus stockleyi Haughton, 1932, and Hyperodapedon huxleyi Lydekker, 1885, but differs from the spatulate atlantal intercentrum of Hyperodapedon gordonii Huxley, 1859. The depth of the intercentrum dorsal concavity decreases posteriorly, and accounts for the greater robustness of the posterior articular facet. The latter is kidney-shaped in posterior view, and ventrally projected.

The unfused atlantal neural arches are about twice as tall as the intercentrum and are erect structures. Their medial surface is concave with a slight inwards curvature in the dorsal portion. The posterior margins of their convex ventral surfaces are slightly elevated relative to the anterior margins, and articulate with faint concavities on the dorsolateral margins of the intercentrum. As described for H. huxleyi (Chatterjee, 1974), the neural arches possess elliptical and obliquely oriented anterior surfaces that are continuous to the semicylindrical cranial facet of the intercentrum. These form the atlantal ring (Romer, 1956) that receives the occipital condyle.

![Image](image_url)
FIGURE 2. Teyumbaita sulcognathus (UFRGS-PV-0232T). Axis and following four cervical vertebrae. A, anterior; B, right lateral; C, left lateral; D, dorsal; E, ventral views. Scale bar equals 2 cm.
The dorsal half of each neural arch is triradiate, including the prezygapophysis anteriorly, the transverse process laterally, and the postzygapophysis posteriorly. The prezygapophysis projects anteromedially towards its counterpart. There is no medial contact, but each almost reaches the ventral portion of the elongated anterior process of the axial neural spine. The postzygapophysis has an anteroposteriorly and mediolaterally tapering proximal tip, and convex medial and lateral surfaces. An excavation immediately dorsal to the articular area forms a rudimentary infrayapophyseal fossa. The postzygapophysis corresponds to a posteroventrally and medially oriented process, the articular facet of which is straight, dorsoventrally elongated, and medioventrally directed. It articulates with the axial neural arch via a lateral excavation located between the transverse process and the anterior process of neural spine. The rod-like transverse process is posterolaterally and dorsally oriented. It is subtriangular in dorsal view, and its posterior margin does not reach the posterior-most limit of the postzygapophysis, from which the transverse process is separated by a short concave surface.

The posterior articular facet of the atlantal intercentrum has a beveled contact with the anterior surface of the axial intercentrum, and the space surrounded by the atlantal neural arches is occupied by the anteriorly projecting atlantal centrum. Because the articulation between the atlantal and axial intercentra is inclined, the dorsal margin of the former lies significantly above the latter. The atlantal centrum (odontoid process) is a very prominent anteriorly directed process, attached to the anteroventral surface of the axis of T. sulcognathus (Benton, 1983), the axial intercentrum of T. sulcognathus is not reduced, and is half the height of the axial centrum.

The axial centrum possesses an oval cross-section, with concave lateral walls, and thick anterior and posterior margins. The articular facets are vertically oriented, so that the centrum is subquadrangular in lateral view. As in postaxial cervical vertebrae, the ventral margin of the axis bears a conspicuous keel along its entire length, a feature also observed in Mesosuchus browni Watson, 1912, and Howesia browni Broom, 1906 (Dilkes, 1995, 1998). The three small bulbous processes present on the anteroventral surface of the axis of H. gordoni (Benton, 1983) are not present in T. sulcognathus or in other hyperodapedontines (Chatterjee, 1974; IGMPT-23, IML-3432, MCNSJ-680). The axial parapophyses apparently correspond to small rugose facets on the anterior middleheight of the lateral surfaces of the centrum. They are dorsally bound by the neural spine and anteroventrally by the contact area between the atlantal centrum and axial intercentrum.

The morphology of the axial neural spine of T. sulcognathus is variable. In the holotype, it bears anterior and dorsal processes, as seen in Rhynchosaurus articeps Owen, 1842 (Benton, 1990), whereas in UFRGS-PV-0298T it is fan-shaped, as in M. browni (Dilkes, 1998), S. stockleyi (Huene, 1938), H. gordoni (Benton, 1983), H. huxleyi (Chatterjee, 1974), and the South American Hyperodapedon (IGMPT-23, IML-3432, MCNSJ-680). That trait in the holotype was used to diagnose the taxon by Azevedo and Schultz (1987), but regarded by Schultz (1991) as a taphonomic artifact. We consider this an inherent feature, and suggest that the region between both processes was capped by cartilage, as proposed for R. articeps and H. gordoni (Benton, 1983, 1990). This variation may result from different ontogenetic stages, sexual dimorphism, or intraspecific variation.

The axial neural arch is twice as tall as the centrum, and the distance between the ventral margin of the transverse process and the base of the anterior process of neural spine accounts for half of its height. The short pyramidal transverse process projects lateroventrally, marking the anteroventral edge of the neural arch. From the dorsal margin of the transverse process, a
lamina projects posterodorsally, reaching the level of the prezygapophysis of the following vertebra.

The anterior process of the neural spine emerges from the anterodorsal margin of the neural arch as a triangular structure. Its dorsal margin is roughly horizontal, but the ventral margin forms an angle of about 40° with the longitudinal plane of the cervical column. The anterior margin of the process does not project further than the anterior margin of the axial intercentrum. The posterior process of the neural spine constitutes about one-third of the neural arch height and its base is much longer than its apex. It projects posterodorsally, with the anterior margin forming an angle of about 140° with the dorsal margin of the anterior process. The posterior margin of the posterior process is more vertical, and bears a dorsoventral sulcus for interspinous ligaments.

Anteroventral to the base of the neural spine, on the lateral margin of the pedicles surrounding the neural canal, a rugose, elliptical, and anteroposteriorly oriented surface corresponds to the articular facet for the atlantal postzygapophysis. The axial postzygapophysis is dorsoventrally compressed, mediodiagonally directed, and linked to the posteroventral margin of the neural spine by a thin lamina.

The presence of a distinct structure on the anterolateral surface of the axial centrum of T. sulcognathus, probably homologous to the parapophysis of the following cervical vertebrae, suggests the occurrence of an axial rib. This is also the case for S. stockleyi (Huene, 1938), the Brazilian Hyperodapedon (IGMPT-19.2; Huene, 1942), and M. browni Dilkes (1998), but not for H. gordoni (Benton, 1983), which seems to lack this rib.

The following four articulated vertebrae and an additional isolated cervical vertebra have similar morphology. Based on the position of the transverse process and the size and shape of the isolated vertebra, it probably represents the seventh cervical. The number of cervical vertebrae is variable in Rhynchosauria. Stenaulorrhynchos stockleyi possesses seven cervical and one transitional vertebra (Huene, 1938); M. browni (Dilkes, 1998); R. articeps (Benton, 1990); Isalorhynchus genovefae Buffetaut, 1983 (Whatley, 2005); H. gordoni (Benton, 1983), and H. luxleyi (Chatterjee, 1974) possess eight cervical vertebral.

By contrast, Huene (1942) suggests the presence of six cervical and one transitional vertebra in the Brazilian Hyperodapedon. The postaxial cervical vertebrae have subelliptical amphicoelous centra. The lateral walls are excavated, markedly on the mediodiagonal portion, forming a well-developed keel ventrally, as in Hyperodapedon (IGMPT-19.2 and -23; Huene, 1942; Chatterjee, 1974; Benton, 1983) and Isalorhynchus (Whatley, 2005). As in all rhynchosauroids, no intercentrum is present in the cervical series.

The transverse processes of cervical vertebrae 3–5 are subtriangular in lateral and dorsal aspects. Moving posteriorly along the preserved cervical series, the processes gradually occupy a more dorsal position on the neural arch. The transverse process of the sixth vertebra is distally rounded and stouter than those of the preceding vertebrae. As in Hyperodapedon (Huene, 1942; Chatterjee, 1974), the parapophyseal facets are not clear in T. sulcognathus. Yet, the position of that structure can be inferred from the laterally thickened anterior margin of the centrum, anteroventral to the transverse process in T. sulcognathus, these articular surfaces are dorsoventrally expanded, but their shape varies from subtrangular in the third cervical, to rounded in the sixth cervical.

The neural arches are partially preserved in the first four postaxial cervical vertebrae. In dorsal view, the prezygapophyses diverge to form an angle of 30° with each other, and the articular facets are nearly vertical. They cover the well-developed postzygapophyses of the preceding vertebra laterally, except the third cervical in which the prezygapophyses cover laterally the extremely reduced axial postzygapophyses at the base of the posterior process of the axis neural arch. The postzygapophyses have a subtriangular shape in lateral view, and their articular facets face laterally, also diverging at an angle of 30° in dorsal view.

The postzygapophyses of the third cervical vertebra possess a well-developed epipophysis on the left side. This structure is crest-shaped, dorsally tapering, and also observed in transverse sections of the postzygapophyses of the fourth and probable ninth presacral vertebrae (for discussion on rhynchosaurian cervical epipophyses, see Whatley, 2005).

Another isolated vertebra of the holotype (Fig. 4F–J) is the best-preserved trunk element of T. sulcognathus. Due to the position of the transverse process and the robustness of the centrum, it probably represents the eighth or ninth vertebra. It shares traits with the articulated and isolated more anterior cervical vertebrae and can be regarded as a ‘transitional’ element, as described for S. stockleyi and the Brazilian Hyperodapedon (Huene, 1938, 1942).

The amphicoelous centrum is rounded in cross-section, the lateral walls of which are slightly excavated. It lacks the typical ventral keel of the cervical vertebrae. On its dorsolateral portion, the neural arch transverse process expands as in the inferred seventh cervical vertebra. As in the cervical vertebrae, the parapophyseal region lies anteroventral to the transverse process. However, the parapophysis is less prominent and further displaced dorsally, showing the beginning of a trend towards convergence with the transverse process in the more posterior trunk vertebrae.

In dorsal view, the prezygapophyses diverge at an angle of 30°. Their dorsomedially oriented articular facets cover the postzygapophysis of the preceding vertebra ventrolaterally. Each postzygapophysis is subtriangular in lateral view, and forms an angle of 30° with the midline. The articular facet is lateroventrally oriented, with a dorsal epipophysis, as in the cervical vertebrae. This is not fully preserved, but the preserved portion suggests that it is less developed than in the cervical vertebrae.

The neural spine is a blade-like element, and is about one-half of the centrum height. It is located posterior to the centrum mid-length, on the posterodorsal region of the neural arch. In lateral view, this process is roughly semicircular in outline, with the anterior edge perpendicular to the main axis of the vertebral column. The anterior edge forms a continuous curve from the base to the dorsal tip. In posterior view, the neural spine possesses a longitudinal groove that is ventrally deeper and wider.

Four more posterior trunk vertebrae are preserved in the holotype. These show two distinct morphologies associated with their positions within the series, as seen in the complete vertebral series of Hyperodapedon (e.g., UFRGS-PV-0408T, IGMPT-19.2 and -23; Chatterjee, 1974; Benton, 1983). The preserved centra are all amphicoelous, and circular in cross-section. They have depressed lateral walls, but no ventral keel as those of H. gordoni (Benton, 1983). No neural arch is complete, but a sinuous suture marks the boundary with the centrum.

The transverse processes are more expanded laterally than in the preceding vertebrae, being located more posteriorly on the centrum. Indeed, the dorsal edges of the transverse processes are located above the level of neural channel roof, following the trend of dorsalization of that structure, already observed in more anterior vertebrae. In anterior view, the dorsal surface of the transverse processes forms a right angle with the neural arch. By contrast, its ventral surface is oblique to the lateral wall of the centrum, so the process has a subtriangular anterior outline (Fig. 5A–F). Conversely, in the two posterior-most preserved trunk vertebrae, the ventral surface is also perpendicular to the lateral surface of the centrum (Fig. 5G–L). The neural spines are damaged and no relevant information is available.

Four trunk vertebrae are preserved in UFRGS-PV-0290T (Fig. 6). Two are middle trunk vertebrae (Fig. 6D–I; trunks c and d of Table 1), whereas the remaining two elements (Fig. 6A–C, J–L) are assumed to be more anteriorly (trunk b of Table 1) and posteriorly (trunk e of Table 1) placed, respectively. Basically,
these share the same morphology as the holotype vertebrae, but their preservation does not allow a detailed comparison. One remarkable difference is their size (Table 1): the centra are about twice the length of other specimens, which is congruent with the size of the respective cranial material (Montefeltro et al., 2010). In addition, the transverse process of the more posterior trunk vertebra of UFRGS-PV-0290T is dorsoventrally narrower than those of the holotype, resulting in its bar-like shape in anterior view. Some trunk elements are also preserved in UFRGS-PV-0298T, but little information can be recovered due to lack of preparation.

There is a significant discrepancy in the estimation of the number of caudal vertebrae in rhynchosaurs varying from approximately 25 to as many as 50 based on relatively complete specimens (Chatterjee, 1974; Carroll, 1976; Benton, 1990; Dilkes, 1995, 1998). The exact number of caudal vertebrae of T. sulcognathus is unknown, but at least 16 elements are partially preserved in the holotype. It is not possible to determine whether the vertebrae form a continuous series. The weakly developed transverse processes suggest a distal position in the tail for all elements (Fig. 7). The centra are rounded in cross-section and are more amphicoelous than the trunk elements. In ventral view, they are as laterally compressed as in H. gordoni and H. huxleyi (Benton, 1983).

The tail neural arches are only partially preserved, but their general morphology is basically the same as that of the cervical and trunk vertebrae. The preserved transverse processes are located at the midheight of the vertebrae and are subtriangular in anterior and posterior views. Ventrally, the elliptical chevron facets are positioned on the posteroventral margin of the centrum.

Rib fragments are preserved in UFRGS-PV-0298T. Their transverse cross-sections are variably circular to elliptical. Two proximal rib portions are partially preserved, and their diaphalous morphology suggests that they are from either a cervical or anterior-most dorsal rib. The gastralia are also partially preserved in the holotype. They are fragile rod-like elements, frequently recognized only by their outlines on small blocks of sediment. These are circular to elliptical in cross-section, but the preservation does not allow recognition of the ‘V’-shaped pattern seen in other rhynchosaur gastralia (Benton, 1983, 1990; Dilkes, 1998).

Forelimb—The holotype includes a complete left and a partial right scapulocoracoid in which the dorsal half of the scapula blade is missing (Figs. 8, 9). Remarkable differences are observed between these elements. The left girdle was described together with the corresponding humerus by Schultz (1986), who proposed an unprecedented forelimb position for the taxon. However,
FIGURE 5. Teyumbaita sulcognathus (UFRGS-PV-0232T). Anterior (A–F) and posterior (G–L) trunk vertebrae. A, D, G, and J, anterior views; B, E, H, and K, left lateral views; C, F, I, and L, posterior views. Scale bar equals 2 cm.

FIGURE 6. Teyumbaita sulcognathus (UFRGS-PV-0290T). Anterior (A–C), median (D–I), and posterior (J–L) trunk vertebrae. A, D, G, and J, anterior views; B, E, H, and K, left lateral views; C, F, I, and L, posterior views. Scale bar equals 2 cm.
TABLE 1. Vertebra measurements (in cm) of *Teyumbaita sulcognathus*.

<table>
<thead>
<tr>
<th>Vertebra</th>
<th>Centrum length</th>
<th>Centrum height</th>
<th>Centrum width</th>
</tr>
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<tbody>
<tr>
<td>UFRGS-PV-0232T</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Axis</td>
<td>1.4</td>
<td>1.6</td>
<td>1.9</td>
</tr>
<tr>
<td>Axis</td>
<td>2.6</td>
<td>2.9</td>
<td>2.7</td>
</tr>
</tbody>
</table>
| Intercen
|              | length         |                |               |
| Intercen
|              | height         |                |               |
| Intercen
|              | width          |                |               |
| Atlas        | 2.6            | 3.4            | 4.0           |
| Anterior trunk a | 1.5          | 3.4            | 2.9           |
| Anterior trunk b | 3.2          | 3.4            | 2.4           |
| Posterior trunk a | 2.6          | 2.8            | 2.2           |
| Posterior trunk b | 2.2          | 2.8            | 2.3           |
| Caudal a      | 2.5            | 2.4            | 2.1           |
| Caudal b      | 2.2            | 2.6            | 1.5           |
| Caudal c      | 2.2            | 2.1            | 1.8           |
| Caudal d      | 2.1            | 2.5            | 1.6           |
| Caudal e      | 2.1            | 1.6            | 1.5           |
| Caudal f      | 2.1            | 2.1            | 1.5           |
| Caudal g      | 2.1            | 1.6            | 1.4           |
| Caudal h      | 2.0            | 1.1            | 1.0           |
| Caudal i      | 1.8            | 1.4            | 1.1           |
| Caudal j      | 1.8            | 1.2            | 1.1           |
| Caudal k      | 1.8            | 1.1            | 0.9           |
| Caudal l      | 1.7            | 1.0            | 1.1           |
| Caudal m      | 1.5            | 1.3            | 0.9           |
| Caudal n      | 1.4            | 0.9            | 0.9           |
| Caudal o      | 1.3            | 1.5            | 1.3           |
| Caudal p      | 1.1            | 0.8            | 0.8           |
| UFRGS-PV-0290 |                |                |               |
| Dorsal a      | 2.8            | 3.3            | 3.6           |
| Dorsal b      | 4.4            | 4.1            | 2.8           |
| Dorsal c      | 3.9            | 4.5            | 3.1           |
| Dorsal d      | 3.4            | 4.7            | 3.9           |

Caudal vertebrae a–p ordered by centrum length.

more detailed preparation revealed that the unique traits of the appendicular skeleton of UFRGS-PV-0298T result from taphonomic distortion (Schultz, 1991). As a consequence, the right scapulocoracoid is considered more reliable for reconstructing the original positions of the various structures. The differences between these elements are stressed in the following description.

Although similar in profile, the scapulocoracoid of *T. sulcognathus* is more slender than those of *Hyperodapedon* (Huene, 1942; Chatterjee, 1974; Benton, 1983) and *I. genovefae* (Whatley, 2005). Based on the left element, the scapula is twice the dorsoventral height of the coracoid (see Table 2). On the medial surface, the scapulocoracoid has a shallow dorsoventrally extending groove, spanning from the ventral one-third of the scapula blade to the dorsal margin of the coracoid foramen. The scapula blade has a gently convex dorsal margin, a sigmoid anterior margin, and a nearly straight posterior margin. The coracoid is semicircular in outline, and lacks the posterior process. The latter condition has been regarded as a hyperodapedontine synapomorphy (Benton, 1984, 1985, 1987, 1990; Dilkes, 1995; Langer and Schultz, 2000; Hone and Benton, 2008; Langer et al., 2010; Montefeltro et al., 2010), shared by *H. gordoni* (Benton, 2005), *H. gordoni* (Benton, 1983), *H. huxleyi* (Chatterjee, 1974), and *Hyperodapedon sanjuanensis* Sill, 1970 (MACN-18185). On the contrary, *M. browni, R. articeps, and S. stockleyi* possess a well-developed posterior process. In anterior view, both scapula and coracoid are mediolaterally concave, following the lateral contour of the ribcage. This curvature is more pronounced in the left scapulocoracoid, but this is a taphonomic distortion. The alleged capping of the entire scapulocoracoid by cartilaginous tissue (Benton, 1983, 1990) cannot be confirmed, but the texture of the dorsal margin of the scapula blade and the ventral margin of the coracoid suggest that these regions were covered by cartilage.

A blunt acromion process is present on the ventral one-third of the scapula anterior margin, dorsal to the level of the glenoid cavity. The acromion is stouter in the left scapula, and occupies a more lateral position, probably due to taphonomic distortions. The right acromion possesses an anterolateral position similar to that of other hyperodapedontines (Chatterjee, 1974; Benton, 1983; UFRGS-PV-0298T, MCNSJ-679, IMGPT-19.2).

The glenoid is deeply excavated at the posterior margin of the scapula-coracoid junction, and is bounded dorsally and ventrally by stout supra- and subglenoid buttresses. On the left side, the glenoid is posteriorly oriented, its borders are robust, and the scapula and coracoid portions form an angle of 90° to one

TABLE 2. Scapulocoracoid measurements (in cm) of *Teyumbaita sulcognathus*, UFRGS-PV-0232T.

<table>
<thead>
<tr>
<th>Dimension</th>
<th>Left</th>
<th>Right</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total height</td>
<td>19.3</td>
<td>—</td>
</tr>
<tr>
<td>Scapula height</td>
<td>14.1</td>
<td>—</td>
</tr>
<tr>
<td>Scapula dorsal length</td>
<td>5.8</td>
<td>—</td>
</tr>
<tr>
<td>Scapula ventral length (just dorsal to glenoid)</td>
<td>5.2</td>
<td>4.8</td>
</tr>
<tr>
<td>Coracoid height</td>
<td>5.2</td>
<td>6.3</td>
</tr>
<tr>
<td>Coracoid length (at ventral portion of glenoid)</td>
<td>6.7</td>
<td>7.5</td>
</tr>
</tbody>
</table>

—, not measurable.
another. On the right side, the cavity is lateroventrally directed, with smooth borders, and the articular facets diverging at an angle of 105°. As mentioned above, the right side seems to be less affected by taphonomic distortion. In both scapulae, the suture between the scapula and coracoid forms a straight line. It extends onto the glenoid cavity asymmetrically, with greater participation of the scapula. This condition differs from those seen in *M. browni* (Dilkes, 1998), *Rhynchosaurus articeps* (Benton, 1990), *S. stockleyi* (Huene, 1938), *I. genovefae* (Whatley, 2005), *H. gordoni* (Benton, 1983), and *H. huxleyi* (Chatterjee, 1974), and represents an autapomorphic trait of *T. sulcognathus*.

The blade-like coracoid is thicker posteriorly than anteriorly. An elongated coracoid foramen pierces its dorsal portion, but enters dorsally into the scapula on the left side. The position of this structure is highly variable among rhynchosaurs and a foramen entirely rimed by the coracoid (as in the right side of *T. sulcognathus*) is seen in *R. articeps* (Benton, 1990), *S. stockleyi* (Huene, 1938), and *H. huxleyi* (Chatterjee, 1974).

The left humerus of the holotype is nearly complete. In UFRGS-PV-0298T, the left humerus lacks the proximal region, and the right one has only its distal portion (Figs. 10, 11). The humerus/femur length ratio (see Tables 3 and 4) is almost 0.7, a plesiomorphic condition also observed in *M. browni* (Dilkes, 1998), *R. articeps* (Benton, 1990), *S. stockleyi* (Huene, 1938), *I. genovefae* (Whatley, 2005), and South American specimens of

<table>
<thead>
<tr>
<th>Specimen</th>
<th>Total length</th>
<th>Proximal width</th>
<th>Distal width</th>
<th>Medium shaft width</th>
</tr>
</thead>
<tbody>
<tr>
<td>UFRGS-PV-0232T</td>
<td>13.9</td>
<td>7.6</td>
<td>5.2</td>
<td>1.9</td>
</tr>
<tr>
<td>UFRGS-PV-0298T</td>
<td>14.8</td>
<td>8.5</td>
<td>—</td>
<td>2.8</td>
</tr>
</tbody>
</table>

—, not measurable.

<table>
<thead>
<tr>
<th>Femur/Tibia</th>
<th>Total length</th>
<th>Proximal width</th>
<th>Distal width</th>
</tr>
</thead>
<tbody>
<tr>
<td>UFRGS-PV-0232T</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Right femur</td>
<td>19.3</td>
<td>5.1</td>
<td>6.3</td>
</tr>
<tr>
<td>Right tibia</td>
<td>14.2</td>
<td>5.8</td>
<td>4.0</td>
</tr>
<tr>
<td>UFRGS-PV-0290T</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Right femur</td>
<td>21.6*</td>
<td></td>
<td>6.6</td>
</tr>
<tr>
<td>Right tibia</td>
<td>20.3</td>
<td>8.6</td>
<td>—</td>
</tr>
</tbody>
</table>

—, not measurable; *, not fully measurable.
Hyperodapedon (UFRGS-PV-0408T, IGMPT-23, MCNSJ-679, MCNSJ-680). In contrast, a ratio of about 1.0 is observed in H. gordonii and H. huxleyi (Benton, 1983, 1990; Dilkes, 1998; Langer and Schultz, 2000; Home and Benton, 2008).

The humerus of T. sulcognathus is more strongly expanded proximally than distally, with the shaft narrow and nearly cylindrical. In proximal or distal view of the holotypic bone, the long axis of the proximal margin (across the deltopectoral crest and the humeral head) is almost perpendicular to that across the distal condyles, a unique condition among rhynchosaurus. However, the left humerus of UFRGS-PV-0298T (Fig. 12) is more similar to that of other hyperodapedontines, and the holotype condition is probably a taphonomic artifact.

In proximal view, the humeral head of the holotype is anteriorly concave due to the anterior inflexion of the deltopectoral crest, forming an almost continuous curve. The shape of the deltopectoral crest is highly variable among rhynchosaurus. That of T. sulcognathus approaches the condition seen in R. articeps (Benton, 1990), I. genovefae (Whatley, 2005), and H. gordoni (Benton, 1983). Conversely, S. stockleyi (Huene, 1938), H. huxleyi (Chatterjee, 1974), and some South American Hyperodapedon specimens (UFRGS-PV-0408T, IGMPT-23, MCNSJ-574, MCNSJ-680, MCNSJ-679) this condition contrasts with that of M. browni (Dilkes, 1998), H. brownii (Dilkes, 1995), S. stockleyi (Huene, 1938), and H. gordonii (Benton, 1983) in which the intertrochanter is continuous with the ventral margin of the articular area. It tapers distally in anterior and posterior views, merging smoothly into the shaft in UFRGS-PV-0290T, but forming a sharper angle in the holotype. The internal trochanter forms the anterior margin of the intertrochanter fossa, which is more anteroposteriorly developed in the holotype, but broader dorsoventrally in the referred specimen. Posteriorly, the fossa is bounded by a faint crest that starts on the posterior surface of the proximal articulation and extends distally to a small tuberosity. This may correspond to the insertion area of part of the M. pubioischiofemoralis externus, as proposed for S. stockleyi (Huene, 1938) and R. articeps (Benton, 1990).

The distal femoral condyles of T. sulcognathus possess the usual rhynchosaurid traits, including the constricted cylindrical shaft and the slight sigmoidal curvature. The articulation areas are expanded, and the ratio between the width across the distal condyles and the length of the bone is greater than 0.3 (see Table 4), giving the femur a stout shape.

Although badly preserved, the capitellum can be partially recognized as a rugose area on the laterodistal portion of the fossa brachialis, as in H. gordonii (Benton, 1983). On the external surface of the ectepicondyle, the supinator process forms a thin crest and a relatively well-developed ligament groove, as also seen in H. sanjuanensis (MACN-18185), H. gordonii (Benton, 1983), and H. huxleyi (Chatterjee, 1974), and differing from the hook-shaped supinator process of S. stockleyi (Huene, 1938; Hunt and Lucas, 1991). Unique among rhynchosaurus, the entepicondyle of T. sulcognathus bears a well-developed lateral groove, flanked by elevated borders, that extends proximally to the proximal portion of the fossa brachialis.

The right femur of both the holotype and UFRGS-PV-0290T are preserved (Figs. 13, 14, 15). The femur of UFRGS-PV-0290T is larger, and the femur/tibia length ratio is significantly smaller than in the holotype (1.35 for the holotype and 1.06 for UFRGS-PV-0290T). The femur of T. sulcognathus possesses the usual rhynchosaurid traits, including the constricted cylindrical shaft and the slight sigmoidal curvature. The articulation areas are expanded, and the ratio between the width across the distal condyles and the length of the bone is greater than 0.3 (see Table 4), giving the femur a stout shape.

The proximal surface of the holotype femur is heavily worn and was probably covered by cartilaginous tissue in life. The well-developed internal trochanter corresponds to a proximodistally oriented crest that is stouter in the holotype than in UFRGS-PV-0290T. This structure is not continuous with the articular area, as also seen in H. huxleyi (Chatterjee, 1974), I. genovefae (Whatley, 2005), and some South American Hyperodapedon specimens (UFRGS-PV-0408T, IGMPT-23, MCNSJ-574, MCNSJ-680, MCNSJ-679) this condition contrasts with that of M. browni (Dilkes, 1998), H. brownii (Dilkes, 1995), S. stockleyi (Huene, 1938), and H. gordonii (Benton, 1983) in which the internal trochanter is continuous with the ventral margin of the articular area. It tapers distally in anterior and posterior views, merging smoothly into the shaft in UFRGS-PV-0290T, but forming a sharper angle in the holotype. The internal trochanter forms the anterior margin of the intertrochanter fossa, which is more anteroposteriorly developed in the holotype, but broader dorsoventrally in the referred specimen. Posteriorly, the fossa is bounded by a faint crest that starts on the posterior surface of the proximal articulation and extends distally to a small tuberosity. This may correspond to the insertion area of part of the M. pubioischiofemoralis externus, as proposed for S. stockleyi (Huene, 1938) and R. articeps (Benton, 1990).

The ventral surface of the femur has a robust adductor crest extending from the distal portion of the internal trochanter to the posterior portion of popliteal area, where it is much reduced. In UFRGS-PV-0290T, the proximal extension of the adductor crest reaches the boundary between the anterior and ventral surfaces of the femur, whereas in the holotype, the entire crest is visible in ventral view. The fourth trochanter is lacking in T. sulcognathus.

The distal femoral condyles of T. sulcognathus conceal the shallow, rhomboidal popliteal fossa. Although not fully preserved in any specimen, it can be inferred that the posterior condyle is more developed than the anterior. Additionally, their worn condition suggests a cartilage capping, as inferred for the other long bones. In the holotype, the distal condyles and the length of the bone is greater than 0.3 (see Table 4), giving the femur a stout shape.
portion less anteroposteriorly expanded. In the holotype, the proximal surface presents an anteroposteriorly directed groove, possibly representing the division of the tibial head for the reception of the femoral condyles, as suggested for *H. gordoni* (Benton, 1983) and *I. genovefae* (Whatley, 2005).

The tibial shaft is ovoid in cross-section and bowed medially away from the fibula, a condition more prominent in UFRGS-PV-0290T. Distal to the proximal head of both specimens, an evident tibial crest extends from the anterolateral to the anteromedial surfaces of the shaft, disappearing before reaching the distal articular end. This crest is absent in *M. browni* and *H. browni* (Dilkes, 1995, 1998), but present in *R. articeps* (Benton, 1983), *S. stockleyi* (Huene, 1938), *H. gordoni* (Hone and Benton, 2008), *H. huxleyi* (Chatterjee, 1974), and *I. genovefae* (Whatley, 2005). In addition, the crest is more prominent at its midlength, where a rounded depression related to the extensor musculature is seen. The elliptical distal end of the tibia is badly preserved in both specimens.

Only the proximal portion of the right fibula of UFRGS-PV-0290T is preserved. It is slender in comparison with the tibia, but is more robust than the fibula of *R. articeps* (Benton, 1990) and *S. stockleyi* (Huene, 1938), and more similar to that of *I. genovefae* (Whatley, 2005), *H. gordoni* (Benton, 1983), and *H. huxleyi* (Chatterjee, 1974). Its proximal margin is slightly expanded, and the articular facet is ovoid. On the proximal two-thirds of the incompletely preserved element (probably the proximal half of the bone), the fibula is laterally bowed, forming a large interosseous space with the tibia. In that portion, the shaft is twisted, as clearly indicated by a lateral crest that extends along the midlength of the preserved portion. The medial surface bears a less prominent crest, which starts opposite the lateral crest.

Only the right astragalus of UFRGS-PV-0290T is preserved. It has the typical features of Late Triassic rhynchosaurs, with four clear articular facets: a dorsomedial facet for the tibia, a dorsolateral facet for the fibula, a ventromedial facet for the...
FIGURE 11. *Teyumbaita sulcognathus* (UFRGS-PV-0232T). Schematic drawings of left humerus. A, anterior view; B, posterior view; C, proximal view; D, dorsal view; E, ventral view; F, distal view. Modified from Schultz (1986). Scale bar equals 3 cm.

centrale, and a ventrolateral facet for the calcaneum (Fig. 18). The element is not as robust as those of similar-sized South American *Hyperodapedon* specimens (e.g., UFRGS-PV-0247T, UFRGS-PV-0408T, MCNSJ-574).

Based on the articular facets of the astragalus, the proximal tarsals can be reconstructed as a row of three elements, with the centrale probably of equivalent size to the astragalus, as in other rhynchosaurs (Benton, 1984, 1985, 1987; Dilkes, 1995, 1998; Hone and Benton, 2008). Additionally, the astragalar facet for the centrale is the largest on the bone, as in other hyperodapedontines (Langer and Schultz, 2000; Langer et al., 2000a; Hone and Benton, 2008). The articular facets for the tibia and fibula are smaller and separated by a thin ridge, as described for

$H. \text{huxleyi}$ (Chatterjee, 1974). The articular facet for the calcaneum faces lateroventrally; but in contrast to *I. genovefae* (Langer et al., 2000b) and *Hyperodapedon* (UFRGS-PV-0247T), this is not ‘U’-shaped. It possesses an excavation bordering the astragalar peg distally, which Langer et al. (2000b) homologized with the astragalo-calcaneal canal of other basal archosauromorphs. However, this excavation is not as clear in *T. sulcognathus* as in *I. genovefae*. The posterior surface of the astragalus has a well-developed transverse boss spanning from the lateral astragalar peg to the medial region of the articular facet for the centrale. It forms a bar that is unique within Rhynchosauria and represents an autapomorphic trait of *T. sulcognathus*.

Three metapodials are preserved in UFRGS-PV-0298T. One is isolated, whereas the others are partially preserved articulated to phalanges. These share the general morphology of other rhynchosaur metapodials, which are rather alike in the fore- and hind limbs. Yet, they are assumed to represent metacarpals, because only elements of the forelimb are preserved in the specimen, but there is not sufficient information to assign them to specific digits. The elements are flat and elongated, with slightly expanded extremities, especially proximally. The shafts are slightly twisted distally counterclockwise, possessing shallow grooves along the middle of both dorsal and ventral surfaces.

The phalanges of UFRGS-PV-0298T form two digits with at least four elements each, probably two of the digits 3–5. The two isolated phalanges of UFRGS-PV-0232T and the nine of UFRGS-PV-0290T are more robust, and may represent pedal phalanges. The latter specimen has four phalanges forming a complete toe, whereas the remaining ones (including two unguals) are isolated. Despite their size discrepancy, the phalanges attributed to *T. sulcognathus* share the same general morphology. The more compact proximal elements have expanded extremities, proximal ends with biconcave facets, and extend beneath the following phalanx. Collateral depressions are seen on the ungual distal ends, indicating the presence of strong collateral ligaments.

The two ungual phalanges have distinct morphologies. One of them is stouter than the other, modestly curved, and has pointed distal tip, whereas the other is strongly compressed lateromedially. This variation can be attributed to different finger positions, because inner digits are usually more laterally compressed, and outer ones tend to have stouter unguals (Benton, 1983, 1990; Hone and Benton, 2008).
FIGURE 13. Teyumbaita sulcognathus (UFRGS-PV-0232). Right femur (A–F) and tibia (G–H). A, dorsal view; B, anterior view; C, ventral view; D, posterior view; E, proximal view; F, distal view; G, medial view; H, lateral view. Scale bar equals 3 cm.

DISCUSSION

In contrast to preview works (Benton, 1983; Langer and Schultz, 2000; Langer et al., 2000b; Montefeltro et al., 2010), the present paper does not support the idea that rhynchosaur all shared a conservative postcranial morphology. In fact, there are important variations in these elements, including autapomorphic characters that can be added to the diagnosis of *T. sulcognathus*. Characters varying among the three specimens of the taxon (e.g., differences in the axial neural spine morphology, the femur/tibia length ratio, and the origin of the femoral internal trochanter) show that postcranial morphology is also variable intraspecifically. In addition, some characters are potentially useful for phylogenetic analyses, which are tested according to the parameters described below.

Phylogenetic Analysis

Six potentially informative characters were added to the data matrix of Langer et al. (2010) in order to test their effect on rhynchosaur phylogeny (see Appendix 1). The resulting data matrix (Appendix 2 and online Supplemental Data) was analyzed using TNT version 1.1 (Goloboff et al., 2008) using the implicit enumeration algorithm. *M. browni* was constrained as the basalmost taxon and *H. browni* as the sister group of Rhynchosauridae sensu Langer et al. (2010). The latter is formed by eight taxa (see Appendix 3), as in the ‘second analysis’ of Langer et al. (2010).
Two most parsimonious trees (MPTs) of 104 steps were recovered, depicting the same structure as that of Langer et al. (2010) (Fig. 19). The only unresolved relationship concerns the position of *Bentonyx sidensis* Langer, Montefeltro, Hone, Whatley, and Schultz, 2010, as the sister taxon of either *Fodonyx spenceri* (Benton, 1990) plus Hyperodapedontinae or *S. stockleyi* plus the ‘Mariante rhynchosaur.’ The inclusion of the newly proposed postcranial characters did not change the previously recovered topology, but two new rhynchosaurid apomorphies (axis with a ventral keel, crest on the anteromedial region of tibial shaft) as well as two delayed synapomorphies for Hyperodapedontinae (postaxial cervical vertebrae with ventral keel, supinator process formed by a low supinator ridge and the ligament groove) were recognized.

A recently discussed issue in rhynchosaur taxonomy concerns the identification of two different genera in the Anisian-aged Otter Sandstone Formation of south Devon, England (Hone and Benton, 2008; Langer et al., 2010). The presence of both *F. spenceri* and *B. sidensis* in the same stratigraphic unit precludes an unambiguous referral of the postcranial specimen EX-EMS 79/1992 to either of the genera (Langer et al., 2010). An exploratory analysis was conducted in order to elucidate...
the affinities of EXEMS 79/1992. It was included as a discrete taxonomic unit in the matrix and the same search protocol was employed. This resulted in 12 MPTs (of 104 steps). The strict consensus tree (Fig. 19) has a less resolved topology, with a highly polytomic Hyperodapedontidae. Yet, when the MPTs are considered separately, a clear pattern emerges. In all trees, EXEMS 79/1992 is placed on the branch leading to Hyperodapedontinae, where it can assume any of the possible positions. In addition, EXEMS 79/1992 is never associated with B. sidensis, even in trees in which the latter has a pro-Hyperodapedontinae placement. Only four characters could be scored for EXEMS 79/1992, and clearly further inquiry and more complete specimens are required. Yet, the present results suggest its association with F. spenceri, rather than B. sidensis.

FIGURE 18. Teyumbaita sulcognathus (UFRGS-PV-0290T). Right astragalus. A, anterior view; B, posterior view; C, proximal view. Scale bar equals 2 cm.

FIGURE 19. Phylogenetic relations of Rhynchosauria. A, strict consensus of the two most parsimonious trees (MPTs) recovered in the analysis including six new postcranial characters; B, strict consensus of the 12 MPTs recovered in the analysis including EXEMS 79/1992. Bootstrap (1000 replicates) and ‘Bremer support’ values of each node are indicated. Arrows designate stem-based taxa according to Langer et al. (2010).

CONCLUSIONS

The comparative study presented here reveals that the postcranial morphology of rhynchosaurs is not as conservative as thought previously (Benton, 1983; Langer and Schultz, 2000; Langer et al., 2000a; Nesbitt and Whatley, 2004). Different levels of variation were recognized, from intraspecific to those defining rhynchosaur internal relationships. This variation in postcranial morphology offers new information on the phylogenetic relationships of rhynchosaurs and emphasizes the importance of considering postcranial data in phylogenetic studies of the group. This corroborates the hyperodapedontine affinity of T. sulcognathus, and also suggests referral of the postcranial specimen EXEMS 79/1992 to that lineage, most likely to F. spenceri.

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LITERATURE CITED


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APPENDIX 1. Character descriptions. The 65 characters used in the phylogenetic analyses are described below (along with the character states). The character-taxon matrix is presented in Appendix 2. Characters 36, 43, and 47 were treated as additive. Characters are either new or adapted from previously published
analyses (acknowledged accordingly). New postcranial characters are at the end of the list (characters 60–65).

1. Skull dimensions: longer than broad (0); broader than long (1) (Benton, 1984).
2. Skull height: <50% of the midline length (0); >50% of the midline length (Hone and Benton, 2008).
3. Orbit orientation: mostly lateral (0); mostly dorsal (1) (Langer and Schultz, 2000).
4. Orbital medial margin: rounded (0); forming a marked angle (1) (Montefeltro et al., 2010).
5. Jugal and maxillary heights below the orbit ventral margin: maxilla higher (0); jugal higher (1) (Benton, 1984).
6. Jugal-lacrimal contact: minimal (0); extensive contact of the jugal anterior process (1) (Whatley, 2005).
7. Jugal lateral surface: anguli oris crest does not reach jugal anterior process (0); anguli oris crest reaches jugal anterior process (1) (Benton, 1984).
8. Jugal surface dorsal to anguli oris crest: lacking a secondary crest (0); with a secondary anguli oris crest (1) (Langer and Schultz, 2000).
9. Lateral overlap of maxilla by jugal: absent or minimally expanded (0); well developed (1) (Whatley, 2005).
10. Jugal subtemporal processes: squamosal ventral process anterior to quadratojugal (0); squamosal ventral process anterior to quadratojugal and overlapping the quadrate (1) (Montefeltro et al., 2010).
11. Postorbital anteroventral process: expanding ventral to the jugal (0); fits dorsal to the jugal (1) (Whitley, 2005).
12. Postorbital-parietal suture: visible in dorsal view (0); hidden in dorsal view (1) (Dilkes, 1998).
13. Parietal body: not expanded laterally at midlength (0); expanded laterally at midlength (1) (Montefeltro et al., 2010).
14. Parietal transverse process: posterolaterally directed (0); laterally directed (1) (Montefeltro et al., 2010).
15. Distal tip of parietal transverse process: not anteriorly curved (0); anteriorly curved (1) (Montefeltro et al., 2010).
16. Squamosal ventral process: thinner than 50% of dorsoventral length (0); broader than over 50% of dorsoventral length (1) (Benton, 1990).
17. Relative position of quadratojugal and squamosal processes: squamosal ventral process anterior to quadratojugal dorsal process (0); squamosal ventral process overlapping the quadrate (1) (Montefeltro et al., 2010).
18. Ventral margin of opisthotic paroccipital process: convex (0); straight (1) (Montefeltro et al., 2010).
19. Pterygoid midline suture length: greater than or equal to the distance between the posterior margin of the suture and the basiptygoid articulation (0); less than the distance between the posterior margin of the suture and the basiptygoid articulation (1) (Whatley, 2005).
20. Elements forming the border of the suborbital fenestra: ectopterygoid, palatine, and maxilla (0); ectopterygoid and palatine only (1) (Dilkes, 1995).
21. Occipital condyle position: anterior to craniomandibular articulation (0); aligned to craniomandibular articulation (1) (Benton, 1984).
22. Basiooccipital and basiphenoid/parasphenoid lengths: basiphenoid/parasphenoid longer (0); basiooccipital longer (1) (Langer and Schultz, 2000).
23. Relative positions of the basiptygoid process of the basiphenoid and the ectopterygoid at the same level (0), basiptygoid process of the basiphenoid posterior to ectopterygoid process of the pterygoid (1) (Dilkes, 1995).
24. Basiptygoid process dimensions (dorsoventral length, anteroposterior width): longer than wide (0); wider than long (1) (Langer and Schultz, 2000).
25. Mandible height: <0.25 of the total length (0); >0.25 of the total length (1) (Benton, 1984).
26. Dentary length: half, or less, than the total mandibular length (0); greater than half of the total mandibular length (1) (Benton, 1990).
27. Medial maxillary groove: absent (0); present but not reaching the anterior half of the maxilla (1); present and reaching the anterior half of the maxilla (2) (Benton, 1984).
28. Maxillary area lateral to main groove: narrower than the medial area (0); same width or broader than the medial area (1) (Benton, 1990).
29. Maxillary cross-section lateral to main groove: crest-shaped (0); cushion-shaped (1) (Langer et al., 2000b).
30. Tooth rows lateral to main maxillary groove: a single clear row (0); more than one clear row (1) (Langer and Schultz, 2000).
31. Number of tooth rows medial to main maxillary groove: two rows and scattered teeth (0); three or more tooth rows (1) (Langer et al., 2000b).
32. Occlusal tooth rows on the anterior half of the maxilla: four or more tooth rows (0); fewer than four tooth rows (1) (Whatley, 2005).
33. Maxillary lingual teeth: absent (0); scattered teeth (1); large number of teeth on the medial surface of the bone (2) (Benton, 1984).
34. Maxillary teeth: only conicals (0); conicals and ‘pyramidal’ (1) (Whatley, 2005).
35. Dentary teeth: only conicals (0); conical and anteroposteriorly compressed (1) (Whatley, 2005).
36. Posterior-most dentary teeth: on the anterior half of dentary (0); on the posterior half of dentary (1) (Langer and Schultz, 2000).
37. Lingual dentary teeth: absent (0); present and forming one row (1); present and forming more than one row (2) (Benton, 1984).
38. Truncal vertebrae with ossified intercentrum: present (0); absent (1) (Evans, 1988).
39. Epipophyses on cervical postzygapophyses: spine-shaped (0); crest-shaped (1) (Whatley, 2005).
40. Second sacral vertebra: with a notch between the iliac articulation and the posterior process (0); posterior process continuous to the iliac articular surface (1) (Dilkes, 1998).
41. Caudal vertebrae neural spines: height twice the length (0); height less than twice the length (1) (Dilkes, 1998).
42. Interclavicle: posterior process longer than twice the lateral processes (0); posterior process shorter than twice the lateral process (1) (Dilkes, 1998).
APPENDIX 2. Character-state matrix. Missing data are marked as ‘?’, nonapplicable characters as ‘-‘, and variable condition under ‘[ ]’.

<table>
<thead>
<tr>
<th>Character</th>
<th>Mesosuchus browni</th>
<th>Howesia browni</th>
<th>Rhynchosaurus spp.</th>
<th>Stenaulorhynchus stockleyi</th>
<th>‘Mariante rhynchosaur’</th>
<th>Bentonyx sidensis</th>
<th>Fodonyx spenceri</th>
<th>Isalorhynchus genovefae</th>
<th>T. sulcognathus</th>
<th>Hyperodapedon spp.</th>
<th>EXEMS 79/1992</th>
</tr>
</thead>
<tbody>
<tr>
<td>Posterior process of the coracoid: present (0); absent (1) (Benton, 1984).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Dorsal margin of the ilium: anterior process &lt;15% of the length of the posterior process (0); anterior process &gt;15% of the length of the posterior process (1) (Dilkes, 1995).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Internal trochanter: continuous with the femoral head (0); separated from femoral head (1) (Whatley, 2005).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Relative size of astragalar articular facets: tibial facet greater than centrale facet (0); centrale facet greater than tibial facet (1) (Langer and Schultz, 2000).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Metatarsal I: longer than broad (0); broader than long (1) (Hone and Benton, 2008).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Axis ventral keel: present (0); absent (1).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Axial parapophysis: present (0); absent (1).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Cervical postaxial vertebrae ventral keel: absent (0); present (1).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Position of the transverse process of cranial truncal vertebrae: at the level of prezygapophysis (0); posteriorly located in the vertebra centrum (1).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Supinator process on the external surface of humeral ectepicondyle: absent (0); present and hook-shaped (1); present and formed by a low supinator ridge and ligament groove (2).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Crest on anteromedial region of tibial shaft: absent (0); present (1).</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

APPENDIX 3. Operational taxonomic units. Ingroup and outgroup operational taxonomic units used in the phylogenetic analysis. Sources of data for coding are listed for each taxon (specimens studied firsthand and published descriptive accounts).

**Outgroup**


**Ingroup**


Stenaulorhynchus stockleyi Haughton, 1932 (IGMPT-317; Huene, 1938).

‘Mariante rhynchosaur’—Taxon not formally described; see Schultz and Azevedo (1990) (UFRGS-PV-0168T, UFRGS-PV-0315T).

Bentonyx sidensis Langer, Montefeltro, Hone, and Schultz, 2010 (BRSUG 27200; Hone and Benton, 2008; Langer et al., 2010).


Isalorhynchus genovefae Buffetaut, 1983 (Whatley, 2005).

Hyperodapedon spp.—H. gordoni Huxley, 1859; H. huxleyi Lydekker, 1885; H. mariensis (Tupi Caldas, 1933); H. sanjuanensis (Sill, 1970); H. huenei Langer and Schultz, 2000 (NHMUK R699, NHMUK G281 FZB-PV-1867, MACN-18185, UFRGS-PV-0132; Sill, 1970; Chatterjee, 1974; Benton, 1983; Langer and Schultz, 2000).

Teyumbaita sulcognathus (Azevedo and Schultz, 1987) (UFRGS-PV-0232T, UFRGS-PV-0298T, and UFRGS-PV-0290T; Montefeltro et al., 2010).