

Terocephalian (Therapsida) and chroniosuchian (Reptiliomorpha) from the Permo-Triassic transitional Guodikeng Formation of the Dalongkou Section, Jimsar, Xinjiang, China

LIU Jun^{1,2} Fernando ABDALA³

(1 Key Laboratory of Vertebrate Evolution and Human Origins of Chinese Academy of Sciences, Institute of Vertebrate Paleontology and Paleoanthropology, Chinese Academy of Sciences Beijing 100044, China liujun@ivpp.ac.cn)

(2 University of Chinese Academy of Sciences Beijing 100039)

(3 Evolutionary Studies Institute, University of the Witwatersrand Private Bag 3, WITS 2050, Johannesburg, South Africa)

Abstract The Guodikeng Formation encompasses the terrestrial Permo-Triassic transition sequence in China. This formation crops out in the Dalongkou section, Jimsar, Xinjiang where remains of the dicynodonts *Jimusaria* and *Lystrosaurus* were found. We are describing here a therocephalian and a chroniosuchian from the Dalongkou section, which are the first records of these groups for the Guodikeng Formation. Diagnostic characters of the new therocephalian, *Dalongkoua fuae* gen. and sp. nov., include maxillary ventral margin strongly concave in lateral view; incisors spatulated and rounded; incisors and canines with faint serrations; coronoid process of the dentary with a marked external adductor fossa; triangular reflected lamina of the angular with two smooth concavities. Chroniosuchians are represented by several postcranial elements and the vertebral morphology is similar to *Bystrowiana* and *Bystrowiella*. These remains are interpreted as representing a Bystrowianidae indeterminate. The new findings increase the diversity of the Guodikeng Formation that is now represented by three or four dicynodonts, one therocephalian and one chroniosuchian. All these groups survived the massive P-T extinction but disappear from the fossil record in the Middle to Upper Triassic.

Key words Dalongkou, Xinjiang; Permian; Triassic; Guodikeng Formation; chroniosuchian; therocephalian

Citation Liu J, Abdala F, 2017. Terocephalian (Therapsida) and chroniosuchian (Reptiliomorpha) from the Permo-Triassic transitional Guodikeng Formation of the Dalongkou Section, Jimsar, Xinjiang, China. *Vertebrata Palasiatica*, 55(1): 24–40

1 Introduction

Dalongkou section, Jimsar, in Xinjiang is a famous terrestrial section with Permo-Triassic transitional sequences. The southern limb Dalongkou anticline section (SLS) has been suggested as a candidate for the global non-marine Permian-Triassic boundary (PTB) reference section (Cheng and Lucas, 1993, Liu, 1994). This section was originally known by its fossil

国家重点基础研究发展计划项目(编号: 2012CB821902)、国家自然科学基金(批准号: 41572019)、现代古生物学和地层学国家重点实验室基金(编号: 20161101)、中国科学院国际人才计划(编号: 2016VBB054)和科技部中德联合研究计划项目(编号: CS08-L02)资助。

收稿日期: 2016-06-30

vertebrate bearings. This formation produced the dicynodonts *Jimusaria sinkianensis* and the first specimen of *Lystrosaurus* from China (Sun, 1963; Yuan and Young, 1934a, b), later the co-occurrence of these taxa on the same horizon of the Dalongkou section were noticed (Cheng, 1993). The lowest occurrence of *Lystrosaurus* at the SLS is at the base of bed 63 or 54 of the measured lithostratigraphic section (Fig. 1), ~65 m below the base of the Jiucaiyuan Formation (Cheng, 1993; Liu et al., 2002). The exact position of the holotype of *Jimusaria sinkianensis* is unknown, but Cheng (in Yang et al., 1986) suggested it could lie within the top 20 m of the Guodikeng Formation. This is perhaps the reason why Kozur and Weems (2011) located the stratigraphic position of this dicynodont 15 m below the base of the Jiucaiyuan Formation. Besides dicynodonts, the procolophonid *Santaisaurus yuani* and the archosauriform *Chasmatosaurus yuani* were discovered from horizons of the Jiucaiyuan Formation at the Dalongkou section (Koh, 1940; Li et al., 2008; Young, 1958, 1963).

The PTB in the Junggar Basin was traditionally positioned at the base of the Jiucaiyuan Formation, but a joint team of the Chinese Academy of Geological Sciences and the Xinjiang Bureau of Geology and Mineral Resources first proposed the PTB within the Guodikeng Formation (Yang et al., 1986). They tentatively located the PTB at the uppermost Guodikeng Formation; one major reason for this decision was the position of the lowest occurrence of *Lystrosaurus*. For a long time, *Lystrosaurus* was a Triassic index fossil, and its lowest occurrence marked (or at least approximate) the base of the Triassic (cf. Lucas, 1998).

Based on palynomorph evidence, Ouyang and Norris (1999) placed the PTB at the base of Unit 3 in the uppermost Guodikeng Formation, ~50 m below the base of the Jiucaiyuan Formation. Hou (2004) discovered overlapping interval of sporomorphs from the middle part of the Guodikeng Formation at SLS, and placed the PTB at the base of Bed 51 of the measured lithostratigraphic section 1 (Fig. 1). Furthermore, Pang and Jin (2004) observed a distinct turnover in the ostracod assemblages between Bed 54 and 55 in the Guodikeng Formation (Fig. 1), approximately 10 m above the PTB suggested by the palynological analysis of Hou (2004). Foster and Afonin (2005) studied the abnormal pollen grains from Russia and China, and suggested that the latest Permian mass extinction level lies at the upper lower Guodikeng Formation, between Bed 63 to 65 on s1. Using conchostracan biostratigraphy, Kozur and Weems (2011) proposed the continental extinction event horizon at the middle part of the Guodikeng Formation at SLS; and they proposed that the first appearance datum (FAD) of *Falsisca verchojanica*, ~25 m below the base of the Jiucaiyuan Formation at SLS, correlates to the PTB. Based on the paleomagnetic data, Li et al. (2003) put the PTB between their Bed 41 and 42 at the SLS. Cao et al. (2008) suggested an isotopically defined PTB at the base of Bed 65 (Fig. 1).

In summary, there are distinct faunal and floral turnovers in the middle part of the Guodikeng Formation. As suggested by Kozur and Weems (2011), their relationships with the end-Permian mass extinction need to be studied carefully, but they must not equate to the PTB.

A recent revision of *Dicynodon* restricted the name to two species from the African

Permian and *Jimusauria*, that was synonymized to *Dicynodon*, is considered as a valid taxon different of *Dicynodon* (Kammerer et al., 2011). The phylogeny presented by Kammerer et al. (2011:fig. 156) found *Jimusauria* as a basal form in a clade of Permian and Triassic dicynodonts. Metcalfe et al. (2009) points out the poor record of *Lystrosaurus* in the Guodikeng Formation in opposition to the rich representation of this taxon in the overlying Jiucaiyuan Formation. Considering the faunal composition of the Permo-Triassic transition in South Africa, they argue that the combined presence of *Lystrosaurus* and the *Jimusauria* is more likely representing levels of the Late Permian, in which the former taxon is poorly represented (Smith and Botha-Brink, 2014). As a result the PTB is interpreted on the top of the Guodikeng Formation, near the base of the Jiucaiyuan Formation. Recently, Gastaldo et al. (2015) reported a dicynodontoid skull of characteristically Permian aspect more than 10 m above the previous last occurrence of *Dicynodon* and a zircon age of (253.48 ± 0.15) Ma from a layer ~60 m below the current vertebrate-defined boundary from Karoo Basin. They

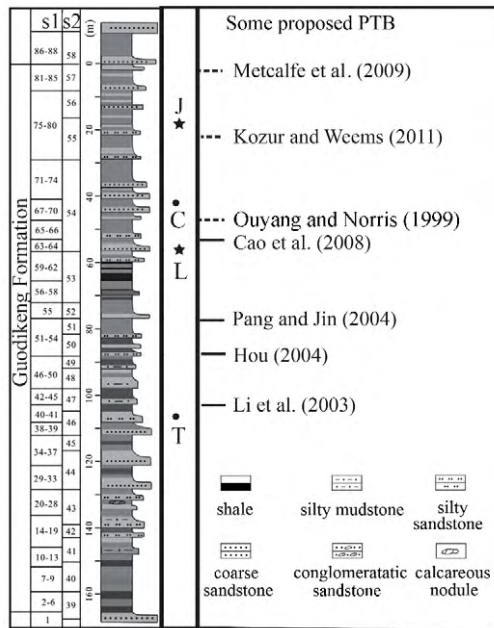


Fig. 1 Position of some vertebrate fossils and proposed PTB from the Guodikeng Formation in the SLS

The stratigraphic column adapted from Cao et al. (2008). Subunits of lithological stratigraphy listed as s1 are from team work of 2000 (published in Li et al., 2003; Hou, 2004 and Pang and Jin, 2004), s2 from Yang et al. (1986)

C. IVPP V 23295, a chroniosuchian; J. the possible position of *Jimusauria sinkiangensis*; L. the lowest occurrence of *Lystrosaurus*; T. IVPP V 23296, the therocephalian *Dalongkoua fuae*. The continuous lines mean the exact position of PTB, while the dashed lines indicate unsure on the position

suggested the PTB should be higher than that recognized by Ward et al. (2005). The work in Karoo Basin suggested an older occurrence of *Lystrosaurus*, and the PTB could be around the base of the Jiucaiyuan Formation in Junggar Basin.

In 2000, some bones were collected from the SLS while digging a trench for paleomagnetic study. One specimen (IVPP V 23296) was produced from a green mudstone close to the yellow sandstone in Bed 40. All bones of this specimen are badly weathered, before being buried. Another specimen came from the red mudstone of Bed 70, including several vertebrae and a nearly complete humerus (IVPP V 23295). These materials were prepared and identified as a therocephalian and a chroniosuchian. Therocephalian therapsids were important component of Middle Permian to Middle Triassic terrestrial faunas (Abdala et al., 2008, 2014; Huttenlocker et al., 2015; Ivakhnenko, 2011). In China, therocephalians are only known from the Triassic, including *Urumchia lii* recovered from the Jiucaiyuan Formation of Xinjiang, which was originally assigned as Late Permian (Li et al., 2008;

Young, 1952). The new discoveries represent, thus, the first report of a Permian therocephalian for China. Chroniosuchians is an enigmatic and a recent addition to the group of tetrapods. It was first reported in the Permian and Triassic of Russia (Golubev, 1998; Novikov and Shishkin, 2000; Shishkin et al., 2014), and later from the Permian of China (Li and Cheng, 1999; Young, 1979), and the Triassic of Kyrgyzstan and Germany (Schoch et al., 2010; Witzmann et al., 2008). The new material here presented is a new Permian record for China, and the first one from Xinjiang.

Abbreviation IVPP, Institute of Vertebrate Paleontology and Paleoanthropology, Chinese Academy of Sciences.

2 Systematic paleontology

Therapsida Broom, 1905

Therocephalia Broom, 1903

Eutherocephalia Hopson & Barghusen, 1986

***Dalongkoua fuae* gen. et sp. nov.**

Holotype IVPP V 23296, incomplete left premaxilla with three teeth, incomplete right premaxillae with four teeth, snout and part of the orbital region with the maxilla on both sides featuring functional and replacement canines, isolated vertebrae, left humerus, two phalanges and some bone fragments (Figs. 2–4).

Etymology ‘Dalongkou’, the name of locality; ‘fu’, dedicated to the preparator Ms. Fu Hua-Lin.

Diagnosis Maxillary ventral margin strongly concave in lateral view; incisors spatulated and rounded; incisors and canines with faint serrations; coronoid process of the dentary with a marked external adductor fossa; triangular reflected lamina of the angular with two smooth concavities.

Description Skull The skull is strongly deformed as a flat plate and broken into small blocks. The anterior portions of the premaxillae are separated from the rest of the skull (Fig. 2).

Both premaxillae are represented as independent elements, with the right being better preserved. It forms a curved plate housing the alveoli for four incisors (Fig. 2C). Posteromedially, it has the process for contacting the vomer. The posterolateral side of the palatal surface curves dorsally, forming the anterior margin of the palatal fenestra for the lower canine (Fig. 2C). There are four preserved teeth, three of them with the main axis of their bases oriented longitudinally, whereas the second is circular in section. The latter tooth, is completely preserved, presenting spatulated crown, concave lingually, and mesiolingual and distal smooth ridges (Fig. 2E). Remaining teeth are incomplete lacking any ridge at their bases, where the external surface is relatively smooth. Based on the preservation, we interpret the probable presence of at least five incisors. Two replacing teeth lie on the lingual side of the first and third functional incisors (Fig. 2C). The crown of the first replacement incisor show faint

serrations on the medial margin. A tiny foramen lies posterior to the first replacing incisor. A long diastema separated the last incisor and the canine. The left premaxilla has three poorly preserved incisors and an extra alveolus for the fourth incisor, without tooth. There are pits with replacement teeth behind the second and third incisors.

The maxilla is a long bone with large facial plate. It extends anteriorly and covers the lateral surface of the premaxilla anterior to the canine. Ventrally, there is a concave ventral step. The maxilla is laterally expanded to encapsulate the canine, and the snout seems

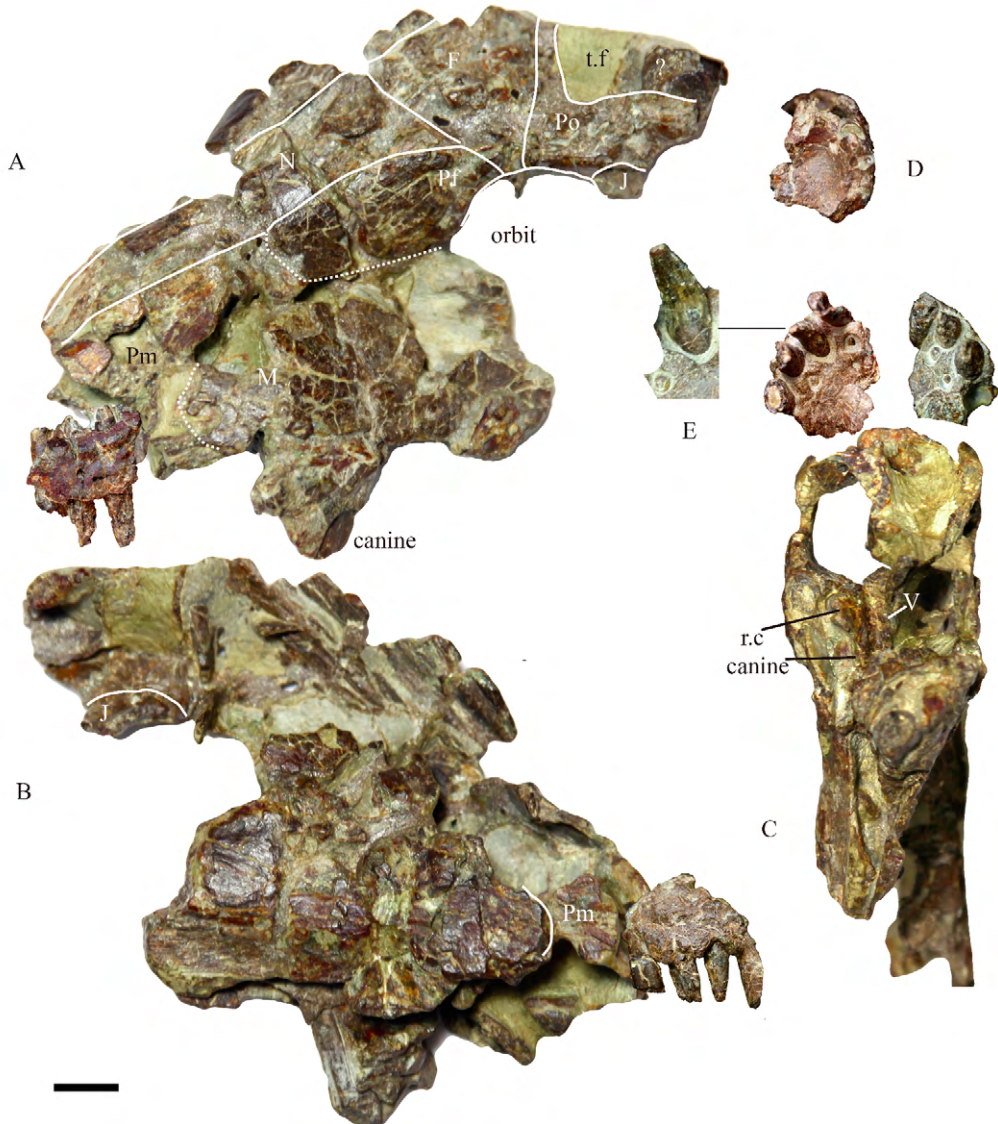


Fig. 2 Holotype of *Dalongkoua fuae* gen. et sp. nov. (IVPP V 23296)

Skull, including premaxilla in left lateral (A), right lateral (B) and ventral (C) views; right premaxilla in dorsal view (D); the second preserved right incisor in lingual view (E)

Abbreviations: F. frontal; J. jugal; M. maxilla; N. nasal; Pf. prefrontal; Pm. premaxilla; Po. postorbital; r.c. replacing canine; t.f. temporal fenestra; V. vomer. Scale bar equals 1 cm

constricted behind the canine. On the right side, the facial plate is folded medially, and a remarkable longitudinal ridge is present, suggesting a well exposed maxillary platform. The platform is posteriorly concave in lateral view as in *Theriognathus microps* (Huttenlocker and Abdala, 2016). The root of the left canine is exposed posteriorly. An erupting replacing canine lies anteromedially to the canine on both sides. The maxillary ventral margin is strongly concave in lateral view. The postcanines are not observed on the skull, but some isolated teeth could be postcanines. Two of them preserved the base of the crown, and it is possible to see strong ridges on two margins without serration. One margin is remarkable concave, the other is straight to slightly concave. One preserved tip of postcanine have faint serrations on both margins of the cusp. The putative preserved postcanines are much smaller than the incisors.

The nasal is long and forms the dorsal surface of the snout. A middle ridge is present on the posterior portion of the nasal. It is sutured with the frontal on the level of the anterior margin of the orbit (Fig. 2A). Laterally, it contacts the maxilla and the prefrontal. The prefrontal, frontal and postorbital form the dorsal margin of the orbit, whereas the postorbital bar is formed by the postorbital and jugal.

Lower jaw **Dentary** The symphyseal region and the posterior part of the left dentary is preserved, as well as the right reflected lamina of the angular (Fig. 3). The articular symphysis with the right dentary is broken, and the lower margin is incomplete (Fig. 3B). The symphyseal region bears a marked constriction lateral to the lower canine where the upper canine rests. The anterior part of this region bears pits and grooves, whereas the surface that will be contacting the medial side of the upper canine is smooth. The posteroventral portion of the dentary forms a thickened lower border that supports the angular in a trough on its medial surface (Fig. 3B). The posteroventral corner of the dentary forms an angle of 137° . The coronoid process is well developed, with a dorsal adductor fossa on the lateral surface (Fig. 3A). The posteroventral margin of the dentary, below the bulged ridge that limits ventrally the adductor fossa, forms a descending flange ending in a sharp ridge. Although the posterodorsal corner is incomplete, the preserved portion shows a nearly straight posterodorsal margin. Six teeth are preserved on the symphyseal region. The biggest one is the canine, and anteromedially there is a replacing canine that is partially wrapped by it (Fig. 3C). Faint serrations developed on the anterior margin of the replacing canine. There are four lower incisors, including three preserved teeth and one alveolus (Fig. 3C). A tiny tip of an erupting tooth lies immediately anterior to the canines. The last incisor is small and partially preserved; whereas two other incisors, medial to the later, are also preserved. One of them is faceted showing a well-developed ridge laterally (the medial side is not exposed). Very fine serrations, only visible under microscope magnification, are present on the margins of the larger two preserved incisors.

Angular The right angular is nearly complete. This bone is flat and elongated posteriorly with a triangular-like reflected lamina (Fig. 3D, E). The large reflected lamina bears a deep ‘U’-shaped dorsal notch which is posterodorsally directed (Fig. 3D). The lateral surface of the reflected lamina is ornamented by fine, radiating ridges and grooves, and is divided into three

parts by large grooves directed ventrally and posteriorly (Fig. 3D).

Prearticular The prearticular lies on the medial side of the angular (Fig. 3E). It is an elongate element forming an anterior rod to contact the splenial. Posteriorly, it is a triangular plate, whose posterior process contacts the articular.

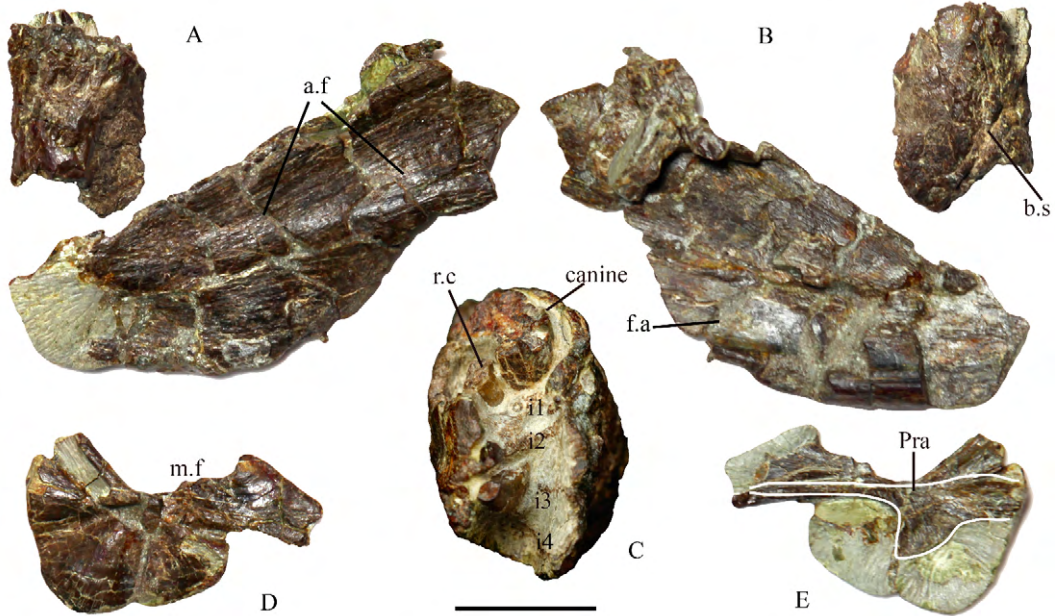


Fig. 3 Holotype of *Dalongkoua fuae* gen. et sp. nov. (IVPP V 23296)

Symphysis and posterior portion of the left dentary in lateral (A) and medial (B) views; left dentary symphysis in dorsal view (C); right angular in lateral (D) and medial (E) views

Abbreviations: a.f. adductor fossa; b.s. broken surface; f.a. facient on the dentary for the angular; i. incisor; m.f. mandibular fenestra; Pra. prearticular; r.c. replacing canine. Scale bars equal 1 cm

Vertebrae Two partial vertebrae including centra and a left partial neural arch are represented. The complete centrum in one of them, interpreted as dorsal vertebra, is amphicoelous, and deeply hollowed. The second element has an incomplete centrum and a clear suture between the centrum and the right partial neural arch shows lack of synostosis between them (Fig. 4A, B). The transverse process extends dorsolaterally and is angled posterolaterally, with a vertical flange on the ventral side. This is similar to the dorsal vertebra of *Promoschorhynchus platyrhinus* (Huttenlocker et al., 2011).

The isolated arch is possible of a cervical vertebra, with its zygapophyses bearing more or less horizontal articulating surfaces (Fig. 4C, D). The transverse process is not preserved and should lie well down on the neural arches. The neural spine is robust and inclines anteriorly. This is similar to the cervical of *Olivierosuchus parringtoni* (Botha-Brink and Modesto, 2011).

Sternum A partial posterior margin of the sternum is preserved (Fig. 4G, H). It is a large plate with convex ventral surface and concave dorsal surface. There is no posterior notch, and a weak midline ridge is developed on the ventral surface, as in *Promoschorhynchus platyrhinus* (Huttenlocker et al., 2011).

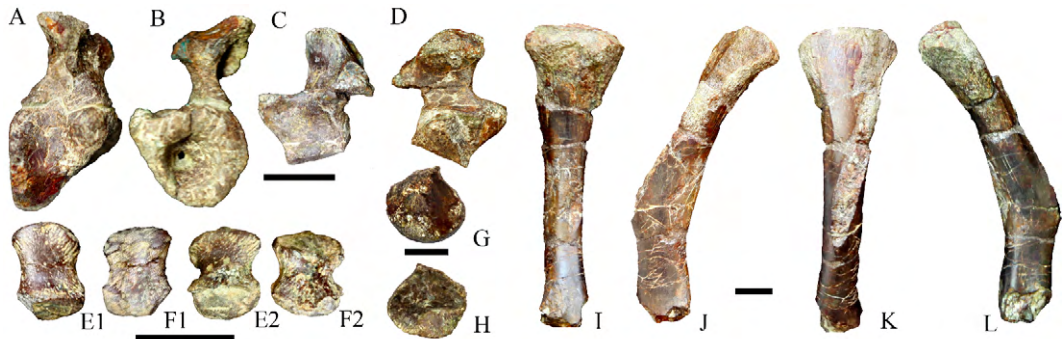


Fig. 4 Holotype of *Dalongkoua fuae* gen. et sp. nov. (IVPP V 23296)

An incomplete dorsal vertebra in lateral (A) and posterior (B) views; an incomplete cervical neural arch in left lateral (C) and medial (D) views; two phalanges in dorsal (E) and ventral (F) views, distal to the top; sternum in ventral (G) and dorsal (H) views; left humerus in dorsal (I), anterior (J), ventral (K) and posterior (L) views. Scale bars equal 1 cm

Humerus The humerus is robust with expanded proximal end, missing the distal end and the deltopectoral crest (Fig. 4I–L). The proximal half is curved dorsally relative to the shaft (Fig. 4J, L). A low ridge lies on the rough posterodorsal side, possible representing part of the wide posteroventral ridge described by Kemp (1986). The long shaft is broken on the place of the entepicondylar foramen.

Phalanges Two phalanges are well preserved (Fig. 4E, F). A longitudinal middle groove develops on the ventral surface. They are nearly as wide as length. Nearly quadrangular phalanges are represented in several therocephalians including the Scylacosauridae *Glanosuchus*, the Akidnognathidae *Olivierosuchus*, the Hofmeyriid *Mirotenthes* and the Whaitsiidae *Theriognathus* (Attridge, 1956; Boonstra, 1964; Botha-Brink and Modesto, 2011; Fourie and Rubidge, 2007; Abdala, pers. obs).

Chroniosuchia Tatarinov, 1972

Bystrowianidae Vjuschkov, 1957

Genus and species undetermined

Referred specimen IVPP V 23295, five vertebrae, rib, gastral scales, partial ilium, left femur, and partial fibula (Figs. 5, 6).

Description Vertebrae Three articulated vertebrae show two intercentra, two isolated pleurocentra are fused to the neural arches, and there are three isolated intercentra. Four larger vertebrae are tentatively identified as dorsals and the smallest as caudal.

The morphology of the vertebrae is very similar to that of *Bystrowiana sinica* or *Bystrowiella schumanni* (Liu et al., 2014; Witzmann et al., 2008) (Fig. 5).

The length of the large pleurocentrum is approximately 14 mm, whereas the length in the other two is 12 and 10 mm. The pleurocentrum are roughly twice as long anteroposteriorly as the intercentra. The height of the pleurocentra is about 10 mm. All pleurocentra are deeply amphicoelous with a round cross-section, and they are not perforated by the notochord. In

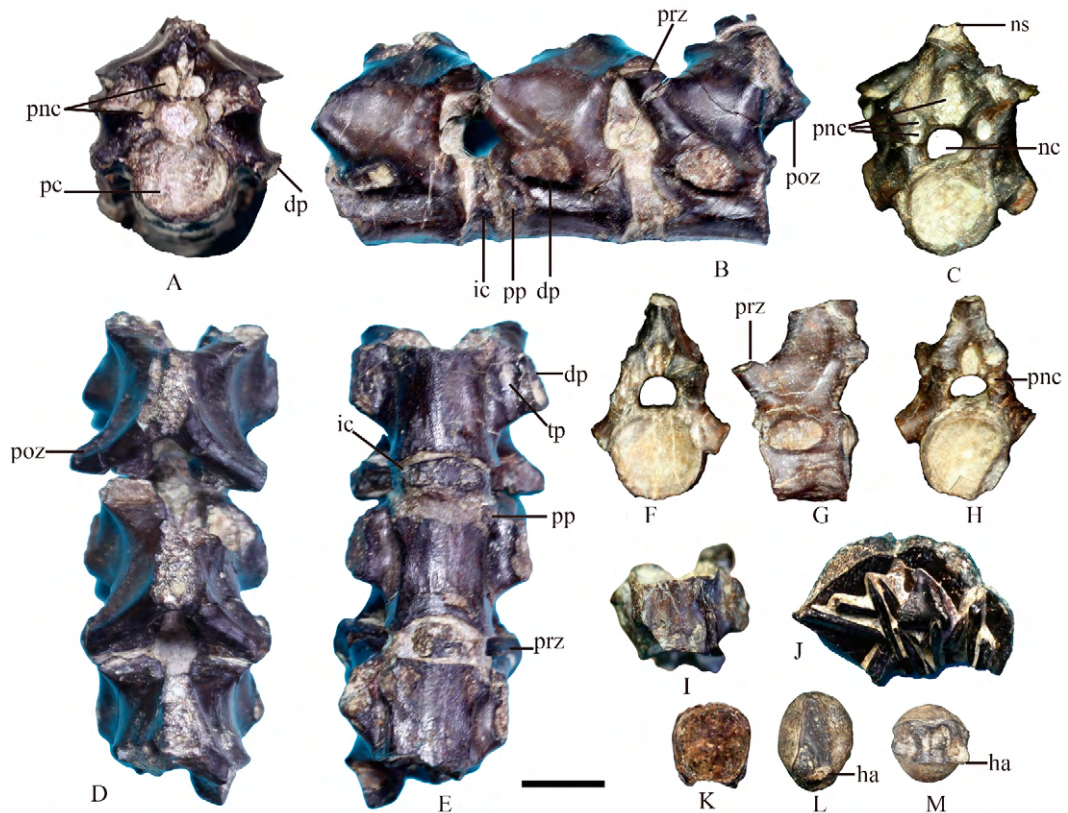


Fig. 5 Chroniosuchian specimen (IVPP V 23295) from Dalongkou section

Three dorsal vertebrae in anterior (A), left lateral (B), posterior (C), dorsal (D) and ventral (E) views; one caudal vertebra in anterior (F), left lateral (G), posterior (H), and ventral (I) views; J. gastral scales; caudal intercentrum in anterior (K), left lateral (L), and ventral (M) views

Abbreviations: dp. diapophysis; ha. haemal arch; ic. intercentrum; nc. neural canal; ns. neural spine; pc. pleurocentrum; pnc. paraneural canal; poz. postzygapophysis; pp. parapophysis; prz. prezygapophysis; tp. transverse process. Scale bar equals 1 cm

lateral view, the anterior and posterior margins of the pleurocentrum are curved but the ventral margin is nearly straight. A small knob (parapophysis) for the capitulum lies on the middle of the pleurocentrum anterior margin (Fig. 5B, E), and a longitudinal faint ridge runs posterior to it. The parapophysis is separated from the diapophysis by an anterodorsally directed incisure. On the larger pleurocentrum, the ventral surface is slightly convex to smooth (Fig. 5E). On the smallest pleurocentrum, two anteroposteriorly directed, low ridges delimit the faintly concave ventral margin (Fig. 5I) as in *Bystrowiella schumanni* (Witzmann et al., 2008).

The intercentra are about as tall as the pleurocentra in lateral view. The articular intercentra show smooth periosteal bone, without haemal arch on the ventral surface. They appear wedge-like in articulation (Fig. 5B, E), but actually are ball-like (Fig. 5K–M). On three free caudal intercentra, the ventral surface consists of slightly concave, smooth periosteal bone, and bears the ventrolateral bases of the hemapophyses as in *Bystrowiella schumanni* (Witzmann et al., 2008). The roughened dorsal surface of the intercentrum is convex and did not articulate

with the neural arch, forming part of the base of the neural canal. The largest intercentrum measures 9.7 mm long, while the others range from 8 to 9 mm. The exposed length of them in ventral view is nearly 5 mm.

All neural arches are fused to the pleurocentrum and the spines only preserved at their base. The zygapophyses are widely spaced (Fig. 5D, E) and the facets slope medially at a small angle to the horizontal plane in anterior view. There are two pairs of well-developed paraneural canals at the base of the zygapophyses, with the anterior openings of the dorsal pair being larger than those of the lateral pair (Fig. 5A). As in *Bystrowiana permira*, the number and shape of the openings on the two sides could be different (Liu et al., 2014). The neural canal is compressed dorsoventrally. The short transverse process lies on the upper half of the pleurocentrum, bearing small, unfinished diapophyses (Fig. 5A–C).

Rib and gastral scales A proximal rib fragment, and some gastral scales are preserved on a bone plate (Fig. 5J). The gastral scales are thin, some ending in a sharp point.

Ilium The preserved acetabular area of the right ilium is similar to that of *Dendrysecos helogenes* or *Chroniosaurus* (Clack and Klembara, 2009; Holmes et al., 1998; Schoch and Milner, 2014). The anterior margin is slightly concave (Fig. 6A). There is a distinct crescent supra-acetabular buttress, and, anteroventrally, a notch on the anterior margin of acetabulum. The iliac blade is only preserved as a narrow base above the acetabular rim. On the medial side, a strong iliac ridge runs dorsoventrally and divides the medial surface in two portions (Fig. 6B).

Femur The slender left femur is well-ossified and nearly complete (Fig. 6C–H). The general shape of the bone is same as that of *Chroniosaurus* (Clack and Klembara, 2009). The length of the bone is 73 mm, and the width is 21 mm for proximal and distal ends. The distal side deflects ventrally relative to the proximal part, so the bone is curved dorsally. The anterior margin is curved while the posterior margin is relatively straight. Differing from *Chroniosaurus dongusensis*, the articular surface on the proximal side is smoothly convex and dorsoventrally flattened (Fig. 6C). The dorsoventral height of the articular surface is near half its anteroposterior length (Fig. 6G). The proximal extensor surface of the femur bears some striations on the posterior side (Fig. 6C), indicating the insertion for the ischiotrochantericus.

On the anteroflexor side, the ventrally directed adductor blade demarcates the anterior side of the deep intertrochanteric fossa (Fig. 6D, E). The broken internal trochanter is separated from the femoral head by a narrow ridge, and it continues distally as a low crest, the fourth trochanter (Fig. 6D). The anterior surface of the adductor blade is rugose with deep muscle scars.

From the distal end of the fourth trochanter, the adductor crest follows a strongly diagonal course posteroventrally to a point distal to the middle of the shaft (Fig. 6E, F). A low ridge continues distally on the posterior side of the shaft ending on the distal posterior corner. This ridge does not seem to be for muscle attachment. In other groups such as captorhinomorphs, seymouriamorphs and temnospondyls, the adductor crest generally runs towards the popliteal area (Fox and Bowman, 1966; Klembara and Bartik, 2000; Pawley, 2007).

The popliteal area is pitted and concave on the flexor side (Fig. 6E). The fibular (posterior)

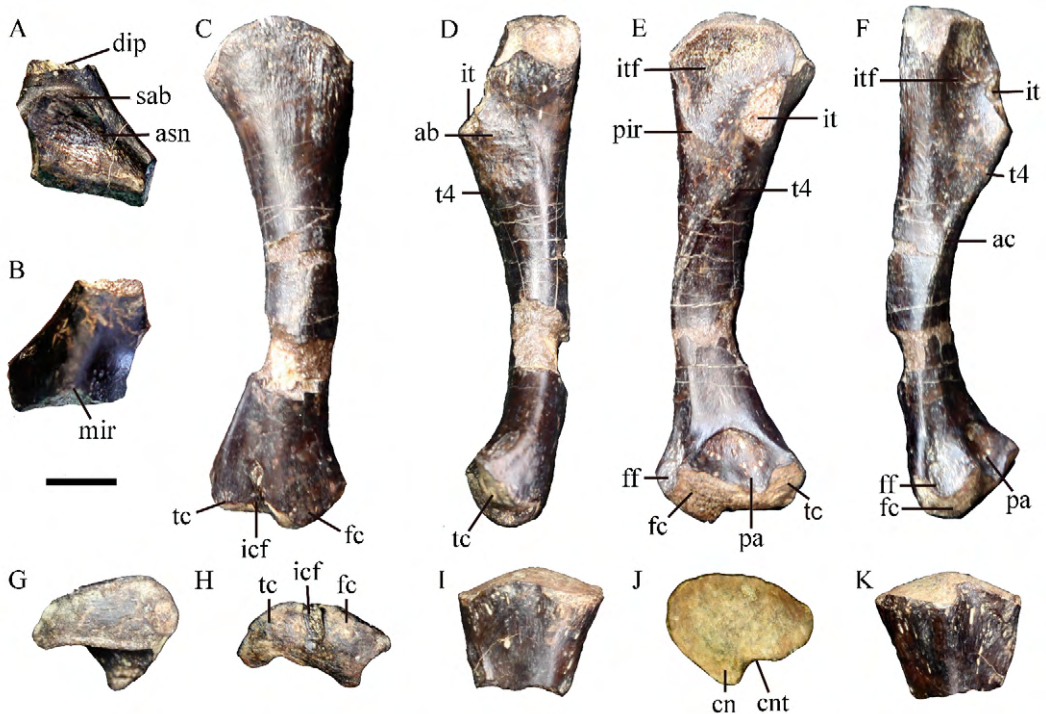


Fig. 6 Chroniosuchian specimen (IVPP V 23295) from Dalongkou section

Right ilium in lateral (A) and medial (B) views; left femur in extensor (C), anterior (D), flexor (E), posterior (F), proximal (G), and distal (H) views; left fibula in flexor (I); proximal (J) and extensor (K) views. Abbreviations: ab. adductor blade; ac. adductor crest; asn. anterior supracetabular notch; cn. cnemial crest; cnt. cnemial trough; dip. dorsal iliac process; fc. fibular condyle; ff. fibula fossa; icf. intercondylar fossa; it. internal trochanter; itf. intertrochanteric fossa; mir. mesial iliac ridge; pa. popliteal area; pir. posterior intertrochanteric ridge; sab. supracetabular buttress; tc. tibial condyle; t4. fourth trochanter. Scale bar equals 1 cm

condyle projects more distally than the tibial (anterior) condyle. Proximal to the fibular condyle is a narrow but deep fibular fossa. In extensor view, a deeply incised intercondylar fossa is oriented posteroventrally towards the midline of the fibular condyle (Fig. 6H).

Fibula The proximal end of the left fibula is preserved (Fig. 6I–K). The head is convex to fit on the concave articular surface of the femur. Small striae distribute around the proximal articular surface (Fig. 6I, K), indicating the presence of cartilage or perhaps ligamentous connective tissue. The extensor surface is incised by a cnemial trough, which is bordered anteriorly by the prominent cnemial crest (Fig. 6J). Another crest is on the posterior surface of the bone. A striate area on the medial side of this ridge, distal to the proximal articular surface, provided attachment for muscles.

3 Discussion

The new therapsid specimen (IVPP V 23296) can be diagnosed as a therocephalian for the presence of palatal fenestra for the lower caniniform confluent with choana, and the

posterventral portion of the dentary forming a thickened lower border that extends below the angular bone and supports the latter in a trough on its medial surface. It has four lower incisors, and can be then diagnosed as an eutheriocephalian (Huttenlocker, 2009). In addition, the shape of the reflected lamina, lacking an anterodorsal rounded portion in front of the notch that makes this structure circular in basal therocephalians is also an indication that the new species is member of Eutheriocephalia.

Dalongkoua fuae show evidence of faint serrations in the margin of a replacement incisor and canine and on the tip of a postcanine. These findings are contrary to the assumed absence of serrations in the dentition of eutheriocephalia (e.g. van den Heever, 1994; Abdala et al., 2008; Huttenlocker, 2009; and Huttenlocker et al., 2015, for incisors and Huttenlocker and Sidor, 2012 for anterior dentition). The serrations in the teeth of *Dalongkoua fuae* are exceedingly faint and it is clear that they will disappear with a minimum wear as demonstrated by the smoothly ridged crown of the preserved elements.

This specimen represents a new species on the basis of the following characters: maxillary ventral margin is strongly concave in lateral view; incisors spatulated and rounded; incisors and canines with faint serrations; coronoid process of the dentary with a marked adductor fossa; triangular reflected lamina with two concavities.

The maxillary ventral margin is convex or nearly straight in most therocephalians, but it is slightly concave in *Moschowiaisia* (Ivakhnenko, 2011), and strongly concave in *Euchambersia* and *Theriognathus* (Broom, 1931; Huttenlocker and Abdala, 2016). Recent studies (e.g., Huttenlocker et al., 2015) indicate that spatulated incisors are restricted to a group of akidnognathids. In *Dalongkoua fuae* the completely preserved second left incisor is spatulated, whereas remaining left incisors, although incomplete, show the crown enough preserved to suggest the presence of rounded (non-spatulated) incisors. The adductor fossa is developed in the dorsolateral portion of the coronoid process, above of the through for the postdentary bones. This fossa is different to the one described on the lateral surface of the dentary in the bauriids *Microgomphodon oligocynus* (Abdala et al., 2014) and in the Russian *Notogomphodon danilovi* (Abdala, pers. obs.), as in these taxa there is no connection between the fossa and the through for the postdentary bones. The incomplete left dentary of *Urumchia lii* also seems to have an adductor fossa, although less developed than in *D. fuae* (Fig. 7). A triangular reflected lamina is also present in the holotype of *Tetracynodon darti*, but the lamina show a more complex lateral surface with several ridges and valley. In the lamina of *D. fuae* there is a posterior deep canal and a dorso-ventrally oriented shallow and wide depression.

Urumchia lii is the only known therocephalian from the Xinjiang, documented in the Jiucayuan Formation (Sun, 1991; Young, 1952). Diagnostic characters in this species are concentrated in the palate and in the postcanine morphology (Li et al., 2008; Sun, 1991). Unfortunately both palate and postcanines are incompletely preserved in *Dalongkoua fuae*, the postcanines only represented by isolated fragments including two crown bases and one tip. There are however a set of characters shared between these two species indicating perhaps

relationships (Fig. 7): medial margin of the frontal bones directed dorsally in *Dalongkoua fuae* indicating the presence of a central ridge in the posterior internasal and interfrontal sutures, the presence of a depressed facial area of the maxilla immediately above of the dental series, also described recently in *Ichibengops* of East Africa by Huttenlocker et al (2015), an apparent development of an adductor fossa, although clearly smaller, in the left dentary of *Urumchia lii*; and the presence of spatulated incisors. The putative relationship between these two taxa should be considered as a working hypothesis, whose validation will require finding of additional material.



Fig. 7 *Urumchia lii* (IVPP V 702)

Skull in dorsal (A) and right lateral (B) views; lower jaws in left lateral view (C). Showing the sharing characters with *Dalongkoua fuae*: 1. the mid-ridge between the posterior portions of the nasals and the anterior portions of the frontals; 2. depressed facial area of the maxilla; 3. adductor fossa on the dentary

Scale bars equal 1 cm

The new specimen of Reptiliomorpha (IVPP V 23295) is referred to chroniosuchian for the characteristic ball-and-socket joint between pleurocentra and intercentra (Ivakhnenko and Tverdokhlebova, 1980). It can be further assigned to Bystrowianidae for paired paraneural canals on the vertebrae (Novikov et al., 2000). The vertebrae show the similar morphology as other bystrowianids, and intercentra is similar to the figured for *Bystrowiella schumanni* (Novikov and Shishkin, 2000; Witzmann et al., 2008). The morphology of ilium, femur, and fibula is consistent with this taxonomic identity (Clack and Klembara, 2009; Ivakhnenko and Tverdochlebova, 1980). Unfortunately, no dermal scute was discovered, and no further identification could be made.

Chroniosuchian specimens in China were first reported from the Upper Permian Jiyuan fauna (Young, 1979), and then from the Middle Permian Dashankou fauna (Li and Cheng, 1999). The new findings here reported represent the latest occurrence of chroniosuchians in

China. This group however is known from the Middle/Upper Triassic Madygen Formation of Kyrgyzstan (Schoch et al., 2010), indicating their continued presence in Laurasia after the Permian extinction.

These new findings allow recognizing that the diversity of tetrapod near the Permo-Triassic boundary in China, and particularly in Xinjiang, was higher than previously recognized. Only dicynodonts were previously reported from the Guodikeng Formation, including *Lystrosaurus*, *Jimsaria*, *Turfanodon*, and possibly *Diictodon* (Li et al., 2008). Both chroniosuchians and therocephalians crossed the P-T boundary; therocephalians become extinct in the Middle Triassic (Abdala et al., 2014), whereas the youngest record of chroniosuchians is in the Middle or Late Triassic (Schoch et al., 2010).

Acknowledgements The authors thank the field crew to Dalongkou in 2000 (Cheng Zheng-Wu, Pang Qi-Qing, Li Yong-An, Sun Dong-Jiang and Jin Xiao-Chi), FU Hua-Lin prepared the specimens, Jozef Klembara kindly provided some photos of *Chroniosaurus*. Adam Huttenlocker and Florian Witzmann provided valuable comments. JL is supported by National Basic Research Foundation (973 grant no. 2012CB821902), NSFC grants (41572019), and State Key Laboratory of Palaeobiology & Stratigraphy (Nanjing Institute of Geology & Palaeontology, CAS) grants (No. 20161101). FA research is supported by the National Research Foundation of South Africa and a research trip to China funded by the Chinese Academy of Sciences President's International Fellowship Initiative Grant (2016VBB054).

新疆吉木萨尔大龙口剖面锅底坑组新发现的兽头类和迟滞鳄类

刘俊^{1,2}

Fernando ABDALA³

(1 中国科学院古脊椎动物与古人类研究所, 中国科学院脊椎动物演化与人类起源重点实验室 北京 100044)

(2 中国科学院大学 北京 100039)

(3 南非金山大学进化研究所 约翰内斯堡 WITS2050)

摘要: 锅底坑组是重要的二叠—三叠系过渡序列, 本组在新疆吉木萨尔大龙口出露广泛, 产出吉木萨尔兽以及水龙兽两类二齿兽类。本文报道了在大龙口剖面首次发现的兽头类和迟滞鳄类, 这也是此二类群在锅底坑组的首次报道。新发现的兽头类被命名为付氏大龙口兽(*Dalongkoua fuae*)。其鉴定特征包括上颌骨犬后齿齿槽外边缘向背向凹入; 切齿有圆形和匙形; 切齿和犬齿有微弱的锯齿; 齿骨冠状突有显著的收肌窝; 反折翼近三角形, 有两个平滑的凹陷。锅底坑组目前有3~4属二齿兽类、1属兽头类和1属迟滞鳄类, 这一发现增加了其多样性。这些类群都在二叠纪末的大灭绝中幸存下来, 直到中晚三叠世才消失。

关键词: 新疆大龙口, 二叠纪, 三叠纪, 锅底坑组, 兽头类, 迟滞鳄类

中图法分类号: Q915.864 **文献标识码:** A **文章编号:** 1000-3118(2017)01-0024-17

References

- Abdala F, Rubidge B S, Van Den Heever J A, 2008. The oldest therocephalians (Therapsida, Eutheriodontia) and the early diversification of Therapsida. *Palaeontology*, 51(4): 1011–1024
- Abdala F, Kammerer C F, Day M O et al., 2014. Adult morphology of the therocephalian *Simorhinella baini* from the Middle Permian of South Africa and the taxonomy, paleobiogeography, and temporal distribution of the Lycosuchidae. *J Paleont*, 88(6): 1139–1153
- Attridge J, 1956. IV.—The morphology and relationships of a complete therocephalian skeleton from the *Cistecephalus* Zone of South Africa. *Proc R Soc Edinburgh B*, 66(1): 59–93
- Boonstra L D, 1964. The girdles and limbs of the pristerognathid Therocephalia. *Ann S Afr Mus*, 48: 121–165
- Botha-Brink J, Modesto S P, 2011. A new skeleton of the therocephalian synapsid *Olivierosuchus parringtoni* from the Lower Triassic South African Karoo Basin. *Palaeontology*, 54(3): 591–606
- Broom R, 1931. Notices of some new genera and species of Karoo fossil reptiles. *Rec Albany Mus*, 4: 161–166
- Cao C Q, Wang W, Liu L J et al., 2008. Two episodes of ¹³C-depletion in organic carbon in the latest Permian: evidence from the terrestrial sequences in northern Xinjiang, China. *Earth Planet Sci Lett*, 270: 251–257
- Cheng Z W, 1993. On the discovery and significance of the nonmarien Permo-Triassic transition zone at Dalongkou in Jimsar, Xinjiang, China. *New Mexico Mus Nat Hist Sci Bull*, 3: 65–67
- Cheng Z W, Lucas S G, 1993. A possible nonmarine GSSP for the Permian-Triassic boundary. *Albertiana*, 12: 39–44
- Clack J A, Klembara J, 2009. An articulated specimen of *Chroniosaurus dongusensis* and the morphology and relationships of the chroniosuchids. *Spec Pap Palaeontol*, 81: 15–42
- Foster C B, Afonin S A, 2005. Abnormal pollen grains: an outcome of deteriorating atmospheric conditions around the Permian-Triassic boundary. *J Geol Soc*, 162(4): 653–659
- Fourie H, Rubidge B S, 2007. The postcranial skeletal anatomy of the therocephalian *Regisaurus* (Therapsida: Regosauridae) and its utilization for biostratigraphic correlation. *Palaeont Afr*, 42: 1–16
- Fox R C, Bowman M, 1966. Osteology and relationships of *Captorhinus aguti* (Cope) (Reptilia: Captorhinomorpha). *Univ Kansas Paleontol Contrib Vertebr*, 11: 1–79
- Gastaldo R A, Kamo S L, Neveling J et al., 2015. Is the vertebrate-defined Permian-Triassic boundary in the Karoo Basin, South Africa, the terrestrial expression of the end-Permian marine event? *Geology*, 43(10): 939–942
- Golubev V K, 1998. Narrow-armored chroniosuchians (Amphibia, Anthracosauromorpha) from the Late Permian of Eastern Europe. *Paleontol J*, 32(3): 278–287
- Holmes R B, Carroll R L, Reisz R R, 1998. The first articulated skeleton of *Dendrerpeton acadianum* (Temnospondyli, Dendrerpetontidae) from the Lower Pennsylvanian locality of Joggins, Nova Scotia, and a review of its relationships. *J Vert Paleont*, 18(1): 64–79
- Hou J P, 2004. Two spore-pollen assemblages of Guodikeng formation and discussion on the Permo-Triassic boundary in Junggar Basin, Xinjiang. *Prof Pap Stratigr Palaeont*, 28: 177–204
- Huttenlocker A, 2009. An investigation into the cladistic relationships and monophyly of therocephalian therapsids (Amniota: Synapsida). *Zool J Linn Soc*, 157(4): 865–891
- Huttenlocker A K, Abdala F, 2016. Revision of the first therocephalian, *Theriognathus* Owen (Therapsida: Whaitsiidae), and implications for cranial ontogeny and allometry in nonmammaliaform eutheriodonts. *J Paleont*, 89(04): 645–664

- Huttenlocker A K, Sidor C A, 2012. Taxonomic revision of Therocephalians (Therapsida: Theriodontia) from the Lower Triassic of Antarctica. *Am Mus Novit*, 3738: 1–19
- Huttenlocker A K, Sidor C A, Smith R M H, 2011. A new specimen of *Promoschorhynchus* (Therapsida: Therocephalia: Akidognathidae) from the Lower Triassic of South Africa and its implications for theriodont survivorship across the Permo-Triassic boundary. *J Vert Paleont*, 31(2): 405–421
- Huttenlocker A K, Sidor C A, Angielczyk K D, 2015. A new eutheriocephalian (Therapsida, Therocephalia) from the Upper Permian Madumabisa Mudstone Formation (Luangwa Basin) of Zambia. *J Vert Paleont*, 35(5): e969400
- Ivakhnenko M F, 2011. Permian and Triassic therocephals (Eutherapsida) of Eastern Europe. *Paleontol J*, 45(9): 981–1144
- Ivakhnenko M F, Tverdochlebova G I, 1980. Systematics, Morphology, and Stratigraphic Significance of the Upper Permian Chroniosuchians from the East of the European Part of the USSR. Saratov: Izdatel'stvo Saratovskogo Universiteta. 1–69
- Kammerer C F, Angielczyk K D, Fröbisch J, 2011. A comprehensive taxonomic revision of *Dicynodon* (Therapsida, Anomodontia) and its implications for dicynodont phylogeny, biogeography, and biostratigraphy. *J Vert Paleont*, 31(sup1): 1–158
- Kemp T S, 1986. The skeleton of a baurioid therocephalian therapsid from the Lower Triassic (*Lystrosaurus* Zone) of South Africa. *J Vert Paleont*, 6(3): 215–232
- Klembara J, Bartik I, 2000. The postcranial skeleton of *Discosauriscus* Kuhn, a seymouriamorph tetrapod from the Lower Permian of the Boskovic Furrow (Czech Republic). *Trans R Soc Edinb- Earth Sci*, 90(4): 287–316
- Koh T P, 1940. *Santaisaurus yuani* gen. et sp. nov., ein neues Reptil aus der unteren Trias von China. *Bull Geol Surv China*, 20: 73–92
- Kozur H W, Weems R E, 2011. Detailed correlation and age of continental late Changhsingian and earliest Triassic beds: implications for the role of the Siberian Trap in the Permian-Triassic biotic crisis. *Palaeogeogr, Palaeoclimatol, Palaeoecol*, 308(1-2): 22–40
- Li J L, Cheng Z W, 1999. New anthracosaur and temnospondyl amphibians from Gansu, China - the fifth report on Late Permian Dashankou lower tetrapod fauna. *Vert PalAsiat*, 37(3): 234–247
- Li J L, Wu X, Zhang F C, 2008. *The Chinese Fossil Reptiles and Their Kin*. 2nd ed. Beijing: Science Press. 1–473
- Li Y A, Jin X C, Sun D J et al., 2003. Paleomagnetic properties of non-marine Permo-Triassic transitional succession of the Dalongkou Section, Jimsar, Xinjiang. *Geol Rev*, 49(5): 525–536
- Liu J, Li J L, Chen Z W, 2002. The *Lystrosaurus* fossils from Xinjiang and their bearing on the terrestrial Permian-Triassic boundary. *Vert PalAsiat*, 40(4): 267–275
- Liu J, Xu L, Jia S H et al., 2014. The Jiyuan tetrapod fauna of the Upper Permian of China–2. stratigraphy, taxonomical review, and correlation with other assemblages. *Vert PalAsiat*, 52(3): 328–339
- Liu S, 1994. The non-marine Permian-Triassic boundary and Triassic conchostracan fossils in China. *Albertiana*, 13: 12–24
- Lucas S G, 1998. Global Triassic tetrapod biostratigraphy and biochronology. *Palaeogeogr, Palaeoclimatol, Palaeoecol*, 143(4): 347–384
- Metcalfe I, Foster C B, Afonin S A et al., 2009. Stratigraphy, biostratigraphy and C-isotopes of the Permian-Triassic nonmarine sequence at Dalongkou and Lucaogou, Xinjiang Province, China. *J Asian Earth Sci*, 36: 503–520

- Novikov I V, Shishkin M A, 2000. Triassic chroniosuchians (Amphibia, Anthracosauromorpha) and the evolution of trunk dermal scutes in bystrowianids. *Paleontol J*, 34(S2): 165–178
- Novikov I V, Shishkin M A, Golubev V K, 2000. Permian and Triassic anthracosaur from Eastern Europe. In: Benton M J, Shishkin M A, Unwin D M et al. eds. *The Age of Dinosaurs in Russia and Mongolia*. Cambridge: Cambridge University Press. 60–70
- Ouyang S, Norris G, 1999. Earliest Triassic (Induan) spores and pollen from the Junggar Basin, Xinjiang, northwestern China. *Rev Palaeobot Palyno*, 106(1–2): 1–56
- Pang Q, Jin X C, 2004. Ostracoda of the Guodikeng Formation and the continental Permo-Triassic boundary of Dalongkou section, Jimsar, Xinjiang. *Prof Pap Stratigr Palaeont*, 28: 205–246
- Pawley K A T, 2007. The postcranial skeleton of *Trimerorhachis insignis* (Temnospondyli: Trimerorhachidae): a plesiomorphic temnospondyl from the Lower Permian of North America. *J Paleont*, 81(5): 873–894
- Schoch R R, Milner A R, 2014. *Temnospondyli I*. München: Verlag Dr. Friedrich Pfeil. 1–150
- Schoch R R, Voigt S, Buchwitz M, 2010. A chroniosuchid from the Triassic of Kyrgyzstan and analysis of chroniosuchian relationships. *Zool J Linn Soc*, 160(3): 515–530
- Shishkin M A, Novikov I V, Fortuny J, 2014. New bystrowianid chroniosuchians (Amphibia, Anthracosauromorpha) from the Triassic of Russia and diversification of Bystrowianidae. *Paleontol J*, 48(5): 512–522
- Smith R M H, Botha-Brink J, 2014. Anatomy of a mass extinction: Sedimentological and taphonomic evidence for drought-induced die-offs at the Permo-Triassic boundary in the main Karoo Basin, South Africa. *Palaeogeogr, Palaeoclimatol, Palaeoecol*, 396: 99–118
- Sun A L, 1963. The Chinese kannemeyerids. *Palaeont Sin, New Ser C*, 17: 1–109
- Sun A L, 1991. A review of Chinese therocephalian reptiles. *Vert PalAsiat*, 29(2): 85–94
- van den Heever J A, 1994. The cranial anatomy of the early Therocephalia (Amniota: Therapsida). *Ann Univ Stellenbosch*, 1994(1): 1–59
- Vjuschkov B P, 1957. New kotlassiomorphs from the Tatarian of the European part of the USSR. *Tr Paleontol Inst*, 68: 89–107
- Ward P D, Botha J, Buick R et al., 2005. Abrupt and gradual extinction among Late Permian land vertebrates in the Karoo Basin, South Africa. *Science*, 307: 709–714
- Witzmann F, Schoch R R, Maisch M W, 2008. A relict basal tetrapod from Germany: first evidence of a Triassic chroniosuchian outside Russia. *Naturwissenschaften*, 95: 67–72
- Yang J D, Qu L F, Zhou H Q et al., 1986. Permian and Triassic strata and fossil assemblages in the Dalongkou area of Jimsar, Xinjiang. Beijing: Geological Publishing House. 1–262
- Young C C, 1952. On a new therocephalian from Sinkiang, China. *Acta Sci Sin*, 2: 152–165
- Young C C, 1958. On the occurrence of *Chasmatosaurus* from Wuhsiang, Shansi. *Vert PalAsiat*, 2(4): 259–262
- Young C C, 1963. Additional remains of *Chasmatosaurus yuani* Young from Sinkiang, China. *Vert PalAsiat*, 7(3): 215–222
- Young C C, 1979. A Late Permian fauna from Jiyuan, Henan. *Vert PalAsiat*, 17(2): 99–113
- Yuan P L, Young C C, 1934a. On the discovery of a new dicynodon in Sinkiang. *Bull Geol Surv China*, 1934: 563–573
- Yuan P L, Young C C, 1934b. On the Occurrence of *Lystrorhynchus* in Sinkiang. *Bull Geol Surv China*, 1934: 575–580