

# Historical Biology: An International Journal of Paleobiology

Publication details, including instructions for authors and subscription information: <u>http://www.tandfonline.com/loi/ghbi20</u>

The dentition of Manidens condorensis (Ornithischia; Heterodontosauridae) from the Jurassic Cañadón Asfalto Formation of Patagonia: morphology, heterodonty and the use of statistical methods for identifying isolated teeth

Marcos G. Becerra <sup>a b</sup> , Diego Pol <sup>a b</sup> , Claudia A. Marsicano <sup>b c</sup> & Oliver W.M. Rauhut <sup>d</sup> <sup>a</sup> Museo Palaeontológico Egidio Feruglio , Fontana 140, 9100, Trelew , Chubut , Argentina

<sup>b</sup> Consejo Nacional de Investigaciones Científicas and Técnicas (CONICET), Buenos Aires, Argentina

<sup>c</sup> Laboratorio de Paleontología de Vertebrados, Departamento de Ciencias Geológicas , Facultad de Ciencias Exactas y Naturales, Universidad de Buenos Aires , Buenos Aires , Argentina

<sup>d</sup> Bayerische Staatssammlung für Paläontologie und Geologie, Department of Earth and Environmental Sciences, LMU München, Richard-Wagner-Str. 10, 80333, Munich, Germany Published online: 22 May 2013.

To cite this article: Marcos G. Becerra, Diego Pol, Claudia A. Marsicano & Oliver W.M. Rauhut (2013): The dentition of Manidens condorensis (Ornithischia; Heterodontosauridae) from the Jurassic Cañadón Asfalto Formation of Patagonia: morphology, heterodonty and the use of statistical methods for identifying isolated teeth, Historical Biology: An International Journal of Paleobiology, DOI:10.1080/08912963.2013.794227

To link to this article: <u>http://dx.doi.org/10.1080/08912963.2013.794227</u>

### PLEASE SCROLL DOWN FOR ARTICLE

Full terms and conditions of use: http://www.tandfonline.com/page/terms-and-conditions

This article may be used for research, teaching, and private study purposes. Any substantial or systematic reproduction, redistribution, reselling, loan, sub-licensing, systematic supply, or distribution in any form to anyone is expressly forbidden.

The publisher does not give any warranty express or implied or make any representation that the contents will be complete or accurate or up to date. The accuracy of any instructions, formulae, and drug doses should be independently verified with primary sources. The publisher shall not be liable for any loss, actions, claims, proceedings, demand, or costs or damages whatsoever or howsoever caused arising directly or indirectly in connection with or arising out of the use of this material.



# The dentition of *Manidens condorensis* (Ornithischia; Heterodontosauridae) from the Jurassic Cañadón Asfalto Formation of Patagonia: morphology, heterodonty and the use of statistical methods for identifying isolated teeth

Marcos G. Becerra<sup>a,b</sup>\*, Diego Pol<sup>a,b</sup>, Claudia A. Marsicano<sup>b,c</sup> and Oliver W.M. Rauhut<sup>d</sup>

<sup>a</sup>Museo Palaeontológico Egidio Feruglio, Fontana 140, 9100 Trelew, Chubut, Argentina; <sup>b</sup>Consejo Nacional de Investigaciones Científicas and Técnicas (CONICET), Buenos Aires, Argentina; <sup>c</sup>Laboratorio de Paleontología de Vertebrados, Departamento de Ciencias Geológicas, Facultad de Ciencias Exactas y Naturales, Universidad de Buenos Aires, Buenos Aires, Argentina; <sup>d</sup>Bayerische Staatssammlung für Paläontologie und Geologie, Department of Earth and Environmental Sciences, LMU München, Richard-Wagner-Str. 10, 80333 Munich, Germany

(Received 6 March 2013; final version received 5 April 2013)

The recently described *Manidens condorensis* is one of the most completely known taxa of the family Heterodontosauridae from the southern landmasses. However, some dental aspects are not well known due to preservational problems in the type material. This contribution reports new isolated teeth found in the Cañadón Asfalto Formation (Early-Middle Jurassic). These teeth are referred to *Manidens condorensis* based on the presence of autapomorphic characters of the unusual dentition of this taxon, such as the highly asymmetric tooth crowns and small crenulations on each denticles. The isolated crowns are well preserved and reveal the presence of undescribed and new autapomorphical features, including apical and basal wear facets on the occlusal surface of isolated crowns and a wear surface also in the caniniform tooth. We carried out statistical analyses (including morphogeometrical and discriminant analyses), using the holotype crowns as a morphological starting point, for characterising shape variation of the crowns along the toothrow and for identifying the position of isolated crowns. These analyses allow defining morphological regions within the postcaniniform toothrow and produce a metrically based discriminant function to predict the hypothetical position of future discoveries, providing a methodological framework that could be applied to other extinct heterodont dinosaurs.

Keywords: Manidens condorensis; teeth; morphometrics; heterodonty

#### 1. Introduction

Among ornithischian dinosaurs, Heterodontosauridae has been regarded as one of the most phylogenetically problematic clades (Butler 2005; Norman et al. 2011; Sereno 2012). The controversy on the affinities of this group probably stems from the presence of morphological similarities between different groups of ornithischian dinosaurs and Heterodontosaurus (the best known heterodontosaurid), and the scarcity of morphological information is due to the incompleteness and rarity of other heterodontosaurid taxa. This situation led to phylogenetic hypotheses that postulated heterodontosaurids either as advanced ornithischians related to Ornithopoda (e.g. Sereno 1986), as sister taxon to Marginocephalia (Xu et al. 2006) or as one of the most basal ornithischian clades (e.g. Butler, Upchurch, et al. 2008, 2010; Zheng et al. 2009; Norman et al. 2011; Pol et al. 2011; Sereno 2012). Despite the debated affinities of this group of small-bodied cursorial dinosaurs, they have long been characterised by numerous apomorphies, such as a conspicuous caniniform and highly modified heterodont dentition in more evolved heterodontosaurids, which was traditionally regarded as an adaptation to herbivory (e.g. Galton 1973; Weishampel 1984; Weishampel and Norman 1989).

Until recently, Heterodontosauridae was considered to include only Early Jurassic forms from the Elliot Formation of South Africa (*Abrictosaurus consors* [Thulborn 1974], *Lycorhinus angustidens* [Haughton 1924; Thulborn 1970; Gow 1990], *Heterodontosaurus tucki* [Crompton and Charig 1962; Santa Luca 1980]), all of which have a very specialised masticatory apparatus. The temporal and geographical range of this dinosaur group was subsequently extended with the

Actually, although several authors recently cast some doubts on the herbivoral diet on heterodontosaurids (Barrett 2000; Butler, Porro, et al. 2008, 2010, 2012), the currently accepted phylogenetical topologies of Heterodontosauridae and the morphological jaw characterisation of each species show a masticatory specialisation leading to a more efficient chewing mechanism within the group (Pol et al. 2011), and more basal species are related to an omnivoral diet and more evolved species to an herbivoral diet. Furthermore, their early appearance in the fossil record (at least in the Early Jurassic and possibly in the Late Triassic; see Báez and Marsicano 2001) places this group in a critical position for understanding the early diversification of Ornithischia and character evolution at the base of this clade (Butler, Upchurch, et al. 2008).

putative heterodontosaurid affinities of Pisanosaurus mertii (Casamiquela 1967; Bonaparte 1976) and the appearance of fragmentary heterodontosaurid material from the Late Triassic of Argentina (Báez and Marsicano 2001), as well as the possible heterodontosaurid affinities of Echinodon becklesii from the Early Cretaceous of England (e.g. Norman and Barrett 2002; Butler et al. 2012; Sereno 2012). Though the heterodontosaurid relationships of these animals were not unambiguously supported in all recent analyses, the discovery of four new species of undisputed heterodontosaurid affinities has modified the current understanding on the evolution of this group. Recent phylogenetic studies on Heterodontosauridae placed Fruitadens haagarorum (Butler et al. 2010, 2012), Tianyulong confuciusi (Zheng et al. 2009) and Manidens condorensis (Pol et al. 2011) as successive sister taxa of South African heterodontosaurids. More recently, Sereno (2012) described a new heterodontosaurid taxon from the Early Jurassic of South Africa, Pegomastax africanus, which was placed as a sister taxon of Manidens condorensis, and revised the taxonomy, evolution and masticatory function of heterodontosaurids. These new studies offered new data that changed our understanding of heterodontosaurid evolution, including the timing of their radiation, their biogeographic history, and the acquisition and development of adaptations to herbivory in their dentition.

The dental anatomy and phylogenetic position of Manidens condorensis within Heterodontosauridae revealed a previously unknown stage on the evolution of this group of ornithischian dinosaurs (Pol et al. 2011). The heterodontosaurid species from the Southern Hemisphere (Manidens, Pegomastax, Abrictosaurus, Heterodontosaurus and Lycorhinus) differ remarkably from the species of the Northern Hemisphere (Fruitadens, Echinodon and Tianyulong) in their dentition and were recently clustered in the subclade Heterodontosaurinae by Sereno (2012). Although Pisanosaurus (Casamiquela 1967; Bonaparte 1976) and the fragmentary Triassic Argentinean material (Báez and Marsicano 2001) are actually regarded as dubious specimens (Sereno 2012), the interpretation of these fragmentary remains as heterodontosaurids could also represent the oldest record for the family (Bonaparte 1976; Báez and Marsicano 2001; Butler, Upchurch, et al. 2008, 2012). Furthermore, the South African heterodontosaurids have been characterised by the extensive and obliquely disposed wear facets that formed a wide occlusal surface (Hopson 1980; Norman et al. 2011; Sereno 2012), a similar feature is also present in Pisanosaurus and the fragmentary Triassic Argentinean material (Bonaparte 1976; Báez and Marsicano 2001). Although Sereno (1991) noted that the wear facets do not form a continuous surface between teeth such as Heterodontosaurus, the mentioned set of features suggests the presence of an orthal jaw movement with a transverse component and/or a rotation of the lower jaw along its long axis during occlusion, reflecting an advanced stage in the evolution of herbivory (Porro 2007; Holliday

and Witmer 2008; Norman et al. 2011; Sereno 2012). Although both *Pisanosaurus* and the Laguna Colorada specimens share several features with heterodontosaurids (Báez and Marsicano 2001; also see Sereno 1991, 2012; Butler, Upchurch, et al. 2008, 2012) and basal ornithischians (e.g. Irmis et al. 2007), further analysis and more complete specimens are needed to clarify their phylogenetic affinities. However, the taxa from the Northern Hemisphere, which include more recent species, have non-overlapping plesiomorphic crowns (i.e. leaf-shaped, mesiodistally symmetrical teeth with denticles disposed along the margin, similar height of crowns along the toothrow and absence of extensive wear facets [Norman and Barrett 2002; Zheng et al. 2009; Butler et al. 2010]).

In this context, the dental anatomy of *Manidens* condorensis has an important unique combination of characteristics. In the holotype MPEF-PV 3211, the lower postcaniniform toothrow has anteroposteriorly heterogeneous crowns in their height-width proportions that are mesiodistally imbricated, indicating an incipient evolutionary stage in the formation of a compact toothrow. This dental arrangement represents an intermediate stage between the plesiomorphic dentition of northern heterodontosaurids and the highly derived (but early appearing) dentition of some South African heterodontosaurids (e.g. Heterodontosaurus) that have a greater specialisation of their dental anatomy. The pattern of dental transformations related to the increasing adaptation to herbivory differs in heterodontosaurids and other ornithischian groups (Pol et al. 2011), underscoring the unique evolutionary path that led to the sophisticated chewing mechanism of heterodontosaurids.

Here, we describe new and well-preserved isolated teeth that have been found at the same locality and horizon as the holotype of Manidens condorensis. The detailed comparison between the new isolated teeth and the lower teeth of MPEF-PV 3211 revealed the presence of shared features (including several autapomorphies) that allow referral of the isolated teeth to this taxon. These new isolated teeth are much better preserved than those of the holotype and therefore provide new anatomical information to characterise the particular dentition of Manidens condorensis. Given the heterogeneous dentition of this taxon within the toothrow (represented by the lower postcaniniform crowns), we carried out statistical analyses to infer the position of each isolated tooth along the toothrow. This approach furthermore provides predictive functions that will be useful for future discoveries of isolated teeth of Manidens condorensis, and a methodological framework that could be applied to other extinct taxa with heterodont dentition.

#### 1.1 Institutional abbreviation

MPEF-PV, Museo Paleontológico Egidio Feruglio, Trelew, Argentina.

### 2. Systematic paleontology Dinosauria Owen, 1842 Ornithischia Seeley, 1887 Heterodontosauridae Kuhn, 1966 Manidens Pol, Rauhut and Becerra, 2011 Manidens condorensis Pol, Rauhut and Becerra, 2011 (Figures 1–2 and 6)

#### 2.1 Holotype

MPEF-PV 3211, partial associated skeleton, including about 80% of the skull and lower jaws, vertebrae of cervical, dorsal and sacral regions, left scapula and coracoid, and almost complete pelvic girdles (Pol et al. 2011).

#### 2.2 Referred material

MPEF-PV 1719, MPEF-PV 1786, MPEF-PV 1718, MPEF-PV 3810 and MPEF-PV 3811 were originally referred to this taxon by Pol et al. (2011) based on general similarities. MPEF-PV 3812, MPEF-PV 3813, MPEF-PV 3814, MPEF-PV 3815 and MPEF-PV 3816 are new referred remains. All of them are isolated teeth.

#### 2.3 Locality and horizon

The holotype and referred material were collected from the microvertebrate locality Queso Rallado (Rauhut et al. 2002; Rougier, Martinelli, et al. 2007), approximately 5.5 km north-west of the Cerro Cóndor village (see also Pol et al. 2011, Supplementary Material, available online). These outcrops have been included within the Cañadón Asfalto Formation and this particular layer is a silicified mudstone within a series of mudstones and limestones, representing a lacustrine environment (Rougier, Garrido, et al. 2007). The Cañadón Asfalto Formation has traditionally been regarded as Callovian-Oxfordian based on the floral content of this unit (e.g. Tasch and Volkheimer 1970; Silva Nieto et al. 2002, 2003); however, recent radioisotopic dates of this unit yielded more ancient ages (ranging from the late Toarcian to the early Bathonian [Salani 2007; Cabaleri et al. 2010; Cúneo et al. 2013]). These older dates have also been supported by recent and detailed studies of the palynoflora of the Cañadón Asfalto (Volkheimer et al. 2009; Zavattieri et al. 2010; for additional discussion of the geology and age, see Cuneo et al., in press). The age of this unit is therefore currently regarded as ranging from the latest Early Jurassic to the Early-Middle Jurassic. Because the Queso Rallado locality is placed in the basal part of the formation, a late Early Jurassic age for *Manidens* is likely.

#### 2.4 Emended diagnosis

Small heterodontosaurid (estimated body length of approximately 50-60 cm) with the following autapomorphies: jugal with strongly developed, dorsally placed lateral boss; dorsal part of the postorbital process of the jugal very slender and flexes abruptly posteriorly at the beginning of the articular facet for the postorbital; forebrain facet on the ventral surface of the frontal enlarged and with significantly raised margins; external mandibular fenestra absent; posterior teeth with asymmetric arrangement of denticles and with a mesial concavity in which the distal margin of the preceding tooth is lodged (as in cerapodan ornithischians; e.g. Chinnery et al. 1998; Rauhut 2001; Galton 2007); the posteriormost dentary tooth 'hand shaped' in lateral view, with only one or two mesial denticles, the most anterior of which diverges mesially from the mesial margin of the crown (acting like a 'thumb' in the 'hand-shaped' crown) and four to five distal denticles apically to distally oriented (acting like the remaining 'fingers' in the 'hand-shaped' crown); the presence of small crenulations along the cutting margin of each denticle, exclusively formed by the enamel (the underlying dentine is unaltered); antorbital fossa reaches the jugal anterior process, deviating below the jugal boss; mesial denticulate margin approximately 60% of the length of the distal margin; apical and basal wear facets on the labial surface of the dentary crowns that are vertically oriented.

#### 2.5 Referral of isolated teeth to Manidens condorensis

The new isolated teeth and those of the dentary of the holotype of Manidens condorensis (MPEF-PV3211) share several diagnostic morphological characters. These include mesiodistally asymmetrical, labiolingually compressed and leaf-shaped crowns that bear a central, conspicuous apicobasal ridge and a few slightly developed apicobasal ridges on both surfaces of the crown. These ridges originate at the base of each denticle, extend basally along the surface of the crown and disappear before reaching its base. The margin of the crowns has morphologically subequal denticles that are apicodistally or distally oriented (except the mesialmost denticle). In addition, the isolated teeth share the dental autapomorphies of *Manidens condorensis* noted by Pol et al. (2011), such as the presence of a mesial groove, delimited by two ridges, denticles asymmetrically disposed on the tooth margin (with one or two mesial denticles and four to six distal denticles), the mesialmost denticle mesially projected above the mesial margin and small enamel crenulations on the margins of the denticles. The presence of these features allows us to refer these crowns to Manidens condorensis (crown pictures showed in Pol et al. 2011, Figure 2(e)-(i) and the presence of wear facets allows us to infer some aspects of the masticatory



Figure 1. Main morphological features of the crowns of *Manidens condorensis*. (A) Occlusal view of MPEF-PV 3813; (B) distal view of specimen MPEF-PV1719; (C) occlusal surface of crown of MPEF-PV3812, boxes show the areas with wear facets with detailed the SEM images (D, F and G); (D) detail of apical wear facet of MPEF-PV 3812, in which the dentine is not exposed; (E) detail of apical planar wear facet of MPEF-PV 1786, both enamel and dentine are worn; (F) detail of apical concave wear facet of MPEF-PV 3812, exposing both enamel and dentine, (G), basal planar wear surface of MPEF-PV 3812; (H and I) lower caniniform tooth of the holotype MPEF-PV 3211. Scale: 1 mm.

movement, in this case predominantly orthal. Finally, due to the absence of a lingual view of dentary crowns and the lack of adequately preserved crowns in the maxilla of MPEF-PV 3211, the identification to the maxillary or dentary toothrow of isolated crowns is not possible at the moment.

## 2.6 New anatomical information on the dentition of Manidens condorensis

The well-preserved isolated teeth provide new information on the dentition of Manidens condorensis. These teeth show that the labial and lingual surfaces of the crown differ in their basal extension, the development of their apicobasal ridges and the basal extension of the apicobasal ridges. In occlusal view, both surfaces of the crown are mesiodistally convex (related to the development of the central ridges; Figure 1(A)), and the surface that is more basally extended is also slightly apicobasally concave in distal view (related to a basally bulbous crown; Figure 1(B)). Finally, and more importantly, the isolated crowns reveal new autapomorphic features related to the presence of unusually developed wear facets. Two distinct types of wear facets are present in the more basally extended surface of the preserved teeth: one is located apically on the crown (on the denticles) and the other is a flat wear facet located at the base of the crown (Figure 1(C)-(G)). The apical wear facets extend on the surface of the denticles and are mostly flat to slightly concave and vertically oriented. The facets expose only the enamel or both enamel and dentine, depending on the stage of wear. The facets of the different denticles of the same crown are coplanar to each other, suggesting that they were produced by tooth-tooth occlusion (Figure 1(C)-(E)). A few facets have the dentine surface slightly apicobasally concave, which may have been produced by differential wear by food abrasion (Figure 1(F); see Costa and Greaves 1981). The basal wear facets are also interpreted as being caused by tooth-tooth occlusion, as they form a simple, laterodistally oriented planar surface (Figure 1(G)). The isolated crowns and the holotype crowns show insufficient evidence to identify maxillary and dentary crowns, and almost all isolated crowns reveal the presence of wear facets (and evidence for dental occlusion) in the apicobasally higher side of the crown. Based on the observed asymmetry in the available teeth and the fact that asymmetric teeth can be inversely oriented in the mandible and maxilla among ornithischians (specular disposition of features on the faces of opposing crowns, being the lingual face of maxillary crowns similar to the labial face of denary crowns and the labial face of maxillary crowns similar to the lingual face of dentary crowns; e.g. Dryosaurus lettowvorbecky Norman, 2004; Hypsilophodon foxii Galton, 1974), here we assume that the apicobasally higher face represents the surface that came into contact with the opposing teeth during the occlusion (because is generally worn), and that there is no marked morphological differentiation between the maxillary and dentary crowns. With this, to save words to refer to each surface of the crown during this study, and counting on that the differentiation of maxillary and dentary crowns remains unsolved (including also the differentiation in lingual and labial views of each crown), the worn surface of the crown is here named as 'occlusal surface', whereas the opposite unworn surface is here called 'non-occlusal surface', rather than labial or lingual. Finally, the Scanning Electron Microscope images of the caniniform tooth MPEF-PV 3211 revealed the presence of a planar and continuous worn surface that is labiomesially oriented. This facet is limited mesially by a smooth border and both basally and apically by a fracture (Figure 1(H)-(I). Apically worn caniniforms have been reported for Lycorhinus and Heterodontosaurus (Gow 1990; Norman et al. 2011), but Sereno (2012) cast doubts on this particular feature. In Manidens, the wear surface is planar, continuous and does not reach the apex of the crown (contrasting with the alleged wear facets of the other taxa). Given that, in other heterodontosaurids, the dentary caniniform tooth is lodged posterior to the premaxillary caniniform tooth during occlusion, this facet might reflect tooth-to-tooth occlusion of the caniniform teeth at least during some part of the mastication cycle.



Figure 2. Dental material preserved in the holotype of *Manidens condorensis* (MPEF-PV 3211). (A) General view, boxes show the crowns detailed in B and C; (B) last five dentary postcaniniform crowns; (C) first three dentary postcaniniform crowns. Scales in A, 1 cm; B and C, 1 mm. Mesial to the right, all images in labial view (mentioned also as occlusal surface).

#### 3. Materials and methods

The specimens used in this study include the best preserved postcaniniform crowns of the dentary of the holotype of *Manidens condorensis* (MPEF-PV 3211; Figure 2), which are the second, third, seventh and ninth postcaniniform teeth, and all the referred isolated crowns (MPEF-PV 1719, MPEF-PV 1786, MPEF-PV 1718, MPEF-PV 3810, MPEF-PV 3811, MPEF-PV 3812, MPEF-PV 3813, MPEF-PV 3814, MPEF-PV 3815 and MPEF-PV 3816; see Supplementary Material, available online).

For the geometric morphometric analysis (GMA), the photos were first treated with MakeFan6 software (Sheets 2003) to draw guidelines to help in the determination of landmarks and semilandmarks. The specimen image files (TPS files) were generated using tpsUTIL 1.46 (Rohlf 2010a). The digitalization and scaling of landmarks and semilandmarks were performed out using tpsDIG 2.16 (Rohlf 2010b). Landmark and semilandmark configurations were superimposed using generalised procrustes analysis (GPA; Goodall 1991; Rohlf 1999) to eliminate the distorting effects caused by different factors (translation, scaling, rotation), following the least squares criterion (the sum of the squared distances between the corresponding points is minimised). The semilandmarks were slid using the minimum bending energy criterion (Bookstein 1997), which consist of sliding the semilandmarks until the strain energy is minimised, softening the deformation between curves (Pérez et al. 2006). The aligned coordinates of landmarks and semilandmarks of all specimens were compared using a thin-plate spline function (Bookstein 1989). The partial deformation (partial warps) and principal components (relative warps) were calculated using tpsRELW 1.49 (Rohlf 2010c). Further details of this analysis are given in the Supplementary Material, available online.

The discriminant analysis (DA) was carried out using the Infostat free version program (Di Rienzo et al. 2011). The measurements of each metrical variable used in the DA were obtained using a digital caliper (accuracy 0.01 mm).

## 4. Morphological variation in the toothrow of *Manidens condorensis*

The heterodont dentition of Manidens condorensis includes a lower caniniform teeth and a morphologically variable postcaniniform toothrow that is almost completely preserved in the dentary of the holotype material. The tooth count is at least 11 in the dentaries, including the hypertrophied anterior caniniform (a low tooth count as in other heterodontosaurids [Pol et al. 2011], including the closely related taxon *Pegomastax* [Sereno 2012]). The caniniform is the largest tooth of the dentary. It is a stout element, with a recurved apex, and is slightly compressed labiolingually. This tooth seems to lack marginal serrations, in contrast to the caniniform teeth of the South African heterodontosaurids Heterodontosaurus, Lycorhinus, Abrictosaurus and Pegomastax (Charig and Crompton 1974; Hopson 1975; Sereno 2012). However, an unserrated carina is present distally, but apparently not mesially, although the apical wear opens the possibility that a short carina might have been present along the apical part. The postcaniniform crowns vary transitionally along the toothrow in shape, size and proportions. Basically, the first three postcaniniform crowns are rhomboidal-shaped small crowns, but distal to these elements the crowns increase in size rapidly and become mesiodistally expanded. The teeth in the middle of the toothrow are higher apicobasally than broad mesiodistally, with the highest tooth crowns being more than 1.5 times higher than broad. Especially, high tooth crowns were traditionally considered to be a synapomorphy of heterodontosaurids (e.g. Weishampel and Witmer 1990; Norman et al. 2004), but the crowns are considerably lower in the basal heterodontosaurids Echinodon, Fruitadens, Tianyulong and an unnamed taxon from the Early Jurassic Kayenta Formation (Norman and Barrett 2002; Zheng et al. 2009; Butler et al. 2010, 2012; Sereno 2012), so that such a high crown height/crown width ratio represents a synapomorphy shared by Manidens and the South African forms (Heterodontosaurinae sensu Sereno 2012). Finally, the last two crowns decrease in height, but not in mesiodistal width. Due to preservation, the maxillary toothrow in MPEF-PV 3211 includes only four apically incomplete crowns, which are labially and mesially worn. The



Figure 3. (A) Definition of landmarks and semilandmarks for the morphogeometrical analysis, including schematic representation of landmarks (grey) and semilandmarks (white) position: 1, basalmost point of the mesial margin; 2, apex of the first denticle; 3, apex of the last denticle; 4, basalmost point of the distal margin; sl1 and sl2, semilandmarks describing the margin between two landmarks. (B) Variables measured on each crown for the DA: hdm, height distal margin; mh, maximum height; hmm, height mesial margin; bw, basal width; mw, maximum width; a, apical; m mesial. The arrows represent a view of the occlusal surface of the crown.

preserved maxillary crowns vary slightly in height. On the basis of the available information, we assume that the observed variation within the dentary toothrow was also present in the maxillary toothrow. Thus, the lower toothrow present in MPEF-PV 3211 is here used as a proxy for representing the shape variation for both maxillary and dentary toothrows.

The variation of these characteristics along the toothrow is also observed in isolated crowns referred to *Manidens condorensis*, and could therefore provide useful information to identify their position. This is of particular interest, given that the isolated crowns are much better preserved than those of the holotype (see Figure 2), so that a detailed identification of the position of these teeth will improve our knowledge on the available morphological information on the peculiar dentition of *Manidens condorensis*. In the following section, we use morphometric data to perform two statistical analyses to test the identification of the position of the new isolated teeth using the postcaniniform mandibular teeth of the holotype as a reference.

## 5. Morphometric characterisation of the toothrow of *Manidens condorensis*

Although we only have 14 teeth (data points) for the two statistical analyses (including the isolated teeth and those of the holotype), our aim is to provide a quantitative approach to characterise the different regions of the postcaniniform toothrow, rather than achieving the values of statistical significance for the inference made upon these variables (for which a much larger sample would be needed). We first conducted an exploratory GMA, which aims to search for the existence of morphological clusters. Second, we conducted DA with metrical variables to reevaluate the nature of the clusters of teeth identified in the GMA. Third, we obtained a canonical function, which will allow identifying the position of future findings of isolated teeth based on the morphometric data.

#### 5.1 Geometric morphometric analysis

The described transitional change within the toothrow of the holotype is reflected in the morphological variation in the proportions of the isolated teeth here reported, as each of these isolated crowns shares features with only a few crowns of the holotype. The differences between specimens were measured by defining landmarks and semilandmarks. The combination of these variables in the studied specimen is used to define a 2D morphospace on the basis of the shape variation (Bookstein 1991, 1997; Richtsmeier et al. 2002). Based on the premise that the similarity between isolated teeth and those attached to the dentary of the holotype will be reflected in the location of the specimens in the morphospace, this method should show morphological clusters that would relate crown morphology and dental position along the toothrow. The GPA method ensures that the variation explained by size is excluded from the total variance, thus reducing the variance to that explained by shape differences.

The landmarks and semilandmarks were defined following a mesiodistal arrangement within the crown. Four landmarks and 10 semilandmarks were used, which summarise the morphological information available from the occlusal (apicobasally higher) surface of the crown (Figure 3(A)). Theoretically, each landmark and group of semilandmarks should represent homologous structures. In this particular analysis, the homology of each point corresponds to a positional homology within the tooth (see Figure 3(A)). Although the isolated crowns are excellently preserved, some of the holotype crowns have a damaged apex and only have their labial (occlusal) surface exposed (Figure 2(B)). Because of this, we did not define a landmark for the tooth apex or on the non-occlusal surface of the crowns.

#### 5.1.1 Results

The results of the GMA show that the first two components explain 89.5% of the total variance among the specimens, determining three different clusters (Figure 4). The first component (Rel1 in Figure 4) explains 76.97% of the total variance. This variation implies towards positive values of this axis, the relative increase of the mesiodistal width (an increase in the distance between landmarks 1 and 4, and 2 and 3), a decrease in the distance between landmarks 1 and 2, and a decrease in the mesial projection of mesial denticle (captured by semilandmarks between landmarks 1



Figure 4. Morphometric analysis results, modified to clarify the three different groups. A–D show the shape of most extreme values of each axis in grey (A and B in axis *X*, C and D in axis *Y*) and the consensus in black (shape in the origin), the vectors show the morphological change: 1, MPEF-1718; 2, MPEF-1719; 3, MPEF-1786; 4, MPEF-3810; 5, MPEF-3811; 6, MPEF-3812; 7, MPEF-3813; 8, MPEF-3814; 9, MPEF-3815; 10, MPEF-3816; H11, holotype, ninth crown; H12, holotype, seventh crown; H13, holotype, third crown; H14, holotype, second crown.

and 2; see Figure 4(A),(B)). On the basis of the information provided only by this component, there is a clearly separated group formed by the anterior teeth of the holotype (H13, H14; Figure 4) and one of the isolated teeth (9; Figure 4). The other teeth of the holotype (H11 and H12; Figure 4) are located towards negative values of this axis, sowing the change from the lower and wider anterior crowns, to the higher and narrower crowns of the medial and posterior portion of the lower toothrow of *Manidens*. Most of the isolated crowns (1–8 and 10) are located along the first component (Rel1) between the mid to posterior teeth of the holotype (H11 and H12).

The second component (Rel2 in Figure 4) explains 12.53% of the total variance. This variation includes the posterior projection of the apical region of the crown (landmarks 2 and 3) with respect to the base of the crown (landmarks 1 and 4) and the increase in the height of the distal margin (hdm) of the crown (progressive separation of landmarks 3 and 4).

#### 5.1.2 Morphological clusters

Based on these results and the location of all teeth along these two axes, the specimens analysed are classified into three different groups or morphological clusters. This clustering is mainly explained by the difference in height of the mesial margin (hmm) and hdm of the crowns, which vary markedly along the toothrow of *Manidens*. In the left sector of the graphic, the crowns are higher than wide and have the mesial margin apicobasally higher than the distal margin (circles in Figure 4). Towards the right, the crowns are wider than high (with a smaller difference between the hdm and hmm; triangles in Figure 4). Finally, the low crowns with distal and mesial margins that are subequal in height are located at the right end of the graphic (squares in Figure 4).

The location of the crowns of the holotype in this graph helps allocating the isolated teeth into three distinct clusters that are morphometrically distinguishable. These clusters are referred herein as anterior crowns (squares in Figure 4), middle crowns (circles in Figure 4), and posteriorly placed crowns (triangles in Figure 4), representing the three regions of the toothrow observed in the crowns of the holotype.

The groups defined from the GMA can be further analysed through DA based on the simple morphometric measures to test if simple measurements can be used to easily identify the position of isolated teeth, which would be useful for the new discoveries of isolated teeth of *Manidens condorensis*.

#### Discriminant analysis 5.2

DA uses the clusters identified through the GMA and a set of simple morphometric measures to test if the latter are informative to infer the position of isolated teeth in the toothrow of Manidens (using as comparative source those teeth found in the dentary of the holotype). Working with more easily measurable variables facilitates expanding these results to determine the position of future discoveries.

The metrical variables of the occlusal surface of the crown used for the DA are basic height and width linear dimensions of different regions of the crown (Figure 3(B)). For the two apically incomplete anterior crowns on the holotype lower toothrow (Figure 2(C)), the values of maximum height (mh) could not be measured. To circumvent this problem, we carried out a sensitivity analysis by varying the estimated mh value of the incomplete crowns extending up to 0.5 mm beyond the preserved apical region, to test how the uncertainty on the height of these elements (20% and 24% of the preserved length of the crown) affects the results of the DA. The results of these measured variations were tested in multiple DAs and we corroborated that it does not affect the significance of the identification and the contribution of each variable to the delimitation of clusters.

On the basis of this result, we used the maximum preserved height, ignoring the apical incompleteness of these two anterior crowns. All new variables were measured in both the holotype and the isolated crowns. The values of each metrical variable are detailed in the Supplementary Material, available online.

#### 5.2.1 Results

The first canonical axis of the DA explains 63.13% of the total variance (eigenvalue, Table 1(C)) and it shows a clear separation of two cloud points: the crowns placed in the middle and anterior region of the holotype toothrow, on the one hand (circles and squares in Figure 5(B)), and the crowns placed in the posterior region of the holotype, on the other hand (triangles in Figure 5(B)). The clustering of the isolated crowns with those of the holotype in the DA (Figure 5B) is consistent with that of the GMA (Figure 4). All the measured variables contribute to the first canonical axis, but the largest contribution is given by maximum width (mw) and mh (see Figure 5(A)). The mw relates directly to the values of this canonical axis, but the mh does so inversely (Figure 5(A)). This means that crowns with high positive values for the canonical axis 1 have the

Table 1. DA results. A. Centroids in the discriminant space Groups Axis 1 Axis 2 -1.073.02 1 2 -1.83-1.26 3 2.84 0.3 B. Canonical discriminant functions 2 1 Constant -1.323.78 -7.48-0.85mh 1.45 1.84 hw mw 7.85 -1.263.2 -1.68hmm 1.97 hdm -2.68C. Eigenvalues % Accumulated % Eigenvalues 63.13 63.13 5.8 3.39 36.87 100 D. Cross-classification table 2 Error (%) Group 1 3 Total 3 0 0 3 0 0 2 6 0 6 0 0 0 3 0 5 5 Total 3 6 5 14 0

Notes: A, centroid coordinates of each cluster in the discriminant space; B, coefficient values for the canonical function; C, variance percentage explained by each canonical axis or eigenvalues; D, resulting table showing how many specimens of each group were correctly assigned. Group 1, 'anterior region' crowns; group 2, 'middle region' crowns; group 3, 'posterior region' crowns.



Figure 5. DA results, each point corresponds to a single specimen, each line indicates the relative contribution of a single variable in proportion. Squares refer to the specimens included in the anterior region; triangles refer to the posterior region; circles refer to the middle region; the black circles refer to the centroid *C* of each group. (A) hdm, height distal margin; mh, maximum height; hmm, height mesial margin; bw, basal width; mw, maximum width. (B) Disposition of isolated tooth and holotype material: 1, MPEF-1718; 2, MPEF-1719; 3, MPEF-1786; 4, MPEF-3810; 5, MPEF-3811; 6, MPEF-3812; 7, MPEF-3813; 8, MPEF-3814; 9, MPEF-3815; 10, MPEF-3816; H11, holotype, ninth crown; H12, holotype, seventh crown; H13, holotype, third crown; H14, holotype, second crown. The symbol ' + ' marks the point where values of both axes are zero.

largest values of maximum crown width and lowest mh (i.e. wide-based low crowns). This morphology characterises the posterior crowns (triangles in Figure 5). In contrast, the crowns placed in the middle and anterior regions are proportionally higher crowns with lower values of mw, or crowns higher than wide (circles and squares in Figure 5). In particular, the mid-positioned crowns (circles in Figure 6) are displaced towards the negative side of this axis.

The second canonical axis accounts for the remaining 36.87% of the total variance (Table 1(C)), and the greatest contribution is given by the hdm (directly related to the axis) and hmm (inversely related to the axis), and by a contribution of the variable basal width (bw; directly related to the axis). Towards higher values of the second axis, the mesial height becomes lower and the hdm and bw become larger (Figure 5(A)). Basically, this axis is related to the changes of mesial and distal margins (anterior crowns; squares in Figure 5(B)) from those with different hdm and hmm (middle and posterior crowns; circles and triangles in Figure 5(B)).

#### 5.2.2 Summary

The combination of the two axes corroborates the distinction of the three pre-defined clusters and places all isolated crowns in the same cluster as in the GMA (Table 1(D)). These results demonstrate the usefulness of simple metrical variables to identify the position of isolated teeth of the toothrow of *Manidens condorensis*. Although it is clear that *Manidens condorensis* has a transitional variation in the crown proportions along the toothrow (as all other heterodontosaurids, except for *Tianyulong* [Zheng et al. 2009]), the statistical analysis here implemented

allows us distinguishing three different regions that are useful for descriptive and comparative purposes, helping in the identification of future discoveries of isolated material, and is based on morphological features that vary along the toothrow.

## 6. Regionalization of postcaniniform toothrow of *Manidens condorensis*

The results obtained help defining three regions in the postcaniniform elements of the toothrow of *Manidens* based on their proportions and the lower dentition of the holotype. We describe here the major morphological features of these three regions, identifiable in the lower jaw and hypothetically present in the maxilla.

#### 6.1 Anterior region

These crowns of the anterior region (Figure 6(A)-(D)) are characteristically higher than wide (DA), their mesial margin is much shorter apicobasally than the total apicobasal height of the crown (in comparison with this proportion in more posterior teeth). In addition, in anterior teeth the mesial margin is slightly higher than the distal margin (GMA; Figure 6(A)-(D)). Three teeth are included in this region, two apically incomplete crowns (i.e. the two mesialmost preserved crowns of the holotype MPEF-PV 3211; Figure 6(A)) and one isolated and well-preserved crown (MPEF-PV 3815; Figure 6(B)-(D)). The crowns of the holotype lack their apex, but the isolated crown MPEF-PV 3815 shows that the apical region is high in comparison with the mesial and distal margins (Figure 6(B)-(D)).

The well-preserved isolated crown also differs morphologically from more posterior crowns in other characters not considered in the morphometric analyses. This anterior crown is labiolingually narrow and the mesial cavity is barely developed, both apically and labiolingually. The mesial cavity does not reach the base of the first denticle, which is consistent with the wellspaced arrangement of anterior crowns in the lower toothrow of the holotype of Manidens. The anterior teeth have two mesial denticles and four distal denticles, less than in middle and posterior teeth. The mesial denticles are less prominent in anterior teeth than in middle and posterior teeth and the mesialmost denticle is projected apically. The apex of the anterior tooth (MPEF-PV 3815) is more clearly separated from the marginal denticles than in middle and posterior teeth (Figure 6(B),(D)). The ridge of the occlusal surface extending basally from the apex is broader than in middle and posterior teeth, approximately occupying the mesial half of the occlusal surface of the crown. Finally, the apical region of the crowns of anterior teeth is not posteriorly recurved as in the middle and posterior regions. These differences help characterising the morphological features of the anterior crowns of Manidens condorensis.

#### 6.2 Middle region

The teeth included in this group have a similarly high ratio of mh to mw (DA) as the anterior crowns, but differ from them in a larger difference between the hmm and hdm (GMA). These teeth also have the largest apicobasal height of the lower toothrow (considering both the maximum crown height and the hmm).

The crowns of the middle region also differ from other teeth in characters not considered by the morphometric analyses, such as having the apicobasally highest crowns and a deeper mesial cavity than all other elements of the toothrow (Figure 6(E)-(G)). The development of a deep mesial cavity is related to the tight marginal contact between adjacent crowns in the middle region of the toothrow. The apicobasally high crowns of the middle region also have more denticles on their distal margin than other teeth. There are five well-developed denticles and an incipiently developed denticle that is partially merged with the apex of the tooth (e.g. MPEF-PV 3813; see Supplementary Material, available online). The mesial margin of the teeth of the medial region bears two denticles, as well as an incipiently developed denticle that is partially merged with the apex of the tooth (e.g. MPEF-PV 3813; see Supplementary Material, available online). The two mesial denticles of these teeth are well spaced from each other and differ in shape and size. The basalmost of these two denticles is more conspicuous in comparison with those of anterior teeth and projects apically (with only a slightly developed mesial deflection), whereas the other denticle is small, as those of anterior teeth.

#### 6.3 Posterior region

The crowns of the posterior region have a lower ratio of crown height/bw than preceding elements of the toothrow (DA). The mesial margin is apicobasally higher than the distal margin, but not as much as in the crowns of the middle region (GMA). The crowns of the posterior region



Figure 6. Pictures of representative specimens of the defined regions using GMA and DA analyses; anterior (A–D), middle (E–H) and posterior region (I–L), including isolated and holotype crowns. Some images were rotated to allow orientation. (A) Third holotype crown preserved, labial view, the punctuated line represents the non-preserved apical region; (B–D) specimen MPEF-3815 in occlusal surface, mesial and non-occlusal surface view, respectively; (E) seventh holotype crown preserved in labial (occlusal surface) view; (F–H) specimen MPEF-1719 in occlusal surface, mesial and non-occlusal surface view, respectively; (I) ninth holotype crown preserved in labial (occlusal surface) view; (J and K), specimen MPEF-1786 occlusal surface and mesial views; (L) specimen MPEF-1718 in non-occlusal surface view. Scales: 1 mm. In mesial view, occlusal surface on right; in non-occlusal and occlusal surface view, mesial on right.

also differ from other elements of the toothrow in characters not considered by the morphometric analyses, such as having the labiolingually broadest mesial cavity (Figure 6(K)). The number of denticles on the distal margin of the posterior teeth is either five or four, and they differ from those of the anterior and middle region in being more distally projected and in having deeper interdenticular slits. There are two well-developed mesial denticles in the posterior teeth. These also differ from those of other regions in having deeper interdenticular slits, and the basalmost denticle of the posteriormost teeth has an apicomesial projection (MPEF-PV 3211; Figure 6(I)). The posterior region of the dentition of Manidens has the strongest imbrication of adjacent crowns, which is favoured by both the remarkably broad mesial cavity (Figure 6(K)) and the anterior projection of the basalmost mesial denticle and the posterior convexity of the distal margin of the crown (Figure 6(L)).

#### 7. Applicability to new discoveries

The statistical analyses carried out help assigning isolated crowns to a hypothetical position within the toothrow. The DA also formulates a discriminant function that maximises the explained variance to form the previously obtained clusters using the defined metrical variables as a starting point. This discriminant function indicates the contribution of each variable to the explained variance in each canonical axis (obtained from the eigenvectors; see Table 1(B)). The simple metric variables used in this analysis can easily be measured in new isolated lower teeth of *Manidens* to determinate its coordinates in each canonical axis from the canonical function (see Table 1). This will provide a quantitative justification for identifying the approximate position along the toothrow.

The component of the variance explained by canonical axis 1 can be determined by the following equation:

$$Z_x = -1.32 - (7.48 \times \text{mh}) + (1.45 \times \text{bw}) + (7.85 \times \text{mw}) + (3.2 \times \text{hmm}) - (2.68 \times \text{hdm}).$$

And the component of the variance explained by canonical axis 2 can be determined by the following equation:

$$Z_y = 3.78 - (0.85 \times \text{mh}) + (1.84 \times \text{bw}) - (1.26 \times \text{mw})$$
$$- (1.68 \times \text{hmm}) + (1.97 \times \text{hdm}).$$

#### 8. Discussion

The morphological variation along the postcaniniform toothrow of *Manidens condorensis* (Becerra et al. 2011;

Pol et al. 2011) is clearly transitional (i.e. anteroposterior variation of height and width of the crowns), but the morphological and statistical analyses carried out here allow to go further and distinguish three morphologically distinct regions. This division of the dentary toothrow is not only useful for descriptive purposes but also enables the assignment of isolated crowns of Manidens to the anterior, middle or posterior regions of the toothrow, based on the morphometric characteristics. Although the use of statistical analyses for identifying the position of isolated crowns provides only a hypothesis based on the currently available evidence, it is worth noting that we have reached consistent results using three different approaches: the identification of discrete morphological characters, the use of landmarks and semilandmarks in geometric morphometrics and simple metric measures in the DA. Furthermore, the DA also provided a canonical function that will facilitate the identification of the position of new isolated teeth of Manidens condorensis. Finally, depending on the statistical method, the three regions of the postcaniniform toothrow of Manidens condorensis can be distinguished by the ratio of mh to mw (DA) or by the difference between hmm and hdm (GMA). This opens the possibility of applying one or the other method depending on the measures or landmarks that are available in incompletely preserved teeth.

Based on these results, working with statistical methods on isolated teeth can increase knowledge of a particular species or even operate as a tool for taxonomic assignments at higher levels. In an example, the study of isolated teeth dinosaur taxa provides difficulties in terms of their taxonomic assignment and, in many cases, such elements can only be referred to higher taxonomic groups (e.g. Currie et al. 1990; Chinnery et al. 1998; Barrett and Wang 2007; Ruiz-Omeñaca et al. 2010; Saegusa and Tomida 2011). However, the highly variable dental morphology of ornithischian taxa allows in some cases (with the corresponding morphological comparison) reaching the identification of isolated crowns up to the species level, although many of these taxa are highly incomplete and either have uncertain phylogenetic affinities or have been subsequently regarded as nomina dubia (e.g. Alocodon kuehnei, Trimucrodon cuneatus, Phyllodon henkeli [Thulborn 1973], Siluosaurus zhanggiani [Dong 1997], Iguanodon ottingeri [Galton and Jensen 1979]). A similar approach with useful results have been conducted by Smith et al. (2005) on identifying higher taxonomic levels for isolated theropod teeth, showing the applicability of this approach even if teeth are not diagnostic at the species level (as in the case of theropods).

Finally, in some ornithischian groups (as well as other groups of extinct archosaurs), there is a marked variation in the proportions and morphology of the crowns along the toothrow (e.g. *Heterodontosaurus*, *Abrictosaurus*, *Lycorhinus*, *Lesothosaurus*, *Gasparinisaura*, *Orodromeus*, *Hyp-* *silophodon*). The quantitative approach developed in this contribution for the study of isolated teeth of *Manidens condorensis*, therefore, has the potential of being applied in other heterodont taxa for inferring the position of isolated crowns. In many cases (such as the one of *Manidens*), this can help achieving a more complete and thorough knowledge on the dentition of some ornithischian dinosaurs, providing critical data for understanding the paleoecology and evolution of one of the two major groups of herviborous dinosaurs.

#### Acknowledgements

The authors of this publication thank their family and friends; Mariano Ramirez and Guillermo Rougier; Leandro Canessa for the careful preparation of the specimens described here; Aluar Aluminio Argentino SAIC for access to the SEM lab as well as the valuable help of Mr Jaime Groizard. This research was supported by the following grants: PICT 1756 and PICT 0808 (DP) from ANPCyT, International Collaboration grant CON-ICET-Chinese Academy of Sciences (DP), DFG RA1012/9-1 (OWMR), the project 'Paleontological Exploration of Patagonia' [Fundación Antorchas/Lampadian Foundation (GWR) and NSF DEB 0946430 and DEB 1068089 (GWR)] and UBACyT 20020100100728 (CAM). This study is CAM's R-83 contribution to the Instituto de Estudios Andinos Don Pablo Groeber (IDEAN-CONICET).

#### References

- Báez AM, Marsicano CA. 2001. A heterodontosaurid ornithischian dinosaur from the Upper Triassic of Patagonia. Ameghiniana. 38:271–279.
- Barrett PM. 2000. Prosauropod dinosaurs and iguanas: speculations on the diets of extinct reptiles. In: Sues HD, editor. Evolution of herbivory in terrestrial vertebrates: perspectives from the fossil record. Cambridge: Cambridge University Press. p. 42–78.
- Barrett PM, Wang XL. 2007. Basal titanosauriform (Dinosauria, Sauropoda) teeth from the Lower Cretaceous Yixian Formation of Liaoning Province, China. Palaeoworld. 16:265–271.
- Becerra MG, Pol D, Marsicano CA. 2011. New isolated teeth of Manidens condorensis from the Cañadón Asfalto Formation: exploring new methods of morphological analysis. Paper presented at: IVCLPV 2011. IV Congreso Latinoamericano de Paleontología de Vertebrados; San Juan, Argentina
- Bonaparte JF. 1976. Pisanosaurus mertii Casamiquela and the origin of the Ornithischia. J Paleontol. 50:808–820.
- Bookstein FL. 1989. Principal warps: thin-plate splines and the decomposition of deformation. IEEE Trans Pattern Anal Mach Intell. 11:423–437.
- Bookstein FL. 1991. Morphometrics tools for landmark data. Geometry and biology. New York: Cambridge University Press.
- Bookstein FL. 1997. Landmark methods for forms without *landmarks*: morphometrics of group differences in outline shape. Med Image Anal. 1:225–243.
- Butler RJ. 2005. The 'fabrosaurid' ornithischian dinosaurs of the Upper Elliot Formation (Lower Jurassic) of South Africa and Lesotho. Zool J Linn Soc Lond. 145:175–218.
- Butler RJ, Galton PM, Porro LB, Chiappe LM, Henderson DM, Erickson GM. 2010. Lower limits of ornithischian dinosaur body size inferred from a new Upper Jurassic heterodontosaurid from North America. Proc R Soc B Biol. 277:375–381. doi: 10.1098/rspb.2009.1494.
- Butler RJ, Porro LB, Galton PM, Chiappe LM. 2012. Anatomy and cranial functional morphology of the small-bodied dinosaur *Fruitadens haagarorum* from the Upper Jurassic of the USA. PLoS One. 7:e31556. doi: 10.1371/journal.pone. 0031556.

- Butler RJ, Porro LB, Norman DB. 2008. A juvenile skull of the primitive ornithischian dinosaur *Heterodontosaurus tucki* from the 'Stormberg' of southern Africa. J Vert Paleontol. 28:702–711. doi: 10.1671/0272-4634(2008)28[702:AJSOTP]2.0.CO;2.
- Butler RJ, Smith RMH, Norman DB. 2007. A primitive ornithischian dinosaur from the Late Triassic of South Africa, and the early evolution and diversification of Ornithischia. Proc R Soc B Biol. 274:2041–2046. doi: 10.1098/rspb.2007.0367.
- Butler RJ, Upchurch P, Norman DB. 2008. The phylogeny of the ornithischian dinosaurs. J Syst Palaeontol. 6:1–40. doi: 10.1017/S1477201907002271.
- Cabaleri N, Volkheimer W, Nieto DS, Armella C, Cagnoni M, Hauser N, Matteini M, Pimentel MM. 2010. U-Pb ages in zircons from Las Chacritas and Puesto Almada members of the Jurassic Cañadón Asfalto Formation, Chubut province, Argentina. Paper presented at: VIISSAGI: 7th South American Symposium on Isotope Geology; Brasilia. Brasil.
- Casamiquela RM. 1967. Un nuevo dinosaurio ornitisquio Triásico (*Pisanosaurus mertii*: Ornithopoda) de la Formación Ischigualasto, Argentina. Ameghiniana. 4:47–64.
- Charig AJ, Crompton AW. 1974. The alleged synonymy of *Lycorhinus* and *Heterodontosaurus*. Ann S Afr Mus. 64:167–189.
- Chinnery BJ, Lipka TR, Kirkland JI, Parrish JM, Brett-Surman MK. 1998. Lower and Middle Cretaceous terrestrial ecosystems. In: Lucas SG, Kirkland JI, Estep JW, editors. Neoceratopsian teeth from the Lower to Middle Cretaceous of North America. Albuquerque, NM: New Mexico Museum of Natural History and Science. p. 297–302.
- Costa RL, Greaves WS. 1981. Experimentally produced tooth wear facets and the direction of jaw motion. J Paleontol. 55:635–638.
- Crompton AW, Charig AJ. 1962. A new ornithischian from the Upper Triassic of South Africa. Nature. 196:1074–1077.
- Cuneo R, Bowring S. 2010. Dataciones Geocronologías preliminares en la Cuenca Cañadón Asfalto, Jurasico de Chubut, Argentina. Implicancias Geológicas y paleontológicas. Paper presented at: XCAPB: X Congreso argentino de Paleontología y Bioestratigrafía; La Plata, Buenos Aires, Argentina
- Cúneo R, Ramezani J, Scasso R, Pol D, Escapa I, Zavattieri AM, Bowring SA. 2013. High-precision U–Pb geochronology and a new chronostratigraphy for the Cañadón Asfalto Basin, Chubut, central Patagonia: Implications for terrestrial faunal and floral evolution in Jurassic. Gondwana Research. http://dx.doi.org/10.1016/j.gr.2013.01.010
- Currie PJ, Rigby JKJr, Sloan RE. 1990. Dinosaur systematics: perspectives and approaches. In: Carpenter K, Currie PJ, editors. Chapter 8: Theropod teeth from the Judith River Formation of southern Alberta, Canada. Cambridge: Cambridge University Press. p. 107–125.
- Di Rienzo JA, Casanoves F, Balzarini MG, Gonzalez L, Tablada M, Robledo CW. 2011. InfoStat versión. Argentina. Available from http://www.infostat.com.ar Grupo InfoStat, FCA, Universidad Nacional de Córdoba.
- Dong Z. 1997. Sino-Japanese silk road dinosaur expedition. In: Dong Z-M, editor. A small ornithopod from Mazongshan area, Gansu Province, China. Beijing: China Ocean Press. p. 24–26.
- Galton PM. 1973. The cheeks of ornithischian dinosaurs. Lethaia. 6:67–89.
- Galton PM. 1974. The ornithischian dinosaur *Hypsilophodon* from the Wealden of the Isle of Wight. Bull Br Mus (Nat Hist) Geol. 25:1–152.
- Galton PM. 2007. Teeth of ornithischian dinosaurs (mostly Ornithopoda) from the Morrison Formation (Upper Jurassic) of the western United States. In: Carpenter K, editor. Horns and beaks. Ceratopsian and ornithopod dinosaurs. Bloomington, IN and Indianapolis, IN: Indiana University Press. p. 17–47.
- Galton PM, Jensen JA. 1979. Remains of ornithopod dinosaurs from the Lower Cretaceous of North America. Brigham Young Univ Geol Stud. 25:1–10.
- Goodall C. 1991. Procrustes methods in the statistical analysis of shape. J R Stat Soc B. 53:285–339.
- Gow CE. 1990. A tooth-bearing maxilla referable to *Lycorhinus* angustidens Haughton, 1924 (Dinosauria, Ornithischia). Ann S Afr Mus. 99:367–380.
- Haughton SH. 1924. The fauna and stratigraphy of the Stormberg Series. Ann S Afr Mus. 12:323–497.

- Holliday CM, Witmer LM. 2008. Cranial kinesis in dinosaurs: intracranial joints, protractor muscles, and their significance of cranial evolution and function of diapsids. J Vert Paleontol. 24:1073–1088.
- Hopson JA. 1975. On the generic separation of the ornithischian dinosaurs *Lycorhinus* and *Heterodontosaurus* from the Stormberg Series (Upper Triassic of South Africa). S Afr J Sci. 71:302–305.
- Hopson JA. 1980. Tooth function and replacement in early Mesozoic ornithischian dinosaurs: implications for aestivation. Lethaia. 13:93–105. doi: 10.1111/j.1502-3931.1980.tb01035.x.
- Irmis RB, Parker WG, Nesbitt SJ, Liu J. 2007. Ornithischian dinosaurs: the Triassic record. Hist Biol. 18:3–22.
- Norman DB. 2004. Basal Iguanodontia. In: Weishampel DB, Dodson P, Osmólska H, editors. The Dinosauria. 2nd ed. Berkeley, CA: University of California Press. p. 413–437.
- Norman DB, Barrett PM. 2002. Ornithischian dinosaurs from the Lower Cretaceous (Berriasian) of England. Spec Pap Palaeontol. 68:161–189.
- Norman DB, Crompton AW, Butler RJ, Porro LB, Charig AJ. 2011. The Lower Jurassic ornithischian dinosaur *Heterodontosaurus tucki* Crompton & Charig, 1962: cranial anatomy, functional morphology, taxonomy, and relationships. Zool J Linn Soc. 163:182–276. doi: 10.1111/j.1096-3642.2011.00697.x.
- Norman DB, Sues H-D, Witmer LM, Coria RA. 2004. Basal Ornithopoda. In: Weishampel DB, Dodson P, Osmólska H, editors. The Dinosauria. 2nd edition. Berkeley, CA: University of California Press. p. 393–412.
- Pérez SI, Bernal V, Gonzales P. 2006. Differences between sliding semilandmark methods in geometrics morphometrics, with an application to human craniofacial and dental variation. J Anat. 208:769–784.
- Pol D, Rauhut OWM, Becerra MG. 2011. A Middle Jurassic heterodontosaurid dinosaur from Patagonia and the evolution of heterodontosaurids. Naturwissenschaften. 98:369–379.
- Porro LB. 2007. Feeding and jaw mechanism in *Heterodontosaurus tucki* using finite element analysis. J Vert Paleontol. 27:131A.
- Rauhut OWM. 2001. Herbivorous dinosaurs from the Late Jurassic (Kimmeridgian) of Guimarota, Portugal. Proc Geologist Assoc. 112:275–283.
- Rauhut OWM, Martin T, Ortiz-Jaureguizar E, Puerta P. 2002. A Jurassic mammal from South America. Nature. 416:165–168.
- Richtsmeier JT, DeLeon VB, Lele S. 2002. The promise of geometric morphometrics. Yearbook Phys Anthropol. 46:63–91.
- Rohlf FJ. 1999. Shape statistics: procrustes method for the optimal superimposition of landmarks. Syst Zool. 39:40–59.
- Rohlf FJ. 2010a. tpsUtil, file utility program. version 1.46. Department of Ecology and Evolution, State University of New York at Stony Brook, Stony Brook.
- Rohlf FJ. 2010b. tpsDig, digitize landmarks and outlines, version 2.16. Department of Ecology and Evolution, State University of New York at Stony Brook, Stony Brook.
- Rohlf FJ. 2010c. tpsRelw, relative warps analysis, version 1.49. Department of Ecology and Evolution, State University of New York at Stony Brook, Stony Brook.
- Rougier GW, Garrido A, Gaetano A, Puerta P, Corbitt C, Novacek MJ. 2007. A new triconodont from South America. Am Mus Nov. 3580:1–17.
- Rougier GW, Martinelli AG, Forasiepi AM, Novacek MJ. 2007. New Jurassic mammals from Patagonia, Argentina: a reappraisal of australosphenidan morphology and interrelationships. Am Mus Nov. 3566:1–54.
- Ruiz-Omeñaca JI, Piñuela L, Garcia-Ramos JC. 2010. El primer diente de ornitopodo del Jurasico Superior de España (Asturias). Geogaceta. 48:83–86.

- Saegusa H, Tomida Y. 2011. Titanosauriform teeth from the Cretaceous of Japan. An Acad Bras Cienc. 83:247–265.
- Salani FM. 2007. Aporte a la edad de la Formación Cañadón Asfalto, Chubut, Argentina. 3º Simposio Argentino del Jurásico. Mendoza, Argentina. Resúmenes. 30:71.
- Santa Luca AP. 1980. The postcranial skeleton of *Heterodontosaurus* tucki (Reptilia, Ornithischia) from the Stormberg of South Africa. Ann S Afr Mus. 79:159–211.
- Sereno PC. 1986. Phylogeny of the bird-hipped dinosaurs (Order Ornithischia). Natl Geogr Res. 2:234–256.
- Sereno PC. 1991. Lesothosaurus, 'fabrosaurids', and the early evolution of the Ornithischia. J Vert Paleontol. 11:168–197.
- Sereno PC. 2012. Taxonomy, morphology, masticatory function and phylogeny of heterodontosaurid dinosaurs. ZooKeys. 226:1–225.
- Sheets HD. 2003. IMP integrated morphometrics package. Buffalo, NY: Department of Physics, Canisius College.
- Silva Nieto DG, Cabaleri N, Salani FM. 2003. Estratigrafía de la Formación Cañadón Asfalto (Jurásico Superior), Provincia del Chubut, Argentina. Ameghiniana, Resúmenes Primer Simposio Argentino del Jurásico. 40:p 46.
- Silva Nieto DG, Cabaleri N, Salani F, González Díaz E, Coluccia F. 2002. Hoja Geológica 4369-27 Cerro Cóndor, Provincia del Chubut. Instituto de Geología y Recursos Minerales, Servicio Geológico Minero Argentino, Buenos Aires. Boletín. 328:1–68.
- Smith JB, Vann DR, Dodson P. 2005. Dental morphology and variation in theropod dinosaurs: implications for the taxonomic identification of isolated teeth. Anat Rec A. 285A:699–736.
- Tasch P, Volkheimer W. 1970. Jurassic conchostracans from Patagonia. Univ Kansas Paleontol Contrib. 50:1–23.
- Thulborn RA. 1970. The systematic position of the Triassic ornithischian dinosaur Lycorhinus angustidens. Zool J Linn Soc. 49:235–245.
- Thulborn RA. 1973. Teeth of ornithischian dinosaurs from the Upper Jurassic of Portugal, with description of a hypsilophodontid (*Phyllodon henkeli* gen. et sp. nov.) from the Guimarota Lignite. Mém Serv Géol Port. 22:89–134.
- Thulborn RA. 1974. A new heterodontosaurid dinosaur (Reptilia: Ornithischia) from the Upper Triassic Red Beds of Lesotho. Zool J Linn Soc. 55:151–175.
- Volkheimer W, Gallego OF, Cabaleri NG, Armella C, Narváez PL, Silva Nieto DG, Páez MA. 2009. Stratigraphy, palynology, and conchostracans of a Lower Cretaceous sequence at the Cañadón Calcáreo locality, Extra-Andean central Patagonia: age and palaeoenvironmental significance. Cretaceous Res. 30:270–282.
- Weishampel DB. 1984. Evolution of jaw mechanisms in ornithopod dinosaurs. Adv Anat Embryol Cell. 87:1–109.
- Weishampel DB, Norman DB. 1989. Paleobiology of the dinosaurs. In: Farlow JO, editor. Vertebrate herbivory in the Mesozoic: jaws, plants, and evolutionary metrics. Boulder: Geological Society of America, Special Paper. p. 87–100.
- Weishampel DB, Witmer LM. 1990. Heterodontosauridae. In: Weishampel DB, Dodson P, Osmólska H, editors. The Dinosauria. Berkeley, CA: University of California Press. p. 486–497.
- Xu X, Forster CA, Clark JM, Mo J. 2006. A basal ceratopsian with transitional features from the Late Jurassic of northwestern China. Proc R Soc B. 273:2135–2140. doi: 10.1098/rspb.2006.3566.
- Zavattieri A, Escapa I, Scasso R, Olivera D, Cúneo NR. 2010. Nuevos aportes al conocimiento palinoestratigráfico y palinofacies de la sección inferior de la Formación Cañadón Asfalto en el Cañadón Lahuincó, depocentro de Cerro Cóndor, Chubut, Argentina. Paper presented at: IVSAJ: IV Simposio Argentino Jurasico y sus Límites; Bahia Blanca, Buenos Aires, Argentina
- Zheng XT, You HL, Xu X, Dong ZM. 2009. An Early Cretaceous heterodontosaurid dinosaur with integumentary structures. Nature. 458:333–336. doi: 10.1038/nature07856.