

Avian-like breathing mechanics in maniraptoran dinosaurs

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In 1868 Thomas Huxley first proposed that dinosaurs were the direct ancestors of birds and subsequent analyses have identified a suite of 'avian' characteristics in theropod dinosaurs. Ossified uncinat processes are found in most species of extant birds and also occur in extinct non-avian maniraptoran dinosaurs. Their presence in these dinosaurs represents another morphological character linking them to Aves, and further supports the presence of an avian-like air-sac respiratory system in theropod dinosaurs, prior to the evolution of flight. Here we report a phylogenetic analysis of the presence of uncinat processes in Aves and non-avian maniraptoran dinosaurs indicating that these were homologous structures. Furthermore, recent work on Canada geese has demonstrated that uncinat processes are integral to the mechanics of avian ventilation, facilitating both inspiration and expiration. In extant birds, uncinat processes function to increase the mechanical advantage for movements of the ribs and sternum during respiration. Our study presents a mechanism whereby uncinat processes, in conjunction with lateral and ventral movements of the sternum and gastral basket, affected avian-like breathing mechanics in extinct non-avian maniraptoran dinosaurs.

Keywords: Aves; theropod; uncinat processes; ventilation; gastralia

1. INTRODUCTION

Uncinat processes extend from the caudal edge of the vertebral ribs. In modern birds (with the exception of the Sphenisciformes and *Apteryx*), the processes are ossified to the thoracic ribs and typically have a bell-shaped base and taper distally (figure 1*a–c*). Avian uncinat processes were originally interpreted as having a primary function in stiffening the rib cage and stabilizing the shoulder musculature (Bellairs & Jenkins 1960). However, recent work has demonstrated that uncinat processes are crucial ventilatory structures in birds, involved in both inspiration and expiration (Codd *et al.* 2005). Morphological adaptations including differences in wing shape, leg morphology and the rib cage are commonplace in birds with adaptations to different forms of locomotion. Interestingly, recent work has indicated that uncinat length also correlates with locomotor mode in birds: being short in walking (figure 1*a*); intermediate length in flying (figure 1*b*); and long in diving species (figure 1*c*; Tickle *et al.* 2007). The appendicocostal muscle attaches the uncinat process to the following rib and is active during inspiration in the giant Canada goose (Codd *et al.* 2005). The uncinat processes also act as a strut for the insertion of the external oblique muscle, which pulls the sternum dorsally during expiration (Codd *et al.* 2005).

Recently, we have developed a geometric model to relate uncinat function to morphological differences

associated with locomotor specializations in birds. Our geometric model indicates that uncinat processes act as levers which improve the mechanical advantage for the forward rotation of the dorsal ribs and therefore lowering of the sternum during respiration in birds (Tickle *et al.* 2007; figures 2–4). Given their lever arm action and site for muscle attachment, the length of the processes is functionally important as it will alter the mechanical advantage and area available for muscle attachment.

The aim of this paper is to review the occurrence and morphology of uncinat processes in non-avian maniraptoran dinosaurs. In particular, we have examined the phylogenetic distribution of these processes to determine whether these are homologous structures. The wider distribution of uncinat processes and analogous structures will also be determined. Furthermore, the occurrence of these processes, in conjunction with movements of the sternum and a kinetic gastral basket, has reconstructed avian-like breathing mechanics in extinct non-avian maniraptoran dinosaurs.

2. UNCINATE PROCESSES IN NON-AVIAN MANIRAPTORAN DINOSAURS

Recent discoveries have improved our understanding of the phylogenetic relationships of theropods and indicate that origin of Aves (*Archaeopteryx* to Neornithes; Padian 1998) is embedded within maniraptoran theropod dinosaurs (Clark *et al.* 2002; figure 2). Within non-avian maniraptoran dinosaurs, ossified uncinat processes are found in oviraptorids and many dromaeosaurs.

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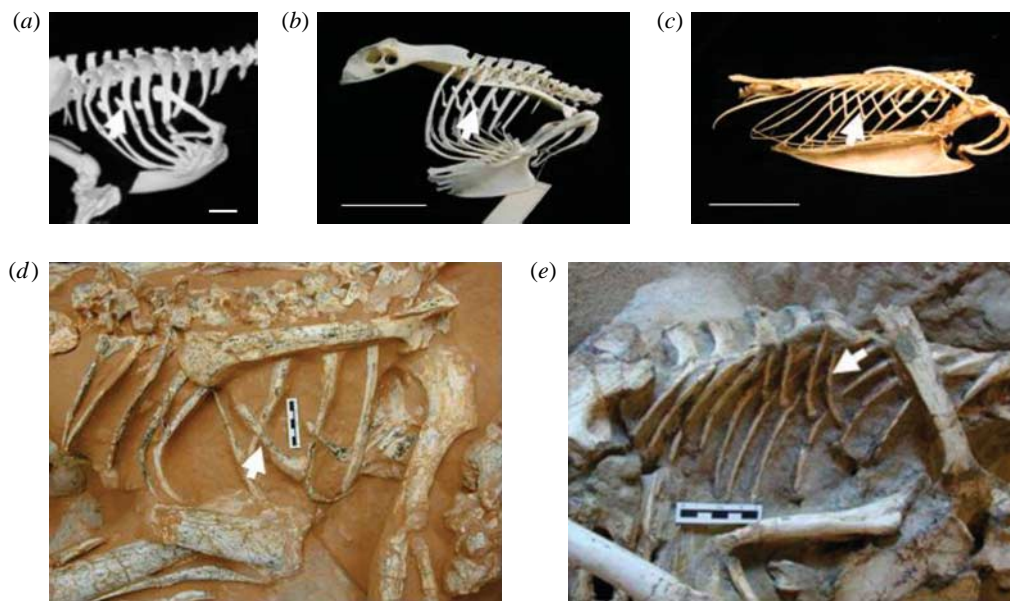


Figure 1. Uncinate processes (arrows) of (a) a running bird, the Cassowary (*Casuaris casuaris*), (b) a flying bird, the Eagle Owl (*Bubo bubo*), (c) a diving bird, the Razorbill (*Alca torda*); canonical variate analyses indicates that the uncinates are shorter in running, long in diving and intermediate in all other birds (Tickle *et al.* 2007), (d) AMNH 6517, (*Oviraptor philoceratops*), (e) MPD 100/25 (*Velociraptor mongoliensis*). Anterior is to the right in all figures. Scale bars, 5 cm. Photographs taken by J.R.C.

Oviraptor philoceratops (AMNH FR 6517) has four ossified uncinates present on the first four thoracic ribs (Clark *et al.* 1999; figure 1d). The processes extend from the caudal edge at the midpoint of the thoracic ribs and overlap the preceding ribs. The uncinates are flat, broad and bell shaped at the base, then taper distally and have a characteristic ‘L-shaped’ morphology, most closely resembling that of flying avian species (figure 1b). The processes on the first and fourth ribs are reduced in length and thickness. However, the uncinates on the second and third ribs exhibit both uniform geometry and morphology. The processes are not ossified to the ribs.

Five long uncinates are present on the left hand side of *Velociraptor mongoliensis* (MPD 100/25; Kielan-Jaworowska & Barsbold 1972; Norell & Makovicky 1999, 2004), extending from the caudal edge at the midpoint of the third to the seventh thoracic ribs to the edge of the second following rib (figure 1e). As described for *Oviraptor*, the processes are not ossified to the rib (figure 1e). In this taxon, the uncinates most closely resemble those of the diving avian species (figure 1c). The processes are flattened and uniform in morphology; they are thin and taper towards the tip. The uncinates on the seventh rib is reduced in size and length and has been displaced from the thoracic rib. Rib scars indicate that the uncinates on the first and second thoracic ribs are missing.

The description of *Deinonychus antirrhopus* (Ostrom 1969) made no mention of uncinates; however, ‘probable ventral ribs’ were figured (see Ostrom 1969, fig. 52). A review of *Deinonychus* (YPM5246, YPM5247 and YPM5250) material has identified these ‘probable’ elements as uncinates (P. L. Manning 2006, personal observation). The ‘articular scars’ on *Deinonychus* ribs (Ostrom 1969) were almost certainly cartilaginous attachment sites between the thoracic rib and proximally expanded articular surface of each uncinates process.

Deinonychus uncinates are typically bell shaped proximally, taper distally and were not ossified to the ribs. The uncinates may have been attached via cartilage as is suggested by corresponding roughened areas. The length of the uncinates indicates that they would have caudal–dorsally spanned several (2–4) ribs. It follows that gastral elements described in other non-avian maniraptoran theropods may also represent misidentified, unfused uncinates processes.

The smallest known non-avian theropods, *Microaptor zhaoianus* (V12330; Xu *et al.* 2000; Hwang *et al.* 2002) and *Microaptor gui* (IVPP V13352; Xu *et al.* 2003), are often well preserved in life positions and have uncinates processes (figure 3). The uncinates processes of *Microaptor* expand from the caudal edge of the ribs to a bell-shaped articulation with a distal tapering shaft and are homologous to, but much smaller than, those identified in this study for *Deinonychus*. Each process spans three ribs at an angle of 55° and was not fused to the ribs.

The occurrence of uncinates processes is patchy among the non-avian maniraptoran dinosaurs (figure 2). In the species so far described (*Velociraptor*, *Microaptor*, *Deinonychus*, *Khaan* and *Citipati*), the morphology of the uncinates processes indicates that they were attached via cartilage as shown by characteristic rough regions on both the ribs and the posterior articular surface of the processes. Cartilage does not readily fossilize as it has a lower structural density than bone (Lyman 1994) meaning the processes may have become detached and lost during disarticulation, weathering or scavenging of the skeleton. Given the diminutive size of these bones, it is also possible that they have been routinely overlooked during collection or simply not recognized as uncinates processes (e.g. Ostrom 1969). The cartilaginous attachment of the process to rib would also make disarticulation and mixing with gastral elements potentially problematic for identification. Uncinates processes have also been described in some basal avian species, *Confuciusornis* (Zhonghe & Lianhai 2002) and may also be present in

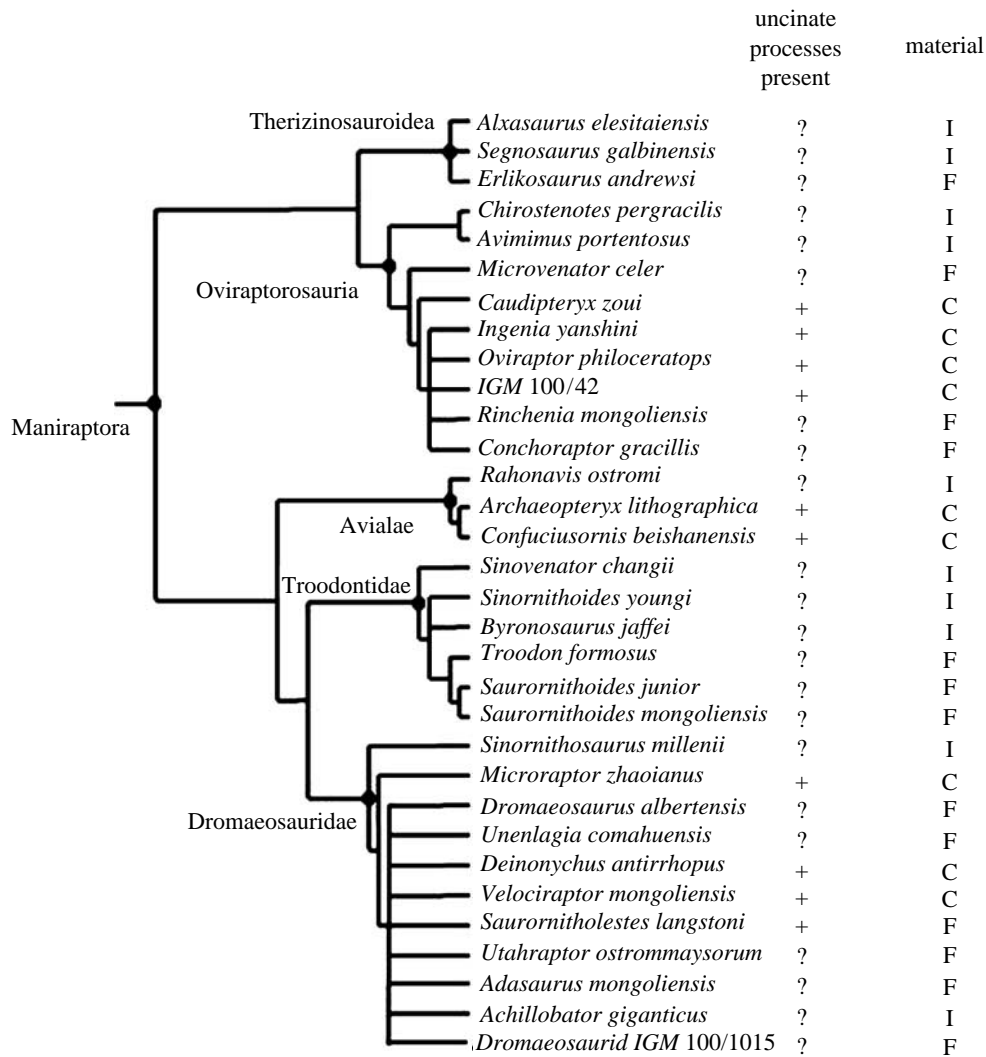


Figure 2. Summary cladogram of maniraptoran dinosaurs showing the occurrence of ossified uncinates. *Uncinate processes*: plus, ossified uncinates present; minus, ossified uncinates absent; question mark, presence of ossified uncinates unknown. *Material*: C, complete postcranial skeleton; I, incomplete postcranial skeleton; F, fragmentary postcranial skeleton. (Adapted from Clark *et al.* (2002).)

Archaeopteryx (Mayr *et al.* 2005; see figure 2b; P. L. Manning 2007, personal observation). Using parsimony, the presence of uncinates in early avian species, their presence in several non-avian maniraptoran groups and their phylogenetic distribution indicate that they are homologous structures.

3. WIDER DISTRIBUTION OF UNCINATE PROCESSES AND ANALOGOUS STRUCTURES

Cartilaginous or ossified uncinates are reported in crocodiles (Hofstetter & Gasc 1969) and the tuatara (Romer 1956); however, their possible role as ventilatory structures has yet to be determined. Well-developed gastralia are also present in both crocodilians and tuatara and have been suggested to play some role during ventilation (Lambe 1917; Perry 1983; Carrier & Farmer 2000; Claessens 2004). Indeed, in tuatara (*Sphenodon punctatus*), the external oblique muscle originates at the margin of the gastral basket and inserts onto the base of the uncinates similar to that described in Aves (Maurer 1896; figure 4).

Rib flanges which are occasionally referred to as uncinates are also well described in early tetrapods (Clack



Figure 3. Uncinate processes (arrows) of IVPP V13352 (*Microraptor gui*). Anterior is to the left. Scale bar, 13 cm. (Adapted with permission from Xu *et al.* (2003).)

2002). For example, *Eryops* and *Dissorophus* possessed small triangular costal expansions on the thoracic ribs (Gregory 1951; Clack 2002). The general morphology of these structures and their location on the ribs, which get progressively more ventral as you move rostrally, are quite different from uncinates found in Aves and the non-avian Maniraptora. The ventilatory mechanics of early tetrapods are unclear; however, air gulping, buccal pumping

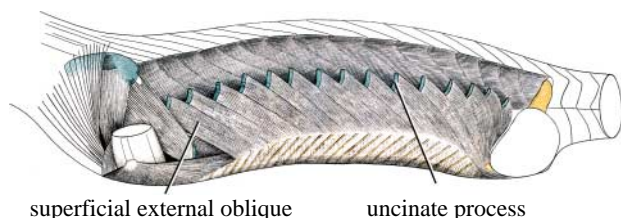


Figure 4. Hypaxial musculature of tuatara (*Sphenodon punctatus*) demonstrating insertion of external oblique muscle from the gastralia to the uncinata processes. Anterior is to the left. (Adapted with permission from Maurer (1896).)

and an early version of rib movement have all been suggested. Costal breathing is thought to have originated in early amniotes (Clack 2002; Perry & Sander 2004). The expanded ribs of early tetrapods may have been associated with the development of the transverse abdominal muscle which assists exhalation during aspiration breathing in tetrapods (Brainerd *et al.* 1993) or alternatively these processes are thought to function in the stiffening of the body during axial locomotion rather than serving any putative respiratory function (Clack 2002; J. Gauthier 2007, personal communication).

Some ornithischian dinosaurs (e.g. *Thescelosaurus*) possessed deep, plate-like intercostal structures attaching successive thoracic ribs (Fisher *et al.* 2000; see figure 1). Uncinates are morphologically distinct structures either fused to the thoracic ribs (Aves) or connected via cartilage (Non-Avian Maniraptora), while the multiple (often more than one per rib) structures observed in ornithischian dinosaurs consist of proximally and distally attached cartilaginous sheets connecting rib to rib. The structures observed in ornithischian dinosaurs are not likely to be homologous to avian uncinates. Furthermore, skeletal pneumatization is absent in ornithischian dinosaurs suggesting that they possessed a less 'bird-like' respiratory system than the theropods (Perry & Sander 2004). However, the cartilaginous plates of ornithischian dinosaurs might have functioned as passive elastic structures assisting expiration, with the recoil energy from inhalation reducing the volume of the lungs by pulling the rib cage back to a neutral position.

4. RECONSTRUCTING THE BREATHING MECHANICS OF NON-AVIAN MANIRAPTORAN DINOSAURS

Indirect evidence must be used to reconstruct the mechanics of breathing in extinct dinosaurs as there are no fossilized soft tissues or lungs. Both crocodiles and birds have a greater diffusing capacity in more proximal than distal regions of their lungs. Therefore, dinosaurs, being phylogenetically bracketed by these two groups (Witmer 1995), have been suggested to also possess a highly heterogeneous lung structure (Perry & Sander 2004). Furthermore, the pattern of regional postcranial pneumatization in theropods is similar to modern birds and has been used as evidence of the presence of a cervical and abdominal air-sac system in non-avian theropods (Britt 1997; Burnham *et al.* 2000; Perry 2001; O'Connor & Claessens 2005). Taken together, these suggest that a flow through respiratory system is a general theropod characteristic (O'Connor & Claessens 2005).

The implications of a putative air-sac system in theropods are that these were highly active animals.

The morphology of the uncinata processes in extant bipedal flightless birds is very different from that of extinct bipedal non-avian maniraptoran dinosaurs (figure 1). The uncinata processes in non-avian maniraptoran dinosaurs are not reduced as in running birds but are of intermediate to long lengths and resemble those of the flying or diving birds (figure 1). Running avian species lack the large breast musculature associated with flight, and therefore have no need for the lever-arm action provided by the uncinata processes to move a large muscle mass. The relatively long processes described for non-avian maniraptoran dinosaurs therefore suggest that an improved mechanical advantage, as demonstrated in extant avian species (Tickle *et al.* 2007), is functionally important. Birds ventilate their air sacs by moving the sternum in a dorsal-ventral plane to facilitate the bellows-like movement of air into and out of the air sacs. Non-avian maniraptoran dinosaurs lack a keeled sternum, but have well-developed gastralia in the ventral belly wall (Claessens 2004), and ossified sterna, attached to the distal ribs by large ossified sternal ribs. Gastralia have been postulated to play some role in respiration by preventing the inward movement of the belly wall (Perry 1983; Claessens 2004). However, more recently the gastral basket has been hypothesized to play a dynamic role in archosaur ventilation (Carrier & Farmer 2000).

The gastralia of non-avian maniraptoran theropods are highly derived, cross the midline and articulate with two gastralia from the opposite side of the body. This arrangement constitutes a lever arm which through association of hypaxial musculature could have narrowed or widened the cuirassal basket (Carrier & Farmer 2000). If the medial ends of the gastralia were drawn caudally by the ischiotruncus muscle, lateral and ventral movement of the belly wall would occur. The ischium functioned as the origin for the ischiotruncus muscle and the expanded foot of the pubis acted to direct the force of the muscle (Carrier & Farmer 2000). In modern birds, the external oblique muscle inserts onto the base of the uncinata processes from the sternum and the process acts as a strut to pull the sternal mass dorsally during expiration (Codd *et al.* 2005). A similar avian-like arrangement of the musculature in non-avian maniraptoran dinosaurs, whereby the external oblique pulled in a primarily dorsal direction, would have provided an attachment site for the insertion of an external oblique muscle originating at the edges of the gastral basket and sternum. This muscle, in conjunction with sternal movements and the narrowing and widening of the gastralia affected by the rectus and ischiotruncus, would have provided a mechanism for facilitating the bellows-like movement of air through a putative air-sac system.

Our study indicates that the presence of uncinata processes, coupled with specialized gastralia, sterna and pelvic girdles, provides a mechanism for facilitating avian-like breathing mechanics in non-avian maniraptoran dinosaurs.

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