First trace and body fossil evidence of a burrowing, denning dinosaur

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A fossil discovery in the mid-Cretaceous Blackleaf Formation of southwest Montana, USA, has yielded the first trace and body fossil evidence of burrowing behaviour in a dinosaur. Skeletal remains of an adult and two juveniles of Oryctodromeus cubicularis gen. et sp. nov., a new species of hypsilophodont-grade dinosaur, were found in the expanded distal chamber of a sediment-filled burrow. Correspondence between burrow and adult dimensions supports Oryctodromeus as the burrow maker. Additionally, Oryctodromeus exhibits features of the snout, shoulder girdle and pelvis consistent with digging habits while retaining cursorial hindlimb proportions. Association of adult and young within a terminal chamber provides definitive evidence of extensive parental care in the Dinosauria. As with modern vertebrate cursors that dig, burrowing in Oryctodromeus may have been an important adaptation for the rearing of young. Burrowing also represents a mechanism by which small dinosaurs may have exploited the extreme environments of polar latitudes, deserts and high mountain areas. The ability among dinosaurs to find or make shelter may contradict some scenarios of the Cretaceous–Paleogene impact event. Burrowing habits expand the known range of nonavian dinosaur behaviours and suggest that the cursorial ancestry of dinosaurs did not fully preclude the evolution of different functional regimes, such as fossoriality.

Keywords: Dinosauria; Ornithopoda; Ichnology; burrow; parental care; Blackleaf Formation

1. INTRODUCTION
Despite the longevity and predominance of dinosaur lineages throughout the Mesozoic, the radiation of nonavian dinosaurs remained morphologically restricted in comparison with the diversity found among Cenozoic mammals (Sereno 1999). Originating from small bipedal cursors, nonavian dinosaurs maintained primarily terrestrial habits throughout their history (Sereno 1999). Even the climbing capabilities hypothesized for the dinosaurian ancestors of birds remain quite controversial (Zhang et al. 2002; Padian 2004). Although skeletal features suggest a digging and tearing habit in the dinosaurs Heterodontosaurus and Mononykus (Norman et al. 2004; Senter 2005) and earthen nest structures represent the products of digging behaviour in the theropod Troodon (Varricchio et al. 1997) and titanosaur sauropods (Chiappe et al. 2004), the nonavian dinosaur radiation has been regarded as devoid of true burrowing forms (Sereno 1999; Robertson et al. 2004). A novel discovery (Museum of the Rockies, MOR, 1636) from the mid-Cretaceous of Montana consists of the skeletal remains of an adult and two juvenile dinosaurs preserved within an infilled burrow (figures 1 and 2). Here, we describe this dinosaur, Oryctodromeus cubicularis, as a new genus and species of hypsilophodont (here considered a grade of basal ornithopods), and present the first trace and body fossil evidence for fossorial behaviour in a nonavian dinosaur. Recognition of such behaviour has implications for dinosaur ecology, reproduction, distribution, evolution and extinction.

2. SYSTEMATIC PALAEONTOLOGY
Ornithischia Seeley, 1887
Ornithopoda Marsh 1881
Euornithopoda Sereno 1986
Oryctodromeus cubicularis gen. et sp. nov.

(a) Etymology
The generic name comes from the Greek words orycto and dromeus, meaning the ‘digging runner’. The species name, cubicularis, translates to ‘of the lair’, referring to the inferred denning habit of the taxon.

(b) Holotype
MOR 1636a consists of the following elements: fused premaxillae; posterior/occipital region of the brain-case; three cervical, six dorsal, sacrum and 23 caudal vertebrae; three dorsal ribs; scapulocoracoid; scapula; coracoid; humerus; ulna; radius; tibiae; distal fibula; and metatarsal IV (figures 3 and 4).

(c) Paratype
MOR 1636b includes cranial and postcranial materials found in association with the holotype and represents two juveniles with most linear dimensions measuring 55% to 65% those of the adult (figure 4).

(d) Locality and horizon
The new specimens come from the upper portion of the mid-Cretaceous Blackleaf Formation near the Lima Peaks, Beaverhead Co., Montana, USA (figure 1). Dyman & Nichols (1988) designate this sequence as the ‘volcaniclastic lithofacies’, a 270 m thick lithologically heterogeneous unit.
Radiometric dates indicate a Cenomanian age for this and the Aspen Formation of western Wyoming and eastern northern outcrops near Great Falls, Montana (Cobban et al. 1988). The upper, volcaniclastic lithofacies may correlate with the Vaughn Member described from more

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...The preserved portion of the structure measures more than 210 cm long and is sinuous and semi-helical. Throughout most of its length, the structure maintains a fairly constant oval cross-section measuring 30–38 cm high × 30–32 cm wide before ending abruptly as a partially preserved expansion. From its upper eroded limit, the structure slopes downwards at 6° for 60 cm, turns sharply right and continues down at 18° for another 70 cm (figure 2a,d). It then turns sharply left and proceeds a short distance (40 cm) before expanding (figure 2b,d). Over its course, the structure drops nearly 50 cm from the horizontal. Erosion had exposed the terminal expansion and its bone content by the time of discovery. Excavation of fossils further destroyed aspects of the structure, particularly the upper portions of the expansion, prior to its recognition and photo documentation (figure 2). Hence, the expansion’s preserved dimensions, 40–50 cm long × 45 cm wide, represents only a portion of its original size. The bone assemblage consisted of a densely packed jumble of adult and juvenile bones concentrated in the basal 7 cm of the lower sandstone just prior to and primarily within the terminal expansion (figure 2d). This concentration is greatest a few centimetres above the base of the structure and the coarsest non-bone clasts. A few small juvenile bones, e.g. caudal centra and cranial elements, occurred at a higher level in this unit. Except for one unidentifiable fragment and those truncated by recent erosion, bones were complete prior to fossilization and assignable to Oryctodromeus.

Small cylindrical features composed of the same sandstone as the large described structure extend horizontally from the main structure at each of the two right-angle turns (figure 2a,c,d). A cluster of five or six

**Figure 1.** Map of southwestern Montana, USA, showing the distribution of mid-Cretaceous rocks of the Blackleaf and Frontier Formations. Star denotes the location of MOR 1636. Modified from Dyman & Nichols (1988).

Consisting of abundant mudstone, siltstone, lithic-rich sandstone and subordinate conglomerate (Zartman et al. 1995). Low in this lithofacies, mudstone is olive green to grey, variably calcareous and structureless. Mudstone of the middle and upper portions may be porcelanitic of various colours (red, grey or green), bentonitic with ‘popcorn-texture’ weathering, or strongly calcareous, non-porcelanitic and grading laterally into micritic limestone (Dyman & Nichols 1988). Siltstone may be tan to grey, rippled, bioturbated and variably calcareous. It occurs as thin discontinuous beds within mudstone or above sandstone in lenticular bodies (Dyman & Nichols 1988). Sandstone is typically fine-grained, variably calcareous and thin to medium bedded. In the Lima Peaks area, sandstone contains a high proportion (as much as 50%) of volcanic and plagioclase grains (Dyman et al. 1988).

These sedimentary rocks represent deposition on a broad flood plain with relatively small fluvial channels at the western, proximal edge of the foreland basin (Dyman & Nichols 1988). The high volcanioclastic content reflects the proximity of nearby volcanism associated with the development of the Idaho batholith (Dyman & Nichols 1988; Dyman et al. 1988). Bivalves, unidentified reptilian bone fragments, leaves and wood, including the genus Tempekiya, represent the only macrofossils previously described for the Blackleaf Formation (Cobban et al. 1976; Dyman & Nichols 1988). The upper, volcanioclastic lithofacies may correlate with the Vaughn Member described from more northern outcrops near Great Falls, Montana (Cobban et al. 1976), the Mowry Shale of Montana and Wyoming and the Aspen Formation of western Wyoming and eastern Idaho (Dyman & Nichols 1988). Palynomorphs and radiometric dates indicate a Cenomanian age for this portion of the formation (Dyman & Nichols 1988; Zartman et al. 1995).

The holotype and paratype material occurs within an unusual sedimentary structure consisting of an isolated, sinuous sandstone body sitting within and crosscutting a succession of three mudrock units (figure 2). The lowermost mudstone is greater than 35 cm in thickness, light greenish grey and weakly calcareous. This unit exhibits extensive dark reddish grey mottling over its uppermost portion. Additionally, carbonate nodules as much as 4.5 cm in diameter form a discontinuous layer of 20–25 cm below the sharp upper contact of the unit. A dark greenish-grey claystone succeeds vertically. This 25 cm thick unit is calcareous with a platy structure and grades upwards into a greenish grey blocky mudstone with some organics. This third unit encloses the upper portion of the sedimentary structure (figure 2a,d) and bears a sharp, irregular contact with an overlying burrowed dark red claystone. A thin (approx. 30 cm thick) sandstone and additional mudrocks represent the remaining strata visible at this outcrop.

The lithology of the structure contrasts sharply with the surrounding host mudrock and facilitates recognition of the structure in the field (figure 2). The lower, bone-bearing portion consists of 25–30 cm of medium to fine, calcareous greenish-grey sandstone, moderately sorted with abundant plagioclase and other volcanogenic grains (figure 2d). Small (3–6 mm), rounded mud clasts are common in the coarser basal portion. This structureless unit fines upwards with an increasing clay content. A thin, 1–2 cm thick, grey claystone separates this lower sandstone from an upper fine sandstone. The bedding parallels that of the host mudrocks. Reduced clay content, grey colour and larger (approx. 10 mm), platy mud clasts distinguish this unit from the otherwise similar, lower sandstone. A thin (less than 1 cm) rim of tan claystone with abundant calcite veins separates the lower portion of the structure from the host mudrocks.

**Figure 1.** Map of southwestern Montana, USA, showing the distribution of mid-Cretaceous rocks of the Blackleaf and Frontier Formations. Star denotes the location of MOR 1636. Modified from Dyman & Nichols (1988).
closely spaced and in some instances doubled cylinders,
each 1×2 cm in cross-section and 15 cm long, originate
from the upper aspect of the main structure near the outside
corner of the uppermost turn. A single, larger feature, 7–
9 cm in diameter and at least 10 cm long, likewise extends
from the main structure at the outside of the lowermost turn.
Unlike the smaller features, this emerges from the lower
aspect of the main structure. No external ornamentation
or internal structures such as menisci are evident in
these features.

(e) Diagnosis
Oryctodromeus cubicularis exhibits the following autapo-
morphies: long paraoccipital processes indicating a skull
with relatively broad proportions; basioccipital with a
steeply sided ventral ‘box’ just rostral to the occipital
condyle (figure 3b); sacrum incorporating seven vertebrae,
including two posterior dorsals and their ribs; large scapula
with a sharply angled and narrow acromion process bearing
a thin-edged laterally projecting scapular spine and a
distinct posterior bend to the scapular blade; ilium with
very short pre-acetabular and long post-acetabular portions; brevis shelf slopes mediolaterally and is visible laterally throughout its length; and long prepubic process with a transversely broad proximal portion possessing an elongate ventral fossa (figure 4).

(f) Description
Medium-sized, hypsilophodont-grade, basal ornithopod; adult humerus length 157 mm and adult tibia 254 mm. Extensive adult fusion (premaxillae, posterior braincase, all neurocentrum sutures, sacrum and scapulocoracoid). Anterior margin and posterior nasal processes of premaxillae angled more posterodorsally (at 45°) than in Zephyrosaurus (Sues 1980). Juvenile lacking jugal boss; adult condition unknown. Cervical centra weakly opisthocoelous. Transverse processes of anterior dorsal vertebrae steeply angled (figure 4a). Paraphyses and diapophysis not occurring together on horizontal plane until possibly dorsal six or seven, as in more advanced ornithopods such as Tenontosaurus and Iguanodon (Scheetz 1999). Sacral ribs forming continuous iliac contact surface from the pubic peduncle through length of brevis shelf. Third sacral vertebra with small abutment for bracing of pubis (figure 4b). Caudal centra developing more hexagonal cross-sections distally with transverse processes diminishing rapidly and absent by 20th caudal. Post-zygapophyses by about caudal-14 consisting simply of lateral depressions at the neural spine base. In more distal caudal vertebrae, elongate pre-zygapophyses simply overlap the shortening neural spine. Few ossified tendons and absence of a tendon lattice in tail. Robust scapula longer than humerus (176 versus 157 mm) and with long, narrow acromion running nearly perpendicular away from scapular blade (figure 4c). As in other hypsilophodonts (Norman et al. 2004), ilium with bulbous antitrochanter (supratrochanteric process) on ischial peduncle. Bone completely encircling obturator foramen. Bowed femur with pendant fourth trochanter and small cleft separating the cranial and greater trochanters. Juvenile hindlimb ratios of 1.19 for tibia/femur and 0.58 for metatarsal III/femur, adult ratios probably similar but slightly lower.

3. DISCUSSION
Phylogenetically, O. cubicularis groups among basal euornithopods with two other Montana hypsilophodonts: Zephyrosaurus schaffi from the Lower Cretaceous Cloverly Formation (Sues 1980) and Orodromeus makelai from the Upper Cretaceous Two Medicine Formation (Horner & Weishampel 1988; Scheetz 1999; figure 5). These three taxa share a prominent scapular spine and direct pubosacral articulation. Three features unite Orodromeus and Zephyrosaurus more closely: inflated postorbital; fossa or groove at the base of pterygoid wing of quadrate; and jugal boss (Scheetz 1999). The adult status of these features in Oryctodromeus remains unknown. Additional features shared by Oryctodromeus and at least one of the other Montana forms include: fused premaxillae (Zephyrosaurus); cranial nerve V passing fully through the prootic; and potentially seven sacral vertebrae (Orodromeus). The last two features are not preserved in Zephyrosaurus (Sues 1980).

The distinct, stratified fill, crosscutting relationship with surrounding horizons and regular form and dimensions indicate that the Oryctodromeus-bearing sedimentary structure represents an infilled burrow (Voorhies 1975; Smith 1987; Groenewald et al. 2001; Haslotes et al. 2004). The varied and bedded rock types that comprise the structure are inconsistent with soft-sediment deformation and instead indicate passive filling of an open structure (figure 2). At least two episodes of filling would be required to account for the distinct sandstone units (Groenewald et al. 2001). The thin, calcite-veined claystone surrounding the lower portion of the sandstone fill could represent initial deposition within the burrow, but more probably represents secondary clay movement and mineral precipitation along the lithological boundary between the sandstone fill and host mudrocks.

Burrows of both terrestrial and marine taxa often house commensal species. For example, a wide variety of both invertebrates and vertebrates reside in or temporarily occupy the burrows of the gopher tortoise (Gopherus polyphemus) and nine-banded armadillo (Dasypus novemcinctus) (Kent et al. 1997; Platt et al. 2004). These taxa may generate their own associated tunnels intersecting the host burrow (Kent et al. 1997; de Gibert et al. 2006; Martin 2006). For the Blackleaf specimen, the small, cylindrical structures that extend from near the two major turns (figure 2a,c) possess the same fill as the main burrow structure. This indicates that they were open until the time of burrow filling. These accessory structures are too small to be generated by Oryctodromeus and, consequently, may represent the products of commensal cohabitants of the overall burrow structure.

The mudrocks surrounding the burrow are typical of those broadly interpreted as representing flood-plain deposition within the Blackleaf Formation (Cobban et al. 1976; Dyman & Nichols 1988). The sharp upper contact, mottled upper zone and carbonate nodule layer clearly indicate that the lower host unit (figure 2d) represents a moderately developed palaeosol (Retallack 1988). They further suggest a calcisol (Mack et al. 1993), a moderately alkaline, well-drained soil formed under modest precipitation as in semi-arid to subhumid climates (Retallack 1990). In contrast, the overlying two units exhibit only minimal soil development. Although they share a gradational contact, a sharp upper contact at the top of the third

Figure 3. Adult cranial elements for the O. cubicularis holotype (MOR 1636a). (a) Fused premaxillae in right lateral view. Premaxillae bears five teeth posterior to a small beak and a transversely expanded and roughened anterior face, f. (b) Ventral view of posterior braincase showing elongate paroccipital processes, poc, and steeply sided ventral 'box', b, just anterior to the occipital condyle, oc. Scale bar, 1 cm.
unit and platy to blocky structures, they lack discrete soil horizons and root traces (figure 2a,d). The grey coloration, alkaline nature and presence of visible organics may link them to relatively immature palaeosols, such as the Halts and Maksini pedotypes described from the Chinle and Two Medicine Formations, respectively (Retallack 1997). These pedotypes are interpreted as representing young, streamside communities with short soil development histories (Retallack 1997).

Burrow architecture consists of a sloping, sinuous tunnel ending with a terminal chamber (figure 2d) and is similar to those of many extant burrowing vertebrates, e.g. various rodents, striped hyena (Hyaena hyaena), aardwolf (Proteles cristatus), gopher tortoise and puffin (Fratercula arctica) (Voorhies 1975; Rieger 1981; Koehler & Richardson 1990; Doonan & Stout 1994; Terres 1995; Hasiotis et al. 2004). The taphonomy, burrow dimensions and anatomical evidence identify Oryctodromeus as the burrow maker. Both adult and juvenile fractions include portions of the cranial, axial and appendicular skeleton. Transport of complete carcasses deep within a burrow by either hydraulic processes or scavengers seems unlikely. The hydraulically disparate mixture of large adult limb bones and delicate, juvenile cranial and axial elements argues against fluvial transport. Furthermore, the absence of bite marks and pre-fossilization breaks is inconsistent with carnivore-generated bone assemblages (Brain 1981).

Based on the preserved vertebrae (n = 39), the adult would measure 2.1 m long with a major portion (approx. 1.25 m) representing tail. Breadth of the trunk as estimated from placement of the scapulocoracoids and dorsal ribs would be 26–30 cm. Tunnel width (approx. 30 cm) and length between turns (approx. 70 cm) thus correspond to torso breadth and length of the adult. Comparable tight fits occur in modern burrowing animals and often serve as a deterrent against predators (Rieger 1981; Koehler & Richardson 1990; Miller et al. 2001). For example, the aardwolf stands 45–50 cm at the shoulder but has tunnels measuring only 20 × 30 cm (Koehler & Richardson 1990); the striped hyena, 60–94 cm at the shoulders, has tunnel diameters of only 40 cm (Rieger 1981). Furthermore, an allometric equation comparing burrow cross-sectional area and body mass for a variety of modern invertebrate and vertebrate burrowers (White 2005) predicts an animal weighing between 22 and 32 kg for the Blackleaf burrow (figure 2a). These figures correspond to weight estimates of medium-sized hypsilophodonts (Pezkis 1994).

Although its hindlimb proportions compare favourably to other hypsilophodonts and denote cursorial habits (Carrano 1999; Norman et al. 2004), Oryctodromeus exhibits features of the snout, shoulder girdle and pelvis consistent with digging. Several of these represent derived features of Oryctodromeus and its closest sister taxa (figure 5), the two other Montana hypsilophodonts. For example, some extant fossorial vertebrates use their snouts to loosen or move dirt (Hildebrand 1985). The fused premaxillae with expanded anterior face of Oryctodromeus could represent such an implement (figure 3a). Among ornithopods, only Zephyrosaurus exhibits similar fused premaxillae (Sues 1980).

Despite similar forelimb proportions and morphology, the scapulocoracoid of Oryctodromeus differs markedly in comparison with those of other hypsilophodont-grade ornithopods, such as Thescelosaurus and Hypsilophodon (Norman et al. 2004). The robust scapula bears a long, narrow acromion that angles sharply away from the scapular blade (figure 4c). A distinct ventrolaterally projecting spine extends the entire length of the acromion. Orodromeus and Zephyrosaurus also possess scapular spines (Scheetz 1999), but that of Oryctodromeus differs, in that the spine projects nearly perpendicular from the flat, lateral surface of the acromion. Further, the anterior margin of the scapular blade bears a small triangular
project. From this point distally, the blade expands markedly and bends sharply posteriorly. In the adult specimen, the left scapula and coracoid are fully fused. Among ornithopods, this condition occurs elsewhere only in the iguanodontian Ouranosaurus (Norman 2004).

The enlarged, fused scapulocoracoid with prominent acromion and scapular spine would increase attachment areas for muscles that used to stabilize and operate digging forelimbs (Hildebrand 1985). The armadillo Dasypus possesses a distal expansion and posterior bend in the scapular blade similar to those of Oryctodromeus. This modification increases the lever arm for the enlarged teres major muscle and facilitates forelimb retraction (Hildebrand 1985; Pough et al. 1999).

In order to effectively apply a force to the ground, a digging animal must counteract the equal opposing force generated. For example, those mammals that dig with their forelimbs brace themselves with their hindlimbs and prop-like tail (Hildebrand 1985). Oryctodromeus exhibits several skeletal features that among extant vertebrates are associated with such behaviour (Hildebrand 1985): an expanded sacrum; greater attachment of pelvis and sacrum; absence of a pelvic symphysis; and expanded proximal caudals (figure 4). The large sacrum has seven vertebrae, including two dorsal vertebrae and their ribs, a long iliac attachment and direct pubosacral contact. Oryctodromeus is unusual in possessing seven sacrals, a number more typical of much larger Iguanodontia and other more advanced ornithopods (Norman 2004). Irregular surfaces on unfused posteriormost dorsals of O. makelai suggest the possibility of a seven-vertebrae sacrum in this Late Cretaceous form. The pubosacral articulation and the scapular spine represent the two synapomorphies that define the clade of Zephyrosaurus, Oryctodromeus and Orodromeus (figure 5). In the tail, the longest transverse processes occur on the third or fourth caudal vertebrae as in Hyspsilophodon and more advanced ornithopods (Scheetz 1999). The tallest neural spines of the vertebral column also occur in this region. Together, these features suggest a robust basal portion of the tail.

Oryctodromeus lacks the extreme skeletal modifications found in subterranean vertebrates, such as the golden mole (Amblysomus sp.), wombat (Lasiorhinus sp.), echidna (Tachyglossus aculeatus) and aardvark (Orycteropus afer). Missing traits include an elongate humeral head, transverse expansion of the distal humerus and prominent olecranon (Hildebrand 1985). Given its hindlimb proportions and modest forelimb modifications, Oryctodromeus is similar but more specialized than various cursorial, facultative diggers, such as the aardwolf (Koehler & Richardson 1990), hyaenids (Riegler 1981), various canids, Patagonian cavy (Dolichotis patagonum; Campos et al. 2001) and the Old World rabbit (Oryctolagus cuniculus; Nowak & Paradiso 1983). The bipedal gait of the dinosaur may have accommodated greater forelimb specializations without compromising terrestrial locomotion.

Although associated burrows are currently unknown for Zephyrosaurus and Orodromeus, the skeletal features shared by all the three Montana hypsilophodonts suggest that the clade may have been specialized for digging. The three small dinosaurs each occur within proximal portions of the Cretaceous foreland basin and in association with variegated, often red and purple, caliche-bearing mudstones (Retallack 1997). Modern tetrapod burrowers prefer drier, upland soils (Voorhies 1975; Groenewald et al. 2001; Hasiotis et al. 2004), and these well-drained seasonally arid palaeosols (Retallack 1997) of the Cretaceous would represent favourable substrates. At least two Orodromeus specimens (MOR 1136 and 1141) consisted of densely packed, three-dimensional bone assemblages similar to that of Oryctodromeus, suggesting within-burrow preservation. Uniformity of sediments may have obscured burrow preservation (Voorhies 1975) in these instances.

Fossorial behaviour represents a marked departure from the strictly cursorial niche envisioned for basal ornithopods (Carrano 1999; Norman et al. 2004). Furthermore, despite their bipedal, cursorial ancestry, dinosaurs evolved a variety of functional modes that now include fossorial, arboreal (Zhang et al. 2002) and several unrelated quadrupedal forms (Sereno 1999).

Vertebrates burrow for a variety of reasons: to find food; escape predators; and avoid harsh environmental conditions (Voorhies 1975; Hildebrand 1985). Among modern digging cursors, burrows serve primarily as dens for the rearing of
young (Rieger 1981; Nowak & Paradiso 1983; Koehler & Richardson 1990; Campos et al. 2001). The occurrence of an adult and two juveniles within the expanded portion of a burrow indicates that Oryctodromeus both burrowed and exhibited extensive parental care. The juveniles possess limb element and centra lengths measuring 60% to 65% those of the adult, exhibit no fusion except in the distal most caudals and would have measured 1.3 m in total length. Although growth rates were probably fast (Padian et al. 2004), the large size of the juveniles suggests that the parent–young bond persisted for a minimum of several months. Parental care of the young has been proposed for several dinosaur taxa (Horner & Makela 1979; Bakker & Bir 2004; Meng et al. 2004), but even the best examples either lack a direct juxtaposition of adult and young (Horner & Makela 1979), consist only of shed teeth (Bakker & Bir 2004) or remain taphonomically ambiguous (Meng et al. 2004). Thus, MOR 1636 uniquely preserves both an adult–young association and direct evidence of denning in the form of a dwelling trace. The presence of parental care in a basal ornithopod supports the hypothesis that such behaviour was ancestral for Dinosauria and, possibly, Archosauria (Meng et al. 2004).

Burrowing behaviour allows vertebrates to escape harsh environmental conditions by taking advantage of equable burrow microclimates (Voorhies 1975; Hildebrand 1985; Miller et al. 2001; Platt et al. 2004). By denning, small dinosaurs could have potentially withstood severe conditions, such as aridity, drought and daily or seasonal temperature extremes. Such behaviour would have allowed dinosaurs to occupy high mountains, desert environments (Loope 2006a) and polar regions and may explain the presence of hypsilophodonts in high-latitude areas, such as the Early Cretaceous of Australia (Rich & Rich 1989). The most similar Mesozoic traces to the one described here include a collection of burrows from the Middle Jurassic Entrada Sandstone (Loope 2006b). Although these burrows lack the sinuosity of the Blackleaf specimen, they share comparable tunnel lengths, slopes and diameters. One even possesses a terminal chamber (Loope 2006b). These taxonomically unassignable traces represent the only other burrows suitably sized to be the product of nonavian dinosaurs. Regardless of the trace maker, these burrows probably represented temporary shelters from harsh desert and dune field conditions of the Middle Jurassic (Loope 2006b).

Survivability of terrestrial vertebrates at the end-Cretaceous event has been attributed to sheltering behaviour, with the extinction of nonavian dinosaurs resulting from their inability to find an appropriate cover (Robertson et al. 2004). Burrowing dinosaurs would challenge this argument, but these are yet to be found in the latest Cretaceous formations. Much of the research on the Cretaceous–Paleogene extinction event has focused on the Hell Creek Formation and similar equivalent units of North America (Hartman et al. 2002). If burrowing dinosaurs existed in the latest Cretaceous, they would more probably be found not in coastal settings like the Hell Creek Formation (Hartman et al. 2002) but rather in more upland environments with better-drained palaeosols.

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