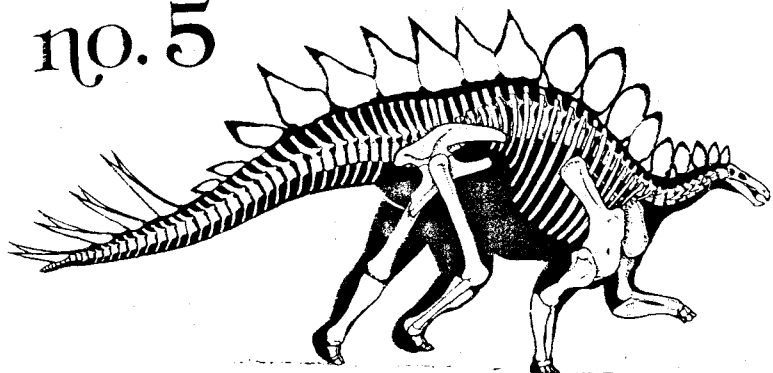


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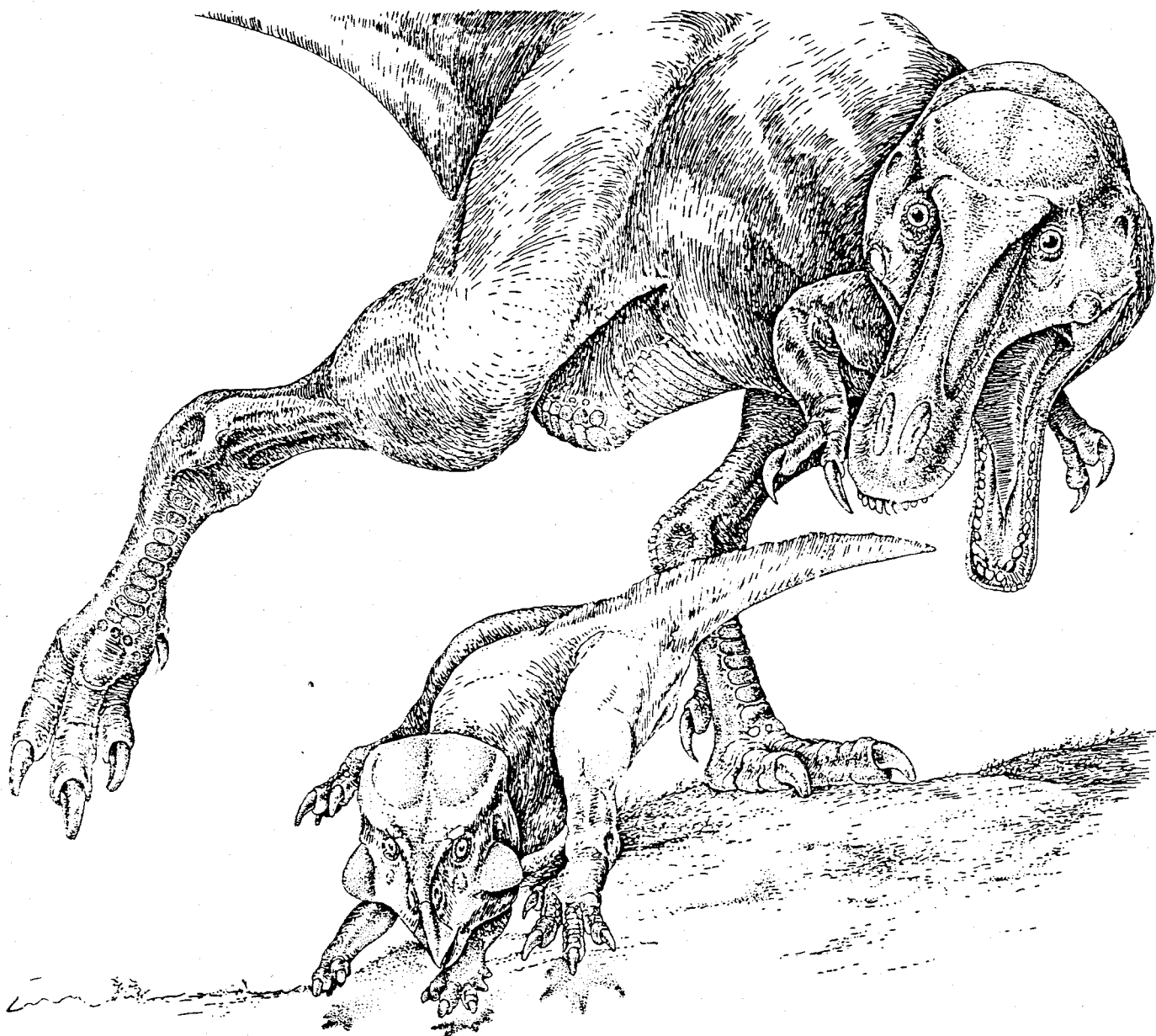


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NANOTYRANNUS, A NEW GENUS OF PYGMY TYRANNOSAUR, FROM THE LATEST CRETACEOUS OF MONTANA



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ABSTRACT

Nanotyrannus, type species *Gorgosaurus lancensis*, is the smallest known tyrannosaurid dinosaur and represents, with *Tyrannosaurus rex*, the last and most advanced members of the Family Tyrannosauridae. *Nanotyrannus* shows unusually strong adaptations for stereoscopic vision — the temporal region is expanded to four times the width of the muzzle, the eyes face forwards, and the head was bent down sharply on the neck. More primitive tyrannosaurs — gorgosaurs and daspletosaurs — lack adaptations for stereoscopy but some other more advanced genera — albertosaurs, tarbosaurus and *Tyrannosaurus* itself — show marked frontal expansion and muzzle constriction, although not to the same degree as in *Nanotyrannus*. The teeth in *Nanotyrannus* were far more primitive than those of all other tyrannosaurs except *Alioremus* in retaining a laterally-compressed form, and the braincase was advanced in its great width and flat ventral surface. Although *Nanotyrannus* shows marked structural convergence with *Tyrannosaurus rex*, the pygmy tyrannosaur probably evolved independently from a very primitive tyrannosaurid. *Nanotyrannus* and *Tyrannosaurus* are the last large land predators of the Age of Dinosaurs, and the convergent evolution of stereoscopy suggests that visual acuity and depth perception were necessary to deal with the contemporaneous herbivores, especially the heavily armed horned dinosaurs.

INTRODUCTION

The dominant group of large land-living predators during the Late Cretaceous was the Tyrannosauridae, and this family is far more diverse than is usually suspected (Bakker, Currie, and Williams, in press). In the last zone of the North American Cretaceous, the Lancian, the most common large predator was *Tyrannosaurus* itself, a genus with very advanced adaptations that included a narrow snout and wide temporal region ("forehead") so that both eyes were directed forward, and an occiput that bent the head sharply downward on the neck. Unlike the earlier tyrannosaurids such as *Gorgosaurus* and *Daspletosaurus*, *Tyrannosaurus* could focus both eyes on an object over a wide visual field and thus acquired the prerequisite for stereoscopic vision. Greater depth perception and visual acuity may well have been required because the most common large Lancian herbivore was *Triceratops*, a five-ton ceratopsian armed with very long, sharp horns mounted on a mobile and powerfully muscled head. Successful attack upon an adult *Triceratops* would require great precision on the part of the predator.

Here we report a new genus of pygmy tyrannosaurid, *Nanotyrannus*, based on a superb skull in the Cleveland Museum of Natural History, CMNH 7541 (Fig. 3; Plate 14). The eyes in *Nanotyrannus* had achieved a degree of forward orientation as great or greater than that of *Tyrannosaurus* and the head was very sharply flexed on the neck (Fig. 1). The anatomy of the braincase in the new genus is very advanced in a curious and unexpected way and suggests that *Nanotyrannus* evolved from an ancestor distinct from that of any well known tyrannosaur (Fig. 2). The type of *Nanotyrannus* was excavated by David Dunkle for the Cleveland Museum during the 1942 field season in the classic Latest Cretaceous outcrops of the Hell Creek Formation in north central Montana. Gilmore (1946) described the type of *Nanotyrannus* as a new species of *Gorgosaurus*, *G. lancensis*, but he never made detailed comparisons of the

teeth, braincase and occiput with those in specimens of *Gorgosaurus*. These structures in *Nanotyrannus* are so different from those in *Gorgosaurus*, *Albertosaurus*, *Tarbosaurus* and *Tyrannosaurus* that the new pygmy tyrannosaur may well represent a tyrannosaurid evolutionary line that branched off before the radiation of the well known large-bodied genera.

Institutional abbreviations — AMNH American Museum of Natural History, New York; CMNH Cleveland Museum of Natural History, Cleveland, Ohio; DMNH Denver Museum of Natural History, Denver, Colorado; FMNH Field Museum of Natural History, Chicago, Illinois; LACM Los Angeles County Museum, Los Angeles, California; MUO Museum of the University of Oklahoma; NMC National Museum of Canada, Ottawa, Ontario; P Tyrrell Museum of Palaeontology, Drumheller, Alberta; PIN Palaeontological Institute, Moscow, USSR; ROM Royal Ontario Museum, Toronto, Ontario; USNM United States National Museum, Washington D.C.; YPM Yale Peabody Museum, New Haven, Connecticut.

SYSTEMATIC PALEONTOLOGY

Nanotyrannus, genus novum

Type species — *Gorgosaurus lancensis* Gilmore, 1946 (*Albertosaurus lancensis* in Russell 1970).

Type specimen — CMNH 7541, complete skull and lower jaws in occlusion. Found by David Dunkle in 1942, in the Hell Creek Formation, SE 1/4, sec. 11, R61E, T4S, Sand Creek, Carter County, Montana; Lancian Faunal Age.

Referred specimen — DMNH uncatalogued, three teeth broken off at the crown-root junction, found by D. Reinheimer in 1924, in the Lancian beds of Corson County, South Dakota, accompanied by five juvenile *Tyrannosaurus* teeth, also broken off at the crown-root junction.

Diagnosis (Fig. 2). Differs from all other tyrannosaurids in having the following derived features: muzzle width greatly constricted to only one fourth the width of the temporal region; extraordinarily wide basicranium between basitubera and basiptyergoid processes; two large pneumatic foramina, placed one behind the other, near the midline of the basisphenoid. Differs from *Daspletosaurus*, *Gorgosaurus* and *Alioramus* and agrees with *Tyrannosaurus*, *Albertosaurus* and *Tarbosaurus* spp. in having the derived features of a marked expansion of the width across the temple, relative to the skull length, and basitubera that are displaced forward towards the basiptyergoid processes. Differs from all other tyrannosaurids, except *Alioramus*, in retaining the primitive characters of a long, low snout and maxillary teeth that are strongly compressed side-to-side. Differs from all other tyrannosaurids in lacking strong ridges and striae along the dorsal surface of the nasal. Differs from *Tyrannosaurus* in retaining the primitive character of a smooth depressed area for muscle origin on the anterior-ventral corner of the lachrimal.

Etymology. *Nano* — Latin "small", plus *tyrannus*, Latin "tyrant".

CHARACTER ANALYSIS

In North America there are three successive Land Faunal Ages that have produced tyrannosaurs: the Judithan, the Horseshoe Canyon Faunal Age, and the Lancian. The Lancian is the last fauna of the Cretaceous. Faunal turnover was rapid, and so each age has its diagnostic fauna of common horned dinosaurs and duckbills. The Judithan Age is characterized by the ceratopsians *Chasmosaurus* and *Centrosaurus* and by the duckbills *Corythosaurus*, *Lambeosaurus* and *Prosaurolophus*. The Horseshoe Canyon Age is characterized by the ceratopsian *Anchiceratops* and the hadrosaurs *Saurolophus* and *Hypacrosaurus*. The Lancian is defined by the horned dinosaur *Triceratops* and the duckbill *Edmontosaurus annectens*. It would be surprising if large predators had a

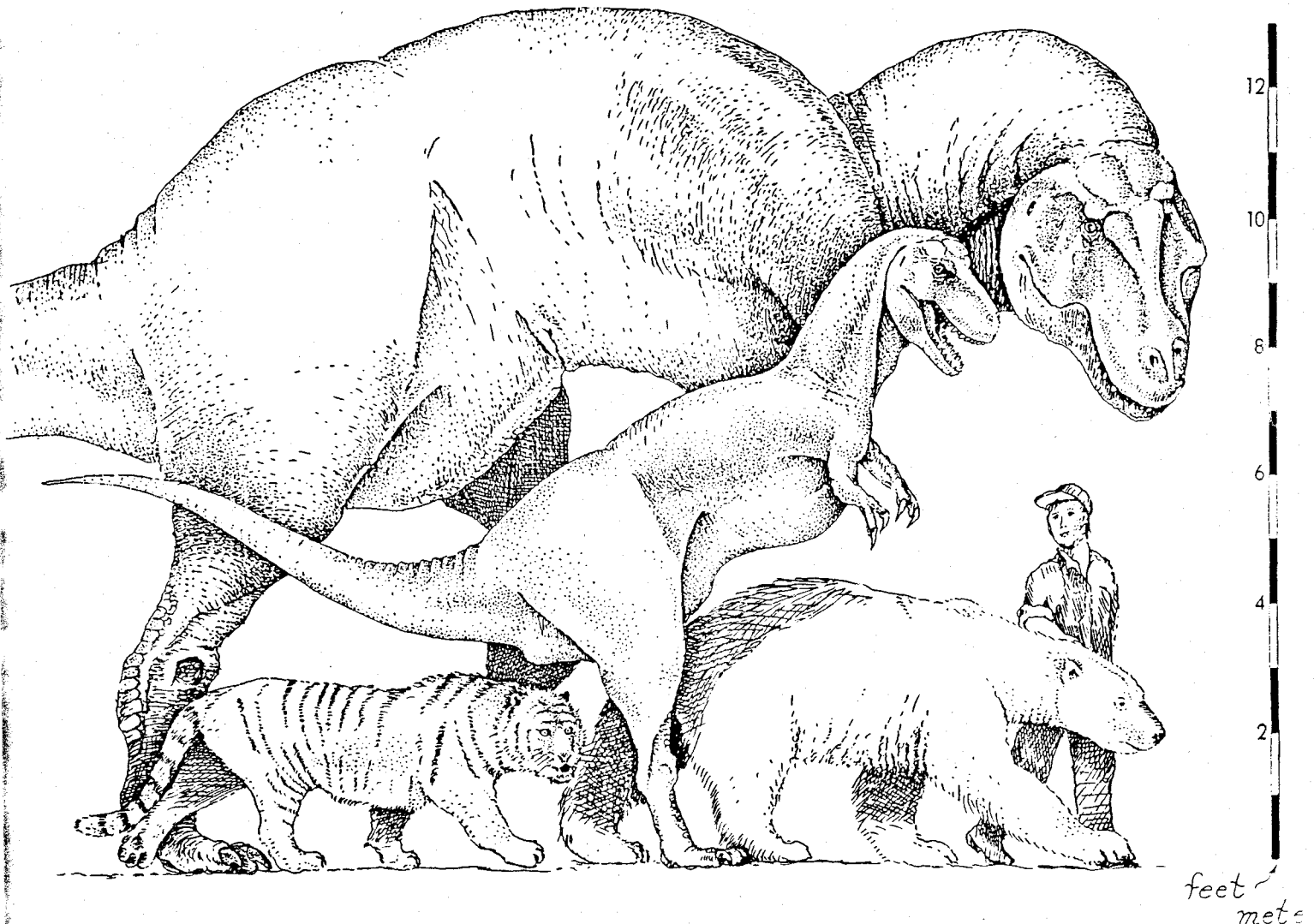
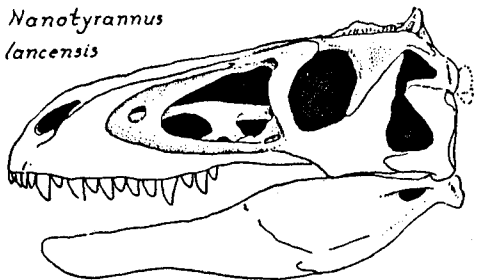
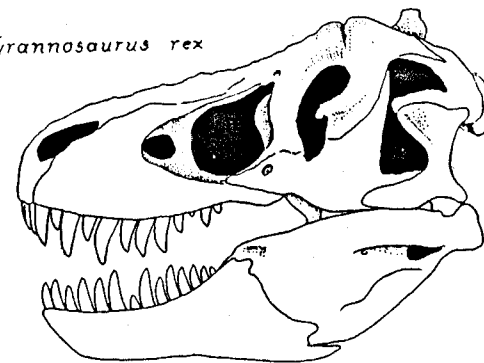


Figure 1—*Nanotyrannus*, the Cleveland Pygmy Tyrannosaur, shown in scale with an adult *Tyrannosaurus rex* and with the largest extant terrestrial meateaters, a Siberian tiger and a polar bear. The body frame of *Nanotyrannus* is taken from the type of *Gorgosaurus sternbergi*, scaled to match the skull size of *Nanotyrannus*.

*Nanotyrannus
lancensis*

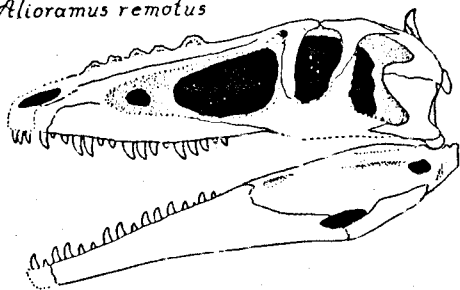


Tyrannosaurus rex

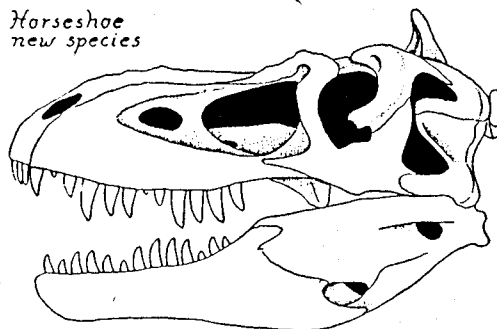


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Alioramus remotus

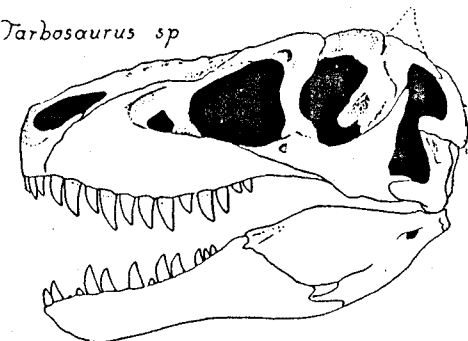


*Horseshoe
new species*



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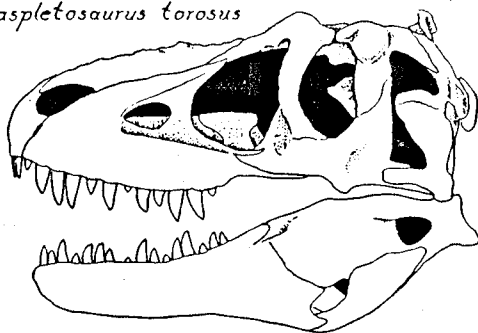
Tarbosaurus sp



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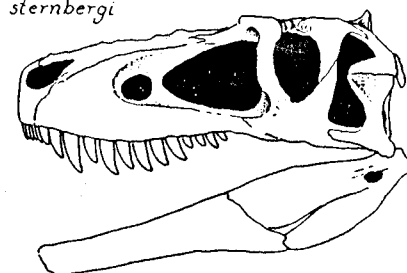
Daspletosaurus torosus



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*Gorgosaurus
G. sternbergi*



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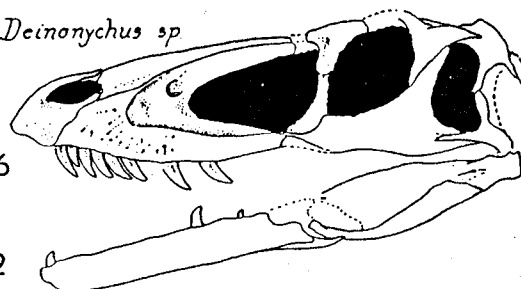
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Deinonychus sp



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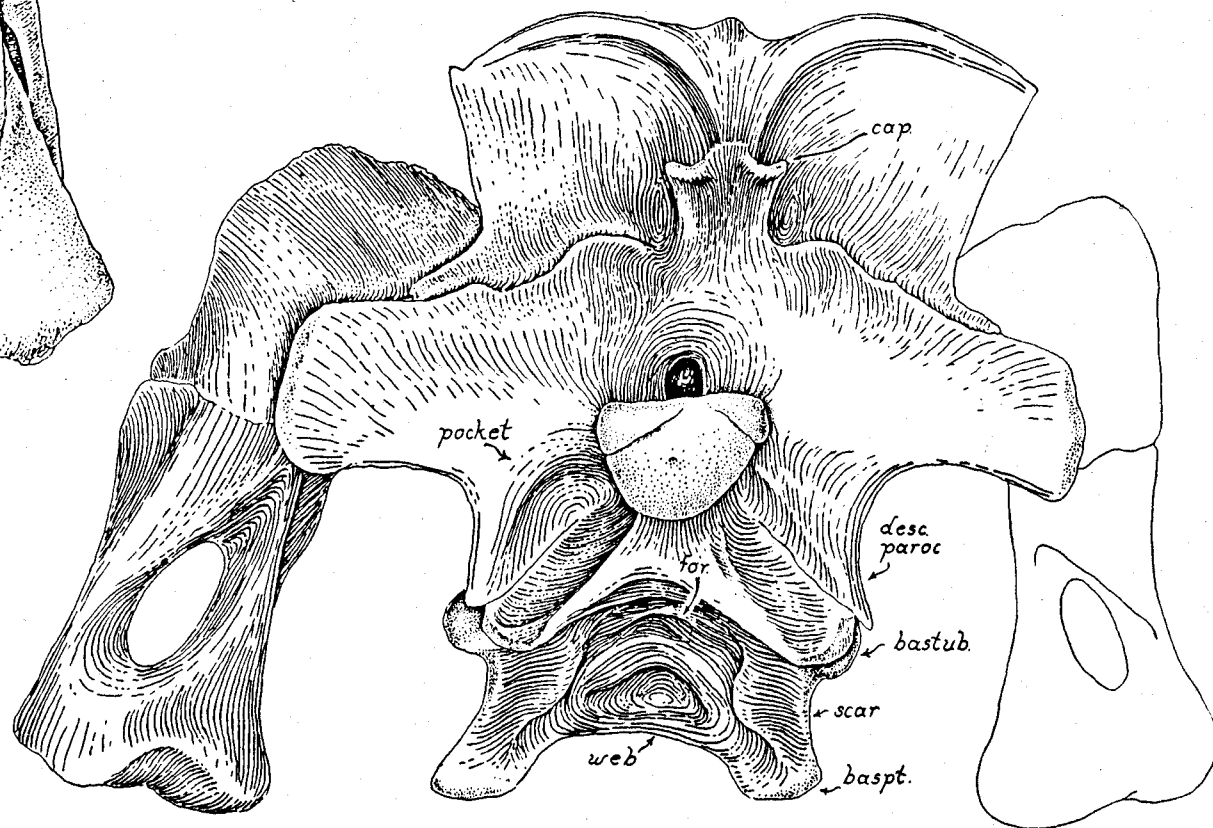
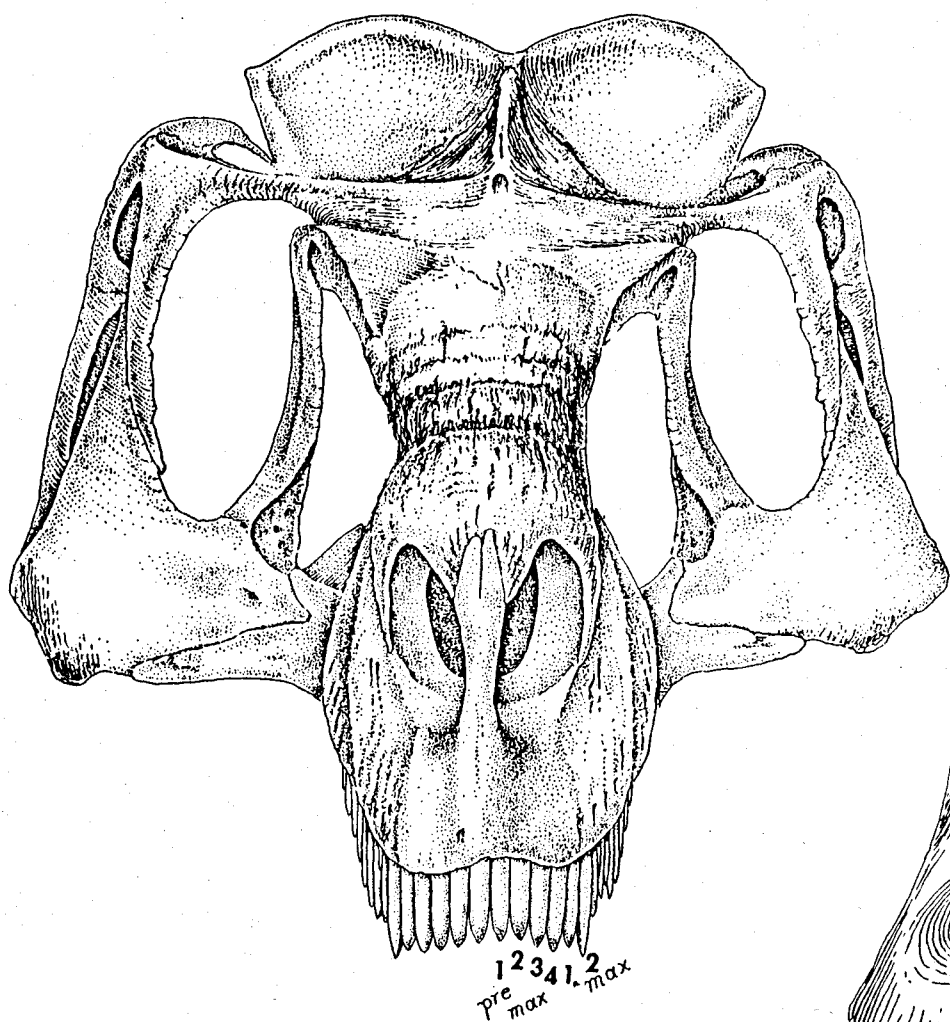
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slower rate of evolutionary turnover, and yet Gilmore's identification of CMNH 7541 as a *Gorgosaurus* would give this genus a time range spanning all three faunal ages. No other genus of large dinosaur lasts from the Judithan through the Lancian.

Figure 2—Branching diagram of the tyrannosaurids and their close allies, with lateral views of the skulls shown in correct stratigraphic sequence. Nodes and the derived characters that define them: 1) Neotheropoda (Late Jurassic-Latest Cretaceous) — premaxillary tooth crowns strongly asymmetrical, with inner (lingual) face nearly flat and outer (buccal) face strongly convex; premax. symphysis U-shaped in dorsal view; intramandibular joint fully developed, with anterior prong of the angular penetrating into the cavity between the dentary and splenial. 2) Ceratosauridae (Late Jurassic) — premaxillary tooth count reduced to three; premaxillary incisors with thick, strong sulci and ridges on the inner face. 3) Advanced neotheropods (Late Jurassic-Latest Cretaceous) — occiput much deeper above the foramen magnum, as seen in posterior view; accessory antorbital fenestra present; posterior shafts of cervical ribs do not overlap one another; presacral column compressed fore-to-aft relative to femur length; scapula blade very narrow throughout its length. 4) Allosauridae (Late Jurassic) — paroccipital process bent downwards strongly; basituber with a deep notch in the posterior-ventral edge for the ilio-costalis cervicis-capitis muscle; sphenethmoid ossification weak. 5) Very advanced neotheropods (Early Cretaceous-Latest Cretaceous) — ascending process of astragulus very tall, wide transversely and thin front-to-back; nasals narrow. 6) Dromaeosauridae (Early Cretaceous-Latest Cretaceous) (*Deinonychus*) — pubis turned backwards; second hindclaw very large and sickle-shaped; distal half of tail encased within basketwork of bony rods developed from chevrons and prezygapophyses. 7) Tyrannosauroidae (Early Cretaceous-Latest Cretaceous) — paroccipital process very deep top-to-bottom at the root; large excavation around the fenestra ovalis and pneumatization of the paroccipital root. 8) Acrocantosaurids (Early Cretaceous) — neural spine of cervicals and dorsals elongated. 9) Advanced tyrannosauroids (Late Cretaceous) — occiput deeper above the supraoccipital wedge; metatarsal bundle very long and compressed side-to-side, with strong pinching of the proximal end of metatarsal III. 10) Ornithomimids + troodontids + birds + ?oviraptorids (?Latest Jurassic-Latest Cretaceous) — periotic region with large depression and highly pneumatic. 11) Tyrannosauridae (Late Cretaceous) — adductor muscle scar developed forward over the frontals to a position opposite the orbits; squamosal-quadratojugal suture very long, straight and nearly parallel to the long axis of the skull, as seen in side view; supraoccipital wedge with two tabs of bone placed in tandem; first maxillary tooth like the four premaxillary teeth; all incisiform teeth very crowded and narrow across buccal face; parietal occipital wings very tall above the supraoccipital; large oval foramen in jugal. 12) *Nanotyrannus* (Latest Cretaceous, Lancian Faunal Age) — very wide basicranial boxwork with flat ventral floor; very wide frontal-orbital region with very narrow snout; parietal wing of occiput with sharp angle between dorsal and lateral edges. 13) Rough-snouted tyrannosaurids (Late Cretaceous) — dorsal surface of nasals very rough, with irregular longitudinal striae and ridges. 14) *Daspletosaurus torosus* (Late Cretaceous, Judithan Faunal Age) — snout and mandible short front-to-back and deep; teeth large and reduced in number; lacrimal horn developed into blunt triangular apex. 15) Tyrannosaurids with anterior pneumatic foramina in basicranial boxwork (Late Cretaceous). 16) *Alioramus* (Late Cretaceous, Nemegt Fauna) — multiple oval hornlets on nasals. 17) Massive snouted tyrannosaurids with anterior basicranial foramina (Late Cretaceous) — snouts and mandibles short and deep; tooth count reduced. 18) New genus and species from the Horseshoe Canyon Formation (Late Cretaceous) — orbit closed off from below by prong of postorbital. 19) Tyrannosaurids with large anterior foramina. 20) *Gorgosaurus* (Late Cretaceous, Judithan Faunal Age) — lacrimal horn developed into apex that is directed forward. 21) Tyrannosaurids with large foramina and wide basicrania (Late Cretaceous) — orbit closed off from below by postorbital; lacrimal and postorbital swollen above orbits; lacrimal swollen around pneumatic foramen; maxillary tooth row curved more strongly; maxillary tooth count reduced; mandible deeper; basicranial boxwork wider; first maxillary tooth enlarged. 22) *Tarbosaurus* (Late Cretaceous, Nemegt Fauna) — tooth crowns swollen and thick for their height. 23) *Tyrannosaurus* (Latest Cretaceous, Lancian Fauna) — teeth strongly procumbent; mandible very deep; lacrimal and postorbital very swollen above and behind orbit; muscle attachment surface, for anterior pterygoideus, at posterior-dorsal corner of antorbital fenestra eliminated by swelling of lacrimal; pneumatic foramen in lacrimal surrounded by grossly swollen bone; basicranium compressed fore-to-aft and basitubera displaced forward against basiptyergoid processes.

The most recent review of the tyrannosaurids in North America is that by Russell (1970), who recognizes two primitive genera, the gracile *Albertosaurus* and more robust *Daspletosaurus*, and one advanced genus, *Tyrannosaurus*. Both primitive genera, according to Russell, occur in the Judithan Age. Judithan tyrannosaurids are well documented by large samples from the Judith River Formation of Alberta. According to Russell's review, all named species of *Gorgosaurus* Lambe 1916 belong within *Albertosaurus* Osborn 1905; *Albertosaurus* and *Daspletosaurus* supposedly persisted into the next youngest sample, that of the Horseshoe Canyon. Finally, in the latest sample, from the Lancian Age, Russell identified two genera — a small species which he called *Albertosaurus lancensis* Gilmore 1946, and the very large *Tyrannosaurus rex*, Osborn 1903. Russell thus gave the genus *Albertosaurus* the time range of Judithan through Lancian, and this anomalously great generic longevity remained unexplained. The Mongolian Late Cretaceous Nemegt Formation has yielded two types of tyrannosaurid — *Tarbosaurus* Maleev 1955, intermediate in structure between *Albertosaurus* and *Tyrannosaurus*, and *Alioramus* Kurzanov 1976, a strange genus with a long snout ornamented with curious excrescences.

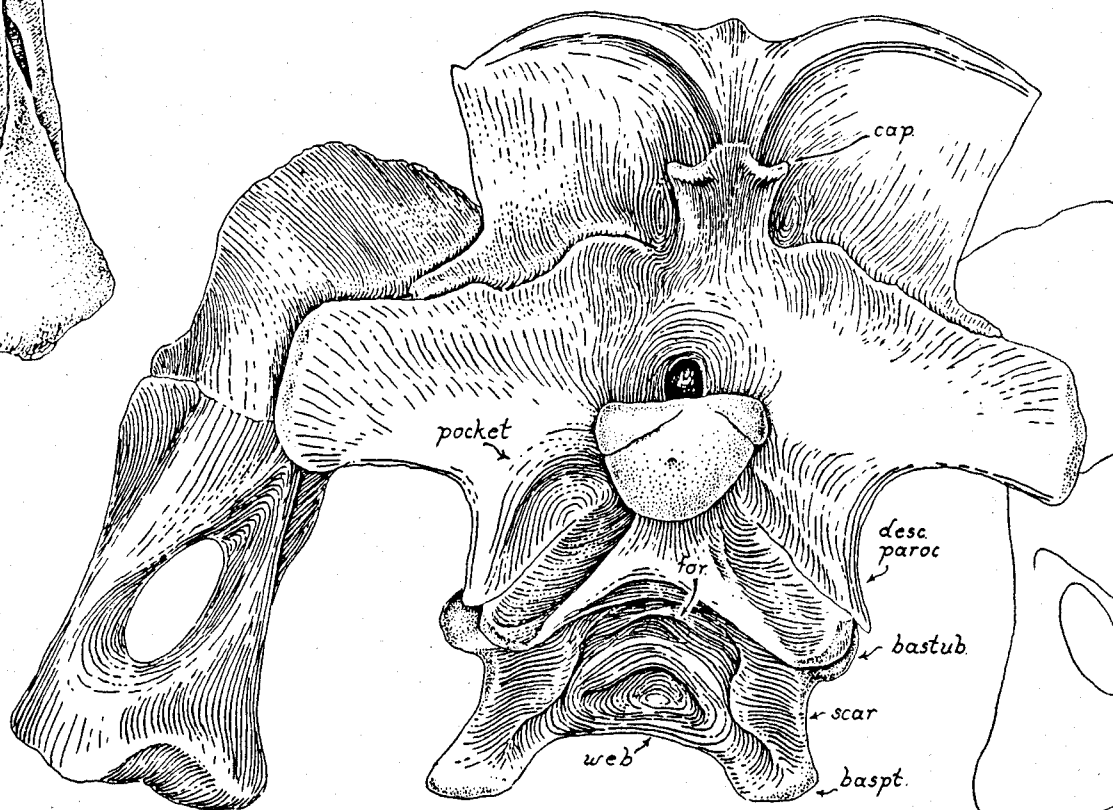
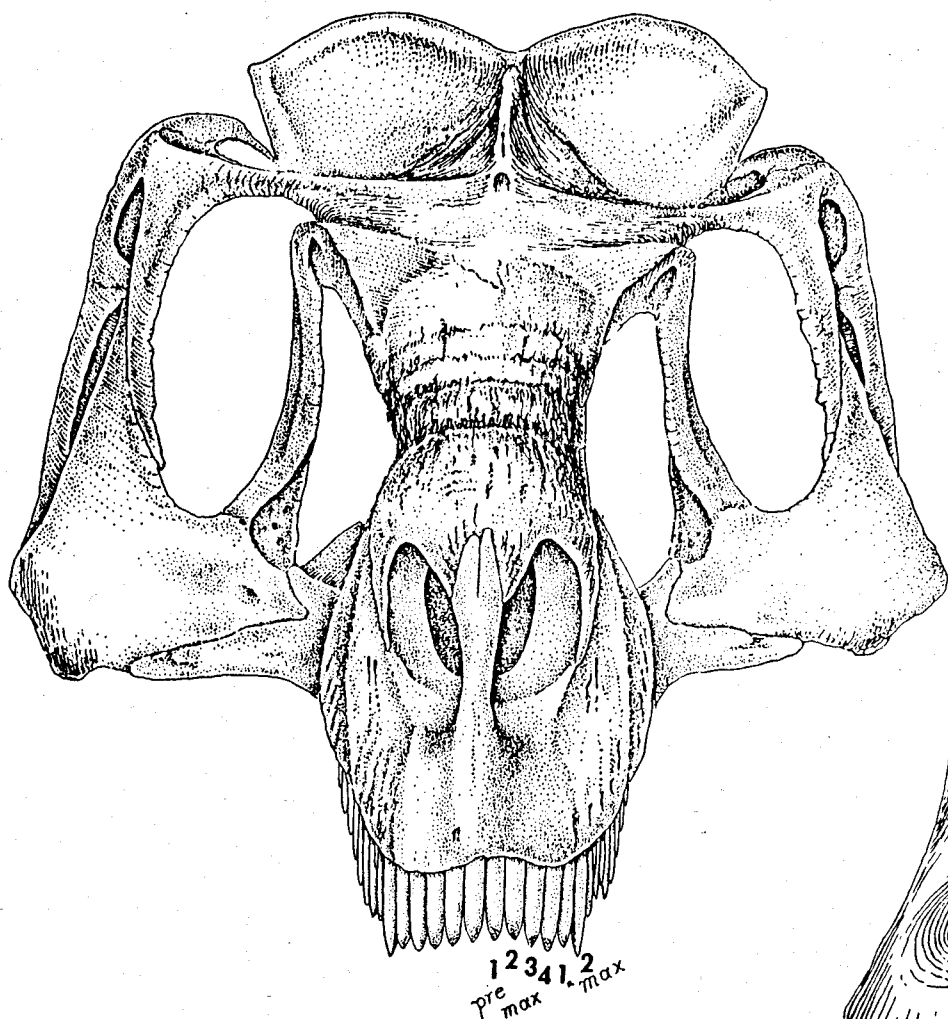
As we shall show in a subsequent publication (Bakker, Currie, and Williams, in press), the genus *Albertosaurus* is distinct from the genus *Gorgosaurus*, and these two genera, plus *Tarbosaurus* and *Tyrannosaurus*, make a clade distinct from *Daspletosaurus*. In the following analysis, we restrict the concept of *Albertosaurus* to the type and paratype specimens NMC 5600 & 5601, both from the Horseshoe Canyon Formation. A great variety of tyrannosaurs have been grouped together in *Gorgosaurus* and we are confident that at least two species can be diagnosed (Fig. 11). In the Judith River sample there is a gorgosaur, represented by FMNH 303, with relatively small teeth and a high tooth count (15 in the maxilla). In addition, there is a Judith River gorgosaur, represented by USNM 12814 and AMNH 5336, with relatively large teeth and a low tooth count (13 in the maxilla). Both of these categories have robust cranial bones and occipital condyles that are depressed top-to-bottom, suggesting that movement of the head on the neck was concentrated in a horizontal plane. Several additional Judithan gorgosaur specimens, such as ROM 1247 and AMNH 5664, have unusually delicate braincases and occipital condyles that are spherical and not depressed. Russell (1970) identified these delicate gorgosaurs as juveniles of the robust species but they may, in fact, represent a distinct species, *Gorgosaurus sternbergi* Matthew and Brown 1922. Unfortunately the type specimen of *Gorgosaurus* is the type of *G. libratus* Lambe 1916, a skeleton that lacks a braincase and is very badly broken in the muzzle, and the correct nomenclature for gorgosaurs is hard to work out. Therefore, in the following discussion, we will use the informal categories "gracile gorgosaur", "small-tooth gorgosaur", and "big-tooth robust gorgosaur". A new genus of gorgosaur-like tyrannosaurid is represented by Tyrrell Museum P 81.3, a nearly complete skull and much of the postcranium from the Horseshoe Canyon Formation. This



CMNH 7541

Nanotyrannus lancensis type

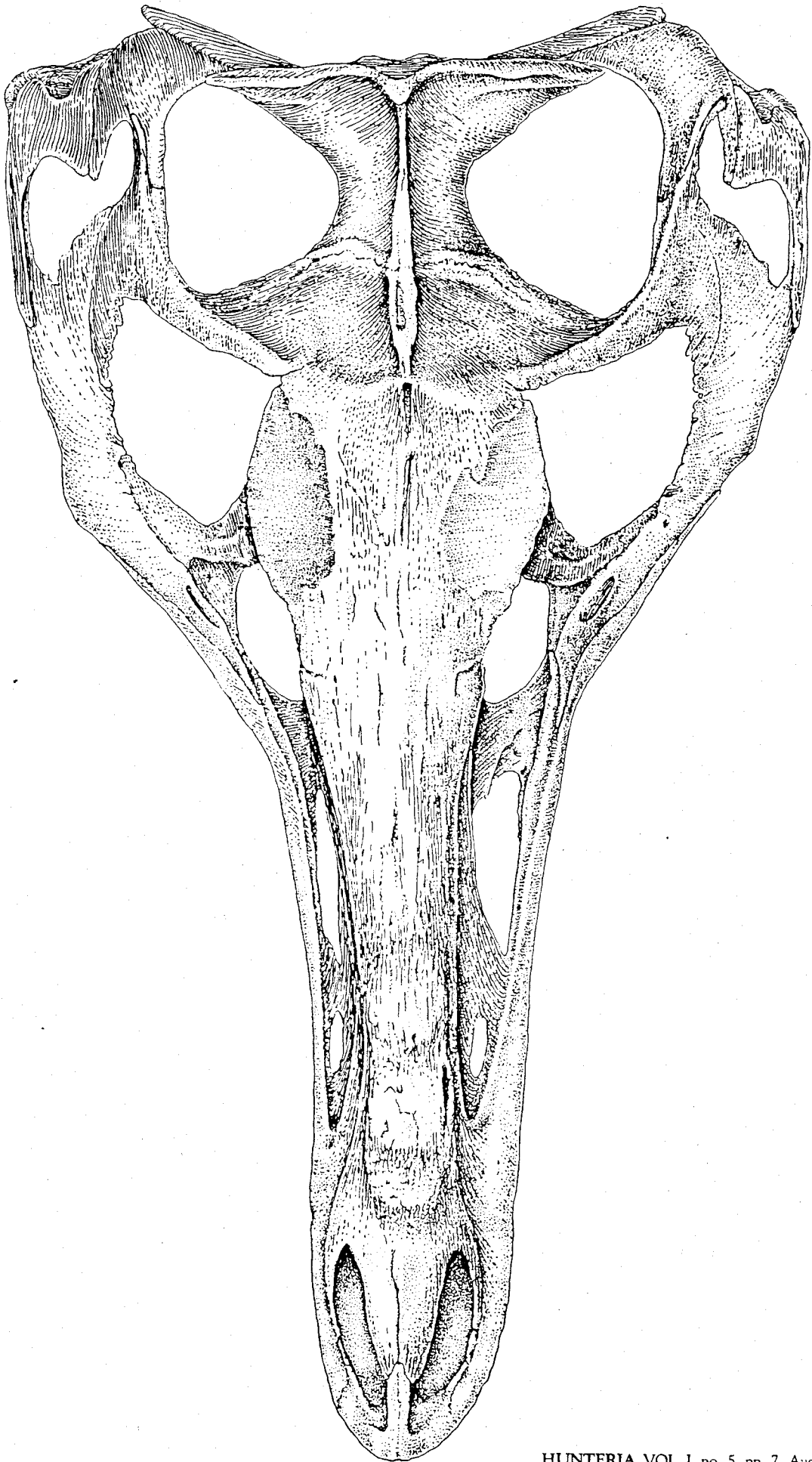
Figure 3—Front and occipital views of the type of *Nanotyrannus lancensis*. The twisting distortion of the muzzle has been corrected. The right posterior corner of the temporal region has been restored from the left side and the forward compression of the quadratojugal has been removed.



CMNH 7541

Nanotyrannus lancensis type

Figure 3—Front and occipital views of the type of *Nanotyrannus lancensis*. The twisting distortion of the muzzle has been corrected. The right posterior corner of the temporal region has been restored from the left side and the forward compression of the quadratojugal has been removed.



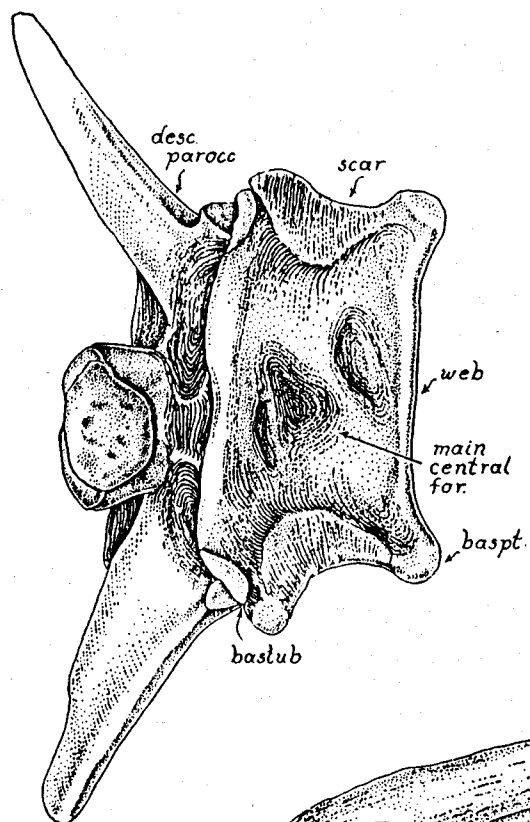
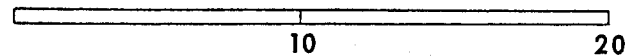
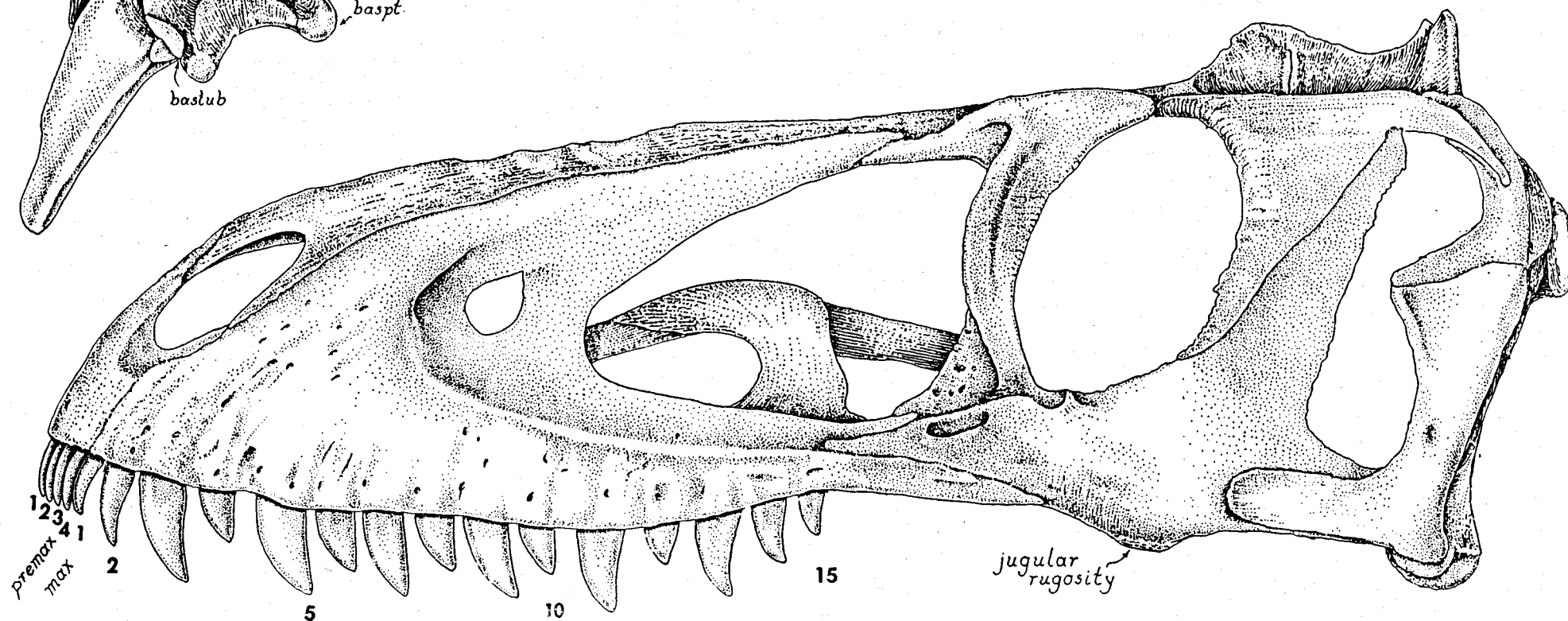


Figure 3 continued—Lateral view of skull and ventral view of basicranium of *Nanotyrannus lancensis*.



skull shows a mixture of features seen separately in *Alioramus*, *Gorgosaurus*, and *Tarbosaurus*.

Basicranium. We introduce here a few descriptive terms that are useful in comparing theropod braincase structures. The transverse web of bone connecting right and left basiptyergoid processes is the "basiptyergoidal web" (bpt.web in Fig. 4). The transverse web of bone connecting right and left basituberal processes is the "basituberal web" (bastub.web in Fig. 4-6). The wall of basisphenoidal bone running fore-to-aft connecting the basiptyergoid web to the basituberal web on either side is the "basicranial boxwork wall" (boxwork wall in Fig. 5). The two boxwork walls and the two transverse webs constitute the "basicranial boxwork".

The most outstanding feature of *Nanotyrannus* is the enormous lateral expansions of the basicranial boxwork, relative to the length of the skull (Fig. 3-5). In primitive theropods, such as *Ceratosaurus*, the width across the basitubera is only about 8% of the skull length (USNM 4735; Fig. 4; Table 1). In daspletosaurs and in most gorgosaurs the basituberal width reaches 12% to nearly 15% of the skull length. In *Tyrannosaurus* the basituberal width is enormously expanded, reaching 19% of the skull length (Fig. 4-6; Table 1). In *Nanotyrannus* the basituberal width is fully 20% of skull length as we restore the skull, and the condition of the specimen strongly suggests that the basicranium has been preserved with no distortion of width. Such lateral expansion of the basitubera as is seen in *Nanotyrannus* would greatly increase the leverage available to the neck muscles for flexing the head sideways.

The basitubera in tyrannosaurs are composed of three units: 1) The main tuberos process (tub. in Fig. 4-5) is a swollen development of the basioccipital at the posterior-outer corner of the basicranial boxwork. In gorgosaurs and daspletosaurs the tuberos process is very thick but in *Albertosaurus*, *Tyrannosaurus* and *Nanotyrannus* it is reduced in bulk, a change that reflects a decrease in the size of the muscle tendon attaching to the process. 2) A line of rugose muscle attachment scars ascends the posterior face of the basioccipital from the tuberos process. This ascending scar complex is well developed in *Nanotyrannus*, as it is in most other tyrannosaurs, and marks the attachment of the longus colli muscle (= rectus capitis anterior of human anatomy). 3) Anterior to the tuberos process, along the ventral-lateral rim of the basicranium, is a large, flat, oval scar (scar in Fig. 4-6) marking the insertion of the iliocostalis cervicis capitis muscle. This oval scar is small and weakly striated in *Nanotyrannus*, compared to the state in other tyrannosaurs, suggesting that the muscle attachment was fleshy rather than tendinous.

In primitive theropods and most tyrannosaurs, the ventral surface of the basisphenoid is excavated by a huge central cavity (cav. in Fig. 5) bounded by the basiptyergoidal web in front, the basituberal web behind, and the basicranial boxwork walls on either side. Foramina of various sorts penetrate the ceiling of this cavity. In primitive theropods the cavity is very deep and a pair of tiny vascular foramina penetrate the anterior part of the cavity, just anterior to the basituberal web, a condition retained in allosaurids, dromaeosaurids and acrocanthosaurs (Fig. 4, 9). Among tyrannosaurids this primitive state is known only in *Daspletosaurus* (Fig. 8). In

TABLE 1

	Skull L = premax- quadrate	Skull length occ con-premax	Bastub W	BastubW/ skull L	Quadrato- jugal W (QQ)	QQ/skull L	Snout W at mid-length	Snout W/ skull L	QQ/ Snout W	Maxilla tooth count
<i>Nanotyrannus lancensis</i> CMNH 7541	572	550	117	0.20	325	0.571	75	0.13	4.33	15
<i>Alioramus remotus</i> PIN 3141	637	78.3		0.123						15++
<i>Gorgosaurus libratus</i> NMC 2120	950e									15
<i>Gorgosaurus sternbergi</i> AMNH 5664	678				264e	0.390				15
<i>Gorgosaurus</i> sp. AMNH 5458	990				330	0.333				
<i>Gorgosaurus</i> sp. AMNH 5336	962	980	122	0.124	380	0.388	232	0.241	1.70	15
<i>Gorgosaurus</i> sp. FMNH 308 = AMNH 5434	1,050		120e	0.114	347e	0.333				15
<i>Gorgosaurus</i> sp. USNM 12814	791		114	0.114	332	0.420	166	0.210	2.00	15
<i>Gorgosaurus</i> sp. ROM 1247	782		92.3	0.118	289e	0.370				15
<i>Daspletosaurus torosus</i> NMC 8506	1,040	1,040	144	0.139	425	0.409	329	0.313	1.31	12
<i>Tyrannosaurus rex</i> AMNH 5027	1,355		256	0.190	835	0.616	411	0.303	2.03	12
<i>Tyrannosaurus rex</i> Tyrrell P81.6.1	1,300e		234	0.18	750	0.577				12
<i>Tyrannosaurus rex</i> LACM 23844	1,475e				981e	0.665				12

the long-snouted Mongolian tyrannosaurid *Alioramus* and in the new genus of tyrannosaurid from the Horseshoe Canyon the paired foramina are displaced forward but are still relatively small. In gorgosaurs the foramina are anterior in position and greatly enlarged. These enlarged foramina lead to a complex system of pneumatic spaces within the braincase. In *Albertosaurus* and *Tyrannosaurus* the foramina are still larger than those in gorgosaurs (Fig. 4-6, 11). *Nanotyrannus* has a unique foraminal pattern — there is one large foramen anteriorly almost on the ventral midline of the basisphenoid and a second large foramen, with an irregular outline, off-center further posteriorly in front of the basituberal web (Fig. 3, 4). Preparation of the matrix filling the foramina is not yet complete, but one or both almost certainly lead inward to the pneumatic spaces. It is not obvious how the *Nanotyrannus* system could evolve from the gorgosaur pattern of bilaterally symmetrical and paired foramina. It is quite possible that the pneumatic system of *Nanotyrannus* evolved independently from the gorgosaur-albertosaur-*Tyrannosaurus* clade.

Most Late Cretaceous theropod dinosaur families show trends for increasingly complex pneumatization of the braincase (Currie, 1985; Bakker, Currie, and Williams, in press). The functional significance of large pneumatopores is not simply to "lighten the skull" because hollow bones have evolved in some birds without the development of large pneumatic foramina. Furthermore, all modern crocodilians have skulls full of pneumatic spaces and yet the pneumatic foramina opening to the outer skull bones are all very small. Clearly large pneumatopores are not necessary for the development of thin-walled, hollow bones. The relatively huge foramina in *Nanotyrannus*, *Gorgosaurus*, *Albertosaurus* and *Tyrannosaurus* suggest that there was easy movement of air through the holes. Some modern predators, such as the cheetah (*Acinonyx jubatus*), incur great exercise heat loads in their heads during running or subduing struggling prey (Taylor et al., 1973). A relatively small brain, such as that possessed by tyrannosaurs, could reach lethal temperature more rapidly than a large brain. It may be the air from the throat was circulated up into the basicranium in tyrannosaurs,

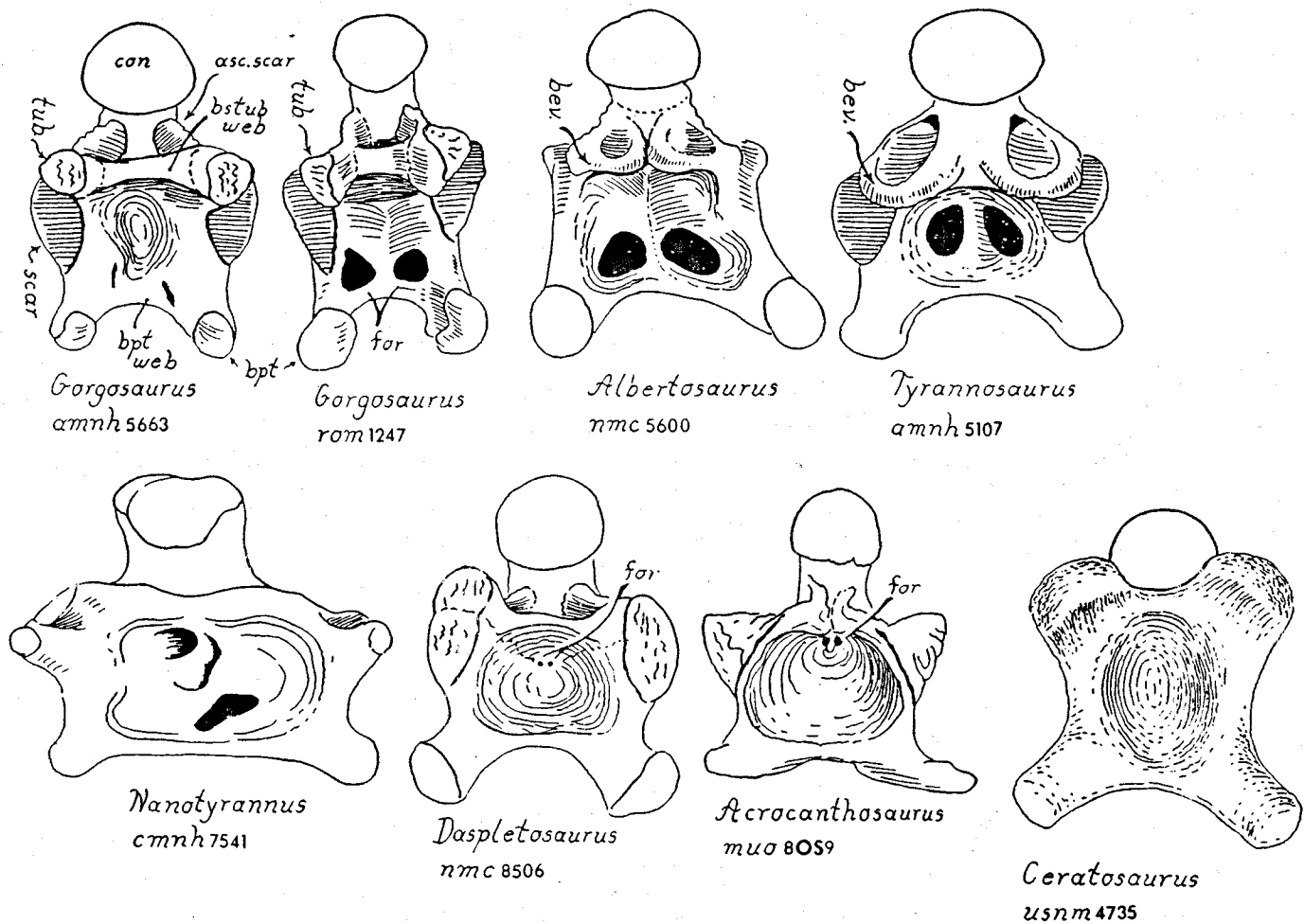


Figure 4—Ventral views of theropod basicrania, drawn to a common distance of the occipital condyle-to-basiptyergoid web.

Osteological abbreviations: asc scar ascending muscle scar on the posterior-inner corner of the basituber. basiocc basioccipital. basph basisphenoid. baspt basiptyergoid processes. basiph scar oval scar on the basisphenoid boxwork wall. bastub basituber. cav midline ventral cavity in the basisphenoid. con occipital condyle. desc paroc descending ventral root of the paroccipital process. mid scar midline scar on the posterior face of the basisphenoid. pa.wing parietal wing of the occipital plate. ser.rug serrated tendon scar on the outer corner of the parietal wing. sup.cap caps on the supraoccipital wedge. web transverse web of bone between basiptyergoid process or basisphenoid processes. wedge supraoccipital wedge.

and the air was cooled before entry by evaporation along the walls of the mouth. Alternatively, the large foramina may be part of adaptations for increasing the sensitivity of the ears to certain frequencies (Currie, 1985). These explanations are

not mutually exclusive, and both cooling and auditory functions may be part of the adaptive design in tyrannosaurs.

In primitive theropods and in gorgosaurs the ceiling of the central basisphenoidal cavity is very tall and the cavity is very

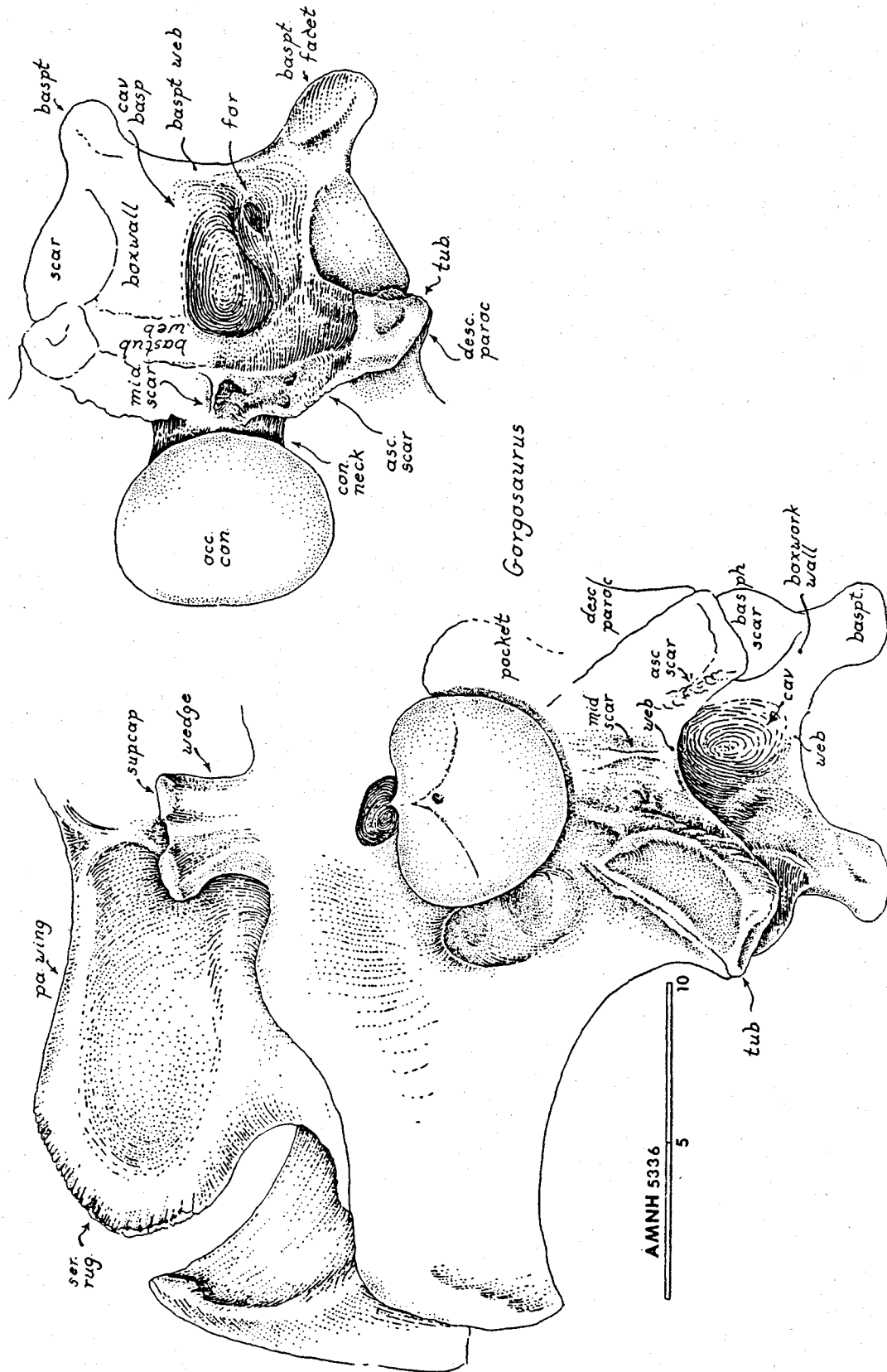


Figure 5—Posterior and ventral views of the braincase of a large-toothed gorgosaurus with a dorsal-ventrally compressed condyle. The pneumatic foramina have been closed into slits by dorsal-ventral distortion incurred after death.

deep top-to-bottom. The cavity remains very deep in daspletosaurs and gorgosaurs, and in gorgosaurs the large pneumatic foramina face mostly inwards from their location on the steep inner surfaces of the boxwork walls (Fig. 4-6, 8).

In *Nanotyrannus* the ventral floor of the basisphenoid is nearly flat between the basituberal and basiptyergoidal webs, a condition seen in *Tyrannosaurus* and, to a lesser extent, in *Albertosaurus*. In the last two genera and, probably, in *Nanotyrannus* as well there are very large pneumatic chambers above the basisphenoidal floor, filling in the space that is occupied by the primitive open cavity in gorgosaurs (Fig. 3-6, 9).

The occipital condyle is carried on a constricted neck in *Nanotyrannus* and the neck faces far more strongly

downward than in gorgosaurs and daspletosaurs, or in ceratosaurs and allosaurs, agreeing with the condition in *Tyrannosaurus*. Moreover, the articular surface on the condyle in *Nanotyrannus* has been displaced downwards and around onto the ventral surface of the neck far more than in gorgosaurs and daspletosaurs, again agreeing with *Tyrannosaurus* (Fig. 6, 9). Clearly the head was flexed far more sharply downwards in *Nanotyrannus* and *Tyrannosaurus* than in more primitive tyrannosaurids. The type and paratype specimens of *Albertosaurus* are badly broken

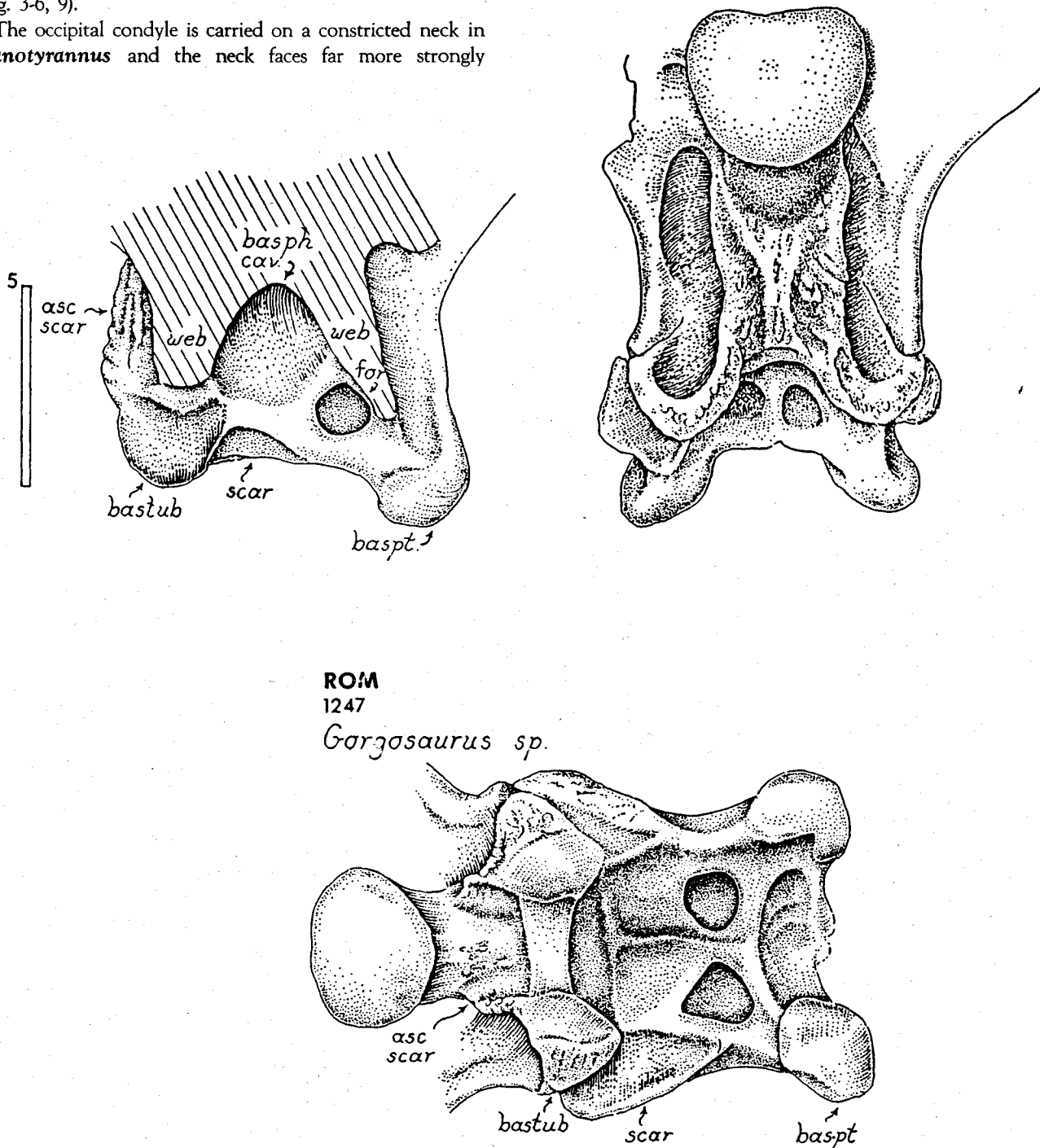


Figure 6—Posterior, ventral, and sagittal cross section of a subadult gorgosaur braincase.

but, as far as is known, *Albertosaurus* approaches *Tyrannosaurus* in these adaptations for head-neck flexure and is distinctly more advanced than *Gorgosaurus*. No complete braincase has been described for the Mongolian *Tarbosaurus* but the available evidence suggests that the structure is intermediate between *Albertosaurus* and *Tyrannosaurus*.

The orientation of the basitubera must change to maintain leverage when there is an evolutionary increase in the flexure at the head-neck joint. In an animal that frequently holds its head out straight, in line with the cervical vertebrae, such as a modern crocodilian, the basitubera must project directly downwards to maximize the leverage of the longus colli pulling directly backwards (Fig. 7, 10). But, in an animal carrying its head flexed at nearly a right angle to the neck, such as an ostrich, the basitubera must be displaced forward along the basicranium to achieve the same leverage. With the head flexed at a right angle to the neck, the basitubera need not project downwards at all, because leverage of the longus colli is controlled entirely by the distance from the center of rotation of the condyle to the floor of the braincase at the base of the basitubera. In primitive tyrannosaurs, such as the gracile gorgosaurs, the basitubera are located far aft, beneath the

occipital condyle, and project strongly downward (Fig. 7, 10). But in *Tyrannosaurus* and *Nanotyrannus*, the basitubera are displaced forward and project less strongly below the floor of the basicranium. Thus these two genera were equipped for powerful head-neck movements in a vertical plane in an arc that was, on average, far more flexed than in gorgosaurs or in primitive theropods generally.

The occipital condyle, as seen from behind, is triangular, with rounded corners, in *Nanotyrannus*, and resembles that of gracile gorgosaurs rather than the dorsal-ventrally depressed condyle of *Tyrannosaurus rex* and some robust gorgosaurs (Fig. 3, 5, 6, 9). The triangular shape suggests that side-to-side movement was more extensive when the head was extended in line with the neck than when it was sharply flexed downwards.

In *Nanotyrannus*, the occipital plate above the foramen magnum is reinforced by a vertical midline wedge of supraoccipital bone (wedge in Fig. 5) that is capped dorsally by two rounded tab-like processes placed side by side. The tandem-cap arrangement is unique to the Tyrannosauridae and found in all genera of the family (Fig. 3, 5, 6, 9). The supraoccipital wedge received the insertion of the short rectus capitis (=

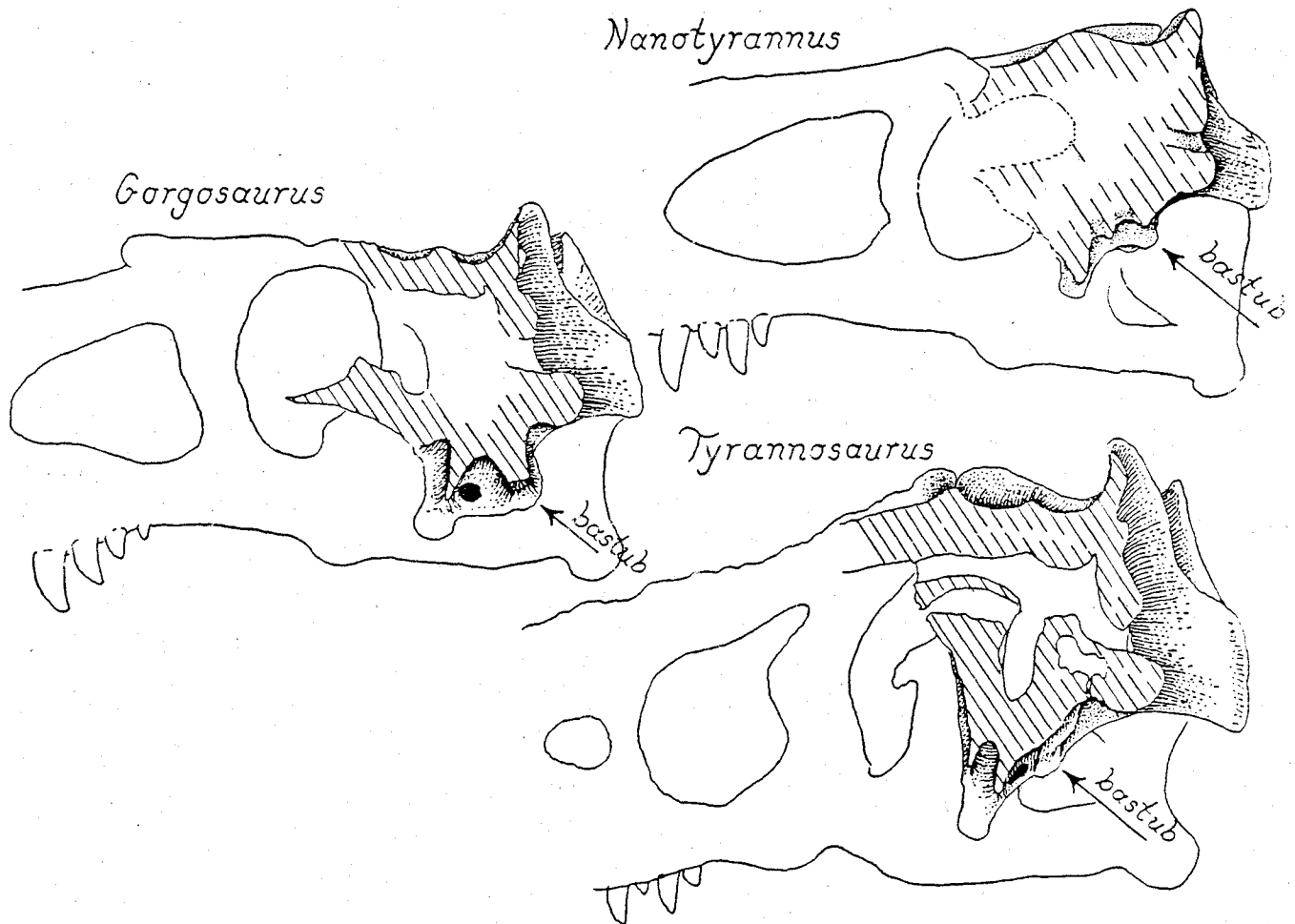


Figure 7—Sagittal sections through the posterior parts of the skulls in tyrannosaurs to show the arrangement of the braincase. The internal arrangement of pneumatic spaces is known only for *Tyrannosaurus*. Arrow shows the ventral apex of the basituber. *Tyrannosaurus* skull from AMNH 5027 (dermal bones and basiptyergoid processes); AMNH 5029 (braincase section) and AMNH 5107 (basitubera). *Gorgosaurus* skull from USNM 12814 (dermal bones) and ROM 1247 (braincase).

rectus capitis posterior of human anatomy) running forward from the top of the axis neural spine; the two caps probably received two tendons of that muscle. Above and either side of the tandem caps in *Nanotyrannus*, as in other tyrannosaurs, is a great expanse of the parietal wings (pa. wing in Fig. 5). The central part of the posterior face of the parietal wing is concave and must have received the insertion of a thick fleshy bundle of the long rectus capitis originating from the tops of the cervical neural spines. The dorsal margin of the parietal wings is thickened and must have received the insertion of the spinalis capitis, the most dorsal cervical muscle. In North American tyrannosaurs other than *Nanotyrannus*, the outer corners of the parietal wings are rounded in contour and are ornamented with serrated rugosities, indicating a tendinous attachment of the outer part of the spinalis capitis muscle. But in *Nanotyrannus* the dorsal margin of the parietal wing meets the lateral margin at a sharply defined angle and the rugosities are poorly expressed, indicating a less tendinous attachment here. Right and left parietal wings meet above the supraoccipital wedge at the dorsal occipital midline and the bone here is thickened and striated in a vertical direction, indicating the attachment of a thick tendinous sheet that must have separated the right and left halves of the spinalis capitis-rectus capitis bundles.

Temporal Region. In *Nanotyrannus* the width of the superior temporal fenestrae and of the entire temporal region is greatly expanded relative to the snout (Fig. 3, 11). The breadth between the jaw joints, measured between quadratojugals, is 57% of the skull length; the breadth of the snout at the midpoint of the main antorbital fenestra is only 13% of the skull length, so that the quadratojugal width/snout width ratio is 4.3 (Table 1). In the best preserved *Tyrannosaurus* skull, AMNH 5027, the quadratojugal width is 62% of skull length but the muzzle is not nearly as constricted as in *Nanotyrannus* and the ratio of quadratojugal width to mid-snout width is only about 2. No gorgosaur or daspletosaur skull known shows a ratio of quadratojugal width to skull length or to muzzle width anywhere nearly as high as in *Nanotyrannus*. The type of *Nanotyrannus* has been subjected to slight postmortem distortion — the occiput is pushed forward slightly, crushing the quadrate forward, and the snout is twisted about its long axis so that the left external naris is rotated to the right. But in general the specimen shows virtually no compression lines or fractures and the extraordinary constriction of the muzzle cannot be ascribed to postmortem deformation.

The seven best Judithan Age gorgosaur specimens are distorted in various directions but not one even approaches the shape of *Nanotyrannus* (Fig. 11, Table 1). All the available evidence indicates that dorsal-ventral compression, suffered by a theropod skull during the processes of burial and sediment compaction, does not expand the skull width at the rear. The type of *Gorgosaurus sternbergi* (AMNH 5664) is compressed from top-to-bottom but this distortion has simply collapsed the skull roof down onto the bones along the jaw margin — there has been no spreading of the skull between

the jaw joints and the quadratojugal breadth is only 39% of the skull length. In AMNH 5336, a robust, big-toothed gorgosaur, the skull has suffered moderate dorsal-ventral distortion that has compressed the parietal wings of the occiput downward but the quadrates remain nearly vertical and the quadratojugal breadth is only 39% of the skull length. In USNM 12814, a gorgosaur skull preserved with slight oblique distortion in a dorsal-lateral direction, the quadratojugal breadth is 42% of the skull length. This skull is probably the least distorted known for a gorgosaur and shows a sharply defined break in the skull contour, as seen in dorsal view, between the muzzle and the temporal region, but the degree of temporal expansion is very weak compared to that seen in *Tyrannosaurus* and *Nanotyrannus*. The type of *Daspletosaurus torosus*, NMC 8506, is preserved with no visible distortion (Russell, pers. comm. 1988); the muzzle is wider than that of gorgosaurs, and the temporal region is expanded only slightly relative to muzzle breadth. In the gracile gorgosaur skull ROM 1247, moderate side-to-side compression has reduced the quadratojugal breadth to 26% of skull length; in life the ratio was probably about 37%. The robust, small-toothed gorgosaur skull FMNH 308 was chosen by Russell (1970) to illustrate the structure of common Judithan species *G. limbatus*, but this specimen is severely distorted side-to-side, with the quadratojugal breadth only 25% of the skull length; in life the ratio was probably similar to that of ROM 1247. The robust, big-toothed gorgosaur skull AMNH 5458 (the specimen mounted in a running pose in the public gallery of the American Museum) shows a little evidence of modest lateral distortion and has a quadratojugal width 33% of skull length.

It is especially important to note that, of the seven best Judithan skulls, three show slight to severe side-to-side compression (FMNH 308, AMNH 5434, ROM 1247) and none show strong top-to-bottom compression. In contrast, the

Daspletosaurus torosus

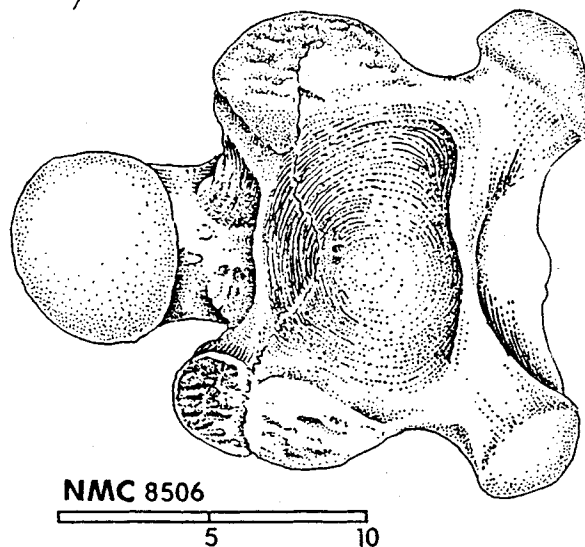


Figure 8—Ventral view of the basicranium of the type of *Daspletosaurus torosus*. Note the presence of a pair of small, anterior foramina within the central cavity.

sample of Lancian tyrannosaurid skulls shows no example of side-to-side distortion — the type of *Nanotyrannus* and the three best *Tyrannosaurus* skulls, AMNH 5027, LACM 23844, and Tyrrell Museum P 81.6, are undistorted or show slight to very slight dorsal-ventral compression and very slight front-to-back shear. Surely the natural shape of the skull will determine the pattern of postmortem distortion — after death, skulls that are relatively tall and narrow will tend to fall on their sides and be so buried, whereas skulls that are wide will tend to rest right-side-up on the substrate.

In the type of *Nanotyrannus* the quadrates have a slight outward-and-downward slant as seen in posterior view (Fig. 3), a condition repeated in *Tyrannosaurus* and in the best preserved gorgosaurus and daspletosaurs (USNM 12814,

NMC 8506). As in all tyrannosaurs, the *Nanotyrannus* quadrate is very wide distally relative to the shaft length so that distal width is only slightly less than maximum quadrate height. In primitive theropods such as podokesaurus and ceratosaurs the quadrate shaft is much longer relative to the distal articular width. Allosaurs have a quadrate that is very close in proportions to that of tyrannosaurs.

As a consequence of the temporal breadth/muzzle breadth ratio in *Nanotyrannus* and *Tyrannosaurus*, right and left eyes have a wide overlap of visual fields. Stereoscopy was probably present over a range of 30 degrees or more. Since the orbits face upwards as well as forwards in *Tyrannosaurus*, the widest possible arc of visual field overlap lies in a plane that is tilted upwards and forwards relative to the muzzle. In

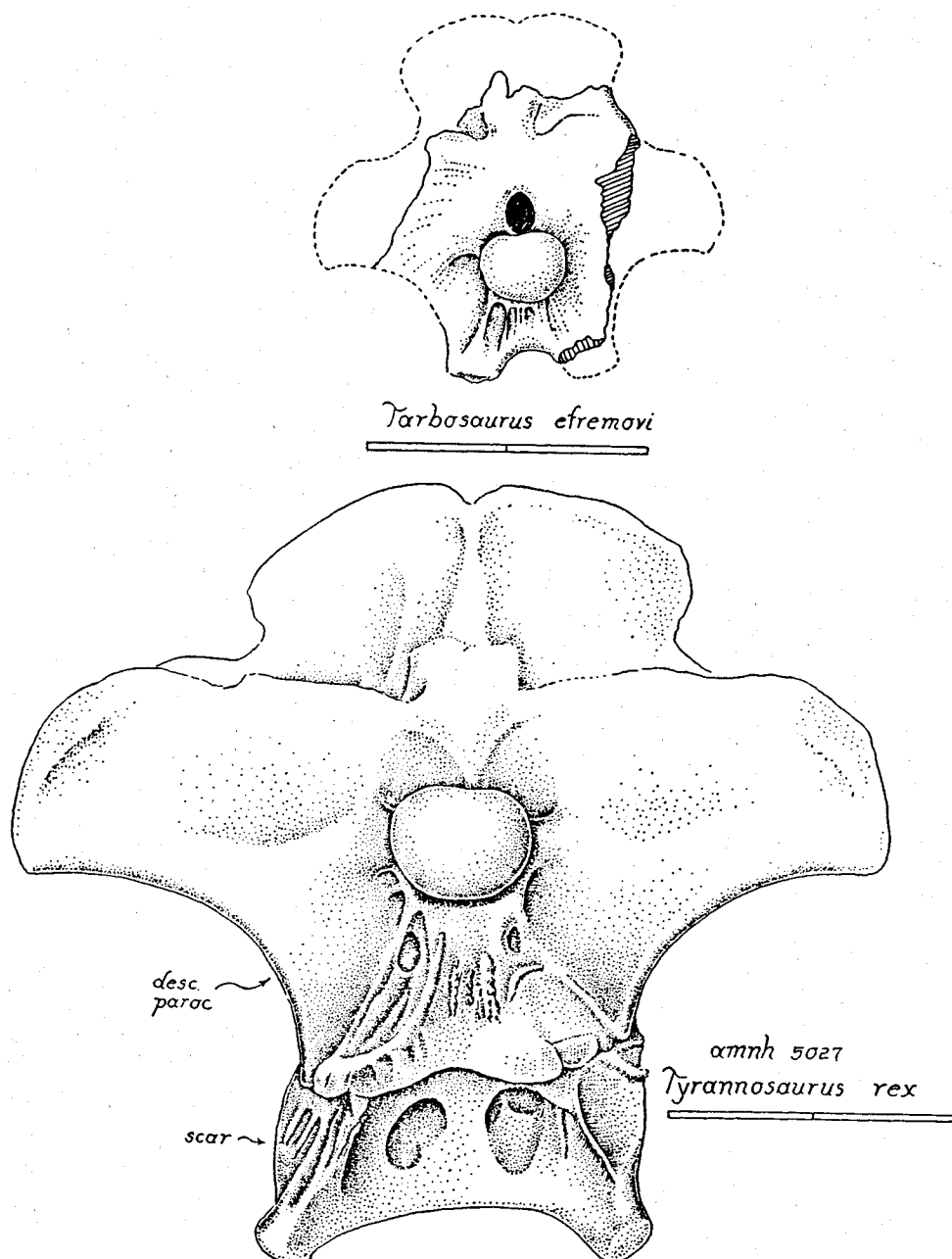


Figure 9—Posterior-ventral views of the braincase in *Tyrannosaurus rex* and *Tarbosaurus* (after Maleev, 1974). Note the great breadth between the basitubera.

Nanotyrrannus the upward orientation of the orbits is less marked, and, because the muzzle is extremely constricted, a wide arc of overlap is possible in a plane almost parallel to the muzzle.

The paroccipital processes of *Nanotyrannus* agree with those of *Tyrannosaurus* in being oriented nearly directly outwards and having much less of a backward orientation than that seen in gorgosaurs and daspletosaurs (Fig. 3, 11). The area

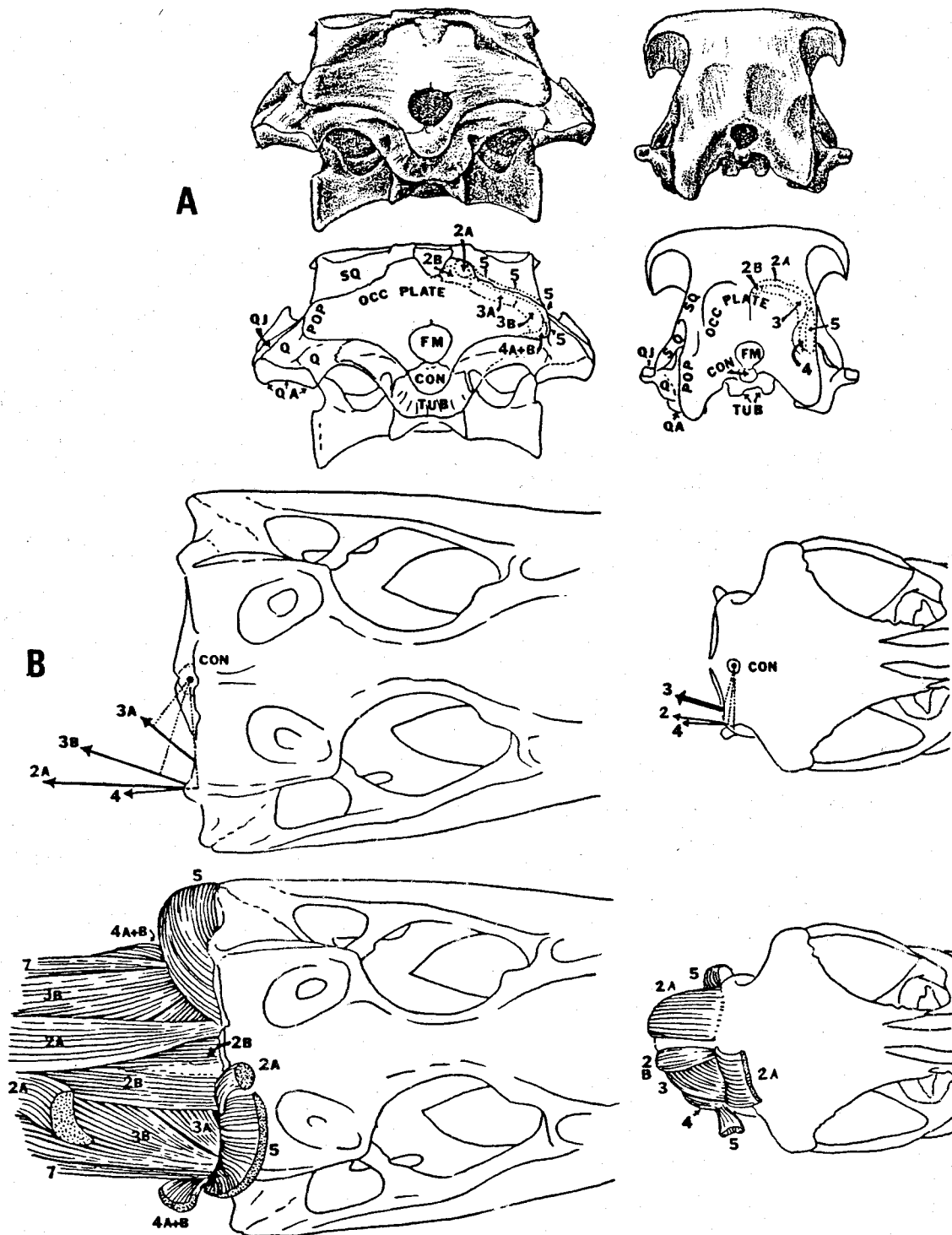


Figure 10—How the head-neck flexure affects the evolution of basitubera. Comparison of the head-neck muscles in: **RIGHT** — an ostrich, and **LEFT** — an alligator. **A** — occipital views of the skull showing muscle attachments. **B** — partial dissection of the head-neck muscles in lateral and dorsal views. **C** — sagittal section and lateral outline showing the leverage available; line of action of the outermost fibers of each muscle shown as an arrow; leverage shown as a dotted line passing from line of action to the center of rotation in the occipital condyle.

Abbreviations: CON occipital condyle. FM foramen magnum. lv leverage. POP paroccipital process. OCC PLATE occipital plate. Q quadrate. QA articular surface for articular. QJ quadratojugal. RET retroarticular process. SQ squamosal. TUB basitubera. TYM tympanum.

Muscles: 2A spinalis capitis. 2B rectus capitis. 3A obliquus capitis. 3B longissimus capitis. 4 transversarius part of longissimus capitis (two divisions in *Alligator*). 5 depressor mandibulae. 6 longus colli and iliocostalis cervicis-capitis. 7 iliocostalis cervicis.

for muscle attachment in the superior temporal fossa is very great in *Nanotyrannus*, as it is in all tyrannosaurids. The right and left temporal muscle scars meet along a very high and very narrow sagittal crest that extends from parietal at the front face of the occipital plate forward along the frontal midline to a point opposite the dorsal midmargin of the orbit. The anterior extension of the temporal scars in tyrannosaurs is developed as an excavated area on the dorsal surface of the frontals and is far larger than that of the large theropods of the Early Cretaceous or Jurassic, such as *Acrocanthosaurus*, allosaurs and ceratosaurs. The sutures between the lachrymal and prefrontal have thoroughly coossified in *Nanotyrannus*, as have the sutures between frontals and prefrontals, and between parietals and frontals. This degree of coossification is not matched in any gorgosaur or daspletosaur specimen and is even greater than in the *Tyrannosaurus* skull AMNH 5027. Without question, the type of *Nanotyrannus* was fully adult and had reached the maximum size the individual would have attained if it had lived longer.

The lower temporal fenestra in the type of *Nanotyrannus*

has suffered slight fore-to-aft compression but appears to have been like that of tyrannosaurs in general, with a long, horizontal quadratojugal-squamosal contact that gives a Σ -shape to the posterior outline of the fenestra. Although the *Nanotyrannus* type is clearly fully adult, there is no coossification at the squamosal-quadratojugal contact, and, since the quadrate has a moveable joint with the squamosal, the entire quadrate-quadratojugal complex can swing outwards to increase the size of the gullet. The quadratojugal in *Nanotyrannus* has a long anterior prong that overlaps onto a deep notch in the posterior process of the jugal. Postmortem compression has pressed the quadratojugal prong forward so that it extends beyond its jugal articular surface. There is no coossification at this joint, suggesting that some twisting and flexing could occur in life, a situation that also occurs in *Tyrannosaurus* and the other genera of the family.

The postorbital bar in *Nanotyrannus*, as in tyrannosaurids generally, is very wide front-to-back with a firm, interdigitating suture between postorbital and jugal (Fig. 3, 11). The postorbital bar in all Jurassic theropods is much narrower. The

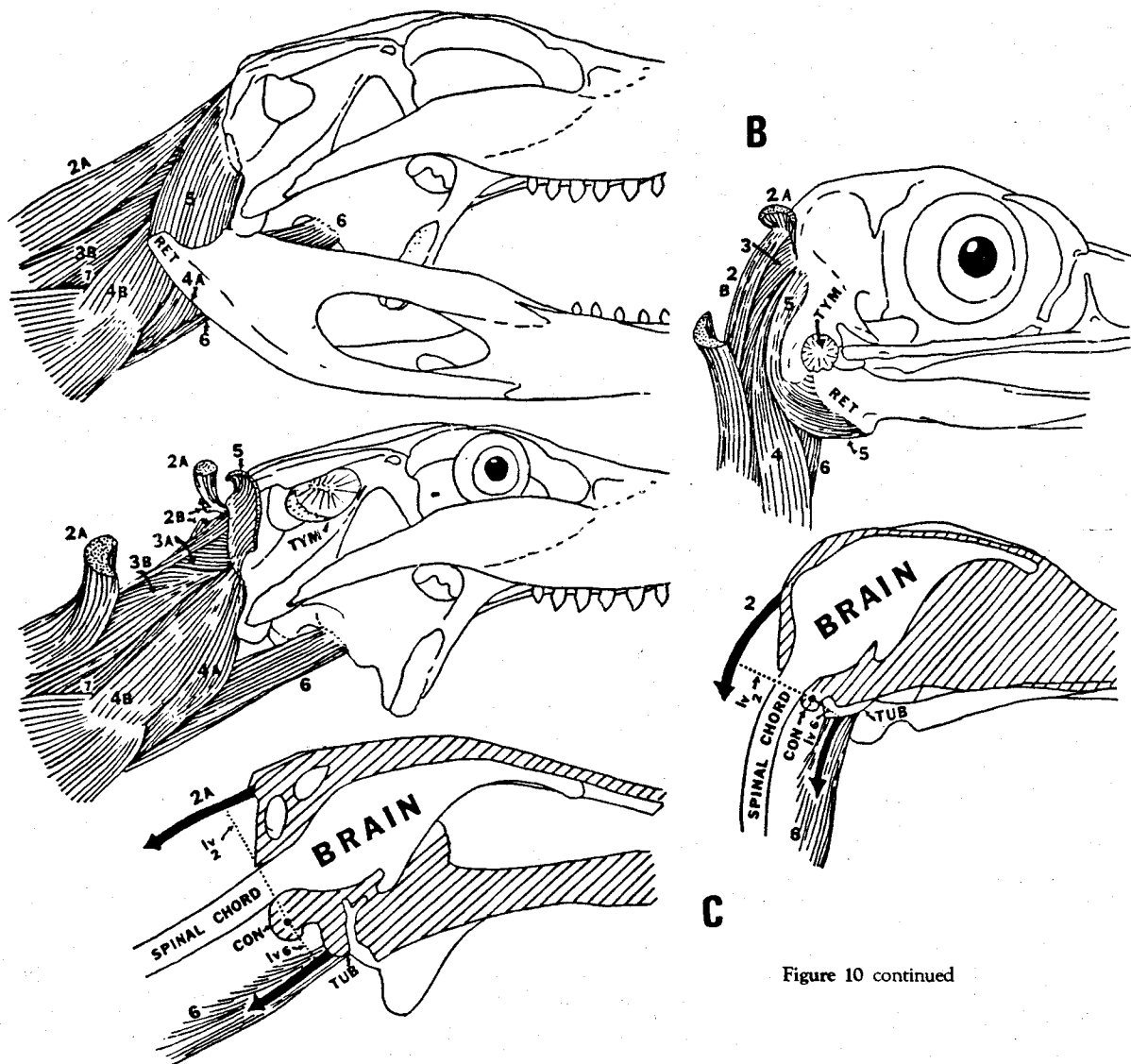
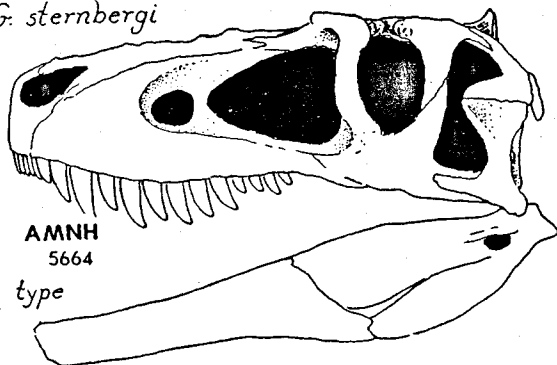


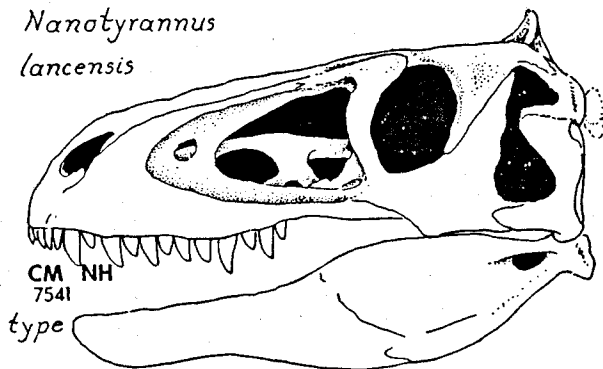
Figure 10 continued

NANOTYRANNUS NEW TYRANNOSAUR

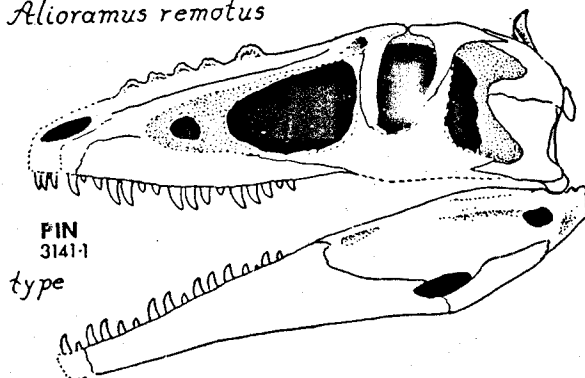
Gorgosaurus
G. sternbergi



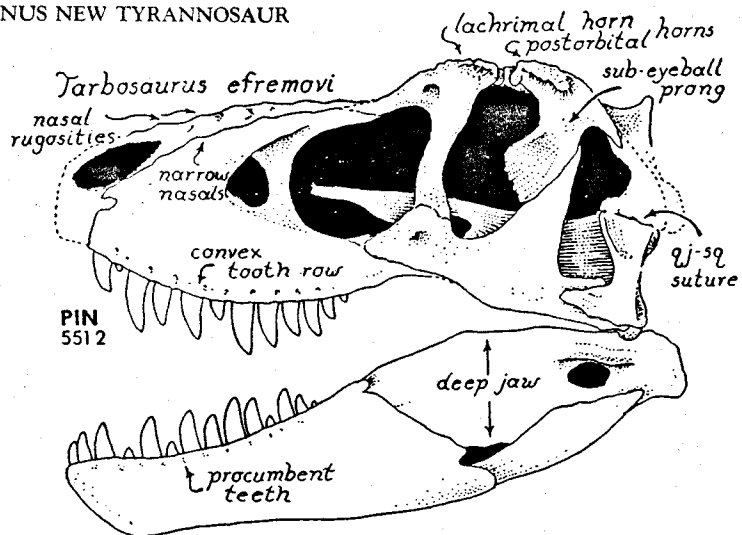
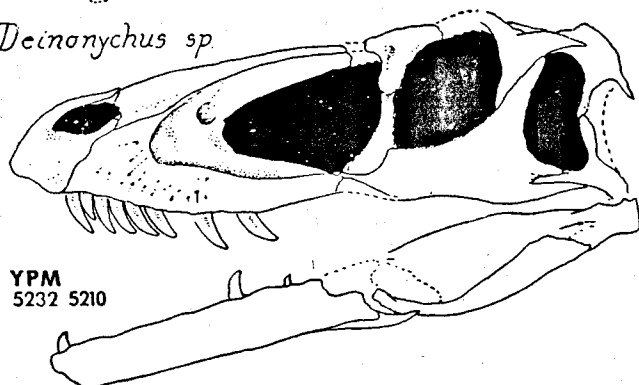
Nanotyrannus
lancensis



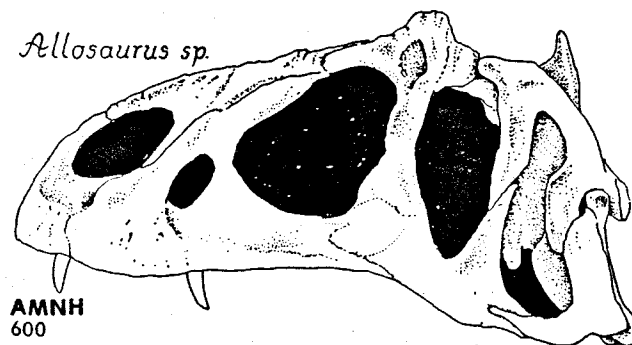
Alioramus remotus



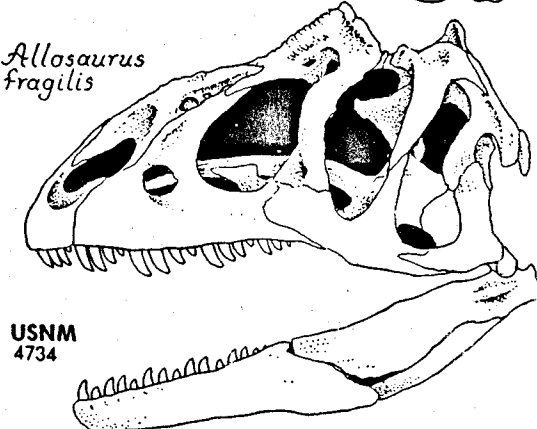
Deinonychus sp



Allosaurus sp.



Allosaurus
fragilis



Ceratopsaurus nasicornis

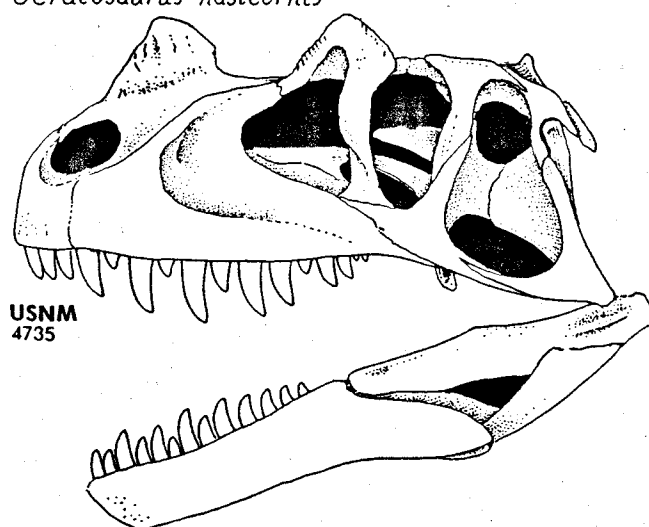
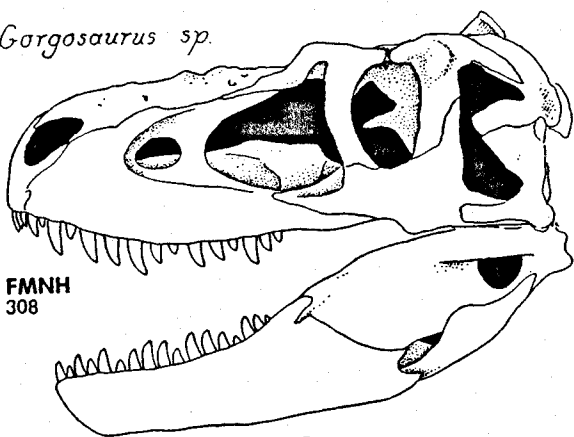


Figure 11

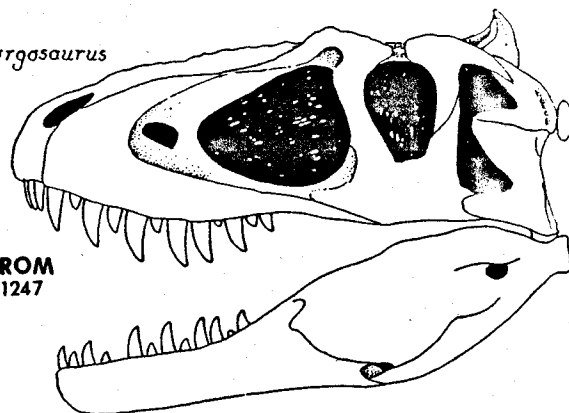
Gorgosaurus sp.

FMNH
308



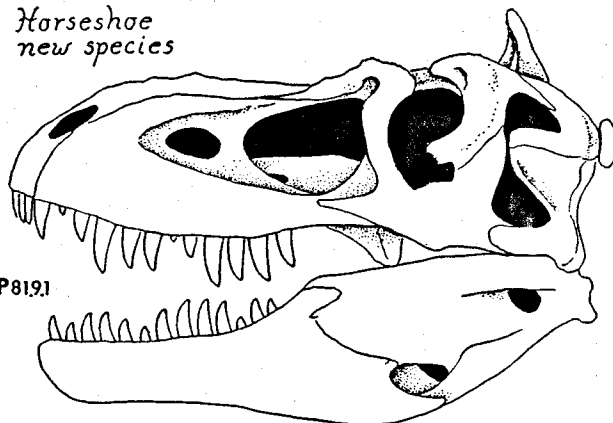
Gorgosaurus
sp

ROM
1247

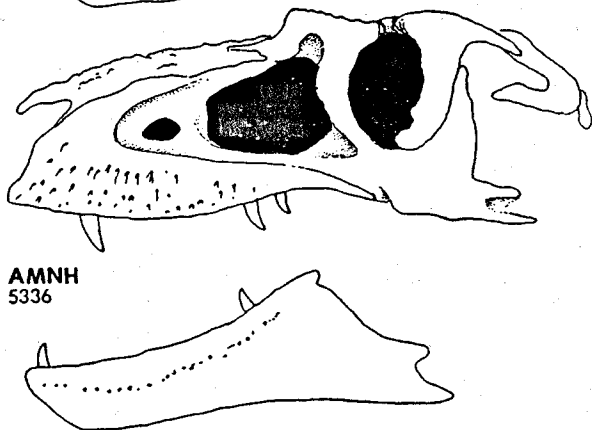


Horseshoe
new species

P8191

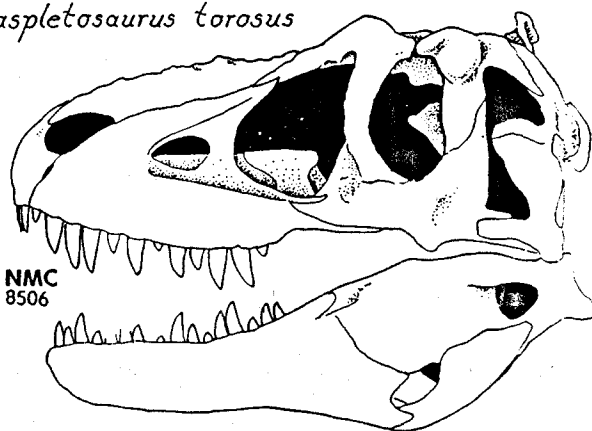


AMNH
5336

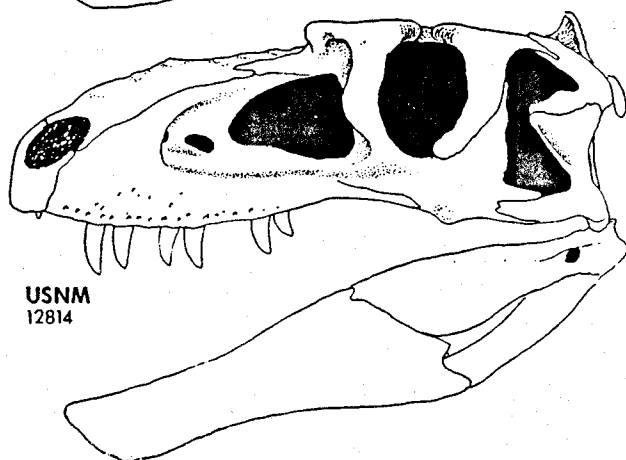


Daspletosaurus torosus

NMC
8506

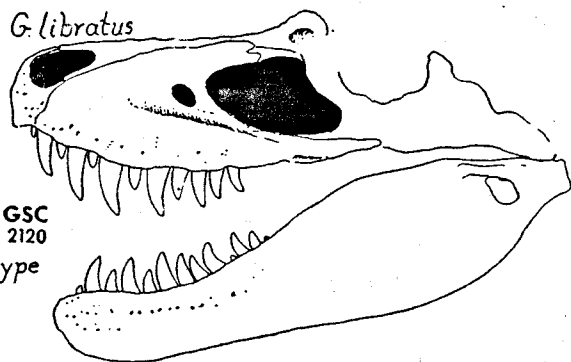


USNM
12814



G. libratus

GSC
2120
type



Tyrannosaurus rex

AMNH
5027

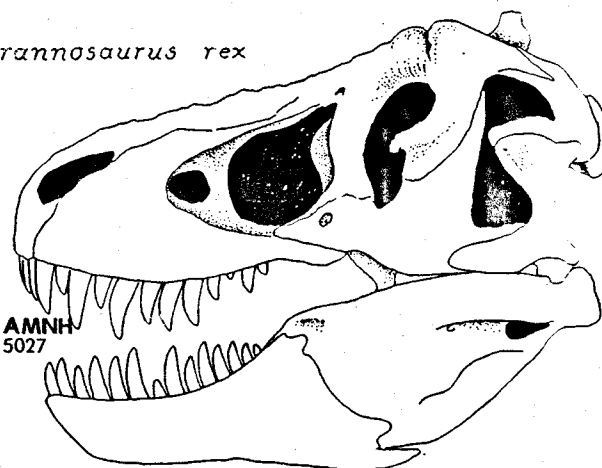
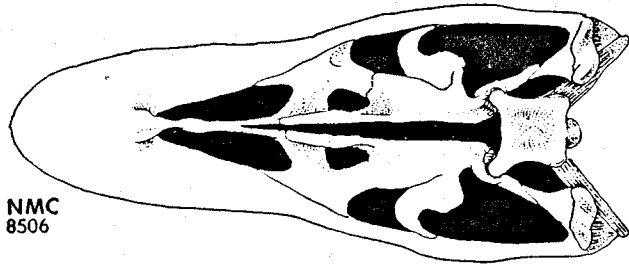


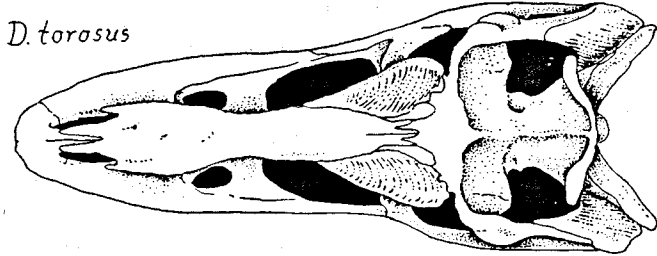
Figure 11—Lateral, dorsal and ventral outlines of the skulls of tyrannosaurs drawn to the same length, except for *Alioramus*. Deinonychid drawn to the same height as *Alioramus*, *Nanotyrannus* and *G. sternbergi*. Jurassic theropods — allosaurs and ceratosaur — drawn to a constant skull depth. The tarbosaur skull is labeled to show the features that distinguish an advanced tyrannosaur from a Jurassic allosaur.

postorbital bar in the troodont *Saurornithoides* is very similar to that of *Nanotyrannus* and the other tyrannosaurids (Barsbold, 1974). Although the tyrannosaurid post-

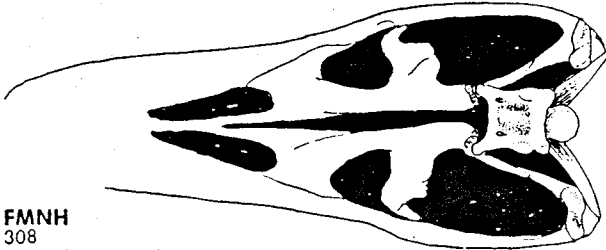
orbital bar is very wide, it is thin side-to-side and some bowing or twisting is possible about the postorbital-jugal sutural zone. Below the posterior-ventral edge of the orbit in



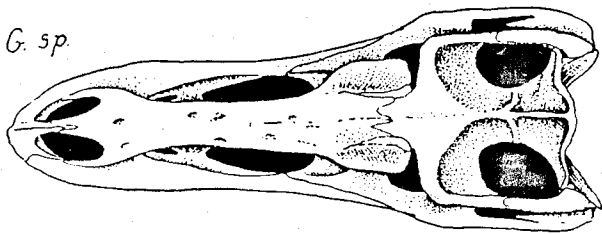
NMC
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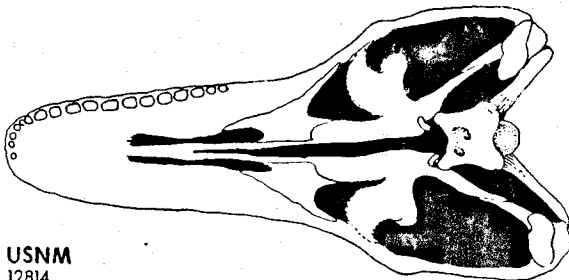
D. torosus



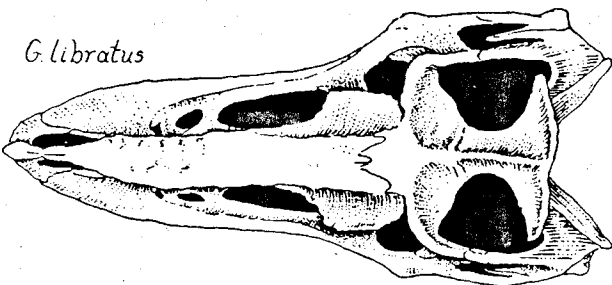
FMNH
308



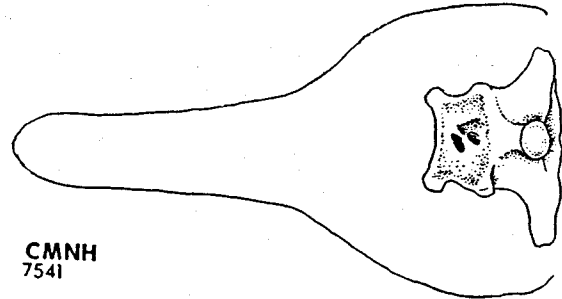
G. sp.



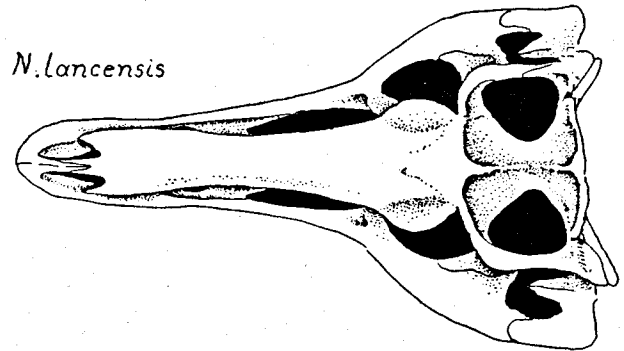
USNM
12814



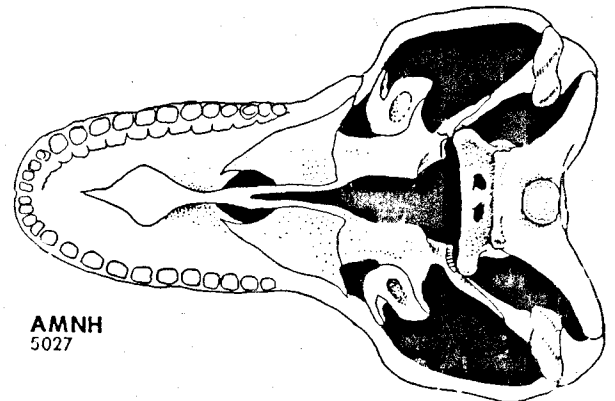
G. libratus



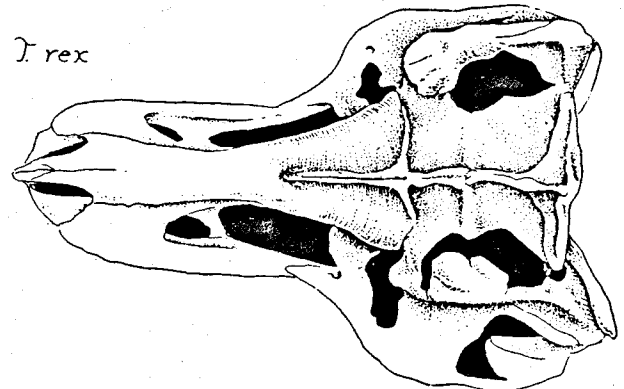
CMNH
7541



N. lancensis



AMNH
5027



T. rex

Nanotyrannus, the jugal bears a slight swelling with a lightly striated surface, indicating the presence in life of a low dermal horn. A horn in this location is present in allosaurs and most Cretaceous theropods of large size but appears to be absent in ceratosaurs and primitive theropods of the Late Triassic and Early Jurassic, such as *Coelophysis* and *Syntarsus*.

Orbit and Muzzle. The lacrimal in *Nanotyrannus* was restored in plaster with a pre-orbital horn of a most unusual configuration (Plate 1-3) — in side view, the horn has an evenly rounded, convex-up profile. When the plaster was removed, the dorsal surface of the lacrimal was seen to lack any clear-cut horn-like growth, although further preparation is necessary to clean down to the bone. At present, it appears that the dorsal preorbital bar is neither greatly swollen nor highly rugose. Ceratosaurs and allosaurs have erect horns with strong rugosities (Fig. 11). All gorgosaurs and daspletosaurs have lacrimal horns with upper edges that are swollen side-to-side and bear coarse grooves and ridges. All the best preserved gorgosaur skulls — e.g. USNM 12814, AMNH 5336, 5664,

FMNH 308 — have horns with apices that are displaced forward. Daspletosaurs and the new, unnamed tyrannosaur from the Horseshoe Canyon have horn apices that rise directly upwards. *Tyrannosaurus* and *Tarbosaurus* have no horn apices whatever and, instead, have dorsal-lateral bars of the lacrimal that are greatly swollen. In *Tarbosaurus* and *Tyrannosaurus* the postorbital part of the orbital rim is also greatly swollen, and, together the postorbital and lacrimal make a low, broad reinforced forehead area adapted for delivering head-butting blows. The apparently weak development of a rugose lacrimal bar or lacrimal horn in *Nanotyrannus* is a condition that cannot be derived easily from the lacrimal device seen in any other tyrannosaur. As we argue below, the ancestor of tyrannosaurs probably was an animal close in grade to a deinonychiid and probably possessed a lacrimal with no horny development other than a slight thickening along the dorsal-lateral edge. Therefore the lacrimal condition in *Nanotyrannus* may well be the most primitive known for the Family Tyrannosauridae.

The ventral-anterior angle of the dorsal bar of the lacrimal in *Nanotyrannus* bears a depressed, smooth surface that

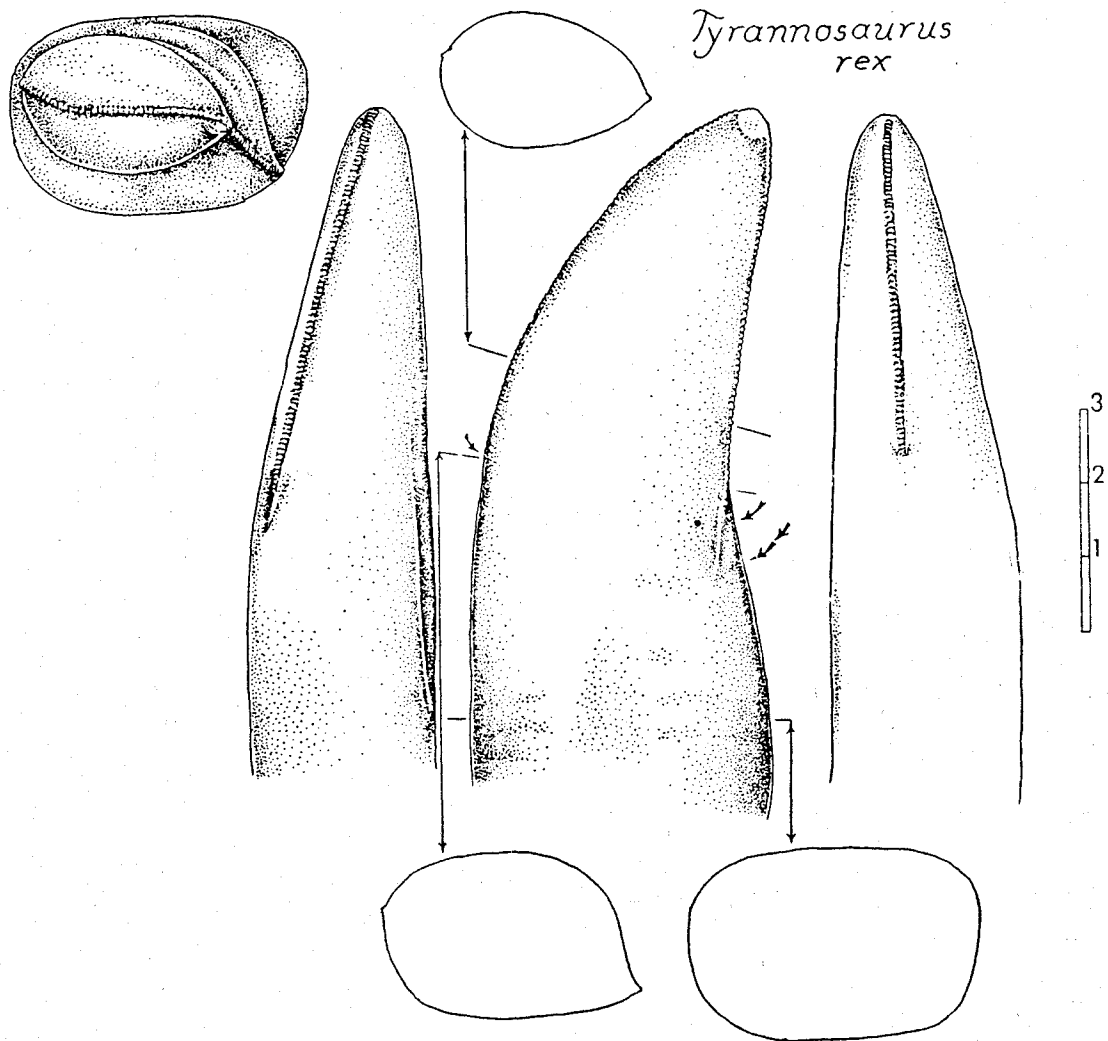
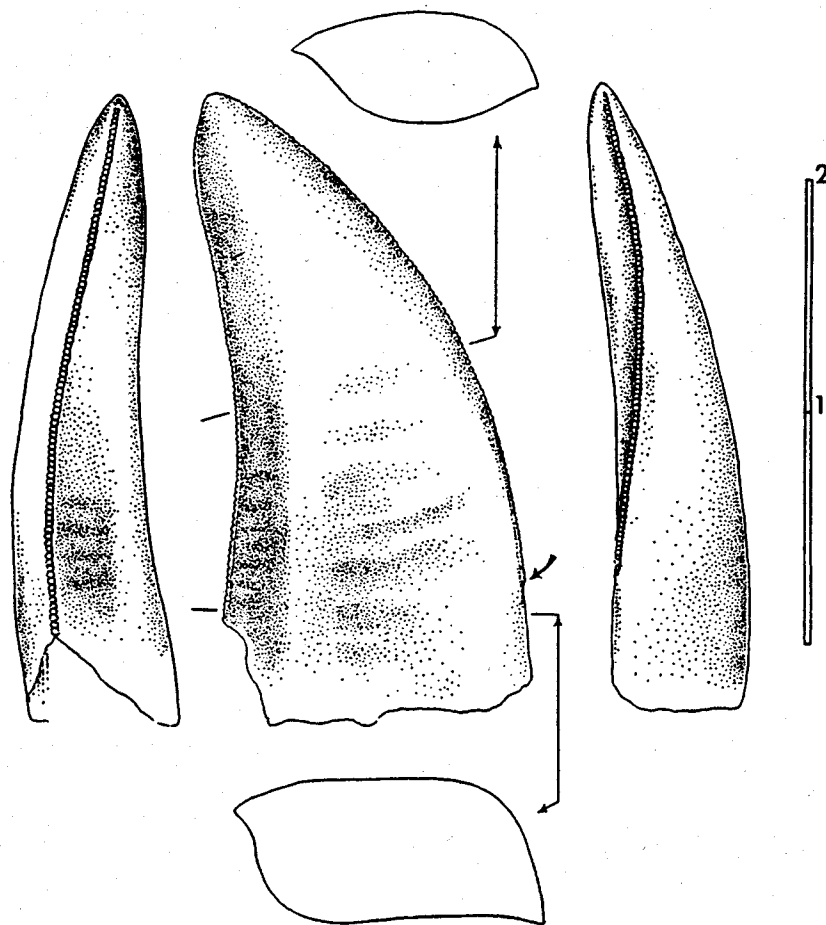


Figure 12—Tyrannosaurid teeth. Left seventh dentary tooth of *Tyrannosaurus* (AMNH 5027). Left-to-right: crown view, posterior view, outer (buccal) view, anterior view. Cross sections given as indicated by arrows. Single heavy arrow shows termination of serrated keel. Double heavy arrow shows termination of unserrated keel.



Nanotyrannus

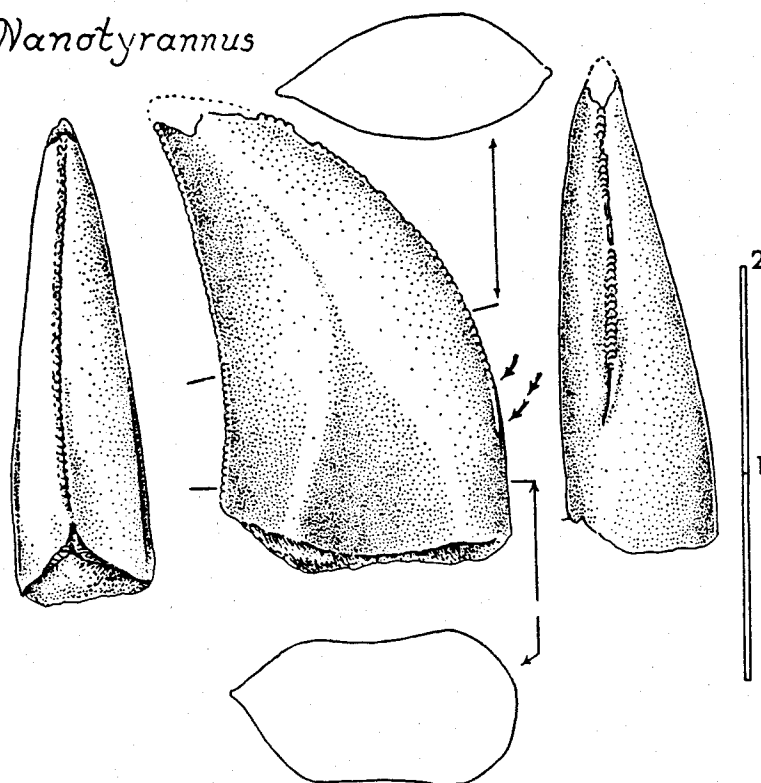


Figure 12 continued—Two isolated teeth ascribed to *Nanotyrannus*, uncatalogued, Corson Co., South Dakota, Denver Museum of Natural History. Left-to-right: posterior view, inner (lingual) view, anterior view. Symbols as in previous figure.

forms the posterior-dorsal corner of the antorbital fenestra (Fig. 3). The outer anterior edge of the vertical limb of the lacrimal bears a similar depressed surface that is continuous with the smooth, depressed zone that borders the ventral edge of the antorbital fenestra on the maxilla. All these depressed borders probably served as attachment areas for the anterior pterygoideus muscle bundles that bulged outwards through the fenestra. Nearly all theropods have a similar pattern. In *Tyrannosaurus* the entire preorbital part of the lacrimal is greatly swollen and the smooth depressed area has disappeared. In most theropods of a ceratosaur grade or higher, there is a pneumatic canal opening near the top of the posterior-dorsal depressed area on the lacrimal. *Nanotyrannus* shows this condition. In *Tyrannosaurus rex* this foramen has become enclosed by swollen lacrimal bone on all sides, a condition also developed in *Tarbosaurus* but to a lesser extreme.

The nasals in *Nanotyrannus* are very narrow, as in all other Late Cretaceous theropods and in contrast with Jurassic taxa. All tyrannosaurs have snouts with some sort of rugosities on the dorsal surface of the nasals, but *Nanotyrannus* displays the weakest development of this character. *Nanotyrannus* shows only a set of shallow striae and ridges running fore-to-aft along the dorsal surface of the nasals (Fig. 3, Plate 1-3). Gorgosaurs, daspletosaurs, tarbosaurus and *Tyrannosaurus* show a much heavier ridge-and-furrow ornamentation that in life must have supported a very rough

horny skin topography. The long-snouted Mongolian tyrannosaur *Alioramus* displays a grotesque set of oval hornlets on the nasal (Fig. 11).

Nanotyrannus agrees with other tyrannosaurs in having a large external foramen, opening upwards, in the jugal.

The antorbital fenestrae of *Nanotyrannus* are very large but very close in pattern to those of the gracile gorgosaurs (Fig. 11). As in all tyrannosaurids, there is a small anterior, accessory fenestra, a character that links tyrannosaurs with dromaeosaurs, troodonts and allosaurs. The snout, in lateral view, is longer and lower in *Nanotyrannus* than in any other tyrannosaurid except *Alioramus* and approaches the condition seen in dromaeosaurs.

Lower Jaw. In the type of *Nanotyrannus* the upper edge of the mandible and the lower teeth are hidden by the skull and upper teeth. On the outer surface of the mandible there is a clear line of separation between the dentary in front and the surangular + angular behind (Plate 1). This line of separation shows no evidence of suturing or firm connection whatever, and undoubtedly there was a loosely articulating intramandibular joint as is found in all other tyrannosaurids. The intramandibular joint in tyrannosaurids is constructed very much like that of troodonts, dromaeosaurs, allosaurs and ceratosaurs (Fig. 13): the dentary and splenial enclose a spacious intramandibular cavity, open to the rear; the angular sends a forwardly directed prong into the cavity; the angular prong

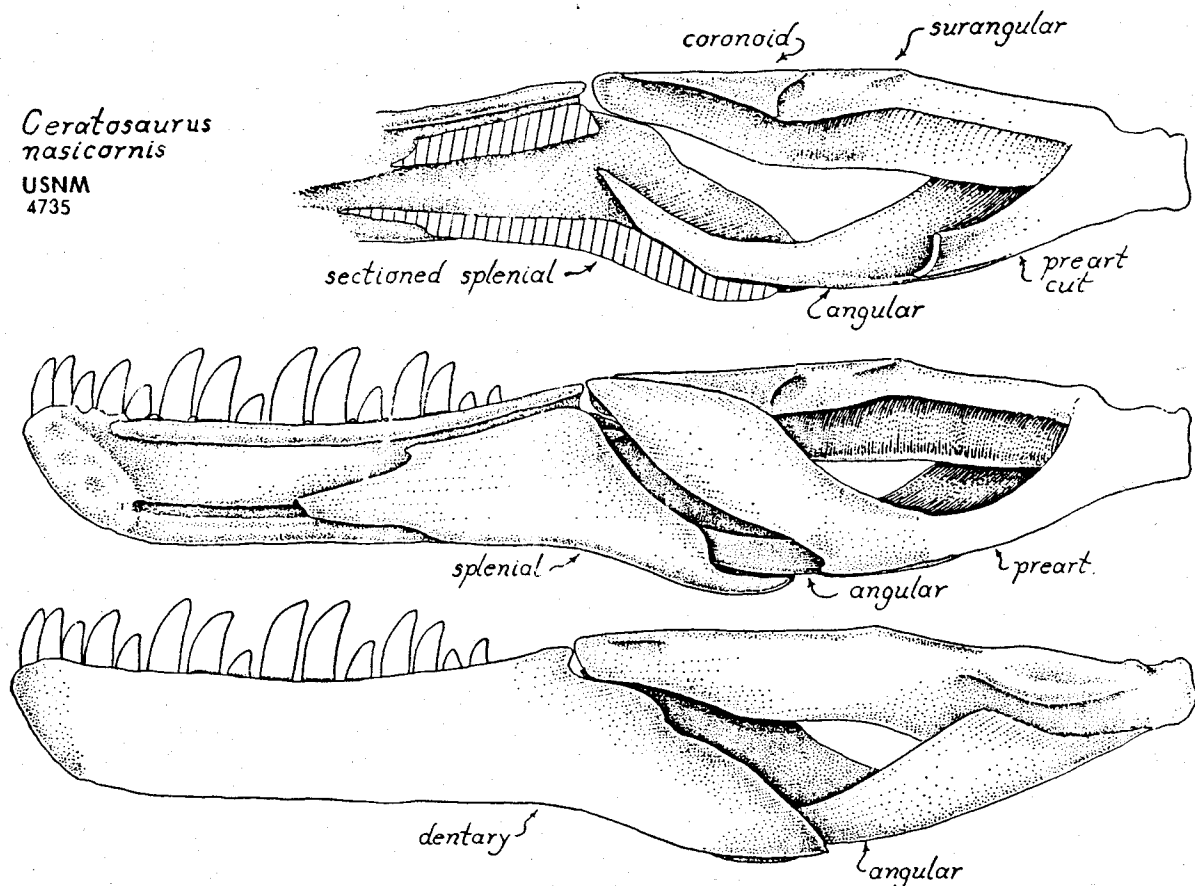


Figure 13—Construction of the intramandibular joint in a primitive neotheropod, *Ceratosaurus*.

has no sutural connection to the inner walls of the intramandibular cavity and can twist and flex in all directions; the anterior edges of the surangular + prearticular have no firm suturing to the posterior edges of the splenial + dentary, so the intramandibular joint can permit a wide range of movement that could expand the gullet or adjust the pitch and position of the mandibular teeth relative to the upper teeth. The podokesaurid theropods of the Triassic and Early Jurassic lack an intramandibular joint and have, instead, a firm suturing of the posterior and anterior mandibular units. The Middle Jurassic megalosaurids probably lacked the joint also, but no complete megalosaurid lower jaw has been recovered. (We restrict the term "megalosaurid" to *Megalosaurus bucklandi*, *Poikilopleuron bucklandi* and closely allied species, as argued by Waldman, 1974).

Teeth. *Nanotyrannus* differs from all other tyrannosaurids, except possibly *Alioramus*, in retaining the primitive theropod condition of maxillary teeth that are compressed strongly from side-to-side, very wide front-to-back relative to the crown height, and very sharp (Fig. 12). None of the teeth in the type of *Nanotyrannus* can be seen from the inner surface, but the three loose teeth in the Denver Museum collection seem to match quite closely with those in the middle and anterior sector of the maxilla of the type (Fig. 12). These loose teeth supply most of the information on tooth shape for the genus. In the loose tooth that is from the mid-maxillary row, the keel along the anterior edge reaches almost as far towards the crown base as does the posterior keel. The entire posterior keel is serrated but the basal part of the anterior keel is free of serrations. The anterior keel moves toward the inside of the tooth as the keel passes to the base; the posterior keel moves towards the outside. In the loose tooth that is from the front of the maxillary row, the crown is concave on the inner surface and the anterior keel is displaced further towards the inner edge. The only other Late Cretaceous theropod with maxillary teeth of a form similar to that of *Nanotyrannus* is the "Jordan theropod" described by Molnar (1978), an animal that may be a large dromaeosaur.

Gorgosaurs, daspletosaurs, *Albertosaurus*, *Tarbosaurus* and *Tyrannosaurus* all have maxillary teeth with cross sections that are much thicker side-to-side relative to their length fore-to-aft in the upper part of the crown, and the crowns are much taller relative to the anterior-posterior length (Fig. 12). The posterior maxillary teeth of *Tyrannosaurus rex* are very blunt and have an overall shape like that of posterior teeth in *Gorgosaurus* and *Daspletosaurus*; the crown cross-section is thick and nearly symmetrical. Clearly the function of these small teeth was different from that of the tall, strong anterior maxillary teeth — perhaps the posterior teeth were used to crack bones. In most of the other theropods — allosaurs, ceratosaurs, megalosaurs — the posterior maxillary teeth are shorter and blunter in the crown than are the anterior maxillary teeth, although the degree of heterodonty is not so marked as in *T. rex*. In *Nanotyrannus* the posterior maxillary teeth are much sharper and more compressed side-to-side than those in the same loci in the other tyrannosaurids.

The sharp, delicate, blade-like teeth of *Nanotyrannus* indicate that the bite delivered a long, narrow wound; surely this genus could not produce the bone-crushing bite of a *Tarbosaurus* or *Tyrannosaurus*. The tooth count in *Nanotyrannus* is 15 maxillary and 4 premaxillary, as in *Gorgosaurus sternbergi*; all other gorgosaurs have 13 maxillary teeth except for FMNH 308 which has 15 maxillary teeth. The high count in the latter specimen strongly suggests that it represents a species distinct from that of USNM 12814 and AMNH 5336.

The four premaxillary teeth in *Nanotyrannus* have outer (buccal) crown faces that are very narrow and crowded close together. The crown cross section must be very deep buccally and probably had a flat inner (lingual) face as in other tyrannosaurids. The inner edges of the teeth cannot be seen and the presence or absence of serrations cannot be determined. The first maxillary tooth has a form like that of the premaxillary teeth. All other gorgosaur species have a similar arrangement of incisiform teeth. *Tarbosaurs* and *Tyrannosaurus* have a large, massive first maxillary tooth. Incisiform premaxillary teeth with asymmetrical crowns are also present in ceratosaurs, allosaurids, dromaeosaurids (especially in *Dromaeosaurus* itself) and troodontids but are not developed in the primitive podokesaurs or megalosaurs. In all the theropods equipped with incisiform teeth, the right and left upper toothrows converge together to make a U-shaped symphysis as seen in dorsal or ventral view. Thus the incisor row is arrayed along a broad anterior arcade useful for cropping meat from bones or grooming skin and pelage. The podokesaurs and megalosaurs have V-shaped symphyses where the right and left tooth rows are nearly linear (Waldman, 1974). Incisiform teeth in a U-shaped arcade and an intramandibular joint define the Neotheropoda (Bakker, 1986). It is only among the tyrannosaurids that the premaxillary teeth become exceptionally crowded and very narrow across the buccal face.

The maxillary tooth row is nearly straight in *Nanotyrannus* and the teeth are set in the maxilla at right angles to the row, probably the primitive condition for the family. Some gorgosaurs and all daspletosaurs, albertosaurs, *Tarbosaurs* and *Tyrannosaurus* have tooth rows that are more strongly convex-down, as seen in lateral view, and the anterior maxillary teeth tend to be procumbent (Fig. 11). Procumbent teeth suggest that the bite was delivered as the head and neck lunged forward at the parts of the prey's body that were gently curved — such as the flank or belly. Procumbent teeth used in this way could bite a chunk of bone and flesh off the rib cage or abdominal wall. The sharp but delicate *Nanotyrannus* teeth, set vertically in the jaw, would be better designed for biting objects with less gently curved surfaces.

ANCESTRY AND EVOLUTION OF NANOTYRANNUS

In order to determine the primitive state of characters within the tyrannosaurids, we must know what groups are the sister clades of the family (Fig. 2). Dromaeosaurs differ from Jurassic dinosaurs and agree with tyrannosaurs, ornithomimids,

troodontids and acrocanthosaurs and possible oviraptorids in possessing the derived characters of a tall, flat, wide ascending process of the astragalus and a dorsal column that is shortened relative to the pelvis. But dromaeosaurs are more primitive than the other groups in lacking extensive pneumatization of the otic region, and in retaining a primitive paroccipital process that is shallow top-to-bottom at its root. Therefore we take dromaeosaurs as the primitive sister group of the clade tyrannosaurs + troodonts + ornithomimids + oviraptorids + acrocanthosaurs. The four Late Cretaceous families — tyrannosaurs, oviraptorids, troodonts and ornithomimids — all share the derived character of a metatarsus that is more elongated and more compact from side-to-side than in acrocanthosaurs and thus probably comprise a clade with an origin in the early Late Cretaceous. Currie (1986) has shown that ornithomimids and troodonts share unusual basicranial characters and probably are more closely related to each other than either is to tyrannosaurs. From the foregoing analysis, it follows that for tyrannosaurs the most primitive sister group among Cretaceous theropods is the dromaeosaurs. Although troodonts and ornithomimids are closer cladistically to tyrannosaurids than are dromaeosaurs, in jaws and dentition and snout the dromaeosaurs probably are better guides for reconstructing the primitive tyrannosaurid condition. Ornithomimids — ostrich dinosaurs — have muzzle and jaws very highly modified as toothless beaks and hence offer little clue to the primitive condition in tyrannosaurs. Troodonts are highly modified in a different direction — the teeth have relatively huge serrations on the trailing edge, unlike the more normal theropod serration pattern seen in tyrannosaurs. Dromaeosaurs show a basic, primitive theropod pattern of compressed, recurved teeth with fine serrations.

The teeth of *Nanotyrannus* are very similar to those of dromaeosaurs in degree of lateral compression. All other tyrannosaurs, with the possible exception of *Alioramus*, have maxillary teeth that are thicker side-to-side, and therefore, on dental evidence alone, *Nanotyrannus* could be argued to be the most primitive North American tyrannosaur.

The morphology of the *Nanotyrannus* skull displays several exceptionally advanced features, all or most of which are related to the development of stereoscopy and a sharper head-neck flexure. Indeed, these two functional shifts may be reflections of a single behavioral reorganization — by holding the muzzle flexed downwards, the reorientation of the eyes achieves the maximum overlap of visual fields. Great side-to-side expansion of the basicranium and temporal regions also increases the size of the jaw-closing muscles and the leverage of the muscles flexing the neck laterally. In *Tyrannosaurus* the expansion of the temporal region occurs without constriction of the muzzle but in *Nanotyrannus* the muzzle has a very narrow width relative to the skull length. Theropod muzzles housed the anterior part of the pterygoideus D muscle, a very important jaw-closing unit, and the temporal expansion in *Nanotyrannus* could be, in part, a compensation for the decrease in muscle volume incurred in the snout.

The type of *Nanotyrannus* is the smallest adult tyrannosaurid skull known, only 572 mm long, fully 13% shorter

than the types of *Alioramus* and *Gorgosaurus sternbergi* (Table 1). And the degree of coossification in the dorsal temporal region in *Nanotyrannus* confirms that the type was fully adult, while, on the other hand, the types of *G. sternbergi* and *Alioramus remotus* may well be subadult in size. Could *Nanotyrannus* evolve by a process of dwarfing from a more typical tyrannosaur of large size? Since *Nanotyrannus* is from the very last faunal age of the Cretaceous, and since all earlier tyrannosaurid skulls that are definitely adult are 40% larger, it is justifiable on stratophenetic grounds to deduce that the small adult size is a late-evolving derived character. On the other hand, as we argue elsewhere (Bakker, Currie, and Williams, in press), the most closely related sister group of tyrannosaurids are the oviraptorid, troodont and ornithomimid theropods, animals of small size, so a cladistic argument could be generated to defend the notion that small size is a primitive character for the Family Tyrannosauridae.

The apparently weak lachrymal horn of *Nanotyrannus* cannot be derived in any straightforward way from the low, grossly thickened lachrymal of *Tyrannosaurus* and *Albertosaurus*, or the more erect, pointed horn of *Gorgosaurus* and *Daspletosaurus*. Neoteny is a possible mechanism for producing a delicate adult horn from an ancestral state that was more swollen and more coarsely sculpted. However, the type of *Gorgosaurus sternbergi*, although probably far from adult size, already has a swollen and rough-textured lachrymal horn. Lachrymal horns of various shapes have evolved many times independently among large theropods (but, strangely enough, never among theropods with skulls less than 300 mm long) and the primitive state for the Family Tyrannosauridae is probably a lachrymal like that of dromaeosaurs — without any erect horn and only slight thickening along the dorsal-lateral border (Bakker, Currie, and Williams, in press). At present we favor the hypothesis that the lachrymal condition of *Nanotyrannus* evolved from a hornless and unswollen ancestral state.

Whatever its evolutionary origins, *Nanotyrannus* seems to have evolved its stereoscopy independently of *Tyrannosaurus*. The low, long snout and primitive teeth of *Nanotyrannus* seem to preclude an ancestry from an animal as advanced as *Gorgosaurus*. The two stereoscopic genera, *Nanotyrannus* and *Tyrannosaurus*, are the only well known large predators from the Lancian Age, and it is noteworthy that both have achieved the highest degree of potential stereoscopy known among large theropods. Small theropods of the Cretaceous also show strong tendencies for frontal reorientation of the orbits — the orbits had moderate forward orientation in the Early Cretaceous dromaeosaurs and very strong forward orientation in ornithomimids and troodonts of the Late Cretaceous. Clearly natural selection was favoring greater precision of depth perception. A very similar trend towards greater stereoscopy can be seen among mammalian predators of the Cenozoic. The earliest predators of the Paleocene and Eocene — didymictids, miacids, hyaenodontids, oxyaenids — all had relatively small orbits that faced mostly outwards. But these archaic groups were replaced eventually by more modern cats, nimravids, dogs, bear-dogs,

and hyaenas, all of which had relatively larger orbits with a greater degree of forward orientation. Thus *Nanotyrannus* seems to have been shaped by an inexorable law of the coevolution between predator and prey, a law that operated to produce ever greater finesse in the choreography of attack behavior.

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CITED REFERENCES

- Bakker, R.T. 1986. *Dinosaur Heresies*. New York: William Morrow. 432 pp.
- Bakker, R.T., Currie, P., and Williams, M. 1988. Revision of the tyrannosaurid dinosaurs, with the description of a new genus from the Horseshoe Canyon fauna of Alberta and a discussion of theropod phylogeny and theropod mass extinctions. *Hunteria*, in press.
- Barsbold, R. 1974. Saurornithoididae, a new family of small theropod dinosaurs from central Asia and North America. *Palaeontologia Polonica* 30:5-22.
- Currie, P. 1985. Cranial anatomy of *Stenonychosaurus inequalis* (Saurischia, Theropoda) and its bearing on the origin of birds. *Canadian Journal of Earth Science* 22:1643-1658.
- Gilmore, C.W. 1946. A new carnivorous dinosaur from the Lance Formation of Montana. *Smithsonian Miscellaneous Collections* 106:1-19.
- Kurzanov, S.M. 1976. A new Late Cretaceous carnosaur from Nogontuva, Mongolia. *Nauka, Trudi, vip. 3, Akademia Nauk USSR (Moskva)* (1976):93-104.
- Lambe, L. M. 1914. A new genus species of carnivorous dinosaur from the Belly River Formation of Alberta. *Ottawa Naturalist* 28:13-20.
- Matthew, W.D., and Brown, B. 1922. Preliminary notices of skeletons and skulls of Deinodontidae from the Cretaceous of Alberta. *American Museum of Natural History Bulletin* 89:1-10.
- Maleev, E.A. 1955. New carnivorous dinosaurs from the Upper Cretaceous of Mongolia. *Dokladi Akademii Nauk, USSR* 104:779-782.
- Maleev, E.A. 1974. Gigantic carnosaur of the Family Tyrannosauridae. *Nauka, Akademia Nauk, USSR* 1974:132-191.
- Molnar, R. 1978. A new theropod dinosaur from the Upper Cretaceous of Central Montana. *Journal of Paleontology* 52:73-82.
- Osborn, H.F. 1905. *Tyrannosaurus* and other Cretaceous carnivorous dinosaurs. *American Museum of Natural History Bulletin* 21:259-265.
- Parks, W.A. 1928. *Albertosaurus arctunguis*, a new species of theropodous dinosaur from the Edmonton Formation of Alberta. *University of Toronto Studies, Geology Series* 25:1-42.
- Rhodestvensky, A.K. 1965. Growth changes in Asian dinosaurs. *Paleontologicheskii Zhurnal* 1965:95-105.
- Russell, D.A. 1970. Tyrannosaurs from the Late Cretaceous of Western Canada. *Publications in Palaeontology of the National Museum of Canada* 1:1-34.
- Stovall, J.W., and Langston, W. 1950. *Acrocanthosaurus atokensis*, a new genus and species of Lower Cretaceous Theropoda from Oklahoma. *American Midland Naturalist* 43:696-728.
- Taylor, R. 1977. Exercise and environmental heat loads: different mechanisms for solving different problems. In: Robertshaw, D. (ed.). *Environmental Physiology, International Review of Physiology* (Baltimore: University Park Press) pp. 1-45.
- Waldman, M. 1974. Megalosaurids from the Bajocian (Middle Jurassic) of Dorset. *Palaeontology* 17:325-340.

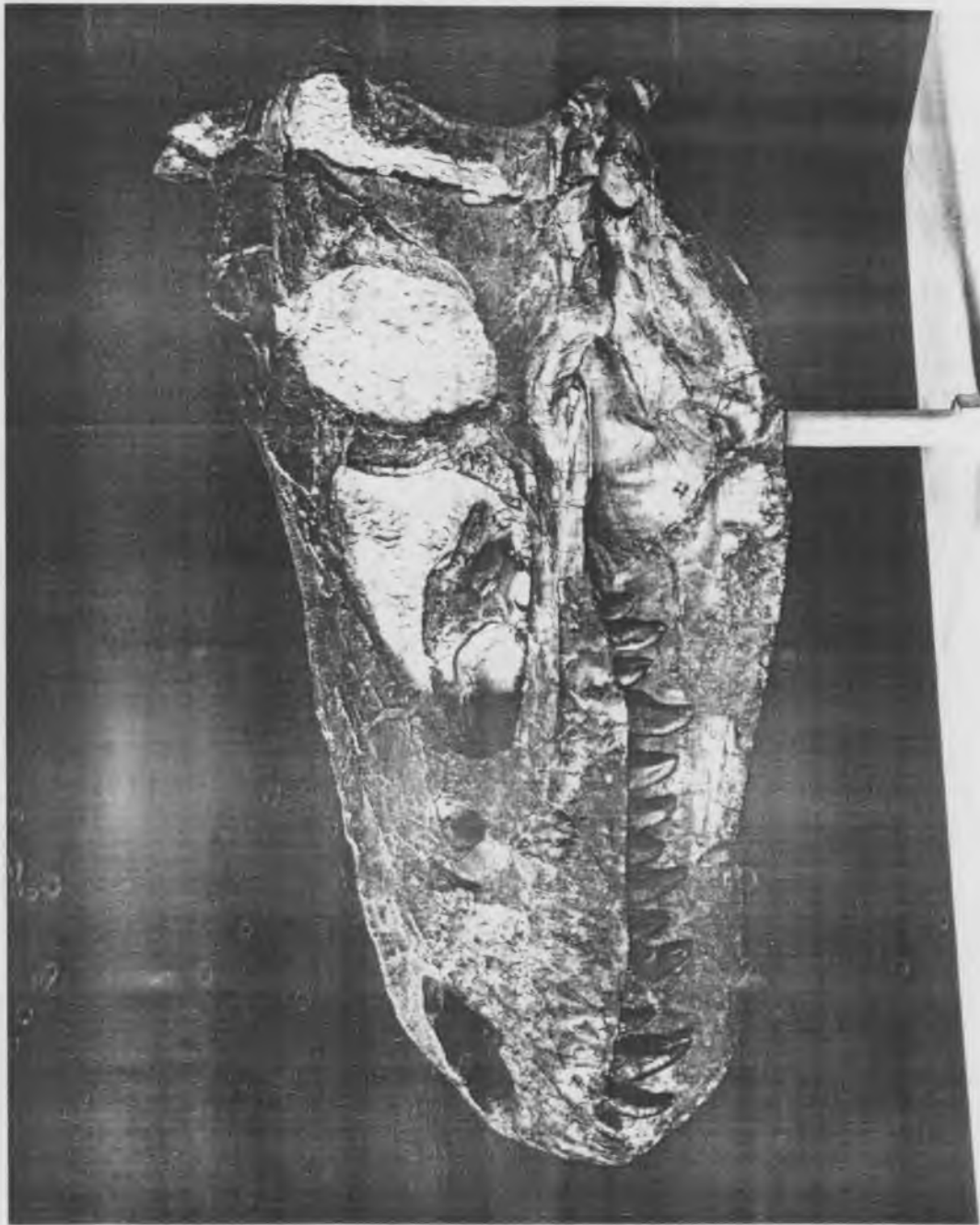


Plate 1. Lateral view of *Nanotyrannus lancensis*, type, CMNH 7541.



Plate 2. Dorsal view of *Nanotyrannus lancensis*, type, CMNH 7541.

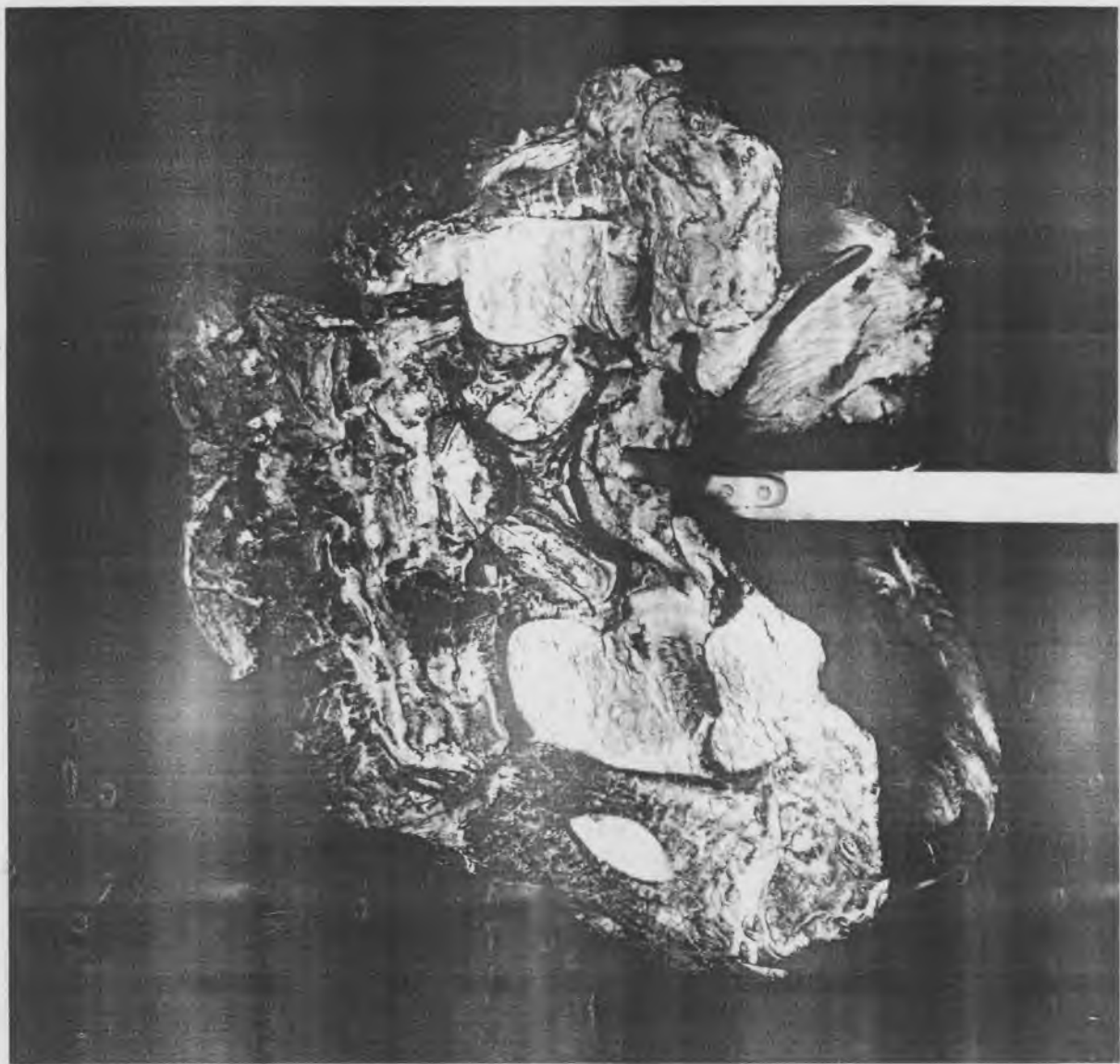


Plate 3. Occipital view of *Nanotyrannus lancensis*, type, CMNH 7541.



Plate 4. Close up of maxilla of *Nanotyrannus lancensis*, type, CMNH 7541.