# New Protocetid Whale from the Middle Eocene of Pakistan: Birth on Land, Precocial Development, and Sexual Dimorphism 

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#### Abstract

Background: Protocetidae are middle Eocene (49-37 Ma) archaeocete predators ancestral to later whales. They are found in marine sedimentary rocks, but retain four legs and were not yet fully aquatic. Protocetids have been interpreted as amphibious, feeding in the sea but returning to land to rest.

Methodology/Principal Findings: Two adult skeletons of a new 2.6 meter long protocetid, Maiacetus inuus, are described from the early middle Eocene Habib Rahi Formation of Pakistan. $M$. inuus differs from contemporary archaic whales in having a fused mandibular symphysis, distinctive astragalus bones in the ankle, and a less hind-limb dominated postcranial skeleton. One adult skeleton is female and bears the skull and partial skeleton of a single large near-term fetus. The fetal skeleton is positioned for head-first delivery, which typifies land mammals but not extant whales, evidence that birth took place on land. The fetal skeleton has permanent first molars well mineralized, which indicates precocial development at birth. Precocial development, with attendant size and mobility, were as critical for survival of a neonate at the land-sea interface in the Eocene as they are today. The second adult skeleton is the most complete known for a protocetid. The vertebral column, preserved in articulation, has 7 cervicals, 13 thoracics, 6 lumbars, 4 sacrals, and 21 caudals. All four limbs are preserved with hands and feet. This adult is $12 \%$ larger in linear dimensions than the female skeleton, on average, has canine teeth that are $20 \%$ larger, and is interpreted as male. Moderate sexual dimorphism indicates limited male-male competition during breeding, which in turn suggests little aggregation of food or shelter in the environment inhabited by protocetids.

Conclusions/Significance: Discovery of a near-term fetus positioned for head-first delivery provides important evidence that early protocetid whales gave birth on land. This is consistent with skeletal morphology enabling Maiacetus to support its weight on land and corroborates previous ideas that protocetids were amphibious. Specimens this complete are virtual 'Rosetta stones' providing insight into functional capabilities and life history of extinct animals that cannot be gained any other way.


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## Introduction

Archaeoceti are basal cetaceans that document the evolutionary transition of whales from land to sea during the Eocene before the appearance of modern Mysticeti and Odontoceti [1]. Archaeocetes are primitive compared to later cetaceans in retaining cheek teeth with shearing crests, flexible elbow joints, and well-formed hind limbs with feet and toes (Figure 1). Skulls with long jaws, pointed incisors, and cuspate cheek teeth indicate that archaeocetes were specialized piscivores.

Two stages of archaeocete evolution, Protocetidae and Basilosauridae, are known from articulated skeletons. Protocetidae are
middle Eocene in age (Lutetian-Bartonian, 49-37 Ma), and comprise fifteen genera and 16 species that range from South Asia and Africa to North America [2]. All are found in marine sedimentary rocks. The best known protocetids are 2 to 3 -meter long primitive forms such as Artiocetus and Rodhocetus represented by skulls and partial skeletons with artiodactyl-like ankle bones [3]. These were amphibious foot-powered swimmers [4] that retained fully developed hind limbs connected to the vertebral column, enabling limb-supported locomotion on land. Maiacetus, described below, was part of this early diversification (Figure 1C, D). Basal archaeocetes have long been interpreted as amphibious, feeding in the sea but returning to land to rest, mate, and give birth $[5,6]$.


Figure 1. Skeletons of the Eocene archaeocete whales Dorudon atrox and Maiacetus inuus in swimming pose. (A, B)- Dorudon atrox $(5.0 \mathrm{~m} ; 36.5 \mathrm{Ma}$ ) based on UM 101222 and 101215 [11] in lateral and dorsal views, respectively. (C, D)- Maiacetus inuus ( $2.6 \mathrm{~m} ; 47.5 \mathrm{Ma}$ ) based on male specimen GSP-UM 3551 in lateral and dorsal views, respectively. doi:10.1371/journal.pone.0004366.g001

Late early Eocene and early middle Eocene pakicetid and ambulocetid archaeocetes, known from less complete remains, were almost certainly semiaquatic like protocetids [7,8,9].

Basilosauridae are late middle Eocene and late Eocene in age (Bartonian-Priabonian, $40-34 \mathrm{Ma}$ ) and were widely distributed in the world's oceans. The best known is the 5 -meter long Dorudon (Figure 1A, B). Basilosauridae retained protocetid-like skulls, but their hind limbs were greatly reduced, making them fully aquatic. Basilosaurids had a powerful tail with a terminal fluke and were clearly tail-powered swimmers like modern whales $[10,11]$. Generalized Eocene protocetid and basilosaurid Archaeoceti such as Maiacetus and Dorudon were on or near the main line of early whale evolution leading to later Mysticeti and Odontoceti, which appeared at the end of the Eocene or beginning of the Oligocene.

Two specimens of a new early middle Eocene (ca. 47.5 Ma ) protocetid whale are described below, providing the first nearly complete skeleton of a protocetid whale, the first remains of a fetal skeleton of an archaeocete, and the first direct evidence for birth,
life history, and sexual dimorphism in early archaeocetes - at a time when whales were still amphibious mammals spending some time on land and some time in the sea.

## Methods

## Field Methods

Fossil specimens described here were located during surface surveys in Pakistan in 2000 and 2004. These were found as articulated skeletons, and each specimen was separated into blocks of manageable size along cracks during excavation. The blocks were encased in plaster jackets and transported to the laboratory for preparation.

## Preparation

The two specimens described here were prepared differently. The three plaster jackets of the adult female (GSP-UM 3475) were opened and cleaned to expose the articulated bones (Figure 2).


Figure 2. Adult female and fetal skeletons (type) of the protocetid Maiacetus inuus. Skull of the adult female (GSP-UM 3475a) is colored beige with brown teeth; her postcranial skeleton is colored red; the fetal skeleton (GSP-UM 3475b) is colored blue with red teeth. Blue dashed lines indicate the contours of the three field jackets and the red dashed line marks the edge of erosion.
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The ventral surface of the cranium of the female whale was stabilized, supported by a fitted fiberglass cradle, and then turned and cleaned to expose the dorsal side of the cranium. Bones of the left forelimb were removed from the second jacket individually. The thorax and lumbus of the mother whale and the fetal skull and skeleton were stabilized in situ in the third jacket with only the side shown here being cleaned (Figure 2).

The nine plaster jackets of the adult male (GSP-UM 3551) were prepared manually with a micro-airscribe. Each block containing bones was molded and cast to create an archive of the position of the bones as found in the field. Each block was then fully prepared to free individual skeletal elements. These associated elements were then assembled into a full skeletal mount for exhibition in the University of Michigan Exhibit Museum (see Figure 1A, B).

## Imaging

The fetal dentition described here was imaged with computed tomography using a Philips Tomoscan AVE1 scanner at the Radiologische Klinik, Universität Bonn (Germany).

## Terminology

Ma, geological age in millions of years before present.
Anatomical Abbreviations in Text. $C$, cervical vertebra; $C a$, caudal vertebra; $L$, lumbar vertebra; $M c$, metacarpal; Mt, metatarsal; $R$, rib; $S$, sacral vertebra; and $T$, thoracic vertebra.

Teeth are numbered sequentially from front to back (mesial to distal), with upper and lower incisors $(I)$, canines $(C)$, premolars $(P)$,
and molars $(M)$ distinguished by superscripts and subscripts, respectively.

Institutional Abbreviations. GSP-UM, Geological Survey of Pakistan-University of Michigan collection, housed in Quetta, provincial capital of Balochistan; UM, University of Michigan Museum of Paleontology, Ann Arbor, Michigan, United States of America.

## Results

## Systematic Paleontology

Mammalia Linnaeus, 1758
Cetacea Brisson, 1762
Archaeoceti Flower, 1883
Protocetidae Stromer, 1908
Protocetinae Stromer, 1908

## Maiacetus gen. nov.

Etymology. Maia, mother, and ketos, whale (Greek): named for the sex and gravid state of the holotype.

Type Species. Maiacetus inuus sp. nov.
Diagnosis. Medium-sized protocetid archaeocete with a skeleton 2.6 m in length and an estimated weight of 280390 kg . Skull has the medium-length rostrum, anteriorlypositioned nares, large mandibular canals, narrow frontal shield, broad cranial base, and large tympanic bullae typical of early protocetids. Differs from contemporary Artiocetus clavis and Rodhocetus balochistanensis in having solidly co-ossified left and
right dentaries. In the ankle, the astragalus has deep proximal and distal trochleae as in Rodhocetus, but differs in having an indented dorsolateral border; cuboid is less deeply notched for the calcaneum than in Artiocetus and more deeply notched than in Rodhocetus. Metacarpals, carpal phalanges, and all hind limb elements are shorter relative anterior thoracic centrum length than comparable ratios in Rodhocetus.

## Maiacetus inuus sp. nov.

Figures 1C-D, 5-9, 10C, 11B
Etymology. Inuus, god of fecundity (Latin): named to acknowledge both the exceptional recovery of a gravid female in the cetacean fossil record, and the importance of life history in mammalian evolution.

Holotype. GSP-UM 3475a, articulated skull, thorax, and left forelimb of an adult female. Holotype contains the skull and partially ossified skeleton of a near-term fetus, GSP-UM 3475b (Figure 2). The mother is a young adult as shown by the complete eruption of the permanent dentition, limited tooth wear, fusion of most epiphyses, and presence of a fetus.
Type Locality. Kunvit, Kohlu District, eastern Balochistan Province, Pakistan (Figure 3). GPS coordinates are $30.0963^{\circ} \mathrm{N}$ and $69.7900^{\circ} \mathrm{E}$ (WGS84 datum).

Referred Specimen. GSP-UM 3551, virtually complete skeleton interpreted as male (see below; Figures 1C, D, 9). Found in Kunvit, Kohlu District, eastern Balochistan Province, Pakistan in the same strata and about 1 km from the holotype (Figure 3). GPS coordinates are $30.1051^{\circ} \mathrm{N}$ and $69.7926^{\circ} \mathrm{E}$ (WGS84 datum).
Age. All specimens are from upper beds of the Habib Rahi Formation, a succession of interbedded marine marls and shales. The marls are approximately $10-20 \mathrm{~cm}$ thick, and the shales are approximately 1 meter thick. The marls thin and the shales thicken toward the top of the section [12]. The age of the Habib Rahi Formation has been established on the east side of the Sulaiman Range at Rakhi Nala 35 km southeast of Kunvit $[13,14]$. The specimens described here were found near the top of the major flooding sequence in the early Lutetian stage of the middle Eocene, calibrated to approximately 47.5 Ma (Figure 4).

Diagnosis. As for the genus.

## Description of Maiacetus inuus

Adult Female Skeleton (GSP-UM 3475a). The adult female skeleton (GSP-UM 3475a) and its associated fetal skeleton (GSPUM 3475b) were collected in three blocks: (1) a cranial block with


Figure 3. Map showing localities of some Eocene whales in eastern Balochistan (Pakistan). Map showing the Kunvit area in the southwestern part of Lakha Kach syncline, northwest of the town of Rakhni. Red diamonds mark type localities for Artiocetus clavis (GSP-UM 3458), Rodhocetus balochistanensis (GSP-UM 3485) [3], Protosiren eothene (GSP-UM 3487) [49], Makaracetus bidens (GSP-UM 3570; from younger beds of the overlying Domanda Formation) [2], and Maiacetus inuus (GSP-UM 3475a, b, female, fetal skeleton). Localities are in the upper part of the Habib Rahi Formation of early Lutetian age [12]. The red circle marks the locality of the referred specimen of Maiacetus inuus (GSP 3551, male), yellow circles show localities of other archaeocete specimens, and white circles show localities of other vertebrate specimens.
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## RAKHI NALA


'Ranikot': lower Rakhi Gaj shales (3125-3135) P4 Latif (1964)
Morozovella aequa, M. angulata, Planorotalites pseudomenardi, etc.

Figure 4. Temporal constraints on the age of Maiacetus inuus. Red rectangle highlights the platy limestones and marls of the upper part of the marine Habib Rahi Formation. Nannoplankton Sphenolithus spininger and Helicosphaera bramlettei indicative of NP zones 14-15 have been reported from the Habib Rahi Formation [13]. The age of Maiacetus inuus is about 47.5 Ma . Figure from [14]. doi:10.1371/journal.pone.0004366.g004
the female skull, cervical vertebrae, some anterior thoracic vertebrae, and partial scapulae; (2) a block containing the left forelimb of the adult female; and (3) a block containing the thorax of the adult mother whale, and the skull and remaining skeleton of the fetus (Figure 2). Blocks 2 and 3 were slightly displaced (3-5 cm) relative to block 1 along a small fault or joint surface after the specimen was buried and fossilized. This movement deformed immediately adjacent bones and may have destroyed parts of bones near the fault. Some bone pieces removed from the fault zone during collection fit onto bones of one block or the other.
The adult female skull and skeleton came to rest on their dorsal surfaces before burial, as is typical for archaeocetes. This stomachup orientation may be caused by a buildup of gases in the abdomen during decomposition. Skulls are seemingly more stable lying on their dorsal surface than in any other position.

The nearly complete skull of the adult female has been exposed in ventral and dorsal views (Figures 5, 6). It measures 56 cm in length (Table 1). Salient features of the skull include a relatively narrow and anteroposteriorly-elongated rostrum, with premaxillae expanded anteriorly in front of the canine teeth. The rostrum widens posteriorly and is relatively broad at the position of the left and right upper molars.

On the dorsal surface of the cranium, the external nares open above the upper canine teeth (Figure 6). The nasals are narrow and extend posteriorly to overlap the frontals. Expanded frontals
form a frontal shield with prominent supraorbital processes. The frontal shield is not as broad relative to the rest of the skull as it is in later protocetids. The jugals are relatively thin and flare posteriorly. The skull is broadest across the left and right squamosals. There is a low sagittal crest and a moderately developed nuchal crest.

On the ventral surface of the cranium the glenoid fossa for articulation of the dentary is open and relatively flat as in other early protocetids. Remnants of the basihyal, tympanohyals, and stylohyal bones can be identified on the ventral surface of the basicranium (Figure 5). The tympanic bullae are large with a sigmoid process well developed laterally and involucrum well developed medially.

The mandibular symphysis is solidly fused from the anterior end of the dentaries to a position beneath $\mathrm{P}_{2}$ (Figure 5). The ventral portions of the dentaries are hollow posteriorly, and the mandibular canal is well developed. The canal cannot be measured, however, due to dorsoventral compression of the dentaries.

Dorsoventral compression of the adult female skull has pushed some of the crowns of lower teeth into the bones of the cranium and some crowns of upper teeth into the left dentary (Figures 5, 6). The upper dental formula is 3.1.4.3 (incisors, canines, premolars, molars). Partial or complete crowns of all upper teeth are exposed in the right premaxilla


Figure 5. Skull of Maiacetus inuus (GSP-UM 3475a; female). Skull in ventral view (A) with interpretive shading and labels (B). Shaded components include skull bones (grey), teeth (brown), hyoids (orange), and postcranial bones (red). Osteological abbreviations: Ba., basihyal; Basicr., basicranium; C, cervical vertebra; Dent., dentary; Exoc., exoccipital; Jug., jugal; L, left; Max., maxilla; R, right, rib; Scap., scapula; Squ., squamosal; St., stylohyal; Stern., sternebrum; T, thoracic vertebra; Th., thyrohyal; Tym., tympanic. doi:10.1371/journal.pone.0004366.g005


Figure 6. Skull of Maiacetus inuus (GSP-UM 3475a; female). Skull in dorsal view (A) with interpretive shading and labels (B). Shaded components include skull bones (grey), teeth (brown), and postcranial bones (red). Osteological abbreviations: Dent., dentary; Front., frontal; Jug., jugal; L, left; Lac., lacrimal; Max., maxilla; Nas., nasal; Par., parietal; Pmax., premaxilla; R, right, rib; Scap., scapula; Squ., squamosal. doi:10.1371/journal.pone.0004366.g006

Table 1. Skeletal measurements (cm) of Maiacetus inuus compared to those of the previously known protocetids Artiocetus clavis and Rodhocetus balochistanensis, which are also from the early middle Eocene of Pakistan [3].

| Measurement | Artiocetus clavis | Rodhocetus balochistan-ensis | Maiacetus inuus | Maiacetus inuus | Maiacetus inuus |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | GSP-UM 3458 (Type) | GSP-UM 3485 (Type) | GSP-UM 3475a <br> (Type, female) | GSP-UM 3551 (Male) | Male/female Ratio ${ }^{2}$ |
| Cranial length | 57.5 |  | 56.0 |  |  |
| Cranial width | 44.0 |  | 27.0 |  |  |
| Tympanic bulla length | 6.5 |  | 6.1 | 6.6 | 1.09 |
| Tympanic bulla width | 4.8 |  | 4.9 | 5.4 | 1.12 |
| Mandibular foramen height | 6.0 |  |  | 7.0 |  |
| Cervical vert. C4 length | 2.8 | 3.2 |  | 2.7 |  |
| Thoracic vert. T4 length | 3.2 | 3.9 | 2.7 | 4.0 | 1.51 |
| Lumbar vert. L4 length | 4.3 |  | 4.5 | 4.7 | 1.04 |
| Caudal vert. C4 length |  | 5.5 |  | 5.0 |  |
| Scapula length | 20.0* |  |  | 21.7 |  |
| Humerus length |  | 21.0* |  | 21.5 |  |
| Ulna length |  |  | 14.6 | 17.5 | 1.20 |
| Ulna olecranon length |  |  | 4.6 | 5.7 | 1.24 |
| Ulna prox. articular breadth |  |  | 1.9 | 2.0 | 1.05 |
| Radius length |  | 11.0 | 10.0 | 11.7 | 1.17 |
| Radius distal width | 2.7 | 3.0 |  | 2.5 |  |
| Radius distal thickness | 2.3 | 1.8 |  | 2.3 |  |
| Carpus length |  | 2.3 | 2.1 | 2.2 | 1.05 |
| Pisiform length |  | 4.7 |  | 3.7 |  |
| Metacarpal length (Mc-III) |  | 7.8 | 6.1 | 6.5 | 1.07 |
| Manual phalanx length (III-1) |  | 6.3 | 5.9 | 5.9 | 1.00 |
| Manual phalanx length (III-2) |  | 3.3* | 2.6 | 3.8* | 1.48 |
| Manual phalanx length (III-3) |  | 1.3* | 1.1 | 1.4 | 1.26 |
| llium length (acetabular margin) | 12.4 |  |  | 12.4 |  |
| Acetabulum diameter | 2.8* | 3.1* |  | 3.1 |  |
| Femur length |  | 19.0 |  | 17.2 |  |
| Femur midshaft diameter (anteroposterior) | 2.2 | 2.3 |  | 2.0 |  |
| Femur midshaft diameter (bilateral) | 2.5 | 3.2 |  | 2.5 |  |
| Patella length | 3.1 | 4.1 |  | 2.9 |  |
| Patella width | 1.9 | 2.6 |  | 1.8* |  |
| Patella thickness | 1.9 | 2.3 |  | 1.6 |  |
| Tibia length |  | 21.0 |  | 18.6 |  |
| Astragalus length | 4.4 | 5.4 |  | 3.8 |  |
| Astragalus width | 2.1 | 2.3 |  | 2.0 |  |
| Astragalus height | 2.1 | 2.9 |  | 2.2 |  |
| Cuboid length | 1.9 | 2.7 |  | 1.9 |  |
| Metatarsus length (Mt-III) |  | 12.2 |  | 8.9 |  |
| Pedal phalanx length (III-1) |  | 9.0 |  | 7.0 |  |
| Pedal phalanx length (III-2) |  | 7.0 |  | 4.7 |  |
| Pedal phalanx length (III-3) |  | 2.3 |  | 1.8 |  |
| Median dimorphism |  |  |  |  | 1.12 |
| ${ }^{1}$ Body weight (kg; estimate) | 400* | 410* | 280* | 390* | 1.39* |

Sexes of the A. clavis and R. balochistanensis specimens are not known.
${ }^{*}$ Asterisk indicates estimated measurement.
${ }^{1}$ Body weight is estimated from the centrum size of available thoracic vertebrae, using multiple regression and a reference set of 13 marine mammal skeletons of known weight ( 8 cetaceans and 5 pinnipeds) spanning a range of 85 to $23,500 \mathrm{~kg}$ (see Figure 13). Weight for female Maiacetus inuus is estimated from male $M$. inuus in two ways: using the $12 \%$ average difference in comparable linear dimensions or the $4 \%$ average difference found for premolars and molars (Table 2).
${ }^{2}$ Ratio based on measurements of male and female adult specimens in Maiacetus inuus.
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and maxilla (Figure 5), except for $\mathrm{M}^{3}$, which is known in the adult male (GSP-UM 3551; see below). All incisors are caniniform with simple conical crowns on a single root. $\mathrm{I}^{1-2}$ are large and $\mathrm{I}^{3}$ is relatively small (Table 2). The canine is moderate in size and simple in form, with a conical crown on a single root. $\mathrm{P}^{1}$ has a simple crown and is single-rooted, while $\mathrm{P}^{2-4}$ have more elongated crowns and double roots. Upper molars have a simple large external paracone, with little development of a metacone. Protocones on upper molars are not exposed in the adult female skull but are well developed in
the adult male (GSP-UM 3551; see below). $\mathrm{P}^{3}$ has the anteroposteriorly longest crown in the upper dentition (Figure 5).

The lower dental formula, like the upper, is 3.1.4.3. $\mathrm{I}_{1-3}, \mathrm{C}_{1}$, and $P_{1-3}$ are present in the left dentary (Figure 6), and the remaining premolar and lower molars are known in the adult male (GSP-UM 3551; see below). Again, all incisors are caniniform with simple conical crowns on a single root. $\mathrm{I}_{1}$ is relatively small (judging from the root), $\mathrm{I}_{2}$ is large, and $\mathrm{I}_{3}$ is small (Figure 6, Table 2). Lower canine $\mathrm{C}_{1}$ is moderate in size and simple in form, with a conical crown on a single root. Lower $P_{1}$ is small, and $P_{2-3}$

Table 2. Measurements ( mm ) of teeth in female and male specimens of Maiacetus inuus.

| Tooth pos. | Upper teeth |  |  |  | Dimorphism <br> M/F | Tooth pos. | Lower teeth |  |  |  | Dimorphism ${ }^{1}$ <br> M/F | All <br> Median M/F dimorphism |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{aligned} & \text { Female GSP-UM } \\ & 3475 \mathrm{a} \end{aligned}$ |  | Male GSP-UM 3551 |  |  |  | Female GSP-UM3475a |  | Male GSP-UM 3551 |  |  |  |
|  | Left | Right | Left | Right |  |  | Left | Right | Left | Right |  |  |
| $1^{1} \mathrm{~L}$ | 16.84 |  |  |  |  | $\mathrm{I}_{1} \mathrm{~L}$ |  |  |  |  |  |  |
| w | 12.28 |  |  |  |  | w |  |  |  |  |  |  |
| H | 29.07 |  |  |  |  | H |  |  |  |  |  |  |
| $\mathrm{I}^{2} \mathrm{~L}$ | 18.26 | 18.17 |  |  |  | $\mathrm{I}_{2} \mathrm{~L}$ | 18.00* | 17.80 |  |  |  |  |
| w |  | 12.61 |  |  |  | w |  | 16.30 |  |  |  |  |
| H | 23.49 | 29.03 |  |  |  | H |  |  |  |  |  |  |
| $\mathrm{I}^{3} \mathrm{~L}$ | 16.01 | 14.91 |  |  |  | $\mathrm{I}_{3} \mathrm{~L}$ | 14.17 |  |  |  |  |  |
| w |  | 9.36 |  |  |  | W | 9.52* |  |  |  |  |  |
| H | 12.60 | 15.24 |  |  |  | H | 12.43* |  |  |  |  |  |
| $\mathrm{C}^{1} \mathrm{~L}$ |  | 19.23 |  | 23.80* | 1.24 | $\mathrm{Cl}_{1} \mathrm{~L}$ | 18.09 |  |  |  |  |  |
| w | 13.21* |  |  | 13.65 | 1.03 | w | 10.95 |  |  |  |  | 1.20 |
| H | 26.00 |  |  | 31.30 | 1.20 | H |  |  |  |  |  |  |
| $\mathrm{P}^{\mathbf{1}} \mathrm{L}$ |  | 16.81 |  | 16.24 | 0.97 | $\mathrm{P}_{1} \mathrm{~L}$ | 15.69 |  |  |  |  |  |
| w |  | 7.76 |  | 8.98 | 1.16 | W | 8.69 |  |  |  |  |  |
| H |  | 16.13 |  | 17.06 | 1.06 | H | 14.10 |  |  |  |  |  |
| $\mathbf{P}^{\mathbf{2}} \mathrm{L}$ |  | 28.60 |  | 29.00 | 1.01 | $\mathrm{P}_{2} \mathrm{~L}$ | 28.28 |  | 28.97 |  | 1.02 |  |
| W |  |  |  | 12.72 |  | w | 11.00* |  | 11.44 |  | 1.04 |  |
| H |  |  |  | 27.25 |  | H |  |  | 27.64 |  |  |  |
| $\mathrm{P}^{3} \mathrm{~L}$ |  | 28.65 |  | 33.79 | 1.18 | $\mathrm{P}_{3} \mathrm{~L}$ |  |  |  |  |  |  |
| w |  |  |  | 16.40 |  | W | 11.60 |  | 12.06 |  | 1.04 |  |
| H |  |  |  | 25.04 |  | H |  |  | 30.42 |  |  |  |
| $\mathrm{P}^{4} \mathrm{~L}$ |  | 25.46 |  | 25.57 | 1.00 | $\mathrm{P}_{4} \mathrm{~L}$ |  |  | 33.15 |  |  |  |
| w |  |  |  | 20.22 |  | w |  |  | 12.20 |  |  | 1.04 |
| H |  |  |  | 23.68 |  | H |  |  | 30.33 |  |  |  |
| $\mathrm{M}^{1} \mathrm{~L}$ |  | 19.93 |  | 20.21 | 1.01 | M ${ }_{\text {L }}$ |  |  | 22.35 |  |  |  |
| w |  |  |  |  |  | W |  |  | 9.74 |  |  |  |
| H |  |  |  |  |  | H |  |  |  |  |  |  |
| $\mathrm{M}^{\mathbf{2}} \mathrm{L}$ |  | 20.00 |  | 21.39 | 1.07 | $\mathrm{M}_{2} \mathrm{~L}$ |  |  | 25.85 |  |  |  |
| w |  |  |  | 17.40 |  | w |  |  | 12.50 |  |  |  |
| H |  |  |  | 15.99 |  | H |  |  | 26.32 |  |  |  |
| $\mathrm{M}^{\mathbf{3}} \mathrm{L}$ |  |  |  | 14.00 |  | M ${ }^{\text {L }}$ |  |  | 24.26 |  |  |  |
| w |  |  |  | 12.16 |  | w |  |  | 12.61 |  |  |  |
| H |  |  |  | 15.22 |  | H |  |  | 23.48 |  |  |  |

## "Asterisk indicates estimated measurement.

${ }^{1}$ Ratios quantifying male-to-female tooth size dimorphism are provided when teeth at the same position can be compared. Median dimorphism in canine size, based on upper canines, is 1.20 , meaning males have canines on the order of $20 \%$ larger than those of females. Median dimorphism in premolar and molar size is 1.04 , meaning that males have premolars and molars only about $4 \%$ larger than those of females.
doi:10.1371/journal.pone.0004366.t002
have anteroposteriorly longer crowns. The longest tooth in the lower dentition was probably $\mathrm{P}_{3}$, but this cannot be determined with confidence from the specimens at hand.

The cervical vertebrae (C1-C7) of the adult female (GSP-UM 3475a) are preserved in articulation in the cranial block, but these are difficult to study or measure individually (Figures 2, 5). The first three thoracic vertebrae and a remnant of the fourth (T1-T4) are preserved in articulation posterior to the cervicals. Nothing has been recovered of T 5 , which was evidently lost during displacement of cranial and thoracic blocks along the intervening fault or joint surface. The vertebral column continues in the thoracic block. T6 is represented by a piece of the centrum, while the remaining thoracic vertebrae (T7-13) and lumbar vertebrae (LlL4 and part of L5) are preserved in articulation (Figure 2).
A sternebrum and some ribs are present in the cranial block (Figures 5, 6). The ribs continue into the thoracic block, where they are preserved for the most part in order. The distal ends of the right ribs lie on top of the vertebral centra, whereas distal ends of the left ribs are displaced more laterally. Ribs in the thoracic block can only be identified by counting backward from the most distal rib (R13).
The left forelimb was preserved in articulation on the left side of the thorax (Figures 2, 7). A fragment of the left scapula lies under the left squamosal, and a fragment of the right scapula lies under the anterior ribs on the right side of the cranium. Much of the left scapula and the proximal end of the left humerus were damaged at the juncture of the cranial and forelimb blocks. The distal end of the humerus is relatively narrow with a shallow trochlea and deep olecranon fossa for articulation with the ulna. The radius and ulna are well preserved although somewhat flattened (Figure 7).

The distal row of carpal bones is alternating rather than serial relative to the proximal row. Metacarpal I is the smallest, metacarpals III and IV are the largest, and metacarpal III is distinctly longer than metacarpal IV. With the exception of the pollex (digit I), all of the proximal phalanges are about the same length (the proximal phalanx of digit I is not known). Similarly, middle phalanges of digits II through V are similar in length, although phalanx III-2 is distinctly more robust. Terminal phalanges of digits I and V are simple and pointed and probably did not bear a hoof, whereas the blunt end of terminal phalanx II3 indicates that it was hoof bearing. Terminal phalanges II-3, III3, and IV-3 in the adult male (GSP-UM 3551) all bore hooves.

Preservation of the vertebral column ends at L5. Nothing remains of L6, the sacrum, or the tail. Very little of a hind limb is preserved in the adult female (Figure 2). The left ilium is represented only by the crescent-shaped anterior margin. The distal left femur, patella, and proximal end of the left tibia are preserved in articulation, but the rest of the left hind limb and all of the right hind limb are missing.
Fetal Skeleton within Adult Female (GSP-UM 3475b). The fetal skeleton (GSP-UM 3475b) is preserved within the ribcage of the type specimen (GSP-UM 3475a) (Figure 2, colored blue). This immature individual is interpreted as a fetal skeleton, rather than an ingested meal, because of the absence of any damage to the skull. Protocetids had shearing molars used to slice and chew their prey. The skull of the fetal skeleton could not have survived such mastication and be as well preserved as it is.

The fetal skeleton preserves the skull, some cervical and thoracic vertebrae, ribs, and portions of the fore- and hind limbs. The skull includes the cranium and left and right lower jaws, with crowns and roots of many deciduous teeth in place in the upper and lower jaws. The surfaces of the fetal bones are porous in contrast to the smoother surfaces of adult bones. The fetal skeleton has a


Figure 7. Left radius, ulna, carpus and manus of Maiacetus inuus (GSP-UM 3475a, female). Open areas are reconstructed. Abbreviations: C, cuneiform; L, lunar; M, magnum; P, pisiform; S, scaphoid; Tr, trapezium; $T r d$, trapezoid; $U$, unciform.
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preserved length of about 33 cm , and probably had a total length approximately twice the preserved length.

The length of the fetal skull, as preserved, is 15 cm , and this probably measured 17 cm when the missing anterior end of the
rostrum is added (Figure 8). Identifiable bones include the left frontal, the left jugal, the occipital with left and right condyles flanking the foramen magnum, and the left and right dentaries.

The fetal dentition is largely intact but interpretation is complicated by bilateral compression of the skull (Figure 8A, B). Left upper teeth are displaced dorsally relative to right upper teeth, and the left dentary is displaced slightly anteriorly relative to the right dentary. Teeth from both left and right sides, being denser and harder than surrounding fetal bone, all show on the exposed left side of the skull.

The largest tooth in each quadrant is the distinctive fourth deciduous premolar. These four premolars (left and right $\mathrm{dP}^{4}$ and $\mathrm{dP}_{4}$ ), serve as landmarks for identification of more anterior and posterior teeth. Anterior to $\mathrm{dP}^{4}$ and $\mathrm{dP}_{4}$, there are partial crowns of all of the third deciduous premolars (left and right $\mathrm{dP}^{3}$ and $\mathrm{dP}_{3}$ ), as well as upper and lower deciduous canines. Teeth anterior to $\mathrm{dP}^{3}$ and $\mathrm{dP}_{3}$ were identified by their morphology and by their stage of development, using an undescribed protocetid skull with a good deciduous dentition for comparison.

Posterior to $\mathrm{dP}^{4}$, there are partial crowns of the left and right first permanent upper molars visible below the left frontal and above the left jugal (Figure 8,). Left $\mathrm{M}^{1}$ is the most clearly exposed and it appears to have the main paracone cusp almost fully formed, with the crown as a whole about one-half mineralized. Crowns of left and right lower permanent molars are not visible but were presumably also partially mineralized.

Cervical and anterior thoracic vertebrae are compressed in articulation, and the ribs are preserved as narrow ribbons of porous bone pressed onto the ventral surfaces of the mother's thoracic vertebrae. Parts of seven long bones are evident. The largest of these are presumably humeri, and there may be remnants of ulnae and radii as well. Seven long bones would exceed the number expected in a pair of protocetid forelimbs, and so some may pertain to the hind limbs. More precise identification is hindered by the absence of ossified epiphyses.

Adult Male Skeleton (GSP-UM 3551). The most complete skeleton of Maiacetus inuus is that of a second adult (Figure 1G, D) found in articulation in the field (Figure 9). The skull lacks the anterior portion of the rostrum, while the vertebral column and fore- and hind limbs are virtually complete. The specimen is interpreted as male because it is larger than the female skeleton described above, the canine teeth are larger relative to the size of other teeth, and the pelvic morphology is male (see below). Comparative measurements are provided in Tables 1-3.

The male skull is similar to that of the female described above. Here again the anterior rostrum is narrow but it expands rapidly at mid length toward the posterior end of the skull. The palate is much broader posteriorly than it is anteriorly. The frontal shield is expanded with broad supraorbital processes, but it is again narrower than that of later protocetids. Lacrimal bones are exposed on the face anterior to the orbits. The high, narrow, and deep nuchal area is bordered ventrolaterally by robust, broadlyspaced parocciptal processes, suggesting the presence of powerful neck muscles. Tympanic bullae are large, robust, and firmly attached to the underside of the endocranium. The convex occipital condyles are oriented posteriorly.

The mandibular symphysis is fused as far posteriorly as the distal margin of $\mathrm{P}_{2}$. The large mandibular foramen leading to the mandibular canal is mediolaterally narrow and dorsoventrally deep (depth estimated as 7 cm ). It opens internally at the posterior end of the dentary. The mandibular condyles are positioned well below the apex of the coronoid process, and the gonial angle of the mandible is compressed and square in outline (viewed laterally; Figure 1A).

Upper and lower incisors are missing in the adult male (GSPUM 3551), but the upper canine ( $\mathrm{C}^{1}$ ) is large and anteroposteriorly expanded. The upper first premolar $\left(\mathrm{P}^{1}\right)$ is small, while $\mathrm{P}^{2}$ is closer in size to $\mathrm{P}^{3}$ and $\mathrm{P}^{4} . \mathrm{P}^{3}$ has the largest crown of the upper cheek teeth. $\mathrm{P}^{3}-\mathrm{M}^{3}$ are closely spaced with no diastemata. A protocone is well developed on $\mathrm{P}^{4}$ and on $\mathrm{M}^{1}$. This primitive cusp decreases in size on $\mathrm{M}^{2}$ and nearly disappears on $\mathrm{M}^{3}$. In the lower dentition, $\mathrm{C}_{1}$ and $\mathrm{P}_{1}$ are represented only by impressions of the crowns and roots. The crown of the lower canine crown is longer anteroposteriorly and the root thicker than in the female (GSPUM 3475a), while $P_{1}$ is small. Lower $P_{2}, P_{3}$, and $P_{4}$ are all relatively large, with $\mathrm{P}_{3}$ or $\mathrm{P}_{4}$ the largest of the lower cheek teeth. The lower molars are closely spaced, with a high anterior cusp (protoconid) and lower posterior cusp (hypoconid).

The vertebral formula (cervicals, thoracics, lumbars, sacrals, caudals) is $7 \mathrm{C}-13 \mathrm{~T}-6 \mathrm{~L}-4 \mathrm{~S}-21 \mathrm{Ca}$ for a total of 51 vertebrae, which differs by only two caudal vertebrae from the formula found in some primitive artiodactyls [15]. Precaudal vertebrae increase in size along the series as in other protocetids (Table 3). The atlas (C1) has broad transverse processes, moderately deep and curved prezygapophyses, and a complex foraminal pathway for the vertebral arteries to accommodate motion of the skull. In C3-C7 the orientation of zygapophyses, the saddle-shape of vertebral body surfaces, and the imbrication of transverse processes suggest that the neck was more capable of dorsiflexion than lateral rotation. Anterior thoracic vertebrae have long spinous processes for muscles and ligaments inserting on the exoccipital region and nuchal area of the skull to stabilize the head. Thoracic T11, the diaphragmatic vertebra, has thoracic-like prezygapophyses and lumbar-like postzygapophyses. T12 is the anticlinal vertebra with a vertical neural spine (Figure 1A). In the lumbus, anterodorsally inclined spinous processes, non-revolute zygapophyses, and the anteroventral orientation of transverse processes indicate the potential for dorsoventral flexion of the trunk and vertebral column in general. Flexion, however, was limited by the anteroposterior width and length of the spinous processes of the lumbar vertebrae.

The sacrum is composed of four vertebrae, the first three fully co-ossified and the fourth with fused pleurapophyses. Robust auricular processes are present on S1 for articulation with ilia of the pelvis. The spinous processes of all sacral vertebrae have massive bases. Presence of a solidly fused sacrum precluded smooth undulatory motion of the body during swimming.

The caudal series is composed of 21 vertebrae (Figures 1C, D, 9). The first 14 caudals have associated hemal arches or chevron bones. Anterior caudals are robust, with substantial spinous processes, metapophyses, and transverse processes. The latter are divided in Ca 4-6. Mid-caudal vertebrae are also large but have more diminutive processes. The size and complexity of vertebrae in the posteriormost part of the tail decrease rapidly following the last chevron-bearing caudal (Figures 1C, D, 9).

The ribs are elongate and moderately curved, with no sign of pachyostosis or osteosclerosis. All but the last few are doubleheaded. The sternum includes eight elements with a T-shaped manubrium, six sternebrae, and an elongate xiphisternum.

The delicate scapula is longer than wide, with a projecting acromion and shallow glenoid; the coracoid process forms a large tubercle above the glenoid. The scapular spine divides the scapular blade unequally into smaller anterior and larger posterior portions. The humerus is relatively long, flattened, and anteriorly curved. Proximally, the head is ovoid with a narrow bicipital groove. Distally, the humerus is narrow with a deep trochlea for the ulna. Posteriorly, there is a deep olecranon fossa. The convex articulation for the radial head is located lateral to the ulnar


Figure 8. Fetal skull of Maiacetus inuus (GSP-UM 3475b). (A)-Photograph showing bones (shaded white) and teeth (shaded brown) in lateral view. (B)-CT image overlain on tooth-bearing part of fetal skull in lateral veiw. Abbreviations: left frontal (L. Front.), left jugal (L Jug.), occipital (Occip.) with left and right condyles flanking the foramen magnum, and left and right dentaries ( $L$ dent., $R$ dent.). Teeth are shaded brown, with darker brown representing enamel, and lighter brown representing roots and exposed dentine. White and black labels identify teeth of the left and right sides, respectively. Dotted lines trace outlines of fully formed crowns of left $\mathrm{dP}^{3}$ and $\mathrm{dP}^{4}$. These crowns are visible on the surface (A) where thin bone of the maxilla is pressed over more rigid underlying crowns, and as denser masses in the CT scan (B). Remaining teeth are identified by size and position relative to $\mathrm{dP}^{3}$ and $\mathrm{dP}^{4}$. Note presence of the developing crown of permanent left $\mathrm{M}^{1}$ posterodorsal to the crown of left $\mathrm{dP}^{4}$ (dorsal to the left jugal and posterior to the left frontal). Partial crown of right $M^{1}$ (unlabeled) is visible just below and posterior to left $M^{1}$. doi:10.1371/journal.pone.0004366.g008


Figure 9. Skeleton of Maiacetus inuus (GSP-UM 3551, male). Skeleton as preserved in left lateral view. Gray shading is marl matrix (see Figure 1C, D for skeletal restoration). doi:10.1371/journal.pone.0004366.g009
trochlea. The medial epicondyle is robust and projecting. Ulna and radius are shorter than the humerus (Figure 7). The ulna is flattened and has a large, blade-shaped olecranon process. The trochlear notch of the ulna projects farther anteriorly, locking into the humeral olecranon fossa during extension. The notch is bifid with a broad articulation distally for the radius. Although the radius and ulna are not coossified, they were fixed in a pronated position. The head of the radius, the shortest forelimb element, has a square contour and saddle-shaped surface. The radial shaft, unlike that of the humerus and ulna, has a subtriangular crosssection. The robust distal end of the radius has a distinct oval, flat articular surface for the carpus.

The carpus is composed of a series of proximal and distal carpals that vary in shape from ovoid to polygonal (Figure 7). Proximal and distal carpal rows are staggered with interlocking articulations. The pisiform projects strongly ventrally, more than required to maintain the aponeurosis of the carpal tunnel, providing a powerful mechanical advantage for the flexor carpi ulnaris. The prominent medial malleolus on the humerus provides the origin for this muscle. The pisiform resembles the fifth metacarpal ( $\mathrm{Mc}-\mathrm{V}$ ) in size and morphology. The carpals are transversely arched providing a deep ventral tunnel for flexor tendons. Digits three and four are dominant and metacarpal length decreases in the order of $\mathrm{Mc}-\mathrm{III}, \mathrm{Mc}-\mathrm{IV}, \mathrm{Mc}-\mathrm{II}, \mathrm{Mc}-\mathrm{V}$, and Mc-I (Table 1). Large, paired sesamoids are located on the ventral surfaces of the head of each metacarpal. Proximal phalanges are long and relatively thin, as are middle phalanges. Distal phalanges
are much shorter and somewhat more robust. The manus is shorter than the pes and has more delicate terminal phalanges.

The innominate is anchored to the sacrum. The ilium is short anterposteriorly relative to the length of the pelvis. The round, well buttressed acetabulum is notched with a pit for the ligamentum teres. The obturator foramen is large. The ischium is large, flattened, and extends farther from the acetabulum than does the ilium (Figure 1A). The pubic notch formed by inferior rami of left and right pubic bones is ' V ' shaped (Figure 10), as expected in a male. The robust femora are short relative to the length of the distal segments of the hind limb. The femoral head is rounded with a deep fovea capitis femora. Robust muscle attachment sites on the femur include prominent greater and lesser trochanters, a deep intertrochanteric fossa, and a distally extended lateral crest for insertion of vastus lateralis. The distal condyles of the femur are offset from one another, which served to rotate the tibia medially during flexion. The relatively long tibia has a robust proximal end and deeply trochleated distal end shaped to receive the astragalus. The relatively long, thin fibula has articular surfaces proximally for the tibia and distally for the calcaneum.

The ankle, preserved in articulation on both sides in GSP-UM 3551, resembles that of larger-bodied Rodhocetus balochistanensis more than that of similar-sized Artiocetus clavis (Figure 11). The astragalus is 'double-pulleyed' with a trough-shaped trochlea for the tibia and an opposing trochlea for the navicular. Other astragalar facets include a narrow convex articular surface for the cuboid adjacent to the deeply notched surface for the navicular, a

Table 3. Vertebral centrum measurements ( mm ) for adult male Maiacetus inuus (GSP-UM 3551; Figure 9).

| Vertebra | Length | Width | Height | Vertebra | Length | Width | Height |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| C1 | 23.5 | 89.5 | 63.3 | S1 | 47.8 | 60.4 | 34.6 |
| C2 | 34.5 | 37.7 | 34.5 | S2 | 44.7 | 58.4 | 32.8 |
| C3 | 28.3 | 42.5 | 34.0 | S3 | 48.5 | 54.1 | 34.4 |
| C4 | 26.6 | 39.3 | 31.9 | S4 | 52.2 | 51.8 | 37.5 |
| C5 | 25.4 | 37.9 | 31.7 | Ca1 | 47.6 | 52.7 | 42.9 |
| C6 | 28.6 | 41.4 | 32.6 | Ca2 | 47.5 | 51.6* | 41.7 |
| C7 | 27.7 | 49.9 | 29.5 | Ca3 | 49.4 | - | 47.9 |
| T1 | 34.4 | 48.6 | 35.2 | Ca4 | 50.2 | - | 48.8 |
| T2 | 35.8 | 50.8 | 35.4 | Ca5 | 52.8 | - | (49.4) |
| T3 | 35.8 | 52.9 | 35.7 | Ca6 | 54.2 | - | (50.5) |
| T4 | 40.1 | 52.7 | 40.0 | Ca7 | 59.8 | - | (49.4) |
| T5 | 35.0 | 54.7 | 39.5 | Ca8 | 61.1 | - | (48.8) |
| T6 | 36.0 | 56.2 | 36.4 | Ca9 | 60.0 | - | (47.0) |
| T7 | 36.4 | 56.8 | (36.7) | Calo | 60.0 | - | (44.0) |
| T8 | 36.5 | 57.1 | (37.0) | Cal1 | 57.4 | - | (41.0) |
| T9 | 36.4 | 58.0 | (37.3) | Cal2 | 51.9 | - | (36.5) |
| T10 | 35.9 | 59.6 | 37.7 | Cal3 | 45.2 | 27.0* | 31.0* |
| T11 | 39.9 | 58.0 | 36.8* | Cal4 | 37.6 | 29.0 | 27.8 |
| T12 | 39.7 | 54.0 | (38.0) | Ca15 | 29.7 | 24.7 | 16.3 |
| T13 | 39.5 | 52.4 | 39.2 | Cal6 | 23.9 | 19.5 | 11.4 |
| L1 | 40.1 | 51.9 | 39.6 | Cal7 | 24.7 | 14.2 | 9.2 |
| L2 | 41.2 | 48.6 | (41.5) | Cal 18 | 20.3 | 8.6 | 5.1 |
| L3 | 46.0 | 54.3 | 42.2 | Cal9 | 19.9 | 6.6 | 3.8 |
| L4 | 46.5 | 60.2 | 40.0 | Ca20 | 16.7 | 6.6 | 5.0 |
| L5 | 46.1 | 65.3 | 40.2 | Ca21 | 11.1 | 4.8 | 3.1 |
| L6 | 52.2 | 65.1 | 37.8 |  |  |  |  |

Abbreviations: $C$, cervical vertebra; Ca, caudal vertebra; $L$, lumbar vertebra; S, sacral vertebra; $T$, thoracic vertebra.
*Asterisk indicates estimated measurement.
( ) indicate interpolated measurement.

- Ca3 through Ca12 are bilaterally compressed, precluding measurement of their width.
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small curved surface laterally for articulation with the ectal facet of the calcaneum, a proximodistally elongated convex facet ventrally for long excursion on the calcaneal sustentacular surface, and a small facet adjacent to the cuboid facet for articulation with the calcaneum. On the calcaneum, the tuber is long and robust; the sustentaculum is oval, small, and shelf-like; the ectal facet is convex and confluent with the curved facet for the fibula; and the distal cuboid facet is slightly concave and obliquely oriented. The deeply concave navicular has a proximal keel for articulation with the astragalus, and distal facets for articulation with the ento-, meso-, and ectocuneiform (Figure 11). The cuneiforms are small and subrectangular with simple articular surfaces for the second and third metatarsals ( $\mathrm{Mt}-\mathrm{II}$ and $\mathrm{Mt}-\mathrm{III}$ ). It is possible that a tiny distal extension of the entocuneiform accommodated a very diminutive (but unrecovered) Mt-I. A pair of proximal articular surfaces on the cuboid include a concave medial portion for the astragalus and an obliquely notched, flatter surface for articulation with the distal surface of the calcaneum. Distally, the cuboid has a larger, flat articular surface for Mt-IV and a smaller articular surface for Mt-V.

Pedal digits III and IV are largest. Lengths decrease in the sequence Mt-IV, Mt-III, Mt-V, and Mt-II. Each metatarsal head has paired facets ventrally for articulating sesamoid bones. Proximal and middle phalanges are long and relatively thin. Ungual phalanges are much shorter, ventrally flattened, and ornamented with dorsal tubercles and sulci to accommodate extensor tendons.

## Discussion

## Extraordinary Fossils

The fossils described here include the first association of adult female and fetal whale skeletons, the latter apparently near term and in birth position (GSP-UM 3475a, b) (Figure 2). The adult male skeleton (GSP-UM 3551) found nearby is exceptionally complete (Figure 9). Together these specimens document a new protocetid whale preserving a complete vertebral column and complete fore- and hind limbs. The vertebral formula of Maiacetus is similar to that preserved in the other protocetids Rodhocetus and Qaisracetus (7C-13T-6L-4S) [14,16], and in addition indicates the presence of 21 caudal vertebrae. As in all living semiaquatic mammals, the limbs are relatively short and pelvic girdles provide a direct connection to the vertebral column for weight-bearing. The ankles are artiodactyl-like, with a double-pulley astragalus, notched cuboid, and curved fibular facet characteristic of artiodactyls [3]. Metapodials are elongated, and the manual and pedal digits were almost certainly webbed.

Maiacetus has a piscivorous dentition and, like other early protocetids, is interpreted as an amphibious, semiaquatic, footpowered swimmer that fed in the sea and came ashore to rest, mate, and give birth, as anticipated by Fordyce [5]. While the hind limbs were capable of bearing the weight of the body on land, the proportions of the limbs and the long phalanges of both hands and feet would have limited terrestrial locomotion and prevented Maiacetus from traveling any substantial distance from water. Maiacetus differs from other early protocetids in having dentaries fused at the mandibular symphysis (Figure 5), ankles of distinct proportions (Figure 11), and limbs slightly different in proportion from those of other early protocetids (Figures 12, 13). These differences suggest that there are feeding and swimming specializations among protocetids that remain to be clarified.

## Birth on Land

Presence of an intact fetal skeleton (GSP-UM 3475b) enables the first unequivocal determination of sex for an archaeocete: GSP-UM 3475a is female. The presence of one fetal skeleton and no trace of a second indicates that birth in Maiacetus involved a single calf. Early archaeocetes thus resemble all extant large semiaquatic mammals, which invest their energy in gestation and parenting of a single infant per breeding event [17].

The rostrum of the fetal skeleton is pointing opposite that of the mother (Figure 2). Posterior orientation of the fetal head occurs in the initial stage of birth in artiodactyls (Figure 14C). However, given the shorter neck in Maiacetus, this may have been the fetal position during all of late gestation. Because the head of the fetal skeleton of Maiacetus is not in the mother's pelvic canal, the fetus may have been near term but not full term. Partially formed M1 crowns in the fetal cranium suggest that the fetus was at least near term.

The fetal skeleton is positioned for head-first birth, a universal birthing posture in large-bodied land mammals, but one that is anomalous in fully-aquatic marine mammals [18,19,20]. A nearterm fetus in an ungulate or whale may rotate about its long axis as it passes through the birth canal, but it cannot turn head-to-tail


## Basilosauridae

Dorudon atrox
36.5 Ma



Figure 12. Proportion-adjusted skeletal profiles of middle and late Eocene archaeocete whales Maiacetus inuus and Dorudon atrox, respectively. (A)—Maiacetus inuus, a semiaquatic foot-powered swimmer from the Middle Eocene. (B)—Dorudon atrox, a fully-aquatic tail-powered swimmer from the Late Eocene. Baseline is mean length of anterior thoracic vertebrae (stippled box). Sacral vertebrae are enclosed in a second stippled box where sacrals can be identified (e.g., by co-ossification). Maiacetus has a profile more like that of mammals capable of supporting their weight on land, whereas Dorudon has the profile of a modern whale. Interpretation of profiles and the method of median serial-multiple-regression estimation of body weights is explained in [50]. Abbreviations: TCNDSFB, longest tooth length, condylobasal cranium length, narial position, dentary length, symphysis position, mandibular foramen height; and bulla length, respectively; SHRC, scapula, humerus, radius, and Mc-III lengths, respectively; IFTT, innominate, femur, tibia, and Mt-III lengths, respectively.
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Altricial-to-precocial birth status in mammals is generally represented on a behavioral spectrum scored from I to IV. Altricial I lies at one extreme ('birth with eyes closed, naked, and nidicolousor nest-dwelling'), and precocial IV at the other ('birth with eyes open, haired, and nidifugous or nest-fleeing') [21]. Dental development provides a parallel indication of birth status
(Table 4). The state of the dentition at birth is a direct indicator of the ability of a newborn to supplement mother's milk with other foods. Natural selection can separately adjust a newborn's ability to feed itself, sense its environment, regulate its temperature, and move [24], yet there is also an overall association of these aspects of newborn maturity. Table 4 shows that mineralization of the


Figure 13. Principal components plot of trunk and limb skeletal proportions for Maiacetus inuus and a representative sample of 50 extant semiaquatic mammals. Maiacetus inuus is similar to Rodhocetus species, but lacks elongation of the manus, hind limb, and pes characteristic of the latter. Maiacetus is about equally aquatic (PC-II) to Rodhocetus but less specialized as a hind-limb swimmer (PC-III). It falls closest to the giant otter (Pteronura brasiliensis) among extant mammals. Background and comparative data for this analysis are documented and explained in [4].
doi:10.1371/journal.pone.0004366.g013
dentition at birth corresponds to the classic categories of neonate maturity. In generalized mammals, crowns of the deciduous dentition appear to be largely mineralized in utero. The permanent dentition, however, shows an enormous range, with some


Figure 14. Head versus tail presentation of near- and full-term calves in a domestic cow and harbor porpoise. Domestic cow (Bos taurus) calf (blue) in (A) ninth and final month of gestation before 'turning' (axially rotating), (B) turned as the birth process begins, and (C) with forelimbs and head partially extruded in the initial stage of birth (from [51, 52]). Harbor porpoise (Phocaena phocaena) (D) with full-term calf (blue) with tail partially extruded in the initial stage of birth (from [18]).
doi:10.1371/journal.pone.0004366.g014
mammals showing no mineralization of permanent teeth at all (e.g., a newborn bear cub [22]), and others having all adult crowns fully formed (e.g., newborn guinea pig [23]). While eruption of teeth through the gums in living mammals is informative, the clearest association with precocial categories II-IV is some degree of mineralization of the permanent dentition, beginning with the first molar and in some cases extending to mineralization of later developing teeth.

The fetal skeleton of Maiacetus described here shows development of the deciduous dentition to have been well underway, with partial mineralization of upper first permanent molars in utero. The fetal dentition of Maiacetus is as developed as that of a newborn fallow deer (Table 4). Thus Maiacetus was clearly a precocial mammal.

Precociality involves a high degree of maternal investment starting from conception and extending through relatively long periods of gestation and lactation to nutritional independence of the calf. Marine mammals living today are all precocial, and energy is channeled to early growth [25]. Semiaquatic young must be able to move soon after birth to avoid predators on land and in the sea, and fully aquatic young must be able to swim to keep pace with their mothers [17]. Larger newborn body size is also an important component of precociality because it reduces the newborn's surface-to-volume ratio and enhances its ability to maintain its body temperature on exposed shorelines and in the water.

Artiodactyls, the group ancestral to whales [3], give birth to young that range from slightly precocial (pigs) to highly precocial (sheep, deer, giraffe). Whales probably evolved from anthracobunid artiodactyls, a stem group that may be closely related to

Table 4. Developmental maturity categories of Langer [21] and mineralization of first molar $M^{1}$ and other permanent teeth in newborn terrestrial mammals compared to the early whale Maiacetus inuus.

| Species | Mineralization of the permanent dentition at birth | Reference |
| :---: | :---: | :---: |
| Category I. Altricial: eyes closed, naked, nidicolous or 'nest-dwelling' |  |  |
| Gray short-tail opossum, Monodelphis domestica | None | [33] |
| Brown bear, Ursus arctos | None | [22] |
| Indian mongoose, Herpestes auropunctatus | None | [34] |
| Ferret, Mustela putorious | None | [35] |
| Domestic dog, Canis familiaris | Trace | [36] |
| Category II. Precocial: eyes open, haired, nidicolous or 'nest-dwelling' |  |  |
| Domestic pig, Sus scrofa | None | [37] |
| Miniature pig, Sus scrofa | M1 cusp tips beginning to mineralize | [38] |
| Collared peccary, Tayassu tajacu | M1 cusp tips formed | [39] |
| Category III. Precocial: eyes open, haired, transported (largely primates) |  |  |
| Human, Homo sapiens | M1 cusp tips beginning to mineralize | [40] |
| Great apes, Pan, Gorilla, and Pongo spp. | M1 cusp tips beginning to mineralize | [40] |
| Gibbon, Hylobates sp. | M1 cusp tips beginning to mineralize | [40] |
| Rhesus macaque, Macaca mulaltta | M1 cusp tips formed | [41] |
| Mongoose lemur, Eulemur mongoz | M1 cusp tips formed | [42] |
| Coquerel's sifaka, Propithecus verreauxi | M1 crown almost complete, M2 near coalescence, M3 trace | [42] |
| Tarsier, Tarsius bancanus | M1 'well differentiated' (some deciduous teeth resorbed in utero) | [43] |
| Category IV. Precocial: eyes open, haired, nidifugous or 'nest-fleeing' |  |  |
| American tapir, Tapirus cf. T. bairdii | M1 anterior cusps connected | [44] |
| Domestic sheep, Ovis aries | M1 cusps connected; crown one-quarter formed | [45] |
| Maiacetus inuus ${ }^{1}$ | M1 crown one-half formed | this study |
| Fallow deer, Dama dama | M1 crown one-half formed | [46] |
| Pygmy hippo, Hexaprotodon liberensis | M1? (many deciduous teeth erupted at birth; large M1 crypt) | [27] |
| Giraffe, Giraffa camelopardalis | M1 crown complete (all deciduous teeth erupted at birth) | [47] |
| Fur seal, Callorhinus ursinus | M1-2 crowns complete (deciduous teeth shed in utero) | [48] |
| Guinea pig, Cavia porcellus | All permanent teeth erupted; roots incomplete | [23] |

Living whales are not included because none have deciduous teeth and many have few or no teeth. Because early whales evolved from artiodactyls [3], Ovis, Dama, Hexaprotodon, and Giraffa are better analogs than primates.
${ }^{1}$ Presence of mineralization of the crown of $M^{1}$ in the fetal skull of Maiacetus inuus indicates precocial development and the probability that newborns were open-eyed, hairy and nidifugous.
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Hippopotamidae [26]. Hippopotamid young lie at the precocial end of the birth spectrum [27,28], and it is reasonable to infer that precociality was primitive for whales and their closest artiodactyl ancestors. Precociality may be a key life history trait that enabled the earliest whales to navigate the land-sea transition.

## Sexual Dimorphism

The type specimen of Maiacetus inuus, GSP-UM 3475a, has a fetus in utero and is clearly female. We interpret the referred adult skeleton of Maiacetus (GSP-UM 3551) as male because it contains no fetus, averages $12 \%$ larger in linear measurements than the known female, and has canine teeth that are 20\% larger (Tables 1, 2). A $12 \%$ difference in linear measurements of male and female Maiacetus yields an expected weight difference of about $39 \%$ (Table 1). Relatively large canines are commonly found in male mammals that use these teeth in threat displays and fighting. In addition, the pelvis of GSP-UM 3551 is well preserved and has a V-shaped pubic notch, formed by the inferior rami of the left and right pubes, which is characteristic of males (Figure 10). The notch
is broader and U-shaped in females that give birth to a single large fetus, such as the fallow deer Dama dama [29].

Dimorphism of body size in marine mammals ranges from females that are larger than males (Hawaiian monk seal and Gervais' beaked whale at the extreme) to males that are much larger than females (sperm whale, southern elephant seal, and Hooker's sea lion at the extreme) $[17,30]$. Here we quantify the difference in terms of body length (Figure 15). The mating system is not known for all marine mammals, but when known, a malefemale length difference greater than about $16 \%$ ( 0.15 on a natural $\log$ scale) indicates a territorial or harem mating system [31]. Intense male-male competition is associated with marked sexual dimorphism and often spatial aggregation of females related to the accessibility of food and shelter. A male-female length difference less than $16 \%$ corresponds to a more generalized 'aquatic,' or 'dispersed', mating system, in which males have less opportunity to guard and monopolize females.

The male-female length difference of $12 \%$ ( 0.11 on a natural log scale) in Maiacetus indicates moderate sexual dimorphism and


Figure 15. Mating systems and sexual dimorphism in marine mammals. Sexual dimorphism of males and females is quantified for 105 species of cetaceans, pinnipeds, sirenians, and the sea otter, expressed as male-minus-female length in natural log units [17,30]. Territorial and harem systems that involve intense male-male competition and spatial aggregation of females have male-female dimorphism greater than about 0.15 (dashed line) [31]. Maiacetus inuus falls in the group of 'dispersed' mating systems with limited aggregation and limited male control of mating. Both cetaceans and pinnipeds span the entire range shown here, although mating systems are known for relatively few species [32].
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probably limited male-male competition for mates. Limited opportunity to monopolize mates suggests in turn that food and shelter were dispersed in protocetid habitats. This is corroborated by the geographically extensive but environmentally uniform shallow marine deposits of the Habib Rahi Formation, which were deposited on a broad, shallow marine shelf that would have provided little spatial aggregation of food or shelter.

Much remains to be learned about mating systems in cetaceans because their behavior is so difficult to study [32]. At present, body size dimorphism is itself sometimes used as a measure of 'contest competition' between males [17]. Maiacetus suggests that sexual dimorphism appeared early in cetacean evolution. Males were larger than females, but the difference was modest on the scale of dimorphism observed in marine mammals today.

## Conclusions

We describe exceptional specimens of a new early middle Eocene (47.5 Ma) protocetid whale, Maiacetus inuus, that include an adult female skeleton, a near term fetal skeleton, and an adult male skeleton. Maiacetus differs in size and proportions from younger, more derived basilosaurids such as Dorudon atrox (Figures 1, 12, 13), but it is close in size and similar in proportions to a composite skeleton of Rodhocetus balochistanensis/kasranii (Figure 13). Maiacetus retains many characteristics of its terrestrial forebears, including shearing cheek teeth with protocones on upper molars, a vertebral formula very close to that of primitive artiodactyls, a forelimb retaining mobile digits, a pelvis and hindlimb anchored on the vertebral column; and a double-pulley astragalus within the ankle. The feet of Maiacetus are not as elongated as those of Rodhocetus, indicating that Maiacetus may have been a slightly less specialized foot-powered swimmer (Figure 13).

Preservation of an intact near-term fetal skull and partial skeleton indicates that birth in early archaeocetes involved a single
calf that was born head-first as in land mammals, not tail-first as in living whales. This, in turn, indicates that birth almost certainly took place on land during this phase of early whale evolution. The presence of partially mineralized permanent first molars in the fetal skull indicates precocial development, which may have been a key life history trait in early whales facilitating the transition from land to sea. Sexual dimorphism in body and canine size are moderate, suggesting limited competition among males for mates.

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## Author Contributions

Performed the experiments: ISZ. Analyzed the data: PDG MuH WvK WJS BHS. Wrote the paper: PDG MuH WvK WJS BHS ISZ. Participated in field work: PDG MuH ISZ. Contributed geological interpretations: PDG ISZ MuH. Developed chronological framework: PDG MuH ISZ. Carried out functional analysis of skeletons: PDG WJS ISZ. Interpreted life history: PDG BHS.

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