



# Sexual Dimorphism and Body Reconstruction of a Hornless Rhinocerotid, *Plesiaceratherium gracile*, From the Early Miocene of the Shanwang Basin, Shandong, China

Xiaokang Lu<sup>1,2,3</sup>, Tao Deng<sup>2,4,5\*</sup>, Xiaoting Zheng<sup>6</sup> and Fuchang Li<sup>7</sup>

<sup>1</sup> Human Anatomy and Histology Department, Henan University of Chinese Medicine, Zhengzhou, China, <sup>2</sup> Key Laboratory of Vertebrate Evolution and Human Origins, Institute of Vertebrate Paleontology and Paleoanthropology, Chinese Academy of Sciences, Beijing, China, <sup>3</sup> State Key Laboratory of Palaeobiology and Stratigraphy, Nanjing Institute of Geology and Palaeontology, Chinese Academy of Sciences, Nanjing, China, <sup>4</sup> CAS Center for Excellence in Life and Paleoenvironment, Beijing, China, <sup>5</sup> University of Chinese Academy of Sciences, Beijing, China, <sup>6</sup> Tianyu Museum of Natural History, Pingyi, Shandong, China, <sup>7</sup> Paleontology Department, Museum of Linqu County, Linqu, China

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### \*Correspondence:

Tao Deng  
dengtao@ivpp.ac.cn

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We investigated the sexual dimorphism and reconstructed the body of *Plesiaceratherium gracile* based on the numerous new specimens from the Early Miocene Shanwang Basin, China. The sexual dimorphism of *P. gracile* is reflected by the size of both the lower incisor i2 [length male/female (M/F) ratio = 2.69, width M/F = 1.63] and the upper incisor I1 (length M/F = 1.63). Other mensural variations reveal their ranges greatly overlapping on both genders, with a low M/F ratio; hence, they cannot be interpreted as sexual dimorphism. The large-sized body and the strong sexual difference of incisors suggest that *P. gracile* is polygynous and has a solitary lifestyle. When we reconstructed the body of *P. gracile*, we found that the head-body length is 2,796–3,117 mm, the shoulder height is 1,476–1,627 mm, and the body weight is about 1,198 kg. All skeletons have the ventral edge of the thorax at the level of the olecranon of the ulna. The spinous processes of the posterior thoracic vertebrae and the lumbar vertebrae are always vertical, the metatarso-femur ratio and the tibio-femur ratio are high, and the two joint angles between the scapula, the humerus, and the radius are 102°–143° and 115°–160°, respectively, indicating that *P. gracile* has a mediportal limb and a horse-gallop running type. Furthermore, the limb-segment comparisons of *P. gracile* and other rhinoceroses demonstrate that the proximal long bones, including the scapula, the humerus, and the femur, are important for the body height, regardless of whether their body is large or small-sized; the distal parts of the limb, including the metacarpus and the metatarsus, are sensitive to body mass changes.

**Keywords:** *Plesiaceratherium*, Early Miocene, Shanwang Basin, Rhinocerotidae, Aceratheriinae, body size, sexual dimorphism

## INTRODUCTION

Intraspecific variation records nearly every aspects of individual difference within a species, the most interesting of which is sexual dimorphism. Pocock (1945) was the first to note that the secondary sexual character of the extant Asiatic rhinoceroses lies in the horns. Groves (1982) recognized more sexual differences in the body size and the skull of the Asiatic rhinoceroses, including several skull measures. Owen-Smith (1988) found that the gender of the African white rhinoceros could be judged based on the difference of the body and horn sizes. Dinerstein (1991) focused on the detailed sexual dimorphism of the greater one-horned rhinoceros, recording the difference of the mandible and the skull. With regard to the fossil rhinocerotids, many studies have revealed that sexual dimorphism lies in the presence of the horn or the size of the lower incisors (Osborn, 1898a,b; Peterson, 1920; Cerdeño and Sánchez, 2000; Mead, 2000; Deng, 2001, 2005; Muhlbachler, 2005, 2007; Chen et al., 2010). However, the sexual dimorphism of the skeleton in fossil rhinoceroses remains controversial. Loose (1975) stated that there is no significant sexual difference in the skull of African species and *Dicerorhinus sumatrensis*. Borsuk-Bialynicka (1973) mentioned that bimodality in the occiput width, maximum skull length, orbit-nuchal crest length, and orbit-nares length suggested the presence of sexual dimorphism in the skull of *Coelodonta antiquitatis*. Voorhies (1990) suggested that sexual dimorphism is evident in the ulna and radius dimensions of Barstovian *Teleoceras medicornutum* and Hemphillian *T. fossiger*, respectively. Muhlbachler (2007) recorded the skeleton of *Menoceras arikareense* and suggested that the greater degree of sexual bimodality of the nasal horn may relate to the small-sized body, but found no sexual difference in body size. Mead (2000) provided a detailed analysis on the skeleton of *Teleoceras major* and pointed out the cross section outline of the concerning long bones has the sexual difference. More recently, the gender of extinct rhinoceroses has been estimated by means of paleoproteomic analyses (Cappellini et al., 2019).

Mass estimation is different based on the equations from different measures and taxonomic levels (Mead, 2000; Qiu and Wang, 2007). Damuth (1990) investigated the linear relation between body weight and the measures of the head-body length and teeth. Janis (1990) focused on the measures of the skull and teeth. Scott (1990) further provided information on the length and width of limb bones. Before Damuth and MacFadden (1990) collected such papers about mass estimation, Legendre (1986) once tried to establish the linear relation between body weight and the crown area of M1, while Anderson et al. (1985) also noted the relation between the circumference of the long bones and the body weight. The equation based on these parameters has been supported by MacFadden and Hulbert (1990) and Mead (2000) for estimating the body weight of rhinoceroses. Cerdeño and Nieto (1995) once used the astragalus to estimate the body size; fortunately, we now have other, more reliable, measures. All these abovementioned equations were established based on data from higher taxa level, such as mammals, all ungulates, or perissodactyls and hyracoids. The earlier work of Freeman and King (1969) tried to establish several equations for the mass

estimation of the extant black rhinoceros based on the measures of the head-body length and the limb bones. In this study, we use such different equations to estimate the body weight of *Plesiaceratherium gracile*.

Size increase is a common feature in nearly all mammals, including the rhinocerotids, but the subfamily Aceratheriinae shows a clear tendency of size decrease (Heissig, 1989; Prothero and Schoch, 1989; Cerdeño and Nieto, 1995; Cerdeño, 1998; Prothero, 2005). *P. gracile*, from the Early Miocene, is a less specialized representative of Aceratheriinae, characterized by the moderately enlarged upper incisor I1, longer nasal bone, deeper nasal notch, slightly complicated cheek teeth, and longer and slender metapodials (Young, 1937; Yan, 1983; Yan and Heissig, 1986). Based on the ratio of height to length of the second molar (Fortelius et al., 2006), the cheek teeth of *P. gracile* are mesodont. The Shanwang Basin is famous for the well-preserved skeletons of *P. gracile* and other large mammals, which were produced from the Early Miocene Shanwang Formation (Young, 1937; Deng et al., 2003). In the Shanwang Formation, there are three fossil beds: the intermediate diatomaceous bed, the underlying sandstone bed, and the overlying sandstone bed. All well-preserved skeletons of *P. gracile* were recovered from the diatomaceous bed, while specimens from the underlying sandstone bed are very rare and always badly preserved (Yan et al., 1983; Deng et al., 2003). Numerous skeletons of *P. gracile* give us a chance to analyze sexual dimorphism, reconstruct the body, and provide new insights on the size evolution of the limb bones.

## MATERIALS AND METHODS

All specimens of *P. gracile* used in this study come from the diatomaceous shale of the Early Miocene Shanwang Basin, with well-preserved bones but with only one side exposed. Therefore, some measures listed in previous studies estimating sexual dimorphism could not be obtained, such as the circumference of the limb bones and the width of the symphysis (Mead, 2000; Chen et al., 2010). Fifty-two specimens are newly reported, except the skull STM 44-68, previously reported by Lu et al. (2016) (**Supplementary Table 1**). Because there were several very badly crushed skeletons, our study is based on the most accessible and informative specimens. To avoid age-related morphological changes, all specimens used in the analysis were of adult individuals, judged by the established criteria for estimating their age (Goddard, 1970; Hitchins, 1978; Hillman-Smith et al., 1986; Deng, 2001). The measurements are according to the criteria established by Guérin (1980) and Qiu and Wang (2007).

For each measure, we calculated the Student's *t*-test and conducted a descriptive analysis, including mean, maximum, minimum, standard deviation, coefficient of variation, kurtosis, skewness, and Shapiro-Wilk test. Following many previous studies, we provided the male/female (M/F) ratio for estimating the sexual difference, which is called dimorphic ratio (*dr*) for the means of each measure. The new specimens included several skeletons with normal posture, so we could measure the shoulder height, the hind-limb height, and the body length. Along with the measures of the skull, teeth, and limb bones, we reconstructed

the skeleton of *P. gracile*. Meanwhile, recognition of sexual dimorphism offered a chance to discuss the social behavior of *P. gracile*.

Body weight was estimated using previously established equations, based on length of the skull, head-body, M1, and m1 (Freeman and King, 1969; Damuth, 1990; Janis, 1990; Scott, 1990; Silva and Downing, 1995). As for the extant rhinoceroses, comparative data are from Guérin (1980), and head-body lengths are from different papers (Groves, 1972; Groves and Kurt, 1972; Laurie et al., 1983; Hillman-Smith and Groves, 1994).

For the limb bone, we postulate the locomotion type and compare limb elements between *P. gracile* and several rhinoceroses. The former was estimated based on the discussion of Gregory (1912) and Slijper (1946), and also the suggestion of Christiansen (2002). The results are determined through line and spine charts.

To avoid long tables, we have provided the abbreviations of the measures here, which are the same as in the text tables, **Supplementary Tables**, and figures.

Institutional abbreviations are IVPP OV, Vertebrate specimens housed in the Institute of Vertebrate Paleontology and Paleoanthropology (IVPP), Chinese Academy of Sciences, Beijing, China; S, prefix of the fossils from the Early Miocene Shanwang Basin housed in the Museum of Linqu County, Linqu, Shandong, China; SNGPM, Shanwang National GeoPark Museum, Linqu, Shandong, China; STM, Shandong Tianyu Museum, Pinyi, Shandong, China.

Abbreviations of parameters listed in the table of the descriptive analysis are min: minimum; n, number of mensural data; SD, standard deviation; cv, coefficient of variation; skew, skewness; skew.2se, significant test of skewness; kurt, kurtosis; kurt.2se, significant test of kurtosis; sw: Shapiro-Wilk test; *p*, significance of Student's *t*-test; d, dimorphic ratio, male mean/female mean.

Abbreviations for skull are DCAO, distance between occipital crest and lacrimal tubercle or anterior rim of orbit; DCPO, distance between occipital crest and postorbital process; DCN, distance between occipital crest and nasal tip; DOM3, distance between occipital condyle and M3; DON, distance between occipital condyle and nasal tip; DONN, distance between nasal notch and orbit; DONT, distance between orbit and nasal tip; DOP, distance between occipital condyle and premaxillary tip; HMC, height of mandible condyle; HZ, height of zygomatic arch between upper edge and lower edge posterior to the orbit; LM, maximum length of mandible; LN, distance between nasal tips and bottom of nasal notch; LS, length of symphysis; LZ, length of zygomatic arch from anterior edge below orbit to posterior edge above pseudoauditory meatus; WV, maximum width of vertical ramus of mandible.

Abbreviations for teeth are DP, deciduous upper premolars; I, upper incisors; i, lower incisors; L, length; M, upper molars; m, lower molars; P, upper premolars; p, lower premolars; W, ventral width.

Abbreviations for postcranial bones are BL, body length; C.V., cervical vertebra; DWMc3, distal width of Mc III; FFA, fore-limb first joint angle; FSA, fore-limb second joint angle; FLP1, maximum length of first phalanx of fore-limb; FLP2, maximum

length of second phalanx of fore-limb; FLP3, maximum length of third phalanx of fore-limb; FPWP1, maximum proximal width of first phalanx of fore-limb; FPWP2, maximum proximal width of second phalanx of fore-limb; FPWP3, maximum proximal width of third phalanx of fore-limb; HFA, hind-limb first joint angle; HLP1, maximum length of first phalanx of hind-limb; HLP2, maximum length of second phalanx of hind-limb; HLP3, maximum length of third phalanx of hind-limb; HPWP1, maximum proximal width of first phalanx of hind-limb; HPWP2, maximum proximal width of second phalanx of hind-limb; HPWP3, maximum proximal width of third phalanx of hind-limb; HSA, hind-limb second joint angle; LF, maximum length of femur; LH, maximum length of humerus; LMc3, maximum length of Mc III; LMT3, maximum length of Mt III; LR, maximum length of radius; LS, maximum length of scapula; LT, maximum length of tibia; L.V., lumbar vertebrae; Mc, metacarpus; Mt, metatarsus; NWMc3, narrowest width of Mc III; PWF, maximum proximal width of femur; PWH, maximum proximal width of humerus; PWMc3, maximum proximal width of Mc III; PWMt3, maximum proximal width of Mt III; PWR, maximum proximal width of radius; PWS, maximum proximal width of scapula; SH, shoulder height; TPW, maximum proximal width of tibia; T.V., thoracic vertebrae.

Abbreviations of body mass estimation are HBL, head and body length; SUML, antero-posterior length of upper molar (M1); SLML, antero-posterior length of lower molar (m1).

## RESULTS

### Sexual Dimorphism

*Plesiaceratherium* is a hornless rhinoceros, so the gender cannot be determined based on the horn difference, which is relatively evident on some horned rhinocerotids. Given that the size of incisors has long been recognized as reflecting sexual dimorphism, we first investigated this feature and found that the length of i2 is the most dimorphic character ( $dr = 2.69$ ), followed by its width ( $dr = 1.63$ ) (**Figures 1–3** and **Table 1**). All accessible specimens for measuring i2 were adult individuals, with moderately worn cheek teeth. The length of i2 has a wider range in the male. Concerning upper incisors, we noted the length measures of 10 I1 of the adult specimens, including two older females and one older male with highly worn M1. The measures of I1 between the two genders are widely separated, and the mean ratio is similar to that of the width of i2 ( $dr = 1.63$ ) (**Figure 3**, **Supplementary Figure 1**, and **Table 1**).

Meanwhile, based on the dimorphism of the incisors, we compared several non-mensural characters between two genders, such as the posterior position of the nasal notch and the presence of the mesostyle of P2. The posterior position of the symphysis is below dp1 or moved backward to p2, bearing an evident intraspecific variation but not being sexually significant. The presence of the first lower cheek teeth dp1 is variable and has no sexual difference or other non-mensural characters (**Supplementary Table 1**).

Among the cheek tooth measures, the lengths of P2 and M1 seem to have a significant Student's *t*-test value,

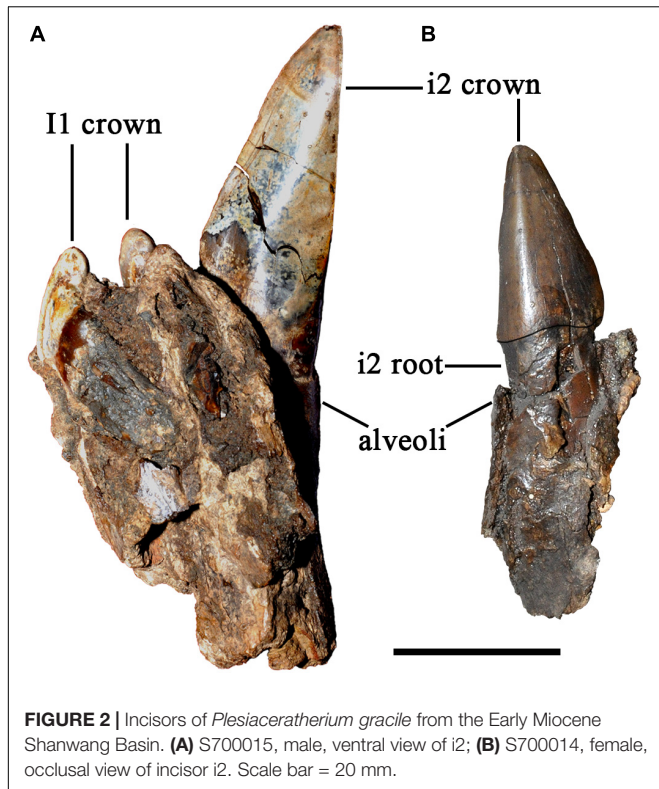


$p < 0.05$  (Table 2). However, the M/F ratio of these two measures is 1.09, much lower than the value of the incisors, and the lengths between two genders are greatly overlapping.

On the skull, the mean of the female length is longer than that of the male in several measures, including DCN (dr = 0.97), LN (DR = 0.93), DONN (dr = 0.89), DONT (dr = 0.95), and LZ (dr = 0.95) (Table 3). The ranges of these measures between two genders are overlapping, and their  $t$ -test values are not significant. On the other hand,

the height of the zygomatic arch varies more widely in the male specimens and has a relatively higher sex ratio (dr = 1.06) (Table 3). Most of these measures are in a normal distribution and have a low value of skewness and kurtosis. Meanwhile, the  $t$ -test value is not significant. The measures of the symphysis on both genders are nearly totally overlapping (Table 4).

Like measures of the skull, the limb bones of the female specimens always have a larger mean than do the male. However, the scapula in the male is longer than that of the female



( $dr = 0.96$ ), and the male has a relatively wider range, 172–205 mm. Furthermore, the female has a longer femur and tibia, but the ratio is lower. The length of the Mt III in the male is slightly longer than in the female ( $dr = 1.01$ ), and the proximal width has a higher M/F ratio, up to 1.2. It is noteworthy that all mean measures of proximal, narrowest, and distal widths of the long bones in the male specimens are larger than those of the female (Table 5).

### Body Size

The direct measure of seven articulated skeletons of *P. gracile* revealed that the body length from the atlas to the ischial tuberosity is 2,148–2,469 mm (Supplementary Table 5). The mean skull length is 649 mm (Table 3) and mean of the head-body length of *P. gracile* is 2,991 mm, with a range of 2,796–3,117 mm. We measured different parts of the spinal column of the different individuals and composed a complete vertebral series. Based on the nearly undeformed pelvis of S 700018, the inclination of the ala-acetabulum axis of the coxa relative to the horizontal plane is about  $56.4^\circ$ . Combining such different parts of the body, including the 7 C.V., 18 T.V., 5 L.V., and the pelvis, the estimated body length is about 2,162 mm and the head-body length (considering the mean head length) is about 2,811 mm, within the range of the direct measure mentioned above (Supplementary Table 6). The spinous process of the T.V. is inclined posteriorly, but sometimes it is vertical in the last two T.V. The L.V. also have a vertical spinous process. In the medium-sized individual STM 44-62, the linear length of the rib is about 830 mm. Regardless of the size, all skeletons have the ventral edge of the thorax flushing with the olecranon of the ulna.

( $dr = 1.04$ ), with the latter having a higher kurtosis. The humerus and the radius have the same M/F ratio ( $dr = 0.98$ ) and a lower cv. The M/F ratio of the Mc III is more evident

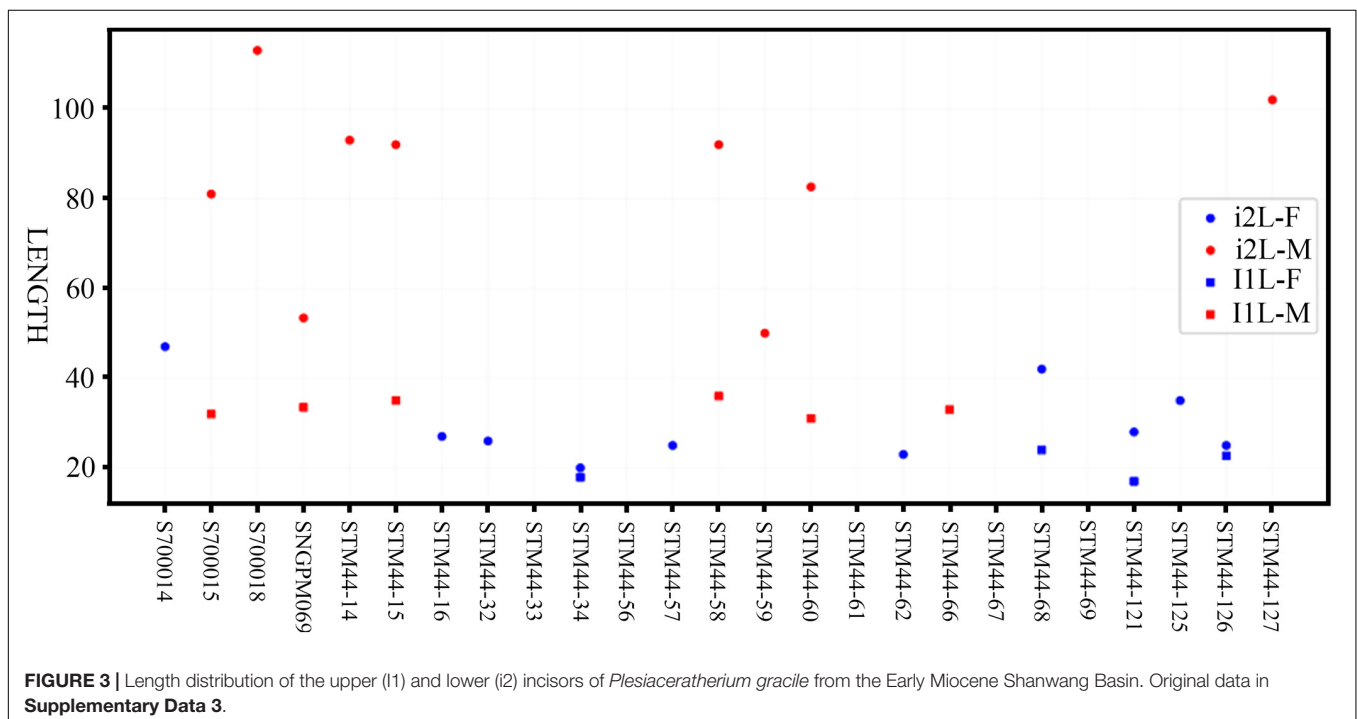


FIGURE 3 | Length distribution of the upper (i1) and lower (i2) incisors of *Plesiaceratherium gracile* from the Early Miocene Shanwang Basin. Original data in Supplementary Data 3.

**TABLE 1** | Statistic of incisor measures of *Plesiaceratherium gracile*.

Measure	Sex	n	Min	Max	Mean	SD	cv	Skew	Skew.2se	Kurt	Kurt.2se	sw	p	dr
L i2	F	10	20	47	29.8	8.73	0.29	0.83	0.6	-0.88	-0.33	0.86	>0.05	2.69
	M	9	50	113	84.33	20.92	0.25	-0.49	-0.34	-1.18	-0.42	0.9		
W i2	F	7	14	28	24.14	4.91	0.2	-1.11	-0.7	-0.29	-0.09	0.79	< 0.05	1.63
	M	5	30	56	40.8	12.28	0.3	0.27	0.15	-2.18	-0.55	0.83		
L I1	F	4	17	24	20.5	3.51	0.17	0	0	-2.38	-0.45	0.86	< 0.01	1.63
	M	6	31	36	33.5	1.87	0.06	0	0	-1.8	-0.52	0.98		

**TABLE 2** | Statistic of teeth measures of *Plesiaceratherium gracile*.

Measure	Sex	n	Min	Max	Mean	SD	cv	Skew	Skew.2se	Kurt	Kurt.2se	sw	p	dr
DP1	F	11	14	31	23.73	4.78	0.2	-0.53	-0.4	-0.71	-0.28	0.94	>0.05	1.05
	M	6	23	26	24.83	1.33	0.05	-0.18	-0.11	-2.05	-0.59	0.81		
P2	F	11	26	34	31	2.49	0.08	-0.42	-0.32	-0.98	-0.38	0.93	< 0.05	1.09
	M	7	31	39	34	2.71	0.08	0.6	0.38	-1.03	-0.33	0.91		
P3	F	12	32	45	36.92	3.7	0.1	0.75	0.59	-0.4	-0.16	0.93	>0.05	0.99
	M	9	32	42	36.89	3.14	0.09	-0.04	-0.03	-1.19	-0.43	0.97		
P4	F	12	36	44	38.75	2.18	0.06	0.97	0.76	0.23	0.09	0.9	>0.05	1.05
	M	10	37	44	40.7	2.41	0.06	-0.2	-0.15	-1.59	-0.6	0.93		
M1	F	12	37	50	44	4.41	0.1	-0.08	-0.06	-1.4	-0.57	0.95	< 0.05	1.09
	M	10	42	52	48	2.87	0.06	-0.61	-0.44	-0.58	-0.22	0.94		
M2	F	12	46	54	49.67	2.5	0.05	-0.07	-0.05	-1.21	-0.49	0.94	>0.05	1.02
	M	10	49	53	50.8	1.23	0.02	0.34	0.24	-1.26	-0.47	0.92		
M3	F	10	45	52	48.7	2.41	0.05	-0.37	-0.27	-1.39	-0.52	0.92	>0.05	1
	M	7	48	50	48.86	0.69	0.01	0.11	0.07	-1.24	-0.39	0.84		
DP1-M3	F	11	229	277	252.55	16.97	0.07	0	0	-1.59	-0.62	0.93	>0.05	1.04
	M	9	229	279	261.67	17.08	0.07	-0.68	-0.48	-1.09	-0.39	0.88		
DP1-P4	F	11	112	140	125.55	8.62	0.07	0.04	0.03	-1.26	-0.49	0.98	>0.05	1.05
	M	9	116	143	131.89	9.28	0.07	-0.5	-0.35	-1.47	-0.53	0.9		
M1-M3	F	11	116	144	130.91	10.55	0.08	-0.13	-0.1	-1.71	-0.67	0.9	>0.05	1.03
	M	10	123	142	135.3	6.99	0.05	-0.48	-0.35	-1.54	-0.58	0.86		
p2	F	9	23	33	28.22	2.91	0.1	-0.2	-0.14	-0.92	-0.33	0.96	>0.05	1.02
	M	8	24	33	28.75	3.45	0.12	0.15	0.1	-1.81	-0.61	0.88		
p3	F	9	30	38	33.78	2.28	0.07	0.24	0.17	-0.71	-0.25	0.96	>0.05	1.01
	M	9	31	36	33.89	1.9	0.06	-0.25	-0.17	-1.81	-0.65	0.87		
p4	F	9	33	39	35.89	1.76	0.05	0.27	0.19	-0.88	-0.31	0.92	>0.05	1
	M	9	32	39	35.89	2.32	0.06	-0.31	-0.22	-1.33	-0.47	0.94		
m1	F	9	38	42	40.33	1.22	0.03	-0.2	-0.14	-0.76	-0.27	0.85	>0.05	1
	M	8	34	45	40.25	3.24	0.08	-0.44	-0.29	-0.62	-0.21	0.94		
m2	F	7	40	44	41.43	1.62	0.04	0.41	0.26	-1.7	-0.54	0.86	>0.05	0.98
	M	8	36	43	40.38	2.67	0.07	-0.43	-0.29	-1.67	-0.56	0.87		
m3	F	9	41	45	42.33	1.32	0.03	0.61	0.42	-0.8	-0.29	0.87	>0.05	0.98
	M	9	36	47	41.44	4.42	0.11	-0.09	-0.07	-1.94	-0.69	0.87		
p2-m3	F	7	209	261	236.29	21.92	0.09	-0.06	-0.04	-2.06	-0.65	0.87	>0.05	0.96
	M	9	139	257	227.11	35.86	0.16	-1.51	-1.05	1.15	0.41	0.75		
p2-p4	F	7	90	124	107.14	12.95	0.12	-0.06	-0.04	-1.91	-0.6	0.93	>0.05	1.05
	M	9	101	128	112.33	9.33	0.08	0.18	0.12	-1.49	-0.53	0.94		
m1-m3	F	8	117	149	129	10.41	0.08	0.69	0.46	-0.94	-0.32	0.92	>0.05	1
	M	10	118	137	128.6	6.7	0.05	-0.29	-0.21	-1.55	-0.58	0.93		

Because sample size must be at least 3, we exclude measures of dp1.

Based on five less distorted skeletons, the shoulder height from the tip of the T.V. to the distal extremity of the fore-limb is about 1,491.8 mm, with a range of 1,438–1,603 mm,

and the height of the hind-limb is about 1,526 mm, with a range 1,476–1,627 mm (**Supplementary Table 5**). Given that all skeletons have curled bodies as a result of drowning in the

**TABLE 3** | Statistic of measures of skull for adult *Plesiaceratherium gracile*.

Measure	Sex	n	Min	Max	Mean	SD	cv	skew	Skew.2se	Kurt	Kurt.2se	sw	sw.p	p	dr
DOP	F	9	522	692	632.78	59.64	0.09	-0.92	-0.64	-0.93	-0.33	0.79	0.02	>0.05	1.01
	M	5	631	645	638.00	5.57	0.01	0.04	0.02	-1.93	-0.48	0.97	0.91		
DON	F	7	524	692	645.86	55.97	0.09	-1.36	-0.86	0.29	0.09	0.71	0.00	>0.05	0.99
	M	5	625	648	636.20	9.36	0.01	0.08	0.04	-1.99	-0.50	0.97	0.86		
DCN	F	7	622	695	659.00	25.41	0.04	0.12	0.07	-1.55	-0.49	0.95	0.75	>0.05	0.97
	M	9	610	696	640.00	31.58	0.05	0.86	0.60	-1.04	-0.37	0.80	0.02		
LN	F	8	190	274	247.13	26.22	0.11	-1.07	-0.71	0.02	0.01	0.85	0.09	>0.05	0.93
	M	9	215	274	230.44	18.56	0.08	1.35	0.94	0.51	0.18	0.77	0.01		
DCPO	F	4	271	305	286.25	17.35	0.06	0.07	0.03	-2.36	-0.45	0.84	0.19	>0.05	0.97
	M	4	254	297	276.25	17.65	0.06	-0.09	-0.05	-1.89	-0.36	0.98	0.88		
DCAO	F	5	325	369	343.60	19.99	0.06	0.28	0.15	-2.15	-0.54	0.85	0.21	>0.05	0.98
	M	4	320	352	337.75	15.76	0.05	-0.10	-0.05	-2.31	-0.44	0.87	0.31		
DONN	F	7	74	93	79.86	7.43	0.09	0.66	0.41	-1.37	-0.43	0.82	0.06	>0.05	0.89
	M	5	61	86	71.40	9.61	0.13	0.39	0.22	-1.62	-0.40	0.96	0.78		
DOM3	F	8	204	298	270.75	33.06	0.12	-1.00	-0.67	-0.67	-0.23	0.79	0.02	>0.05	1.03
	M	5	264	292	279.40	12.24	0.04	-0.14	-0.08	-2.09	-0.52	0.92	0.51		
DONT	F	5	286	334	315.40	18.05	0.06	-0.63	-0.34	-1.36	-0.34	0.89	0.36	>0.05	0.97
	M	4	283	330	304.75	23.30	0.08	0.06	0.03	-2.35	-0.45	0.87	0.29		
LZ	F	5	237	338	266.20	41.56	0.16	0.90	0.49	-1.13	-0.28	0.76	0.04	>0.05	0.95
	M	7	232	304	252.71	23.51	0.09	1.36	0.86	0.33	0.10	0.69	0.00		
HZ	F	5	54	65	61.40	4.39	0.07	-0.79	-0.44	-1.27	-0.32	0.84	0.15	>0.05	1.06
	M	8	57	71	66	6.00	0.09	-0.09	-0.06	-2.03	-0.68	0.83	0.07		

**TABLE 4** | Statistic of mandible for adult *Plesiaceratherium gracile*.

Measure	Sex	n	Min	Max	Mean	SD	cv	Skew	Skew.2se	Kurt	Kurt.2se	sw	sw.p	p	dr
LM	F	10	474	582.2	550.22	-1.32	-0.96	0.93	0.35	0.83	0.04	-1.32	-0.96	>0.05	0.99
	M	9	522	576	546	0.35	0.24	-1.09	-0.39	0.98	0.94	0.35	0.24		
WV	F	7	126	192.3	155.04	0.29	0.19	-1.03	-0.32	0.93	0.57	0.29	0.19	>0.05	1
	M	7	146	164	155.71	-0.32	-0.20	-1.86	-0.59	0.87	0.19	-0.32	-0.20		
HMC	F	6	230	256	242.33	0.02	0.01	-2.24	-0.64	0.79	0.05	0.02	0.01	>0.05	0.97
	M	6	216	253	234.83	-0.04	-0.02	-1.63	-0.47	0.98	0.94	-0.04	-0.02		
LS	F	8	118	161.5	137.44	15.61	0.11	0.19	0.13	-1.66	-0.56	0.95	0.67	>0.05	1.01
	M	5	112	168	136.6	0.35	0.24	-1.09	-0.39	0.98	0.94	0.35	0.24		

lake, the shoulder height probably has been underestimated, and it is also difficult to precisely predict the joint angle between the adjacent long bones. Among numerous skeletons, only STM 44–60 has a nearly straight fore-limb; its shoulder height is 1,603 mm, and the angle between the scapula and the humerus is 143° and between the humerus and the radius is 160°. In the slightly curled body S 700018, the first angle of the fore-limb is 128° and the second is 131°. In the most curled skeleton STM 44–61, the first angle of the fore-limb is 102°, the second is 115°; the angle of the hind-limb between the coxa and the femur is 101.2° and between the femur and the tibia is 91.8°. We could not find a skeleton with an intact hind-limb, but S 700018 is just slightly curled and could be used for body reconstruction: the first angle is 133.1° and the second is 137.8°. All accessible skeletons show a slightly longer hind-limb although they are deformed in a different degree. In the best-preserved S 700018, the hind-limb from the tip of the sacrum to the distal extremity is 1,532 mm, slightly longer than the fore-limb.

The reconstruction of the joint angle of the fore-limb is based on STM 44–60, while that of the hind-limb is based on the S 700018. Given that the curled body and limb would inevitably reduce the body height, we used the larger value for the skeleton reconstruction (**Figure 4**).

Body weight based on the mean head-body length (HBL) was calculated as 832.1 kg following the equations for mammals and as 996.6 kg following the equations for all ungulates (**Table 6**). Using the equations for the black rhinoceroses, the estimation based on the mean HBL is 1,198.1 kg, and that from the largest HBL is 1,362.8 kg. Using Janis's (1990) equation for perissodactyls and hyracoids, the estimation based on the total skull length (TSL) is 1,130.3 kg. According to the weight record of the extant rhinoceroses, using the equations for all ungulates established by Damuth (1990), the estimated weight based on the molar size is high and even exceeded the heaviest weight records; but, for *P. gracile*, using Janis's (1990) equations, the more reasonable molar-based estimations provided a range of 1,092.9–2,244.5 kg.

**TABLE 5** | Statistic of limb measures for adult *Plesiaceratherium gracile*.

	Sex	n	Min	Max	Mean	SD	cv	Skew	Skew.2se	Kurt	Kurt.2se	sw	p	dr
LS	F	3	371	424	393.67	27.32	0.07	0.26	0.11	-2.33	0.00	0.94	> 0.05	1.04
	M	4	382	432	411.00	21.69	0.05	-0.34	-0.17	-1.97	-0.38	0.95		
LH	F	7	372	420.1	389.44	16.40	0.04	0.69	0.44	-0.96	-0.30	0.91	> 0.05	0.98
	M	9	349	404	380.22	18.40	0.05	-0.22	-0.15	-1.36	-0.49	0.94		
PWH	F	4	146	181	161.75	15.00	0.09	0.22	0.11	-2.01	-0.38	0.98	> 0.05	1.08
	M	4	162	186	174.00	9.83	0.06	0.00	0.00	-1.89	-0.36	0.98		
LR	F	7	335	373.9	356.03	12.74	0.04	-0.10	-0.06	-1.22	-0.38	0.93	> 0.05	0.98
	M	9	322	363	347.67	11.72	0.03	-0.86	-0.60	-0.02	-0.01	0.90		
LMc3	F	6	180	209.5	194.92	11.84	0.06	0.22	0.13	-1.85	-0.53	0.87	> 0.05	0.96
	M	11	172	205.2	186.47	8.78	0.05	0.53	0.40	-0.31	-0.12	0.94		
PWMc3	F	6	37	52	46.17	5.64	0.12	-0.53	-0.31	-1.55	-0.44	0.92	> 0.05	1.08
	M	11	45	57	49.73	3.80	0.08	0.32	0.24	-1.07	-0.42	0.94		
NWMc3	F	4	37	40	38.25	1.26	0.03	0.42	0.21	-1.82	-0.35	0.89	> 0.05	1.1
	M	7	35	46	42.14	4.34	0.10	-0.68	-0.43	-1.52	-0.48	0.81		
DWMc3	F	5	51	62	54.80	4.27	0.08	0.79	0.43	-1.24	-0.31	0.83	> 0.05	1.09
	M	8	56	66	59.75	3.41	0.06	0.53	0.35	-1.25	-0.42	0.91		
FLP1	F	4	33	40	37.75	3.20	0.08	-0.70	-0.35	-1.72	-0.33	0.75	> 0.05	1.1
	M	9	34	52	41.67	5.36	0.13	0.47	0.33	-0.81	-0.29	0.94		
FLP2	F	4	20	28	23.25	3.95	0.17	0.18	0.09	-2.24	-0.43	0.86	> 0.05	1.13
	M	9	21	31	26.22	3.67	0.14	-0.27	-0.19	-1.61	-0.57	0.91		
FLP3	F	5	32	43	38.40	4.39	0.11	-0.37	-0.20	-1.81	-0.45	0.94	> 0.05	1.04
	M	8	36	45	39.75	2.92	0.07	0.40	0.26	-1.18	-0.40	0.97		
LF	F	8	453	498	480.05	18.05	0.04	-0.48	-0.32	-1.73	-0.58	0.84	> 0.05	0.97
	M	8	429	490	467.38	20.63	0.04	-0.66	-0.44	-1.16	-0.39	0.90		
PWF	F	5	124	187	167.20	24.91	0.15	-0.91	-0.50	-1.10	-0.27	0.78	> 0.05	1.02
	M	4	146	196	170.75	21.56	0.13	0.02	0.01	-2.08	-0.40	0.99		
LT	F	6	343.8	368	354.30	9.20	0.03	0.25	0.15	-1.80	-0.52	0.93	> 0.05	0.99
	M	8	325	368	350.46	14.93	0.04	-0.42	-0.28	-1.38	-0.47	0.91		
LMt3	F	5	169	184	177.20	6.69	0.04	0.00	0.00	-2.09	-0.52	0.89	> 0.05	1.01
	M	8	172	187	178.38	5.13	0.03	0.26	0.17	-1.46	-0.49	0.96		
PWMt3	F	4	35	45	38.25	4.57	0.12	0.68	0.33	-1.73	-0.33	0.79	> 0.05	1.2
	M	5	37	52	45.80	8.04	0.18	-0.29	-0.16	-2.25	-0.56	0.71		
HLP1	F	3	39	45	43.00	3.46	0.08	-0.38	-0.16	-2.33	0.00	0.75	> 0.05	1.04
	M	5	41	51	44.80	3.90	0.09	0.55	0.30	-1.49	-0.37	0.89		
HLP2	F	3	26	45	32.33	10.97	0.34	0.38	0.16	-2.33	0.00	0.75	> 0.05	0.96
	M	4	26	37	31.00	4.55	0.15	0.24	0.12	-1.88	-0.36	0.96		

## DISCUSSION

### Sexual Dimorphism and Behavior

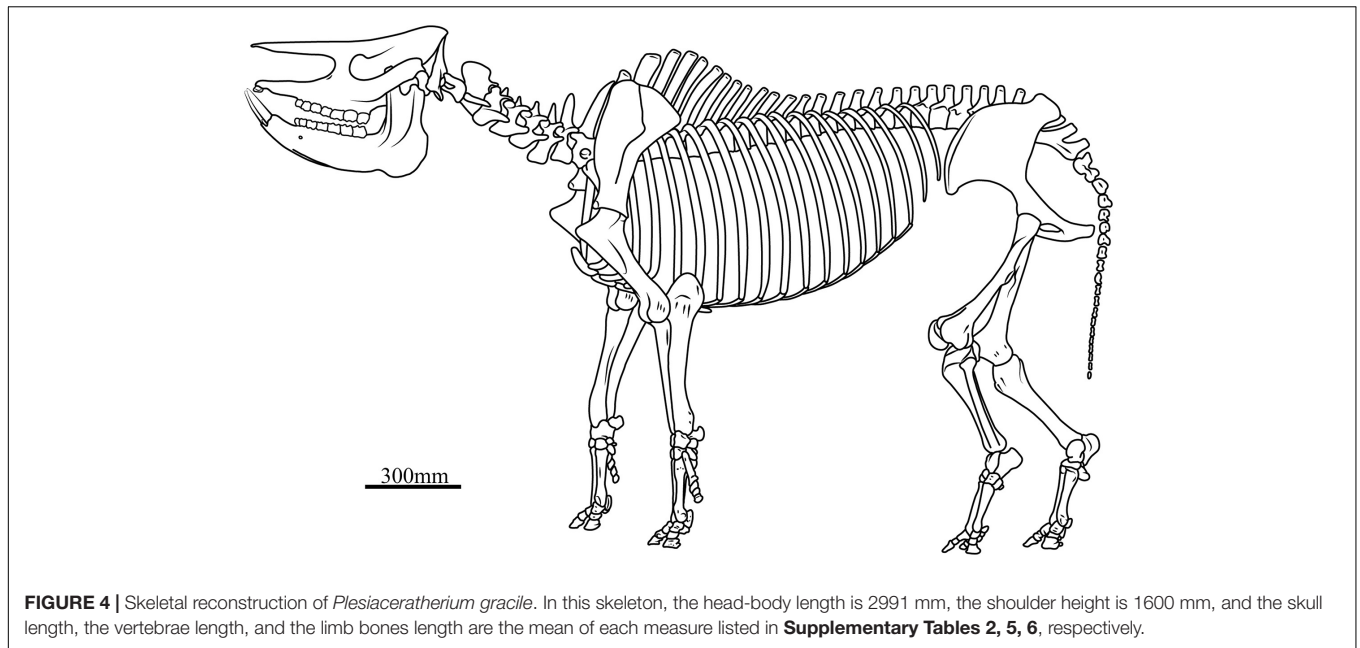
#### Sexual Dimorphism

The size difference of the lower incisor i2 is the strongest sexual dimorphic character in rhinocerotids. The enlarged tusk i2 has occurred since the Eocene rhinocerotids *Teletaceras* and *Trigonias* (Prothero and Schoch, 1989) and was specialized to a great level in the Oligocene rhinocerotid *Mesaceratherium* and *Aprotodon* (Beliajeva, 1954; Borissiak, 1954; Heissig, 1969; Qiu and Xie, 1997; Deng, 2013). The sexual difference of the incisor i2 has been demonstrated in many Neogene rhinocerotids, *Menoceras*, *Aphelops*, *Alicornops*, *Chilotherium*, and *Teleoceras* (Osborn, 1898a,b; Cerdeño and Sánchez, 2000; Mead, 2000; Muhlbachler, 2005, 2007; Chen et al., 2010). The measures of *P. gracile* demonstrated that not only the tusk i2 but also the

upper incisor I1 is sexually different and almost does not involve size overlapping between both genders (Table 1). This has not yet been noted in other rhinoceroses.

In our analysis, measures of the skull and limbs of *P. gracile* show a clear possibility that the females have a longer skull and body than do the males, which have a more robust skull and body. However, the M/F ratio of such measures is much lower than that of the incisors; more importantly, measures between the two genders are greatly overlapping. Mead (2000) listed many significant sexual dimorphisms in limbs of *Teleoceras major*, especially the cross-section of the radius (dr = 1.34) and the femur (dr = 1.19), with widely separated ranges. Chen et al. (2010) also suggested that the male specimens are larger than the female in mean values of the width of the mastoid process on the occipital area, and both genders have a nearly totally separated range. Additionally, Chen et al. (2010) revealed that the width of the





**TABLE 6 |** Mass estimation of *Plesiaceratherium gracile* and comparisons with extant rhinos (weight/kg).

	<i>P. gracile</i>		<i>D. bicornis</i>		<i>C. simum</i>		<i>D. sumatrensis</i>		<i>R. unicornis</i>	
	Value	Weight	Value	Weight	Value	Weight	Value	Weight	Value	Weight
HBL <sup>^</sup>	2991	730.4	2885	651.7	3666	1389.4	2767	571.1	4120	2009.4
HBL <sup>*</sup>	2991	863.2	2885	761.3	3666	1753.2	2767	658.37	4120	2632.4
HBL <sup>**</sup>	2.991	1198.1	2.885	1070.4	3.666	2261.5	2.767	939.6	4.12	3256.1
TSL <sup>†</sup>	64.83	1130.3	58.42	848.8	79.68	1993.4	53.98	682.9	67.49	1262.5
SUML <sup>^</sup>	46.11	2244.5	54.48	3758.1	55.54	3988.7	47.94	2531.3	53.25	3502.0
SLML <sup>^</sup>	40.32	1710.6	46.95	2746.5	49.22	3180.9	37.86	1406.5	46.75	2710.3
SUML <sup>†</sup>	4.611	1361.4	5.448	2208.4	5.554	2335.3	4.794	1524.1	5.325	2066.9
SLML <sup>†</sup>	4.032	1092.9	4.695	1728.3	4.922	1992.2	3.786	904.3	4.675	1706.2
LH <sup>#</sup>	38.425	1049.8	41.04	1313.4	41.82	1400.3	36.43	875.6	46.09	1949.3
LR <sup>#</sup>	35.184	378.2	37.27	445.6	38.65	494.2	30.36	248.6	39.2	514.5
LMc3 <sup>#</sup>	19.07	99.2	17.68	85.7	18.46	93.2	16.49	74.9	20.91	118.5
PWMc3 <sup>#</sup>	4.795	257.6	6.107	488.9	6.854	663.7	5.333	341.4	7.116	733.0
LF <sup>#</sup>	47.37	848.7	46.52	796.8	50.8	1082.8	42.1	562.6	55.4	1464.8
LT <sup>#</sup>	35.238	232.4	34.4	213.2	37.41	287.9	31.06	147.9	40.71	389.7
LMT3 <sup>#</sup>	17.78	71.2	15.7	54.6	16.84	63.4	14.92	49.0	19.26	84.5
PWMT3 <sup>#</sup>	4.203	253.2	4.891	388.7	5.58	564.2	4.78	364.3	6.13	735.9

Equation of <sup>^</sup> from Damuth (1990), measure value mm; <sup>\*</sup> from Silva and Downing (1995), measure value mm; <sup>\*\*</sup> from Freeman and King (1969), measure value meter; <sup>†</sup> from Janis (1990), measure value cm; <sup>#</sup> from Scott (1990), measure value cm.

mandibular symphysis that related with the size of i2 is sexually different, with moderately overlapping ranges. However, studies on the extant rhinoceroses show a wide overlap between genders on relevant skull measures, such as the occipital-nasal length or the nasal width in *C. simum*, *R. unicornis*, and *D. sumatrensis* (Heller, 1913; Guérin, 1980; Groves, 1982). Owen-Smith (1988) determined the gender of the African white rhinoceros mainly based on the size difference of the horn, and then postulated the sexual difference of the body size.

It remains controversial whether a size difference between two genders extensively exists in rhinoceroses. The most evident

sexual dimorphism in rhinoceroses is reflected in the horns, incisors, or features directly related to them, such as the nasal and symphysis widths. The sexual dimorphism should be easily identified independently and also be as unambiguous as possible. In *P. gracile*, our study reveals that sexual dimorphism is easily recognized from the size of the incisors (i2 and I1) but not from the size of the skull and limb bones.

### Social Behavior

Trivers (1972) suggested that the stronger the dimorphism in a species, the more likely it is to be polygynous. The

extant *R. unicornis*, *D. sumatrensis*, and *C. simum* have a great degree of sexual dimorphism in the horn or incisor size and are polygynous (Pocock, 1946; Owen-Smith, 1988; Dinerstein, 1991; Hillman-Smith and Groves, 1994). *Chilotherium wimani* and *Teleoceras* are two advanced rhinoceroses with very large i2, whose dimorphic ratio is 1.73 and 2.02, respectively (Mead, 2000; Chen et al., 2010). In *P. gracile*, the lengths of the lower incisor i2 do not overlap between the two genders. Such a difference is not weaker than that of *Ch. wimani* and *Teleoceras*, indicative of the presence of polygyny.

From the behavioral and ecological perspective, Cerdeño (1998) established three anatomic types of rhinoceroses: small and cursorial, medium-large size with intermediate gracility, and large and strongly brachypodial. The gregarious lifestyle of the small and cursorial species has an advantage in protecting them from predators and ensuring the continuation of the species (Cerdeño and Nieto, 1995). Size increase enabled them to defend themselves against predators and occasionally constitute small groups. The extant rhinos have a large-sized body, and they are all solitary or form “small groups” of mother-offspring units (Groves, 1972; Groves and Kurt, 1972; Laurie et al., 1983; Hillman-Smith and Groves, 1994). *P. gracile* belongs to the second type, having the advantage of living a solitary life.

However, the large number of *P. gracile* specimens from the Early Miocene Shanwang Basin could be interpreted as a consequence of their gregarious social behavior. In this basin, the badly crushed fragments of *P. gracile* from the sandstone bed are rare, but the well-preserved skeletons from the diatomaceous bed are numerous (Yan et al., 1983; Deng et al., 2003). In fact, the extensive amount of rhinoceros material could not be due to an abnormal natural catastrophic event; the alternative possibility is that they drowned in the lake progressively through time. On the other hand, the specimens from the sandstone beds indeed were deposited through a normal burial process, but this is rare in the Shanwang Formation of the Shanwang Basin, indicative of a small number of *P. gracile* living around the deposition area. After all, mass mortality resulting from an abnormal natural catastrophe would generate more specimens from the sandstone beds than from the diatomaceous, as is the case of the large number of *Chilotherium* specimens from the Linxia Basin in western China, where an extensive drought occurred during the Late Miocene and *Chilotherium* remains formed many lentoid fossil concentrations (Liang and Deng, 2005). It is clear that the number of skeletons of *P. gracile* from the Shanwang Basin has nothing to do with a gregarious way of life.

## Postcranial Bone Analysis

### Locomotion Adaptation

Slijper (1946) suggested that the spinous process of the T.V. and L.V. inclines to a different direction in mammals that have a different running type, including the leap gallop and the horse gallop. In equids, in order to perform the “horse gallop,” the longissimus dorsi is very developed, and the spinous processes

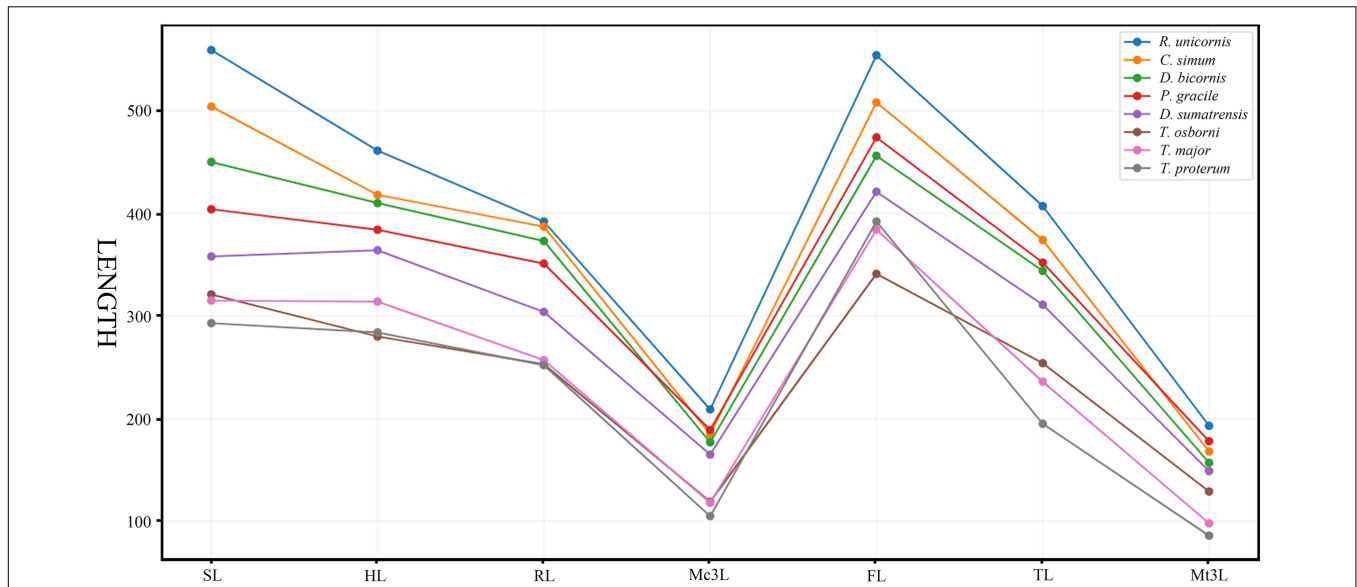
of the posterior 4–5 T.V. and the L.V. incline forward to forming the most efficient angle with the longissimus dorsi (Sisson, 1953). In the skeleton reconstruction of the primitive rhinoceroses *Trigonias* and *Juxia*, the posterior 4–5 T.V. and the L.V. are vertical, still having the ability of “horse gallop” (Scott, 1941; Qiu and Wang, 2007). In the extant *R. unicornis* (IVPP OV 1383) and *C. simum*, the spinous processes of the posterior T.V. and most of the L.V. are vertical or inclined posteriorly, and the running efficiency of the longissimus dorsi is declined, indicative of probably weakened running ability (Slijper, 1946; Groves, 1972). In the skeleton of *P. gracile*, the spinous processes of all T.V. are inclined anteriorly, or only the last two T.V. and the L.V. have a vertical spinous process (Figure 4). Thus, we conclude that *P. gracile* would have a horse gallop locomotion style and a running ability that is close to that of *Trigonias* and *Juxia*.

Gregory (1912) divided mammals into four types based on locomotion: cursorial, subcursorial, mediportal, and graviportal, using the limb segments ratio to estimate the locomotion type, namely the metatarso-femoral ratio and tibio-femoral ratio. He considered the four extant rhinoceroses as mediportal and *Teleoceras* as graviportal. According to this hypothesis, *P. gracile* would be mediportal, with values of 0.37 and 0.49 for the respective ratios, close to those of four extant rhinoceroses, but higher than those of *Teleoceras proterum*, which has a massive body and graviportal limb (Supplementary Table 7).

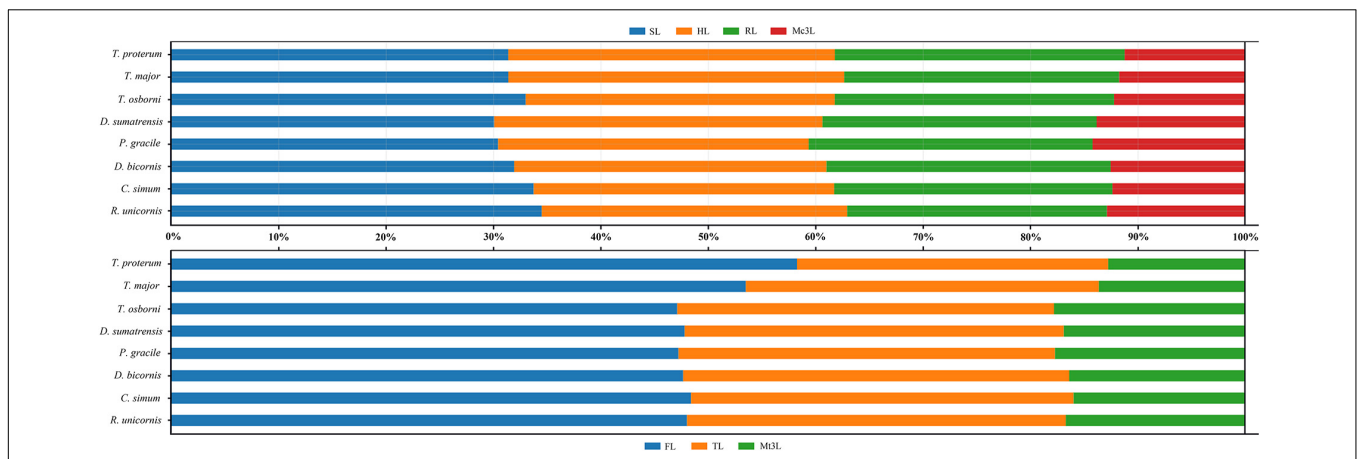
The bent of the limb joint in horses or other cursorial mammals is correlated with a bounding gallop or trotting gait (Gregory, 1912). Graviportal ungulates have evolved a vertical shaft to transmit the body weight and enlarge the stride (Gregory, 1912). In the graviportal rhinoceros *Teleoceras* from the late Miocene, the first and second angles of the fore-limb was found to be 139° and 160°, respectively, which supports this hypothesis (Osborn, 1898a; Gregory, 1912). In the mediportal large-sized rhinoceros *C. simum*, the first and the second angles of the fore-limb was 123° and 135°, respectively (Groves, 1972). *P. gracile* is close to the extant *C. simum* in terms of the joint angle of the long bones, being a mediportal type.

### Limb Length

The Early Miocene *P. gracile* has a simple occlusal pattern of upper cheek teeth, moderately enlarged incisors, and long and narrow metapodials; it is less specialized than the Late Miocene species of *Chilotherium* and *Teleoceras* (Young, 1937; Yan, 1983; Yan and Heissig, 1986). To investigate the condition of the limb segments in *P. gracile*, we compared the measures and the proportions of the limb bones with the primitive rhinoceros *Trigonias osborni* from the Paleogene, the massive *Teleoceras major* and *T. proterum* from the Late Miocene, and the four extant rhinoceroses. The result reveals an obvious tendency of size increase from *Tr. osborni* to *R. unicornis* in the length of the limb, and each limb segment contributes to this elongation. The proximal long bones of the fore-limb show a similar increasing tendency during the progress of the body size increase among these rhinoceroses (Figure 5). However, the proportion of each limb segment relative to the limb length is different, and



**FIGURE 5 |** Comparative length of the limb bones in *Plesiaceratherium gracile* and other rhinos. Original data in **Supplementary Table 7**. See abbreviations in text. Data for three extant rhinos from Guérin (1980); data for *Trigonias osborni* and *Teleoceras proterum* from Prothero (2005).



**FIGURE 6 |** Comparative limb bone proportion in *Plesiaceratherium gracile* and other rhinos. Original data in the **Supplementary Table 7**. See abbreviations in text.

*P. gracile* and *D. sumatrensis* show an increased proportion of the metacarpus when compared with other rhinoceroses (**Figure 6**). The limb segments of the hind-limb have a pattern of elongation distinct from the fore-limb. The elongation of the metatarsals of *P. gracile* is proportionally similar to that of the femur with increasing size. However, relative to *Tr. osborni*, all other rhinoceroses have a proportional increase of the femur but a decrease of the metatarsals (**Figures 5, 6**).

The analyses of *T. major* and *T. proterum* confirm that the shortening of both the proximal and distal parts of the limb contributes to the decreasing of the body size, particularly the metatarsus. With further shrinking of the body size in *T. proterum* relative to *T. major*, the tibia has a more evident decrease of the length than does the radius, consistent with the analysis of relevant limb-segments in terms

of the sexual difference of the limb bones in *Teleoceras* by Mead (2000). However, the femur of *T. proterum* is not shortened and has a proportional increase relative to other limb bones. The significance of the distal limb parts in supporting body weight is further documented by the large and heaviest extant rhinoceros *C. simum*, in which both the metacarpus and the metatarsus are proportionally smaller than other large- or medium-sized rhinoceroses listed in **Figures 5, 6**.

### CONCLUSION

The mensural intraspecific variations in *P. gracile* are wide ranging. We recognize the size of i2 and I1 as significant sexual dimorphism of *P. gracile*. With respect to other features, the

female skull and limb have a larger mean value than those of the male, but the female has a relatively narrower proximal width of the long bones. Meanwhile, the mean length of P2 and M1 of the male is larger than that of the female. However, all these measures show a high degree of overlap; therefore, they do not individually contribute to determining the gender.

Concerning body size, many well-preserved skeletons of *P. gracile* reveal that its head-body length varies between 2,796 and 3,117 mm, and the shoulder height between 1,476 and 1,627 mm. The mass estimation suggests a body weight of about 1,198 kg, with the highest value being 2,244 kg. The bend of the limb and spinal column indicates a mediportal limb, and the running type would be horse gallop. Based on the length of the limb bones among rhinoceroses with different sizes and degrees of evolution, we conclude that the proximal long bones, including the scapula, the humerus, and the femur, are important for a longer limb, and the distal part, namely the metapodials, is important for bearing the body mass. *P. gracile* has a relatively large size, which, together with the strong dimorphic incisors, allows the inference of polygynous behavior and a solitary lifestyle for this rhinocerotid species.

## DATA AVAILABILITY STATEMENT

All datasets presented in this study are included in the article/**Supplementary Material**.

## AUTHOR CONTRIBUTIONS

All authors listed have made a substantial, direct and intellectual contribution to the work, and approved it for publication.

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## SUPPLEMENTARY MATERIAL

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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