

# *First diatomyid rodent from the Early Miocene of Arabia*

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# First diatomyid rodent from the Early Miocene of Arabia

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**Abstract** The Asian family Diatomyidae is known from the Early Oligocene to the present. Among living rodents, this group comprises only the recently discovered *Laonastes aenigmamus* from Laos. Fossil diatomyids are known from only a few sites, in which they are often rare. The discovery of *Pierremus explorator* gen. nov. sp. nov. in the Lower Miocene of As-Sarrar (Saudi Arabia) raises to ten the number of extinct diatomyid species recognized. *Pierremus explorator* is the first record of a diatomyid from the Afro-Arabian plate. This discovery provides evidence that, together with other rodents (ctenodactylids, zapodids...), the diatomyids took advantage of the corridor that was established between Afro-Arabia and Eurasia in Early Miocene times.

**Keywords** Diatomyidae Saudi Arabia Miocene Dam Formation Paleobiogeography

## Introduction

The Diatomyidae Mein and Ginsburg 1997 is a family of hystricomorphous and sciurognathous rodents that originated from the Paleogene ctenodactyloid stock (Flynn 2007; Dawson et al. 2006; Marivaux et al. 2004b). As presently

understood, they comprise nine fossil species in four genera (*Fallomus razae* Flynn et al. 1986, “*F.*” *ladakhenis* Nanda and Sahni 1998, *F. ginsburgi* Marivaux and Welcomme 2003, *F. quraisyi* Marivaux and Welcomme 2003, *Diatomys shantungensis* Li 1974, *D. liensis* Mein and Ginsburg 1985, *D. chitaparwalensis* Flynn 2006, *Marymus dalanae* Flynn 2007, *Willmus maximus* Flynn and Morgan 2005). The amazing discovery in 2005 of the modern *Laonastes aenigmamus* Jenkins et al. 2005 was recognized as a member of the family Diatomyidae by Dawson et al. (2006).

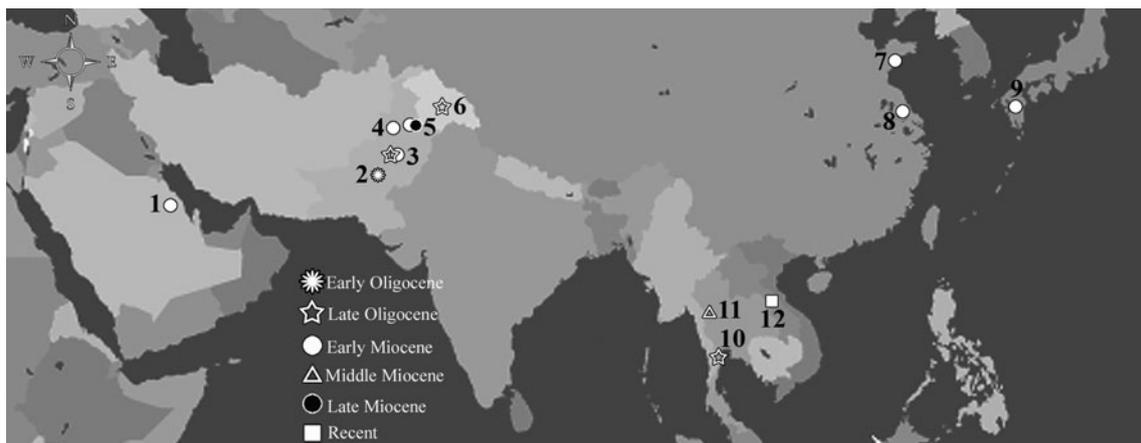
The Diatomyidae are usually uncommon in the fossil record. They are known from the Early Oligocene to the Late Miocene in Pakistan (Flynn 2006, 2007; Flynn and Morgan 2005; Flynn et al. 1986; Marivaux and Welcomme 2003), India (Nanda and Sahni 1998), Thailand (Ducrocq et al. 1995, Marivaux et al. 2004a; Mein and Ginsburg 1985, 1997), China (Ducrocq et al. 1995; Li 1974) and Japan (Kato and Otsuka 1995) (Fig. 1). *Diatomys* sp. was also mentioned from the Lower Miocene of Jebel Zelten, Libya (Savage 1990). However, it is most likely a misidentification because all the remains collected at Jebel Zelten during the geological and paleontological field campaigns carried out by Fejfar in 1982–1983 and by El Arnauti-Daams in 1997 were later studied in detail and no material of Diatomyidae was found amongst them (Wessels et al. 2003, 2008; Fejfar and Horáček 2006; Fejfar pers. comm. 2010). In this work, I report the first record of a diatomyid from the Afro-Arabian plate.

## Acronyms

The following acronyms were used: AS, As-Sarrar (Saudi Arabia); MNHN, Muséum national d’Histoire naturelle (Paris, France); TF, Thai fossil (Department of Mineral Resources, Bangkok, Thailand); SHM-CP, Srisuk House

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**Fig. 1** Distribution of fossil and recent Diatomyidae localities. 1 As-Sarrar (Saudi Arabia). 2 Bugti (Pakistan). 3 Zinda Pir Dome (Pakistan). 4 Banda Daud Shah (Pakistan). 5 Potwar Plateau (Pakistan).

6 Ladakh (India). 7 Shanwang (Shandong Province, China). 8 Xiacaowan (Jiangsu Province, China). 9 Kyushu (Japan). 10 Phetchaburi (Thailand). 11 Li Basin (Thailand). 12 Kammouhan (Laos)

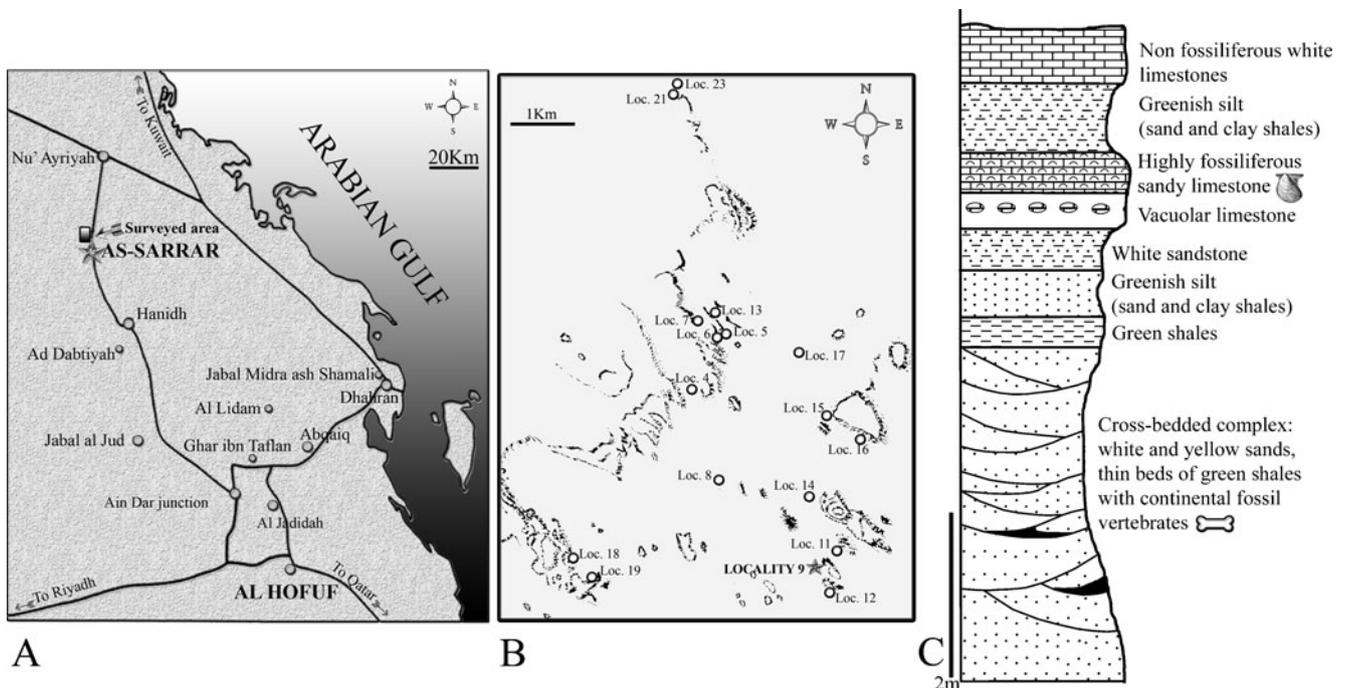
Museum (Cha Prong, Thailand); WIMF, Museum of Wadia Institute of Himalayan Geology (Dehra Dun, India).

**Geological context**

The fossiliferous localities of the As-Sarrar region are located about 10 km N-NW of As-Sarrar (26° 59' N; 48° 23' E), less than 90 km from the coast of the Arabian Gulf and about 250 km NW of Al Hofuf (Thomas 1982; Thomas

et al. 1982) (Fig. 2a). Twenty sites numbered 4 to 9 and 11 to 24 are situated in the Dam Formation (Fig. 2b). Only four of them have yielded rodent remains and among them only at locality 9 the family Diatomyidae is represented.

The Dam Formation in the As-Sarrar area displays interbedded continental and marine facies indicating the proximity of the seashore (Fig. 2c). Thus, during deposition of the Dam Formation, the area of As-Sarrar was a transitional zone, particularly sensitive to environmental changes (Thomas and Battail 1980).



**Fig. 2** a Location of As-Sarrar in the Eastern Province of Saudi Arabia. b Detail of the surveyed area near the city of As-Sarrar with the location of the fossiliferous localities. c Stratigraphical section of locality 9. Data after Thomas et al. (1982)

All the vertebrate localities of the Dam Formation in the As-Sarrar area are considered roughly contemporaneous (Thomas et al. 1982). Thomas et al. (1982:132) and Thomas (1984:267–268) presented the negative and positive evidence for a late Early Miocene age. The results of the study of some rodent families from As-Sarrar (López-Antoñanzas 2004; López-Antoñanzas and Sen 2004, 2005, 2006) are consistent with a late Early Miocene age for this site.

**Pierremus gen. nov.**

Order Rodentia Bowdich, 1821

Family Diatomyidae Mein and Ginsburg 1997

Genus *Pierremus* gen. nov.

**Etymology** Named after the great paleontologist Pierre Mein for his substantial contribution to the understanding of rodent evolution, plus *mus* (Latin for mouse).

**Diagnosis** Diatomyid rodents with a trilophodont pattern on the lower teeth; characterized by protruding cusps and cuspids; having strong to indistinct ectostylid and lacking accessory styles or stylids; lower teeth with the entoconid anteriorly displaced with respect to the hypoconid and with a prominent development of the hypoconulid; the third lower molar being the largest of the cheek teeth.

**Differential diagnosis** Differs from *Fallomus*, *Diatomys*, *Marymus*, *Willmus* and *Laonastes* in having a trilophodont pattern on the lower teeth, the entoconid anteriorly displaced with respect to the hypoconid and the m3 much larger than the m1 and the m2.

**Type species** *Pierremus ladakhensis* (Nanda and Sahni 1998) (Fig. 3a)

**Referred species** *Pierremus explorator* sp. nov. (Fig. 3b).

***Pierremus ladakhensis*** (Nanda and Sahni 1998) (Fig. 3a).

**Synonymy**

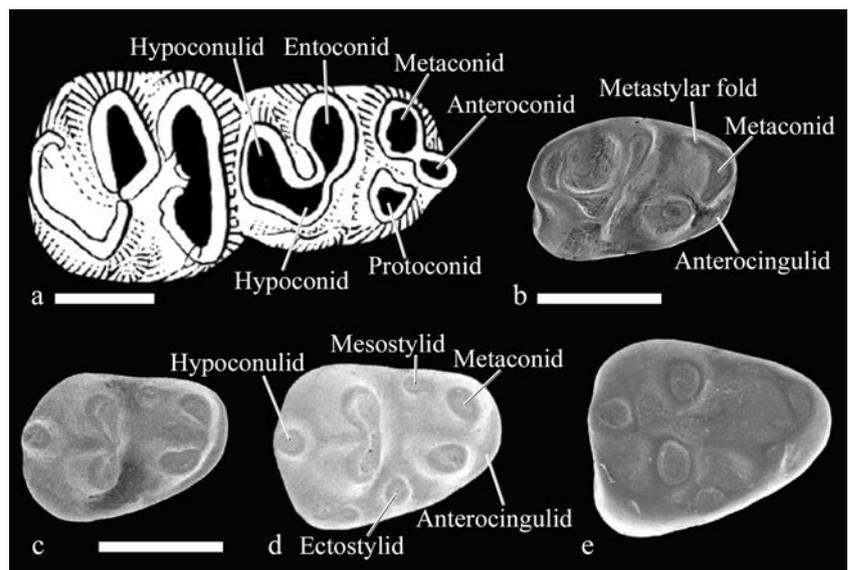
*Fallomus ladakhensis* Nanda and Sahni 1998

*Fallomus razae* (partim) Flynn et al. 1986

**Referred material** WIMF/A 1701–1703, WIMF/A 1705 (Kargil area, India; Nanda and Sahni 1998) and SHM-CP 353, SHM-CP 250, TF 6181–6182 (Phetchaburi Province, Thailand; Marivaux et al. 2004a).

Nanda and Sahni (1998) erected the new species *Fallomus ladakhensis* on the basis of two isolated teeth (second and third lower molars) from the Oligo-Miocene Kargil Formation of the Ladakh Molasse Group of the Kargil area (India). In the same work, these authors described from the same locality five isolated teeth that they identified as *F. razae*: one first lower molar (WIMF/A 1701), three second lower molars (WIMF/A 1702–1704) and one upper molar (WIMF/A 1705). However, these teeth are larger and high crowned and have a quite different morphology from the material from the Bugti area (e.g., they have a trilophodont dental pattern, the entoconid is more anteriorly located than the hypoconid) for them to be considered to belong to *F. razae*. In fact, all the specimens but WIMF/A 1704, which is smaller (but still too large to pertain to *F. razae*), could belong to *F. ladakhensis*. In addition, *F. ladakhensis* (Fig. 3a) does not match the diagnosis given by Flynn et al. (1986) for the genus *Fallomus* in having higher crowned teeth, a trilophodont dental pattern, the entoconid anteriorly displaced with respect to the hypoconid and the third lower molar larger than the second one. Additional material of *Pierremus ladakhensis* comes from the Late Oligocene

**Fig. 3** Diatomyid dental terminology used in this paper. **a** *Pierremus ladakhensis*, right mandible with p4-m1. **b** *Pierremus explorator* sp. nov., right p4. **c** *Fallomus razae*, right p4. **d** *Fallomus ginsburgi*, right p4. **e** *Fallomus quraishyi*, right p4. Scale bar equals 1 mm



deposits of the Nong Ya Plong coal mine, which is located in peninsular Thailand (Phetchaburi Province; Marivaux et al. 2004a). The material of the former *F. ladakhensis* has been described in details by Nanda and Sahni (1998) and Marivaux et al. (2004a)

***Pierremus explorator* sp. nov.**

*Etymology Explorator*, Latin for explorer, for being the first known diatomyid that “dared” to travel out of the Eurasian plate.

*Holotype* MNHN-AS9-22 (Fig. 3), a right p4, the single specimen found to date. It is housed in the palaeontological collections of the Museum national d’Histoire naturelle (Paris, France).

*Locality, horizon, and age* The single specimen of *Pierremus explorator* has been recorded from the late Early Miocene locality 9 of the As-Sarrar area (Saudi Arabia).

*Diagnosis* Diatomyid rodent with brachydont dentition; having a trilophodont dental pattern with large cuspids; the metaconid anteriorly located, the protoconid posteriorly displaced with respect to the metaconid, the entoconid much anteriorly displaced with regard to the hypoconid, and lacking accessory stylids.

*Differential diagnosis* MNHN-AS9-22 differs from the p4 of all species belonging to the family Diatomyidae (except for *Pierremus ladakhensis*) in having a trilophodont dental pattern and the entoconid anteriorly displaced with respect to the hypoconid. It differs from *P. ladakhensis* in being more brachydont, smaller, in having a large hypoconid and in lacking the anteroconid. This species is smaller than *Fallomus quraishyi*, *Diatomys liensis*, *D. shantungensis*, *Marymus dalanae*, *Willmus maximus*, and the extant *Laonastes aenigmamus*.

**Description**

The holotype is a worn brachydont permanent premolar (MNHN-AS9-22:1.661 × 1.241 mm) with the trigonid slightly narrower than the talonid. Its occlusal outline is oval, longer than wide. This tooth is identified as a permanent premolar rather than a deciduous one mainly due to the lack of anteroconid and the presence of massive main cuspids. The presence of the anteroconid and sharp principal cuspids and crests are indeed usually characteristics of milk teeth (Hartenberger 1982:23; Marivaux and Welcomme 2003:422). MNHN-AS9-22 has an anterocingulid and it is characterized by a trilophodont dental pattern with large

metaconid, protoconid, entoconid, hypoconid, and hypoconulid. The metaconid is located on the anterolingual border of the tooth. There is a short, but distinct, metastylar fold (posterior arm of the metaconid). The protoconid is isolated and it is located much further back than the metaconid. Both metalophulid I and metalophulid II (anterior and posterior arms of the protoconid, respectively) are absent. The entoconid is much more anteriorly positioned than the hypoconid. The tooth is damaged in its posterobuccal side and therefore it is not possible to know if it had the ectostylid, but additional stylids are absent. A longitudinal valley between the buccal (protoconid) and lingual (metaconid) cuspids runs posterobuccally. The anterior side of the hypoconid and the entoconid are linked by the hypolophid. The enlarged hypoconulid forms a third posterior lophid by contact with the hypoconid.

**Discussion**

*Fallomus*

The genus *Fallomus* was originally interpreted as a Chapatimyidae *incertae sedis* (Flynn et al. 1986). After the works of Flynn et al. (1986) and Mein and Ginsburg (1997), it has been included in the Diatomyidae (Marivaux and Welcomme 2003; Marivaux et al. 2002, 2004a). Flynn et al. (1986) characterized the genus as follows: cheek teeth transversely bilophodont, lacking well-developed longitudinal crests, high, inclined cusps; upper cheek teeth with four major cusps, lacking metaconule and paraconule; lower teeth with five major cusps including hypoconulid; enterostyle and ectostylid present in high frequency; with the second molars being the largest cheek teeth of the tooth row; four-rooted M1–M3 and m2. These authors named the type and only species known then *F. razae* on the basis of a large sample of specimens from locality Y417 (south of Dera Bugti, Balochistan, Pakistan). An Early Oligocene age is currently assigned to the site (Welcomme et al. 2001; Marivaux and Welcomme 2003; Métais et al. 2009).

Marivaux and Welcomme (2003) named two new species of *Fallomus* (*F. quraishyi* and *F. ginsburgi*) on the basis of numerous specimens from the Early Oligocene Paali Nala C2 locality of the Chitarwata Formation (Bugti Hills, Balochistan, Pakistan) (Fig. 3d–e). From the same locality, remains of *F. razae* (Fig. 3c) were found (Marivaux and Welcomme 2003). Thus, three species belonging to the genus *Fallomus* are known to date: *F. razae*, *F. quraishyi*, and *F. ginsburgi* (Fig. 3c–e).

Additional material of *Fallomus razae* has been found from the Late Oligocene Zinda Pir Dome localities Z144 and Z108 (Flynn 2006). In this latter locality, *F. razae* was recorded along with *F. quraishyi* and *F. ginsburgi*.

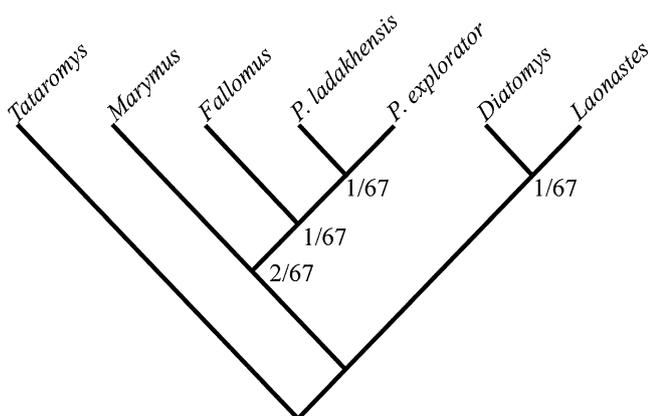
## Pierremus

As seen above, the new genus *Pierremus* is here named to include the former *Fallomus ladakhensis*, which did not match the diagnosis of the genus *Fallomus* proposed by Flynn et al. (1986). The new species from Saudi Arabia, *P. explorator*, shares with *P. ladakhensis* the following synapomorphies: a trilophodont dental pattern and the entoconid anteriorly displaced with respect to the hypoconid. These two species form a clade with sister-group relationships with *Fallomus* (Fig. 4 and supporting online material).

*Fallomus* and *Pierremus* share some characters with primitive chapattimyid ctenodactyloids such as the presence of large submolariform premolars, large third molars, and the retention of the hypoconulid on the lower molars. However, the genus *Pierremus* is more derived than *Fallomus* in having a trilophodont dental pattern and in having the entoconid anteriorly displaced.

The single specimen of *P. explorator* has retained another primitive ctenodactyloid feature: the presence of an anterocingulid, which is absent in *P. ladakhensis*. Thus, *P. explorator* may be more primitive than *P. ladakhensis*. Furthermore, regarding the size and height of the crown, *P. ladakhensis* appears more derived than *P. explorator* and *Fallomus*. In fact, the former species is much larger and has higher crowned molars than the latter taxa. These characters, together with a more derived incisor enamel microstructure, have been thought to mean that *P. ladakhensis* is more advanced morphologically than *Fallomus* (Marivaux et al. 2004a).

Regarding the pattern of premolar replacement, the three species of *Fallomus* known to date show the primitive ctenodactyloid condition of normal premolar replacement.



**Fig. 4** Single most parsimonious tree generated by the cladistic analysis performed in this paper (matrix in supplementary data). Bremer and relative Bremer support indices (Goloboff and Farris 2001) of each clade are indicated at the appropriate nodes. The tree has a length of 15 steps and a low level of homoplasy (CI=0.867 and RI=0.833)

The deciduous premolars of *Fallomus* (the most ancient genus of the diatomyid rodents) are easily recognizable, even though they have not a very different morphology from that of the permanent premolars (Marivaux and Welcomme 2003; Flynn 2007).

This is not the case for more modern genera. With regard to the pattern of premolar replacement in *Pierremus*, little can be state with certainty. The only known two premolars of *P. ladakhensis* are insufficient to settle the question. It is worth mentioning that the only two specimens of p4 of *P. ladakhensis* found (Marivaux et al. 2004a) have a developed anteroconid. According to Marivaux and Welcomme (2003), the deciduous premolars of *Fallomus* are distinguishable from the permanent ones thanks to the development of anteroconids and anterocones, a suggestion with which I agree. This could suggest that the two lower premolars of *P. ladakhensis* are deciduous premolars. However, the two mandibles of *P. ladakhensis* (SHM-CP 250 and SHM-CP 353) from Thailand do not appear to correspond to juvenile individuals (the third molar is fully erupted and worn). Furthermore, the careful examination of this material did not show any trace of erupting permanent premolars (Marivaux pers. comm. 2010). Thus, it is possible that *P. ladakhensis* retained the deciduous premolars throughout its lifetime as was the case of *Diatomys* (Mein and Ginsburg 1997), probably that of *Marymus* (Flynn 2007), both of which also show an anteroconid on the premolars, and that of the extant *Laonastes* (Hautier and Saksiri 2009). If so, *P. ladakhensis* would be derived with respect to *Fallomus* also in the lack of premolar replacement. The discovery of additional material of *P. ladakhensis* is necessary in order to confirm or refute this hypothesis. On the other hand, the finding of a single premolar of the Arabian species does not allow inferring the pattern of premolar replacement in *P. explorator*.

## Conclusion

The new diatomyid species found at the late Early Miocene locality 9 of As-Sarrar belongs to a new genus, *Pierremus*, to which is reallocated *Fallomus ladakhensis*. The latter species does not belong to the genus *Fallomus* because of the trilophodont dental pattern of its lower cheek teeth and the entoconid anteriorly displaced with respect to the hypoconid. These characters are shared with the new Arabian species, *P. explorator*, and suggest that *Pierremus* is more derived than *Fallomus*. In addition, *P. ladakhensis* is more derived than *P. explorator* and *Fallomus* spp. in being larger and in having higher crowned teeth. Furthermore, *P. ladakhensis* is more derived than *Fallomus* spp. in the incisor enamel microstructure and, perhaps, in suppressing the permanent premolars, which is a derived condition

among the members of this family. *Laonastes*, the sole extant member of the Diatomyidae, is also characterized by the retention of the deciduous premolars (Hautier and Saksiri 2009). If the recovery of more material of *Pierremus* confirms that it lacked premolar replacement, the acquisition of this derived condition within the family Diatomyidae would have occurred during the Late Oligocene.

Except for the Early Oligocene Paali Nala locality (Pakistan) and the Middle Miocene Li Basin (Thailand), the diatomyids are rare in the fossil record so that each find has significance for the understanding of their evolution, which, for the time being, is quite enigmatic. It has been postulated that about 18.5 million years ago, the counter-clockwise rotation of the Afro-Arabian plate brought it into contact with Eurasia, from which it was formerly separated by the Tethys Seaway (e.g., Rögl 1998, 1999a, b). The obstruction of the Tethys and the establishment of passageways between the two land masses enabled continental faunal interchanges between the Afro-Arabian and Anatolian areas. The existence of these corridors has been substantiated by the fossil record of various mammalian groups including rodents (e.g. Koufos et al. 2005; López-Antoñanzas 2004, 2009; López-Antoñanzas et al. 2004; Wessels 2009). By the beginning of the Miocene the emerged lands of the Arabian plate experienced the arrival of rodents from various origins. Previous works on the Early Miocene rodents from As-Sarrar (López-Antoñanzas 2004; López-Antoñanzas and Sen 2004, 2006) provided us with examples of long distance dispersal from Asia, as evidenced by the ctenodactylid *Sayimys assarrarenis* and the zapodid *Arabosminthus isabellae*. *Pierremus explorator*, a rodent with incontestable Asian origin, proves that diatomyids also took advantage of this corridor. The dispersal of ctenodactylids and zapodids from Asia to North Africa (Fejfar and Horáček 2006; Wessels 2009; Wessels et al. 2003, 2008) through the Arabian Peninsula suggests that the absence of diatomyid remains in Africa could be due to a bias in the Early Miocene fossil record of this continent.

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