# A NEW RODENT FROM QUATERNARY DEPOSITS OF THE CANARY ISLANDS AND ITS RELATIONSHIPS WITH NEOGENE AND RECENT MURIDS OF EUROPE AND AFRICA 

by<br>Rainer HUTTERER*, Nieves LOPEZ-MARTÍNEZ** and Jacques MICHAUX***

## SOMMAIRE

Page
Abstract, Résumé, Resumen ..... 242
Introduction ..... 242
Description of localities and materials ..... 243
Methods ..... 244
Description of the new rodent ..... 246
Character analysis and phylogenetc affinities ..... 250
Conclusions ..... 254
Aknowledgments ..... 255
Tables 1 to 5 ..... 256
Bibliography ..... 260
Legends of plates ..... 262

* Zoologisches Forschungsinstitut und Museum Alexander Koenig, Adenauerallee 150-164. D-5300, BONN, West Germany.
** Departamento y UEI Paleontologia. Instituto de Geología Económica CSIC. Facultad de C.Geologicas. Universidad Complutense, 28040 MADRID, Spain.
*** U.R.A. 327, Institut des Sciences de l'Evolution (C.N.R.S.) et Laboratoire de Palcontologie des Vertébrés de I'E.P.H.E., U.S.T.L., Place E. Bataillon. 34060 MONTPELLIER Cedex, France.

Key-words: Rodents, Muridae, Holocene, Canary Islands, Spain, Phylogeny, Island evolution.
Mots-clés: Rongeurs, Muridae, Holocène, Iles Canaries, Espagne, Phylogénie, Evolution insulaire.


#### Abstract

A peculiar new rodent, Malpaisomys insularis nov. gen., nov. sp., is described from subfossil deposits of the eastern Canary Islands. The species shows some highly specialized skull features although its molars exhibit a mixture of primitive and derived characters among which a partial stephanodonty is most notable. A comparison of the new rodent with several Miocene to Holocene Muridae shows that Malpaisomys possibly shares a common ancestor with Acomys and Uranomys.


## RESUME

Malpaisomys insularis nov. gen., nov. sp. est un rongeur Muridé original trouvé dans des dépôts récents des îles Canaries orientales. Il possède des structures craniennes très spécialisées et ses molaires présentent une combinaison de caractères primitifs et dérivés dont, parmi ces derniers, une structure stéphanodonte. La comparaison de ce nouveau rongeur avec des Muridés néogènes et actuels montre qu'il partage probablement un ancêtre commun avec Acomys et Uranomys.

## RESUMEN

Malpaisomys insularis nov. gen., nov. sp. es un roedor Múrido peculiar hallado en depósitos recientes de las Islas Canarias orientales. Pose estructuras craneanas muy especializadas y sus molares presentan una combinación de caracteres primitivos y derivados, entre estos ultimos, una estructura estefanodonta. La comparación del nuevo roedor con los Múridos neógenos y actuales muestra que posiblemente comparte un antecesor común con Acomys y Uranomys.

## INTRODUCTION

The Canary Islands comprise seven larger islands and a number of islets in the Atlantic Ocean west of the coast of southern Morocco. They are of volcanic origin and date back from 2 to 35 Mio. years (Abdel-Monem et al., 1967). The mammalian fauna of these islands is not yet fully known, and only recently a new species of shrew was discovered (Hutterer et al., 1987). Among the rodents, two species of extinct giant rats were described: Canariomys bravoi CRUSAFONT-PAIRO \& PETTER, 1964, and Canariomys tamaranis LOPEZ-MARTÍNEZ \& LOPEZ-JURADO, 1987. The former lived on Tenerife, the second on Gran Canaria. No fossils of endemic rodents have ever been found before in one of the eastern Canary Islands, Lanzarote and Fuerteventura. It was therefore a great surprise when in 1979 an archaeological excavation of a Prehispanic site in Fuerteventura (Garralda et al., 1981) yielded numerous remains of an unknown rodent apparently not related to any of the Canariomys species. Recent fieldwork by the first author, together with S. Lenné and F. Hutterer, added seven new localities in Fuerteventura, Lanzarote and the islet Graciosa where remains of the same rodent were found. This first account serves to describe and name the new murid and to evaluate its possible origin.

## DESCRIPTION OF LOCALITIES AND MATERIALS

The known localities of the new murid are shown in fig. 1. A short description of each of them follows including notes on the associated mammal fauna. The material obtained at each site is shown in brackets and expressed as the "Minimum Number of Individuals" (MNI); the MNI was calculated from the left and right maxillae and mandibulae; the actual number of specimens is therefore five to ten times larger.


Fig. 1. Map of the Canary Islands with the localities where remains of Malpaisomys nov. gen. were found.

Locality 1. Fuerteventura, Cueva Villaverde (MNI $=944+$ uncounted material). A cave of volcanic origin which was inhabited by Prehispanic men. The cave deposits were excavated by a team of Spanish archaeologists from 1979 to 1987 (see Garralda et al., 1981; Meco Cabrera et al., 1982; Hernández \& Sanchez, 1986) and their activities yielded an extremely rich vertebrate fauna. Sixty samples containing microfauna were obtained, 40 from inside the cave, and 20 from outside. Seven levels of different age have been distinguished in the cave. Radiocarbon dates are available for the uppermost level (surface; $1070 \pm 50$ B.P.) and for one of the deeper levels (level 3; $1730 \pm 50$ B.P.). Additional small mammals in this site are Crocidura canariensis and Mus musculus. The house mouse is dominant in the upper levels but almost absent in the lowermost levels (Carrascosa \& López-Martínez, 1988).

Locality 2. Fuerteventura, Malpais de Arena ( $\mathrm{MNI}=20$ ). A small section of a collapsed lava tube with a compact sediment filling of 5 to 20 cm thickness. A section of $1 \times 1$ m was screened in 1987 by R. Hutterer and S. Lenné (also loc. 3 and 4). It contained also bones of Crocidura canariensis, Mus musculus and Oryctolagus cuniculus. Goat and pieces of pottery were also found. No datation of this site is yet available, but the presence of Oryctolagus indicates a younger age than locality 1.

Locality 3. Fuerteventura, Malpais Grande ( $\mathrm{MNI}=7$ ). Bones were screened from the sediment which filled small caves in the malpais, some were found on the surface. Crocidura and Mus were also present. The age of these recent lava flows is around a few thousand years (Afonso Perez, 1980).

Locality 4. Fuerteventura, Jandia Peninsula, north coast at 150 m a.s.l.; (MNI $=2$ ). A hanging dune consisting of compressed white sand yielded a rich bird fauna but only few rodent bones. No other mammal was found in this site. Judged from the geological situation this site might be older than the preceding ones.

Locality 5. Lanzarote, Malpais de Corona ( $\mathrm{MNI}=10$ ). Bones screened from sediment in fissure and cave fillings in the malpais by R. and F. Hutterer in 1987 (all Lanzarote localities). Access to sediment was only possible at road cuts. Other mammals found are Crocidura, Mus, Rattus and Oryctolagus. The presence of Rattus and Oryctolagus indicates a young age of this site. The age of the Malpais de Corona was estimated as ca. 3000-5000 years (Bravo, 1964; Hernandez-Pacheco, 1909).

Locality 6. Lanzarote, Arrieta, ca. 3 m a.s.l. $(\mathrm{MNI}=18)$. Fissure fillings of an ancient malpais covered by about 2 m of alluvial sediments. The site was accessible for a few days during the construction of a building pit. The small mammal fauna includes Crocidura and Mus.

Locality 7. Lanzarote, Casa de la Caldera ( $\mathrm{MNI}=4$ ). A few bones were screened from a cave sediment in a remnant malpais area. Crocidura, Mus and Oryctolagus were also present.

Locality 8. Graciosa, Montaña Amarilla (MNI = 2). Bones were screened from a fissure filling. Crocidura, Mus and Oryctolagus were also present.

In addition to the mapped localities (fig. 1), we found small samples of Malpaisomys nov. gen. at La Vegueta and at Costa Teguise, Lanzarote. Further material from Mozaga, Lanzarote, was studied by us in the Museo Insular, Santa Cruz de Tenerife. Additional material was also collected in Fuerteventura by A. Alcover (pers. comm.)

## METHODS

For the evaluation of the characters of Malpaisomys insularis nov. gen., nov. sp., we selected 10 recent and 13 extinct species of murids as follows:

Apodemus sylvaticus (LINNÉ, 1758)
Rattus rattus (LINNÉ, 1758)
Mus musculus (LINNÉ, 1758)
Acomys cahirinus (DESMAREST, 1819)
Arvicanthis niloticus (DESMAREST, 1822)
Dasymys incomtus (SUNDEVALL, 1847)
Grammomys dolichurus (SMUTS, 1832)
Oenomys hypoxanthus (PUCHERAN, 1855)
Thamnomys venustus THOMAS, 1907
Uranomys ruddi DOLLMAN, 1909
Canariomys bravoi CRUSAFONT \& PETTER, 1964
Canariomys tamarani LOPEZ-MARTÍNEZ \& LOPEZ-JURADO, 1987
Progonomys cathalai SCHAUB, 1938
Occitanomys adroveri (THALER, 1966)
Occitanomys montheleni AGUILAR \& MICHAUX, 1986

Stephanomys donnezani (DEPÉRET, 1890)
"Paraethomys" miocaenicus JAEGER, MICHAUX \& THALER, 1975
"Paraethomys" meini (MICHAUX, 1969)
Paraethomys darelbeidae JAEGER, 1975
Parapodemus lugdunensis SCHAUB, 1938
Karnimata darwini JACOBS, 1978
"Parapodemus" vireti SCHAUB, 1938
Anthracomys majori SCHAUB, 1938
Initial comparisons were also made with specimens of Castillomys crusafonti MICHAUX, 1969, C. magnus SEN, 1977, Occitanomys brailloni MICHAUX, 1969, Occitanomys sp. from Calta (Sen, 1977), Paraethomys cf. filfilae PETTER, 1968, P. cf.anomalus DE BRUIJN, DAWSON \& MEIN, 1970, P. ras JAEGER, 1975, and Valerymys ellenbergeri (THALER, 1966). For the final analysis we used only 17 taxa. The initial comparison started with 4 mandibular and 46 dental characters, of which only 20 were used for the phylogenetic analysis to reduce the number of highly convergent characters.


Fig. 2. Abreged nomenclature of dental structures : a diagram of upper and lower molars in Apodemus. Upper molars : cusps are numbered according to Miller's (1912) scheme and cusps (or tubercules) are referred to in the text with the prefix t ; $\mathrm{pc}:$ posterior cingulum; arrows indicate the position respectively of $t$ lbis and $t$ bis. Lower molars: L1, L2, L3 are respectively the first, the second and the third (or posterior) lobes; alc : anterolabial cusp on second lower molar; cl : cusplet on the side of postero external cusp of first and second lower molar; c accessory cusplets on the external border of the lower molar; pc : posterior cingulum.

We used a cladistic approach for phylogenetic analysis. Cladograms have been obtained using a Phylip 2.8. program of J. Felsenstein (1985), and a manual procedure of a stepwise elimination of the most convergent characters (López-Martínez, 1978). Some tables have been made with the MacClade 2.1 program of W . Maddison and D. Maddison (1987).

Nomenclature used for dental description is given in fig. 2. Occlusal structures of murid teeth have received various names. For upper molars (fig. 2A) the nomenclature is the one proposed by Michaux (1971) which is derived from Miller's nomenclature. Lower molars (fig. 2B) which play a lesser role in phylogenetical studies than the upper ones are here described using a simplified nomenclature derived from the one of Michaux (ibid.). The accessory cusps in front of the anterior lobe (L1) of the first lower molar is called the tma, and c1 is the cusplet located on the side of the external cusp of the posterior lobe or third lobe (L3) of the first and second lower molars.

## DESCRIPTION OF THE NEW RODENT

## RODENTIA: MURIDAE

## MALPAISOMYS NOV. GEN.

Type species: Malpaisomys insularis nov. sp.
Diagnosis: As for the type species.
Etymology: The generic name is formed by combining the Greek mys for mouse with "malpaís", the Canarian name for rough lava fields.

Malpaisomys /nsularis nov. sp.
Holotype: Partial skull of an adult specimen with the anterior part well preserved but lacking most of the braincase (fig. 3), CV-188.1, deposited in the Department of Paleontology, Universidad Complutense, Madrid.
Type locality: Cueva Villaverde near La Oliva ( $22^{\circ} 38^{\prime} 10^{\prime \prime} \mathrm{N}, 10^{\circ} 12^{\prime} 30^{\prime \prime} \mathrm{W}$ ), Fuerteventura, Canary Islands. The age of the type level (CV-A1 surface) is $1070 \pm 50$ B.P.


Fig. 3. Upper M1-3 of the holotype of Malpaisomys insularis nov. gen., nov. sp. (CV. 188-1).

Paratypes: Numerous cranial and skeletal elements from level 1 to 3 of the type locality representing 944 individuals. Paratypes are deposited in the Departamento de Paleontologia, Universidad Complutense, Madrid; in the Museo Nacional de Ciencias Naturales, Madrid; the Museo Canario, Las Palmas de Gran Canaria; the Museo Insular de Ciencias Naturales de Santa Cruz de Tenerife; the Departamento de Zoologia, Universidad de La Laguna, Tenerife; the Museum National d'Histoire Naturelle, Paris; the Laboratoire de Paléontologie des Vertébrés, the Université de Montpellier; and the Museum Alexander Koenig, Bonn.
Other material: Specimens representing 65 individuals from Fuerteventura (localities 2-4), Lanzarote (loc. 5-7) and Graciosa (loc. 8), deposited in the respective collections of Bonn, Montpellier, Madrid, Santa Cruz de Tenerife, and La Laguna de Tenerife.
Diagnosis: A small-bodied terrestrial rat characterized by the following combination of characters: skull stout with wide zygomatic arches and an extremely narrow interorbital constriction; incisive foramina long, their distal tips reaching the intermolar space at the middle of M1 sup.; palate extending posteriorly and covering most of the mesopterygoid fossa with bone; bulla small; mandible with a prominent alveolar crest which passes almost to the distal part of the condyle; the condyle sigmoid-shaped in lingual view; first upper molar with cusp t1 placed in a rather distal position; cusps t6 and t 9 separated far from each other; cusps $\mathrm{t} 1-\mathrm{t} 5$ and $\mathrm{t} 6-\mathrm{t} 9$ frequently connected by a crest.
Measurements: Tables 1-4.
Etymology: Named because the species is only known from islands.
Distribution: Known from subfossil deposits in Fuerteventura, Lanzarote and Graciosa, Canary Islands. The species got extinct in historical times.
Description: The skull and dentary of Malpaisomys insularis are shown in figs. 4 and 5. Its distinct shape, namely the wide zygomatic arches and the extremely narrow interorbital, recalls at once the skull of Nesoryzomys, a genus of lava-dwelling rats endemic to the Galapagos Islands (see Patton \& Hafner 1983, fig. 2). The dorsal profile of the skull is convex; the rostrum and nasals are of moderate size, the nasals are slightly broadened in their proximal parts; the zygomatic arches are very wide, their


Fig. 4. Skull of Malpaisomys insularis nov. gen., nov. sp. in dorsal, ventral and lateral view, based upon an almost complete specimen from Fuerteventura (loc. 2).
1


## 3 <br> 

4






TTT

Fig. 5. Mandibular fragments of Progonomys, Occitanomys and Paraethomys compared with a dentary of Malpaisomys.

1. Progonomys cathalai (Montredon sup., France).
2. "Paraethomys" miocaenicus (Hellin, Spain, see Calvo et al., 1978).
3. Occitanomys adroveri (Algora, Spain)
4. Occitanomys montheleni (Mt Hélène, France)
5. Paraethomys meini (Mt Hélène, France)
6. Paraethomys ras (Doukkala-2, Marocco)
7. Malpaisomys insularis (Cueva Villaverde, Fuerteventura)

Specimens in the collections of Montpellier and Madrid.
width being $57 \%$ of the total skull length; the zygomatic plate is wide and its anterior margin is almost vertical; the frontal bones are very narrow so that the interorbital breadth counts $10,8 \%$ of the total skull length; the frontal and parietal ridges are very faint; the braincase is high, its height being almost constant from the middle of the frontal to the end of the parietal bones, then at the relatively small interparietal bone the braincase gently slopes down to the occipital bone; the incisive foramina are long and wide and extend past the middle of the first upper molars; the palatal bridge is narrow but extends far beyond to the back of the third molars, palatinum and pterygoid fossa are fused so that most of the mesopterygoid fossa is covered by bone (a similar condition is known among murids in Acomys and Uranomys, see Ellerman, 1941); the bullae are relatively small and of triangular shape, each is separated from the squamosum by a wide postglenoid vacuity; the bone of the eustachian tube is rather long; the mandible is massive; its dorsal curvature of the diastema is sharply angulated (fig. 5); the mental foramen is situated externally to the line of the molar row; the upper margin of the articular process is of sigmoid shape; the coronoid process is small; the angular process is short and rounded; a prominent alveolar ridge is running from the distal end of the alveolar row to the lower margin of the condyle.

The teeth of Malpaisomys insularis are brachyodont (plate 1). The first upper molar shows three roots, the internal one being a large one looking as if it was made of two fused roots. Two roots anchor the first lower molar.

The first upper molar is relatively stout, wider for example than the first upper molar in Stenocephalemys. Molars weakly overlap each other. The second upper molar is more or less as wide as long and the third upper molar is not very reduced. The first upper molar consists of three transverse rows of cusps linked together by crests. This tooth is widest at the level of cusp t1. The central cusps are slightly larger than the inner and outer ones. There are two inner and three outer cusps. Cusp t7 and posterior cingulum are absent on both first and second upper teeth. A gentle and shallow fold of the crown can nevertheless be observed on first upper molar between cusp t 9 and cusp t8.

Cusp t1 occupies a rather posterior position by comparison with cusps t 2 and t 3 . In this manner, Malpaisomys resembles more Occitanomys than Paraethomys. There is no cusplet $t 1$ bis. On most of the molars a narrow crest of variable length links the cusp tl and t 5 following the lingual flank of cusp t5. A short posterior crest can be seen on cusp t3, like often in Apodemus. The second row of cusps ( $t 4, t 5, \mathrm{t} 6$ ) runs more or less parallel to the first one. Cusp t6 slopes weakly in the anterior direction so it offers a long anterior flank. This situation contrasts sharply with the more vertical cusp t3. The relative size difference between cusps $t 9$ and $t 6$ is small and connection between them occurs in half of the cases. Stephanodonty is not completely established in Malpaisomys. The second upper molar possesses a large cusp $t 1$ and a small cusp t 3 . Cusp t1 shows a beak directed labially. In most of the cases, cusps t6 and t9 are isolated and there is no incipient crest linking cusps $t 1$ and $t 5$ as observed on the first upper molar. The third upper molar has a triangular cusp t1 with a short labial beak. This cusp is as big as cusp t4. Cusp t3 is absent. Behind cusp t4 there are two cusps more or less at the same transversal level. The third upper molar is more or less square-shaped. The central cusps, $\mathrm{t} 4, \mathrm{t} 5$ and t 6 form an asymmetric row.

The lower first molar shows in half of the cases a rather small anteromedian cusp. First and second lower molars have a relatively reduced posterior cingulum. The external row of cusplets on the first lower molar is weak or absent. The anterior lobe of the first lower molar is wide and shows two equally well differentiated cusps which are centrally linked to the median lobe. No connection occurs between the median and the posterior lobe. The same situation is present on the second lower molar. The anterolabial cusp is rather small on the second lower molar and absent on the third one.

The posterior lobe of the third lower molar is isolated and widened labially.
Measurements of 100 mandibles from Cueva Villaverde (table 2) show a trimodal distribution of mandible length. Distribution has been separated into three age classes: 1 (molars slightly worn), 2 (molars moderately worn) and 3 (molars heavily worn). Bimodality appears separately in both 2 and 3 age classes. This indicates that dimorphism not related with age is present in the mandibular length of Malpaisomys. The most probable explanation is the sexual dimorphism. There are no indications of a dimorphism in tooth size. However, there is a trend towards smaller molar measurements in the Lanzarote samples, notably in the upper M 1 (table 3).

Numerous remains of the postcranial skeleton have been recovered. At first sight no special features have been observed which would indicate special adaptations. Some elements are shown in plate 2 and measurements are given in table 4. Apparently there exist significant differences in the size of some skeletal elements between some populations. First, within Fuerteventura, the population from locality 4 seems to be larger. Secondly all the Fuerteventura populations show higher measurements than the Lanzarote populations. As the samples may differ in geological age, it is difficult at this moment to evaluate this variation.

## CHARACTER ANALYSIS AND PHYLOGENETIC AFFINITIES

We have selected 20 dental characters for the phylogenetic analysis out of 51 features used in the preliminary comparison. Characters listed below have been rejected because information for all taxa was not available (1) or because they were redundant in relation to other selected characters (2).

1 - Alveolar ridge of mandible (1)
2 - Articular process (1)
3 - Mental foramen (1)
4 - Diastema of the mandible (1)
5 - Overlap of the upper molars (1)
6 - Slope of the molar cusps (1)
7 - Roots of the lower first molar (2)
8 - Shape of the upper second molar (2)
9 - Posterior cingulum of the second upper molar (2)
10 - Relative size of $t 9$ in the first upper molar (2)
11 - Cusp t 7 on upper molars (not present in the compared species)
12 - Connection between t6-19 in second upper molar (2)
13 - Connection between t1-t5 in second upper molar (2)
14 - Connection between t 3 - 55 in second upper molar (2)
15 - Relative position of t6 and t8 on the third upper molar (1)
16 - Posterior cingulum of the second lower molar (2)
17 - Width of the anterior lobe of the first lower molar (2)
18 - Position of the anterior cusps of the first lower molar (2)
19 - Connection between the two lobes of the second lower molar (2)
20 - Antero-labial cusp of the third lower molar (2)
21 - Width of the posterior lobe of the third lower molar (1)
22 - Number of cusps at the posterior part of the third lower molar (1)
23 - Shape of the third upper molar (1)
24 - Size of cusp $t 1$ in the third upper molar (1)
25 - Shape of cusp t 1 in the third upper molar (1)
26 - Shape of the medial lobe in the third upper molar (1)

Remaining characters have been analysed previously with the Lequesne method in search of the a priori less convergent characters. The number of combinations of character states was controlled for each pair of characters in all the selected species. The presence of the four possible combinations of the two main states (primitive and derived) of two characters indicates that one of the features, at least, is convergent or, in the Lequesne terminology, has not been uniquely derived.

After statistical calculation, the characters concerning the stephanodonty showed the smallest number of combinations, indicating that they are a priori less frequently subject to convergence. On the contrary, the characters listed below, and frequently cited in the literature as relevant features for the phylogenetic and systematic studies, showed the highest number of combinations. Therefore, they have been rejected because they introduce noise in the phylogenetic information. The remaining features may have also convergences, but they show a priori not a random distribution.

1 - Relative size of cusp t9 in the second upper molar
2 - Anterio-external beak of cusp t 1 in the second upper molar
3 - Divergence of cusps t3 and t6 in the first upper molar
4 - Size of cusp t 3 in the second upper molar
5 - Presence of cusp $t 3$ in the third upper molar.
The characters selected for the phylogenetic analysis are defined below. Code (0) indicates our view of the primitive state. The species Antemus chinjiensis JACOBS, 1978 has been used as the outgroup for this purpose.

1 - Tooth morphology: brachyodont (0) or semi-hypsodont (1)
2 - Enamel differentiation: thickness of the enamel slightly differentiated (0); almost not differentiated (1) strongly differentiated (2). The two evolved states have probably derived independently.
3 - Width of the first upper molar: position of the maximum width at $\mathrm{t4}(0)$; similar width at the t 4 and t 1 (1); or maximum width at t 1 (2)
4 - Number of roots of the first upper molar: lower than 5 (0) or 5 or more (1)
5 - Position of the t 1 at the first upper molar: strongly backwards as in Acomys (0); less backwards as in Apodemus (1) or at the level of t3 like in Dasymys (2)

6 - Accessory cusps t1bis and t3bis on the anterior lobe of the first upper molar: absent (0), variable (1) or present (2)
7 - Connection of $t 1$ with $t 2$ or $t 4$ with $t 5$ on the first upper molar: absent ( 0 ) or present (1)
8 - Cusps t2-t3 and t5-t6 of the first upper molar: connected but clearly individualized (0) or fused (1)
9 - Connection of t6 with 19 at the first upper molar: absent (0), variable (1) or present (2)
10 - Connection of t 1 with t 5 at the first upper molar: absent ( 0 ), incomplete (1) or present (2)
11 - Connection between cusps t3-t5 at the first upper molar: as in number 10
12 - Connection between t4-t8 at the first upper molar: as in number 9
13 - Posterior cingulum at the first upper molar: large (0), small (1) and absent (2)

14 - Relative size of the third upper molar: ratio length M3/ length M2/ between 0.7 and $0.8(0)$; ratio lower than 0.7 (1) or above 0.8 (2). Values have been calculated for the two morphotypes of Jacobs (1978, p. 79) the intermediate state being considered as primitive
15 - Anterior median cusps (tma) at the first lower molar: absent (0), variable (1) and present (2)
16 - Posterior cingulum at the first lower molar: large (0), reduced (1) or absent (2)

17 - External cingular margin at the first lower molar: with one or two additional cusps (0); with no cusps (1) or with more than two cusps (2). The cusp named C1 and another anterior one are usually present in murids and therefore this condition is considered to be primitive. Derived states are divergent
18 - Anterior lobe of the first lower molar: asymmetrical weak connection as in Stephanomys and Progonomys (0); isolated as in Rattus (1), or strong central connection like in Mus (2). The only well-preserved M1 of Antemus chinjiensis (see Wessels et al. 1982, pl. 2, fig. 7) presents the character state most common among murids, and therefore regarded as primitive. Derived states are divergent
19 - Posterior lobe of the first lower molar: isolated from the median lobe (0); incompletely linked (1) or completely linked (2)
20 - Anterolabial cusp of the second lower molar: large (0), or reduced or absent (1).

The data matrix of the character states' distribution in the selected taxa is shown in table 5. The aim of the phylogenetic analysis is to select one of the possible trees constructed by manual or automatic technics. The phylogenetic tree must have as few convergences as possible; but the problem of character reversion has to be considered.

In the Camin-Sokal procedure, reversions are not allowed; all the evolutionary steps are considered irreversible. On the contrary, the Wagner procedure allows reversions for all characters; steps $0>1$ and $1>0$ are equally probable.

In the case of murids, reversions have been documented in some species lineages. For example, Stephanomys donnezani-S. minor illustrate reversion of the character 17 (Bachelet, pers. comm.); Apodemus gudrunae-A. jeanteti illustrate reversion of the character 15 (Michaux, 1967). For other characters, as the enamel differentiation ( $\mathrm{n}^{\circ} 1$ ), acquisition of the semi-hypsodonty ( $\mathrm{n}^{\circ} 2$ ) and tooth-size reduction ( $\mathrm{n}^{\circ}$ 14), it seems unlikely that reversion has occurred in the evolution of mammals in general. The characters 13 and 16 are considered as not reversible.

Fig. 6 shows one of the cladograms constructed under these above mentioned conditions. The total number of steps for this cladogram is 86 , and includes as much as 54 convergences and 6 reversions. The high number of neccesary convergences indicates that it cannot be taken as a definitive phylogenetic tree. Nevertheless it shows some clusters which were previously recognized by some authors using also characters other than molars, whereas our cladogram is exclusively based on molar characters.

Malpaisomys groups with Acomys and Uranomys; all three share the derived characters 16 and 18 . This cluster is close to the group including all the highly stephanodont taxa: Stephanomys, Occitanomys, "Paraethomys" darelbeidae, and Oenomys. Malpaisomys, Acomys and Uranomys share one additional character in the extension of the palatal bridge (see fig. 4), a unique feature among murids, according to Ellerman (1941). Also the incisive foramina extend far past the middle of the first upper molars in these three genera, which is a character not frequently encountered in murids. On the other hand, Malpaisomys shows a sharply angulated diastema in the mandible
(fig. 5), a character not shared by the two other genera.
Another cluster is formed by Arvicanthis, Rattus and "Canariomys" tamarani. Karnimata darwini cannot be clustered because of its low number of derived characters. At the basis of the tree, the branches are supported by few derived characters, but the general grouping seems congruent with the "rat-mouse" dichotomy of Jacobs (1978, p. 83, fig. 35). Progonomys appears related to the group including Mus and Acomys, as Jacobs proposed (1978, p. 83, fig. 37).

This group includes also several large-sized murids like Canariomys bravoi, and Oenomys. Both share a high degree of stephanodonty and the presence of some accessory cusps in the lower first molar (characters 15 and 17).

The two species of Canariomys (López-Martínez \& López-Jurado, 1987) appear in two separate clusters. This is mainly the consequence of different states in the characters numbers 3,4 and 20. At this stage of our study it seems premature to draw conclusions, but our result might indicate that the two species are not as closely related as previously thought. The same applies to the two species of Paraethomys included in our study. The high number of taxa described in the genus Paraethomys (Jaeger, 1975, Michaux, 1969; de Bruijn et al., 1970; Jaeger, 1975, Jaeger et al., 1975) deserves a careful re-examination before any phylogenetic interpretation can be drawn.


Fig, 6. One cladogram obtained from the data matrix given in table 5. The first three branchings are based on one derived character.

Other cladograms gave sometimes different results in the position of Malpaisomys, indicating a closer relationship with either Paraethomys or Mus. However, none of these alternative solutions fits with additional skull characters like the Malpaisomys - Acomys - Uranomys hypothesis, which we therefore prefer.

## CONCLUSIONS

As mentioned earlier, the skull of Malpaisomys (fig. 7) is highly specialized in its extremely narrow interorbital region and the wide zygomatic arches. Such a skull configuration is also known from other island endemics, like Nesoryzomys of the Galapagos Islands (Patton \& Hafner, 1983). The species of Nesoryzomys are large-eyed, soft-furred nocturnal rats (see Brosset, 1963, for a photograph of Nesoryzomys narboroughi) which live in all available habitats, from the coasts to the summits of the volcanoes (four out of five island populations are extinct; Clark, 1984). Much of the habitat available in the Galapagos corresponds to the malpaís of the Canary Islands. A further example of convergent skull features are the giant rats of the two archipelagos. The Galapagos Megaoryzomys (see Steadman \& Ray, 1982, for an illustration) is very similar in skull shape to the Canarian Canariomys (see López-Mártinez \& López-Jurado, 1987, for an illustration). In addition, the skulls of


Fig. 7. Dorsal aspects of the skull of Nesoryzomys narboroughi, a cricetid, and of Malpaisomys insularis, a murid, to show similarities in skull shape which are due to parallel evolution. Both rodents shared the same type of habitat (volcanic islands). Illustrations are based on specimens in the Museum Alexander Koenig, Bonn. The scales represent 5 millimetres each.
both resemble Crateromys, a genus of arboreal giant rats of the Philippines (Musser et al., 1985). Apparently the three-dimensional creviscular malpais claims largely the same morphological adaptations as thick bush or trees.

The above-mentioned genera belong to different families. Therefore the skull features they have in common certainly developed by parallel evolution. This indicates a rather long, independent evolution of Malpaisomys in the Canary Islands. In our search for its phylogenetic relationships we therefore disregarded these peculiar features. It is for this reason that we tried to use mainly dental characters and cranial features which we supposed not to be adaptive. However, the phylogenetic study showed that even within these selected characters convergences occur rather frequently. For that reason we finally used only 20 characters out of 51 . We arrived at the conclusion that Malpaisomys may share a common ancestor with two African genera, Acomys and Uranomys. Many species of Acomys, most of them of unsettled status, occur in Africa, Arabia, and Asia (Honacki et al., 1982; Petter, 1984). East Africa houses the highest number of species. Along the Atlantic coast of NW-Africa only members of the A. cahirinus group occur. The distributional range of Uranomys is limited by southern Senegal in the north and by Mozambique in the south. Only one species, $U$. ruddi, is currently recognized in this genus (Honacki et al., 1982).

There exists no fossil record of Uranomys, and that of Acomys is very poor. Wesselman (1984) referred a single first upper molar from the Shungura Formation, Upper Pliocene of Ethiopia, to Acomys sp., which was quite similar to the modern species of the genus. Material from Laetoli, Pliocene of Tanzania, also exhibits modern traits (Denys, 1987). Further fossils of Acomys were reported by De Graaf (1961a, b) from Makapansgat and Kromdraai, Pleistocene of South Africa. All the fossil material of Acomys resembles modern species and therefore contributes little to the understanding of the evolution of that genus. Jacobs (1978) postulated that Acomys derived from the "Progonomys stock". According to our results, we must claim that the common ancestor of Malpaisomys, Acomys and Uranomys lived between 11Ma (branching of the Progonomys -"mouse group") and Pliocene times.

## ACKNOWLEDGEMENTS

A team of archaeologists led by F. Hernández and M.D. Sánchez obtained the major part of the fossils from Fuerteventura. S. Lenné and F. Hutterer participated in the fieldwork in Fuerteventura and Lanzarote. M. Martin and F. Garcia-Talavera showed us further material from Lanzarote in the Museo Insular de Tenerife, and G. Storch gave access to the collection of fossil murids in the Senckenberg-Museum Frankfurt. M. Hitzeler and I. Perez aided in the sorting of the fossils. Scanning electron micrographs were made in the Paleontological Institute Bonn by R. Olesinski, and photographic prints by E. Schmitz and J. Schicke. We thank all of them for their invaluable help.
B. Bachelet, C. Denys, J.-J. Jaeger, F. Petter and G. Storch generously shared their experiences with fossil murids with us. For the final upwriting of this report we met at the Laboratoire de Paléontologie des Vertébrés et de Paléontologie Humaine, Universite de P. et M. Curie. We thank all the members of this institute for the facilities they offered to us during that week, including personal computers and coffee, and for their patience. Typewriting of the manuscript and photographs have been respectively done by G. Jean and J. Martin in Montpellier.

Grants were provided by the Museum Alexander Koenig and the French and Spanish cooperation program ( $\mathrm{n}^{\circ} 124-51 / 62$ ).

TABLES 1 to 5

| Locality : | 2 | 2 | 2 | 1 | 1 | 2 | 1 | 1 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Age class : | 1 | 1 | 1 | 2* | 2 | 3 | 3 | 3 |
| Measurement : |  |  |  |  |  |  |  |  |
| Greatest length of skull | - | - | - | - | - | 31,5 | - | - |
| Zygomatic breadth | - | - | - | - | - | 18,0 | - | - |
| Breadth of braincase | - | - | - | - | - | 14,1 | - | - |
| Height of braincase | - | - | - | - | - | 10,3 | - |  |
| Interorbital breadth | 3,5 | 3,6 | 3,6 | 3,6 | 4,1 | 3,4 | 3,5 | 3,2 |
| Length of nasals | 12,2 | 13,4 | - | 12,2 | - | 13,9 | 13,4 | - |
| Breadth of zygomatic plate | 3,9 | 4,0 | 4,1 | 3,9 |  | 4,1 | 4,0 | 4,1 |
| Length of diastema | 8,1 | 7,7 | 8,2 | 8,1 | 7,3 | 8,0 | 7,7 | 8,2 |
| Length of palate | 15,2 | 15,1 | 16,2 | 15,2 | 14,8 | 16,2 | 15,1 | 16,2 |
| Length of incisive foramina | 8,3 | 8,1 | 8,5 | 8,3 | 7,4 | 8,3 | 8,1 | 8,5 |
| Breadth of incisive foramina | 3,1 | 2,9 | 2,5 | 3,1 | 2,4 | 3,0 | 2,9 | 2,5 |
| Breadth of palatal bridge at M1/ | 2,6 | 2,4 | 2,4 | 2,6 | 2,6 | 3,0 | 2,4 | 2,4 |
| Breadth of palatal bridge posterior to M3/ | 4,4 | 4,5 | 4,1 | 4,4 | 4,1 | 4,5 | 4,5 | 4,1 |
| Length of bulla | - | - | - | - | - | 7,2 | - | - |
| Alveolar length of M1-3/ | 5,9 | 6,1 | 6,7 | 6,0 | 6,1 | 7,0 | 6,6 | 6,7 |
| Crown length of M1-3/ | 5,4 | 5,6 | 5,8 | 5,3 | 5,6 | - | 5,8 | 5,8 |

* holotype

Table 1. Malpaisomys insularis n. g., n. sp., skull measurements of specimens from Fuerteventura (localities 1 and 2; see fig. 1).
Locality $\quad$ Age class 1 $\quad$ Age class 2 $\quad$ Age class 3

## Upper toothrow length

Fuerteventura

Loc. 1
Loc. 2
Loc. 4
Lanzarote

Loc. 5
5,52 $\pm 0,19$ (4)
5,65
(2)

5,90
Loc. 6
$5,62 \pm 0,05$ (4)
$5,67 \pm 0,05$ (7)
$5,67 \pm 0,13$
Lower toothrow length
Fuerteventura

Loc. 1
Loc. 2
Loc. 3
Loc. 4
Lanzarote
Loc. 5
Loc. 6
Loc. 1
$5,37 \pm 0,14$ (25)
$5,39 \pm 0,15$ (25)
$5,34 \pm 0,17$
$5,35 \pm 0,13$ (4)
$5,49 \pm 0,12$
(7)
$5,37 \pm 0,17$ (16)
$5,44 \pm 0,19$ (25)
$5,44 \pm 0,16$ (50)
$5,90 \pm 0,10$
$5,60 \pm 0,08$ (4)
$5,61 \pm 0,13$ (6)
5,85

5,30
(2) 5,50
(1)
$5,60 \pm 0,26$
5,60
5,20
(1)
$(6)$

5,40
(2) 5,20
(2)

Mandible length
Fuerteventura
Loc. 1
Loc. 2
$16,68 \pm 0,60$
$16,75 \pm 0,45$
(25) $18,18 \pm 0,62(50)$
$18,58 \pm 0,52$
Loc. 3
17,00
(4) $17,26 \pm 0,44$
(5)
$18,00 \pm 0,75$
$17,07 \pm 0,92$ (4)
Loc. 6
-

Table 2. Malpaisomys insularis n.g., n. sp.; crown lengh of upper and lower toothrow and length of mandible without incisor. Measurements separate for age class 1 (no or little wear), 2 (teeth moderately wom) and 3 (heavily wom). Mean, standard deviatian and sample size indicated.

| Locality | M1／Length | M1／Width | M／1 Length | M／1 Width |
| :---: | :---: | :---: | :---: | :---: |
| Fuerteventura |  |  |  |  |
| Loc． 1 | 2，80土0，10（70） | 1，96 $\pm 0,08$（59） | 2，46 $\pm 0,08$（33） | 1，65 $\pm 0,07$（33） |
| Loc． 2 | 2，82 $\pm 0,24$（21） | 1，96 $\pm 0,09$（21） | 2，50 $\pm 0,08$（24） | 1，67 $\pm 0,09$（25） |
| Loc． 3 | 2，80 $\pm 0,14$（3） | 1，92土0，11（3） | 2，48 $\pm 0,09$（9） | 1，61 $\pm 0,07$（9） |
| Loc． 4 | 2，90（2） | 1，90（2） | 2，56（1） | 1，68（1） |
| Lanzarote |  |  |  |  |
| Loc． 5 | 2，72 $\pm 0,06$（9） | 1，91 $\pm 0,10$（10） | 2，44 $\pm 0,10$（9） | 1，56 $\pm 0,06$（9） |
| Loc． 6 | 2，69 $\pm 0,08$（16） | 1，90 50,08 （16） | 2，45 $\pm 0,06$（21） | 1，62 $\pm 0,08$（21） |
| Loc． 7 | 2，75 $\pm 0,02$（3） | 1，95 $\pm 0,06$（3） | 2，56（2） | 1，64（2） |
| Loc． 8 | 2，68（2） | 1，84（2） | － | － |
| Locality | M2／Length | M2／Width | M／2 Length | M／2 Width |
| Fuerteventura |  |  |  |  |
| Loc． 1 | 1，88 $\pm 0,07$（71） | 1，87 $\pm 0,06$（71） | 1，69 $\pm 0,08$（33） | 1，66 $\pm 0,06$（33） |
| Loc． 2 | 1，90 $\pm 0,08$（20） | 1，86 $\pm 0,07$（20） | 1，72 $\pm 0,09$（26） | 1，71 $\pm 0,09$（26） |
| Loc． 3 | 1，96（1） | 1，88（1） | 1，73 $\pm 0,03$（9） | 1，63 $\pm 0,07$（9） |
| Loc． 4 | 1，85（2） | 1，88（2） | 1，80（1） | 1，76（1） |
| Lanzarote |  |  |  |  |
| Loc． 5 | 1，87 $\pm 0,09$（9） | 1，81 $\pm 0,07$（9） | 1，68 $\pm 0,06$（8） | 1，61 $\pm 0,07$（8） |
| Loc． 6 | 1，85土0，07（16） | 1，83土0，06（16） | 1，76 $\pm 0,07$（20） | 1，65 $\pm 0,07$（20） |
| Loc． 7 | 1，90（2） | 1，82（2） | 1，83 $\pm 0,04$（4） | 1，73 $\pm 0,11$（4） |
| Loc． 8 | 1，84（2） | 1，78（2） | － | － |
| Locality | M3／Length | M3／Width | M／3 Length | M／3 Width |
| Fuerteventura |  |  |  |  |
| Loc． 1 | 1，43 $\pm 0,08$（69） | 1，36 $\pm 0,07$（69） | 1，32 $\pm 0,05$（33） | 1，31 $\pm 0,06$（33） |
| Loc． 2 | 1，43 $\pm 0,06$（14） | 1，32 $\pm 0,06$（14） | 1，28 $\pm 0,07$（17） | 1，30 $\pm 0,06$（15） |
| Loc． 3 | － | － | 1，30 $\pm 0,05$（5） | 1，23 $\pm 0,07$（5） |
| Loc． 4 | 1，50（2） | 1，40（2） | 1，40（1） | 1，44（1） |
| Lanzarote |  |  |  |  |
| Loc． 5 | 1，42 $\pm 0,08$（7） | 1，34 $\pm 0,06$（7） | 1，21 $\pm 0,02$（6） | 1，27 $\pm 0,03$（6） |
| Loc． 6 | 1，44土0，06（12） | 1，32 $\pm 0,05$（12） | 1，35 $\pm 0,09$（13） | 1，29 $\pm 0,06$（13） |
| Loc． 7 | － | － | 1，36（1） | 1，36（1） |

Table 3．Malpaisomys insularis n．g．，n．sp．，molar measurements；mean，standard deviation and sample size given．

| Locality | Total length | Length without <br> epiphysis | Minimum <br> diameter |
| :--- | :---: | :---: | :---: |

Humerus

| Fuerteventura |  |  |  |
| :---: | :---: | :---: | :---: |
| Loc. 1 | 16,47 $\pm 0,60$ (9) | $14,75 \pm 0,78$ (32) | 1,31 $\pm 0,10$ (41) |
| Loc. 2 | 16,00 $\pm 0,62$ (3) | $14,67 \pm 1,34$ (10) | 1,32 $\pm 0,13$ (13) |
| Loc. 3 | - | 15,10 $\pm 0,17$ (3) | 1,30 $\pm 0,10$ (3) |
| Loc. 4 | - | 15,8 (1) | 1,30 (1) |
| Lanzarote | - | $14,50 \pm 0,63$ (4) | 1,42 $\pm 0,10$ (4) |
| Loc. 6 | 14,9 (1) | 14,62 $\pm 0,49$ (4) | 1,32 $\pm 0,08$ (5) |

Femur
Fuerteventura

| Loc. 1 | 24,56 $\pm 0,99$ (8) | $21,67 \pm 1,13$ (61) | $1,84 \pm 0,13$ (69) |
| :---: | :---: | :---: | :---: |
| Loc. 2 | 23,05 (2) | $21,26 \pm 1,00$ (11) | 1,76 $\pm 0,13$ (13) |
| Loc. 3 | $\bigcirc$ | $20,95 \pm 1,47$ (4) | 1,85 $\pm 0,06$ (4) |
| Loc. 4 | - | $24,15 \pm 1,14$ (6) | 1,95 $\pm 0,08$ (6) |
| rote |  |  |  |
| Loc. 5 | - | 19,5 (2) | 1,65 (2) |
| Loc. 6 | - | $19,19 \pm 1,22$ (10) | 1,75 $\pm 0,07$ (10) |
| Loc. 7 | - | 18,8 (1) | 1,6 (1) |
| Loc. 8 | - | 20,8 (1) | 2,1 (1) |

Ulna
Fuerteventura
Loc. 1
$19,41 \pm 1,38(30) \quad-$
$19,06 \pm 1,69(11) \quad-$
$0,66 \pm 0,06$ (30)
Loc. 2
$19,06 \pm 1,69$ (11) -
$0,67 \pm 0,04$ (11)
Tibia
Fuerteventura

| Loc. 1 | - | $24,81 \pm 1,65(30)$ | $1,23 \pm 0,09(30)$ |
| :--- | :--- | :--- | :--- |
| Loc. 2 | $24,84 \pm 2,45(9)$ | $1,20 \pm 0,11$ (9) |  |

Table 4. Malpaisomys insularis n. g., n. sp. ; measurements of some postcranial elements. Mean, standard deviation and sample size indicated.

| Taxa | $\left\lvert\, \begin{array}{ccc} 1 & 35791113151719 \\ 2468101214161820 \end{array}\right.$ |
| :---: | :---: |
| Can. bra |  |
| Can. tamaran | 11011001000022201 |
| Mus. musculus | 00200010000021001 |
| Rat. rattus | 011121100000200000 |
| Mal. insularis | 002010101211201112 |
| Gra. dolichurus | 0021101021020020 |
| Oe. hypoxanthus | 02211210212112202 |
| Arv. niloticus | 01012011000022020 |
| Ac. cahirinus | 00200010000110010201 |
| Uranomys ruddi | 00200010000020010 |
| P. darelbeidae | 00001010211220020 |
| P. meini | 00101010200010010 |
| Occ. adroveri | 00200210211210012110 |
| Occ. monthele | 00200210221210012110 |
| Pro. cathalai | 00101110000001002100 |
| Ste. donnezan |  |
| Kar. darwini | 000010100000020001 |

Table 5. Matrix of the taxa and characters used for the phylogenetic analysis.

## BIBLIOGRAPHY

ABDEL-MONEM, A., WATKINS, N.D. \& GAST, P.W., 1967. - Volcanic history of the Canary Islands. Amer. Geophys. Union Trans., 48: 226-227.
AFONSO PEREZ, D.L. (Editor), 1980. - Atlas basico de Canarias. Editorial Interinsular Canaria, Barcelona, 80 p .
BRAVO, T., 1964. - El Volcán y el malpais de la Corona. La "Cueva de los Verdes" y los "Jameos". Publ. Cabildo Insular de Lanzarote, Arrecife: 1-21.
BROSSET, A., 1963. - Statut actuel des mammifères des îles Galapagos. Mammalia, 27: 323-338.
BRUIJN, H. de, DAWSON, M. \& MEIN, P., 1970. - Upper Pliocene Rodentia, Lagomorpha and Insectivora (Mammalia) from the Isle of Rhodes (Greece). I, II, III. Kon. Ned. Akad. van Wettensch., Proc. ser. B, 73 (5): 535-584, Amsterdam.
CALVO, J.P., ELIZAGA, E., LOPEZ-MARTÍNEZ, N., ROBLES, F. \& USSERA, J., 1978. - El Mioceno superior continental del Prebético Externo: Evolución del Estrecho Nordbético. Bol. Geol. Min., 89: 407-426.
CARRASCOSA, M.C. \& LOPEZ-MARTÍNEZ, N., 1988. - The house mouse from a prehistoric site in Fuerteventura (Canary Islands, Spain). Bonn. zool. Beitr., 39 (in press).
CLARK, D.A., 1984. - Native land mammals. In: R. PERRY (Ed), Key environments Galapagos: 225-231. Pergamon Press, Oxford.
CRUSAFONT-PAIRO, M. \& PETTER, F., 1964. - Un muriné géant fossile des îles Canaries Canariomys bravoi gen. nov., sp. nov. (Rongeurs, Muridés). Mammalia, 28: 608-611.

De GRAAF, G., 1961a. - A preliminary investigation of the mammalian microfauna in Pleistocene deposits of caves in the Transvaal system. Paleont. Afr., 7: 59-118.
De GRAAF, G., 1961b. - On the fossil mammalian microfauna collected at Kromdraai by Draper in 1895. S. Afr. J. Sci., 57: 259-260.

DENYS, C., 1987. - Rodentia and Lagomorpha 6.1. Fossil rodents (other than Pedetidae) from Laetoli. In: M.D. LEAKEY \& J.M. HARRIS (Eds.), The Pliocene site of Laetoli northern Tanzania: 118-170. Oxford University Press.
ELLERMAN, J.R., 1941. - The families and genera of living rodents. Vol. 2, p. 1-690.
FELSENSTEIN, J., 1985. - "Phylip", phylogeny inference package (version 2.8), University of Washington, Seattle.
GARRALDA, M. D., F. HERNANDEZ \& D., SANCHEZ, 1981. - El enterramiento de la cueva de Villaverde (La Oliva, Fuerteventura). Anuario de Estudios Atlánticos, 27: 673-690.
HERNANDEZ, F. \& SANCHEZ VELAZQUEZ, M.D., 1986. - Fuerteventura: Estado de la cuestión con especial referencia a la cueva de Villaverde. Congreso de Cultura de Canarias, Sec. Arg. Prehistoria, Publ. Gob. Aut. Canarias.
HERNANDEZ-PACHECO, E., 1909. - Estudio geológico de Lanzarote y de las Isletas Canarias. Mem.R. Soc. Esp. Hist. nat., 6: 107-342.
HONACKI, J.H., KINMAN, K. E. \& KOEPPL, J.W., 1982. - Mammal species of the world. Allen Press and A.S.C., Lawrence, 694 p.
HUTTERER, R., LOPEZ-JURADO, L.F. \& VOGEL, P., 1987. - The shrews of the eastern Canary Islands: a new species (Mammalia: Soricidae). J. nat. Hist., 21: 1347-1357.
JAEGER, J.J., 1975. - Les Muridae (Mammalia, Rodentia) du Pliocène et du Pléistocène du Maghreb. Origine; Evolution; Données biogéographiques et paléoclimatiques. Thèse Doct. ès Sciences, Université de Montpellier II (U.S.T.L.), 124 p.
JAEGER J.J., MICHAUX, J. \& THALER, L., 1975. - Présence d'un rongeur muridé nouveau, Paroethomys miocoenicus nov. sp., dans le Turolien supérieur du Maroc et d'Espagne. Implications paléogéographiques. C.R. Acad. Sc., Paris, 280 (D): 1673-1676.
JACOBS, L.L., 1978. - Fossil rodents (Rhizomyidae and Muridae) from Neogene Siwalik deposits, Pakistan. Museum of Northern Arizone Press, Bull. Ser., 52: 1-103.
LOPEZ-MARTÍNEZ, N., 1978. - Cladistique et paléontologie. Application à la phylogénie des Ochotonidés européens (Lagomorpha, Mammalia). Bull. Soc. géol. France, 20: 821-830.
LOPEZ-MARTÍNEZ, N. \& LOPEZ-JURADO, L.F., 1987. - Un nuevo murido gigante del Cuaternario de Gran Canaria Canariomys tamarani nov. sp. (Rodentia, Mammalia). Doñana, Publ. ocas., 2: 1-66.
MADDISON, W. \& MADDISON, D., 1987. - Mac Clade, version 2.1. A phylogenetics computer program distributed by the authors. Museum of Comparative Zoology, Harvard University.
MECO CABRERA, J., HERNANDEZ, F. \& SANCHEZ VELAZQUEZ, D., 1982. - La cueva de Villaverde (Fuerteventura) y su mastologia (Nota preliminar). Libro homenaje a Jesús Arencibia, Servicio de Publicaciones de la Escuela Universitaria de Formación del Profesorado de E.G.B. de Las Palmas, Las Palmas de Gran Canaria: 187-194.
MICHAUX, J., 1967. -- Origine du dessin dentaire Apodemus (Rodentia, Mammalia). C.R. Acad. Sc., Paris, 264 (D): 711-714.
MICHAUX, J., 1969. - Muridae (Rodentia) du Pliocène supérieur d'Espagne et du Midi de la France, Palaeovertebrata, Montpellier, 3 (1): 1-24.
MICHAUX, J., 1971. - Muridae (Rodentia) néogènes d'Europe sud-occidentale. Evolution et rapports avec les formes actuelles. Paléobiologie continentale, Montpellier, 2 (1): 1-67.
MUSSER, G., HEANEY, L.R. \& RABOR, D.S., 1985. - Philippine rats: a new species of Crateromys from Dinagat Island. Amer. Mus. Novit., no. 2821, p. 1-25.
PATTON, J.L. \& HAFNER, M.S., 1983. - Biosystematics of the native rodents of the Galapagos archipelago, Ecuador. In: R.I. BOWMAN, M. BERSON \& A.E. LEVITON (Eds.), Patterns of Evolution in Galápagos Organisms: 539-568. Amer. Association for the Advancement of A Science, Pacific Division.

PETTER, F., 1984. - Eléments d'une révision des Acomys africains. Un sous-genre nouveau, Peracomys PETTER \& ROCHE, 1981 (Rongeurs, Muridés). Ann. Kon. Mus.Mid. Afr., Zool. Wetensch., 237: 109-119.
SEN, S., 1977. - La faune de Rongeurs pliocènes de Calta (Ankara, Turquie). Bull. Mus. natn. Hist. nat., Paris, 3ème sér., n ${ }^{\circ} 465$, Sci. Terre 91: 65-75.
STEADMAN, D.W. \& RAY, C.E., 1982. - The relationships of Megaoryzomys curioi, an extinct cricetine rodent (Muroidea: Muridae) from the Galápagos Islands, Ecuador. Smiths. Contr. Paleobiol., 51: 1-23.
WESSELMAN, H.B., 1984. - The Omo micromammals. Systematics and paleoecology of early man sites from Ethiopia. Contr. Vertebr. Evol., 7: 1-219.
WESSELS, W., BRUIJN, H. de, HUSSAIN, S.T. \& LEINDERS, J. H.-M., 1982. - Fossil rodents from the Chinji Formation, Banda Daub Shah, Kohat, Pakistan. Kon. Ned. Akad. van Wetensch., Proc. series B, 84 (1): 71-99.

## LEGENDS OF PLATES

PLATE 1
Upper and lower molars of Malpaisomys insularis n. g., n. sp.

1. Upper toothrow (Lanzarote, loc. 6)
2. Upper toothrow (Fuerteventura, loc. 2)
3. Upper toothrow (Fuerteventura, loc. 2)
4. Lower toothrow (Fuerteventura, loc. 2)
(specimens in the Museum Bonn).

## PLATE 2

Some skeletal elements of Malpaisomys insularis n. g., n. sp. from Fuerteventura (loc. 2).

1. Atlas ; a, caudal view; $b$, dorsal view
2. Axis; a, cranial view; b, dorsal view
3. Lumbar vertebra; $a$, caudal view; $b$, lateral view
4. Sacrum ; a, dorsal view; $b$, ventral view
5. Scapula; a, distal view; b, dorsal view
6. Humerus ; $a$, anterior view; $b$, medial view
7. Ulna; $a$, lateral view; $b$, anterior view
8. Pelvis; lateral view
9. Femur ; $a$, anterior view; $b$, lateral view
10. Tibia; $a$, lateral view; $b$, posterior view (Specimens in the Museum Bonn).


