

## Exploring the landscape and climatic conditions of Neanderthals and anatomically modern humans in the Middle East: the rodent assemblage from the late Pleistocene of Kaldar Cave (Khorramabad Valley, Iran)

Ivan Rey-Rodríguez, Juan Manuel Lopez-García, Blain Hugues-Alexandre, Emmanuelle Stoetzel, Christiane Denys, Mónica Fernández-García, Laxmi Tumung, Behrouz Bazgir

#### ▶ To cite this version:

Ivan Rey-Rodríguez, Juan Manuel Lopez-García, Blain Hugues-Alexandre, Emmanuelle Stoetzel, Christiane Denys, et al.. Exploring the landscape and climatic conditions of Neanderthals and anatomically modern humans in the Middle East: the rodent assemblage from the late Pleistocene of Kaldar Cave (Khorramabad Valley, Iran). Quaternary Science Reviews, 2020, 236, pp.106278. 10.1016/j.quascirev.2020.106278. hal-02568534

## HAL Id: hal-02568534 https://hal.science/hal-02568534

Submitted on 10 Nov 2020

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial - NoDerivatives 4.0 International License

1 Exploring the landscape and climatic conditions of Neanderthals and

2 anatomically modern humans in the Middle East: the rodent assemblage

#### from the late Pleistocene of Kaldar Cave (Khorramabad Valley, Iran)

3 4

Iván Rey-Rodríguez <sup>a,b</sup>\*, Juan-Manuel López-García <sup>c,d</sup>, Hugues-Alexandre
 Blain <sup>c,d</sup>, Emmanuelle Stoetzel <sup>a</sup>, Christiane Denys <sup>e</sup>, Mónica Fernández-García
 <sup>b,a</sup>, Laxmi Tumung <sup>c,d,a</sup>, Andreu Ollé <sup>c,d</sup>, Behrouz Bazgir <sup>d</sup>

8

<sup>a</sup> HNHP UMR 7194, CNRS / Muséum national d'Histoire naturelle / UPVD / Sorbonne
 Universités, Musée de l'Homme, Palais de Chaillot, 17 place du Trocadéro, 75016 Paris, France
 <sup>b</sup> Sezione di Scienze Preistoriche e Antropologiche, Dipartimento di Studi Umanistici, Università
 degli Studi di Ferrara, C.so Ercole I d'Este, 32 - 44121 Ferrara, Italy

<sup>c</sup> Institut Català de Paleoecologia Humana i Evolució Social (IPHES). Zona Educacional 4,
 Campus Sescelades URV (Edifici W3) 43007 Tarragona, Spain

<sup>d</sup> Área de Prehistòria, Universitat Rovira i Virgili. Facultat de Lletres, Avinguda Catalunya 35,
 43002 Tarragona, Spain

<sup>e</sup> ISYEB UMR 7205, CNRS / Muséum national d'Histoire naturelle / UPMC / EPHE / Sorbonne
 Universités, Paris, France

\* corresponding author: ivan.rey-rodriguez@edu.mnhn.fr / ivanreyrguez@gmail.com (lván
 Rey-Rodríguez)

21

#### 22 Abstract

The Middle East, specially the Zagros region, lies in a strategic position as a crossroads between Africa, Europe and eastern Asia. The landscape of this region that prevailed around the Neanderthal and anatomically modern human occupations is not well known. Only a few sites have been studied in detail in this area, often providing only a faunal list. These reveal that Neanderthals and anatomically modern humans lived in a landscape mainly composed of dry steppes.

Here we extend the data obtained from Kaldar Cave through a systematic study of the rodent assemblage. The site provided evidence of a Pleistocene occupation attested by lithic tools associated with the Middle and Upper Palaeolithic, but it was also occupied during the Holocene, as evidenced by Neolithic artefacts. First excavations have revealed small vertebrates in Layer 4 (sub-layer 5 and 5II), belonging to the Upper Palaeolithic, and Layer 5 (sublayers 7 and 7II), belonging to the Middle Palaeolithic.

The rodent assemblage of Kaldar Cave is mainly composed of six arvicoline, two cricetine, one glirid, one dipodid, one gerbilline and two murine species.

This assemblage shows that during the Late Pleistocene the environment around the site was mainly composed of open dry steppes, as indicated by the most abundant taxa, *Microtus*, *Ellobius* and *Meriones*. However, murine species indicate the presence of a vegetation cover. The palaeoclimatic conditions are characterised by lower temperatures and also less precipitation than at present.

The results obtained with the rodent assemblages show that there is no major palaeoenvironmental or palaeoclimatic change that would explain the 46 cultural shift between Layer 5 (Middle Palaeolithic) and Layer 4 (Upper47 Palaeolithic).

48

49 **Keywords**: Rodentia, Middle East, taxonomy, palaeoecology, human 50 occupations

- 51
- 52

#### 53 **1. Introduction**

54

Small vertebrates are recognized to be good palaeoecological indicators due 55 56 to their rapid evolution, the limited geographic range of a species and unique niche requirements, as well as their frequent preservation in archaeological and 57 palaeontological sites. They might be excellent indicators of palaeovegetation 58 type and provide a high-resolution proxy for palaeoenvironmental changes (e.g. 59 60 Belmaker and Hovers, 2011). Studies of small vertebrates from archaeological sites in the Middle East are still scarce (Belmaker and Hovers, 2011; Belmaker 61 et al., 2016; Demirel et al., 2011; Fernández-Jalvo, 2016a; Kandel et al., 2017; 62 Maul et al., 2015a, 2015b; Smith et al., 2015; Weissbrod and Zaidner, 2014), 63 64 and they mainly propose preliminary lists of taxa. On the other hand, some 65 studies of extant owl pellet assemblages have been performed in the Middle East (Abi-said et al., 2014; Darvish et al., 2000; Haddadian Shad et al., 2014; 66 Kopij and Liven-schulman, 2013; Obuch and Khaleghizadeh, 2012; Rey-67 Rodríguez et al., 2019; Shehab et al., 2013), providing taxonomic and some 68 taphonomic reference data in Iran, Lebanon, Israel, Turkey and Syria. 69

The main palaeoecological and palaeoclimatic reconstructions based on faunal assemblages relate to the unique niche requirements of a particular species. Belmaker *et al.* (2011) pointed out that their interpretations, in contrast, are based on analyses of community composition. Analyses of persistence *versus* change over time should take into account species presence-absence, rank abundance and proportional abundance (Belmaker and Hovers, 2011).

The Middle East is a prime location to study Neanderthal and anatomically 76 77 modern human (AMH) interactions during the Middle to Upper Palaeolithic transition (Belmaker et al., 2016). However, sites yielding evidence of both 78 Neanderthal and AMH occupations in the same sequence are rare, only five 79 sites were well excavated (Bazgir et al., 2014; Bazgir et al., 2017). In this 80 framework, Kaldar Cave is a key archaeological site for evaluating the influence 81 of environment on cultural changes in both human species because of its 82 location and the faunal and lithic remains. The main objective of this paper is to 83 infer the landscape composition and the climatic conditions in which 84 Neanderthal and AMH populations pursued their activities through the study of 85 small mammals. We identify and describe the rodent fossils recovered from 86 Kaldar Cave in order to establish the main criteria for further identifications in 87 this region and provide a preliminary basis for future systematic studies. 88

89

90

#### 91 2. Kaldar Cave

92

Kaldar Cave is located in the northern part of the Khorramabad valley, 93 Lorestan Province, western Iran (48º17'35" E, 33º33'25" N) at 1290 m a.s.l. 94 95 (Fig. 1a,b). The cave was discovered in 2007, and the first archaeological intervention was undertaken by an Iranian-Spanish team in 2011-2012, 96 revealing the great potential of the cave (Bazgir et al., 2014). A second 97 excavation was performed in 2014-2015. The trench exposed an approximately 98 2m-thick section of sedimentary deposits, characterized by 5 main layers, 99 100 themselves divided into several sub-layers (Fig. 1c).

Layer 5 (including sub-layers 7 and 7II) consists of an extremely cemented reddish-brown sediment with some small angular limestone blocks and Middle Palaeolithic artefacts with Levallois elements. So far, no chronometric data are available for this layer (Bazgir et al., 2017).

Layer 4 (including sub-layers 5, 5II, 6 and 6II) consists of a silty but compact 105 dark-brown sediment with cultural remains attributed to the Upper and early 106 Upper Palaeolithic. In the uppermost parts of this layer, two fireplaces made of 107 108 clay were recovered and dated through thermoluminescence, yielding ages that 109 ranged from 23,100 ± 3 300 to 29,400 ± 2 300 BP (Bazgir et al., 2017). The dates obtained show that these fireplaces were made or re-used from existing 110 older sediment from the upper part of this layer in the later stages of the Upper 111 Palaeolithic. AMS radiocarbon dates of 38,650-36,750 cal BP, 44,200-42,350 112 113 cal BP, and 54,400–46,050 cal BP have been obtained from charcoal material located below this layer (Bazgir et al., 2017; Becerra-Valdivia et al., 2017). 114

Layers 1 to 3 (including sub-layers 4 and 4II) consist of ashy sediments of a 115 blackish-green colour containing both thick and thin angular limestone clasts. 116 117 These layers varied in thickness from 60 to 90 cm and contained many phases dating to the Holocene: Islamic and historical eras, Iron Age, Bronze Age, 118 Chalcolithic and Neolithic. However, due to the presence in these layers of 119 bioturbation mainly by burrowing animals, the phases were recognized only by 120 121 a preliminary study of the potsherds, metal artefacts and some diagnostic lithic 122 artefacts from the lower lavers.

This site provides evidence for the Middle to Upper Palaeolithic transition in the Middle East (Bazgir et al., 2017). Moreover, Kaldar Cave has yielded the oldest evidence of *Prunus* spp. in the area, through the study of thirty charcoal remains (Allué et al., 2018), that helps us to reconstruct the environmental conditions, evidencing a wooded vegetation.

128

#### 129 **3. Material and methods**

130

The rodent fossil remains used in this study, 1112 molars, come from the archaeological excavation campaigns carried out in Kaldar Cave in 2011-2012 and 2014-2015. The samples comprise disarticulated bones and isolated teeth that were collected in the field by water screening using superimposed 5 and 0.5 -mm mesh screens.

137

#### 138 **3.1. Taxonomy**

139

In this article we only focus on rodents, because these are one of the most useful tools for palaeoenvironmental and palaeoclimatic reconstructions of archaeological sites, and the other material, as amphibians, squamates, birds, shrews and bats, will be studied later. The taxonomic identification of the rodent remains is based mainly on molar morphology and measurements of the lower m1, the taxonomically most diagnostic tooth.

The remains were identified at a specific level whenever possible, thanks to 146 147 comparisons with reference collections from the Natural History Museum of London (NHM) and comparative morphological and biometric data from the 148 literature, notably for Microtus (Coskun, 2016; Kryštufek and Shenbrot, 2016; 149 Kryštufek and Vohralík, 2009; López-García, 2011; Rusin, 2017; Shenbrot, 150 Kryštufek et al. 2016; Tesakov, 2016), Cricetulus (Bogicevic et al., 2011; 151 Kryštufek et al., 2017; Sándor, 2018), Mesocricetus (Kryštufek and Vohralík, 152 2009), Meriones (Coşkun, 1999; Darvish, 2011; Dianat et al., 2017; Kryštufek 153 and Vohralik, 2009; Souttou and Denys, 2012; Stoetzel et al., 2017), Allactaga 154 (Karami et al., 2008; Shenbrot, 2009), Myomimus (Gerrie and Kennerley, 2017; 155 Karami et al., 2008; Kennerley and Kryštufek, 2019), Apodemus (Amori et al., 156 157 2016; Bogicevic et al., 2011; Knitlová and Horáček, 2017; Kryštufek and Vohralík, 2009; López-García, 2008) and Mus (Darviche et al., 2006; Kryštufek 158 and Vohralík, 2009; Siahsarvie and Darvish, 2008). 159

Quantification of taxonomic frequencies was based on standard measures in zooarchaeological analyses, including the number of identified specimens (NISP) and minimum number of individuals (MNI) (Weissbrod and Zaidner, 2014). This latter was estimated using the most abundant skeletal element present in the assemblage (molars in our case).

165

#### 166 **3.2. Taphonomic remarks**

167

A preliminary study was performed on the Kaldar Cave micromammal 168 remains. This was based on the systematic descriptive method that examines 169 the modifications of prey bones induced by predation, focusing on the degree of 170 171 digestion observed in teeth during the identification (Andrews, 1990; Fernández-Jalvo et al., 2016). This study showed a significant number of 172 digested teeth (Bazgir et al., 2017), indicating that predation activity was the 173 174 origin of at least part of the accumulation. According to the different degrees of digestion (mostly light, moderate and, heavy in a few cases) observed in the 175 176 molars, the predator responsible for this accumulation could be a category 3 177 predator such as the tawny owl (Strix aluco) or the Eurasian eagle owl (Bubo *bubo*). Both species are currently present in the area, have opportunistic
hunting habits, and are sedentary, so their prey spectrum is assumed to be a
good representation of the ecosystem in which they live.

- 181
- 182 183

185

#### 3.3. Palaeoenvironmental and palaeoclimatic reconstructions

184 3.3.1. Habitat weighting method

186 Palaeoecological interpretations derived from faunal data are based on analyses of community composition (Belmaker and Hovers, 2011). The method 187 188 used for the palaeoenvironmental reconstruction is the habitat weighting method, also named the taxonomic habitat index (Evans et al., 1981; Andrews, 189 2006; modified by Blain et al., 2008; López-García et al., 2011), which is based 190 on the current distribution of each taxon in the habitat(s) where it can be found 191 192 nowadays. We assume that the Kaldar Cave species had equivalent ecological requirements to their present-day relatives, because the Late Pleistocene 193 194 species are the same as those today and no extinct fossil species are found in the assemblage. We adapt the method to our studied area, differentiating the 195 196 following types of habitats: Forest (Fo), a large area covered with trees; 197 Shrubland (Sh), vegetation dominated by shrubs; Grassland (Gr), an open area 198 covered with grass; Desert (De), an area with little precipitation and no vegetation cover; Wetland (We), an area where water covers the soil; Steppe 199 (St), a dry grassy plain; and Rocky (Ro), a rocky or stony substrate. Each 200 201 species has a score of 1.00, which is divided between the habitats where the 202 species can be found at present (Table 1), all the species ranges were taken **IUCN** Red Threatened 203 from the List of Species (https://www.iucnredlist.org/resources/spatial-data-download). 204

205

207

206 3.3.2. Bioclimatic Model

In order to reconstruct the climate at Kaldar Cave, we applied the *bioclimatic* 208 209 model (BM), which was developed by Hernández-Fernández (2001) on the basis of the faunal spectrum, assuming that small- and large-mammal species 210 can be ascribed to ten different climates (Hernández-Fernández, 2001; 211 Hernández-Fernández and Peláez-Campomanes, 2003; Hernández-Fernández 212 et al., 2007). It was first necessary to calculate the climatic restriction index 213  $(CRI_{=1/n}, where i is the climatic zone where the species appear and n is the$ 214 215 number of zones where the species is present) and the bioclimatic component  $(BC_i = (\Sigma CRI_i) 100/S$ , where *i* is the climatic zone and S is the number of 216 species). To BC a mathematical model is applied as a regression that allows us 217 to calculate climatic parameters (see Appendix 1). 218

The different climatic groups defined by Hernández-Fernández (2001) and Hernández-Fernández et al. (2007) are: I *equatorial*; II *tropical with summer rains*; II/III *transition tropical semiarid*; III *sub-tropical arid*; IV, *subtropical* with winter rains and summer droughts; V, *warm temperate* with not very severe winters but high humidity; VI, *typical temperate* with winters that are cold but not very long, but summers that are cool; VII, *arid-temperate* with large temperature contrasts between winter and summer; VIII, *cold-temperate* with cool summers and long cold winters (boreal) and IX, *artic* (Table 2).

By means of the *BM* we were able to estimate the mean annual temperature (MAT), the mean temperature of the coldest month (MTC), the mean temperature of the warmest month (MTW) and the mean annual precipitation (MAP). This method also allowed us to assess whether there were changes in the temperature and precipitation regimes during the Middle to Upper Palaeolithic transition at Kaldar Cave.

- 233
- 234 4. Results and discussion
- 235

#### 236 4.1. Taxonomic identifications

237

243

244 245

246

249

254

In total 264 minimum number of individuals (MNI) were identified (Table 3 and Fig. 2). Some differences were found with respect to the preliminary list previously published by Bazgir et al. (2017). We notably found some new specimens and reconsidered the identifications for *Ellobius talpinus*, *Calomyscus* sp. and *Dryomys* cf. *nitedula*.

#### Order Rodentia Bowdich, 1821 Rodentia indet.

There are 14 teeth (11 in Layer 4 and 3 in Layer 5) that have not been identified to the family or genus level because they were broken.

# 250Family Cricetidae Fisher, 1817251Subfamily Arvicolinae Gray, 1821252Genus Microtus Schrank, 1798253Microtus spp.

Material: 554 isolated teeth. Layer 1-3: four isolated teeth; three left lower
m1 and one right lower m1. Layer 4: 258 isolated teeth; 81 m2; 101 m3, 13
right upper M1, 31 right lower m1; four left upper M1 and 28 left lower m1.
Layer 5: 292 isolated teeth; 119 m2; 53 m3; nine left upper M1, 51 left lower
m1; 10 right upper M1; 50 right lower m1.

260 261

262

### Microtus socialis Pallas, 1773

Material: 74 isolated teeth. Layer 1-3: three isolated teeth, three right lower m1. Layer 4: 36 isolated teeth; 20 right lower m1 and 16 left lower m1. Layer 5: 35 isolated teeth; 20 right lower m1 and 15 left lower m1.

266	
267	<u>Microtus irani Thomas, 1921</u>
268	
269	Material: 15 isolated teeth. Layer 1-3: one isolated tooth, one right lower m1.
270	Layer 5: 14 isolated teeth; nine right lower m1 and five left lower m1.
271	
272	Microtus guentheri Danford & Alston, 1880
273	
274	Material: 17 isolated teeth. Layer 1-3: one isolated tooth, one left lower m1.
275	Layer 4: 16 isolated teeth; nine right lower m1 and seven left lower m1.
276	Description and discussion: all the molars are hypsodont and arhizodont
277	with crown cementum in the re-entrant angles. The enamel differentiation is
278	Microtus-like. Six extant species of Microtus have been identified in Iran:
279	Microtus arvalis, Microtus irani, Microtus kermanensis, Microtus socialis,
280	Microtus transcaspicus and Microtus guentheri (Aşan Baydemir and Duman,
281	2009; Golenishchev et al., 2019; Firouz, 2005).
282	Microtus m1 are characterized by four buccal and five lingual re-entrant angles
283	(Kryštufek and Vohralík, 2009; Tsytsulina et al., 2017) with a posterior lobe
284	(PL), seven triangles (T) and an anterior cap (AC). We identified the Kaldar
285	specimens on the basis of their measurements (Table 4), the arrangement and
286	morphology of the triangles, and the AC (which may or may not be connected
287	with the other elements).
288	In our sample we identified Microtus spp., Microtus socialis, Microtus guentheri
289	and Microtus irani.
290	In <i>Microtus socialis</i> , the triangles from T1-T5 are closed, but T6-T7 are parallel
291	and broadly confluent with the AC. Microtus guentheri is distinguishable from
292	Microtus socialis by the rounded shape of the AC, which is of an arhombomorph
293	type (Aşan Baydemir and Duman, 2009), and also by T6-T7, which are
294	separated from the AC. In accordance with the measurements given in Aşan
295	Baydemir et al. (2009), the size of Microtus guentheri and Microtus socialis is
296	similar. In <i>Microtus irani</i> , triangles T1-T5 are closed, but T6-T7 are open and not
297	parallel with one another; moreover, Microtus irani is clearly larger than the
298	other species considered in our reference collection and also in our sample.
299	Finally, we include in <i>Microtus</i> spp. all the elements that we cannot attribute to
300	one species or another, in particular because of their bad state of preservation
301	(due to digestion, breakage).
302	Habitat and distribution: Microtus irani occupies grasslands in Iran, Turkey
303	and the Caucasus (Kryštufek and Kefelioğlu, 2001).
304	Microtus socialis is found in Russia, Ukraine, the Caucasus, Transcaucasia,
305	Turkestan, Iran and Afghanistan (Tsytsulina et al., 2017), although the range is
306	fragmented. It is a highly colonial species, found in steppe habitats and also in
307	agricultural lands, but extending also to bushy scrubs and uncultivated
308	mountain valleys, as well as to open oak forests on dry hillsides (Tsytsulina et
309	al., 2017).

*Microtus guentheri* occurs from the southeastern Balkans and Turkey through Syria, Lebanon and Israel, with an isolated range segment in northern Libya, Iran and some parts of Europe (Aşan Baydemir and Duman, 2009). This species is present in dry grasslands with sparse vegetation (Amr, 2015).

314

315

316 317

#### <u>Genus Chionomys Miller, 1908</u> <u>Chionomys nivalis Martins, 1842</u>

Material: 25 isolated teeth. Layer 1-3: two isolated teeth; one left lower m1 and one right lower m1. Layer 4: 17 isolated teeth; 12 right lower m1; five left lower m1. Layer 5: six isolated teeth; three left lower m1; three right lower m1.

**Description and discussion:** in our specimens it can be observed that the 321 first lower molars display five triangles and that the morphology of the 322 anteroconid complex (AC) is characteristic of the *nivalis* morphotype, where 323 324 triangles T6 and T7 are absent and the anterior cap is of an arrowhead or oval shape, inclined towards the labial part. The enamel is of the *Microtus* type with 325 cement (Krystufek, 2017; Kryštufek and Vohralík, 2009; López-García, 2011). 326 These specimens differ from Microtus in that they only have five triangles, and 327 328 the morphology of the AC is the other main characteristic of this species that 329 helps us to differentiate it from *Microtus*.

Habitat and distribution: *Chionomys nivalis* has a global distribution extending from southwestern Europe through southeastern Europe to the Caucasus, Turkey, Israel, Lebanon, Syria and Iran. In Iran it is distributed in the north and also in the west (Shenbrot and Krasnov, 2005; Krystufek, 2017). This is the only *Chionomys* species occurring today in Iran, but two other species (*C. gud, C. roberti*) are represented a little further north (northeastern Turkey, southern Georgia).

Regarding its habitat, it is present in open rocky areas, typically above the tree line and with scarce vegetation cover (Amori, 1999).

#### Genus Ellobius Fischer, 1814 Ellobius spp.

Material: 173 isolated teeth. Layer 1-3: two isolated teeth; two left lower m1.
Layer 4: 91 isolated teeth; seven indet.; eight right lower m1; five left lower m1;
11 upper M1; 43 m2; 17 m3. Layer 5: 80 isolated teeth; seven left lower m1;
three upper M1; 45 m2; 25 m3.

347 348

349

339 340

341

342

#### Ellobius fuscocapillus Blyth, 1843

Material: 11 isolated teeth. Layer 1-3: one isolated tooth; one right lower m1. Layer 4: five isolated teeth; two right lower m1; three left lower m1. Layer 5: five isolated teeth; four left lower m1 and one right lower m1.

353

354 355

Material: 20 isolated teeth. Layer 4: 10 isolated teeth; seven right lower m1 and three left lower m1. Layer 5: 10 isolated teeth; four left lower m1 and six right lower m1.

359 **Description and discussion:** 204 isolated teeth show the typical traits of the 360 genus Ellobius (Miller, 1896; Hinton, 1962, Kretzoi, 1969; Coşkun, 2016; Kryštufek & Shenbrot, 2016; Kryštufek and Vohralík, 2009; Rusin, 2017; 361 Shenbrot et al., 2016; Tesakov, 2016). Ellobius molars are notably 362 characterized by the presence of roots that are well visible in adults and old 363 364 individuals, but not always apparent in young specimens (Coskun, 2016). Moreover, Ellobius molars lack cement in the re-entrant angles. The Ellobius 365 m1 is composed of the anterior cap (AC), five triangles (T) with three buccal and 366 367 four lingual salient angles, and one posterior lobe (PL). Both M3 and m3 are 368 reduced and smaller than the other molars, with three triangles on the labial side and two triangles on the lingual side. M1 has three inner and outer folds, 369 370 and M2 and M3 have two inner and two outer re-entrant folds. The first and second upper molars have three triangles on the lingual and labial sides. 371 372 However, the re-entrant angle between the first and the second triangles on the 373 lingual side is more superficial (Gharkheloo and Kıvanç, 2003).

In Iran, the genus *Ellobius* is currently represented by three species: *Ellobius fuscocapillus*, *Ellobius lutescens* and *Ellobius talpinus* (Firouz, 2005; Kryštufek & Shenbrot, 2016; Kryštufek and Vohralík, 2009; Moradi Gharkheloo, 2003; Rusin, 2017; Shenbrot et al., 2016). The m1 is quite similar among these species, but there are some differences that can help us to differentiate them.

The AC is broad in *Ellobius lutescens*, narrow in *Ellobius talpinus* and elongated in *Ellobius fuscocapillus* (Maul et al., 2015b). The distance between T4 and T5 (W) and the total length (L) differ among the species (Table 5). We have observed in modern specimens from the Natural History Museum of London that *Ellobius fuscocapillus* is the largest and *Ellobius talpinus* the smallest.

In our sample we identify as Ellobius spp. those m1s that are too broken or 384 385 digested to distinguish, as well as all the m2, m3, M1, M2 and M3 because of 386 the lack of discriminant characters. The identification of Ellobius lutescens and Ellobius fuscocapillus in the Kaldar material is based on the morphology of the 387 AC and the measurements (L and W). We do not identify Ellobius talpinus in our 388 sample, as no specimen displays morphological features characteristic of this 389 species, such as the narrow AC, and because the measured specimens are 390 larger than E. talpinus. 391

Habitat and distribution: *Ellobius* species frequent steppes, grasslands and
semi-deserts in eastern Europe and central Asia; these fossorial species are
specialised in subterranean life (Coşkun, 2016; Kryštufek and Vohralík, 2009). *Ellobius lutescens* is a Palaearctic species distributed across Iran, Iraq,
Azerbaijan, Armenia, Transcaucasia and East Anatolia (Thomas, 1905;
Ellerman and Morrison-Scott, 1951; Darlington, 1957; Osborn, 1962; Walker,

1964; Lay, 1967; Hassinger, 1973; Roberts, 1977; Corbet, 1978; Corbet and
Hill, 1991; Wilson and Reeder, 2005; Coşkun, 1997, 2016; Nowak, 1999;
Kryštufek & Shenbrot, 2016). In Iran, this species is found in mountain
grasslands, sandy semi-deserts and steppe areas (Kryštufek & Shenbrot, 2016;
Tesakov, 2016). *Ellobius fuscocapillus* shows a wide range across eastern Iran,
Turkmenistan, Afghanistan and Pakistan. In Iran it is found in open steppes with
loose soil (Shenbrot, Kryštufek & Molur, 2016).

- 405 406
- 407

## 407

409

#### Subfamily Cricetinae Fischer, 1817 Genus Cricetulus Milne-Edwards, 1867 Cricetulus migratorius Pallas, 1773

Material: 19 isolated teeth. Layer 1-3: one isolated tooth; one right lower m3.
Layer 4: 13 isolated teeth; three left lower m1, two left upper M2; three right
lower m1, three right upper M1 and two right upper M2. Layer 5: five isolated
teeth; three left lower m1 and two right lower m1.

Description and discussion: the first molars (m1 and M1) are brachyodont 414 and cuspidate, with two longitudinal series of cusps. Each series of cusps 415 416 consists of three pairs. The m1 and M1 are the largest and the m3/M3 the 417 smallest. The lower m3 only has two pairs of cusps (Kryštufek and Vohralík, 2009). We identify Cricetulus migratorius in all the layers of Kaldar Cave in 418 419 accordance with the measurements and identification keys for molars based on the morphology and arrangement of the tubercles and cusps provided by 420 421 Kryštufek and Vohralík (2009). We also draw comparisons with the reference 422 collection from Iran, Afghanistan and Azerbaijan housed in the Natural History Museum of London. The grey hamster, or migratory hamster, is the smallest 423 hamster species (Bogicevic et al., 2011; Sándor, 2018). 424

Habitat and distribution: *Cricetulus migratorius* extends from eastern Europe through Russia and central Asia to Mongolia and western China (Kryštufek et al., 2017; Kryštufek and Vohralík, 2009). In Iran, this species is found all over the country. The habitats of this species are mostly dry grasslands, steppes and semi-deserts. Arid areas with relatively sparse vegetation are preferred (Kryštufek et al., 2017; Maul et al., 2015a).

431

432

- 433
- 434

#### <u>Genus Mesocricetus Nehring, 1898</u> Mesocricetus brandti Nehring, 1898

Material: 10 isolated teeth. Layer 4: three isolated teeth; one left upper M1,
one right lower m1 and one right upper M1. Layer 5: seven isolated teeth; four
left upper M1, one right upper M2 and two right upper M1.

438 **Description and discussion**: the specimens from Kaldar Cave are 439 attributed to *Mesocricetus brandti* on the basis of the size and morphology of 440 the teeth. The molars present a similar morphological pattern to *Cricetulus* 441 *migratorius*, but are significantly larger in size. The first molars have six tubercles, the second and third molars only four. The largest molars are m1 and
M1, whereas m2/M2 and m3/M3 are reduced (Kryštufek and Vohralík, 2009). In
Iran we also find *Mesocricetus raddei*, which presents a similar morphology of
the teeth to *Mesocricetus brandti*, but compared with the NHM reference
collection of the latter, *Mesocricetus raddei* is bigger

Habitat and distribution: Mesocricetus brandti has the largest distributional 447 448 area of the species belonging to the genus *Mesocricetus*, ranging from Anatolia, Transcaucasia (Armenia, Georgia and Azerbaijan) and southeast Dagestan to 449 450 northwest Iran (Qazvin in the east, Lorestan in the south; Lay, 1967). This species is found at altitudes from sea level up to 2,600 m. However, the primary 451 452 range is from 1,000-2,200 m. Mesocricetus brandti is found in arid and semi-453 arid steppe habitats in lowlands and in mountainous areas (Kryštufek et al., 2015; Kryštufek and Vohralík, 2009; Neumann et al., 2017). 454

455

456 457

458

459

460

#### <u>Family Muridae Illiger, 1811</u> <u>Subfamily Gerbillinae Gray, 1825</u> <u>Genus Meriones, Illiger 1811</u> <u>Meriones cf. persicus Blanford, 1875</u>

Material: 157 isolated teeth. Layer 1-3: three isolated teeth; two left upper
M1 and one right lower m1. Layer 4: 65 isolated teeth; 21 m2; four m3; 17 left
lower m1; eight right upper M1, five left upper M1; 10 right lower m1. Layer 5:
88 isolated teeth; eight m3; 21 m2; 18 left lower m1, 10 left upper M1, 16 right
upper M1 and 15 right lower m1.

Description and discussion: the genus *Meriones* is one of the most diverse
among the tribe Gerbillini in the Palaearctic region, particularly in arid regions of
Asia (Darvish et al., 2011; Denys, 2017). The *Meriones* species reported in Iran
are: *Meriones crassus*, *Meriones hurrianae*, *Meriones lybicus*, *Meriones meridianus*, *Meriones persicus*, *Meriones tristrami*, *Meriones vinogradovi* and *Meriones zarudnyi* (Darvish, 2011; Dianat et al., 2017; Kryštufek and Vohralík,
2009; Souttou and Denys, 2012).

473 The material from Kaldar Cave attributed to the genus Meriones displays the 474 typical morphology of this group, including semi-hypsodont molars with prismatic enamel triangles linked by a longitudinal crest and with no trace of 475 cusps. In our sample, we identify first upper molars (M1) with three roots, which 476 is characteristic of Meriones persicus and Meriones tristrami. Unfortunately, the 477 dental morphology of *Meriones persicus* and *Meriones tristrami* is very similar; 478 479 there are three roots in m1, the second molars have two transverse plates and 480 two roots, whereas the third molars are simple and rounded with a single root (Coşkun, 2016; Kryštufek and Vohralík, 2009). 481

Given its current distribution and the morphological traits observed in the reference collection from Iran, Azerbaijan and Pakistan housed in the Natural History Museum of London, we provisionally attribute our specimens to *Meriones* cf. *persicus*, especially in the light of the number of roots and the 486 morphology of M1 and m1, pending a revision of the Middle Eastern species of487 the genus.

Habitat and distribution: the genus *Meriones* is distributed across North 488 Africa, Central Asia, Transcaucasia, Turkey and Pakistan (Darvish et al., 2014; 489 Stoetzel et al., 2017). It lives mostly in dry steppes of short or tall grass, on 490 491 open hillside, among rocky outcrops in desolate steppes, or in open dry 492 meadows. The distribution of Meriones persicus ranges from the Caucasus (including the southeastern foothills of the Lesser Caucasus and the Talysh 493 494 Plateau in Azerbaijan) in the west, through northeastern Iraq and Iran to Turkmenistan, Afghanistan (Habibi, 2004) and Pakistan, where it is widely 495 496 distributed. It generally occurs in arid, rocky or mountainous regions (Kryštufek and Vohralík, 2009; Molur & Sozen, 2016). 497

498

499

500

501

502

<u>Genus Allactaga Cuvier, 1837</u> <u>Allactaga sp.</u>

Family Dipodidae Fischer, 1817

503 **Material:** two isolated teeth. **Layer 4:** one indet. **Layer 5:** one right lower m3. 504 **Description and discussion:** this rodent group is poorly known in the 505 Middle East (Shenbrot, 2009) and sometimes there is a size overlap between 506 the species. In our sample, we only found two items attributed to *Allactaga* sp. 507 on the basis of the complete lower m3, which presents a morphology with two 508 inner folds and one outer fold; the other tooth is broken, which prevents any 509 precise identification.

Habitat and distribution: in Iran four species are currently present: Allactaga elater, Allactaga euphratica, Allactaga firouzi and Allactaga hotsoni (Karami et al., 2008). The different species are morphologically very close and remain poorly studied. They are mainly found in steppe vegetation and semidesert areas (Shenbrot, 2009).

Family Gliridae Thomas, 1897

Genus Myomimus Ognev, 1924

Myomimus sp.

515

516

517

518 519

520 **Material:** five isolated teeth, **Layer 5:** five isolated teeth, one right lower m1, 521 three right lower m2 and one left lower m2.

**Description and discussion:** the genus *Myomimus* is present in Iran with 522 523 two species, Myomimus personatus and Myomimus setzeri (Firouz, 2005; Gerrie & Kennerley, 2017; Karami et al., 2008; Kennerley & Kryštufek, 2019). 524 Regarding the remains found at Kaldar, the m1 is of a trapezium-like shape, 525 and its anterior part tends to be narrower than the posterior part. The m1 has 526 three roots, two in the anterior part and one in the posterior part. The m2 is 527 528 subrectangular, and its occlusal pattern is simpler than in the m1. There are 529 three roots in total, two in the anterior part and one in the posterior part (Kaya

12

and Kaymakçı, 2018; Kryštufek and Vohralík, 2009). The two species are
similar to one another and poorly studied. Consequently, we were unable to
identify the Kaldar material precisely. Moreover, the reference collection of the
Natural History Museum of London only houses *Myomimus personatus*specimens, which could therefore not be compared with *Myomimus setzeri*.

Habitat and distribution: these species are not well known as regards their
distribution. They are mainly found in desert areas (Gerrie & Kennerley, 2017;
Kennerley & Kryštufek, 2019).

## 538

539 540

541

542

#### Family Muridae Illiger , 1811 Genus Apodemus Kaup, 1829 <u>Apodemus sp.</u>

543 **Material:** 13 isolated teeth. **Layer 1-3:** two isolated teeth; one left lower m1 544 and one left lower m2; **Layer 4:** six isolated teeth; three left lower m1, two right 545 lower m1 and one left upper M1. **Layer 5**: five isolated teeth; two left lower m1, 546 one right lower m1, one right upper M1 and one m2.

Description and discussion: the first lower molar (m1) can be seen to 547 548 present a low occlusal surface with six main cusps. The anterolabial and 549 posterolabial cusps of m1 converge in an X-shape. The posterior cusp of m1 is low, rounded and well developed, with two or three secondary cusps in the 550 labial part, and a mesial tubercle. We attribute the remains found in Kaldar 551 Cave to Apodemus sp. because they present the traits characteristic of the 552 553 genus Apodemus (Amori et al., 2016; Bogicevic et al., 2011; Knitlová and Horáček, 2017; Kryštufek and Vohralík, 2009; López-García, 2008). 554

Habitat and distribution: five Apodemus species are currently recognized
in Iran: A. hyrcanicus, A. flavicollis, A. witherbyi, A. avicennicus and A. *uralensis*, all belonging to the Sylvaemus sub-genus (Jangjoo et al., 2011).
Apodemus has a large distribution range extending from Great Britain across
much of continental Europe to the Urals. It also extends east through Turkey to
western Armenia, the Zagros Mountains of Iran and south to Syria, Lebanon
and Israel. It inhabits a variety of woodland habitats (Amori et al., 2016).

#### 562

563

- 564
- 565

#### <u>Genus Mus Linnaeus, 1758</u> Mus cf. musculus Linnaeus, 1758

Material: three isolated teeth. Layer 4: one isolated tooth; one left upper M1.
Layer 5: two isolated teeth; one indeterminate and one left lower m1.

**Description and discussion**: as in other murines, the first upper molar (M1) has three rows of tubercles: the first (t1, t2, t3) and second (t4, t5 and t6) groups situated in the anterior part have three tubercles, and the third group (t7 and t9) has two tubercles (Darviche et al., 2006). In Iran (Kryštufek and Vohralík, 2009) both *Mus musculus domesticus* and *Mus macedonicus* may be found.

The specimens from Kaldar Cave fit well with *M. musculus* as regards the upper 573 M1 morphology: the lingual row of cusps (t1 and t4) is shifted posteriorly; cusp 574 t7 is reduced to an enamel ridge; distal cusps t8 and t9 leave no space for a 575 576 posterior cingulum or posterolabial cusp t12 (Siahsarvie and Darvish, 2008). The dental ends of the mesial and central cusps on m1 fuse early; the 577 578 mesiolabial cusp is small. Upper molars normally have three roots each, one 579 lingual and two labial, whereas lower molars are two-rooted (one anterior and one posterior) (Kryštufek and Vohralík, 2009). Despite these observations, 580 581 bearing in mind the high intra-specific morphological variability, the low quantity of material and the bad state of preservation of some teeth (Fig. 2.13), we only 582 583 attribute the Kaldar material to M. cf. musculus.

Habitat and distribution: Mus musculus is a commensal species, well 584 distributed throughout the world: it is present over all the continents except 585 Antarctica. It is found in a wide range of habitats but tends not to be found in 586 forest and deserts. Mus musculus is ecologically highly opportunistic but a weak 587 competitor; it can cope with aridity and can also expand into the desert 588 589 (Kryštufek and Vohralík, 2009). It is also found in arid habitats along the border with Syria and Iraq, in desert landscapes and near the Euphrates River 590 591 (Kryštufek & Vohralík, 2009; Denys, 2017).

- 592
- 593

#### 4.2. Past landscape and climate at Kaldar Cave

594

595 The small-mammal assemblages from Layers 4 and 5 of Kaldar Cave are 596 dominated by the genus *Microtus* (60 individuals in Layer 4 and 79 in Layer 5), 597 followed by Ellobius in Layer 4 (17 individuals) and Meriones cf. persicus (17 individuals in Layer 4 and 18 in Layer 5). These species indicate that the 598 environment in the area was mainly composed of open dry and steppe areas, 599 600 although we also found Apodemus sp. and Mus cf. musculus, which are related rather to a dense vegetation cover (including trees/bush). All the species 601 602 identified at Kaldar Cave still occur in the area today.

Layers 1-3 do not yield enough material to draw palaeoclimatic inferences 603 604 (MNI < 30). For Layers 4 and 5, the *bioclimatic model* shows similar results 605 (Table 6). It should be pointed out that some species are not included in the BM data matrix, such as Myomimus (present in Layer 4) and Microtus guentheri 606 607 (Layer 5), because their values did not appear in the BM and could not be used in the analysis. The BM based on the small mammals from Kaldar Cave 608 suggests lower temperatures and lower precipitation than at present in both 609 Layers 4 and 5 (Table 6). The MAT is around 6°C lower, and there is a higher 610 temperature amplitude between maximum and minimum mean temperatures. 611 This indicates that Layers 4 and 5 of Kaldar Cave were deposited during a 612 613 colder period, which is not in agreement with the preliminary interpretations of Bazgir et al. (2017) suggesting a temperate "interstadial". 614

The climate in Khorramabad nowadays is warm an temperate. The average mean annual temperature is 16.90°C, the maximum mean temperature occurs in July with 29.60°C and the minimum mean temperature is in January with 5°C
 (<u>https://en.climate-data.org/asia/iran/lorestan/khorramabad-764550/</u>). This
 contrast with the climactic conditions inferred form the *BM* on Kaldar Cave with

620 lower temperatures and drier conditions.

Applying the *habitat weighting method*, our data show a palaeoenvironment mainly composed of steppes in both levels. In Layer 4, grasslands are also well represented, notably through the presence of *Microtus guentheri*. In Layer 5, deserts are the second most represented habitat, thanks to the considerable presence of *Myomimus* sp. The proportion of rocky areas is also significant, indicated by the presence of *Chionomys nivalis* and *Meriones* cf. *persicus* (Figure 3).

- There thus seems to be no major palaeoenvironmental or palaeoclimatic change that can explain the cultural shift between Layer 5 (Middle Palaeolithic) and 4 (Upper Palaeolithic). However, Layer 5 appears slightly colder and drier than Layer 4, with a higher proportion of desert habitats.
- 632 We can compare these results with previous studies carried out in Kaldar Cave using other palaeoenvironmental proxies. Charcoal analyses (Allué et al., 633 2018; Bazgir et al., 2017) indicate that there were active water sources or flows, 634 635 with specific plant communities characteristic of open forest growing in cool and 636 dry conditions (Allué et al., 2018). The presence of forested areas is also supported by the large mammals, with the presence of Sus scrofa, Capreolus 637 sp. and Cervus elaphus (Bazgir et al., 2017). Several amphibian and squamate 638 species have also been found (Bazgir et al., 2017): a toad (Bufo sp.), an agamid 639 640 lizard (Agamidae indet.), a gecko (Gekkonidae indet.), a skink (Scincidae indet.), a lacertid (Lacertidae indet.), a glass lizard (Pseudopus sp.), a sand boa 641 (Eryx sp.), possibly six types of colubrine snakes (Colubrinae indet.), a cobra 642 (Elapidae indet.), and a viper (Viperidae indet.). Most of these taxa (Agamidae, 643 644 Eryx and Elapidae) live in savannahs, steppes and deserts, with a way of life always linked with warm arid areas in rocky or sandy environments. Pseudopus 645 lives in dry and bushy environments, sometimes in open woodlands, but avoids 646 dense forest areas (Bazgir et al., 2017). 647

648 Our results based on small mammals, combined with previous studies of other proxies, thus suggest a mosaic landscape alternating between dry-steppe 649 areas and wooded patches. Nowadays, the vegetation in the Zagros is a 650 montane grassland-woodland characterised by steppe of grassland and herbs 651 with occasional to fairly common trees. The principal local trees are deciduous 652 oaks and junipers, with maples, walnut, almond and ash at middle elevations, 653 and Pistacia and Olea in drier areas (Fiacconi and Hunt, 2015). Thus, the 654 landscape during the Middle and Upper Pleistocene at Kaldar Cave was similar 655 to that today, despite a colder climate. 656

657

#### 658 **4.3. Kaldar Cave in the Middle Eastern context**

659

Kaldar Cave is situated in the Khorramabad Valley, which played a 660 significant role in human adaptation and dispersal during the Quaternary. Some 661 other caves close to Kaldar, such as Gilvaran, Ghamari and Gar Arjene rock 662 663 shelter, have yielded Mousterian and Aurignacian occupations (Bazgir et al., 2014). Moreover, surveys in Kermanshah have also documented Middle and 664 665 Upper Palaeolithic sites (Heydari-Guran and Ghasidian, 2020). None of them 666 has given rise to small-mammal studies. However, Gilvaran Cave has been studied from a palaeoenvironmental point of view through charcoal analyses 667 (Allué et al., 2018), showing a similar pattern to Kaldar, where Prunus is also 668 present and indicates the presence of open forest areas. Similarly, other proxies 669 670 such as pollens and charcoals have been analysed at Shanidar (Campana and Crabtree, 2019; Fiacconi and Hunt, 2015) and Gilvaran (Allué et al., 2018), 671 indicating similar environmental conditions to Kaldar. 672

673 Other studies based on small mammals from the Middle East have been performed in Qesem Cave (Maul et al., 2015a; Smith et al., 2015), Hummal 674 (Maul et al., 2015b), Azokh Cave (Fernández-Jalvo, 2016), Aghitu-3 Cave 675 (Frahm, 2019; Kandel et al., 2017; Nishiaki and Akazawa, 2018), Nesher Ramla 676 (Weissbrod and Zaidner, 2014), Amud Cave (Belmaker and Hovers, 2011), 677 678 Dzudzuana Cave (Belmaker et al., 2016) and Karain Cave (Demirel et al., 679 2011). These studies did not use HW or the BM for their palaeoclimatic and palaeoenvironmental reconstructions, and the sites 680 are not alwavs 681 contemporary with Kaldar, except Amud Cave (Belmaker and Hovers, 2011), Aghitu-3 (Djamali et al., 2008; Kandel et al., 2017) and Dzudzuana Cave 682 683 (Belmaker et al., 2016) (Table 7).

Most of these studies highlight problems in identifying Middle Eastern rodent 684 species, as faced in the present work. There are some studies of present-day 685 taxonomy and systematics in Iran, Turkey, Israel, Jordan, Lebanon and Syria 686 687 (Abi-said et al., 2014; Darvish et al., 2000; Haddadian Shad et al., 2014; Kopij and Liven-schulman, 2013; Obuch and Khaleghizadeh, 2012; Shehab et al., 688 2013), but these are mostly based on skull morphology and thus not applicable 689 in archaeological or palaeontological contexts where the remains are broken. 690 691 Very few studies provide descriptions of molars with comparative elements, 692 allowing for the correct identification of fossil specimens. In this context, comparison with modern specimens from museum collections of genetically 693 typed specimens is crucial to establish a correct taxonomic reference. 694

Palaeoenvironmental inferences have been drawn on the basis of small-695 mammal studies, as well as palynological and anthracological analyses. During 696 697 the Upper Pleistocene, small-mammal compositions differ from one site to another (Belmaker et al., 2016; Kandel et al., 2017) (Table 8), probably because 698 699 the sites are not exactly contemporaneous and are likely to belong to different eco-regions. It is noteworthy that the genera Microtus, Chionomys, Cricetulus, 700 701 Mesocricetus, Ellobius and Allactaga are represented in all the sites, allowing 702 us to reconstruct the species communities of this region. Kaldar is the most diverse site in terms of small-mammal species. 703

We applied the *habitat weighting method* at other Middle Eastern sites where 704 small-mammal studies have been performed, namely Aghitu-3 and Dzudzuana 705 Cave, but these are located in other eco-regions (Figure 1a). Aghitu-3 is located 706 707 in Armenia, at an elevation of 1601 m a.s.l. and the current climate is continental with considerable seasonality in temperature. Dzudzuana Cave is 708 located in Georgia, at an elevation of 560 m a.s.l., and the current climate is 709 710 warm and temperate. The estimations obtained with the habitat weighting method (Figure 4) show similar environmental conditions between Kaldar Cave 711 and Aghitu-3, where the landscape is dominated by steppe (indicated by the 712 presence of Ellobius lutescens, Mesocricetus brandti, Cricetulus migratorius 713 714 and *Allactaga*), with a relatively high percentage of grassland and forests.

To ascertain the palaeoclimatic conditions, we calculated and directly 715 compared our results with Aghitu-3 and Dzudzuana Cave using the *bioclimatic* 716 method (Table 9). The results obtained for Aghitu-3 show higher temperatures 717 718 and precipitation than at present, whereas Dzudzuana Cave shows the same 719 pattern as Kaldar Cave, with drier conditions and lower temperatures than 720 today. For Amud Cave, it was not possible to calculate HW and BM, but data 721 from the literature (Belmaker and Hovers, 2011) indicate a grassland 722 environment similar to the one that we reconstructed at Dzudzuana Cave.

723

#### 724 **5. Conclusions**

725

726 This work represents the first study of a Late Pleistocene rodent assemblage 727 the Middle East with palaeoenvironmental and palaeoclimatic from 728 reconstructions, using and adapting the habitat weighting method and the bioclimatic model to this area. We identified 1112 rodent remains, 729 corresponding to a minimum number of 264 individuals. The rodent assemblage 730 731 is composed of 13 taxa: six arvicoline (Microtus socialis, Microtus irani, Microtus guentheri, Chionomys nivalis, Ellobius fuscocapillus and Ellobius lutescens), 732 two cricetine (Cricetulus migratorius and Mesocricetus brandti), one glirid 733 (Myomimus sp.), one gerbilline (Meriones cf. persicus), one dipodid (Allactaga 734 735 sp.) and two murine species (Apodemus sp. and Mus cf. musculus). Augmenting the preliminary analysis of the material (Bazgir et al., 2017), new 736 species were identified, such as Microtus socialis, Microtus irani, Microtus 737 guentheri, Ellobius fuscocapillus and Meriones cf. persicus. We also 738 reconsidered the identification of Ellobius talpinus, Calomyscus sp. and 739 Dryomys cf. nitedula, based on modern specimens from museum collections 740 741 and measurements; this is why they do not appear in the new faunal list of Kaldar Cave. 742

Given the scarcity of studies in this biogeographical region, we encountered some difficulties in the identification of species, which could slightly affect the palaeoenvironmental and palaeoclimatic interpretations. In order to counter these potential errors, further studies of small mammals in this region are necessary, as well as research on discriminant characters in molars, using
 reference collections including genetically typed specimens.

The palaeoecological analysis of the rodents from Kaldar Cave revealed 749 lower temperatures and lower precipitation than present-day conditions, and an 750 environment mainly composed of dry steppes with patches of forested areas. 751 752 The results obtained are supported by palynological, anthracological and large-753 mammal studies. At a broader geographical scale, colder conditions were also inferred at Dzudzuana Cave. The genera Microtus, Chionomys, Cricetulus, 754 Mesocricetus, Ellobius and Allactaga are present at all the Middle East sites 755 (Aghitu-3, Dzudzuana and Amud) during the Upper Pleistocene, that have 756 757 yielded small mammals, allowing us to reconstruct the rodent communities of 758 the area.

Considering all the results from Kaldar Cave and other contemporaneous sites, we can conclude that both Neanderthals and AMH lived in dry steppes with patches of forested areas, without major environmental changes occurring between the Middle and the Late Palaeolithic. Climatic shifts during the MIS 4-3 transition were of a magnitude that did not have a major impact on small mammals in the region, suggesting that climate change may not have had the hypothesized effect on the Neanderthal extinction in the Levant.

766

767 Acknowledgments: I. Rey-Rodriguez is the beneficiary of a PhD scholarship funded by the Erasmus Mundus Program (IDQP). J.M. López-García was 768 supported by a Ramón y Cajal contract (RYC-2016-19386) with financial 769 770 sponsorship from the Spanish Ministry of Science, Innovation and Universities. M. Fernández-García is the beneficiary of a PEJ grant (PEJ2018-005222-P) 771 funded by the Spanish National Youth Guarantee System and the European 772 773 Social Fund. This work was developed within the framework of the projects 774 2017SGR859, 2017SGR840 and 2017SGR1040 (AGAUR, Generalitat de 775 Catalunya), and 2018PFRURVB291 (Univ. Rovira i Virigli). We thank the head 776 of the Research Institute of Cultural Heritage and Tourism (RICHT) (Dr. B. Omrani) and the head of the Iranian Center for Archaeological Research (ICAR) 777 778 (Dr. R. Shirazi) for providing us with the necessary support and permissions in 779 studying the materials. We thank the head of the Lorestan Cultural Heritage. Handicraft and Tourism Organization (Mr. A. Ghasemi) for all his support. We 780 781 also thank the head of International Collaboration and Ties of the RICHT (Mrs. M. Kholghi) for all her cooperation and help. B. Bazgir received his PhD 782 scholarship from the Fundación Atapuerca, for which he is grateful. Laxmi 783 Tumung received her PhD scholarship from the Erasmus Mundus Programme 784 (IDQP). We would like to thank Roberto Portela Miguez, Senior Curator in 785 Charge of Mammals, for his help with the reference collection in the Natural 786 787 History Museum of London. We would like to thank Rupert Glasgow for reviewing the English language of the manuscript. We also want to thank to the 788 789 Editor Prof. José S. Carrion and the two anonymous reviewers for their

comments and suggestions that strongly improve the final version of themanuscript.

792

#### 793 References

794

Abi-Said, M.R., Shehab, A.H., Amr, Z.S., 2014. Diet of the Barn Owl. Jordan
 Journal of Biological Sciences, Vol. 7, Nº 2, 109–112.

- Allué, E., Expósito, I., Tumung, L., Ollé, A., Bazgir, B., 2018. Early evidence of *Prunus* and *Prunus* cf. *amygdalus* from Palaeolithic sites in the Khorramabad
  Valley, western Iran. Comptes Rendus Palevol 17, 335–345.
  https://doi.org/10.1016/j.crpv.2018.01.001
- Amori, G. 1999. *Chionomys nivalis*. In: A. J. Mitchell-Jones, G. Amori, W.
  Bogdanowicz, B. Kryštufek, P. J. H. Reijnders, F. Spitzenberger, M. Stubbe, J.
  B. M. Thissen, V. Vohralík and J. Zima (eds), *The Atlas of European Mammals*,

Academi Press, London, UK

- 805Amori, G., Hutterer, R., Kryštufek, B., Yigit, N., Mitsain, G. & Palomo, L.J., 2016.806Apodemus flavicollis. The IUCN Red List of Threatened Species 2016:807e.T1892A115058023.8235.
- 808 https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 809 3.RLTS.T1892A22423256.en
- Amr, A., 2015. *Microtus guentheri*, *The IUCN Red List of Threatened Species* 2016: e.T13463A22349143.

Andrews, P., 1990. Owls, Caves and Fossils, Owls, Caves and Fossils. https://doi.org/10.2307/3889096

Andrews, P., 2006. Taphonomic effects of faunal impoverishment and faunal
mixing. Palaeogeography, Palaeoclimatology, Palaeoecology 241, 572–589.
https://doi.org/10.1016/j.palaeo.2006.04.012

- Aşan Baydemir, N., Duman, L., 2009. Molar patterns in *Microtus guentheri*(Danford and Alston, 1880) (Mammalia: Rodentia) from Kirikkale province.
  Journal of Applied Biological Sciences 3, 47–54.
- Bazgir, B., Ollé, A., Tumung, L., Becerra-Valdivia, L., Douka, K., Higham, T., Van 820 821 Der Made, J., Picin, A., Saladié, P., López-Garciá, J.M., Blain, H.-A., Allué, E., 822 Fernández-Garciá, M., Rey-Rodríguez, I., Arceredillo, D., Bahrololoumi, F., Azimi, M., Otte, M., Carbonell, E., 2017. Understanding the emergence of 823 824 modern humans and the disappearance of Neanderthals: Insights from Kaldar 825 Cave (Khorramabad Valley. Western Iran). Scientific Reports 7. https://doi.org/10.1038/srep43460 826
- Bazgir, B., Otte, M., Tumung, L., Ollé, A., Deo, S.G., Joglekar, P., López-García,
  J.M., Picin, A., Davoudi, D., van der Made, J., 2014. Test excavations and initial
  results at the middle and upper paleolithic sites of Gilvaran, Kaldar, Ghamari
  caves and Gar Arjene Rockshelter, Khorramabad Valley, western Iran. Comptes
- 831 Rendus Palevol 13, 511–525. https://doi.org/10.1016/j.crpv.2014.01.005
- Becerra-Valdivia, L., Douka, K., Comeskey, D., Bazgir, B., Conard, N.J., Marean, C.W., Ollé, A., Otte, M., Tumung, L., Zeidi, M., Higham, T.F.G., 2017.

- Chronometric investigations of the Middle to Upper Paleolithic transition in the
  Zagros Mountains using AMS radiocarbon dating and Bayesian age modelling.
  Journal of Human Evolution 109, 57–69.
  https://doi.org/10.1016/j.jhevol.2017.05.011
- Belmaker, M., Bar-Yosef, O., Belfer-Cohen, A., Meshveliani, T., Jakeli, N., 2016.
  The environment in the Caucasus in the Upper Paleolithic (Late Pleistocene):
  Evidence from the small mammals from Dzudzuana cave, Georgia. Quaternary
  International 425, 4–15. https://doi.org/10.1016/j.quaint.2016.06.022
- Belmaker, M., Hovers, E., 2011. Ecological change and the extinction of the
  Levantine Neanderthals: Implications from a diachronic study of micromammals
  from Amud Cave, Israel. Quaternary Science Reviews 30, 3196–3209.
  https://doi.org/10.1016/j.quascirev.2011.08.001
- Blain, H.-A., Bailon, S., Cuenca-Bescós, G., 2008. The Early–Middle Pleistocene
  palaeoenvironmental change based on the squamate reptile and amphibian
  proxies at the Gran Dolina site, Atapuerca, Spain. Palaeogeography,
  Palaeoclimatology, Palaeoecology 261, 177–192.
  https://doi.org/10.1016/j.palaeo.2008.01.015
- Bogicevic, K., Nenadic, D., Mihailovic, D., Lazarevic, Z., Milivojevic, J., 2011. Late
  Pleistocene rodents (Mammalia: Rodentia) from the Baranica cave near
  Knjaževac (eastern Serbia): Systematics and palaeoecology. Rivista Italiana di
  Paleontologia e Stratigrafia, Vol. 117, Nº2, 331-346.
- Campana, D. V., Crabtree, P., 2019. Evidence for skinning and craft activities
  from the Middle Paleolithic of Shanidar Cave, Iraq. Journal of Archaeological
  Science: Reports 25, 7–14. https://doi.org/10.1016/j.jasrep.2019.03.024
- Corbet, G.B., 1978. The Mammals of the Palearctic Region: A Taxonomic
  Review. London, Brit. Mus.(Nat. Hist.). Cornell Univ. Press. 117-118.
- Corbet, G.B. and Hill, J.E., 1991. A World List of Mammalian Species. 3<sup>rd</sup> ed.
   London, UK: British Museum (Natural History).
- Coşkun, Y., 2016. Review of unique odd chromosome-numbered underground
  rodent species of the Palearctic region: *Ellobius Lutescens* Thomas 1897
  (Rodentia: Cricetidae). Turkish Journal of Zoology 40, 831–841.
  https://doi.org/10.3906/zoo-1509-53
- Coşkun, Y., 1999. Morphological characteristics of *Meriones tristrami* Thomas,
  1892 (Rodentia: Gerbillinae) from Diyarbakır, Turkey. Turk. J. Zool 23, 345–
  355.
- Coşkun,Y., 1997. *Ellobius lutescens* Thomas, 1897 (Rodentia: Cricetidae) Turk.
  J. of Zoology. 21, 349-354.
- Darlington, P.J., 1957. Zoogeography: The Geographical Distribution of Animals.
  New York, NY, USA: John Wiley and Sons.
- Barviche, D., Orth, A., Michaux, J., 2006. *Mus spretus* et *M. musculus* (Rodentia,
  Mammalia) en zone méditerranéenne: Différenciation biométrique et
  morphologique: Application à des fossiles marocains pléistocènes. Mammalia
  70, 90–97. https://doi.org/10.1515/MAMM.2006.010
- Darvish, J., Mohammadi, Z., Mahmoudi, A., Siahsarvie, R., 2014. Faunistic and

- taxonomic study of Rodents from northwestern Iran. Iranian Journal of Animal
  Biosystematics (IJAB) 10, 119–136.
- Darvish, J., 2011. Morphological comparison of fourteen species of the genus *Meriones* Illiger, 1811 (Rodentia: Gerbillinae) from Asia and North Africa.
  Iranian Journal of Animal Biosystematics 7, 49–74.
- Darvish, J., Ghiyasi, R., Khosravi, M., 2000. Recognition of rodents of Robat
  Sharaf pellets owl by morphological and neontological studies. Journal of
  Sciences 12.
- Denys, C., 2017. Subfamily Deomyinae, Gerbillinae, Leimacomyinae,
  Lophiomyinae species accounts. Pp. 598-650. in : Wilson, D.E., Lacher, T.E.,
  Jr. & Mittermeier, R.A. eds. Handbook of the Mammals of the World. Vol.7.
  Rodents II. Lynx Edicions, Barcelona.
- Demirel, A., Andrews, P., Yalçinkaya, I., Ersoy, A., 2011. The taphonomy and
  palaeoenvironmental implications of the small mammals from Karain Cave,
  Turkey. Journal of Archaeological Science 38, 3048–3059.
  https://doi.org/10.1016/j.jas.2011.07.003
- Dianat, M., Darvish, J., Cornette, R., Aliabadian, M., Nicolas, V., 2017.
  Evolutionary history of the Persian Jird, *Meriones persicus*, based on genetics,
  species distribution modelling and morphometric data. Journal of Zoological
  Systematics and Evolutionary Research 55, 29–45.
  https://doi.org/10.1111/jzs.12145
- Djamali, M., de Beaulieu, J.L., Shah-hosseini, M., Andrieu-Ponel, V., Ponel, P.,
  Amini, A., Akhani, H., Leroy, S.A.G., Stevens, L., Lahijani, H., Brewer, S., 2008.
  A late Pleistocene long pollen record from Lake Urmia, NW Iran. Quaternary
  Research 69, 413–420. https://doi.org/10.1016/j.ygres.2008.03.004
- Ellerman, J.R, and Morrison-Scott, T.C.S., 1951. Checklist of Palearctic and Indian Mammals, 1780 to 1946. Brit. Mus. Nat. Hist. London, 1-810.
- Evans, E.M.N., Van Couvering, J.A.H., Andrews, P., 1981. Palaeoecology of
  Miocene sites in western Kenya. J. Hum. Evol. 10, 99-116.
- Fernández-Jalvo Y., King T., Yepiskoposyan L. & Andrews P. (eds.), 2016a.
  Azokh Cave and the Transcaucasian Corridor, Vertebrate Paleobiology and
  Paleoanthropology Series, Springer
- Fernández-Jalvo, Y., Andrews, P., Denys, C., Sesé, C., Stoetzel, E., MarinMonfort, D., Pesquero, D., 2016b. Taphonomy for taxonomists: Implications of
  predation in small mammal studies. Quaternary Science Reviews 139, 138–
  157. https://doi.org/10.1016/j.quascirev.2016.03.016
- Fiacconi, M., Hunt, C.O., 2015. Pollen taphonomy at Shanidar Cave (Kurdish
  Iraq): An initial evaluation. Review of Palaeobotany and Palynology 223, 87–93.
  https://doi.org/10.1016/j.revpalbo.2015.09.003
- 917 Firouz, E., 2005. The complete fauna of Iran. Ed. I.B. Tauris & Co Ltd
- Frahm, E., 2019. Upper Palaeolithic Settlement and Mobility in the Armenian
  Highlands: Agent-Based Modeling, Obsidian Sourcing, and Lithic Analysis at
  Aghitu-3 Cave.
- 921 Gerrie, R., Kennerley, R., 2017. Myomimus personatus. The IUCN Red List of

 922
 Threatened
 Species
 2017:
 e.T14088A22222124
 8235.

923 https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2017-

924 2.RLTS.T14088A22222124.en

Gharkheloo, M., Kivanç, E., 2003, A study on the morphology, karyology and
distribution of *Ellobius* Fischer, 1814 (Mammalia: Rodentia) in Iran. Turkish
Journal of Zoology 27(4):281-292

Golenishchev, F.,Malikov, V., Nazari, F., Vaziri, A.,Sablina, O., Polyakov, A.,
2002. New species of vole of *"guentheri "* group (Rodentia, Arvicolinae, *Microtus*) from Iran. Russia Journal of Theriology 1 (2), 117-123

- Golenishchev, F., Malikov, V., Petrova, T., Bodrov, S., Abramson, N., 2019.
  Toward assembling a taxonomic puzzle: Case study of Iranian gray voles of the
  subgenus *Microtus* (Rodentia, Cricetidae). Mammalian Biology 94, 98–105.
  https://doi.org/10.1016/i.mambio.2018.06.007
- Habibi, K. 2004. Mammals of Afghanistan. Zoo Outreach Organisation/USFWS,
  Coimbatore, India.
- Haddadian Shad, H., Darvish, J., Mohammadian, T., Mahmoudi, A., Alaie Kakhki,
  N., Ghanbarifardi, M., Molavi, F., Barani-Beiranvand, H., 2014. Preliminary
  study of rodents using pellets of predatory birds in Iran. Iranian Journal of
  Animal Biosystematics 10, 36–50.
- Hassinger, J.D., 1973. A survey of the mammals of Afghanistan: Resulting from
  the 1965 Street Expedition (Excluding Bats). Fieldiana Zoology, Vol. 60.
  Chicago, IL, USA: Field Museum of Natural History.
- Hernández Fernández, M., 2001. Bioclimatic discriminant capacity of terrestrial
  mammal faunas. Global Ecology and Biogeography 10, 189–204.
  https://doi.org/10.1046/j.1466-822x.2001.00218.x
- Hernandez Fernández, M., Peláez-Campomanes, P., 2003. The bioclimatic
  model: A method of palaeoclimatic qualitative inference based on mammal
  associations. Global Ecology and Biogeography 12, 507–517.
  https://doi.org/10.1046/j.1466-822X.2003.00057.x
- Hernández Fernández, M., Álvarez Sierra, M.Á., Peláez-Campomanes, P., 2007. 951 Bioclimatic analysis of rodent palaeofaunas reveals severe climatic changes in 952 953 Southwestern Europe during the Plio-Pleistocene. Palaeogeography, 954 Palaeoclimatology. Palaeoecology 500-526. 251. https://doi.org/10.1016/j.palaeo.2007.04.015 955
- Heydari-Guran, S., Ghasidian, E., 2020. Late Pleistocene hominin settlement patterns and population dynamics in the Zagros Mountains: Kermanshah region. Archaeological Research in Asia 21, 100161. https://doi.org/10.1016/j.ara.2019.100161
- Hinton, M.A.C., 1926. Monograph of the Voles and Lemmings (Microtinae) Livingand Extinct. London: British Museum (Natural History). Vol.1. 488 p.
- Jangjoo, M., J.Darvish & J. D. Vign 2011. Application of outline analysis on fossil
  and modern specimens of *Apodemus*. Iranian J. Anim. Biosystem., 7 (2): 143155.
- Kandel, A.W., Gasparyan, B., Allué, E., Bigga, G., Bruch, A.A., Cullen, V.L.,

- Frahm, E., Ghukasyan, R., Gruwier, B., Jabbour, F., Miller, C.E., Taller, A.,
  Vardazaryan, V., Vasilyan, D., Weissbrod, L., 2017. The earliest evidence for
  Upper Paleolithic occupation in the Armenian Highlands at Aghitu-3 Cave.
  Journal of Human Evolution 110, 37–68.
  https://doi.org/10.1016/j.jhevol.2017.05.010
- Karami, M., Hutterer, R., Benda, P., Siahsarvie, R., Kryštufek, B., 2008.
  Annotated check-list of the mammals of Iran. Lynx (Praha) 39, 63–102.
- Kaya, F., Kaymakçı, N., 2018. Systematics and dental microwear of the late
  Miocene Gliridae (Rodentia, Mammalia) from Hayranlı, Anatolia: Implications for
  paleoecology and paleobiodiversity. Palaeontologia Electronica 16.
  https://doi.org/10.26879/385
- Kennerley, R., Kryštufek, B., 2019. *Myomimus setzeri*, The IUCN Red List of
  Threatened Species 2019: e.T14089A22222049. 8235.
  https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2019-
- 980 1.RLTS.T14089A22222049.en
- Knitlová, M., Horáček, I., 2017. Late Pleistocene-Holocene paleobiogeography of
  the genus *Apodemus* in Central Europe. PLoS ONE 12, 1–23.
  https://doi.org/10.1371/journal.pone.0173668
- Kopij, G., Liven-Schulman, I., 2013. Zoology in the Middle East Diet of the Lesser
  Kestrel, *Falco naumanni*, in Israel 7140.
  https://doi.org/10.1080/09397140.2012.10648914
- Kretzoi, M., 1969. Skizze einer Arvicoliden Phylogenie -Stand 1969 // Vertebrata
  Hungarica. Vol.11. No.1–2. P.155–193.
- Kryštufek, B., Bukhnikashvili, A., Sozen, M., Isfendiyaroglu, S., 2017. *Cricetulus migratorius*. The IUCN Red List of Threatened Species 2016:
  e.T5528A115073390
  8235.
- 992 https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 993 3.RLTS.T5528A22391440.en
- Kryštufek, B., Yigit, N., Amori, G., 2015. *Mesocricetus brandti*, Brandt 's
  Hamster. The IUCN Red List of Threatened Species 2008: e.T13220A3421550.
  8235.
- https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2008.RLTS.T13220A3421550.
   en
- Kryštufek, B., Shenbrot, G., 2016. *Ellobius lutescens*, Transcaucasian Mole
  Vole. The IUCN Red List of Threatened Species 2016: e.T7655A22340006.
  8235. https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 1002 2.RLTS.T7655A22340006.en
- Krystufek, B., 2017. *Chionomys nivalis*, European Snow Vole. The IUCN Red
  List of Threatened Species 2016: e.T4659A115069366 8235.
  https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 1006 3.RLTS.T4659A22379147.en
- 1007 Kryštufek, B., Kefelioğlu, H., 2001. Redescription and species limits of *Microtus* 1008 *irani* Thomas, 1921, and description of a new social vole from Turkey 1009 (Mammalia: Arvicolinao).

- Kryštufek, B., Vohralík, V., 2009. Mammals of Turkey and Cyprus, Rodentia II:
  Cricetinae, Muridae, Spalacidae, Calomyscidae, Capromyidae, Hystricidae,
  Castoridae. https://doi.org/10.1644/10-MAMM-R-221.1
- Lay, D.M., 1967. A Study of the Mammals of Iran, Resulting from the Street Expedition of 1962-63. Fieldiana Zool., 54: 168-171.
- López-García, J.M., 2011. Los micromamíferos del Pleistoceno superior de la
   Península Ibérica: Evolución de la diversidad taxonómica y cambios
   paleoambientales y paleoclimáticos. Editorial Académica Española
- López-García, J.M., Blain, H.-A., Cuenca-Bescós, G., Alonso, C., Alonso, S.,
  Vaquero, M., 2011. Small vertebrates (Amphibia, Squamata, Mammalia) from
  the late Pleistocene-Holocene of the Valdavara-1 cave (Galicia, northwestern
  Spain). Geobios 44, 253–269. https://doi.org/10.1016/j.geobios.2010.10.001
- Maul, L.C., Bruch, A.A., Smith, K.T., Shenbrot, G., Barkai, R., Gopher, A., 2015a.
  Palaeoecological and biostratigraphical implications of the microvertebrates of
  Qesem Cave in Israel. Quaternary International.
  https://doi.org/10.1016/j.quaint.2015.04.032
- Maul, L.C., Smith, K.T., Shenbrot, G., Bruch, A.A., Wegmüller, F., Le Tensorer,
- 1027 J.M., 2015b. Microvertebrates from unit G/layer 17 of the archaeological site of
- Hummal (El Kowm, Central Syria): Preliminary results. Anthropologie (France)
- 1029 119, 676–686. https://doi.org/10.1016/j.anthro.2015.10.010
- 1030 Miller, G.S., 1896. Genera and subgenera of voles and lemmings// North 1031 American Fauna. Vol.12. P.1–85.
- Molur, S. & Sozen, M., 2016. *Meriones persicus*, Persian Jird. The IUCN Red List
  of Threatened Species 2016: e.T13166A22433231. 8235.
  https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 1035 2.RLTS.T13166A22433231.en
- Moradi Gharkheloo, M., 2003. A Study on the Morphology, Karyology and Distribution of *Ellobius* Fischer, 1814 (Mammalia: Rodentia) in Iran. Turkish Journal of Zoology 27, 281–292.
- Neumann, K., Yiğit, N., Fritzsche, P., Çolak, E., Feoktistova, N., Surov, A.,
  Michaux, J., 2017. Genetic structure of the Turkish hamster (*Mesocricetus brandti*). Mammalian Biology 86, 84–91.
  https://doi.org/10.1016/i.mambio.2017.06.004
- 1043 Nishiaki, Y., Akazawa, T., 2018. The Middle and Upper Paleolithic Archeology of 1044 the Levant and Beyond. Replacement of Neanderthals by Modern Humans
- 1045
   Series 35–47. https://doi.org/10.1007/978-981-10-6826-3\_3
- 1046 Nowak, R.M., 1999. Walker's Mammals of the World. 6th ed. Baltimore, MD,1047 USA: The Johns Hopkins University Press.
- Obuch, J., Khaleghizadeh, A., 2012. Spatial Variation in the Diet of the Barn OwlTyto alba in Iran 6, 103–116.
- Osborn, D.J., 1962. Rodents of subfamily Microtinae from Turkey. J. Mammalia,43: 515-529.
- Rey-Rodríguez, I., Stoetzel, E., López-García, J.M., Denys, C., 2019.
   Implications of modern Barn owls pellets analysis for archaeological studies in

- 1054the Middle East. Journal of Archaeological Science 111, 105029.1055https://doi.org/10.1016/J.JAS.2019.105029
- 1056 Roberts, T.J., 1977. The Mammals of Pakistan. London, UK: Ernst Benn Ltd.
- 1057Rusin, M., 2017. Ellobius talpinus , Northern Mole Vole. The IUCN Red List of1058ThreatenedSpecies2016:e.T7656A115085720.8235.1059https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 1060 3.RLTS.T7656A22339917.en
- Sándor, A.D., 2018. Rediscovered after half a century: A new record of the grey
   dwarf hamster, *Cricetulus migratorius* (Mammalia: Cricetidae), in Romania.
   Turkish Journal of Zoology 42, 495–498. https://doi.org/10.3906/zoo-1712-15
- 1064 Shehab, A., Daoud, A., Kock, D., Amr, Z., 2013. Zoology in the Middle East Small 1065 mammals recovered from owl pellets from 7140. 1066 https://doi.org/10.1080/09397140.2004.10638061
- Shenbrot, G., Kryštufek, B., Molur, S., 2016. Ellobius fuscocapillus, Southern 1067 1068 Mole Vole. The **IUCN** Red List of Threatened **Species** 2016: e.T7654A22339730. 8235. 1069
- 1070 https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 1071 2.RLTS.T7654A22339730.en
- 1072 Shenbrot, G., 2009. On the conspecifity of *Allactaga hotsoni* Thomas, 1920 and 1073 *Allactaga firouzi* Womochel, 1978 (Rodentia: Dipodoidea). Mammalia 73, 231– 1074 237. https://doi.org/10.1515/MAMM.2009.043
- Shenbrot, G.I. and Krasnov, B.R. 2005. An Atlas of the Geographic Distribution of
  the Arvicoline Rodents of the World (Rodentia, Muridae: Arvicolinae). Pensoft
  Publishers, Sofia.
- Siahsarvie, R., Darvish, J., 2008. Geometric morphometric analysis of Iranian
  wood mice of the genus *Apodemus* (Rodentia, Muridae). Mammalia 72, 109–
  115. https://doi.org/10.1515/MAMM.2008.020
- Smith, K.T., Christian, L., Flemming, F., Barkai, R., Gopher, A., 2016. The
   microvertebrates of Qesem Cave : A comparison of the two concentrations.
   Quaternary International 1–13. https://doi.org/10.1016/j.quaint.2015.04.047
- Souttou, K., Denys, C., 2012. Small Mammal Bone Modifications in Black Shouldered Kite *Elanus caeruleus* Pellets from Algeria: Implications for
   Archaeological Sites. Journal of Taphonomy 10, 1–19.
- Stoetzel, E., Cornette, R., Lalis, A., Nicolas, V., Cucchi, T., Denys, C., 2017.
  Systematics and evolution of the *Meriones shawii/grandis* complex (Rodentia,
  Gerbillinae) during the Late Quaternary in northwestern Africa: Exploring the
  role of environmental and anthropogenic changes. Quaternary Science Reviews
  164, 199–216. https://doi.org/10.1016/j.quascirev.2017.04.002
- Tesakov, A.S., 2016. Early Middle Pleistocene *Ellobius* (Rodentia, Cricetidae,
   Arvicolinae ) from Armenia. Russian J. Theriol. 15 (2): 151–158.
- Thomas, O., 1905. *Ellobius woosnami*. Abstr. Proc. Zool. Soc. 23; Proc. Zool.
  Soc. 526 London.
- Tsytsulina, K., Kryštufek, B., Yigit, N., Bukhnikashvili, A., Shenbrot, G., 2017.
   *Microtus socialis*, Social. The IUCN Red List of Threatened Species 2016:

- 1098 e.T13458A115114745.
- 1099 8235.https://doi.org/http://dx.doi.org/10.2305/IUCN.UK.2016-
- 1100 3.RLTS.T13458A22348936.en
- 1101 Walker, E.P., 1964. Mammals of the World. Vol. 2. Baltimore, MD,USA: The 1102 Johns Hopkins Press.
- 1103 Weissbrod, L., Zaidner, Y., 2014. Taphonomy and paleoecological implications of
- 1104fossorial microvertebrates at the Middle Paleolithic open-air site of Nesher1105Ramla, Israel.QuaternaryInternational331,115–127.
- 1106 https://doi.org/10.1016/j.quaint.2013.05.044
- Wilson, D.E. and Reeder, D.M., 2005. Mammal Species of the World.
  Washington, DC, USA: Smithsonian Institution Scholarly Press.
- 1109

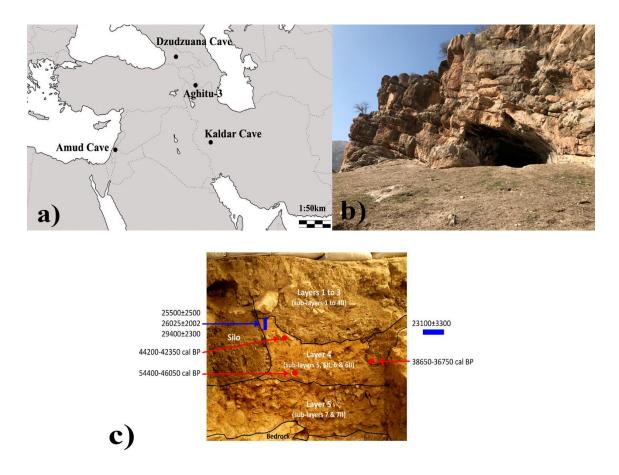


Figure 1: a) Locations of the Late Pleistocene sites with rodent studies in the Middle East, including Kaldar Cave. b) General view of the entrance to Kaldar Cave. c) Stratigraphy of the eastern section of Kaldar Cave, including the location and results of the dated samples, from A. Ollé and B. Bazgir (Bazgir et al., 2017).



Figure 2: Some rodent species identified at Kaldar Cave in comparison with equivalent (analogue) modern specimens from the NHM of London. 1) Microtus irani: 1.1-Kaldar Cave, 2014, Layer 5 (sub-layer 7II), E7, 150-156, right lower m1, number 447. 1.2-modern, NHM205207, Iran (Shiraz), right lower m1; 2) Microtus guentheri: 2.1-Kaldar Cave, 2014, Layers 1-3 (sub-layer 4II), F6, 97-107, left lower m1, number 17; 3) Microtus socialis: 3.1-Kaldar Cave, 2014, Layer 4 (sub-layer 5II), F7, 115-118, right lower m1, number 106. 3.2modern, NHM631799, Turkey (Amasya); 4) Chionomys nivalis: 4.1-Kaldar Cave, 2014, Layer 4 (sub-layer 5), E6, 94-104, left lower m1, number 542. 4.2-modern, NHM71820, Syria, left lower m1; 5) Ellobius fuscocapillus: 5.1-Kaldar Cave, 2014, Layer 4 (sub-layer 5II), E6, 125-130, right lower m1, number 157. 5.2-modern, NHM86101513, Afghanistan, right lower m1; 6) Ellobius lutescens: 6.1-Kaldar Cave, 2014, Layer 5 (sub-layer 7II), F6, 130-140, right lower m1, number 319. 6.2-NMH916416, Turkey, right lower m1; 7) Cricetulus migratorius: 7.1-Kaldar Cave, 2014, Layer 4 (sub-layer 5II), E6, 135-140, left lower m1, number 259. 7.2modern, NHM773029, Iran, right lower m1; 8) Mesocricetus brandti: 8.1-Kaldar Cave, 2014, Layer 5 (sub-layer 7II), F6, 135-145, left upper M1, number 549. 8.2- modern, NHM193461226, Caucasus, left upper M1; 9) Meriones cf. persicus: 9.1-Kaldar Cave, 2014, Layer 5 (sub-layer 7II), F6, 156-166, left lower m1, number 551. 9.2-modern, NHM510434, Iran, left lower m1; 10) Allactaga sp.: 10.1-Kaldar Cave, 2014, Layer 5 (sub-layer 7II), E7, 141-147, right lower m3, number 203; 11) Myomimus sp.: 11.1- Kaldar Cave, 2014, Layer 5 (sub-layer 7II), E7, 141-147, left lower m1 and m2. 11.2-modern, Myomimus personatus, NHM67623, Turkey, left lower mandible; 12) Apodemus sp.: 12.1-Kaldar Cave, 2014, Layer 4 (sub-layer 5), E6, 94-104, right m1 and m2, number 547; 13) Mus cf. musculus: 13.1-Kaldar Cave, 2014, Layer 5 (sub-layer 7II), 166-174, left lower m1, number 495. Scale 1 mm.

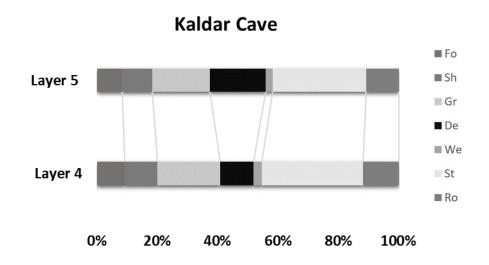


Figure 3: Results of the *habitat weighting method* for Kaldar Cave (Layer 4 and Layer 5). Forest (Fo), Shrubland (Sh), Grassland (Gr), Desert (De), Wetland (We), Steppe (St) and Rocky (Ro).

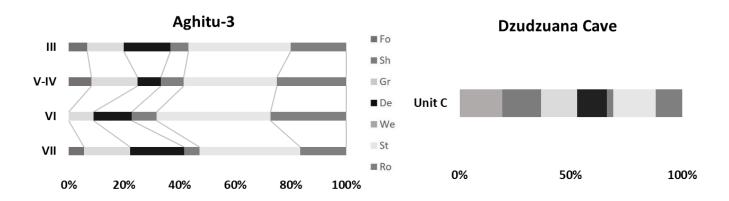


Figure 4: Results of the *habitat weighting method* for Aghitu-3 and Dzudzuana Cave. Forest (Fo), Shrubland (Sh), Grassland (Gr), Desert (De), Wetland (We), Steppe (St) and Rocky (Ro).

Species	Fo	Sh	Gr	De	We	St	Ro
Ellobius lutescens			0.33	0.33		0.33	
Ellobius fuscocapillus						1	
Microtus irani			1				
Microtus guentheri			1				
Microtus socialis		0.5				0.5	
Chionomys nivalis							1
Cricetulus migratorius			0.33	0.33		0.33	
Mesocricetus brandti						1	
Meriones persicus		0.33	0.33				0.33
Allactaga sp.				0.5		0.5	
Apodemus sp.	1						
Mus musculus		0.33	0.33		0.33		
<i>Myomimus</i> sp.				1			

Table 1: scores attributed to each rodent species found at Kaldar Cave according to its ecological requirements, used for the *habitat weighting method*: Forest (Fo), Shrubland (Sh), Grassland (Gr), Desert (De), Wetland (We), Steppe (St) and Rocky (Ro).

Species	I	II	11/111	Ш	IV	V	VI	VII	VIII	IX
Ellobius lutescens								1		
Ellobius fuscocapillus				0.33	0.33			0.33		
Microtus irani					1					
Microtus socialis					0.5			0.5		
Chionomys nivalis					0.25		0.25		0.25	0.25
Cricetulus migratorius					0.33		0.33	0.33		
Mesocricetus brandti					1					
Meriones persicus				0.5	0.5					
Allactaga sp.				0.5				0.5		
Apodemus sp.							1			
Mus musculus	0.11	0.11	0.11	0.11	0.11	0.11	0.11	0.11	0.11	

Table 2: scores attributed to each rodent species found at Kaldar Cave for the bioclimatic model (in accordance with Hernández-Fernández, 2001; Hernández-Fernández et al., 2007). See text for the significance of the Roman numerals corresponding to the climatic groups.

	Layer 5				Layer	4	Layers 1-3			
Taxon	NISP	MNI	%	NISP	MNI	%	NISP	MNI	%	
Indet.	3	-	-	11	-	-	-	-	-	
Microtus spp.	292	51	52.33	258	31	48.31	4	3	20	
Microtus irani	14	9	2.51	0	0	0.00	1	1	5	
Microtus guentheri	0	0	0.00	16	9	3.00	1	1	5	
Microtus socialis	35	20	6.27	36	20	6.74	3	3	15	
Chionomys nivalis	6	3	1.08	17	12	3.18	2	1	10	
Ellobius spp.	80	7	14.34	91	8	17.04	2	2	10	
Ellobius fuscocapillus	5	4	0.90	5	3	0.94	1	1	5	
Ellobius lutescens	10	6	1.79	10	7	1.87	-	-	-	
Cricetulus migratorius	5	3	0.90	14	3	2.62	1	1	5	
Mesocricetus brandti	7	4	1.25	3	1	0.56	0	0	0	
Meriones cf. persicus	88	18	15.77	65	17	12.17	3	2	15	
Allactaga sp.	1	1	0.18	1	1	0.19	0	0	0	
<i>Myomimus</i> sp.	5	3	0.90	0	0	0.00	0	0	0	
<i>Apodemus</i> sp.	5	2	0.90	6	3	1.12	2	1	10	
Mus cf. musculus	2	1	0.36	1	1	0.19	0	0	0	
Total	558	132	100.00	534	116	100.00	20	16	100.00	

Table 3: Representation of the Kaldar Cave rodent species in terms of number of identified specimens (NISP), minimum number of individuals (MNI) and percentage of the NISP (%).

Species NS		L (Length)		La (Width T4)		Li (W	'idth T5)	Total Width	
		Mean	Min-Max	Mean	Min-Max	Mean	Min-max	Mean	Max-Min
<b>Reference collection</b>									
Microtus irani	4	3.06	2.98-3.23	0.42	0.40-0.44	0.64	0.62-0.67	1.06	1.03-1.08
Microtus socialis	69	2.78	2.29-3.34	0.42	0.32-0.51	0.6	0.44-0.73	1.02	0.78-1.19
Microtus arvalis	25	2.72	2.45-2.96	0.39	0.32-0.46	0.6	0.50-0.70	1	0.83-1.13
Microtus guentheri	70	2.58	2.2-3						
Kaldar Cave									
Microtus guentheri	17	2.52	2.18-3.01	0.35	0.27-0.43	0.54	0.40-0.68	0.90	0.70-1.10
Microtus irani	15	2.89	2.52-3.39	0.41	0.37-0.50	0.62	0.54-0.70	1.05	0.92-1.18
Microtus socialis	74	2.54	2.06-3.01	0.37	0.28-0.48	0.55	0.40-0.70	0.92	0.70-1.16

Table 4: Measurements of *Microtus* specimens (in mm): NS, number of specimens; L, length; La, T4 width; Li, T5 width.

			Ellobius fu	scocapillus		Ellobius	lutescens	Ellobius talpinus		
		Layer 5 NISP=5	Layer 4 NISP=5	Layer 1-3 NISP=1	NHM NISP=12	Layer 5 NISP=9	Layer 4 NISP=10	NHM NISP=14	NHM NISP=14	
	Min-Max	3.29-3.84	3.73-3.74	-	3.06-4.05	3.08-3.32	3.05-3.4	3.07-3.3	2.86-3.35	
L	Mean	3.45	3.74	-	3.53	3.28	3.29	3.21	3.12	
W	Min-Max	1.32-1.51	1.40-1.54	-	1.3-1.59	1.20-1.45	1.13-1.48	1.11-1.52	1.09-1.72	
vv	Mean	1.45	1.48	1.5	1.41	1.34	1.32	1.34	1.24	

Table 5: Measurements of *Ellobius* specimens (in mm): L, total length; W, width. NHM: Natural History Museum of London. NISP: number of identified specimens.

	Layer 5	SD	Layer 4	SD	Current values
Mean annual temperature	10.90 °C	3.39	11.73ºC	3.39	16.90 °C
Maximum mean temperature	23.45 °C	4.77	23.10°C	4.77	29.60 °C
Minimum mean temperature	-1.69 °C	4.66	0.25°C	4.66	5°C
Mean annual precipitation	258.68mm	533.24mm	104.08mm	533.24mm	488mm

Table 6: Bioclimatic model estimates for Kaldar Cave. SD, standard deviation. Current values obtained from: <u>https://en.climate-data.org/asia/iran/lorestan/khorramabad-764550/</u>.

Site/Author	Layer	Human culture	Chronology	Environmental conditions	Rodent
Amud Cave	B4	Neanderthals	68.5 ± 3.4ka	Grassland vegetation	Microtus guentheri
(Belmaker and Hovers,					
2011)	B2-B1	Neanderthals	$56.5 \pm 3.5, 57.6 \pm 3.7,$ respectively	Woodland	Apodemus cf. mystacinus and Mus macedonicus
Dzudzuana Cave	Unit C	Modern	27-24ka cal BP	Mild and humid	Microtus, Ellobius, Arvicola,
(Belmaker et al., 2016)		human			Apodemus and Allactaga.
Aghitu-3 (Djamali et al., 2008; Kandel et al., 2017)	Level VII-III	Modern human	The oldest layer is VII with a chronology between 39- 36,000 cal BP and the most recent is layer III, between 29-24,000 cal BP	This sequence documents that the climate becomes colder	<i>Microtus</i> spp., <i>Chionomys</i> sp., <i>Arvicola amphibius</i> and <i>Ellobius lutescens</i>

Table 7: Comparison between Middle Eastern sites with small-mammal studies

	Ka		Aghitu-3	Dzudzuana				
Taxon	Layers 1-3	Layer 4	Layer 5	VII	VI	V-IV	III	Unit C
Microtus spp.	Х	Х	Х	Х	Х	Х	Х	
Microtus cf. arvalis								Х
Microtus irani	X		Х					
Microtus guentheri	X	Х						
Microtus socialis	X	Х	Х					
Chionomys nivalis	-	Х	Х	Х	Х	Х	Х	Х
Clethrionomys glareolus								Х
Ellobius spp.	X	Х	Х					
Ellobius fuscocapillus	X	Х	Х					
Ellobius lutescens	-	Х	Х	Х				Х
Cricetus cricetus								Х
Cricetulus migratorius	X	Х	Х	Х		Х	Х	Х
Mesocricetus brandti		Х	Х	Х	Х	Х	Х	Х
Meriones cf. persicus	X	Х	Х					
Allactaga sp.		Х	Х	Х	Х		Х	Х
Myomimus sp.			Х					
Apodemus sp.	X	Х	Х					Х
Mus musculus		Х	Х					Х
Arvicola amphibius				Х	Х	Х	Х	Х

Table 8: comparison of the small-mammal lists from several Upper Palaeolithic sites in the Middle East.

		Aghitu-3							Dzudzuana Cave		
	VII	VI	V-IV	III	SD	Current	Unit C	SD	Current		
						Values			Values		
Mean annual Temperature	15.46 °C	19.44 °C	19.21 ⁰C	17.87 ⁰C	3.386	8.5 °C	5.09 °C	3.39	9.7 °C		
Maximum mean temperature	23.59 °C	24.44 °C	24.09 °C	24 °C	4.772	20.2 °C	18.55 ⁰C	4.77	20.3 °C		
Minimum mean temperature	7.31 ⁰C	14.67 °C	14.72 °C	11.96 ⁰C	4.656	-3.8 °C	-8.26 ⁰C	4.66	-1.6 °C		
Mean annual precipitation	1169.04 mm	1729.48 mm	1825.99 mm	1484.60 mm	533.236	532 ⁰C	194.98 mm	533.24	868 mm		

Table 9: Estimates using the *bioclimatic method* for Aghitu-3 and Dzudzuana Cave. SD, standard deviation. Current values obtained from Kandel et al. (2017) for Aghitu-3 and <u>https://es.climate-data.org/asia/georgia/imereti/jria-414325/?amp=true</u> for Dzudzuana Cave.