New insights in Neanderthal palaeoecology using stable oxygen isotopes preserved in small mammals as palaeoclimatic tracers in Teixoneres Cave (Moia, northeastern Iberia)

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Abstract
The northeastern region of Iberia constitutes a natural pass-area for arriving populations into the peninsula and becomes a key area to understand Neanderthal resilience to changing environmental conditions experienced during Marine Isotope Stage 3 (MIS 3; 60–30 ka). Short-term but repeated occupations by Neanderthal groups occurred in Teixoneres Cave (Moia, Barcelona) in alternation with large and small carnivores during MIS3. Abundant small-mammal remains accumulated in units III and II of this fossiliferous deposit, providing local climatic and environmental information. This work focuses on the taphonomic history of small-mammal faunas, which a is clue to validate previous palaeoecological interpretations. As was observed with leporids and bird remains, raptors are considered the major source of small-mammal remains. The most likely accumulator is an opportunistic predator, the eagle owl, with very rare inputs by mammalian carnivores. In parallel, high-resolution palaeoclimatic data are provided through oxygen isotope analyses (δ18O) of rodent teeth from four subunits (IIIb to IIa), which are compared with independent methods of palaeotemperature estimations. According to air temperatures estimated from δ18O rodent teeth, cooler conditions than present day (−1.6/−0.5 °C) are recorded along the sequence, but homogenous (<1 °C). Complementary methods also explain higher rainfall than present day (+44/+682 mm). Only slight changes between units III and II show climatic instability, which could be related to palimpsests of stadial-interstadial events. Climatic stable conditions are reported from coeval isotopic and palaeodiet analyses from northeastern Iberia in agreement with the palynological records that underline how the Mediterranean area could have sustained rich ecosystems that assured the Neanderthal subsistence during the abrupt climatic pulsations of the Late Glacial.

Keywords Middle Palaeolithic · Southwestern Europe · Palaeoenvironment · Geochemistry · Taphonomy

Introduction
Southern European peninsulas are key areas to understand Neanderthal resilience to changing environmental conditions experienced during Marine Isotopic Stage 3 (MIS 3; ca. 60–30 ka). This period is characterized by intense climatic fluctuations, reflected in global ice and marine proxies (Fletcher and Sánchez-Goñi 2008; Fletcher et al. 2010a; Wolff et al. 2010; Moreno et al. 2014; Rasmussen et al. 2014), and coincides with the most extensive Neanderthal occupations reflected by the archaeological record in Iberia, along with their substitution by anatomical modern humans at the end of this period (Maroto et al. 2012; Higham et al. 2014; Wood et al. 2014; Marín-Arroyo et al. 2018). Climate, even if not the unique cause, should have an influence on the exploitation and occupation of their territory, conditioning hunting areas and the availability of vegetable resources (Zilhão 2000, 2006; D’Errico and Sánchez Goñi 2003; Finlayson et al. 2006; Finlayson and Carrión 2007). The Iberian Peninsula is a particularly interesting region considering its buffered climatic conditions compared to northern latitudes, but at the same time sheltering highly diverse micro-environments derived from their complex orography (Hewitt 2000; Sommer and Nadachowski 2006; Stewart et al. 2010; Jones 2021). Due to its geographic position, the northeastern
region of the Iberian Peninsula became a natural pass-area for arriving populations and the primary connection with the rest of Eurasia through the Mediterranean corridor. Evaluating the degree of adaptation of the Neanderthal groups in this region is essential to understand how they could face future human population arrivals. To provide reliable local-to-regional palaeoclimatic reconstructions of Middle Palaeolithic sites, especially in quantifiable terms that allow regional-to-global comparisons, it is essential to reach a deep knowledge in the ecological conditions faced by the Neanderthals during the millennia before the arrival of anatomically modern humans.

Teixoneres Cave preserves a well-studied fossiliferous deposit that can help to approach specific climatic conditions when Neanderthal populations were still spreading across Iberia. Neanderthal groups recurrently occupied Teixoneres Cave (Moia, Barcelona) during MIS 3, as it is attested by the Mousterian lithics recovered in the site. Carnivores such as hyenas, bears, and other small mammals, and birds of prey both nocturnal and diurnal, also frequented the cave and played an active role in the development of this sequence. These occupational dynamics were interrupted by short and sporadic Neanderthal settlements (Rosell et al. 2008, 2010a, b, 2017; Rufà et al. 2014, 2016; Sánchez-Hernández et al. 2014, 2016). Along this fossiliferous sequence, abundant small-mammal remains were accumulated, providing local climatic and environmental information. Owls, which probably roosted inside the cave over decades and centuries, are preliminarily assumed to be the main agent involved in a large number of small mammals (insectivores, bats, and rodents) in 2 sediments. However, until now, no complete taphonomic analysis was performed on this group and only some preliminary remarks were made (López-García et al. 2012a, 2014).

Accurate ecological inferences based on small mammals require complete knowledge of the agents involved in their accumulation (e.g. Andrews 1990), especially for deposits from which carnivores have frequently contributed, as is the case of Teixoneres Cave (see synthesis in Rosell et al. 2017). In this work, taphonomic analyses of small-mammal samples recovered from four subunits (IIIb, IIIa, IIb, and Ia) are presented. Apart from the identification of the main taphonomic agent involved in their accumulation, post-depositional alterations have also affected these small-mammal remains until they were recovered. A multidisciplinary team working at the site has recently provided extensive data concerning issues that can affect small-mammal assemblages, such as the origin of small prey (leporids and birds), the seasonality of the production of these remains, and the evaluation of time periods of human occupations, that should be addressed (Rufà et al. 2014, 2016; Sánchez-Hernández et al. 2014, 2016, 2020a, b; Rosell et al. 2017; Picin et al. 2020; Zilio et al. 2021).

Recent excavations in unit III have also provided interesting findings with ecological implications, such as the occasional presence of the woolly mammoth and the woolly rhinoceros, which are two cold-adapted fauna infrequent in Iberia (Álvarez-Lao et al. 2017). Moreover, previous absolute dates (Tissoux et al. 2006) and relative chronology inferred from small-vertebrate faunas (López-García et al. 2012a) have been adjusted through new radiocarbon dates (Talamo et al. 2016). Also, with special relevance, a new and complete palynological study has been just published (Ochando et al. 2020). These new findings, in addition to expanding archaeo-palaeontological material during the ongoing excavations, allowed us to review the initial palaeoenvironmental reconstruction based on small vertebrates. This work complements previous reconstructions performed by using 40 oxygen isotope analysis ($\delta^{18}$O) of rodent incisors from subunits IIIb to Ia. The relationships between $\delta^{18}$O preserved in mammal body tissues with $\delta^{18}$O of meteoric waters and mean annual temperatures allowed us to quantify the magnitude of the climatic changes through the sequence (Royer et al. 2013a, 2014; Fernández-García et al. 2019, 2020). Thus, considering all the recent research on the site, this work aims to reevaluate the ecological conditions during this sequence formation, unravelling the origins of the small-mammal remains and post-depositional processes. In addition, it presents an attempt to estimate air palaeotemperatures, based on the oxygen isotope analysis of rodent remains, and a comparison with independent methods for estimating climatic parameters. These environmental and climatic reconstructions are framed in this work within the regional-to-global ecological conditions experienced by Neanderthal populations during MIS 3 in Iberia.

**Teixoneres Cave**

**Archaeological general information**

Teixoneres Cave is located 5 km east of the village of Moià (Barcelona, Spain) (Fig. 1A) and belongs to the 2 km-long Toll Caves’ karstic system, formed by the drainage system of the Torrent del Mal, which modelled the Neogene limestone, called Collsuspina Formation. The karst coordinates are 2°09’02”E and 41°48’25”N at 760 m a.s.l. (Rosell et al. 2008). It is located in the highlands between the two main rivers that are the Llobregat and the Ter, connecting the central region of Catalonia with the Mediterranean coast (Rosell et al. 2017). The Toll karstic complex was discovered in the 1950s by a local speleological group. Several archaeo-palaeontological deposits from different chronologies are contained within this complex of caves, of which Teixoneres Cave and Toll Cave are those that contain Palaeolithic assemblages (Rosell et al. 2014). Several interventions were carried out by different research teams until the 1990s in the Toll karstic complex, bringing to light an important Holocene and Late
Fig. 1  A Location of the Teixoneres Cave in Iberia; B current monthly temperatures (continuous line; T; in °C) and precipitations (bars; P; in mm) (Climate-data.org) and present-day oxygen isotope composition of meteoric waters (dash black; $\delta^{18}O_{mw}$2017) of Moià; C synthetic stratigraphic sequence of chamber X in Teixoneres Cave, modified from Zilio et al. (2021); U-Th series (Tissoux et al. 2006) and radiocarbon dates (Talamo et al. 2016); D ground plan of unit III of Teixoneres Cave showing the 3D-position of the recovered items: lithics (blue), faunal remains (yellow), charcoals (black), and coprolites (brown). Squares sampled for taphonomic analyses of small mammals correspond to red (unit III) and green (unit II) stars. Drawing modified from Rosell et al. (2017)
Pleistocene palaeontological record. Since 2003, systematic excavations are annually undergone under the leadership of a multidisciplinary team of the Institut Català de Paleoecologia Humana i Evolució Social (IPHES) excavating in extension the whole surface of the Teixoneres Cave, ~250 m² (Rosell et al. 2008, 2010a, b, 2014).

Teixoneres Cave is composed of three main chambers (X, Y, and Z) that together form a 30 m length U-shape, with two access points from chambers X and Z (Fig. 1D). Chamber X is the biggest gallery, which is 30-m-long and filled with a 6-m-thick sediment package containing eight archaeopalaeontological units, including two speleothems (unit I and IV) (Rosell et al. 2008, 2010a; Zilio et al. 2021). The dates of unit I (ca. 14–16 ka) and unit IV (ca. 100.3 ± 6.1 ka) have been obtained through U-series dating of stalagmite layers (Fig. 1C) (Tissoux et al. 2006). More recently, radiocarbon dates adjusted the chronology of unit III, from >51,000 14C BP to 44,210 cal BP, and unit II, which extends from 44,210 to 33,060 cal BP (Talamo et al. 2016). This work focuses on these two units (II and III) of Chamber X, which have been the focus of recent excavations (Rosell et al. 2008, 2010b; Rufà et al. 2016). Both are units comprising sandy lutites and limestone blocks. A distinction between two archaeostratigraphic subunits (IIa and IIb) has been drawn due to the presence of large limestone blocks at the base of IIa. In turn, at least two different subunits (IIIA and IIIb) can be appreciated in unit III, distinguished by an increased presence of reddish clays in IIIb, limestone blocks at the base of IIIa, and the increased abundance of archaeological material in IIIb.

The sediment comes from allochthonous colluvial clays and silts, which entered the cave through the main entrance and a chimney located on the northeast side of the cave. Those sediments imbricate at the center of the main gallery. An autochthonous component is formed by limestone blocks falling from the walls and the roof of the cave.

Most previous studies concluded that there were short-term human occupations of the cave, in alternance with use by carnivorous predators, ruling out temporal contact between them (Rosell et al. 2010a, b; Rufà et al. 2014, 2016; Sánchez-Hernández et al. 2014, 2016; Bustos-Pérez et al. 2017; Picin et al. 2020; Zilio et al. 2021). Moreover, a bimodal spatial distribution is observed, of which human groups tend to concentrate their activities at the entrance of the cave, whereas carnivores and other predators preferentially use the inner areas (Fig. 1D). Among large mammals, a wide diversity of species was recorded in the cave, where both ungulates (Equus ferus, Equus hydruntinus, Cervus elaphus, Capreolus capreolus, Bos primigenius, Sus scrofa, Capra pyrenaica, and Rupicapra pyrenaica) and carnivores (Ursus spelaeus, Crocuta crocuta, Canis lupus, Vulpes vulpes, Lynx spelaea, Meles meles) are frequent. Medium-sized vertebrates were also identified as tortoise (Testudo hermanni), leporids (Oryctolagus cuniculus and Lepus sp.), and birds (Corvidae, Phasianidae, Strigidae and Columbidae) (Rosell et al. 2008, 2010a, b; Rufà et al. 2014, 2016). The presence of two cold-adapted species has also been recorded, Mammutthus primigenius and Coelodonta antiquitatis (Álvarez-Lao et al. 2017). Both hominin and carnivore activities were documented among larger fauna. Human butchering activities, mainly concerning ungulates, were confirmed from the detection of cut marks, fresh fractures, and burning (Rosell et al. 2008, 2010a, b). Analyzed leporid and bird remains are not primarily related to human activity, but with mammalian carnivores and raptors inputs (Rufà et al. 2014, 2016). However, sporadic consumption of leporids by humans was detected. The presence of Neanderthal groups is confirmed at the site by the discovery of Mousterian lithic tools. Raw materials have a local or semi-local origin (<15 km) and consist mainly of quartz, followed by chert, agates, and limestones. Reduction sequences are highly fragmented, consisting in flakes and final products, with some retouched tools (mainly scrapers) and in subunit IIIb also bone retouchers, linked to the short-term occupation patterns (Rosell et al. 2010a, b; Talamo et al. 2016; Bustos-Pérez et al. 2017; Mateo-Lomba et al. 2019). In unit III, Picin et al. (2020) proposed a differentiation between in situ knapping activities in local quartz and the mobile toolkit transported at the site. It was thus interpreted as a combination of hunting tools and cutting artefacts used during the primary butchery of the hunted preys in their punctuated visits to the cave. Some combustion structures characterized by sediment areas with rubefaction or hearths were mainly detected at the entrance in unit III (Rosell et al. 2010b, 2014). Moreover, human remains are represented by three deciduous teeth corresponding to at least two different individuals younger than 7 years old along with a molar of an adult.

Environmental context: past and present

The Moià region is nowadays included in the continental Mediterranean climate, at the boundary between the humid and sub-humid areas inside the Catalanian climatic subdivisions (Servei Meteorològic de Catalunya 2018), near the boundary of the Mediterranean and Eurosiberian regions. The climate is generally wet and temperate (mean annual temperatures around 12 °C), but with hot summers (18–21 °C) and cold winters (5–6 °C), along with a consequent thermic annual oscillation around 15 °C. Mean annual rainfall is 700–800 mm. Precipitation is abundant throughout the year, with autumn and spring being the wettest periods and winters being slightly drier (Climate-data.org 2018; Servei Meteorològic de Catalunya 2018). There is consistent correspondence between average monthly temperatures and monthly oxygen isotope compositions of meteoric water estimated for the area (OIPC data; Bowen 2017; Climate-Data.
org 2018) (Fig. 1B). The area is in the supramediterranean belt (Rivas-Martínez 1987; Rivas-Martínez et al. 2007), dominated by forests (mainly box, pine, and oak) adapted to mild summers and cold winters (Ochando et al. 2020).

Different approaches to reconstruct the palaeoecological conditions of Teixoneres Cave were performed. The specific attribution of the small-vertebrate remains (amphibians, squamates reptiles, and small mammals) was performed by López-García et al. (2012a, b, 2014). Small-vertebrate assemblage abundances underline a clear predominance of forest components (> 60%), especially in unit III, with some faunal component changes between Mediterranean and Mid-European taxa, from base to top. These data, in combination with pollen, charcoal, and large-mammal dental wear analyses, describe an open forest landscape, with temperate and humid conditions during the formation of unit III, and cooler and drier conditions during the formation of unit II (López-García et al. 2012a; Sánchez-Hernández et al. 2016, 2020a, b). Palynological data confirm a landscape dominated by woodlands composed of pines and oaks and some open areas (herbaceous taxa such as Poaceae, Chenopodiacea, and Astereaceae), always showing high proportions of arboreal pollen (> 50%). The most recent and complete palynological study (Ochando et al. 2020) described a mixed oak-pine forest with a high diversity of woody taxa that combines evergreen trees (Quercus), deciduous trees (Quercus suber; Juniperus), and the occurrence of Corylus, Castanea, Betula, Fraxinus, Buxus, Olea, Populus, Salix, Abies, Taxus, Cedrus, Acer, Alnus, Celtis, Juglans, Fagus, Ulmus, Calicotome, Ceratonia, Cistus, Ephedra fragilis, Myrtus, Pistacia, Phillyrea, Rhamnus, and Viburnum. Notably, high peaks of oak pollen are noticed in units III to I. Charcoal remains are scarce in the cave, but four taxa are recorded (Buxus sempervirens, Pinus pinea/pinaster type, Pinus sylvestris type, and Quercus sp.) (López-García et al. 2012a). The morphometric variation of the first lower molars of two vole species abundant at the site, Microtus agrestis and Microtus arvalis, indicates that the former decreased from subunit IIIa to IIb. In contrast, M. arvalis increased from IIIa to IIb (Luzi et al. 2017). The authors related these changes to better water management given the decrease in humidity from lower to upper units.

**Materials and methods**

**Taphonomic analysis**

The small-mammal remains included in this analysis consist mainly of isolated teeth and bones recovered from sediment samplings (organized in 5–10 cm splits) from the entire excavated surface of Chamber X (divided in 1 x 1 m), completed during excavation seasons from 2008 to 2016. The remains were collected by washing and sieving the sediment, with two superimposed meshes of 5 mm and 0.5 mm, and selected by subsequent sorting. Remains were observed under light microscopes (Exacta-Optech LFZ 10x-90x; Olympus SZ-PT 18x-50x). Due to the high amount of small-mammal material available at the site, specific squares were selected for the taphonomic analysis by considering the subunit divisions (IIb, IIIa, IIb, and IIa) to evaluate potential taphonomical difference between these subunits (Fig. 1D). For unit II, square J14 was selected, which is the site area with the most abundant small-mammal remains. Not enough material was available in unit III from this square, and a random selection including squares N8, O8, N10, O10, and M12 was chosen from the exterior part of the cave, where the human occupations were more intense.

The identified remains were counted (NISP) and grouped using the minimum number of individuals (MNI) method, determined by counting the most highly represented diagnostic element, considering laterality. Dental remains were considered independently whether they are in situ or isolated (e.g. each molar on a mandible is counted as a single remain). The MNI was adapted according to the taphonomic sampling performed by considering the minimum number of elements (MNE) of each subunit. The taphonomic study was based on the observation and description of the superficial modifications of skeletal elements (Andrews 1990; Fernández-Jalvo and Andrews 2016; Fernández-Jalvo et al. 2016) including the differentiation between predation and post-depositional damage. The taphonomic analysis included all rodent elements in anatomical representation (recount and indexes explanation detailed in Appendix 1) and incisors, molars, femora, and humeri of rodents were included in analyses of breakage, digestion, and post-depositional modifications. The analyses of soricids, talpids, and chiropters were also considered in the final interpretation.

**Oxygen isotope analysis**

Oxygen isotope compositions (δ18O) were measured on 40 rodent teeth from subunits IIIb, IIIa, IIb, and IIa. To avoid possible interspecific variability, Apodemus sylvaticus fossil lower incisors were preferentially selected, with thirty-four incisors corresponding to at least twenty-two individuals (Table 1). However, well-preserved incisors were not always available, and so, four teeth of the subfamily Arvicolineae in subunit IIb (n = 1) and IIa (n = 3) and two teeth of the species Eliomys quercinus in subunit IIIb and IIa were also included. In total, ten samples from each subunit were selected in order to gather a representative range of samples (Lindars et al. 2001; Gehler et al. 2012; Royer et al. 2013b; Peneycad et al. 2019). Oxygen isotope analysis performed on present-day material coming from the Moià area is also included. The remains came from one pellet recovered at the end of
August from a barn owl (Tyto alba) nest, located at 7.7 km from Teixoneres Cave at 1000 m a.s.l (Table 2; Fernández-García et al. 2019). These remains consist of three lower incisors from M. (T.) duodecimcostatus and M. musculus. One extra lower incisor of M. (T.) duodecimcostatus was also included from scattered pellets on the nest. The season in which this last element was produced is unknown.

The wet chemistry and the mass spectrometer measurements were developed in the Laboratoire de Géologie of Université Claude Bernard (Lyon, France). The rodent teeth were ultrasonicated and enamel fragments were selected by hand-picking and the basal part of the enamel was discarded. Enamel was homogenized by crushing in an agate mortar. The enamel tooth samples were treated following the wet
chemistry procedure described by Crowson et al. (1991) and Lécuyer et al. (1993), adapted for small-sample weights (Bernard et al. 2009). This protocol is based on the isolation of phosphate ions (PO₄³⁻) from apatite as silver phosphate (Ag₃PO₄) crystals using acid dissolution and anion-exchange resin. For each sample, around 1–3 mg of enamel powder from the resin with 6 ml 0.1 M 24 h, the solution was removed, and phosphate was eluted by centrifugation and the solution was neutralized with 1 ml 2 M KOH; then 1.5 ml of Amberlite™ anion-exchange resin was added to separate the PO₄³⁻ ions. After 24 h, the solution was removed, and phosphate was eluted from the resin with 6 ml 0.1 M NH₄NO₃. After 4 h, 0.1 ml NH₄OH and 3 ml of AgNO₃ were added to the solution and the samples were placed in a thermostatic bath at 70 °C for 6 h, enabling the precipitation of Ag₃PO₄ crystals. Four standard samples of natural phosphorite (NBS120c) were included during each batch of wet chemistry. Oxygen isotope compositions were measured using a varioPYROcube™ elemental analyser interfaced in continuous flow mode with an isotopic ratio mass spectrometer Isoprime™ (EA-Pyro-CF-IRMS technique performed at CNRS UMR 5276 LGL; Lécuyer et al. 2007; Fourel et al. 2011). Measurements were calibrated with the standard samples of NBS120c phosphorite (21.7‰; V-SMOW; Lécuyer et al. 1993), and NBS127 barium sulfate (9.3‰; V-SMOW; Hut 1987). Samples were measured usually in five replicates, but in some cases just three. Based on phosphate chemical yields measured during the wet chemistry procedure, clustered phosphorus pentoxide (P₂O₅) contents close to 35 wt% indicate that the original stoichiometry of the teeth was most likely preserved (Navarro et al. 2004; Hérant et al. 2010; Royer et al. 2013a). The average standard deviation of δ¹⁸O for the Teixoneres samples is 0.31 ± 0.03‰ (n = 40) and NBS120c quality control standards included in precipitation provide 21.5 ± 0.3‰ (n = 16), homogenous results close to their consensus value. 

### Palaeotemperature estimations

Usually, air temperatures are estimated from δ¹⁸Op by a two-step procedure involving two linear equations: (1) the measured δ¹⁸Op values of the fossil rodent teeth allow for the estimation of the δ¹⁸O values of local meteoric water (δ¹⁸Omw), and (2) the calculated δ¹⁸Omw values can be used to estimate past air temperatures. This method assumes that temperature is the dominant factor controlling the δ¹⁸Omw values recorded at mid- and high latitudes (Dansgaard 1964; Rozanski et al. 1993). However, secondary effects such as latitude, distance to sea, altitude, and distinct humid air mass trajectories in the past should also be considered. In parallel, oxygen isotope compositions of vertebrate biogenic apatite phosphate are linearly correlated to the oxygen isotope composition of their environmental waters (e.g., Longinelli 1984; Luz et al. 1984), with the physiological of each species being recorded in the oxygen isotope fractionation equation empirically established by the use of present-day individuals.

In this work, mean annual temperatures (MAT) were calculated from δ¹⁸O analysis of each subunit, following the strategy proposed for δ¹⁸Op from rodent teeth accumulated in the Iberian Peninsula (Fernández-García et al. 2019), that employed: (1) the empirically determined oxygen isotope fractionation equation established for extant rodent species for transforming δ¹⁸Op into δ¹⁸Omw [Royer et al. 2013a: δ¹⁸Op = 1.21 (±0.20) x δ¹⁸Omw + 24.76 (±2.70)]; and (2) a linear regression model relating δ¹⁸Omw and air temperatures based in IAEA Iberian stations climatic records [MAT (°C) = 2.388(±0.10) x δ¹⁸Omw + 28.19(±0.58)]. Indeed, this proposal takes into account the possible temperature seasonal bias derived from a preferent spring–summer rodent accumulation by predation [Fernández-García et al. 2019, 2020: δ¹⁸Omw (annual mean) = 1.04 (±0.05) x δ¹⁸Omw (March–June 2022)].

### Table 2  Oxygen isotope composition of tooth enamel phosphate (δ¹⁸Oₚ, ‰ V-SMOW) from rodent lower incisors from modern pellets of Moià. The table includes the provenance of remains, identified taxa, and the conversion to the oxygen isotope composition of meteoric waters (δ¹⁸Oₚ, ‰ V-SMOW) following the Royer et al. (2013a) oxygen isotope fractionation equation. SD standard deviation

| Origin          | MA-12          | MA-4          | MA-6          | MA-5          |
|-----------------|----------------|---------------|---------------|---------------|
| State           | Disaggregated  | Semi-dry      | Semi-dry      | Semi-dry      |
| Recovering      | Unknown        | End of August | End of August | End of August |
| Taxon           | M. (T.) duodecimcostatus | M. (T.) duodecimcostatus | M. musculus | M. musculus  |
| Laterality      | Right          | Left          | Right         | Left          |
| δ¹⁸Oₚ (‰ V-SMOW)| 16.1           | 18.6          | 19.2          | 19.5          |
| δ¹⁸Oₚ (‰ V-SMOW)| 0.4            | 0.4           | 0.3           | 0.3           |
| SD              | -7.2           | -5.1          | -4.6          | -4.4          |
mean)—0.86(±0.28) and also corrected the ice-sea level fluctuations that impact the δ¹⁸O of seawater (Schrag et al. 2002), which is the source of humid air masses (estimated in +0.6‰ for MIS 3 sea level descent).

In parallel, MAT and precipitations (mean annual precipitations; MAP) are compared with two qualitative and independent palaeoenvironmental methods: the mutual eco-geographic range (MER) and the bioclimatic model (BM) (detailed explanation in Appendices 3 and 4). The former was previously calculated for the Teixoneres site by López-García et al. (2012a, b, 2014), but this work applies the BM method for the first time (Hernández Fernández 2001; Hernández Fernández et al. 2007; Royer et al. 2020). The current temperature data recorded in the Teixoneres area (Moia) are taken from Climate-Data.org (https://es.climate-data.org/). Present-day δ¹⁸O_mw values for the Teixoneres area were obtained using the Online Isotopes in Precipitation Calculator (OIPC; Bowen 2017) based on datasets collected by the Global Network for Isotopes in Precipitation (IAEA/WMO, 2018).

Results and discussion

Origin of small mammals

The taphonomic analysis was completed on 2642 remains (Table 3; Figs. 2 and 3). The most frequent taxonomic group is the order Rodentia (NISP = 1536; 58.0%), followed by chiropters (NISP = 163; 6.0%), talpids (NISP = 11; 0.5%), and soricids (NISP = 37; 1.4%) in lower proportions (Appendix 1). Higher relative abundances of skeleton elements have been recorded in unit III (around 55%) than in unit II (IIb = 29%; IIa = 42%). However, all subunits shared a similar pattern regarding the skeletal part conservation. According to indexes comparing cranial and postcranial elements, these are almost in equilibrium, but slightly better preservation of dental elements (molars and, especially, incisors) is noticed, except for subunit IIa (Table 3). A preferential destruction of distal over proximal extremities is noticed. Breakage of mandibles and maxillae can be deduced, especially for lower subunits. Most recovered teeth in these assemblages are isolated (>90% of incisors and >75% of molars). Molar breakage is low, around 15–20% in subunits IIb and IIb, and 8–10% in subunits IIa and IIa. On the contrary, total breakage in incisors is around 65% for unit III and around 45% for unit II. All humeri and femora found in the sample were broken.

Digestion percentages ranged between 31% in subunit IIb and 43% in subunit IIa; subunits IIb and IIIa had an almost equal proportion of total digested elements (Table 3; Fig. 2). Different degrees of digestion are observed from light to heavy, except for subunit IIIb where extreme degrees are also detected. At least 35% of the elements in unit III exhibit digestion marks, which show a practically equal pattern in the digestion of incisors (50%) and molars (around 25%). Unit II is not far from these estimations, but there is a certain disturbance within their subunits, as both present an equivalent number of digested molars and incisors: around 30% in subunit IIb and around 40% in subunit IIa. Postcranial digestion accounts for around 40% of the elements, except for subunit IIb (<30%). Occasionally, intra-mandibular digestion of incisors was identified in all subunits. Digestion was occasionally detected in bats (subunit IIb and IIIb) and also in soricids (IIIb), for which extreme degrees of digestion are reached. Bat remains were preliminarily considered a result of both predation and in situ death, because the species has been recorded to frequently inhabit caves (Kowalski 1995; Palomo et al. 2007; López-García et al. 2012a).

More than 30% of elements in each subunit were affected by digestion, and skeleton parts spectra are fragmentary, confirming that predation was the main cause of small-mammal accumulation. Breakage and proportional representation indexes are not coincident with any predator pattern (Andrews 1990), even sometimes showing values included in Categories 4 and 5 of predators. It is common for indexes in archaeological contexts to differ from those of modern collections, because palimpsests and long-term post-depositional processes can occur. The predominance of remains without digestion marks, light alterations being the most frequent degree of digestion, low frequency of heavy digestions, and the absence of extreme digestion (except in subunit IIb) throughout the sequence point to nocturnal raptors as the main agent. They do not support the possibility of intervention by predators included in Categories 4–5 (Andrews 1990), such as diurnal raptors or small carnivores, as the observed fragmentation rates could initially suggest (Andrews 1990; Fernández-Jalvo and Andrews 1992; Terry 2007; Lloveras et al. 2008a, b, 2012; Comay and Dayan 2018). The occasional extreme digestions recorded in subunit IIb (2% of rodents, but also detected in some Soricidae remains) can be consistent either with unfrequented damage inflicted by owls related to different physiological or ethological factors (Andrews 1990; Royer et al. 2019) or most likely consequence of the punctual intervention of small carnivores in the cave.

Globally, the progressive proportion of degrees of digestion across each anatomical element and the correlation between proportion and degrees of digestion between the analyzed skeleton elements could indicate that a single type of predator is the most likely cause (Bennásar et al. 2016). The fraction of digestion remains, and the degree of digestion attained, besides medium–high relative abundance of elements and molar breakage recorded by small mammals analyzed at all subunits, fits with an intermediate modification predator (Andrews 1990), such
as *Bubo bubo* or *Strix aluco*. Distinguishing between these two owls is not simple, as both inflict similar damages on small-mammal remains. Subunits IIIa and IIIb, which provide similar digestion patterns in each skeletal element, present some characteristics more consistent with patterns expected from *B. bubo* than those from *S. aluco* (Andrews

| Taphonomic Variables Analyzed, Including Recounts (NR number of remains; NISP number of identified specimens; MNI minimum number of individuals of order Rodentia considered in skeletal representation), alterations caused by predation (skeletal representation, proportional representation indexes, breakage, and digestion), and post-depositional alterations | IIIb | IIIa | IIb | Ila |
|---|---|---|---|---|
| **Recount** | | | | |
| NR (n) | 782 | 988 | 388 | 484 |
| NISP (n) | 573 | 791 | 323 | 402 |
| Rodent MNI (n) | 22 | 31 | 23 | 22 |
| **Skeletal representation** | | | | |
| Relative Abundance Index (%) | 58.5 | 62.1 | 33.6 | 47 |
| SD of Relative Abundance Index | 42.2 | 42.2 | 14.9 | 47 |
| **Proportional representation indexes** | | | | |
| Postcranial/Cranial Index | 46.8 | 47.6 | 45.7 | 78.8 |
| Humerus + Femur/Maxilla + Mandible Index | 54.0 | 51.6 | 49.3 | 55.4 |
| Radius + Tibia/Humerus + Femur Index | 40.5 | 35.2 | 33.3 | 31.3 |
| Isolated teeth/Empty Alveolus Index | 67.2 | 74.8 | 60.6 | 54.8 |
| **Breakage** | | | | |
| Incisors fracture (%) | 68.1 | 68 | 45.8 | 42 |
| Molars fracture (%) | 11.7 | 8.3 | 17.1 | 6.9 |
| Postcranial fracture (%) | 100 | 100 | 100 | 100 |
| **Digestion** | | | | |
| Total digested elements | 93 | 35.6 | 134 | 36.3 | 48 | 31.4 | 78 | 43.3 |
| Light degree | 63 | 24.1 | 105 | 28.5 | 36 | 23.5 | 56 | 31.1 |
| Moderate degree | 17 | 7 | 22 | 6 | 10 | 7 | 15 | 8 |
| Heavy degree | 9 | 3.4 | 7 | 1.9 | 2 | 1.3 | 7 | 3.9 |
| Extreme degree | 4 | 2 | 0 | 0 | 0 | 0 | 0 | 0 |
| Digested incisors | 49 | 49.5 | 76 | 50 | 16 | 29.5 | 21 | 43.8 |
| Light degree | 32 | 32.3 | 61 | 40.1 | 11 | 21.6 | 14 | 29.2 |
| Moderate degree | 12 | 12.1 | 13 | 8.6 | 3 | 5.9 | 5 | 10.4 |
| Heavy degree | 3 | 3 | 2 | 1.3 | 2 | 3.9 | 2 | 4.2 |
| Extreme degree | 2 | 2 | 0 | 0 | 0 | 0 | 0 | 0 |
| Digested molars | 36 | 26.9 | 39 | 22.5 | 23 | 29.5 | 44 | 44.4 |
| Light degree | 25 | 18.7 | 29 | 16.8 | 16 | 20.5 | 29 | 29.3 |
| Moderate degree | 3 | 2.2 | 5 | 2.9 | 7 | 9 | 10 | 10.1 |
| Heavy degree | 6 | 4 | 5 | 3 | 0 | 0 | 5 | 5 |
| Extreme degree | 2 | 1 | 0 | 0 | 0 | 0 | 0 | 0 |
| Digested postcrania | 8 | 28.6 | 19 | 43.2 | 9 | 37.5 | 13 | 39.4 |
| Light degree | 6 | 21.4 | 15 | 34.1 | 9 | 37.5 | 13 | 39.4 |
| Moderate degree | 2 | 7.1 | 4 | 9.1 | 0 | 0 | 0 | 0 |
| Heavy degree | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Extreme degree | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Post-depositional agents | | | | |
| Stiations | 295 | 79.1 | 381 | 79 | 167 | 79.5 | 171 | 77.4 |
| Fissures | 104 | 27.9 | 141 | 29.3 | 34 | 16.2 | 46 | 20.8 |
| Cracks | 33 | 8.8 | 56 | 11.6 | 10 | 4.8 | 4 | 1.8 |
| Cementation | 20 | 5.4 | 57 | 11.8 | 2 | 1 | 3 | 1.4 |
| Manganese oxide pigmentation | 175 | 46.9 | 177 | 36.7 | 121 | 57.6 | 108 | 48.9 |
| Chemical corrosion—plant activity | 126 | 33.8 | 105 | 21.8 | 34 | 16.2 | 35 | 15.8 |
| Burning | 7 | 1.9 | 2 | 0.4 | 2 | 1 | 0 | 0 |

Bold is used to mark the main indexes and alterations considered in the analysis.
This owl tends to produce higher relative abundances of elements (around 60%), a low incidence of heavy digestion, while it introduces in its diet medium-sized preys such as leporids (some of which have been already related to this predator in unit III, Rufà et al. 2014). Digestion patterns moreover are highly in agreement with those reported for *B. bubo* by Lebreton et al. (2020) and those summarized by Fernández-Jalvo et al. (2016).

This pattern is clear for unit III, but less so for unit II, for which some clarifications should be made. Considering the scarcity of remains with moderate and heavy digestion and the absence of tooth marks, diurnal raptors and carnivores can be ruled out as the main cause (Andrews 1990). Thus, the main action of a predator of intermediate modification...
can be confirmed, but considering present-day reference collections (Andrews 1990; Fernández-Jalvo et al. 2016), none of the nocturnal birds of prey inflicted equal damage to molars and incisors. The presence of this equal damage in these subunits is most likely attributed to a combination of different groups of owls, which produce low to intermediate modifications. It is possible that *B. bubo* continued to contribute to unit II, but other predator inputs obscure the results. In addition, the low quantity of remains involved in the analysis of these subunits and the higher presence of the *Arvicolinae* subfamily should also be considered. The enlarged sample presented in this work provides enough remains to rule out the preliminary attribution to Category I predators (such as *Tyto alba*) as the main small-mammal accumulator (López-García et al. 2012a, 2014) in any of the subunits, at least in exclusivity.

Previous studies devoted to small-prey remains from the site also demonstrated a high degree of breakage of leporids in unit III (>80%) and birds for the full sequence (>70%). Anatomical profiles of leporids demonstrated proportions closer to accumulations produced by diurnal raptors or mammal carnivores (such as foxes, lynx, or wolves) (Rufà et al. 2014, 2016). However, as well as it is observed in small mammals, the percentages of digested remains (leporids: 9%; birds: 19–35%) and the predominance of light degrees of digestion and in the case of birds also anatomical representation are unlike patterns established for small carnivores and more related to nocturnal raptors. In both cases, a similarity with digestion patterns of *Bubo bubo* was suggested. Anecdotally, one skeletal remain of *Bubo bubo* was recovered in subunit IIa while Strigidae family remains were found in subunits IIIa, IIb, and IIa (Rufà et al. 2016). However, a low percentage of modifications made by carnivores in some avian remains was also observed throughout the sequence. Small carnivore alternations were also detected in leporid accumulations from unit III (16%), as low percentages of heavy and extreme digestion, which were mainly related to the action of foxes (*Vulpes vulpes*). In the case of small mammals analyzed here, the possible intervention of carnivores through digestion traces has only been detected in subunit IIIb. However, it cannot be rejected that their action incremented breakage rates and biased anatomical profiles in the rest of the sequence.

In summary, these results indicate that the small-mammal assemblages of the Teixoneres Cave were mainly produced by nocturnal raptors and, at least for unit III, most likely by the eagle owl, whose nest or roosting site might have been located near or even inside the cave. This species has been present in the Iberian Peninsula since the Lower Pleistocene and has been the sole species of the genus *Bubo* since the Middle Pleistocene. It is assumed that their past distribution was similar to the present but was probably more southern during stadial periods (Arribas 2004). The eagle owl is a sedentary species that usually nests in cave entrances, cliffs ledges, or rock crevices, using the same place for several years (Mikkola 1983; Andrews 1990; Manzanares 2012; IUCN 2018). It is an opportunistic hunter, including in its diet whatever is available in its hunting territory, which normally extends to around 10 km². It feeds mostly on mammals, but its prey assemblages are highly diverse and reflect the local ecological variability of its nesting area, and include reptiles, amphibians, fish and larger insects. This owl species is the largest in Iberia (2–4 kg) and, thus, tends to include larger preys in its diet, frequently consuming leporids and birds (such as herons and buzzards) (Olsson 1979; Andrews 1990; Manzanares 2012, IUCN 2018). It is a valuable predator for palaeoecological reconstructions, since it generates representative and coherent accumulations.

**Fig. 3** Taphonomic variables related to predation observed on rodents from subunits IIIb–IIa. A) Skeletal profile based on relative abundances of maxillae (MX), mandibles (MD), molars (MO), incisors (IN), humeri (HU), radii (RA), ulnae (UL), femora (FE), and tibia-fibula (TF); B) breakage rates; C) digestion percentages; D) total degree reached.
from the ecosystem it exploits (Andrews 1990). In addition, this bird may provide some environmental information due to its common preference by wooded habitats despite being observed in a wide range of environments (Mikkola 1983; IUCN 2018).

Post-depositional alterations

Post-depositional alterations occurred after small-mammal accumulation but could engender sizable modification in bone conservation and affect skeletal profile quantifications. A common pattern was detected in the post-depositional history of small-mammal remains along the sequence (Table 3; Fig. 2; Appendix 1). Most of the analyzed remains show striations (>75%), fissures (15–30%), and cracks (2–12%) most likely related to humidity changes. Between 37% (subunit IIIa) and 58% (subunit IIb) of the remains are affected by manganese oxide pigmentation. Another common alteration is chemical corrosion, mainly related to the presence of roots, which affected more of the lower subunits (from 34 to 22%) than the upper ones (around 16%). Occasionally, remains from subunits IIIb and IIIa were covered by isolated cementations (5% and 12%, respectively), being rare in unit II (<2%). Therefore, all features recorded indicated a wet fossiliferous environment where humidity levels underwent some fluctuation, in keeping with cave taphonomic patterns. Finally, few remains in subunits IIIb, IIIa, and IIb were burned and reached carbonization and calcination degrees. The presence of hearths was confirmed at the site, mainly at the entrance of the cave in unit III. In unit II, sediment alterations in form of rubefactions were related to thermal exposure (Rosell et al. 2010a, 2014). Small mammals trapped in the substrate would have been affected by unintentional thermal alterations as a consequence of the surrounding combustion activities (Fernández-Jalvo et al. 2018), suggesting also that a part of the material in direct contact with fire could have disappeared by burning.

The absence of rounding and polish produced by water abrasion supports the idea that small-mammal remains were not affected by water streams active in the surroundings of the site, supporting previous interpretations based on the absence of round-angle gravel accumulation (Rosell et al. 2010a). Considering the frequency of manganese oxide coating, however, punctual events of flooding cannot be rejected. The taphonomic analysis performed on birds revealed some cases of weathering and trampling (Rufà et al. 2016). In the last case, its apparent absence in small-mammal remains should be clarified in the future with the help of a scanning electron microscope study. The absence of weathering is common in cave deposits where the exposure to light is highly limited. This may confirm the in situ deposition of the small-mammal assemblages and the potentially faster burial of small-mammal remains than that was expected for larger remains considering the low sedimentation rates defined for this cavity (Rosell et al. 2010a, b; Rufà et al. 2016). As was previously observed, the anatomical profiles and breakage rates obtained in each subunit differ from those produced by owls. Post-depositional modifications should be considered as a possible cause of these fractures. Considering the absence of trampling and weathering among small mammals and a low incidence of this damage in medium-size vertebrates (Rufà et al. 2014, 2016), it is complex to consider these agents as the main cause of higher fragmentation rates. Sediment compactions could also play a role in this sense.

Oxygen isotope compositions of rodent remains

The oxygen isotope composition of the rodent incisor enamel from the Teixoneres sequence ranges from 15.5 to 21.3‰, representing a variation range of 5.8‰ (SD = 1.3) (Fig. 4; Table 1; Table 4). Both mean and median δ18O values show little variation among different subunits, ranging from 17.7‰ (subunit IIIa) to 18.5‰ (subunit IIb) and from 17.9‰ (subunit IIIa) to 18.5‰ (subunit IIa), respectively. Differences in mean and median δ18O values are around 0.1–0.3‰, showing a homogenous distribution of data inside each subunit. The oxygen isotope compositions of remains from the subfamilies Glirinae and Arvicolinae...
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included in subunits IIa and IIb are consistent with the general δ18Op range obtained from Murinae remains in the site and have no influence either on the mean or median values. However, in subunit IIa, two of the three values related to Arvicolinae correspond to extreme δ18Op values increasing the intra-level range. If these samples are not considered, no differences are noticed in average measurements, but the amplitude of this subunit is largely reduced (1.7; SD = 0.6). All sedimentary layers show a normal distribution of data (Shapiro–Wilk’s test, p-value > 0.01).

If the δ18Omw values were estimated from the δ18Op, the amplitudes recorded inside each subunit of Teixoneres Cave would be lower than the current seasonal amplitude of δ18Omw values recorded at Moià (8.3‰; OIPC data; Bowen 2017; Fig. 1C; Table 4) and the average seasonal amplitude of δ18Omw values for Iberia (8‰; OIPC data; Bowen 2017; Fernández-García et al. 2019). Tentatively, it can be suggested that analyzed rodent remains most likely accumulated during a preferential period of the year. Indeed, there is a higher probability of accumulating prey by owls during the warm season in Iberia (Chaline 1974; Salamolard et al. 2000; Norrdahl and Korpima 2002; Manzanares 2012; Lagos 2019), as it is indicated by the higher number of murid specimens and the observed oxygen isotope composition of contemporaneous Iberian rodent tooth samples (see discussion in Fernández-García et al. 2019; Royer et al. 2013a, b). Mean and median δ18Omw fossil values are notably higher (from −5.2 to −5.9‰) than the current mean annual δ18Omw values documented at Moià (−7.3‰; Bowen 2017), reinforcing this hypothesis. The minimum value from the sequence, in subunit IIIa, expressed in δ18Omw equivalent to −7.7‰ (Table 2), is not reaching the present-day winter δ18Omw (from −8.8 to −10.5‰; Bowen 2017; Fig. 1B). In contrast, the maximum δ18Op value record in the subunit IIIB (δ18Op = 21.3‰; δ18Omw = −2.9‰) corresponds to present-day summer oxygen compositions (from −2.6 to −3.8‰; Bowen 2017; Fig. 1B). δ18O values in all subunits do not indicate predation during the cold season but rather coincide with lower mean-medians and higher ranges-SDs in unit III. These results could be either a consequence of climatic instability (if rodent remains are produced in the same period), stadial-interstadial mixings of remains (if mixed faunal communities occurred), and/or changes in seasonality patterns or in the altitude of hunting of the predator. However, it should be considered that the comparison between estimated and modern δ18Omw values is highly speculative considering that several actualistic assumptions should be accepted, including similar humid air masses circulation and similar correspondence between oxygen isotopic values and seasonality patterns.

The eagle owl (B. bubo) has been identified in this work as the main cause of small-mammal accumulation in the Teixoneres sequence at least in unit III, in keeping with results obtained for birds and leporids (Rufà et al. 2014, 2016). Today, its breeding operates between January and March in the Iberian Peninsula, incubation lasts about 5 weeks, and the offspring can fly about 2 months after (Manzanares 2012). Consequently, a preferential accumulation is proceed during spring and beginning of summer, knowing that its foraging is usually increment coinciding with their breeding period (Salamolard et al. 2000; Norrdahl and Korpima 2002; Manzanares 2012; Lagos 2019). Taphonomic studies on avian remains, which probably share their origin with small mammals, could help to infer the

| Table 4 Minimum, maximum, mean, median, standard deviation, and range of oxygen isotope composition of incisor enamel phosphate (δ18Op, ‰ V- SMOW) from fossil rodents recovered from Teixoneres Cave samples; conversion to the oxygen isotope composition of meteoric waters (δ18Omw, ‰ V- SMOW), including minimum, maximum, mean, median, and range; mean annual temperature estimations (MAT; °C), including seasonality and sea level corrections of δ18Omw. Present-day samples from Moià are also incorporated to calculate their average temperature, without applying corrections
| \( \delta^{18}O_p \) | \( \delta^{18}O_{mw} \) | MAT \( °C \) |
|---|---|---|
| IIa | IIb | IIIa | IIIb | Pellet |
| n | 10 | 10 | 10 | 10 | 3 |
| Min | 16.6 | 16.8 | 15.5 | 16.0 | 19.0 |
| Max | 20.4 | 20.3 | 19.5 | 21.3 | 19.5 |
| Mean | 18.4 | 18.5 | 17.7 | 18.2 | 19.1 |
| Median | 18.5 | 18.2 | 17.9 | 18.0 | 19.2 |
| Range | 3.8 | 3.5 | 4.0 | 5.2 | 0.5 |
| SD | 1.0 | 1.0 | 1.3 | 1.7 | 0.3 |
| \( \delta^{18}O_{mw} \) (seasonality correction) | −6.7 | −6.6 | −7.7 | −7.2 | −4.8 |
| \( \delta^{18}O_{mw} \) (sea level correction) | −3.6 | −3.7 | −4.3 | −2.9 | −4.3 |
| MAT \( °C \) | 11.8 | 11.3 | 10.7 | 10.8 | 17.3 |
| SD | 1.3 | 1.3 | 1.3 | 1.3 | 2.3 |
| Error margin | 2.6 | 2.6 | 2.6 | 2.6 | 4.5 |
seasonality of human and carnivore/raptor occupations. Few medullary bones among avian remains (*Pica pica* in level IIa and *Pyrrhocorax* spp. in level IIIb) have been observed, tentatively suggesting that at least part of the bird assemblage was accumulated during spring (Rufà et al. 2016).

Moreover, zooarchaeological and taphonomic analyses reflect the presence of carnivores (mainly cave bear and hyena) alternated with Neanderthal occupations, and the evidence associated with humans and carnivores does not usually overlap temporally (Rosell et al. 2010a, b, 2017). Regarding carnivores, bears are known to use the cave mainly for hibernation during winter months, but patterns of hyenas and small carnivores are unknown (Rosell et al., 2017). From a combination of tooth microwear, cementum analysis, and dental eruption patterns in cervids and equids from the site, Sánchez-Hernández et al. (2014, 2016, 2020b) suggested that there were repeated short-term hominin occupations at the site within the same season of the year, probably in summer for subunit IIa; in autumn and early winter for subunit IIb; summer and winter, in subunit IIIb; and repeated and short occupations in all seasons for subunit IIIa. Equids, however, were mainly hunted during late spring and summer. These periods of frequentation of the cave may be related to an abundance of cervids and equids in the territory (Sánchez-Hernández et al. 2014, 2016; Rosell et al. 2017; Picin et al. 2020).

Based on these interpretations of short-term human occupations and the preferential use of the cave by owls in spring, in addition to noted carnivore dynamics, in general, no coexistence between humans and owls and/or carnivores can be supported. This reinforces the absence of temporal contact between them underlined by previous taphonomic and zooarchaeological studies, which demonstrate long periods without human occupation of the site (Rosell et al. 2010a, b, 2017; Rufà et al. 2014, 2016; Sánchez-Hernández et al. 2014).

Reconstruction of past temperatures

Mean annual temperature (MAT) was calculated based on the median δ18Op of each sedimentary subunit (Fig. 5). The estimated MATs vary between 10.7 ± 2.6 °C (subunit IIIa) and 11.8 ± 2.6 °C (subunit IIa). These temperatures are lower than the current MAT recorded at Moià (12.3 °C; Climate-Data.org; Fig. 1B), with differences ranging from −0.5 to −1.6 °C. Along the sequence, MAT differences are small (≤1.1 °C). Therefore, based on oxygen isotope compositions, a homogenous and milder climate without notable changes is indicated; however, a slight increase in temperatures from lower subunits (10.8–10.7 °C) to upper ones (11.3–11.8 °C) has been detected.

Additionally, oxygen isotope compositions of the remains recovered in the modern pellet from the Moià region provide homogenous values of 19.1 ± 0.3‰ (−4.6 ± 0.4‰ in δ18Omw), which is especially relevant considering those remains belong to two different species (Table 4). Considering δ18Omw at the place of sample recovery (at 1000 m a.s.l.; OIPC data; Bowen 2017), the pellet records could be coincident with May–June (−5.4/−4.2‰) or with September (−4.5‰). The pellet was recovered in August in a semi-dry state. Thus, the most plausible explanation is that this pellet recorded information from May to July, considering that *Apodemus* and *Microtus* incisors reflected the one or two last months of life of the animal (Kleve zal et al. 1990; Klevezal 2010; Royer et al. 2013a; Fernández-García et al. 2019). The mean δ18Omw value of the pellet equals an average temperature of 17.3 °C, close to the current mean temperature recorded for June (Climate-Data.org). This mean

![Fig 5: Mean annual temperature (MAT) and mean annual precipitation (MAP) estimations for the Teixoneres Cave sequence with respect to present-day climate in Moià (MAT = 12.3 °C; MAP = 749 mm; Climate-data.org), considering the bioclimatic model (in blue) and mutual ecogeographical range (in green) methods, and the oxygen isotope compositions (in black) of rodent phosphates samples (error is presented in grey fringe).](image-url)
temperature is notably higher than that for the fossil samples of the Teixoneres Cave but is generally consistent with temperature and $\delta^{18}$Omw annual variations for the region, validating established correlations in that region. The incisor recovered from an unknown accumulation of disintegrated pellets recorded 16.1‰ and does not correspond to the same season of accumulation, reporting lower $\delta^{18}$Omw values ($−7.2‰$), probably out of the warm season.

The MATs calculated from the $\delta^{18}$Op of Teixoneres samples are compared with two other methods of palaeoclimatic reconstruction usually used for small-mammal assemblages: the bioclimatic model (BM) and the mutual ecogeographic range (MER) (Fig. 5; Appendixes 3 and 4). Both methods provide lower temperatures than present day: the BM method from $−2.6$ to $−4.4$ °C, whereas MER estimations from $−3.3$ to $−5.3$ °C. $\delta^{18}$O air temperature estimations offer the highest temperatures, recording similar trends as MER temperature reconstructions, but with notably higher estimations than this method ($+1.4/+3.7$ °C with respect to MER). Thus, all the three methods show highly in agreement trends and confirm lower MATs than present day in Moià, a pattern enforced when mean oxygen isotope values are considered (Table 4). In general, homogenous temperature patterns are observed along the sequence with independence of the method considered, although some discrepancies can be observed in subunit IIIa. Despite these differences, BM and MER method estimations generally fall within the confidence interval of $\delta^{18}$O estimations and inter-level differences considering equivalent methods are small. Whereas the MER method records mean annual precipitations (MAP) higher than present day in Moià ($+44/+682$ mm), the BM method supports the opposite trend, with all levels lower than nowadays ($−271$ to $−305$ mm). However, both methods underline slight pluviometry changes along the sequence, with homogenous climatic conditions again.

Discrepancies between these methods can be explained by several factors. BM and MER are both qualitative methods, just transforming species composition changes and the into quantifiable climatic terms. This operation is mainly based on the modern biogeographic distribution of species and linking them to specific biomes or climatic parameters, respectively. The geographical scale considered from both methods either the climatic quantifiable parameters between both methods differed, and this can lead to inconsistencies between methods estimations. BM is based on biome classification (and their derived climatic information) from species spread through Eurasian, whereas MER considers species distribution on Iberia, which directly links to regional temperatures and precipitations. Moreover, small-mammal geographical distribution can be related to climatic factors but also can be a consequence of other causes, such as anthropic pressure, human modification on the ecosystems, species competition, predation, or just the existence of non-analogue faunal associations in the past (Blain et al. 2016; Lyman 2017; Royer et al. 2020). Otherwise, stable isotope estimations are expected to be less susceptible to archaeological biases, but the sampling method and actualistic assumptions linked to the empirical correlations as well as correlation selection according to the taxonomic group and geographical scale of the dataset considered may condition the final temperature estimation (detailed discussion on Fernández-García et al., 2019; 2020). This means that the comparison between palaeoclimatic reconstruction methods, as well as the comparison with other environmental proxies of a site, is key to reach a reliable interpretation.

**Teixoneres sequence and Neanderthal ecological context**

Short-term Neanderthal occupations occurred in 2, in alternation with mammalian carnivore and raptor activities in the site. Competition and cohabitation theories between carnivores/raptors and humans have been highly discussed (e.g., Rosell et al. 2010a, b, 2017; Picin et al. 2020). It is constantly evaluated inside the Middle Palaeolithic, how climatic conditions should play a decisive role in the subsistence strategies and mobility patterns of these human populations and their migration routes or if in some degree this strengthens their adaptive behaviour. The calibrated radiocarbon dates for the Teixoneres Cave sequence (Talamo et al. 2016) allow us to approach global climatic data and the contemporaneous deposition of these units and evaluated how and which of these climatic fluctuations have been recorded at the local scale. MIS 3 (ca. 60–30 ka) is a period characterized by a continuous oscillation of climatic events, including strong and brief cold episodes, such as stadials and Heinrich Events (e.g., Fletcher and Sánchez-Goñi 2008; Fletcher et al. 2010b; Wolff et al. 2010; Rasmussen et al. 2014). However, the southern Europe peninsulas, and particularly the Mediterranean basin, are usually less prone to major climate changes resulting in significant remobilizations of animal communities.

This research has shown that climatic conditions in the Teixoneres Cave surroundings were generally homogenous and always cooler and wetter than nowadays, but slight differences found between lower subunits (unit III) and upper subunits (unit II) from the analysis of small vertebrates of Teixoneres Cave should be addressed. Small variations in the fauna spectrum, in terms of presence/absence and abundances, suggest slight environmental changes notably between subunits (López-García et al. 2012a). For instance, the presence of the Mediterranean species, *Microtus (T.) duodecimcostatus*, is higher (12–14%) in unit III and lower in subunit IIb (<3%), disappearing next in upper subunit IIa. The relative abundance of *A. sylvaticus*, a generalist species with a preference for woodland environments, is remarkably high for subunit IIb (41%) (Appendix 2). The progressive
are interpreted as indicators of cold and dry pulsations. This identified in units III and II (Álvarez-Lao et al. 2017), whichcies are predominant and cold-adapted taxa are very scarce. reflect a faunal mixture in which temperate ungulate spe-

causes are primarily rejected as the main explanation formost small-mammal inputs, so taphonomicditions were relatively temperate and humid (López-García corrediscrepancies are found between methods and with previous
test (− 4.4 °C) and driest (− 305 mm) conditions than present
day in the region, but not for the MER method. Thus, some
discrepancies are found between methods and with previousenvironmen
tal studies, particularly for unit III which was
tentatively related to cooler conditions than in upper subunits,or at least to punctuated climatic instability. Based on the
geochemical record, MATs estimated are the lowest of thesequence (IIIb: 10.8 ± 2.6 °C; IIIa: 10.7 ± 2.6 °C). Moreover,
unit III contents reflect drier conditions than for unit IIwhen calculated by the MER method, while the BM methodprovidedit notably lower precipitation rates for all four sub-
units. Contrary to unit III, rodent teeth oxygen isotopecompositions of the upper subunits (IIb–IIa) have higher δ18O means and medians (18.2–18.5‰), a lower intra-
level amplitude (3.5–3.8‰), and lower δ18O averages mean values, ten-
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may be the most plausible explanation that these subunits are the result of slow sedimentation rates that promoted theoverlap between different climatic events, precluding theisolation of different stadial and interstadial events from thefaunal spectra. A mix of species dependent on either temper-
ate or cold conditions points to the sedimentary recording ofpunctual cold events in agreement with the occurrence of thewoolly mammoth and the woolly rhinoceros, but that undercurrent small-mammal analysis is not possible to isolate,giving a general faunal spectrum that seems globally temper-
ate. Probably, small mammals from unit III are recording a
long-term environmental trend based on temperate condi-
tions and woody landscape, which is coincident with thepalyнологical sequence (Ochando et al. 2020).

Similarly, the cooler and drier conditions detected bysmall-mammal assemblages in unit II were tentatively asso-
ciated with cold events and even to Heinrich Event peaks ofMIS 3 (López-García et al. 2012a, 2014), which could beconsistent with the punctual occurrence of C. antiquitatis in this level (Álvarez-Lao et al. 2017). But considering thenew radiometric dates of this unit (ca. 33–44 ka) (Talamo et al. 2016), it should correspond to HE4 if this correlation is accepted (Hemming 2004). However, this climatic correlation with an exceptional cold and dry event cannot be fully supported by the results of this work. Moreover, small mammals are not directly linked to human presence in thecave and their accumulation is produced in other seasons than human visits to the cave. This is a complex issue to
| Unit | Subunits | Pollen and charcoal sources (López-García et al., 2012a, b) | New data pollen (Ochando et al., 2020) | Large mammals and palaeodiet (Álvarez-Lao et al., 2017; Carlos-Sánchez et al., 2020a, 2020b) | Small vertebrates (López-García et al., 2012a, b) | Oxygen isotope analysis (This work) | Small mammals’ predator (This work) | Past temperatures (this work & López-García et al., 2012a, b) | Past precipitations (this work & López-García et al., 2012a, b) |
|------|----------|----------------------------------------------------------------|---------------------------------|---------------------------------------------------------------|--------------------------------------------------|---------------------------------|--------------------------------|--------------------------------|--------------------------------|
| II (ca. 33–44 ka) | Ia | Open woodland and meadows. Dry conditions in relation to steppe taxon *Artemisia* (Ia), *Pinus*, *Quercus* and angiosperm charcoals | Mix oak-pine forest. AP > 65%, with evergreen *Quercus* (16–47%), *Pinus nigra-sylvestris* (8–17%), and *Juniperus* (1–17%). Herbs: *Poaceae*, *Artemisia* are dominant in NAP | Abundant temperate-adapted species (mainly *Cervus elaphus* followed by equids), presence of *Coelodonta antiquitatis*, and higher presence of *Rupicapra Capra pyrenaica* and *Sus scrofa*. Open forest and rocky landscape | Homogenous, low amplitude, and the highest median. Stable period | Nocturnal raptors; digestion = 43% | BM = 9.4 ºC/ MER = 9 ºC/ δ18O = 11.8 ºC | BM = 477 mm/ MER = 981 mm |
|      | Iib | Mix oak-pine forest. AP = 58–74%, dominated by evergreen *Quercus* (17–52%), *Juniperus* (2–13%), *Pinus* (9–20%). Herbs: *Poaceae*, *Artemisia*, Amaranthaceae | Abundant Mid-European species (*M. agrestis*, *M. arvalis*), scarce presence of forest dwellers (*A. sylvaticus*), and absence of Mediterranean species. Cooler and drier phase related to stadial. Open woodland | Homogenous, the lowest amplitude, and high median. Stable period | Nocturnal raptors; digestion = 31% | BM = 7.9 ºC/ MER = 8.8 ºC/ δ18O = 11.3 ºC | BM = 444 mm/ MER = 950 mm |
| Unit     | Subunits | Pollen and Charcoals (López-García et al., 2012a, b) | New data pollen (Ochando et al., 2020) | Large mammals and palaeodiet (Álvarez-Lao et al., 2017; Carlos-Sánchez et al., 2020a, 2020b) | Small vertebrates (López-García et al., 2012a, b) | Oxygen isotope analysis (This work) | Small mammals’ predator (This work) | Past temperatures (this work & López-García et al., 2012a, b) | Past precipitations (this work & López-García et al., 2012a, b) |
|----------|----------|------------------------------------------------------|----------------------------------------|---------------------------------------------------------------------------------------------|-------------------------------------------------|-----------------------------------|---------------------------------|-------------------------------------------------|-------------------------------------------------|
| III (ca. 44–51 ka) IIIa | Open woodland and meadows. Relative moisture, due to high representation of Corylus and Pteridophyta spores. Buxus charcoals | Mix oak-pine forest. AP> 87%. Evergreen Quercus is outstanding (7–56%); Pinus halepensis-pinea (2–31%), Juniperus (1–30%), and Pinus nigra-sylvestris (14–16%). Several fungal spore | Abundant temperate-adapted (mainly Cervus elaphus and some equids) but scarce cold-adapted species (Mammuthus primigens in IIIa and Coelodonta antiquitatis in IIIb). Open forested landscape with patchy areas. Increase of wild ass browser behaviour indicates wetter conditions in IIb, whereas increasing in grass consumption by equids and cervids indicates open areas and worsening conditions for IIIa | Mediterranean species (M. (T.) duodecimcostatus, I. calberae), Mid-European species (M. arvalis, M. agrestis, C. nivalis, M. (T.) gerbeii), and forest dwellers (A. sylvaticus). Temperate and humid phase related to interstadial. Open woodland but increase in humid meadows and water | Heterogenous, medium amplitude, the lowest median. Unstable | Bubo bubo; digestion=36% | BM=9.2 °C/ MER=7 °C/ δ18O =10.7 °C | BM=476 mm/ MER=925 mm |
| IIIb | Mix oak-pine forest. AP=60%; Pinus (11–26%); Quercus (14–46%); Juniperus (3–8%), Castanea, Betula, Corylus. Herbs: Poaceae, Artemisia, Amaranthaceae | Abundant presence of forest dwellers (A. sylvaticus), presence of Mediterranean (M. (T.) duodecimcostatus), and Mid-European (M. arvalis, M. agrestis) species. Temperate and humid phase related to interstadial. Open woodland | Heterogenous, low median, and the largest amplitude. Unstable | Bubo bubo+carnivores occasional inputsdigestion=36% | BM=13.4 °C/ MER=9.7 °C/ δ18O =10.8 °C | BM=463 mm/ MER=840 mm | BM=463 mm/ MER=840 mm |
be addressed for climatic interpretation because it cannot be evaluated if it is just in different seasons from not very distant years. Considering that there is not a clear spatial or stratigraphical distinction between small mammals and archaeological remains in the cave sediments, we assumed that small mammals and remains derived from human occupations even if being introduced in different seasons or seasons from different years are coincident in few-millennial sedimental accumulation. Thus, if abrupt climatic changes are experienced could be detected from both approaches. Nevertheless, only a complete spatial distribution study and direct radiocarbon dating on small mammals remains, as well as a comparison with biochemical analyses on hunted ungulate, could give us some clues on the overlapping and the chronological resolution.

The present study underscores that despite these punctual cold episodes marked by the presence of small- and large-mammal cold-adapted species, the local-to-regional ecological conditions experienced by Neanderthal groups in this region were milder and highly conservative; differences of less than 1 °C between all subunits are estimated considering oxygen isotope compositions of rodent teeth and with mean standard deviations < 1 °C for MER and BM methods. Recent works point to the Late Pleistocene “tundra-steppe” biome extension to southern latitudes as one of the main drivers in C. antiquitatis spread waves (Puzachenko et al. 2021; Rodriguez-Almagro et al. 2021). However, palynological studies of the Teixoneres Cave sequence (Ochando et al. 2020) confirmed that the global trend is a dense oak-pine forest in all subunits, with arboreal pollen oscillating between 60 and 90% during the whole sequence and with unfrequented high values of oak pollen, considering that pine forests usually dominated in the Mediterranean area (Allué et al. 2017; Carrión et al. 2018; Ochando et al. 2020, 2021). Cold component pollen taxa (such Artemisia, Poaceae, Amaranthaceae, Erica, Ephedra fragilis) experienced punctuated increases from subunits IIb and IIIb but are anecdotal. Indeed, the woodland component is always the dominant type of landscape considering small vertebrates in spite of faunal frequencies (López-García et al. 2012a).

These results are in agreement with regional interpretations of MIS 3 equivalent small-mammal studies performed in Abric Romani rock-shelter (Burjachs et al. 2012; Fernández-García et al. 2018, 2020) and Xaragalls cave (López-García et al. 2012a; Fernández-García et al. 2019). In both sequences, a detailed climatic characterization based on small-mammal assemblages has been accomplished through a combination of taxonomy, taphonomy, and oxygen isotope studies. Oxygen isotope compositions of rodent bioapatite phosphate are usually homogenous with higher and less scattered values than equivalent studies from northern latitudes (Royer et al. 2013a, b, 2014). Environmental reconstruction proposed for these sites points always to wetter, cooler, and homogenous climatic cycles, and forested areas next to the settlements. A general trend is observed from northeastern Iberia small-mammal communities (López-García et al. 2014; Fernández-García et al. 2016), but also extent to the Mediterranean area, as documented in Arbreda Cave (López-García et al. 2015), Terrasses de la Riera dels Canyars (López-García et al. 2013), Cova del Gegant (López-García et al. 2012b), Cova del Coll Verdaguera (Daura et al. 2017), El Salt (Fagoaga et al. 2018, 2019), or Gorham’s Cave (López-García et al. 2011). Pollen and charcoal analyses conducted in some of these sites agree on a predominance of forests composed of conifers (mainly Pinus and Juniperus) and evergreen and deciduous species (González-Sampérez et al. 2010; Burjachs et al. 2012; Carrión 2012; Allué et al. 2017; Daura et al. 2017). Stable climatic conditions, which prevailed during MIS3, sheltered a high biodiversity during the Late Pleistocene as also proposed for the Levantine area (including the MIS3 records from Cova de les Malladetes and Cova de les Cendres) based on charcoal, pollen, and faunal studies (Real et al. 2021).

These findings are in line with research undergone on large-mammal faunas palaeodiet traces. Sánchez-Hernández et al. (2020a) reconstruct the Neanderthal settlement patterns for several archaeological sites in the Mediterranean area (Arbreda, Teixoneres, Abric Romaní, and El Salt), based on tooth wear and cementum analysis. From this research can be indirectly inferred patched areas with comparable proportions of open and wooded areas. Environmental oscillations are buffered for these sites that are continuously visited by human groups and are not expected to alter logistic models of settlement and mobility of Neanderthal groups, based on short-seasonal movements. Moreover, from the comparison on feeding behaviours of ungulates (Cervus elaphus, Equus ferus, and Equus hydruntinus) from Teixoneres and Arbreda caves, Sánchez-Hernández et al. (2020b) observed a maintenance of the same dietary behaviour of red deer and horses in both sites pointing to steady local environmental conditions over a long-term scale, independently of altitude. The dietary features observed for ungulates correlate well with the relative temperate and humidity conditions for the studied assemblage and from regional small-mammal trends already mentioned.

Among the less abrupt climatic conditions widely explained that Iberia experienced during the Last Glacial, Teixoneres Cave as well as other archaeological sites of northeastern Iberia underline how the Mediterranean corridor allows optimal conditions for the maintenance of Neanderthal populations. The high environmental and climatic variability offered by the Iberian Peninsula not only by its southern position but also by the high variability of micro-environments provides the recurrent mosaic landscape that oscillated from more open to close forests (Allué et al. 2017;
Fernández-García et al. 2020; Ochando et al. 2021; Real et al. 2021). These Neanderthal populations take advantage of the temperate forest provided by the Mediterranean area. This allows long-term preservation of a high diversity of resources available to be exploited by Neanderthal groups, which even when fluctuating in location or abundance probably did not pose a problem for Neanderthal subsistence considering their behavioural plasticity. It does not exclude, however, that punctual colder pulsations were experienced and affected past ecosystems, as cold-adapted large and small faunal episodes have revealed. But, probably, the mosaic landscapes sustained in the Mediterranean area allowed maintenance of these human group subsistence strategies, as well as their preys and their usual plant resources.

Conclusions

The analyzed small mammals accumulated in the cave mainly due to predation by nocturnal raptoress. In agreement with taphonomic analyses performed on bird and leporid remains, the main agent corresponds to the eagle owl (Bubo bubo), at least in unit III. Bubo bubo—type predators probably maintained their roosting or nesting place in the Teixoneres Cave during the formation of unit II, but other owls may have contributed with small-mammal inputs as well. Consistent with other small-prey studies in the sequence, the punctual intervention of mammal carnivores can be suggested for level IIIb. Among post-depositional agents, plant activity, cracking, and manganese oxide coating are common, indicating a wet fossiliferous microenvironment with humidity fluctuations, where abundant breakage rates are observed. Some burned remains have been detected in relation to unintentional burning.

The medium to low intra-level ranges of oxygen isotope values point to a preferential accumulation of rodents, probably during the spring season. This period of frequen-
tation by owls is in accordance with previous seasonality inferences based on several medullary bones of birds, which share a common origin with small mammals, and results generally differed from use-wear and dental eruption analyses on ungulates, mainly related to hominin contributions to the cave. According to δ18O estimations of temperatures, slightly cooler climatic conditions than present were recorded (−1.6/−0.5 °C). The observed trend is globally in agreement with MER estimations, which recorded colder MATs (−2.9/−5.3 °C), but differ in unit III with BM, which recorded MATs similar to present-day levels (+0.1/+1.1 °C). Both methods calculated wetter conditions than during present day (+44 mm/ +682 mm).

Only slight changes are observed between lower unit III and unit II, which may tentatively be related to some climatic instability. This trend is not fully in agreement with previous environmental interpretations based on small-mammal relative abundances but may be in keeping with the occasional occurrence of cold-dependent large and small-mammal species. These inconsistencies between methods could point to a palimpsest where stadial and interstadial events overlapped, especially in subunit IIIa. However, oxygen isotope compositions of analyzed rodent teeth underscore the presence of rather stable climatic conditions across the sequence, with little variation between subunits, including those expressed in MAT (<1 °C). A favorable and milder climatic scenario is confirmed for the Mediterranean area in combination with constant presence of open forests that potentially sustained the faunal and vegetal resources preferred by Neanderthal groups.

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Author contribution Conceptualization, investigation, material preparation, data collection, and formal analysis were mainly performed by M. Fernández-García. J. M. López-García, A. Royer, and C. Lécuyer were actively involved in the conceptualization, formal analysis, investigation, resources, and writing and editing of the original draft. R. Blasco, F. Rivals, J. Rosell, and A. Rufà provide guidance in the conceptualization, resources, and reviewing-editing of the manuscript. The first draft of the manuscript was written by M. Fernández-García, and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

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Declarations

Conflict of interest  The authors declare no competing interests.

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References

Allué E, Solé A, Burguet-Coca A (2017) Fuel exploitation among Neanderthals based on the anthropological record from Abric Romani (Capellades, NE Spain). Quat Int 431:6–15. https://doi.org/10.1016/j.quaint.2015.12.046

Álvarez-Lao DJ, Méndez M (2016) Latitudinal gradients and indicator species in ungulate paleoassemblages during the MIS 3 in W Europe. Palaeogeogr Palaeoclimatol Palaeoecol 449:455–462. https://doi.org/10.1016/j.palaeo.2016.02.050

Álvarez-Lao DJ, Rivals F, Sánchez-Hernández C et al (2017) Ungulates from Teixoneres Cave (Moia, Barcelona, Spain): presence of cold-adapted elements in NE Iberia during the MIS 3. Palaeogeogr Palaeoclimatol Palaeoecol 466:287–302. https://doi.org/10.1016/j.palaeo.2016.11.040

Álvarez-Lao DJ, Ruiz-Zapata MB, Gil-García MJ et al (2015) Palaeoenvironmental research at Rexidora Cave: new evidence of cold and dry conditions in NW Iberia during MIS 3. Quat Int 379:35–46. https://doi.org/10.1016/j.quaint.2015.04.062

Andrews P (1990) Owls, Caves and Fossils. Predation, preservation and accumulation of small mammal bones in caves, with an analysis of the Pleistocene Cave Faunas from Westbury-sub-Mendip, Somerset, UK. The University of Chicago, Chicago

Arribas Ó (2004) Fauna y paisaje de los Pirineos en la Era Glaciar. Lynx, Barcelona

Bennásar M (2010) Tafonomía de micromamíferos del Pleistoceno Inferior de la Sierra de Atapuerca (Burgos): Sima del Elefante y Gran Dolina. Universitat Rovira i Virgili (Tarragona)

Bennásar M, Cáceres I, Cuenca-Bescós G (2016) Palaeoecological and microenvironmental aspects of the first European hominids inferred from the taphonomy of small mammals (Sima del Elefante, Sierra de Atapuerca, Spain). Comptes Rendus - Palevol 15:635–646. https://doi.org/10.1016/j.palevol.2015.07.006

Bernard A, Daux V, Lécuyer C et al (2009) Pleistocene seasonal temperature variations recorded in the δ18O of Bison priscus teeth. Earth Planet Sci Lett 283:133–143. https://doi.org/10.1016/j.epsl.2009.04.005

Blain HA, Lozano-Fernández J, Agustí J et al (2016) Refining upon the climatic background of the Early Pleistocene hominin settlement in western Europe: Barranco León and Fuente Nueva-3 (Guadix-Baza Basin, SE Spain). Quat Sci Rev 144:132–144. https://doi.org/10.1016/j.quascirev.2016.05.020

Bowen GJ (2017) The Online Isotopes in Precipitation Calculator, Version 3.1 (4/2017). http://waterisotopes.org

Burjachs F, López-García JM, Allué E et al (2012) Palaeoecology of Neanderthals during Dansgaard-Oeschger cycles in northeastern Iberia (Abric Romaní): from regional to global scale. Quat Int 247:26–37. https://doi.org/10.1016/j.quaint.2011.01.035

Bustos-Pérez G, Chacón MG, Rivals F et al (2017) Quantitative and qualitative analysis for the study of Middle Paleolithic retouched artifacts: unit III of Teixoneres cave (Barcelona, Spain). J Archaeol Sci Reports 12:658–672. https://doi.org/10.1016/j.jasrep.2017.02.021

Carrión JS (2012) Paleoflora y paleovegetación de la Península Ibérica e Islas Baleares: Plioceno-Cuaternario. Ministerio de Economía y Competitividad - Universidad de Murcia, Murcia

Carrión JS, Ochando J, Fernández S et al (2018) Last Neanderthals in the warmest refugium of Europe: palynological data from Vanguard Cave. Rev Palaeobot Palynol 259:63–80. https://doi.org/10.1016/j.revpalbo.2018.09.007

Chaline J (1974) Les proies des rapaces. Doin Éditeurs, Paris

Climate-Data.org. (2018): https://es.climate-data.org/

Comay O, Dayan T (2018) Taphonomic signatures of owls : new insights into micromammal assemblages. Palaeoegor Palaeoclimatol Palaeoecol 492:81–91. https://doi.org/10.1016/j.palaeo.2017.12.014

Crowson RA, Showers WJ, Wright EK, Hoering TC (1991) Preparation of phosphate samples for oxygen isotope analysis. Anal Chem 63:2397–2400. https://doi.org/10.1021/ac001829320058-W

D’Errico F, Sánchez Goñi MF (2003) Neandertal extinction and the millennial scale climatic variability of OIS 3. Quat Sci Rev 22:769–788. https://doi.org/10.1016/S0277-3791(03)00009-X

Dansgaard W (1964) Stable isotopes in precipitation. Tellus XVI:436–468

Daura J, Sanz M, Allué E et al (2017) Palaeoenvironments of the last Neanderthals in SW Europe (MIS 3): Cova del Coll Verdager (Barcelona, NE of Iberian Peninsula). Quat Sci Rev 177:34–56. https://doi.org/10.1016/j.quascirev.2017.10.005

Fagoaga A, Laplana C, Marquina-blasco R et al (2019) Palaeoecological context for the extinction of the Neanderthals : a small mammal study of stratigraphic unit V of the El Salt site, Alcoi, eastern Spain. Palaeogeogr Palaeoclimatol Palaeoecol 530:163–175. https://doi.org/10.1016/j.palaeo.2019.05.007

Fagoaga A, Ruiz-Sánchez FJ, Laplana C et al (2018) Palaeoecological implications of Neanderthal occupation at Unit Xb of El Salt (Alcoi, eastern Spain) during MIS 3 using small mammals proxy. Quat Int 401:101–112. https://doi.org/10.1016/j.quaint.2017.10.024

Fernández-García M, López-García JM, Bennásar M et al (2018) Palaeoenvironmental context of Neanderthal occupations in northeastern Iberia: the small-mammal assemblage from Abric Romani (Capellades, Barcelona, Spain). Palaeoegor Palaeoclimatol Palaeoecol 506:154–167. https://doi.org/10.1016/j.palaeo.2018.06.031

Fernández-García M, López-García JM, Lorenzo C (2016) Palaeoecological implications of rodents as proxies for the Late Pleistocene-Holocene environmental and climatic changes in northeastern Iberia. CR Paleol 15:707–719. https://doi.org/10.1016/j.crpy.2015.08.005

Fernández-García M, López-García JM, Royer A et al (2020) Combined palaeoecological methods using small-mammal assemblages to decipher environmental context of a long-term Neanderthal settlement in northeastern Iberia. Quat Sci Rev 228:106072. https://doi.org/10.1016/j.quascirev.2019.106072

Fernández-García M, Royer A, López-García JM et al (2019) Unraveling the oxygen isotope signal (δ18O) of rodent teeth from northeastern Iberia, and implications for past climate reconstructions. Quat Sci Rev 218:107–121. https://doi.org/10.1016/j.quascirev.2019.04.035
Fernández-Jalvo Y (1992) Tafonomía de microvertebrados del complejo cártico de Atapuerca (Burgos). Universidad Complutense de Madrid

Fernández-Jalvo Y, Andrews P (1992) Small mammal taphonomy of Gran Dolina, Atapuerca (Burgos), Spain. J Archaeol Sci 19:407–428. https://doi.org/10.1016/0305-4403(92)90058-B

Fernández-Jalvo Y, Andrews P, Denys C et al (2016) Taphonomy for taxonomists: Implications of predation in small mammal studies. Quat Sci Rev 139:138–157. https://doi.org/10.1016/j.quascirev.2016.03.016

Fernández-Jalvo Y, Andrews PJ (2016) Atlas of taphonomic identifications: 1001+ images of fossil and recent mammal bone modification. Springer, Dordrecht

Fernández-Jalvo Y, Tormo L, Andrews P, Marin-Monfort MD (2018) Taphonomy of burnt bones from Wonderwerk Cave (South Africa). Quat Int 495:19–29. https://doi.org/10.1016/j.quaint.2018.05.028

Finlayson C, Carrión JS (2007) Rapid ecological turnover and its impact on Neanderthal and other human populations. Trends Ecol Evol 22:213–222. https://doi.org/10.1016/j.tree.2007.02.001

Finlayson C, Giles Pacheco F, Rodríguez-Vidal J et al (2006) Late Quaternary environmental reconstructions. PLoS ONE 7:16–27. https://doi.org/10.1371/journal.pone.0001583

Fletcher WJ, Sánchez-Goñi MF (2008) Orbital- and sub-orbital-scale climate impacts on vegetation of the western Mediterranean basin over the last 48,000 yr. Quat Res 70:451–464. https://doi.org/10.1016/j.yqres.2008.07.002

Fletcher WJ, Sánchez-Goñi MF, Peyron O, Dormoy I (2019) Abrupt climate changes of the last deglaciation detected in a Western Mediterranean forest record. Clim past 6:245–264

Fletcher WJ, Sánchez-Goñi MF, Allen JR et al (2010) Millennial-scale variability during the last glacial in vegetation records from Europe. Quat Sci Rev 29:2839–2864. https://doi.org/10.1016/j.quascirev.2009.11.015

Fourel F, Martineau F, Lécuyer C et al (2011) 180O/16O ratio measurements of inorganic and organic materials by elemental analysis–pyrolysis–isotope ratio mass spectrometry continuous-flow techniques. Rapid Commun Mass Spectrom 25:2691–2696. https://doi.org/10.1002/rcm.5056

Gehler A, Tütken T, Pack A (2012) Oxygen and carbon isotope variations in a modern rodent community - implications for palaeoenvironmental reconstructions. PLoS ONE 7:16–27. https://doi.org/10.1371/journal.pone.0001583

González-Sampériz P, Leroya. G, Carrión JS, S et al (2010) Steppes, Gehler A, Tütken T, Pack A (2012) Oxygen and carbon isotope variations in a modern rodent community - implications for palaeoenvironmental reconstructions. PLoS ONE 7:16–27. https://doi.org/10.1371/journal.pone.0001583

Hewitt GM (2000) The genetic legacy of the Quaternary ice ages. Nature 405:907–913. https://doi.org/10.1038/35016000

Higham T, Douka K, Wood R et al (2014) The timing and spatiotemporal patterning of Neanderthal disappearance. Nature 512:306–309. https://doi.org/10.1038/nature13621

Hut G (1987) Consultants’ group meeting on stable isotope reference samples for geochemical and hydrological investigations, 16–18 Sep. 1985. International Atomic Energy Agency, Vienna

IAEA/WMO, 2018. Global network of isotope precipitation. The GNIP Database. Accessible at WISER (Water Isotope System for Data Analysis, Visualization and Electronic Retrieval): https://nucleus.iaea.org/wiser.

Jones EL (2021) What is a refugium? Questions for the Middle–Upper Palaeeolithic transition in peninsular southern Europe. J Quat Sci 1–6. https://doi.org/10.1002/jqs.3274

Klevezal GA (2010) Dynamics of incisor growth and daily increments on the incisor surface in three species of small rodents. Biol Bull 37:836–845. https://doi.org/10.1134/S1062359010080078

Klevezal GA, Ek MPUC, Sukhovskaja LI (1990) Incisor Growth in Europe. Quat Sci Rev 29:2839–2864. https://doi.org/10.1016/0277-3791(90)90085-W

Kowalski K (1995) Taphonomy of bats (Chiroptera). Geobiology 18:251–256. https://doi.org/10.1016/S0016-0022(95)80172-3

Lagos P (2019) Predation and its effects on individuals: from individual to species. Elsevier Ltd.

Lécuyer C, Fourel F, Martineau F et al (2007) High-precision determination of 18O/16O ratios of silver phosphate by EA-pyrolysis-IRMS continuous flow technique. J Mass Spectrom 42:36–41

Lécuyer C, Grandjean P, O’Neill JR et al (1993) Thermal excursions in the ocean at the Cretaceous-Tertiary boundary (northern Morocco): 818O record of phosphatic fish debris. Palaeogeogr Palaeoclimatol Palaeoecol 105:235–243. https://doi.org/10.1016/0016-7037(93)90085-W

Lindars ES, Grimes ST, Mattey DP et al (2001) Phosphate d18O determination of modern rodent teeth by direct laser fluorination: an appraisal of methodology and potential application to palaeoclimate reconstruction. Geochim Cosmochim Acta 65:2535–2548

Lloveras L, Moreno-García M, Nadal J (2008) Taphonomic study of leporid remains accumulated by the Spanish Imperial Eagle (Aquila adalberti). Geobiol 41:91–100. https://doi.org/10.1016/j.geobios.2006.11.009

Lloveras L, Moreno-García M, Nadal J (2012) Feeding the foxes: an experimental study to assess their taphonomic signature on leporid remains. Int J Osteoarchaeol 22:577–590. https://doi.org/10.1002/oja.1280

Lloveras L, Moreno-García M, Nadal J (2008) Taphonomic analysis of leporid remains obtained from modern Iberian lynx (Lynx pardinus) scats. J Archaeloe Sci 35:1–13. https://doi.org/10.1016/j.jas.2007.02.005

Longinelli A (1984) Oxygen isotopes in mammal bone phosphate: a new tool for paleohydrological and paleoclimatological research? Geochim Cosmochim Acta 49:781–788. https://doi.org/10.1016/0016-7037(84)90259-X

López-García JM, Blain H-A, Bennásar M et al (2013) Heinrich event 4 characterized by terrestrial proxies in southwestern Europe. Clim past 9:1053–1064. https://doi.org/10.5194/cp-9-1053-2013

López-García JM, Blain H-A, Burjachs F et al (2012) A multidisciplinary approach to reconstructing the chronology and environment of southwestern European Neanderthals: the contribution of Teixoneres cave (Moïà, Barcelona, Spain). Quat Sci Rev 43:33–44. https://doi.org/10.1016/j.quascirev.2012.04.008

López-García JM, Blain HA, Bennásar M, Fernández-García M (2014) Environmental and climatic context of neanderthal occupation in southwestern Europe during MIS3 inferred from the small-vertebrate assemblages. Quat Int 326–327:319–328. https://doi.org/10.1016/j.quaint.2013.09.010

López-García JM, Blain HA, Sanz M, Daura J (2012b) A coastal reservoir of terrestrial resources for neanderthal populations in north-eastern Iberia: Palaeoenvironmental data inferred from the
small-vertebrate assemblage of Cova del Gegant, Sitges, Barcelona. J Quat Sci 27:105–113. https://doi.org/10.1002/jqs.1515

López-García JM, Cuenca-Bescós G, Finlayson C et al (2011) Palaeoenvironmental and palaeoclimatic proxies of the Gorham’s cave small mammal sequence, Gibraltar, southern Iberia. Quat Int 243:137–142. https://doi.org/10.1016/j.quaint.2010.12.032

López-García JM, Soler N, Maroto J et al (2015) Palaeoenvironmental and palaeoclimatic reconstruction of the Latest Pleistocene of L’Arbreda Cave (Serinyà, Girona, northeastern Iberia) inferred from the small-mammal (insectivore and rodent) assemblages. Palaeogeogr Palaeoclimatol Palaeoecol 435:244–253. https://doi.org/10.1016/j.palaeo.2015.06.022

Luz B, Kolodny Y, Horowitz M (1984) Fractionation of oxygen isotopes between mammalian. Geochim Cosmochim Acta 48:1689–1693

Luzi E, López-García JM, Blasco R et al (2017) Variations in Microut arvalis and Microut agrestis (Arvicolinae, Rodentia) dental morphologies in an archaeological context: the case of Teixoneres Cave (Late Pleistocene, North-Eastern Iberia). J Mamm Evol 24:495–503. https://doi.org/10.1007/s10914-016-9355-8

Lyman RL (2017) Paleoenvironmental reconstruction from faunal remains: ecological basics and analytical assumptions. J Archaeol Res 1–57. https://doi.org/10.1007/s10814-017-9102-6

Manzanares A (2012) Aves rapaces de la Península Ibérica, Baleares y Canarias. Ediciones Omega, Barcelona

Marín-Arroyo AB, Rios-Garaiar J, Straus LG et al (2018) Chronological reassessment of the Middle to Upper Paleolithic transition and Early Upper Paleolithic cultures in Cantabrian Spain. PLoS ONE 13:1–20. https://doi.org/10.1371/journal.pone.0194708

Maroto J, Julià R, López-García JM, Blain H-A (2012) Chronological and environmental context of the Middle Pleistocene human tooth from Mollet Cave (Serinyà, NE Iberian Peninsula). J Hum Evol Evol 62:655–663. https://doi.org/10.1016/j.jhevol.2012.01.009

Mateo-Lomba P, Rivals F, Blasco R, Rosell J (2019) The use of bones as retouchers at Unit III of Teixoneres Cave (MIS 3; Moià, Barcelona, Spain). J Archaeol Sci Reports 27:101980. https://doi.org/10.1016/j.jasrep.2019.101980

Mikkola H (1983) Owls of Europe. Buteo Books, Sussex

Moreno A, Svensson A, Brooks SJ et al (2014) A compilation of Western European terrestrial records 60–8kaBP: towards an understanding of latitudinal climatic gradients. Quat Sci Rev 106:167–185. https://doi.org/10.1016/j.quascirev.2014.06.030

Navarro N, Lécuyer C, Montuire S et al (2004) Oxygen isotope compositions of phosphate from arvicoline teeth and Quaternary climatic changes, Gigny, French Jura. Quat Res 62:172–182. https://doi.org/10.1016/j.yqres.2004.06.001

Norrdahl K, Korpima E (2002) Seasonal changes in the numerical responses of predators to cyclic vole populations. Ecography (cop) 25:428–438

Ochando J, Amorós G, Carrión JS, et al (2021) Iberian Neandertals in forests and savannahs. J Quat Sci 1–28. https://doi.org/10.1002/jqs.3339

Ochando J, Carrión JS, Blasco R, et al (2020) Neandertals in a highly diverse, mediterranean-Eurosiberian forest ecotone: the pleistocene pollen record of Teixoneres Cave, northeastern Spain. Quat Sci Rev 241. https://doi.org/10.1016/j.quascirev.2020.106429

Olsson V (1979) Studies on a population of Eagle owls in southeast Sweden. Viltrevy 11:1–99

Palomo LJ, Gisbert J, Blanco C (2007) Atlas y libro rojo de los mamíferos terrestres de España. Organismo Autónomo Parques Nacionales, Madrid

Penneyd E, Candy I, Schreve DC (2019) Variability in the oxygen isotope compositions of modern rodent tooth carbonate: implications for palaeoclimatic reconstructions. Palaeogeogr Palaeoclimatol Palaeoecol 514:695–705. https://doi.org/10.1016/j.palaeo.2018.11.017

Picin A, Chacón MG, Gómez de Soler B et al (2020) Neandertal mobile toolkit in short-term occupations at Teixoneres Cave (Moia, Spain). J Archaeol Sci Reports 29:102165. https://doi.org/10.1016/j.jasrep.2019.102165

Rasmussen SO, Bigler M, Blockley SP et al (2014) A stratigraphic framework for abrupt climatic changes during the Last Glacial period based on three synchronized Greenland ice-core records: Refining and extending the INTIMATE event stratigraphy. Quat Sci Rev 106:14–28. https://doi.org/10.1016/j.quascirev.2014.09.007

Real C, Martínez-varela CM, Carrión Y, et al (2021) Could the central-eastern Iberian Mediterranean region be defined as a refugium? Fauna and flora in MIS 5–3 and their implications for Palaeolithic human behaviour. J Quat Sci 1–17. https://doi.org/10.1002/jqs.3285

Rivas-Martínez S (1987) Memoria del Mapa de Series de Vegetación de España. ICONA, Serie Tecnica, Madrid

Rivas-Martínez S, Asensi A, Diez -Garretas B, et al (2007) Mapa de series, geoseries y geoprocessaries de vegetación de España. In: Memoria del Mapa de Series de Vegetación de España. Parte I. Itineraria geobotánica, pp 5–436

Rosell J, Blasco R, Cebrià A, et al (2008) Mosseguedes i Levallolis: les noves intervencions a la Cova de les Teixoneres (Moia, Bages). Trib d’Arqueologia 29–43.

Rosell J, Blasco R, Rivals F et al (2010) Las ocupaciones en la Cova de les Teixoneres (Moia, Barcelona): relaciones espaciales y grado de competencia entre hienas, osos y neandertales durante el Pleistoceno superior. Zo Arqueol 13:392–403

Rosell J, Blasco R, Rivals F et al (2010) A step along the way: the role of neanderthal groups at level III of Teixoneres cave (Moia, Barcelona, Spain). Quaternaire 21:139–154

Rosell J, Blasco R, Rivals F et al (2017) A resilient landscape at Teixoneres Cave (MIS 3; Moia, Barcelona, Spain): the Neandertals as disrupting agent. Quat Int 435:195–210. https://doi.org/10.1016/j.quascirev.2015.11.077

Rosell J, Blasco R, Rivals F et al (2014) Cova del Toll y Cova de les Teixoneres Moia, Barcelona. In: Sala Ramos R (ed) Pleistocene and Holocene hunter-gatherers in Iberia and the Gibraltar Strait: the current archaeological record. Universidad de Burgos, Burgos, pp 302–307

Royer A, García Yelo BA, Laffont R, Hernández Fernández M (2020) New bioclimatic models for the quaternary palaearctic based on insectivore and rodent communities. Palaeogeogr Palaeoclimatol Palaeoecol 560:110040. https://doi.org/10.1016/j.palaeo.2020.110040

Royer A, Lécuyer C, Montuire S et al (2013) What does the oxygen isotope composition of rodent teeth record? Earth Planet Sci Lett 361:258–271. https://doi.org/10.1016/j.epsl.2012.09.058

Royer A, Lécuyer C, Montuire S et al (2014) Summer air temperature, reconstructions from the last glacial stage based on rodents from the site Taillis-des-Coteaux (Vienne), Western France. Quat Res 82:420–429. https://doi.org/10.1016/j.yqres.2014.06.006

Royer A, Lécuyer C, Montuire S et al (2013) Late Pleistocene (MIS 3–4) climate inferred from micromammal communities and δ18O of rodents from Les Pradelles, France. Quat Res 80:113–124. https://doi.org/10.1016/j.yqres.2013.03.007

Royer A, Montuire S, Gilg O, Leroulandie V (2019) A taphonomic investigation of small vertebrate accumulations produced by the snowy owl (Bubo scandiacus) and its implications for fossil studies. Palaeogeogr Palaeoclimatol Palaeoecol 514:189–205. https://doi.org/10.1016/j.palaeo.2018.10.018

Rozanski K, Araguz-araguz L, Gonfiantini R (1993) Isotopic patterns in modern global precipitation. In: Swart PK, Lohman KC, McKenzie J, Savin S (eds) Climate Change in Continental Isotopic Records. Washington, pp 1–36

Rufà A, Blasco R, Rivals F, Rosell J (2014) Leporids as a potential resource for predators (hominins, mammalian carnivores, raptors): an example of mixed contribution from level III of
Teixoneres Cave (MIS 3, Barcelona, Spain). CR Paleol 13:665–680. https://doi.org/10.1016/j.crpa.2014.06.001

Rufà A, Blasco R, Rivals F, Rosell J (2016) Who eats whom? Taphonomic analysis of the avian record from the Middle Palaeolithic site of Teixoneres Cave (Moia, Barcelona, Spain). Quat Int 421:103–115. https://doi.org/10.1016/j.quaint.2015.06.055

Rufí I, Solés A, Soler J, Soler N (2018) Un diente de cría de mamut (Mammuthus primigenius Blumenbach 1799, Proboscidea) precedente del Musterien de la Cueva de la Arbedna (Serinyà, NE de la Península Ibérica). Estud Geològics 74:e079. https://doi.org/10.3989/egel.43110.478

Salomoland M, Butet A, Leroux A, Bretagnolle V (2000) Responses of an avian predator to variations in prey density at a temperature latitude. Ecology 81:2428–2441

Sánchez-Hernández C, Gourichon L, Blasco R, et al (2020a) High-resolution Neanderthal settlements in mediterranean Iberian Peninsula: a matter of altitude? Quat Sci Rev 247. https://doi.org/10.1016/j.quascirev.2020.106523

Sánchez-Hernández C, Gourichon L, Soler J et al (2020b) Dietary traits of ungulates in northeastern Iberian Peninsula: did these Neanderthal preys show adaptive behaviour to local habitats during the Middle Palaeolithic? Quat Int 557:47–62. https://doi.org/10.1016/j.quaint.2020.01.008

Sánchez-Hernández C, Rivals F, Blasco R, Rosell J (2014) Short, but repeated Neanderthal visits to Teixoneres Cave (MIS 3, Barcelona, Spain): a combined analysis of tooth microwear patterns and seasonality. J Archaeol Sci 49:317–325. https://doi.org/10.1016/j.jas.2014.06.002

Sánchez-Hernández C, Rivals F, Blasco R, Rosell J (2016) Tale of two timescales: combining tooth wear methods with different temporal resolutions to detect seasonality of Palaeolithic hominin occupations of Teixoneres Cave (Moia, Barcelona, Spain). Radiocarbon 58:247–256. https://doi.org/10.1017/RDC.2015.19

Terry RC (2007) Inferring predator identity from skeletal damage of small-mammal prey remains. Evol Ecol Res 9:199–219

Tissoux H, Falguères C, Bahain J et al (2006) Dation par les séries de l’uranium des occupations moustériennes de la grotte des Teixoneres (Moia, province de Barcelone, Espagne). Quaternaire 17:27–33

IUCN, 2018. The IUCN Red List of Threatened Species [WWW Document]. Version 2018-2. URL www.iucnredlist.org

Wolff EW, Chappellaz J, Blunier T et al (2010) Millennial-scale variability during the last glacial: the ice core record. Quat Sci Rev 29:2828–2838. https://doi.org/10.1016/j.quascirev.2009.10.013

Wood RE, Arrizabalaga A, Camps M et al (2014) The chronology of the earliest Upper Palaeolithic in northern Iberia: new insights from L’Arbreda, Labeko Koba and La Viña. J Hum Evol 69:91–109. https://doi.org/10.1016/j.jhevol.2013.12.017

Zilhão J (2006) Chronostratigraphy of the middle-to-upper palaeolithic transition in the Iberian Peninsula. Pyrenees 37:7–84

Zilhão J (2000) The Ebro frontier: a model for the late extinction of Iberian Neanderthals. In: Stringer CB, Barton RNE, Finlayson C (eds) Neanderthals on the edge:150th anniversary conference of the Forbes’ Quarry discovery, Gibraltar. Oxbow Books, Oxford, pp 111–121

Zilio L, Hammond H, Karampaglidis T et al (2021) Examining Neanderthal and carnivore occupations of Teixoneres Cave (Moia, Barcelona, Spain) using archaeostratigraphic and intra-site spatial analysis. Sci Rep 11:1–20. https://doi.org/10.1038/s41598-021-83741-9

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