



Integrative approaches to the study of animal management practices during the Neolithic of South Iberian Peninsula: the case of El Toro cave (Antequera, Málaga, Spain)

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Abstract

The introduction and adoption of livestock played a pivotal role in shaping subsistence strategies of populations in the southern Iberian Peninsula during the Neolithic. However, there is lack of information regarding animal management strategies, such as grazing areas and changes in foddering strategies, and their correlation with the environmental characteristics, type of site and use of the settlement. The study of feeding strategies of domesticates provides a crucial information about the interaction between the management of the environment, husbandry systems and the exploitation of animal products. In this study, we use the archaeozoological data and the $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ stable isotope composition of the faunal bone collagen to understand herding systems and management strategies during the Neolithic in Phases IV and IIIB at El Toro cave (Antequera, Málaga). Archaeozoological and isotopic results revealed diverse husbandry practices and feeding strategies in El Toro cave during the Neolithic. The variability in $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values suggests the access of domesticates to different grazing areas and foddering strategies. This study contributes new insights into husbandry practices during the Neolithic and opens new perspectives for analysing animal management in mountain areas.

Keywords Archaeozoology · $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ stable isotopes · Husbandry practices · Foddering strategies · Bulk collagen

Introduction

The introduction of livestock during the early stages of the Neolithic in the Iberian Peninsula brought about significant changes in the subsistence strategies of populations. Currently, available archaeological data suggest a high degree of variability in animal management practices during the Neolithic in the Iberian Peninsula (Altuna and Mariezkurrena 2009; Aura et al. 2005; Castaños 2004; Martín-Socas et al. 2018; Morales and Riquelme 2004; Navarrete and Saña 2017; Navarrete et al. 2017; 2019; 2023; Pérez-Ripoll 2013, 2016; Saña 2013; Saña et al. 2020; Valente 2016). These management strategies include factors such as herd composition, size and production goals (i.e. meat, wool, milk, traction), often involving some level of human control over animal diet to ensure adaptation to local environmental conditions. In this sense, the study of feeding strategies can provide insight into the relationship between environmental management, husbandry systems and the exploitation of animal products.

Archaeological evidence of Neolithic occupation in the south and southeast of the Iberian Peninsula dates back to

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the second half of the sixth millennium cal BC, suggesting complex and diversified socio-economic dynamics (Aura et al. 2005; Camalich and Martín-Socas 2013; Martín-Socas et al. 1998, 2018; Morales and Martín 1995; Riquelme 1998; Salvatierra 1994). Different management and exploitation strategies are evidence depending on the region, diversity of biogeographic environments, site type (cave or open-air) and function. However, available archaeozoological data in the south of the Iberian Peninsula point to a successful ecological adaptation of domesticates (Martín-Socas et al. 2018). The dichotomy between wild and domestic species during the Neolithic is evident. During the early Neolithic, domestic animals were predominant in caves and open-air sites (e.g. Los Castillejos and La Molaina, Riquelme 1998; El Toro cave, Watson et al. 2004; Nerja cave, Morales and Martín 1995; El Retamar, Cáceres 2002) although high percentages of wild species are documented in some cave sites (e.g. Cueva Chica de Santiago, Pellicer and Acosta 1982; Acosta 1995). Among domesticates recovered from early Neolithic sites, domestic caprines (*Ovis aries* and *Capra hircus*) have a higher quantitative representation than other domestic species (*Bos taurus* and *Sus domesticus*), with sheep being more abundant than goats. In certain cave sites, such as El Toro cave, it is suggested that domesticates and people cohabited (Égüez et al. 2016; Martín-Socas et al. 2004). From the Late Neolithic, the high predominance of domestic species, with a significant increase in cattle and pig remains is documented (e.g. La Dehesilla, Acosta 1987; Acosta and Pellicer 1990; El Toro cave, Watson et al. 2004; Los Castillejos, Riquelme 1998). However, sites such as Cueva del Nacimiento and Valdecuevas still exhibited a significant presence of wild fauna (Morales and Riquelme 2004).

The success in integrating domesticates into economic practices in the south of the Iberian Peninsula during the early Neolithic is evident. Slaughter pattern results suggest a primary focus on obtaining meat, although secondary product exploitation is proposed from the early Neolithic (Acosta and Pellicer 1990; Riquelme 1998; Watson et al. 2004). Evidence also points to complementarity between agricultural and livestock practices (Buxó 1997; Peña-Chocarro et al. 2013; Rovira 2007; Zapata et al. 2005). However, there is a lack of information regarding animal management strategies during the Neolithic, such as grazing areas or changes in feeding and foddering strategies and their correlation with environmental characteristics of the area, site type and settlement use. For example, in the Antequera region, Neolithic sites primarily documented in caves (Martín-Socas et al. 2004, 2018; Perdigüero 1981; Sanchidrián and García 1987, with one exception being the open-air site Piedras Blancas I: García and Wheatley 2009), were associated with seasonal occupations of highly mobile communities (Martín-Socas et al. 2004, 2018). In this context, studies conducted at El Toro cave (Antequera, Málaga) suggested the synchronous

cave use for domestic activities and caprines stabling during the Neolithic (Égüez et al. 2016; Martín-Socas et al. 2004). The sequence of the cave's stratigraphy, featuring two well-defined Neolithic occupations, allows us to study changes in animal feeding strategies across different periods within the Neolithic.

Stable isotope analysis can be used to study aspects of pastoral management strategies, such as grazing and foddering strategies. In general, there is a consensus that the variable contribution of animal and plant macronutrients to animal diet can potentially reflect husbandry practices in the past (Balasse et al. 2015; Fuller et al. 2012; Halley and Rosvold 2014; Madgwick et al. 2012; Hamilton and Thomas 2012; Hammond and O'Connor 2013; Minagawa et al. 2005; Müldner and Richards 2005; Pechenkina et al. 2005). Previous studies have demonstrated the potential of bulk collagen carbon ($^{13}\text{C}/^{12}\text{C}$) and nitrogen ($^{15}\text{N}/^{14}\text{N}$) stable isotopes to explore different animal management practices in various contexts, for example, reconstructing strategies used in pig management during the prehistory (Balasse et al. 2015, 2016; Madgwick et al. 2013; Navarrete et al. 2017), characterizing livestock strategies in mountain environments (Villalba-Mouco et al. 2018; Navarrete et al. 2023) or inferring feeding strategies of the herds (Madgwick et al. 2023; Makarewicz 2023; Zavodny et al. 2014). Nevertheless, this approach has not yet been applied to explain husbandry practices in the south Iberian Peninsula during the Neolithic.

In this study, we use the archaeozoological data and the $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ stable isotopes data of the faunal remains to characterize herding systems and management strategies in El Toro cave (Antequera, Málaga) during the Neolithic. These will also contribute to evaluating the availability of herd maintenance areas and the implication of livestock strategies in feeding herds.

Materials

The archaeological site of El Toro cave (Antequera, Málaga)

El Toro cave is located in the Sierra del Torcal (Antequera, Málaga) at 1190 m a.s.l. (Fig. 1). Sierra del Torcal is a wide karstic mountain range of calcareous rocks and diastrophic systems that separates Mediterranean Andalusia from the Sub-Baetic System in the southern Iberian Peninsula. The stratigraphic sequence of El Toro cave has been associated with four chrono-cultural phases, of which the lower two correspond to the Neolithic occupations: Phase IV and Phase III (Santana et al. 2019) (Fig. 2).

During Phase IV (Early Neolithic, 6200–5980 BP: 5280–4780 2σ cal BC), the composition and spatial

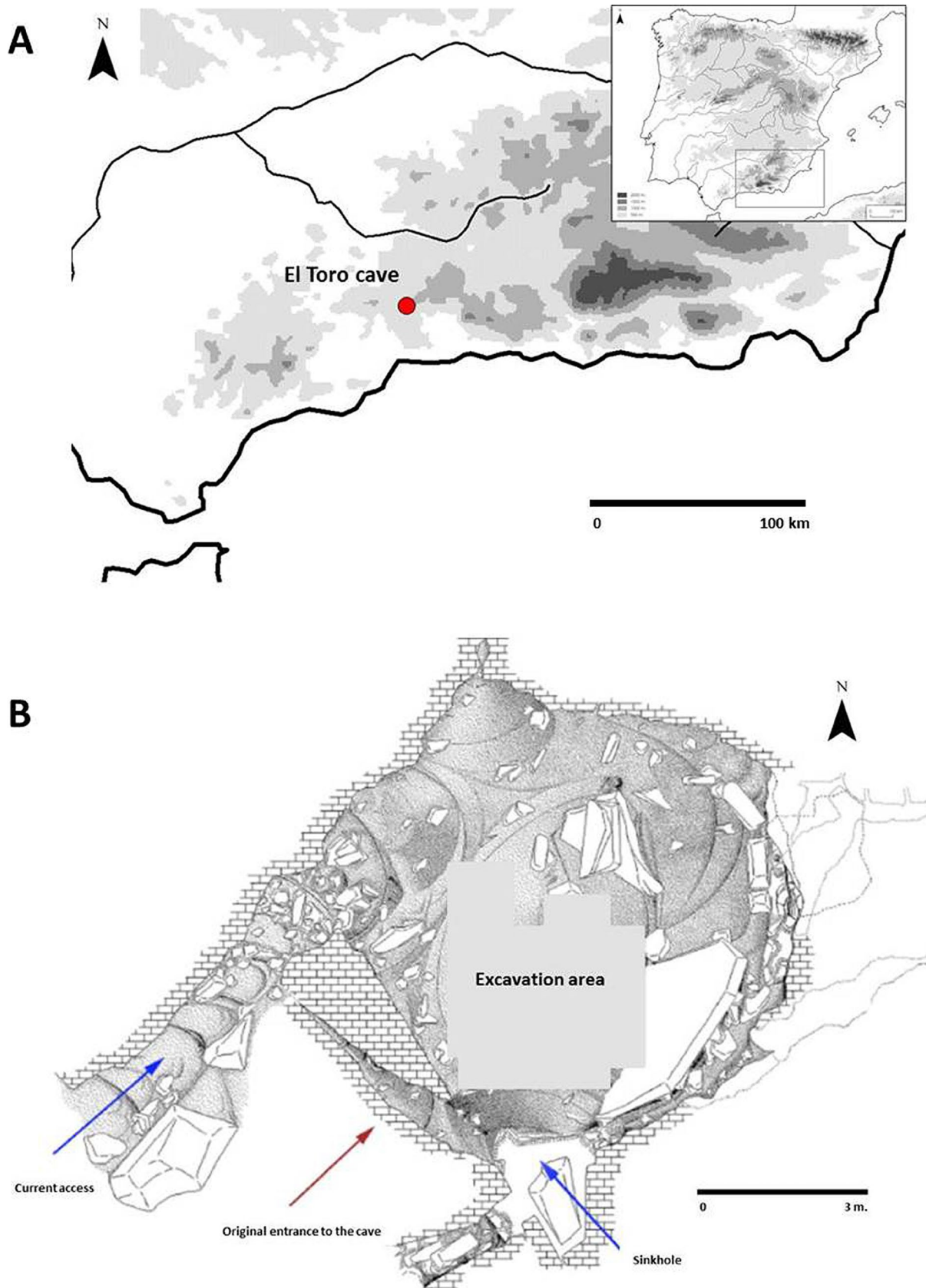


Fig. 1 A Location of El Toro cave (Antequera, Málaga). B General topography of the site of El Toro cave (Antequera, Málaga)

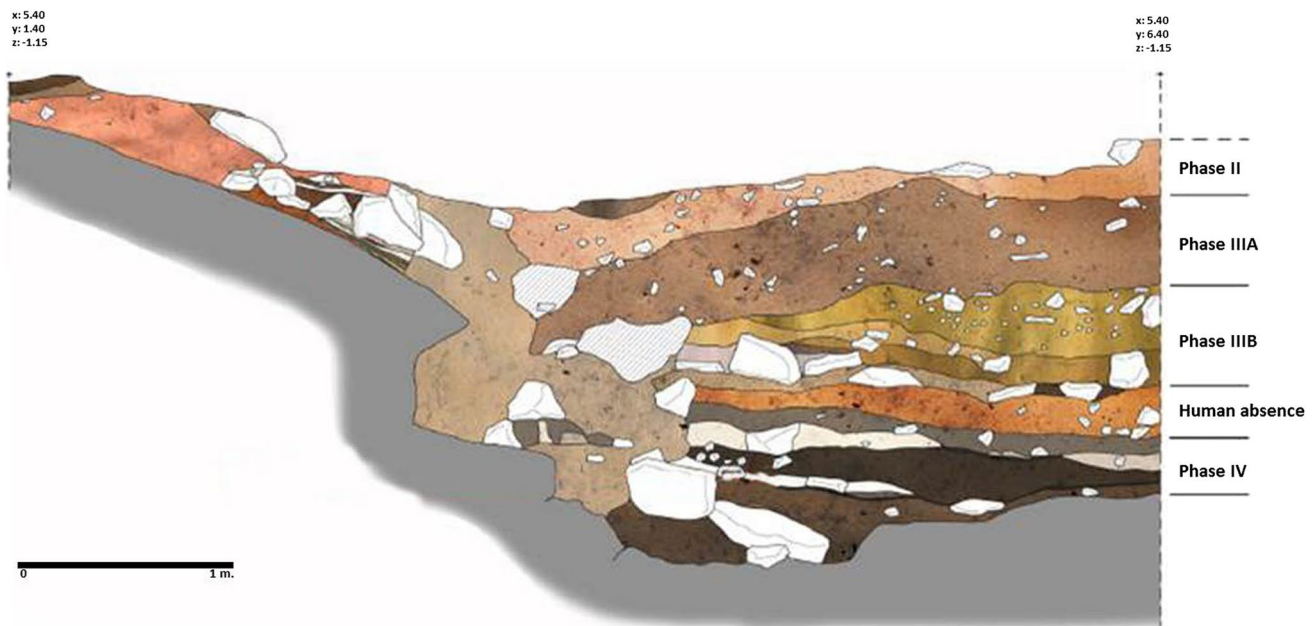


Fig. 2 Stratigraphic layers (phases) identified in the site of El Toro cave (Antequera, Málaga)

distribution of archaeological remains indicate domestic activities associated with plant and meat processing. Additionally, there is evidence of craft activities such as animal skin treatment and ceramic manufacturing (Martín-Socas et al. 2004; Rodríguez-Rodríguez et al. 1996, 2004). Camalich and Martín-Socas (2013) and Martín-Socas et al. (2004, 2018) interpret Phase IV as a seasonal and periodic occupation. This hypothesis is reinforced by the presence of lithic remains reaching the cave as a finished product (Rodríguez-Rodríguez 2004) and the presence of marine mollusks (Cuenca-Solana et al. 2021). Concerning animal resource management, the predominance of domestic animals over wild is observed. Among domesticates, caprines are predominant, specifically sheep over goats, followed by pigs and cattle (Watson et al. 2004). Micromorphological analysis has identified coprolites from caprines, suggesting that sheep and goats were stabled in the cave (Égüez et al. 2016). The archaeobotanical results record the predominance of wild plants like acorn and wild olives compared to domestic resources like naked wheat (*Triticum aestivum*), hulled barley (*Hordeum distichum*), lentils (*Lentil* sp.), broad beans (*Vicia faba*) and grass pea (*Lathyrus sativus*) (Buxó 1997).

Phase III (Late Neolithic, 5320–5170 BP: 4250–3950 2σ cal BC) is subdivided into phases IIIB (base) and IIIA (top) according to the differences in the intensity and spatial distribution of the archaeological remains. At the end of Phase IIIB, structural changes in the cave and the sealing of the main entrance caused by a collapse of the karst system are observed. Micromorphological analysis suggests the synchronous use of the cave as a domestic space and a stable (Égüez et al.

2016). The interior part of the cave would be used for stabling livestock, while the area closer to the entrance was used for domestic and craft activities (Égüez et al. 2016).

Radiocarbon data indicate a more stable and continuous occupation than in Phase IV, which is supported by micromorphological studies (Égüez et al. 2016; Martín-Socas et al. 2004). Domestic cereals such as naked wheat (*Triticum aestivum*), hulled barley (*Hordeum distichum*), barley (*Hordeum vulgare*) and legumes such as lentils (*Lentil* sp.) and broad beans (*Vicia faba*) become more representative during this period (Buxó 1997). Regarding livestock, Watson et al. (2004) observed an increase in butchery marks on faunal remains, suggesting a higher level of use and exploitation of animal carcasses. They also noted changes in the slaughter patterns of domestic caprines and could indicate changes in the production and exploitation practices of sheep and goats.

Between both chrono-cultural phases, a sterile deposit was identified and interpreted as a period of site abandonment. Based on dates obtained on short-lived samples, this sterile deposit is estimated to have formed in around 600 years (Martín-Socas et al. 2018).

The selection of faunal remains from Phase IV (early Neolithic) and Phase III (late Neolithic) of El Toro cave

Selection of faunal remains

For this work, the macrofaunal remains of Phase IV and Phase IIIB were reviewed and analysed. A total of 10924

faunal remains were analysed. Considering the main objective of this study is to characterize husbandry strategies, this paper focuses only on the number of identified specimens for Phase IV (NISP = 2132) and Phase IIIB (NISP = 1260) (Table 1).

Selection of faunal samples for isotopic analysis

A total of 82 samples for $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ isotopic analysis in bulk collagen were selected. A range of herbivores (Phase IV $n = 28$; Phase IIIB $n = 21$), including domestic (*Ovis aries*, *Capra hircus* and *Bos taurus*) and wild animals (*Cervus elaphus*, *Capra pyrenaica*, *Capreolus capreolus*, *Rupicapra rupicapra*), omnivores (Phase IV $n = 23$; Phase IIIB $n = 8$) (*Sus domesticus* and *Sus scrofa*) and carnivores (Phase IV $n = 1$; Phase IIIB $n = 1$) (*Vulpes vulpes*, *Canis familiaris*) were selected to establish the local $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ isotopic baselines. Samples comprised only adult specimens and included the diaphysis of bones, maxilla and mandibular diastema. Whenever possible, specimens were selected to represent adult individual animals by sampling the same-sided portion of a specific element.

Methods

Archaeozoological analysis

The bones were analysed and identified anatomically and taxonomically in the Museo de Málaga (Spain) using the

Table 1 Taxonomic representation of animal remains recovered in Phase IV and Phase IIIB of the El Toro cave (Antequera, Málaga) (NISP number of identified specimens; MNI minimum number of individuals)

	Phase IV		Phase IIIB	
	NISP	MNI	NISP	MNI
<i>Bos taurus</i>	69	3	48	3
<i>Ovis aries</i>	222	13	136	11
<i>Capra hircus</i>	146	9	105	10
<i>Ovis/Capra</i>	1195	8	661	13
<i>Sus domesticus</i>	380	19	210	8
<i>Canis familiaris</i>	0	0	1	1
<i>Cervus elaphus</i>	22	2	16	2
<i>Capreolus capreolus</i>	16	3	5	1
<i>Capra pyrenaica</i>	17	1	19	2
<i>Rupicapra rupicapra</i>	3	1	3	1
<i>Sus scrofa</i>	3	1	1	1
<i>Equus sp.</i>	0	0	1	1
<i>Oryctolagus cuniculus</i>	58	5	54	4
<i>Vulpes vulpes</i>	1	1	0	0

studies by Boessneck (1980); Fernandez (2001); Gillis et al. (2011); Halstead and Collins (2002); Payne (1985); Prummel and Frisch (1986); Salvagno and Albarella (2017); Zeder and Lapham (2010); and Zeder and Pilaar (2010) to distinguish *Rupicapra*, *Ovis*, *Capra* and *Capreolus*. The distinction between *Sus domesticus* and *Sus scrofa* was carried out on standardized criteria published by von den Driesch (1976) and Payne and Bull (1988). The analysis focused on the representation frequencies of different species and estimate of the age of death of the domesticates (*Ovis aries*, *Capra hircus*, *Sus domesticus* and *Bos taurus*). The number of identified specimens (NISP) and the minimum number of individuals (MNI) were used to quantify the assemblages. The MNI was calculated by considering the estimated age, anatomical representation and laterality. The relative frequency or presence/absence of taxa was used to estimate taxonomic variability. The age of the animals was estimated according to the stages of tooth wear (Grant 1982; Jones 2006; Lemoine et al. 2014; Payne 1985, 1987), eruption sequences and epiphyseal fusion in postcranial elements (Amorosi 1989; Barone 1976; Silver 1969; Zeder 2006).

Collagen extraction and stable isotope analysis

The extraction of collagen and analysis of stable isotopes were conducted at the Unitat d'Antropologia Biològica (Department of Animal Biology, Plant Biology and Ecology) and Laboratori d'Arqueozoologia (Department of Prehistory) at the Autonomous University of Barcelona (Spain). The bones were cleaned mechanically to remove the surface, and the extraction followed a modified Longin (1971) method (Brown et al. 1988). In short, shards of bones (ca. 200 to 300 mg) were demineralized using 0.6 M HCl at 4°C for several days, then rinsed with ultrapure water (Milli-Q®) and gelatinized with 0.001 M HCl at 80°C for 48 h. Samples were then ultrafiltered (30 kDa, Amicon® Ultra-4 centrifugal filter units; Millipore, MA, USA), frozen and freeze-dried. Collagen samples (0.3 mg) were analysed in duplicate using a Thermo Flash 1112 elemental analyser (EA) coupled to a Thermo Delta V Advantage isotope ratio mass spectrometer (IRMS) with a ConFlo III interface at the Institute of Environmental Science and Technology (ICTA) at the Autonomous University of Barcelona (Spain). Isotopic values are expressed as δ . The international laboratory standard IAEA 600 (caffeine) was used as a control. The average analytical error was $<0.2\text{‰}$ (1σ), as determined from the duplicate analyses of $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$. Comparison between $\delta^{13}\text{C}$ values of wild and domestic herbivores, omnivores and carnivores was performed using one-way ANOVA ($\alpha = 0.05$) after checking for normal distribution with the Shapiro–Wilk

test for normality ($\alpha = 0.05$). The null hypothesis that the data were normally distributed was rejected for the $\delta^{15}\text{N}$ values. Thus, the $\delta^{15}\text{N}$ values were compared using the Kruskal–Wallis test ($\alpha = 0.05$). All statistical tests were performed in PAST 4.11 (Hammer et al. 2001).

Results

Herd composition

A total of 2132 faunal remains from Phase IV and 1260 faunal remains from Phase IIIB were analysed. The remains of domesticated animals constituted more than 94.4 % (Phase IV, NISP = 2012) and 92.1 % (Phase IIIB, NISP = 1161) of the recovered assemblage, and wild species represented 5.6 % (Phase IV, NISP = 120) and 7.9 % (Phase IIIB, NISP = 99). Among domestic species, in Phase IV and Phase IIIB, caprines were more important quantitatively and in MNI than pig, followed by cattle (Table 1). The results of the comparison of the percentages of each domestic taxon in the Phase IV and Phase IIIB did not present fluctuations over the period studied.

Mortality profiles were calculated based on the MNI. Pigs and cattle exhibited a similar and homogeneous slaughter pattern across Phase IV and Phase IIIB. Pigs were primarily slaughtered between 14 and 21 months old (Phase IV, 43.5%; Phase IIIB, 57.5%). The number of slaughtered pigs declined after 21 months old (Phase IV, 33.1%; Phase IIIB, 34.4%), and only two individuals documented between 36 and 42 months old in Phase IV (19.5%). Less frequently, neonatal and infantile individual pigs up to 7 months old were documented (Phase IV, 3.9%; Phase IIIB, 8.1%). Regarding cattle, although the total recovered remains were limited, individuals were slaughtered at their meat-optimum (15–30 months old) (Phase IV, 66.7%; Phase IIIB, 66.7%). In both phases, an older individual of more than 48–72 months old was documented (Phase IV, 33.3%; Phase IIIB, 33.3%). The slaughtering patterns of caprines showed slight variations between phases. In both phases, caprine individuals slaughtered between 6 and 24 months old were predominant (Phase IV, *Ovis aries* 58.9% and *Capra hircus* 74.4%; Phase IIIB, *Ovis aries* 62.1% and *Capra hircus* 78%). However, in Phase IIIB, there was a significant increase in sheep individuals of 2–6 months old (24.4%). The presence of neonates younger than 2 months and infantile of 2–4 months old was also documented in *Ovis/ Capra* during Phase IV (49.8%) and Phase IIIB (40.7%). Distinguishing between sheep and goats in these immature specimens was not feasible due to the high degree of bone fragmentation and the absence of bone development in diagnostic elements.

Faunal stable isotopes $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values

Collagen preservation

The isotope data and collagen quality indicators are reported in Table 2 and Fig. 3. Out of a total of 82 specimens, collagen was successfully extracted from 69 (84.1%). Collagen yields ranged from 0.3 to 0.4 mg.

The C% and N% ranged from 22 to 45% and 11 to 16%, respectively, with C:N ratios ranging from 3.2 to 3.6 and falling within the values proposed by DeNiro (1985) and Van Klinken (1999) for preserved collagen. We also applied a 13% and 4.8% cut-off for C% and N%, respectively, as recommended by Ambrose (1990), Ambrose and Norr (1993).

$\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values

The average $\delta^{13}\text{C}$ values of herbivores ranged from -19.9 ± 1.2 ‰ (Phase IV; $n = 23$) to -20 ± 1.1 ‰ (Phase IIIB; $n = 18$). Although there were no statistically significant differences between phases ($p = 0.8345$), high intra-range variability was documented: -21.15 to -16.48 ‰ in Phase IV and -20.88 to -16.48 ‰ in Phase IIIB. The widest amplitude of variation of $\delta^{13}\text{C}$ values was recorded in domestic herds (4.85‰) compared to wild animals (1.57‰).

The average $\delta^{15}\text{N}$ values ranged from $+5.2 \pm 1$ ‰ (Phase IV; $n = 23$) to $+5 \pm 1.2$ ‰ (Phase IIIB; $n = 18$). These differences were statistically indistinguishable ($p = 0.554$). No significant differences were found between the $\delta^{13}\text{C}$ values of wild and domestic herbivores at Phase IV ($p = 0.8408$) and Phase IIIB ($p = 0.4115$). But there were significant statistical differences between $\delta^{15}\text{N}$ values of wild (average = 4.6 ± 1 ‰; $n = 10$) and domestic herbivores (average = 5.6 ± 0.7 ‰; $n = 13$) at Phase IV ($p < 0.009$). Among the domesticated, $\delta^{15}\text{N}$ values were found to be significantly different between cattle and sheep ($p < 0.01$) and between cattle and goats ($p < 0.03$). Cattle were significantly ^{15}N -enriched by 1.3‰ compared to the domestic herbivores ($p = 0.007$) and ^{15}N -enriched by 2.1‰ compared to the wild herbivores ($p = 0.019$). In Phase IIIB, there were also significant statistical differences in $\delta^{15}\text{N}$ values between domestic (average 5.4 ± 1.1 ‰; $n = 13$) and wild herbivores (average 4.05 ± 0.6 ‰; $n = 5$) ($p = 0.0274$). Domesticates were significantly ^{15}N -enriched by 1.35‰ compared to the wild herbivores. Among the domesticated, the only cattle had a $\delta^{15}\text{N}$ value of 6.89‰. Although there were no statistically significant differences between sheep and goats ($p = 0.1366$), the $\delta^{15}\text{N}$ values of sheep ranged from 7.04 ‰ to 3.95 ‰ ($n = 9$), while the $\delta^{15}\text{N}$ values of goats ranged from 4.52 to 4.31‰ ($n = 3$) (Figs. 3 and 4).

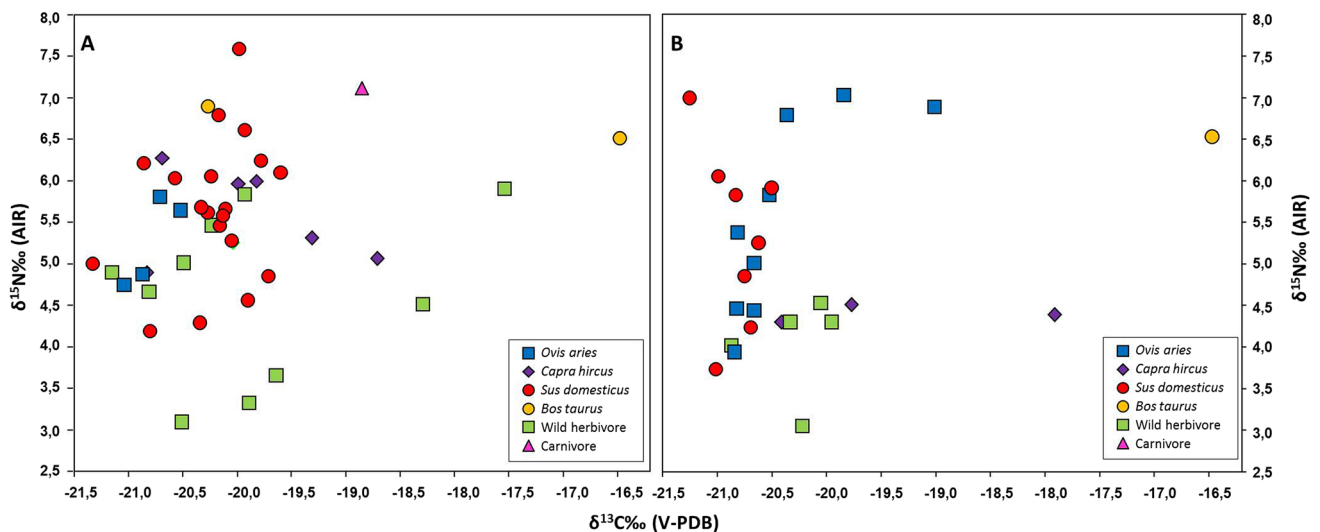
The average $\delta^{13}\text{C}$ values of pigs ranged from -20.2 ± 0.4 ‰ (Phase IV; $n = 19$) to -20.8 ± 0.2 ‰ (Phase IIIB; $n = 8$), and these differences were statistically significant ($p <$

Table 2 Results from carbon and nitrogen isotopic analysis of faunal samples from Phase IV and Phase IIIB of El Toro cave (Antequera, Málaga)

Site	Specimen ID	Phase	Species ID	Skeleton part	%C	%N	$\delta^{13}\text{C}(\text{‰})$	$\delta^{15}\text{N}(\text{‰})$	C:N
TCT	1	IV	<i>Sus domesticus</i>	Radius	41.69	15.14	-19.71	4.89	3.2
TCT	6	IV	<i>Sus domesticus</i>	Radius	44.95	16.00	-20.11	5.70	3.3
TCT	7	IV	<i>Sus domesticus</i>	Tibia	40.32	14.61	-20.16	5.50	3.2
TCT	8	IV	<i>Sus domesticus</i>	Mandible	32.02	11.53	-19.98	7.63	3.2
TCT	10	IV	<i>Sus domesticus</i>	Radius	42.39	15.31	-20.34	4.33	3.2
TCT	12	IV	<i>Sus domesticus</i>	Radius	44.83	16.30	-20.24	6.09	3.2
TCT	13	IV	<i>Sus domesticus</i>	Talus	44.02	15.41	-20.80	4.23	3.3
TCT	14	IV	<i>Sus domesticus</i>	Phalanx	44.87	16.19	-20.13	5.62	3.2
TCT	16	IV	<i>Sus domesticus</i>	Radius	42.37	15.14	-19.60	6.14	3.3
TCT	17	IV	<i>Sus domesticus</i>	Mandible	44.98	16.14	-20.27	5.66	3.2
TCT	18	IV	<i>Sus domesticus</i>	Mandible	39.07	14.09	-19.93	6.65	3.2
TCT	19	IV	<i>Sus domesticus</i>	Mandible	42.04	14.63	-20.86	6.25	3.4
TCT	20	IV	<i>Sus domesticus</i>	Metapodial	41.49	13.58	-20.57	6.07	3.6
TCT	21	IV	<i>Sus domesticus</i>	Phalanx	33.89	11.81	-21.33	5.04	3.3
TCT	37	IV	<i>Sus domesticus</i>	Phalanx	44.87	16.36	-20.33	5.72	3.2
TCT	48	IV	<i>Sus domesticus</i>	Radius	38.24	13.94	-19.90	4.60	3.2
TCT	44	IV	<i>Sus domesticus</i>	Humerus	41.88	15.06	-19.78	6.28	3.2
TCT	57	IV	<i>Sus domesticus</i>	Phalanx	44.60	15.66	-20.05	5.32	3.3
TCT	62	IV	<i>Sus domesticus</i>	Talus	44.23	15.98	-20.17	6.83	3.2
TCT	26	IV	<i>Ovis aries</i>	Humerus	42.91	15.68	-20.71	5.80	3.2
TCT	45	IV	<i>Ovis aries</i>	Humerus	40.96	14.86	-20.87	4.87	3.2
TCT	50	IV	<i>Ovis aries</i>	Metacarpus	41.06	14.99	-21.04	4.74	3.2
TCT	55	IV	<i>Ovis aries</i>	Humerus	42.81	15.62	-20.52	5.64	3.2
TCT	25	IV	<i>Capra hircus</i>	Humerus	42.95	15.52	-18.71	5.06	3.2
TCT	27	IV	<i>Capra hircus</i>	Radius	42.28	15.38	-20.04	5.25	3.2
TCT	28	IV	<i>Capra hircus</i>	Radius	37.29	13.65	-20.69	6.27	3.2
TCT	43	IV	<i>Capra hircus</i>	Radius	40.40	14.67	-19.82	5.99	3.2
TCT	49	IV	<i>Capra hircus</i>	Radius	40.68	14.87	-19.31	5.31	3.2
TCT	51	IV	<i>Capra hircus</i>	Humerus	45.03	16.25	-19.99	5.96	3.2
TCT	56	IV	<i>Capra hircus</i>	Humerus	43.53	15.83	-20.83	4.89	3.2
TCT	32	IV	<i>Bos taurus</i>	Phalanx	42.89	16.05	-20.34	6.89	3.4
TCT	33	IV	<i>Bos taurus</i>	Phalanx	44.13	16.16	-16.48	6.51	3.2
TCT	30	IV	<i>Cervus elaphus</i>	Radius	44.51	16.11	-20.34	5.90	3.2
TCT	31	IV	<i>Cervus elaphus</i>	Radius	42.68	15.12	-20.51	3.09	3.3
TCT	60	IV	<i>Capreolus capreolus</i>	Humerus	42.74	15.41	-20.23	5.46	3.2
TCT	61	IV	<i>Capreolus capreolus</i>	Humerus	45.49	16.50	-21.15	4.89	3.2
TCT	22	IV	<i>Capreolus capreolus</i>	Humerus	43.49	15.87	-18.29	4.51	3.2
TCT	35	IV	<i>Capreolus capreolus</i>	Radius	42.22	15.34	-19.93	5.83	3.2
TCT	36	IV	<i>Capra pyrenaica</i>	Metacarpus	42.88	15.54	-20.81	4.66	3.2
TCT	40	IV	<i>Capra pyrenaica</i>	Tibia	42.32	15.40	-20.49	5.01	3.2
TCT	59	IV	<i>Capra pyrenaica</i>	Metacarpus	41.04	14.79	-19.64	3.65	3.2
TCT	54	IV	<i>Capra pyrenaica</i>	Ulna	37.36	13.68	-19.89	3.32	3.2
TCT	39	IV	<i>Vulpes vulpes</i>	Scapula	42.58	15.62	-18.85	7.11	3.2
TCT	65	IIIB	<i>Sus domesticus</i>	Maxilla	40.34	14.98	-20.63	5.30	3.2
TCT	66	IIIB	<i>Sus domesticus</i>	Maxilla	41.19	15.02	-20.70	4.28	3.2
TCT	78	IIIB	<i>Sus domesticus</i>	Radius	41.89	15.26	-21.26	7.04	3.2
TCT	83	IIIB	<i>Sus domesticus</i>	Radius	42.39	15.36	-20.51	5.96	3.2
TCT	91	IIIB	<i>Sus domesticus</i>	Radius	42.98	15.39	-20.84	5.87	3.2
TCT	92	IIIB	<i>Sus domesticus</i>	Radius	44.23	16.20	-20.76	4.90	3.2

Table 2 (continued)

Site	Specimen ID	Phase	Species ID	Skeleton part	%C	%N	$\delta^{13}\text{C}(\text{‰})$	$\delta^{15}\text{N}(\text{‰})$	C:N
TCT	101	IIIB	<i>Sus domesticus</i>	Radius	40.42	14.60	-21.02	3.78	3.2
TCT	102	IIIB	<i>Sus domesticus</i>	Maxilla	41.17	14.88	-21.00	6.10	3.2
TCT	68	IIIB	<i>Ovis aries</i>	Radius	38.44	13.94	-20.83	4.47	3.2
TCT	71	IIIB	<i>Ovis aries</i>	Humerus	42.08	15.21	-20.67	4.45	3.2
TCT	73	IIIB	<i>Ovis aries</i>	Metacarpus	42.57	15.53	-19.02	6.90	3.2
TCT	74	IIIB	<i>Ovis aries</i>	Humerus	43.35	15.89	-20.37	6.80	3.2
TCT	75	IIIB	<i>Ovis aries</i>	Humerus	43.12	15.70	-20.82	5.39	3.2
TCT	79	IIIB	<i>Ovis aries</i>	Humerus	38.87	14.20	-20.53	5.84	3.2
TCT	80	IIIB	<i>Ovis aries</i>	Radius	41.46	14.98	-20.67	5.02	3.2
TCT	81	IIIB	<i>Ovis aries</i>	Metacarpus	43.33	15.60	-19.85	7.04	3.2
TCT	99	IIIB	<i>Ovis aries</i>	Radius	44.98	16.40	-20.85	3.95	3.2
TCT	93	IIIB	<i>Capra hircus</i>	Ulna	44.56	16.18	-17.92	4.40	3.2
TCT	72	IIIB	<i>Capra hircus</i>	Ulna	42.98	15.62	-19.78	4.52	3.2
TCT	77	IIIB	<i>Capra hircus</i>	Ulna	40.99	14.91	-20.42	4.31	3.2
TCT	84	IIIB	<i>Bos taurus</i>	Phalanx	39.33	14.42	-16.48	6.54	3.2
TCT	94	IIIB	<i>Cervus elaphus</i>	Tibia	44.90	16.36	-19.96	4.31	3.2
TCT	96	IIIB	<i>Cervus elaphus</i>	Tibia	44.36	16.08	-20.34	4.31	3.2
TCT	103	IIIB	<i>Cervus elaphus</i>	Metacarpus	44.27	16.10	-20.23	3.06	3.2
TCT	69	IIIB	<i>Capra pyrenaica</i>	Radius	41.71	15.11	-20.06	4.54	3.2
TCT	82	IIIB	<i>Capra pyrenaica</i>	Radius	43.26	15.68	-20.88	4.03	3.2

**Fig. 3** Plot of bone collagen $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values of faunal remains from El Toro cave (Antequera, Málaga). **A** Phase IV, **B** Phase IIIIB

0.0009). The average $\delta^{15}\text{N}$ values ranged from $+5.7 \pm 0.9 \text{‰}$ (Phase IV; $n = 19$) to $+5.4 \pm 1.1 \text{‰}$ (Phase IIIIB; $n = 8$) and were statistically indistinguishable ($p = 0.435$). In Phase IV, there were no statistically significant differences between pig and wild and domestic herbivores' $\delta^{13}\text{C}$ values (wild herbivore, $p = 0.202$; domestic herbivore, $p = 0.377$). The pig $\delta^{15}\text{N}$ values were enriched by 0.01‰ compared to the wild herbivores ($p = 0.005$). Between pig and domestic herbivores,

the statistical results of $\delta^{15}\text{N}$ values were indistinguishable ($p = 0.772$). In Phase IIIIB, $\delta^{13}\text{C}$ values between pigs and wild herbivores were statistically different ($p = 0.025$). The $\delta^{15}\text{N}$ values of pigs were significantly higher by an average of 1.3‰ compared to wild herbivores ($p = 0.025$). In contrast, there were no statistically significant differences in $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values between pigs and domestic herbivores ($p = 0.890$ and $p = 0.925$, respectively) (Figs. 3 and 4).

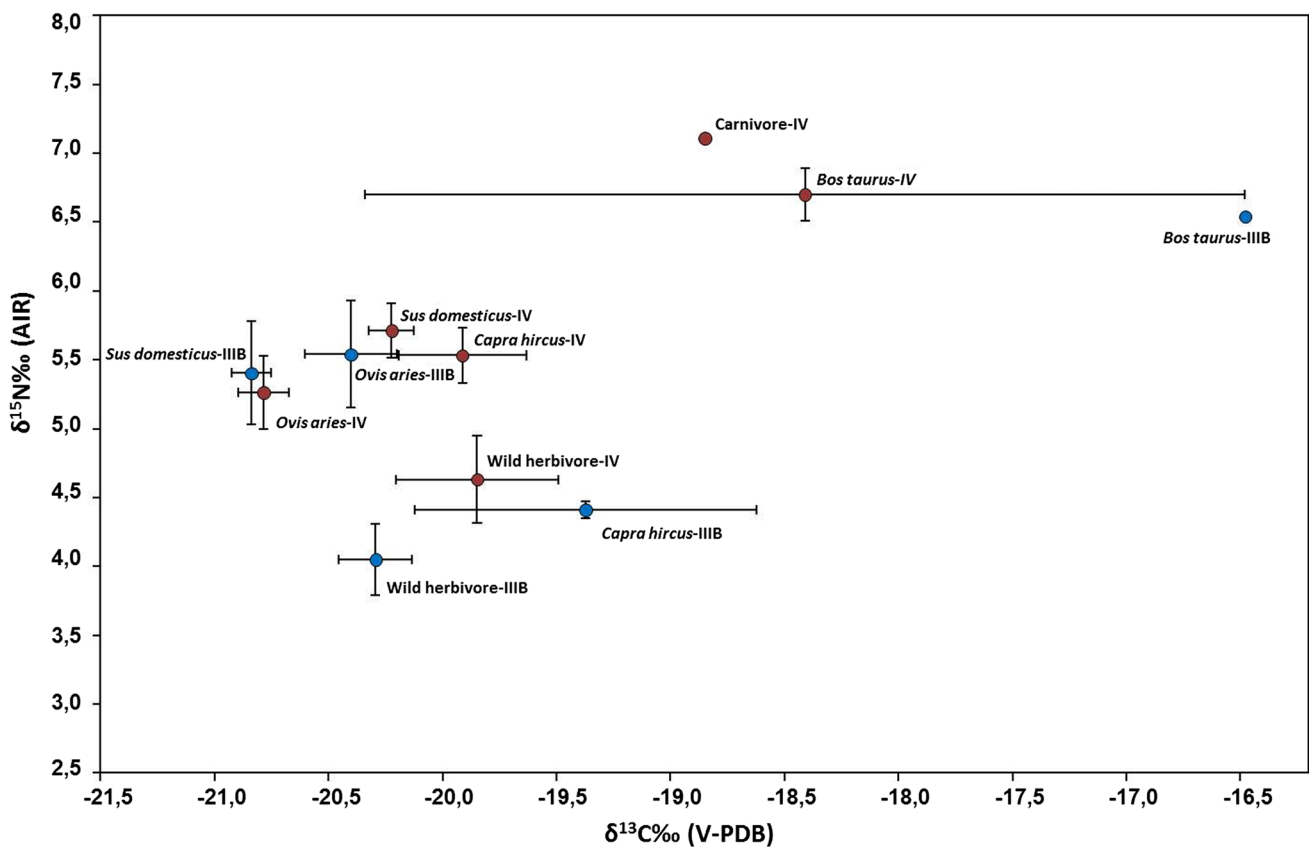


Fig. 4 Plot of mean $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values ($\pm 1\sigma$ s.d.) of faunal remains from Phase IIIIB (blue) and Phase IV (red) of El Toro cave (Antequera, Málaga)

Finally, carnivores were only represented by one individual (*V. vulpes*) from Phase IV. The $\delta^{13}\text{C}$ value was -18.85‰ , and the $\delta^{15}\text{N}$ value was 7.11‰ .

Discussion

Cattle and domestic caprine management strategies

The $\delta^{13}\text{C}$ values of the specimens analysed in El Toro cave ranged between -21.15 and -16.48‰ , reflecting the dominance of C_3 carbon in their diets. As expected, the herbivore $\delta^{13}\text{C}$ values suggest a terrestrial environment dominated by open C_3 plants (DeNiro and Epstein 1978; O'Leary 1988). According to a carbon isotope fractionation of $\sim 5\text{‰}$ between whole plants and consumer's bone collagen (Ambrose and Norr 1993) and a correction $\sim +1.5\text{‰}$ due to the fossil fuel effect when applied to pre-industrial ecosystems (Friedli et al. 1986; Hellevang and Aagaard 2015), $\delta^{13}\text{C}$ values close or higher than -19‰ in herbivorous species could result from the consumption of drought-resistant or dry vegetation, such as shrubs (Filella and Peñuelas 2003). In contrast, plants grown in closed environments produce lower

$\delta^{13}\text{C}$ values (O'Leary 1981; Tieszen 1991). During the early Holocene, the deciduous mesophilic trees such as birch (*Betula pendula*) and oak (*Quercus robur*), with more elements of the xerothermophilic maquis and pinewoods coastward, dominated the landscape of the Antequera region (López-Sáez et al. 2011).

The variation of $\delta^{13}\text{C}$ values $\delta^{15}\text{N}$ values observed between cattle and domestic caprines serves as an indication that these animals fed on plant resources with diverse isotopic composition. Specifically, cattle exhibit a $\delta^{13}\text{C}$ range of 4.67‰ and a ^{15}N -enriched by 2.3‰ compared to the wild herbivores. Wild herbivores are generally less restricted in their ecological niche. In Phase IV, cattle have a wider range of $\delta^{13}\text{C}$ values (-21.1 to -16.48‰) than in Phase IIIIB (-16.48‰). However, the amplitude of variation of $\delta^{15}\text{N}$ values between phases is minimal (0.16‰). The specimens of Phase IV (ID=33) and Phase IIIIB (ID= 84) form a distinct isotopic group (Figs. 3 and 4). Two hypotheses can be considered. Firstly, the low representation percentages and MNI of cattle (Phase IV, NISP = 69; MNI = 3; and Phase IIIIB, NISP = 48; MNI = 3) compared to the rest of the domestic species suggest that cattle would not be a frequently consumed resource in the cave. The narrow

$\delta^{13}\text{C}$ range of wild herbivores could indicate that these cattle could be tentatively associated with non-local animals, raised elsewhere and brought later to the site. Martín-Socas et al. (1999, 2004) suggest that El Toro cave might have been part of a regional trading network. The authors support this hypothesis by pointing to the recovery of malacofauna, ichthyofauna, raw materials and plant remains that were not sourced from the immediate area of the cave. For example, the recovery of ichthyofauna or malacofauna suggests frequent movements between the mountainous area and the coastal zone, which is approximately 35 km south in a straight line (Martín-Socas et al. 2004). Moreover, shrubs and branches collected over a wide geographical area, around the cave to the basal areas of the mountains, were documented inside the cave (Buxó 1997; Lopez-Saez et al. 2011; Martín-Socas et al. 2004; Rodríguez-Rodríguez et al. 1996). For example, *Pistacia* sp. and heather (*Calluna vulgaris*) require decalcified soils or mastic tree that requires areas sheltered from the cold or warmer areas than those existing in the Sierra del Torcal (Martín-Socas et al. 2004).

Secondly, the cattle may have had access to nearby pastures, and the change in $\delta^{13}\text{C}$ values could be a consequence of the changing environmental conditions. The first signs of anthropization in Sierra de El Torcal are detectable from the second half of the sixth millennium cal. BC and are associated, in many cases, with deforestation caused by fire (Lopez-Saez et al. 2011). In the fifth millennium cal. BC (Phase IIIB), livestock practices significantly transformed the landscape of Sierra de El Torcal, resulting in vegetation degradation and the proliferation of understory and scrub species (Martín-Socas et al. 2018; Rodríguez-Ariza 1996, 2004). Similar transformations have been observed in Cueva de Los Murciélagos (Córdoba) and Polideportivo de Martos (Jaén) (Lizcano 1999; Rodríguez-Rodríguez et al. 1996). Conversely, ^{15}N -enriched in animals may indicate consumption of ^{15}N -enriched plants fertilized with manure (Bogaard et al. 2007; Yousfi et al. 2010; Makarewicz 2014). This ^{15}N -enriched may suggest the consumption of agricultural by-products that were fertilized with manure, a phenomenon documented in various Neolithic contexts across Europe (Bogaard et al. 2013; Fraser et al. 2011). The high values of $\delta^{15}\text{N}$ could indicate cattle accessing cultivated fields. However, the impact of animal dung on plant $\delta^{15}\text{N}$ values is highly variable and depends on factors such as manuring frequency and type of fertilizer (Fraser et al. 2011; Szpak 2014). In El Toro cave, cereals such as naked wheat, naked barley and dressed barley and legumes such as lentils, broad beans and pebbles have been documented (Buxó 1997). However, the number of cattle samples from Phase IV (NISP=2) and Phase IIIB (NISP=1) is too small to lead to conclusive interpretations.

Domestic caprines are the most common taxon in El Toro cave during Phase IV (*Ovis/Capra* NISP = 1195;

sheep NISP = 222; goat NISP = 146) and Phase IIIB (*Ovis/Capra* NISP = 661; sheep NISP = 136; goat NISP = 105), with sheep slightly outnumbering goats. Slaughtering patterns of caprines in Phase IV and Phase IIIB suggest a primary focus on meat production, evidenced by animals being killed between 6 and 24 months old. Still, a significant percentage of animals killed before 6 months old. Such early slaughtering of animals is commonly associated with either milk production for human consumption or the acquisition of high-quality tender meat (Oueslati 2006). Furthermore, in El Toro cave, evidence of dairy consumption has been provided (Tarifa-Mateo et al. 2019).

As already pointed out by Watson et al. (2004), the archaeozoological and microstratigraphic analysis of the site confirmed that caprine penning inside the cave during Phase IV and Phase IIIB (Égüez et al. 2016). The intensity of the cave's occupation during Phase IV is evidenced by abundant microscopic fragments of charred seeds and ceramics and burning events commonly found in caprine penning contexts (Égüez et al. 2016).

The isotopic analysis of domestic caprines shows a wide range in $\delta^{13}\text{C}$ values, particularly in sheep (Phase IV = 3.18‰; Phase IIIB = 1.83‰). Goats show differentiated ranges between phases (Phase IV = 0.17‰; Phase IIIB = 2.50‰). A wide range in $\delta^{15}\text{N}$ values is also documented. The ^{15}N -enriched values of sheep (0.63‰) and goats (0.36‰) were lower than those documented for sheep in Phase IIIB (1.49‰). Conversely, goats maintained a consistently low value in Phase IIIB (0.9‰). These wide ranges in $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ may result from various factors, including different grazing areas, herd movement, foddering strategies or management strategies of stabling areas. If there were seasonal herd movements, they might not be discernible from the analysis of carbon and nitrogen stable isotopes in bone collagen. Three hypotheses are considered. Firstly, the access of flocks to diverse pastures enriched in ^{15}N by manure might explain the variability in $\delta^{15}\text{N}$ values observed between Phase IV and Phase IIIB. During Phase IV, wild plants constituted 62%, and cultivated plants only constituted 39% of the human diet (Buxó 1997). Phase IIIB represents a systematic occupation of the cave (Martín-Socas et al. 2004), with an intensification of agriculture and an increase in the importance of cultivated species (cereals as wheat, hulled barley, barley and legumes such as lentils and broad beans) versus wild species (acorns) in the cave (Buxó 1997). Differences in isotopic values between sheep and goats could be conditioned by their distinct dietary preferences and needs (Vaiglova et al. 2014), as well as documented grazing habits during early Neolithic at La Draga site (Navarrete et al. 2019). The recorded values for El Toro cave are lower than those documented in the European context (Bogaard et al. 2013). However, it aligns with the values

recorded for the northeast of the Iberian Peninsula (Navarrete et al. 2017, 2023).

Secondly, variability $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values could be due to foddering strategies. $\delta^{15}\text{N}$ values could reflect the consumption of agricultural by-products such as crop stubble and other plants used as fodder. Moreover, the increase in ^{15}N is not homogenous in plants, especially with manured cereal straws potentially being depleted in ^{15}N compared to grains and legumes (Bogaard et al. 2007, 2013). The documented presence of collected bushes and branches such as *Pistacia* sp. and heather inside the cave (Buxó 1997) suggests that these resources might have complemented the diet of herds. Additionally, evidence of domestic activities within the cave, including processing and preparation of cereals and legumes for human consumption (Égüez et al. 2016; Martín-Socas et al. 2004) supports the hypothesis that the agricultural by-products were provided to the flocks. The neonate and infantile individuals recovered in the El Toro cave (30% of total of domestic caprines) suggest that animal foddering practices likely played a crucial role in maintaining the nutritional health of sheep mothers, vital for the survival of the lambs (Mellor and Stafford 2004).

Thirdly, the ^{15}N -enriched values could result from confined feeding environment, such as a stable. This enrichment may occur due to the use of fodder and the presence of manure. Livestock manure is typically ^{15}N -enriched compared to fodder (Bogaard et al. 2007; Commisso and Nelson 2006). Continuous use of the area where animals are housed can potentially enrich the soil with ^{15}N (Commisso and Nelson 2006). In this sense, micromorphological analyses at El Toro cave indicate caprine stabling practices inside the cave (Égüez et al. 2016).

Pig management strategies

Pigs are likely omnivores and opportunistic feeders (Macdonald and Barrett 1993; Schley and Roper 2003). As omnivores, in the composition of collagen pig bone, nitrogen isotopes can come from both animal and plant proteins. Traditional pig husbandry may have involved home-based systems with complete to partial stabling near settlements or extensive herd management in semi-free to free-range regimes. This practice has been observed in modern traditional communities in northern Mediterranean (Albarella et al. 2007; Hadjikoumis 2012) and is also documented for prehistoric groups in Europe (Balasse et al. 2016) and in Northeast and Central Pyrenees of the Iberian Peninsula (Navarrete et al. 2017; Villalba-Mouco et al. 2018).

At El Toro cave, the amplitude of variation for $\delta^{13}\text{C}$ values of pigs is similar to that of wild herbivores (1.73‰), and their absolute $\delta^{13}\text{C}$ values generally overlap with the herbivore data. There are no significant difference in average $\delta^{15}\text{N}$ values between phases (Phase IV, 4.63‰; Phase IIIB, 4.05‰), and the ^{15}N -enriched values (Phase IV, 3.26‰;

Phase IIIB, 3.40‰) do not comprise the variation range of strict and wild herbivores at the site (2.8‰) (Figs. 3 and 4).

The diversity in ^{15}N -enriched values indicates fluctuations in local livestock management practices (Navarrete et al. 2017). This suggests the pig diet may have mainly consisted of plant products, animal protein and agricultural by-products (cereals, pulses). Two management strategies align with these conditions. One involves household-level regimes or enclosed pigs. These pigs would mainly feed on plants (ground cereals and legumes), potentially supplemented with domestic leftovers, including animal products. This could result in higher $\delta^{15}\text{N}$ values in pigs than local herbivores, though also depending on the quality and proportion of protein in their diets (Balasse et al. 2016; Madgwick et al. 2012; Müldner and Richards 2005). The high $\delta^{15}\text{N}$ values (~ 6.5–7.6‰) in some pig specimens could indicate some degree of manuring effect, which could then transfer to pig collagen through consuming animal products and waste. High values of $\delta^{15}\text{N}$ are also documented in pigs from Can Sadurní cave and Reina Amàlia-Caserna de Sant Pau sites (northeast Iberian Peninsula) during the Neolithic (Navarrete et al. 2017).

Second, pigs reared in semi-free or free-range systems would obtain most of their nutrients from available plants, although this does not exclude some consumption of small animals or insects and worms, as well as human food waste (Hamilton and Thomas 2012). Within this management strategy, we might expect pig collagen $\delta^{15}\text{N}$ values to be similar or close to local herbivores (Madgwick et al. 2012, 2013). Similar interpretations have been proposed for prehistoric pigs possibly raised at the household-level with a mostly herbivorous diet (Balasse et al. 2016) and for modern pigs raised in traditional farming communities (Hadjikoumis 2012). Plant-based diets have been documented in the northeast Iberian Peninsula in sites of El Frare cave, La Draga and Serra de Mas Bonet (Navarrete et al. 2017) and in Central Pyrenees (Villalba-Mouco et al. 2018).

The pigs in El Toro cave present high variability in $\delta^{15}\text{N}$ values, but this is not directly related to size. While there is considerable variability in pig size in the Iberian Peninsula (Navarrete and Saña 2017), the pigs from El Toro cave have a size associated with the domestic form according to the standardized criteria published by von den Driesch (1976) and Payne and Bull (1988). In addition, mortality profiles that these animals were mostly slaughtered between 14 and 21 months old and between 36 and 42 months old, when their reproductive capacity begins to decline. This mortality pattern has been documented in extensive pig-rearing livestock models of ethnographic societies (Hadjikoumis 2012). In extensive regimes, pigs are slaughtered at around 12 and 24 months old, while in intensive regimes, they are slaughtered younger, a consequence of faster fattening due to their strictly controlled diet (Hadjikoumis 2012). Even

though the cave was seasonally occupied, during Phase IV and more intensively in Phase IIIB, and caprine penning inside the cave has been documented, there is no evidence of this practice in the case of pigs (Égüez et al. 2016). In this sense, the variability of isotopic data and archaeozoological data could indicate several feeding strategies for pigs.

Conclusion

Husbandry practices played an essential role in the economic strategies of El Toro cave during the Neolithic. Domestic species show a high percentage of representation compared to wild species. Among domesticates, caprines were most common, followed by pigs and cattle. Slaughtering patterns suggest meat was the focus of animal exploitation during Phase IV and Phase IIIB, especially for pigs and cattle. For caprines, meat-producing specimens dominated during Phase IV and Phase IIIB, although a significant increase in neonatal and infantile remains was recorded in Phase IIIB. This could suggest caprine birth in the cave, and the occurrence of slaughtering in the first 2 months is often associated with milk production (Vigne and Helmer 2007). Phase IIIB coincided with an intensification of agricultural practices with the synchronic use of the cave of humans and animals cohabitation (Martín-Socas et al. 1999, 2004).

Isotopic results from domesticates exhibit an extensive range of $\delta^{15}\text{N}$ and $\delta^{13}\text{C}$ values and provide evidence for diverse husbandry strategies and foddering regimes in El Toro cave during the Neolithic. The wide-ranging herbivore $\delta^{15}\text{N}$ values from Phase IV and Phase IIIB suggest grazing on vegetation with a variety of $\delta^{15}\text{N}$ values, potentially reflecting different manuring degrees on the fields. The isotopic heterogeneity in cattle suggests that the individuals are likely to come from disparate areas rather than just the immediate locality. Pigs were subjected to various foddering regimes, with high $\delta^{15}\text{N}$ that may indicate a diet with animal protein and/or agricultural by-products. In addition, the age of killed pigs suggest a possible extensive regime. Homogeneity in their $\delta^{13}\text{C}$, similar to wild herbivores, may indicate their management at the local level.

In sum, this study provides new insights into Neolithic husbandry practices and presents new perspectives for analysing animal foddering strategies in mountain areas. In the future, the integration of these results with analyses of oxygen and strontium isotopes will help us understand the nature of mobility and the provenance of domesticates.

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Data availability All relevant data generated or analysed for this study are included in this published article.

Code availability Not applicable

Declarations

Ethics approval Not applicable

Consent to participate Not applicable

Consent for publication Not applicable

Competing interests The authors declare no competing interests.

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