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### 1 Neanderthal and Carnivore Occupations in Unit II from the Upper Pleistocene Site of 2 Ramandils Cave, (Port-la-Nouvelle, Aude, France).

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### 8 ABSTRACT

9 The northwestern Mediterranean coast includes varied environment with abundant faunal and lithic resources exploited 10 during the Upper Pleistocene. In this area, the Ramandils Cave (Port-la-Nouvelle, Aude, France) records Middle Palaeolithic 11 levels dating to Marine Isotope Stage (MIS) 5. Evidence of Neanderthal occupations are observed throughout the sequence; 12 with numerous lithic industries, burnt faunal remains, marine shells, and few human bones. In unit II, the richest one, lithic 13 exploitation strategies are linked to the availability of major areas of flint supply implying a defined mobility pattern. Several 14 archaeostratigraphic levels have been individualised, with ungulate remains affected by human and carnivore predation, 15 involving alternated occupations of the cave during different periods. By confronting studies on origin and management of 16 lithic raw material with taphonomical and zooarchaeological data of large mammals, we attempt to gain a better 17 understanding of the site function, occupation duration, and regional mobility pattern in unit II of Ramandils Cave.

18 Keywords: Neanderthal occupations, Upper Pleistocene, large mammals, lithic industry, mobility pattern, site function

#### 19 **1. Introduction**

20 The northwestern Mediterranean coast records several Middle Palaeolithic sites, correlated 21 between Marine Isotopic Stage (MIS) 5 to 3, with recurrent Neanderthal occupations, such as 22 Ramandils cave, Crouzade cave, Tournal cave, and the rockshelter of Les Canalettes 23 (Bischoff et al., 1988; Cochard et al, 2012; Daujeard et al, in Djindjian et al, 2018; Magniez, 2010; Rusch et al., 2019; Saos, 2003; Saos et al, 2016, 2017; Valladas & Joron, 1993; 24 25 Yokoyama et al., 1987a). Faunal associations reflect climatic conditions between these 26 transition periods as well as human and carnivore behaviours, frequently recorded in the same 27 assemblages during Upper Pleistocene (Brugal et al., 2017; Costamagno et al., 2005; 28 Daujeard et al, 2016, Discamps et al., 2012; Magniez, 2010; Rendu, 2007; Saladié et al., 2018, among many others). This involves representation, with different intensities and 29 30 frequencies, of zooarchaeological and taphonomic markers on ungulate carcasses, implying 31 various subsistence behaviors of these two types of predators. Occupation and Mobility 32 patterns are accessible through regional studies (Daujeard & Moncel, 2010; Daujeard et al, 33 2016; Grégoire, 2012; Lebègue et al, 2010; Patou-Mathis, 2000; Szmidt, 2003; Valensi & 34 Psathi, 2004; among many others), by taking into consideration raw material resources available in the territory, as well as faunal remains and lithic tools analyses found in sites. 35 36 Three major occupation modes are discerned in this chronological period, long-term residential camps with predominantly anthropogenic accumulation, short-term regular camps 37 38 that can be occupied alternately with carnivore occupations, and brief stopover as bivouacs, 39 such as highlighted in South of France around Mediterranean coast and in the Rhône Valley 40 (Daujeard & Moncel, 2010; Daujeard et al., 2016; Magniez, 2010; Saos et al., 2016).

41 The coastal context of Ramandils cave (Port-la-Nouvelle, Aude, France) with the 42 proximity of various ecosystem and major areas of flint supply point out this site as a key 43 sequence for mobility studies during upper Pleistocene. Last excavations (Boutié et al, 1998) 44 have yielded several occupation levels with both Neanderthal and carnivore activities, mainly 45 represented by numerous lithic tools, burnt bones, marine shells, and coprolite deposits. 46 Recent studies on large mammal remains, palynological analyses of coprolites and new 47 radiometric dating were conducted (Rusch et al., 2019), and allow to correlate the sequence to 48 the end MIS 5. The results highlighted temperate climate with forest, open (grasslands), and 49 rocky landscapes. The unit II, the richest one of Ramandils Cave, is chosen here for the study 50 of the successives occupations on this strategic place. Archaeostratigraphic approach in 51 progress permitted to individualise several occupation levels in this unit, with evidence of 52 human and carnivore activities. Thereby, thanks to archaeozoological parameters, using 53 mortality patterns, seasonality by large mammal spectrum, anatomical distribution, and 54 taphonomic traces studies, in addition with lithic raw material supply modalities and lithic 55 production technological processes, we will attempt to interpret subsistence strategies and 56 economical scheme of the lithic production. These feature combinations will allow us to gain 57 a better understanding of mobility patterns and site function.

### 58 **2. Site presentation**

Ramandils Cave (Port-la-Nouvelle, Aude department, southern France), is located 30 km south of Narbonne (fig. 1a), altitude 1.5 m, and separated from the current Mediterranean shoreline by 1,400 m of lagoon deposits (fig. 1b). This east-facing cavity, towards the sea, is formed at the bottom of an Urgo-Aptian limestone hill, called Cap Romarin, east of the first

63 Corbières massif foothill. This cave nowadays has a collapsed ceiling on the most part.



64

65 66 Fig. 1: Ramandils Cave locations : (a) raised-relief map with upper Pleistocene sites and (b) aerial photography (modified according www.geoportail.gouv.fr)

67 The site was discovered in 1925 by Théophile Héléna (Grégoire, 2011). Planned yearly 68 excavations were directed by Paul Boutié between 1983 and 1994 (Boutié *et al*, 1998), on 69 more than 50 m<sup>2</sup>, according to systematic methods and based on a grid reference to record 70 Cartesian data (fig. 2).





Fig. 2: (a) Grid reference with excavated areas in gray, each one-meter squared area is distinguished by a letter (E to P) correlated with a number (11 to 24). (b) East view of the cave (modified according to Boutié, 1990 and Rusch et al, in press, photo credit: P. Boutié, Centre Européen de Recherches Préhistoriques, Tautavel).

75 The cave, more than 5 m high, was completely filled with sediment, partially eroded in the lower part by an active natural spring (fig. 3). The stratigraphic sequence firstly records, at the 76 bottom, a marine level with marine shells dated by Electron Spin Resonance (ESR) method to 77 78 128 ±15 ky (Yokoyama et al., 1987) to the last Thyrrhenian transgression. Secondly, 79 continental deposits are individualised (Kabiri, 1993) into five stratigraphic units (numbered I 80 to V), containing among others, equids teeth, dated by ESR/U-Th (Uranium-Thorium) 81 method between 94  $\pm$  7 ky (unit V) and 77  $\pm$  6 ky (unit I) to the end of Marine Isotope Stage (MIS) 5 (Rusch et al., 2019). 82





Mousterian occupations are recorded for all continental levels, mostly represented by lithic industries (29,994 pieces) corresponding to a Typical Mousterian (Molès, 2008). The peculiarity of this lithic collection is that all pieces have very small dimensions. The unit II levels have 15,832 lithic pieces, of which 68.4% are less than 15 mm and thus generally uncoordinated. This is the richest stratigraphic unit of the site.

89 This collection contains 0.7% of cores, 20.9% of flakes, 5.9% of blades. The production modes are mainly Levallois and SSDA (39.7% each). Among the Levallois systems, the 90 91 Lineal (52.7%) is the most represented followed by the orthogonal recurrent modality (24.6%) 92 and the recurrent centripetal ones (20.6%). The cores on flakes (kombewa production) 93 represents 6.9% and testify to a secondary chaine opératoire on the initial productions already 94 of small dimensions (tab. 1) as a clear ramified flint exploitation strategy (Bourguignon et al., 95 2004). Discoids are rare (3.4%). The remainings 10, 3 % are indeterminate knapping systems. 96 The reduction sequence on flint is totally represented into the site from the first steps of 97 preparing cores, through the production of blanks by various methods (Levallois dominant, 98 followed by SSDA and discoide methods) until the retouching process to produce tools on flakes or blades and ramification of a part of the Levallois production, resulting in a 99 100 microlithisation of the production. The larger flakes were selected to produce flake tools. Tool 101 percentage is low 4.8%. Flake tools are mainly side-scrapers (48%), followed by denticulates 102 (20 %) and notches (13 %). A large part of the flake production was not retouched on tools.

|                      | PRINCIPAL<br>RAW<br>MATERIAL   | ORIGIN  | MATRIX                               | INPUT<br>MODALITY                 | SECONDARY RAW<br>MATERIAL         | ORIGIN                 | MATRIX                      | INPUT<br>MODALITY  |
|----------------------|--|---|--------------------------------------|-----------------------------------|-----------------------------------|------------------------|-----------------------------|--------------------|
| Ramandils<br>Unit II | flint  | semi-local  | blocks, slabs                        | raw blocks,<br>prepared<br>blocks | quartzite,<br>chert,<br>limestone | semi local             | pebble                      | whole              |
|                      | REDUCTION<br>SEQUENCE  | DEFICIT<br>PRODUCT                                    | PRODUCTION<br>GOAL                   | SECONDARY<br>METHOD               | RAMIFICATION                      | IMPORTANCE<br>OF TOOLS | DOMINANT<br>TOOL            | ASSOCIATED<br>TOOL |
|                      | complete<br>for flint<br>partial for<br>other raw                                      | retouched<br>tools on<br>flakes                       | flakes (micro)                       | SSDA                              | yes                               | low                    | scraper                     | denticulate        |
|                      | REDUCTION<br>SEQUENCE<br>complete<br>for flint<br>partial for<br>other raw<br>material | DEFICIT<br>PRODUCT<br>retouched<br>tools on<br>flakes | PRODUCTION<br>GOAL<br>flakes (micro) | SECONDARY<br>METHOD               | RAMIFICATION                      | IMPORTANCE<br>OF TOOLS | DOMINANT<br>TOOL<br>scraper | ASSO<br>TC<br>dent |

<sup>103</sup> Tab. 1: Main techno-economic characteristics of the unit II lithic assemblage (according to data from Moles, 2008).

Faunal remains (14,927 remains) are represented by large mammals, (Gerber, 1973; Banès 1996, 1998; Rusch *et al.*, 2019), birds (Vilette in Boutié, 1993), lagomorphs and microvertebrates (Percie-du-Sert, 1992) with terrestrial (André in Boutié, 1993) and marine shells (Cataliotti-Valdina in Boutié, 1990; Celiberti & Moigne in Saos *et al.*, 2016). In unit II, three fragmented human teeth were discovered (Chevalier & Bécam in Saos *et al.*, 2016), without specific attribution or burial context.

## 110 **3. Material and methods**

111 All of the material studied corresponds to coordinated objects that comes from Boutié's 112 excavations, recorded according to the grid reference and stratigraphic subdivision in unit II.

## 113 3.1. Archaeostratigraphy

The Ramandils Cave has delivered an important archaeological material consisting of, unburnt and burnt bones, lithic tools, stones, shells, coprolites, ... representing about 31,000 coordinated objects. The first step is to create a database from the excavation notebooks and to validate the data before adding the faunal and lithic remains. The second step is to realize vertical sections of the objects along the transverse and longitudinal axes of the cave. In Ramandils Cave, the archaeological remains are so numerous that profiles were made every 10 cm. It is then possible to begin the individualization of archaeological levels.

The profiles will be cross-referenced before highlighting the lower and upper limits of each occupation level. For the study of an excavation zone, of a square meter, it is thus proceeded with a hundred of cross-checks. Once, the limits of occupation levels are defined, each object is assigned a soil name (Pois, 1998, 1999). In unit II 6 levels are defined in unit II: S8, S9, S10, S11 and S12.

# 126 3.2. Study of lithic Raw Material and their Origin

127 The raw materials of the whole unit II were petrographically determined on a macroscopic 128 basis and classified by category of rocks. Flints were characterized by the description of their microfacies with binoculars and then close to the known petrographical facies of the 129 130 Languedoc-Roussillon repository of sedimentary siliceous rocks. This repository (Grégoire et 131 al., 2010), set up in the framework of a PCR program, available at the European Center for 132 Prehistoric Research at Tautavel was used as a reference for identifying sources. Following 133 this method,-categories of rocks were identified in unit II and have been linked to potential 134 sources (Boutié et al, 1998; Grégoire, 2000). From these identifications, a territory can be 135 drawn and hypotheses of supply strategies can be proposed in relation to the settlement 136 systems of territories by these nomadic peoples (Grégoire, 2012).

## 137 3.3. Zooarchaeology

Taxonomic and anatomical identifications were made from the reference collection of the 138 139 European Center for Prehistoric Research at Tautavel, following the relevant literature 140 (Barone, 1999; Pales & Garcia, 1981; Schmid, 1972; Varela & Rodriguez, 2004, among many 141 others). The Number of Remains (NR), the Number of Identified Specimens (NISP), and the 142 Minimum Number of Individuals (NMI) by combination were calculated (Grayson, 1984; 143 Klein & Cruz-Uribe, 1984; Lyman, 1994, 2008; Moigne, 1983; Poplin, 1981). The assessment 144 of post-depositional modifications (Lyman, 1994) was conducted for diagenetic processes 145 study (tab. 2) analysing steps of concretion, oxidation (manganese or iron impregnations), 146 cortical bone alteration, and cracking.

|               |                          |     |      |      | Unit II | levels | 5    |      |      |
|---------------|--------------------------|-----|------|------|---------|--------|------|------|------|
|               |                          | S   | 510  | S    | 11      | S      | 512  | То   | tal  |
|               | NR Total                 | 896 | 100% | 2233 | 100%    | 548    | 100% | 3677 | 100% |
| Legil         | ble bone total           | 819 | 91%  | 2039 | 91%     | 489    | 89%  | 3347 | 91%  |
| Illegi        | ble bone total           | 77  | 9%   | 196  | 9%      | 61     | 11%  | 334  | 9%   |
|               | Legible bones (C1)       | 224 | 25%  | 612  | 27%     | 152    | 28%  | 988  | 27%  |
| Concretion    | Illegible bones (C2 +C3) | 39  | 4%   | 101  | 5%      | 38     | 7%   | 178  | 5%   |
|               | Total                    | 263 | 29%  | 713  | 32%     | 190    | 35%  | 1166 | 32%  |
| Oxi           | <b>des</b> (Mn, Fe)      | 474 | 53%  | 739  | 33%     | 122    | 22%  | 1335 | 36%  |
| Cortical hono | Legible bones (D1 +D2)   | 620 | 69%  | 1355 | 61%     | 198    | 36%  | 2173 | 59%  |
| altoration    | Illegible bones (D3)     | 39  | 4%   | 100  | 4%      | 26     | 5%   | 165  | 4%   |
| alleration    | Total                    | 659 | 73%  | 1455 | 65%     | 224    | 41%  | 2338 | 63%  |
| Cracki        | ng (F1 +F2 +F3)          | 136 | 15%  | 263  | 12%     | 96     | 17%  | 495  | 13%  |

147

7 *Tab. 2:* Climatic and edaphic markers on bones remains (NR) from unit II levels.

Concretions are described with 3 steps. The first one (C1) is slight and isolated spots; the second one (C2) is a fine cover on the complete bone and the third one (C3) corresponds to bone being inside the concretion. Only remains with the first step have apparent cortical bone, accounting for around 27 % of NR. Remains with step 2 and 3, for between 4 and 7% of NR, have not legible surface and are therefore not eligible for study other taphonomic markers.

153 Oxide deposits, mainly manganese, are observed on more than a quarter of remain in S11 154 and S12, and on more than half in S10. This low pigmentation of remains reveals bone 155 surface and other taphonomic markers.

156 Cortical bone alteration due to water circulation is divided-with 3 steps. The first one (D1) 157 with thin dissolution cups and the second one (D2) with slight desquamation areas 158 corresponds to bones with legible surface, noticed for 36% of NR in S12 to 69% of NR in 159 S10. The third step, with desquamation affecting a large part of cortical surface is not eligible 160 for study biological markers, representing about 4% of NR.

161 Cracking is related to the fissuring process, also due to the water circulation. The first step 162 is just a thin hair-like fissure, the second one has pronounced splits but the size of bones is not 163 affected, and the third one has deep or severe cracking. All these steps reveal cortical bones 164 with legible surface for around 13% of total NR

165 Legible bones, accounting for 91% of NR, are sorted according step criteria of concretion166 and cortical bone alteration, that can be combined on the same bone fragment.

Material fragmentation was noticed according to size class, in millimetres, and nature of remains (fig. 4). In unit II, mainly pieces are less than 40 mm length, especially for bones remains, taken into account in the study of human and carnivore impacts. For the long bones of mains species, size class and the circumference of diaphysis fragments were noted as well as the distribution of partial remains, with complete epiphysis, and whole bones (Bunn, 1983; Villa and Mathieu, 1991).



173 174 175

Fig. 4: Material length according class size (mm) and nature of remains from unit II, for determinate (dark grey) and indeterminate (white) bones, teeth (light grey) and coprolites (black).

176 A biostratinomic alteration study was carried out on damage caused by roots, and rodents 177 (scraping) or carnivores tooth marks (pitting, furrowing, puncturing and scoring), with 178 digestion processes products by carnivores (Binfor, 1978, 1981; Blumshine, 1986; 179 Dominguez-Rodrigo & Piqueras, 2003; Fosse 1996; Hayne, 1980; Klein and Cruz-Uribe, 180 1984). A part of coprolites is attributed to cave hyena, following morphological description 181 (Jouy-Avantin et al., 1999, 2003; Vivent, 1989), during palynological and parasitological 182 analyses (Lartigot-Campin & Moné, 2017). Coprolites occurrence in unit II levels was used to 183 identify the zooarcheological sequence and carnivore activities in the site (Diedrich, 2012; 184 Discamps et al., 2012; Pineda, 2017).

Anthropic modifications of bones were studied (green fractures, cut marks, percussion marks, and technological applications of bones—essentially retouchers) according the relevant literature (Binford 1981; Daujeard *et al.*, 2018; Mallye *et al.*, 2012; Noe-Nygaard 1989; Potts & Shipman 1981; Shipman & Rose 1983; Villa & Mathieu, 1991) and a microscope was used. Burnt remains were recorded for colour and burning localisation (according Lebon, 2008).

Mortality profiles (Stiner, 1990) were based on age-at-death, established on eruption, replacement, and dental wear, or epiphysis fusion. Seasonality was determined from the current dates of ungulates birth (Azorit *et* al., 2002; Cornevin & Lesbre, 1894; Guadelli, 1998; Hayssen et al. 1993; Hillson, 1986; Magnell, 2007; Mariezkurrena, 1983; Pérez Ripoll, 1988). In this regard, we only considered those species in which the age-at-death is estimated in months corresponding to equids, bovids, cervids, and suids.

### 197 **4. Results**

198 4.1. Archaeostratigraphy

The study highlighted 26 occupation levels for the entire filling (fig. 5). In this article, we focus on stratigraphic unit II, the richest level which delivered almost 20,000 remains. Six occupation levels were defined in this unit, namely S8, S9, S10, S11, S12 and S13. However, the limits between units I and II and between units II and III still need to be clarified. The archaeological levels we are interested in for this study are mainly levels S10, S11 and S12.



 $\begin{array}{c} 204 \\ 205 \end{array}$ 

Fig. 6: Vertical section with S11 (grey), S12a (black) and S12 (white) in band 21. S12a corresponds to a coprolites bed. For an easier lecture, only thirty centimeters (between 0 and 30 cm) are represented for occupation levels S11 and S12

S10, S11, and S12 are the richest levels, they contain respectively 4,267, 7,893 and 3,696
remains (tab. 3). The major part of the archaeological material is distributed into bands 21 to
24 (fig. 7). It is also in these three levels that coprolites are concentrated. Coprolites are
spatially distributed in specific areas, especially in S12a.

| Unit | Occupation | Fauna | Coprolites | Shells | Lithic | Stones | Human | Others | Total |
|------|------------|-------|------------|--------|--------|--------|-------|--------|-------|
|      | level      |       |            |        | tools  |        | bones |        |       |
|      | S8         | 158   | 2          | 10     | 119    | 27     | 0     | 2      | 318   |
|      | S9         | 943   | 5          | 48     | 834    | 158    | 0     | 14     | 2002  |
|      | S10        | 1988  | 7          | 55     | 1977   | 122    | 0     | 23     | 4172  |
|      | S11        | 3795  | 39         | 99     | 3637   | 91     | 2     | 131    | 7794  |
| 11   | S12a       | 6     | 27         | 0      | 0      | 0      | 0     | 0      | 33    |
|      | S12        | 1772  | 43         | 44     | 1665   | 50     | 0     | 56     | 3630  |
|      | S13a       | 0     | 12         | 0      | 0      | 0      | 0     | 0      | 12    |
|      | S13        | 145   | 3          | 2      | 148    | 15     | 0     | 7      | 320   |
|      | Total      | 8807  | 138        | 258    | 8380   | 463    | 2     | 233    | 18731 |

213

Tab. 3: Total archaeological material coordinated from stratigraphic unit II-



### 214 215 216

Fig. 7: Spatial distribution of the archaeological material in occupation level S11. Most of the remains are concentrated in band 21 to band 24.

The archaeological approach allows a finer study of the filling than that carried out from the stratigraphic units (Pois, 2000). At the level of the unit II, it was, indeed, impossible to perceive these coprolites beds. However, it is interesting to see these coprolite beds at the limit of human occupations. It should also be noted the presence of coprolites within an anthropic level that may reflect a short abandonment of the site by humans.

### 4.2. Lithic Assemblages: Raw Material

The lithic collection from the unit II were studied by Moles (Molès, 2008) with a techno-223 224 typological approach and by Grégoire (Boutié et al., 1998; Grégoire, 2000) with a 225 petroarcheological point of view. In this article, we have chosen to use the contribution of the 226 lithic industries, especially raw materials to feed the multidisciplinary discussion about the function site and the mobility pattern of human groups. However, unlike fauna, the results 227 228 from previous studies were not analysed according to the archaeostratigraphic division 229 proposed here for unit II. A homogenization work is actually underway but general trends will 230 therefore be proposed and discussed in relation with the archaeozoological data, offering at 231 this stage a better study resolution.

Ten different raw materials were used in the stratigraphic unit II, the richest one (tab. 4). This is the level with the most lithological diversity. Flint is the most widely used rock with

234 91.5%. Other kinds of rocks are used like quartzite, chert, limestone, quartz, lydian,
235 sandstone, jasper, schist and hornfels.

| Unit      | II     |
|-----------|--------|
| Flint     | 91.5 % |
| Quartzite | 4.0 %  |
| Chert     | 1.9 %  |
| Limestone | 1.2 %  |
| Quartz    | 0.8 %  |
| Hornfels  | 0.2 %  |
| Jasper    | 0.1 %  |
| Sandstone | 0.1%   |
| Schist    | 0.1 %  |
| Lydian    | 0.1 %  |
| Total     | 100 %  |

236 237 238

The flints are generally very alterated with a white or beige and total thick patina. Burnt artefacts are abundant (39.6%) and make it difficult to identify flint categories.

Essentially represented by flint, siliceous rocks are largely in the majority (94%). Quartzite is the second most used rock after flint, followed by limestone and quartz. Non-siliceous rocks (6%) are represented by various raw materials, collected in the form of pebbles. These pebbles came from alluvial deposits, but also from the marine environment because of the proximity of the Mediterranean shore.

Thanks to some flint artefacts little altered or without any thermal impact, several types were recognized. All of them shows the main characteristics of the Miocene and more rarely the Oligocene regional facies, well known (Grégoire, 2010) on the Tertiary Narbonne-Sigean Basin around the site.

The study of the microfacies of these flints shows two main groups which correspond to both the stratigraphic upper Oligocene facies and the lower Miocene facies (Aquitanian).

The upper Oligocene facies is characterized by a mudstone texture slightly fossiliferous consisting of cryptocristallin micritic silica with hematites and frequent ferruginous impregnations as well as rare bioturbations. The Miocene flint shows a wackestone / packstone texture with charophytes (boiled stalks, oogones), gastropods and ostracods.

The sedimentary texture and the inclusions are reliable discriminating characters to distinguish the two stratigraphic facies and pinpoint their source in the current landscape within the Narbonne-Sigean basin (fig. 8).

Tab. 4: proportion of lithic raw material categories in Ramandils Cave unit II.

259 Concerning the most frequent flint, the poor quality of blanks used for debitage production, as well as the presence of charophytes remains visible at a macroscopic level, allows to 260 eliminate the exploitation of high quality oligocene flints provided by the outcrops of 261 262 Roquefort des Corbières (fig. 8, flint source n°4), Indeed, Neanderthals have rarely used this 263 rock which is attested in the site by only a few artefacts. The miocene flints (fig. 8, flint 264 sources  $n^{\circ}1$ , 2, 3, 5), the most used, probably come from mostly tectonized blocks of the 265 Presqu'île du Doul, Portel-des-Corbières or Hameau du Lac Miocene sources. Some cherts 266 coming from the Cretaceous Corbières formation, at the north-west of the site (fig. 8, flint 267 source n°6), and from the nearest Jurassic formation at the north of the tertiary basin (fig. 8, 268 flint source  $n^{\circ}7$ ), were also used in the unit II.



*Fig. 8:* Location of the lithic raw material and faunal sources on the local and semi-local territory around Ramandils Cave. 1 : Presqu'île du Doul (Miocene flint), 2: Portel des Corbières (Miocene flint), 3: Hameau du Lac (Miocene flint), 4: Roquefort des Corbières (Oligocene flint), 5 : Sigean (Miocene flint), 6 : Plan du Couloubret (chert); 7 : Hauts de Narbonne (chert), 8 : Aude alluvial deposits (quartz, quartzite, limestone, hornfels, lydian, sandstone, schist), 9 : Têt alluvial deposits (jasper).

## 4.3. Large Mammals Remains

Unit II of Ramandils Cave counts 4,058 NR of large mammals (without considering coprolites), with 91% distributed in levels S10, S11 and S12. Faunal study will allow to define the species spectrum and to characterise these three occupation levels.

# 278 *4.3.1. Species Composition*

The three levels count 3,682 total NR, corresponding to 653 NISP (tab. 5) with S11 as the richest level (2235 NR, 321 NISP). Faunal spectrum is mainly composed of ungulates (94% NISP), dominated by red deer, Cervus elaphus (33% total NISP), especially in S11 (36% NISP) and S12 (33% NISP), with horse, Equus f. germanicus (28% total NISP), prevalent in S10 (33% NISP). Moreover, European wild ass, Equus hydruntinus, aurochs, Bos primigenius, and wild boar, Sus scrofa, are well represented in these levels (each by 8% of total NISP).

Carnivores are generally rare (6% NISP), particularly represented by cave hyena, Crocuta c. spelaea (2 % NISP), cave lynx, Lynx spelaeus (1% NISP), fox, Vulpes vulpes (1% NISP), wolf, Canis lupus (0.6% NISP), and cave beer, Ursus spelaeus (0.6% NISP). Between S11 and S12, S12 a level is composed of carnivore remains (cave hyena, badger, Meles meles, and unspecified bear) with only one ungulate bone attributed to the horse.

At last, a significant number of indeterminate bones are recorded in S10, S11 and S12 mainly classify as small ungulates (2804 NR) according size and thickness criteria, that could indicate high bone fragmentation.

|                                  |     | S10   | S    | 11    | S | 512a | 9   | 512   | Тс   | otal  |
|----------------------------------|-----|-------|------|-------|---|------|-----|-------|------|-------|
| Total carnivores (NISP)          | 11  | 5.9%  | 18   | 5.6%  | 3 | 75%  | 8   | 5.6%  | 40   | 5.9%  |
| U. arctos                        | 1   | 0.5%  | 1    | 0.3%  | - | -    | -   | -     | 2    | 0.3%  |
| U. spelaeus                      | 1   | 0.5%  | 2    | 0.6%  | - | -    | 1   | 0.7%  | 4    | 0.6%  |
| M. meles                         | -   | -     | 1    | 0.3%  | 1 | 25%  | -   | -     | 2    | 0.3%  |
| Lutrinae indet.                  | 1   | 0.5%  | -    | -     | - | -    | 1   | 0.7%  | 2    | 0.3%  |
| C. lupus                         | -   | -     | 1    | 0.3%  | - | -    | 3   | 2.1%  | 4    | 0.6%  |
| V. vulpes                        | 1   | 0.5%  | 4    | 1.3%  | - | -    | 1   | 0.7%  | 6    | 0.9%  |
| C. c. spelaea                    | 3   | 1.6%  | 6    | 1.9%  | 2 | 50%  | 2   | 1.4%  | 13   | 2.0%  |
| L. spelaeus                      | 4   | 2.1%  | 2    | 0.6%  | - | -    | -   | -     | 6    | 0.9%  |
| F. silvestris                    | -   | -     | 1    | 0.3%  | - | -    | -   | -     | 1    | 0.2%  |
| Unspecified carnivores (NR)      | 2   | -     | 6    | -     | 1 | -    | -   | -     | 9    | -     |
| Total ungulates (NISP)           | 176 | 94.1% | 302  | 94.4% | 1 | 25%  | 135 | 94.4% | 613  | 93.9% |
| P. antiquus                      | -   | -     | 6    | 1.9%  | - | -    | -   | -     | 6    | 0.9%  |
| S. hemitoechus                   | 4   | 2.1%  | 2    | 0.6%  | - | -    | 1   | 0.7%  | 7    | 1.1%  |
| E. f. germanicus                 | 61  | 32.6% | 86   | 26.9% | 1 | 25%  | 38  | 26.8% | 186  | 28.5% |
| E. hydruntinus                   | 17  | 9.1%  | 24   | 7.5%  | - | -    | 12  | 8.5%  | 53   | 8.1%  |
| Bison sp.                        | -   | -     | 1    | 0.3%  | - | -    | -   | -     | 1    | 0.2%  |
| B. primigenius                   | 22  | 11.8% | 20   | 6.3%  | - | -    | 11  | 7.7%  | 53   | 8.1%  |
| C. caucasica                     | 1   | 0.5%  | 8    | 2.5%  | - | -    | 4   | 2.8%  | 13   | 2.0%  |
| C. elaphus                       | 52  | 27.8% | 116  | 36.3% | - | -    | 48  | 33.8% | 216  | 33.1% |
| D. dama                          | 3   | 1.6%  | 7    | 2.2%  | - | -    | 1   | 0.7%  | 11   | 1.7%  |
| R. tarandus                      | -   | -     | 2    | 0.6%  | - | -    | 1   | 0.7%  | 3    | 0.5%  |
| C. capreolus                     | 4   | 2.1%  | 5    | 1.6%  | - | -    | 2   | 1.4%  | 11   | 1.7%  |
| S. scrofa                        | 12  | 6.4%  | 25   | 7.8%  | - | -    | 16  | 11.3% | 53   | 8.1%  |
| Unspecified large ungulates (NR) | 36  | -     | 58   | -     | - | -    | 29  | -     | 123  | -     |
| Unspecified small ungulates (NR) | 646 | -     | 1808 | -     | - | -    | 350 | -     | 2804 | -     |
| Indeterminate large mammal (NR)  | 25  | -     | 41   | -     | - | -    | 27  | -     | 93   | -     |
| Total NISP                       | 187 | 100%  | 320  | 100%  | 4 | 100% | 142 | 100%  | 653  | 100%  |
| Total NR                         |     | 896   | 2233 |       |   | 5    |     | 548   | 3    | 682   |

Tab. 5: Large Mammals composition in unit II levels of Ramandils Cave, by NISP (black) and NR (grey).

### 295 *4.3.2. Human and Carnivore impacts*

Biological markers are noticed in the unit II levels, produced by various agents (tab. 6).

297 Human activities have the highest impact on more half remains, especially with burning 298 processes increasing from the base, 52% NR affected in S12, to the top, 79% NR affected in 299 S10. Butchery activities are stable all along the levels with cut marks, on 13 to 20% of 300 remains, and green fractures related to bone breakage, on 9 to 15% of remains, despite rare 301 percussion marks, on less than 3% of remains. At last few bone retouchers were found mainly 302 on long bone fragment of cervids and unspecified small ungulates. Finally, the red deer 303 remains are the most affected among all species by human impacts, while for the horse, 304 butchery marks are scarcer.

Damage caused by carnivore activities are clearly more reduced, with less than 4% of remains affected by tooth marks and rare digested bones. These markers are observed on main species with close values, constant for the horse and with a low increase for the red deer in S11.

309 Other biological impacts are recorded. Rodent impact is insignificant in unit II levels with

310 less than 1% of remains affected by scraping. Root marks increase from the base, with less

311 than 5% NR in S12, to the top, not exceeding 10.4 % NR, inducing plant cover on the cave

312 floor.

| F      |                |     | S1    | 0        |             |      | <b>S1</b> : | 1         |             |     | S1       | 2         |             |      | Tot   | al       |             |
|--------|----------------|-----|-------|----------|-------------|------|-------------|-----------|-------------|-----|----------|-----------|-------------|------|-------|----------|-------------|
|        |                | _   | -     | M<br>spe | ain<br>cies |      |             | Ma<br>Spe | ain<br>cies |     |          | Ma<br>spe | ain<br>cies | Tota | al NR | M<br>Spe | ain<br>cies |
|        |                | 101 |       |          | 1           | lota |             |           | 1           | lot | Total NR |           | 1           |      |       |          | M           |
|        | NR Total       | 896 | 100%  | 52       | 61          | 2235 | 100%        | 116       | 86          | 550 | 100%     | 48        | 39          | 3681 | 100%  | 216      | 186         |
| Le     | gible bones    | 819 | 91%   | 49       | 54          | 2039 | 91%         | 111       | 59          | 489 | 89%      | 47        | 31          | 3347 | 91%   | 207      | 154         |
|        | Burning        | 710 | 79.2% | 40       | 27          | 1706 | 76.3%       | 88        | 55          | 283 | 51.5%    | 39        | 18          | 2699 | 73.3% | 167      | 100         |
| n      | Cut marks      | 173 | 19.3% | 12       | 2           | 441  | 19.7%       | 36        | 4           | 70  | 12.7%    | 13        | 1           | 684  | 18.6% | 61       | 7           |
| uma    | Fractures      | 133 | 14.8% | 21       | 4           | 278  | 12.4%       | 28        | 5           | 50  | 9.1%     | 10        | 4           | 461  | 12.5% | 59       | 13          |
| Ī      | Percussion     | 14  | 1.6%  | 3        | -           | 34   | 1.5%        | 9         | -           | 13  | 2.4%     | 5         | -           | 61   | 1.7%  | 17       | -           |
|        | Retouchers     | 2   | 0.2%  | -        | -           | 3    | 0.1%        | 1         | -           | 3   | 0.5%     | 1         | -           | 8    | 0.2%  | 2        | -           |
| ivore  | Tooth<br>marks | 31  | 3.5%  | 2        | 4           | 80   | 3.6%        | 9         | 4           | 21  | 3.8%     | 2         | 4           | 132  | 3.6%  | 13       | 12          |
| Carr   | Digestion      | 9   | 1.0%  | 3        | 2           | 7    | 0.3%        | 1         | -           | 7   | 1.3%     | 2         | 1           | 23   | 0.6%  | 6        | 3           |
| Rodent | Scraping       | 3   | 0.3%  | -        | -           | 4    | 0.2%        | -         | -           | 3   | 0.5%     | 1         | -           | 10   | 0.3%  | 1        | -           |
| Plant  | Root marks     | 93  | 10.4% | 12       | 6           | 192  | 8.6%        | 21        | 6           | 27  | 4.9%     | 7         | 1           | 312  | 8.5%  | 40       | 13          |

Tab. 6: Biologic and anthropic markers on bones remains (NR) from unit II levels with main species representation (NISP)

### 314 - Carnivore Activities

315 Direct evidence of carnivore occupation are noticed in unit II levels by a low percentage of 316 carnivore remains, 6% NISP (tab. 5), with the cave hyena and the lynx relatively more 317 represented. Scarce marks on large mammal bones, less than 4% NISP in each level (tab. 6), 318 can be attributed to carnivore impact. Without considering unspecified ungulate remains 319 which have the highest values, traces are mainly recorded on main species, red deer and 320 horse. Most affected remains corresponding to burnt bones with butchery marks, such as 321 green fractures, indicating carnivore scavenging during human abandonment periods of the 322 cave. However, unburnt bones are affected (fig. 9), especially horse remains with whole long 323 bones, which can be transported by large predators such as cave hyena. Further studies such 324 as on the size of tooth marks will allow us to determine the specific agents affecting large 325 mammal bones (Dominguez-Rodrigo & Piqueras, 2003).





Fig. 9: Carnivore marks on unburnt ungulate remains: (a) epiphysis gnawing on a right radio-ulna of horse, (b) puncturing marks on a left talus of horse, (c) epiphysis gnawing on a second phalanx of horse, (d) epiphysis gnawing on a right humerus of ibex, (e) digested metatarsal of red deer (photos L. Rusch and D. Dainat, Centre Européen de Recherches Préhistoriques, Tautavel).

In unit II levels, 116 coprolites are coordinated, with some attributed to cave hyena in S12 (Lartigot-Campin & Moné, 2017). In these levels anthropic impact evidence (lithic industry and burnt bones) are recorded, that confirm recurrent carnivore occupations during short abandonment periods of the site by human.

336 Two unit II levels are mainly constituted of coprolite beds, S12a and S13a, between 337 occupation levels with human activity evidence (fig. 6). Archaeostratigraphical studies permit 338 to observe coprolite concentrations in \$12a, between the bands 21 to 22, especially in areas 339 M, N and O (fig. 10), with four carnivore remains. Indeed, two deciduous tooth of young cave 340 hyena, one badger tibia and one sesamoid of unspecified bear are recorded. Likewise, one 341 ungulate remain was found in S12a corresponding to a whole horse radio-ulna with gnawing 342 marks (fig. 9a). Spatial distribution shows a coprolite concentration more or less circular in 343 the squares N21 and O21 that suggests a latrine area (Bearder and Randall, 1978; Kruuk, 344 1972; Pineda et al., 2017), as well as in S13a, located in the square I21. Further study will 345 confirm the presence or absence of latrines. However, it is interesting to see these coprolite 346 beds at the limit of human occupations.



Fig. 10: Spatial distribution of levels S12a (black) and S13a (grey). Two areas with a coprolites concentration look like latrines.

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## 351 - Skeletal Part and Coprolite Distribution

Fragments of indeterminate long bone (2,469 of total NR corresponding to 72% total NR without coprolites) are predominant in the distribution of anatomical elements (tab. 7) with isolated teeth (385 of total NR as 10% of Total NR without coprolites) attesting to high taphonomic impact. However, fragile elements are recorded in the sequence as cranium (76 total NR), vertebrae (46 total NR), and ribs (36 total NR), adding carnivore coprolites (135 total NR) distributed in the three levels and in S12a.

Main species are less than 100 total NISP, except for the red deer in S11 (116 total NISP) with prevalence of isolated teeth and long bones, such as metapodials, in each level (tab. 7). Almost all skeletal elements of red deer are recorded in S11 and S12, whereas axial elements and bones of hind quarter are missing in S10. For the horse, skeletal distribution seems reversed compared to red deer remains with almost all skeletal elements in S10 and S11, whereas hind leg bones are lacking in S12. For secondary species partial skeletal distribution is recorded in each level with mainly teeth and long bones.

|                                |             | S10    |         |             | \$11   |        | S12a        |             | S12    |        |             | Total |         |
|--------------------------------|-------------|--------|---------|-------------|--------|--------|-------------|-------------|--------|--------|-------------|-------|---------|
|                                |             | Main s | species |             | Main s | pecies |             |             | Main s | pecies |             | Mains | species |
|                                | Total<br>NR | R      |         | Total<br>NR | R      | 3      | Total<br>NR | Total<br>NR | R      | RT     | Total<br>NR | R     | M       |
| Antler/horn                    | 13          | 9      | -       | 11          | 7      | -      | -           | 5           | 1      | - 1    | 29          | 17    | 0       |
| Cranium                        | 16          | 1      | 2       | 43          | 5      | 2      | -           | 17          | 1      | 1      | 76          | 7     | 5       |
| Maxilla                        | 6           | -      | -       | 3           | -      | -      | -           | 3           | -      | -      | 12          | 0     | 0       |
| Mandible                       | 4           | -      | -       | 9           | 1      | 2      | -           | 7           | 3      | -      | 20          | 4     | 2       |
| Tooth                          | 124         | 13     | 37      | 170         | 31     | 51     | 2           | 89          | 12     | 24     | 385         | 56    | 112     |
| Vertebra                       | 11          | -      | 2       | 19          | 2      | 3      | -           | 16          | 2      | 1      | 46          | 4     | 6       |
| Rib                            | 4           | -      | 2       | 22          | 3      | 5      | -           | 10          | -      | 1      | 36          | 3     | 8       |
| Scapula                        | 1           | 1      | -       | 2           | -      | 1      | -           | 2           | -      | -      | 5           | 1     | 1       |
| Humerus                        | 2           | 1      | -       | 8           | 5      | -      | -           | 4           | 1      | 2      | 14          | 7     | 2       |
| Radio-ulna                     | 9           | 7      | 1       | 12          | 7      | 2      | 1           | 3           | 1      | 1      | 25          | 15    | 5       |
| Carpus                         | -           | -      | -       | 2           | -      | 1      | -           | 1           | 1      | -      | 3           | 1     | 1       |
| Metacarpus                     | 11          | 7      | 4       | 24          | 13     | 6      | -           | 6           | 5      | 1      | 41          | 24    | 11      |
| Pelvis                         | 1           | -      | -       | 6           | 1      | 4      | -           | 4           | -      | 3      | 11          | 1     | 7       |
| Femur                          | 1           | -      | -       | 9           | 6      | -      | -           | 5           | 2      | 2      | 15          | 8     | 2       |
| Tibia                          | 2           | -      | 2       | 4           | 3      | -      | 1           | 6           | 4      | -      | 13          | 7     | 2       |
| Tarsus                         | 2           | -      | 1       | 5           | 1      | 1      | -           | 3           | 2      | -      | 10          | 3     | 2       |
| Metatarsus                     | 19          | 8      | 5       | 34          | 20     | 5      | -           | 11          | 10     | -      | 64          | 38    | 10      |
| Metapodial                     | 4           | 1      | 1       | 3           | 1      | -      | -           | 2           | 1      | -      | 9           | 3     | 1       |
| Sesamoid                       | 2           | 1      | -       | 8           | 3      | 1      | 1           | 2           | 1      | -      | 13          | 5     | 1       |
| Phalanx                        | 17          | 4      | 4       | 21          | 7      | 2      | -           | 6           | 1      | 2      | 44          | 12    | 8       |
| Long bone                      | 580         | -      | -       | 1596        | -      | -      | -           | 293         | -      | -      | 2469        | -     | -       |
| Cancellous bone                | 62          | -      | -       | 208         | -      | -      | -           | 48          | -      | -      | 318         | -     | -       |
| Flat bone                      | 5           | -      | -       | 14          | -      | -      | -           | 5           | -      | -      | 24          | -     | -       |
| Coprolite                      | 7           | -      | -       | 39          | -      | -      | 27          | 43          | -      | -      | 116         | -     | -       |
| Total NR                       | 903         |        |         | 2272        |        |        | 32          | 591         |        |        | 3798        |       |         |
| Total NR<br>without coprolites | 896         | 52     | 61      | 2233        | 116    | 86     | 5           | 548         | 48     | 38     | 3682        | 216   | 186     |

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Tab. 7: Distribution of anatomical elements in unit II levels (NR) with main species representation (NISP)

367 Fragmentation

368 In unit II levels, 99% of long bones (without considering phalanges) are broken. The 369 diaphysis lengths are generally less than 50 mm and the diaphysis circumferences are either less than half of their original dimensions, or complete in rarer cases (as for the few whole 370 371 bones).

372 For main species, long bones are not represented with the same size distribution (fig. 11). 373 Red deer bones are highly fragmented with most remains measuring less than 50 mm in the 374 three levels (fig. 11), corresponding to less than <sup>1</sup>/<sub>4</sub> diaphysis lengths. One complete epiphysis 375 was discovered in S11, an unfused proximal extremity of the right tibia, and no whole elements are recorded in unit II levels. This profile can be explained by the prevalence of 376 377 anthropic impacts on red deer remains (tab. 6).

378 Throughout the sequence, the horse is the only large mammal with complete long bones, 379 excavated in unit II levels and corresponding to metapodials in S10 and S11 (fig. 11) with a 380 radio-ulna in S12a (fig. 9a). Furthermore, two complete metatarsal epiphyses, with their 381 diaphysis portion, have been reassembled in S10 and two complete metacarpal epiphyses with 382 their diaphysis portion have likewise been reassembled in S11. These remains are counted as 383 partial element in NISP but form two actual whole metapodials in addition to the other 384 complete long bones. Considering this low fragmentation with unburnt whole bones, human 385 activities seem less affect horse remains, supporting by carnivore tooth marks on unburnt 386 elements (fig. 9a, b, c).



Fig. 11: Fragmentation indexes of long bones. Comparison of NISP between red deer and horse by Size class (mm), partial elements with a complete epiphysis (Part), and complete long bones (Whole)

### 390 Anthropic Activities

391 Human impact (tab. 6) is mainly attested with various burning processes affecting faunal 392 remains mostly for red deer (tab. 8). Degree 1 (brown colour), is the most represented bone 393 modification (47 to 61% of affected NR) decreasing from the base to the top. This first step 394 applies to heated bones at low temperature less than 300°C (Lebon, 2009). Degree 4 (grey 395 colour) is well represented (16 to 27% of affected NR), increasing between levels. This 396 burning step, for temperatures between 400 and 550°C, and partly explain the high 397 fragmentation of faunal material. In contrast, degree 3 (black colour) and degree 6 (white 398 colour) with less than 5% of affected NR are the least represented. Double colouration is 399 observed on 5 to 8% of affected NR.

400

|      |                      |     |        |      |         | C11 C12 |              |      |         |     |        |          |              |      |       |        |        |  |  |  |
|------|----------------------|-----|--------|------|---------|---------|--------------|------|---------|-----|--------|----------|--------------|------|-------|--------|--------|--|--|--|
|      |                      |     | 5      | 510  |         |         | <b>S</b> 1   | 11   |         |     | S      | 512      |              |      | Т     | otal   |        |  |  |  |
|      |                      |     |        | Main | species |         |              | Main | species |     |        | Main     | Main species |      |       | Main s | pecies |  |  |  |
|      |                      | То  | tal NR |      | M       | Tot     | al NR        | T    | M       | То  | tal NR | <b>H</b> | <b>X</b>     |      | al NR | R TH   | 1      |  |  |  |
| 1    | Brown                | 336 | 47.3%  | 24   | 19      | 866     | <b>50.8%</b> | 45   | 38      | 173 | 61.1%  | 22       | 13           | 1375 | 50.9% | 91     | 70     |  |  |  |
| 2    | Brown black          | 52  | 7.3%   | 4    |         | 135     | 7.9%         | 7    | 2       | 12  | 4.2%   | 1        |              | 199  | 7.4%  | 12     | 2      |  |  |  |
| 3    | Black                | 21  | 3.0%   | 1    | 3       | 67      | 3.9%         | 4    | 3       | 22  | 7.8%   | 1        | 2            | 110  | 4.1%  | 6      | 8      |  |  |  |
| 4    | Dark grey<br>/black  | 193 | 27.2   | 7    | 4       | 392     | 23.0%        | 19   | 5       | 44  | 15.6%  | 8        | 2            | 629  | 23.3% | 34     | 11     |  |  |  |
| 5    | Light grey<br>/white | 46  | 6.5    | 1    |         | 103     | 6.0%         | 5    | 1       | 13  | 4.6%   | 3        | 1            | 162  | 6.0%  | 9      | 2      |  |  |  |
| 6    | White blue           | 8   | 1.1%   | 1    |         | 39      | 2.3%         | 2    |         | 5   | 1.8%   |          |              | 52   | 1.9%  | 3      | 0      |  |  |  |
| Dout | ole coloration       | 54  | 7.6%   | 2    | 1       | 104     | 6.1%         | 6    | 6       | 14  | 4.9%   | 4        |              | 172  | 6.4%  | 12     | 7      |  |  |  |
|      | Total                | 710 | 100%   | 40   | 27      | 1706    | 100%         | 88   | 55      | 283 | 100%   | 39       | 18           | 2699 | 100%  | 167    | 100    |  |  |  |

401

Tab. 8: Heating (1) and burning (2 to 6) degrees in unit II levels of Ramandils Cave (NR), with main species representation (NISP).

402 Heating or burning processes are observed on each skeletal element in unit II levels, 403 mainly on teeth and indeterminate long bones. For red deer from the three main levels (fig. 404 12), heating is likewise generally localised on teeth and long bones, especially on metapodials 405 and radio-ulna, followed by cranium and antler fragments. These distributions support the 406 hypothesis of bone marrow exploitation with long bone preparation by heating process.

407 Bones bearing cut marks (13 to 19% NR), or green fractures (9 to 15% NR), with higher 408 percentages in S11 and S10 (tab. 6) are mainly concentrated on unspecified small ungulates 409 as well as on red deer. These anthropic modifications present similar distribution on red deer 410 skeletal (fig. 13), mostly located on long bones, especially on metapodials and radio ulna, 411 likewise for heated processes (fig. 12).

412



413 414

Fig. 12: Skeletal distribution (NISP) of heated bones (degree 1) for red deer.





Fig. 13 (a) Cut mark and (b) green fracture distribution on red deer skeletal (NISP) in unit II levels of Ramandils Cave

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418 Several butchery step were noticed, generally affecting burnt remains, based on 419 morphology, localization and distribution of traces over the bone surface (Binford 1981).

420 Cut marks on burnt cranium of red deer in S11 (fig. 13a) and on one horse phalange in S10 421 could indicate skinning process, as well as for wild boar, with a cranium and phalanges 422 likewise affected in these same levels. However, unlike the wild boar and fallow deer, third 423 phalanges are missing for all ungulates, which may support the hypothesis of an out-site 424 skinning process on main species, or show a differential treatment such as burning process. 425 Incisions on articular parts are observed on carpus and mandible, thoracic vertebrae and long 426 bones extremities (humerus, femur and metapodials) for the red deer in S11 and S12 (fig. 427 13a), interpreted as disassembling step, likewise recorded for aurochs in S11. Traces are 428 recorded in the three levels, on mandibles, ribs, or diaphysis of meat-rich long bones for main 429 species (fig. 13a), showing defleshing processes, likewise for secondary species, essentially 430 aurochs (in S10 and S11) and for wild boar (in the three levels).

Scraping marks are noticed on diaphysis of red deer metapodials (fig. 13a). These meatpoor elements are affected by long fine parallel striation to the longitudinal bone axe (fig. 14a) or short and close parallel marks perpendicular to the bone shaft (fig. 14c) caused by the removal of periosteum. This cleaning process allows to facilitate bone breakage. Moreover, green fracture on mandible and long bones, mainly on red deer metapodials (fig. 13a, fig. 14a), represents the bone marrow removing step, abundant inside these anatomical parts. Some of these remains show percussion marks related to this process.

Carnivores are also affected by butchery marks noticed on heated remains. Incisions are noticed on a wolf occipital in S12 (fig. 14d) and a cave bear phalange in S11 (fig. 14f), which can be explained by a disassembling process for the first and a skinning step for the second. Remains of brown bear, *Ursus arctos*, are likewise heated in unit II, with one phalange in S12 and one fibula bearing cut traces could interpreted as skinning marks, in S13 (fig. 14e).

444 At last, technological applications of bones are identified in the three levels, corresponding 445 to scarce bone retouchers (tab. 6), mainly on heated long bones of cervids (fig. 14b).



Fig. 14: Anthropic marks on large mammal remains: (a) scraping marks and green fracture on an unburnt metacarpal of red deer, (b) bone retoucher on a brown heated metatarsal of roe deer, (c) scraping marks on blue burnt metacarpal of red deer, (d) cut marks on an heated brown temporal of wolf, (e) cut marks on brown heated fibula of brown bear, (f) cut marks on brown heated phalanx of cave bear (photo by L. Rusch).

### 4.3.3. Mortality profiles and seasonality

452

A low MNI is defined for each species (tab. 9) which not exceeds seven individuals, with red deer in S11. Closer count is recorded in S11 between main species, red deer (7 individuals) and horse (6 individuals), and secondary species are well represented in this level by European wild ass (5 individuals), aurochs (5 individuals) and wild boar (4 individuals). Furthermore, fallow deer, with 5 individuals, counts higher MNI than in other levels. In S10 equids are greater represented, by horse and European wild ass, with 6 individuals for each,

|                  |   | S10 |    |    |   |     |   | s  | 11 |    |   | S12A |     |   | s | 12 |    |   |     |   | Тс | otal |    |   |     |
|------------------|---|-----|----|----|---|-----|---|----|----|----|---|------|-----|---|---|----|----|---|-----|---|----|------|----|---|-----|
|                  | F | Y   | SA | А  | 0 | MNI | F | Υ  | SA | А  | 0 | MNI  | MNI | F | Y | SA | А  | 0 | MNI | F | Y  | SA   | А  | 0 | MNI |
| Total carnivores | - | 2   | -  | 5  | - | 7   | - | 2  | -  | 7  | - | 9    | 3   | - | - | -  | 4  | 1 | 5   | - | 6  | -    | 16 | 1 | 24  |
| U. arctos        | - | 1   | -  | -  | - | 1   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | -  | - | 0   | - | 1  | -    | 1  | - | 2   |
| U. spelaeus      | - | -   | -  | 1  | - | 1   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | 1  | - | 1   | - | -  | -    | 3  | - | 3   |
| M. meles         | - | -   | -  | -  | - | 0   | - | -  | -  | 1  | - | 1    | 1A  | - | - | -  | -  | - | 0   | - | -  | -    | 2  | - | 2   |
| Lutrinae indet.  | - | -   | -  | 1  | - | 1   | - | -  | -  | -  | - | 0    | -   | - | - | -  | 1  | - | 1   | - | -  | -    | 2  | - | 2   |
| C. lupus         | - | -   | -  | -  | - | 0   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | 1  | - | 1   | - | -  | -    | 2  | - | 2   |
| V. vulpes        | - | -   | -  | 1  | - | 1   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | 1  | - | 1   | - | -  | -    | 3  | - | 3   |
| C. c. spelaea    | - | 1   | -  | -  | - | 1   | - | 2  | -  | -  | - | 2    | 2Y  | - | - | -  | -  | 1 | 1   | - | 5  | -    | -  | 1 | 6   |
| L. spelaeus      | - | -   | -  | 2  | - | 2   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | -  | - | 0   | - | -  | -    | 3  | - | 3   |
| F. silvestris    | - | -   | -  | -  | - | 0   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | -  | - | 0   | - | -  | -    | 1  | - | 1   |
| Total herbivores | 0 | 10  | 3  | 10 | 1 | 24  | 2 | 14 | 4  | 13 | 6 | 39   | 1   | 1 | 8 | 3  | 7  | 0 | 19  | 3 | 32 | 10   | 30 | 7 | 83  |
| P. antiquus      | - | -   | -  | -  | - | 0   | - | 1  | -  | -  | - | 1    | -   | - | - | -  | -  | - | 0   | - | 1  | -    | -  | - | 1   |
| S. hemitoechus   | - | 1   | -  | -  | - | 1   | - | 1  | -  | 1  | - | 2    | -   | - | 1 | -  | -  | - | 1   | - | 3  | -    | 1  | - | 4   |
| E. f. germanicus | - | 3   | 1  | 2  | - | 6   | - | 3  | 1  | 2  | - | 6    | 1A  | - | 2 | 1  | 1  | - | 4   | - | 8  | 3    | 6  | - | 17  |
| E. hydruntinus   | - | 2   | 1  | 3  | - | 6   | 1 | 2  | -  | 2  | - | 5    | -   | - | - | -  | 2  | - | 2   | 1 | 4  | 1    | 7  | - | 13  |
| Bison sp.        | - | -   | -  | -  | - | 0   | - | -  | -  | -  | 1 | 1    | -   | - | - | -  | -  | - | 0   | - | -  | -    | -  | 1 | 1   |
| B. primigenius   | - | 2   | -  | 1  | - | 3   | - | 3  | -  | 1  | 1 | 5    | -   | - | 3 | -  | -  | - | 3   | - | 8  | -    | 2  | 1 | 11  |
| C. caucasica     | - | -   | -  | 1  | - | 1   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | 1  | - | 1   | - | -  | -    | 3  | - | 3   |
| C. elaphus       | - | 1   | 1  | 1  | - | 3   | - | 2  | 1  | 2  | 2 | 7    | -   | 1 | 1 | 1  | 2  | - | 5   | 1 | 4  | 3    | 5  | 2 | 15  |
| D. dama          | - | -   | -  | -  | 1 | 1   | - | 1  | 1  | 1  | 2 | 5    | -   | - | 1 | -  | -  | - | 1   | - | 2  | 1    | 1  | 3 | 7   |
| R. tarandus      | - | -   | -  | -  | - | 0   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | -  | - | 0   | - | -  | -    | 1  | - | 1   |
| C. capreolus     | - | -   | -  | 1  | - | 1   | - | -  | -  | 1  | - | 1    | -   | - | - | -  | 1  | - | 1   | - | -  | -    | 3  | - | 3   |
| S. scrofa        | - | 1   | -  | 1  | - | 2   | 1 | 1  | 1  | 1  | - | 4    | -   | - | - | 1  | -  | - | 1   | 1 | 2  | 2    | 2  | - | 7   |
| Total            | - | 12  | 3  | 15 | 1 | 31  | 2 | 16 | 4  | 20 | 6 | 48   | 4   | 1 | 8 | 3  | 11 | 1 | 26  | 3 | 38 | 10   | 48 | 8 | 107 |

459 compared to red deer (3 individuals) and aurochs (3 individuals).

460 Tab. 9: Age groups of large mammals (MNI by combination) in unit II levels of Ramandils Cave: (F) foetus, (Y) young, (SA) sub-adult, 461 (A) adult, and (O) old individuals.

462 Carnivores are scarcer, not exceeding 2 individuals by species, and relatively more 463 represented by cave hyena, with 6 individuals of total MNI. Young individuals are only 464 recorded for brown bear in S11 and cave hyena in S11 and S12a (tab. 9), and one senile cave 465 hyena is noticed in S12. 466 For ungulates, the mortality patterns indicate that all age groups are present in unit II, and 467 particularly among dominant species. Adults are the most abundant, closely followed by 468 young individuals (infants or juveniles) and to a lesser extent by sub-adults. This last age 469 group corresponds to animals which have nearly reached their maximum size, meaning that 470 hunting a sub-adult prey could have involved the same diet resources as an adult animal. One 471 senile fallow deer is recorded in S10 while six senile prey are represented in S11 472 corresponding to one bison, one aurochs, two red deer and two fallow deer. We determined 473 three foetus teeth in unit II, for wild boar and European wild ass in S11 and for red deer in 474 S12, meaning that female adult animals were targeted, however no criteria on bones remains 475 allow us to distinguish them.

476 Ungulates were generally slaughtered several times during the year (fig. 15), with spring, 477 autumn and winter always recorded in the three levels. Prey selection followed a seasonal 478 pattern for cervids, systematically occurring during autumn, in the three levels and during 479 spring in S12. In the same way, in S11 and S10, horse was generally hunted between spring 480 and summer, European wild ass rather during winter, and wild boar during summer, while in 481 S12, seasonal recording is not represented for these species. Moreover, aurochs were 482 slaughtered during various periods, hunted during spring in S11 whereas this bovid is 483 represented during autumn in S10 and S12, as well as during winter in S12.



484

Fig. 15: Large mammals saisonality in unit II levels of Ramandils Cave

### 485 **5. Discussion**

The results obtained in this study contribute relevant details to knowledge of Neanderthal and carnivore occupations in unit II of Ramandils Cave (tab. 10), and therefore on Mediterranean coast during the Upper Pleistocene. The three more abundant unit II levels record several occupation periods occurred, the densest of which is undoubtedly observed in level S11.

491 5.1. Occupations in Unit II Levels

For the material accounted, S8, S9, S13 and S14 are the lower levels of unit II, except for coprolites levels S12a and S13a. Further studies will refine these four occupation levels for allow us to characterise unit III and unit I limits with unit II levels.

495 - S13a

S13a level constitute a coprolite deposit in a concentrated area more or less circular. This
distribution could be interpreted as a latrine deposit, usually attributed to cave hyena layer
(Bearder and Randall, 1978; Crezzini *et al.*, 2015; Fernandez Rodriguez *et al.*, 1995; Fosse,
1997; Fourvel, 2012; Kruuk, 1972; Pineda *et al.*, 2017), indicating at least one carnivore
occupation during long periods of human abandonment. However, further studies will allow
us to characterise this level.

502 - S12

Anthropic impact is well attested in S12 level, especially with lithic industry and burnt faunal bones. Red deer and horse remains are prevalent for large mammals, in NISP and MNI. Secondary species are recorded mainly corresponding to aurochs, European wild ass, and wild boar, with lower values of NISP and MNI. Several carnivore species were identified, for less than 6% NISP, relatively represented by wolf and cave hyena, with a low MNI.

508 Almost all skeletal parts are recorded for the red deer, explained by a whole carcasses 509 transport, but hind limb bones were not found for the horse, which could be explained by 510 partial carcass transport likewise for secondary species. However, the lack of third phalanges 511 for all species may indicate a previous skinning step out-site or differentiated impacts for this 512 anatomical elements such as with burning processes. Several butchery steps are noticed, 513 mainly on long bones of red deer. Firstly, disassembling and defleshing marks indicate a meat 514 exploitation, likewise noticed for one wolf remain for the first process. Secondly, scraping 515 marks, resulting of bone cleaning processes, mainly on metapodials, with green fractures, and 516 some percussion marks induce a bone marrow consumption, supported by the high material 517 fragmentation.

Then, high values of heated to burnt bones are highlighted in this level, without any hearths recorded in the sequence. These structures may be located in non-excavated area or might not be preserved, especially in the eroded area on the front of the cave, caused by the active natural spring. Heating and burning in the set may suggest various processes and could partly explain the high material fragmentation. Scarce double colouration on same bone surface indicates differential heat exposure and could suggest possible roasting of the meat, prior to 524 defleshing (Rosell et al., 2012). However, the high bone fragmentation can affect evidence of this meat preparation process, especially for diaphysis fragments. Low calcination can 525 526 indicate a bone preparation to facilitate their fractures (Cáceres et al., 2002) or preparing bone marrow for removal (Oliver, 1993; Speth et al., 2012), while high calcination could show 527 528 possible cleaning activities. The low representation of cancellous bones (articular portions, 529 limb extremities or axial skeleton) and the small number of percussion fragments does not 530 exclude the hypothesis of bone grease exploitation or the use of bones as fuel (Daujeard, 2008; Lebon, 2008; Morin, 2007, 2010). Further studies would allow us to explain these 531 532 proportions.

Young and adult ungulates, with close MNI, seem to be slaughtered without any age selection. Despite low MNI of large mammals, studies show occupations occurred during almost each seasonal period of the year. This faunal studies could indicate specialised seasonal hunting, especially focus on cervids (red deer and fallow deer) during spring and on aurochs during winter, with a mixed period for red deer and aurochs during autumn, other species such as horse not estimating in months.

539 Moreover, low carnivore occurrence is noticed within S12 level. Few carnivore remains 540 are identified, especially wolf and cave hyena. Scarce tooth and digestion marks, mainly 541 affecting heated and fractured ungulate bones, and coprolites induce scavenging during short 542 human abandonment of the cave.

543 - S12a

544 Another coprolite layer, noted S12a, was described during this study, implying recurrent 545 carnivore occupation of Ramandils cave in unit II. In addition to a richer coprolite bed with a 546 concentrated area more or less circular, S12a records few carnivore remains, especially two 547 young cave hyena (2 MNI), as well as one gnawed long bone of horse, the only one ungulate 548 remains. These results could be interpreted as a latrine deposit and explained by the use of the 549 cave as a den by the cave hyena, usually generating this type of distribution (Kruuk, 1972; 550 Bearder & Randall, 1978; Pineda et al., 2017). However, further studies will allow us to 551 characterise this layer.

552 - S11

553 S11 is the richest unit II level with anthropic activities such as lithic industry and butchery 554 processes. The same main species are recorded than in S12 with the prevalence of red deer 555 and similar distribution is noticed for secondary species. Carnivores are more varied for less 556 than 6% NISP, relatively represented by hyena and canids with a low MNI.

557 Almost all skeletal parts are recorded for the main species, explained by a whole carcasses 558 transport, whereas, as in S12, a partial transport is interpreted for secondary species. 559 However, the lack of third phalanges for ungulates, except for wild boar and fallow deer, may 560 still indicate a previous skinning step out-site or differentiated impacts for this anatomical 561 elements such as with burning processes. The same butchery steps are noticed, mainly on long 562 bones of red deer, with the addition of skinning marks for red deer and wild boar, likewise on 563 a cave beer phalange. High values of heating and burning processes have the same 564 interpretation as for \$12.

565 Young and adult ungulates, with close MNI, seem to be slaughtered without any age 566 selection. Despite low MNI of large mammals, seasonality studies could indicate longer or 567 more frequent occupations than in S12. Specialised seasonal hunting seems focus on cervids 568 during autumn and on bovids during spring, with a regular slaughtering of suids and equids 569 (horse and European wild ass) for almost each seasonal periods of the year. European wild ass 570 seems rather hunted during winter, as well as suids, likewise killed during summer.

571 Moreover, low carnivore occurrence is still noticed within S11 level with the same 572 scavenging marks during short human abandonment of the cave. However, considering the 573 preservations of few whole long bones and tooth marks on unburnt remains, not affected by 574 human activities, carnivore may have transported some of their own prey into the cave ().

575 - S10

576 No coprolites level is noticed between S11 and S10 levels as opposed between S13 and 577 S12 with S13a level and between S12 and S11 with S12a level. In S10, recording human 578 occupation evidence, horse is prevalent compared to red deer with the same faunal spectrum 579 and close values for secondary species as in S12 and S11. Carnivores are fewer diversified, 580 represented in lesser extend by cave hyena and lynx.

581 Almost all skeletal parts are recorded for the horse, explained by a whole carcasses 582 transport, but axial elements and bones of hind quarter were not found for the red deer, which 583 could be explained by partial carcass transport likewise for secondary species. However, the 584 lack of third phalanges for ungulates, except for wild boar and fallow deer, may indicate a 585 previous skinning step out-site or differentiated impacts for this anatomical elements such as 586 with burning processes. Several butchery marks are noticed, mainly on long bones of red deer. 587 Skinning step is highlighted for horse and wild boar. Disassembling is not recorded in S10 but 588 other butchery steps are the same as in S11 and S12, likewise for heating and burning 589 processes with the same interpretations as for other levels.

Young and adult ungulates, with close MNI, seem to be slaughtered without any age selection as seen in other levels. Seasonality studies show occupations occurred each seasonal period of the year. Despite low MNI of large mammals, this faunal studies could indicate specialised seasonal hunting, similar to the previous levels, for Cervids and bovids which seem rather killed during autumn, as seen in S12, whereas equids were slaughtered during almost each season, with a focus on European wild ass during winter, as notice in S11. Therefore, Suids seem killed during summer likewise in S11 level.

597 Carnivore occurrence is still noticed within S10 level with the same scavenging marks 598 during short human abandonment of the cave. However, considering the preservations of few 599 whole long bones and tooth marks on scarce unburnt remains, not affected by human 600 activities, carnivore may have transported some of their own prey into the cave.

|      |  |  | Human a  | ctivities               | 5                        |                                     |                    | Ca   | rnivores a  | ctivities          |            |
|------|--|--|--|-------------------------|--------------------------|-------------------------------------|--------------------|--|---|--------------------|------------|
|      | Hunted prey  | Transport  | Butchery steps   | Fire                    | Exploitation             | Saisonality                         | Bone<br>industry   | Main<br>species                                  | Activities  | Coprolites         | Occupation |
| S10  | Main:<br>- Horse<br>- Red deer<br>Secondary:<br>- Aurochs<br>- European wild ass<br>- wild board | Whole<br>Partial<br>Mainly skull<br>and<br>limb ends               | - Skinning<br>- Defleshing<br>- Breakage   | Heating<br>+<br>Burning | Meat<br>+<br>Bone marrow | Seasonal<br>selective<br>slaughters | Rare<br>retouchers | Cave hyena<br>(young)<br>+<br>Lynx (adult)       | Scavenging<br>+<br>Few gnawing<br>remains no<br>affected by<br>human            | yes                | occasional |
| S11  | Main:<br>- Red deer<br>- Horse<br>Secondary:<br>- Wild board<br>- European wild ass<br>- Aurochs | Whole<br>Partial<br>Selection<br>of pieces<br>according to<br>taxa | - Disassembling<br>- Defleshing<br>- Breakage<br>- Skinning<br>- Disassembling<br>- Defleshing<br>- Breakage | Heating<br>+<br>Burning | Meat<br>+<br>Bone marrow | Seasonal<br>selective<br>slaughters | Rare<br>retouchers | Cave hyena<br>(young)<br>+<br>Fox (adult)        | Scarce:<br>Scavenging<br>+<br>Few gnawing<br>remains no<br>affected by<br>human | yes                | occasional |
| S12a |  |  |  |                         |                          |                                     |                    | Cave hyena<br>(Young)                            | Gnawing on<br>one whole<br>radio-ulna<br>of horse                               | Latrine<br>level ? | Den?       |
| S12  | Main:<br>- Red deer<br>- Horse<br>Secondary:<br>- Wild board<br>- European wild ass<br>- Aurochs | Whole<br>Partial<br>Mainly skull<br>and<br>limb ends               | - Disassembling<br>- Defleshing<br>- Breakage<br>- Defleshing<br>- Breakage                                  | Heating<br>+<br>Burning | Meat<br>+<br>Bone marrow | Seasonal<br>selective<br>slaughters | Rare<br>retouchers | Cave hyena<br>(young / old)<br>+<br>Wolf (adult) | Scavenging  | Yes                | occasional |
| S13a |  |  |  |                         |                          |                                     |                    | None   |   | Latrine<br>level?  | occasional |

601

Tab. 10: Synthesis of human and carnivore activities in unit II levels of Ramandils cave

In S10 S11, S12 levels, the material accumulation is mainly due to human input representing various activities during several occupations periods recorded within unit II but also in the whole sedimentary filling of the cave.

Even if the archeological interpretation from paleoenvironments to settlement pattern depends on the temporal resolution of the data (Carbonell *et al*, 2012), this first and preliminary multidisciplinary approach allow us to propose and discuss various hypotheses on the general human mobility pattern, the site function and its status in the territory.

- 609 5.2. Mobility pattern
- 610 Economic and territorial pattern

611 Even if, at that stage of the study, lithics and faunal remains are not analysed at the same 612 stratigraphic resolution level, we could however begin the first discussion about the economic 613 general pattern of the unit II occupations.

614 Generally, Ramandils settlements seems to be linked to the exploitation of the tertiary flint 615 sources of the Narbonne-Sigean Basin (fig. 16). Raw material origins are mainly from the 616 semi-local area (between 5 and 20 km) around the site. Excepted the jasper, the unique raw 617 material imported from the remote area (35 km), every raw material came from the 618 surrounding of the site between 7 km and 17 km. Flint were principally extracted from the 619 tertiary outcrops of Sigean, Presqu'île du Doul, Portel des Corbières, Hameau du lac. More 620 rarely some cherts were exploited a little further from the cretaceous and jurassic outcrops and 621 the others kind of raw material were collected in the Aude alluvial deposits (quartz, quartzite 622 and sandstone...) at the north of the site or sometimes on the beach deposits, just near the 623 cave.

Despite this semi-local procurement pattern, which would rather indicate a strong and well organized exploitation in this area, the punctual arrival of jasper from the Roussillon plain, at 35 km south of the site, seems to indicate that groups had an important mobility on the territory and they import punctually materials of interest from these more remote areas that they browse probably for others activities.

629 Moreover, the faunal acquisition is more oriented towards all the available environments around the sites (Rusch et al., 2019). Main species, dominated by red deer and horse, 630 631 highlight grassland with forests, which could be situated less than 5 km around the cave. 632 Marine shells recorded throughout the sequence involve a transport into the site of less than 633 1,4 m, compared to the current coast. Further studies will allow us to understand the use or 634 consummation strategies of these remains. Furthermore, ibex remains found in this unit 635 involve a rocky environment, located at least in the semi local area between 5 and 20 km (fig. 636 8), because we could hypothesize that ibex were situated on the nearest reliefs of the 637 Corbières massif.

638 The local area seems to be exploited for its subsistence resources while semi local area 639 provided more than 99 % of the lithic raw material used and occasionally this area is 640 exploited for the acquisition of ibex (fig. 16).

This pattern is different from the regional Mousterian sites like Crouzade Cave on the cretaceous Clape massif near Narbonne (Saos *et al.*, in press) and Hortus site on the Montpellier region (Lebègue *et al.*, 2010) which are characterised by a mainly local and semilocal flint and animal resources supply but completed with an additional allochthonous lithic raw material generally coming from more than 80 km away. These occupation sites are interpreted as short terms hunting camp or seasonal camps (Grégoire, 2012).





648 649 650

Fig. 16: Economic exploitation territory divided in: local-0 to 5km, semi-local-5 to 20km, and remote area-more than 20 km, including lithic sources and potential location of the main species exploited as food resources.

651 We must go to the Ardèche region, more than 250 km north of Ramandils Cave, at Saint Marcel Cave, to find a Middle Palaeolithic site where the economic mobility pattern is similar 652 653 (Moncel et al, 2004, Daujeard & Moncel, 2010), with a semi-local supply, oriented on flint 654 procurement, that not exceed 10 km. The different occupations phases of this site, more or 655 less contemporaneous than the Ramandils settlements, are qualified by the authors as "long term residential camp" thanks to the exploitation and consumption strategies, the lithic 656 657 remains density and the topographic situation for example.

658 In the case of Ramandils cave, arrival of allochthonous raw material testify probably of a 659 logistic mobility (Binford, 1980) from areas exploited for their own resources, to a base-camp 660 (Ramandils?) where natural shelter, water source, faunal abundance from various ecosystems (grassland, forest, mountain, and sea with marine shells), proximity of flint sources and of the 661 662 shore, makes this place attractive and justify the frequency and the persistence of the 663 settlement. This mobility pattern indicates a territorial economic organization where Ramandils site would take a central place and could work like an aggregation site (Kuhn, 664 1995) within an economic territory of a 35 km radius. 665

The large quantity of lithic material and the complete reduction sequence on flint testify to an intense production activity on the site during the unit II settlement. The abundance of small artefacts resulting from (i) a first massive production of small flakes, recycled in many cases as matrix for (ii) a second phase of flake production, tend to indicate that Ramandils cave seems to be a production site dependent on nearby flint sources which could be interpreted as a long term settlement because of the production strategies.

Indeed, from the techno-economic point of view, a general overview of the unit II allows to say that the groups strongly occupied the local and semi-local territory (fig. 16) and, at the same time, carried out intensive lithic production activities on the cave. This intensive activity is also noticeable thanks to the "ramification" process (Bourguignon *et al.*, 2004) in the flake production. These facts, added to the previous one, are also in favour of the long term residential camp.

678 On the other hand, the incomplete reduction sequence for almost nine raw materials other 679 than flint, but less represented, as well as the very punctual contribution of jasper by artefacts 680 at the end of the reduction sequence, seems to demonstrate, that lithic pieces were circulating 681 easily on the territory, and that the production, with these secondary raw materials, was very 682 fragmented in space and time (Bourguignon et al., 2006) which is to say that people circulate 683 a lot between different points of the territory. These facts could be in favour of a logistic 684 mobility from the residential camp, to more specialised camps and support the assumption 685 that Ramandils cave could be used as a long term residential camp connected with others 686 specialized sites, one of which could be in the jasper supply area.

687 5.3. Site function in unit II

688 By taking account of this first results, on faunal and lithic exploitation, intrinsic to the 689 Mediterranean coastal context (climate, topography, ecosystems, raw material available, 690 human groups, etc.), it is possible to propose two hypotheses of occupation site modes (tab. 691 11). Indeed, several criteria tend to support long-term camps, nevertheless, some data seem to 692 favour in short-term repeat occupation, compared to Upper Pleistocene sequences from 693 southern France such as those in northwestern Mediterranean coast and in Rhone Valley 694 (Daujeard & Moncel, 2010; Daujeard et al, 2011, 2012, 2016; Magniez, 2010; Saos et al., 695 2016).

696 In unit II levels, humans are the principal agents of the input and exploitation of ungulate 697 carcasses, likewise in long-term occupations, such as in Saint-Marcel, Baume des Peyrards 698 and Abri du Maras. However, large mammal spectrum highlights two main species, attributed 699 to red deer and horse, with less than 50% NISP, and various secondary ungulates with 700 European wild ass, aurochs, and wild boar. Moreover, carnivore activities are recorded within 701 (S10, S11, S12) and between (S12a, S13a) these levels, with young individuals, coprolites 702 deposits, butchery and carnivore tooth-marks found on the same species (red deer and horse), 703 which could support the hypothesis of short-term occupations.

704 Faunal studies indicate hominin occupations during each seasonal period of the year, with 705 varied acquisition strategies according to the taxa and more frequent slaughtering during 706 autumn and spring. In all levels, cervids were systematically hunted during autumn, whereas 707 equids were killed during almost each season (in S11 and S10), with a focus on European 708 wild ass during winter. Seasonal acquisition strategies are recorded in several Pleistocene sites 709 such in the Rhone Valley (Daujeard & Moncel, 2010; Daujeard et al, 2016), at Baume des 710 Peyrards, where ibex and most of the secondary species were hunted during the cold season 711 and summer, whereas horse was the only one killed during spring, or at Saint-Marcel, where 712 the fallow deer from level u seems frequently hunted during spring and autumn. Periodic 713 recurrence, according to taxa in Ramandils sequence, could imply a planned seasonal 714 exploitation of the territory, mainly concentrated on local and semi local area, also observed 715 by the studies on raw material procurement.

In unit II levels, adult and young ungulates are the highest classes, followed by sub-adult 716 717 group, while older individuals have lower rates. Slaughtering is therefore generally aimed at 718 the most profitable prey that could suggest the capture of females (adult individuals) and their 719 young and can be interpreted as a human characteristic of primary and immediate access to 720 the animals (Gaudzinski & Roebroeks, 2000; Klein & Cruz-Uribe, 1984; Stiner, 1994). 721 Nevertheless, it is important to note that only young and young adults were considered for 722 estimating seasonality. There is a chance that adults could have been hunted during different events and therefore, different seasons. However, age at death was mainly estimated through 723 724 dental eruption pattern and almost only on isolates teeth.

725 As at Payre, at Baume Flandin and Le Figuier, (Daujeard & Moncel, 2010; Daujeard et al., 726 2016), several transport strategies and butchery processes have been observed for the various 727 ungulates. A differential degree in carcass treatment according to the species is observed in 728 unit II levels of Ramandils Cave. Red deer, reveal most of the hominins action, rarest for 729 secondary ungulates. Main species were almost systematically brought back whole, 730 dismembered, filleted, and broken for an intensive bone marrow recovery, whereas secondary 731 species are mainly represented by skull and limb end. Skinning process is generally scarce 732 and the lack of third phalanges (except for wild boar and fallow deer) could indicate a first 733 butchery step on the kill site. Furthermore, the scarcity of retouchers in these levels show a 734 low technological applications of bones, usually higher for long term residential camps 735 (Daujeard et al., 2018).

Combustion structures are not recorded while numerous burnt flint and bones, with high degrees, show fire was intensely used as well as for long-term occupation sites. Hearths could be located in not excavated area or might not be preserved, especially in the eroded area on the front of the cave, caused by an active natural spring (fig. 2). Heated and burnt remains could therefore result of secondary deposits from hearth cleaning activities. The hypothesis of bone grease exploitation or the use of bones as fuel is not exclude, further studies would allow us to explain the low proportion of cancellous bone and percussion marks.

743 Lithics evidence are rather in favour of the hypothesis of the long term pattern with the 744 high quantity of lithic remains, the intensive flake production in situ with a complete flint reduction process, added to the mainly semi-local supply that requires travel between 7 and 17 745 km for supplies and replenishment (tab. 11). However, the short term pattern cannot be totally 746 747 dismissed supporting by some criteria which indicate a great mobility in the territory like the 748 incomplete reduction sequence for 9 secondary raw materials and the punctual arrival of 749 jasper from 35 km away (fig. 8). In order to go further, it would be necessary to refine lithic 750 studies with the same stratigraphic resolution level as for faunal remains.

751 All these criteria (tab. 11) led us to discuss the status of the site in terms of duration of 752 occupation according to the episodes identified in unit II by archeostratigraphy. All the 753 criteria carrying an information about seasonality, about intensive presence on the site or on 754 the contrary about displacement on the territory were used to evaluate arguments in favour of 755 the two hypotheses tested on this work: the long term residential camp and the short term 756 regular hunting camp. These two categories were chosen to be compared with other studies 757 conducted in South of France, in the same region (Lebègue et al., 2010; Magniez, 2010; 758 Grégoire, 2012; Saos et al., in press) or in the near Rhône valley (Daujeard & Moncel., 2010; 759 Daujeard et al., in Djindjian et al., 2018).

760 However, the site function can also be thought of in terms of activities. Through the lithic 761 toolkit of unit II it is difficult to identify any specialised activities because of the low 762 percentage of flake tools stayed on the site and the absence of traceology study. On the other 763 hand, the large quantity of non-retouched small debitage products and the identification of several and ramified reduction sequences, testify to an intense activity of flint blanks 764 765 production of which a large part could be exported in the form of retouched tools. The non-766 immediate proximity of the tertiary flint sources could testify that the site was selected for its 767 position in relation to these lithic resources to develop easily a flake production activity but 768 also for the immediate proximity of various subsistence resources for the group supplying, 769 during the flint exploitation and production activity. Indeed, despite the relative prevalence of 770 red deer and horse, faunal studies do not show specialised hunting in unit II levels according 771 to taxa or age class. Carcasses were intensely exploited for meat and especially for bone 772 marrow with the high use of fire for various burning degree. These data tend to indicate a 773 consumption of available faunal resources around the cave, supporting by leporid (Morin et 774 al, 2019) and marine shell exploitation (Cataliotti-Valdina in Boutié, 1990; Celiberti & 775 Moigne in Saos et al., 2016).

776

| Hypothesis<br>of sites function                   | Long-term<br>Residential camps   | Short-term<br>regular hunting camps   |
|---|--|---|
| Agents of bone<br>accumulations                   | Accumulation of herbivores is mainly due to hominin activity and carnivore tooth-marks are always less than 4% of the total remains.   | Butchery and carnivore tooth-marks are found on<br>the same species: mainly red deer and horse.<br>Coprolite levels in <b>S13a</b> and <b>S12a</b>  |
| Animal spectrum                                   | -  | Main species < 50% NISP,<br>with various secondary species  |
| Seasonality                                       | All year round (especially in <b>S11</b> and <b>S10</b> ), with most slaughters occurring during the autumn and spring.  | Varied acquisition strategies according to the taxa.  |
| Acquisition<br>strategies for<br>animal resources | Slaughtering is mainly made up of young animals<br>and adults, may be Killed in herds and corresponding to<br>females and their young (most profitable prey)                 | -   |
| Carcass transport<br>strategies                   | -  | Transport strategies and access to carcasses vary<br>according to the taxa.<br>- Main ungulates: whole<br>- Secondary ungulates: partial with poorest pieces<br>(essentially skull and limb ends)   |
| Butchery  | Intensive and generalised butchery sequence for consumption (meat and bone marrow)   | More complete butchery sequence for big and medium size ungulates and scarce human occurrences for poorest pieces of some others (carnivores share ?)   |
| Fire  | Numerous heated (low degree) and burnt bones (high degrees) with burnt lithic artefacts.   | -   |
| Bone industry                                     | -  | Scarce retouchers   |
| Origin of the lithic<br>raw material              | Mainly semi-local, between 7 and 17 km   | Punctual arrival of jasper from 35 km away  |
| Chaîne opératoire<br>according to raw<br>material | -Complete for flint, partial for other raw material<br>-Ramification process on flint  | partial reduction sequence for secondary raw material   |
| % of tools  | Low  | -   |
| Lithic remains<br>density                         | high   | -   |
| Specialization<br>degree of lithic<br>production  | Low  | -   |
| Instance for<br>Middle Palaeolithic<br>sites      | Moula (XV, XIV), Abri du Maras, Saint-Marcel,<br>Baume Néron, Baume Vallée, Mandrin (Daujeard &<br>Moncel, 2010, Daujeard <i>et al.</i> , in Djindjian <i>et al.</i> , 2018) | Orgnac 3, Saint-Anne, Payre, Baume Flandin, Baume<br>de Gigny, Figuier, Baume Moula VIII, IV (Daujeard <i>et al.</i> ,<br>in Djindjian <i>et al.</i> , 2018)<br>Tournal (Magniez, 2010) Hortus (Lebègue <i>et al.</i> ,<br>2010), Crouzade (Saos <i>et al</i> , in press) |

*Tab. 11:* Zooarchaeological and lithic criteria to define types of occupation in unit II from Ramandils with instance of Middle Palaeolithic site in Mediterranean coast and the Rhône Valley (Daujeard & Moncel, 2010; Daujeard *et al.*, in Djindjian *et al.*, 2018; Lebègue *et al.*, 2010; Magniez, 2010; Saos *et al.*, 2016, in press).

Our attempt with this paper was to present the first results of zooarchaeological studies in unit II, subdivided by an archaeostratigraphical approach, into several levels. These spatial analyses allowed us, for the first time, to individualize coprolite beds, produced by carnivores, between human occupation levels, permitting to indicate recurrent periods when Neanderthals did not occupy the site and then allow us to better identify periods of anthropic settlements.

A rich archaeological material is highlighted in three archaeostratigraphic levels with numerous lithic industries and high proportion of burnt bones. Periods of continuing human presence have therefore influenced the taxonomic profile, reflecting planned exploitation of seasonally available resources, as well as more opportunistic behaviours represented by the procurement of mainly semi local raw materials within a 20 km radius territory. This apparent opportunism could be explained by the fact that the tertiary flint, despite an ordinary quality, had to perfectly correspond to the needs of the hominins.

793 The intensive human activities on these anthropic levels (tab. 11), could indicate long term 794 occupations all year round long with slaughtering of most profitable prey, various butchery 795 steps and numerous lithic artefacts, burnt flint and bones with high burning degree. 796 Nevertheless, several features tend to represent recurrent short-term occupations (tab. 11) with 797 various prey slaughtered, seasonal hunting according to taxa, a differential degree in carcass 798 treatment according to the species, the partial reduction sequences for 9 lithic raw materials. 799 Despite the rare occurrence of carnivore marks, less than 4% of total NR, tooth marks affect 800 the same species which are exploited by humans, inducing prompt scavenging during human 801 abandonment of the cave. Moreover, coprolite deposits within and between unit II levels, on 802 well-defined areas, highlighted by archaeostratigraphical studies, might likewise indicate 803 periods when human groups were not occupying the cave.

This multidisciplinary approach permits to highlight the difficulty, for the unit II level of Ramandils cave, of deciding between two hypotheses of function site, related to the duration of occupation and the activities carried out. Indeed, arguments in favour of the two patterns were identified and others seems to be necessary to conclude.

808 To go further, raw material and lithic artefacts studies, performed first on the whole unit II, 809 without fine subdivisions, need to be refined according to the archaeostratigraphical levels, 810 with the aim to define, for each anthropic occupation, the exploited territory and the 811 functional or technical objectives of the settlement. However, a mobility pattern is highlighted 812 by this study indicating an economic organization within a 35 km radius territory of littoral 813 plain, where Ramandils site take probably a central place on Mediterranean coast according to 814 the high human presence recorded on the site and the economic and subsistence modalities 815 observed.

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