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1 2 3	INFLUENCE OF POROSITY ON LIPID PRESERVATION INTO THE WALL OF ARCHAEOLOGICAL POTTERY
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19 20	
20 21	ABSTRACT
22	ADSTRACT
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24	Porosity of archaeological pottery is a key parameter to assess its ability to trap lipids during the use
25	of the pot and to preserve them overtime. Mercury intrusion porosimetry and gas chromatography
26	were used to study the distribution of porosity and the preservation of lipids in different chrono-
27	cultural contexts. The data obtained show that the porosity pattern, related to the raw materials and
28	the <i>savoir-faire</i> of the potters, influences the amount of lipids accumulated in the pottery. A
29	significant overall porosity together with high level of small pores is generally favourable to the
30	preservation of lipids, but variations related to the environmental context are observed.
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33	KEYWORDS
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36	Organic residue analysis; Mercury intrusion porosimetry; Organic matter degradation; Pottery use
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39	INTRODUCTION
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42	Organic molecules have proven to be absorbed inside the walls of unglazed pottery during their use
43	and to be preserved over millennia in archaeological vessels (Charters et al., 1997; Evershed, 2008;
44	Evershed et al., 1995; Regert, 2011). Thanks to the development of analytical techniques such as gas
45	chromatography, mass spectrometry and isotopic ratio mass spectrometry, a wide diversity of

- 46 natural substances was detected in archaeological pottery. The study of these organic residues
- 47 provided valuable data to understand past communities and their changes over time by investigating
- 48 pottery function (Drieu et al., 2018; Fanti et al., 2018; Salque et al., 2013) and the exploitation of
- 49 natural resources (e.g. Craig et al., 2011; Debono Spiteri et al., 2016; Evershed et al., 2008b).
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51 52 The porous nature of unglazed ceramics is known to be one of the key parameters allowing both 53 absorption and preservation of organic products inside pottery walls (Evershed et al., 2008a; 54 Shepard, 1956, p. 126). The absorption of liquid or semi-liquid substances inside the pores of 55 unglazed pottery is easily understood, and the analysis of non-porous vessels generally yields poor 56 results (Evershed, 2008), except if micro-cracks exist on pottery surface (Pecci et al., 2015, 2016). The 57 absorption of organic molecules inside the porous ceramic matrix is thought to be an essential factor of their preservation over time (Eglinton et al., 1991; Heron and Evershed, 1993). Indeed, when 58 59 trapped into the pores, the organic molecules are protected from the surrounding sediment, in an 60 environment generally less favourable to the growth of microorganisms (pH conditions, availability of 61 oxygen, water, and nutriments; Aillaud, 2001, pp. 4-5; Evershed, 1993, 2008). Furthermore, part of 62 the organic compounds is located in small-size pores that strongly limit their access for fungi and 63 microorganisms (Evershed, 1993, 2008; Heron et al., 1991; Heron and Evershed, 1993). Thus, 64 porosity characteristics are thought to play an important role in both the absorption and

- 65 preservation of organic molecules within the pottery walls.
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68 Porosity of ceramic vessels has been investigated for studying pottery manufacture and firing 69 (Carvalho et al., 2006; Gomart et al., 2017; Maniatis and Tsirtsoni, 2002; Morariu et al., 1977; 70 Sanders, 1973; Sobott et al., 2014; Thér, 2016; Volzone and Zagorodny, 2014), physical properties 71 related to pottery function (Lapp, 2012; Moraru and Szendrei, 2010; Rice, 1987, pp. 350–354), and 72 resistance to degradation during burial (Bronitsky, 1986, pp. 225–231). However, although several 73 authors mention relationships between porosity and organic products trapped into pottery walls, 74 only few have investigated porosity in this view (Correa-Ascencio and Evershed, 2013; Matlova et al., 75 2017; Namdar et al., 2009) and none have specifically studied the properties of absorption and 76 preservation of organic molecules detected in archaeological pottery using porosity analysis. 77 From a methodological perspective, porosity of archaeological pottery has been studied by producing 78 2D images using Scanning Electron Microscope (SEM; Carvalho et al., 2006; Correa-Ascencio and 79 Evershed, 2013; Maniatis and Tsirtsoni, 2002; Moraru and Szendrei, 2010) or 3D images by micro 80 tomography (Gomart et al., 2017; Kulkova and Kulkov, 2016; Sobott et al., 2014). Both techniques 81 have the advantage of being non-destructive, but quantitative study of the distribution of porosity 82 requires specific data processing (image analysing software or mathematical treatments of the 83 tomography data; Adrian De la Fuente and Vera, 2015; Aprile et al., 2014; Reedy et al., 2014, Thér, 84 2016). Mercury Intrusion Porosimetry (MIP), a technique specifically adapted to the quantitation of 85 the porosity that allows both the measurement of the total volume of pores and an estimation of the 86 pores distribution depending on their diameter (Giesche, 2006; Sobott et al., 2014), was used to 87 study the porosity of archaeological ceramics (Matlova et al., 2017; Morariu et al., 1977; Namdar et al., 2009; Sanders, 1973; Sobott et al., 2014; Volzone and Zagorodny, 2014), but up to now, the 88 89 question of the influence of porosity on the preservation of organic molecules has not yet been 90 considered in this way

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93 94	With mercury intrusion porosimetry, the present study investigates the relationship between the amount of lipids preserved in archaeological potsherds and the porosity of the ceramic matrix
95	(quantity of pores and distribution). This issue is addressed by studying the porosity of potsherds
96	taken from three different chrono-cultural contexts and having preserved various quantities of lipids
97	as determined by gas chromatography analysis. By improving our understanding of the impact of
98	porosity on the amount of lipids extracted from archaeological sherds, this work aims, in the long
99	term, to (i) determine if it is possible to build a selection grid for optimising sampling of
100	archaeological pottery for lipid analysis; and (ii) better understand lipid absorption and preservation
101	in pottery walls for in-depth interpretation of pottery use.
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104	MATERIALS AND METHODS
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107	Archaeological potsherds
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110	Three archaeological sites, having different chronological and geographical locations, were chosen to
111	represent different contexts of preservation which were either more, or less favourable to
112	biochemical (anoxic vs. oxic conditions, basic vs. acid sediment) and physical degradation (various
113	degrees of waterflow, wetlands).
114 115	The Pendimoun rock shelter was chosen to represent a context favourable to biochemical degradation processes, with oxic conditions of burial and basic sediments related to the calcareous
115	geological environment. In this settlement, archaeological artefacts were also exposed to intensive
117	leaching, due to the location of the site at a stratification joint between Jurassic limestones and
118	Cretaceous marl-limestone (Binder et al., 1993). This site, situated in the south-east of France near
119	the Italian border (Castellar, France), revealed archaeological remains from the Epipaleolithic to the
120	Middle Ages (Binder et al., 1993). Thirteen samples dating from the Early Neolithic period (<i>Impressa</i>
121	ware, 5720 to 5470 BCE; Binder et al., 2017) were selected for analysis. Pottery from this period was
122	recovered in high amounts and is made of three main fabrics: granitoid, glauconitic and mixed
123	granitoid and glauconitic earths (Binder & Sénépart, 2010; Gabriele, 2014, pp. 223–225). Due to the
124	heterogeneity and complexity of mixed earths, only granitoids and glauconitic potsherds were
125	sampled for the present study.
126	The site of Cuciurpula was selected because of its acidic sediment, unfavourable to microorganism
127	activity (DeLaune et al., 1981; Drieu et al., 2018; Moucawi et al., 1981). Twelve potsherds were
128	sampled from this open-air site dating from the end of the Bronze Age to the first half of the Iron Age
129	and situated in the mountainous region of central south Corsica (Serra-di-Scopamena and
130	Sorbollano, France; Peche-Quilichini et al., 2015). In-depth analysis of the ceramic fabric has not yet
131	been carried out but most of the pottery seems to be made of local granitic earth with high quantity
132	of non-plastic inclusions of variable grain size (Peche-Quilichini, 2010).
133	Clairvaux XIV was selected for its wet and anoxic environment, very unfavourable to biochemical
134	degradation of lipids (Den Dooren De Jong et al., 1961; Eglinton et al., 1991). This site is one of a
135	number of settlements surrounding the Lake of Clairvaux (Jura, France) ranging from the Neolithic to

136 137 138 139	the Bronze Age (Pétrequin and Pétrequin, 2016a) and is dated to the first half of the 4th millennium BC (Pétrequin, 2016). For the present study, we selected thirteen potsherds, made of marl clay tempered with ground limestone or calcite (Pétrequin and Pétrequin, 2016b).
140 141 142 143 144 145	To limit excessive destruction of archaeological artefacts, a single fragment of pottery (between 2.5 and 4 g) was sampled on each ceramic vessel investigated. Unspecific samples (non-refitting sherds, absence of decoration or chrono-cultural markers) were preferentially selected. Each sample was then divided into two subsamples for lipid and mercury intrusion porosimetry analyses.
146 147 148 149	Lipids extraction and analysis
150 151	Sample treatment and analysis were carried out following Evershed et al. (1990) with slight modifications. When visible residues existed on the surface of potsherds, they were sampled using a
152 153	clean scalpel blade. Following the vast majority of the literature on lipid residue analysis, the surface of the potsherds (around 1 mm) was then removed using a clean scalpel blade to eliminate any
154 155	exogenous lipid. Around 2 g of ceramic sherd and between 20 and 100 μ g of visible residue were then crushed with a mortar and pestle and 20 μ L of internal standard was added for quantitation (<i>n</i> -
156	tetratriacontane, 1 mg/mL in <i>n</i> -hexane). Lipids were extracted with 10 mL of
157 158	dichloromethane/methanol (DCM:MeOH; 2:1, v/v) by sonication (2 × 15 min). After centrifugation, the supernatant was evaporated to dryness under a gentle stream of nitrogen and dissolved in
158	500 μ L of DCM/MeOH to obtain the total lipid extract (TLE). An aliquot of the TLE (100 μ L) was
160	evaporated to dryness and treated with 50 μ L of N,O-bis(trimethylsilyl)trifluoroacetamide containing
161	1% trimethylchlorosilane (BSTFA; 70°C, 1h). The excess BSTFA was evaporated under nitrogen and
162	the sample recovered in 25 to 2000 μL of cyclohexane (depending on the amount of lipid matter
163	preserved) for high temperature gas chromatography analysis (HT GC).
164	1 μ L of sample was introduced into an Agilent Technologies 7890A gas chromatograph, via an on-
165	column injector. The analysis was performed using a 15 m x 0.32 mm i.d. fused silica capillary column
166	(DB5-MS, 0.1 μ L film thickness, Agilent J&W), and helium as carrier gas. The temperature programme
167	consisted of an increase from 50°C to 100°C at 15°C.min ⁻¹ , and then from 100°C to 375°C at
168	10°C.min ⁻¹ .
169	Gas-chromatography mass-spectrometry was used to determine the structure of the molecular
170	compounds, in particular to distinguish between molecules originating from the pottery use and
171	modern contaminants. These analyses were performed on a Shimadzu GC 2010 PLUS chromatograph
172	coupled to a Shimadzu QP 2010 ULTRA mass spectrometer. An aliquot of sample (1 μ L) was injected
173 174	on a high temperature non-polar column (DB5-HT, 15 m x 0.322 mm i.d., 0.1 μm film thickness, Agilent J&W) via a splitless injector. The GC temperature programme was as follows: 1 min
175	isothermal hold at 50°C followed by an increase to 100°C at 15°C.min ⁻¹ , then to 240°C at 10°C.min ⁻¹
175	and to 380°C.min ⁻¹ and a final isothermal hold for 7 min. The GC-MS interface temperature was
177	maintained at 300°C. The mass spectrometer was used in electron ionization mode (EI, 70 eV) and
178	mass spectra were acquired over the range m/z 50–950.

The lipid concentration was calculated by summing the area of all the peaks and comparing it withthe area of the internal standard peak. Modern contaminants, such as phthalates (easily identified by

- 181 the characteristic m/z 149 fragment), were excluded from the overall calculation.
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183 184 This study is part of a wider project of lipid analysis, in which large sets of samples were analysed for 185 each site (Drieu et al., 2018, submitted). For the present study a subset of samples was selected from 186 each of these assemblages to represent the complete range of lipid preservation inside the ceramic 187 walls at each site: the samples with the best preservation of lipids of each site were selected and 188 compared with samples with medium and very low preservation (Table 1). Potsherds from Clairvaux 189 XIV and Cuciurpula originate from a large diversity of ceramic types: cooking pots, serving and storing 190 vessels. The functional study of the pots, based on their content, shape and traces of use have been 191 published elsewhere (Drieu et al., 2018, submitted). Most of the samples from the site of Pendimoun 192 originate from pots of unknown function, especially because it was not possible to reconstruct the 193 form, and they did not display any traces of use (carbonised residues, traces of soot, etc.). 194 195 196 Table 1: List of samples analysed during the study and lipid yield. * two distinct residues removed 197 from the internal and external surfaces of the pot. 198 199 200 Mercury Intrusion Porosimetry 201

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203 Mercury intrusion porosimetry analyses were performed using microporosimeter Autopore IV 204 (Micromeritics). The pressure range of the technique was from sub ambient up to 400 MPa, in order 205 to cover a pore diameter range from approximately 360 µmm to 3 nm. Archaeological samples were 206 broken into fragments having a small volume between 0.5 and 1 cm³ (0.30 to 1.75 g), dried overnight 207 in an oven at 45 °C and introduced alone or in pairs in the analysis cell. A contact angle of 130° was 208 used for the calculation of the entrance diameter of pores, a value already used for mercury intrusion

- 209 porosimetry on archaeological samples (Sanders, 1973).
- 210 Considering equation (1) where P is the apply pressure; γ is the surface tension of the liquid
- 211 (mercury); θ is the contact angle between the liquid and the substrate; we can determine r, which is
- the radius of the pores.
- 213 $r = (-2 \gamma \cos \theta) / P$ (1)
- 214 This method of characterisation is often used to study the porosity of different types of materials.
- However, some assumptions need to be considered before discussing the results, such as the aging
- of the samples, the presence of mineral post-deposits, but also the influences of the pressure on the
- 217 microstructure, the presence of residual air or humidity into the pores, the use of a cylindrical pore
- 218 model, etc. (as well explained by A.J. Klemm [2009]). In this study, the preparation of the samples
- 219 was designed to remove most of the remaining humidity without inducing any damage in the
- 220 microstructure, by gently drying the samples. The analytical conditions were tested on modern
- 221 pieces of ceramics in order to rule out the risk of damages due to the pressure. The presence of
- visible carbonated deposits on some sherds from the site of Pendimoun is not correlated with a
- reduced porosity, suggesting that they have limited impact on the post-depositional evolution of the

224	porosity, contrarily to was stated in the literature (Bronitsky, 1986, pp. 228; Rice, 1987, pp. 353).
225	Finally, these analyses are semi-quantitative due to the use of the cylindrical pore model and the
226	presence of potential residual air or humidity. However, they provide useful comparative results
227	because they were carried out on similar ceramic materials and in the exact same analytical
228	conditions.
229	It is noteworthy to emphasize that this protocol does not specifically measure the porosity of the
230	internal surface of the pot. This is because mercury not only enters the samples through the pores of
231	the internal and external surface of the pot, but also enters through the fractured edges. The effect
232	of surfaces treatments, such as burnishing and the application of a slip is thus not measured. As
233	these treatments heterogeneously affect the surface properties of the pottery, we think that a
234	specific research project should be entirely dedicated to the study of the influence of the surface
235	properties of the samples. Consequently, as a first approach of the topic, we decided to restrict the
236	scope of this paper to the study of the influence of the core properties (microstructure) of the
237	ceramics.
238	To evaluate the ability of microorganisms to penetrate the pottery walls, the data of Total Intrusion
239	Volume were subdivided for each sample. Two types of pores were distinguished based on their
240	diameter: pores of diameter larger than 1 μ m, accessible to microorganisms and pores with smaller
241	diameters («small pores»), where microorganisms cannot penetrate due to their size (generally in
242	the range of the micrometer; Davis et al., 1990, p. 21).
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245	RESULTS
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248	Porosity analysis
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251	Figure 1 details the results obtained on the pore distributions of the samples. The percentage of
252	overall porosity ranges from 14 to 36% of the volume of the sherd, and the percentage of the total
253	intrusion volume relative to «small pores» (diameters < 1 μ m) ranges from 23 to 78% of the overall
254	porosity (Table 2).
255	
256 257	The samples from Pendimoun are distributed in three different porosity groups (Table 2). Samples
258	from the first group (AP1; LD10700a, LD10702a, LD10703a, LD10716a, LD10718a, LD10720a and
258	LD10723) are characterised by:
260	- an overall porosity between 22 and 25%,
261	- between 48 and 67% of pores having a diameter < 1 μ m («small pores»),
262	- three main types of pores having a diameter around 10 μ m, around 4 μ m and between 0.03
263	and 0.15 μ m (Figure 1a),
264	- less than 7 μ g.g ⁻¹ of lipids preserved.
265	A second group (AP2; LD10711, LD10724, and LD10731) which included three samples made of
266	granitoid paste with:
267	- much smaller overall porosity, between 16 and 19%,

268	- a smaller quantity of «small pores» having a diameter < 1 μ m (44 to 45%),
269	- one main pore type with a diameter between 0.5 and 4 μ m (Figure 1b),
270	 between 1.2 and 34.1 μg.g⁻¹ of lipids preserved.
271	The third group (AP3; LD10710, LD10713, and LD10714; Figure 1c) comprised samples with:
272	 a small percentage of overall porosity (14 to 20%),
273	 between 23 to 40% of «small pores» having a diameter < 1 μm,
274	- two main types of pores having a diameter around 4 μm and around 10 μm,
275	 less than 6.6 μg.g⁻¹ of lipids preserved.
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278	Figure 1: Differential distribution curves measured by mercury intrusion porosimetry for the
279	archaeological potsherds. a) Group AP1; b) Group AP2; c) Group AP3; d) Group Cuci1; e) Group Cuci2;
280	f) Group Cuci3; g) Group Cl1; h) Group Cl2. Dotted lines indicate the main types of pores discussed in
281	the text.
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284	The percentage of overall porosity is more homogeneous for the samples from Cuciurpula (21.8 to
285	28.7%). Three groups of samples can nonetheless be distinguished based on the distribution of the
286	pore diameters (Figure 1d, e, f). The first group (Cuci1; LD10659, LD10665, and MR2707) is defined
287	by samples having:
288	- between 49 and 45% of «small pores» with diameters < 1 μ m
289	- two main types of pores with diameter of around 10 μm and 4 μm, and a smaller number of
290	pores between 0.1 and 1 μm (Figure 1d)
291	- low preservation of lipids (0.94 to 15.1 μg.g ⁻¹ of lipids).
292	The second group (Cuci2; MR2696, MR2698, MR2711, and MR2717) is characterised by:
293	 a percentage of «small pores» higher than the group Cuci1 (55 to 57%)
294	- three main types of pores with diameters close to 10 μ m, 4 μ m and between 0.01 and 0.1 μ m
295	(Figure 1e).
296	- overall low preservation of lipids except one sample that yielded 316.4 μg.g ⁻¹ of lipids.
297	A third group (Cuci3; LD10656, LD10667, LD10676, MR2699 and MR2701) includes samples
298	characterised by:
299	- between 46 and 51% of «small pores» with diameters < 1 μ m
300	 homogeneous porosity profile without clear peaks (Figure 1f)
301	- diverse preservation of lipids: two samples yielded respectively 0.2 and 25.0 µg.g ⁻¹ while the
302	remaining three preserved between 153.3 and 856.2 μg.g ⁻¹ of lipids.
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305	Samples from Clairvaux XIV were divided in two groups, mainly defined by their percentage of
306	porosity (Figure 1g, h). The first group (Cl1; LD10100, LD10101, LD10103, LD10114, LD10126, and
307	LD10133) comprised samples having:
308	- overall porosity between 25 and 36%
309	- less than 65% of «small pores»
310	- two main types of pores with diameters between 6 and 200 μ m and between 0.02 and 1 μ m
311	(Figure 1g).
312	 less than 22 μg.g⁻¹ of lipids preserved.

313	The second group is composed of the seven samples (Cl2; LD10104, LD10113, LD10117, LD10125,
314	LD10129, LD10130, and LD10131) displaying:
315	 overall porosity between 23 and 30%
316	 large quantity of «small pores» (more than 65%).
317	- one main type of pores with a diameter between 0.02 and 1 μ m (Figure 1h)
318	- excellent preservation of lipids (between 95.1 and 901.9 μg.g ⁻¹ , except for LD10104, yielding
319	3.5 μg.g ⁻¹ of lipids.
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321	
322	Table 2: Table summarising the results obtained by lipid and mercury intrusion porosimetry analyses
323	for all groups analysed.
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326	Comparison between lipids preservation and porosity
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329	Figure 2: Lipid concentration (TLE, μ g.g ⁻¹) versus overall porosity (a) and «small pores» (diameters < 1
330	μm) (b).
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333	All the samples preserving lipids have specific porosity values (between 18 and 30% of overall
334	porosity and 43 and 78% of «small pores» of diameters < 1 μ m; Figure 2), potentially suggesting a
335	range of % overall porosity and % of pores of «small pores» (diameters < 1 μ m) that favours the
336	preservation of the lipids absorbed into the ceramic matrix.
337	Particularly, the quantity of «small pores» (diameters < 1 μ m) seems to be related to lipid
338	preservation (Figure 3). This relationship is notably seen for extreme cases: for example only one of
339	the eight samples displaying less than 45% of «small pores» with diameters < 1 μ m yielded more
340	than 20 μg.g ⁻¹ of lipids (Figure 2b). On the contrary, among the six samples with more than 70% of
341	«small pores» of diameter < 1 μ m, five preserved lipids (Figure 2b). The samples having significant
342	percentage of overall porosity (more than 26%; LD10100, LD10103, LD10114, LD10133, LD10659,
343	MR2707, LD10665, LD10126), seem to better preserve lipids when their porosity is mostly made up
344	of «small pores» (more than 65% of pores of diameter < 1 μ m).
345	The near absence of lipids in some samples which have a large quantity of «small pores» (LD10104,
346	LD10700, LD10702, LD10703, LD10716, LD10718, LD10720, LD10723, MR2698, MR2699, MR2711,
347	MR2717) could be related to other parameters such as chemical degradation or specific use of the
348	vessels for non-fatty substances. This latter hypothesis is confirmed for 4 samples from Cuciurpula
349	(MR2698, MR2699, MR2711, and MR2717) bearing visible surface residues that yielded very low
350	amounts of lipids (between 3 and 84 μ g.g ⁻¹ , when other visible residues at the site contain several
351	hundred μ g.g ⁻¹ ; Table 1 and Drieu et al., 2018).
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354	Figure 3: Degree of preservation of lipids depending on «small pores» < 1 μm. Samples with TLE > 20
355	μ g.g ⁻¹ are considered as well preserved and those with TLE < 20 μ g. g ⁻¹ are considered as badly
356	preserved. The percentage indicated corresponds to the % of well-preserved samples for each

357 category of size distribution of «small pores» < 1 μ m.

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359 360	The impact of «small pores» is mainly seen in potsherds from Pendimoun and Clairvaux XIV (Figure
361	2b), as they display extreme cases of porosity: a very low quantity of «small pores» in Pendimoun
362	and a very high quantity in Clairvaux XIV. At Pendimoun, samples with less than 40% of «small pores»
363	(LD10713 and LD10714) did not preserved any lipids, while the only sample yielding more than 20
364	μ g.g ⁻¹ of lipids displays such pores, even in small quantities (LD10731, 44% of pores < 1 μ m). Samples
365	from Clairvaux XIV with both a high quantity of «small pores» and absence of pores of large diameter
366	(more than 10 μ m; Group Cl2) seem to preserve the highest quantity of lipids: 85% of samples
367	yielding more than 20 μ g.g $^{-1}$ are also characterised by more than 65% of «small pores» with
368	diameters smaller than 1 μ m. At Cuciurpula, the porosity range seems to have a lesser impact on the
369	preservation of lipids into pottery walls, as lipids are better preserved inside samples with a very
370	homogenous distribution of pores (Group Cuci3).
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373	DISCUSSION
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376	Diversity of the porosity pattern and making of the pottery
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379	Pottery from the three sites investigated presents various patterns of porosity (Figure 4): sherds from
380	Clairvaux XIV are the most porous (mean value of overall porosity of 28%) with the highest level of
381	«small pores» < 1 μ m (mean value of «small pores» of 66.%) while the opposite if observed for the
382	site of Pendimoun (mean value of overall porosity of 25%; mean value of «small pores» of 50.%). The
383	site of Cuciurpula presents intermediate values (mean value of overall porosity of 21%; mean value
384	of «small pores» of 66%). This diversity reflects different manufacturing chaînes opératoires between
385	sites: differences in terms of the nature of the clay, quantity, nature and granulometry of non-plastic
386	inclusions, intensity of handling, and firing conditions (Bronitsky, 1986, p. 226; Rice, 1987, p. 351;
387	Rye, 1981, pp. 40 and 122; Shepard, 1956, p. 126; Skibo, 2013, p. 40). For example, at Pendimoun,
388	the small percentage of overall porosity of some samples (groups AP2 and AP3) and the absence of
389	very small pores (less than 0.1 μ m) are to be compared with petrographic information. Sherds
390	investigated are all made of granitoid earth and comprised very large quantities of non-plastic
391	inclusions and a very small amount of clay matrix (Binder & Sénépart, 2010). This specificity of the
392 393	ceramic paste could explain the absence of very small pores, that are probably related to clay structures and cannot exist if no clay fine fraction is present. At Clairvaux, the porosity seems to be
394 395	related to the level of <i>savoir-faire</i> (Pétrequin and Pétrequin, 2016b): the best made ceramics generally have low overall porosity and high amount of «small pores».
395 396	generally have low overall porosity and high amount of «small pores».
390 397	
398	Figure 4: Overall porosity versus «small pores» (diameter < 1 μ m) depending on the sites
398 399	investigated.
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401	Relationship between porosity and preservation of lipids
	Relationship between porosity and preservation of lipids

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- Given the diversity of environmental contexts considered in this study, the diameter of the poresappears to impact different types of lipids degradation processes.
- 407 First, our results confirm that the presence of «small pores» limits the degradation of lipids by
- 408 microorganisms, as supposed in the literature. Generally larger than 1 μ m, (Davis et al. 1990, p. 21)
- 409 most of bacteria and fungi are not able to enter pores smaller than this size. Thus, the lipids within
- the ceramic matrix are protected from microbial degradation (Evershed, 1993, 2008; Heron et al.,
- 411 1991; Heron and Evershed, 1993; Matlova et al., 2017; Saboyainsta and Maubois, 2000). This
- 412 property has been exploited in food science for microfiltration: porous ceramic fabrics with pore
- diameters under 1.4 μ m are used as effective milk microfiltration devices, removing up to 99.75% of
- 414 microorganisms responsible for the biochemical degradation of milk (Saboyainsta and Maubois,
- 415 2000; Trouvé et al., 1991). The impact of porosity on the degradation of lipids by biochemical
- degradation was probably important at the site of Pendimoun, but limited at Clairvaux XIV and
- 417 Cuciurpula respectively due to the anoxic context of Clairvaux and the acidic sediments of Cuciurpula
- 418 (Drieu et al., 2018, submitted) considerably reducing microbial activity.
- 419 Other degradation processes therefore need to be considered to explain the impact of porosity on
- 420 the preservation of lipids at Clairvaux. In this wet environment, reduced transport of water through
- 421 porosity is more likely to explain good preservation of lipids in «small pores». Following Poiseuille
- 422 equation, for a laminar flow of water, the smaller the radius of the pore the slower the flow rates
- 423 through it. In «small pores», the slowing of the water flow probably limits the leaching of absorbed
- 424 lipids, and possibly reduces hydrolysis. A similar hypothesis has also been suggested to explain the
- 425 poor preservation of chlorine in Late Neolithic salt moulds (Ard and Weller, 2012). This type of
- 426 degradation process may have also occurred at Pendimoun. Indeed, despite the limited rainfalls of
- 427 the Mediterranean climate, this rock shelter is exposed to a significant water flow related to its
- 428 geological location, as evidenced by the substantial calcium carbonate deposits on the potsherds
- from the site (Binder et al., 1993). At Cuciurpula, leaching is probably limited as the site is open-air,
- 430 without rock walls favouring water flow.
- 431
- 432

The effect of other parameters on lipid degradation in archaeological samples should also beexamined in the future.

- 435 First, the impact of pottery surface treatment (slipping, burnishing) on the absorption and the
- 436 degradation of organic matter has never been studied in-depth, to the best of our knowledge (Reber,
- 437 2007). However, this type of treatment considerably modifies the surface of the pots by promoting
- 438 smaller pores at the surface of the walls, probably affecting both the preservation of lipids and the
- 439 amount absorbed during pottery use (Correa-Ascencio and Evershed, 2013; Reber, 2007). A protocol
- 440 for coating the edges of potsherds with polymer resin is under development to specifically measure
- the porosity of the internal surface of the vessels. This investigation will be particularly useful to
- better understand some samples from Pendimoun that display evidence of deep burnishing (Binder
- 443 et al., 1993; Binder & Maggi, 2001), not reflected in the core porosity measured in the present study.
- 444 Secondly, hydrolysis is not the only chemical reaction responsible for the degradation of lipids in
- 445 archaeological samples: radical oxidation, photo-oxidation, and reduction also affect the lipid content
- 446 (Aillaud, 2001; Heron and Evershed, 1993) and porosity has probably no impact on it.

Finally, the actual use of pottery is a key parameter influencing the amount of lipids absorbed andpreserved in archaeological potsherds. The absence of lipids in numerous samples from Pendimoun

may not be due to their degradation but to specific use of pottery in this early phase of the Neolithic

450 in the western Mediterranean, to transform cereals or to store water, for example, as suggested by

451 other researchers (Spiteri et al., 2017).

- 452 453
- 454 CONCLUSION

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This article has revealed that when potsherds are exposed to normal microbial activity (as at the site of Pendimoun) or to water (at Clairvaux XIV, and to a lesser extent at Pendimoun), the range of porosity inside the pottery walls is of importance for the preservation of lipids. When trapped in «small pores» (less than 1 μ m of diameter), lipids are protected from microorganisms, but also from leaching. A high abundance of «small pores» of diameter < 1 μ m in pottery walls favours thus the preservation of lipids in archaeological samples, by reducing biological (microorganism activity), physical (leaching) and possibly chemical (hydrolysis) degradation.

- In very specific environments when leaching is limited, and microorganism activity reduced, porosity
 appears to have a limited impact on the preservation of lipids. The site of Cuciurpula considered in
- this study meets these criteria, but other types of archaeological contexts, such as arid or frozen
- 467 environments, should be considered and studied in the future.
- These first results need now to be confirmed with new analyses of larger sample sets from a range of
- 469 environmental contexts (acid and basic soils, arid, frozen and submerged environments). A better
- understanding of the preservation of lipids in porous ceramic matrices could lead to more selectivesampling of pottery in the future, avoiding the destruction of archaeological potsherds unlikely to
- 472 have preserved lipids. Secondly, mercury intrusion porosimetry can be used as a new tool for
- 473 functional analysis allowing a better understanding of the process of penetration of organic
- 474 substances into the porous walls of ceramic vessels and the degradation likely to have occurred. In
- 475 particular, this technique can be used to understand the use of ceramic vessels yielding small476 amounts of lipids, such as in the Mediterranean.
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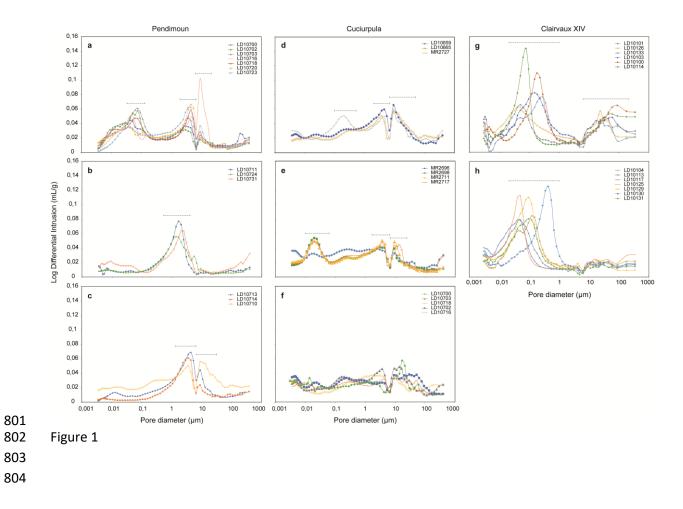
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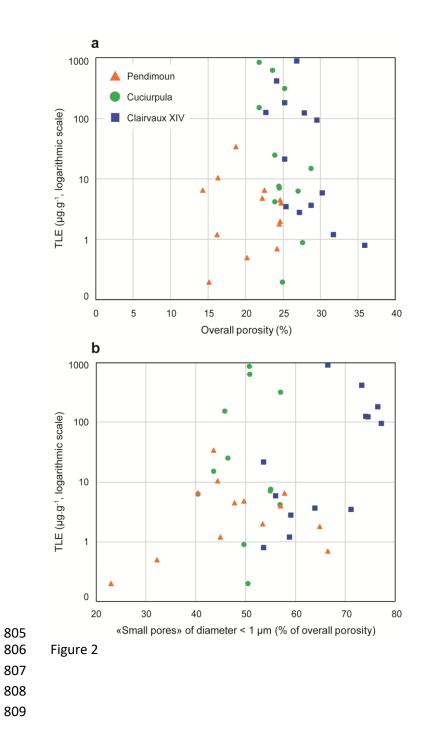
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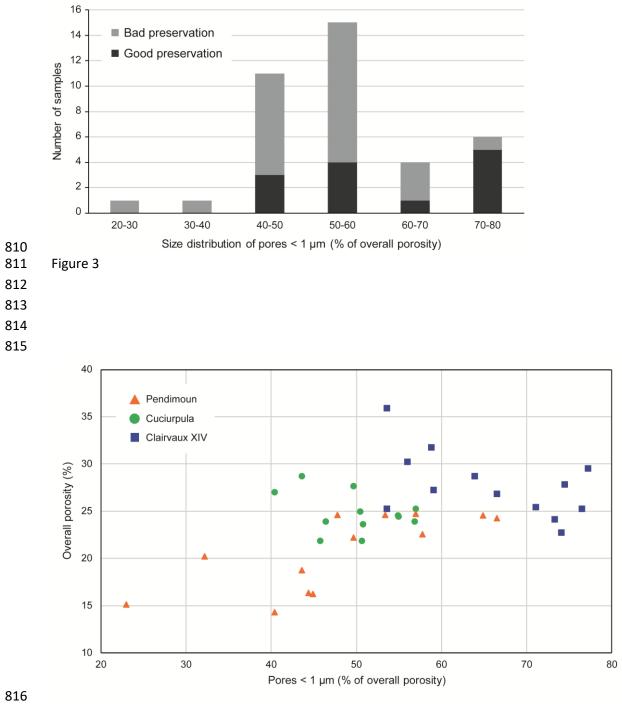
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799	FIGURES
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817 Figure 4