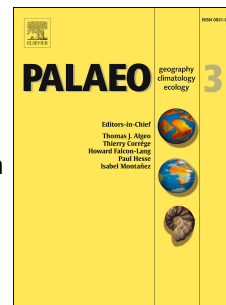


# Journal Pre-proof

A multidisciplinary analysis of shell deposits from Saltés Island (SW Spain): the origin of a new Roman shell midden

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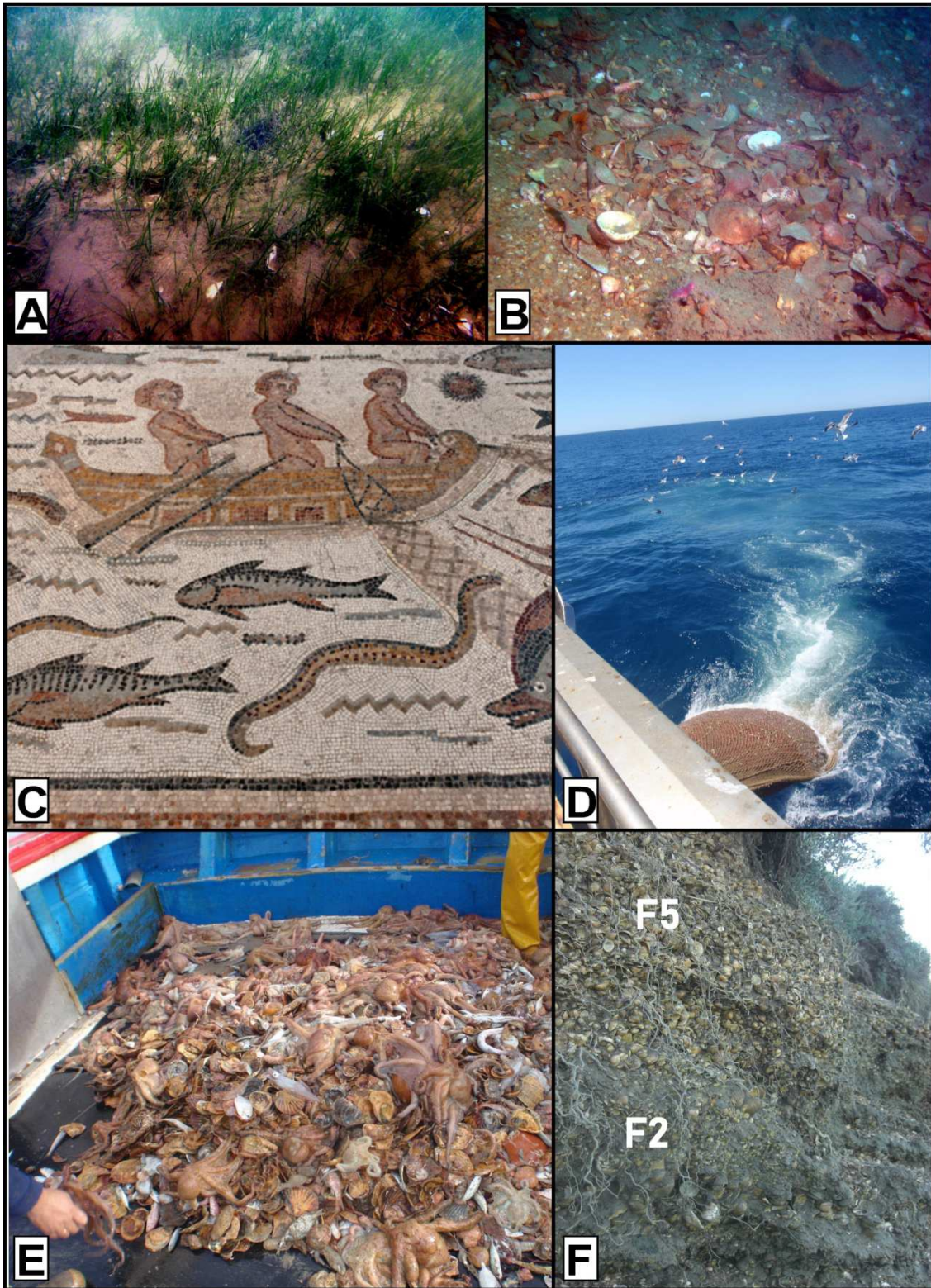
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GRAPHICAL ABSTRACT



1 A multidisciplinary analysis of shell deposits from Saltés Island (SW Spain):  
2 the origin of a new Roman shell midden

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30 ABSTRACT

31 The analysis of shell deposits eroded by a ebb-tide channel on Saltés  
32 Island (Tinto-Odiel estuary, SW Spain) resulted in the identification of a new  
33 shell midden, associated with the activity of a nearby Roman factory over the  
34 4th-5th centuries CE. This midden differs from other old shell deposits (sandy  
35 tidal flats, cheniers, washover fans) in several features: a) its malacological  
36 content, dominated by edible species (mainly the bivalve *Glycymeris nummaria*)  
37 and differentiated by statistical analysis; b) a partial selection and better  
38 conservation of *Glycymeris nummaria* (Linnaeus), its most abundant species; c)  
39 the absence of microfauna, which implies a previous washing to its final deposit;  
40 and d) an age concordant with the one deduced from the Roman amphoraic  
41 remains found in this area and subsequent to the washover fans on whom it was  
42 deposited. All these features, together with the absence of both anthropic  
43 fractures or cooking, would indicate that this Roman shell midden was the end  
44 result of a trawling on subtidal *Glycymeris*-rich sandy bottoms with adjacent  
45 grasslands, where the gastropod *Bittium reticulatum* (da Costa) was the most  
46 abundant mollusc. This gastropod is the dominant species in the remaining shell  
47 deposits.

48

49 *Keywords:* molluscs; taphonomy; statistics; microfauna; archaeology; Roman  
50 period.

51

52 **1. Introduction**

53           The prehistoric and historical evidences of mollusc consumption are very  
54 abundant and started as far back as the Lower Palaeolithic (ca. 300 ka; Colonese  
55 et al., 2011). One of the most widespread results are coastal shell middens,  
56 characterized by massive concentrations of mollusc shells that also contain  
57 frequent bone remains, charcoal and ash that denotes collection, hunting and  
58 food processing (e.g. Álvarez et al., 2011; Thompson et al., 2016). These shell  
59 accumulations are present in numerous coastal environments around the world  
60 (see Fig. 1 for examples).

61           The species richness of each coastal shell midden depends on the  
62 disponibility and paleobiogeographical distribution of edible species in a certain  
63 area for a defined period of time. Among bivalves, some of the most commonly  
64 reported groups are oysters (Lewis, 2011; Lulewicz et al., 2017), mussels  
65 (Jerardino et al., 2008; Erlandson et al., 2009a, b), clams (Bailey, 1977; Hallmann  
66 et al., 2009) or cockles (Dupont et al., 2007; López-Dórigal et al., 2019). The  
67 gastropod record includes marine, brackish and even terrestrial species, such  
68 as strombids, muricids or helicids (Schapiera et al., 2006; Douka et al., 2014;  
69 Magee et al., 2017). In an individual shell midden, the most consumed species  
70 may vary with time due to anthropogenic overexploitation, climatic effects, sea-  
71 level oscillations or geomorphological changes (Branch et al., 2014; García-  
72 Escárzaga et al., 2017).

73           In the geological record of these littoral scenarios, shell middens coexist  
74 with natural shell deposits, such as beach ridges, cheniers, washover fans or  
75 bioclastic tidal facies. An important body of research has focused on the  
76 differences between them, from which it can be deduced that middens shells  
77 are mainly characterized by: 1) Texture: absence or very low percentages of  
78 matrix and grain size selection in edible species; 2) Stratigraphy: no evidence for  
79 internal stratification; 3) Fossil record: low diversity, dominant edible species of  
80 macrofauna (e.g. bivalves, gastropods, crustaceans, bones of various groups)

81 and absence or scarcity of microfossils in relation to natural formations; 4)  
82 Taphonomy: dominant disarticulation, selection of adults, acute fractures of  
83 shell margins, rupture of the last turn of the loop or presence of burnt  
84 evidences; 5) Archaeology: presence of hearth stones or artefacts in some cases;  
85 and 6) Chronology: consistent with the human history in most cases. This  
86 general overview should be treated with caution, because numerous problems  
87 have been described in the individual study of each shell midden (Bailey, 1977;  
88 Gill et al., 1991; Attenbrow, 1992; Carter, 1999; Saunders and Russo, 2011; Betts  
89 and Hrynck, 2017 and references therein).

90 In southern Spain, evidence of the mollusc exploitation has been  
91 highlighted at around 150-120 ka during the Last Interglacial and continued  
92 until the Holocene (Fa, 2008; Cortés et al., 2011; Jordá et al., 2011). During the  
93 Roman period (3rd century BCE-5th century CE), numerous halieutic sites (so-  
94 called *cetariae*) were present along the southwestern Atlantic coast of Spain  
95 (Bernal et al., 2014; Campos et al., 2015a). Associated shell middens are still  
96 poorly studied (e.g. Ramos et al., 2011) and require extensive future research.

97

98

## (FIGURE 1)

99

100 This paper analyzes the main faunistic characteristics of the sedimentary  
101 facies present in the northwestern sector of Saltés Island (Fig. 2, A-B: Tinto-Odiel  
102 estuary, SW Spain) to define their main assemblages and to detect possible shell  
103 middens based on a multidisciplinary analysis (macrofauna, microfauna,  
104 taphonomy, statistics, chronology) of them.

105

## 106 2. Regional setting

## 107 2.1. *The Tinto-Odiel estuary*

108 In southwestern Spain, the Tinto and Odiel Rivers make up a large joint  
109 estuary, with two main channels (Fig. 2, B-C: Padre Santo channel and Punta  
110 Umbría channel) and numerous ebb-tide channels that delimit a set of islands  
111 and marshes (Fig. 2, C: Saltés, Bacuta, Enmedio). This geomorphological  
112 architecture is protected by two sandy spits (Fig. 2, B-C: Punta Umbría and  
113 Punta Arenillas). Hydrodynamic processes are controlled by the tidal regime,  
114 waves and, to a lesser extent, the fluvial dynamics. Tidal regime is mesotidal  
115 (mean range: 2.15 m; Borrego et al., 1993) and the dominant waves come from  
116 the southwest (Borrego, 1992). Fluvial inputs are limited during most of the year  
117 due to the scarce flow of the Tinto and Odiel Rivers, but they are very polluted  
118 by heavy metals (Cu, Pb, Zn) owing to millennial mining activities (Nieto et al.,  
119 2007).

## 120 2.2. *Saltés Island*

121 This small island (37°6'28"-37°13'12"N; 6°50'30"-6°58'30"W) has an  
122 elongated morphology in NW-SE direction and includes a set of cheniers (Fig. 2,  
123 C: El Almendral, El Acebuchal, La Cascajera) separated by extensive marshes and  
124 tidal channels (Fig. 2, D: e.g. Estero de Los Difuntos). These cheniers and some  
125 adjacent washover fans have been formed by the action of the equinoxial tides  
126 of spring and autumn and winter storms, with a migration towards the north of  
127 these bioclastic ridges on a previous tidal sandy plain (Morales et al., 2014;  
128 Cáceres et al., 2018).

129

130

**(FIGURE 2)**

131

132 In the last decades, the tidal action of the Estero de los Difuntos ebb-tide  
133 channel has caused the erosion of La Cascajera chenier, the most important of  
134 this island (Fig. 2, D). Several multidisciplinary studies on the materials now  
135 exposed has made it possible to differentiate five main sedimentary facies in  
136 this area (Fig. 3; modified from Cáceres et al., 2018; González-Regalado et al.,  
137 2019a, b), with a relatively well developed overlying edaphic profile.  
138 Synthetically, its main characteristics are the following: a) facies 1 (F1: silty tidal  
139 flat: 0.5-1 m a.s.l.): sandy blue-gray silts, intensely bioturbated by annelids; b)  
140 facies 2 (F2: washover fan: 0.7-2.5 m a.s.l.): alternating medium to coarse  
141 bioclastic sands, with N-NO orientation and avalanche faces dipping more than  
142 30°, derived from the storm action; c) facies 3 (F3: sandy tidal flat: -0.5-0.5 m  
143 a.s.l.): fine to medium sands with an abundant bioclastic record and reddish  
144 intercalations rich in organic matter; and d) facies 4 (F4: chenier; 0.5-1 m a.s.l.):  
145 well selected bioclastic gravels with planar cross-stratification; and e) facies 5  
146 (F5: high-energy deposits; 0.75-2.25 m a.s.l.): massive shell-supported  
147 accumulations of molluscs (mainly bivalves), locally arranged on facies 2 (Fig. 3,  
148 section C) and initially attributed to storms (Cáceres et al., 2018).

### 149 2.3. Geological evolution of Saltés Island and human occupation

150 The archaeological record of Saltés Island presents an evolution linked to  
151 its own geological formation, which began with the emersion of the first  
152 northern cheniers on a previous sandy tidal plain (Fig. 2, C-E: El Almendral). This  
153 emersion occurred approximately 3,200 years ago (Suárez Bores, 1971) and a  
154 Roman salting factory (*cetaria*) was built later on this chenier (2nd century BCE),  
155 with the presence of pools and massive accumulations of bivalves (mainly  
156 *Glycymeris*; Ponsich, 1988). It was part of a significant number of coastal  
157 settlements distributed along the entire southern Iberian Atlantic coast (Fig. 2,  
158 E).

159 New cheniers subsequently emerged to the south (Fig. 2, C: La Cascajera),  
160 which were partially eroded by the action of storms that caused the deposit of  
161 washover fans between the 1st century BCE and the 3th century CE (González-  
162 Regalado et al., 2018). Recently, a new Roman *cetaria* has been discovered in  
163 the northern end of this chenier, very close to the study area (Campos et al.,  
164 2015). Its first evidence of occupation occurred during the 4th century CE and  
165 they lasted until the end of the 5th century CE or early 6th century CE.  
166 Consequently, this occupation occurred between two and three centuries after  
167 the deposit of washover fans.

168

### 169 **3. Material and methods**

#### 170 *3.1. Field methods*

171 The erosive action of the Estero de los Difuntos ebb-tide channel on the  
172 northwestern sector of La Cascajera chenier has allowed the current exposure of  
173 various profiles. One of them was selected due to the presence of all the facies  
174 defined above (facies 1-5; Cáceres et al., 2018) and three sections about 3.5 m in  
175 height were sampled (Fig. 2, D), with the extraction of forty-one samples (Fig. 3).

176

177

### (FIGURE 3)

178 Samples were selected from the different vertical sedimentary units that  
179 make up the facies mentioned above (Fig. 3). Before taking each sample, the  
180 surface was cleaned and a small vertical profile was made, to avoid possible  
181 contamination of overlying or adjacent areas. Samples were collected with a  
182 spatula and placed in self-sealing plastic bags.

183 In addition, an underwater exploration of the adjacent littoral has been  
184 carried out to find the possible origin of the molluscs present in these facies.

185

186 *3.2. Laboratory methods*187 *3.2.1. Macrofauna*

188 In each sample, two hundred grams of sediment were separated for the  
189 malacological analysis, which were sieved through a 2 mm mesh sieve. The  
190 residue was dried in an oven at a constant temperature of 40 °C for a period of  
191 not less than one day. Once dry, they were weighed and all mollusc specimens  
192 were extracted, weighed and counted. A valve of a bivalve was considered as a  
193 specimen if it retained the hinge and at least half of the valve, and as a  
194 fragment if this element had disappeared. On the other hand, the vast majority  
195 of gastropods have remained almost complete and they have only been  
196 considered as fragments if they did not keep the last body whorl. Complete and  
197 fragmentary specimens of identified species or genera has been counted (e.g.  
198 NISP: Number of Identified Specimens), a common way of quantifying faunal  
199 remains in shell middens (Heritage New Zealand Pouhere Taunga, 2014;  
200 Lambacher et al., 2016).

201 Given that the great majority of samples (> 85%) did not exceed 160  
202 shells, the sample size (NISP: 50 individuals/sample) was calculated for this  
203 number with a confidence level of 95% and an accuracy of 5%, assuming that  
204  $p=0.5$  (50%). These levels are similar to others applied in the study of these  
205 organisms (eg, Paolucci, 2010). The study of 50 shells per sample is the basis of  
206 this work.

207 To select them randomly, the content of each sample was spread on a  
208 tray subdivided into nine squares. The shells were collected by grids. The order  
209 of choice of the grids was established for each sample by random numerical  
210 series from 1 to 9. In the samples with less than 50 shells the total of these were  
211 taken into account.

212 If possible, specimens were identified to the species level, according to  
213 Gómez (2015) and the World Register of Marine Species (WoRMS). We  
214 established: a) the percentage by weight of the molluscs, the number of  
215 molluscs per sample and the species richness; b) the total percentages of each  
216 species in each sample; and c) the range and mean percentages of the main  
217 species in the different facies and the upper soil.

218

### 219 3.2.2. *Edible species*

220 The percentages of the edible species in all the samples were also  
221 calculated, as well as their minimum, maximum and average for the facies and  
222 the upper edaphic profile. They are exclusively bivalves, such as *Glycymeris*,  
223 *Acanthocardia*, *Cerastoderma*, *Chamelea*, *Donax*, *Ruditapes*, *Spisula* and the  
224 family Ostreidae, in accordance with current standards. These genera are  
225 frequently consumed today in southwestern Spain (Junta de Andalucía, 2001)  
226 and they are also some of the most abundant in numerous shelf middens of  
227 southwestern Spain (e.g. Martín and Campos, 1995; Bernal et al., 2014; Campos  
228 et al., 2015b; Bernal et al., 2015). Consequently, it is reasonable to assume that  
229 these species were also appreciated by the Romans.

230

### 231 3.2.3. *Statistical analysis*

232 A statistical analysis was applied to the percentages of the eighteen most  
233 abundant species. In a first phase, the correlation coefficient matrix between  
234 them was calculated to obtain a global view of the main groups of taxa. In  
235 addition, a Q-mode cluster analysis was applied to: a) the percentages of these  
236 eighteen main species; and b) the percentages of edible species in each sample.  
237 Five methods were initially applied (complete linkage clustering; single; average;

238 Mcquitty; Ward) and the first one was selected since it reflects the number of  
239 groups that are deduced in most of the others.

240

#### 241 3.2.4. *Taphonomy*

242 To define the state of preservation of the valves, a classification  
243 composed of four categories was made: perfect condition (A); small erosions (B);  
244 perforations and umbo/broken apex (C); and fragments (D). In a later step, the  
245 percentage of each category in the samples was calculated, as well as their  
246 minimum, maximum and average percentages in each facies and the upper  
247 edaphic profile.

248 Some additional taphonomical features have been tested: a) the  
249 percentages of bioeroded valves and the types of bioerosion; b) the fractures of  
250 the bivalve margins or the gastropod shells with a possible anthropic origin; and  
251 c) evidence of cooking.

252 A morphometric analysis of *Glycymeris nummaria*, the most abundant  
253 species among bivalves, has been made. The anterior-posterior axis of all its  
254 valves was measured and four intervals or growth stages were established with  
255 equal amplitude from 12.1 to 52.8 mm. Next, the percentage of each stage was  
256 calculated in fourteen samples selected and representative of the different  
257 facies and the upper edaphic profile, as well as its minimum, maximum and  
258 average in each sedimentary facies.

259

#### 260 3.2.5. *Microfauna*

261 Forty samples were selected for microfaunistic analysis. In each sample,  
262 ten grams of sediment were separated, an adequate amount according to other  
263 microfaunal studies in archaeological sites (Lilley et al., 1999). These samples

264 were levigated through a 125  $\mu\text{m}$  mesh sieve. The residue was dried in an oven  
265 at a constant temperature of 40  $^{\circ}\text{C}$  for a period of not less than one day. The  
266 total populations of foraminifera and ostracods were extracted and the average  
267 abundance of the main species of both groups was calculated in each facies.

268

### 269 *3.3. Archaeology*

270 The archaeological survey surface focused on the border areas of the  
271 sections studied, with the delimitation of eight sectors (Fig. 11, I-VIII) with a  
272 variable area between 6,500  $\text{m}^2$  (sector III) and 13,000  $\text{m}^2$  (sector I). The  
273 recovered objects were classified, accounted for and divided into ceramic or  
274 construction material (Tab. 10).

275

### 276 *3.4. Dating*

277 The age of the different facies were extracted from Cáceres et al. (2018)  
278 and González-Regalado et al. (2019b), with the application of the reservoir effect  
279 correction ( $-108 \pm 31$   $^{14}\text{C}$  yr) calculated by Martins and Soares (2013) in this  
280 area. Results are presented as calibrated ages for  $2\sigma$  intervals (Tab. 11). In  
281 addition, the  
282 time interval of production of different amphoras found in the upper soil has  
283 been taken into account.

284

## 285 **4. Results**

### 286 *4.1. Malacofauna*

#### 287 *4.1.1. Global analysis*

288 Malacofauna constitutes an important percentage of the total sediment  
289 weight in most of the studied samples, with a facies average that ranges from  
290 2.5% (F1) to 73.8% (F5) (see supplementary data). Mollusc shells are abundant in  
291 F3 (M: mean; M: 84 specimens/sample), F4 (M: 80 specimens/sample) and  
292 especially F5 (50-404 specimens/sample; M: 156 specimens/sample). Species  
293 richness is also greater in these three facies, with 8-11 species in most of their  
294 samples. On the contrary, molluscs are rare in F1 (2 species/sample) and in  
295 some sandy, non-bioclastic levels of F2 (2-3 species/sample).

296

**(TABLE 1)**

297

298

**(FIGURE 4)**

299

300

301 In total, 3,566 specimens were collected and 1,724 of the projected 2,050  
302 have been studied, because in 12 of the samples the number of shells was less  
303 than the calculated sample size (50). Forty-eight taxa have been identified,  
304 reaching the specific level for thirty-two of them (Table 1). Bivalves (Fig. 4;  
305 50.96%) dominates slightly over gastropods (Fig. 5, except R; 48.81%), whereas  
306 scaphopods are poorly represented (Fig. 5, R; 0.23%). The former are  
307 represented by twenty-two species, although the most abundant are *Glycymeris*  
308 *nummaria* (Fig. 4, M; M: 22.81%), *Chamelea gallina* (Linnaeus) (Fig. 4, G; M:  
309 10.21%), *Ostrea* sp. (Fig. 4, O; M: 5.46%), *Glycymeris glycymeris* (Linnaeus) (Fig. 4,  
310 L; M: 3.89%) and *Loripes orbiculatus* Poli (Fig. 4, X; M: 3.83%).

311

**(FIGURE 5)**

312

313

314           Gastropods have a lower diversity with twelve identified species, among  
315 which *Bittium reticulatum* (Da Costa) (Fig. 5, B) constitutes almost half of the  
316 studied specimens (45.39%). The rest of gastropods appear very sporadically  
317 and do not exceed 0.65% in any case. *Calyptraea chinensis* (Linnaeus) (Fig. 5, E;  
318 M: 0.64%) and *Rissoa* sp. (Fig. 5, N; M: 0.58%) approach this value.

319

320

#### 321 4.1.2. Species richness and sedimentary facies

322           The percentages of the species present vary considerably depending on  
323 the sedimentary facies that contain them (Table 2 and supplementary data). Silty  
324 tidal flat (F1) is characterized by the smaller number of individuals (2) and taxa  
325 (2), with scarce specimens of the bivalve *C. gallina* and the gastropod *B.*  
326 *reticulatum*. The faunal abundance of washover fans (F2) varies remarkably  
327 between the medium, non-bioclastic sands (6-25 specimens/200 g) and the  
328 bioclastic levels (45-195 specimens/200 g), although the dominant species are  
329 similar. *B. reticulatum* is the dominant species (M: 61.3%), with the bivalves *G.*  
330 *glycymeris* (M: 10.2%) and *C. gallina* (M: 9.2%) as main secondary species.

331           Bioclastic sands of sandy tidal flats (F3) differ from the previous facies in a  
332 higher average percentage of bivalves, as well as by the replacement of *G.*  
333 *nummaria* (M: 17.6%) by *G. glycymeris* as dominant bivalve. The total  
334 percentage of bivalves slightly increases, although *B. reticulatum* continues to  
335 be the most abundant mollusc in most samples (M: 52.8%). On the other hand,  
336 the bioclastic cheniers of La Cascajera (F4) have similar percentages of bivalves  
337 and gastropods. Among the former, *C. gallina* (M: 19.5%) and *G. nummaria* (M:  
338 15.7%) are dominant, although their average percentages are much lower than  
339 those of *B. reticulatum* (M: 46.7%).

340

341

**(TABLE 2)**

342

343 F5 presents a singular malacological content, being the bivalves (> 60%)  
344 dominant over the gastropods in all the samples. The main species is *G.*  
345 *nummaria* (M: 47.6%), which can represent up to 78% of all molluscs. It is  
346 followed in abundance by *B. reticulatum* (M: 20.2%), while ostreids and *C.*  
347 *gallina* appear frequently represented (M> 6.5%). Occasionally, the bivalves  
348 *Anomia ehippium* Linnaeus, *Cerastoderma glaucum* (Bruguière) and *L.*  
349 *orbiculatus* can reach 10% or more. The two studied samples of the edaphic  
350 profile closely resemble this facies in their faunal composition, with high  
351 percentages of *G. nummaria* (35-66%) and *B. reticulatum* (14-40%). *C. gallina* (2-  
352 10%) is the main secondary species in these samples.

353

354 *4.1.3. Statistical analysis*

355 The correlation matrix allows to distinguish three groups of species (Table  
356 3). The first group (*A. ehippium* -*Cerastoderma edule* -*C. glaucum* -*Ostrea* sp.) is  
357 positively correlated with *G. nummaria*, while these species have negative  
358 correlations with *B. reticulatum*. This group and *G. nummaria* are abundant in F5  
359 (40-84%), the upper edaphic profile (35-72%) and some basal samples of F4  
360 included in section C (24-64%). Their percentages are low in F3 (4-12%). A  
361 second group (*C. gallina*-*Corbula gibba*-*Dentalium* sp.) shows a very irregular  
362 distribution, reaching 50% in some samples of F1 and F4 and not exceeding  
363 32% in F2. The third group (*L. orbiculatus*-*Calyptrea chinensis*-*Rissoa* sp.) is  
364 scarcely represented in F2 (<10% in most samples), F3 (M: 6%), F4 (M: 5.4%) and  
365 F5 (<8% in most samples), although it reaches 24% in sample C-10 (F5).

366

367

**(TABLE 3)**

368

369 Cluster analysis allows us to differentiate two large groups of samples  
370 (Fig. 6, A: groups 1 and 3) and detects an outlier (group 2: sample B-3),  
371 identified by the predominance of *G. glycymeris* (52%) on *B. reticulatum* (26%)  
372 and ostreids (8%). Group 1 (16 samples) includes all samples of F5 and some  
373 samples of F4 below them in section C, as well as the soil samples and two  
374 samples of F2 located below it in sections A and B (Fig. 6, B). The majority of  
375 these samples are characterized by the predominance of *G. nummaria*, although  
376 three subgroups can be distinguished. Subgroup 1.1 (5 samples) is characterized  
377 by the highest percentages of *G. nummaria* (Fig. 6, C: 54-78%; M: 61.2%), with *B.*  
378 *reticulatum* (12-30%; M: 18%) and *C. gallina* (2-20%; M: 8.4%) as main secondary  
379 species. Subgroup 1.2 (3 samples) is exclusively represented in F5, with *G.*  
380 *nummaria* as the most representative species but with lower percentages than  
381 in the previous subgroup (38-44%; M: 42%). Other differentiating characters of  
382 this group are the high percentages de ostreids (16-26%; M: 22.67%), as well as  
383 the highest values of *A. ephippium* (4-12%, M: 7.33%) and *C. glaucum* (2-10%,  
384 M: 5.33%) of all the differentiated subgroups. Subgroup 1.3 (8 samples) shows  
385 very similar percentages of *G. nummaria* (24-42%, M: 33.32%) and *B. reticulatum*  
386 (28-41.3%, M: 34.83%), together with moderate percentages of *C. gallina* (4-  
387 22%, M: 12.17%).

388 Group 3 (24 samples) consists of samples from F1 to F4, all dominated by  
389 the gastropod *B. reticulatum*. Four subgroups can be distinguished, defined  
390 mainly by the decreasing abundance of this taxon. The main feature of  
391 subgroup 3.1 (7 samples) is the extremely abundance of *B. reticulatum*, the  
392 highest amongs all the subgroups determined (74-86.7%; M: 79.5%). *C. gallina*

393 (0-10%; M: 5,06%) is the main secondary species in these samples, the majority  
394 of which belong to F2. Subgroup 3.2 (9 samples) includes seven samples of F2  
395 and one sample of both F3 and F4. Their percentages of *B. reticulatum* are lower  
396 than the previous subgroup (52-70%; M: 64.29%), being accompanied by *C.*  
397 *gallina* (0-14; M: 7.86%), *L. lucinalis* (0-17%; media: 5.83%) and *G. glycymeris* (0-  
398 16%; M: 5.36%). Subgroup 3.3 (2 samples) presents similar percentages of *B.*  
399 *reticulatum* (44.44-50%; M: 47,22%) and *C. gallina* (38.89-50%; media: 44.44%).  
400 The last subgroup (6 samples) includes samples of F2, F3 and F4, which are  
401 distinguished from the previous three subgroups by the lower percentages of *B.*  
402 *reticulatum* (27.3-54%; M: 45.87%), as well as by the moderate values of *C.*  
403 *gallina* (10-30%; M: 18.79%), *G. nummaria* (2-20%; M: 12.18%) and sometimes  
404 appreciable values of *L. orbiculatus* (up to 12%).

405

406

**(FIGURE 6)**

407

408

409

#### 410 4.1.4. Facies and edible species

411 The most representative edible species are *G. glycymeris*, *G. nummaria*, *C.*  
412 *gallina* and oysters. In the sandy tidal flat (F3), these edible species represent an  
413 average of 35% of the malacofauna (range: 18-66%), with *G. glycymeris* (M:  
414 17.6%) and *C. gallina* (M: 10%) as the most representative species and very low  
415 percentages or absence of *G. nummaria* (0-4%). In F2 (washover fans) and F4  
416 (chenier), *C. gallina* also has moderate percentages (M: 10-19.6%), while *G.*  
417 *nummaria* (M: 10-15.7%) replaces *G. glycymeris* as main low-quality culinary  
418 species.

419 The highest percentages of edible species are found in F5 (M: 68.4%) and  
420 in soil (M: 62.5%). F5 includes the largest number of edible species of all the  
421 facies studied (Table 4: 11 species), some of them exclusively present in this  
422 facies (e.g. *Acanthocardia* spp. -3 species-, *P. maximus*, *R. decussatus*). In section  
423 C, the vertical evolution of this facies allows to differentiate three horizons: a)  
424 basal horizon (samples C-5 and C-6), with high percentages of *G. nummaria* (54-  
425 78%); b) intermediate horizon (samples C-7 to C-10), with lower percentages of  
426 *G. nummaria* (38-54%); and c) upper horizon (samples C-11 and C-12), with an  
427 important diversification, significant percentages of oysters (up to 26%) and the  
428 highest percentages of cockles (4-10%) of all the samples studied. The upper  
429 edaphic profile resembles F5, with a high percentage of edible species (M:  
430 62.5%) dominated by *G. nummaria* (M: 50.5%).

431

432

**(TABLE 4)**

433

434 Cluster analysis dendrogram in Q-mode allows to differentiate five  
435 groups of samples, according to their percentages of edible species (Fig. 7, A).  
436 Group E.1 (2 samples) is characterized by the highest percentages of *G.*  
437 *nummaria* of all the samples studied (66-78%, average: 72%). These samples  
438 belong to F5 and to the soil of section B.

439 Group E.2 (14 samples) have lower percentages of *G. nummaria* than  
440 group E.1. This species is usually accompanied by *C. gallina* and ostreids. Two  
441 subgroups can be differentiated, each of them composed of seven samples.  
442 Subgroup E.2.1 includes samples from F4, F5 and the soil of section C, as well as  
443 samples of the upper part of the three sections belonging to F2 (Fig. 7, B). They  
444 have moderate percentages of edible species (45-66.7%), with dominance of *G.*  
445 *nummaria* (24-38%; M: 32.08%) and *C. gallina* (10-16.67%; M: 13.34%). Most of

446 these samples included in subgroup E.2.2 belong to F5 and contain percentages  
447 of *G. nummaria* higher than those of the previous subgroup (38-54%; M:  
448 47.14%). Ostreids (4-26%; M: 12.57%) and *C. gallina* (2-20%; M: 7.71%) are well  
449 represented in this subgroup.

450 Group E.3 (one sample) is distinguished by the domain of *G. glycymeris*  
451 (Fig. 7, C: 52%), with ostreids (8%) and *C. gallina* (4%) as main secondary  
452 species. This group coincides with the outlier detected in the previous cluster  
453 analysis (see section 4.1.3). This last species is the dominant edible species of  
454 group E.4 (30-50%; M: 39.63%), which includes three samples from F4 (section  
455 B) and F1 (section A).

456

457

#### (FIGURE 7)

458

459 Group E.5 (21 samples) includes samples from F2, F3 and F4 separated  
460 in three subgroups, whose common character is the absence or the very low to  
461 moderate percentages of edible species (0-50%). Subgroup E.5.1 (sample A-4:  
462 F2) presents a very low number of specimens (see supplementary data), with  
463 isolated valves of *G. nummaria* and *Ostrea* sp. The common characteristic of  
464 subgroup E.5.2 (14 samples) is a percentage of edible species lower than 25%  
465 in all cases. These samples from F2, F3 and F4 have very low to low percentages  
466 of *C. gallina* (0-14%, M: 6.44%) and *G. glycymeris* (0-16%; M: 4.39%). Subgroup  
467 E.5.3 is constituted by six samples from facies 2, 3 and 4 with intermediate  
468 percentages of edible species (30-50% of the total malacofauna). Its main  
469 components are *C. gallina* (10-22.73%, M: 16.46%) and *G. nummaria* (2-20%; M:  
470 11.85%).

471

472 4.1.5. *Taphonomy and sedimentary facies*

## 473 4.1.5.1. Preservation

474 The defined taxonomic categories are represented unequally in the  
475 different facies. The largest number of well-preserved specimens (category A) is  
476 found in F5 (Table 5; M: 27.97%), which also contains another 50.87% of  
477 specimens with small erosions (category B). F3 and soil are as follows in order of  
478 conservation, with 67-71% of shells between categories A and B.

479 On average, the facies with the worst conserved specimens are, in  
480 descending order, F2 (C + D ~ 36.8%) and, above all, F4 (C + D > 39%). In  
481 addition, the first one has the smallest number of specimens in category A  
482 (<10%).

483

484 **(TABLE 5)**

485

486 Bioerosion is very rare in all facies studied, affecting 1.75% of all  
487 specimens (7 of 400) in F5 and 1.74% of them (23 of 1,324) in the remaining  
488 facies. It is mainly represented by traces of predatory gastropods (Fig. 8, A:  
489 *Oichnus*) or branching networks of galleries formed by sponges (Fig. 8, B:  
490 *Entobia*).

491

492 **(FIGURE 8)**

493

494 Numerous bivalves and gastropods present fractured margins and  
495 broken shells, but the fracture surfaces are rounded and no evidence of human  
496 manipulation has been found. In addition, no vestiges of cooking have been  
497 observed.

498 4.1.5.2. *Morphometric analysis of Glycymeris nummaria*

499

500 According to the length of the antero-posterior axis of the valves, a  
501 predominance of the smallest sizes (4.4-28.6 mm) over the larger sizes (28.6-  
502 52.8 mm) is observed. However, a more detailed analysis allows to differentiate  
503 three groups of facies (see supplementary data files for percentages). Group M1  
504 (F3-F4) is characterized by a significant dominance of sizes smaller than 16.5  
505 mm ( $\geq 61\%$ ), while specimens with an anterior-posterior length greater than  
506 28.6 mm are on average between 5.5% and 9%. Group M2 (F2) presents the  
507 highest percentages of the 16.5-28.6 mm interval (59.75%) and almost the entire  
508 population of *G. nummaria* (>95%) has an antero-posterior length lower than  
509 28.6 mm. In the remaining facies (group M3: F5-soil), the approximate half of  
510 the population is between 16.5 mm and 28.4 mm, while the sizes greater than  
511 28.6 mm are more abundant than in the previous group (26-36%). The largest  
512 average size is found in F5, which also has the smallest percentages of sizes  
513 below 16.5 mm (M: 10.67%). The differences between these groups are reflected  
514 in Fig. 9.

515

516

**(FIGURE 9)**

517

518 4.2. *Microfauna*

519 Both density and diversity of microfauna are very variable in the selected  
520 samples, even within the same sedimentary facies (see supplementary data).  
521 Foraminifera and ostracods are frequent in F3 (26-114 specimens/sample; M:  
522 66.4 specimens/sample) and the basal samples of section C belonging to F4 (up  
523 to 166 specimens/sample), which also have the greatest diversity (30  
524 species/sample). Conversely, both groups disappear in all samples of F5 and soil

525 and an isolated sample of F4. In total, 1,116 specimens belonging to 60 species  
526 were collected from twenty-nine samples. In these samples, foraminifera  
527 (68.19%) dominates clearly over ostracods (31.81%). The former are represented  
528 by thirty-three taxa, with *Ammonia beccarii* (17.2%), *Elphidium crispum* (9.5%),  
529 *Ammonia tepida* (7.08%), and *Elphidium advenum* (4.12%) as main benthic  
530 species. Planktonic species are very rare (4 species; 0.63%).

531

**(FIGURE 10)**

533

534 Ostracods have a lower diversity (27 species), with *Urocythereis britannica*  
535 (11.11%) as most representative species. Only other four species of this group  
536 surpass 2% of the total microfauna (*Bairdia mediterranea*, *Cytherois fischeri*,  
537 *Paracypris polita*, *Xestoleberis communis*).

538 The benthic foraminiferal population of F1 to F4 is very similar, being  
539 composed mainly of *Ammonia beccarii*, *Ammonia tepida* and *Elphidium crispum*  
540 (Table 6). The first species is dominant in F2 (M: > 6 specimens/sample) and F3  
541 (M: ~10 specimens/sample), while the second has its largest number of  
542 individuals in F4 (M: >5 specimens/sample).

543

**(TABLE 6)**

545

546 **4.3. Archaeology**

547 A total of 840 fragments have been recovered in the eight sectors studied  
548 (see supplementary data), almost all of Roman times. Two main types of Roman  
549 materials have been recognized, according to their functionality and tipology: a)

550 construction material, including complete copies and fragments of bricks  
551 (pedalis, semipedalis), imbrices, tegulae and plates (Fig. 11, D-E-F); and b)  
552 ceramic material, which includes several types of amphoras (e.g. III/Keay 25-X,  
553 LRA1, Keay XIX-C, Keay XXXV) and fine tableware (76-type and 91-type of  
554 African Red Slipe Ware-ARSW).

555

556

### (FIGURE 11)

557

558 Density of construction materials is higher in the north-western sectors  
559 closer to the sections studied (Fig. 11, A-B: sectors I-II) and decrease towards the  
560 southeast. On the contrary, the highest concentrations of ceramic materials  
561 were found in the central sectors (IV-V). Both types of materials are rare in the  
562 southeast sectors (VII-VIII).

563

### 564 4.3. Dating

565 According to Cáceres et al. (2018), age of washover deposits (F2) is  
566 between the 1st century BCE and the 2nd century CE (Table 7). A dating  
567 obtained in F5 (sample C-10) has provided a more recent age (370 CE-510 CE).

568

569

### (TABLE 7)

570

571 The amphoric materials of La Cascajera provides an additional dating on  
572 the soil covering the three sections studied. Some of them were produced from  
573 the middle of the 3rd century CE until the 5th century CE (e.g. Keay XIX-C), while

574 the production of others are restricted to the 5th century CE (e.g. Keay XXXV,  
575 LRA1, ARSW-types) (Keay, 1984).

576

## 577 **5. Discussion**

### 578 *5.1. Facies 5: a coastal shell midden?*

579 According to the previous results, F5 has a series of distinctive features in  
580 relation to the rest of facies although very similar to those of the upper edaphic  
581 horizon:

582 1. Faunal content. It presents the highest percentages in total weight of  
583 malacofauna, as well as the largest average number of specimens per sample  
584 (Tab. 1). It is composed overwhelmingly of shells with a very scarce sediment  
585 matrix. This facies and the upper soil stand out for the abundance of the  
586 *Glycymeris nummaria* and other edible species associated with it (see  
587 correlation matrix and cluster analysis), whereas the malacofauna of the  
588 remaining facies are mainly dominated by the gastropod *Bittium reticulatum*.  
589 This characteristic points to a shell deposit created by human activity, similar to  
590 those observed in northern Australia (SMD, *sensu* Holdaway et al., 2017). In  
591 addition, it does not include non-molluscan food remains. These features are  
592 frequent in numerous coastal middens as opposed to other shells deposits like  
593 cheniers (e.g. Beaton, 1985; Sullivan and O'Connor, 1993).

594 *Glycymeris nummaria* is very common in coastal middens of  
595 southwestern Spain in Roman *cetariae*. These factories also processed other  
596 edible species of bivalves (*Ruditapes*, *Chamelea*, *Ostrea*) or very specific species  
597 of gastropods that were dedicated to the production of imperial purple, such as  
598 *Bolinus brandaris* (Campos et al., 2014; Bernal et al., 2015). The presence of  
599 some samples of F2 and F4 with similar abundances of *Glycymeris nummaria*

600 (Fig. 6, B: group 1) may be due to the dropping of valves into the lower  
601 sedimentary layers from F5 or the soil.

602         2. Edible species. F5 is also differentiated by the statistical analysis of  
603 edible species (Fig. 7: group E1 except sample C-6), because it has the highest  
604 percentages and diversity of these species (see supplementary data files). This  
605 characteristic differentiates F5 from the soil, which also has contents higher than  
606 60% in edible species. This high abundance is one of the most widespread  
607 features among shell middens (Bailey, 1977; Rosendahl, 2005).

608         3. Taphonomy. Valves found in both F5 and soil have have a better state  
609 of conservation than those of the other facies, which have undergone an  
610 important transport by traction derived from the action of storms (F2: washover  
611 fans) or fair-weather refracted waves (F4: cheniers; Morales et al., 2014).  
612 However, the absence of both acute fractures and warming evidence rule out  
613 direct consumption of the molluscs found in both F5 and soil. Therefore, the  
614 abundance of edible species must be explained by an alternative accumulation  
615 mechanism (see next chapter).

616         4. Population size of *G. nummaria*. The populations of *Glycymeris* in F5  
617 and soil are larger than those found in other facies. This population distribution  
618 implies a selective process stronger than tides, waves or high-energy events can  
619 perform (e.g. F1 to F4). It is a typical feature of many coastal middens, which  
620 denotes a selective anthropic action (Baker, 1981).

621         5. Microfauna. The absence or extreme shortage of the marine  
622 microfauna in relation to other shell deposits has been cited in several coastal  
623 middens (Lilley et al., 1999; Rosendahl et al., 2007), although it has not been  
624 studied in most of them. In this case, this absence of foraminifera and ostracods  
625 can point to a wash prior to its subsequent accumulation.

626           6. Geological setting. Facies 5 overlaps the rest of facies in the northwest  
627 sector of La Cascajera (Fig. 3: section C; Fig. 12, E) and manifests as a  
628 malacological accumulation that follows the slope towards the Estero de los  
629 Difuntos ebb-tide.

630           7. Age. The datings obtained from an isotopic dating of F5 and the  
631 deduced from the ceramic materials show a high degree of coincidence (e.g.,  
632 5th century CE) and they are concordant with the oldest ages obtained in the  
633 underlying materials (F2: 1st century BCE-2nd century CE).

634           These results reveal that F5 meets the characteristics of a midden shell,  
635 according to the main criteria normally used to distinguish them from natural  
636 shell deposits (see reviews in Bailey, 1977; Gill et al., 1991; Attenbrow, 1992;  
637 Szabó, 2017, among others)

638

### 639 5.2. *The origin of F5*

640           For all the above, it is considered that F5 has an anthropic origin, as a  
641 consequence of the industrial activity of the next Roman *cetariae*. The  
642 taphonomic features and the absence of microfauna point to: a) a wash of the  
643 valves; b) a partial selection by size; and c) its accumulation and the creation of a  
644 landfill in favor of the slope of the ebb-tide channel (Fig. 12, E). Other similar  
645 landfills in haleutic contexts have been described in southwestern Spain,  
646 associated with Roman *cetariae* (Bernal, 2011). It is classified as a bivalent shell  
647 midden, since it does not include molluscs dedicated to the production of  
648 Imperial purple (Bernal, 2011).

649           In Mediterranean and Atlantic shell middens of Spain, the abundant  
650 presence of *Glycymeris* has been attributed to: a) its direct consumption, with  
651 different levels of previous cooking; b) the preparation of sauces, mixed with  
652 viscera and unappreciated parts of fishes (e.g. garum); c) discards and shellfish

653 remains, product of the use of non-selective fishing gear type nets; d) an  
654 ornamental use; e) weight for fishing nets; or f) bait fishing, in archaeological  
655 sites where this genus is in low proportions (Campos et al., 2002; Álvarez et al.,  
656 2011; Bernal et al., 2015; Nadal et al., 2015; Pascual and García, 2015; Valente  
657 and Martins, 2015).

658         The abundance of *B. reticulatum* and *G. nummaria*, the different  
659 taphonomical features mentioned above and a review of the recent underwater  
660 environments of the Huelva littoral point to a discard produced by a non-  
661 selective trawling as the origin of F5. *B. reticulatum* feeds by grazing on small  
662 algae and grasslands of phanerogams (Fig. 12, A; Borja, 1986; Augier 2007). It  
663 feeds on the organic matter that covers the beam of the leaves and the soil of  
664 these meadows (Luque and Templado, 2004). In these meadows, *G. nummaria*,  
665 the main species of F5, is a frequent infaunal filtering species that inhabits the  
666 sandy bottoms of these areas (Fig. 12, B), usually associated with other bivalves,  
667 such as *Acanthocardia* spp., *C. gallina*, *Donax* spp., *Dosinia lupinus*, *Loripes*  
668 *lucinalis*, *Ruditapes decussatus*, *Spisula* spp. or *Tellina* spp. (Péres and Picard,  
669 1964; Pérès, 1982; D'Amico et al., 2013).

670         In the Huelva littoral, these vegetated sandy sediments with this mollusc  
671 assemblage are mainly found at depths between 2 and 20 m, although some of  
672 these species can occupy a wider bathymetric range (Gómez, 2015). These  
673 areas would be the main sources of shells for the deposit of the sedimentary  
674 facies that constitute La Cascajera barrier, with the transport of *B. reticulatum*  
675 and, to a lesser extent, of *G. nummaria*, to the tidal plains and the inner areas of  
676 the Tinto-Odiel estuary due to waves, tides or storms. This tractive transport of  
677 these and other species of molluscs would explain the worst state of shells in F2,  
678 F3 and F4 than in F5 or the soil (see chapter4.1.5.1.).

679 On the other hand, the abundance of *G. nummaria* in F5 would be  
680 explained as the final result of trawling on these grasslands, associated with the  
681 activity of the next Roman *cetaria*. Different features support this claim:

682 1. Mollusc assemblages. The very high percentages of *G. nummaria* and  
683 the secondary presence of *B. reticulatum* and other bivalve species are  
684 consistent with an extraction method of this type in these environments. Other  
685 edible species (*P. maximus*, *R. decussatus*) are also collected with *G. nummaria*  
686 in these subtidal sandy bottoms and consumed on this coast currently (Junta de  
687 Andalucía, 2001). However, other edible species of F5 come from other habitats,  
688 involving a secondary collection in other sedimentary environments. Edible  
689 species also include cockles and inhabitants of hard substrates, such as oysters.  
690 Among the cockles, the lagoon cockle *C. glaucum* inhabits small, often isolated  
691 euryhaline non-tidal basins (Tarnowska et al., 2012), while the common cockle *C.*  
692 *edule* is widely distributed on tidal flats of estuaries and bays (Dabouineau and  
693 Ponsero, 2009).

694 2. Taphonomy. This type of fishing involves at least four types of selection  
695 in relation to the remaining facies:

696 a. First selection, derived from the action of the fishing nets on a certain  
697 type of bottom with abundant shells of *G. nummaria* and an important  
698 proportion of *B. reticulatum* coming from the adjacent meadows. This action  
699 favors the preservation of thick shells like *G. nummaria* in comparison with other  
700 thinner species. This differential preservation or fragmentation possibility has  
701 been revealed in the study of shell middens (Muckle, 1985).

702 b. Second selection, caused by the drag of the fishing nets on the  
703 bottom, which would cause a first wash of the sandy matrix, as well as the loss  
704 of specimens with a length or diameter less than its mesh diameter. In this  
705 selection, the larger size of *G. nummaria* is a positive collection factor in relation

706 to *B. reticulatum*. This selection would also explain the largest average size of *G.*  
707 *nummaria* in this facies.

708

709

### (FIGURE 12)

710

711 c. Third selection, as a consequence of the traction of the net for a new  
712 cleaning of the catches (Fig. 12, D). These last two selections explain the absence  
713 of foraminifera and ostracods in F5, since the elimination of the sands that  
714 contained them has taken place. This absence or a very pronounced shortage  
715 has been tested in different shell middens in comparison with other shell  
716 deposits, as mentioned above (see review in Rosendahl, 2005).

717 d. Fourth selection, which eliminates empty valves and shells, non-edible  
718 species or those edible species that are not subject to industrial activity (Fig. 12,  
719 E). These four selections explain the mollusc assemblage extracted from F5 (Fig.  
720 12, F), as well as the best conservation of the valves and shells in relation to the  
721 rest of the facies.

722 3. This type of fishing has been described during this period (3rd-5th  
723 centuries CE) in the Roman factories, with the use of mobile networks to which  
724 clay and lead weights were associated (e.g. Fig. 12, C; Oliver, 1982; Campos and  
725 Vidal, 2004). In addition, the ages of both shells and archaeological remains are  
726 coincident and point to a creation of F5 as a landfill of the next Roman factory  
727 (see chapter 4.3).

728

729

730 **6. Conclusions**

731 1. The study of shell deposits exposed in the northern sector of Saltés Island  
732 has allowed to determined thirty-three species of mollucs, with a predominance  
733 of the gastropod *Bittium reticulatum* in old sandy tidal flats (F3), cheniers (F4)  
734 and washover fans (F2). Other two facies (F5 and soil) are dominated by the  
735 bivalve *Glycymeris nummaria*, while molluscs are poorly represented in the old  
736 muddy tidal flats (F1).

737 2. A multidisciplinary analysis of F5 allows to classify it as a shell midden,  
738 according to its very high percentages and diversity of edible species (clams,  
739 oysters, cockles), a partial selection by size, a better state of conservation of the  
740 *Glycymeris* valves and the absence of both foraminifera and ostracods.

741 3. This facies represents a bivalent shell midden, with a mollusc content derived  
742 from waste of a next Roman factory (*cetariae*). Soil of this factory has similar  
743 features to those observed in this shell midden. Age of this midden (4th-5th  
744 centuries CE) is similar to that deduced from the amphoric archaeological  
745 remains extracted in this soil.

746 4. The autoecological analysis of mollusc species and the revision of littoral  
747 environments of the adjacent areas indicate a process of collecting malacofauna  
748 (mainly *Glycymeris nummaria*) in shallow sandy bottoms of the Iberian Atlantic  
749 shelf with phanerogam meadows. The presence of oysters or common cockles  
750 also means an additional collection in rocky environments or tidal flats,  
751 respectively.

752 5. This shell midden comes from trawling carried out on these environments,  
753 with up to four selective processes involved in the final result.

754

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763

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1064 **FIGURE CAPTIONS**

1065 **Figure 1.** Examples of coastal shell middens, their most abundant species (B: bivalve; G:  
 1066 gastropod) and their ages. 1: Hallmann et al. (2009); 2: Erlandson et al. (2009a); 3: Erlandson et  
 1067 al. (2009b); 4: Carbotte et al. (2004); 5: Lulewicz et al. (2017); 6: Herrera and Solís (2011); 7:  
 1068 Schapiera et al. (2006); 8: Belknap and Sandweiss (2014); 9: Latorre et al. (2017); 10: Vasconcellos  
 1069 et al. (2014); 11: Zubimendi (2017); 12: Hood and Grovdal (2016); 13: Waddington et al. (2003);  
 1070 14: Lewis (2011); 15: Dupont et al. (2007); 16: García-Escárzaga et al. (2017); 17: López-Dórigal et  
 1071 al. (2019); 18: Wijnen (1981); 19: Douka et al. (2014); 20: Barusseau et al. (2010); 21: De Sapor  
 1072 (1971); 22: Bar-Yosef and Beyin (2009); 23: Martínez (1976); 24: Jerardino et al. (2008); 25: Sivan  
 1073 et al. (2006); 26: Bailey et al. (2013); 27: Magee et al. (2017); 28: Biagi (2014); 29: Cooper (1997);  
 1074 30: Tieng (2013); 31: Cassidy and Vostretsov (2007); 32: Habu et al. (2011); 33: Jing and Liang  
 1075 (2002); 34: Thiel (1986-1987); 35: Morse (1993); 36: Ward et al. (2016); 37: Bailey (1977); 38:  
 1076 Frankland (1990); 39: Flores (2009); 40: Puch (1974).

1077 **Figure 2.** A-B: location of Saltés Island (courtesy: Google maps); C: geomorphological map of  
 1078 Saltés Island and the adjacent areas of the Tinto-Odiel estuary; D: Photograph of the study area,  
 1079 with situation of the three sections analyzed; E: Location of the main Roman *cetariae* along the  
 1080 south-western Iberian coast (courtesy: Google maps).

1081 **Figure 3.** Sedimentary logs of the three studied sections, including the forty-one selected  
 1082 samples and details of the different facies.

1083 **Figure 4.** Class Bivalvia. A-B: *Acanthocardia* sp.; C: *Acanthocardia tuberculata* (Linnaeus, 1758);  
 1084 D: *Anomia ephippium* Linnaeus, 1758; E: *Cerastoderma glaucum* (Bruguière, 1789); F:  
 1085 *Cerastoderma edule* (Linnaeus, 1758); G: *Chamelea gallina* (Linnaeus, 1758); H: *Clausinella*  
 1086 *fasciata* (da Costa, 1778); I: *Donacilla cornea* (Poli, 1791); J: *Donax trunculus* Linnaeus, 1758; K:  
 1087 *Donax venustus* Poli, 1795; L: *Glycymeris glycymeris* (Linnaeus, 1758); M: *Glycymeris nummaria*  
 1088 (Linnaeus, 1758); N: *Limaria tuberculata* (Olivi, 1792); O: *Ostrea* sp.; P: *Panopea glycymeris* (Born,  
 1089 1778); Q: *Pecten maximus* (Linnaeus, 1758); R: *Ruditapes decussatus* (Linnaeus, 1758); S: *Spisula*

1090 *solida* (Linnaeus, 1758); T: Family Cardidae; U: Family Cardiidae; V: *Corbula gibba* (Olivi, 1792); W;  
 1091 *Dosinia lupinus* (Linnaeus, 1758); X: *Loripes orbiculatus* Poli, 1795: *Striarca lactea* (Linnaeus,  
 1092 1758); Z: *Tellina* sp.

1093 **Figure 5.** Class Gastropoda (A-Q)-Class Scaphopoda (R). A: *Bela* sp.; B: *Bittium reticulatum* (da  
 1094 Costa, 1778); C: *Calliostoma* sp.; D: *Calliostoma zizyphinum* (Linnaeus, 1758); E: *Calyptrea*  
 1095 *chinensis* (Linnaeus, 1758); F: *Steromphala umbilicalis* (Linnaeus, 1758); G: *Steromphala varia*  
 1096 (Linnaeus, 1758); H: *Jujubinus striatus* (Linnaeus, 1758); I: *Mangelia* sp.; J: Family Nassaridae; K:  
 1097 *Tritia nitida* (Jeffreys, 1867); L: *Ocenebra erinaceus* (Linnaeus, 1758); M: *Peringia ulvae* (Pennant,  
 1098 1777); N: *Rissoa* sp.; O: *Tritia incrassata* (Strøm, 1768); P: *Tritia pfeifferi* (Philippi, 1844); Q: *Tritia*  
 1099 *pygmaea* (Lamarck, 1822); R: *Dentalium* sp.

1100 **Figure 6.** A: Q-mode cluster analysis applied to the eighteen main species; B: distribution of  
 1101 groups and subgroups in the sections studied; C: Mean percentages of the main species in the  
 1102 different subgroups obtained.

1103 **Figure 7.** Statistical analysis of edible species. A: Q-mode cluster analysis applied to the edible  
 1104 species (see Tab. 4); B: distribution of groups and subgroups in the sections studied; C: Mean  
 1105 percentages of the edible species in the different subgroups obtained. Bold: main species of  
 1106 each subgroup.

1107 **Figure 8.** Evidences of bioerosion. A: *Oichnus*; B: *Entobia*. Scale: 1 cm.

1108 **Figure 9.** *Glycymeris nummaria*: mean percentages of the valve sizes in the different  
 1109 sedimentary facies.

1110 **Figure 10.** Foraminifera (A-H)-Ostracoda (I-K). A: *Ammonia beccarii* (Linnaeus, 1758); B:  
 1111 *Ammonia tepida* (Cushman, 1926); C: *Elphidium crispum* (Linnaeus, 1758); D: *Elphidium advenum*  
 1112 (Cushman, 1922); E: *Haynesina germanica* (Ehrenberg, 1840); F: *Planorbulina mediterraneensis*  
 1113 D'Orbigny, 1826; G: *Quinqueloculina seminulum* (Linnaeus, 1758); H: *Triloculina oblonga*  
 1114 (Montagu, 1803); I: *Urocythereis britannica* Athersuch, 1977; J: *Cythereis fischeri* (Sars, 1866); K:  
 1115 male of *Loculicytheretta pavonia* (Brady, 1866).

1116 **Figure 11.** Archaeology. A-B: location of the eight sectors studied. C: Shore of section C (sector  
 1117 II) with ceramic remains; D: Fragment of brick (sector I; scale: 5 cm); E: Fragment of brick (base of  
 1118 section C, sector II); F: *Semipedalis* (sector I).

1119 **Figure 12.** The formation of a shell midden (F5). A-B-D-E: Huelva littoral. A. Phanerogam  
 1120 meadows; B: Adjacent sea bottom with numerous valves of *G. nummaria*; C. Roman Mosaic,

1121 Hippolytus Museum, Alcalá de Henares (Spain); D: Drag net cleaning; E: Discard from a trawl; F.

1122 Shell midden of La Cascajera (F5) located on washover fans (F2).

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### 1130 **TABLE CAPTIONS**

1131 **Table 1.** Total percentages and sample distribution of mollusc species. ES: edible species. If  
1132 possible, 50 individuals were studied in each sample (NISP).

1133 **Table 2.** Distribution of the main species of molluscs (in %) in the different sedimentary facies:  
1134 mean and range of each species in each facies. White numbers: most abundant species. See the  
1135 total number of specimens per sample in the supplementary data files.

1136 **Table 3.** Coefficient correlation matrix of the most abundant species. Bold:  $p < 0.01$ ; Underlined:  
1137  $p < 0.05$ .

1138 **Table 4.** Percentages of edible species in each facies: mean and range.

1139 **Table 5.** Mean percentages and range of each taphonomic category in each facies. White  
1140 numbers: average maximum values of each taphonomic category.

1141 **Table 6.** Main species of foraminifera and ostracods in each facies: mean and range (number of  
1142 individuals per sample). See the total number of specimens counted per sample in the  
1143 supplementary data files.

1144 **Table 7.** Database of  $^{14}\text{C}$  and U/Th samples and results. Reservoir effect corrected ( $-108 \pm 31$   
1145 years BP; Martins and Soares, 2013).

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1148 **SUPPLEMENTARY DATA FILES: CAPTIONS**

1149 **File 1.** General data of the malacofauna, including its total percentage by weight, the total  
1150 number of individuals collected and studied and the number of species per sample.

1151 **File 2.** Percentages of the determined valve sizes in selected samples of the different  
1152 sedimentary facies and mean percentages of each valve size in each facies.

1153 **File 3.** Total percentages of foraminifera and ostracod species and number of individuals in each  
1154 sample. \*: reworked Neogene species.

1155 **File 4.** Archaeology. Distribution of archaeological remains in the eight sectors studied.

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FACIES	1		2		3		4		5		SOIL	
SPECIES	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range
<b>BIVALVIA</b>												
<i>Anomia ephippium</i>	0	0	0.61	0-4.35	1.6	0-2	0.85	0-4	3.11	0-12	0	0
<i>Chamelea gallina</i>	50	50	9.2	0-22.73	10	4-20	19.55	3.23-38.89	6.67	2-12	6	2-10
<i>Cerastoderma glaucum</i>	0	0	0	0	0	0	0.57	0-2	2	0-10	0	0
<i>Corbula gibba</i>	0	0	0.5	0-0.5	1.2	0-2	2.26	0-5.56	0.11	0-1	2.5	0-5
<i>Glycymeris glycymeris</i>	0	0	4.57	0-16.67	17.6	6-52	2.72	0-5.56	0.67	0-2	3	0-6
<i>Glycymeris nummaria</i>	0	0	10.21	0-33.33	1.6	0-4	15.73	0-54	47.56	36-78	50.5	35-66
<i>Loripes orbiculatus</i>	0	0	3.97	0-16.67	4.8	0-12	4.36	0-8.51	2.89	0-16	1	0-2
<i>Ostrea sp.</i>	0	0	4.5	0-16.67	4.8	2-8	2.75	0-6	9.33	0-26	3	0-6
<b>GASTROPODA</b>												
<i>Bittium reticulatum</i>	50	50	61.33	27.27-86.67	52.8	26-74	46.72	12-80.65	20.22	2-36	27	14-40
<i>Calyptrea chinensis</i>	0	0	0.12	0-2	0.4	0-2	0.59	0-2.13	1.56	0-6	0	0
<i>Rissoa sp.</i>	0	0	1.46	0-4	0.4	0-2	0.46	0-3.23	0.44	0-2	0	0
<b>SCAPHOPODA</b>												
<i>Dentalium sp.</i>	0	0	0.39	0-4.55	0.4	0-2	0.79	0-5.56	0	0	0	0

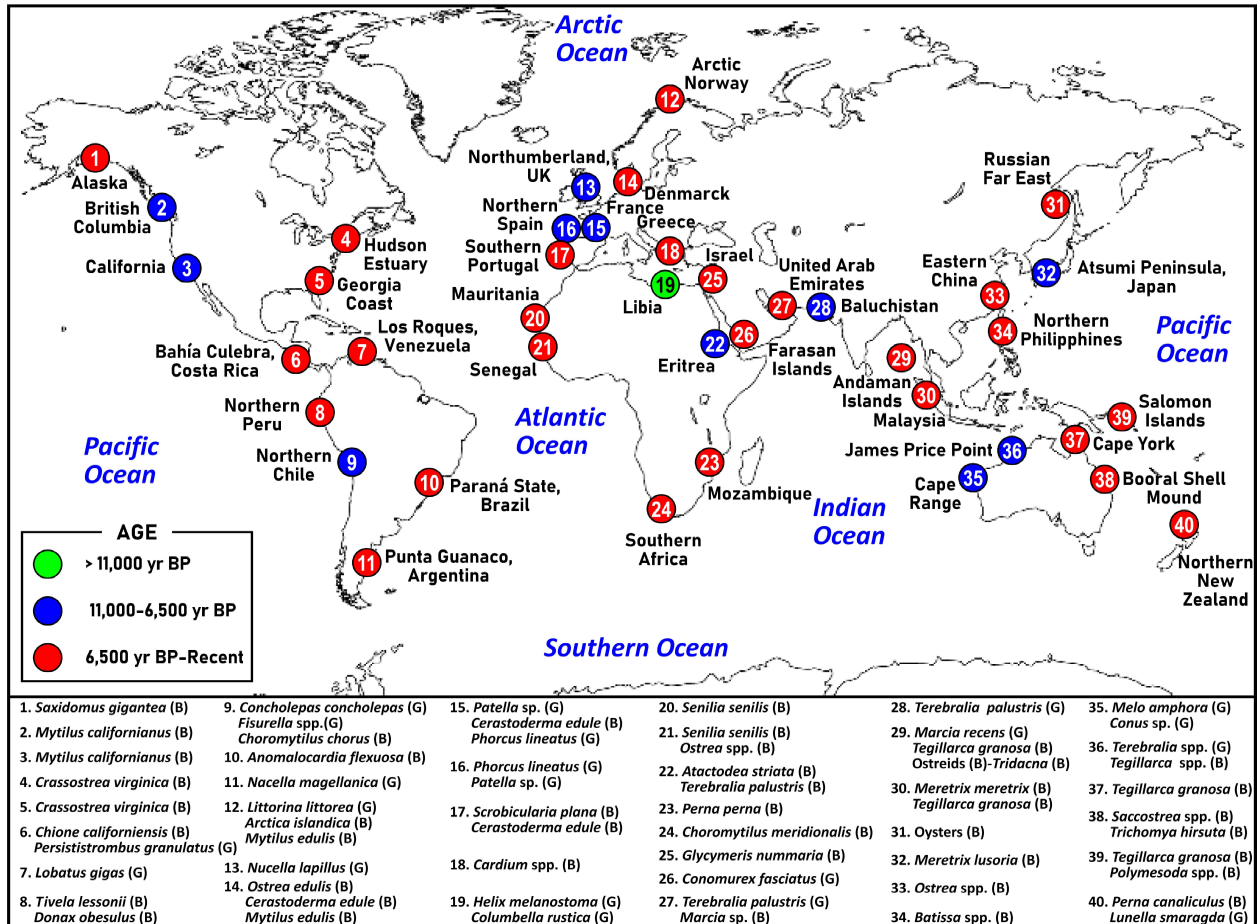
		AE	CH	CE	CG	CO	GG	GN	Lo	OE	BT	CZ	CC	GI	NA	RI	SU	TP	DS
BIVALVIA	<i>Anomia ephippium</i> (AE)	1																	
	<i>Chamelea gallina</i> (CH)	0.106-	1																
	<i>Cerastoderma edule</i> (CE)	<u>0.397</u>	0.140-	1															
	<i>Cerastoderma glaucum</i> (CG)	<b>0.531</b>	0.142-	0.185	1														
	<i>Corbula gibba</i> (CO)	0.121-	0.269	0.127-	0.099-	1													
	<i>Glycymeris glycymeris</i> (GG)	0.064-	0.030-	0.144-	0.160-	0.061	1												
	<i>Glycymeris nummaria</i> (GN)	0.301	0.199-	0.215	<u>0.327</u>	0.222-	0.253-	1											
	<i>Loripes orbiculatus</i> (LO)	0.142-	0.179-	0.192-	0.177-	0.039	0.033-	0.228-	1										
<i>Ostrea</i> sp. (OE)	<b>0.687</b>	0.280-	<u>0.338</u>	<b>0.723</b>	0.178-	0.077-	0.223	0.177-	1										
GASTROPODA	<i>Bittium reticulatum</i> (BT)	<b>0.526-</b>	0.089-	0.189-	<b>0.497-</b>	0.008	0.072-	<b>0.796-</b>	0.157	<b>0.434-</b>	1								
	<i>Calliostoma zizyphinum</i> (CZ)	0.013-	0.024	0.129	0.030-	0.200-	0.103-	0.230	0.094	0.187-	0.135-	1							
	<i>Calyptrea chinensis</i> (CC)	0.219	0.138-	0.129	0.050	0.043	0.175-	0.145	<b>0.535</b>	0.023	0.207-	0.209	1						
	<i>Gibbula</i> sp. (GI)	0.018-	0.149-	0.108-	0.103-	0.177-	0.126-	0.056	0.019	0.033	0.054	0.122-	0.014-	1					
	<i>Nassarius</i> sp. (NA)	0.159-	0.092-	0.223	0.088-	0.151-	0.091-	0.009-	0.185-	0.050-	0.168	0.182	0.129-	0.217	1				
	<i>Rissoa</i> sp. (RI)	0.148-	0.229-	0.153-	0.145-	0.145-	0.081-	0.182-	<u>0.354</u>	0.220-	0.243	0.224	0.302	0.010-	0.130-	1			
	<i>Steromphala umbilicalis</i> (SU)	0.044-	0.111-	0.108-	0.103-	0.114	0.126-	0.193-	0.166	0.090-	0.254	0.122-	0.121	0.156	0.092-	0.164	1		
<i>Tritia pfeifferi</i> (TP)	0.090-	0.126-	0.052-	0.050-	0.085-	0.029-	0.148-	0.127-	0.048-	0.230	0.059-	0.073-	0.052-	0.044-	<b>0.380</b>	0.052-	1		
SCAPHOPODA	<i>Dentalium</i> sp. (DS)	0.169-	<u>0.380</u>	0.098-	0.093-	<b>0.552</b>	0.060	0.234-	0.125-	0.050-	0.017	0.111-	0.137-	0.098-	0.084-	0.138-	0.098-	0.047-	1

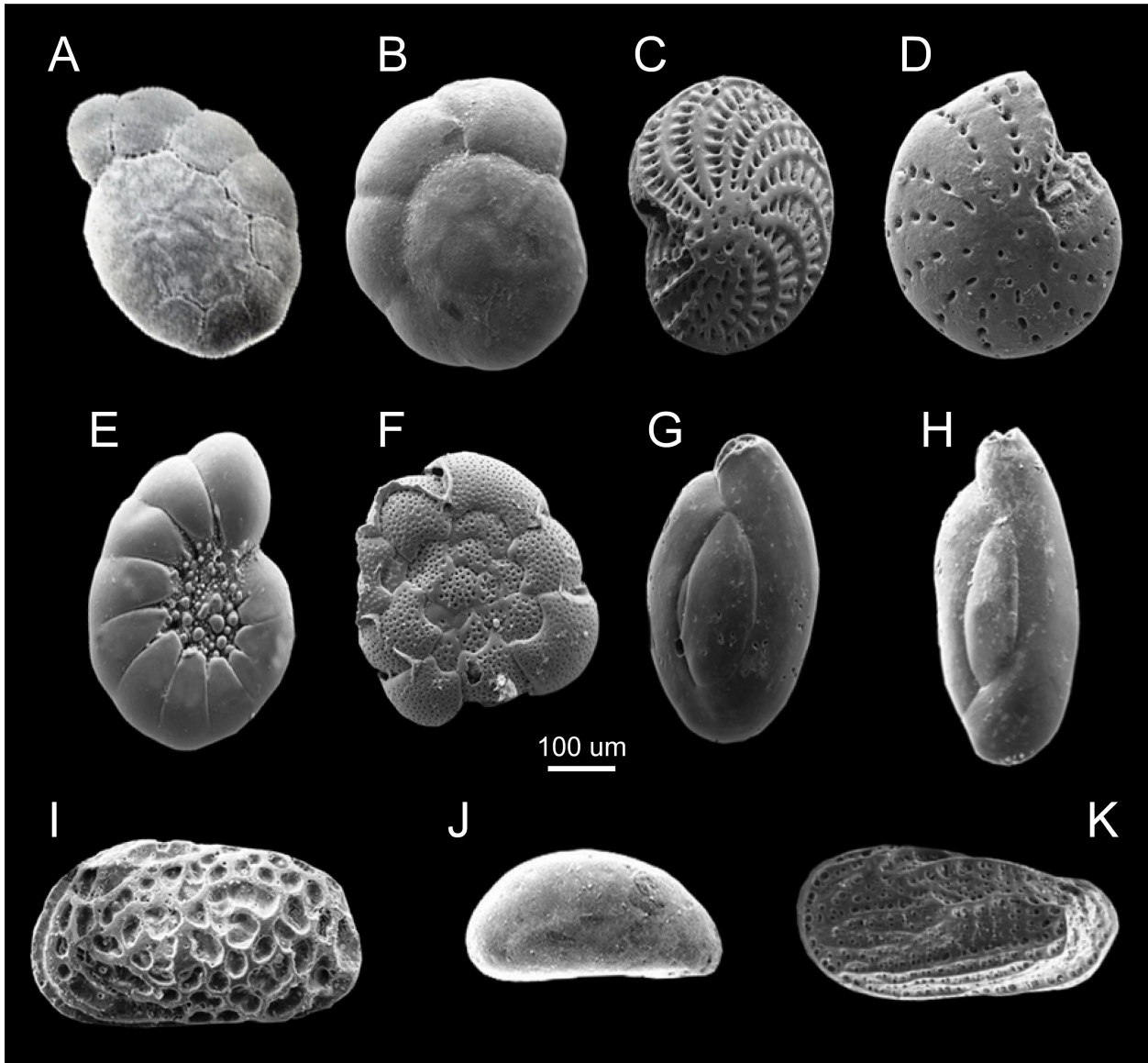
EDIBLE SPECIES/FACIES	1		2		3		4		5		SOIL	
	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range
<i>Acanthocardia aculeata</i>	0	0	0	0	0	0	0	0	0.22	0-2	0	0
<i>Acanthocardia echinata</i>	0	0	0	0	0	0	0	0	0.22	0-2	0	0
<i>Acanthocardia</i> sp.	0	0	0	0	0	0	0	0	0.44	0-2	0	0
<i>Chamelea gallina</i>	50	50	10.42	0-22.7	10	4-20	19.56	10-38.9	6.67	2-12	6	2-10
<i>Cerastoderma edule</i>	0	0	0.12	0-2	0	0	0	0	0.67	0-2	0	0
<i>Cerastoderma glaucum</i>	0	0	0	0	0	0	0.57	0-2	2	0-10	0	0
<i>Donax trunculus</i>	0	0	0.12	0-2	0.4	0-2	0	0	0	0	0	0
<i>Donax venustus</i>	0	0	0	0	0	0	0.29	0-2	0	0	0	0
<i>Glycymeris glycymeris</i>	0	0	4.57	0-16.7	17.6	6-52	2.72	0-5.56	0.67	0-2	3	0-6
<i>Glycymeris nummaria</i>	0	0	10.21	0-33.33	1.6	0-4	15.73	0-54	47.46	36-78	50.5	35-66
<i>Ostrea</i> sp.	0	0	4.5	0-16.7	4.8	2-8	2.74	0-6	9.33	0-26	3	0-6
<i>Pecten maximus</i>	0	0	0	0	0	0	0	0	0.22	0-2	0	0
<i>Ruditapes decussatus</i>	0	0	0	0	0	0	0	0	0.44	0-2	0	0
<i>Spisula solida</i>	0	0	0.24	0-2	0.4	0-4	0	0	0	0	0	0
<i>Spisula</i> sp.	0	0	0.26	0-2.22	0	0	0	0	0	0	0	0
<b>% TOTAL EDIBLE SPECIES</b>	<b>50</b>	<b>50</b>	<b>29.2</b>	<b>0-66.67</b>	<b>35.2</b>	<b>18-66</b>	<b>41.61</b>	<b>19.2-80</b>	<b>68.44</b>	<b>50-88</b>	<b>62.5</b>	<b>45-80</b>
<b>NUMBER OF EDIBLE SPECIES</b>	<b>1</b>	<b>1</b>	<b>8</b>	<b>8</b>	<b>6</b>	<b>6</b>	<b>6</b>	<b>6</b>	<b>11</b>	<b>11</b>	<b>4</b>	<b>4</b>

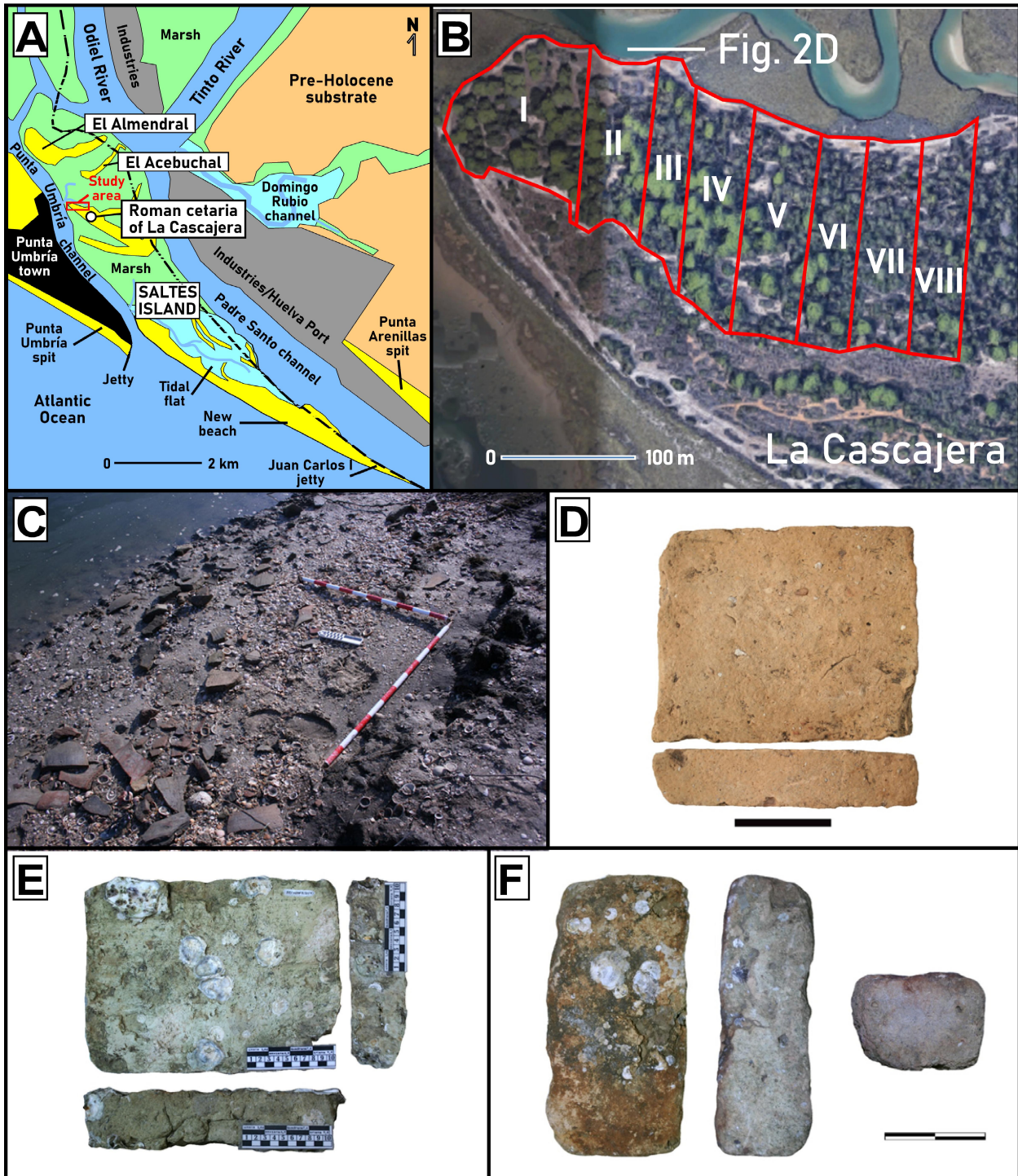
FACIES/CATEGORY	A		B		C		D	
	Mean	Range	Mean	Range	Mean	Range	Mean	Range
2	9.92	0-20	53.27	28-72.72	24.69	8-38	12.11	0-33.33
3	17.39	10.2-29.41	50.54	40.81-64.58	27.65	18.75-40.81	4.42	0-10
4	11.29	0-23.91	49.36	30-74.19	21.2	15.22-28	18.16	3.22-38.89
5	27.97	7.84-50	50.87	38.46-60.78	14.57	3.85-21.57	6.58	0-10
SOIL	4.99	3.85-6.12	65.22	61.22-69.23	15.19	3.85-26.53	14.6	6.12-23.08

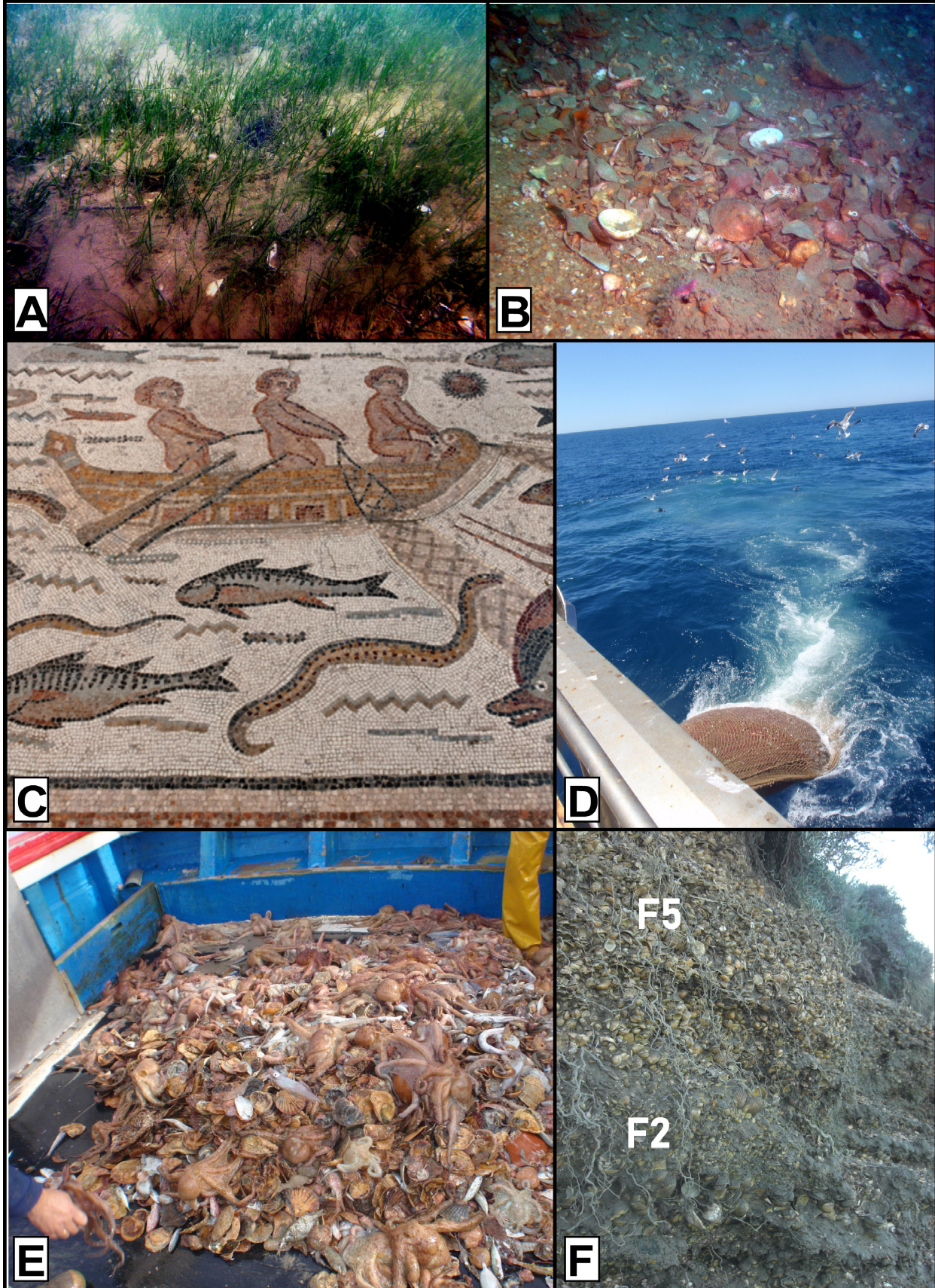
GROUP		FACIES		1		2		3		4		5		SOIL	
		SPECIES		Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range
FORAMINIFERA	BENTHIC FORAMINIFERA	<i>Ammonia beccarii</i>	6	6	6.35	0-13	9.8	5-14	4.14	0-11					
		<i>Ammonia inflata</i>	3	3	0.94	0-8	2	0-4	0.14	0-1					
		<i>Ammonia tepida</i>	3	3	0.53	0-3	5.8	0-10	5.43	0-20					
		<i>Elphidium advenum</i>	4	4	0.82	0-8	4.2	2-8	1	0-4					
		<i>Elphidium complanatum</i>	4	4	0.53	0-3	2	0-6	1.29	0-7					
		<i>Elphidium crispum</i>	7	7	3.35	0-9	4.8	1-7	2.57	0-7					
		<i>Haynesina germanica</i>	3	3	0.59	0-4	1.8	0-2	1.43	0-9					
		<i>Neoconorbina orbicularis</i>	0	0	0	0	0.8	0-4	3	0-9					
		<i>Planorbulina mediterranensis</i>	0	0	0.29	0-3	1.6	0-5	2.57	0-13					
		<i>Rosalina globularis</i>	1	1	0.41	0-4	1.6	0-8	2	0-4					
OSTRACODA		<i>Bairdia longivaginata</i>	0	0	0	0	0	0	2	0-8					
		<i>Bairdia mediterranea</i>	0	0	0	0	0.4	0-2	3.7	0-21					
		<i>Cytherois fischeri</i>	3	3	0.65	0-4	1.2	0-4	1	0-7					
		<i>Hiltermannicythere sphaerulolineata</i>	0	0	0.06	0-1	0.8	0-4	0.71	0-3					
		<i>Palmoconcha laevata</i>	0	0	0	0	0.6	0-3	0.71	0-3					
		<i>Paracypris polita</i>	3	3	0.71	0-5	0.8	0-2	0.86	0-3					
		<i>Pontocythere elongata</i>	0	0	0	0	0.8	0-4	1.43	0-4					
		<i>Semicytherura sulcata</i>	0	0	0.18	0-2	0.4	0-2	0.57	0-3					
		<i>Urocythereis britannica</i>	6	6	2.65	0-9	6.8	0-25	5.57	0-25					
<i>Xestoleberis communis</i>	1	1	0.12	0-2	0	0	4.14	0-19							

Facies	Sample code	Material ( <i>Glycymeris</i> )	Laboratory code	<sup>14</sup> C age	Error	<sup>14</sup> C Calibrated age	Median probability	U/Th	References
F2	HUCA-1301	Shell	CNA-2817	2263	31	170 BCE - 70 CE	48 BCE		Cáceres et al. (2018)
F2	HUCA-1401	Shell	CNA-2820	2210	32	110 BCE - 130 CE	15 CE		Cáceres et al. (2018)
F2	HUCA-1303	Shell	CNA-2819	2189	31	85 BCE - 150 CE	40 CE		Cáceres et al. (2018)
F2	HUCA-1405	Shell	CNA-2823	2172	32	65 BCE - 180 CE	59 CE		Cáceres et al. (2018)
F2	HUCA-1407	Shell	CNA-2825	2170	32	60 BCE - 185 CE	61 CE		Cáceres et al. (2018)
F2	HUCA-1406	Shell	CNA-2824	2150	33	30 BCE - 200 CE	85 CE		Cáceres et al. (2018)
F5	C-10	Shell	CNA-2818					370 CE - 510 CE	This paper

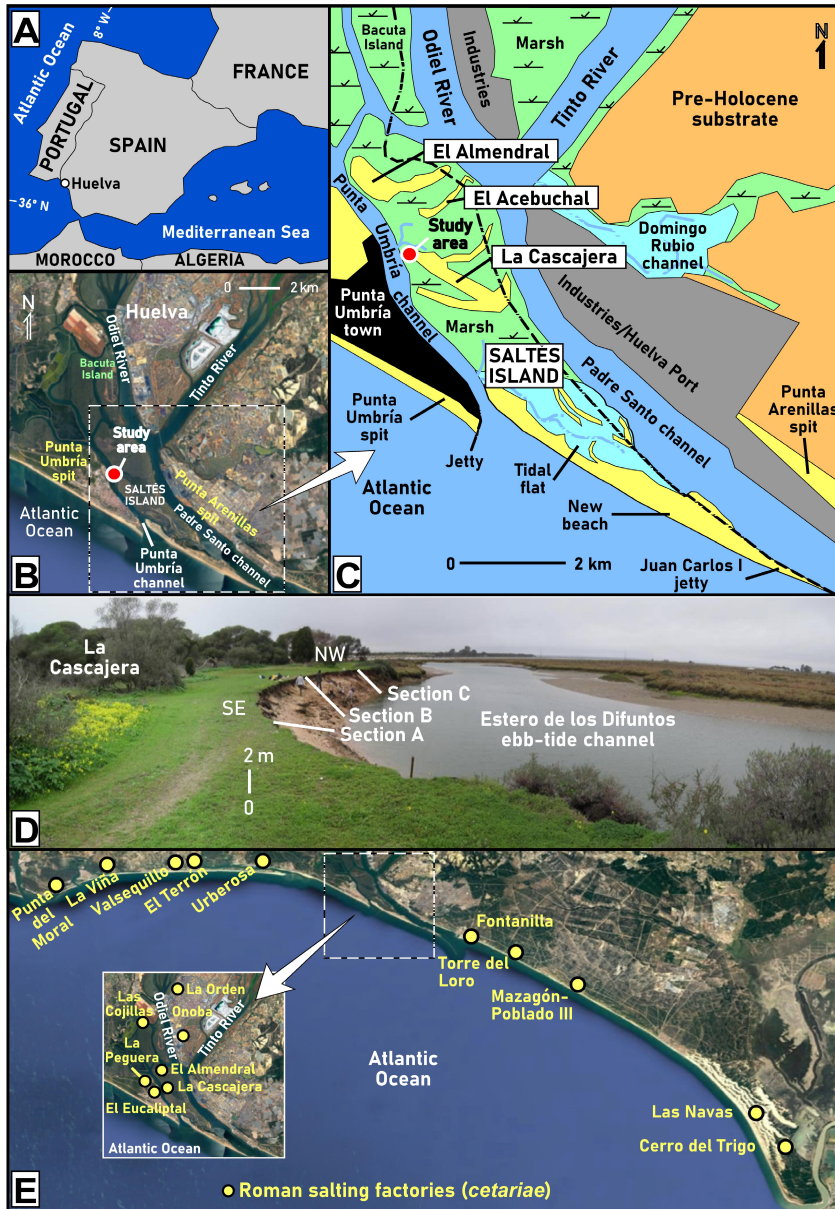


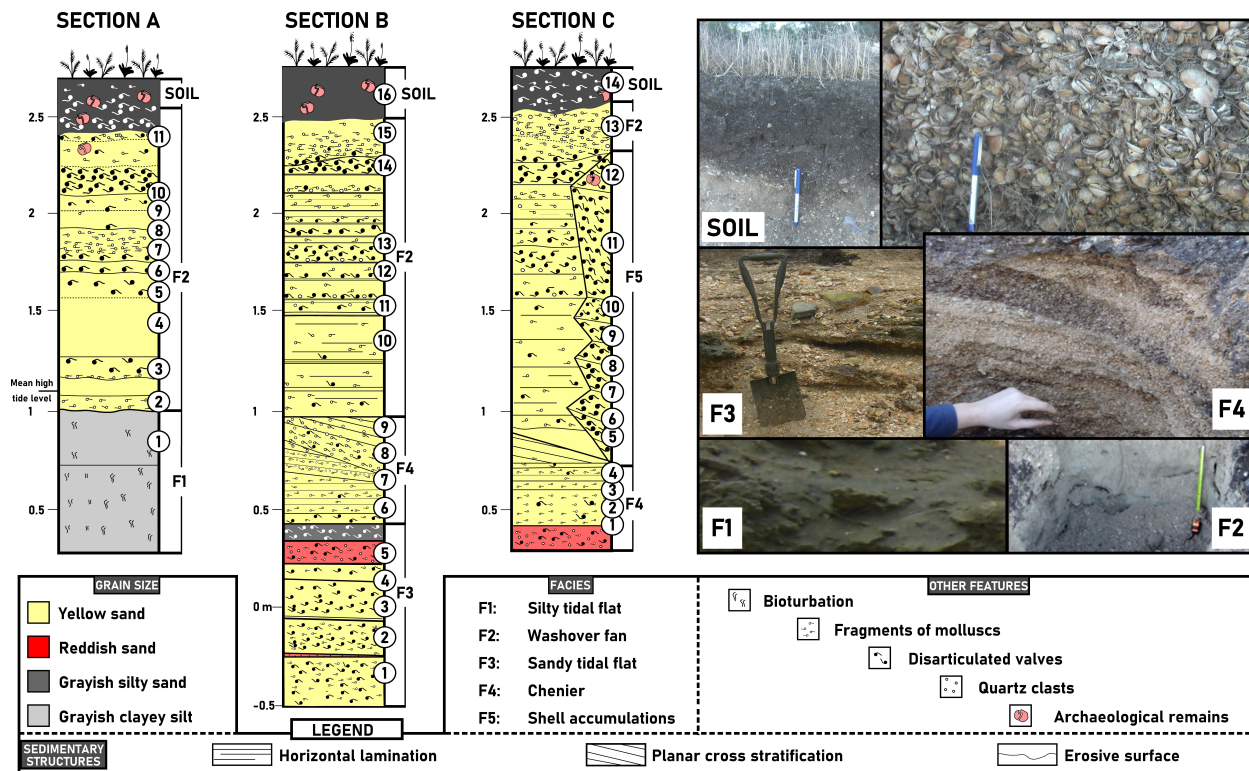






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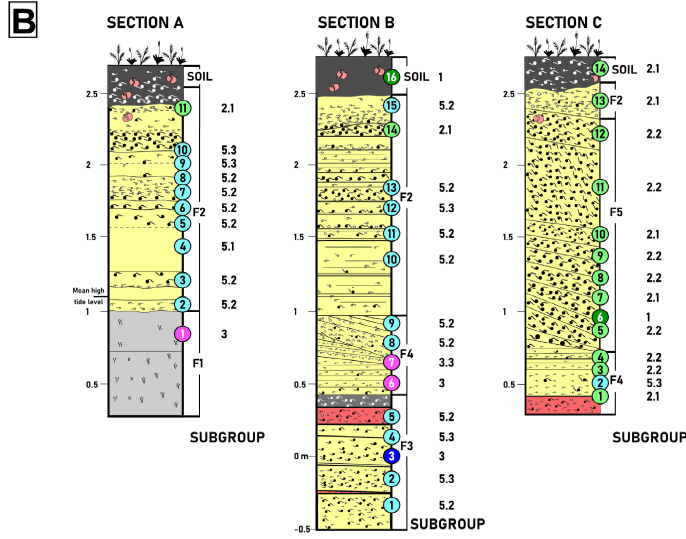
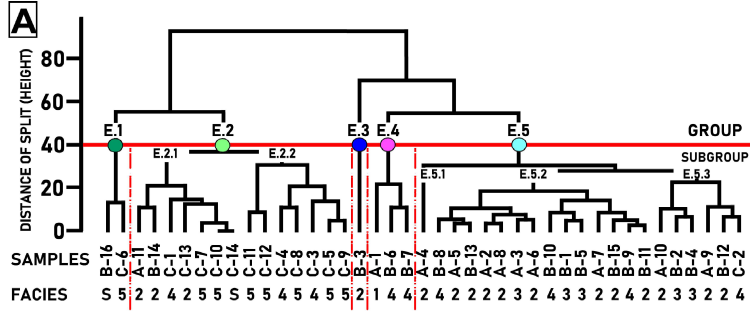






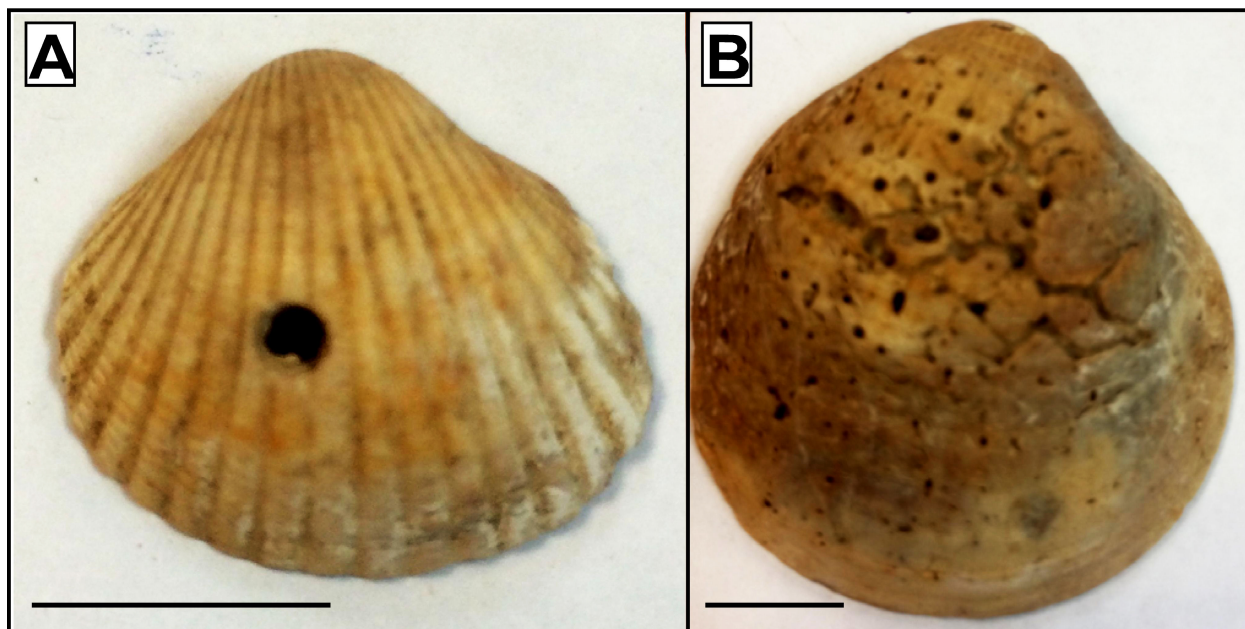




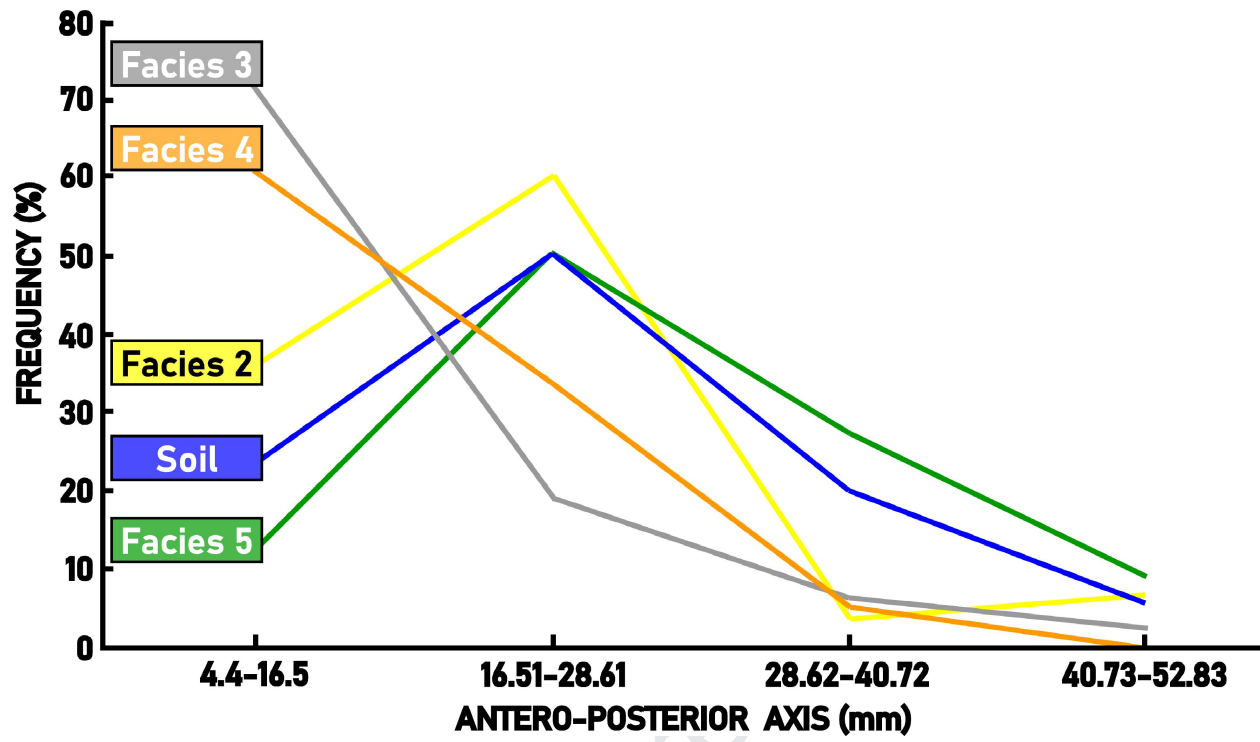


**C**

SPECIES	E.1	E.2		E.3	E.4	E.5		
		E.2.1	E.2.2			E.5.1	E.5.2	E.5.3
<i>Acanthocardia</i> spp.	0.3	0	0.3	0	0	0	0	0
<i>Chamelea gallina</i>	3	13.3	7.7	4	39.6	0	6.4	16.5
<i>Cerastoderma edule</i>	0	0	0.9	0	0	0	0.1	0
<i>Cerastoderma glaucum</i>	0	0.3	2.9	0	0	0	0	0
<i>Donax</i> spp.	0	0	0	2	0	0	0.1	0.3
<i>Glycymeris glycymeris</i>	4	4.4	0.6	52	3.2	0	4.4	5.2
<i>Glycymeris nummaria</i>	72	32.1	47.1	0	2	16.7	2	11.9
<i>Ostrea</i> sp.	3	2.7	12.6	8	0.7	16.7	2.9	4.8
<i>Pecten maximus</i>	0	0	0.3	0	0	0	0	0
<i>Ruditapes decussatus</i>	0	0	0.6	0	0	0	0	0
<i>Spisula</i> spp.	0	0	0	0	0	0	0.7	0.3



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HIGHLIGHTS RUIZ ET AL.

- Multidisciplinary analyzes permit to distinguish shell middens from other shell deposits.
- Percentages of edible species, taphonomy or shell size delimit shell middens.
- Statistics permits to delimitate shell middens from cheniers, washover fans or tidal flats.
- Autoecology of the species delimits the fishing spaces of a Roman factory and the the fishing techniques used.
- Age of shell midden is consistent with that of the nearby archaeological remains.

**Untangling a Roman Atlantic estuary midden:**

**multidisciplinary analysis of shell deposits from Saltés Island (SW Spain)**

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**CONFLICT OF INTEREST**

The authors declare that there is no conflict of interest in the publication of this paper.