

Palaeoenvironmental reconstructions during the Meso- to Neolithic transition (9.2–5.3 cal. ka BP) in Northwestern France: Palynological evidences

Lambert Clément^{1,*}, Vidal Muriel¹, Penaud Aurélie¹, Le Roy Pascal¹, Goubert Evelyne², Pailler Yvan³, Stephan Pierre⁴, Ehrhold Axel⁵

¹ Laboratoire Géosciences Océan (UMR 6538), IUEM, CNRS, Université de Bretagne Occidentale (UBO), France

² Laboratoire Géosciences Océan (UMR 6538), IUEM, CNRS, Université de Bretagne Occidentale (UBO), Université Bretagne Sud, France

³ Grand-Ouest, INRAP, France

⁴ LETG Brest GEOMER, IUEM, UMR 6554, CNRS, Université de Bretagne Occidentale (UBO), France

⁵ Géosciences Marines, Centre de Brest, IFREMER, France

* Corresponding author : Clément Lambert, email address : clement.lambert24@gmail.com

Abstract :

Sedimentological, palynological, and micropalaeontological studies carried out throughout the first half of the Holocene, during the Mesolithic/Neolithic transition in the Bay of Brest (i.e. 9200–9000 and 6600–5300 cal. BP) and in the Bay of Douarnenez (i.e. 9200–8400 cal. BP), allowed characterizing coastal environmental changes under the increasing influence of the relative sea-level rise. The gradual flooding of the two studied sites implied a transition from river valleys to oceanic bays as revealed by the gradual retreat of salt marsh environments, as detected through palynological analysis. In addition, these high-resolution studies highlight the regional imprint of the North Atlantic millennial climate variability in north-western coastal environments. Two cold climate events are indeed suggested to have been locally marked by a moisture increase, mainly detected by increases in *Lingulodinium machaerophorum*, *Corylus*, and *Alnus* percentages at 8550 cal. BP in the Bay of Douarnenez and at 6250 cal. BP in the Bay of Brest. Moreover, regarding the Neolithic transition timing in the Bay of Douarnenez, large pollen grains of Poaceae (i.e. Cerealia-type pollen grains) have been detected at around 8600 cal. BP, that is, 1500 years before the general accepted cereal cropping appearance in Western France. These results, consistent with other palynological studies conducted in the French Atlantic coast, could underline a Mesolithic 'proto-agriculture' in Brittany.

Keywords : benthic foraminifera, climate variability, dinoflagellate cysts, human impacts, palaeoenvironments reconstructions, pollen grains

1. Introduction

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The Holocene is characterized by a gradual decrease in summer air temperatures and seasonality (seasonal thermal amplitude) in line with the continuing summer insolation decrease at 65°N (*Berger and Loutre, 1991*). Sub-millennial climate variability is superimposed on this long-term climate trend, as shown by numerous studies carried out on marine (e.g. *Bond et al., 1997, 2001; Mayewski et al., 2004*), terrestrial (e.g. *Davis et al., 2003*) and glacial records (e.g. *O'Brien et al., 1995*), highlighting iterated abrupt climate events also referred as “Bond events” (*Bond et al., 1997, 2001*) or RCC (i.e. “Rapid Climate Change”; *Mayewski et al., 2004*). These events are often associated with significant climate cooling in North Atlantic surface waters and over northern Europe, as well as increasing aridity in the tropics (*Mayewski et al., 2004; Hammarlund et al., 2005; Wanner et al., 2011*). Finally, on decadal to multi-decadal timescales, climate over northern Europe is forced by the combined influences of atmospheric and oceanic natural oscillations, themselves driven by different teleconnections and physico-chemical exchanges at the air/ocean interface (*Knight et al., 2006; Tréguer et al., 2014; McCarthy et al., 2015; Ruprich-Robert and Cassou, 2015*).

The geographical exposure of the coast to the ocean and North Atlantic climate hazards thus makes the Northwest coast of France a favorite environment for the study of current and past climate. In addition to these factors, coastal areas are constantly changing shaped by different dynamic agents. The sea level-rise occurring across the deglaciation has also impacted coastal environments from a geomorphological, sedimentological and ecological point of view. On a regional scale, recent studies reconstructed and quantified the post-glacial sea-level rise on Brittany’s coasts (*Goslin et al., 2013, 2015; Stéphan et al., 2015; García-Artola et al., 2018*), estimated from about 10 to 15 mm/year between the end of the last glacial period and 9,000 years cal BP to around 4.6 mm/year between 7,500 and 6,500 years cal BP, and less than 1

69 mm/year after 6,000 years cal BP. Moreover, palaeoecological data recently acquired on
70 Holocene Brittany's coastal sequences discussed past environmental variations (vegetation
71 changes, palaeo-storm dynamics, precipitation regimes) and linked these rapid coastal
72 changes with climate dynamics as well as human occupation on watersheds (*Fernane et al.,*
73 *2014, 2015*). These studies especially focused on the Neolithic, a period when the
74 development of agro-pastoral societies became more and more pronounced (*Visset and*
75 *Bernard, 2006; Pailler et al., 2011*).

76 In this study, new sequences retrieved in two sites from the westernmost part of Brittany (NW
77 France), the Bay of Brest (BB) and the Bay of Douarnenez (BD), cover the Mesolithic to
78 Neolithic transition, a still fairly unknown period so far. They allow a multidisciplinary
79 approach based on pollen, dinoflagellate cyst (dinocyst) and foraminiferal analyses, as well as
80 on stable isotopes and sedimentological data. Pollen analysis provides information regarding
81 surrounding vegetation changes on BB and BD watersheds through time. Indeed,
82 palynological studies carried out on modern BB sediments (*Lambert et al., 2017*), as well as
83 across the last 150 years (*Lambert et al., 2018*), evidenced the robustness of studying
84 fossilized pollen grains in BB sediment archives to discuss both natural and anthropogenic
85 forcings. These include: hydrodynamics, fluvial discharges, pollination rates, or agricultural
86 watershed policy. Dinoflagellates are phytoplanktonic organisms that play an important role
87 in the trophic network. Numerous studies carried out on modern marine sediments showed
88 specific patterns regarding the spatial distribution of fossilizable dinocysts according to sea-
89 surface temperature and salinity, sea-ice cover duration, inshore-offshore gradient and/or
90 nutrient concentration (*Morzadec-Kerfourn, 1977, 1979; Dodge et Harland, 1991; Rochon et*
91 *de Vernal, 1994; Mudie et al., 2001; Marret and Zonneveld, 2003; Zonneveld et al., 2013; de*
92 *Vernal et al., 2013*). These palynomorphs have been observed with extremely low
93 concentrations and a poor species richness in modern BB sediments (*Lambert et al., 2017*) but

94 have proved to be excellent markers for reconstructing past BB sea-surface conditions across
95 the last century (*Lambert et al., 2018*). Finally, benthic foraminifera are particularly sensitive
96 to various environmental factors (temperature, salinity, oxygenation, pH, hydrodynamism,
97 organic matter export, water depth) and are frequently used to reconstruct bottom water-
98 column conditions (*Goody, 2003; Jorissen et al., 2007*). In coastal environments,
99 foraminiferal species allow reconstructing natural or human-induced environmental changes
100 (*Debenay et al., 2006; Delaine et al., 2015*), and among the rare studies conducted in the BB
101 with this proxy, *Stéphan (2008)* applied it to discuss palaeobathymetric evolution in salt-
102 marsh environments related to the Holocene sea-level rise (*Stéphan and Goslin, 2014 ;*
103 *Stéphan et al., 2015; García-Artola et al., 2018*).

104 In this paper, we thus aim at discussing the combined influences of local (i.e. watersheds)
105 *versus* regional (i.e. northern European climate) factors driving paleoenvironmental changes
106 across the first half of the Holocene in western Brittany (from 9,200 to 5,300 cal BP), thanks
107 to a cross-correlated pollen-dinocyst-foraminiferal analysis and a pluri-decadal timescale
108 resolution. Furthermore, the studied period covers the Mesolithic to the Neolithic transition,
109 where the environmental changes thanks to the Neolithization must be detectable in
110 paleoenvironmental sequences.

111 **2. Environmental contexts**

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113 **2.1. Present and past sedimentological contexts of study sites**

114 The cores selected for this study are located in the westernmost part of Brittany, in marine
115 coastal bays located on both sides of the Crozon peninsula (Fig. 1a).

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117 **The Bay of Douarnenez (BD)**

118 The BD is a large circular bay (350 km²) limited by the Crozon peninsula to the North and the
119 Cap Sizun to the South, largely connected to the Iroise Sea by a passage of about 9 km wide
120 (Fig. 1b). Its bathymetry reveals a concave morphology characterized by a water depth of
121 about 20 m, reaching 30 m in the center of the bay (*Augris et al., 2005*; Fig. 1b). The
122 geological basement of the BD and its surroundings corresponds to Brioverian/Palaeozoic
123 sandstones and schist in the North, and metamorphic/plutonic formations in the South (*Mélou
124 and Plusquellec, 1974; Augris et al., 2005*). A major NW-SE trending fault network
125 (Kerforne fault system) affects the BD, inherited from the Hercynian orogeny and reactivated
126 during the North Atlantic opening (*Lefort, 1973; Ballèvre et al., 2013*). Coasts are mainly
127 shaped by cliffs up to 100 m high in the Crozon peninsula and 85 m towards the Cap Sizun.
128 Some valleys and small estuaries drain very restricted watersheds (40 km² for the larger),
129 sometimes open between cliffs, protecting wetlands and marshes behind the shoreline (*Augris
130 et al., 2005*). The BD does not receive a large amount of freshwater, implying a strong marine
131 influence with a maximum tidal amplitude of 8 m and a relatively stable salinity (around
132 35‰; *Augris et al., 2005*).

133 The BD sedimentary cover mainly consists of gravels and coarse sands in the central part of
134 the bay and fine sands in sheltered areas (*Hinschberger and Pauvret, 1968; Augris et al.,
135 1988*), reaching a thickness of 18 m in palaeo-valley axes, but remaining relatively thin over
136 the whole BD (2 to 4 m; *Augris et al., 2005*). The substratum study reveals a dendritic palaeo-
137 channel network incising Brioverian basement and flowing into a main valley in the center of
138 the bay named the Ys palaeo-valley (Fig. 1b). These channels are connected to the main
139 continental rivers surrounding the BD (*Musset, 1934; Guilcher, 1948; Hallegouët, 1989;
140 Jouet et al., 2003*). The sedimentary infilling history of the BD is based on the analysis of
141 seismic units (U1 to U6; Fig. 2) combined with analysis of numerous sediment cores (*Jouet et*

142 *al.*, 2003; *Le Roy and Jouet*, 2005). While the BD was totally emerged during the Last Glacial
143 Maximum (around 20,000 years BP), its flooding, initially confined to palaeo-valleys
144 (10,000-8,000 years BP, meander bar units U3 and U4; Fig. 2), then spread out to the rest of
145 the BD (8,000-5,000 years BP, fine estuarine sedimentation U5; Fig. 2). The sea level was
146 stabilized at about 6,000-5,000 years BP (*Goslin et al.*, 2015) and, since then, reworked
147 marine sands form the upper unit of the BD sediment cover (U6; Fig. 2).

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149 **The Bay of Brest (BB)**

150 The BB is located in north-western Brittany (NW France; Fig. 1c) and consists of a shallow
151 semi-enclosed basin of 180 km² surrounded by a 230 km long coastline. Its basement
152 corresponds to Proterozoic igneous rocks in the north and Brioverian/Palaeozoic sediments in
153 the south and east. Present-day low reliefs (few hills reach 330 m high) are inherited from the
154 peneplanation of the Hercynian chain (*Chauris and Plusquellec*, 1980; *Ballèvre et al.*, 2009).
155 The study area has been subsiding since the Eocene and still today (0.02 to 0.04 mm/yr ;
156 *Ziegler*, 1992; *Bonnet et al.*, 2000; *Goslin*, 2014), but it can be considered as negligible at the
157 Holocene timescale (40 mm/10,000 years ; *Goslin et al.*, 2015). The river system is
158 established since the Tertiary (*Hallegouët et al.*, 1994). Today, the BB is characterized by a
159 macrotidal influence with a maximum tidal amplitude of 8 m (*Troadec and Le Goff*, 1997).
160 Grain size analyses of modern sediments reflect specific hydrodynamic conditions mainly
161 related to tidal currents (*Gregoire et al.*, 2016). In its westernmost part, the BB is connected to
162 the Atlantic Ocean through the “*Goulet*” (Fig. 1c), a strait of about 1.8 km wide and 50 m
163 deep. In its easternmost part, the BB receives main freshwater supplies from the Aulne and
164 Elorn Rivers (both contributing up to 85% of the total river discharges; *Delmas and Treguer*,
165 1983), as well as from the smaller Daoulas River (Fig. 1c). BB watersheds are characterized
166 by 2,000 km of waterways and most of their runoff flows into the BB through the Aulne River

167 (114 km long; 1,224 km² of watershed and 20.40 m³/s of annual debit; *Troadec and Le Goff,*
168 *1997*).

169 The bathymetric map of the BB highlights submarine channels that attest to palaeo-fluvial
170 systems (Fig. 1c; *Gregoire et al., 2016*). Palaeo-channels of the two main current rivers,
171 Aulne and Elorn, are about 30 and 15 m deep respectively, and converge in the west at a
172 trough of about 50 m deep (*Troadec and Le Goff, 1997*). Nevertheless, at present, the depth of
173 the BB does not exceed 10 m deep on approximately 60% of its surface (*Monbet and*
174 *Bassoulet, 1989*). The last transgressive episode corresponds to the complex and fragmented
175 sedimentary infilling history of the BB (*Gregoire et al., 2017*), because of i) the non-
176 morphological uniformity of the substratum, ii) the non-linear post-glacial sea level rise,
177 combined with iii) strong and complex hydrodynamical features (*Gregoire et al., 2017*).
178 Palaeo-valleys became flooded according to different steps of large fluvial terraces (*Gregoire*
179 *et al., 2017*), considerably decreasing surfaces occupied by intertidal salt-marshes and
180 mudflats (i.e. typical environments submitted to intertidal dynamics); the shallowest parts of
181 the BB being flooded between 9,000 and 7,000 years BP. Today, the centre of the BB is
182 predominantly under marine influence, main river mouths and intertidal areas having
183 migrated further east.

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185 **2.2. Climatic Context**

186 Brittany is subjected to a temperate oceanic climate regime characterized by the influence of
187 the westerlies and by low seasonal thermal amplitudes with mean annual temperatures of
188 about 10-11°C (*Belleguic et al., 2012*). Annual prevailing winds (with speeds that can exceed
189 100km/h during 5 to 15 days per year) have mainly a south-west origin (*Troadec and Le Goff,*
190 *1997*). Since Brittany is submitted to regular oceanic rainfalls, annual cumulative precipitation

191 data ranges from 600 mm/year to more than 1,600 mm/year in the inner continental part
192 (*Troadek and Le Goff, 1997; Belleguic et al., 2012*). The climate of Brittany is due to the
193 combined influences of atmospheric (North Atlantic Oscillation, NAO) and oceanic (Atlantic
194 Multidecadal Oscillation, AMO) circulations detailed thereafter (*Tréguer et al., 2014,*
195 *Ruprich-Robert and Cassou, 2015; Lambert et al., 2018*). On decadal to multi-decadal
196 timescales, variations in North Atlantic sea-surface temperature (SST) control a large part of
197 the climate variability reconstructed on the continent (*Deser et al., 2010; Knight et al., 2006;*
198 *McCarthy et al., 2015; Ruprich-Robert and Cassou, 2015*), already discussed in the BB
199 watersheds (*Lambert et al., 2018*).

200

201 **2.3. Cultural evolution and territorial occupation dynamics**

202 The Mesolithic is poorly documented in western Brittany due to the scarcity of archeological
203 data. The French Mesolithic period is divided into two sub periods, the first Mesolithic (10th,
204 9th, and 8th millennia BC) and the second Mesolithic (7th, 6th and early 5th millennia BC,
205 depending on the studied region) according to cultural criteria based on lithic industries
206 (*Marchand, 2014*). Across the Mesolithic, traces of shellfish consumption as well as of
207 microlithic industries on the coastline suggest the presence of small communities irregularly
208 spread over western Brittany (perhaps 25,000 to 50,000 people; *Giot et al., 1998*). Also, lithic
209 material resulting from debitage has been found in Finistère (western Brittany), evidencing
210 human settlements 20 kilometers from the current coast. Human groups may have moved
211 seasonally between coasts, riversides and the inner land (*Gouletquer et al., 1994, 1996*), while
212 recent studies rather suggest perennial settlements of restricted communities in some coastal
213 areas (*Schulding and Richards, 2001; Marchand, 2005*). Previous pollen studies have not
214 shown significant environmental impacts in Brittany due to human hunter-gatherer
215 populations (*Morzadec-Kerfourn, 1974*).

216 The Neolithic "revolution" then reached western Europe around 5,500 BC (7,450 BP) and the
217 Armorican Massif around 5,000 BC (6,950 BP; *Blanchet et al., 2010*) via Danubian
218 agricultural populations, evidenced by the western extension of the Linear Pottery Culture
219 (i.e. *Blicquy-Villeneuve-Saint-Germain, in Marcigny et al., 2010*), and particularly in
220 Southern Finistère (*Marchand et al., 2006; Marcigny et al., 2010; Tinévez et al., 2015*)
221 through a diffusion model estimated at 1 km/year (*Giot et al., 1998*). The contact between
222 Mesolithic and Neolithic societies remains poorly documented, and predation practices
223 gradually shifted to a production economy thanks to the domestication of animals and the
224 emergence of agriculture. For many years, these cultural changes have been debated as the
225 result of i) the acculturation of indigenous hunter-gatherer populations, or ii) a population
226 replacement by Neolithic societies coming from the East. Recent genetic studies conducted on
227 European populations suggest a Mesolithic population replacement, nevertheless highlighting
228 a mosaic of *scenarii* according to regions, and a non-uniform demographic transition across
229 Europe with few evidences of cultural adoption (*Sampietro et al., 2007 ; Bramanti et al., 2009*
230 ; *Haak et al., 2010; von Cramon-Taubadel and Pinhasi, 2011 ; Skoglund et al., 2012*).
231 Western Europe population increased sharply as a result of increasing food production.
232 Perennial settlements of Neolithic villages are observed in western France from the first
233 centuries of the 5th millennium BC (around 7,000 to 6,700 BP) (*Marchand, 2014*).

234 Considering the emergence of agriculture, palynological and palaeobotanical data are very
235 rare in Finistère. Some studies carried out over Brittany are mainly based on lithic industries
236 and funeral practices (*Marchand, 2005*), and the beginning of the neolithisation thus cannot
237 be precisely dated, such as in other parts of Europe (*Kirleis et al., 2012; McClatchie et al.,*
238 *2014*). In the coastal Morbihan (Locmariaquer, SE Brittany), *Visset et al. (1996)* suggested
239 ancient cereal farming occurred around 7,243-5,800 BC (9,193-7,750 BP) because of cereal
240 pollen grain occurrences in Mesolithic levels, observed concomitantly with increases of

241 ruderal plant and *Corylus* percentages, as well as a marked decrease of other arboreal pollen
242 taxa. Other palynological studies also discussed the presence of cultivated taxa since the
243 Mesolithic in SE Brittany and around the Loire valley (*Barbier and Visset, 1997; Ouguerram*
244 *and Visset, 2001; Visset et al., 2002; Joly and Visset, 2005*). However, the absence of
245 agricultural tools in archaeological sites in the direct vicinity of the above-mentioned sites did
246 not allow confirming these assumptions based on palynological observations (*Marchand,*
247 *2005*). Since the early Neolithic, anthropogenic environmental disturbances and evidences of
248 cereal-cropping became more and more prevalent (*Visset, 1979; Marguerie, 1991*). In
249 Finistère, the cereal cultivation practice is attested from 3,500 BC (5,450 BP) by the presence
250 of cereal grains in the archeological site of Molène island (*Dréano et al., 2007*). On the other
251 hand, human impact on the environment becomes noticeable from the first half of the 5th
252 millennium BC with the first appearance of grindstones and more particularly with traces
253 detected on lithic furniture typical from the cutting of plant fibers (*Giot et al., 1998; Guéret et*
254 *al., 2014*).

255 **3. Material and methods**

256 **3.1. Study sediment cores**

257 Pictures and RX radiography were carried out at the *Géosciences Marines* laboratory
258 (IFREMER, Plouzané, France) for the three study cores (Fig.1), and description,
259 granulometric analyses (laser granulometer “MASTERSIZER 2000”), as well as sampling for
260 AMS ¹⁴C dates and palynological analyses were carried out at the *Géosciences Océan*
261 laboratory (LGO-IUEM, Plouzané, France). All dates (Table 1) have been calibrated with the
262 CALIB 7.1 program using the Marine13 calibration curve (*Stuiver and Reimer, 1993; Reimer*
263 *et al., 2013*) and a ΔR of -40 ± 23 years (*Mangerud et al., 2006*). In the manuscript, when
264 referred to a precise age, the mention to “Cal.” will not systematically be written, and ages in

265 “years BP” or “years BC” will often be both specified so as to facilitate the lecture of the
 266 results for paleoenvironmental or archeological communities, respectively.

267 Within the BD, core “VC2012-08-PQP” (48°10.2’N, 04°26.4’W; 28 m depth; 486 cm length;
 268 Fig. 1b, 3, 4) was recovered with a vibrocorer by the R/V *Pourquoi Pas?* during the Proteus-
 269 Dunes cruise (Shom, 2012). Within the BB, core “A” (48°19.2’N, 4°31.8’W; 8.2 m depth;
 270 418 cm length in total but 318 cm available for this study: the top 100 cm were entirely used
 271 in 2003-2004 by biologists and bio-geochemists) was retrieved in the Bay of Roscanvel (i.e.
 272 small bay in the western part of the BB; Fig. 1c, 5) also with a vibrocorer by the R/V *Côtes de*
 273 *la Manche* during the “*Défis Golfe de Gascogne*” program (IFREMER, LEMAR-IUEM,
 274 2003). Finally, core “KS-24” (48°19.3’N, 4°31.4’W; 26 m depth; 181.5 cm length) was
 275 retrieved in the Bay of Roscanvel (BR; Fig. 1c, 5) with a gravity corer by the R/V *Thalia*
 276 during the “*SERABEQ 3*” cruise (IFREMER, 2015).

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| Lab code | Depth (cm) | material | Age ¹⁴ C | Error | Age Cal. BP Min-max (mean) | Age Cal.AD/BC |
|------------------|------------|-----------------------------|---------------------|-------|----------------------------------|------------------|
| Core VC08 | | | | | | |
| Poz-85161 | 20 | Gastropod | 1210 | 30 | 702-894 (798) | 1152 AD |
| SacA47754 | 23.5 | Bivalve | 635 | 30 | 257-420 (338.5) | 1611.5 AD |
| SacA47756 | 103 | Bivalve | 7905 | 45 | 8313-8524 (8418.5) | 6468.5 BC |
| SacA47755 | 158 | Bivalve | 8025 | 40 | 8401-8628 (8514.5) | 6564.5 BC |
| SacA43117 | 240 | <i>Acanthocardia</i> sp. | 8030 | 35 | 8410-8625 (8517.5) | 6567.5 BC |
| SacA43118 | 266.5 | Bivalve | 8060 | 35 | 8433-8688 (8560.5) | 6610.5 BC |
| SacA43119 | 337.5 | Bivalve | 8090 | 30 | 8483-8737 (8610) | 6660 BC |
| SacA49420 | 345 | Bivalve | 8105 | 35 | 8492-8774 (8634.5) | 6684.5 BC |
| SacA43120 | 375.5 | <i>Bittium</i> sp. | 8580 | 60 | 9091-9435 (9263) | 7313 BC |

| | | | | | | |
|------------------|-----------------|---|-------------|-----------|-------------------------------|------------------|
| SacA43115 | 405 | <i>Spisula</i> sp. | 8960 | 35 | 9532-9827 (9679.5) | 7729.5 BC |
| SacA43121 | 408 | <i>Ostrea</i> sp. | 8950 | 35 | 9527-9810 (9668.5) | 7718.5 BC |
| Core KS24 | | | | | | |
| Poz-78151 | 102.5 | Gastropod | 8410 | 50 | 8938-9247 (9092.5) | 7142.5 BC |
| Poz-78152 | 173.5 | Gastropod | 8530 | 50 | 9045-9383 (9214) | 7264 BC |
| Core A | | | | | | |
| Poz-42799 | 18-19 | <i>Venerupis senegalensis</i> | 4930 | 35 | 5221-5445 (5333) | 3383 BC |
| SacA49424 | 46 | <i>Turritella</i> sp. | 5390 | 30 | 5704-5899 (5801.5) | 3851.5 BC |
| SacA41585 | 70 | Bivalve | 5375 | 30 | 5686-5890 (5788) | 3838 BC |
| SacA43114 | 109-110 | <i>Venus</i> sp. | 5540 | 40 | 5873-6114 (5993.5) | 4043.5 BC |
| Poz-42840 | 114.5- 116.5 | <i>Venerupis senegalensis</i> | 5580 | 35 | 5911-6149 (6030) | 4080 BC |
| Poz-42841 | 161 | <i>Dosinia lupinus</i> | 5690 | 40 | 6005-6258 (6131.5) | 4181.5 BC |
| Poz-42842 | 187 | <i>Patia rhomboides</i> | 5790 | 40 | 6159-6365 (6262) | 4312 BC |
| SacA41583 | 190-191 | Bivalve | 5715 | 30 | 6059-6273 (6166) | 4216 BC |
| SacA41584 | 190-191 | Organic matter For the reservoir age | 5350 | 60 | 5993-6280 (6136.5) | 4186.5 BC |
| Poz-42843 | 229-231 | <i>Venerupis senegalensis</i> | 5900 | 40 | 6264-6457 (6360.5) | 4410.5 BC |
| SacA41581 | 248-249 | Bivalve | 5990 | 30 | 6335-6557 (6443) | 4496 BC |
| SacA41579 | 258 | Bivalve | 5960 | 30 | 6303-6508 (6405.5) | 4455.5 BC |
| SacA41580 | 297 | Bivalve | 6105 | 30 | 6464-6677 (6570.5) | 4620.5 BC |
| Poz-42844 | 316 | <i>Venerupis senegalensis</i> | 6155 | 35 | 6510-6744 (6627) | 4677 BC |

278 Table 1: carbon dates made on VC-08, A and KS-24 cores. The 190 cm level on core A, dated
279 from continental organic matter, is not considered in the age model. It was duplicated from
280 the level 190 cm dated on bivalve to estimate the reservoir age in the Bay of Brest.

281 **3.2. Palynological analyses**

282 Palynological preparations were carried out at the EPOC laboratory (University of Bordeaux,
283 Talence), following the procedure described by *de Vernal et al. (1999)* and using chemical
284 (cold HCl and cold HF) and physical (sieving through a 10 µm nylon mesh screen) treatments
285 in order to remove the mineral fraction and to concentrate palynomorphs (cf.
286 http://www.epoc.u-bordeaux.fr/index.php?lang=en&page=eq_paleo_protocoles). The final
287 residue was mounted between slide and coverslip with glycerin. Pollen and dinocysts were
288 determined using an optical microscope Leica DMC 2900 at X630 magnification.

289 In this study, 89 samples (46 for core “A”, 4 for core KS-24 and 39 for core VC-08) were
290 analyzed allowing to obtain a study resolution of about 20 to 30 years. Palynomorph
291 identification followed *Beug (1961)*, *Faegri and Iversen (1989)*, *Moore et al. (1991)* and
292 *Reille (1995)* for pollen and *Rochon et al. (1999)* for dinocysts. For each analyzed sample, a
293 minimum of 300 pollen grains and 150 dinocysts have been counted in order to provide robust
294 assemblages from a statistical point of view (*Fatela and Taborda, 2002*). Percentages were
295 calculated on a sum of total pollen grains, or a sum of total dinocysts, without any exclusion.
296 Concentrations (number of specimens/cm³) were obtained thanks to the *Lycopodium* spore
297 method (*Mertens et al., 2009*). Through a camera connected to the optical microscope,
298 measurements were also performed on the size of the annulus and of the diameter of Poaceae
299 grains along the VC-08 core so as to differentiate *Cerealia* from wild grasses (*Leroyer et al.,*
300 *2004; Joly et al., 2007*). Finally, other palynomorphs were counted, including microalgae,
301 foraminiferal linings, spores and copepod eggs.

302

303 **3.3. Foraminiferal analyses**

304 Prior to the palynological treatments, bulk sediments were sieved at 150 μm , and the fraction
305 $>150 \mu\text{m}$ was used for foraminiferal analyses. After dividing the largest samples with a
306 micro-splitter, benthic foraminiferal assemblages were analyzed with a LEICA M60 binocular
307 stereo zoom microscope at X60 magnification. Identification followed *WoRMS Editorial*
308 *Board (2017)* (www.marinespecies.org). Data were expressed in percentages of total benthic
309 foraminiferal counts (at least 100 specimens counted per level). 16 samples (12 for core “A”
310 and 4 for core KS-24) were analyzed allowing a resolution of about 150 years.

311 Monospecific stable oxygen and carbon isotopes were also measured for core “A” on the
312 *Elphidium aculeatum* benthic species, in the same levels than benthic foraminiferal
313 assemblages. About five specimens were hand-picked in the 150-250 μm sediment fraction,
314 cleaned in a methanol ultrasonic bath for a few seconds, then roasted under vacuum at 380 $^{\circ}\text{C}$
315 for 45 min to remove organic matter, prior to isotopic analyses (*Duplessy, 1978*). The $\delta^{18}\text{O}$
316 and $\delta^{13}\text{C}$ (expressed in ‰ VPDB) were measured at the PSO (IUEM, BREST) using the
317 IRMS platform: a Delta V mass-spectrometer coupled with a GasBench II preparation line for
318 benthic species. The external reproducibility (1σ) of an internal standard calibrated with
319 NBS19 is $\pm 0.03 \text{ ‰}$ and 0.06 ‰ for $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ respectively.

320 **4. Sedimentary context and palynological results**

321 **4.1. Bay of Douarnenez, BD (core VC-08)**

322 *Sedimentary facies and age model*

323 The sedimentology, granulometry and magnetic susceptibility analyses (Fig. 3, 4), core VC-
324 08, taken from a BD palaeo-valley (Fig. 1b) highlight different sedimentary deposits that can
325 be related to the sediment units defined in the framework of the study of the sedimentary
326 infilling history of the BD (*Le Roy and Jouet, 2005*; Fig. 2).

327 The base of core VC-08 (from 474 to 425 cm) is made of coarse sands (mean granulometry of
328 about 1,500 μm) with high magnetic susceptibility (MS) values illustrating a strong detrital
329 terrigenous component (Fig. 4a), that can be related to units 1 and 2 (U1 and U2 of *Le Roy*
330 *and Jouet, 2005*; Fig. 2, 4b) and interpreted as fluvial deposits during the last glacial period.
331 Between 425 and 340 cm, granulometry oscillates from coarse silts to very fine sands marked
332 by a large grain-size variability (from 5 to around 1,500 μm ; Fig. 4a,b), and a fairly high
333 concentration of broken bivalve shells, therefore probably reflecting high energetic
334 conditions. The drop of MS values is related to major environmental change from continental
335 to marine influences, with the sea starting to reach the westernmost part of the channels
336 around 10,000 years BP. This interval can be associated to units 3 and 4 (U3 and U4 of *Le*
337 *Roy and Jouet, 2005*; Fig. 2, 4b). Between 340 and 30 cm, clayey sediments (mean
338 granulometry of 10 μm ; Fig. 4a,b) and tidal laminae dominate sedimentary facies. After a
339 small increase, MS values gradually decrease from 250 cm to the top (Fig. 4a). This very
340 thick deposit constitutes most of the sedimentary infilling of BD channels and corresponds to
341 unit 5 (U5 of *Le Roy and Jouet, 2005*; Fig. 2, 4b). The last sedimentological deposit (30 cm
342 upwards), delimited at its base by an erosive surface, is characterized by marine sands (mean
343 granulometry of 100 μm , three main modes between 10 and 1,000 μm ; Fig. 4a,b) reworked
344 and remobilized under the action of waves. Extremely low MS values suggest the scarcity of
345 fluvial detrital inputs within the bay. This deposit is related to the last unit 6 (U6 of *Le Roy*
346 *and Jouet, 2005*; Fig. 2, 4b).

347 Radiocarbon dates have been obtained on bivalve and gastropod shells (Table 1). The age
348 model (Fig. 6a) was then established from two linear regressions and 9 stratigraphic pointers
349 considering two parts obviously distinct considering the description of the core (Fig. 4b, 6a).
350 A significant change in sedimentation rates is therefore observed around 8,600 years BP (i.e.
351 340 cm), related to the Holocene sedimentary infilling history, with 0.06 cm/yr calculated

352 before 8,600 years BP and 1.2 cm/yr calculated after (Fig. 4, 6a). Two additional dates
353 obtained in the uppermost part of the core (i.e. 20 and 23.5 cm, 798 and 338.5 BP,
354 respectively; Fig. 4b) confirm the recent set up of the last unit made of reworked marine sands
355 (U6 in *Le Roy and Jouet, 2005*). These two dates were therefore not used to build the age
356 model (Fig. 6a). The palynological study is conducted on the fine intertidal sedimentation
357 section of the core, corresponding to units U4 and U5 described by *Le Roy and Jouet (2005)*
358 (Fig. 3, 4b) and ranging from 390 to 30 cm (i.e. 9,400 to 8,400 years BP or 7,450 to 6,450
359 years BC), thus providing information on the final Mesolithic.

360

361 **Dinocysts**

362 A total of 18 taxa were recognized with an average of nine different taxa per slide. Only main
363 taxa (i.e. greater than 2 % at least once in palynological spectra) were plotted on Fig. 7a along
364 with concentrations (cysts/cm³) and specific richness (number of different taxa per slide).
365 Taken as a whole, *Spiniferites bentorii* (50 % in average) and *Lingulodinium*
366 *machaerophorum* (11 %) are the most abundant taxa as it is usually described on Brittany's
367 coasts (*Morzadec-Kerfourn, 1977; Lambert et al., 2017*). The variations of the dinocyst
368 assemblages enable to highlight three main zones, with very low diversity values at the base
369 of the record (VC-A, 390-345cm), followed by a slow increase up to 250 cm (VC-B, 345-250
370 cm), and higher values over the entire second half of the core (VC-C, 250-30 cm). Diversity
371 then shifts from around 5 to 12 different taxa per slide. In addition to the low dinocyst specific
372 richness, VC-A palynozone is defined by atypical elevated percentages of *Achomosphaera*
373 spp., suddenly replaced by *S. bentorii*. The VC-A / VC-B boundary is well marked in dinocyst
374 assemblages by the occurrences of *Spiniferites mirabilis*, *Spiniferites ramosus* and *Spiniferites*
375 *delicatus*. Moreover, palynozone VC-B corresponds to increasing percentages of *L.*
376 *machaerophorum* and a slow decrease of *S. bentorii* ones. Finally, palynozone VC-C

377 corresponds to a disappearance of *S. mirabilis* simultaneously observed with decreasing
378 percentages of *Selenopemphix quanta*, and the first occurrence of *Spiniferites lazus* found in
379 the study core (Fig. 7a). Moreover, these biozones are highly related to the variations of
380 dinocyst concentrations.

381 Pollen

382 A total of 33 taxa were recognized with an average of 12 different taxa per slide. This rather
383 low diversity due to the forest taxa dominance differs from the greater diversity observed in
384 more recent Bay of Brest pollinic sequences (e.g. Lambert *et al.*, 2017, 2018) and resulting
385 from the herbaceous diversification accompanying the significant anthropogenic landscape
386 opening. In addition to the taxa characterized by percentages found at least once above 2% in
387 assemblages, some scarce but meaningful elements of the vegetation (i.e. *Alnus*, *Cerealia*-
388 type) are plotted on Fig. 7b. In general, percentages of tree pollen grains are relatively stable
389 then representing the most abundant vegetation group (average of 90 %). They are
390 predominantly represented by *Quercus* (42 %) and *Corylus* (33 %), both taxa being then
391 logically anti-correlated all along the core, and accompanied by low percentages of *Pinus*,
392 *Ulmus*, *Betula* and *Alnus*. Among herbaceous plants, Chenopodiaceae are clearly dominant (7
393 %) as well as Poaceae (2 %).

394 The overall diagram displays weak variations along the core except for total pollen
395 concentrations and diversity that allow identifying two parts corresponding first to both VC-A
396 and VC-B palynozones described from the dinocysts (that we will refer to palynozone VC-
397 A&B in the following description) and, second, to VC-C palynozone (Fig. 7b). The limit
398 between VC-A&B and VC-C is more specifically marked by a sharp increase in pollen
399 concentrations and specific richness. Palynozones VC-A&B also correspond to a decreasing
400 trend of *Corylus*, correlated to a *Pinus* increase, then replaced by *Quercus*. In parallel, the
401 pollen specific richness tends to increase. The palynozone VC-C is marked by oscillations of

402 the two major taxa, *Corylus* and *Quercus*, up to 30%, contrasting with the lower part of the
403 diagram. Concentration peaks (dotted lines in Fig. 7b) correspond to increases of *Corylus*
404 percentages and related decreases of *Quercus* ones. In addition, a decrease of *Ulmus* is
405 noticed throughout the sequence, along with a slight increase of herbaceous plants (e.g.
406 Poaceae, Asteraceae, Brassicaceae). Also, *Cerealia*-type are more pronounced in palynozone
407 VC-C. We can note again the overlap between biozones and variations in pollen
408 concentration, as for dinocysts.

409

410 **Towards a Holocene sedimentary infilling history reconstruction of the BD**

411 The sedimentological and palynological analyses conducted on core VC-08 (depicted in depth
412 in Fig. 7 and in age in Fig. 8) allow to clarify the sedimentary infilling model discussed by
413 *Jouet et al. (2003)* and *Le Roy & Jouet (2005)*.

414 From 474 to 420 cm (Fig. 4b), sediments are too coarse to be sampled for palynological
415 analysis. This part could be related to the bedrock incision by rivers during the end of the last
416 glacial period (U1 and U2; Fig. 2), as also evidenced in our study by high magnetic
417 susceptibility (MS) values (Fig. 4a). Palynological results highlight a high Pollen/Dino ratio
418 from 390 to 340 cm (8,630 years BP) (palynozone VC-A; Fig. 8) suggesting a major pollen
419 contribution of surrounding watersheds and therefore predominant fluvial *versus* marine
420 inputs, corresponding then to the fluvial deposits (fluvial accretion bars of U3-U4 in *Le Roy*
421 *and Jouet, 2005*; Fig. 2 and 4b). The Chenopodiaceae content (Fig. 7b) is likely a clue of the
422 already settled salt-marshes. Still discrete marine influences do not allow dinoflagellates to
423 occur in large number during palynozone A, the study core being located upstream in the
424 palaeo-river network incising the substratum of the BD (Fig. 1b). Unfortunately, the ecology

425 of *Achomosphaera* spp. is not precisely known and the taxa thus cannot be connected to
426 particular ecological conditions at that time.

427 From 340 to 250 cm (8,630 to 8,550 BP, palynozone VC-B, Fig. 7 and 8, lower part of U5 in
428 Fig. 4b) the marine influence is first marked by the increased dinocyst diversity, including
429 euryhaline taxa such as *L. machaerophorum* and *S. belerius*, together with taxa more
430 characteristic of marine environments such as *O. centrocarpum*, *S. delicatus*, *S. mirabilis* and
431 *S. ramosus* (Morzadec-Kerfourn, 1977, 1979). In parallel, decreasing percentages of the
432 typically coastal species *S. bentorii* is probably more related to increasing percentages of the
433 other taxa. The lower part of U5 corresponds to a thick transitional sequence between
434 continental and marine influences, submitted to the relative sea-level rise as observed through
435 the continuous decrease of the Pollen/Dino ratio (Fig. 8). The already mentioned breakdown
436 in sedimentation rates at 340 cm (limit between U4 and U5; Fig. 4, 6a) corresponds to this
437 major transition towards an estuarine sedimentation. Variations in particle size and in
438 sediment facies are also obvious through the sudden jump in MS values, indicating greater
439 continental intakes from 8,600 years BP (Fig. 4a, 10).

440 From 250 cm (palynozone VC-C in Fig. 7 and 8, U5 upper part in Fig. 4b), marine influences
441 are well established with a low Pollen/Dino ratio (Fig. 8), a high dinocyst diversity and the
442 persistence of typically coastal (i.e. *S. bentorii*) and marine (i.e. cysts of *P. dalei*, *O.*
443 *centrocarpum*, *S. membranaceus*, *S. delicatus*) dinocyst taxa, already observed in the previous
444 zone (Fig. 7a). *L. machaerophorum*, often associated to fluvial and estuarine brackish
445 environments (Marret and Zonneveld, 2003), displays percentages that exceed the average of
446 its whole dataset (Fig. 7a) and are anti-correlated with *S. bentorii* and/or *O. centrocarpum* (i.e.
447 more related to the oceanic domain; Morzadec-Kerfourn, 1976, 1979) ones.

448 Palynological studies conducted on core VC-08 therefore confirm the environment sketches
449 proposed by *Le Roy and Jouet (2005)* on the basis of sedimentological and seismic data.
450 Furthermore, an obvious major limit can be defined within the tidal facies of U5 at 8,550
451 years BP, with a lower transitional part under increasing marine influences, and an upper part
452 characterized by well-established marine conditions in a perennial flooded bay.

453

454 **4.2. Western Bay of Brest (A and KS-24 cores Bay of Roscanvel)**

455 **Sedimentary facies**

456 Core “A” (Fig. 5a) is characterized by clayey sediments and numerous shell debris (Fig. 5b)
457 as well as bivalve and gastropod shells in life position often sampled for dating. The age
458 model (Fig. 6b) was established through a linear regression between 13 AMS ^{14}C dates (Table
459 1) allowing to consider an interval of about 1,400 years (6,700 to 5,300 BP or 4,750 to 3,350
460 BC) and mean sedimentation rates of about 0.26 cm/yr (Fig. 6b). Palynological analyses have
461 been conducted along the 318 cm of the core. It is worth noting that in one sample, the 190-
462 191 cm level (Table 1), continental organic matter (vegetal fibers) has been dated in parallel
463 with a bivalve for obtaining an estimation of the age reservoir within the Bay of Brest. When
464 calibrating the ^{14}C ages with IntCal13 without assuming any age reservoir effect on the
465 marine carbonate material, we obtain a difference of about 357 years between both dates, thus
466 indicating a ΔR of -43 years, very close to the one calculated by *Mangerud et al. (2006)* off
467 the Sein Island (i.e. ΔR of -40 ± 23 years) (<http://calib.org/marine/>).

468 Regarding core KS-24 (Fig. 5a), sampled in a deeper but closed site, the basal part is
469 characterized by clayey sediments sampled for palynological analyzes, and the upper part
470 consists in coarse sands and shell debris (Fig. 5c). These two parts of the core are separated by
471 a well-marked erosive surface at around 100 cm depth. Two AMS ^{14}C dates carried out at

472 173.5 and 102.5 cm allow dating the clayey section (i.e. interval sampled for palynological
473 analyzes; Fig. 5c) between 9,214 and 9,092 years BP (7,264 and 7,142 years BC).

474

475 *Dinocysts*

476 In both cores, 24 different taxa were identified, with an average of three different taxa per
477 slide for core KS-24 and 12 for core “A”. Similar to core VC-08, only main taxa were plotted
478 on Fig. 9a along with dinocyst concentrations and the specific richness. These two cores
479 constitute two temporal windows (i.e. 9,200-9,000 years BP for core KS-24, and 6,700-5,300
480 years BP for core “A”) on the same study site (i.e. Bay of Roscanvel, westernmost part of the
481 Bay of Brest; Fig. 1c). Both cores are described together, in their stratigraphical order, core
482 KS-24 representing a palynozone by itself.

483 Core KS-24 is characterized by the largely dominant *S.bentorii* species (69 %; Fig. 9a),
484 similar to the BD at the same period (Fig. 7a). Co-occurring with cysts of *P. dalei*, *O.*
485 *centrocarpum*, *S. membranaceus* and *S. quanta* at the base, *S. bentorii* becomes monospecific
486 at the top of core KS-24. From the base to the top of core “A”, dominant taxa are first
487 characterized by cysts of *P. dalei*, then replaced by *L. machaerophorum* that reach about 80
488 %. Decreasing percentages of cysts of *P. dalei* and *S. ramosus* are consequently observed at
489 the same time. Variations in percentages of major species and in total concentrations allow us
490 to delimit three palynozones (two of them being subdivided in two sub-palynozones). First
491 palynozone A-A (from 318 to 250 cm, Fig. 9a) displays a significant increasing trend of cysts
492 of *P. dalei*, a decreasing trend of *L. machaerophorum* and high percentages of *S. ramosus* and
493 *S. membranaceus*. At the limit between A-A and A-B, total cyst concentrations as well as *L.*
494 *machaerophorum* percentages show their lowest values, while cysts of *P. dalei* reach its
495 highest percentages (i.e. around 50 %). Second palynozone A-B (250-100 cm) is well marked
496 by noticeable opposite trends, and especially by increasing percentages of *L.*

497 *machaerophorum* and decreasing ones of cysts of *P. dalei*. Then *S. bentorii* displays higher
498 values than in the previous zone, while *S. ramosus* displays lower percentages. Moreover,
499 palynozone A-B corresponds to the strongest occurrences of *S. lazus*. This palynozone has
500 been divided into two sub-palynozones. Within zone A-B1, *L. machaerophorum* percentages
501 are the lowest while those of cysts of *P. dalei* are the highest of their respective whole
502 datasets. *L. machaerophorum* occurrences gradually rise in zone A-B2 and *S. bentorii*
503 percentages stabilize with a plateau of high percentages during this interval. The third
504 palynozone A-C (from 100 cm onwards; Fig. 9a) corresponds to a huge increase of *L.*
505 *machaerophorum* percentages while those of cysts of *P. dalei* sharply decrease, together with
506 a drop of *S. ramosus*. An additional subdivision is proposed to highlight the abrupt increase in
507 dinocyst concentrations at 50 cm (limit between A-C1 and AC-2). Furthermore, within zone
508 A-C1, weak but significant occurrences of *S. membranaceus* and *S. mirabilis* are observed,
509 while percentages of cysts of *P. dalei*, *S. mirabilis*, but also *S. ramosus*, and *S. lazus* drop to
510 extremely low values within zone A-C2. Conversely, *O. centrocarpum* and *S. belerius* slightly
511 increase in last zone A-C2.

512

513 **Pollen**

514 In both cores, 34 different taxa were recognized with an average of 11 different taxa per slide
515 for core KS-24 core and 12 for core “A” (Fig. 9b). While the two major taxa are the same for
516 both cores (i.e. *Corylus* and *Quercus*), pollen concentrations are much higher in core KS-24
517 (9,200-9,000 years BP) than in core “A” (6,700-5,300 years BP). Furthermore, trees are much
518 more present in core “A” than in the core “KS-24 (i.e. 93 % vs. 79 %). Among the herbaceous
519 in the core KS-24, Chenopodiaceae are particularly represented with an average of 12 %.

520 In core “A”, increasing percentages of *Alnus* (from 0 % to 15 %), and decreasing trends of
521 *Ulmus* and *Tilia*, allow to delimit the same palynozones than those described for dinocysts
522 (Fig. 9a). In general, herbaceous plants are very rare but a slow diversification is observed
523 towards the top, with the appearance of some minor taxa (e.g. Ranunculaceae, *Mercurialis*,
524 *Plantago* spp., Caryophyllaceae). *Corylus* and *Quercus* are anti-correlated throughout the
525 core. More specifically, the first palynozone A-A (318 to 250 cm; Fig. 9b) corresponds to low
526 percentages of *Corylus* that gradually increase at the end of the palynozone and, conversely,
527 to high percentages of *Quercus* that tend to gradually decrease. During this interval, *Tilia* and
528 Poaceae represent a meaningful part of the pollen content. Second palynozone A-B (250 to
529 100 cm) displays the occurrence and rise of *Alnus*, as well as the noticeable decrease of both
530 Poaceae and *Tilia*. Furthermore, at the start of palynozone A-B, *Ulmus* and *Pinus* percentages
531 exhibit stable values when compared with palynozone A-A, while they suddenly drop around
532 200cm, allowing to discriminate sub-palynozones A-B1 (from 250 to 200 cm) and A-
533 B2 (from 200 to 100 cm). Conversely, *Alnus* percentages strongly increase in sub-palynozone
534 A-B2. Palynozone A-C (from 100 cm onwards) highlights a new threshold with a marked
535 decrease of *Ulmus* together with a strong increase in *Alnus* percentages, the latter reaching its
536 highest values within zone A-C2 in parallel with relatively continuously high pollen
537 concentrations. Interestingly, most of the main peaks observed with pollen concentrations
538 (dotted lines in Fig. 9b) also correspond to increases of *Alnus* percentages along with high
539 values of *Corylus*, as already noticed for the VC-08 record.

540

541 **Micropalaeontological results**

542 Benthic foraminiferal assemblages are presented in depth for cores “A” and KS-24 (Fig. 10),
543 in parallel to the isotopic data measured on *Elphidium aculeatum*. Since *Elphidium* spp. are
544 the dominant taxa, all other taxa percentages have been calculated on a main foraminiferal

545 sum that excludes *E. crispum* and *E. aculeatum*. Similar to palynological data, only species
546 occurring at least once with percentages above 2% have been plotted in Fig. 10.

547 Along core KS-24, foraminiferal concentrations are low and assemblages are dominated by
548 *Haynesina germanica* (average of about 50 %), with few *Elphidium* spp. and important
549 variations of *Ammonia* spp., here strictly assigned to the species *A. tepida*.

550 Along core A, an increase of total benthic foraminiferal concentrations is observed throughout
551 the sequence and epifauna are the most represented in benthic foraminiferal assemblages, with
552 *Elphidium aculeatum* and *E. crispum* both accounting for 70%. Zones previously described
553 for dinocysts and pollen grains have been reported in the foraminiferal diagram, so as to
554 facilitate the cross-correlated approach between all fossil bio-indicators. Zone A-A is mainly
555 characterized by *Elphidium* spp. that displays an increasing trend, an important content of
556 Miliolidae, and an abrupt decrease of *Cibicides* spp. In addition, *Ammonia tepida* and
557 *Planorbulina mediterranensis* are observed with significant percentages. Palynozone A-B
558 corresponds to still high *Elphidium* spp. percentages and occurrences of *A. becarii*. Within
559 this interval, two subzones correspond first (i.e. A-B1, 250 to 200 cm) to decreasing values of
560 *Cibicides* and Miliolidae, together with the occurrence of *Lagena* spp., and then (i.e. A-B2,
561 200 to 100 cm) to increasing percentages of *Cibicides* (reaching 20%) and drastically low
562 *Lagena* percentages. Within palynozone A-C (from 100 cm onwards), *A. becarii* is replaced
563 by *A. tepida*, while *Cibicides* spp., *P. mediterranensis* and epiphytes (i.e. with a flat face that
564 allow them to be attached to sediments or plants) show approximately a same increasing
565 profile, opposite to the *Elphidium* spp. trend.

566 Regarding isotopic analyzes (Fig. 10), $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ show opposite trends with a slight
567 decrease towards lighter values for the $\delta^{18}\text{O}$ signal (amplitude of 1‰ between minimal and
568 maximal values in the dataset, ranging from 1.8‰ to around 0.8‰), and a slight increase

569 towards heavier values for the $\delta^{13}\text{C}$ signal (amplitude of 2‰ between minimal and maximal
570 values in the dataset, ranging from -1.5‰ to around 0.5‰).

571

572 **Environmental evolution in the Bay of Roscanvel (BR) under the rising sea level influence**

573 Environments of the BR were greatly impacted by the relative sea-level rise between 9,000
574 and 5,000 years BP. The general palaeoenvironmental evolution of the BR can be
575 reconstructed thanks to two Holocene windows obtained on core “A” (6,700-5,400 years BP)
576 and core KS-24 (9,200-9,090 years BP) and with averaged palynological (pollen and
577 dinocysts) and foraminiferal results compiled in pie charts (Fig. 11).

578 Throughout core KS-24, dinocyst assemblages show a low general diversity with major
579 occurrences of *S. bentorii* (Fig. 9a, 11), arguing for a strictly coastal environment. Among
580 pollen assemblages (Fig. 9b, 11), core KS-24 displays an important content of *Corylus*, a
581 species considered as pioneer during the post-glacial temperate forest (mixed oak) reconquest
582 that mainly marked north-western European landscapes around 9,000 years BP (Huntley,
583 1993; Tinner and Lotter, 2001). Among herbaceous plants, Chenopodiaceae are significantly
584 present, certainly coming from nearby salt-marshes. In addition, foraminiferal assemblages
585 are dominated by *Haynesina germanica*, *Ammonia tepida*, and *Elphidium williamsoni*. *A.*
586 *tepida* and *E. williamsoni* are closely related to intertidal sheltered mudflat environments,
587 while *H. germanica* is related to continental organic matter inputs (Debenay et al., 2006;
588 Rossi et al., 2011). These three species indicate a highly tidal-influenced environment and
589 local small runoffs rich in continental organic matter (Redois, 1996; Debenay et al., 2006;
590 Perez-Belmonte, 2008; Rossi et al., 2011). However, the noticeable absence of *L.*
591 *machaerophorum* and *Alnus* suggests low major freshwater supplies. The surrounding
592 continental context may then correspond to small watersheds, while the Aulne River was still

593 flowing to the north of core KS-24 location taking into account the low sea level position at
594 that time (i.e. -26 m; *Gregoire et al., 2017*; Fig. 11). In this context, the marine influence
595 remains confined to the axis of the main channels of the Aulne and Elorn Rivers (*Gregoire et*
596 *al., 2017*). The higher river terraces (i.e. where core “A” was taken; Fig. 1c, 5c) are entirely
597 emerged at that time, while the lower site where core KS-24 has been retrieved (Fig. 1c, 5c) is
598 characterized by the development of abundant maritime marshes (schorre and slikke) with
599 halophilous plants (i.e. *Chenopodiaceae*) and a rather weak fluvial dynamic. Considering the
600 current 26 m depth location of core KS-24 and its 1.81 m length, it allows deducing a relative
601 sea-level between 27 and 28 meters under the current sea-level which is very similar to the 26
602 meters proposed at 9,000 years BP by *Gregoire et al. (2017)*, taking into account the
603 macrotidal context of north-western Brittany.

604 Throughout core “A”, dinocyst assemblages display a general higher diversity than in core
605 KS-24 with 24 taxa, mostly oceanic to coastal, along with occurrences of about 35% of *L.*
606 *machaerophorum* (estuarine taxon). Predominant marine influences, favorable for diversified
607 dinoflagellate blooms, are associated to the BR marine flooding in a context of superimposed
608 fluvial influences, as evidenced by *L. machaerophorum*, mainly associated to Aulne River
609 inputs, here perceptible compared to the previously marsh environment described with core
610 KS-24. Concerning pollen assemblages, the mixed temperate forest is settled (90 % of tree
611 pollen grains, Fig. 9b and 11) with a higher representation of *Quercus* when compared to core
612 KS-24. This indicates a connection of the BR to the main stream of the Aulne River due to the
613 flooding of the BB, thus highlighting a more regional signature (i.e. BB watersheds) in core
614 “A”, also corresponding to the colonization of temperate tree taxa in western Europe during
615 the mid-Holocene (*Ruddiman and McIntyre, 1981; Brewer et al., 2002*). The *Chenopodiaceae*
616 signal is less pronounced in core “A” due to the disconnection of the study site from marsh
617 environment, as also confirmed by the drastic reduction in marsh-related foraminiferal species

618 (i.e. *Haynesina germanica*, *Ammonia tepida*, and *Elphidium williamsoni*). Also, benthic
619 foraminiferal assemblages display a stronger diversity in core “A” suggesting a greater marine
620 influence in agreement with dinocyst results. Moreover, the presence of some epiphytic and
621 fixed species (i.e. *Cibicides* spp. and *Planorbulina mediterraneensis*) suggests a higher
622 hydrodynamism. In addition, the absence of benthic infauna could indicate a lack of very
623 important organic matter inputs. The environment would therefore be characterized by a
624 sufficient organic matter supply to allow the development of algal coverings (allowing the
625 presence of epiphytes), and hydrodynamism would disperse this organic material, thus
626 oxygenating the bottom. In summary, around 6,000 years BP the relative sea level was
627 relatively close to the present-day one (*Goslin et al., 2015*) and the BB as a whole (and thus
628 the BR) is flooded (Fig. 11). The environment is then marked by a clearer marine influence
629 and the remoteness of salt-marshes areas. Tidal currents erode sediments on terrace slopes
630 from the main channels, leading to erosive surfaces separating muddy and sandy facies (i.e.
631 upper part of KS-24 core from 100 cm onwards not sampled for palynological analysis, Fig.
632 5c), as also identified by *Gregoire et al. (2017)*.

633

5. Climate impacts and land-use changes

5.1. Local imprints of the 9,000-8,000 and 6,000-5,000 years BP RCC events

Superimposed on the palaeoenvironmental reconstitutions discussed in the context of the rising sea-level context, climate trends are well expressed in our palynological records due to our high resolution study carried out on cores characterized by high sedimentation rates.

In the Bay of Douarnenez, from 8,550 to 8,400 years BP (zone VC-C; Fig. 7b and 12), *Corylus* becomes clearly dominant. Moreover, this interval displays a slight decrease of *Ulmus*, a disappearance of the thermophilous taxon *S. mirabilis*, and significant occurrences of *S. lazus*, a dinocyst associated with cold SST (Zonneveld *et al.*, 2013). These palynological evidences may support a cooling event. Indeed, this cooling associated with the appearance of a strong seasonal contrast between 8,855 and 8,000 years BP observed by Naughton *et al.* (2007) favors the development of *Corylus* woodlands and contributes to the decrease of deciduous *Quercus* forest expanse. In the same time, the obvious increase of pollen fluxes, coeval with a marked increase of *L. machaerophorum* percentages, and *Alnus* ones to a lesser extent, may argue for increasing fluvial discharges (as previously discussed in Lambert *et al.* 2017) and thus of humidity. It is worth noting that low percentages of *Alnus* may here be ascribed to its later settlement in the vegetal reconquest succession, as observed in the Armorican Massif by David (2014). Each *L. machaerophorum* increase (Fig. 7a and 12), also concomitant to the withdrawal of coastal (*S. bentorii*) and marine (*O.centrocarpum*) species, coincides with increasing percentages and concentrations of *Corylus* (Fig. 7b and 12).

In the Bay of Roscanvel, dinocyst assemblages were dominated by *S. ramosus* (oceanic taxa; Morzadec-Kerfourn, 1977) and cysts of *P. dalei* before 6,000 years BP, and are characterised after this limit (between zones A-B and A-C) by an obvious and sharp increase of *L. machaerophorum* percentages from 30 to 70 % (Fig. 9a and 12), a species usually

658 encountered in estuarine and fluvial environments of Brittany's coasts (*Morzadec-Kerfourn,*
659 *1977*). This could emphasize the establishment of stratified waters related to increasing fluvial
660 inputs. Furthermore, *L. machaerophorum* is accompanied by increasing percentages of the
661 heterotrophic taxa *S. quanta* (Fig. 9a) that could support increasing fluvial nutrient inputs to
662 the BB. Moreover, since *Alnus* began to appear on the territory, we now observe an increase
663 of this tree in parallel with *L. machaerophorum*. As previously observed for core VC-08,
664 *Corylus* increases are concomitant with peaks of pollen concentrations and of *L.*
665 *machaerophorum* percentages. Finally, higher percentages of *H. germanica* and *A. tepida*
666 (Fig. 10), benthic foraminiferal species subordinate to environments under fluvial influences
667 (*Debenay et al. 2006; Perez-Belmonte, 2008; Estournès et al. 2012*), confirm previous
668 observations of increasing fluvial discharges probably related to increasing moisture.

669 For both cores VC-08 and "A", increasing fluvial inputs would correspond to a local detection
670 of the large-scale Holocene millennial-scale climatic variability (i.e. "Bond events", *Bond et*
671 *al. 2001*; or RCC for "Rapid Climate Change", *Mayewski et al., 2004*). Most of these cold
672 Holocene events, initially identified in North Atlantic sedimentary cores by detrital grains
673 drained by glaciers (Fig. 12a), correspond to Scandinavian glacier advances (*Nesje et al.,*
674 *2001*), colder northern hemisphere temperatures and drier conditions in the tropics (*Mayewski*
675 *et al., 1997, 2004; Meeker and Mayewski, 2002*). Also, the strengthening of the westerlies
676 (*Bradbury et al., 1993; Mayewski et al., 2004*) may have resulted in recurrent positive modes
677 of the NAO, leading to increased precipitations in north-western Europe. Interestingly,
678 *Mojtahid et al. (2013)* associated the 6,000-5,000 interval to persistent positive NAO
679 conditions in the Bay of Biscay. A solar irradiance decline may be responsible for this 6,000-
680 5,000 years BP event (*Steinhilber et al., 2009; Fig. 12a*), also leading to a North-Atlantic SST
681 decrease (*Jiang et al., 2015; Fig. 12a*). Thus, cores VC-08 around 8,550 years BP, and core
682 "A" around 6,000 years BP (Fig. 12c), may then evidence for the first time the regional

683 imprint in western Brittany's coasts of two RCC events through significant increases in
684 moisture and fluvial discharges.

685

686 **5.2. Local signal of *Cerealia*-type pollen grains during the Mesolithic: coastal Poaceae or** 687 **early traces of a proto-agriculture?**

688 Between 8,600 and 8,300 years BP, VC-08 core shows a very homogeneous laminated
689 facies which attests to a stable environment aggradation related to a deltaic alluvial plain
690 characterized by intertidal regimes (*Le Roy and Jouet, 2005; Fig. 4*). Constant percentages of
691 Chenopodiaceae all along the record, associated with few and constant percentages of
692 Poaceae (*Fig. 7b*), suggest a coastal marsh (slikke and schorre-type) environment, confirming
693 the aggradation of intertidal deposits. In this sedimentary context, the palynological analysis
694 interestingly records iterative occurrences of large pollen grains of Poaceae (*Fig. 13*).

695 Cereal pollen identifications are usually based on the diameter criteria (grain and *annulus*)
696 following *Beug (1965), Leroyer et al. (2004)* and *Joly et al. (2007)*, because of the
697 impossibility to separate wild grasses and *Cerealia*-type pollen grains using exine sculpture
698 types (*Beug, 2004*). In western Europe, the commonly accepted thresholds of 45 μm for the
699 grain diameter and 8 μm for its *annulus* (that we will refer as the 45-8 μm criteria)
700 theoretically enable the distinction between cereal pollen grains and those of indigenous wild
701 grasses from the French western Atlantic coast (*Leroyer et al., 2004*). Considering these
702 criteria along the whole sequence, 7 levels delivered cereal-type grains with one to 3
703 occurrences in palynological slides. These 12 large pollen grains recognized in total, and
704 distributed all along the core (325, 295, 220, 170, 140, 120 and 65 cm, *Fig. 13*), have been
705 carefully measured, providing average values of 46 μm for the grain diameter and 9.9 μm for
706 the *annulus*. Nevertheless, the "45-8 μm " criteria have been criticized (*Joly et al., 2007*). The

707 statistical study conducted by *Joly et al. (2007)* on modern plants from the Atlantic coast
708 reveals that a “45-10 μm ” criteria makes it possible to identify 91 % of cereals (but 9 % of
709 pollen grains are from wild grasses). In their study, *Joly et al. (2007)* then proposed to
710 increase the thresholds to a most discriminant level of “47-11 μm ”, thus avoiding to consider,
711 as much as possible, larger pollen grains of coastal grasses from the Atlantic coast. The “47-
712 11 μm ” criteria then make it possible to discriminate 96% of cereal pollen grains (i.e. 4% of
713 wild poaceae fall within this criterion) (*Joly et al., 2007*). However, with these new criteria,
714 the sensibility to detect *Cerealia* pollen grains significantly declines. More precisely, among
715 100 modern *Cerealia* pollen grains, only 59 would be detected as such with the “47-11 μm ”
716 criteria, the other grains being then supposed as wild Poaceae. Despite this low sensibility and
717 the fact that some cereals are probably not taken into account with these thresholds, the
718 applied “45-11 μm ” criteria to our data confirm discreet peaks of *Cerealia*-type pollen grains,
719 from around 8,600 to 8,300 years BP (at 65, 120 and 170 cm; Fig. 13).

720 Considering that indigenous coastal grasses may have larger pollen grains than continental
721 species, it is also worth noting that these weak occurrences observed in the BD record do not
722 co-occur in parallel with variations of other Poaceae or Chenopodiaceae, suggesting no
723 significant changes in landscape and/or pollen transport at that time. Moreover, to be more
724 confident about these atypical and early observations, many other clues must be considered.
725 Some plants often associated with crops (e.g. adventitious taxa) are necessary to discuss
726 anthropogenic signatures in the vegetation cover (*Behre, 1981; Willcox, 2005*). It is precisely
727 interesting to note that slight increases in *Plantago lanceolata*, *Rumex* sp. and Brassicaceae
728 percentages are observed in VC-08 core during main occurrences of *Cerealia*-type (Fig. 8 and
729 14). A slight long-term decrease can also be highlighted in tree percentages between 8,600
730 and 8,300 years BP, perhaps arguing for an early start of a small human-made landscape
731 opening (Fig. 8 and 14).

732 Our findings are however in marked opposition with the regional neolithisation model
733 admitted by the archaeological community and based on clues related to changes in cultural
734 practices identified by lithic (and ceramic) industries (*Cassen et al., 1998; Marchand, 2005,*
735 *2007; Marchand et al., 2006; Blanchet et al., 2006; Paillet et al., 2007, 2014; Hamon, 2008*).
736 In addition, studies on Mesolithic archeological sites in Brittany have highlighted traces of
737 plant cuttings, but tools did not show the classical polishes associated with cereal cuttings
738 (*Guéret, 2013; Guéret et al., 2014*). Even if our results raise questions and encourage caution,
739 they are consistent with other palynological studies carried out in western France that also
740 evidenced early discreet appearances of *Cerealia*-type pollen grains (Fig. 15). As early as
741 1996, analyses conducted by *Visset et al.* opened the debate about the existence of a very
742 ancient cereal farms as highlighted by low occurrences of *Cerealia*-type pollen between 7,200
743 and 5,800 years BC (9,200-7,800 years BP), concomitantly with increases of *Corylus* and
744 ruderal plant percentages and decreasing *Quercus* percentages in the coastal Morbihan region
745 (SE Brittany; Fig. 15). Other studies along the Loire River revealed the presence of cereal
746 pollen taxa as well as *Juglans* between 6,400 and 5,900 years BC (8,400-7,900 years BP;
747 *Ouguerram and Visset, 2001*) and between 6,600 and 5,800 years BC (8,500-7,800 years BP;
748 *Carcaud et al., 2000*). Moreover, *Joly and Visset (2005, 2009)* reported cereal and ruderal
749 pollen grains on the southern Atlantic coast (Vendée) between 7,500 and 6,300 years BC
750 (9,500-8,200 years BP) and between 7,500 and 6,200 years BC (9,400-8,100 years BP).
751 Beyond Brittany, some early *Cerealia*-type occurrences have already been observed as co-
752 occurring with ruderal plants between 6,400 and 5,800 years BC (8,400-7,800 years BP) in
753 the northern Pyrenean region by *Galop and Vaquer (2004)* or between 6,600 and 5,800 years
754 BC (8,600-7,800 years BP) in the Parisian Basin (*Leroyer and Allenet, 2006*). In addition,
755 occurrences of *Cerealia*-type pollen grains have been reported from many Swiss sites
756 between 7,750 and 5,800 Cal. BC (*Erny-Rodmann et al., 1997; Lotter, 1999; Tinner et al.,*

757 1999, 2007; Beckmann, 2004; Fig. 15). Taking into account our data and the numerous
758 references cited above, we underline the great timing coherence of large Poaceae pollen grain
759 appearances, with sizes usually interpreted as resulting from a cultivated origin. Among the
760 reservations that are often opposed to these detections (e.g. reliability of dating, long-distance
761 transport or laboratory contaminations; Behre, 2007), we can consider minimal errors in
762 carbon datings (see the core stratigraphy) or in laboratory treatments that are identical for all
763 cores (i.e. Roscanvel “A” core, without any large Poaceae pollen grain). Regarding the long-
764 distance transport of pollen grains from the eastern areas where cereals were already
765 cultivated, as already mentioned by Tinner *et al.* (2007), it is surprising that we do not detect
766 pollen grains from other exotic species with a better pollen dispersion than cereals (e.g.
767 *Quercus ilex*, *Pistacia*). Moreover, the BB and BD watersheds are too small to receive long-
768 transported pollen grains from eastern or southern sites. We can thus suggest two hypotheses:

769 1. These atypical large pollen grains are not inherited from cultures and represent a signature
770 of littoral grasses. Indeed, some coastal Poaceae on the Atlantic coast may have pollen grains
771 that can reach large sizes (Joly *et al.*, 2007). The size threshold which is considered by
772 palynologists as a robust identification criteria for the determination of cereal pollen grains,
773 may be revised again (cf. Joly *et al.*, 2007) in light of the particular pollen sizes of some
774 regional littoral grasses. Regarding this first hypothesis, it still appears curious not to detect
775 any significant increase in total Poaceae or Chenopodiaceae when the most relevant peaks of
776 *Cerealia*-type pollen grains are recorded (Fig. 7b). Furthermore, in the Bay of Roscanvel
777 records (A and KS-24 cores, Fig. 9b), studied from 9,200 to 5,300 years BP in same
778 environmental sedimentary context, the total absence of these large pollen grains of Poaceae
779 before 8,600 years BP implies no peculiarity of size among pollen grains from littoral grasses.

780 2. A discreet proto-agriculture signal would be recorded for the first time during the
781 Mesolithic in western Brittany. Discussed by the Mesolithic archaeological community, this

782 type of “agriculture” would be characterized by its discretion in the landscape and the absence
783 of specific hard rock made agricultural tools (*Marchand, 2005*). The work of plants seems
784 indeed to grown toward the end of the Mesolithic, but regional use-wear analysis on
785 archeological material suggested that wear traces result from scrapping of wood and soft
786 plants, with however no traces of cereal plant work (e.g. *Gassin et al., 2013*; Beg an Dorchen,
787 SW Brittany). In the previous palynological studies mentioned above (Fig. 15), *Cerealia*-type
788 pollen grain occurrences are always scarce and isolated in space and time (*Visset et al., 2002*).
789 Our study, conducted on a core taken in the marine coastal domain, enables to record a
790 temporal BD watershed signal with iterative occurrences of these large pollen grains (8,482;
791 8,440; 8,395 years BP or 6,532; 6,490; 6,445 years BC) associated with some other cultural
792 indicators as *Plantago lanceolata* and *Rumex* spp. (Fig. 14). The scarcity of Mesolithic data
793 on the territory, and the difficulties to identify proto-agriculture evidence, both with pollen or
794 lithic data, does not allow to discuss the local introduction of this proto-agriculture imprint as
795 a result from a meridional migration movement (Retzian culture *via* the Loire estuary; *Visset*
796 *et al., 2002*) or from the main Neolithic cultural in Brittany from a Danubian origin
797 (*Dubouloz, 2003*; *Gomart et al., 2015*; Fig. 15).

798 In order to progress, future studies must be conducted i) to newly establish the size
799 variation of the current littoral wild grass pollen grains, ii) to confirm the timing of the first
800 large pollen grain detection at a broader regional-scale.

801

802 **6. Conclusion**

803 Our results allowed characterizing Holocene coastal palaeoenvironments of NW France (Bay
804 of Brest and Bay of Douarnenez, western Brittany) over two given periods (9,200-8,400 and
805 6,600-5,300 years BP). Various factors forced coastal environmental changes during the early
806 to mid-Holocene especially including the relative sea-level rise that modified sedimentation
807 processes and the post-glacial recolonization of temperate trees. Sedimentological and
808 palynological analyses carried out in this study allowed us to characterize and specify the
809 environmental variations that impacted coastal environments of western Brittany. The relative
810 sea-level rise influenced the sedimentary infilling history of shallow marine environments that
811 transited from river valleys to oceanic bays. In addition, high resolution studies enabled us to
812 detect the regional response to more global events arising from the millennial-scale climate
813 variability in the North Atlantic. Thus, around 8,600 and 6,000 years BP, onsets of two major
814 abrupt climate events are locally detected by a moisture increase marked by strong
815 occurrences of *Corylus*, *Alnus*, and *L. machaerophorum* dinocyst taxon. In addition, the
816 gradual decrease of continental summer temperatures is marked by the withdrawal of
817 thermophilous species *Ulmus* and *Tilia*. Moreover, during this time period also marked by the
818 appearance of the neolithisation, *Cerealia*-type pollen grains have been detected in the Bay of
819 Douarnenez, about 1,500 years before the advent of agriculture commonly accepted. These
820 local results, while being consistent with other palynological studies conducted on the French
821 Atlantic coast, must be taken with caution and open the way to further studies so as to confirm
822 these "pre-domestic" agriculture indices.

7. Acknowledgements

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824
825 This study was supported by the French CNRS and is a contribution to the 2015-2016 INSU
826 project EC2CO-LEFE: « *CAMOMI : Convergences / Approches croisées des signaux*
827 *MOléculaires et Mlcropaléontologiques pour décrypter les forçages anthropiques et*
828 *climatiques en milieu côtier (Rade de Brest)* » and the UBO-BQR project : « *PARADE :*
829 *Signature PALéoenvironnementale des séquences holocènes en RADE de Brest* ». This work
830 was supported by the «Laboratoire d'Excellence» LabexMER (ANR-10-LABX-19) and co-
831 funded by a grant from the French government under the program « *Investissements*
832 *d'Avenir* ». We thank the UMR CNRS 5805 EPOC (Talence) for palynological laboratory
833 procedures (Muriel Georget), the UMS 2572 LMC14 (Saclay) for carbon dating via Artemis
834 project fundings, the sedimentology laboratory of the Shom (« *Service Hydrographique et*
835 *Océanographique de la Marine*») for the VC-08 core taken from the Bay of Douarnenez
836 (PROTEUS-DUNES cruises; 2012), the laboratory IFREMER-Marine Geosciences
837 (Plouzané) for the KS-24 core collected in the Bay of Brest (SERABEQ cruises; G. Gregoire
838 PhD thesis, 2016) and the LEMAR laboratory (« *Laboratoire des Sciences de l'Environnement*
839 *Marin*»; IUEM, Plouzané) for the A core taken from the Bay of Brest (Défis Golfe de
840 Gascogne cruise; 2003). Main issues of this project are integrated within the theme
841 "Dynamics of Human Settlement and Paleoenvironments » of the Zone Atelier Brest Iroise
842 (ZABrI, INEE-CNRS).

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1302

1303 **Figure caption**

1304

1305 Figure 1: a) Location of study sites in North Western France; location of sediment cores on
1306 the bathymetric maps of the (b) Bay of Douarnenez (black star) and of the (c) Bay of Brest
1307 (white stars).

1308 Figure 2: Schematic scenario of the Bay of Douarnenez sediment infilling (after Le Roy and
1309 Jouet, 2005). Four sketches related to the sedimentary units defined from seismic and
1310 sedimentological analyses (cf details *in* Le Roy and Jouet, 2005).

1311 Figure 3: a) Pictures and b) XR radiography of the VC-08 core. The red strips represent the
1312 sampled core portion to perform palynological analyzes.

1313 Figure 4: Sedimentological log of the VC-08 core (b), grain size evolution and magnetic
1314 susceptibility along the core (a). For the different sedimentary units (identified by color
1315 strips), a graph representing the grain percentages distribution by size was added. ¹⁴C dates
1316 are represented by red stars on the log.

1317 Figure 5: Cores of the Bay of Brest, “A” and KS-24, and their location on a MNT of the Bay
1318 of Roscanvel with their bathymetric position (c). Sedimentological logs, photographs and X-
1319 ray radiography for the “A” (a) and KS-24 cores (b). The ¹⁴C dates are represented on the logs
1320 by red stars. The red strips represent the sampled core portion to perform palynological
1321 analyzes.

1322 Figure 6: a) VC-08 (Bay of Douarnenez) and b) “A” core (Bay of Brest) age models with their
1323 sedimentological logs., linear regression lines (in black) and sedimentation rates in cm/yr.

1324 Figure 7: Graphs against depths with major taxa (greater than 2%) of dinocysts (a) and pollen
1325 (b) along the VC-08 core, Bay of Douarnenez. The ¹⁴C dates are represented by red stars. The
1326 dotted lines represent the pollen concentration peaks.

1327 Figure 8: Diagram depicted in ages, from the VC-08 core, grouping together data from
1328 palynological counting (Pollen, dinocyst and spore concentrations, percentages of trees,
1329 *Corylus*, *Quercus*, Poaceae, Brassicaceae, *Rumex*, *Cerealia*, the dinocyst *L.machaerophorum*
1330 and the total Pollen / dinocysts ratio), and the magnetic susceptibility, with respect to
1331 sedimentary units defined in Figure 4.

1332 Figure 9: Diagram in depth grouping the major taxa (greater than 2%) of dinocysts (a) and
1333 pollen (b) for the “A” and KS-24 cores, both taken from the Bay of Roscanvel and plotted one
1334 above the other according to the time periods covered by the two sequences. The ¹⁴C dates are
1335 represented by red stars. For the “A” core, the most significant minor pollen taxa (greater than
1336 1%) were also reported. The dotted lines represent the pollen concentration peaks.

1337 Figure 10: Major benthic foraminiferal taxa data plotted against depths (greater than 2%) in
1338 the “A” and KS-24 cores. ¹⁴C dates are represented by red stars. *Ammonia spp*, *Cibicides spp*,
1339 *P. mediterraneensis*, Epiphytic species, *Lagena spp* and Miliolideae are represented in
1340 percentages according to a main sum excluding the *Elphidium* species, major taxa that greatly

1341 tainted the individual signal of each one. The isotopic data measured on shells of *E.*
1342 *aculeatum* species are added to the graph.

1343 Figure 11: Mean palynological (pollen and dinocysts) and micropalaeontological (benthic
1344 foraminifera) data for the “A” and KS-24 cores represented in pie charts (a). Diagrams
1345 representing two models of the landscape evolution in the Bay of Roscanvel following the sea
1346 level rise (b). On each model, the typical foraminifera encountered in the different
1347 environments according to their ecology were added.

1348 Figure 12: a) Published reconstituted palaeoclimatic data across the Holocene in the North
1349 Atlantic region (solar irradiance (dTSi) by *Steinhilber et al. 2009*, Sea surface temperature
1350 (SST) by *Jiang et al., 2015* and Northern Hemisphere summer insolation by *Berger and*
1351 *Loutre, 1991*) and the detrital Hematite Stained Grains by *Bond et al., 2008*. b) Diagram
1352 showing the palynological data of the “A” core between 6,600 and 5,400 years Cal. BP. (from
1353 left to right: pollen and dinocyst concentrations, tree percentages, *Corylus*, *Alnus*, percentages
1354 and the *L. machaerophorum* / *S. bentorii* ratio). c) Diagram showing the palynological data of
1355 the VC-08 core between 8,700 and 8,400 years Cal. BP with the same succession of
1356 palynological proxies. The temporal extent of each core is represented by a black rectangle in
1357 the a) part and the temporal extent of the graphs b) and c) is represented by a blue rectangle.
1358 The limits of palynozones A-B / A-C and VC-B / VC-C have also been reported in the a) part
1359 of the figure.

1360 Figure 13: Pollen grains photographs of wild Poaceae (a) and cereals (b), with grain and
1361 *annulus* diameter measurements for each one, on 4 levels of the VC-08 core.

1362 Figure 14: Diagram in depth from the VC-08 core, representing variations of some
1363 herbaceous taxa percentages (i.e. *Cerealina*-type, other Poaceae, *Brassicaceae*, *Rumex* sp.) and
1364 the occurrences of *Plantago lanceolata*.

1365 Figure 15: Map showing the first occurrences of *Cerealia* in published pollen records, with
1366 related dates. The two circled areas (Teviecién and Retzien) delimit the cultural specificities
1367 of the end of the Mesolithic after *Marchand (2005)*. The gray arrows correspond to the
1368 different neolithisation paths reaching Western Europe.

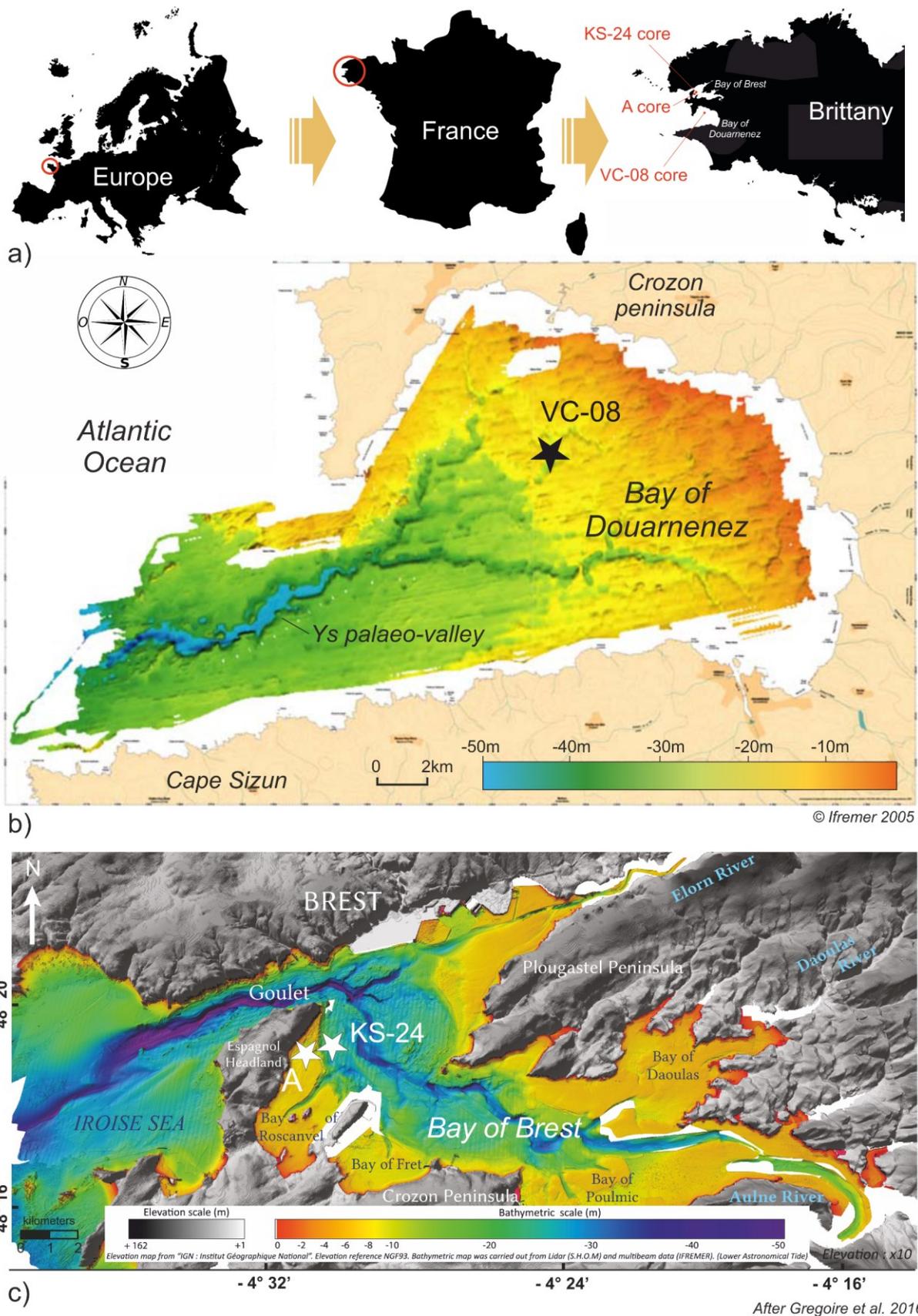


Figure 1

Schematic scenario of Bay of Douarnenez sedimentary infilling
(after Le Roy and Jouet, 2005)

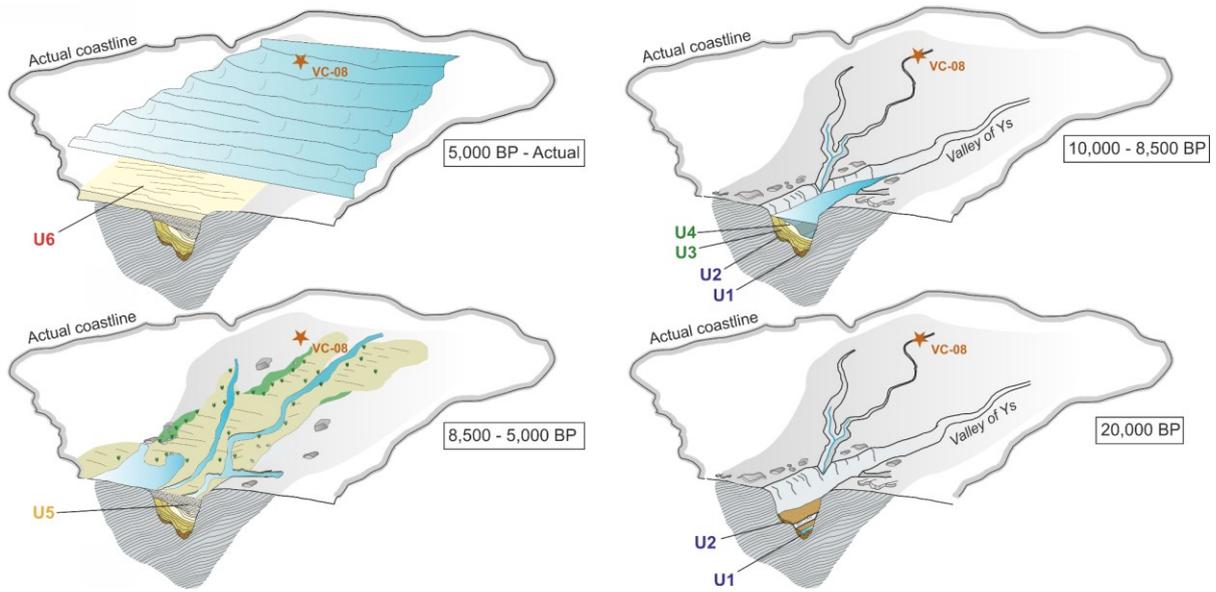


Figure 2

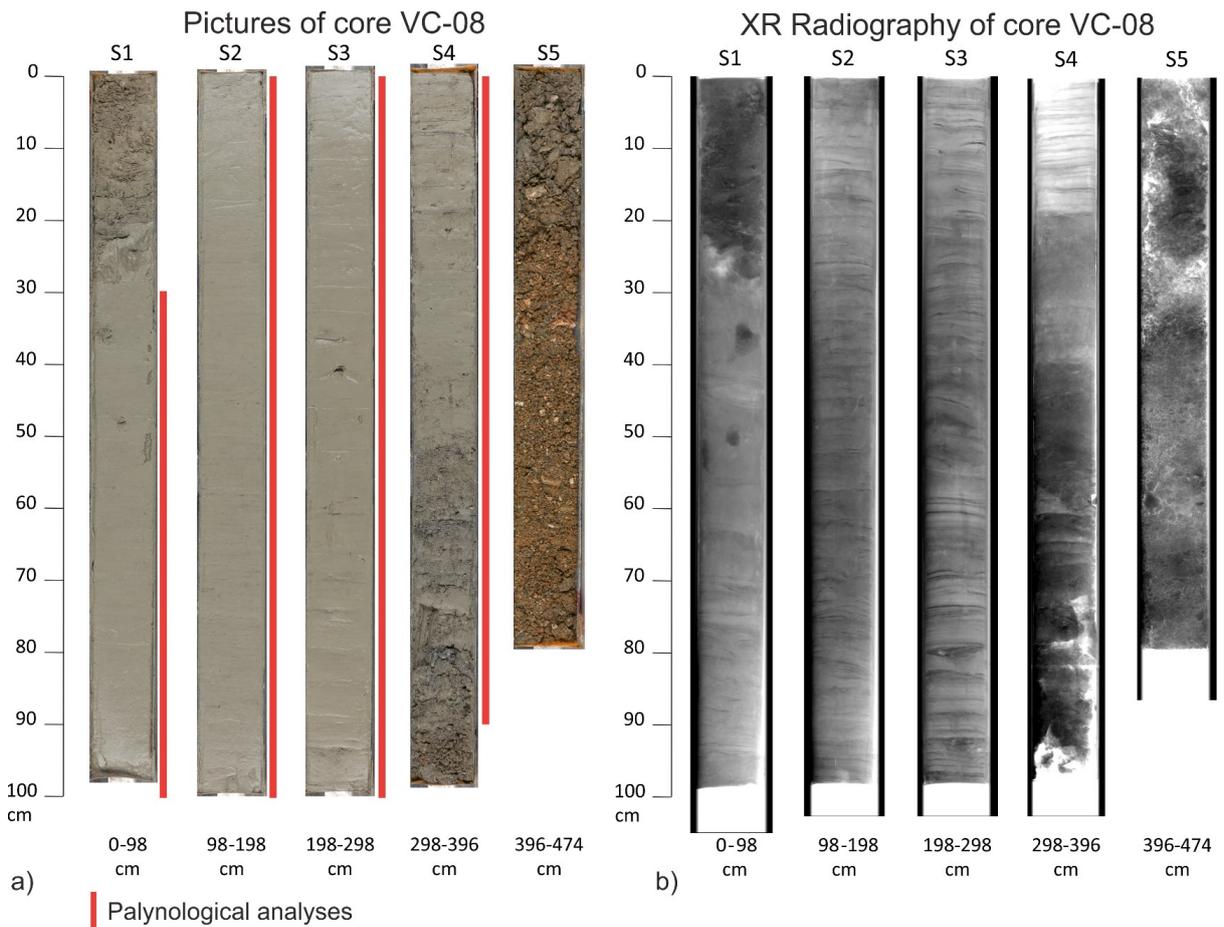


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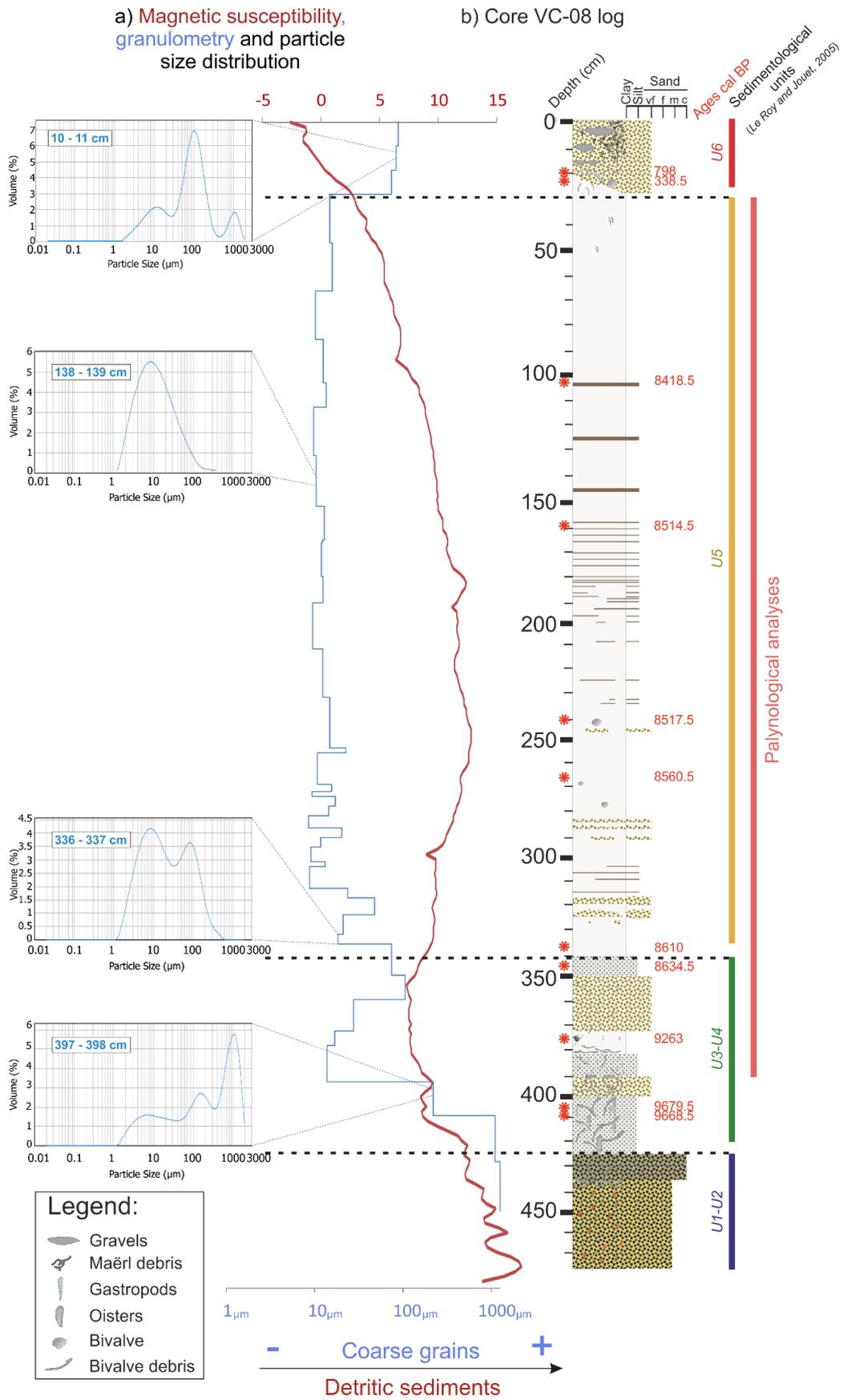


Figure 4

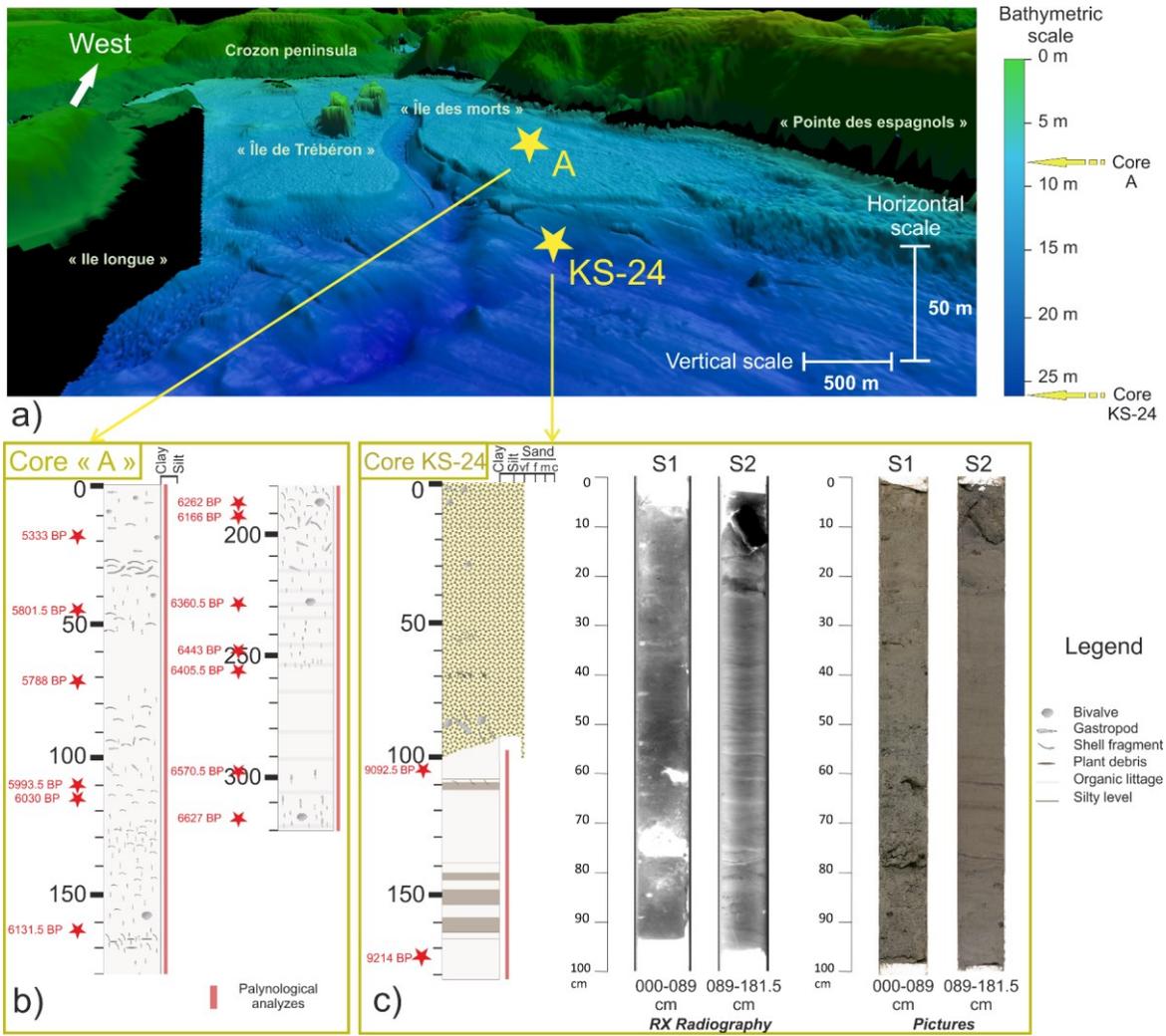


Figure 5

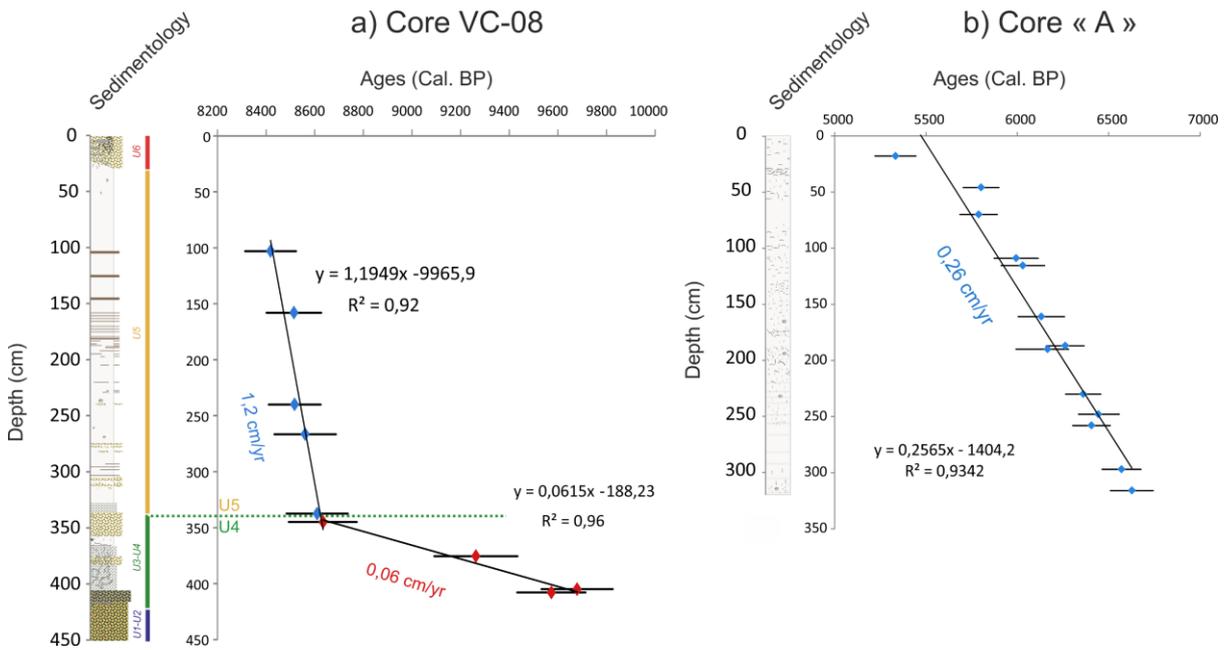


Figure 6

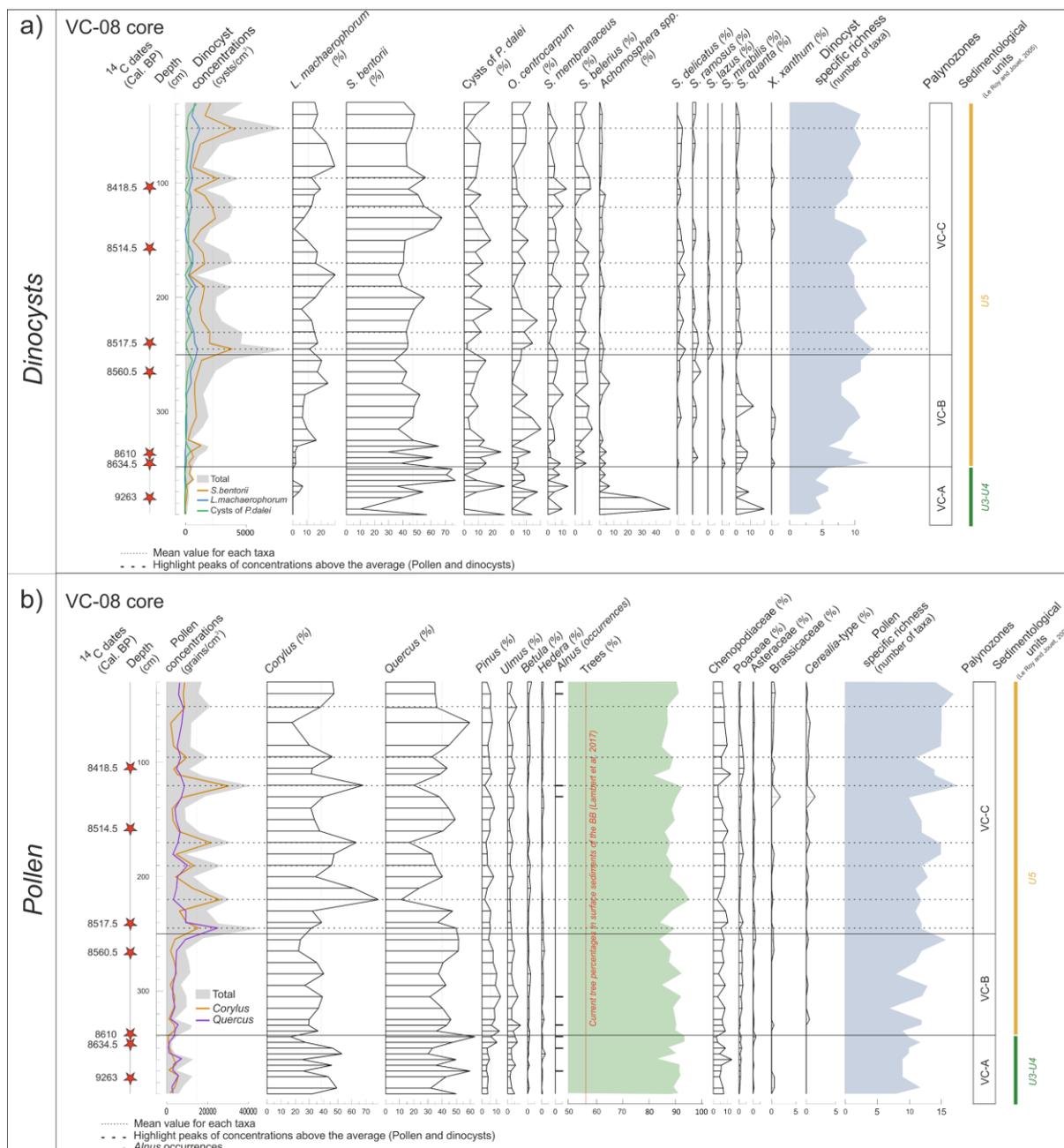


Figure 7

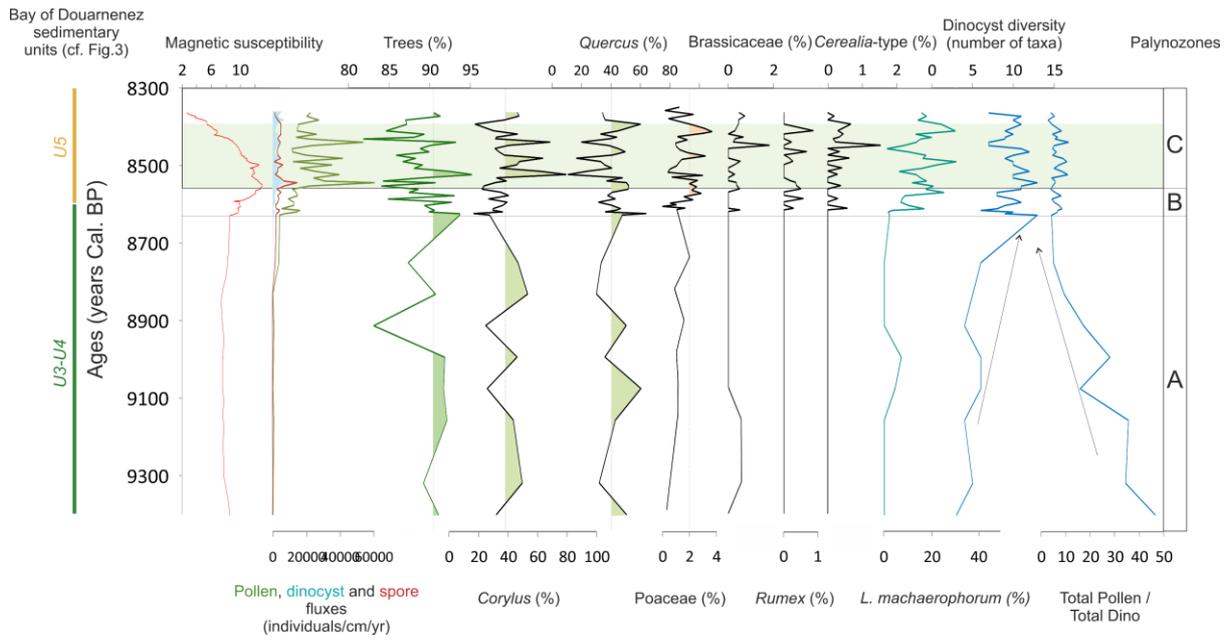


Figure 8

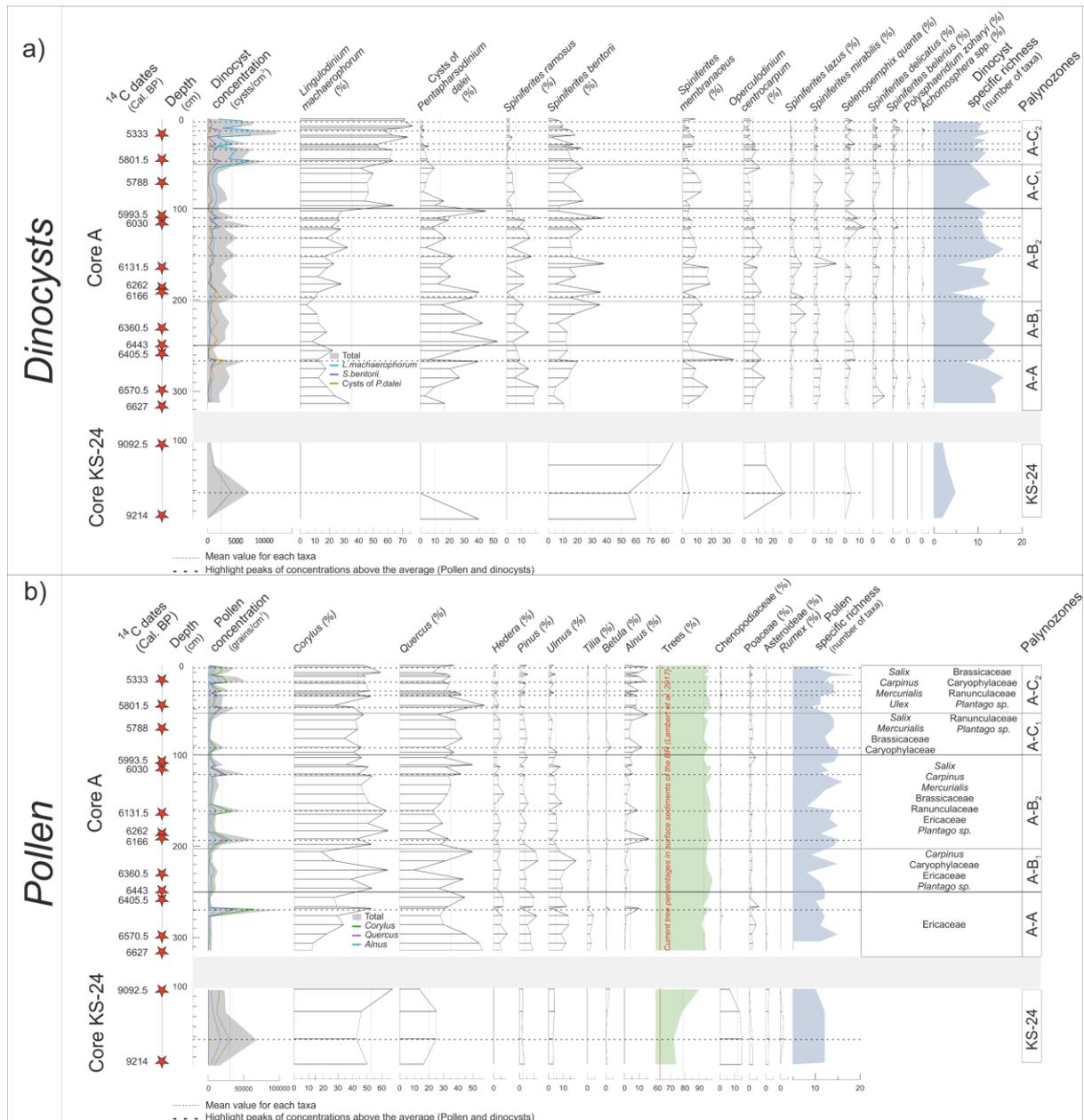


Figure 9

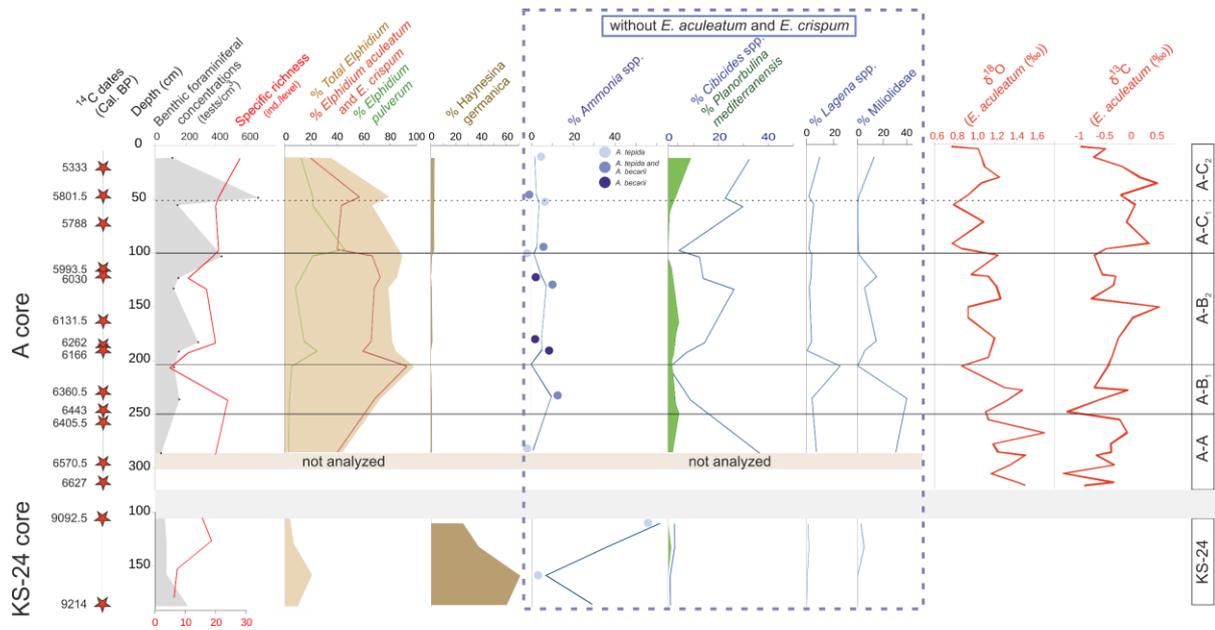


Figure 10

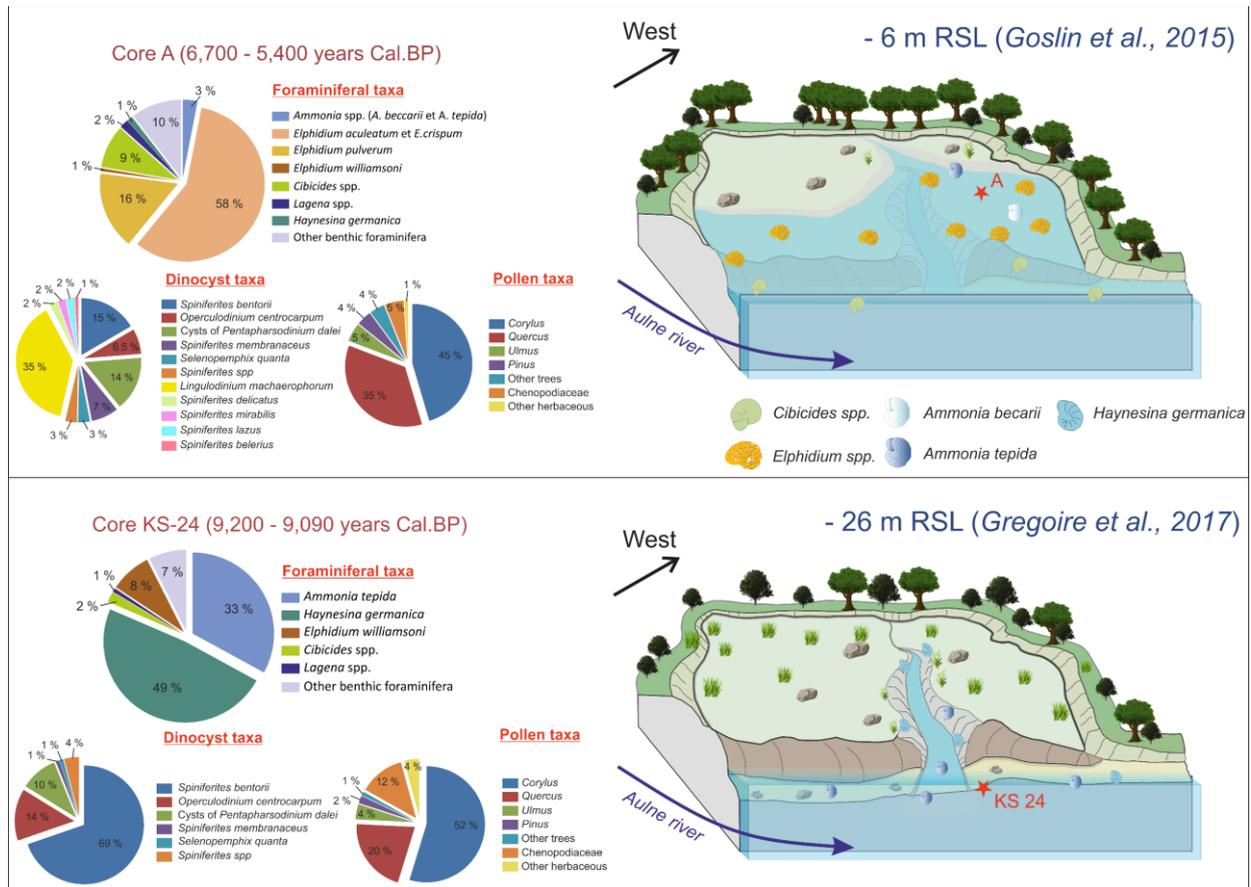


Figure 11

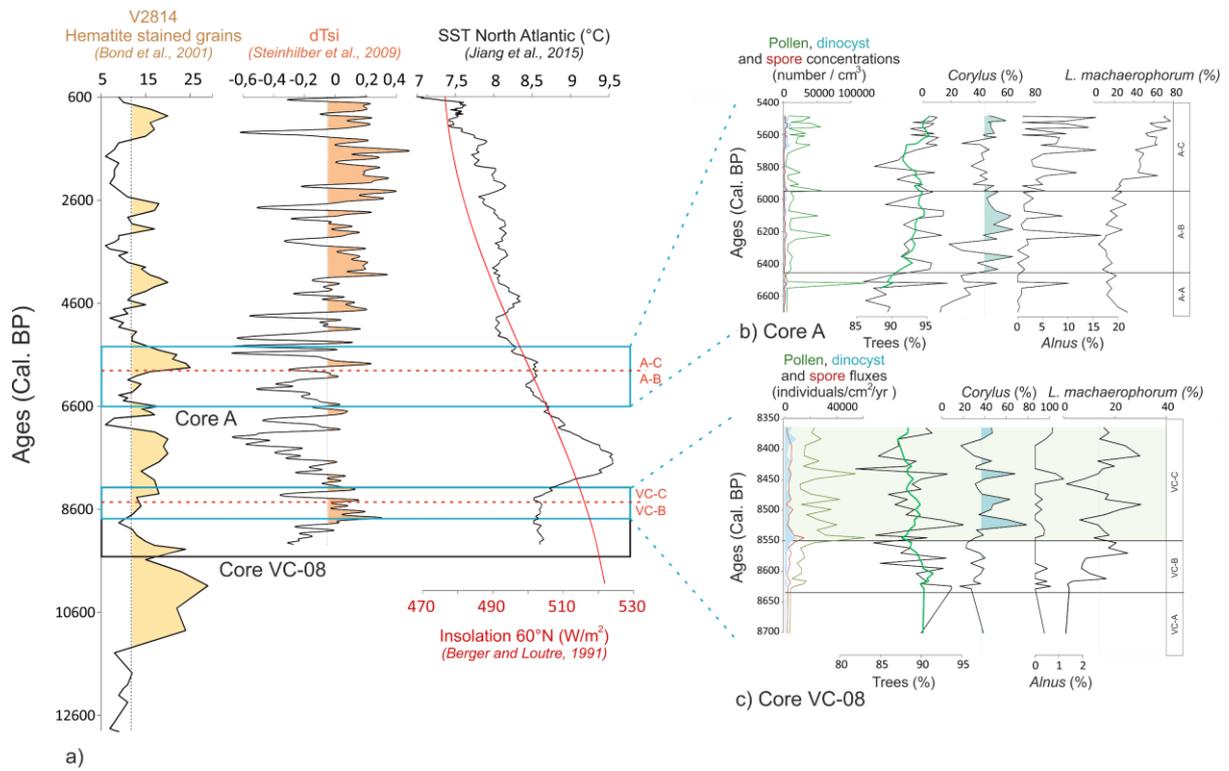


Figure 12

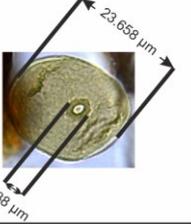
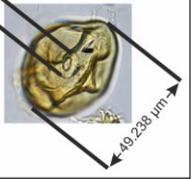
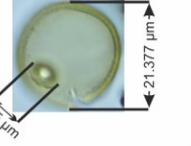
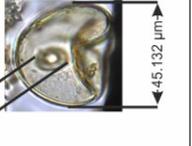
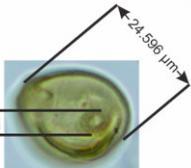
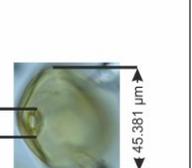
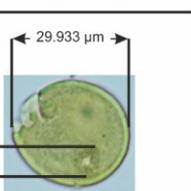
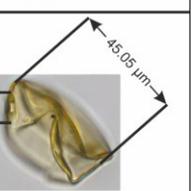
| Core VC-08 | a) Poaceae | b) Cerealia-type |
|-------------------------|---|--|
| 120 cm 8441 years BP |  |  |
| 140 cm 8457 years BP |  |  |
| 220 cm 8524 years BP |  |  |
| 325 cm 8612 years BP |  |  |

Figure 13

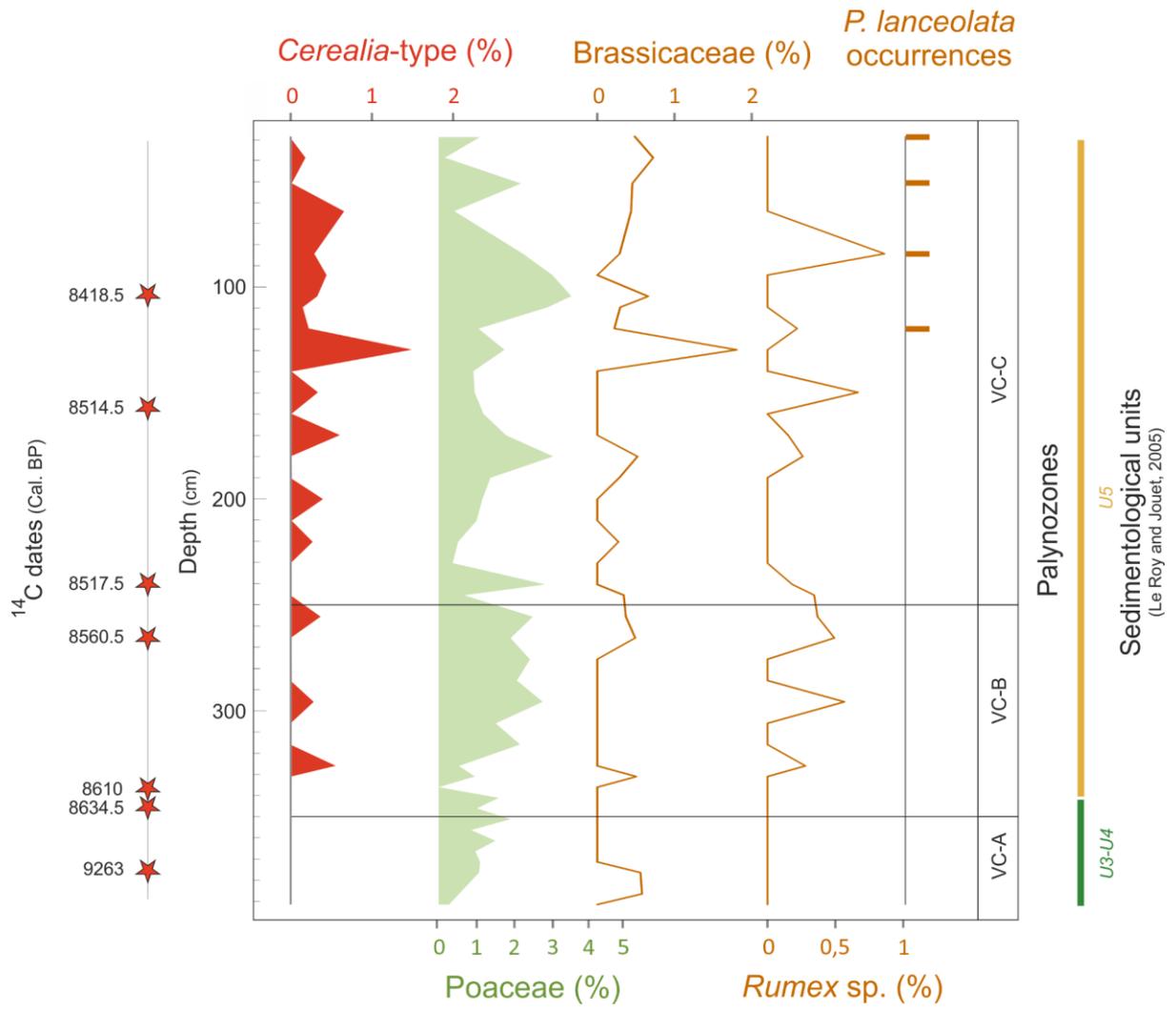


Figure 14



1. Cupillard et al., 1994 ; 2. Nielsen, 2003 ; 3. Lotter, 1999 and Tinner et al., 2007 ; 4. Beckmann, 2004 ; 5. Erny-Rodmann et al., 1997 ; 6. Tinner et al., 1999 ; 7, 8. Triat-Laval, 1978 ; 9. Puertas, 1999 ; 10. Guenet, 1995 ; 11. Jalut and Vernet, 1989 ; 12. Amat, 1995 ; 13. Galop and Vaquer, 2004 ; 14. Legigan and Marambat, 1993 ; 15. Joly and Visset, 2005 ; 16. Joly and Visset, 2009 ; 17. Visset et al., 1996 ; 18. Ouguerram and Visset, 2001 ; 19. Visset et al., 2001 ; 20. Carcaud et al., 2000 ; 21, 22. Leroyer and Allenet, 2006 ; 23, 24. Ruffaldi, 1999 ; 25. This study

Figure 15