

1 **Deglacial and Holocene vegetation and climatic changes at the southernmost tip of the Central**
2 **Mediterranean from a direct land-sea correlation.**

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19 **Abstract**

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21 Despite a large number of studies, the long-term and millennial to centennial-scale climatic
22 variability in the Mediterranean region during the last deglaciation and the Holocene is still debated,
23 **in particular in the southern Central Mediterranean.** In this paper, we present a new marine pollen
24 sequence (MD04-2797CQ) from the Siculo-Tunisian Strait documenting the regional vegetation and
25 climatic changes in the southern Central Mediterranean during the last deglaciation and the
26 Holocene.

27 The MD04-2797CQ marine pollen sequence shows that semi-desert plants dominated the
28 vegetal cover in the southern Central Mediterranean between **18** and 12.3 ka **BP** indicating prevailing
29 dry conditions during the deglaciation, even during the Greenland Interstadial **(GI)-1. Such** arid
30 conditions likely restricted the expansion of the trees and shrubs despite the GI-1 climatic
31 amelioration. Across the transition Greenland Stadial (GS)-1 – Holocene, Asteraceae-Poaceae steppe
32 became dominant till **10.1ka.** This record underlines with no chronological ambiguity that even
33 though temperatures increased, deficiency in moisture availability persisted into the Early
34 Holocene. **Temperate** trees and shrubs with heaths as oak forest understorey or heath maquis

35 expanded between 10.1 and 6.6 ka, while Mediterranean plants only developed from 6.6 ka
36 onwards. These changes in vegetal cover show that the regional climate in southern Central
37 Mediterranean was wetter during Sapropel 1 (S1) and became drier during the Mid- to Late
38 Holocene. Wetter conditions during S1 were likely due to increased winter precipitation while
39 summers remained dry. We suggest, in agreement with published modelling experiments, that the
40 increased melting of the Laurentide Ice Sheet between 10 to 6.8 ka in conjunction with weak winter
41 insolation played a major role in the development of winter precipitation maxima in the
42 Mediterranean region in controlling the strength and position of the North Atlantic storm track.

43 Finally, our data provide evidences of centennial-scale vegetation and climatic changes in the
44 southern Central Mediterranean. During the wet Early Holocene, alkenones-derived cooling episodes
45 are synchronous to herbaceous composition changes that indicate muted changes in precipitation. In
46 contrast, enhanced aridity episodes, as detected by strong reduction in trees and shrubs, are
47 recorded during the Mid- to Late Holocene. We show that the impact of the Holocene cooling events
48 depend on the baseline climate states insolation and ice sheet volume, shaping the response of the
49 mid-latitude atmospheric circulation.

50

1. Introduction

The Mediterranean region received increasing attention during the past decade when it was identified as a major “climate change hot spot” in future climate modeling projections as a result of the global increase in greenhouse gases (Giorgi, 2006; IPCC, 2007). Sensitivity of the Mediterranean climate and ecosystems to global climate change during glacial and interglacial periods has been demonstrated by a numerous paleoclimatic studies. Holocene Mediterranean records show that the long-term Holocene climatic evolution in response to orbital forcing can be described by three phases: a humid phase from 11.5 to 7 ka, a transition phase from 7 to 5.5 ka and an aridification phase from 5.5ka to present-day (Jalut et al., 2009; Finné et al., 2011; Pérez-Obiol et al., 2011). However, Holocene environmental conditions in the Mediterranean region remain a matter of debate. Davis and Brewer (2009) even hypothesized that higher Early Holocene moisture availability was not related to increased precipitation but decreased temperature. The respective contribution of climate and human activities to environmental changes, the contrasting and complex information provided by different proxy records and the heterogeneity and complexity of this region contribute to the existing uncertainties on the Holocene climate evolution in the different regions of the Mediterranean (Tzedakis, 2007; Roberts et al., 2011a). This is particularly true for the southern Central Mediterranean as shown by an increasing number of lake level, pollen and speleothem records depicting contrasting signals. Some paleorecords from Sicily show wetter conditions (Sadori and Narcisi, 2001; Magny et al., 2011b; Peyron et al., this volume-b) while others indicate drier conditions during the Early Holocene (Tinner et al., 2009; Calò et al., 2012). Note that “Early Holocene” is defined here as the 11.7 to 6.5 ka interval and “Mid- to Late Holocene” the interval from 6.5 ka to present-day. On the southern rim of the Mediterranean Sea, peat sequences from Kroumirie (Ben Tiba and Reille, 1982; Ben Tiba, 1995; Stambouli-Essassi et al., 2007) give a synthetic view of the Holocene vegetation history in the humid mountainous NW Tunisia. More detailed sequences in this area and in the semi-arid plains are needed to get a complete description of the Holocene climatic changes in northern Tunisia. In addition, delayed forest expansion appears to be recorded by few sites from the southernmost areas of the Mediterranean region suggesting that dry conditions may have persisted into the Holocene (Tzedakis, 2007). A clearer picture of the long-term climatic changes in the southern Central Mediterranean is a crucial step towards a mechanistic understanding of the regional hydroclimatic changes in response to large-scale orbital climatic changes.

Deglacial vegetation changes in the Central Mediterranean are based on few pollen sequences mainly located in central Italy and Adriatic Sea (Watts et al., 1996; Combourieu-Nebout et

85 al., 1998; Magri, 1999; Magri and Sadori, 1999). These records clearly show the afforestation
86 associated to the climatic improvement of the last deglaciation as well as the response of the
87 Mediterranean forest to millennial-scale climatic changes. However, the low temporal resolution of
88 available records (Ben Tiba and Reille, 1982; Brun, 1983) does not allow to evaluate the impact of the
89 millennial-scale variability during the last deglaciation. Since the identification of abrupt changes in
90 the North Atlantic during the Holocene (Bond et al., 1997), a number of publications have shown that
91 such changes affected the Mediterranean Sea and borderlands (e.g. synthesis papers and references
92 therein: Magny et al., 2007a; Fletcher and Zielhofer, 2011; Peyron et al., 2011; Roberts et al., 2011b;
93 Sadori et al., 2011). The complexity of the Mediterranean climate as well as the interpretation of
94 proxies and chronological uncertainties remains a major limitation to our understanding of abrupt
95 climatic changes. Records from Central Mediterranean provide evidences of increased dryness during
96 the Holocene cooling episodes which contrasts with the increase in humidity in the European mid-
97 latitudes, above 40°N (Magny et al., 2011b). Recently, a change in nature and tempo of centennial-
98 scale variability during the mid-Holocene has been identified in the SW Mediterranean (Fletcher et
99 al., in press). However, key questions remain about the impact of Holocene cooling events on the
100 Mediterranean climate and ecosystems and the processes responsible for the propagation and
101 modulation of the North Atlantic variability into the Mediterranean region.

102 In this paper, we present pollen data from a marine record retrieved from the Siculo-Tunisian
103 strait providing a regional picture of climate and vegetation changes in the southernmost Central
104 Mediterranean during the last deglaciation and the Holocene. Our record based on a direct land-sea
105 correlation will bring new insight into regional long-term trend and millennial to centennial-scale
106 climatic changes. We compare our results with other records from Central and Western
107 Mediterranean to show a coherent pattern between records. Finally, we discuss the potential links
108 between the Mediterranean climate evolution and the low and mid-latitude atmospheric circulation
109 features in relation with insolation and ice sheet volume forcings.

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2. Environmental setting and potential pollen source

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114 Core MD04-2797CQ (36°57'N 11°40'E, 771 m water-depth) was retrieved from the Siculo-
115 Tunisian Strait at ~50km east off Cape Bon at the tip of the north-eastern Tunisian coast and at ~110
116 km off Sicily (Fig.1). The strait bottom morphology is characterized by a sill and two troughs
117 separating the Sicilian and Tunisian continental shelf areas. The upper water column is occupied by
118 the Modified Atlantic Water (MAW). This water mass is fed by the Atlantic Tunisian Current (ATC)
119 which is the southern branch of the MAW flowing eastward along the Tunisian coast through the
120 strait while the northern branch flows along the Sicilian coast (Fig. 1) (Béranger et al., 2004). The
121 upper water mass above the core site is influenced by the ATC. At depth, waters originating from the
122 Eastern Mediterranean flowing westward across the Siculo-Tunisian Strait are composed of Levantine
123 Intermediate Water and transitional Eastern Mediterranean Water (Astraldi et al., 2002). The surface
124 ocean circulation is driven by the atmospheric circulation dominated by westerly to north-westerly
125 winds (Pinardi et al., 2005).

126 The present-day climate in northern Tunisia and Sicily is of Mediterranean-type,
127 characterized by warm and dry summers and cool and wet winters. The orography and maritime
128 influences modulate the climate in both regions. An West to East humidity gradient exists in northern
129 Tunisia between the humid to sub-humid northwestern mountains and the semi-arid northeastern
130 lowlands while in southern Tunisia arid conditions prevail (INRF, 1976). The Sicily climate ranges from
131 semi-arid to sub-humid at the highest elevations of the island (Drago, 2005). The autumn-winter-
132 spring seasonal precipitation mainly derives from moisture advection from the Atlantic Ocean. The
133 western Mediterranean Sea is another source of moisture for the surrounding landmasses through
134 evaporation and atmospheric circulation. Summer dryness results from the subsidence development
135 associated with the northward displacement of the Hadley cell (Lionello et al., 2006).

136 The vegetation distribution in the Central Mediterranean is controlled by temperature and
137 precipitation gradients and seasonality. The potential vegetation cover of the Medjerda watershed is
138 dominated by *Olea-lentiscus* maquis in the eastern semi-arid areas and by holm oak-aleppo pine
139 forest in the southern upper semi-arid High Tell and Tunisian Dorsal. In the most north-western
140 Kroumirie region of the Medjerda watershed, cork oak with *Erica arborea* underbrush and high
141 altitude zeen oak forest prevail due to humid and sub-humid conditions in the mountains. Cape Bon
142 considered as the “backbone” of the Tunisian Dorsal is the closest landmass to our core site where
143 moisture from maritime influence leads to semi-arid to sub-humid conditions. Vegetation of Cape
144 Bon is dominated by *Oleo-lentiscus* maquis, kermes oak (*Quercus coccifera*) and Thuya (Gausson and
145 Vernet, 1958; Posner, 1988; El Euch, 1995; Boussaid et al., 1999).

146 In addition, numerous permanent or temporary wetlands are located in Tunisia (Posner,
147 1988) mostly in the humid, sub-humid and upper semi-arid bioclimatic zones (INRF, 1976). Wetlands
148 from northern Tunisia such as Garaet el Ichkeul are characterized by the dominance of Cyperaceae in
149 the marsh zone although recently replaced by Chenopodiaceae due to decreasing freshwater inflow
150 as a consequence of dams and pipeline constructions (Ramdani et al., 2001). In temporary wetlands
151 such as Sebkhet Ariana located in the semi-arid region, north-east of Tunis, vegetation is dominated
152 by halophyte plants in particular Chenopodiaceae (Posner, 1988).

153 In Sicily, vegetation is distributed along humidity and temperature gradients resulting from
154 the orography. *Oleo-ceratonion* maquis dominates the lowlands in the warmest and driest areas of
155 the SE Sicily. At higher elevations, the potential vegetation has been assigned to mesophilous
156 Mediterranean evergreen woods (*Quercetum ilicis*). Increased humidity in the submontane areas
157 allows the development of deciduous woodlands (*Quercetalia pubescenti-petraeae*) while at highest
158 altitudes of Sicily, *Geranio versicoloris-Fagion* association is favoured (Morinello, 1999).

159
160 A number of studies have shown that marine pollen signal provides an integrated picture of
161 the vegetation from the nearby continents (Heusser and Balsam, 1977; Naughton et al., 2007).
162 Studies on present-day pollen deposition in marine sediments show that pollen grains are fluvial- and
163 wind-transported from the adjacent continent and that agglomeration, flocculation and zooplankton
164 feeding processes enable rapid sinking of pollen through the water column (Heusser and Balsam,
165 1977; Heusser, 1978; Chmura et al., 1999; Dupont and Wyputta, 2003; Mudie and McCarthy, 2006).
166 Further, cores such as MD04-2797 CQ located near continental regions with well-developed
167 hydrographic basins and prevailing offshore winds, mainly recruit pollen from rivers (Heusser, 1978;
168 Turon, 1984; Dupont and Wyputta, 2003). Pollen grains belonging to the fine-particle fraction have a
169 similar behaviour as fine sediments during the sedimentary processes (Muller, 1959; Chmura and
170 Eisma, 1995). Medjerda and Miliane oueds are the two most important and permanent water
171 streams in Tunisia discharging into the Gulf of Tunis and providing sediments to the shelf area (Fig.
172 1). The Medjerda oued is the main Tunisian stream flowing SW-NE between the Northern Tell and
173 High Tell mountains, from eastern Algeria to the Gulf of Tunis. Floods are recurrent in northern
174 Tunisia which receives most rainfall in winter and spring. Floods enhance material transport and
175 contributed to the recent filling in of the Gulf of Tunis (Zahar et al., 2008). The riverine system of
176 Sicily is not well developed and climate varies from semi-arid in the low land to sub-humid upland.
177 Wind-driven sediment input to the shelf might be important in semi-arid regions. Given the
178 dominant Westerly winds and surface ocean circulation, the bathymetry of the strait and large river
179 systems of northern Tunisia, the southern Central Mediterranean and most probably also northern
180 Tunisia are likely be main source of pollens to our core site. Therefore, even though long-range

181 transport of pollen cannot be ruled out, we interpret the pollen signal of core MD04-2797CQ as
182 reflecting vegetation changes in this region.

183

184 **3. Material and methods**

185

186 Sediment core MD04-2797CQ was subsampled for pollen analysis every 2 to 10 cm over the
187 first 590 cm. The sample preparation technique followed the procedure described by de Vernal et
188 al.(1996). Exotic pollen (*Lycopodium*) was added to each sample to determine pollen concentrations.
189 After chemical and physical treatments (cold HCl, cold HF, cold KOH and sieving through 10 µm nylon
190 mesh screens), the residue was mounted unstained in glycerol. Pollen was counted using a light
191 microscope at 400 and 1000 (oil immersion) magnifications. 100 to 160 pollen grains without *Pinus*
192 and a minimum of 20 taxa were counted in each of sample. 10 out of 94 samples analysed were
193 considered as sterile due to extremely low pollen concentration, in particular in the upper 80 cm of
194 the core. Pollen percentages for each taxon are based on the main pollen sum excluding *Pinus*
195 because of its over-representation in marine deposits (Heusser and Balsam, 1977; Turon, 1984).
196 *Pinus* percentages were calculated from the main pollen sum including *Pinus*. Spores and aquatic
197 pollen percentages were obtained from the total pollen sum that includes all pollen grains plus
198 aquatic plant pollen grains, Pteridophyta spores and indeterminable pollen grains.

199 Quantitative climatic reconstruction based on the Modern Analogue Technique (MAT) was
200 performed from pollen data. The MAT have been used to construct here, annual, winter and
201 summer temperatures and precipitation by considering the proportion of pollen assemblages (Guiot,
202 1990). Methods and results are presented and discussed in details in Peyron et al. (this volume-a).

203 Direct land-sea correlation is obtained in comparing pollen to marine proxies from MD04-
204 2797CQ sediments. Dinocyst and planktic foraminifera assemblages and derived SSTs, planktic
205 foraminifera $\delta^{18}\text{O}$ and SST obtained from the C_{37} alkenones unsaturation index have been published
206 earlier (Essallami et al., 2007; Rouis-Zargouni et al., 2010). In this paper, we compare our pollen data
207 with the new high resolution alkenone-derived SST record (Sicre et al., this volume).

208

209 **4. Results and Interpretations**

210 **4.1. Chronological framework and pollen analysis results**

211 Core MD04-2797CQ chronology was established from 13 radiocarbon dates among which 6
212 are from the upper 4 m of the core corresponding to the Holocene interval (Table 1). Except for the
213 date at 0 cm, all radiocarbon dates have been replicated, thus increasing the reliability of the age-
214 depth model. Averages of the ^{14}C date duplicates were corrected for a reservoir age of 400 yr for the

215 Holocene, GS-1 and LGM intervals, of 560 yrs for GI-1 (level 470 cm) and of 800 yrs for Heinrich
216 Stadial 1 (HS1) (level 510 cm), according to Siani et al. (2001). Corrected dates were calibrated using
217 the calibration curve INTCAL 09 (Reimer et al., 2009). The age-depth model was obtained using linear
218 interpolation between dated levels (Fig. 2). The upper 80 cm represents the last 5800 years and is
219 only chronologically constrained by two radiocarbon dates (5830 and 670 yr cal BP). A strong change
220 in sedimentation rate occurred in between the two dated levels increasing age uncertainties for this
221 interval. Here after, ka and kyr will be used for ages in cal BP.

222 According to the age-depth model (Fig.2), the MD04-2797CQ pollen record encompasses the
223 last 18, 000 years. Pollen analyses were performed at high temporal resolution (less than 100 years)
224 for the Holocene interval between 12 and 5.5 ka. Unfortunately, the low pollen richness and
225 sedimentation rate did not allow a high resolution study of the upper 80 cm. Most of the samples of
226 the upper 40 cm were even pollen barren making the pollen record only reaches 2.2 ka. The
227 alkenone-SST record also displays the highest temporal resolution (20 to 75 years) over the first half
228 of the Holocene (Sicre et al., this volume).

229 Pollen zones (PZ) were identified visually from the pollen diagram shown in Figure 3 and
230 confirmed using constrained hierarchical clustering analysis based on Euclidean distance between
231 samples using *chclust* function from the R package *Rioja* (Juggins, 2009). Clusters were determined
232 from filtered pollen data at 1percent taxa presence in at least 5 percent of the samples using the R
233 package *PaleoMas* (Correa-Metrio et al., 2011). A summary of the pollen diagram description is
234 presented in Table 2.

235

236 4.2. Deglacial vegetation and climatic changes

237 The core MD04-2797CQ pollen record depicts the deglacial and millennial-scale vegetation
238 changes from southern Central Mediterranean, a region for which few terrestrial paleodata are
239 available. In particular, pollen records from Tunisia do not resolve the deglacial millennial-scale
240 variability because of low temporal resolution or hiatuses (Ben Tiba and Reille, 1982; Brun, 1985;
241 Stambouli-Essassi et al., 2007).

242 Our marine pollen record shows that a steppe with semi-desert plants (*Artemisia*,
243 *Ephedradistachya*, *Ephedrafragilis*, Chenopodiaceae) dominates the vegetal cover up to 12.3ka,
244 indicating prevailing dry conditions during the last deglaciation (PZ-1-2-3; Fig. 3 and 4). Such arid
245 environments likely characterized most of Tunisia during this period, from at least 34°N (Brun, 1985)
246 to the northern Tunisian coast (this work). However, the moister mountainous NW part of Tunisia
247 provided a favourable habitat for an open oak forest and a conifer forest of pine-cedar-fir in the
248 highest altitudinal vegetation belts (Ben Tiba and Reille, 1982; Stambouli-Essassi et al., 2007).

249 Two intervals of maximal semi-desert plant expansion, coeval with cold SSTs occurred
250 between 18.2 and 14.7ka (PZ-1) and 13.1 and 12.3ka (PZ-3) (Fig. 3 and 4) broadly at the time of HS 1
251 and GS-1, respectively. These time intervals are major abrupt climatic changes interrupting the last
252 deglaciation that are widely documented as strong cooling in the North Atlantic, Europe and over
253 Greenland (Johnsen et al., 1992; Walker, 1995; Bard et al., 2000; Peterson et al., 2000; Naughton et
254 al., 2007). These events presumably result from the abrupt reduction in Atlantic Meridional
255 Overturning Circulation (AMOC) due to massive iceberg discharges or freshwater releases in the
256 North Atlantic (MacAyeal, 1993; Bond and Lotti, 1995; McManus et al., 2004; Broecker, 2006). Our
257 sequence indicates that in southern Central Mediterranean, climate shifted toward cold and dry
258 conditions in response to HS 1 and GS-1 (Fig. 4). Prevailing arid conditions in Tunisia during GS-1 are
259 confirmed by large aeolian terrigenous supply as detected by high illite, palygorskite and zirconium
260 contents in our core (Bout-Roumazeilles et al., this volume). Similar climatic variations were reported
261 from Western Mediterranean marine pollen and SST records (Cacho et al., 2001; Fletcher and
262 Sánchez Goñi, 2008; Combourieu Nebout et al., 2009). Despite extremely arid conditions, our record
263 indicates that deciduous tree populations, mainly deciduous *Quercus*, probably persisted in some
264 remaining humid areas, and that the conifers *Cedrus* and *Pinus* were present in the regional
265 vegetation. Some deciduous taxa pollen grains may originate from Sicily or even from southern Italy
266 where scattered woodlands have been detected (Magri and Sadori, 1999; Allen et al., 2002).
267 However, most of the tree pollen grains recovered at our site during HS 1 and GS-1 were likely
268 transported from NW Tunisia uplands by wind or river (Medjerda) where open oak and conifer
269 forests were present during the last glacial and deglaciation (Ben Tiba and Reille, 1982; Stambouli-
270 Essassi et al., 2007). Note the increase in *Cedrus* abundances during HS1 and GS-1 with a corresponding
271 oak decrease (Fig. 3). A similar pattern has been described in marine pollen records from the Alboran
272 Sea documenting cedar variations in the Moroccan Rif (Fletcher and Sánchez Goñi, 2008;
273 Combourieu Nebout et al., 2009). As discussed in Fletcher and Sanchez Goñi (2008), increases in
274 cedar pollen abundances could reflect either 1) enhanced wind-driven pollen supply due to wind
275 strengthening and opening of the vegetation or 2) maintenance of some degree of moisture since
276 cedar is generally considered as a cold tolerant-moisture demanding conifer. Colder conditions and
277 moisture availability at mid to high altitudes during HS 1 and GS-1 could have favoured cedar
278 expansion at the expense of deciduous oak in NW Africa, from the Moroccan Rif to the High Tell. The
279 Mediterranean region presents a wide range of habitats due to highly variable topography and local
280 climate conditions which enables refugia zones during the last glacial period (Médail and Diadema,
281 2009). This work supports the idea that tree populations maintained in the moist sub-humid region of
282 North Africa even during the most severe episodes of the last deglaciation.

283 During GI-1, SSTs increased, semi-desert plants and cedar reduced while some deciduous and
284 evergreen oak woodlands and Mediterranean shrubs (*Olea*) developed (PZ-2; Fig. 4). This finding is
285 indicative of relatively warmer and wetter climatic conditions than during HS 1 and GS-1. Despite the
286 temperature rise, the semi-desert plants remained dominant and tree and shrubs expansion limited
287 compared to terrestrial pollen records from Italy (Magri and Sadori, 1999; Allen et al., 2002) or
288 western Mediterranean Sea pollen records (Fletcher and Sánchez Goñi, 2008; Combourieu Nebout et
289 al., 2009). This result can be explained by lower water availability in the semi-arid areas of southern
290 central Mediterranean. Terrigenous supply to the Siculo-Tunisian Strait remained characterized by a
291 high content of aeolian-driven particles, such as illite and palygorskite, which also indicate prevailing
292 dry conditions during GI-1 (Bout-Roumazeilles et al., this volume). Strikingly, our pollen data do not
293 indicate a tree and shrub expansion maximum at the beginning of the GI-1 but rather slightly higher
294 values at the end of the interstadial at ~13.4 ka following a long HS1 to GI-1 transition. This feature
295 contrasts with the sharp temperature rise and early GI-1 optimum in Greenland ice cores (NGRIP
296 members, 2004)(Fig. 4) and has already been identified from speleothem records from northern
297 Tunisia, southern France and southern Germany (Genty et al., 2006) as well as from pollen records
298 from the Alboran Sea and southern Alps (Combourieu Nebout et al., 2009; Finsinger et al., 2011). This
299 North-South contrast may be linked to the North Atlantic oceanic and atmospheric circulations and
300 sea ice cover (von Grafenstein et al., 1999; Genty et al., 2006). In the Southern Alps, the gradual
301 expansion of *Quercus* have been attributed to increasing summer temperatures and longer growing
302 season probably related to insolation (Finsinger et al., 2011). Stacked temperature records over the
303 latitudinal bands 30-0°S and 0-30°N (Shakun et al., 2012) indicate lower temperature during GI-1
304 than during the Holocene and display the same peculiar increasing temperatures over the course of
305 the interstadial. Since tropical ocean temperature and associated evaporation regulate the
306 atmospheric moisture content feeding the North Atlantic storm track (Braconnot et al., 2007),
307 stronger inland moisture transport toward the end of GI-1 may be associated with the tropical
308 climate in relation with orbital forcing.



309 310 **4.3. Holocene long-term vegetation and climate changes**

311 **4.2.1. Vegetation and climate in and off Tunisia**

312 In core MD04-2797CQ, the GS-1 - Holocene transition is marked by a sharp decrease in semi-
313 desert plants at ~12.3ka while SST-alkenones and summer SSTs derived from planktonic foraminifera
314 assemblages increased by 9°C (Essallami et al., 2007) (PZ-4; Fig. 3 and 4). These vegetation and SST
315 changes seem to precede by 600 years the sharp rise in temperature recorded in Greenland
316 (Rasmussen et al., 2006). One can attribute this time difference to age uncertainties of the records.
317 However, this difference is quite large and may suggest earlier changes in temperature and

318 precipitation in the southern Mediterranean region compared to the northern latitudes. Open
319 vegetation with Asteraceae, Poaceae and some semi-desert plants (mainly Chenopodiaceae and
320 *Ephedra fragilis*-type) remained dominant until 10.1 ka when the open oak forest expands. Although
321 Mediterranean and temperate trees and shrubs slightly increased at 12.3ka, they remained restricted
322 at some woodlands/scrubs despite high SST-alkenones (Fig. 4). This vegetation change indicates a
323 shift to warmer conditions and increased humidity during the Early Holocene but moisture
324 availability remained too weak before 10.1ka to enable a forest development. In the southern
325 Mediterranean, temperature influences forest composition but moisture availability is overall critical
326 for forest development (Quezel, 2002). During this interval, decreasing semi-desert plants and
327 increasing Cyperaceae and Ericaceae values suggest a gradual increase in moisture availability (Fig.
328 4). Therefore, the abrupt tree and shrub expansion observed at 10.1ka would likely result from
329 crossing ecological thresholds during gradual climatic change. The gradual decrease in aridity before
330 10 ka is supported by mineralogical and geochemical data found at core site MD04-2797CQ (Bout-
331 Roumazeilles et al., this volume).

332 From 10.1 to 6.6ka, open deciduous oak forest with heath understorey and maquis
333 developed along with a maximal sedge expansion (PZ-5). Mediterranean plants (mainly *Pistacia*,
334 evergreen *Quercus* and *Olea*) remained limited over this interval and only start expanding after 6.6 ka
335 at the expense of heath maquis (PZ-6). At present-day, in the Central Mediterranean and in particular
336 in Tunisia, Ericaceae are mainly represented by *Erica arborea*, *E. scoparia* and *E. multiflora* (Posner,
337 1988; Ojeda et al., 1998). Although *E. arborea* and *E. scoparia* have a wide range of distribution, they
338 are related to humid climate and particularly atmospheric humid conditions (Jalut et al., 2000). This is
339 well illustrated by the Ericaceae distribution in northern Tunisia. Both species are widely distributed
340 in the sub-humid zones of NW Tunisia and Cape Bon as understorey of *Quercus canariensis* and
341 *Quercus suber* or as maquis of heather, while *E. multiflora* mainly grows in the *Olea-lentiscus* maquis
342 or with aleppo pine and holm oak association in the semi-arid areas of northern Tunisia (Posner,
343 1988). Therefore, the development of Ericaceae along with deciduous trees suggests that the most
344 humid conditions of the Holocene occurred during this phase. Another line of evidence supporting
345 this interpretation is the concurrent maximal development of sedges. Cyperaceae is a family
346 comprising numerous species that can grow under a wide range of ecological conditions although
347 most of them develop in humid areas (Brun, 1985). Therefore, increased precipitation can potentially
348 favour their contribution in the Asteraceae-Poaceae steppes. Sedge expansion can also reflect
349 enhanced humid habitats, such as inland or coastal wetlands that are common features in northern
350 Tunisia. For instance, extensive sedge marshland surrounded Garaet el Ichkeul lake until dam
351 constructions dramatically decreased water supply (Ramdani et al., 2001). Part of the strong sedge
352 representation in our record might also result from coastal environment modifications related to

353 post-glacial sea-level rise up to 6.6ka. The highest sedge percentages are also seen in MD95-2043
354 and ODP 976 sites during the same period, although lower than in our sequence. In these Alboran
355 Sea records, sedges are associated with maximum forest development sustained by the most humid
356 conditions of the Holocene (Fletcher and Sánchez Goñi, 2008; Combourieu Nebout et al., 2009). Our
357 pollen record therefore suggests that the 10-6.6 ka interval is the wettest Holocene phase. However,
358 in NE Tunisia water availability was never high enough, even during the most humid phase to sustain
359 the regional development of a closed forest. Decreasing Ericaceae and Cyperaceae between 6.6 and
360 5 ka (PZ-6) suggest a transition toward drier conditions. After 5 ka, the expansion of Mediterranean
361 plants such as evergreen *Quercus* and *Olea* suggests lower available summer moisture and warmer
362 winter.

363 Precipitation estimates derived from our pollen record support the earlier interpretation of
364 highest rainfall between 10 and 6.6 ka, occurring mainly in winter, with mean annual precipitation of
365 ~670 mm (Fig. 5, Peyron et al., this volume-a). This period coincides with the highest pollen
366 concentrations (Fig. 4) possibly due to both strong terrigenous inputs and increase biomass. After 5
367 ka, while dryness increased on land, the SSTs show moderate warming, although different time-scale
368 variability resulting from different temporal resolution may introduce bias. This warming seems
369 however to be confirmed by increasing abundances of warm water dinocysts (Rouis-Zargouni et al.,
370 2010).

371 Interpretation of our pollen record may appear contradictory regarding the gradual increase
372 of temperate trees and shrubs all along the Holocene. Changes in post-glacial sea level may have
373 influenced our marine pollen signal by modifying the dominant pollen transport agent from fluvial to
374 aeolian. Mediterranean post-glacial sea-level reconstructions suggest that Cape Bon coastline and
375 Medjerda river mouth were closer to our core site until 6.5 ka when the Laurentide ice sheets (LIS)
376 melting was completed (Lambeck et al., 2004). This could have led to enhance the representation of
377 lowland relative to upland vegetation up to 6.5 ka and that of the upland open forest afterwards.
378 However, a shift to drier climate could have promoted aeolian transport. Although pollen transport
379 changes may have modified the pollen signal, they do not alter the main vegetation changes. In
380 particular, changes in the lowland shrub communities such as the replacement of heath maquis by
381 *oleo-lentiscus* scrubs remain reliable to evaluate the main traits of the Holocene vegetation and
382 climate history. Sedimentological analyses of core MD04-2797CQ agree with our interpretation of
383 the pollen record. High sedimentation rates, decrease of aeolian-driven clay particles (palygorskite
384 and illite), enhanced contribution of smectite likely reflecting fluvial supply and higher Zr/Al ratio
385 indicate increased sediment supply to the shelf. Higher riverine discharge are likely sustained by
386 more precipitation between 6 and 9.5 ka (Bout-Roumazielles et al., this volume).

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4.3.2. Climatic or human-induced vegetation changes

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The respective role of climatic change and anthropogenic pressure since the Neolithic has been a long standing debate in our understanding of the Mediterranean vegetation changes recorded by pollen sequences (De Beaulieu et al., 2005; Jalut et al., 2009). In Sicily, human impact appears to be detected in paleorecords since the Bronze Age and not since the Neolithic (Sadori and Giardini, 2008), and intense land use began with the Greek and Roman periods (Tinner et al., 2009). Human impact on the Tunisian environments became noticeable from the Phoenician and Carthaginian epochs (from XIIth to IIth century BC) with a strong agricultural development essentially based on cereal cultivations (Brun, 1983). However, our pollen record does not show this cereal signal. It is known that the dispersal of the large sized cereal pollen grains is poor (Vuorela, 1973), making difficult to get a reliable representation of cultivated Poaceae in marine pollen records. The Late Holocene *Olea* expansion is clearly detected in our record at around 3.2 ka that may result from human activities during the Bronze Age. Olive tree expansion is also detected at that time in sequences from the Adriatic Sea, Italy and Sicily (Sadori and Narcisi, 2001; Di Rita and Magri, 2009; Combourieu Nebout, this volume). Management of olive tree populations for fruit production started as early as the Bronze Age in the Western Mediterranean (Terral et al., 2004). However, as shown by our record, the expansion of *Olea* in southern Italy is accompanied by a widespread increase of other Mediterranean trees and shrubs that may thus reveal a concomitant vegetation response to climate during the Bronze Age (Di Rita and Magri, 2009). Because of the low chronological constraints in our sequence between 600 and 5800 years BP, *Olea* increase may also have occurred during Roman times. Expansion of olive trees at around 2000 years cal BP during Roman occupation is detected in southern Italy and Sicily (Terral et al., 2004; Di Rita and Magri, 2009). Roman developed an intensive olive tree cultivation for fruits and oil in the Mediterranean region (Terral et al., 2004). Therefore, human influence cannot be discarded since the Bronze Age but beforehand our pollen record primarily reflects climatic conditions.

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4.3.3. Major climatic changes in the southern Central Mediterranean

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Land-sea correlation from core MD04-2797CQ indicates without chronological ambiguity that Holocene expansion of tree populations in southern Central Mediterranean lagged temperature increase by two millennia. Delayed or slow afforestation due to moisture deficiency in the southernmost areas of the Mediterranean has been previously suggested from few terrestrial records (Tzedakis, 2007; Tinner et al., 2009). Dryness in Sicily before 10 ka is supported by lake level and pollen data (Fig. 6g-h-i) (Sadori and Narcisi, 2001; Tinner et al., 2009; Magny et al., 2011b). Close examination of the Tunisian pollen records from Kroumirie (Stambouli-Essassi et al., 2007) and from Gulf of Gabes (Brun, 1979, 1985) shows that the deciduous oak forest expansion in NW Tunisia and

423 of *Olea-Pistacia* scrubs in central eastern appears delayed by at least one millennium with respect to
424 the beginning of the Holocene (Fig. 6c). Despite loose chronological framework and temporal
425 resolution of the Tunisian pollen records, dry conditions of the Early Holocene derived from our
426 sequence appear to be a robust feature in northern Tunisia. In addition, the pollen and terrigenous
427 records from core MD04-2797CQ parallel the $\delta^{13}\text{C}$ signal of the Tunisian speleothem record from La
428 Mine cave (Fig. 6d) (Genty et al., 2006). Both records suggest a gradual climatic improvement and
429 vegetation development in southern Central Mediterranean immediately after the GS-1 cooling up to
430 the climate optimum around 10 ka. This is in contrast with Lake Preola record showing an abrupt
431 rather than gradual change in lake level (Magny et al., 2011b). This contrasting feature could be
432 season-related with a gradual change in winter precipitation from GS-1 to 10 ka and an abrupt
433 change in summer precipitation at 10 ka.

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435 Increased moisture availability during the Early Holocene, between 10 ka and 6.6ka, is also
436 consistent with available Tunisian and Sicilian records (Fig. 6). This accounts for example for the
437 expansion of forest upland at Lake Pergusa and Majen M'Hida (Sadori and Narcisi, 2001; Stambouli-
438 Essassi et al., 2007) and of *Pistacia* scrubs at the western Sicilian coastal sites Lake Preola and Gorgo
439 Basso (Tinner et al., 2009; Calò et al., 2012). After 6.6 ka, our data and most of the Sicilian and
440 northern Tunisian records show similar long-term climate changes. Pollen and sedimentary $\delta^{18}\text{O}$
441 records and pollen-derived precipitation estimates from Lake Pergusa, lake level record from Lake
442 Preola and stalagmite record from Grotta di Carburangeli suggest a transitional period of decreasing
443 humidity or high amplitude variability in rainfall in Sicily between the wet 10 to 7 ka interval and the
444 pronounced drier period from 5-4 ka to present-day (Frisia et al., 2006; Sadori et al., 2008; Magny et
445 al., 2011a; Peyron et al., this volume-b). Similarly, increasing dry conditions by 4-5 ka is reflected by
446 the replacement of the zeen oak forest by a cork oak forest with heath understorey at Dar Fatma in
447 the humid NW Tunisia (Ben Tiba and Reille, 1982). Alluvial records from the Medjerda floodplain (Fig.
448 6e) showing increased fluvial dynamics with recurrent floods since 5ka also suggest prevailing arid
449 conditions over the end of the Holocene (Zielhofer and Faust, 2008).

450 In contrast, the pollen records from the Sicilian coastal lakes Gorgo Basso, Lake Preola and
451 Biviera di Gela also show a pronounced vegetation change at **approx. 7-6.5 ka**. This change has been
452 interpreted as a second step of humidity increase caused by a decrease in summer dryness (Noti et
453 al., 2009; Tinner et al., 2009; Calò et al., 2012). In addition, in western Sicily, no shift is recorded by 5
454 ka, the evergreen forest declines only around 2.8-2.2 ka likely because of intensified anthropic
455 pressure on Mediterranean ecosystems (Tinner et al., 2009; Calò et al., 2012). However, the long-
456 term climatic variations derived from vegetation changes at Biviera di Gela are not unequivocal and
457 may as well be consistent with our results. Although maximum Mediterranean forest expansion

458 between 7 and 5 ka is interpreted as a response to moister climate at Biviera di Gela (Noti et al.,
459 2009), the reduction of mesophilous trees over this interval might also be indicative of a rather
460 contrasting drying trend. Furthermore, the partial shift from evergreen forest to *Pistacia*-dominated
461 shrublands at this site became pronounced by 5 to 4 ka, concurring with a Mid to Late Holocene
462 aridification reinforced by increasing human impact (Noti et al., 2009). A number of hypotheses,
463 although not entirely satisfactory, have been proposed to reconcile these coastal Sicilian pollen
464 records with Lake Preola lake-level record and Lake Pergusa pollen sequence, such as spatial
465 heterogeneity of landscape and climate (Tinner et al., 2009) or changes in seasonal and interannual
466 variability of drought (Magny et al., 2011b). Possible mis-assignment of *Pistacia* pollen taxa to the
467 evergreen species *P. lentiscus*, while pollen grains of the evergreen and deciduous species (*P.*
468 *terebinthus* and *P. atlantica*) are indistinguishable is another plausible explanation (Sadori et al.,
469 2011). Recently, Calo et al. (2012) also suggested that discrepancies between pollen and lake level
470 records at the Sicilian coastal sites would come from seasonal biases of each proxy.

471 Rainfall seasonality in response to orbital forcing is an important issue. Lower or higher
472 spring/summer dryness in southern Central Mediterranean between 10 and 7 ka than during the Late
473 Holocene has been deduced from continental records (Magny et al., 2011a; Calò et al., 2012) while
474 our marine record shows that summer precipitation (*Psum*) at regional scale remained low (50 to 100
475 mm) all over the Holocene, with no significant change (Fig. 5; Peyron et al., this volume-a). However,
476 even when *Psum* are estimated higher, they remained < 100 mm (Magny et al., 2011a). This denotes
477 that dry summers remained a characteristic of the Holocene climate in that region. Subtle variations
478 in *Psum* from semi-arid regions are particularly difficult to assess from pollen records since summer
479 moisture availability for plants not only depend on summer precipitations but also on groundwater
480 recharge during fall and winter that is critical to sustain vegetation during summer drought (Ez
481 2002; Fletcher et al., in press), on continental evaporation related to summer temperatures and
482 cloudiness, atmospheric humidity, storminess or soil absorption/retention. In contrast, Tunisian and
483 Sicilian pollen and speleothem records (Frisia et al., 2006; Magny et al., 2011a) agrees with our
484 observations (Fig. 5; Peyron et al., this volume-a) underlying that the strongest winter and annual
485 rainfall in the southern Central Mediterranean occurred during the 10.1-6.6 ka interval, followed by a
486 decrease toward the Late Holocene.

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488 **4.3.4. Relationships with large-scale climatic changes**

489 The Mediterranean climate results from complex interactions between low and mid- latitude
490 atmospheric circulations and local features such as orography or land sea-contrast (Brayshaw et al.,
491 2011). The winter Mediterranean hydroclimate depends on the Mediterranean storm activity (and
492 associated precipitation) which is strongly influenced by the position and strength of the mid-latitude

493 North Atlantic (NA) storm track (Brayshaw et al., 2010), while summer dryness is tightly linked to the
494 summer extension and strength of the Hadley cell and associated subtropical subsidence (Lionello et
495 al., 2006). Holocene changes of both NA storm track and tropical convection in response to insolation
496 forcing are therefore critical factors to explain the long-term climatic evolution in Central
497 Mediterranean and seasonal precipitation changes.

498 The drier conditions persisting into the Early Holocene up to 10.1 ka are also detected in the
499 SW and NE Mediterranean with a West to East gradient (Dormoy et al., 2009). The level of annual
500 moisture availability always remained higher in the SW probably due to the stronger Atlantic
501 influence, allowing Mediterranean plants to expand during the first millennia of the Holocene
502 (Fletcher and Sánchez Goñi, 2008; Combourieu Nebout et al., 2009). Such drier conditions are
503 potentially due to the boreal summer insolation maximum of the Early Holocene (Tzedakis, 2007)
504 resulting in deficiency in effective moisture availability due to strong evaporation (Renssen and
505 Isarin, 2001). However, limited winter rainfall due to reduced influence of westerly flow-induced
506 cyclones on the Mediterranean region is also probably involved (Kotthoff et al., 2008). The gradual
507 shift from the GS-1 dry-cold situation to the Holocene warm-humid conditions might also be the
508 result of the progressive reorganisation of the global atmospheric circulation associated with the
509 retreat of northern hemisphere ice sheets. The LIS which disappeared around 6.8 ka (Carlson et al.,
510 2008) was large enough to impact on the atmospheric circulation (Montero-Serrano et al., 2010). The
511 Fennoscandian Ice Sheet (FIS) which persisted up to 9 ka could have influenced the response of the
512 European hydroclimate to insolation forcing in maintaining the NA storm track in a relatively
513 southern position (Magny et al., 2003). However, the NA storm track remained narrow enough not to
514 bring high humidity in the Mediterranean.

515
516 The period comprised between 10.1 and 6.6 ka encompasses to the Saproel 1 (Mercone et
517 al., 2000). Increased precipitation in southern Central Mediterranean is coeval with maximum
518 monsoon activity as shown by tropical African paleohydrological records (Lézine et al., 2011).
519 However, a number of evidences clearly show that the African monsoon never reached the
520 Mediterranean region (Tzedakis, 2007). Our data thus confirm these earlier results since the
521 estimated increase in annual precipitation remains limited on average to 700 mm/an in our southern
522 region and mainly due to a change in winter/fall precipitation (Fig.5). In contrast, monsoonal
523 intensification during summer could have played a predominant role on the Mediterranean summer
524 dry season. Limited summer rainfall in southern Central Mediterranean has been attributed to the
525 development and persistence of subtropical anticyclones as a result of the reinforcement of the
526 Hadley cell related to the African monsoon (Tinner et al., 2009; Gaetani et al., 2011). According to
527 (Harrison et al., 1992), the summer northern position and higher strength of the North Atlantic

528 Subtropical High in response to high summer insolation could have blocked the eastward convection
529 and even generate an offshore flow in southern Europe. This pattern would explain summer dryness
530 over the whole Central Mediterranean region and the northern progression of summer dryness as
531 observed in pollen and lake level records (Magny et al., 2007a; Finsinger et al., 2010; Peyron et al.,
532 2011).

533 The winter rainfall maximum from 10.1 to 6.6 ka, observed in the southern Central
534 Mediterranean, appears to be a broad feature of the climate in the Western, Central and North-
535 eastern Mediterranean regions (Fletcher and Sánchez Goñi, 2008; Kotthoff et al., 2008; Combourieu
536 Nebout et al., 2009; Dormoy et al., 2009; Jalut et al., 2009; Pérez-Obiol et al., 2011; Peyron et al.,
537 2011). The spatial extent and relative simultaneity of maximum winter suggests common
538 mechanisms related to remote controlling factors. Simulations using global and regional models
539 HadSM3 and HadRM3, suggest that increased winter precipitation during the Early Holocene would
540 be linked to shifting NA storm track in response to orbital forcing (Brayshaw et al., 2010; Brayshaw et
541 al., 2011). The Early-Mid Holocene orbital configuration imposes a weaker winter latitudinal
542 insolation gradient that would drive the northern tropical Hadley cell to be narrower and in a more
543 southern position in winter. Consequently, the NA storm-track would weaken and shift to the south,
544 thus enhancing Mediterranean cyclogenesis and winter precipitation (Brayshaw et al., 2010).

545 A contrasting hypothesis involving the Arctic Oscillation/North Atlantic situation (AO /NAO)
546 has been proposed by Davis and Brewer (2009). According to these authors, increase in moisture
547 availability in the Mediterranean region would be caused by lower evaporation due to cooler
548 conditions (in contrast with our data from the Central Mediterranean). Cool and dry southern
549 conditions would be explained by a more dominant positive AO (Davis and Brewer, 2009) and more
550 northern NA storm track. Bonfils et al. (2004) showed that NAO does not support the winter
551 temperatures and precipitation distribution during the mid-Holocene as shown by European
552 paleodata. They suggest instead an insolation-induced southern entry of the NA storm track in the
553 European mid-latitudes than during the Late Holocene.

554 The Early Holocene deglaciation of the LIS may have also contributed to increase winter
555 precipitation in the Mediterranean region. The rapid retreat and increased melting of the LIS notably
556 occurred during the wettest phase of the Mediterranean Holocene climate, between 10 and 6.8 ka
557 (Carlson et al., 2008). The above mentioned modelling experiments performed by Brayshaw et al.
558 (2010) do not show a significant impact of the LIS on the NA storm track. Increased cyclogenesis is
559 simulated only in the western North Atlantic due to the increased temperature gradient resulting
560 from the LIS cooling effect. Another modelling experiment with the fully coupled AOGCM ModelE-R
561 has been performed to simulate the 9ka-climate with a more realistic LIS topography and
562 appropriated freshwater routing (Carlson et al., 2008). ModelE-R outputs showed a stronger impact

563 of the LIS on the ocean and atmospheric circulation over the North Atlantic region. Suppression of
564 the Labrador Sea Water (LSW) formation, substantial reduction of the AMOC (by 15%), and SST
565 cooling in the NW Atlantic and temperature decrease in NE North America are simulated in
566 agreement with paleodata (Carlson et al., 2008). If effective AMOC reduction is debatable (Renssen
567 et al., 2005; Hoogakker et al., 2011), oceanic circulation reorganisation occurred when LIS retreat had
568 completed. At ~ 6.5 ka, deep water convection in the Nordic Seas reduced and LSW formation
569 reactivated (Hoogakker et al., 2011). According to the ModelE-R experiment, the impact of the LIS
570 melting on atmospheric and North Atlantic circulation induced a southern and stronger NA storm
571 track providing slightly more winter precipitation to the mid and southern Europe (Carlson et al.,
572 2008 (SI: Fig. 4)).

573 We propose that during the Early Holocene, insolation and ice sheets, in particular the LIS
574 decay, modulated winter precipitation in the Western to Central Mediterranean regions by altering
575 oceanic and atmospheric circulation patterns of the North Atlantic, and strengthening the
576 Mediterranean cyclogenesis. After the final demise of the LIS, increase in winter insolation and
577 latitudinal insolation gradient probably contributed to the long-term decrease in winter precipitation
578 and seasonality in the Western and Central Mediterranean by gradually shifting northward the NA
579 storm track, which thus affected less frequently the Mediterranean region.

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581 **4.4. Impact of the Holocene centennial-scale climatic changes on the temperature and** 582 **hydroclimate in the southern Central Mediterranean**

583 The pollen record from core MD047-2497CQ displays a series of eight vegetation centennial-
584 scale changes superimposed to the Holocene long-term evolution (11.5-11 ka, 10.3-9.9 ka, 9.5-9.1 ka,
585 8.5-7.9 ka, 7.3-6.7 ka, 6.4-6 ka, and ~ 5.5 ka and 4.5 ka).

586 During the Early Holocene dry phase, two centennial-scale dry and cold events at ~ 11.2 ka
587 and 10.1 ka are clearly marked by increases of semi-desert plants with lower tree and shrubs
588 percentages. During the first event, a large amplitude SST cooling is synchronous to vegetation
589 changes, while no SST change is detected during the second one. Further, three cooling events of the
590 SST record at ~ 9.3 ka, 8.2 ka, 7.0 ka punctuated the wettest Holocene phase (10.1-6.6ka) with the
591 last one at ~ 6.2 ka, at the onset of the transition toward drier Mid- to Late Holocene. All these
592 events are coeval with a change in herbaceous composition as shown by the strong reductions of
593 Cyperaceae, most of the time replaced by Asteraceae (*Taraxacum*). Strikingly, they do not coincide
594 with trees and shrubs reductions as we would have expected (Fig.7). For instance, the Mediterranean
595 forest percentages remain stable during the ~ 9.3 ka and 6.2 ka events while they decline at the end
596 of the 7.0 ka event. Different SST and Mediterranean forest variations have also been reported from
597 the Alboran Sea record MD95-2043 (Fletcher et al., in press) underlying the possible asynchronism

598 between Mediterranean SST millennial-scale cooling and tree development. A seasonal bias towards
599 summer for SST derived from alkenones and winter for the Mediterranean forest is proposed to
600 explain this divergent pattern (Fletcher et al., in press). However, alkenone-SSTs from core MD04-
601 2797CQ during the interval 10.1-6.6ka do not parallel those found in core MD95-2043, suggesting
602 that they might be controlled by different processes. We will see hereafter that centennial SST
603 coolings from the Siculo-Tunisian Strait are associated to drier conditions, as revealed by the
604 herbaceous layer changes.

605 Centennial-scale variations in sedge percentages may have different causes. They could
606 result from variations in the extent of coastal humid areas due to minor sea level changes. Such
607 centennial-scale sea level changes during the Early Holocene may be produced by deglacial pulses,
608 sea-water temperature changes through thermal water expansion or AMOC variations (Levermann et
609 al., 2005; Lombard et al., 2005; Tornqvist and Hijma, 2012). Flooding can also favour sedge
610 development in estuary zones and pollen export to the ocean (Bernhardt et al., 2012). However, high
611 sedge percentages do not coincide with increased periods of flooding as recorded in the fluvial
612 floodplains of northern Tunisia (Zielhofer and Faust, 2008). Regarding the whole pollen assemblage,
613 the most straightforward interpretation is restricted humid areas due to drier conditions. The
614 concomitant *Taraxacum* increase may witness low moisture availability although relatively weak
615 because semi-desert plants did not expand. In modern pollen assemblages, the highest *Taraxacum*
616 percentages are found in dry environments of Greece and Morocco, where mean annual
617 temperatures span from 12 and 17°C and precipitation from 0 to 250 mm (Leroy, 1997). Therefore, a
618 weak decrease in precipitation and slight cooling as shown by SSTs remaining above 16°C would
619 explain why oak populations were not affected by abrupt climate changes during that period. In
620 contrast, over the last 6 kyrs, our pollen record shows two strong abrupt forest reductions around
621 5.5 ka and 4.5 ka likely resulting from enhanced dryness, even though no SST decreases are recorded.

622 All the events detected before 6.6 ka parallel the widely documented abrupt coolings
623 punctuating the Early Holocene: the Preboreal Oscillation (PBO) at ~11.2 ka (Björck, 1996), the Boreal
624 Oscillation at ~10.1 ka (Björck et al., 2001; Boch et al., 2009), the 9.3 ka event (Rasmussen et al.,
625 2007), the “8.2-event” (Alley et al., 1997) and a final event at ~7.4 ka (Bond et al., 2001) which likely
626 corresponds to the drier episode detected at ~7.0 ka bearing in mind the chronological uncertainties
627 (Fig. 7). Our pollen record confirms a decrease in precipitation in the Mediterranean region during
628 the Early Holocene cooling episodes (Combourieu Nebout et al., 2009; Dormoy et al., 2009; Pross et
629 al., 2009; Fletcher et al., 2010). In addition, the centennial drier events in our record match the
630 higher lake level episodes in the European mid-latitudes (Magny and Bégeot, 2004) (Fig. 7). This
631 finding supports the Early Holocene contrasting pattern of the hydrological changes between the
632 mid-latitudes and southern Europe (Magny and Bégeot, 2004; Fletcher et al., in press). These cooling

633 events have been associated with deglacial freshwater outbursts in the North Atlantic from proglacial
634 lakes such as Lake Agassiz (Teller et al., 2002), Lake Labrador-Ungava (Jansson and Kleman, 2004),
635 Lake Superior (Yu et al., 2010) or Baltic ice Lake (Nesje et al., 2004), and with solar activity minima
636 (Bond et al., 2001; Magny and Bégeot, 2004). Early Holocene meltwater pulses and solar forcing have
637 been shown to alter the AMOC and modify the atmospheric circulation in the mid-latitudes but also
638 in the tropics (Bond et al., 2001; Magny and Bégeot, 2004; Marchitto et al., 2010). A southern entry
639 point of the Atlantic jet and associated cyclones in Europe and a stronger zonal flow due to enhanced
640 temperature gradient between low and high latitudes, are suggested from the European hydrological
641 reconstructions (Magny et al., 2003; Magny and Bégeot, 2004; Magny et al., 2007b).

642 Abrupt forest reductions around 5.5 ka and 4.5 ka may be attributed to major widespread
643 cooling events dating to 5.6-5 ka and ~4.2 ka, respectively (Magny and Haas, 2004; Drysdale et al.,
644 2006; Magny et al., 2009). This coincidence might be fortuitous because the age control and time
645 resolution of our sequence is low over the last 5.8 kyrs. Nevertheless, drier conditions in the
646 southern Mediterranean at ~5.5 and 4.2 ka are also suggested by lake-level decreases (Magny et al.,
647 2007a; Magny et al., 2011b). An episode of arboreal biomass reduction at ~4.2 ka is recorded in the
648 Central Mediterranean. It has been associated with climate change despite the possible human
649 impact at the beginning of the Bronze Age (Magri and Parra, 2002; Sadori et al., 2011). Both events at
650 ~5.5 and 4.2 ka are associated with higher lake levels at mid-latitudes in Europe and in Central Italy in
651 response to North Atlantic cooling and decrease in solar activity (Magny, 2004; Magny et al.,
652 2007a)(Fig. 7). However, hydrological changes at 4.2 and 5.5 ka are complex (Magny et al., 2006;
653 Magny et al., 2009; Magny et al., 2011b). For instance, the short-lived dry event in Central Italy
654 probably coeval to the “4.2 ka event” occurred during a cooler and wetter interval between 4.5 and
655 3.8 ka (Magny et al., 2009). In contrast, forest reduction episodes in Western Mediterranean at ~5.4-
656 4.5 ka and 3.7-2.9 ka do not match those events (Fig. 7). They have been associated to dry conditions
657 in the mid-latitudes and enhanced wind strength in the northern latitudes (Fletcher et al., in press).
658 However, it is not clear if these forest reduction episodes actually correspond to warm or cold events
659 in the North Atlantic because of a different timing and periodicity of the centennial scale variations
660 recorded in the Alboran Sea pollen record and in the North Atlantic IRD record (Fletcher, pers. com.).

661 Therefore, given the limitations of the chronology of our core, forest reductions
662 reconstructed from our record may be linked either (1) to higher lake level in the mid-European
663 latitude and lower lake level at Lake Preola, or (2) to the forest reduction in the Western
664 Mediterranean. Answer to this question is crucial to determine the factors controlling the centennial-
665 scale variability during the Mid- to Late Holocene and their impact on the Mediterranean region.
666 From 6 ka onwards, dry episodes in the southern Central Mediterranean could involve (1) wetter
667 conditions above 40°N due to a southern and stronger Atlantic jet like during the Early Holocene

668 cooling events (Magny et al., 2004) or (2) drier conditions in the Mediterranean and Mid European
669 latitudes due to an intensification and northward shift of the NA storm track and subsequent limited
670 penetration of storms in the Mediterranean region (Fletcher et al., in press).

671 Apart from this, our record shows that changes on land are of weaker amplitude during the
672 Early Holocene, in particular during Sapropel 1. Weaker changes in Mediterranean forest in the SW
673 Mediterranean also characterize the Early Holocene as compared to the Mid- to Late Holocene as
674 shown by the MD95-2043 pollen sequence (Fig. 7). The notable weak expression of the 8.2 event in
675 the Central Mediterranean has already been emphasized from the Accesa or Preola sequences
676 (Magny et al., 2011b; Peyron et al., 2011). This event is even missing in Moroccan and Tunisian
677 alluvial sequences (Zielhofer et al., 2008). This finding emphasizes that impact of cooling events in
678 the Early Holocene was relatively weak on the Western to Central Mediterranean hydroclimate while
679 it appears to have been stronger during the Mid to Late Holocene. Baseline climate states, such as
680 insolation and ice sheet volume, exert a strong control on the atmospheric circulation in the North
681 Atlantic and thus likely modulate the impact of abrupt climate changes on the Mediterranean region.

682 During the humid Early Holocene, insolation and melting LIS resulted in a mean southern NA
683 storm track, with cyclones becoming stronger and more zonal during cooling events, enhancing
684 precipitation at mid-latitudes and restricting them in southern Europe. Despite increased dryness,
685 paleodata suggest that precipitations remained relatively sustained in the Western to Central
686 Mediterranean. NA storms were still affecting southern Europe during the Early Holocene cooling
687 events, though to a lesser extent than during the bracketing warm phases. In contrast, during the
688 Mid to Late Holocene, in the absence of ice sheet forcing, the NA storm tracks were likely more
689 northern, thus less frequently affecting southern Europe. Therefore, centennial-scale variations may
690 have induced stronger aridity in the Mediterranean region.

691

692 **5. Conclusions**

693 The deglacial and Holocene climate of the southern Central Mediterranean are examined at
694 orbital and millennial to centennial time-scale using a marine pollen record (MD04-2797CQ). Our
695 reconstruction provides an integrated picture of the regional vegetation changes and main regional
696 climate features over the last 18,000 years:

697 - Prevailing dry conditions during the deglaciation are indicated by the dominance of semi-
698 desert plants. Conditions remained arid even during the GI-1, restricting the expansion of the trees
699 and shrubs despite climatic amelioration.

700 - Our land-sea correlation shows with no chronological ambiguity that even though
701 temperatures increased at the GS-1 – Holocene transition in the southern Central Mediterranean,

702 tree and shrub expansion was strongly limited by moisture deficiency persisting into the Early
703 Holocene up to 10.1 ka.

704 - Temperate trees and shrubs with heaths as oak forest understorey or heath maquis
705 expanded between 10.1 and 6.6 ka while Mediterranean plants only developed from 6.6 ka to the
706 Late Holocene. Climate in southern Central Mediterranean was wetter during the Early Holocene
707 interval corresponding to Sapropel 1 and became drier during the Mid- to Late Holocene, mainly
708 because of a change in winter precipitation while summers remained dry.

709 Our findings suggest, in agreement with modelling experiments, that the Holocene long-term
710 winter precipitation change in the Mediterranean region depends on the mid-latitude atmospheric
711 circulation which is basically controlled by insolation and ice sheet volume. In particular, increased
712 melting of the LIS between 10 and 6.8ka combined with weak winter insolation played a major role in
713 the Mediterranean winter precipitation maxima during the Early Holocene. This feature is likely
714 related to more frequent NA storm tracks in a mean southern position, feeding the Mediterranean
715 cyclogenesis, than during the Mid to Late Holocene. Summer dryness along the Holocene suggests a
716 northern progression that may be related to the development of subtropical subsidence in summer
717 in relation with the Early Holocene monsoonal intensification.

718 Finally, our data provide evidences on the impact of Holocene centennial-scale climate
719 changes on the Central Mediterranean vegetation (11.5-11 ka, 10.3-9.9 ka, 9.5-9.1 ka, 8.5-7.9 ka, 7.3-
720 6.7 ka, 6.4-6 ka, and at around 5.5 ka and 4.5 ka). The lack of response of the Mediterranean forest
721 but detection of herbaceous composition changes during the wet Early Holocene indicate muted
722 changes in precipitation contrasting with episodes of enhanced aridity during the Mid- to Late
723 Holocene. We suggest that the impact of the Holocene centennial-scale variability results from
724 baseline climate states, insolation and ice sheet volume, shaping the response of the mid-latitudes
725 atmospheric circulation.

726

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734

735 **Figure captions**

736 Figure 1: Map of Central Mediterranean with location of the studied core MD04-2797CQ and the
737 pollen sequences mentioned in the text from the southern Central Mediterranean (DF: Dar Fatma
738 (Ben Tiba and Reille, 1982), MH: Majen Ben M'Hida (Stambouli-Essassi et al., 2007), GB: Gorgo Basso
739 (Tinner et al., 2009), LP: Lake Pergusa (Sadori and Narcisi, 2001), BG: Biviera di Gela (Noti et al., 2009)

740

741 Figure 2: Age-depth model for core MD04-2797CQ.

742

743 Figure 3: Percentages of major pollen taxa versus depth, Core MD04-2797CQ, Siculo-Tunisian Strait.
744 Boundaries of pollen zones are based on visual inspection of data and application of clustering
745 analysis (see text for details).

746

747 Figure 4: Vegetation and climatic changes on land over the last 18,000 years. From the bottom to the
748 top: tree and shrubs (grey) and total (white) pollen concentrations, pollen percentages of selected
749 pollen taxa and ecological groups, SST derived from alkenones (Sicre et al., this volume) from core
750 MD04-2797CQ, $\delta^{18}\text{O}$ from NGRIP ice core on GGC05 time-scale (converted to BP)(Rasmussen et al.,
751 2006; Vinther et al., 2006). Long-term Holocene vegetation evolution has been detected by applying
752 a smoothing cubic spline method to selected pollen taxa data. Pollen zones (bottom) and
753 stratigraphical framework (top) are indicated. Note that we will name the Younger Dryas as
754 Greenland Stadial 1 (GS-1) following the INITIMATE recommendations (Lowe et al., 2008) but we will
755 keep using HS 1 in the text (as defined by Sánchez Goñi and Harrison (2010)) because the isotopic
756 changes during GS-2 are difficult to reconcile from one Greenland ice core to another (Rasmussen et
757 al., 2008) and the correlation between GS-2a and HS 1 has to be confirmed.

758

759 Figure 5: Box-Whisker plots of estimated annual, summer and winter temperatures (TANN, MTWA,
760 MTCO) and precipitation (Pann, Psum, Pwin) derived from the pollen data of core MD04-2797CQ
761 (Peyron et al., this volume-a).

762

763 Figure 6: Comparison of the pollen data from core MD04-2797CQ with other records from southern
764 Central Mediterranean and with insolation and ice sheet forcing. **Forcings:** a) Early Holocene ice
765 sheet deglaciation: disappearance of the Fennoscandian Ice Sheet (FIS) at 9.5 kyr cal BP (Lundqvist
766 and Saarnisto, 1995)and enhanced Laurentide Ice Sheet (LIS) retreat between 10 and 6.8 kyr cal BP
767 with only small ice caps remaining up to 5.5 ka (Carlson et al., 2008); b) Summer (JJA) and winter
768 (DJF) latitudinal insolation gradient calculated from insolation values at 60° and 30°N (Berger and
769 Loutre, 1991); **Tunisia:** c) Peat pollen sequence Majen Ben M'Hida: digitalized data *versus depth*

770 (Stambouli-Essassi et al., 2007), asterisks: original dated levels (65 cm: 985 ± 30 yr BP; 100 cm: $2500 \pm$
771 50 yr BP; 205 cm: 4894 ± 40 yr BP; 250 cm: 8190 ± 50 yr BP; 350 cm: 9480 ± 100 yr BP) and calibrated
772 ages using Calib 6.11 (903 (873-959); 2578 (2365-2740); 5629 (5585-5716); 9144 (9014-9287); 10782
773 (10504-11143) yr cal BP); d) $\delta^{13}\text{C}$ speleothem record from La Mine cave (Genty et al., 2006); e)
774 Shaded bar representing the increased fluvial dynamics since 5 ka with recurrent floods associated to
775 drier conditions, based on alluvial sequences from arid to semi-arid Central and North Tunisian
776 floodplains(Zielhofer and Faust, 2008); **Siculo-Tunisia Strait**: f) MD04-2797CQ pollen data : Total
777 trees and shrubs percentages including Ericaceae with smoothing cubic spline (bold green line) and
778 ratio of Mediterranean plants to Ericaceae; **Sicily**: g)Total trees and shrubs percentages from Gorgo
779 Basso pollen sequence (Tinner et al., 2009); h) Arboreal pollen percentages and arboreal pollen grain
780 concentrations from Lake Pergusa sequence (Sadori and Narcisi, 2001; Sadori et al., 2011); i) Relative
781 changes in lake level (LL) from Lake Preola (Magny et al., 2011b); j) Interval corresponding to
782 Sapropel 1 after Mercone et al. (2000). The colored vertical bars represent the successive intervals as
783 described in the text: yellow for the dry interval at the beginning of the Early Holocene, blue for the
784 wet Early Holocene phase, grey for the Mid-Holocene transitional interval and orange for the Mid- to
785 Late Holocene phase of increasing dryness.

786

787 Figure 7: Comparison of the centennial-scale variability recorded in core MD04-2797CQ with changes
788 detected in the Western Mediterranean, middle latitudes of Europe and North Atlantic. From left to
789 right: Mediterranean forest pollen percentages from core MD95-2043 (Fletcher and Sánchez Goñi,
790 2008): 3-points moving average (bold green line) with forest reduction episodes indicated by the grey
791 bars as in Fletcher et al. (in press); MD04-2797CQ pollen percentages and alkenone-SSTs (bold lines:
792 cubic smoothing splines with a degree of freedom adjusted to highlight the centennial-scale changes)
793 with abrupt vegetation and SST changes as described in the text; Mid-European lake level (LL)
794 changes (Magny, 2004); Stacked Hematite Stained Grains (HSG) from North Atlantic records (Bond et
795 al., 2001).

796

Depth (cm)	¹⁴ C age (year BP)	Depth (cm)	Mean ¹⁴ C age from paired duplicates (year BP)	1 σ error (year)	Reservoir corrected ¹⁴ C age (year BP)	Lower cal age (year)	Upper cal age (year)	Calibrated ages (year cal BP)	1 σ error (year)
0	1105 ± 20			20	705	661	674	668	9
80	5425 ± 30	80	5493	95	5093	5725	5929	5827	144
80	5560 ± 25								
160	6760 ± 30	160	6700	85	6300	7156	7325	7241	120
160.5	6640 ± 30								
199	7580 ± 30	199	7523	81	7123	7917	8017	7967	71
199.5	7465 ± 30								
239.5	8170 ± 40	240	8113	81	7713	8419	8557	8488	98
240	8055 ± 35								
329.5	8810 ± 35	330	8888	110	8488	9398	9556	9477	112
330	8965 ± 30								
410	10885 ± 40	410	10863	32	10463	12458	12528	12493	49
410.5	10840 ± 40								
469.5	12850 ± 50	470	12728	173	12168	13786	14246	14016	325
470	12605 ± 40								
510.5	14000 ± 60	510	13900	141	13100	15495	16373	15934	621
511	13800 ± 100								
610	15590 ± 50	610	15590	50	15190	18463	18986	18725	370
610.5	16160 ± 60								
700	18180 ± 60	700	17660	70	17260	20287	20541	20414	180
700	17660 ± 70								
939.5	23300 ± 100	940	23415	163	23015	27645	28113	27879	331
940.5	23530 ± 120								
1029.5	26100 ± 130	1030	26095	7	25695	30379	30600	30490	156
1030	26090 ± 150								

Table 1: Radiocarbon ages, corrected ages for age reservoir according to Siani et al. (2001) and calibrated ages using INTCAL 09 (Reimer et al., 2009)

Pollen Zones	Interval (cm)	Pollen zone age (calyr BP)	Description of pollen zones	Vegetation changes	Inferred climatic changes
PZ-7	70-24	5,300–2,200	Highest values of evergreen <i>Quercus</i> and from 40 cm, <i>Olea</i> and low percentages of Cyperaceae and Ericaceae. Peaks in deciduous <i>Quercus</i> of 20-25%	Open Mediterranean forest with increased Mediterranean plants contribution	Decreased humidity
PZ-6	120-75	6,600–5,300	Decreasing values of Cyperaceae and Ericaceae Increasing percentages of Mediterranean taxa (evergreen <i>Quercus</i> , <i>Olea</i> and <i>Pistacia</i>) and deciduous <i>Quercus</i>	Expansion of an open Mediterranean forest with Asteraceae-Poaceae steppe	Transition to a period with less rainfalls
PZ-5	344-125	10,100–6,600	Increase in AP percentages although AP representation is weak, mainly deciduous <i>Quercus</i> and occurrences of Mediterranean taxa (evergreen <i>Quercus</i> , <i>Olea</i> and <i>Pistacia</i>) High <i>Taraxacum</i> -type percentages. Highest values of Ericaceae and Cyperaceae. Recurrent oscillations in Cyperaceae percentages.	Open oak forest with heath underbrush or maquis and Asteraceae-Poaceae-Cyperaceae steppe	Increased humidity Centennial oscillations
PZ-4	404-350	12,300 – 10,100	Fall of semi-desert taxa percentages and rise of <i>Taraxacum</i> -type. Decreasing trend in semi-desert percentages (25 to 10 %, mainly <i>Ephedra fragilis</i> -type and Chenopodiaceae). Decrease in <i>Pinus</i> and virtual absence of <i>Cedrus</i> . Minor increase in deciduous <i>Quercus</i> . Gradual increase of Ericaceae and Cyperaceae values	Steppe with Asteraceae, Poaceae and some semi-desert plants and development of oak woodlands with heath underbrush or maquis	Warmer conditions and gradual increase in humidity, although limited precipitations
PZ-3	430-408	13,100– 12,300	Rise in <i>Artemisia</i> , <i>Ephedra distachya</i> -type, <i>Cedrus</i> and Cupressaceae values	Steppe with ragweed, chenopods, <i>Ephedra</i> and scarce cypress lowland Cedar-pine conifer forest upland	Cold and dry
PZ-2	480-440	14,700– 13,100	Decrease in <i>Artemisia</i> , <i>Ephedra distachya</i> -type although semi-desert plant remains dominant, <i>Cedrus</i> and Cupressaceae values Small but significant increase in AP taxa, in particular Mediterranean tree taxa (evergreen <i>Quercus</i> and	Steppe with semi-desert plants and scarce Mediterranean woodlands Pine conifer forest upland	Warmer and less dry (limitation in water availability, decreasing trend).

			<i>Olea</i>), and Ericaceae Increase in <i>Taraxacum</i>		
PZ-1	590-500	18,200–14,700	Dominance of NAP taxa, in particular semi-desert taxa (<i>Artemisia</i> , Chenopodiaceae, <i>Ephedra distachya</i> -type, <i>Ephedra fragilis</i> -type) Continuous presence of deciduous <i>Quercus</i> High <i>Pinus</i> percentages and peaks of <i>Cedrus</i> and Cupressaceae	Steppe with ragweed, chenopods, Ephedra and scarce cypress lowland Cedar-pine conifer forest upland	Cold and dry

Table 2: Description of the MD04-2797CQ pollen record.

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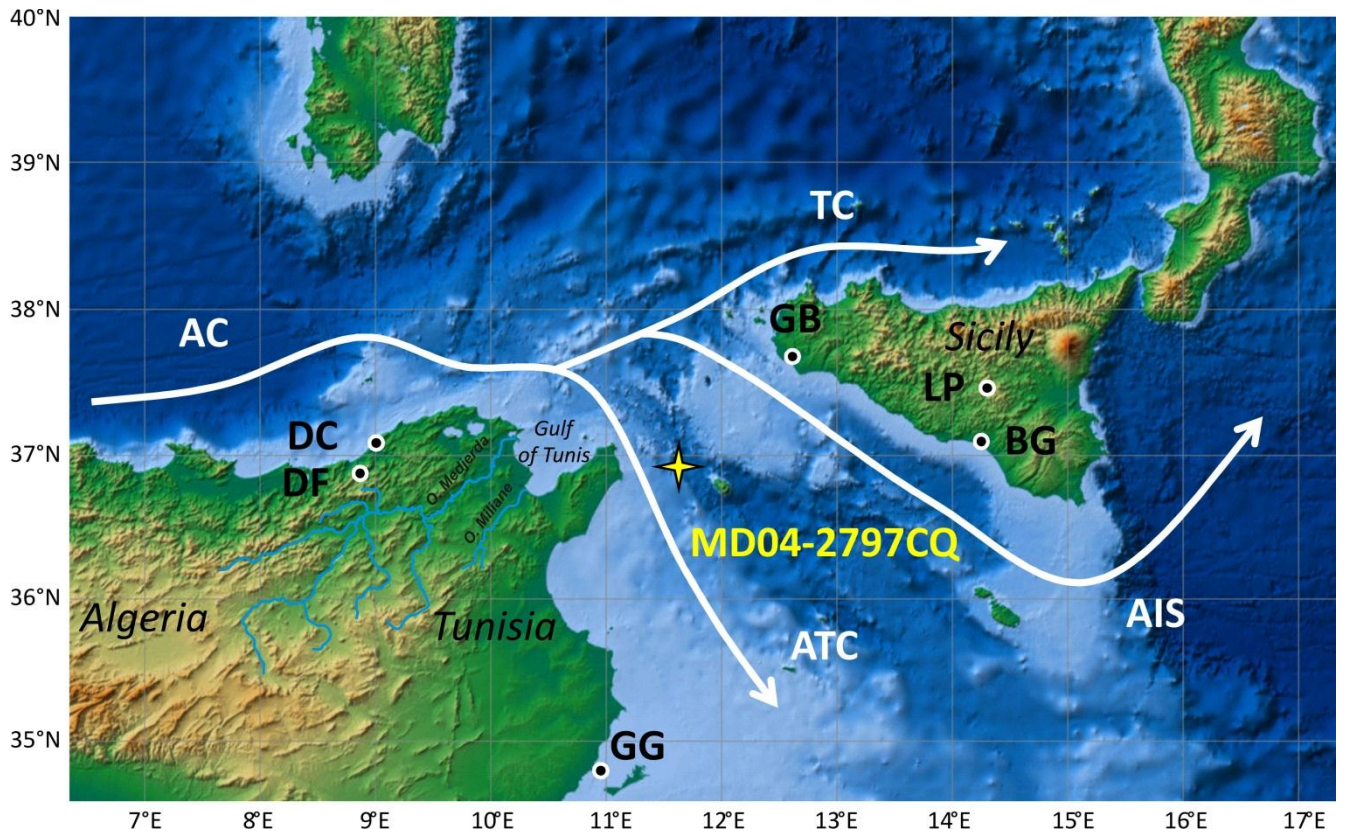


Figure 1

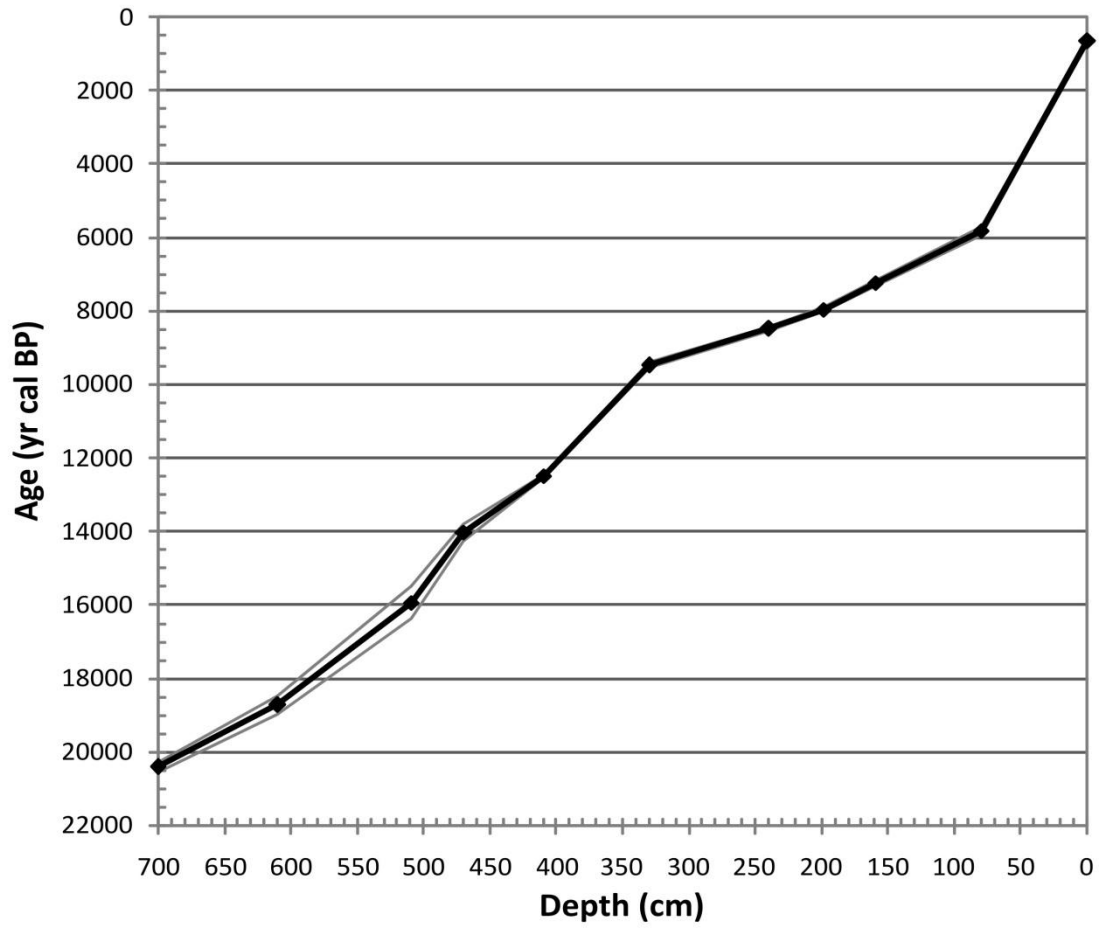


Figure 2

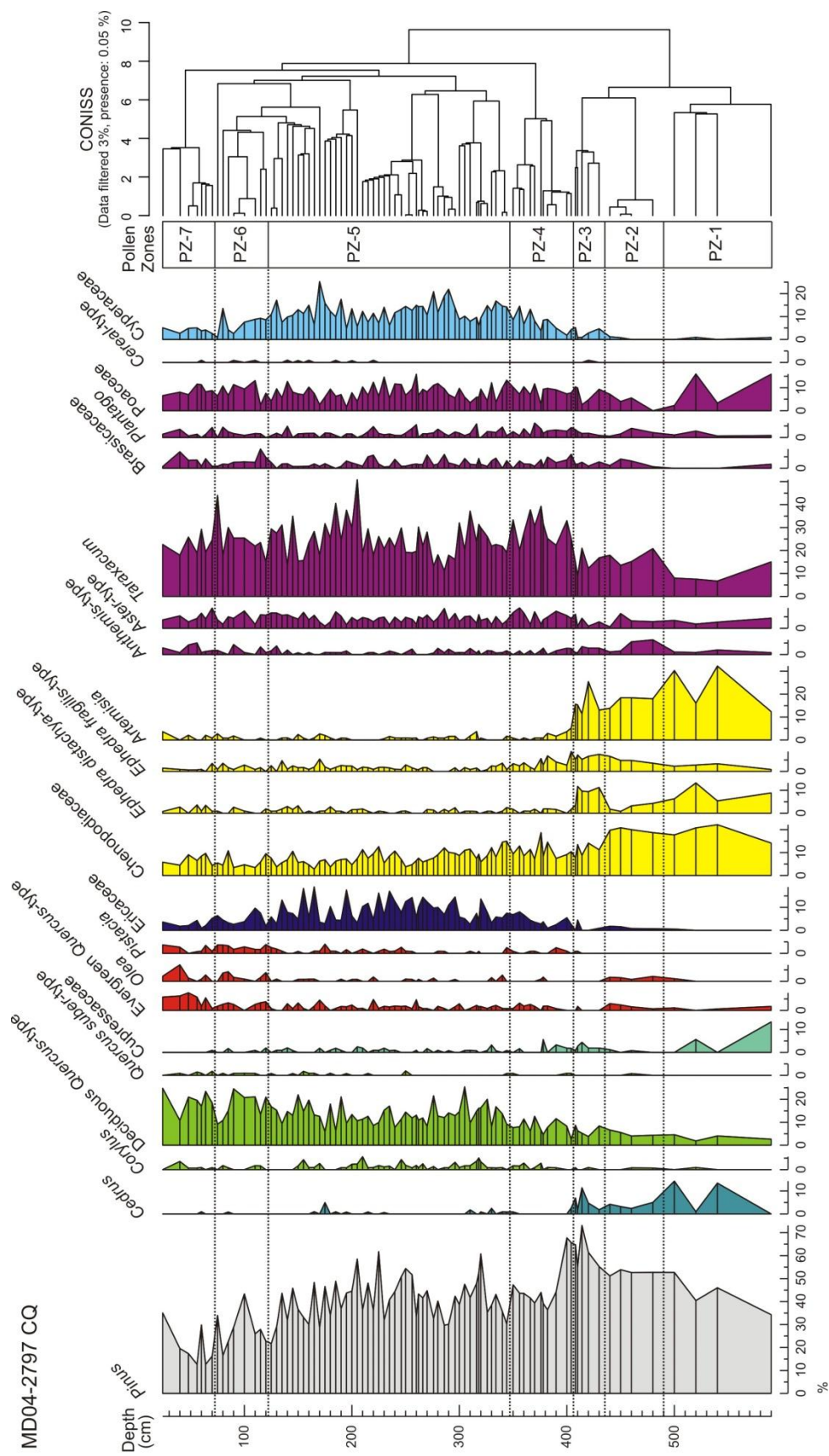


Figure 3

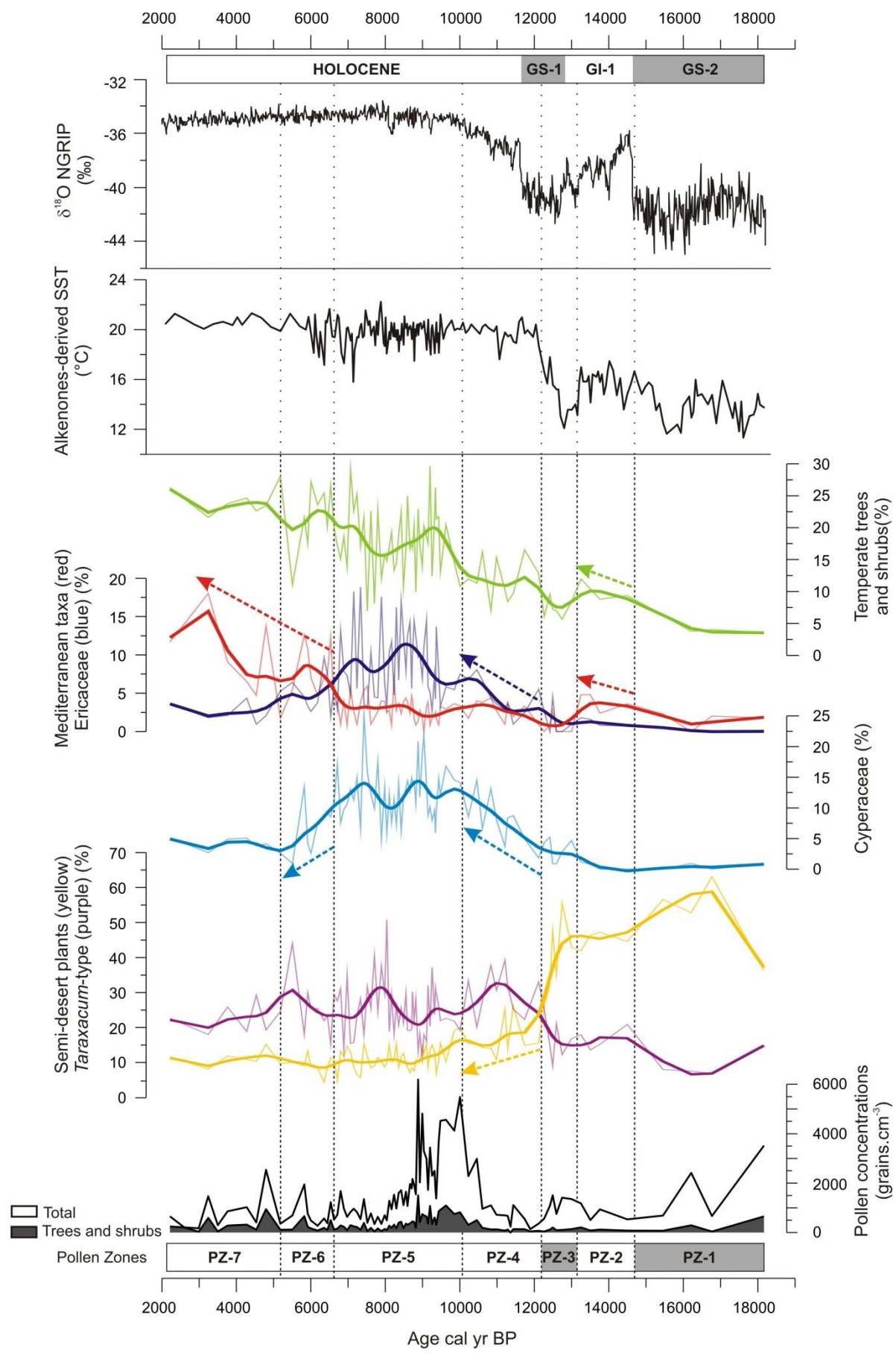


Figure 4

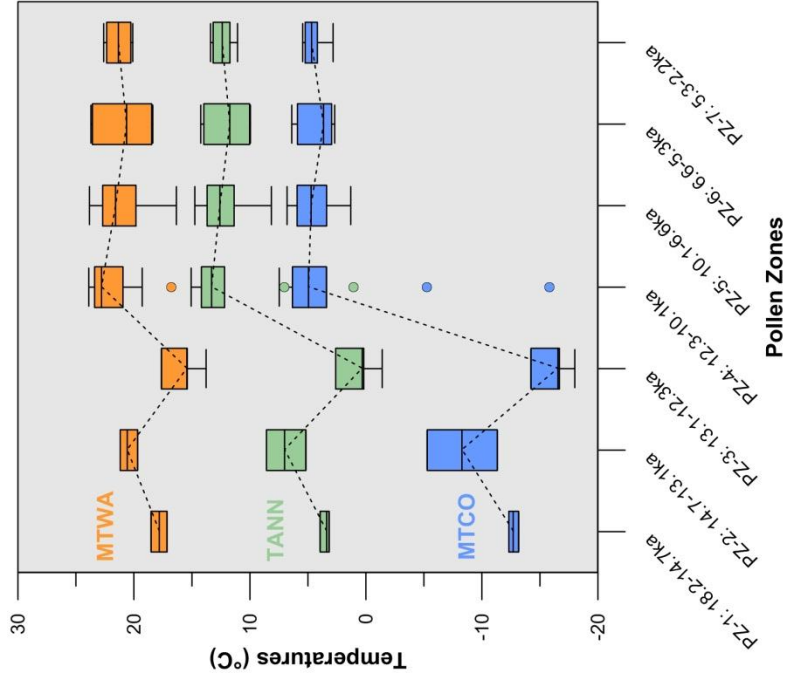
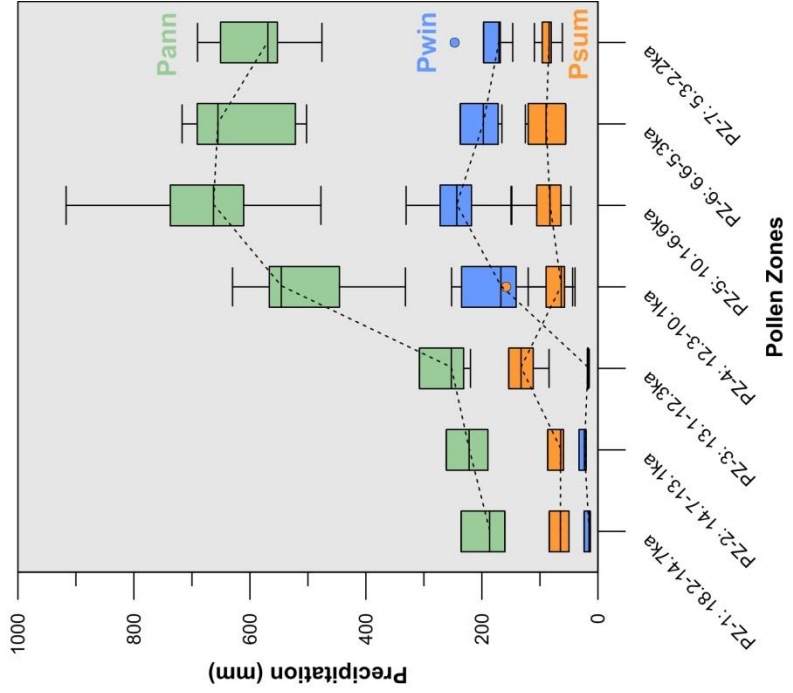


Figure 5

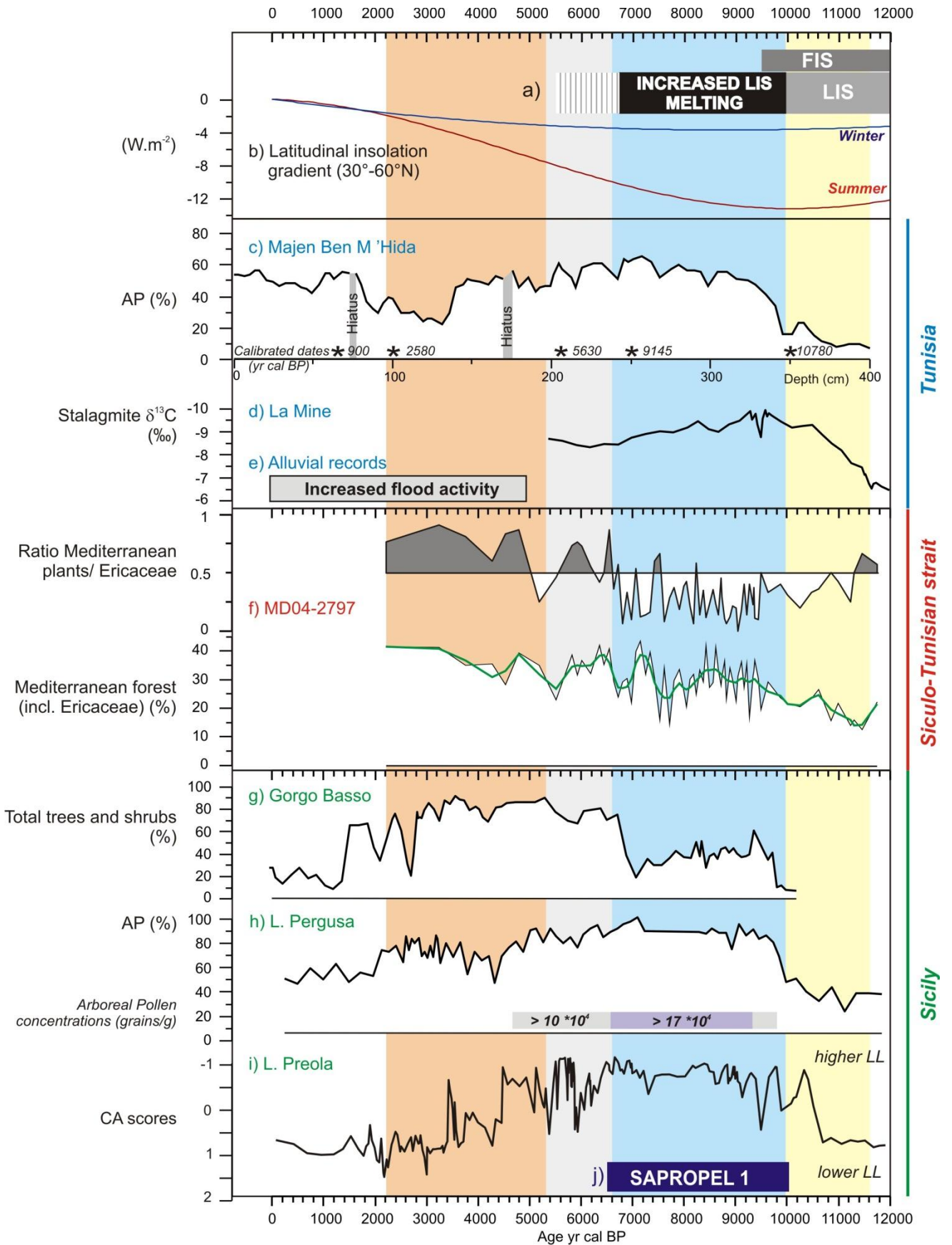


Figure 6

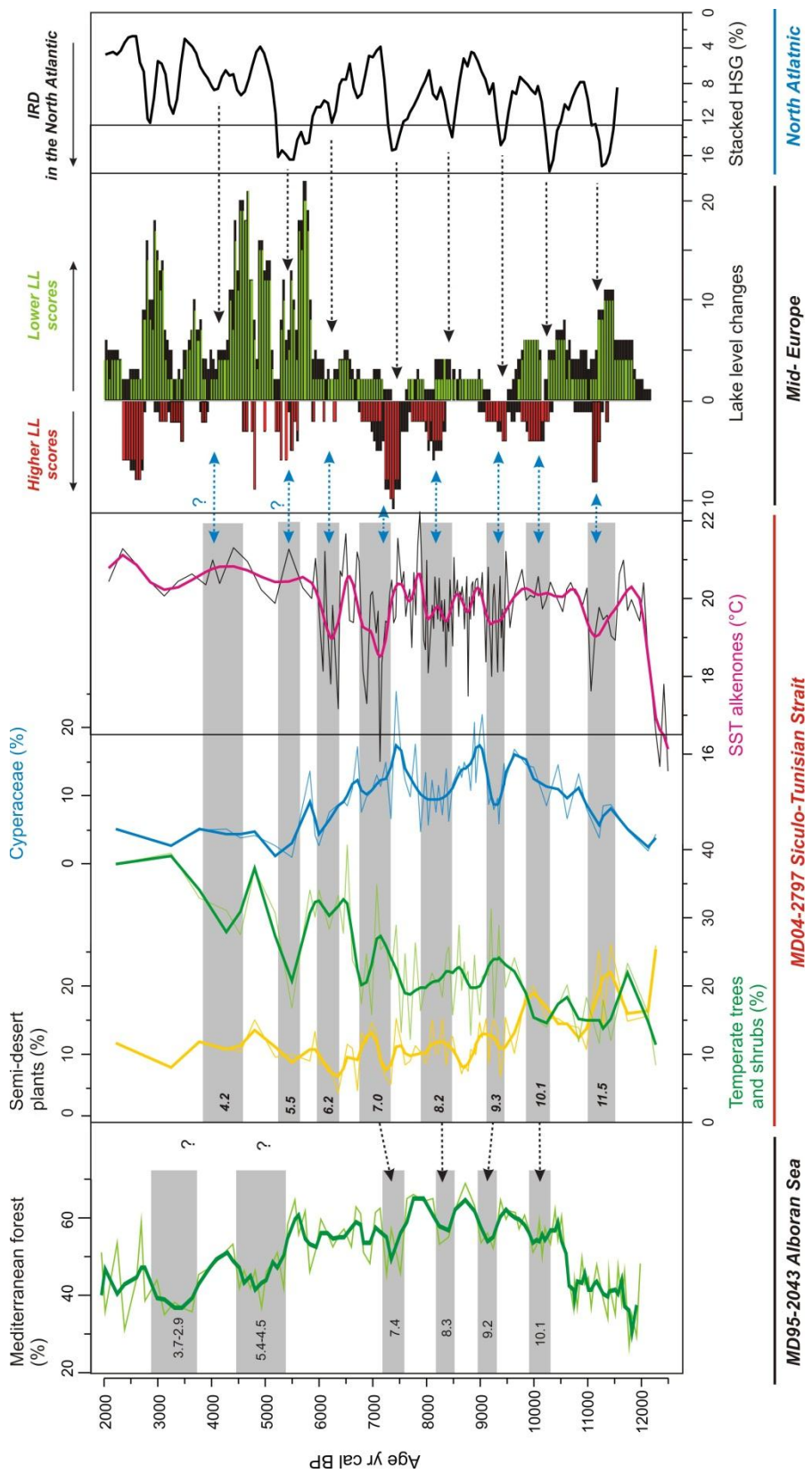


Figure 7