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Climate variability of the last ~2700 years in the Southern Adriatic Sea:

Coccolithophore evidences

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18 **Abstract**

19 New information on paleoenvironmental conditions over the past ~2700 years in the Central
20 Mediterranean Sea have been acquired through the high-resolution study of calcareous
21 nanofossils preserved in the sediment core SW104ND14Q recovered in the Southern
22 Adriatic Sea (SAS) at 1013 m water depth. The surface water properties at this open SAS site
23 are sensitive to atmospheric forcing (acting both at local and regional scale) and the North
24 Ionian Sea driven inflowing waters. Our data show a relationship between reworked coccolith
25 abundances, flood frequency across the Southern Alps and the North Atlantic Oscillation
26 (NAO) confirming their value as indicator of runoff/precipitation. Changes in the abundance
27 of the opportunistic (r-strategist) species *Emiliania huxleyi* and deep dweller taxa
28 *Florisphaera profunda* were used to reconstruct the upper water column stratification and
29 associated changes in coccolithophorid productivity. The negative correlation between
30 reworked coccoliths and the N-Ratio ($r=-0.44$; $p=6^{-7}$) suggest that fresh water induced
31 stratification is a controlling factor of the SAS coccolithophorid production. High
32 coccolithophorid productivity levels occurred during dry periods and/or time intervals of
33 inflowing salty and nutrient-rich Levantine Intermediate Waters (LIW) favouring convection
34 while lower levels took place during high freshwater discharge, mainly during the Little Ice
35 Age (LIA) and two centennial scale intervals of weakest NAO around 200 BCE and 500 CE.

36

37 **Keywords**

38 Coccolithophores; reworked coccoliths; coccolithophorid primary productivity; South
39 Adriatic Sea; central Mediterranean; last millennia.

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42 **Introduction**

43 Coccolithophores (calcareous nanoplankton) and their fossil remains (calcareous
44 nannofossils) are valuable source of information for paleoclimatic studies (Baumann et al.,
45 2005). Coccolithophores are single cell calcareous algae whose ecology and vital functions
46 are driven by environmental parameters within the ocean euphotic zone (e.g., temperature,
47 salinity, sunlight, and nutrient supply). Therefore, abundances of selected taxa have been used
48 to reconstruct variations of physical and environmental parameters and their relation with
49 climate change and human activity. Their skeletons composed of tiny calcareous platelets
50 (coccoliths) are highly abundant in marine sediments making them ideal fossils to produce
51 high-resolution time series (Baumann et al., 2005). These microorganisms are usually
52 considered to prefer warm, stratified, oligotrophic waters of low and middle latitude regions
53 (e.g., Honjo and Okada, 1974; Ziveri et al., 2004). However, local oceanic features such as
54 coastal currents, gyres, eddies, upwelling, and river runoff are known to regionally affect their
55 productivity (Guerreiro et al., 2013). In addition, reworked coccoliths (i.e., the nannofossils
56 which have been removed from their original sedimentary layer and redeposited in a younger
57 layer) can provide information on sediment transport (Bonomo et al., 2014; Ferreira et al.,
58 2008; Ferreira and Cachão, 2005) and used to reconstruct regional scale runoff and/or
59 precipitation changes (Bonomo et al., 2016a; Incarbona et al., 2010; Sprovieri et al., 2006).

60 Understanding the trends and variability of the Mediterranean climate at local and regional
61 scales has been subject of intense research. (Bonomo et al., 2016a; Cacho et al., 1999; Frigola
62 et al., 2007; Martrat et al., 2004; Pérez-Folgado et al., 2004; Rodrigo-Gámiz et al., 2011;
63 Rohling et al., 2002, 2015; Sbaffi et al., 2001; Sierro et al., 2005; Sprovieri et al., 2003, 2006;
64 Triantaphyllou et al., 2009, 2016a). Shelf sediments of the Adriatic Sea (AS) provide ideal
65 natural archives for high-resolution paleoclimatic investigations because of expanded
66 Holocene sedimentary sequences and possible use of recurrent tephras for geochronological

67 control (Jalali et al., 2018; Lowe et al., 2007; Marchini et al., 2014; Matthews et al., 2015;
68 Siani et al., 2013). Terrestrial and marine paleoclimate proxy data (e.g., calcareous plankton,
69 lipid biomarkers, palynomorphs, stable isotopes, lake levels, and speleothems) have shown
70 the occurrence of abrupt climate changes during the Holocene (warmer/colder and
71 drier/wetter periods) at decadal, centennial to millennial time scales in the Mediterranean
72 basin (e.g., Bini et al., 2019; Cisneros et al., 2016; Di Bella et al., 2014; Gogou et al., 2016;
73 Goudeau et al., 2015; Grauel et al., 2013; Jalali et al., 2016, 2018; Kouli et al., 2012; Lirer et
74 al., 2013, 2014; Margaritelli et al., 2016, 2018; Piva et al., 2008; Sicre et al., 2016; Skampa et
75 al., 2019; Triantaphyllou et al., 2009, 2010, 2016b).

76 Many studies have focussed on (late) Holocene climate variability and its impact on the
77 environment and human activity in the SAS (Caroli and Caldara, 2007; Combourieu-Nebout
78 et al., 2013; Di Rita and Magri, 2009; Giunta et al., 2003; Grauel and Bernasconi, 2010; Jalali
79 et al., 2018; Leider et al., 2010; Oldfield et al., 2003; Piva et al., 2008; Sangiorgi et al., 2003;
80 Siani et al., 2013; Sicre et al., 2016). The recent study of Jalali et al. (2018) in the SAS
81 highlighted the links between the centennial scale variability of SSTs and local climatic and
82 oceanographic features, and notably the role of the Bimodal Oscillating System (BiOS) of the
83 Ionian Sea and North Atlantic Oscillation (NAO). Although there has been a substantial
84 number of publications on the investigated area, very few studies have explored calcareous
85 nannofossils as a proxy of past climate and environmental changes (e.g., Giunta et al., 2003;
86 Narciso et al., 2012; Sangiorgi et al., 2003). Narciso et al. (2012) studied a gravity core close
87 to our site between 13000 and 5500 BP, thus focused on the Greenland Stadial 1/Younger
88 Dryas, Pre-Boreal, and Sapropel 1 equivalent periods. Giunta et al. (2003) and Sangiorgi et al.
89 (2003) reported data from 18000 to 2300 yrs BP at a more southern site, documenting the
90 distribution of calcareous nannofossils during the Sapropel S1. As far as living
91 coccolithophores are concerned, the only study carried out in the SAS is that of Balestra et al.
92 (2008) describing assemblages in the water column and surface coastal sediments of the Gulf

93 of Manfredonia (SAS). Other very recent data were restricted to the Mid and North Adriatic
94 Sea (e.g., Cerino et al., 2017; Godrijan et al., 2018; Skejić et al., 2018, and references there
95 in) or are part of phytoplankton biomass and productivity assessments aiming at providing
96 rough estimates of coccolithophore distribution in open SAS (e.g., Fonda Umani, 1996;
97 Ljubimir et al., 2017, and references therein).

98 The aim of this work is to evaluate the reliability of Coccolithophores/calcareous nannofossils
99 as a proxy of environment and climate variability over the last three millennia in the Central
100 Mediterranean. For this purpose, we carried out a high-resolution study from a deep-sea
101 gravity core recovered in the SAS and from a second shallow coastal gravity core (C5
102 Composite) from the Tyrrhenian Sea (Gulf of Gaeta) obtained within the framework of the
103 NEXTDATA Project (<http://www.nextdataproject.it>). Our data evidence major changes in
104 nutricline depth as well as variations of river runoff and precipitation. We explore the cause of
105 the observed changes by comparing our results to alkenone derived Sea Surface Temperatures
106 (SSTs) and terrestrial inputs derived from higher plant biomarkers (Jalali et al., 2018). We
107 also use other indicators of past precipitation changes in the Mediterranean basin: i.e. the
108 flood activity in the Southern Alps (Wirth et al., 2013), the reworked coccolith record from a
109 Southern Tyrrhenian sea core (Bonomo et al., 2016b), the XRF record from lake sediments of
110 the Iberian Peninsula (Moreno et al., 2012) and the reconstruction of the forested fraction of
111 usable land in Central and Western Europe (Kaplan et al., 2009).

112

113 **Oceanographic setting of the study area**

114 The AS is a semi-enclosed basin located between the Italian Peninsula and the Balkans,
115 connected to the Mediterranean Sea through the Strait of Otranto (Fig.1a). The North Adriatic
116 (NA) is primarily influenced by the southeast Europe climate, while the SAS experiences
117 more arid conditions typical of Mediterranean and Northern Africa climates (Ilijanić et al.,
118 2014). The general surface circulation of the AS is cyclonic (Fig.1b,c) (Sellschopp and

119 Álvarez, 2003) and consists of a northward current flowing along the eastern Adriatic coast
120 (i.e., the Eastern Adriatic Current, EAC) balanced by southward current flowing along the
121 western coast (i.e., the Western Adriatic Current, WAC). The intermediate layer mainly
122 present in the southern and mid AS is occupied by the Levantine Intermediate Water (LIW)
123 (Artegiani et al., 1997). The deep circulation is characterized by the Adriatic Deep Water
124 (ADW) a dense water mass formed by the mixing of the Northern Adriatic Dense Water
125 (NADW) and Southern Adriatic Dense Water (SADW) (Manca et al., 2002).

126 The SAS is a sub-basin (South Adriatic Pit, SAP, 1260 m max depth) characterized by a
127 quasi-permanent cyclonic circulation, i.e. the South Adriatic Gyre (SAG; Gačić et al., 1997)
128 (Fig. 1b,c). The physical and chemical properties of SAS surface waters depend on the
129 characteristics of inflowing waters into the basin, the strength of SAG, as well as wind stress
130 and river discharges. Inflowing waters consist mainly of WAC and NADW from the North,
131 the LIW and occasionally Modified Atlantic Water (MAW) from the South. The WAC is
132 strongly influenced by river runoff mostly from the Po River, making it fresher and nutrient
133 rich. The inflow of LIW and MAW depends on the variability of the North Ionian Gyre (NIG)
134 (Fig. 1b,c). According to the BiOS (Bimodal Oscillating System) model, the NIG circulation
135 may either be cyclonic or anticyclonic (Civitarese et al., 2010; Gacic et al., 2010). This
136 mechanism is sustained by internal processes driven by the density of the ADW outflowing
137 the Otranto Strait. When the circulation in the NIG is cyclonic, saltier and warmer LIW enters
138 the SAS promoting deep convection and the formation of a denser ADW. In an anticyclonic
139 NIG mode, fresher and colder MAW enters the SAS leading to the production of lower
140 density ADW. However, some studies invoke the role of more complex driving mechanisms
141 involving the whole Ionian Sea circulation and not just its northern sector (Reale et al., 2016;
142 Simoncelli et al., 2016; Theocharis et al., 2014). The intensity of the SAG depends on local
143 wind intensity and properties of advected waters from the Ionian Sea (Shabrang et al., 2016).
144 Shabrang et al. (2016) reported a significant negative correlation between the NAO index and

145 local wind intensity. However, they did not find unequivocal relationship between the NAO
146 and SAG variability because of additional effects of the advection of the Ionian waters,
147 suggesting that the BIOS mode does not depend on NAO. Nevertheless, Pinardi et al. (2015)
148 reported a sustained BIOS anticyclonic circulation during the period of positive NAO in
149 1987-1996.

150 The surface waters of the SAS are more oligotrophic than those of the NA (e.g., Civitarese et
151 al., 1998). The influence of Po River and secondary Apennines rivers flowing into the western
152 AS on the nutrient budget of the SAS seems rather weak and limited to a narrow coastal
153 current flowing over the Italian shelf (Faganeli et al., 1989). The nutrient supply to the SAS
154 occurs mainly via the inflow of LIW lying at about 300 m in the Adriatic Sea (Gačić et al.,
155 2002). Nevertheless, according to Civitarese et al. (2010) larger amounts of nutrients are
156 advected by the MAW during periods of anticyclonic BIOS.

157 **Methods**

158 *Core SW104-ND14Q*

159 Core SW104-ND14Q (17°37'3.612''E; 41°17'2.4''N) was recovered at 1013m water depth in
160 the SAS (Fig. 1). The sedimentary sequence was retrieved with a SW104 gravity corer
161 system, which preserves the water-sediment interface and allowed the recovery of 116 cm of
162 undisturbed and uncompressed homogeneous brown-grey hemipelagic sediments. The
163 magnetic susceptibility measured on board with a Bartington Instrument M2 revealed three
164 tephra layers (Fig. 2). The age model used here is from Jalali et al. (2018) and has been
165 constructed combining radionuclides ages (^{210}Pb activity-depth profile and ^{137}Cs activity) for
166 the last ca. 150 years and the additional dates derived from the correlation of three tephra
167 layers with well-dated volcanic events onland [Pompeii eruption (79 CE); Pollena eruption
168 (472 CE); 1631 CE] (see Jalali et al. 2018 for details on tephrostratigraphy). Linear
169 interpolation between the tie-points has been used to construct the age-depth profile from the
170 top down to the base core documenting a mean Sed. Rate of 0.04 cm/y (Fig. 2). Based on the

171 age model, core SW104-ND14Q ranges from 700 BCE to 2003 CE and has a mean temporal
172 resolution of ~26 yrs.

173 *Core C5 Composite*

174 To investigate the reliability of the reworked coccoliths as a regional proxy of precipitation,
175 we also used the central Tyrrhenian Sea shallow sequence C5 Composite (C5Comp) (Fig.1).
176 The location of this site in front of Volturno River mouth makes it particularly suitable for
177 reconstructing runoff variability and for comparing coastal and open sea sites (Bonomo et al.,
178 2016). The core C5Comp is a composite marine sequence consisting of two cores: the
179 SW104-C5 and core C5 (710 cmbsf length) both recovered in the Gulf of Gaeta, at 93 m
180 water depth (see Margaritelli et al., 2016 for details). Calcareous nannofossils of the core
181 SW104-C5 (back to 1630 CE) was already published by Bonomo et al. (2016). In this work
182 we extended their reconstruction back to ~ 400 CE. The chronology used is that of
183 Margaritelli et al. (2016) and has been assembled combining radionuclides ages (^{210}Pb
184 activity-depth profile and ^{137}Cs activity) for the last ca. 150 years, planktonic foraminiferal
185 event, tephrostratigraphy and oxygen stable isotope correlation with other marine sites (for
186 details see Margeritelli et al., 2016). The age-depth profile has been constructed by a linear
187 interpolation between the tie-points showing a progressive decrease in sedimentation rate
188 from the top down to the base core.

189 The analysed time interval of core C5Comp covers the period between ~ 400 and 2013 CE
190 with a mean temporal resolution of ~10 yrs.

191 *Calcareous Nannofossils*

192 116 samples of the SW104-ND14Q core and 108 of the C5Comp were prepared as standard
193 smear slides (Bown, 1998) and analyzed with a transmitted light microscope at x1250
194 magnification. Some samples of SW104-ND14Q core were analysed with a scanning electron
195 microscope (SEM) in order to solve taxonomic identification for smaller placoliths difficult to
196 achieve by light microscope (e.g., *Emiliana huxleyi*). The relative abundance of *in situ*

197 species was estimated only in the SW104-ND14Q core based on the count of at least 600
198 specimens. The abundance of reworked nannofossils was estimated in the SW104-ND14Q
199 and C5Comp as the number of reworked specimens encountered during the count of the *in*
200 *situ* coccoliths. All abundances are expressed in percentages. SW104-ND14Q coccolith
201 species abundances were also used to calculate the N-ratio as defined by Flores et al. (2000)
202 to assess the nutricline depth fluctuations. The N-ratio is based on the absolute abundances of
203 the main surface r-strategist species (in our record *E. huxleyi* and small placoliths) over that of
204 *F. profunda* (lower photic zone taxon). High values of the N-ratio indicate shallow
205 nutricline/thermocline (relatively high surface coccolithophorid productivity) while low
206 values indicate deep nutricline/thermocline (relatively low surface coccolithophorid
207 productivity). As small placoliths, we counted the placoliths not confidently recognizable as
208 *E. huxleyi* and *Reticulofenestra* spp.

209 Finally, the reworked coccoliths (RC) group includes taxa from different stratigraphic
210 intervals (Mesozoic, early Cenozoic) and Cenozoic long-range taxa showing poor
211 preservation (etching and/or overgrowth). Raw data are shown in supplementary material.

212

213 *Ecology of selected taxa*

214 *E. huxleyi* tolerates a wide range of ecological conditions and is therefore abundant in nearly
215 all oceanic environments (Schwab et al., 2012). This species is considered an opportunistic (r-
216 strategist) taxon capable to quickly respond to nutrient availability in both eutrophic and
217 oligotrophic areas (e.g., Balestra et al., 2008; Broerse et al., 2000; Dimiza et al., 2008, 2015;
218 Haidar and Thierstein, 2001). *E. huxleyi* is generally more abundant in temperate (cold) mixed
219 surface waters (e.g., Hagino et al., 2000; Malinverno et al., 2003), but may also be found in
220 stable regimes in terms of vertical mixing with relatively high nutrient availability (Andruleit
221 et al., 2005). Ausín et al. (2015) further postulated that *E. huxleyi* (size >4 µm) can find
222 optimal conditions for its development in cold water that are also low-salinity.

223 The lower photic zone species *F. profunda* has a more constrained habitat and has thus been
224 widely used to monitor past changes in nutricline-depth and induced changes in surface
225 productivity (Beaufort, 1997). The abundance of *F. profunda* increases with respect to other
226 coccolithophores when the nutricline is deep and overlaid by a nutrient-depleted upper photic
227 layer (Balestra et al., 2008; Bown et al., 2009; Dimiza et al., 2015; Incarbona et al., 2008,
228 2010). These conditions generally reveal stable, stratified, oligotrophic surface waters during
229 summer months (Baumann et al., 2005; Malinverno et al., 2009) that can be disrupted under
230 increased wind stress and / or upwelling and divergence circulation (Bown et al., 2009).
231 Hernández-Almeida et al. (2019), using *F. profunda* relative abundance vs MODIS (Moderate
232 Resolution Imaging Spectroradiometer) chlorophyll- α , show a pronounced temperature
233 sensitivity of *F. profunda* and no correlation with surface net primary production at latitudes
234 higher than 30°N–30°S, such as Mediterranean area. Contrary, Grelaud et al. (2012) showed a
235 strong anticorrelation ($R = -0.76$) between *F. profunda* % and chlorophyll- α in the Aegean
236 Sea (eastern Mediterranean Sea).

237 *Biomarker analyses*

238 Sea surface temperature and TERR-alkane reconstructions along the SW104-ND14Q core
239 have been published by Jalali et al (2018). The method used for biomarker analyses have been
240 described by (Sicre et al., 2002). Fatty alcohol biomarker data were used to calculate the C₂₆
241 fatty alcohol / C₂₉ n-alkane + C₂₆ fatty alcohol ratio ($C_{26OH}/(C_{26OH}+C_{29})$). This ratio was
242 determined along the core to infer information on water oxygenation as proposed by Cacho et
243 al. (2000). High values of this ratio presumably correspond to low ventilation and *vice versa*.

244

245 **Results**

246 The coccolithophore assemblages in the SW104-ND14Q core are generally well preserved
247 and abundant. *E. huxleyi* dominates the assemblages with an average abundance of ~80%. *F.*
248 *profunda* is also well represented with an average abundance of ~10%. Other taxa are largely

249 subordinated with percentages ranging between ~1- 3% (e.g. *Syracosphaera*, *Rhabdosphaera*
250 and *Calciosolenia*) and no significant variations (not shown). Reworked specimens are always
251 present and are found in higher amounts in the upper part of the core. *E. huxleyi*, *F. profunda*,
252 RC, and the N-ratio data shown in Figure 3 are used for the discussion.

253 *E. huxleyi* abundance range from 65 to 90 % (Fig. 3a). Its downcore distribution pattern can
254 be divided into two major intervals. A first one that includes the late Iron Age (IA) and the
255 almost entire Roman Period (RP; between ~700 BCE and ~400 CE) with abundance above
256 80%. This period is followed by a decline to lower values (< 65%) between ~400 and ~800
257 CE, i.e. from the late RP throughout the Dark Age (DA). *E. huxleyi* returns to moderately
258 higher abundances (65 - 75%) at the late DA and during the Medieval Climate Anomaly
259 (MCA) (800 to 1100 CE). Then, values remain approximately at these levels but with
260 superimposed short-lived oscillations especially during the upper LIA. The distribution of *F.*
261 *profunda* (Fig. 3b) reveals three main intervals: the first one runs from the bottom of the core
262 till ~400 CE and is characterized by fluctuating values between 7 and 9 %. Over the second
263 interval, from 400 to 1200 CE, the taxon abundances increase almost continuously, except for
264 two time spans of strong decrease centred at 600 and 900 CE. From 1200 CE, *F. profunda*
265 declines till 1550 CE and rises again to Present day values. As shown in Fig. 3d, the N-ratio
266 shows similar trends as *E. huxleyi*, but with more pronounced fluctuations especially in the
267 upper half of the core. During the first 1200 years (700 BCE - 400 CE) the N-ratio value is >
268 0.9. At ~400 CE a sharp drop sets the beginning of a long-term decreasing trend till Present
269 that suggests a progressive reduction of coccolithophorid productivity.

270 RC percentages (%RC) along the core range from ~3 to ~25%, and depict a steady increase
271 from the bottom core to ~800 CE. Then, after a period of lower values around 900 CE and
272 1300 CE, %RC increases up to Present with the highest values (17-25%) during the LIA
273 (~1400 -1800 CE) (Fig. 3c). In the C5Comp core, %RC ranges from ~14 to ~79% with lowest
274 levels found between ~400 and ~1350 CE.

275

276 **Discussion**

277 *Reworked coccoliths and runoff fluctuations*

278 The NAO is one of the dominant atmospheric mode of variability in the North Atlantic sector
279 that has a considerable influence on winter temperature/precipitation in Europe including the
280 Mediterranean region (Hurrell, 1995). In the central Mediterranean, positive NAO conditions
281 result in colder and drier winters than average, while winters are warmer and wetter during
282 negative phases of NAO (Benito et al., 2015; López-Moreno et al., 2011a, 2011b; and
283 references within). Bonomo et al. (2016) were able to evidence a negative correlation between
284 the NAO index of Trouet et al. (2009) and the %RC in the Central Tyrrhenian Sea core
285 SW104-C5 over the last 400 years. Our results show that this relationship could have
286 persisted back to ~700 BCE (Fig. 4). The resemblance between the %RC short and long term
287 trends the flood frequency in Southern Alps (Wirth et al., 2013) and the Southern Tyrrhenian
288 marine record (Gulf of Salerno; Lirer et al. 2013) seems to confirm the link between the %RC
289 and runoff/precipitation in the region on longer time span (Bonomo et al., 2016a; Incarbona et
290 al., 2010; Sprovieri et al., 2006). This finding is supported by the slight negative correlation
291 between the NAO index of Trouet et al. (2009) ($r=-0.4$ $p=5^{-33}$, $n=34$) and Olsen et al. (2012)
292 ($r=-0.2$ $p=0.01$, $n=82$) and the %RC along the SW104 record. Our data agree with the
293 negative correlation between NAO and winter precipitation, for the 1950–2006 period,
294 reconstructed over large areas of Morocco and Tunisia, most of the Iberian Peninsula,
295 southeastern France, Italy, the Balkan Peninsula, and large areas of central and northern
296 Turkey (López-Moreno et al., 2011a). Notwithstanding the age models accuracy of the
297 different cores, the main drier spells recorded in the SAS, in the Central and Southern
298 Tyrrhenian as shown by red dots in Fig. 4 might be considered synchronous as well to the
299 XRF Si fluctuations found in lake sediments of Iberian Peninsula (Moreno et al., 2012). A
300 noteworthy result is the high %RC (RC Acme event) during the late LIA, between ~1600 and

301 ~1850 CE, that coincides with a long standing interval of negative NAO and is consistent
302 with a regional scale humid period already documented in marine and continental sedimentary
303 sequences of the Western and Central Mediterranean (e.g., Barrera-Escoda and Llasat, 2015;
304 Goudeau et al., 2015; Vallefucio et al., 2012; Moreno et al. 2012) .
305 Jalali et al. (2018) highlighted similarities between the TERR–alkane record in SW104-
306 ND14Q and the forested fraction of usable land (FF) in Central and Western Europe (Fig.4 f,
307 i) (Kaplan et al., 2009). Considering that FF fluctuations are indicative of anthropogenic
308 deforestation (Kaplan et al., 2009), they concluded that TERR–alkane at SW104-ND14Q
309 reflects primarily human activity rather than climate fluctuations. Since the RC signal does
310 not match with either the TERR–alkanes or FF index but with the flood activity
311 reconstruction and the $C_{26OH}/(C_{26OH}+C_{29})$ ratio (Fig.4), we suggest that RC reflect
312 precipitation changes that are also seen in other Mediterranean RC records overall supporting
313 the hypothesis that %RC is a reliable index of past runoff/precipitation changes in the region.

314

315 *N-ratio and South Adriatic hydrology*

316 Highest N-ratio values almost all along the RP indicate shallow nutricline (surface productive
317 waters) during this period considered as generally mild (Figs. 3, 5). This is in contrast with
318 the LIA showing deep nutricline (lower surface productivity levels) (Figs. 3, 5), a cold period
319 that one would expect to be favourable to water column mixing and growth of r-strategy taxa
320 *E. huxleyi*. Comparable results has been recorded in the North Aegean Sea during the last
321 1500 years (Gogou et al., 2016; Skampa et al., 2019). In particularly, in the North Aegean Sea
322 Gogou et al.(2016) and Skampa et al. (2019) recorded periodic occurrence of “*E. huxleyi*
323 dominance” intervals indicating strong water column convection coupled with NAO positive
324 shifts , EMT-like events (Incarbona et al., 2016), cool spells, and enhanced continental inputs
325 as well. In contrast, the occurrence of *F. profunda* dominance intervals may be linked to
326 enhanced stratification of the upper water column and warm surface waters, potentially

327 associated with increased lower salinity Black Sea Water intrusion. During the RP, alkenone-
328 derived SSTs show cold oscillations that do not seem to have any relationship with the N-
329 ratio (Fig. 5 a, d). Local atmospheric and hydrological conditions (i.e. properties of inflowing
330 waters into the basin and strength of SAG) play an important role in the stratification of the
331 upper water column and associated changes in productivity (Civitarese et al., 2010; Ljubimir
332 et al., 2017; Vilibić et al., 2012). Several studies in open sea SAS waters have linked high
333 abundances of coccolithophorids with the inflow of saltier Ionian waters (Fonda Umani,
334 1996; Totti et al., 2000). In contrast, Ljubimir et al. (2017) reported higher abundances of
335 coccolithophorids in lower salinity SAG waters during years of anticyclonic mode of the
336 BiOS and their absence during cyclonic BiOS years. However, despite the lack of significant
337 correlation between salinity and total coccolithophore abundances, increased abundance of *E.*
338 *huxleyi* has been often related to the inflow of LIW or eastern Mediterranean surface waters
339 (Malinverno et al., 2003; Skejić et al., 2018). Advection of saltier LIW by promoting deep
340 convection (Gačić et al., 2014) would favour the development of *E. huxleyi* known to rapidly
341 respond to increased nutrient supply to the photic zone (Fig.5 g) (Malinverno et al., 2003).
342 Conversely, reduced inflow of LIW, or enhanced input of less salty waters (mainly the WAC,
343 and occasionally the MAW), and a weak SAG, would lead to higher surface water buoyancy
344 and stratified conditions (Fig.5 f, g). The consequent deepening of the nutricline would thus
345 favour *F. profunda* growth (Fig. 5 f). This conceptual scheme is in agreement with the slight
346 negative correlation ($r = -0.44$; $p = 6^{-7}$) between the N-ratio and %RC values. For instance,
347 higher values of %RC associated with sustained negative NAO during the LIA are coherent
348 with higher precipitation and runoff (Bonomo et al., 2016a; Incarbona et al., 2010; Sprovieri
349 et al., 2006) and the Po River flood record (Camuffo and Enzi, 1996). Rising
350 $C_{26OH}/(C_{26OH}+C_{29})$ ratio to their highest values suggests an abrupt reduction of water
351 oxygenation that is also compatible with stratified conditions caused by the large freshwater

352 discharge during the LIA and lowest N-ratios. Similar observation can be made for two
353 intervals of weaker NAO, i.e. around 200 BCE and around 500 CE.
354 Regarding nutrient supply, our results also support the idea of a limited influence of the Po
355 River (and secondary Apennines rivers) on the nutrient budget of the open SAS surface
356 waters and coccolithophore productivity, as the nutrients are usually rapidly consumed during
357 their transport within the WAC. The same have been observed around the eastern Adriatic
358 coasts (Vilibić et al., 2012). Overall, our findings suggest that fresh water input due to
359 increased precipitation and river runoff impact essentially on buoyancy and subsequent
360 stratification in the SAG.

361 Apart from the LIA and these two major short time intervals that all took place during
362 prolonged negative NAO, other N-ratio fluctuations cannot robustly be attributed to NAO and
363 high river discharge (Fig. 5). Under weaker freshwater forcing, other factors such as the BIOS
364 circulation may have been a more important controlling factor on the SAG dynamics and
365 productivity, but this question will need further investigations to be addressed.

366

367 **Conclusion**

368 This high-resolution study of calcareous nannofossils from the sediment core SW104-ND14Q
369 was used to provide information on paleoceanographic and climatic conditions in the SAS,
370 over the past ~2700 years. Based on the distribution of *E. huxleyi*, *F. profunda*, the N-ratio,
371 and reworked coccoliths we were able to evidence hydrological variability and related
372 coccolithophore production changes in the SAG.

373 One outstanding result is the good correspondence we found between the % reworked
374 coccoliths in the SAS and Tyrrhenian Sea cores and flood activity across the Southern Alps,
375 highlighting the value of %RC as a proxy for reconstructing regional scale precipitation and
376 runoff.

377 We also showed that lowest N-ratio took place during extended weakest NAO phases, i.e.
378 primarily the LIA and two other intervals (200BCE and 500CE), as a result of large fresh
379 water discharge and subsequent stratified surface ocean reducing nutrient supply and
380 production of coccolithophorids in the SAG. Outside these periods of strong negative NAO,
381 whether and to what extent other factors such as the BIOS may have played a role on the
382 hydrology and productivity of the SAG remains an open question.

383

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392

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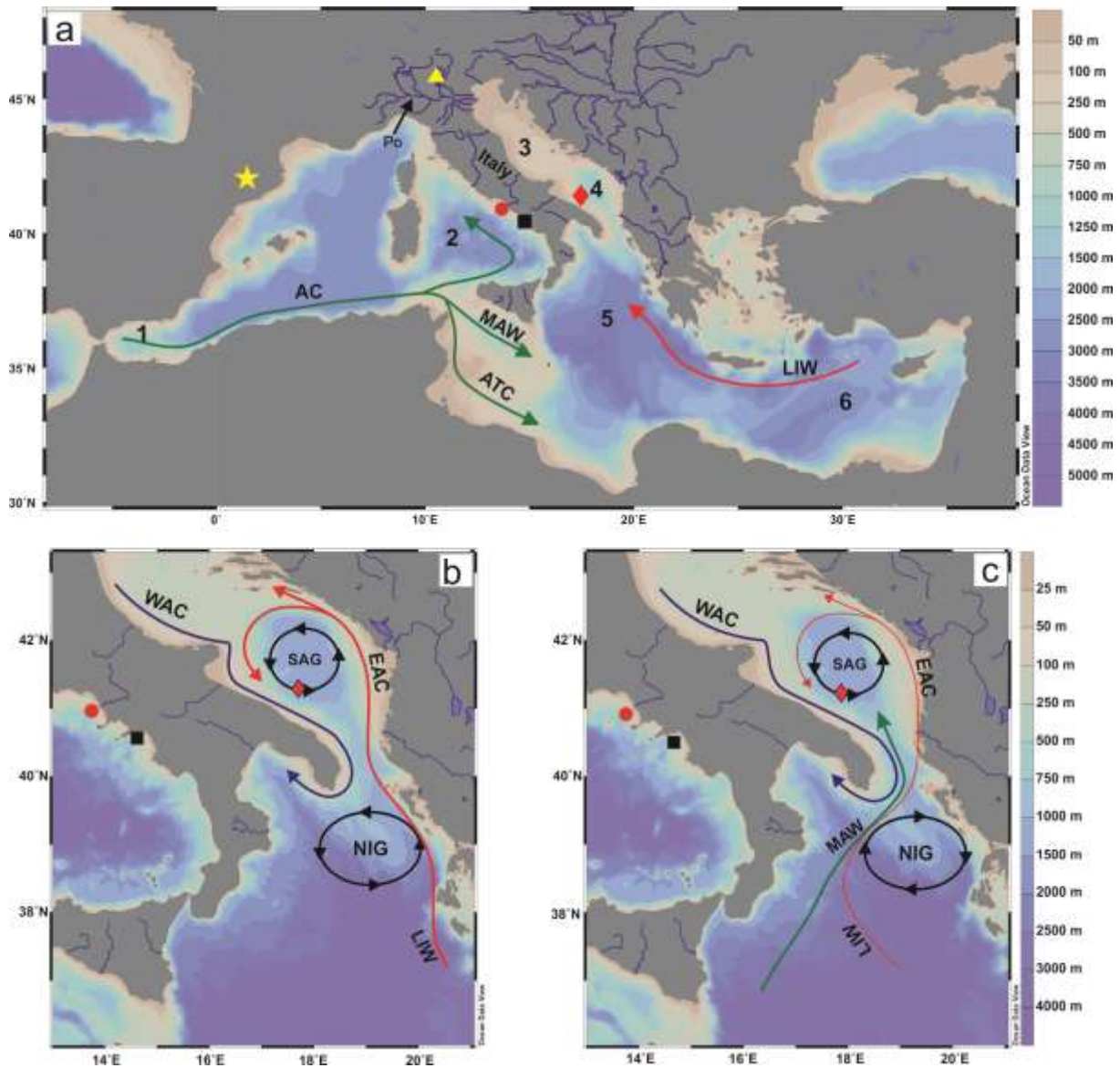
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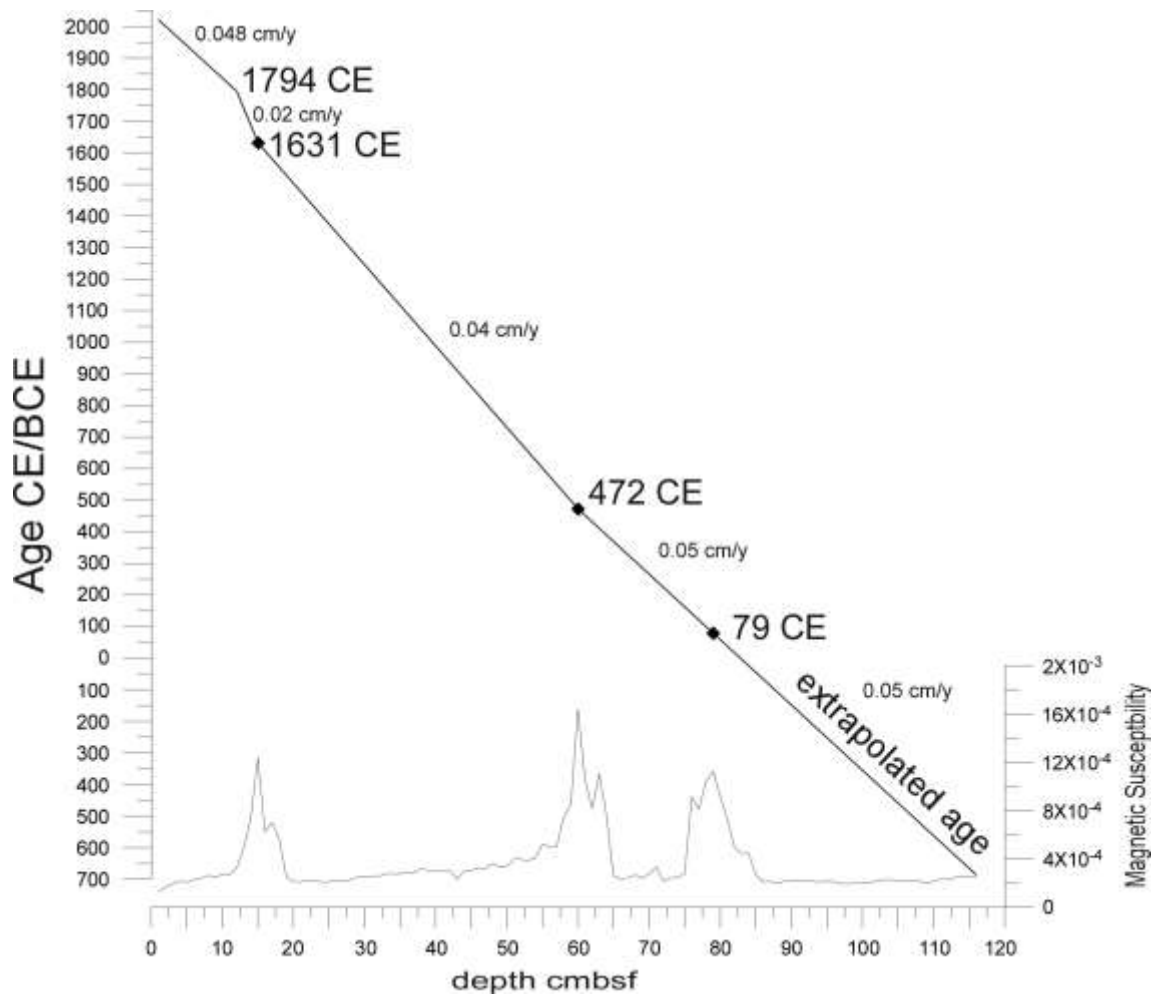
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745 **Figure 1.** Location of the cores SW104-ND14Q (diamond), C5 Composite (dot), C90 (square, Lirer et al., 2013), Basa de la Mora (star, Moreno et al., 2012), and Ledro (triangle, Wirth et al., 2013). (a): bathymetric map of the Mediterranean Basin and main surface (green arrow) and intermediate circulation pattern (red arrow). AC: Algerian Current; MAW: Modified Atlantic Water ; ATC: Atlantic Tunisian Current; LIW: Levantine Intermediate Water. Numbers 1-6: 1-Alboran Sea; 2-Tyrrhenian Sea; 3-Adriatic Sea; 4-South Adriatic Pit, SAP; 5-Ionian Sea; 6 Levantine Sea. The main catchment basins of river flowing into the Adriatic Sea are reported (blue thick lines). (b) and (c): bathymetric map and main circulation pattern of South Adriatic Sea and North Ionian Sea during cyclonic (b) and anticyclonic (c) mode of the BiOS; WAC: Western Adriatic Current; EAC: Eastern Adriatic Water; LIW: Levantine Intermediate Water; MAW: Modified Atlantic Water; SAG: South Adriatic Gyre; NIG: North Ionian Gyre.

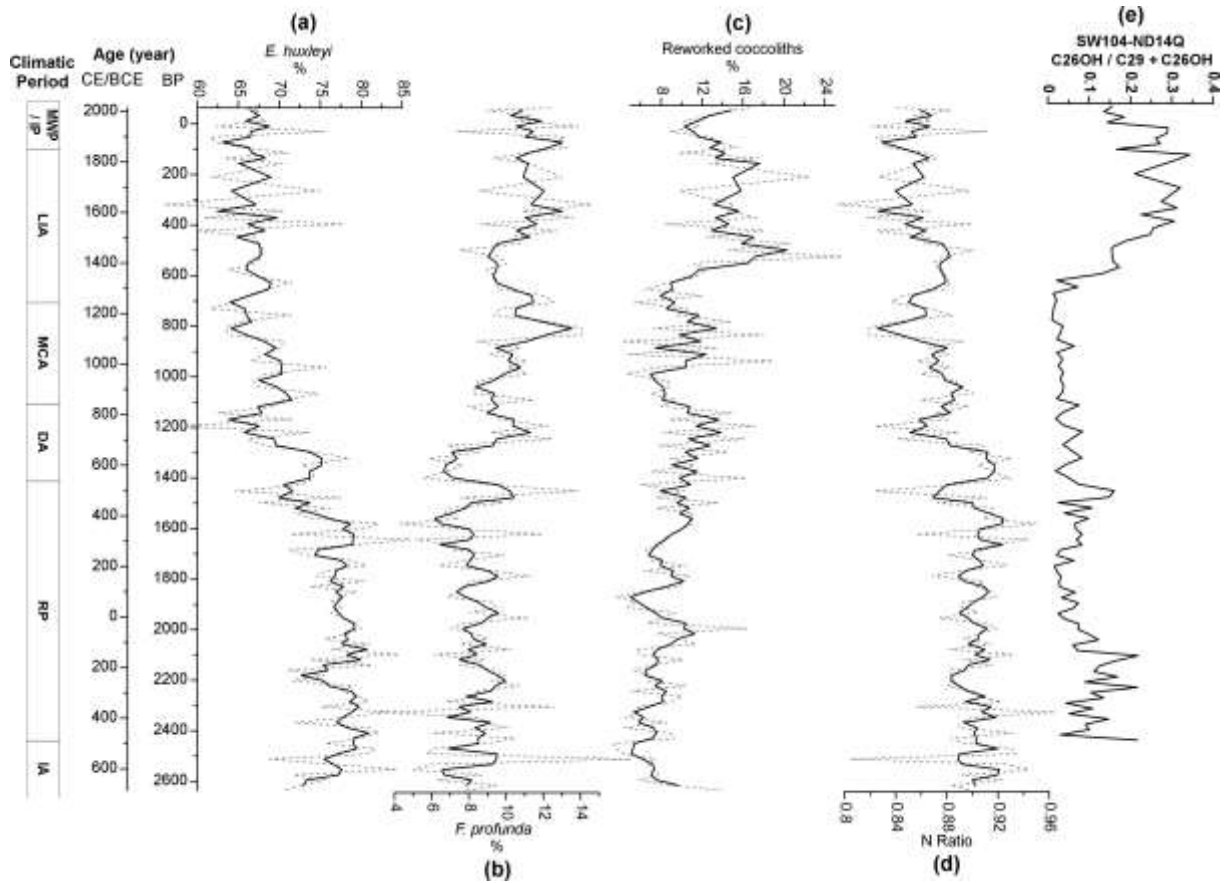
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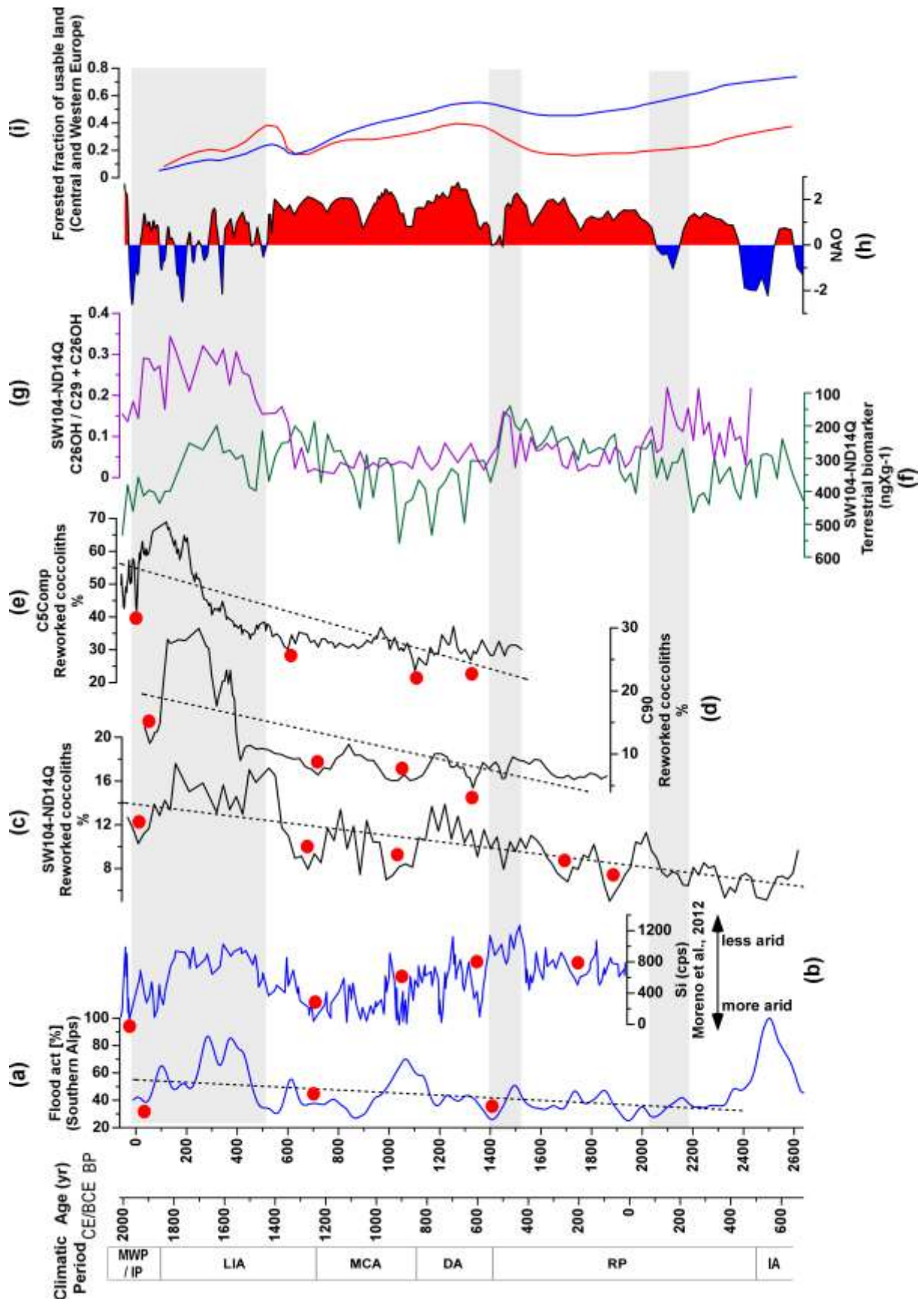
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Figure 2. SW104-ND14Q age-depth model and magnetic susceptibility signal. Sedimentation rate and tephra layers (diamond) were reported.



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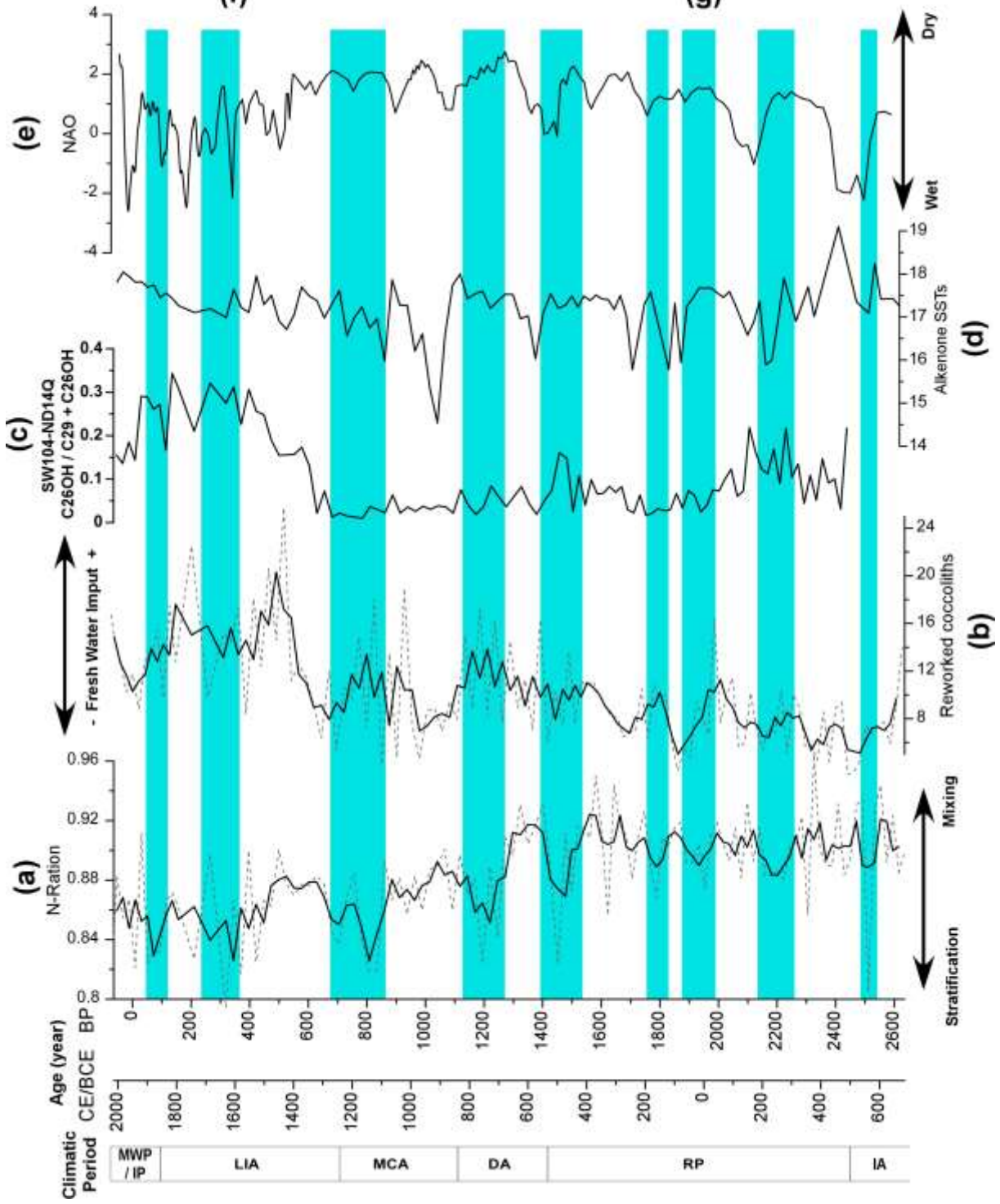
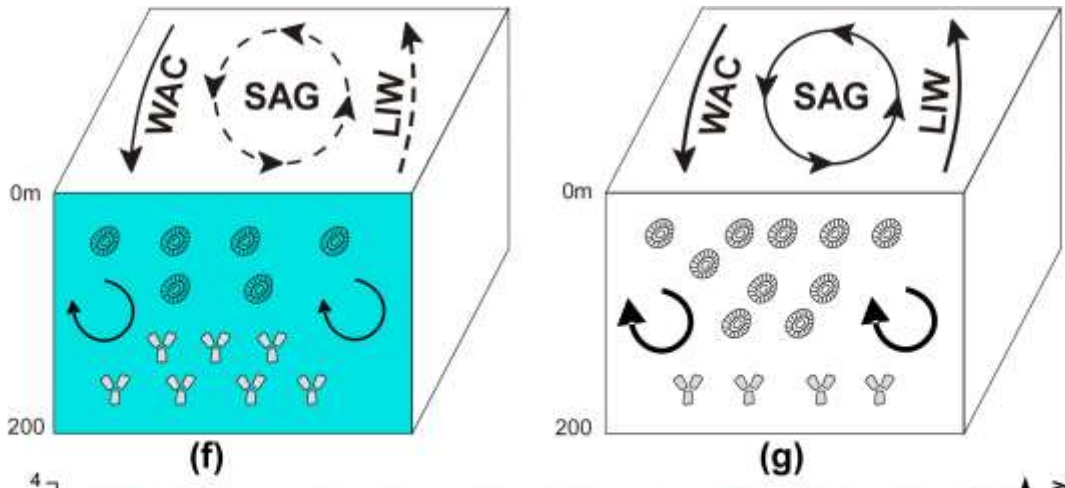
Figure 3. Time domain distribution of (a) *E. huxleyi*, (b) *F. profunda*, (c) RC, (d) N-ratio, and (e) $C_{26OH}/C_{29}+C_{26OH}$ ratio in core SW104-ND14Q. Raw and three points running average data are reported in grey dashed and black full lines, respectively. The age model is from Jalali et al. (2018) and the climate period intervals are those of Margaritelli et al. (2016).



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776 **Figure 4.** Comparison in time domain between (c) SW104-ND14Q, (d) C90, and (e) C5Comp
 777 reworked coccoliths, (g) SW104-ND14Q $C_{26OH}/C_{29}+C_{26OH}$ ratio, (b) Si fluctuations (Moreno
 778 et al., 2012), and (a) Flood frequency reconstruction from Southern Alps (Wirth et al., 2013).
 779 (f) Terrestrial biomarker concentration (Jalali et al., 2018) and (i) Forest fraction of usable
 780 land (Kaplan et al., 2009) are reported. The dots mark the dry spells identified in the records.

781 The bands highlight the relationship between $C_{26OH}/C_{29}+C_{26OH}$ ratio and negative (h) NAO
782 states. The climate periods are from Margaritelli et al. (2016).
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789 **Figure 5.** Schematic view of inferred relationship between (a) SW104-ND14Q N-Ratio and
790 (f, g) SAS hydrology. (b) %RC fluctuations, (c) $C_{26OH}/C_{29}+C_{26OH}$ ratio, (d) SSTs fluctuations,
791 and (e) winter NAO index (Olsen et al., 2012; Trouet et al., 2009) are reported. The bands
792 highlight the N-Ratio during periods of stratified surface water (diagram f). The climate
793 periods are from Margaritelli et al. (2016).