

# The Valanginian Weissert Event on the south Tethyan margin: a dynamic paleoceanographic evolution based on the study of calcareous nannofossils.

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Spangenberg, A. El-Samrani

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- 1 The Valanginian Weissert Event on the south Tethyan margin: a dynamic paleoceanographic
- 2 evolution based on the study of calcareous nannofossils
- 3 M. Shmeit<sup>1,2\*</sup>, F. Giraud<sup>1</sup>, E. Jaillard<sup>1</sup>, S. Reboulet<sup>3</sup>, M. Masrour<sup>4</sup>, J.E. Spangenberg<sup>5</sup>, A. El-
- 4 Samrani<sup>2</sup>
- 5 1. Univ. Grenoble Alpes, Univ. Savoie Mont Blanc, CNRS, IRD, Univ. Gustave Eiffel,
- 6 ISTerre, 38000 Grenoble, France
- 7 2. Lebanese University, Doctoral School of Science and Technology, Laboratory of
- 8 Geosciences Georesources and Environment L2GE, EDST/PRASE, Beirut, Lebanon
- 9 3. Univ Lyon, UCBL, ENSL, UJM, CNRS, LGL-TPE, F-69622, Villeurbanne, France
- 10 4. Université Ibn Zohr, Faculté des Sciences, Département de Géologie, BP 8106, Cité
- 11 Dakhla, Agadir, Morocco
- 12 5. University of Lausanne, Institute of Earth Surface Dynamics (IDYST), CH-1015 Lausanne,
- 13 Switzerland
- 14 \* corresponding author: majd.homaidan-shmeit@u-bourgogne.fr
- 15 Highlights:
- Paleoceanographic reconstruction around the Weissert Event in the southern Tethys
- High surface-water fertility and nannofossil production during the Weissert Event
- Major impact of sea-level and river clastic input on the nannofossil community
- Sea level fall caused the nannoconid decline before the Weissert Event
- Nannoconid recovery was associated with high surface-water fertility

#### 21

## **ABSTRACT:**

22 Paleoceanographic conditions across the Valanginian Weissert oceanic anoxic event 23 were reconstructed for the first time on the central Moroccan margin from a quantitative-based 24 calcareous nannofossil study. Two onshore successions in the Essaouira-Agadir Basin and one 25 offshore from DSDP Hole 416A were studied, providing a proximal-distal transect on the 26 central Moroccan margin. The paleoceanographic conditions were reconstructed in an accurate 27 chronostratigraphic framework. The sedimentary evolution of the Essaouira-Agadir Basin 28 shows that sea-level variations controlled the paleogeographic context, clastic input, and 29 nutrient availability. A thick photic zone and mesotrophic conditions favored the development 30 of a diverse nannofossil community before the Weissert Event in the early Valanginian. In all 31 studied successions, a collapse of the nannofossil community was recorded during the onset of 32 the Weissert Event at the early-late Valanginian transition, and was caused by a major sea-level 33 fall. The nannofossil community gradually recovered and reached its highest nannofossil 34 production associated with high surface-water fertility during the Weissert Event in the late 35 Valanginian. This eutrophication is coeval with the high and stable  $\delta^{13}C_{carb}$  values of the carbon-36 isotope positive shift. After the Weissert Event, surface-water fertility and nannofossil 37 production decreased but remained higher than during the early Valanginian. Additionally, the 38 nannoconid decline started in the early Valanginian and was linked to a low sea-level. Their recovery in the early late Valanginian occurred during conditions of high sea-level and of high 39 40 surface-water fertility. The central Moroccan margin is integrated into a global 41 paleoceanographic reconstruction.

42 Keywords: Carbon isotope excursion; Central Moroccan margin; Biostratigraphy;
43 Sedimentary evolution; Surface-water fertility

#### 44 **1. Introduction**

45 The Valanginian stage (137.7 – 132.6 Ma; Gale et al., 2020) records the first positive 46 carbon-isotope excursion (CIE) of the Early Cretaceous (Lini et al., 1992; Weissert and Erba, 47 2004), termed the Weissert oceanic anoxic event or the Weissert Event (Erba et al., 2004; Kujau 48 et al., 2012). The amplitude of the CIE ranges between 1.5–2 % in bulk marine carbonate rock 49 and 1.5-3 % in marine organic matter (Lini et al., 1992; Hennig et al., 1999; McArthur et al., 50 2007; Charbonnier et al., 2013). In addition, the positive CIE was recognized in terrestrial fossil 51 plants with a magnitude of ~4 %, implying a perturbation of both the oceanic and atmospheric 52 Earth systems (Gröcke et al., 2005; Nunn et al., 2010). Martinez et al. (2015) identified three phases for the CIE, a first phase of rapid  $\delta^{13}C_{carb}$  increase with a duration of 0.60 myr, a second 53 phase of stable  $\delta^{13}C_{carb}$  values lasting 1.48 myr, and a final phase of smooth decrease in  $\delta^{13}C_{carb}$ 54 55 values lasting 3.77 myr and extending to the early-late Hauterivian boundary. The onset of the 56 positive CIE was observed just below the early-late Valanginian boundary (Duchamp-Alphonse 57 et al., 2007; McArthur et al., 2007; Gréselle et al., 2011; Martinez et al., 2015). In the Tethyan 58 Realm, the positive CIE was dated from the late NK3A calcareous nannofossil Subzone and 59 extends to the NC4A calcareous nannofossil Subzone in some localities (Lini et al., 1992; Sprovieri et al., 2006; Duchamp-Alphonse et al., 2007; Bornemann and Mutterlose, 2008; 60 Charbonnier et al., 2013; Aguado et al., 2018). Calibrated with ammonite biostratigraphy, the 61 62 onset of the positive CIE was identified in the lower part of the Karakaschiceras inostranzewi 63 Standard Zone (uppermost early Valanginian) and the highest values occur during the 64 Saynoceras verrucosum Standard Zone (lowermost late Valanginian; McArthur et al., 2007; 65 Gréselle et al., 2011; Martinez et al., 2015; Aguado et al., 2018). The aforementioned zones follow the standard ammonite zonation that is suitable for the Mediterranean Province of the 66 67 Mediterranean-Caucasian Subrealm (Tethyan Realm; Reboulet et al., 2018). In addition to the 68 Tethyan Realm, records of the positive CIE extend to a wide range of geographic locations that include the Atlantic Ocean (Brenneke, 1978; Cotillon and Rio, 1984), the northwestern Pacific
Ocean (Bartolini, 2003), the Boreal Realm (Price and Mutterlose, 2004; Meissner et al., 2015;
Morales et al., 2015), and the southern Hemisphere (Aguirre-Urreta et al., 2008; Gómez-Dacal
et al., 2018).

73 The positive CIE has been associated with a global increase in marine surface-water 74 primary productivity (Erba et al., 2004). During the Valanginian, calcareous nannofossils and 75 dinoflagellates were the major marine planktonic primary producers (Bown et al., 2004). 76 Several studies on calcareous nannofossil assemblages have shown enhanced surface-water 77 trophic conditions occurring before and during the Weissert Event, inferred from increasing 78 abundances of high-fertility taxa such as Biscutum, Discorhabdus, Diazomatolithus, and small 79 Zeugrhabdotus spp. (Tethyan Realm: Bersezio et al., 2002; Erba and Tremolada, 2004; Duchamp-Alphonse et al., 2007; Mattioli et al., 2014; Atlantic Ocean: Bornemann and 80 Mutterlose, 2008; Kessels et al., 2006; Pacific Ocean: Lozar and Tremolada, 2003; and Boreal 81 82 Realm: Williams and Bralower, 1995; Melinte and Mutterlose, 2001; Kessels et al., 2006). 83 Nevertheless, paleoceanographic reconstructions based on calcareous nannofossils from the 84 south Tethyan margin have not been documented yet, such that it needs to be further 85 investigated. The causes of the increase in marine primary productivity differ in coastal and 86 distal pelagic settings. Eutrophication in the coastal settings is hypothesized to have been caused 87 by a greenhouse climate, intensification of the hydrological cycle, and consequently higher 88 weathering rates and detrital input (Lini et al., 1992). Climate reconstructions demonstrated 89 intensified hydrolyzing conditions culminating around the early-late Valanginian transition 90 observed from clay mineral distributions (Duchamp-Alphonse et al., 2011; Charbonnier et al., 91 2020) and spore-pollen ratios (Kujau et al., 2013). Eutrophication in pelagic settings could be 92 explained by the introduction of limiting bio-metals from hydrothermal vents during the break-93 up of Gondwana (Lini et al., 1992; Weissert et al., 1998; Erba et al., 2004). Sea-level changes

might have also influenced the fertilization of the ocean. (1) Enhanced continental runoff 94 95 increases nutrient fluxes to the proximal parts of the ocean (gradual fertilization) during low 96 sea level. (2) An increase in nutrients within the outer shelf realm may be linked to upwelling 97 (sharp fertilization) during high sea level. In the late Valanginian, a global sea-level rise was 98 documented after an all-time minimum of the Early Cretaceous occurred in the mid-99 Valanginian (Haq, 2014). The different hypotheses that could cause the fertilization of the 100 ocean need to be tested for the Weissert positive CIE. A biocalcification crisis in both platform 101 and pelagic settings has been inferred from carbonate platform drowning around the early-late 102 Valanginian (Weissert et al., 1998; Wortmann and Weissert, 2000; Föllmi et al., 2006), and a 103 nannoconid decline predating the positive CIE has been identified (Channell et al., 1993; 104 Bersezio et al., 2002; Erba and Tremolada, 2004; Gréselle et al., 2011; Barbarin et al., 2012).

105 This study aims to reconstruct the paleoceanographic conditions around the Weissert 106 Event on the central Moroccan margin, which was located on the south Tethyan margin during 107 the Cretaceous. Calcareous nannofossil abundances and accumulation rates together with their 108 assemblages were used to quantify changes in surface-water nannofossil production and fertility 109 conditions. Nannofossil samples were collected from three geological successions located on a 110 proximal-distal transect: two onshore sections (Zalidou and Aït Hamouch belonging to the 111 Essaouira-Agadir Basin, EAB) and an offshore DSDP Hole 416A (east Atlantic Ocean). The 112 paleogeographic context of the onshore sections was defined from a sedimentological analysis 113 done at the scale of the EAB. The time framework was based on ammonite (only for the onshore 114 sections) and calcareous nannofossil biostratigraphy, and carbon-isotope stratigraphy. The 115 transect allowed to better constrain the variations of continental nutrient flux, upwellings that 116 were present on the NW African margin during the Early Cretaceous due to active trade winds 117 considering atmospheric circulation models from Price et al. (1995) and Poulsen et al. (1998), 118 and calcareous nannofossil production.

119 **2. Geological setting** 

120 The Essaouira-Agadir Basin (EAB) is located in southwest Morocco and corresponds 121 to the westernmost part of the High Atlas fold belt, bounded by 9°55' to 9°20' W and 30°30' to 122 31°05' N. The basin is bordered to the north by the Jebilet Massif and Doukkala Plateau 123 (Paleozoic basements), to the south by the plain of Souss (Neogene basin), and to the east by 124 the High Atlas (Fig. 1). The EAB presently constitutes part of the Atlantic passive margin which 125 is considered one of the oldest existing passive margins (Stets and Wurster, 1982; Sahabi et al., 126 2004). The EAB extends offshore until the western limit of the continental margin (Frizon de 127 Lamotte et al., 2008). During the Cretaceous, the central Moroccan margin which includes the 128 EAB occupied a paleogeographic location on the south Tethyan margin, between paleo-129 latitudes ~ 15°N and ~ 22°N (Fig. 2). Until the opening of the South Atlantic (Valanginian) and 130 its connection with the Central Atlantic (early Albian), the Tethys Ocean included what is 131 presently the Central Atlantic. Therefore, the geological evolution of the Atlantic-Tethyan 132 passive margin controlled the development of the EAB (Ellouz et al., 2003). Two sections in 133 the EAB, namely Zalidou and Aït Hamouch, were selected for the study of calcareous 134 nannofossils. The Zalidou section is located 45 km north of Agadir (30°54'23"N; 9°39'48"W; 135 Fig. 1B), whereas the Aït Hamouch section is located ~25 km NNW of Agadir (30°36'54"N; 136 9°42'14"W). The Aït Hamouch section is composed of two adjacent sections, only 0.6 km apart 137 from each other. The primary section, denoted "AH", was not accessible in its middle part 138 (between 32 and 42 m). Accordingly, the nearby "ATH" section was studied and correlated 139 using ammonite biostratigraphy and sedimentological discontinuities. Moreover, calcareous nannofossils were studied in the offshore succession DSDP Leg 50 Hole 416A (Fig. 1A). 140 141 Additional sections were used in this paper to improve the precision of the ammonite 142 biostratigraphy and the sedimentological framework (Fig. 1B), although these sections will not 143 be discussed in detail herein as this is beyond the scope of this paper.



Fig. 1. Location of the study area in SW Morocco, modified from Ouajhain et al. (2009). (A) Location of the central Moroccan margin including the Essaouira-Agadir Basin and DSDP Hole 416A (not to scale). Approximate location of the Mesozoic shelf margin from Tari and Jabour (2013). (B) Geological sketch including the location of the sections used in this study (square shape). The successions studied for their calcareous nannofossil assemblages (Zalidou, Aït Hamouch, and Hole 416A) are identified in "bold" font and by a "star" sign.



151 -180 -150 -120 -90 -60 -30 0 30 60 90 120 150 180
152 Fig. 2. Paleogeographic map of the Valanginian (136 Ma) showing the location of the central
153 Moroccan margin (star shape). The "white circles" correspond to the location of
154 basins/successions where calcareous nannofossils of the Weissert Event were investigated:
155 Tethyan Realm, (1) Vocontian Basin (France) (Duchamp-Alphonse et al., 2007, 2014; Gréselle
156 et al., 2011; Barbarin et al., 2012; Mattioli et al., 2014), (2) Lombardy Basin (Italy) (Bersezio
157 et al., 2002; Erba and Tremolada, 2004), and (3) Moesian Platform (south and east Carpathians158 Romania) (Melinte and Mutterlose, 2001); western and central Atlantic, (4) DSDP 534A (Blake

Bahama basin, NE Florida) and (5) DSDP 603B (east Florida) (Bornemann and Mutterlose,
2008), and (6) DSDP 535 (SE Gulf of Mexico) (Kessels et al., 2006); Boreal Realm, (7) ODP
638 (NW Iberian Peninsula) (Kessels et al., 2006), (8) BGS 81/43 (southern North Sea Basin,
east England) (Williams and Bralower, 1995; Kessels et al., 2006), and (9) east Greenland Basin
(NE Greenland) (Pauly et al., 2012); western Pacific, (10) ODP 1149B (Nadezhda Basin)
(Lozar and Tremolada, 2003). Construction from http://www.odsn.de (made from the datafiles
of Hay et al. (1999)).

#### 166 **2.1 Ammonite zonal framework for the late Berriasian-early Hauterivian interval of the**

167 **EAB** 

168 The systematic revision and identification of ammonites sampled at Zalidou and Aït 169 Hamouch sections are being prepared as a separate paper and were part of a chapter in Shmeit's 170 PhD thesis (2021), in which paleontological comments and ranges of ammonite taxa are 171 provided. The standard ammonite zonation for the Mediterranean Province (Tethyan Realm, 172 Reboulet et al., 2018) is applied here. Only two regional units are added, namely Neocomites subtenuis and Busnardoites campylotoxus local subzones, to make more accessible the 173 174 comparisons with previous works made in this Moroccan area by Wippich (2001, 2003) and 175 Ettachfini (2004). The ammonite (sub-)zones are interval (sub-)zones defined by the first 176 appearance datum (FAD) of two successive ammonite-indices. Hereafter will be used the 177 following abbreviations: standard zone = StZ, standard subzone = StSz, and local subzone = 178 LSz.

179 The base of the *Thurmanniceras pertransiens* StZ, and therefore the lower boundary of 180 the Valanginian, is placed by the FAD of the index-species (Kenjo et al., 2021). As it is 181 relatively rare at Zalidou and Aït Hamouch, the base of the zone can be placed at the first records 182 of Neocomites premolicus (Ettachfini, 2004; Reboulet et al., 2018). Due to the rarity of the 183 index-species, the base of the N. neocomiensiformis StZ is identified at Zalidou and Aït 184 Hamouch sections by the first occurrence (FO) of *Neocomites subtenuis*, as its appearance is 185 contemporaneous with that of *N. neocomiensiformis* in SE Spain (Company and Tavera, 2015). 186 Above the *N. subtenuis* LSz, the upper part of the *N. neocomiensiformis* StZ is characterized by

187 the Busnardoites campylotoxus LSz, for which the base is placed at the FO of its relatively 188 abundant index-species at Zalidou and Aït Hamouch. The base of the Karakaschiceras 189 inostranzewi StZ is placed at the FO of its index species. In both sections, Saynoceras 190 verrucosum, index species of the S. verrucosum StZ, was not found. Other ammonite markers 191 of the basal part of this zone, as Valanginites ventrotuberculatus, were also not observed. 192 Consequently, at Zalidou, the base of the S. verrucosum StZ, and so the base of the late 193 Valanginian, are placed at the FO of Karakaschiceras pronecostatum that is the index-species 194 of the second subzone of the S. verrucosum StZ. At Aït Hamouch, very few specimens of some 195 taxa that could be restricted to the S. verrucosum StZ are recorded in a single layer (AH89) with 196 the FO of *Neocomites peregrinus* allowing to identify the base of the *N. peregrinus* Stz. Thus, 197 the ammonite assemblage could be interpreted as a result of a condensed lithology. In this layer, there are also few doubtful specimens of K. inostranzewi?; their preservation could indicate 198 199 that they have been reworked. At Zalidou, N. peregrinus was also observed allowing to place 200 the base of the zone. In this section, the Olcostephanus nicklesi StSz is identified by the 201 occurrence of the genus Himantoceras (see Reboulet, 1996). The O. nicklesi StSz is not 202 recognized at Aït Hamouch. The base of the Criosarasinella furcillata StZ and that of the 203 Tescheniceras callidiscum StSz are identified in both sections. The base of the Acanthodiscus 204 radiatus StZ, and thus the base of the Hauterivian (see Mutterlose et al., 2020), are placed at 205 the FO of Breistrofferella castellanensis and that of Acanthodiscus radiatus at Zalidou and Aït 206 Hamouch, respectively.

## 207 2.2 DSDP Hole 416A

DSDP Leg 50 Hole 416A was drilled ~180 km NW of Essaouira (32°50'10.7"N; 10°48'03.6"W) at a water depth of 4201 m in the eastern Atlantic (Fig. Supp. 1). The Berriasian to Hauterivian sediments consist of a turbidite succession deposited in a lowermost continental rise environment (Lancelot et al., 1980). The Valanginian interval was identified onboard from 212 cores 48 to 14, between -1540 and -1222 meters below seafloor (mbsf) (Čepek and Gartner, 213 1980; Sliter, 1980). The detailed lithology of Hole 416A was described by Lancelot et al. 214 (1980). Briefly, the Berriasian and early Valanginian were assigned to lithological unit VII (from -1624 to -1430 mbsf, cores 57 to 37), characterized by alternating siliciclastic and 215 216 carbonate turbidites. The lithology displays quartz-rich siltstone and mudstone cycles 217 alternating with hard micritic and calpionellid-rich limestones. A prominent change in lithology 218 above -1430 mbsf characterizes the shift to the overlying lithological unit VI (from to -1430219 to -880 mbsf, cores 36 to 7). This change is identified by an abrupt decrease in carbonate 220 deposition and a subsequent increase in terrigenous input, with siliciclastic terrigenous 221 turbidites replacing the calcareous turbidites. The lithology change was regarded as a classical 222 example of the mid-Valanginian platform drowning (Schlager, 1980). Lithological unit VI 223 spans the rest of the Valanginian and builds up to the Hauterivian (Lancelot et al., 1980; 224 Schlager, 1980). The lithology shows graded calcareous and siliciclastic cycles, from siltstone 225 or fine sandstone to mudstone.

**3. Materials and methods** 

227 The onshore successions (Zalidou and Aït Hamouch) span the late Berriasian to early 228 Hauterivian covering the entire Valanginian. The offshore DSDP Leg 50 Hole 416A spans the 229 Valanginian stage (Čepek and Gartner, 1980). Samples for calcareous nannofossils, calcium 230 carbonate content, and carbon-isotope analyses were collected from different lithologies 231 (except sandstones). The sampling interval was on average 0.3 m in Zalidou, 0.5 m in Aït 232 Hamouch, and 5 m in Hole 416A. Both macrofossils and nannofossil smear slides are curated 233 at the Collections de Géologie de l'Observatoire des Sciences de l'Univers de Grenoble 234 (OSUG), with an appropriate UJF-ID number. "OSUG-COLLECTIONS" is a database of 235 rocks, minerals, and fossils (https://web.collections.osug.fr, OSUG, UGA; doi:10.5072/OSUG-236 COLLECTIONS.all).

237 **3.1 Sedimentological analysis** 

238 Sedimentological analysis was based on sections located in the southern and central 239 parts of the EAB (Fig. 1B). Observations were made both in the field and under microscope. 240 All sedimentary features (lithology, peculiar surfaces, sedimentary figures, faunal content, and 241 bioturbation) were observed and recorded in the field. Limestone beds, mainly in the lower 242 Valanginian series, were also sampled for microfacies analysis. We used the classification 243 defined by Dunham (1962).

### 244 **3.2 Calcareous nannofossils**

Biostratigraphic and paleoenvironmental analyses of the Zalidou and Aït Hamouch sections were based on 85 and 54 samples (Supplementary Table), respectively, from calcareous claystone to limestone. For Hole 416A, 44 samples were investigated.

248 Smear slides were prepared using a random settling technique developed by Beaufort et 249 al. (2014) and modified by Menini et al. (2019). Only 17 samples from the Aït Hamouch "AH 250 section" were prepared using the random settling technique of Geisen et al. (1999) (see 251 Supplementary Table). Both techniques allow the determination of the nannofossil absolute 252 abundance (nannofossils/gram of rock) using the equations presented in the Supplementary 253 Table. A comparison between both techniques for absolute abundance quantification has not 254 been done yet. Studies comparing other slide preparation techniques have shown that 255 nannofossil proportions can be reliably determined regardless of the technique (Bordiga et al., 256 2015; Lupi et al., 2016).

Nannofossils were observed using a polarizing light microscope (Leica "DM2500 P")
at 1000x magnification. Five hundred specimens per slide, both coccoliths and nannoliths, were
generally counted in a variable number of fields of view (a mean of 52 for Zalidou, 80 for Aït
Hamouch, and 39 for DSDP Hole 416A). All nannofossils with at least more than half of the
specimen preserved were counted. In few samples (4 from Zalidou and 3 from Aït Hamouch),

characterized by low abundance of nannofossils (approx. 1–2 specimens/4 fields of view), ~
350–400 specimens were counted (Supplementary Table). In addition, two longitudinal random
transverses (200 fields of view per transverse) were scanned after each counting for rare species
that could be of biostratigraphic importance. To improve the biostratigraphic schemes whenever
needed, scanning ~8–12 random longitudinal transverses was done on additional samples not
counted (5 from Zalidou, 1 from Aït Hamouch, and 11 from Hole 416A).

The taxonomic frameworks of Perch-Nielsen (1985), Bown and Young (1998), and 268 269 Nannotax3 (Young et al., 2017) were followed. For nannoconids, the work of Deres and 270 Achéritéguy (1980) was followed. The calcareous nannofossil zonation applied to the 271 investigated sections were the NC and NK zones of Bralower (1987) and Bralower et al. (1989, 272 1995), modified from Roth (1978). This zonation scheme was selected because of its applicability to low-latitude Tethyan continental margins and Atlantic oceanic settings 273 274 (Bralower et al., 1995; Bown, 2005; Bornemann and Mutterlose, 2008). The biostratigraphic 275 scheme (CC zones and subzones) of Sissingh (1977), modified by Perch-Nielsen (1979, 1985) 276 and Applegate and Bergen (1988), was also applied to the studied sections because it allows 277 subdividing the early Valanginian. This zonal scheme uses taxa with cosmopolitan distribution 278 and Tethyan affinities, and it relies on the first occurrence (FO) of Eiffelithus taxa (E. windii 279 and *E. striatus*) for subzone divisions during the Valanginian stage.

Nannofossil preservation was evaluated following the classes defined by Roth and Thierstein (1972) and modified by Roth (1983). Nannofossil absolute abundances are controlled by the nannoplankton biomass, preservation, and sedimentation rate. Nannofossil accumulation rates (nannofossils/m<sup>2</sup>/yr) were calculated to cancel the sedimentation rate effect (see formula in Supplementary Table). The duration of the different phases of the positive CIE from Martinez et al. (2015) were used to determine the sedimentation rates in the studied sections. These authors provided an improved age model for the Valanginian-Hauterivian boundary using astrochronology, and acquired absolute age durations for the different phases of the positive CIE. The  $\delta^{13}C_{carb}$  values and the ammonite zonation of the onshore sections (Zalidou and Aït Hamouch) were correlated with those of Martinez et al. (2015).

290 For each nannofossil taxon (or group of combined taxa) the absolute abundance (AA), 291 the relative abundance (RA), and the accumulation rate (AR, only for the Zalidou and Aït 292 Hamouch sections) were calculated. In addition, the nannofossil total absolute abundance 293 (NTAA) and the nannofossil total accumulation rate (NTAR, for Zalidou and Aït Hamouch 294 sections) were calculated, corresponding to the entire nannofossil assemblage. To consider the 295 inherent counting uncertainty, binomial proportion confidence intervals for relative abundances 296 were computed based on the 'exact' Clopper-Pearson method using the PAST software 297 (Hammer et al., 2001; Suchéras-Marx et al., 2019). Additionally, diversity indices were 298 calculated. *Nannoconus* spp. were excluded from the calculation of sample diversity indices, 299 absolute/relative abundances, and accumulation rates of other taxa because of their uncertain 300 biological affinity (Aubry et al., 2005). The counting and the different calculation formulae are 301 given in the Supplementary Table.

302 **3.3 Calcimetry and carbon isotopes** 

The calcium carbonate content was measured on 117 and 56 bulk rock samples from Zalidou and Aït Hamouch sections, respectively (Supplementary Table). For Aït Hamouch, analyses were done on samples from the "AH" section. Additionally, 88 samples from DSDP Hole 416A were measured (Supplementary Table). The CaCO<sub>3</sub> content was determined using the carbonate bomb technique (Müller and Gastner, 1971). For the calculation of the calcium carbonate percentage, see Appendix A in Peybernes et al. (2013).

Carbon stable isotope analysis was performed on the same samples as for the calcium
carbonate content at the stable isotope laboratories of the Institute of Earth Surface Dynamics
of the University of Lausanne, using a Thermo Fisher Scientific Gas Bench II carbonate

312 preparation device connected to a Delta Plus XL isotope ratio mass spectrometer. The CO<sub>2</sub> 313 extraction was done at 70°C. The carbon stable isotope ratios were reported in the delta ( $\delta$ ) 314 notation as the per mil (%) relative to the Vienna Pee Dee belemnite standard (VPDB), where  $\delta = (R_{sample} - R_{standard})/R_{standard}$  and  $R = {}^{13}C/{}^{12}C$ . The  $\delta^{13}C_{carb}$  values were standardized relative 315 316 to the international VPDB scale by calibration of the reference gases and working standards with international reference materials NBS 18 (carbonatite,  $\delta^{13}C = -5.01 \%$ ) and NBS 19 317 (limestone,  $\delta^{13}C = +1.95 \%$ ). Analytical uncertainty (1 sigma), monitored by replicate analyses 318 319 of the international calcite standard NBS 19 and the laboratory standard Carrara Marble ( $\delta^{13}$ C = +2.05 %) was not greater than  $\pm 0.05\%$  for  $\delta^{13}$ C. 320

**4. Results** 

## **322 4.1 Summary of the Valanginian sedimentary evolution**

323 Geological sections indicated in "bold" font hereafter, correspond to sections that have 324 been investigated for calcareous nannofossils in the present study. Detailed analysis allowed 325 the identification of several sequences. Three main stages can be recognized in the Valanginian 326 sedimentary evolution of the EAB.

327 1. The top of the Berriasian carbonate shelf is marked by a karstified surface in the northern sections. The overlying lower Valanginian succession (8-15 m) is dominated by 328 329 marlstone and thin limestone beds (Fig. 3). It shows a thinning-upward, then thickening-upward 330 trend. In the thinning-upward lower part, ammonites, belemnites, plicatulids, and brachiopods 331 dominate. They are also associated with ostreids, *Trichites* sp., pectinids, and with sea-urchins 332 at the base (Fig. 3). Microfacies show mudstone to wackestone textures containing mainly 333 echinoids, bivalves, and agglutinated foraminifers. The fauna indicates an outer shelf 334 depositional environment that represents the maximum depositional depth for the early 335 Valanginian period. In the upper part of the lower Valanginian succession, limestone beds are 336 more abundant and sandstone beds appear in the south (Aït Hamouch). The macrofauna is 337 composed of ammonites, belemnites, brachiopods, oysters, and plicatulids, with subordinate 338 regular and irregular sea-urchins, crinoids, serpulids, and locally corals and bone fragments 339 (Fig. 3). Compared with the lower part, this assemblage indicates a shallower depositional 340 depth. Microfacies analysis reveals wackestone textures containing agglutinated foraminifers, 341 brachiopods, echinoids, and benthic foraminifers. Several discontinuities (erosion, karsted 342 surfaces) indicate the existence of three or four depositional sequences, and support a marked 343 shallowing-upward trend that culminated in the late early Valanginian. The Obbay section 344 exhibits a very reduced lower Valanginian succession (3 m), mainly made of limestone beds 345 alternating with thin marlstone interbeds (Fig. 3). Limestones show a wackestone texture 346 containing abundant phosphate-rich glauconitic oolites and are frequently capped by iron-rich 347 hardgrounds. The causes of condensation of this series are discussed in section 5.1.

348 2. The lower part of the upper Valanginian series (5-15 m) is mainly made of marlstone, 349 thin limestone beds, sandy marlstone, and sandy limestone. The sandy beds are more abundant 350 to the north and east (Fig. 3). The succession shows a thinning-upward trend, locally overlain 351 by a thickening-upward series, interrupted by a major erosion period (D7) that removed one 352 depositional sequence in many sections (Fig. 3). It contains an outer shelf to hemipelagic 353 macrofauna (ammonites, belemnites, brachiopods, and plicatulids), although small bivalves and 354 wood fragments are locally found. The upper Valanginian succession is interpreted as a 355 transgressive-regressive sequence, the top of which is frequently eroded. The maximum 356 depositional depth reached a rather deep outer shelf environment during the "middle" late 357 Valanginian.

358 3. The uppermost Valanginian series (8-22 m) is mainly composed of marlstone, thin 359 limestone beds, sandy limestone to the south, and of sandy marlstone and calcareous sandstone 360 to the north. The succession exhibits an overall thinning-, then thickening-upward trend. The 361 macrofauna is dominated by ammonites, belemnites, brachiopods, plicatulids, and serpulids, 362 locally associated with gastropods and wood fragments. To the north and east of the basin, the 363 uppermost Valanginian deposits are made of sandstone (**Zalidou**) locally showing current 364 ripples, Hummocky Cross Stratification (HCS), and low angle lamination (Ida w Iddar) (Fig. 365 3), interpreted as foreshore deposits. This succession is thus interpreted as a transgressive-366 regressive sequence reaching an outer shelf environment to the south. The thick overlying 367 Hauterivian marlstone represent a new, major sea level rise (Ferry et al., 2007).





371 **4.2 Calcareous nannofossil biostratigraphy** 

The identified biozones of Bralower et al. (1995) and Sissingh (1977) are illustrated for the Zalidou section (Fig. 4), Ait Hamouch section (Fig. 5), and Hole 416A (Fig. 6). Secondary nannofossil bioevents were also observed (Figs. 4-6). Nannofossil zonal and subzonal markers are illustrated in Figure 7. A gradual size transition was observed between *Eiffelithus windii* and *E. striatus* (Applegate and Bergen, 1988; Bralower et al., 1989; Bown and Concheyro, 2004), making a proper differentiation difficult. Coccoliths with the length of their long-axes greater than 6.0  $\mu$ m and with a relatively wide central area (almost 2/3 or greater of coccolith length) were identified in this study as *E. striatus* (Fig. 7).

381 Additionally, a new calcareous nannofossil biostratigraphy is proposed for Hole 416A. Čepek and Gartner (1980) dated the interval between cores 48-1 and 7-1 (~-1530 to -1000 382 383 mbsf) to the early Valanginian-late Hauterivian based on calcareous nannofossil 384 biostratigraphy. The LO of *R. wisei* was observed in this study in core 11-4 (-1200 mbsf; Fig. 385 6). In the Zalidou and Aït Hamouch sections, the LO of *R. wisei* was observed in the early late 386 Valanginian (lower and upper N. peregrinus StZ in Zalidou and Aït Hamouch, respectively; 387 Figs. 4, 5). Also, our results demonstrate that the Valanginian stage extends to core 9-3 (-1122) 388 mbsf) based on the LO of T. verenae (top NK3B Subzone). In the onshore sections, the top of 389 the NK3B Subzone ends in the late Valanginian (uppermost *N. peregrinus* StSz). The LO of *T*. verenae was also observed at a similar stratigraphic interval both in the Vocontian Basin 390 391 (Duchamp-Alphonse et al., 2007; Gréselle et al. 2011) and in the Subbetic Basin (Aguado et al. 392 2018). Therefore, we can assume that the top of the studied interval at Hole 416A records the 393 late Valanginian, but not the latest Valanginian.



Fig. 4. Calcareous nannofossil and geochemical results in the Zalidou section. Sample locations are indicated by the arrow signs. The recognized stages are plotted with, the ammonite biostratigraphy (zones/subzones) following the ammonite standard zonation of the Mediterranean Province

- 397 from Reboulet et al. (2018), calcareous nannofossil biostratigraphy (zones/subzones) following the (a) NC and NK zonation of Bralower et al.
- 398 (1989, 1995) and (b) CC zonation of Sissingh (1977) modified by Applegate and Bergen (1988), sedimentary discontinuities (D0 to D8), and
- 399 lithology. Calcareous nannofossil first and last occurrences are included; taxa indicated in "bold" font are used in the biostratigraphic schemes (a)

and (b), and those in "normal" font correspond to the secondary bioevents. Binomial proportion confidence intervals are included for the nannofossil
relative abundances using the 'exact' Clopper–Pearson method. The proportion of narrow- and wide-canal nannoconids within the nannoconid
population are also shown, "other" nannoconids include specimens whose central-canal was not recognizable. Abbreviations: Berr., Berriasian;
Haut., Hauterivian; *T. pertra.*, *T. pertransiens*; *N. prem.*, *N. premolicus*; *N. neocomiensi.*, *N. neocomiensiformis*; *B. camp.*, *B. campylotoxus*; *K. i.*, *K. inostranzewi*; *S. ver.*, *S. verrucosum*; *K. pro.*, *K. pronecostatum*; *T. c.*, *Tescheniceras callidiscum*; NTAA, nannofossil total absolute abundance
and NTAR, nannofossil total accumulation rate.



407 Fig. 5. Calcareous nannofossil and geochemical results in the Aït Hamouch section, same caption and lithology legend as for Fig. 4. Calcareous
 408 nannofossil results from "AH" section are separated from "ATH" section (see section 2).



410 Fig. 6. Calcareous nannofossil and geochemical results in the DSDP Hole 416A. Sample locations are indicated by the arrow signs. The recognized 411 stage is plotted with, the calcareous nannofossil biostratigraphy (zones/subzones) following the (a) NC and NK zonation of Bralower et al. (1989, 412 1995) and (b) CC zonation of Sissingh (1977) modified by Applegate and Bergen (1988), and lithology. Calcareous nannofossil first and last 413 occurrences are included; taxa indicated in "bold" font are used in the biostratigraphic schemes (a) and (b), and those in "normal" font correspond 414 to the secondary bioevents. Binomial proportion confidence intervals are included for the nannofossil relative abundances using the 'exact' 415 Clopper-Pearson method. The proportion of narrow- and wide-canal nannoconids within the nannoconid population are also shown, "other" 416 nannoconids include specimens whose central-canal was not recognizable. Abbreviations: Berr., Berriasian; magnetic anom., magnetic anomalies; 417 and NTAA, nannofossil total absolute abundance.



418

Fig. 7. Photomicrographs of the selected nannofossil species (XPL: cross-polarized light; PPL: 419 420 plane-polarized light). 1, Calcicalathina oblongata, side view, XPL, sample Za43c; 2, Calcicalathina oblongata, top view, XPL, sample Za49b; 3, Rucinolithus wisei, XPL, sample 421 422 Za27a; 4, Rucinolithus wisei, XPL, sample Za27a, same specimen rotated ~90°; 5, Tubodiscus 423 verenae, XPL, sample 416A 21R2w 103-104; 6, Lithraphidites bollii, XPL, sample AH112a; 424 7, Eiffellithus windii, XPL, sample 416A 10R2w 144-145; 8, Eiffellithus striatus, XPL, sample 425 Za53a; 9, Polycostella parvistellatus, XPL, sample 416A 49R1w 124-125; 10, Hagius 426 circumradiatus, XPL, sample ATH2d; 11, Watznaueria barnesiae, XPL, sample Za28a; 12, 427 Watznaueria fossacincta, XPL, sample Za39a; 13, Diazomatolithus lehmanii, XPL, sample 428 Za32b; 14, Zeugrhabdotus erectus, XPL, sample 416A 9R3w 135-136; 15, Biscutum ellipticum, 429 XPL, sample Za32a; 16, Watznaueria biporta, XPL, sample Za32a; 17, Watznaueria 430 britannica, XPL, sample Za39c; 18, Discorhabdus ignotus, XPL, sample Za39d; 19, Assipetra *infracretacea*, XPL, sample Za32a; 20, *Rotelapillus crenulatus*, PPL, sample 416A 47R1w 7677; 21, *N. steinmannii subsp. steinmannii*, XPL, sample ATH2h; 22, *N. steinmannii subsp. steinmannii*, PPL, sample ATH2h, same specimen; 23, *N. kamptneri subsp. minor*, XPL, sample
ATH2f; 24, *N. kamptneri subsp. minor*, PPL, sample ATH2f, same specimen; 25, *N. globulus subsp. globulus*, XPL, sample ATH2f; 26, *N. globulus subsp. globulus*, PPL, sample ATH2f,
same specimen.

#### 437 **4.3 Calcareous nannofossil assemblages**

The majority of studied samples show good to moderate preservation of calcareous nannofossils with minor dissolution and recrystallization (Fig. 8). Note that the delicate structures of coccoliths are still visible. Highly dissolved and recrystallized coccoliths were limited to 14–18 % of all the samples.

442 Following the abundance classes defined by Erba and Quadrio (1987), calcareous 443 nannofossils are common (10-20 specimens per field of view) in the majority of the studied 444 samples. The calcareous nannofossil assemblage is dominated by eight and nine groups of taxa 445 in the onshore and offshore successions, respectively, illustrated in Figure 7. Altogether, they 446 constitute on average 81 % of the total assemblage at Zalidou, 88 % at Aït Hamouch, and 79 % 447 Hole barnesiae/fossacincta, Diazomatolithus at 416A. They are: Watznaueria lehmanii/subbeticus, small Zeugrhabdotus spp., Biscutum ellipticum, nannoconids, and to a 448 lesser extent Discorhabdus ignotus, other Watznaueria species (W. biporta and W. britannica), 449 450 and Assipetra infracretacea. Rotelapillus crenulatus are negligible in both onshore sections, 451 but not at Hole 416A. The reader is referred to the Supplementary Table for more details.

In this study, *Diazomatolithus lehmanii* and *Diazomatolithus subbeticus* were combined since they are considered as morphological variants (Grün and Allemann, 1975). The small *Zeugrhabdotus* spp. in this study correspond to *Zeugrhabdotus* with long axis  $\leq$ 5 µm combined with *Zeugrhabdotus erectus*. Finally, *W. barnesiae* and *W. fossacincta* were merged since they are considered to be morphological variants (Bornemann and Mutterlose, 2006).

457 The nannoconid group is mainly represented in all studied successions by three taxa, *N*.
458 *steinmannii* corresponding to narrow-canal, and *N. kamptneri* and *N. globulus* both

459 corresponding to wide-canal (see Supplementary Table). These three taxa constitute on average
460 ~80 % of the total nannoconid population in both onshore sections and ~94 % at Hole 416A.
461 Additionally, *N. steinmannii* is the dominant taxon within the total nannoconid population in
462 all three successions (see Supplementary Table).



463

464 Fig. 8. Plot of W. barnesiae/fossacincta relative abundance (RA, %) and taxa richness in

Zalidou, Aït Hamouch, and Hole 416A samples. Preservation classes follow the classes defined
by Roth and Thierstein (1972) and modified by Roth (1983). N is the total number of samples,

467 r and  $r^2$  are the correlation coefficients.

#### 468 4.3.1 Calcareous nannofossil absolute abundance and accumulation rate

Four and three intervals with different sedimentation rates were identified throughout the Valanginian stage in Zalidou and Aït Hamouch, respectively (see Supplementary Table). Thus, allowing the calculation of the nannofossil total accumulation rate (NTAR, nannofossils/m<sup>2</sup>/year). It was not possible to calculate sedimentation rates, and then NTAR, for Hole 416A because the large scatter in  $\delta^{13}C_{carb}$  (Fig. 6) hindered identifying possible correlation points with the  $\delta^{13}C_{carb}$  curve from Martinez et al. (2015).

475 The average nannofossil total absolute abundance (NTAA, nannofossils/gram of rock) 476 is higher offshore compared to onshore, and the average NTAR is higher at Zalidou than at Aït 477 Hamouch (Figs 4-6, see also Supplementary Table). In both Zalidou and Aït Hamouch 478 successions (Figs. 4, 5), the NTAR is high in the early Valanginian. Afterwards, NTAR 479 decreases and minimal values are recorded around the early-late Valanginian transition (K. 480 inostranzewi to S. verrucosum StZ, upper NK3A and upper CC3b subzones). Upwards, in the 481 late Valanginian, NTAR gradually increases in both successions. It reaches maximal values 482 within the *N. peregrinus* StZ at Zalidou (Fig. 4), followed by a slight decrease during the rest 483 of the succession. At Aït Hamouch, NTAR values higher than that of the early Valanginian are 484 recorded within the C. furcillata StZ (Fig. 5). In both successions, NTAR reflects an 485 amplification of NTAA with very similar trends (Figs. 4, 5), which implies that the 486 sedimentation rate does not significantly affect the NTAA. At Hole 416A, NTAA is low in the 487 lower NK3A and CC3a subzones, and increases during the mid-NK3A and upper CC3a to lower 488 CC3b subzones (Fig. 6). Thereafter, the NTAA gradually decreases to minimal values in the 489 NK3B and the CC3b-CC4a subzones transition.

In this study, only the taxa that presented significant variations in all studied successions
are considered (Figs. 4-6). Thereby, the following groups are not described: other *Watznaueria*species (*W. biporta* and *W. britannica*), *A. infracretacea*, *D. ignotus*, and *R. crenulatus*. For
Zalidou and Aït Hamouch sections, the absolute abundance (AA, nannofossils/gram of rock) of

nannofossil taxa are shown in Figs. Supp. 2 and 3, respectively (see also Supplementary Table).
The average accumulation rate (AR, nannofossils/m<sup>2</sup>/year) of each considered nannofossil
taxon is generally higher at Zalidou (Fig. 4) compared to Aït Hamouch (Fig. 5), except for
nannoconids their average AR is higher at Aït Hamouch than at Zalidou (Supplementary Table).

The AR of nannofossil taxa in the onshore sections generally decreases during the *B*. *campylotoxus* LSz and *K. inostranzewi* StZ (NK3A and lower CC3b subzones), earlier than the decrease of NTAR. Their AA in Hole 416A decreases during the same calcareous nannofossil subzones. Notably, the reductions in the NTAR and the AR of taxa are observed earlier in Aït Hamouch (*B. campylotoxus* LSz) with respect to Zalidou (*K. inostranzewi* StZ). During the rest of the successions, the AR or AA of nannofossil taxa generally follow the same trend as that of the NTAR or NTAA, respectively (Figs 4-6).

## 505 4.3.2 Calcareous nannofossil diversity indices and relative abundances

506 Taxa richness and equitability are plotted for Zalidou (Fig. 4), Aït Hamouch (Fig. 5), 507 and Hole 416A (Fig. 6). The offshore succession is characterized by higher nannofossil 508 diversity indices compared to those of the onshore sections. At Zalidou (Fig. 4), the taxa 509 richness shows a long-trend decrease from the early to late Valanginian. The equitability is 510 highest in the early Valanginian. At Aït Hamouch (Fig. 5), the taxa richness decreases to lowest 511 values in the late early Valanginian and in the early late Valanginian. The lowest equitability 512 values are recorded in the late early Valanginian. At Hole 416A (Fig. 6), both taxa richness and 513 equitability increase during the lower NK3A and CC3a to lower CC3b subzones. Upwards, the 514 equitability remains high in the rest of the Valanginian, whereas the richness decreases.

At Zalidou (Fig. 4), the RA of *W. barnesiae/fossacincta* is high in the early Valanginian and around the early-late Valanginian transition, then it decreases during the early late Valanginian. At Aït Hamouch (Fig. 5), its RA increases during the late early Valanginian and during the late Valanginian. The RA of *D. lehmanii/subbeticus* increases at Zalidou in the late 519 Valanginian and remains high upsection. Whereas, its RA at Aït Hamouch decreases in the late 520 early Valanginian, followed by a re-increase in the early late Valanginian. In both onshore 521 sections, the RA of small Zeugrhabdotus spp. increases to highest values in the late 522 Valanginian. The RA of *B. ellipticum* decreases in both onshore sections to minimal values 523 from the late early Valanginian. Its RA re-increases only at Aït Hamouch in the late 524 Valanginian. The RA of nannoconids declines to minimal values in Zalidou (Fig. 4) from the 525 late early Valanginian to the early late Valanginian. Whereas, at Aït Hamouch (Fig. 5), the RA 526 of nannoconids is low in the early Valanginian and increases to maximal values in the early late 527 Valanginian. Change in the proportion of narrow-canal nannoconids within nannoconid 528 population is observed at Aït Hamouch. Their proportion is high from the early Valanginian to 529 the early late Valanginian compared with the rest of the Valanginian (Fig. 5).

At Hole 416A (Fig. 6), the RA of *W. barnesiae/fossacincta* increases during the upper NK3A and CC3b subzones. The RA of *D. lehmanii/subbeticus* is relatively stable throughout the Valanginian. The RA of small *Zeugrhabdotus* spp. increases during the upper NK3A and CC3b subzones. The RA of *B. ellipticum* decreases during the same biozones. The RA of nannoconids decreases in the upper NK3A to NK3B and upper CC3b to CC4a subzones. The proportion of narrow-canal nannoconids within the nannoconid population is generally lower in the lower NK3A and CC3a subzones compared with the rest of the succession.

537 **4.4 Calcium carbonate content and carbon isotopes** 

The CaCO<sub>3</sub> (%) contents range between 15 - 93 (avg. 49 %) at Zalidou, 14 - 88 (avg. 41 %) at Aït Hamouch, and 4 - 92 (avg. 23 %) at Hole 416A. At both onshore sections (Figs. 4-5), the highest CaCO<sub>3</sub> content is recorded in the early Valanginian. At Zalidou, the decrease in CaCO<sub>3</sub> content is concomitant with a lithological change to more clastic-rich marlstonelimestone beds. At Hole 416A (Fig. 6), the CaCO<sub>3</sub> content decreases and exhibits fewer fluctuations in the upper NK3A to NK3B and upper CC3b to CC4a subzones, compared withdown-section.

545 The  $\delta^{13}C_{\text{carb}}$  values range between -2.8 and +1 (avg. -0.8 %) at Zalidou, -3.3 and +0.4 (avg. -1.5 %) at Aït Hamouch, and between -4.9 and +1.9 (avg. -1.0 %) at Hole 416A. At 546 547 Zalidou (Fig. 4), the  $\delta^{13}C_{carb}$  increases from -2.4 % to +1 % between the latest early 548 Valanginian (K. inostranzewi StZ, upper NK3A, and upper CC3b subzones) and the latest 549 Valanginian (upper O. nicklesi StSz, upper NK3B, and upper CC4a subzones). At Aït Hamouch 550 (Fig. 5), the  $\delta^{13}C_{carb}$  increases from ~ -2.5 % to ~ -0.4 % between the latest early Valanginian 551 (K. inostranzewi StZ, upper NK3A, and upper CC3b subzones) and the early late Valanginian 552 (lower N. peregrinus StZ, upper NK3A, and lower CC4a subzones). Accordingly, the carbon-553 isotope positive shift is of magnitude 3.7 % and 2.3 % at Zalidou and Aït Hamouch, respectively. Then, the  $\delta^{13}C_{carb}$  smoothly decreases in the rest of the late Valanginian in both, 554 Zalidou (~ +0.1 ‰) and Aït Hamouch (~ -1.1 ‰). At Hole 416A, the  $\delta^{13}C_{carb}$  does not show 555 556 any positive shift (Fig. 6).

557 **5. Interpretation** 

#### 558 **5.1 Sedimentary paleoenvironment**

559 Except in the lower Valanginian condensed strata of the Obbay section, we did not 560 observe oolites in the limestone beds. This indicates a rather low energy depositional 561 environment in the EAB during most of the Valanginian. As a matter of fact, since trade winds 562 blew to the west, and because the EAB is located on the western coast of Africa, it was little 563 affected by intense tropical storms. These storms may have been generated in the Atlantic 564 Ocean, but were pushed to the west by the trade winds and to the north by the Coriolis force 565 (see Trabucho-Alexandre et al., 2011; Pohl et al., 2019; Jaillard et al., 2021). Therefore, the 566 EAB can be considered to have been a low energy ramp gently sloping toward the ocean. However, the numerous oolites observed on the Obbay swell (Fig. 3) indicate that the 567

568 shallowest part of the water column was moderately agitated, at least in the early Valanginian. 569 Actually, in the sandstone and sandy limestone beds of late early and late Valanginian age, 570 ripples and scarce trough cross beds indicate a moderate wave activity, while planar parallel 571 laminations suggest tidal processes. Also, the occurrence of thin-bedded HCS in shallow marine 572 beds suggests the occurrence of storms of low intensity. During the late early Valanginian and 573 late Valanginian, the moderate hydrodynamic activity of the surficial part of the water column 574 also accounted for the occurrence of sandstone or sandy limestone beds above the surfaces 575 interpreted as sequence boundaries (Fig. 3). This may be due to the erosion occurring during 576 low sea-level periods, which removed the shale particles and cleared up quartz grains from the 577 sediments. The quartz grains were then deposited as sandy beds during the subsequent 578 transgression.

The scarcity of corals and algae, the abundance of agglutinated foraminifers, and the presence of bryozoans in the calcareous beds (mainly early Valanginian) suggest temperate to cold sea water temperatures. Because of the activity of trade winds, the EAB was probably subjected to upwellings (Price et al., 1995; Poulsen et al., 1998). Oceanic upwellings may explain both the cold to temperate waters, and the common occurrence of phosphate in shallow deposited beds.

585 The abundance of ostreids and plicatulids throughout the section suggests mesotrophic 586 conditions, while the presence of agglutinated foraminifers in the early Valanginian suggests, 587 additionally, an oxygen depletion at that time (e.g., Kuhnt et al., 1996; Reolid et al., 2008; 588 Kender and Kaminski, 2017). This interpretation is supported by the abundance of pyrite in the 589 lower Valanginian marlstone of the southern sections. Oxygen depletion, however, must have 590 been mild or sporadic since echinoids are common at that time. Mesotrophic conditions were 591 most likely related to the terrigenous influx of fine-grained clastic particles, more abundant 592 toward the north and east (Fig. 9). This is supported by the sporadic occurrence of wood and

leaf fragments (Zalidou) in Valanginian deposits. Actually, a combination of high terrigenous
supply, temperate to cold temperature, and the presence of upwellings favors the development
of heterozoan carbonate systems (Michel et al., 2018).

## 596 5.2 Paleogeography

597 In the early Valanginian, the northern sections were marked by open marine, oyster-rich 598 shallow carbonate shelf facies exhibiting several karstified surfaces. Whereas, the southern ones 599 presented distal facies marked by thick marlstone intervals, abundant cephalopods, and the lack 600 of emergence evidence (Fig. 9). During the late Valanginian, the northern sections recorded the 601 progradation of a clastic sedimentary system (Fig. 9), which indicates that the clastic supply 602 proceeded from the Central High Atlas (Ferry et al., 2007). Clastic supply was interrupted by a 603 significant erosion period (D7, Figs. 3, 9), likely related to an emergence episode. Furthermore, 604 the uppermost Valanginian deposits end up with foreshore deposits and are capped by an 605 emergence surface to the northeast (Ida w Iddar). Conversely, the coeval sediments in Zalidou 606 bear cephalopods and plicatulids, and end up with a submarine erosional surface to the 607 southwest (Aït Hamouch). The south-westward sloping topography is supported by the onlap 608 of the lower upper Valanginian sequence to the north or northeast (Fig. 3). Although sandstone 609 prograded in the northern part of the basin during the late Valanginian, the southernmost section 610 (Aït Hamouch) displays sandy layers in the late early to earliest late Valanginian (Fig. 9). This 611 suggests that the course of rivers that supplied quartz grains to the EAB was changing through 612 time.

In the Obbay section, the lower Valanginian succession is very condensed (Fig. 3). Two explanations may be proposed. First, the presence of strong upwelling currents in the early Valanginian (Price et al., 1995; Poulsen et al., 1998) could have accounted both for the condensation and the abundance of phosphate and glauconite (Föllmi, 1996; Puhfal and Groat, 2017). Second, incipient movements of the Amsittene diapir (e.g., Tari and Jabour, 2013) or

618 compressional tectonics in Early Cretaceous times (Bertotti et al., 2012; Fernández-Blanco et 619 al., 2020) may have locally uplifted this area. If so, the area became a shallow swell reaching 620 the Fair-Weather Wave Base, on which a high energy regime caused condensation. A 621 combination of both upwelling and uplift is likely. The fact that condensation disappears during 622 the late Valanginian sea level rise suggests that either, upwelling currents were less active, or 623 more probably, their activity was distributed and mitigated in the thicker water column.



Fig. 9. Paleogeographic sketch of the Essaouira-Agadir Basin showing the distribution of
sandstone and limestone in the (A) lower Valanginian and the (B) upper Valanginian deposits.
Reconstruction based on data from this study (black squares) and from Ferry et al. (2007) (black
dots).

#### 629 **5.3 Preservation of a calcareous nannofossil primary signal**

624

630 The relative abundance (RA) of the solution-resistant W. barnesiae/fossacincta taxa is 631 used as an indication of the preservation state. Roth and Krumbach (1986) suggested that 632 relative abundances of W. barnesiae > 40 % indicate high degree of diagenesis and dissolution, 633 while Williams and Bralower (1995) suggested that even higher proportions (> 70 %) indicate 634 assemblage alteration. In the studied samples, the RA of W. barnesiae/fossacincta is on average 635 41 % at Zalidou, 43 % at Aït Hamouch, and 37 % at Hole 416A. Additionally, RA ranges of 636 *W. barnesiae/fossacincta* in this study are not very different from those recorded in other basins: 637 ~50 – 90 % in Lombardian and Belluno Basins-northern Italy (Erba and Tremolada, 2004); 48 638 - 78 % in the Vocontian Basin, SE France (Reboulet et al., 2003); 18 - 77 % in DSDP Site 535,

Gulf of Mexico (Kessels et al., 2006); and 22 – 59 % in DSDP Hole 603B, northern Atlantic
(Bornemann and Mutterlose, 2008). Also, *W. barnesiae/fossacincta* dominate the assemblage
in the Subbetic Domain (SE Spain) although no quantification was done (Aguado et al., 2018).
Most of these studies come from sections having experienced low levels of diagenesis,
supporting that the dominance of *W. barnesiae/fossacincta* is not a local effect.

644 The correlation between the RA of W. barnesiae/fossacincta and the taxa richness was 645 tested in all the studied successions (Fig. 8). Despite the fact that a negative correlation can be 646 suspected (r = -0.37,  $p_{uncorr}$  = 0.0005, n = 85 in Zalidou; r = -0.45,  $p_{uncorr}$  = 0.0006, n = 54 in Aït 647 Hamouch; and r = -0.41,  $p_{uncorr} = 0.005$ , n = 44 in Hole 416A), the preservation classes show that 648 samples with strong dissolution and recrystallization do not necessarily correspond to high 649 proportions of W. barnesiae/fossacinta (Fig. 8). These samples are homogenously distributed 650 for any given proportion of W. barnesiae/fossacinta. Moreover, some samples with good and 651 moderate preservation present a high percentage (> 40 %) of *W. barnesiae/fossacincta*, which 652 implies that the high proportions of W. barnesiae/fossacincta can also be part of the original 653 calcareous nannofossil assemblage during the Valanginian.

Thereby, based on all aforementioned evidence, we assume that the nannofossil assemblages of both the Essaouira-Agadir Basin and the DSDP Hole 416A are slightly affected by diagenetic alteration but still hold a primary signal that can be used for the paleoceanographic reconstruction.

## 658 **5.4 Time succession in calcareous nannofossil assemblages**

The nannofossil total accumulation rate (NTAR) and nannofossil total absolute abundances (NTAA) along with the accumulation rate (AR), absolute abundance (AA), and relative abundance (RA) of the major calcareous nannofossil taxa demonstrate variations from the early to late Valanginian. This, associated with the sedimentology results, will allow the reconstruction of the paleoceanographic conditions on the central Moroccan margin and to

| 664 | better constrain the evolution of paleoceanographic conditions around the Weissert Event. As                  |
|-----|---|
| 665 | the late Berriasian <i>p.p.</i> was only recognized at Zalidou and the early Hauterivian <i>p.p.</i> was only |
| 666 | recognized at Aït Hamouch, the discussion will be focused on paleoceanographic conditions                     |
| 667 | during the Valanginian.   |
|     |   |

- 668 For the paleoceanographic reconstruction, the NTAR (and NTAA for Hole 416A) is
- 669 used as a proxy to estimate the nannofossil production. The accumulation rates and abundances

670 (absolute and relative) of the selected nannofossil taxa are used as a proxy for surface-water

- 671 fertility levels. The ecological affinity of calcareous nannofossil taxa used for this purpose,
- along with the associated literature, are summarized in Table 1.

673 Table 1. Main ecological affinity of the calcareous nannofossil taxa used for paleoceanographic reconstruction, following the specified literature. 1. Bersezio et al. (2002); 2. Bornemann et al. 674 (2003); 3. Coccioni et al. (1992); 4. Crux (1989); 5. Erba (1992); 6. Erba et al. (1992); 7. Erba 675 (1994); 8. Eleson and Bralower (2005); 9. Giraud et al. (2003); 10. Herrle (2002); 11. Herrle et 676 al. (2003); 12. Lees (2002); 13. Lees et al. (2005); 14. Mattioli et al. (2014); 15. Mutterlose 677 (1991); 16. Mutterlose (1992a); 17. Mutterlose (1992b); 18. Mutterlose and Kessels (2000); 19. 678 679 Mutterlose et al. (2003); 20. Mutterlose et al. (2005); 21. Noël et al. (1987); 22. Pauly et al. 680 (2012); 23. Pittet and Mattioli (2002); 24. Premoli Silvá et al. (1989); 25. Roth (1981); 26. Roth 681 (1989); 27. Roth and Bowdler (1981); 28. Roth and Krumbach (1986); 29. Street and Bown 682 (2000); 30. Thierstein (1980); 31. Tremolada et al. (2006); 32. Watkins (1989); 33. Williams 683 and Bralower (1995).

| Taxa                     | Fertility of surface waters   | Ecological strategy   | Paleoceanography   |
|--------------------------|---|---|--|
| B. constans              | eutrophic <sup>5, 18, 19, 24, 25, 26, 27, 28, 29, 32, 33</sup>  | low diversified assemblages <sup>18</sup>                               | upwelling <sup>5, 18, 24, 25, 26, 27, 28</sup>                             |
|                          | meso-eutrophic <sup>8</sup>   | eurytopic <sup>17</sup>   | shelf surface-waters <sup>21,</sup><br>29                                  |
| B. ellipticum            | eutrophic <sup>9, 27</sup><br>meso-eutrophic <sup>16</sup>  | r-selected <sup>18</sup>  | upwelling <sup>6, 12, 30</sup>   |
| D. lehmanii              | eutrophic <sup>1, 4, 14, 20, 31, 33</sup>   |   | transgressive<br>intervals <sup>4</sup><br>upwelling <sup>33</sup>         |
| D. subbeticus            | eutrophic <sup>14</sup>   |   | distal environments <sup>14</sup>  |
| nannoconids              | oligotrophic <sup>1, 2, 3, 7, 11, 18</sup><br>meso-oligotrophic <sup>22</sup>                                   | r-selected <sup>29</sup>  | neritic <sup>10, 25, 27, 28, 29</sup><br>lower photic zone <sup>2, 7</sup> |
| W. barnesiae             | oligotrophic <sup>8, 10, 11, 26, 28, 30, 31, 32</sup><br>mesotrophic <sup>23</sup><br>eurytrophic <sup>14</sup> | eurytopic <sup>13, 15, 21, 29</sup><br>r-selected <sup>12, 23, 30</sup> | oceanic setting <sup>27</sup>  |
| W. barnesiae/fossacincta | <b>v</b> 1  | eurytopic <sup>13</sup>   |  |
| Z. erectus               | eutrophic <sup>5, 9, 10, 11, 24, 25, 26, 27, 28, 29, 32, 33</sup>   | r-selected <sup>6, 12, 23</sup><br>low diversified                      | upwelling <sup>5, 24, 25, 26, 27, 28</sup>                                 |
|                          | more eutrophic than <i>B</i> . <i>constans</i> <sup>6</sup>   | assemblages <sup>18</sup>   | shelf-surface water <sup>18</sup>  |
| Small Zeugrhabdotus      | eutrophic <sup>6, 32</sup>  |   |  |

684

*B. ellipticum* is considered as a morphotype of *B. constans* (Bornemann and Mutterlose, 2006)

#### 685 **5.4.1 The Zalidou section**

686 During the early Valanginian (N. premolicus StSz to upper B. campylotoxus LSz, NK3A 687 and upper CC3a to lower CC3b calcareous nannofossil subzones), except for small fluctuations 688 the nannofossil production did not vary and was relatively high (Fig. 4). During the N. 689 premolicus StSz and N. subtenuis LSz (lower NK3A and CC3a to lower CC3b subzones), 690 favorable surface-water conditions allowed the development of the nannofossil community as 691 suggested by the highest diversity indices. Additionally, local mesotrophic conditions are 692 suggested by the abundant high-fertility taxa. Afterwards, a decoupling between NTAR, 693 diversity indices, AR, and RA of the selected taxa is recorded. In the B. campylotoxus LSz 694 (within the NK3A and CC3b subzones), surface-water conditions became less favorable, 695 leading to a decrease in the AR of all nannofossil taxa and in the diversity. Only the eurytopic W. barnesiae/fossacincta remained high during this time interval, and the nannofossil 696 697 production was maintained (Fig. 4).

In the latest early Valanginian (*K. inostranzewi* StZ, upper NK3A and upper CC3b subzones), the nannofossil production collapsed as supported by both, minimal NTAR and minimal AR of all nannofossil taxa (Fig. 4). Excluding poor preservation, this collapse could be explained by major paleoenvironmental changes.

702 Afterwards, in the late Valanginian (S. verrucosum to upper O. nicklesi StSz, upper 703 NK3A to NK3B and uppermost CC3b to lower CC4a subzones), the nannofossil production 704 gradually increased, supported by the re-increasing NTAR and AR of all nannofossil taxa (Fig. 705 4). The end of this recovery phase was marked by a high nannofossil production and meso-706 eutrophic conditions. The meso-eutrophic conditions are interpreted from the important 707 increase in the small Zeugrhabdotus spp. during the late Valanginian (upper N. peregrinus and 708 upper O. nicklesi StSz, NK3B and CC4a subzones). This time interval was also characterized 709 by the recovery of nannoconids (Fig. 4).

Lastly, in the remaining late Valanginian, the nannofossil production slightly decreased with similar magnitudes as those observed in the early Valanginian. However, despite a decrease in surface-water fertility, the fertility remained higher with respect to the early Valanginian as supported by the higher RA of the small *Zeugrhabdotus* spp. (Fig. 4).

714

## 5.4.2 The Aït Hamouch section

715 During the early Valanginian (N. premolicus StSz to N. subtenuis LSz, lower NK3A 716 and CC3a to lowermost CC3b calcareous nannofossil subzones), the nannofossil production 717 was high with values higher compared to the Zalidou section (Figs. 4, 5). A decoupling between NTAR, diversity indices, AR, and RA of the selected nannofossil taxa is also observed (Fig. 5). 718 719 During the N. premolicus StSz (lower NK3A and CC3a subzones), mesotrophic conditions in 720 surface waters and a thick photic zone allowed a diverse calcareous nannofossil community to 721 flourish. Also, these conditions allowed the development of benthic carbonate producers, such 722 as agglutinated foraminifers, ostreids, and plicatulids (see section 4.1). Afterwards, during the 723 *N. subtenuis* LSz, the surface-water conditions became less favorable for both high-fertility taxa 724 and nannoconids. As at Zalidou, the eurytopic W. barnesiae/fossacincta remained high during 725 this time interval and nannofossil production was maintained (Fig. 5).

726 During the late early Valanginian (B. campylotoxus LSz to K. inostranzewi StZ, NK3A 727 and CC3b subzones), the nannofossil production gradually declined and collapsed (Fig. 5). 728 Excluding poor preservation, the trigger of the collapse can be also explained by major 729 paleoenvironmental changes. Decline in the nannofossil production is suggested from the 730 drastic decrease in accumulation rate and abundance of the dominant taxa, W. 731 barnesiae/fossacincta. Nannofossil production remained low during the early late Valanginian 732 (lower *N. peregrinus* StZ, upper NK3A Subzone and CC3a-CC3b subzones transition; Fig. 5). 733 Decreasing percentages of *W. barnesiae/fossacincta* within the nannofossil assemblage explain that all other taxa show increasing percentages (in particular nannoconids) despite decreasingAR.

736 Afterwards, in the late Valanginian (upper N. peregrinus to C. furcillata StZ, upper 737 NK3A to NC4A and CC4a subzones), the nannofossil production first recovered before 738 reaching higher magnitudes than that of the early Valanginian (Fig. 5). This is attested by the 739 increasing NTAR, and AR of all taxa. In parallel, enhanced surface-water trophic levels are 740 demonstrated by the increasing abundances of both D. lehmanii/subbeticus and small 741 Zeugrhabdotus spp.. The nannoconid AR is maximal, and is associated with decreasing 742 proportion of narrow-canal forms (Fig. 5). Lastly, in the latest Valanginian (T. callidiscum StSz, 743 uppermost NC4A and CC4a subzones), the nannofossil production and trophic levels decreased 744 to magnitudes slightly lower than that of the early Valanginian.

#### 745 **5.4.3 DSDP Hole 416A**

746 The lower NK3A and CC3a calcareous nannofossil subzones are characterized by an 747 increase in the nannofossil production, as suggested by both increasing NTAA and abundances 748 of meso-eutrophic taxa (Fig. 6). Marine conditions were also favorable for the development of 749 nannoconids, as their absolute abundance is higher compared with both onshore sections (Fig. 750 6, and Figs. Supp. 2-3). In addition, the nannoconids were mainly represented by narrow-canal 751 forms with the large dominance of N. steinmannii (Fig. 6). The assumption that narrow-canal 752 *Nannoconus* inhabited the deep photic zone under stratified waters and a deep nutricline (Erba, 753 2004, 1994) has been challenged by Mattioli et al. (2014), who showed that N. steinmannii 754 could have affinity for destratified surface waters. Hole 416A is characterized by turbiditic 755 sequences and low calcium carbonate contents within the Valanginian sediments. The repeated 756 input of clastic material would prevent the stratification of the water column.

757 Upsection, in the upper NK3A-NK3B and CC3b-CC4a subzones, the NTAA and AA
758 of all taxa decrease (Fig. 6). Reduced surface-water fertility during this time interval is shown

by the decreasing relative abundance of high-fertility taxa. Nannoconids decline slightly after
the fertility decreased in the uppermost NK3A and upper CC3b subzones. These are interpreted
as unfavorable surface-water conditions, similar to those observed in the onshore EssaouiraAgadir Basin.

The onset of the nannofossil production recovery, similar to that of NTAR observed onshore, seems to occur in the topmost samples (Fig. 6). However, the recovery phase itself as recorded in the onshore settings seems to be missing at Hole 416A.

## 766 **5.5 The carbon isotope record and identifying the Weissert Event**

At Zalidou, we identify the Weissert Event from the base of the carbon-isotope (CI) positive shift (16.25 m) until the climax (32.8 m; Fig. 10), following the formal definition of Erba et al. (2004). The CI positive shift is observed from the lower part of the *K. inostranzewi* StZ to the upper part of the *O. nicklesi* StSz (upper NK3A to upper NK3B and upper CC3b to upper CC4a calcareous nannofossil subzones). The top of the Weissert Event is recorded just below the LO of the nannofossil biomarker *T. verenae*.

773 At Aït Hamouch, the S. verrucosum StZ and the O. nicklesi StSz are not recognized (see 774 section 2.1). Additionally, the N. peregrinus StZ is reduced (~7 m) compared with Zalidou (~14 775 m). This can be also observed from the reduced NK3B calcareous nannofossil Subzone in Aït 776 Hamouch (1.5 m) compared with Zalidou (12.5 m; Fig. 10). Moreover, the smooth increase in 777  $\delta^{13}C_{carb}$  within the CI positive shift is suspected to be missing from Aït Hamouch. This phase 778 is observed during the O. nicklesi StSz in Zalidou (Fig. 10). Thereby, considering both the 779 biostratigraphy of ammonites and calcareous nannofossils along with the chemostratigraphy 780 from carbon-isotope analysis, it is highly possible that part of the late Valanginian could be 781 missing from Aït Hamouch. Furthermore, the paleoceanographic reconstruction from 782 calcareous nannofossils at Zalidou showed a collapse in nannofossil production just after the 783 onset of the identified Weissert Event interval (K. inostranzewi StZ; Fig. 10). The collapse was

followed by an increase in both nannofossil production and surface-water fertility, and then by
slightly decreasing nannofossil production and fertility during the rest of the Weissert Event
interval. Using all the above-mentioned information, the Weissert Event interval at Aït
Hamouch is cautiously identified from 32.2 to 39.2 m (Fig. 10). This identified Weissert Event
interval starts just before the nannofossil production collapse, and terminates when both
nannofossil production and surface-water fertility recover and prior to the LO of *T. verenae*.
The LO of *T. verenae* demonstrates that it is a reliable event.

791 At Hole 416A, the measured  $\delta^{13}C_{carb}$  values do not show any positive shift (Fig. 6). Wortmann and Weissert (2000) measured  $\delta^{13}C_{org}$  in samples from Hole 416A and identified a 792 793  $\delta^{13}$ Corg positive excursion from -1420 to -1300 mbsf, which is not recognized in this study (Fig. 794 6). In both onshore sections, the base of the Weissert Event is roughly near to the FO of H. 795 *circumradiatus*, although we acknowledge its occurrence is very sporadic. Therefore, we 796 tentatively consider the interval corresponding to the Weissert Event from -1303 to -1122 mbsf 797 in Hole 416A (Fig. 10). Its base is after the onset of the decline in nannofossil production and 798 surface-water fertility, similar to the observation from the onshore sections. The explanation 799 for the absence of a CI positive shift at Hole 416A is unclear. The supposed positive shift might 800 have been diluted by the highly reworked material and terrigenous supply, or concealed by the 801 graded turbidite cycles and/or by the high sedimentation rates known for this interval (Lancelot 802 et al., 1980; Schlager, 1980). Indeed, the Valanginian sediments at Hole 416A are more than 803 400 m thick (Lancelot et al., 1980). Furthermore, the carbonate sediments in all studied 804 successions, including Hole 416A, generally present a similar order of magnitude to the  $\delta^{13}C_{carb}$ 805 values (~ −1.0 %*o*).

In the Essaouira-Agadir Basin (EAB), the calibration of the Weissert Event with calcareous nannofossils allows the assignment of the CI positive shift to the upper NK3A-upper NK3B subzones (upper CC3b to "middle" CC4a). The base of the Event is close to the FO of 809 *H. circumradiatus*, whereas its end coincides roughly with the LO of *T. verenae* (Fig. 10). The 810 Zalidou section is the most appropriate section to study the Weissert Event in the EAB. 811 Although sedimentological discontinuities and related hiatuses are present, it is the most 812 complete section and well dated by ammonite and nannofossil biostratigraphy (Fig. 10). The 813  $\delta^{13}C_{carb}$  clearly shows a carbon-isotope positive shift similar to that observed elsewhere.

## 814 **5.6 Basin-wide paleoceanographic reconstruction**

815 In the EAB, a high calcareous nannofossil production and mesotrophic surface-water 816 conditions occurred in the early Valanginian before the Weissert Event interval. This time-817 interval was also characterized by a low energy depositional environment and a mild oxygen-818 depletion (see section 5.1), which could infer stratification of the water-column. However, some 819 differences existed between the different settings due to their respective paleogeographic 820 location (Fig. 9). The nannofossil production was higher at Aït Hamouch with respect to 821 Zalidou in the early Valanginian (Fig. 10). Aït Hamouch was more distal from the shoreline 822 with respect to Zalidou (see section 5.2; Fig. 9). Consequently, the greater depth of deposition 823 at Aït Hamouch and its location farther from the coastline allowed the development of a large 824 and diverse nannofossil community in a thicker photic zone. Also, trophic levels in surface 825 waters were higher at Aït Hamouch than at Zalidou (Fig. 10). Nutrient input in the EAB was 826 generally maintained from both, clastic river input and oceanic upwelling (see sections 5.1 and 827 5.2). During the early Valanginian, the northern part of the EAB (Zalidou section) was less 828 influenced by clastic river influx with respect to the southern part (Aït Hamouch; see Fig. 9). 829 Moreover, Aït Hamouch was potentially fed by nutrients from the nearby upwelling system 830 observed in the Obbay section. Although, the highest nannofossil production was in Hole 416A, 831 being offshore with the thickest photic zone (Fig. 6, and Figs. Supp. 2-3). Fertility levels 832 decreased prior to the Weissert Event interval in the late early Valanginian in all successions 833 (Fig. 10). Nonetheless, a high nannofossil production was maintained by the eurytopic W. 834 barnesiae/fossacincta. The fertility decrease was probably related to a major sea level fall 835 recorded on the central Moroccan margin, and culminated with the expression of the D4 and 836 D5 sedimentary discontinuities (Figs. 3, 10). The sea level fall first impacted the nannofossil 837 community of the distal Aït Hamouch section with the possible ocean-ward migration of the 838 upwelling system away from this site, which should have consequently reduced nutrient input. 839 At the same time, storm deposits and coarse clastic input are recorded at Aït Hamouch (section 840 5.1). Additionally, more turbid surface-water due to riverine-clastic input must have favored 841 the proliferation of the eurytopic W. barnesiae/fossacincta. Afterwards, the entire nannofossil 842 community collapsed at the onset of the Weissert Event and in its lower part, around the early-843 late Valanginian transition (Fig. 10). Offshore, in Hole 416A, the onset of the surface-water 844 fertility decline (upper NK3A and CC3b subzones) and the later collapse of the nannofossil 845 community (NK3B and CC3b-CC4a subzones) seem to have occurred slightly later than in the 846 EAB (Fig. 10). We similarly relate these changes to the maximum drop of sea level expressed 847 by the D4 and D5 discontinuities in the EAB. The collapse is observed later in the distal Hole 848 416A possibly due to the hiatuses related to emergences observed on the platform which are 849 not recorded in a deep setting. The nannofossil production gradually recovered during the 850 Weissert Event in the early late Valanginian and was associated with a rising sea-level (section 851 4.1, Fig. 10). Higher riverine clastic input is recorded in the north (Zalidou) with respect to the 852 south (Aït Hamouch; see Fig. 9). High input of clastics and associated nutrients could explain 853 the early recovery at Zalidou in the nannofossil production and in the abundance of high fertility taxa. At Aït Hamouch, the recovery occurred later, and can be explained by the nutrient input 854 855 that started arriving from the upwelling system. Maximal nannofossil production along with 856 eutrophic surface-waters are recorded only at Zalidou during the upper part of the Weissert 857 Event interval in the late Valanginian (upper *N. peregrinus* to upper *O. nicklesi* ammonite StSz, 858 NK3B and lower CC4a subzones; Fig. 10). Eutrophic surface-waters occurred during maximal

859 flooding and continuous continental influx (section 5.1, Fig. 10). Lastly, a decrease in both the 860 nannofossil production and surface-water fertility occurred after the Weissert Event interval in 861 the rest of the late Valanginian (Fig. 10). Although, trophic levels were higher compared with 862 the early Valanginian. An abrupt increase in the nannofossil production and surface water 863 fertility is observed only at Aït Hamouch during the late Valanginian (C. furcillata StZ, NC4A 864 and upper CC4a subzones) after the Weissert Event. At this time, the depositional environment 865 in Aït Hamouch corresponded to an outer shelf (Fig. 9), possibly with a thick photic zone for 866 the nannofossils to flourish. Whereas, the depositional environment in Zalidou corresponded to 867 a clastic shelf with higher turbidity responsible for a reduced photic zone.

868 The nannoconids in the EAB and Hole 416A, mainly represented by N. steinmannii 869 (narrow-canal form), generally show a distal-to-proximal gradient with highest abundances in 870 Hole 416A followed by Aït Hamouch and Zalidou, successively. Also, the nannoconids were 871 "thriving" during periods of relatively high fertility in surface-waters in all studied successions 872 (Fig. 10). Initially, this is observed in the early Valanginian onshore (N. premolicus StSz to lowermost N. subtenuis StSz in Aït Hamouch and N. subtenuis LSz in Zalidou section, lower 873 874 NK3A and CC3a subzones) and in the lower part of the Valanginian interval offshore. Then, a 875 nannoconid decline is recorded onshore in the early Valanginian (N. neocomiensiformis to 876 lower N. peregrinus StZ, NK3A to lower NK3B subzones, CC3b to lower CC4a subzones), 877 earlier in the southernmost section Aït Hamouch (Fig. 10). Offshore, in Hole 416A, the decline 878 occurred slightly later in the uppermost NK3A to lower NK3B and upper CC3b to lower CC4a 879 subzones. On the central Moroccan margin, this decline can be explained by the drop of sea-880 level that negatively impacted the entire nannofossil community. A fall of sea-level reduces the 881 size of the photic zone (thickness and geographic extent), and can thus explain the observed 882 decrease within the nannoconid population. DSDP Hole 416A was impacted later due to its 883 distal setting and thicker photic zone. Afterwards, the nannoconids recovered in the late 884 Valanginian (upper N. peregrinus StZ, NK3A-NK3B and lower CC4a subzones) during the Weissert Event interval. The nannoconids recovery occurred during periods of relatively high 885 886 surface-water fertility in all studied successions (Fig. 10). These time-intervals were 887 characterized by a high sea-level, which allowed the development of a thicker photic-zone and 888 consequently various habitats to be occupied. Moreover, during the late early and late 889 Valanginian, the occurrence of thin-bedded HCS restricted to shallow marine beds suggests the 890 occurrence of storms of low intensity (section 5.1). This could suggest the absence of 891 stratification in the water-column.

892 Different positions within the photic zone were proposed for narrow- and wide-canal Nannoconus (Table 1). At Aït Hamouch, the proportion of narrow-canal forms (mainly N. 893 894 steinmannii) within the nannoconid population reached its highest value during the sea-level 895 fall interval and coincided with maximum clastic input (Fig. 5). Afterwards, although the clastic input was reduced, the proportion of narrow-canal forms within the nannoconid population 896 897 decreased all along the succession. Then, as shown for Hole 416A, high abundance of N. 898 steinmannii was not associated with a stratified water column but occurred during high clastic 899 input (see section 5.4.3). This implies a shallow nutricline. Our results confirm the assumption 900 of Mattioli et al. (2014) suggesting that N. steinmannii inhabited de-stratified surface waters.



Fig. 10. Synthesis of the paleoceanographic conditions on the central Moroccan margin. Gray shading refers to the Weissert Event interval, for more details see section 5.5. Striped-correlation refers to a possible missing interval in Aït Hamouch (section 5.5). Simplified schemes of nannofossil production, fertility, and nannoconids abundance are comparable between Zalidou and Aït Hamouch since they are based on the accumulation rate results. Whereas, those of Hole 416A are based on the absolute abundance results. Also added are the transgression (T) – regression (R) cycles (sections 5.1 and 5.2).

907 5.7 Integrating the central Moroccan margin in a global paleoceanographic
908 reconstruction

In order to globally reconstruct the fertility in surface-waters during the Weissert Event, the paleoceanographic conditions (reconstructed from calcareous nannofossil data) of the central Moroccan margin are compared with those of other basins. All of the considered studies used the following taxa (small *Zeugrhabdotus* spp., *B. ellipticum*, *D. ignotus*, and *D. lehmanii*) as indicators of high surface-water fertility.

914 In the early Valanginian, high surface-water fertility was suggested in the western 915 Atlantic in DSDP 535 (Gulf of Mexico; Kessels et al., 2006). In the western and northern 916 Altantic (DSDP 534A/603B), Bornemann and Mutterlose (2008) suggested that surface-water 917 productivity was already increasing from the Berriasian-Valanginian boundary prior to the 918 Weissert Event. Whereas, in the Tethys (Vocontian Basin, SE France), generally oligotrophic 919 conditions were recorded in the earliest Valanginian (lowermost NK3 Zone). These 920 oligotrophic conditions were followed by more mesotrophic surface-waters prior to the 921 Weissert Event in the early Valanginian (lower NK3 Zone; Duchamp-Alphonse et al., 2007). 922 In the EAB, and in Hole 416A, the nannofossil production and surface-water fertility were 923 relatively high in the early Valanginian (N. premolicus StSz to N. subtenuis LSz, lower NK3 924 and lower CC3 subzones).

At the early-late Valanginian transition and/or during the incipient Weissert Event, increasing surface-water fertility was observed in several basins. In the Pacific (Site 1149B) and Tethys oceans (Lombardian Basin, northern Italy), higher fertility and primary productivity was demonstrated in the upper NK3 Zone to the NC4A Subzone (Erba et al., 2004; Erba and Tremolada, 2004). In the Vocontian Basin (SE France), increasing calcareous nannofossil fluxes resulted from an increase in surface-water fertility during the Weissert Event, with highest values recorded within the lower NK3B calcareous nannofossil Subzone (upper *B*. 932 Campylotoxus to lower S. verrucosum ammonite zones; Gréselle et al., 2011; Mattioli et al., 933 2014). In the western and northern Atlantic (DSDP 534A and 603B), a further increase in 934 surface-water primary productivity was observed during the NK3A Subzone and just prior to 935 the carbon-isotope excursion (Bornemann and Mutterlose, 2008). Also, in the western Atlantic 936 (DSDP 535), Kessels et al. (2006) interpreted higher surface-water fertility in the early late 937 Valanginian. The central Moroccan margin complements these previous records, and similarly 938 demonstrates increasing surface-water fertility and nannofossil production during the Weissert 939 Event on the south Tethyan margin.

940 In the rest of the late Valanginian, during the end or after the Weissert Event, surface-941 water fertility was observed to globally decrease. In the western Atlantic (DSDP 535), trophic 942 conditions decreased in the late Valanginian (Kessels et al., 2006). In the northern Tethys, a 943 decrease in surface-water fertility was observed in the Lombardian Basin (northern Italy), during the NC4A Subzone when  $\delta^{13}C_{carb}$  is decreasing (Erba and Tremolada, 2004). In the 944 945 Vocontian Basin (SE France), a decrease in both nannofossil fluxes and fertility indicators was 946 observed from the upper part of the Weissert Event (N. peregrinus Zone and upper NK3-lower 947 NC4 subzones; Gréselle et al., 2011; Mattioli et al., 2014). Nonetheless, in the Lower Saxony 948 Basin (northern Germany) surface-water fertility increased during the latest Valanginian 949 (Möller et al., 2020). In the EAB, surface-water fertility and nannofossil production decreased 950 at the top of the Weissert Event and after it, as seen in the most complete section of Zalidou. 951 Although, the fertility remained higher than in the early Valanginian. In the Aït Hamouch 952 section, an abrupt increase in surface-water fertility and nannofossil production related to a 953 thick photic zone in an outer shelf setting was observed in the latest Valanginian (upper C. 954 furcillata StZ, NC4A and upper CC4a subzones).

However, in other studied basins, marine surface-water fertility was stable during the
Valanginian. In the northern-central Atlantic (DSDP 638, NW Spain), high surface-water

957 fertility was observed throughout the Valanginian (Kessels et al., 2006). Further north in NE 958 Greenland, which was a gateway between the Arctic Ocean and the northern Tethys, 959 oligotrophic conditions were observed during the Valanginian (Pauly et al., 2012). In the eastern 960 Tethys (south and east Carpathians), generally oligotrophic surface-waters were observed 961 during the Berriasian-Valanginian interval, supported by stable nannofossil diversity and 962 abundance (Melinte and Mutterlose, 2001). From the available data, the eastern and 963 northernmost Tethys seem to have been characterized by unchanging paleoceanographic 964 conditions during the Valanginian. This might have been linked to the physiographic 965 characteristics of these basins and the associated atmospheric/oceanic circulation currents (see 966 Price et al., 1995; Poulsen et al., 1998).

#### 967 **5.8 The nannoconid decline**

968 The decrease in the nannoconid abundances is known as the "Nannoconid decline", and 969 is globally recorded to start before the Valanginian positive CIE and spans the Weissert Event 970 (Channell et al., 1993; Bersezio et al., 2002; Erba and Tremolada, 2004; Pauly et al., 2012; 971 Möller et al., 2015). In the western Atlantic, the decrease in the nannoconid abundances was 972 observed from the NK3A to NC4A calcareous nannofossil subzones (DSDP 534A; Bornemann 973 and Mutterlose, 2008). In the Vocontian Basin (SE France), a more complicated decline was 974 observed, such that a first decline was recorded from the NK3A to lower NK3B calcareous 975 nannofossil subzones. This was followed by a brief recovery, then by another decline from the 976 upper NK3B Subzone to NC4 Zone (Barbarin et al., 2012). On the central Moroccan margin, 977 the nannoconid decline similarly started prior to the CI positive-shift. It is generally recorded 978 from the NK3A to lower NK3B subzones (CC3b to lower CC4a subzones). Several authors 979 interpreted the decline to be due to enhanced continental-clastic influx and increased 980 nannofossil primary productivity, and possibly under higher oceanic  $pCO_2$  levels (Erba and 981 Tremolada, 2004; Barbarin et al., 2012; Duchamp-Alphonse et al., 2014). Increasing surface-

982 water fertility induces the breakdown of the water-column stratification, and then the decrease 983 in the nannoconid population. Various authors proposed that the distribution of narrow-canal 984 versus intermediate and wide-canal Nannoconus is controlled by fluctuations in nutricline 985 dynamics. In particular, a nutricline rise from the lower to the upper photic zone would lead to 986 the decrease in the narrow-canal forms, these forms are interpreted as deep-dwellers under 987 stratified waters (Erba, 1994, 2004; Duchamp-Alphonse et al., 2014). Since different habitats 988 within the photic zone seem plausible for the big calcifying nannoconids, a deep-dwelling 989 behavior for N. steinmannii can be challenged in this study (see section 5.6) confirming the 990 hypothesis of Mattioli et al. (2014). Furthermore, in the Vocontian Basin, high proportions of 991 narrow-canal forms (mainly represented by N. steinmannii) correlate with high coccolith 992 abundance (NK3B to lower NC4A calcareous nannofossil subzones; Angles section; Duchamp-993 Alphonse et al., 2007, 2014). In addition, the highest proportion of narrow-canal nannoconids 994 are recorded within the Weissert Event in the composite section of Vergol/La Charce, when the 995 values of nannofossil fluxes are the highest (Gréselle et al., 2011; Barbarin et al., 2012). On the central Moroccan margin, the nannoconid decline was probably caused by a sea-level fall that 996 997 gave rise to higher turbidity in the surface-waters. Higher turbidity probably attenuated light 998 penetration, reducing the photic zone and the development of the total nannoconid population. 999 However, enhanced clastic input and associated nutrients led to a shallow nutricline that did not 1000 particularly impact N. steinmannii.

1001 Moreover, the nannoconid recovery was observed to coincide with the high  $\delta^{13}C_{carb}$ 1002 values in the Tethys (Vocontian Basin, France) and in the western and northern Atlantic (DSDP 1003 534A and 603B; Erba and Tremolada, 2004; Bornemann and Mutterlose, 2008; Barbarin et al., 1004 2012). This agrees with observations in the EAB, since nannoconids recover during the 1005 Weissert Event when the  $\delta^{13}C_{carb}$  is high and under conditions of both, high nannofossil production and surface-water fertility. This suggests that nannoconids do not particularlydisfavor such surface-water conditions.

#### 1008 **6.** Conclusions

The quantitative analyses of calcareous nannofossil assemblages from two onshore sections belonging to the Essaouira-Agadir Basin (SW Morocco) and one offshore succession (DSDP Hole 416A, east Atlantic Ocean) allowed to reconstruct the paleoceanographic conditions on a proximal-distal transect across the Valanginian Weissert Event. This reconstruction was done in a well-constrained ammonite (for onshore sections only) and nannofossil biostratigraphic and chemostratigraphic (carbon-isotopes) framework, along with a detailed sedimentological characterization.

1016 The Weissert Event interval is identified in the onshore sections from the 1017 Karakaschiceras inostranzewi to upper Neocomites peregrinus Standard Zone (StZ) or from 1018 the upper NK3A to upper NK3B, and upper CC3b to "middle" CC4a calcareous nannofossil 1019 subzones. The Valanginian stage demonstrated variations in surface-water fertility and 1020 nannofossil production. A general paleoceanographic scheme is developed: (1) mesotrophic 1021 and favorable surface-water conditions for the nannofossil community prevailed before the 1022 Weissert Event time interval in the early Valanginian. (2) Surface-water fertility levels 1023 decreased prior to the Weissert Event interval, and a high nannofossil production was 1024 maintained by the eurytopic Watznaueria barnesiae/fossacincta. (3) A collapse of the 1025 nannofossil community during the onset of the Weissert Event interval at the early-late 1026 Valanginian transition was caused by a major sea-level fall. (4) A recovery followed in the early 1027 late Valanginian linked to the rising sea-level. (5) The recovery was followed by maximal 1028 nannofossil production associated with high surface-water fertility, recorded in the northern 1029 section during the upper part of the Weissert Event interval in the late Valanginian; or in the 1030 southern section after the Weissert Event interval in the latest Valanginian. Finally, (6) a decrease in both surface-water fertility and nannofossil production occurred after the Weissert
Event interval, although they remained high compared with the early Valanginian. Moreover,
the "nannoconid decline" occurred from the early Valanginian to the early late Valanginian.
The nannoconid recovery occurred during conditions of high sea-level and high surface-water
fertility.

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