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1 **Isotopic record of Aptian third-order sea-level trends in platform margin carbonates:**  
2 **implications for sequence stratigraphic analysis**

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13 **ABSTRACT**

14 In ancient carbonate systems, establishing relationships among sea-level fluctuations, carbonate  
15 factory productivity and carbonate geochemistry is challenging due to complex depositional and  
16 diagenetic overprinting. The Aptian platform carbonate succession from the western Maestrat  
17 Basin, in Spain, serves as an ideal example for potentially linking these processes, particularly  
18 establishing the relationships between isotopic records ( $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$ ) and third-order sea-level  
19 trends with implications for sequence stratigraphy. This succession is biostratigraphically well-  
20 constrained and comprises two depositional sequences that were controlled by a major relative  
21 sea-level fall and a subsequent rise. These depositional sequences exhibit stratal terminations and  
22 stacking patterns, enabling the establishment of a well-defined sequence stratigraphic framework  
23 comprising four systems tracts and their key bounding stratigraphic surfaces. The analytical

24 results reveal that both  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  values outline distinct temporal trends, which can be  
25 correlated with specific third-order stages of relative sea-level fluctuations. The transgressive and  
26 highstand systems tracts exhibit the most positive  $\delta^{13}\text{C}$  values (up to +5‰) and the least negative  
27  $\delta^{18}\text{O}$  values (up to -1.8‰). This range, similar to values of carbonates in equilibrium with  
28 Cretaceous seawater,  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from +2‰ to +5‰ and from -2‰ to -5‰,  
29 respectively, likely reflects the marine influence on the isotope values during the stage of high  
30 relative sea level. In contrast, the forced regressive and lowstand intervals exhibit less positive  
31  $\delta^{13}\text{C}$  values (reaching +0.5‰) and significantly more negative  $\delta^{18}\text{O}$  values (around -6.1‰),  
32 interpreted as the influence of soil-derived organic matter and meteoric waters, respectively,  
33 during stages of lower relative sea level. Furthermore, the sedimentary succession records a  
34 decreasing trend of  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values towards the sequence boundary, which marks the  
35 lowest point of relative sea level. This study underscores the potential of carbonate platforms to  
36 record geochemical signals that can be directly correlated with different third-order stages of  
37 relative sea-level variations. In addition, this framework allows for linking and predicting the  
38 potential fluid-rock interaction processes in each systems tract. The proposed approach could  
39 offer a predictive framework for characterizing carbonate reservoirs and other carbonate  
40 platform successions with poorly defined sequence-stratigraphic frameworks elsewhere.

41 **Keywords:** Stable Isotopes; Geochemistry; Sequence Stratigraphy; Sea-level Fluctuation;  
42 Carbonates

## 43 INTRODUCTION

44 Third-order relative sea-level fluctuations (sensu Vail et al., 1991) play a pivotal role in  
45 controlling physical and ecological accommodation and influencing the nature and architecture  
46 of depositional facies in platform carbonates (Eberli, 2000; Anselmetti et al., 2000). The

47 interplay between sedimentation and accommodation defines the sequence stratigraphic  
48 framework and exerts control over the extent and distribution of depositional and diagenetic  
49 alterations that impact the sedimentary succession during and after deposition (Schlager, 2005;  
50 Catuneanu et al., 2009). Given that many factors influencing carbonate geochemistry, such as  
51 pore water chemistry, depositional setting, and burial/exposure, are also sensitive to changes in  
52 relative sea level, isotope geochemistry can be directly linked to sequence stratigraphy (Tucker,  
53 1993; Weissert et al., 1998; Spezzaferri et al., 2002; Burla et al., 2008; Mehrabi et al., 2022).  
54 Correlations between carbon isotopic values and changes in relative sea level, with implications  
55 for sequence stratigraphy, have already been noticed in contributions studying pelagic and  
56 shallow-water carbonates (Föllmi et al., 1994; Vahrenkamp, 1996; Eberli, 2000; Saltzman, 2002;  
57 Spezzaferri et al., 2002; Immenhauser et al., 2003; Krull et al., 2004). However, in some case  
58 studies, including the Holocene to Early Miocene record of the Bahamas,  $\delta^{13}\text{C}$  values of platform  
59 and periplatform sediments were unrelated to synchronous shifts in the  $\delta^{13}\text{C}$  of the open oceans.  
60 This was interpreted to indicate that  $\delta^{13}\text{C}$  trends of these sediments did not correlate with the  
61 ocean-climatic system (or the global  $\delta^{13}\text{C}$  signal) over this time period (Swart and Eberli, 2005).  
62 In the Arabian Gulf, however, the  $\delta^{13}\text{C}$  patterns of Lower Cretaceous shallow-water carbonates  
63 were controlled by third-order fluctuations in relative sea level and correlated with isotopic  
64 profiles reported elsewhere in Tethyan pelagic limestones of the same age (Föllmi et al., 1994;  
65 Vahrenkamp, 1996). This led to the suggestion that the isotopic proxy was an effective tool for  
66 characterizing relative sea-level fluctuations in the lower Cretaceous record of the Arabian  
67 (Tethyan?) region (Vahrenkamp, 1996). Similar conclusions, suggesting comparable  $\delta^{13}\text{C}$  shifts  
68 in carbonate lithofacies from different water depths, were obtained in Mississippian carbonates  
69 from western USA (Saltzman, 2002, 2003). The later authors placed the carbonate  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$

70 variations in a sequence stratigraphic framework to evaluate the presence of exposure surfaces  
71 and to distinguish between primary and secondary stable isotope patterns (Saltzman, 2002).

72 Despite this good correlation between isotope stratigraphy and relative sea-level fluctuations in  
73 carbonate environments, linking chemostratigraphic data to sequence stratigraphic concepts have  
74 proven effective mostly in siliciclastic systems (Haq et al., 1987; Posamentier et al., 1992; Morad  
75 et al., 2000, 2010, 2013; Al-Ramadan et al., 2005). This is due to the wide variety and  
76 complexity of carbonates resulting from the interaction among relative sea-level changes, local  
77 tectonics, climate, type of biota, diagenesis, and heterogeneous mineralogy and texture (Gischler  
78 et al., 2009; Pomar and Haq, 2016). Besides, recognizing or interpreting sequence stratigraphic  
79 attributes in ancient carbonate environments can also be challenging due to poor exposure  
80 conditions, variable rates of erosion and sedimentation, and superposition of different  
81 stratigraphic surfaces (Immenhauser et al., 2002; Christ et al., 2012). In such cases, data gaps are  
82 addressed by incorporating interpreted surfaces that introduce greater uncertainty to the  
83 interpreted stratigraphic model (Catuneanu et al., 2009).

84 Carbonate platform margins represent the best setting to estimate fluctuations in relative sea  
85 level and to evaluate the global versus local controls on platform evolution (Pomar et al., 2005).

86 A modern setting, often applied as analogue of ancient platform carbonates, comes from the  
87 Great Bahama Bank (GBB) and its subsurface sediments (Eberli, 2000; Anselmetti et al., 2000;  
88 Eberli et al., 2002; Spezzaferri et al., 2002; Reuning et al., 2006; Reijmer et al., 2009; Murray  
89 and Swart, 2017; Murray et al., 2021). However, the question that commonly arises is whether  
90 the results, in terms of facies distribution, mineralogy and geochemistry, are applicable to all  
91 geological time periods (Swart et al., 2009). In the case of ancient carbonates, the Guadalupian  
92 outcrops of the Permian Basin (USA) provide a classic example of a large and continuous

93 exposure illustrating the architecture and sequence stratigraphy of a self-to-basin section (Harris  
94 and Saller, 1999).

95 In this contribution, we focus on the Aptian “Las Mingachas” section in the western marginal  
96 part of the Maestrat rift basin (Iberian Chain) in Spain. This section stands out as one of the very  
97 few outcrops worldwide that preserves stacking patterns and stratal terminations that permit one  
98 to define sequence stratigraphic surfaces that bound different systems tracts (Bover-Arnal et al.,  
99 2009, 2022). In general, Las Mingachas section exhibits stratigraphic continuity, with its age  
100 calibrated using Sr-isotope stratigraphy, as well as ammonite, orbitolinid, and rudist  
101 biostratigraphies (Moreno-Bedmar et al., 2010; Bover-Arnal et al., 2016). Furthermore, Las  
102 Mingachas section offers an exceptionally well-preserved platform-to-basin transition, showing  
103 sedimentary evidence of the influence of third-order relative sea-level fluctuations, including a  
104 major sea-level fall and a subsequent rise, on carbonate sedimentation across the system (Bover-  
105 Arnal et al., 2009, 2022). Thus, drawing upon this well-established sequence stratigraphic  
106 framework, this contribution provides carbonate geochemical measurements at Las Mingachas  
107 section aiming to: i) test if the stable carbon and oxygen isotopes of the Aptian platform  
108 carbonates are in line with the interpreted sequence-stratigraphic framework and the reported  
109 third-order fluctuations in relative sea level, especially considering the nature of the major sea-  
110 level fall, interpreted to be of global significance with a dominant eustatic contribution; ii) verify  
111 if this geochemical record enables the identification of the attributes of a four systems tracts-  
112 based sequence stratigraphic framework (Hunt and Tucker, 1992), which is a complete  
113 framework not commonly preserved in the stratigraphic record; and (iii) to corroborate if the new  
114 provided data allows us to discern between divergent interpretations concerning part of the  
115 stratigraphic framework reported in the study area (Bover-Arnal et al., 2009, 2022; Peropadre et

116 al., 2013; Pomar and Haq, 2016; Pomar, 2020). Thus, this case study exemplifies an ideal  
117 scenario to assess the robustness of stable isotope proxies in identifying Aptian third-order sea-  
118 level trends and geochemically characterizing the complete “four-systems-tract” sequence  
119 stratigraphic model of Hunt and Tucker (1992).

## 120 **GEOLOGICAL SETTING**

121 The studied Aptian carbonate succession is located in the eastern part of the Iberian Chain in  
122 Spain (Fig. 1), which is an intraplate fold-and-thrust belt that resulted from the Alpine (late  
123 Eocene-early Miocene) tectonic inversion of the Mesozoic Central Iberian Rift System (Salas &  
124 Casas, 1993; Salas et al., 2001; Guimera et al., 2004). Late Jurassic and Early Cretaceous rifting  
125 stages in Iberia were linked to the opening of the North Atlantic Ocean and the Bay of Biscay,  
126 respectively. These two rifting stages gave rise to major NW-SE, E-W and NE-SW normal faults  
127 that controlled the formation of rapidly subsiding basins, including the Maestrat Basin, which is  
128 the focus of this study (Salas et al., 2001). The selected outcrops are located in a western  
129 marginal area of the Maestrat Basin known as the Galve sub-basin (Fig. 1) (Liesa et al., 2006;  
130 Salas et al. in Martín-Chivelet et al., 2019). The stratigraphic interval analyzed corresponds to a  
131 platform-to-basin transition area known as "Las Mingachas" (Figs. 1 & 2) and spans the late  
132 early-earliest late Aptian time slice (Moreno-Bedmar et al., 2010; Bover-Arnal et al., 2016,  
133 2022). This marine succession postdates the early Aptian oceanic anoxic event (OAE1a) (Bover-  
134 Arnal et al., 2010). The Las Mingachas section comprises the upper part of the long-term T-R  
135 Sequence II and the lower part of T-R Sequence III described in Bover-Arnal et al. (2010, 2016).  
136 The following description of facies and the sequence stratigraphic framework is based on Bover-  
137 Arnal et al. (2009, 2022).

### 138 *Sequence Stratigraphic Framework*

139 The transition zone from platform to basin at Las Mingachas was interpreted within a sequence  
140 stratigraphic framework consisting of four systems tracts, encompassing two depositional  
141 sequences: A and B. Depositional Sequence A includes a Highstand Systems Tract (HST) and a  
142 Forced Regressive Wedge Systems Tract (FRWST). In contrast, Depositional Sequence B is  
143 composed of a Lowstand Prograding Wedge Systems Tract (LPWST), a Transgressive Systems  
144 Tract (TST), and the subsequent HST (Figs. 2 & 3).

#### 145 **HST (Depositional Sequence A).---**

146 The first HST consists of a carbonate platform comprising corals and rudists belonging to the  
147 Villarroya de los Pinares Formation, which shows aggrading and subsequent prograding patterns.  
148 Towards the basin, located in the SW of the study area, the platform top limestones pass to slope  
149 deposits, which exhibit downlapping stratal terminations and laterally pass into the marls of the  
150 Forcall Formation (Fig. 2). The downlap surface separates the upper limestones with corals and  
151 rudists from lower, deeper basinal marls, thus representing the maximum flooding surface (MFS)  
152 of Depositional Sequence A (Bover-Arnal et al., 2009). The basinal marls, equivalent to the  
153 platform's top limestones, are interpreted as part of the highstand normal regressive unit, whereas  
154 the marly deposits below the MFS are considered a transgressive succession. These basinal  
155 marly deposits correspond to the Forcall Formation.

#### 156 **FRWST (Depositional Sequence A).---**

157 The FRWST is represented by a high energy regressive wedge constituted by cross-bedded  
158 grainstones occurring basinwards, at the toe of highstand slopes. These calcarenite deposits  
159 belong to the Villarroya de los Pinares Formation, and are made up of an autochthonous  
160 carbonate factory dominated by oysters, and of allochthonous lithoclasts that resulted from

161 erosion of the exposed carbonate platform during a sea-level fall that was estimated to be of at  
162 least 60 meters (Bover-Arnal et al., 2009). The lower boundary of the calcarenite deposits, i.e.,  
163 the basal surface of forced regression (BSFR), is represented by a sharp surface separating the  
164 underlying basinal highstand marls of the Forcall Formation from the overlying shallower forced  
165 regressive wedge. In turn, the upper boundary of the regressive wedge is represented by the  
166 sequence boundary (SB), which separates depositional sequences A and B and marks the lowest  
167 point of relative sea level. The SB is characterized by a subaerial unconformity that becomes  
168 younger seawards, passing to a marine correlative conformity (CC) over the FRWST (Fig. 2).

#### 169 **LPWST (Depositional Sequence B).---**

170 The LPWST consists of a small carbonate platform dominated by corals and rudists that shows a  
171 progradational stacking pattern followed by aggradation (Fig. 2). The lowstand platform, which  
172 developed during the subsequent base-level rise, is also part of the Villarroya de los Pinares  
173 Formation. The platform top facies pass into slope deposits basinwards. The upper boundary of  
174 the lowstand unit is marked by the transgressive surface (TS), which corresponds to a  
175 hardground with *Lithophaga* borings and iron mineralization. Landwards, the TS superimposes  
176 the SB and therefore, the subaerial unconformity that separates the two depositional sequences  
177 becomes a composite sequence boundary (i.e., SB + TS).

#### 178 **TST (Depositional Sequence B).---**

179 In Las Mingachas, the TST corresponds to retrograding platform top carbonates with rudists and  
180 corals that onlap the composite sequence boundary between Depositional Sequences A and B  
181 (SB + TS) towards the land (Fig. 2). The backstepping limestones change basinwards into marly  
182 deposits of the lower part of the Benassal Formation, and also upwards in the succession after

183 drowning of the Villarroya de los Pinares platform. The maximum flooding surface (MFS) of  
184 Depositional Sequence B is interpreted to be located at the top of the thickest marl bed of the  
185 lower part of the Benassal Formation, coinciding with the downlap surface of overlying coral-  
186 bearing deposits.

### 187 **HST (Depositional Sequence B).---**

188 The succeeding highstand deposits also belong to the Benassal Formation (Fig. 2) and are  
189 characterized by platform carbonates with corals, rudists and nerineid gastropods. Above the  
190 MFS of Depositional Sequence B, the basal part of this highstand unit is characterized by a  
191 marly horizon with irregular and massive corals, alternating with limestone beds of carbonate  
192 shed from platform top settings.

### 193 **METHODOLOGY**

194 This study includes stable isotope measurements ( $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values) of the Aptian carbonate  
195 platform cropping out at Las Mingachas section. Previous research established the facies  
196 architecture, as well as the paleoenvironmental and sequence stratigraphic interpretation,  
197 including the reconstruction of the relative sea-level stages based on stratigraphic data (Bover-  
198 Arnal et al., 2009, 2022). This previous research guided the selection of outcrops and samples to  
199 conduct geochemical analyses. The selected outcrop corresponds to the platform-to-basin  
200 transition area of Las Mingachas section, where the sedimentary succession includes four  
201 systems tracts encompassing two depositional sequences (Fig. 2). In this area, a representative  
202 log was measured, and facies variations, along with key stratigraphic surfaces, were mapped on a  
203 panoramic view and correlated with the surfaces described in previous studies of the area (Fig.  
204 3). Ninety samples were collected in the field following two sampling strategies for comparison.

205 “LM samples” cover laterally the full platform to basin transition area shown in Figures 2 and 3  
206 (i.e., covering the entire platform and throughout all systems tracts), whereas “vertical log  
207 samples” were taken vertically next to the stratigraphic log (Fig. 3). Thus, collected samples  
208 cover along dip and along strike the full outcrop from the uppermost slope and towards the basin,  
209 where the subaerial unconformity, representing the composite sequence boundary, passes to its  
210 marine correlative conformity (Fig. 2A).

211 LM samples were prepared for stable isotope measurements ( $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$ ) of bulk carbonates  
212 and separate carbonate components (micritic matrix, grains, and cements), whereas, in vertical  
213 log samples, only bulk carbonates were analyzed. In total, this study provides fifty-nine analyses  
214 of bulk carbonates and thirty-one analyses of separate carbonate components.

215 Bulk carbonates and separate carbonate components (micritic matrix, grains, and cements) of  
216 LM samples were sampled using a microdrill to extract carbonate powder (n=49). Powdered  
217 samples (50-60  $\mu\text{g}$ ) were reacted at 70°C with 102–105% phosphoric acid before  $\text{CO}_2$  gas was  
218 liberated to the automated purification line. Then, the product  $\text{CO}_2$  was purified using cryogenic  
219 traps to eliminate contaminant gases, such as water vapor and non-condensable gases produced  
220 during the acid digestion of the carbonate samples. Purification was performed automatically  
221 using a Kiel IV carbonate device. The  $\text{CO}_2$  gas released was analyzed with a Thermo Scientific  
222 MAT 253+ mass spectrometer at the Stable Isotope Laboratory, King Fahd University of  
223 Petroleum and Minerals (Dhahran, Saudi Arabia) with an overall precision of 0.08‰ for  $\delta^{18}\text{O}$   
224 and 0.04‰ for  $\delta^{13}\text{C}$ , as measured by blind sample replicates. The oxygen and carbon isotope  
225 data are presented in  $\delta$  notation relative to the Vienna Pee Dee Belemnite (VPDB) standard via  
226 standard reference materials obtained from the US National Institute of Standards and  
227 Technology and the International Atomic Energy Agency.

228 The measurements on a mass spectrometer drift over time for several reasons, including ion  
229 source performance (D'Autry et al., 2010) or the presence of contaminants. The drift was  
230 corrected by plotting the raw  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  data against the internal laboratory carbonate  
231 standard (KIS; KFUPM Iceland Spar) with the accepted isotopic values of 2.25‰ and -6.23‰  
232 (VPDB) for  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$ , respectively.

233 Bulk vertical log samples, encompassing limestones, marly limestones, and marls were also  
234 analyzed for  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  measurements (n=41). For each analysis, 60-70  $\mu\text{g}$  of material was  
235 used for limestones, while marls and marly limestones were analyzed with quantities ranging  
236 from 100-1000  $\mu\text{g}$ . To extract carbonate powder from limestones and avoid large skeletal  
237 components, corals, microencrusters or diagenetic calcite veins, a microdrill was employed. The  
238 samples underwent treatment with  $\text{H}_3\text{PO}_4$  (100%) at 70°C, and the evolved  $\text{CO}_2$  was analyzed  
239 using a Thermo Finnigan MAT-252 stable isotope ratio mass spectrometer at the Scientific and  
240 Technological Centers of the University of Barcelona. The isotope results are expressed in ‰  
241 relative to the VPDB standard, with a precision of  $\pm 0.03$  for  $\delta^{13}\text{C}$  and  $\pm 0.06$  for  $\delta^{18}\text{O}$ .

## 242 RESULTS

### 243 *Sedimentary succession logged and sampled*

244 The analyzed stratigraphic section is representative of the two identified depositional sequences,  
245 A and B, and the three sedimentary formations: Forcall, Villarroya de los Pinares and Benassal  
246 (Fig. 3). This section commences with alternating marls, marly limestones and limestones, which  
247 are attributed to the uppermost part of the Forcall Formation (Fig. 3 & 4A). These marly deposits  
248 between meters 0 and 15 represent the highstand deposits of Depositional Sequence A (Bover-  
249 Arnal et al., 2009, 2022). The marly limestones and limestones exhibit mudstone to packstone

250 textures and contain skeletal components such as *Palorbitolina lenticularis* (Fig. 4B), calcareous  
251 dinoflagellate cysts, textularids, miliolids, ammonoids, sponge spicules, brachiopods, and  
252 echinoids, whereas the non-skeletal components mainly consist of peloids and quartz grains.  
253 Towards the north (i.e., landwards, in higher topographic areas), this marly interval passes to an  
254 aggrading to prograding carbonate platform that exhibits downlap stratal terminations. These  
255 highstand platform carbonates are attributed to the Villarroya de los Pinares Formation and  
256 consist of limestones showing floatstone to rudstone textures made up of rudist and corals in life  
257 position.

258 Above the marly deposits of the Forcall Formation, there are packstone to grainstone limestones,  
259 displaying low-angle crossbedding. These rocks are characterized by the occurrence of conical  
260 *Orbitolinopsis* (Fig. 4C), calcareous algae *Permocalculus* and *Marinella lugeoni*, fragments of  
261 corals, and intraclasts. These deposits of the Villarroya de los Pinares Formation, between metres  
262 15 and 19, constitute the FRWST of Depositional Sequence A (Fig. 3; see Bover-Arnal *et al.*,  
263 2009, 2022). At meter 19, a sharp stratigraphic surface marks the top of the FRWST. This  
264 surface, which is downlapped by lowstand slope deposits has been interpreted as the lowest point  
265 of relative sea level, marking the sequence boundary (SB) (Bover-Arnal *et al.*, 2009, 2022).  
266 Landwards, the SB is truncated and exhibits paleokarst features and thus, it is classified as a  
267 subaerial unconformity (SU). Within the uppermost slope and towards the basin, the SB does not  
268 show evidence of subaerial exposure and thus, it is classified as a marine correlative conformity  
269 (CC). Between meters 19 and 43 the Villarroya de los Pinares Formation succession is made up  
270 of an alternation between marls, which contain scleractinian colonies in growth position, and  
271 limestones with rudstone to floatstone textures with rudist shells, corals, and *Permocalculus* (Fig.  
272 4D). Limestones show a prograding and then an aggrading pattern and exhibit downlap stratal

273 terminations. Consequently, Bover-Arnal *et al.* (2009, 2022) interpreted this stratigraphic  
274 interval as a LPWST belonging to Depositional Sequence B (Fig. 3).

275 The upper part of the Villarroya de los Pinares Formation, between meters 43 and 46,  
276 corresponds to limestones showing a floatstone to rudstone texture (Fig. 4E). These strata are  
277 dominated by colonial corals, rudists such as *Toucasia carinata* and *Polyconties hadriani*,  
278 *Chondrodonta*, *Permocalculus*, and nerineid gastropods. This interval represents the early TST  
279 of Depositional Sequence B (Figs. 2 & 3; Bover-Arnal *et al.*, 2009, 2022), and exhibits a  
280 backstepping geometry that is topped by a hardground with borings made by lithophagid  
281 bivalves and ferruginous stains (see Figs. 8E-F in Bover-Arnal *et al.*, 2022). Above, from meter  
282 46 until the end of the measured section, a marly succession characterized by the presence of  
283 large gastropods of the species *Trochoneritas gigas* overlies the platform carbonates of the  
284 Villarroya de los Pinares Formation. This marly stratigraphic interval was interpreted as the  
285 upper part of the TST of Depositional Sequence B (Bover-Arnal *et al.*, 2009, 2022) and  
286 constitutes the lower part of the Benassal Formation (Figs. 2 & 3). Given the absence of  
287 sedimentary or biotic indicators of bathymetry, the maximum flooding surface (MFS) of  
288 Depositional Sequence B was placed at the top of the thickest marl unit within the TST. Above  
289 this surface, the marly deposits evolve into slope facies, characterized by marls containing  
290 isolated coral colonies. The slope deposits pass upwards into platform top carbonates made up of  
291 floatstones and rudstones with corals, rudists, and nerineid gastropods. These slope to platform  
292 top facies represent the HST of Depositional Sequence B (Bover-Arnal *et al.*, 2009).

### 293 *Carbon and oxygen isotopes*

294 Stable carbon and oxygen isotopic values of the studied Aptian platform carbonates vary widely  
295 from +0.5‰ to +5.3‰ and from -6.2‰ to -1.8‰, respectively (Table 1 and Figs. 5 & 6).

296 Overall, the  $\delta^{13}\text{C}$  curve displays a distinctive pattern characterized by two intervals of persisting  
297 positive carbon isotopic values ( $+4\text{‰} \pm 1\text{‰}$  approx.) separated by a negative carbon isotopic  
298 excursion of around 4‰. Likewise, with a greater dispersion of values, the  $\delta^{18}\text{O}$  curve shows a  
299 negative oxygen isotopic excursion, reaching -6‰, separating two intervals of less negative (but  
300 widely varying) oxygen values (Figs. 5 & 6).

301 Furthermore, despite the slight overlap between some samples, each studied systems tract  
302 exhibits a distinctive range of isotopic values. This range does not vary laterally within the same  
303 systems tract (i.e., we do not observe along strike isotopic patterns within the same genetic type  
304 of deposits). Instead, the isotopic trends vary vertically (i.e., isotopic values increasing or  
305 decreasing from one systems tract to another), as explained below.

#### 306 **HST (Depositional Sequence A).---**

307 In LM samples, bulk carbonates from the first HST exhibit  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values ranging from  
308  $+2.6\text{‰}$  to  $+4.7\text{‰}$  and from  $-5.3\text{‰}$  to  $-3.2\text{‰}$ , respectively, whereas carbonate components vary in  
309  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from  $+3.6\text{‰}$  to  $+4.7\text{‰}$  and from  $-5.7\text{‰}$  to  $-4.4\text{‰}$ , respectively (Table 1 and  
310 Figs. 5 & 6). In vertical log samples, these HST bulk carbonates show  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from  
311  $+3.2\text{‰}$  to  $+5.1\text{‰}$  and from  $-3.7\text{‰}$  to  $-3.1\text{‰}$ , respectively.

#### 312 **FRWST (Depositional Sequence A).---**

313 Compared to the HST, the FRWST interval shows an overall trend towards less positive and  
314 more negative carbon and oxygen isotope values, respectively (Table 1 and Figs. 5 & 6). That is,  
315 LM bulk carbonate samples from this interval yield average  $\delta^{13}\text{C}$  values of  $+0.9\text{‰}$  and  $\delta^{18}\text{O}$   
316 values of  $-5.7\text{‰}$ , whereas carbonate components have  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from  $+0.5\text{‰}$  to  
317  $+3.5\text{‰}$  and from  $-6.0\text{‰}$  to  $-5.6\text{‰}$ , respectively. These values are within the range of values

318 obtained in FRWST bulk carbonates from vertical log samples, where  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values vary  
319 from +0.8‰ to +4.2‰ and from -6.0‰ to -3.6‰, respectively. In the  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  profiles of  
320 the FRWST interval, both carbon and oxygen isotopic values generally decrease vertically  
321 (upwards) from the lower (BSFR) to the upper (CC) stratigraphic surface (Figs. 5 & 6). This  
322 means that carbon and oxygen values progressively shift from around +4.0‰ and from -4.0‰,  
323 respectively in the BSFR, to +0.5‰ and -6‰, respectively, in the CC.

#### 324 **LPWST (Depositional Sequence B).---**

325 The LPWST has a similar range of values to the FRWST interval (Table 1 and Figs. 5 & 6). That  
326 is, bulk carbonates in LM samples display  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from +1.7‰ to +4.1‰ and from  
327 -5.2‰ to -4.7‰, respectively, whereas carbonate components show  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from  
328 +1.2‰ to +4.5‰ and from -6.1‰ to -3.7‰, respectively. Again, bulk vertical log samples in  
329 LPWST yield similar  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values, between +1.3‰ and +4.1‰ and between -6.1‰ and  
330 -3.5‰, respectively. Overall,  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values progressively increase vertically from the  
331 lower to the upper surface of the LPWST interval, that is, from the CC towards the TS (Figs. 5 &  
332 6).

#### 333 **TST (Depositional Sequence B).---**

334 LM Bulk carbonate samples from the TST exhibit  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from +2.5‰ to +5.1‰  
335 and from -5.9‰ to -3.7‰, respectively, whereas carbonate components vary in  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$   
336 values from +3.5‰ to +5.3‰ and from -6.2‰ to -2.3‰, respectively (Table 1 and Figs. 5 & 6).  
337 Similar values have been measured in bulk vertical log samples for TST, which yield  $\delta^{13}\text{C}$  and  
338  $\delta^{18}\text{O}$  values between +1.3‰ and +5.2‰ and between -5.7‰ and -2.4‰, respectively.

#### 339 **HST (Depositional Sequence B).---**

340 This carbonate interval has only been sampled in LM samples (Table 1 and Figs. 5 & 6). The  
341 bulk carbonate sample shows  $\delta^{13}\text{C}$  values of +3.6‰ and  $\delta^{18}\text{O}$  values of -1.8‰, whereas  
342 carbonate components have  $\delta^{13}\text{C}$  values of +4.4‰ and  $\delta^{18}\text{O}$  values of -3.9‰.

## 343 DISCUSSION

### 344 *Linking third-order sea-level trends to carbonate stable isotopes*

345 The sequence stratigraphic framework of the Aptian platform margin studied here was arranged  
346 into two depositional sequences (A and B) that formed as a result of a major fall and a  
347 subsequent rise in relative sea level, respectively (Fig. 2; Bover-Arnal et al., 2009, 2022).  
348 Carbonate production and accumulation during the late early-early late Aptian was closely linked  
349 to these relative sea-level fluctuations (Bover-Arnal et al., 2009; Gratacós et al., 2021).  
350 Consequently, the isotopic patterns recorded in these platform carbonates may be used to  
351 reconstruct not only the depositional setting but also the superposed changes in relative sea level  
352 and its associated fluid-rock interactions (e.g., Swart & Eberli, 2005; Vahrenkamp, 1996).  
353 The two observed intervals in the  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  profiles showing the most positive carbon  
354 values (+4‰  $\pm$  1‰ approx.) and the less negative (and less varying) oxygen values (-5.5‰ to -  
355 3‰) coincide with two specific stages of relative sea-level high: HST of Depositional Sequence  
356 A and TST-HST of Depositional Sequence B (Figs. 5 & 6). These more positive  $\delta^{13}\text{C}$  and less  
357 negative  $\delta^{18}\text{O}$  values are within the range of carbonates in equilibrium with Cretaceous seawater,  
358 which typically display  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values ranging from around +2‰ to +5‰ and from -2‰  
359 to -5‰, respectively (Veizer et al., 1999). The correspondence between isotopic values recorded  
360 in highstand and transgressive intervals, and those typical values of contemporaneous seawater is  
361 interpreted to reflect the marine influence on the carbonate isotope values during the maximum

362 relative sea-level high (D'Argenio et al., 2004; Föllmi et al., 2006). During periods of high  
363 relative sea level, the continental shelf is flooded and the carbonate factory produces carbonate at  
364 its maximum activity, leading to high amounts of organic carbon burial (Magaritz, 1989). In turn,  
365 the increase in carbon burial rates enhances removal of light carbon from the ocean-atmosphere  
366 system by burial of  $^{13}\text{C}$ -depleted organic matter (Jenkyns, 1996; Gröcke et al., 1999; Jarvis et al.,  
367 2001, 2002). A high rate of carbonate production during the HST of Depositional Sequence A is  
368 evidenced by the abundance of lime mud present, the diversity of carbonate producers, including  
369 corals and rudists in life position and nerineid gastropods, and the metric thickness of beds that  
370 characterize this systems tract (Bover-Arnal et al., 2009). The high carbonate accumulation rate  
371 during the slowing sea-level rise, typical of the HST, notably reduced accommodation in the late  
372 HST. This interpretation is supported by the observed prograding stacking pattern exhibited by  
373 the HST. In contrast, during the TST of Depositional Sequence B, the carbonate accumulation  
374 rate was outpaced by the rate of relative sea-level rise. As a result, the carbonate platform,  
375 displaying a retrograding pattern, was eventually drowned. This is evidenced by the sedimentary  
376 succession passing upwards into marly deposits (Fig. 3). The observed geochemical behavior  
377 agrees with similar conclusions elsewhere suggesting that positive carbon isotope excursions are  
378 associated with maximum relative sea-level highs and vice versa (D'Argenio et al., 2004; Swart  
379 and Eberli, 2005; Föllmi et al., 2006).

380 The second observed trend in the isotopic profiles shows a decrease in  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values in  
381 the FRWST interval of Depositional Sequence A (i.e., coinciding with the stage of base-level  
382 fall). In this regressive interval,  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values progressively decrease from the BSFR  
383 towards the SB (correlative conformity), shifting from +4‰ to around 0‰ and from -4‰ to -  
384 6‰, respectively (Figs. 5 & 6). Therefore, this lowering isotopic values during the falling

385 relative sea level is here interpreted to reflect the gradual exposure of the platform top to  
386 subaerial conditions (Figs. 2-3 & 5-7). Through water-rock interaction under subaerial  
387 conditions: (i) the  $\delta^{18}\text{O}$  values are usually lowered by isotopically light meteoric fluids, which  
388 have lower  $\delta^{18}\text{O}$  values than seawater values, whereas the  $\delta^{13}\text{C}$  values are lowered by the  
389 increasing influence of  $\text{CO}_2$  derived from the soil zone and the oxidation of organic matter  
390 (Allan and Matthews, 1977; Popp et al., 1986; Sattler et al., 2005); and (ii) the previously  
391 deposited organic-rich marine sediments were likely reworked, thus returning the isotopically  
392 light carbon to the marine reservoir (Jarvis et al., 2002). Further evidence supporting the  
393 relationship between the sea-level drop and the lowering isotopic values comes from the  
394 characteristic surface in the isotopic profiles defining the lowest carbon values ( $< 2\text{‰}$ ) and  
395 lowest oxygen values (approaching  $-6\text{‰}$ ). This surface, recognized by a marked negative carbon  
396 and oxygen isotope excursion, coincides with the correlative conformity that represents the  
397 sequence boundary, and thus, with the lowest point of the relative sea level (Figs. 5 & 6).  
398 Furthermore, landwards, the sequence boundary is characterized by a subaerial unconformity,  
399 evidenced by truncation and paleokarst features, which further confirm the subaerial exposure of  
400 the platform (Bover-Arnal et al., 2009, 2022). This interpretation agrees with investigations in  
401 the NE margin of the Arabian plate that also demonstrated that erosional surfaces can be  
402 geochemically identified by characteristic negative isotope excursions (Rahimpour-Bonab et al.,  
403 2013; Mehrabi et al., 2022). Furthermore, comparable relationships between stable isotopes  
404 (mostly  $\delta^{13}\text{C}$  values) and fluctuations in relative sea level have been also reported from the  
405 Oxfordian sedimentary record from central and southern Europe (Jenkyns, 1996), the  
406 Cenomanian-Turonian rocks from southern England (Jarvis et al., 2001), and the Campanian  
407 successions from northern Tunisia (Jarvis et al., 2002).

## 408           **Significance of the carbonate geochemical imprints.---**

409   It is remarkable that the timing and magnitude of the Aptian relative sea-level fall recorded in  
410   Depositional Sequence A align with the global sea-level event known as “KAp2” (Haq, 2014).  
411   This sea-level event is coeval with forced regressive wedges and deep incised valleys in the  
412   Tethys realm and in other Aptian carbonates worldwide (Rameil et al., 2012; Maurer et al., 2013;  
413   Horner et al., 2019). This lowering of relative sea level in the Tethys and elsewhere has been  
414   tentatively linked to glacio-eustasy (Dujoncquoy et al., 2018; Ray et al., 2019; Bover-Arnal et  
415   al., 2024), and thus, was likely triggered by changes in continental ice volume during an early  
416   Aptian cooling event (Stoll and Schrag, 1996; Hochuli et al., 1999; Steuber et al., 2005; Miller et  
417   al., 2005; Dujoncquoy et al., 2018; Davies et al., 2020). In the Galve Sub-basin, where the  
418   studied section is located, this cooling event has been reported from  $\delta^{18}\text{O}$  values of well-  
419   preserved bivalves sampled along the Aptian succession (Fig. 1) (Bonin et al., 2016). Therefore,  
420   a discussion is presented next to explore if the reported isotope values in this study have  
421   paleoclimatic significance.

422   The isotopic trends discussed earlier in this section diverge from the typical isotopic patterns  
423   reported in studies linking paleoclimate and carbonate geochemistry (Bonin et al., 2016). In  
424   particular, sea-level drops caused by the formation of widespread continental ice sheets during  
425   climate cooling events are commonly interpreted from the correlation between sea-level  
426   regressions and positive  $\delta^{18}\text{O}$  excursions (Stoll and Schrag, 2000; Puc at et al., 2003; Steuber et  
427   al., 2022). This is because, during cooling periods, the lighter isotope ( $^{16}\text{O}$ ) is readily taken by  
428   the glacial ice and the heavier isotope ( $^{18}\text{O}$ ) is left behind, increasing the concentration of  $^{18}\text{O}$   
429   and the  $\delta^{18}\text{O}$  value in the ocean (Kumar and Verma, 2021). In the current study, however, the

430 observed trend is contrary, and the forced regressive deposits record a negative  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$   
431 isotope excursion.

432 In paleoclimate studies, samples are commonly chosen to encompass homogeneous lithologies,  
433 generally mudstones, deposited in pelagic environments or in relatively deep-water settings. This  
434 selection aims to avoid isotopic modifications arising from relative sea-level fluctuations,  
435 subaerial exposures, and diagenesis (Stoll and Schrag, 2000; Vimperc et al., 2023). Alternatively,  
436 well-preserved foraminifers or other carbonate fossils that offer robust biostratigraphic  
437 constraints, are commonly used to extract paleoclimatic information (Pucéat et al., 2003; Bonin  
438 et al., 2016). In contrast to this, the stratigraphic interval analyzed here represents a well-  
439 preserved carbonate platform margin, where the carbonate facies, architecture, and likely the  
440 geochemical signature were mainly controlled by the relative sea-level fluctuations (Bover-Arnal  
441 et al., 2009, 2022; Gratacós et al., 2021). Because the shallow marine carbonates were exposed  
442 during the relative sea-level fall, the isotopic values of the regressive deposits are not interpreted  
443 as reflecting a climatic pattern (i.e., cooling event). Instead, they are interpreted to reflect the  
444 depositional environment and associated processes resulting from base-level fall during the  
445 subaerial exposure of the platform (Allan and Matthews, 1982) (Fig. 7).

446 In conclusion, our findings, which are in line with the interpreted third-order sea-level fall  
447 recorded in the Aptian carbonate platform studied, supports one of the previously debated  
448 sequence stratigraphic interpretations for the studied succession. Specifically, the basin floor  
449 wedge in the study area, interpreted by Bover-Arnal et al. (2009) as a forced regressive unit, was  
450 later reinterpreted as the upper part of a transgressive systems tract (Peropadre et al., 2013;  
451 Pomar and Haq, 2016; Pomar, 2020), with the latter authors arguing against evidence for a  
452 falling relative sea level during the deposition of this sedimentary wedge. However, our

453 carbonate geochemical data align with the interpreted exposure of the carbonate platform and  
454 thus, the relative sea-level drop proposed by Bover-Arnal et al. (2009). This interpretation is  
455 further corroborated by recent studies in the eastern Iberian Chain, where this major base-level  
456 fall was regionally recorded, either as forced regressive wedges or deep incised valley developed  
457 on highstand carbonate platforms (Gratacós et al., 2021; Bover-Arnal et al., 2022, 2024).  
458 According to our results, the high variation of C and O isotopic values recorded in the FRWST,  
459 together with the decreasing trend of values from the lower to the upper boundary of this unit,  
460 contrast with the relative homogeneous (and more positive) values characterizing the highstand  
461 deposits of both studied depositional sequences (Figs. 5 & 6). Therefore, we suggest that these  
462 contrasting geochemical patterns are the expression of different genetic types of deposits, which  
463 result from distinct stages of relative sea level (HST and FRWST, respectively). Moreover, a  
464 deeply incised subaerial unconformity (SU) located at the highstand platform top of Depositional  
465 Sequence A was correlated by Bover-Arnal et al. (2009) with the correlative conformity (CC)  
466 capping the forced regressive wedge (Fig. 2). The evidence of exposure in the SU, including  
467 local karst development, agrees with the geochemical signal of the CC displaying the lowest  
468 obtained C and O isotopic values. Thus, these observations are compatible with a sea-level  
469 regression.

470 *Implications for linking sequence stratigraphy and carbonate geochemistry*

471 In sequence stratigraphic interpretations, analyzing suitable outcrop analogs commonly furnishes  
472 detailed information below seismic resolution. This enables the recognition and characterization  
473 of stratal stacking patterns, facies architecture, and depositional systems (Catuneanu et al., 2009;  
474 Ruppel and Ward, 2013) as has been previously done in the studied Aptian section (Bover-Arnal  
475 et al., 2009, 2022). In subsurface reservoir studies for energy exploration, where the geological

476 interpretations and the scale of observations are required below the vertical seismic resolution,  
477 spatial and temporal discontinuities may compromise the resolution and the quality of the  
478 sequence stratigraphic interpretations (Carter et al., 1998). Accordingly, in cases where stratal  
479 patterns and their bounding surfaces are not well defined, the isotope records can serve as a  
480 complementary proxy to bridge specific data gaps in sequence stratigraphic analysis during the  
481 characterization of specific systems tracts and the subsequent reconstruction of relative sea-level  
482 changes.

483 In the study area, the geochemical data reveal that the isotope signatures recorded in the  
484 investigated Aptian platform carbonates were significantly influenced by depositional and  
485 environmental processes (such as changes in the origin of fluids) during fluctuations in relative  
486 sea level. The isotope trends reported here fall in line with the systems tracts and sequence-  
487 stratigraphic interpretation already defined by the evolution of facies and their stacking patterns  
488 (Figs. 6 & 7). Overall, despite the overlap of isotopic values displayed by different systems  
489 tracts, the forced regression and lowstand units display the widest range of  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values  
490 obtained in this study, varying from around 0.0‰ to +4.0‰, and from -6.0‰ to -3.5‰,  
491 respectively. Despite the similar isotopic range, the FRWST can be recognized in the  
492 chemostratigraphic profiles by a general trend of decreasing  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values from samples  
493 taken at the BSFR, which was formed when relative sea level was still high, to samples collected  
494 at the correlative conformity (SB), representing the lowest point of the relative sea level (Figs. 2,  
495 5 & 6). By contrast, the LPWST shows an opposite pattern marked by a consistent increase in  
496 the  $\delta^{13}\text{C}$  and a varying increase in the  $\delta^{18}\text{O}$  values from samples close to the SB to samples  
497 collected at the TS. Thus, where the FRWST is preserved, the sequence boundary, which is  
498 represented by a subaerial unconformity that passes basinward to its marine correlative

499 conformity, and thus developed during the lowest point of relative sea level, correlates with a  
500 marked negative carbon and oxygen isotope excursion (Figs. 6 & 7). This likely suggests the  
501 progressive influence of the meteoric environment (and the exposure of the platform) during  
502 relative sea-level falling.

503 In contrast, the intervals of higher relative sea level (transgressive and highstand carbonate  
504 deposits) are geochemically recognized by a characteristic trend towards the highest  $\delta^{13}\text{C}$  values  
505 documented in this study ( $>3.0\%$ ) and the less varying (and less negative)  $\delta^{18}\text{O}$  values ( $-5.5\%$  to  
506  $-3\%$ ) registered throughout the succession investigated. This reflects, as discussed in the  
507 previous section, the maximum relative sea-level peak recorded in each depositional sequence  
508 and thus, the marine influence on the isotopic values.

509 Therefore, as proposed in this study, the application of isotope stratigraphy (i.e.,  
510 chemostratigraphy) to carbonate rocks, within a sequence stratigraphic framework, enables the  
511 recognition of characteristic geochemical patterns in the vertical stratigraphic profile. In turn,  
512 these isotopic patterns linked to sequence stratigraphy might be correlatable among coeval  
513 carbonate successions from different geographical locations (Rahimpour-Bonab et al., 2013;  
514 Swart, 2015), enabling the determination of specific stages of relative sea level (systems tracts)  
515 within a poorly defined sequence stratigraphic framework.

#### 516 *Diagenetic and geochemical considerations for isotope stratigraphy*

517 Diagenesis of marine carbonates commonly results in geochemical modifications of their  
518 original isotopic signature. With increasing burial depth, carbonate  $\delta^{18}\text{O}$  values tend to decrease  
519 reflecting alteration of primary marine values under the influence of diagenetic fluids and  
520 increasing temperatures (Killingley, 1983). Therefore, using the geochemical data, this section

521 discusses if potential post-depositional effects could have compromised the stratigraphic  
522 significance of the measured isotope values. In particular, the  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values measured in  
523 the studied Aptian succession show that, with only a few exceptions, bulk carbonates and  
524 separate carbonate components (matrix, grain, cement) from the same sample provided similar  
525 (or slightly varying) range of isotope values (Figs. 5 & 6). Thus, this could either indicate that  
526 the carbonate components are genetically related, that the isotope values were homogenized  
527 during carbonate deposition and early diagenesis, or that the original values were reset  
528 (overprinted) during later burial diagenesis (Swart et al., 2009; Swart, 2015). In this regard,  
529 characteristic geochemical patterns have been recognized in the vertical profile of the  $\delta^{13}\text{C}$  and  
530  $\delta^{18}\text{O}$  values (Figs. 5 & 6). These geochemical profiles, as discussed in the previous section,  
531 define isotopic trends that broadly correlate with specific systems tracts. Therefore, although a  
532 certain extent of post-depositional modification of the isotope values cannot be disregarded, the  
533 geochemical trends of the studied platform carbonates have isotope stratigraphic significance  
534 considering that: (i) the isotopic values seem to vary according to the different systems tracts and  
535 their stratigraphic boundaries (Droxler et al., 1983; Roth and Reijmer, 2005); (ii) the overall  
536 observed trends, with positive carbon excursions matching rising sea levels and vice versa,  
537 follow the arguments and interpretations widely discussed in the literature (Föllmi et al., 1994;  
538 Vahrenkamp, 1996); and (iii) the different systems tracts, particularly the HST and TST from  
539 both depositional sequences, show little isotopic variation from the uppermost slope and towards  
540 the basin (as observed in LM samples, Fig. 5). Regarding this latter statement, if strong  
541 diagenesis had overprinted our isotopic values, HST and TST samples located respectively  
542 below and above the subaerial unconformity (SU), would have differing isotopic values likely  
543 resulting from meteoric vs marine diagenesis, respectively. Similarly, HST and TST samples

544 located near the uppermost slope (around the SU), would have different diagenetic overprint to  
545 HST and TST samples located around the correlative conformity (towards the basin), but again,  
546 we do not observe a different isotopic pattern that could indicate a strong diagenetic overprint.  
547 This also means that the TST and HST intervals display similar variation of C and O isotopic  
548 values, irrespective of the type of facies, which is an observation already made in modern  
549 carbonates from the Great Bahama Bank (Swart et al., 2009).

550 It is also noteworthy that the previous contributions linking geochemistry to sequence  
551 stratigraphy (Vahrenkamp, 1996; Jenkyns, 1996; Jarvis et al., 2001) have mainly used carbon  
552 isotope stratigraphy as a proxy for relative sea-level variation. This is because, in shallow marine  
553 carbonate settings, carbon isotopes tend to be rock-dominated (rather than fluid-dominated) and  
554 are less sensitive to temperature fractionation and diagenetic alteration compared to oxygen  
555 isotopes (Banner and Hanson, 1990; Marshall, 1992; Banner and Kaufman, 1994). This can  
556 explain, for instance, the greater scattering of the O isotope values in each systems tract with  
557 respect to the C isotope values in the same interval (Fig. 5). Nevertheless, the  $\delta^{18}\text{O}$  values of  
558 carbonates are controlled not only by temperature but also by the  $\delta^{18}\text{O}$  values of the formation  
559 (or diagenetic) fluids (because of the dominant oxygen reservoir of the fluids) (O'Neil et al.,  
560 1969; Muñoz-López et al., 2022; Cruset et al., 2023). In contrast to the negative  $\delta^{18}\text{O}$  values of  
561 meteoric waters, for instance, marine fluids exhibit more positive values. Consequently, both  
562 isotope proxies ( $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$ ) in the studied platform carbonates serve as valuable tools for  
563 sequence stratigraphic analysis.

## 564 CONCLUSIONS

565 This study is an example of how isotope geochemistry and chemostratigraphy, when applied to  
566 ancient carbonate systems, may help in characterizing sequence-stratigraphic attributes. In the

567 studied Aptian platform carbonate succession, the  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  values show a strong correlation  
568 with a previously reported sequence stratigraphic framework. This allowed us to discriminate  
569 between divergent sequence stratigraphic interpretations previously proposed for the study area.

570 The systems tracts associated with relative sea-level highs (i.e., HST and TST) are characterized  
571 by a trend towards the highest  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values obtained in this study, reaching up to +5.1‰  
572 and up to -1.8‰, respectively. Among these, HST exhibits the most uniform values. In contrast,  
573 the systems tracts associated with stages of lower relative sea-level (i.e., FRWST and LPWST)  
574 show a greater isotope variability, with  $\delta^{13}\text{C}$  values ranging from +0.5 to +4.5‰ and  $\delta^{18}\text{O}$  values  
575 from -6.1 to -3.5‰. Within the FRWST, both isotope proxies progressively increase from the  
576 sequence boundary (SB) towards the BSFR, whereas in the LPWST, isotope values increase  
577 from the sequence boundary towards the TS. This implies that the lowest  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values  
578 documented in this study are found at the SB (correlative conformity), coinciding with the lowest  
579 point of relative sea level. Furthermore, the results show that, with only a few exceptions,  
580 sampling bulk carbonates and separate carbonate components (micritic matrix, grain, cement)  
581 also yield similar isotope patterns.

582 The interpretations link carbonate isotope variations to a well-constrained sequence stratigraphic  
583 framework. These isotope variations are the expression of changing depositional and  
584 environmental processes during third-order Aptian sea-level fluctuations and can serve as  
585 analogue for: (i) carbonate platform successions characterized by poor (bio)stratigraphic  
586 resolution and weak to moderate diagenetic overprinting, and (ii) for potentially identifying and  
587 correlating systems tracts (specific stages of relative sea level) and their bounding surfaces in  
588 other locations.

589

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#### 599 **DATA AVAILABILITY STATEMENT**

600 The data that support the findings of this study are available from the corresponding author upon  
601 reasonable request.

#### 602 **CONFLICT OF INTEREST**

603 The authors declare that they have no known conflict of interest

#### 604 **CREDIT AUTHORSHIP CONTRIBUTION STATEMENT**

605 **D. Muñoz-López:** Conceptualization, Data curation, Formal analysis, Investigation,  
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607 **A. Koeshidayatullah:** Conceptualization, Data curation, Formal analysis, Writing – review and  
608 editing. **T. Bover-Arnal:** Conceptualization, Formal analysis, Investigation, Writing – review  
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958

## FIGURE CAPTIONS

959 **Fig. 1.** From right to left: simplified structural and isopach map of the Maestrat Basin (located in  
960 the eastern Iberian Chain, eastern Iberian Peninsula) during the Late Jurassic-Early Cretaceous  
961 rifting, and geological map of the central Galve Sub-basin. The schematic transect A-A' is found  
962 in Fig. 2A and the red star indicates the location of the study area at Las Mingachas section. The  
963 geological map was modified from Canérot et al. (1979), Gautier (1980) and Bover-Arnal et al.  
964 (2022), whereas the structural map of the Maestrat Basin was modified from Salas et al. in  
965 Martín-Chivelet et al. (2019) and Bover-Arnal et al. (2022).

966

967 **Fig. 2.** Sequence stratigraphic framework of Las Mingachas section. (A) Simplified and  
968 schematic cross section showing the platform to basin transition setting cropping out in the study  
969 area. See Fig. 1 for the location of the section A-A'. The dashed red square represents the  
970 panoramic photo showed in Figs. B and C. (B) Panoramic view and (C) sequence stratigraphic  
971 interpretation of Las Mingachas. The grey bar represents the location of the stratigraphic log

972 (Fig. 3). The stages of relative sea-level fluctuation are also included. Modified from Bover-  
973 Arnal *et al.* (2009, 2022).

974

975 **Fig. 3.** Location of samples in outcrop (upper) and in the stratigraphic log of the studied Aptian  
976 sedimentary succession (lower). The stratigraphic log shows the chronostratigraphic and  
977 sequence stratigraphic analysis. Ammonite data come from Moreno-Bedmar *et al.* (2010) and  
978 Bover-Arnal *et al.* (2016). Samples were collected considering two sampling strategies for  
979 comparison. LM samples (LM1-LM18) covers laterally the full studied outcrop, whereas vertical  
980 log samples (samples 131-174) runs parallel to the stratigraphic log. Colored dots link the  
981 samples to the specific stages of the relative sea-level curve (systems tracts).

982

983 **Fig. 4.** (A) Outcrop image showing the location of representative samples where the sedimentary  
984 facies of the section were logged and sampled. (B) Wackestone to packstone limestones of the  
985 upper part of the Forcall Formation showing *Palorbitolina lenticularis* (P) (HST, Depositional  
986 Sequence A). (C) Forced regressive deposits of the Villarroya de los Pinares Formation  
987 characterized by a grainstone texture with specimens of *Orbitolinopsis* (O) (FRWST,  
988 Depositional Sequence A). (D) Lowstand deposits of the Villarroya de los Pinares Formation  
989 exhibiting a rudstone texture with sections of corals and calcareous algae (LPWST, Depositional  
990 Sequence B). (E) Transgressive limestones of the Villarroya de los Pinares Formation showing a  
991 rudstone texture with fragments of rudist shells, corals and calcareous algae (TST, Depositional  
992 Sequence B).

993

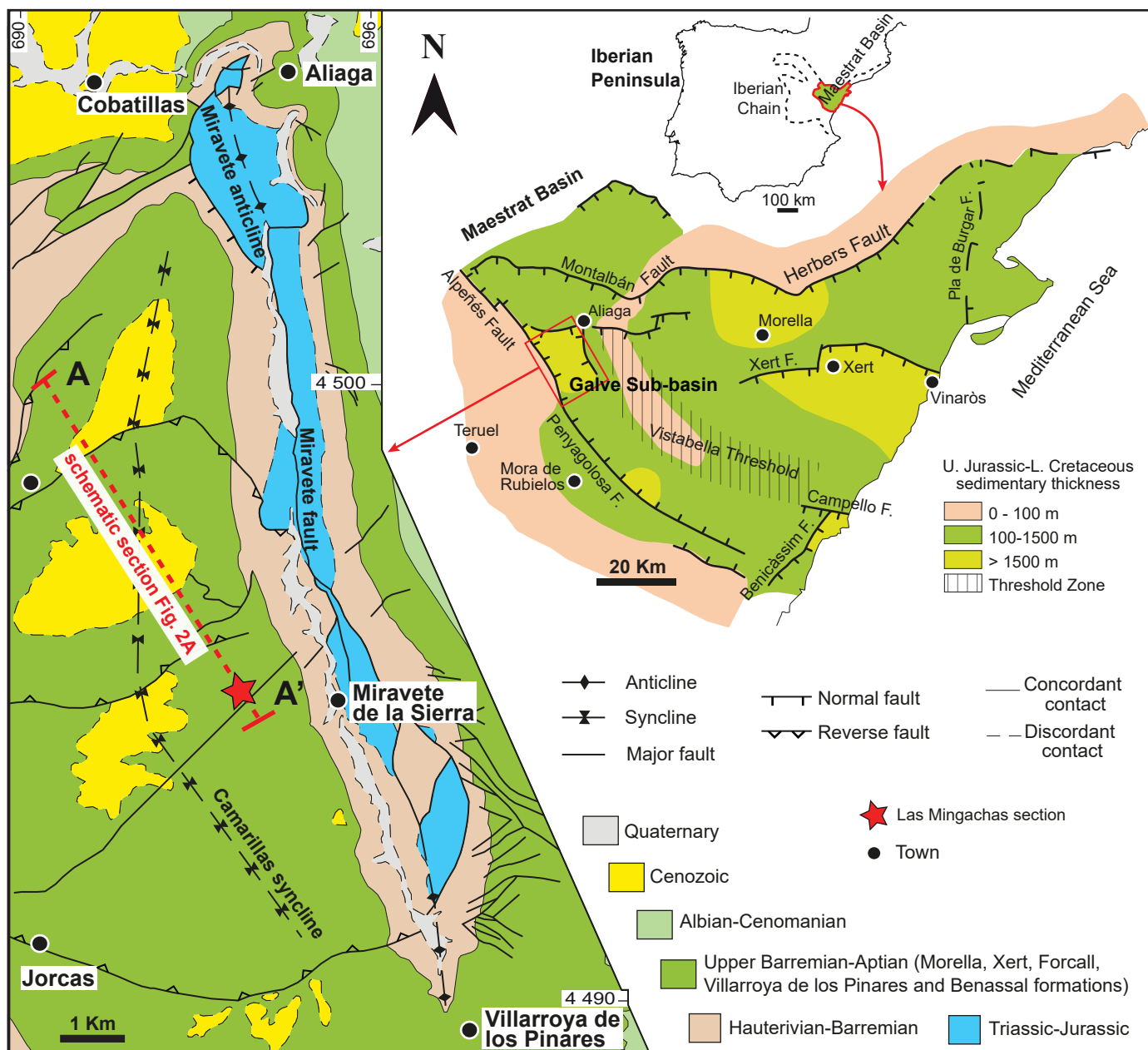
994 **Fig. 5.** Stratigraphic log of the Aptian sedimentary succession studied at Las Mingachas  
995 including the sequence stratigraphic analysis and the carbon and oxygen isotope profiles  
996 measured. LM samples include bulk and separate carbonate components, whereas vertical log  
997 samples consist of bulk carbonates. Legend of symbols referring to sedimentary structures and  
998 sample numbers and references are explained in Fig. 3.

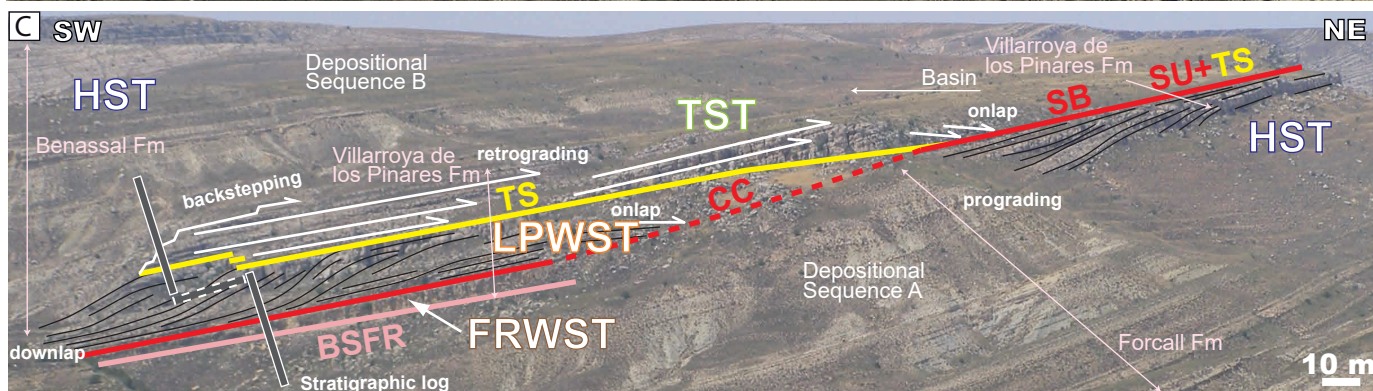
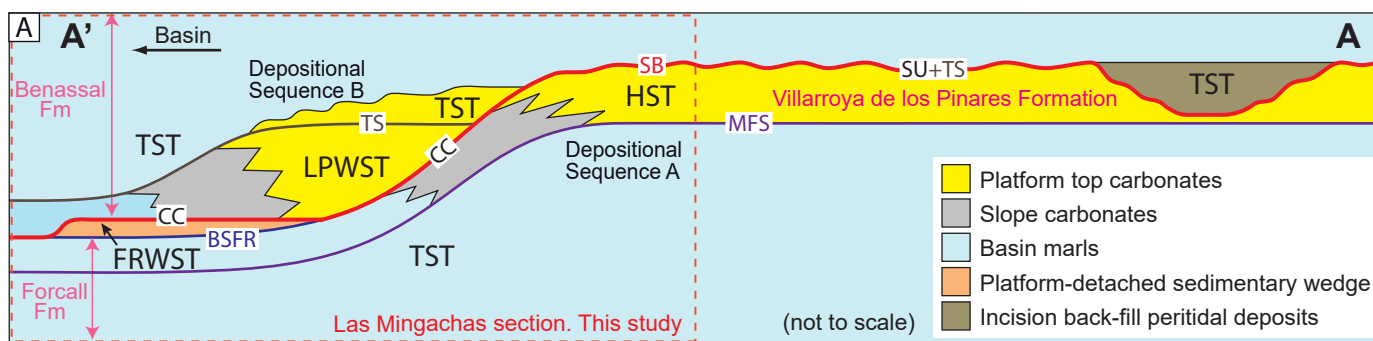
999

1000 **Fig. 6.** (A) Crossplot showing the  $\delta^{18}\text{O}$  vs  $\delta^{13}\text{C}$  values of the studied platform carbonate  
1001 succession. The colored box indicates the typical isotope values of Lower Cretaceous marine  
1002 carbonates. (B) Evolution of  $\delta^{13}\text{C}$  values through time linked to the depositional sequences and  
1003 the corresponding systems tract. The  $\delta^{13}\text{C}$  profile is characterized by an episode of lowermost C  
1004 values separating two intervals of more positive C values. The red boxes represent isotope values  
1005 from the closest samples to the sequence boundary, which coincide with the lowermost C and O  
1006 isotopic values.

1007

1008 **Fig. 7.** Schematic diagram showing the interpreted relationship among relative sea-level  
1009 fluctuations, sedimentation, sequence stratigraphic analysis, and the observed geochemical  
1010 patterns of the studied Aptian platform carbonates. Note that the specific stages of relative sea-  
1011 level high (coinciding with highstand and transgressive system tracts) show a general trend  
1012 towards more positive and less negative  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values, respectively. In contrast, the  
1013 stages of lower relative sea level (forced regressive and lowstand system tracts) yield less  
1014 positive  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values. The correlative conformity representing the sequence boundary is  
1015 marked by a negative C and O isotopic excursion.

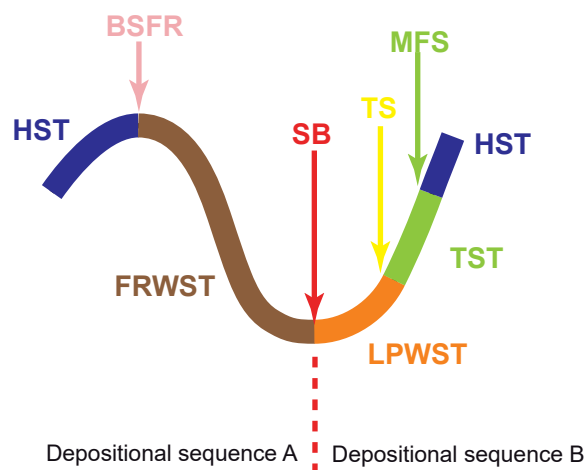


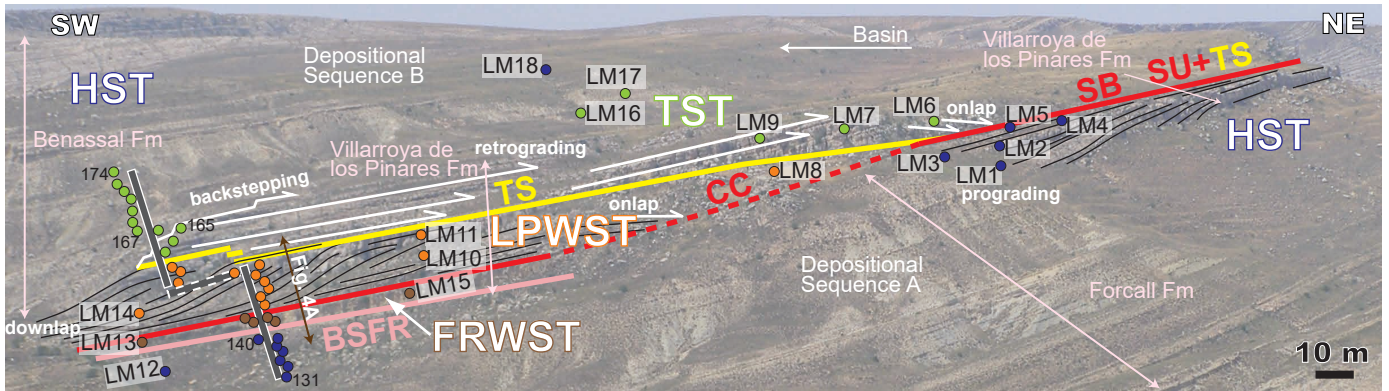


**Key**

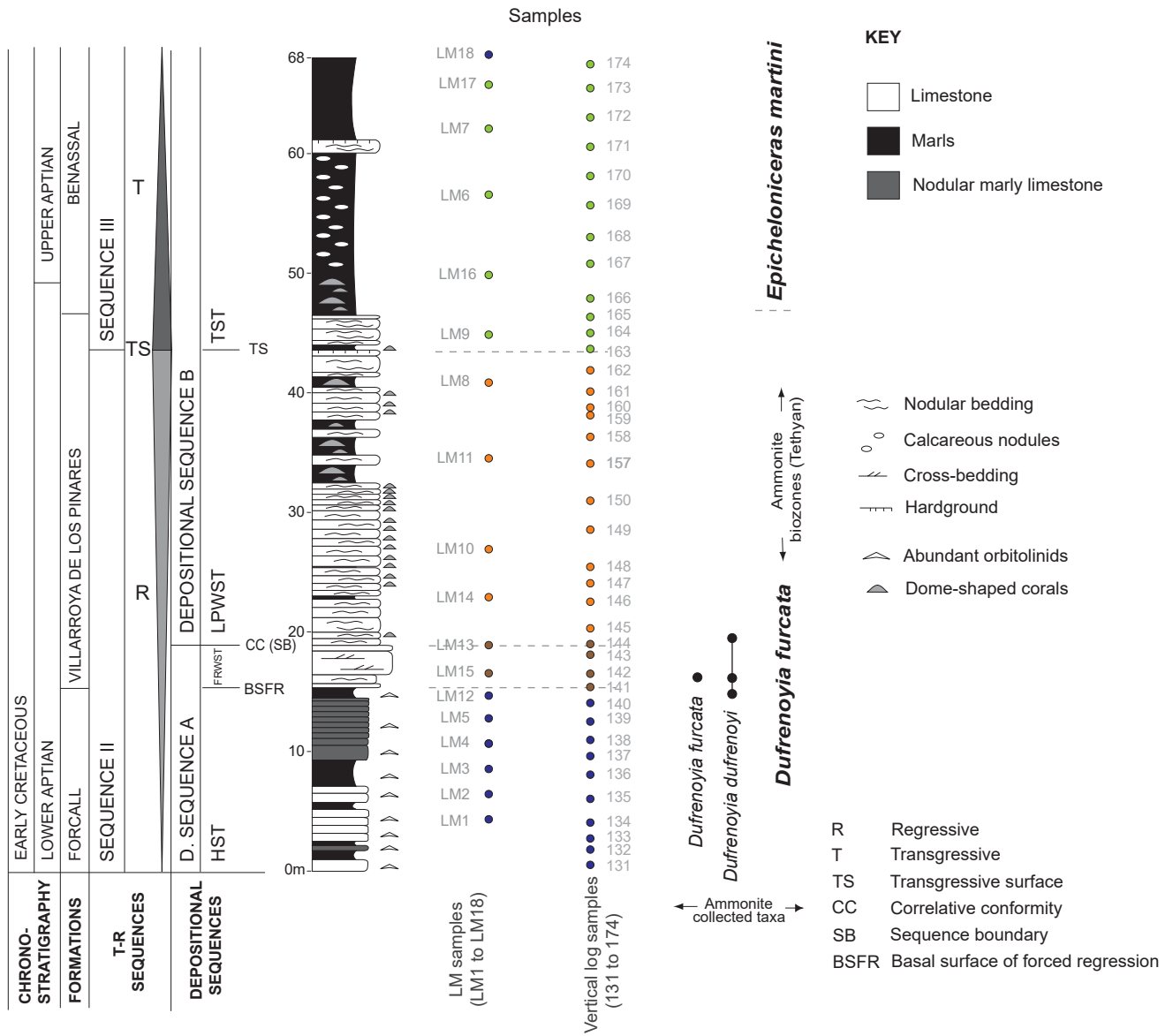
- HST: Highstand Systems Tract
- FRWST: Forced Regressive Wedge Systems Tract
- LPWST: Lowstand Prograding Wedge Systems Tract
- TST: Transgressive Systems Tract
- BSFR: Basal Surface of Forced Regression
- SB: Sequence Boundary
- TS: Transgressive Surface
- MFS: Maximum Flooding Surface
- CC: Correlative Conformity
- SU: Subaerial Unconformity

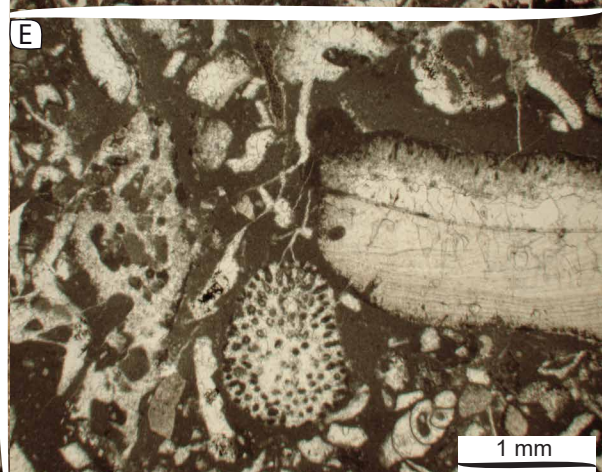
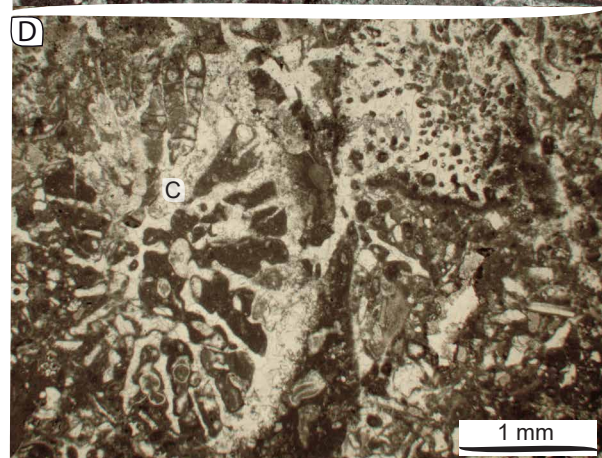
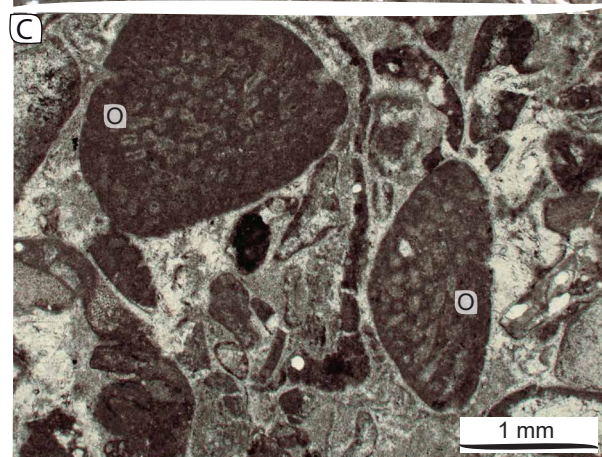
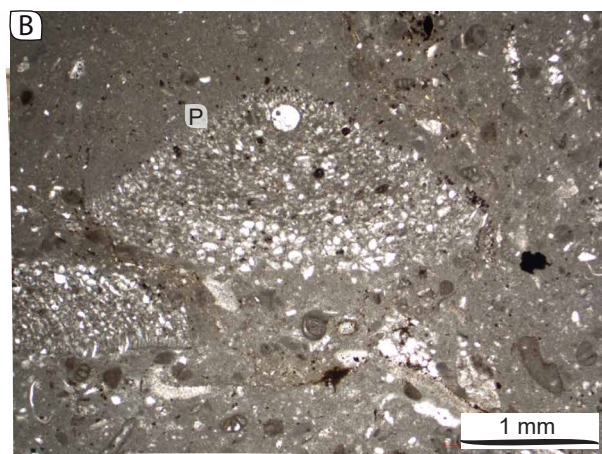
**Relative sea-level fluctuation**

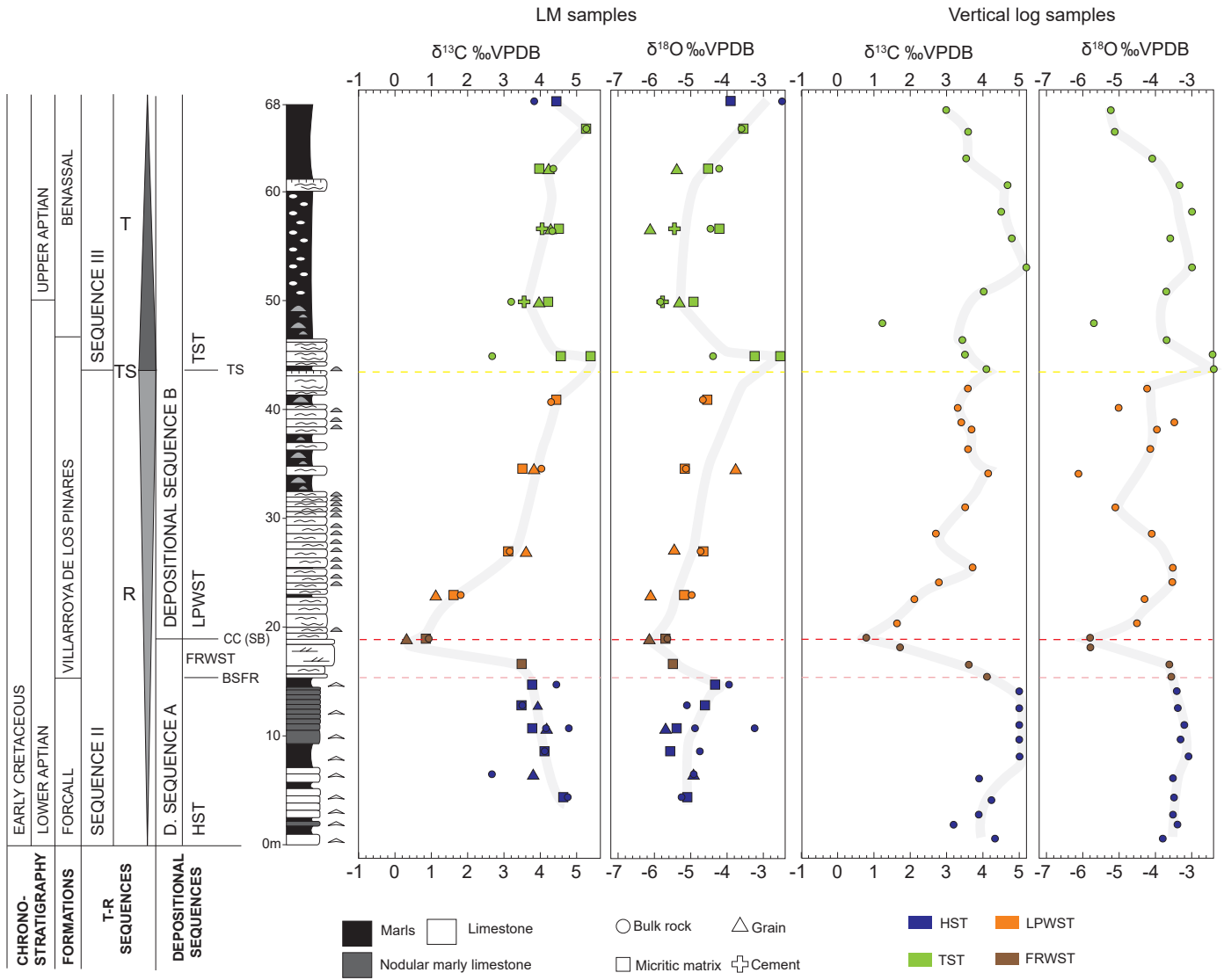


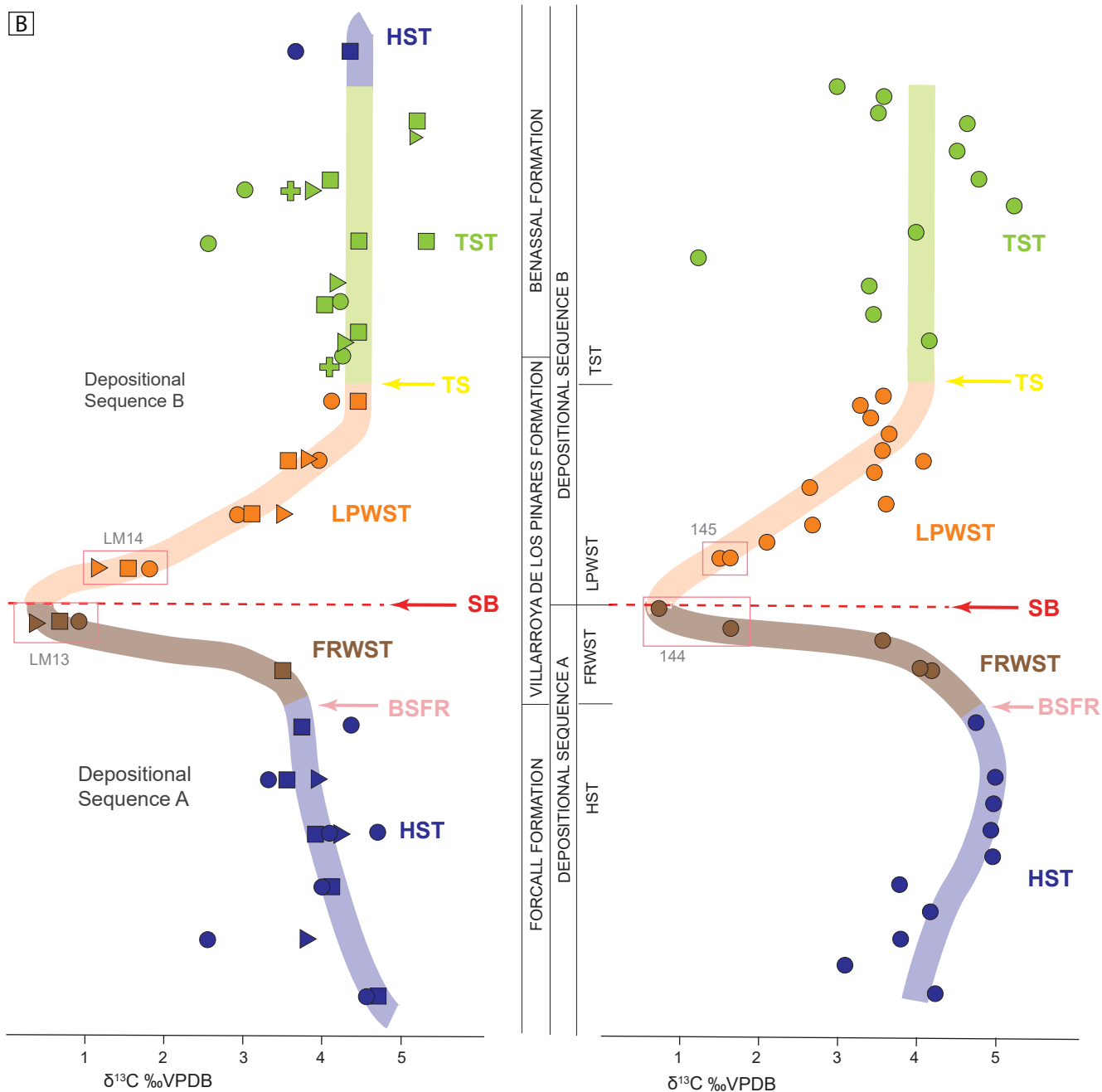
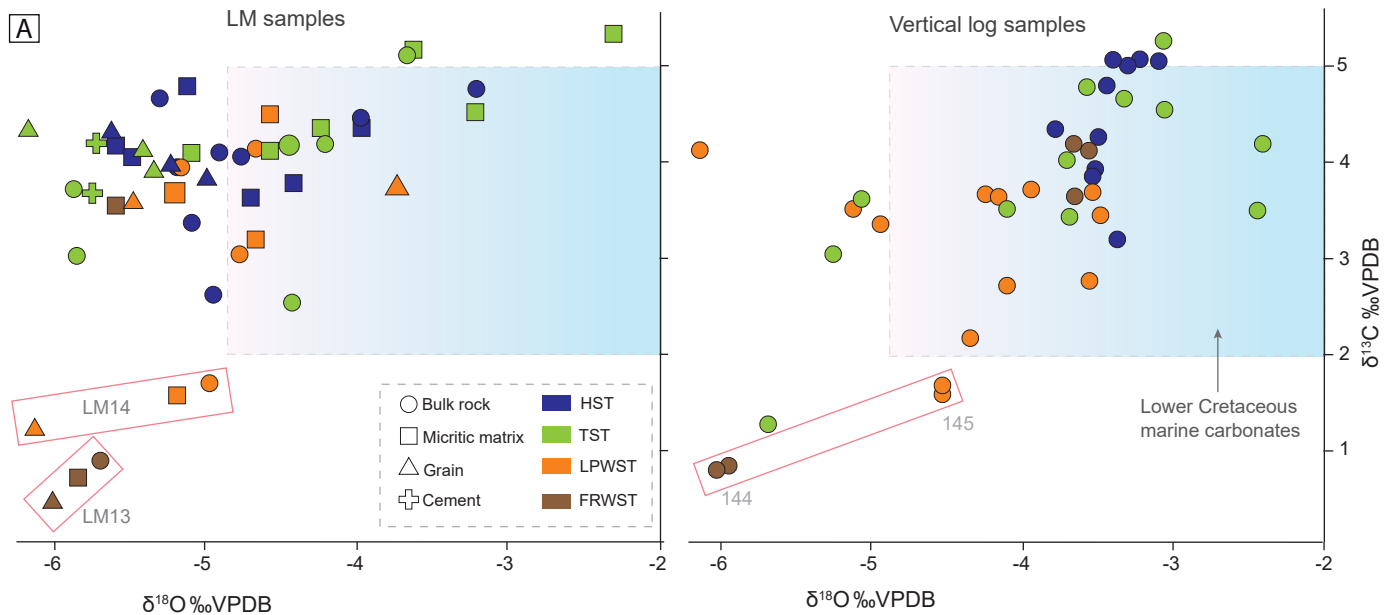


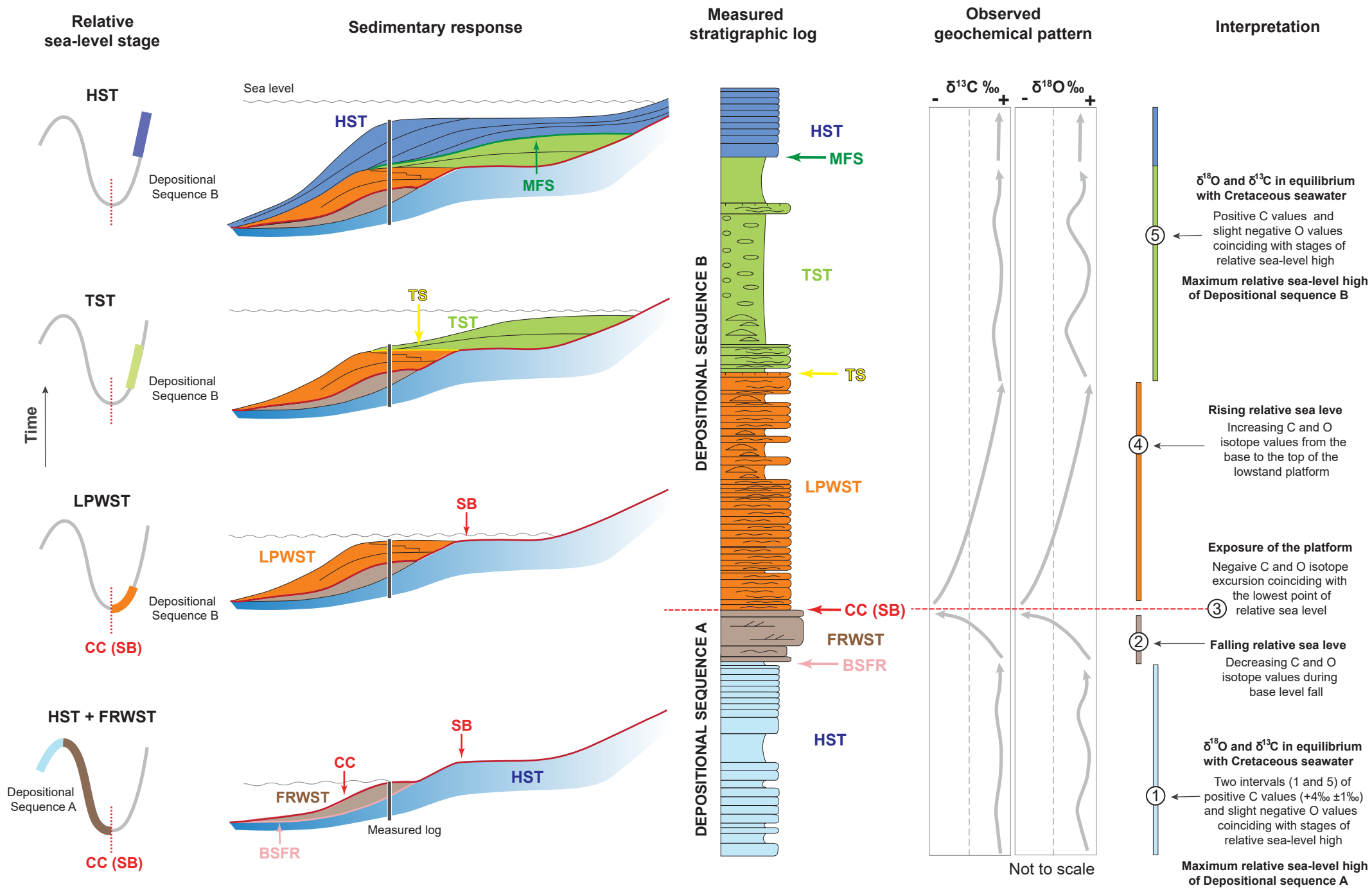
Stratigraphic log and vertical log samples











**Table 1.**  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values of the studied Aptian platform carbonates. These values are arranged according to Depositional sequences A and B, their associated systems tracts, and sampling strategy (LM samples vs vertical log samples).

Depositional sequence	Systems tract	Sample reference	$\delta^{13}\text{C}$	$\delta^{18}\text{O}$	Sample reference	$\delta^{13}\text{C}$	$\delta^{18}\text{O}$
			(‰VPDB)	(‰VPDB)		(‰VPDB)	(‰VPDB)
			LM samples		Vertical log samples		
A	HST	LM1-B	4.7	-5.3	131-B	4.3	-3.7
		LM1-M	4.7	-5.1	132-B	3.2	-3.4
		LM2-B	2.6	-4.9	133-B	3.9	-3.5
		LM2-G	3.8	-4.9	134-B	4.3	-3.5
		LM3-B	4.1	-4.8	135-B	3.9	-3.5
		LM3-M	4.2	-5.6	136-B	5.1	-3.1
		LM4-B	4.1	-4.9	137-B	5.0	-3.3
		LM4-B	4.7	-3.2	138-B	5.1	-3.2
		LM4-G	4.2	-5.7	139-B	5.1	-3.4
		LM4-M	4.1	-5.5	140-B	4.8	-3.4
		LM5-B	3.4	-5.1			
		LM5-G	3.9	-5.1	141a-B	4.2	-3.7
		LM5-M	3.6	-4.7	141b-B	4.1	-3.6
		LM12-B	4.4	-3.9	142-B	3.6	-3.6
		LM12-M	3.8	-4.4	143-B	1.7	-5.8
A	FRWST	LM15-M	3.5	-5.6	144a-B	0.8	-6.0
		LM13-B	0.9	-5.7	144b-B	0.8	-5.9
		LM13-M	0.7	-5.8	145a-B	1.6	-4.5
		LM13-M+G	0.5	-6.0	145b-B	1.6	-4.5
		LM14-B	1.7	-4.9	146-B	2.2	-4.3
B	LPWST	LM14-G	1.2	-6.1	147-B	2.8	-3.6
		LM14-M	1.6	-5.2	148-B	3.7	-3.5
		LM10-B	3.0	-4.8	149-B	2.7	-4.1
		LM10-G	3.5	-5.5	150-B	3.5	-5.1
		LM10-M	3.2	-4.7	157-B	4.1	-6.1
		LM11-B	3.9	-5.2	158-B	3.6	-4.2
		LM11-G	3.7	-3.7	159-B	3.7	-4.0
		LM11-M	3.7	-5.2	160-B	3.4	-3.5
		LM8-B	4.1	-4.7	161-B	3.4	-5.0
		LM8-M	4.5	-4.6	162-B	3.6	-4.2
B	TST	LM9-B	4.2	-4.5			
		LM9-CC(AH)	4.2	-5.7			
		LM9-G	4.3	-6.2			
		LM9-M	4.3	-4.2	163-B	4.2	-2.4
		LM16-B	4.2	-4.2	164-B	3.5	-2.4
		LM16-G	4.1	-5.4	165-B	3.4	-3.7
		LM16-M	4.1	-4.6	166-B	1.3	-5.7
		LM6-B	2.5	-4.4	167-B	4.0	-3.7
		LM6-M1	5.3	-2.3	168-B	5.2	-3.1
		LM6-M2	4.5	-3.2	169-B	4.8	-3.6

		LM7-B	3.7	-5.9	170-B	4.5	-3.1
		LM7-B	3.0	-5.9	171-B	4.7	-3.3
		LM7-CC	3.7	-5.8	172-B	3.5	-4.1
		LM7-G	3.9	-5.3	173-B	3.6	-5.1
		LM7-M	4.0	-5.1	174-B	3.2	-5.3
		LM17-B	5.1	-3.7			
		LM17-M	5.2	-3.6			
B	HST	LM18-B	3.6	-1.8			
		LM18-M	4.4	-3.9			