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4
5 **Constraining the paleoclimate and paleoecology of the Selandian**
6 **– Thanetian transition in the Lower Wilcox, Texas Gulf Coast**

7
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23 **ABSTRACT**

24 The Selandian–Thanetian Transition Event (STTE) represents a relatively understudied interval of
25 carbon cycle instability and environmental disruption during the Paleocene. This study presents a
26 comprehensive sedimentological, palynological, and geochemical record of the STTE based on
27 two shallow marine wells (Moczygamba VT #11 and Vogelsang Frieda #1) from the Texas Gulf
28 Coast. Our multi-proxy approach integrates palynology and nannofossil biostratigraphy to
29 establish age constraints and track ecological changes, organic carbon isotopes to detect regional
30 preservation of the STTE, and use elemental ratios to infer chemical weathering intensity and mean
31 annual precipitation. Results indicate that the STTE along the Gulf Coast was characterized by
32 elevated weathering rates under semi-arid to humid climatic conditions. Palynological
33 assemblages show a marked increase in *Thomsonipollis magnificus*, suggesting the expansion of
34 mangrove or coastal swamp taxa during inferred warming phases. Notably, negative excursions in
35 organic carbon isotopes, along with increased terrestrial input of organic matter in both cores,
36 confirm the regional expression and impact of the STTE. These findings provide critical
37 constraints on the Selandian–Thanetian Transition Event by linking carbon cycle perturbations to
38 continental-scale source-to-sink responses across North America, and demonstrate how deltaic
39 systems along the Gulf Coast recorded and responded to mid-Paleocene global warming.

40

41 **INTRODUCTION**

42 The Paleocene Epoch (66–56 Ma) was characterized by repeated, short-lived perturbations in the
43 global carbon cycle, culminating in the well-documented Paleocene–Eocene Thermal Maximum
44 (PETM; [Zachos et al., 2008](#)). Prior to the PETM, other transient warming events have been
45 recognized in both marine and terrestrial records. One such event is the early late Paleocene

46 hyperthermal (ELPE; [Bralower et al., 2002](#)), also referred to as the mid-Paleocene biotic event
47 (MPBE; [Bernaola et al., 2007](#)). This climatic perturbation occurred near the boundary of the
48 Selandian and Thanetian stages and has been identified in multiple deep-sea cores including
49 Shatsky Rise (ODP Leg 198), Walvis Ridge (ODP Leg 208), Maud Rise (ODP Leg 113), and
50 Black Nose (ODP Leg 171) as well as in several onshore and terrestrial sections across Europe,
51 North Africa, and the Southern Hemisphere (e.g., Zumaia, Contessa Road, Tejerouine, Ouled
52 Abdoun, Naqb El-Rufuf, Mead Stream, Cerro Bayo) ([Bralower et al., 2002](#); [Zachos et al., 2004](#);
53 [Petruzzo, 2005](#); [Bernaola et al., 2007](#); [Westerhold et al., 2011](#); [Coccioni et al., 2012](#); [Karoui-](#)
54 [Yaakoub et al., 2014](#); [Kocsis et al., 2014](#); [Littler et al., 2014](#); [Pujalte et al., 2014, 2016](#); [Hyland et](#)
55 [al., 2015](#); [Hewaidy et al., 2019](#)).

56 In the Tethyan realm, biotic and environmental changes across this interval have been collectively
57 termed the Selandian–Thanetian Transition Event (STTE) by [Coccioni et al. \(2012\)](#). However, the
58 precise stratigraphic and causal relationships among the STTE, ELPE, and MPBE remain poorly
59 understood ([Coccioni et al., 2019](#)).

60 The STTE is marked by negative carbon isotope excursions (CIEs), biotic turnover, and
61 oceanographic reorganization, suggesting transient climate warming and disruption of the global
62 carbon cycle ([Bornemann et al., 2009](#)). Proposed mechanisms include volcanic outgassing,
63 methane hydrate destabilization, and oxidation of organic carbon ([Sluijs et al., 2007](#)). While the
64 magnitude of the STTE was smaller than that of the PETM, which involved $>5^{\circ}\text{C}$ global warming
65 and major ecosystem restructuring ([McInerney & Wing, 2011](#)), it perhaps reflects preconditioning
66 processes in the Earth system ([Maufrangeas et al., 2020](#)).

67 Recent studies using high-resolution carbon isotope stratigraphy have documented multiphase
68 carbon cycle perturbations during the late Selandian–early Thanetian, pointing to complex

69 interactions between carbon sources and sinks (Westerhold et al., 2011; Mafrangeas et al., 2020).
70 Paleooceanographic proxies suggest parallel changes in ocean circulation, oxygenation, and
71 productivity that may have modulated the climatic response (Bornemann et al., 2009).

72 Both marine and terrestrial records now provide growing evidence for the global extent of the
73 STTE. In marine settings, changes in nannofossil and foraminiferal assemblages along with shifts
74 in carbonate content and magnetic susceptibility are observed at sites such as Contessa Road (Italy)
75 and Naqb El-Rufuf (Egypt) (Coccioni et al., 2019; Hewaidy et al., 2019). On land, paleosol carbon
76 isotope excursions and changes in vegetation indicate transient warming and hydrological shifts.
77 At Cerro Bayo (Argentina), the STTE is preserved as a two-stage hyperthermal event (Hyland et
78 al., 2015), while the Lairière section in the Pyrenees reveals CIEs within fluvio-lacustrine strata
79 correlated to global events (Mafrangeas et al., 2020).

80 In this study, we evaluate whether the Selandian–Thanetian Transition Event is recorded in a
81 shallow marine delta system along the Texas Gulf Coast. We test the hypothesis that carbon cycle
82 instability is reflected in negative $\delta^{13}\text{C}$ excursions, alongside shifts in vegetation, weathering
83 intensity, and precipitation. Using a multi-proxy approach that integrates palynological, calcareous
84 nannofossil, and geochemical data, we examine two shallow marine wells (Moczygemba VT #11
85 and Vogelsang Frieda #1). Although coastal-deltaic environments are susceptible to sedimentary
86 discontinuities, our results reveal a coherent STTE signal. Given the Gulf Coast's role in North
87 American sediment routing, this study offers insights into source-to-sink dynamics and how
88 marginal marine systems recorded mid-Paleocene climate perturbations at a continental scale.

89

90 **GEOLOGICAL SETTING, STRATIGRAPHY AND AGE CONSTRAINTS**

91 **Sediment routing along the Texas Gulf Coast**

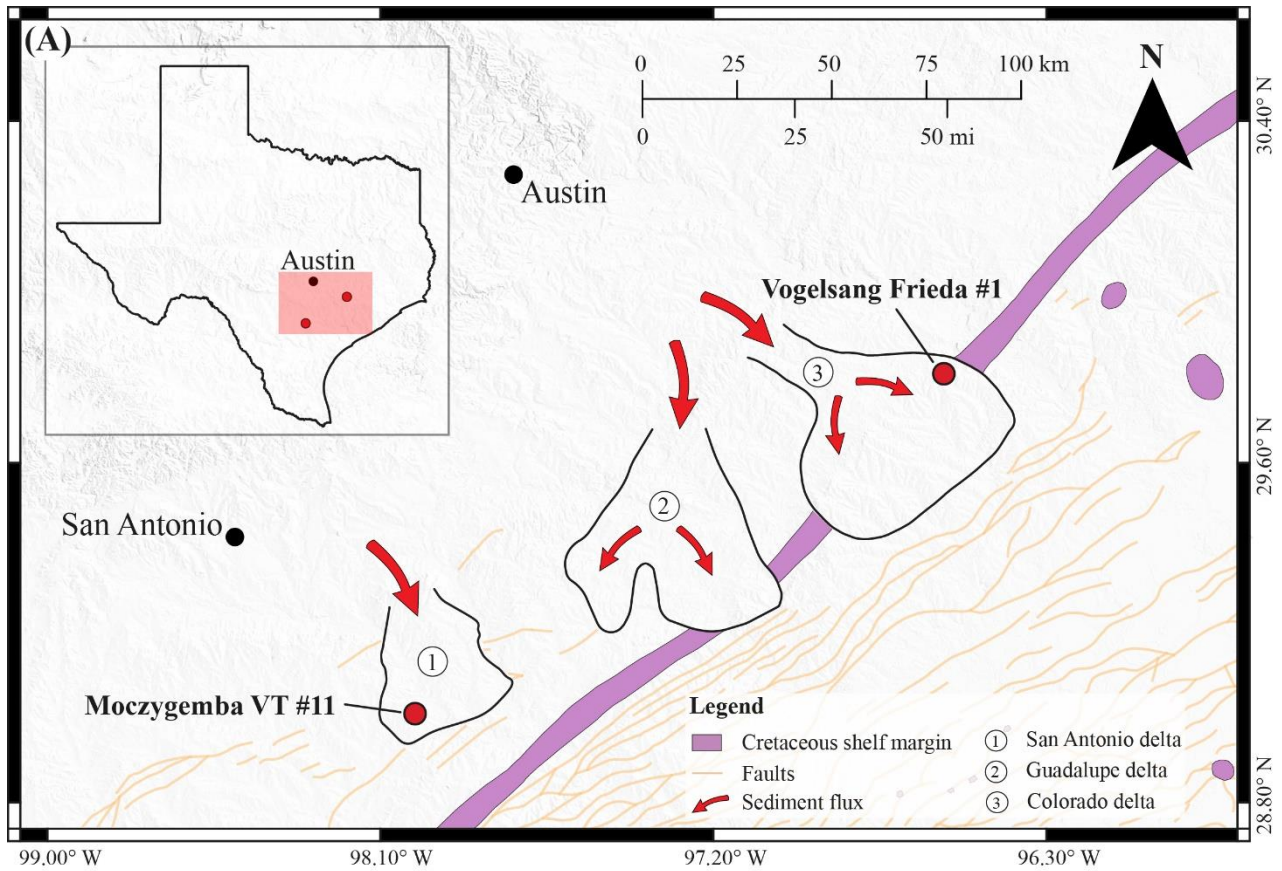
92 During the Selandian-Thanelian, large-scale fluvial systems primarily controlled sediment routing
93 along the Texas Gulf Coast, transporting siliciclastic sediments from the North American
94 continental interior to the Gulf of Mexico (Galloway et al., 2000; Galloway et al., 2011; Hessler
95 et al., 2017). The Laramide Orogeny in NW-USA during the Paleogene played a key role in
96 reshaping North American drainage patterns, generating significant hinterland relief and initiating
97 continental-scale sediment transport (Blum and Pecha, 2014; Sharman et al., 2017). A limited
98 number of major fluvial axes delivered these sediments, each forming large prograding deltaic
99 depocenters in the Gulf Basin (Galloway et al., 2011).

100 The paleo-Colorado rivers were dominant transport systems, with the Colorado axis serving as a
101 primary Laramide-era drainage route during Wilcox time. The paleo-Colorado River, with an
102 estimated 2×10^6 km² catchment area and sediment load of ca. 290 Mt/yr, supplied material to
103 deltaic depocenters in the Rockdale delta system within the Houston embayment (Fisher and
104 McGowen, 1967; Sweet and Blum, 2012). Detrital zircon U-Pb data confirm sediment sources
105 from the Southern Rockies, Cordilleran arc, and northern Mexico (Galloway et al., 2011).

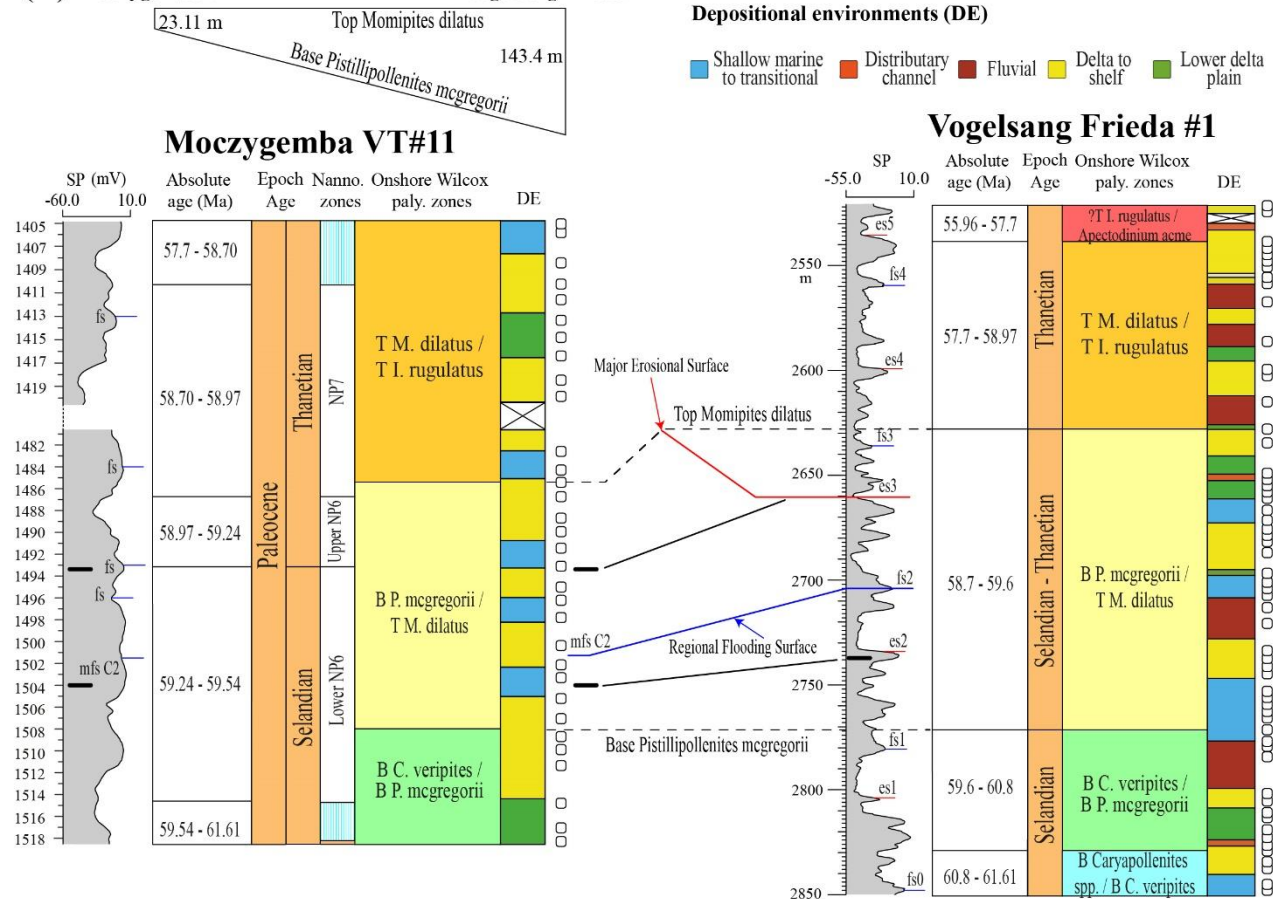
106 The Texas Wilcox Group represents the first major influx of Laramide-derived siliciclastic
107 sediments into the Gulf Basin, marking a shift in sedimentation as fluvial systems expanded
108 southward and bypassed overfilled Laramide basins (Galloway et al., 2011). This influx was
109 significant, with sediment flux exceeding 150,000 km³/Ma, one of the highest multi-million-year
110 depositional episodes recorded for the Gulf of Mexico (Galloway et al., 2000). The Wilcox Group
111 is subdivided into the Lower, Middle, and Upper Wilcox depositional episodes, with the Middle

112 Wilcox being overlain by the broad marine Yoakum Shale, commonly associated with the
113 Paleocene-Eocene boundary (Galloway, 2008; Vimpere et al., 2023).

114 The Rockdale delta system acted as a major depocenter, with multiple distributary channels
115 transporting material across a broad shelf setting under greenhouse conditions (see figure 1 of
116 Zeng and Olariu, 2024). Sediment supply fluctuations, eustatic sea-level changes, and subsidence
117 rates influenced transgressive–regressive (T-R) cycles, generating complex stratigraphic stacking
118 patterns (Zhang et al., 2018). Paleoclimatic shifts also played a key role in modulating sediment
119 supply, with geochemical proxies indicating variations in hinterland weathering intensity. Warmer
120 intervals were associated with intensified chemical weathering and increased clay input, while
121 fluctuations in riverine discharge affected delta progradation and shoreline migration (Hessler et
122 al., 2017; Vimpere et al., 2023).



(B) Moczygamba VT#11 Vogelsang Frieda #1



124 **Figure 1. Map, depositional framework and stratigraphy of the study area.** (A) Regional map
125 of Texas highlighting the Moczygemba VT#11 (San Antonio Delta) and Vogelsang Freida #1
126 (Colorado Delta) wells. Major sediment flux pathways (red arrows), the Cretaceous shelf margin
127 (purple line), and major growth fault zones (orange lines) are shown. Growth faults dip southeast.
128 (B) Stratigraphic sections with self-potential logs, absolute ages, biozones (nannofossil and
129 palynological), and interpreted depositional environments (shallow marine, transitional, deltaic
130 distributary and fluvial channels, delta-shelf transition, and lower delta plain). SP log trends
131 (coarsening- or fining-upward) help define transgressive and regressive intervals, marked by
132 flooding surfaces (FS), maximum flooding surfaces (MFS), and erosional surfaces (ES). Dotted
133 lines indicate biostratigraphic boundaries; black solid lines mark key surface correlations (FS, ES);
134 blue line = MFS C2; red line = major erosional surface (ES3) at the base of the Colorado Delta.
135 White circles mark sample locations. Vertical scales differ between wells.

136

137 **Stratigraphic arrangement**

138 The Moczygemba VT #11 and Vogelsang Frieda #1 cores provide valuable records of the Lower
139 Wilcox Rockdale Delta system, which consists of eight depocenters reaching thicknesses of up to
140 1500 m (Fisher and McGowen, 1967; Zeng and Olariu, 2024). The Vogelsang Frieda #1 core
141 sampled the Colorado Delta, while the Moczygemba #11 core is within the San Antonio Delta
142 (Fig. 1). The biostratigraphic correlation is defined by time lines represented in Figure 1 by the
143 base of *Pistillipollenites mcgregorii* and the top of *Momipites dilatatus*. The biostratigraphic
144 correlation clearly indicates that there is an expanded stratigraphic section toward the north east
145 associated with the Colorado Delta implying higher sedimentation rates in this region.

146 The Vogelsang Frieda #1 core, spanning 685 m, records 14 transgressive–regressive cycles, with
147 individual cycle thickness ranging from 18 to 110 m (Zhang et al., 2022). These cycles document
148 alternating phases of delta progradation and marine transgression, driven by changes in sediment
149 supply, relative sea level, and coastal dynamics. Wave-dominated deposits constitute the largest
150 proportion (48%) of the Vogelsang core stratigraphy, followed by river-dominated deposits (40%)
151 and tide-influenced facies (12%). Each T-R cycle progresses from wave- or river-dominated delta-
152 front deposits (9–75 m thick) to fluvial or fluvial-tidal channel fills (8–33 m thick), capped by
153 wave- or tide-dominated transgressive deposits (3–30 m thick) (Zhang et al., 2022) (Fig. 1).

154 Lithologically, both cores consist of interbedded sandstones, siltstones, and mudstones with
155 organic-rich horizons and bioturbated intervals. Sandstone intervals feature sedimentary structures
156 such as hummocky cross-stratification (HCS), swaley cross-stratification (SCS), and wave ripples,
157 indicating storm-wave reworking of deltaic and shoreface sands. Thin tidal rhythmites within
158 heterolithic deposits suggest intermittent tidal influence in estuarine settings (Galloway et al.,
159 2000; Zhang et al., 2022).

160 The supplementary material accompanying this manuscript provides further details on the
161 depositional environments of Moczygemba VT #11 and Vogelsang Frieda #1 cores.

162

163 **Age model**

164 Age control for both wells is provided by nannofossil biostratigraphy (Moczygemba VT #11 only)
165 and palynological biostratigraphy (both Moczygemba VT #11 and Vogelsang Frieda #1).

166 Estimated absolute ages are based on Geologic Time Scale 2012 (Gradstein et al. 2012) ages for
167 nannofossil events and based on Crabaugh and Elsik (2000), Elsik and Crabaugh (2001), and Zarra
168 et al. (2019) for palynological events. The calcareous nannofossil biozonation follows Martini

169 (1971) and the palynological biozonation follows [Zarra et al. \(2019\)](#). Absolute ages for T
170 *Insulapollenites rugulatus* and B *Pistillipollenites mcgregorii* in [Zarra et al. \(2019\)](#) were estimated
171 based on their height relative to the chronostratigraphic column. An age model based on the
172 biostratigraphic data is provided in the Supplementary Materials. We provide absolute age
173 estimates as minimum and maximum ages for each sample; ages with higher uncertainties appear
174 in parentheses.

175

176 MATERIAL AND METHODS

177 Sample collection

178 Core samples were collected for palynological and geochemical analysis from both the
179 Moczygemba VT #11 and Vogelsang Frieda #1 wells (Fig. 1, Supplementary Material).

180

181 Palynology

182 All palynological samples were subjected to standard acid preparation involving treatment with
183 warm hydrochloric acid (HCl) to remove carbonates and warm hydrofluoric acid (HF) to dissolve
184 silicates. Processing techniques are similar to those described by [Traverse \(2007\)](#). If sufficient
185 organic matter was present, three slides were produced per sample, the first containing an un-
186 sieved and unoxidized fraction for kerogen analysis, and the other two oxidized and sieved with a
187 10 µm aperture nylon mesh for palynological analysis. For lower abundance samples (<300
188 palynomorphs in a slide), all specimens were counted; for higher abundance samples, a modified
189 version of the cascading count technique described by [Styzen \(1997\)](#) was used to estimate the total
190 number of palynomorph specimens present on the slide.

191 For Moczygamba VT #11, the full palynological counts were simplified for linear discriminant
192 analysis (LDA) using the PAleontological STatistics (PAST) software (Hammer et al. 2001).
193 Vogelsang Frieda #1 was not analyzed using LDA because of generally lower palynomorph
194 abundances which introduce more statistical noise. Due to low dinoflagellate cyst abundances in
195 Moczygamba VT #11, all dinoflagellate cyst taxa have been combined into a single type (“total
196 dinoflagellate cysts”). Fungal hyphae have been excluded from the analysis and all other fungal
197 remains have likewise been combined into a single type (“fungal remains”). Algal remains aside
198 from dinoflagellate cysts and acritarchs have been excluded from the analysis. Pollen and plant
199 spore taxa with less than 200 total occurrences in Moczygamba VT #11 have been excluded from
200 analysis. These rare taxa were excluded to ensure that there were a higher number of samples than
201 variables (i.e., taxa) in the matrix, and because abundance counts of rare taxa are inherently more
202 prone to random variability. The simplified palynological counts for both wells and detailed results
203 for the linear discriminant analysis are provided in the Supplementary Materials.

204

205 **Micropaleontology**

206 A total of 28 samples from the Vogelsang Frieda #1 core were analyzed for foraminifera. The
207 original samples and disaggregated residues were weighted, and all samples were washed in water
208 over a 63 μm sieve and dried in an oven. All specimens were selected, systematically counted, and
209 mounted on a microslide. The absolute abundance of foraminifera was obtained by standardizing
210 to 1 gram of sediment (forams/1 gr Sed).

211

212 **Rock-Eval pyrolysis**

213 The quality and quantity of organic matter (OM) was analyzed in 97 bulk sample powders using a
214 Rock-Eval 6 instrument at the Institute of Earth Sciences, University of Lausanne (ISTE-UNIL),
215 following the methodology outlined by Behar et al. (2001) and utilizing the IFP 160 000 standard.
216 Sample aliquots were placed in an oven and initially heated to 300°C under an inert atmosphere
217 before undergoing gradual pyrolysis up to 650°C. After pyrolysis, the samples were transferred to
218 another oven and progressively heated to 850°C in the presence of air, with CO₂ and hydrocarbon
219 (HC) concentrations monitored throughout the process. The calculated parameters included total
220 organic carbon content (TOC, wt.%), hydrogen index (HI in mg HC g⁻¹ TOC), oxygen index (OI
221 in mg CO₂ g⁻¹ TOC), and T_{max} (°C), following the methods of Espitalié et al. (1985) and Behar et
222 al. (2001).

223

224 **Organic carbon isotopes**

225 The carbon isotope composition of organic matter ($\delta^{13}\text{C}_{\text{org}}$) in 155 samples was determined at the
226 stable isotope laboratory of the Institute of Earth Surface Dynamics, University of Lausanne
227 (IDYST-UNIL). Samples were first decarbonated using 10% v/v HCl, then thoroughly washed
228 with deionized water and dried at 40°C for 48 hours. $\delta^{13}\text{C}_{\text{org}}$ measurements were conducted using
229 a Carlo Erba 1100 (Fisons Instruments, Milan, Italy) elemental analyzer, coupled to a Thermo
230 Fisher Scientific Delta V Plus isotope ratio mass spectrometer, both operating under continuous
231 helium flow. The measured $\delta^{13}\text{C}$ values were calibrated and normalized using international
232 reference materials and in-house standards (Spangenberg, 2006, 2016) and are reported in per mil
233 (‰) relative to the Vienna Pee Dee Belemnite limestone standard (VPDB). The precision of $\delta^{13}\text{C}_{\text{org}}$
234 values was better than 0.1‰.

235

236 **X-ray Fluorescence (XRF) analysis**

237 Major element concentrations were determined using a Bruker Tracer IV ED-XRF instrument
238 equipped with an Rh x-ray tube and pXRF technology and excitation voltage of 15 kV, at the
239 Bureau of Economic Geology, UT Austin, Texas. The instrument operated with a He purge and
240 utilized a filter composed of 0.006-inch Cu and 0.001-inch Ti. Before analysis, the core was
241 cleaned with hot water and a plastic brush to eliminate potential contaminants like salts that might
242 have crystallized over time. The pXRF scans were regularly checked for Cl peaks, which could
243 interfere with the Compton peak used in peak normalizations. Samples showing excessive Cl were
244 cleaned again and reanalyzed. Calibration of the pXRF measurements was conducted using
245 semiquantitative methods described by [Rowe et al. \(2012\)](#).

246
247 **Paleo-weathering**

248 To evaluate weathering trends, we applied established weathering proxies that have been used on
249 both terrestrial and marine sediments (e.g., [Hessler et al., 2017](#); [Deng et al., 2022](#); [Sharma et al.,](#)
250 [2024](#)), including the Chemical Index of Alteration (CIA, in %, Eq. 1) and the Chemical Index of
251 Weathering (CIW, in %, Eq. 2), originally proposed by [Nesbitt and Young \(1982\)](#) and [Harnois](#)
252 [\(1988\)](#), respectively.

253 The CIA was used to quantify the degree of weathering by calculating the molar ratio of immobile
254 Al_2O_3 to the mobile oxides CaO, Na_2O , and K_2O within the silicate fraction. The CIA is also used
255 to evaluate environmental factors such as temperature, precipitation, elevation, and slope, which
256 influence silicate weathering.

257
$$\text{CIA} = \frac{\text{Al}_2\text{O}_3}{\text{Al}_2\text{O}_3 + \text{Na}_2\text{O} + \text{CaO}^* + \text{K}_2\text{O}} \times 100 \quad (1)$$

258 Where CaO* represents the CaO incorporated in the silicate fraction and is determined as: CaO*
259 = CaO – (10/3 × P₂O₅).

260 Under intense weathering, mobile oxides are removed, leading to an increased concentration of
261 Al₂O₃, with CIA values approaching nearly 100 wt.%. Conversely, weak weathering results in
262 lower CIA values (<50%) due to the prevalence of mobile oxides.

263 For comparison, we also applied the CIW, which is a modified version of the CIA, designed to
264 account for potassium metasomatism in sediments (e.g., [Stein et al., 2021](#)).

$$265 \quad \text{CIW} = \frac{\text{Al}_2\text{O}_3}{\text{Al}_2\text{O}_3 + \text{Na}_2\text{O} + \text{CaO}^*} \times 100 \quad (2)$$

266

267 **Mean annual precipitation**

268 Mean annual precipitation (MAP, in mm yr⁻¹, Eq. 3) was estimated using the CIW (Eq. 2) and is
269 based on the equation proposed by [Sheldon et al. \(2002\)](#), with a standard error of ±182 mm yr⁻¹.

$$270 \quad \text{MAP} = 221e^{0.0197(\text{CIW})} \quad (3)$$

271

272 **Uncertainty estimates**

273 All data presented in this study include uncertainties expressed as the standard error of the mean
274 (SE), calculated using the formula $\text{SE} = \text{SD}/\sqrt{n}$, where SD represents the standard deviation and n
275 is the number of replicates analyzed. Although uncertainty estimates are provided, they are often
276 not clearly visible on the plots due to their small magnitude relative to the plot scale.

277

278 **RESULTS**

279 **Biostratigraphy**

280 Palynological biostratigraphy is largely based on the biostratigraphic scheme developed by [Zarra](#)
281 [et al. \(2019\)](#) for the deepwater Wilcox. However, it was found that an event-based biostratigraphy
282 was more suitable to the studied onshore Lower Wilcox ([Smith et al., 2025; in press](#)) wells than
283 the largely assemblage-based zonation in [Zarra et al. \(2019\)](#). An alternative palynological zonation
284 for the onshore Lower Wilcox, with the boundaries defined by individual events, has therefore
285 been used.

286 Aside from the events “*T Momipites dilatus*” and “*T Caryapollenites* spp.,” the other zonal
287 boundaries in our alternative zonation are used as markers in [Zarra et al. \(2019\)](#). “*T Momipites*
288 *dilatus*” and “*B Caryapollenites* spp.” (considered synonymous with “*Carya* spp. <30 µm Base”)
289 were used by [Crabaugh and Elsik \(2000\)](#) and [Elsik and Crabaugh \(2001\)](#) in their onshore Wilcox
290 Group zonation. “*T Momipites dilatus*” is early Thanetian in age, near the top of the lower Wilcox
291 as defined by [Crabaugh and Elsik \(2000\)](#). “*T Caryapollenites* spp.” is earliest Selandian in age,
292 slightly above the Midway Group/Wilcox Group boundary. For the purposes of this study, we have
293 only used previously published biostratigraphic events.

294 The top analyzed sample in Moczygamba VT #11 at 1405.13 mbs (meters below surface) is
295 Thanetian in age, in the “*T Momipites dilatus*/*T Insulapollenites rugulatus*” zone (Fig. 1).
296 Nannofossil recovery in the top three analyzed samples in Moczygamba VT #11 was insufficient
297 for age determination. The highest sample with sufficient nannofossil recovery is at 1410.74 mbs
298 and is in nannofossil zone NP7. “*T Momipites dilatus*” was observed at 1485.60 mbs, slightly
299 above the NP6/NP7 boundary at 1486.81 mbs. The lower NP6/upper NP6 boundary at 1492.81
300 mbs is an approximate marker for the Selandian/Thanetian boundary. The primary palynological
301 marker event “*B Pistillipollenites mcgregorii*” was observed at 1508.61 mbs, slightly above the
302 questionable base of NP6 at 1513.23 mbs (Fig. 1). The sample at 1516.79 mbs contains a probably

303 NP5 age nannofossil assemblage and the observed base of multiple palynomorph species,
304 including the primary marker event “B *Caryapollenites veripites*.” A large (ca.300 m) sample gap
305 between 1516.79 mbs and 1843.13 has been left unzoned. However, this large unsampled section
306 is constrained to be Selandian in age based on the presence of *Caryapollenites* spp. in downsection
307 samples. Nannofossil recovery in the bottom section below the large sample gap is insufficient for
308 age determination, but based on the presence of *Caryapollenites imparalis* in the bottom sample
309 (1851 mbs), the bottom section is early Selandian in age, in the “B *Caryapollenites* spp./B
310 *Caryapollenites veripites*” zone.

311 Vogelsang Frieda #1 was not sampled for nannofossil analysis and biostratigraphic age control
312 relies on palynology. Although the recovery and preservation of palynomorphs was generally
313 better in the Moczygemba VT #11 samples, the relatively high and continuous sampling resolution
314 in Vogelsang Frieda #1 makes it an excellent reference well for palynological biostratigraphy.
315 Near the top of the analyzed section at 8328.50 mbs, the primary marker “T *Insulapollenites*
316 *rugulatus*” was observed, possibly below its true stratigraphic top. The top two samples (at 8272
317 mbs and 8282 mbs) are questionably in the Thanetian “T *Insulapollenites rugulatus/Apectodinium*
318 acme” zone, the uncertainty resulting from the rarity of the marker taxon *I. rugulatus* near its top
319 and the fairly low abundances in the top two samples. “T *Momipites dilatus*,” an early Thanetian
320 marker, was observed at 8622 mbs. Palynological biostratigraphy is not able to precisely delineate
321 the Selandian/Thanetian boundary, but it lies within the “B *Pistillipollenites mcgregorii*/T
322 *Momipites dilatus* zone (8622-9092.50 mbs) (Fig. 1). The primary marker “B *Caryapollenites*
323 *veripites*” was observed at 9275 mbs, at the same depth as the base of *Spinaepollis spinosa* and
324 *Momipites triradiatus* type, possibly suggestive of an unconformity between 9275.00 – 9281.50
325 mbs. Below 9275 mbs, palynological abundances are generally lower than upsection, reducing the

326 confidence in the placement of bioevents below 9275 mbs. The earliest Selandian event “B
327 *Caryapollenites* spp.” was questionably observed near the base of the analyzed section at 10488
328 mbs; only a single downsection sample was analyzed, and the absence of *Caryapollenites* spp. in
329 that sample may be due to low general palynomorph abundance rather than true regional absence.

330

331 **Foraminiferal assemblages**

332 The foraminiferal assemblages studied in the Vogelsang Frieda #1 core determined that in all the
333 samples analyzed only benthic foraminifera were found, of which practically all are agglutinated
334 foraminifera such as *Haplophragmoides* spp., *Reophax* sp, and *Ammobaculites* sp. Most of the
335 analyzed foraminifera display poor preservation characterized by pyritized tests, in which sutures
336 and chambers are not recognizable.

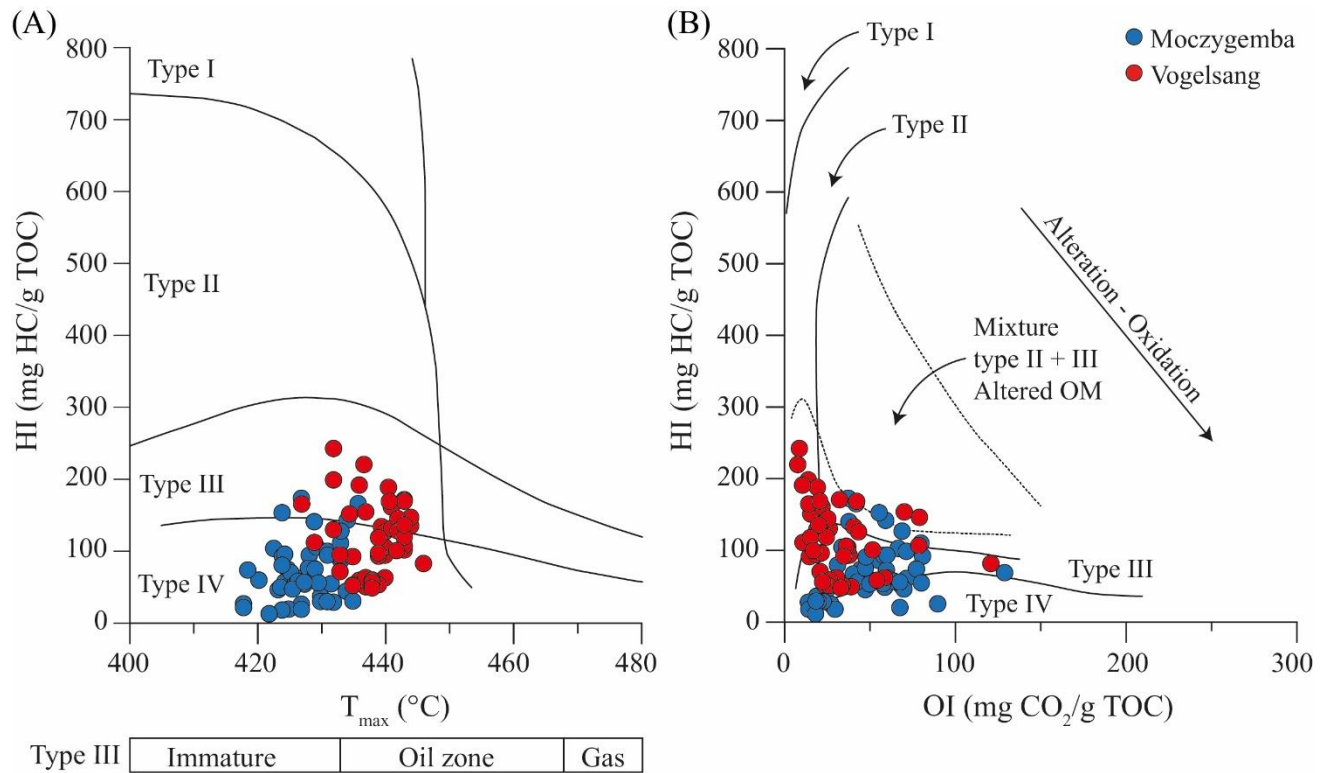
337

338 **Organic matter content and characterization**

339 The total organic carbon (TOC) content in both sections has an average value of 1.75 ± 0.09 wt.%
340 (N = 155). In the Moczygamba section, TOC values range from 0.24 ± 0.09 to 4.95 ± 0.09 wt.%,
341 averaging 1.78 ± 0.09 wt.% (N = 109). In contrast, TOC values in the Vogelsang section range
342 from 0.67 ± 0.22 to 6.76 ± 0.22 wt.%, with an average of 1.83 ± 0.22 wt.% (N = 46).

343 To classify organic matter (OM) by type (origin) and thermal maturity, T_{\max} values and hydrogen
344 index (HI)/oxygen index (OI) ratios were analyzed (Espitalié et al., 1985). T_{\max} values range from
345 418 to 446°C, suggesting that the preserved OM is predominantly immature or within the oil
346 window (Fig. 2). HI values in both Moczygamba and Vogelsang are generally below 200 mg HC/g
347 TOC (average 96.25 mg HC/g TOC), indicating that the OM is primarily Type III and IV,
348 characteristic of a strong terrestrial plant input.

349 OI values are lower compared to HI, generally remaining below 100 mg CO₂/g TOC (average 42.5
 350 mg CO₂/g TOC), except two samples which have values up to 150 mg CO₂/g TOC. Overall, the
 351 Rock-Eval parameters suggest that the organic matter in both sections (Moczygamba and
 352 Vogelsang) originates from a recycled and/or terrestrial source.



353 Type III Immature Oil zone Gas

354 **Figure 2. Kerogen type and thermal maturity assessment of organic matter.** (A) Hydrogen
 355 Index (HI) vs. T_{max} plot showing the classification of kerogen types (I–IV) and thermal maturity
 356 for Moczygamba (blue circles) and Vogelsang (red circles) samples. Most samples classify as
 357 Type III or Type IV (immature and oil zone) organic matter. (B) Hydrogen Index (HI) vs. Oxygen
 358 Index (OI) plot illustrating the kerogen type distribution and the effects of oxidation and alteration.
 359 The majority of samples cluster in the Type III–IV range.

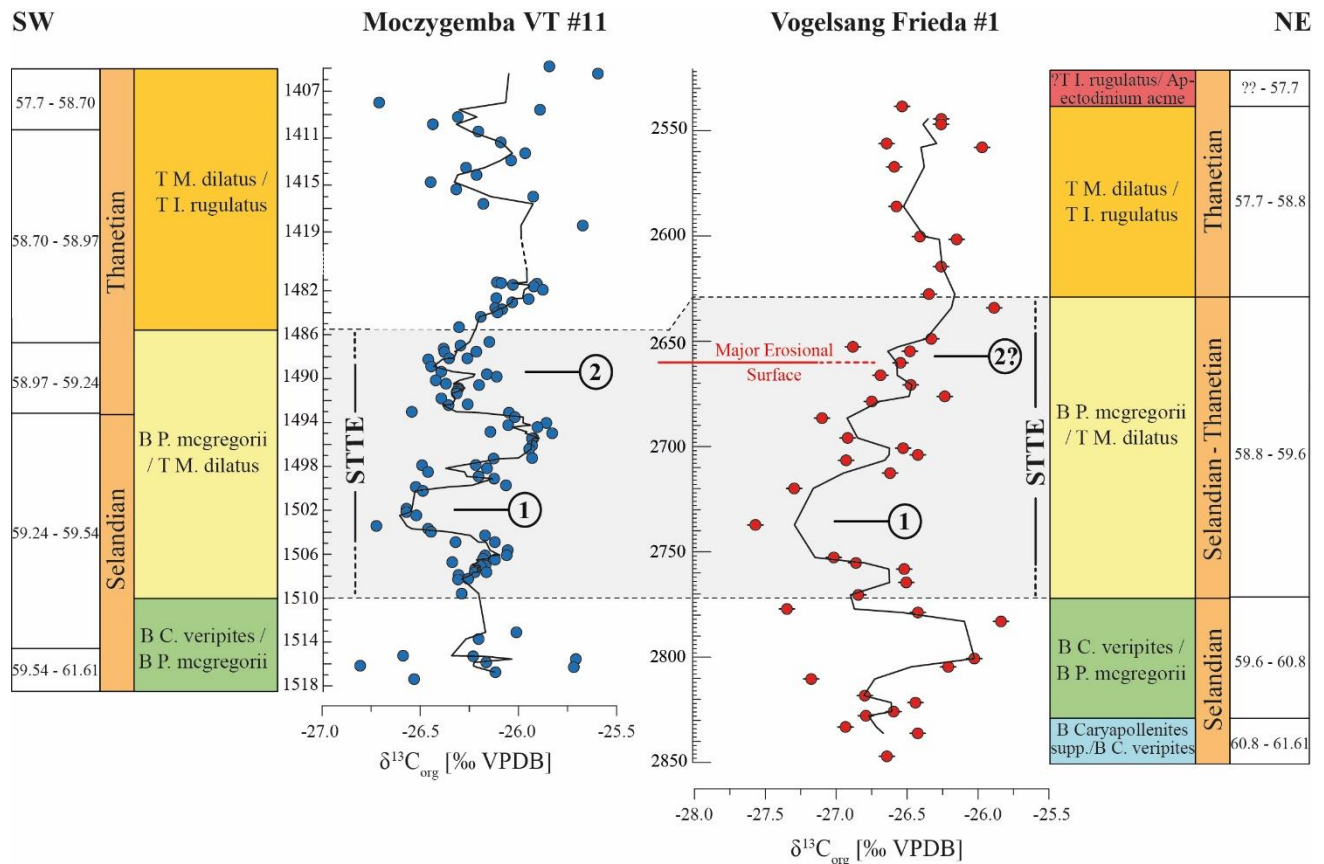
360

361 **Organic carbon isotopes**

362 The organic carbon isotope ($\delta^{13}\text{C}_{\text{org}}$) records from the Moczygamba VT #11 and Vogelsang Frieda
363 #1 cores provide insights into the carbon cycle dynamics across the Selandian-Thanetian transition.
364 These data offer valuable constraints on paleoenvironmental conditions and potential climatic
365 perturbations during the Paleocene.

366 The $\delta^{13}\text{C}_{\text{org}}$ records from the two cores exhibit distinct yet comparable trends across the studied
367 stratigraphic intervals. In Moczygamba VT #11, values range between -27.0 ± 0.02 and $-25.5 \pm$
368 0.02‰ VPDB (average value of $-26.2 \pm 0.02\text{‰}$; $N = 109$), displaying notable fluctuations. A
369 pronounced negative carbon isotope excursion (CIE) of 0.5‰ magnitude is identified between
370 depths 1498–1504 m, followed by a subsequent recovery until 1494 m and a second negative CIE
371 of 0.4‰ magnitude between depths 1486 – 1494 m. These negative shifts suggest a perturbation
372 in the global carbon cycle, or may reflect changes in carbon sources or burial processes (Fig. 3).

373 In Vogelsang Frieda #1, $\delta^{13}\text{C}_{\text{org}}$ values range between -28.0 ± 0.06 and $-25.5 \pm 0.06\text{‰}$ VPDB
374 (average value of $-26.6 \pm 0.06\text{‰}$; $N = 46$), with slightly more negative values compared to
375 Moczygamba VT #11. A negative CIE of ca. 1‰ magnitude is evident in the lower part of the
376 section (2700–2750 m) spanning the Selandian-Thanetian transition, closely corresponding to the
377 negative excursion observed in Moczygamba VT #11. A major erosional surface in Vogelsang
378 Frieda #1 coincides with another possible negative shift of 0.2‰ magnitude, noted between depths
379 2650–2670 m, though its exact stratigraphic correlation remains uncertain (Fig. 3).



380

381 **Figure 3. Organic carbon isotope records across the Selandian-Thanetian transition.** $\delta^{13}C_{org}$

382 profiles from the Moczygemba VT #11 (blue circles) and Vogelsang Frieda #1 (red circles) cores

383 plotted against depth, with biostratigraphic and lithostratigraphic frameworks. Key biozones and

384 transitions between the Selandian and Thanetian stages are highlighted, with notable negative

385 carbon isotope excursions (CIEs) indicated with a 5-point moving average. A major erosional

386 surface observed in the Vogelsang Frieda #1 core is also marked (see Figure 1 for more detail).

387

388 **Weathering and annual precipitation estimates**

389 The Chemical Index of Alteration (CIA) profiles for the Moczygemba VT #11 and Vogelsang

390 Frieda #1 cores reveal distinct weathering patterns across the Selandian–Thanetian interval. In

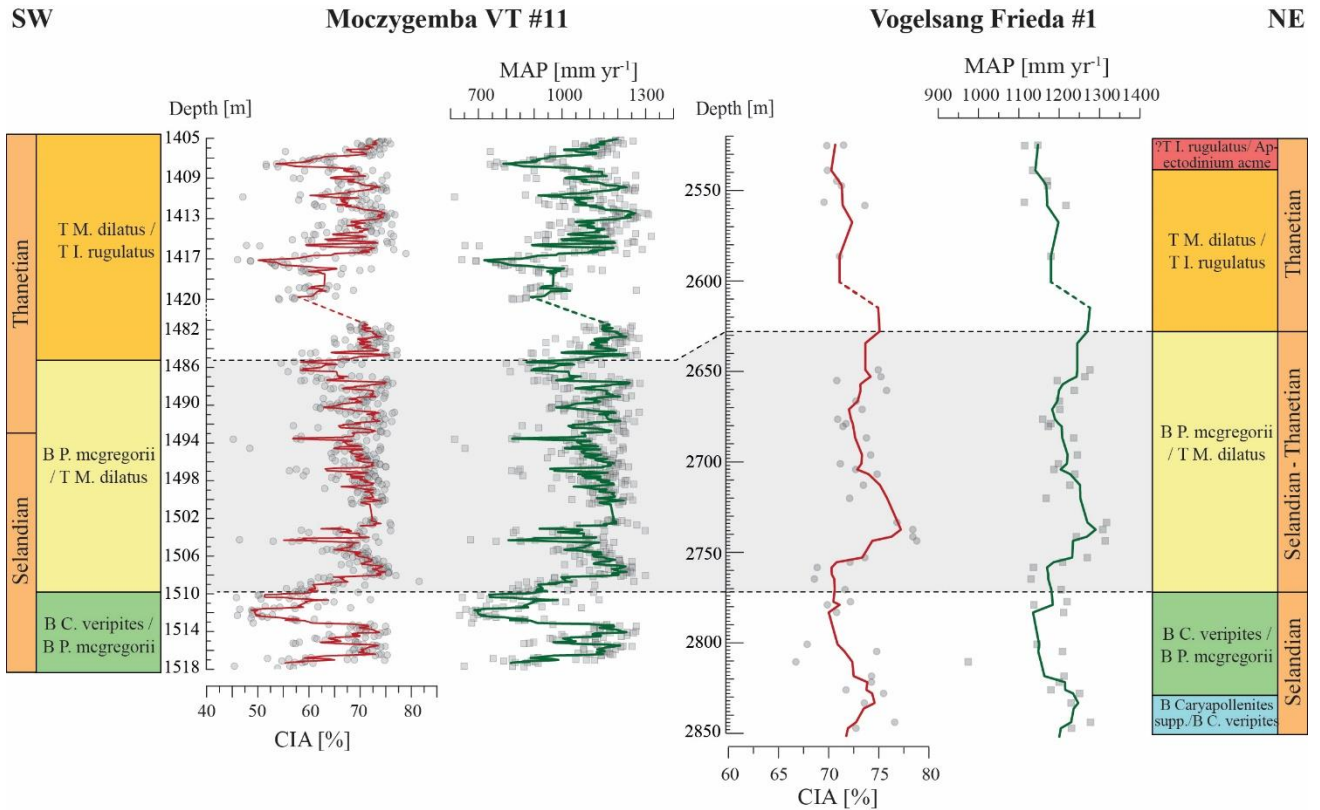
391 Moczygemba VT #11, CIA values range from 45 to 82%, with an average of 67%, while values

392 in Vogelsang Frieda #1 range from 61 to 79%, averaging 72% (Fig. 4). These values are broadly
393 indicative of moderate to intense chemical weathering and, if accurate, suggest a sustained
394 prevalence of such weathering along the Gulf Coast during this interval. CIA trends across the
395 broader Paleogene succession support this interpretation. During the Paleocene, values remain
396 relatively stable between 69 and 77, approaching but generally staying just below the chemical
397 alteration threshold (CIA = 77). At the Paleocene–Eocene boundary, values increase and cross this
398 threshold, reaching 77 to 83, indicating intensified weathering likely driven by global warming.
399 This peak is followed by a notable decline during the Oligocene, with CIA values falling to 45–
400 65, reflecting diminished weathering intensity under cooler climatic conditions. The CIA values
401 from both cores are consistent with late Paleocene to early Eocene estimates reported by [Hessler](#)
402 [et al. \(2017\)](#) and highlight the strong link between chemical weathering intensity and major
403 Paleogene climate transitions.

404 Mean Annual Precipitation (MAP) estimates from the Gulf Coast cores indicate persistently humid
405 conditions during the late Paleocene to early Eocene, with values ranging from 615 to 1330 mm
406 yr⁻¹ (average 1070 mm yr⁻¹) in the Moczygemba VT #11 core and from 960 to 1380 mm yr⁻¹
407 (average 1200 mm yr⁻¹) in the Vogelsang Frieda #1 core (Fig. 4). These estimates are broadly
408 consistent with those from southwestern Wyoming, where paleobotanical data indicate MAP
409 values between 1300 and 1500 mm yr⁻¹ during most of the late Paleocene to early Eocene, followed
410 by a decline to approximately 800 mm yr⁻¹ under more seasonal and arid conditions later in the
411 interval ([Wilf, 2000](#)). High-resolution climate modeling by [Sewall and Sloan \(2006\)](#) similarly
412 reconstructs a humid, tropical to subtropical monsoonal climate across south-central North
413 America during this time, including the southern coastal plain and much of the Laramide foreland.
414 Their simulations suggest that precipitation in these regions was predominantly convective or

415 orographic and sourced from the paleo-Gulf of Mexico. Together, these proxy- and model-based
416 records support the interpretation of warm, wet conditions across much of the continent during the
417 late Paleocene to early Eocene, though with pronounced regional variation controlled by
418 paleogeography, elevation, and atmospheric circulation patterns.

419 The overall trends in both cores suggest an initial phase of moderate chemical weathering that
420 intensified across the Selandian–Thanetian transition. This shift is mirrored by the highest MAP
421 estimates, indicating a period of enhanced precipitation. Notably, these peaks in weathering and
422 precipitation coincide with negative carbon isotope excursions (CIEs) recorded in both the
423 Moczygamba VT #11 and Vogelsang Frieda #1 cores, linking increased hydroclimatic intensity to
424 short-term carbon cycle perturbations. Following this interval, both CIA and MAP values return
425 to more moderate levels.



426

427 **Figure 4. Chemical weathering and precipitation trends.** Lateral correlation of CIA values and
 428 Mean Annual Precipitation (MAP) estimates, represented using a 5-point moving average, for the
 429 Moczygamba VT #11 and Vogelsang Frieda #1 cores. The CIA profiles (left panels) indicate
 430 variations in chemical weathering intensity, while MAP estimates (right panels) indicate changes
 431 in precipitation from the Selandian to the Thanetian. Key biostratigraphic markers are indicated
 432 and shaded intervals highlight correlative sections between the two cores.

433

434 **DISCUSSION**

435 **Paleoecology**

436 *Palynomorphs*

437 It would normally be expected that palynomorphs produced by more thermophilic organisms
 438 would become more common during hyperthermal events, with the caveat that various other

439 factors (differing hydrodynamic properties of palynomorph types, differing susceptibility to
440 degradation by oxidation, etc.) may also influence relative palynomorph abundances. We
441 investigated changes in relative palynomorph abundance by splitting the palynological samples
442 from Moczygamba VT #11 into three groups: a group above the carbon isotope excursion (CIE)
443 near the Selandian-Thanetian boundary (1405-1482 m), a group in the CIE in the Selandian and
444 Thanetian (1482-1505 m), and a group below the CIE in the Selandian (1505-1851 m). Because
445 of the high diversity in palynomorph taxa, the raw specimen counts have been simplified as
446 described in the methods section and analyzed using LDA (Fig. S3, Supplementary Material).

447 A plot of the first two axes of the LDA for the Moczygamba VT #11 samples is shown in the
448 Supplementary Material files (Fig. S3). The three groups have been outlined with convex hulls
449 and show significant separation. Biplots of the taxa are overlain on the discriminant axis scores for
450 the samples, the length and direction of the vectors indicating the significance of particular taxa in
451 the LDA. The samples in the CIE plot with generally positive discriminant axis scores for both
452 axis 1 and axis 2; only one sample depth in the CIE (at 1486.81 m) falls inside the convex hull of
453 another group. Aside from this outlier, all CIE samples have more positive axis 2 values than the
454 non-CIE samples. A jackknifed confusion matrix (provided in the Supplementary Materials)
455 successfully classified 55% of the samples.

456 The two taxa with the longest biplot vectors are *Thomsonipollis magnificus* and
457 Betulaceae/Myricaceae type pollen. The biplot vector of *T. magnificus* is nearly vertical upwards,
458 which suggests that higher abundances of *T. magnificus* are associated with the CIE. The
459 Betulaceae/Myricaceae type biplot vector is angled towards the top left quadrant of the graph and
460 suggests that higher abundances of Betulaceae/Myricaceae type pollen are associated with samples
461 above the CIE in the Thanetian. Other taxa with biplots angled towards positive axis 2 values

462 include *Momipites coryloides* (20-27 μm), *Tricolpites hians*, and fungal remains. Taxa with very
463 short biplot vectors, suggesting no significant differences in relative abundance between CIE and
464 non-CIE samples, include *Momipites waltmanensis*, *Classopollis classoides*, and *Deltoidospora*
465 *microadriennis*.

466 Although changes in taxa abundance through time are not necessarily related to paleoclimatic
467 change, the LDA results are considered tentative evidence for paleoclimatically induced changes
468 in palynomorph assemblages related to the Selandian/Thanetian hyperthermal. In particular, *T.*
469 *magnificus* may represent pollen from the Rubiaceae family (Elsik 1968) and has previously been
470 associated with mangrove paleoenvironments (Lenz et al. 2021). Higher abundances of *T.*
471 *magnificus* may represent an expansion of mangrove paleoenvironments during the CIE.
472 *Momipites coryloides* (20-27 μm) is a form species for pollen morphologically similar to modern
473 *Engelhardia* pollen (Nichols and Ott 1978). Although the extant genus *Engelhardia* is mainly
474 restricted to tropical montane environments, the paleoecological preferences of plants producing
475 *Momipites* pollen in the Paleogene United States appear to be broader, and *Momipites* pollen is
476 abundant in many lowland swamp paleoenvironments in Paleogene strata (Frederiksen 1985). The
477 nearly vertical upwards biplot for *M. coryloides* (20-27 μm) suggests an association with the CIE
478 samples.

479 The Betulaceae/Myricaceae type is considered to mainly represent pollen from the Betulaceae and
480 Myricaceae families, although identification to modern genera in the Paleogene is problematic
481 (e.g., Jardine 2011). Harrington (2008) observed very high abundances of Betulaceae/Myricaceae
482 type pollen in samples interpreted as representing swamp paleoenvironments in the Paleocene and
483 Eocene of Alabama and suggested that the ecology may be similar to extant coastal strands of
484 *Myrica* shrubs along the Atlantic coast of the United States (Crawford and Young 1998).

485 Betulaceae/Myricaceae type pollen is associated with post-CIE samples in the LDA graph and may
486 indicate this pollen type is derived from less thermophilic swamp vegetation. *Tricolpites hians*
487 pollen may have an affinity with the Platanaceae (Pocknall and Nichols 1996) but may have been
488 produced by other plant families; in the LDA graph it appears associated more with CIE and post-
489 CIE samples. The biplot vector for fungal remains is angled in the direction of the CIE samples
490 and suggests an association with the CIE.

491 In order to confirm the significance of the taxa biplots, average relative abundances of selected
492 taxa were calculated for the three sample groups in Moczygamba VT #11 (pre-CIE, CIE, and post-
493 CIE) (see Supplementary Material). In general, these abundances confirm the interpretations from
494 the LDA graph. Average relative abundances of *T. magnificus*, for example, were ca.7.4% in the
495 post-CIE samples, ca.12.7% in the CIE samples, and ca.6.3% in the pre-CIE samples. Relative
496 abundances of *T. magnificus* (relative only to the pollen and plant spore assemblage) have been
497 graphed for both Moczygamba VT #11 and Vogelsang Frieda #1 (Figure S1). *T. magnificus*
498 abundances in Moczygamba VT #11 increase during the Selandian/Thanetian CIE, even exhibiting
499 a double peak at approximately the same depths as the double peak in the carbon isotope data.
500 Although the biplot vector for total dinoflagellate cysts in Moczygamba VT #11 is small and
501 angled towards the upper left quadrant, the raw counts do appear to show two peaks in the CIE
502 section at approximately the same depths as the double peak in *T. magnificus* abundances and
503 carbon isotope ratios (Figure S3), suggesting a potential rise in relative sea level during these
504 hyperthermal events. This relationship is not apparent from the LDA graph (Figure S3), likely
505 because, aside from these two peaks in the CIE section in Moczygamba VT #11, dinoflagellate
506 cyst counts are quite low. An increase in *T. magnificus* abundances is not clearly observed in the
507 CIE of Vogelsang Frieda #1; this may be related to low total pollen abundances in that well

508 introducing more statistical noise, or this could be related to differences in pollen source areas
509 between the two wells.

510 The relative abundance of fungal remains (calculated relative to the total fungal, pollen, and plant
511 spore assemblage) is higher in the pre-CIE samples than the CIE in Moczygamba VT #11. A
512 notable increase in abundance of fungal spores was observed in a Paleocene-Eocene Thermal
513 Maximum (PETM) section from the southern Gulf of Mexico in the Chicxulub impact crater and
514 interpreted as an indicator for more humid conditions (Smith et al. 2020); in both Moczygamba
515 VT #11 and Vogelsang Frieda #1 there is a general trend towards higher fungal abundances
516 downsection in the Selandian (Figure S3) but there is no clear acme associated with the CIE. There
517 is an overall trend towards decreasing relative abundances of *Classopollis classoides* in the
518 younger Thanetian sections of both Moczygamba VT #11 and Vogelsang Frieda #1, which could
519 be interpreted as a regional contraction of arid coastal paleoenvironments, possibly related to sea
520 level change. However, the decline and eventual extinction of the Cheirolepidiaceae (the plant
521 family which produced *Classopollis* pollen) in the Paleogene globally suggests the decline in
522 *Classopollis* abundance in these wells may be related to competition by angiosperm plants for
523 similar ecological niches rather than a local change in paleoenvironmental conditions (Smith et al.,
524 2024). In summary, the clearest palynological assemblage change during the CIE in Moczygamba
525 VT #11 is an increase in *T. magnificus*, possibly indicating an expansion of mangrove
526 paleoenvironments in response to global warming.

527

528 *Foraminifera and Nannofossils*

529 The foraminiferal assemblages in Vogelsang Frieda #1 are entirely agglutinated (Figure S1),
530 showing extremely low diversities and strong dominance of *Haplophragmoides* spp.,

531 *Ammobaculites* sp., and *Reophax* sp. These taxa are associated with an infaunal life mode, which
532 inhabits below the sediment surface and can tolerate low-oxygen conditions and high nutrient
533 availability (Kaminski and Gradstein, 2005). The shallow marine to deltaic environment in this
534 section suggests that the agglutinated foraminifera were limited for short time periods and adapted
535 to restricted conditions (different from the normal marine shelf), where the main limiting factors
536 were probably the depleted oxygen-concentration and/or nutrient availability in bottom waters
537 throughout the Selandian-Thanetian transition.

538

539 **Carbon isotopic record**

540 The observed negative $\delta^{13}\text{C}_{\text{org}}$ excursions may be attributed to several paleoenvironmental and
541 carbon cycle processes. These excursions could be linked to transient warming events, potentially
542 associated with smaller-scale perturbations in the mid to late Paleocene such as the STTE and/or
543 ELPE. The decrease in $\delta^{13}\text{C}_{\text{org}}$ values suggests an increased contribution of isotopically lighter
544 carbon, which could be derived from enhanced terrestrial organic matter influx due to weathering
545 and soil erosion, methane hydrate destabilization, or shifts in oceanic productivity affecting
546 organic carbon burial (e.g., Arthur et al., 1988; Dickens et al., 1995; Meyers, 1997; Pancost et al.,
547 2007, 2013).

548 The lateral comparison of $\delta^{13}\text{C}_{\text{org}}$ profiles from Moczygamba VT #11 and Vogelsang Frieda #1
549 reveals regionally consistent carbon cycle perturbations across the Selandian–Thanetian transition.

550 The presence of negative excursions, supported by biostratigraphic markers, points to
551 paleoenvironmental changes that warrant further investigation. The observed negative carbon
552 isotope excursions (CIEs) align with coeval records from both marine and terrestrial settings. For
553 instance, a -0.6% CIE was identified in the shallow marine Contessa Road section (Gubbio, Italy)

554 (Coccioni et al., 2019), while significantly larger excursions were reported from terrestrial sections
555 such as Lairière in the northern Pyrenees (-3.7‰ ; Maufrangeas et al., 2020) and Cerro Bayo in
556 Argentina (-2.9‰ and -4.5‰ ; Hyland et al., 2015). These terrestrial CIEs are roughly 3–4 times
557 the magnitude of those in shallow marine records, a discrepancy that may reflect differences in
558 carbon reservoir size (Beerling, 2000) and isotopic fractionation between marine and terrestrial
559 organic matter (Sheldon and Tabor, 2009). Alternatively, or additionally, these differences could
560 result from intensified oceanic carbon ventilation and increased petroleum generation in the Gulf
561 of Mexico, analogous to mechanisms proposed for the Early Eocene Climatic Optimum (Hyland
562 et al., 2013).

563 The $\delta^{13}\text{C}_{\text{org}}$ values can also serve as indicators of paleoecology and paleoclimate (Kohn, 2010). C_3
564 plants, which include trees, most shrubs, and cool-season grasses, have $\delta^{13}\text{C}$ values ranging from
565 -37‰ to -20‰ and have historically dominated terrestrial vegetation (Kohn, 2010). This broad
566 range of $\delta^{13}\text{C}$ values in plants is influenced by factors such as temperature, altitude, latitude, and
567 mean annual precipitation (MAP) (Schulze et al., 1996; Kohn, 2010). Non-water-stressed C_3 plants
568 are enriched in ^{12}C , resulting in more negative $\delta^{13}\text{C}$ values, typically below -26‰ . In contrast,
569 higher $\delta^{13}\text{C}$ values ($> -26\text{‰}$) are associated with plants growing under water-limited conditions
570 and low soil transpiration rates ($\text{MAP} < 500 \text{ mm yr}^{-1}$) (e.g., Cerling and Quade, 1993; Kohn, 2010).
571 Measured $\delta^{13}\text{C}_{\text{org}}$ values in both cores suggest the presence of C_3 vegetation during the mid to late
572 Paleocene, confirmed by previous studies such as Cerling and Quade (1993) and Jacobs et al.,
573 (1999). While the local climate remains mainly humid during the deposition of Vogelsang Frieda
574 #1, in Moczygamba VT #11, variations in climate can be observed over the depositional period.
575 Humid conditions correspond to negative CIE (lower $\delta^{13}\text{C}_{\text{org}}$ values) while higher $\delta^{13}\text{C}_{\text{org}}$ values
576 correspond to a dry ecosystem ($\text{MAP ca. } 600 \text{ mm yr}^{-1}$).

577 Apart from paleoecology and paleoclimate interpretations, the presence of isotope excursions
578 allows the lateral correlation of stratigraphic intervals highlighting the use of stable isotopes as a
579 correlation tool in notably difficult to date sub-surface sedimentary successions.

580

581 **Primary versus diagenetic signals**

582 Changes in composition of the preserved organic matter can influence the observed trends in
583 organic carbon isotope composition and it is as such necessary to evaluate potential diagenetic and
584 secondary overprint on the original geochemical signature before paleoenvironmental
585 interpretations can be carried out.

586 For this, first, we assessed the relationship between $\delta^{13}\text{C}_{\text{org}}$ and TOC values (Brasier et al., 1996).

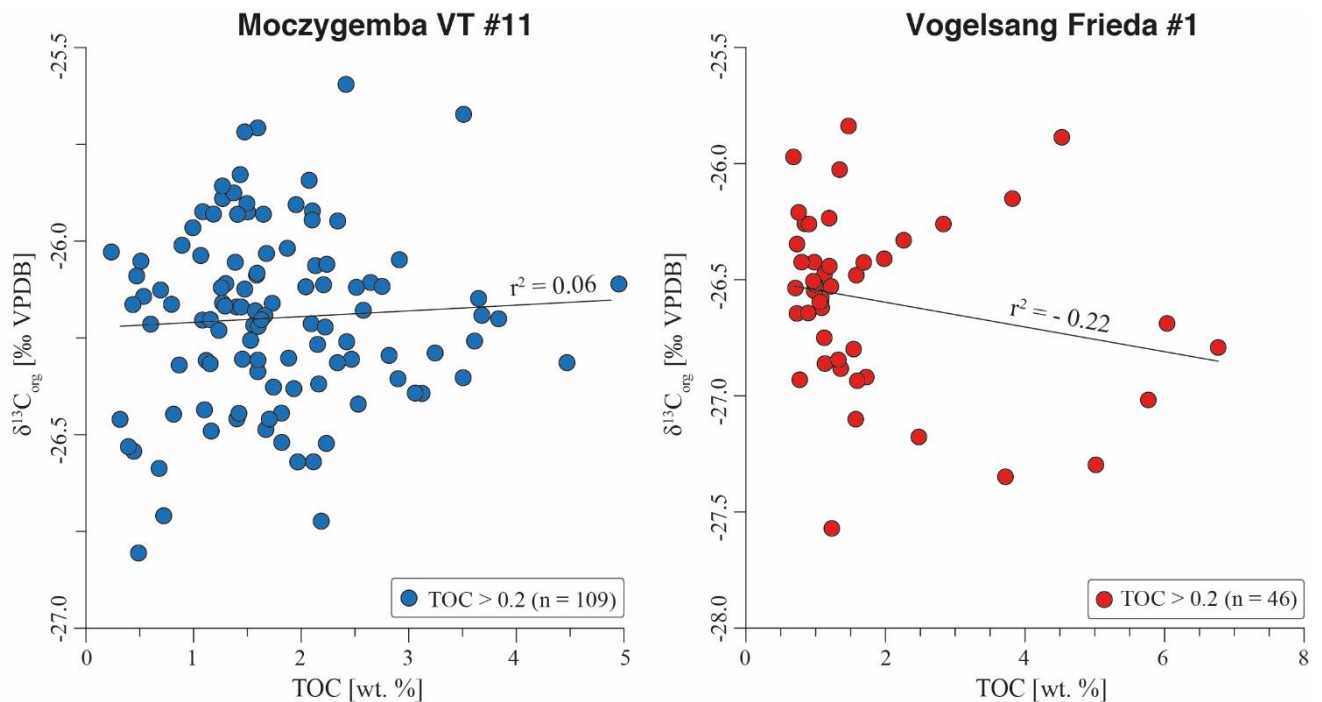
587 Generally, Pearson correlation coefficient (r) less than 0.6 suggest no significant correlation
588 between the isotopic signature and organic matter content (e.g., Fio et al., 2010). In both sections,
589 much lower coefficients were observed ($r < 0.3$) hence there is no significant correlation (Fig. 5).

590 The absence of correlation indicates that the observed negative CIE and the isotopic compositions
591 in general are not due to changes in the TOC content and can therefore be used for
592 paleoenvironmental interpretations (Fig. 5).

593 Another indicator of diagenesis is the maximum temperature (T_{max}) reached during Rock-Eval
594 pyrolysis (S2), which reflects the thermal maturity of organic matter (OM). In samples with
595 relatively high OM content (TOC > 0.5 wt.%; S2 > 0.2), the measured T_{max} values were below
596 440°C (Fig. 5, check Supplementary Material), corresponding to the onset of the oil window
597 (ca.60°C; Espitalié et al., 1985).

598 These two approaches collectively indicate that the primary isotopic signal is largely preserved in
599 both sections. The data presented can therefore be reliably used to reconstruct paleoclimatic

600 conditions and compare the data with available global isotopic records from the pre-PETM
601 hyperthermal events.



602
603 **Figure 5. Correlation between TOC and $\delta^{13}\text{C}_{\text{org}}$ in Moczygamba VT #11 and Vogelsang**
604 **Frieda #1.** Scatter plots showing the correlation between total organic carbon (TOC) content
605 (wt.%) and $\delta^{13}\text{C}_{\text{org}}$ (% VPDB) for Moczygamba VT #11 (left, blue circles) and Vogelsang Frieda
606 #1 (right, red circles). The coefficient of determination (r^2) for each dataset is displayed, indicating
607 a non-significant positive correlation in Moczygamba VT #11 ($r^2 = 0.06$) and a non-significant
608 negative correlation in Vogelsang Frieda #1 ($r^2 = -0.22$).

609
610 **Weathering conditions and regional climate**

611 The observed CIA trends provide valuable insights into the prevailing weathering regimes and
612 regional climate conditions during the Selandian-Thanelian transition. The increasing CIA values
613 in both cores indicate a shift towards more intense chemical weathering, typically associated with
614 warmer and wetter conditions. This trend is consistent with climate reconstructions during global

615 warming periods such as the PETM, which suggest enhanced hydrological cycling in response to
616 global warming and rapid release of carbon (e.g., [Chen et al., 2016](#); [Izumi et al., 2018](#)).

617 The slight disparity in CIA and MAP estimates between the two cores may reflect differences in
618 local depositional environments and sediment flux. The consistently higher CIA and MAP values
619 in Vogelsang Frieda #1 suggest a more humid climate regime than Moczygamba VT #11,
620 potentially due to basin topography and proximity to moisture sources. Alternatively, the elevated
621 values in Vogelsang could result from increased detrital sediment input (higher sedimentation
622 rates), creating a more active depositional environment and thus higher CIA and MAP values. The
623 observed differences might also be similar to the Holocene Gulf Coast fluvial systems that span a
624 climate gradient from semi-arid in the southwest to humid in the northeast ([Milliken et al., 2017](#)).

625 The CIA values estimated here are slightly lower than those estimated for the PETM, which range
626 between 75 to 85 % ([Stokke et al., 2021](#)), but supports the assumption of intense weathering during
627 periods of global warming. The prevailing understanding of Earth's carbon cycle suggests that the
628 negative silicate weathering feedback intensifies in response to rising atmospheric pCO₂ ([Colbourn
629 et al., 2015](#); [Penman et al., 2020](#)). Although, the magnitude of this feedback is largely influenced
630 by various local and regional environmental factors, including temperature, precipitation,
631 geomorphology, and lithology (e.g., [Richey et al., 2020](#); [Deng et al., 2022](#)). Our observed
632 weathering intensities also align with the predicted semi-arid to humid ecosystem inferred from
633 $\delta^{13}\text{C}_{\text{org}}$ values, further supporting our interpretation of a locally semi-arid climate and an enhanced
634 hydrological system along the Gulf Coast during the mid to late Paleocene.

635 The observed regional differences in weathering intensity and precipitation estimates underscore
636 the importance of local paleoenvironmental conditions in modulating broader climatic trends. In
637 conclusion, the combined CIA and MAP reconstructions indicate a trend of increasing weathering

638 intensity and precipitation from the Selandian into the Thanetian, with spatial variations
639 highlighting the interplay between global climate forcing and regional paleoenvironmental factors.
640 These results contribute to a growing body of evidence supporting a warm and humid Middle to
641 late Paleocene climate, with implications for silicate weathering feedbacks and carbon cycle
642 dynamics during this critical interval in Earth's history.

643

644 **CONCLUSIONS**

645 This study presents the first detailed characterization of the Selandian–Thanetian Transition Event
646 (STTE) within a shallow marine delta system along the Texas Gulf Coast, integrating
647 sedimentological, palynological, and geochemical evidence. Geochemical proxies reveal negative
648 excursions in organic carbon isotopes, intensified chemical weathering, and increased precipitation
649 during the STTE. Palynological data indicate dynamic shifts in coastal vegetation, including the
650 expansion of mangrove-like taxa during inferred hyperthermal phases. While local stratigraphic
651 discontinuities limit signal continuity, comparison with coeval global records supports interpreting
652 this Gulf Coast section as a regional expression of broader Paleocene carbon cycle perturbations.
653 These findings refine the local stratigraphic framework and, more importantly, underscore the
654 sensitivity of marginal marine systems to short-lived hyperthermals, highlighting the value of
655 nearshore archives in complementing both deep-marine and terrestrial isotope records.

656

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663

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