

## The Paleocene-Eocene Thermal Maximum

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# <sup>1</sup> **The Paleocene-Eocene Thermal Maximum:** <sup>2</sup> **How much carbon is enough?**

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3 The Paleocene-Eocene Thermal Maximum (PETM),  $\sim 55.53$  million years  
4 before present, was an abrupt warming event that involved profound changes  
5 in the carbon cycle and led to major perturbations of marine and terrestrial  
6 ecosystems. The PETM was triggered by the release of a massive amount  
7 of carbon, and thus the event provides an analogue for future climate and  
8 environmental changes given current anthropogenic CO<sub>2</sub> emissions. Previ-  
9 ous attempts to constrain the amount of carbon released have produced widely  
10 diverging results, between 2000 and 10000 gigatonnes carbon (GtC). Here  
11 we use the UVic Earth System Climate Model in conjunction with a recently  
12 published compilation of PETM temperatures [*Dunkley Jones et al., 2013*]  
13 to constrain the initial atmospheric CO<sub>2</sub> concentration as well as the total  
14 mass of carbon released during the event. Thirty-six simulations were ini-  
15 tialized with varying ocean alkalinity, river runoff, and ocean sediment cover.  
16 Simulating various combinations of pre-PETM CO<sub>2</sub> levels (840, 1680, and  
17 2520 ppm) and total carbon releases (3000, 4500, 7000, and 10 000 GtC),  
18 we find that both the 840 ppm plus 7000 GtC and 1680 ppm plus 7000-10000  
19 GtC scenarios agree best with temperature reconstructions. Bottom waters  
20 outside the Arctic and North Atlantic Oceans remain well oxygenated in all  
21 of our simulations. While the recovery time and rates are highly dependent  
22 on ocean alkalinity and sediment cover, the maximum temperature anomaly,  
23 used here to constrain the amount of carbon released, is less dependent on  
24 this slow acting feedback.

## 1. Introduction

25 The Paleocene-Eocene Thermal Maximum (PETM),  $\sim 55.53$  million years before present  
26 [*Westerhold et al.*, 2007], is arguably the most intensively studied abrupt warming event  
27 in the geologic record (e.g., *Kennett and Stott* [1991]; *Dickens et al.* [1995]; *Thomas and*  
28 *Shackleton* [1996]; *Thomas et al.* [2002]; *Zachos et al.* [2003, 2005]; *Sluijs et al.* [2007a]).  
29 Thousands of gigatonnes (Gt) of carbon were released into the atmosphere and ocean  
30 over less than 20 000 years leading to profound changes in climate, the carbon cycle, and  
31 ocean chemistry, as well as major perturbations in marine and terrestrial ecosystems (e.g.  
32 *Kelly et al.* [1996]; *Dickens et al.* [1997]; *Sloan and Thomas* [1998]; *Thomas* [1998]; *Bains*  
33 *et al.* [1999]; *Crouch et al.* [2001]; *Bralower* [2002]; *Wing et al.* [2005]; see *Sluijs et al.*  
34 [2007b] and *McInerney and Wing* [2011] for thorough reviews). The event is of particular  
35 interest because it may provide an analogue for future climate and environmental change  
36 if anthropogenic CO<sub>2</sub> emissions continue on their current trajectory (e.g., *Ridgwell and*  
37 *Schmidt* [2010]; *Zeebe and Zachos* [2013]).

38 The PETM is associated with substantial warming of sea surface and deep waters,  
39 based on interpretation of several proxies: oxygen isotopes and Mg/Ca ratios measured  
40 in foraminifera and the lipid-based TEX<sub>86</sub> proxy [*Kennett and Stott*, 1991; *Thomas and*  
41 *Shackleton*, 1996; *Thomas et al.*, 2002; *Zachos et al.*, 2003; *Sluijs et al.*, 2007a]. Recon-  
42 structed temperatures show a significant amount of variation resulting from proxy dis-  
43 crepancies, local environmental effects and foraminiferal recrystallization (e.g. *Bralower*  
44 *et al.* [1995]; *Zachos et al.* [2003]; *Kozdon et al.* [2013]). Recently *Dunkley Jones et al.*  
45 [2013] compiled and compared the available temperature data and assessed the reliability

46 of proxy temperature estimates. They estimated the global mean surface temperature  
47 anomaly to be within the range of 4 to 5°C and the intermediate water temperature  
48 anomaly to be ~5°C.

49 Although the PETM has received vigorous study by the paleoclimate community, the  
50 pre-existing climate and the magnitude of the perturbation remain poorly resolved. For ex-  
51 ample, late Paleocene (pre-PETM) atmospheric CO<sub>2</sub> concentration estimates vary widely.  
52 *Pearson and Palmer* [2000] found late Paleocene concentrations exceeding 2400 ppm based  
53 on boron-isotope ratios of planktonic foraminiferal shells. This is in stark contrast with es-  
54 timates of concentrations below 300 ppm and 400 ppm, based on modeled carbon isotope  
55 gradients [*Hilting et al.*, 2008] and leaf stomatal indices [*Royer et al.*, 2001], respectively.  
56 A recent study by *Schubert and Jahren* [2013] constrained the range of late Paleocene  
57 carbon dioxide concentrations to 674-1034 ppm. Since the radiative forcing of an atmos-  
58 pheric CO<sub>2</sub> change depends on the background concentration, a given release of carbon  
59 will cause more warming in a low-CO<sub>2</sub> atmosphere than in a high-CO<sub>2</sub> atmosphere. In  
60 other words, the higher the initial CO<sub>2</sub> concentration, the larger the carbon release that  
61 is required to explain the same amount of warming. A better understanding of pCO<sub>2</sub> lev-  
62 els immediately before the PETM is therefore essential in order to reconstruct the event  
63 itself.

64 Estimates of the total amount of carbon released during the PETM also vary signifi-  
65 cantly. *Zachos et al.* [2005] estimated a total release of > 4500 GtC based on the extent  
66 of sea-floor carbonate dissolution. *Panchuk et al.* [2008] refined this number to > 6800  
67 GtC, based on dissolution estimates simulated with the GENIE-1 model. Reconstruction

68 of Arctic hydrology also supported the high release estimate [*Pagani et al.*, 2006a]. At  
69 the other end of the spectrum, *Zeebe et al.* [2009] constrained the initial carbon release  
70 to  $< 3000$  GtC, based on simulations of carbonate dissolution and the magnitude of the  
71 carbon isotope excursion using the carbon cycle model LOSCAR. More precise estimates  
72 of the magnitude and rate of carbon release are vital to determine the source of  $\text{CO}_2$  that  
73 fueled the PETM (e.g. *Dickens et al.* [1997]; *Kurtz et al.* [2003]; *Panchuk et al.* [2008]), to  
74 understand the magnitude of potential positive feedbacks in the natural climate system  
75 during the event, and to constrain changes in ocean chemistry and marine and terrestrial  
76 ecosystems resulting from future anthropogenic  $\text{CO}_2$  emissions.

77 The lack of consensus on the background  $\text{CO}_2$  levels and the magnitude of the carbon  
78 pulse necessitates an independent approach. Here, we use a climate model of intermediate  
79 complexity (the UVic Earth System Climate Model [*Weaver et al.*, 2001]) in conjunc-  
80 tion with a recently published compilation of temperature reconstructions [*Dunkley Jones*  
81 *et al.*, 2013] to constrain the atmospheric  $\text{CO}_2$  concentration prior to the PETM as well  
82 as the amount of carbon released during the event.

## 2. Methods

83 The UVic Earth System Climate Model (UVic ESCM) consists of an ocean general  
84 circulation model (Modular Ocean Model, Version 2, [*Pacanowski*, 1995]) coupled to a  
85 vertically integrated two dimensional energy-moisture balance model of the atmosphere,  
86 a dynamic-thermodynamic sea ice model, a land surface scheme, a dynamic global veg-  
87 etation model [*Meissner et al.*, 2003], and a sediment model [*Archer*, 1996; *Meissner*  
88 *et al.*, 2012]. It also includes a fully coupled carbon cycle [*Matthews et al.*, 2005; *Meissner*

89 *et al.*, 2003; *Schmittner et al.*, 2008]. The marine ecosystem/biogeochemical model is an  
90 improved NPZD (nutrient, phytoplankton, zooplankton, detritus) model with a param-  
91 eterization of fast nutrient recycling due to microbial activity [*Schartau and Oschlies*,  
92 2003]. It includes two phytoplankton classes (nitrogen fixers and other phytoplankton),  
93 one zooplankton class, two nutrients (nitrate and phosphate), oxygen, dissolved inorganic  
94 carbon and alkalinity as prognostic tracers. Carbonate production is calculated as a fixed  
95 proportion of primary production, which is indirectly a function of temperature through  
96 the Eppley function [*Eppley*, 1972]. A complete description of the ecosystem model can  
97 be found in *Schmittner et al.* [2008]. The ocean biogeochemical model calculates carbon  
98 fluxes to the sediments as well as their rain ratios. Sediment processes are represented  
99 using a model of deep ocean sediment respiration [*Archer*, 1996; *Meissner et al.*, 2012].  
100 This model assumes oxic conditions, therefore all incoming organic carbon is assumed to  
101 dissolve. The remaining  $\text{CaCO}_3$  is added to the first sediment layer, eventually passes  
102 through the pore layers to be added to more stable layers and finally the lithosphere.  
103 Weathering fluxes are either based on atmospheric  $\text{CO}_2$  concentrations or a combination  
104 of surface atmospheric temperature and net primary productivity [*Meissner et al.*, 2012].  
105 The UVic ESCM is computationally very efficient and has been developed to address  
106 scientific questions related to climate variability on time scales of hundreds of years to  
107 millennia (e.g. *Meissner et al.* [2008]; *Eby et al.* [2009]).

108 For the present study, we integrated four control simulations for over 10 000 years  
109 with Eocene paleogeography, bathymetry and wind fields [*Sijp et al.*, 2011]. Orbital  
110 parameters and the solar constant were set to present day values. Simulations were

111 integrated with atmospheric CO<sub>2</sub> concentrations held constant at 280, 840, 1680 and  
112 2520 ppm. The three warmer simulations were then forced with carbon emission pulses of  
113 3000, 4500, 7000 and 10000 GtC over one year. While pulse scenarios are commonly used  
114 in the modelling community for simplicity and ease of comparison between models (e.g.,  
115 *Cao et al.* [2009]; *Eby et al.* [2009]), they likely overestimate the short-term temperature  
116 and atmospheric carbon dioxide response. Recently *Wright and Schaller* [2013] proposed  
117 that the PETM was indeed triggered by an instantaneous release based on proxy and  
118 sedimentary data, but this interpretation has been disputed (e.g., *Zeebe et al.* [2014]).  
119 The long-term climate response appears to be independent of the rate at which CO<sub>2</sub> is  
120 emitted (e.g., *Eby et al.* [2009]; *Meissner et al.* [2012]). In addition, one set of gradual  
121 release scenarios was integrated with emissions of 1 GtC per year for 4500 years to simulate  
122 a slower release scenario (e.g. *Cui et al.* [2011]). Each scenario was integrated twice  
123 with differing weathering parameterizations [*Meissner et al.*, 2012], based on either a  
124 combination of surface atmospheric temperature and net primary productivity [*Lenton*  
125 *and Britton*, 2006] or on atmospheric CO<sub>2</sub> concentrations [*Zeebe et al.*, 2008], termed LB  
126 and ZL hereafter (see Table 1 for a list of all simulations).

127 During model spin-up, total alkalinity and DIC are conserved by balancing sedimen-  
128 tary CaCO<sub>3</sub> deposition with the alkalinity and DIC fluxes from river discharge [*Meissner*  
129 *et al.*, 2012]. Calcium and magnesium ion concentrations are assumed constant and equal  
130 to modern concentrations when solving for saturation state. Sediments, ocean biogeo-  
131 chemistry and the global carbon cycle then adjust to the given pre-defined global mean  
132 ocean alkalinity. During transient simulations, the weathering fluxes are calculated prog-



133 nostically and global alkalinity and DIC are free to evolve. Figure 1a-c shows the model  
134 percent dry weight  $\text{CaCO}_3$  at the end of the three warmer control simulations integrated  
135 with present day global mean ocean alkalinity ( $2.429 \text{ mol/m}^3$ ). Sediment cover is low  
136 in these high- $\text{CO}_2$  scenarios, which likely leads to an underestimation of the sediment-  
137 alkalinity feedback during the recovery period. To address this issue, we integrated three  
138 additional control simulations with 1680 ppm  $\text{CO}_2$  and global mean alkalinity increased  
139 by a factor of 2 ( $4.858 \text{ mol/m}^3$ , 1680\_Alk2), 1.5 ( $3.644 \text{ mol/m}^3$ , 1680\_Alk15), and 1.2  
140 ( $2.915 \text{ mol/m}^3$ , 1680\_Alk12). Simulated percent dry weight  $\text{CaCO}_3$  at the end of these  
141 simulations is shown in Figure 1d, 1e, 1f, and 1b respectively; the global mean equals 77%  
142 (1680\_Alk2), 44% (1680\_Alk15), 19% (1680\_Alk12) and 7% (1680). Each high-alkalinity  
143 simulation was then forced with a carbon emission pulse of 7000 GtC. 1680\_Alk15 was  
144 also forced with pulses of 3000, 4500, and 10000 GtC (Table 1).

145 Model results were compared to a compilation of SST and deep water temperature  
146 estimates for the interval immediately preceding the PETM and the peak of the event  
147 [*Dunkley Jones et al.*, 2013]. The compilation includes estimates based on  $\delta^{18}\text{O}$ , Mg/Ca  
148 and  $\text{TEX}_{86}$  temperature proxies.

### 3. Results

#### 3.1. Late Paleocene climate conditions

149 All six equilibrium simulations form deep water in the North Pacific, while there is  
150 little to no deepwater formation in the North Atlantic. In all simulations Southern Ocean  
151 sourced bottom water is the dominant water mass (similar to earlier studies, e.g. *Bice and*  
152 *Marotzke* [2001]; *Thomas et al.* [2003]; *Winguth et al.* [2012]), and the Tethys Ocean forms

153 warm and saline deep water which sinks to a depth of  $\sim 1000\text{m}$ . Figure 2 shows simulated  
154 annual sea surface temperatures (SSTs) together with reconstructed SSTs compiled by  
155 *Dunkley Jones et al.* [2013] at Deep Sea Drilling Project (DSDP) Site 527, Ocean Drilling  
156 Program (ODP) Sites 690, 865, 1172 and 1209, and Leg 174AX Bass River and Wilson  
157 Lake cores from on-shore New Jersey, and Integrated Ocean Drilling Program (IODP)  
158 Leg 302 ACEX Core 4A from the Lomonosov Ridge (Arctic Ocean). Where multiple  
159 proxies exist at a single site, the median SST was plotted. Figure 3a shows all pre-  
160 PETM temperature proxies along with estimates for proxy calibration uncertainty and  
161 one standard deviation of data variability (see *Dunkley Jones et al.* [2013] for details).

162 From both figures it is clear that none of our simulations is able to reproduce the warm  
163 SST proxy interpretations at the high-latitude ACEX core (tagged with [1] in Figure 2a)  
164 and Site 1172 [8]. The warmer simulations (840 ppm, 1680 ppm and 2520 ppm) are all  
165 three in agreement with the third high-latitude Site 690 [7] as well as with the mid-latitude  
166 cores Bass River [2] and Wilson Lake [3]. Sites 1209 [4] and 865 [5] in the tropics align  
167 best with the 840 ppm simulation, whereas SSTs from Site 527 [6] in the South Atlantic  
168 are consistent with the two warmest simulations (1680 ppm and 2520 ppm).

169 We quantified the model-proxy disparity for each site as the absolute difference between  
170 modelled SST (interpolated to the correct location) and the closest proxy reconstruction.  
171 If modelled SST fell within the error bounds of a proxy, the disparity was defined as  
172 0. Figure 3b shows this metric plotted for each core. Other than for the previously  
173 discussed high-latitude ACEX core [1] and Site 1172 [8], the three warmer simulations  
174 always achieve a model-proxy disparity below  $2.5^\circ\text{C}$ . Given that the proxy data themselves

175 present important discrepancies for a given site, we calculate the median disparity (which  
176 by definition gives less weight to outliers than the mean disparity) to establish which  
177 control simulation is closest to reconstructions. Overall, the median disparity across all  
178 proxies is lowest for the 840 ppm simulation at 0°C. Simulation 1680 has the second lowest  
179 median disparity (0.025°C). Thus either the 840 or 1680 ppm simulations are plausible,  
180 especially if pre-PETM  $\delta^{18}\text{O}$  proxies are regarded as minimum estimates [*Dunkley Jones*  
181 *et al.*, 2013].

182 When we calculate the model-proxy disparity for each proxy separately, we find that  
183 both  $\delta^{18}\text{O}$  and Mg/Ca proxy SSTs agree best with the 840 ppm simulation (median  
184 disparities of 0°C); TEX proxy SSTs are closest to the 2520 ppm simulation (median  
185 disparity 1.8°C). Although the calculation of Mg/Ca-based temperatures relies on various  
186 assumptions about seawater Mg/Ca, the calibrations used best-fit independent estimates  
187 of Mg/Ca<sub>sw</sub> and are in reasonable agreement with  $\delta^{18}\text{O}$ -derived temperatures from sites  
188 with excellent planktic foraminiferal preservation (see discussion in *Dunkley Jones et al.*  
189 [2013]). For the  $\delta^{18}\text{O}$  proxy-based disparity calculation, we disregarded oxygen isotope data  
190 from Sites 527, 865 and 1209 based on the large discrepancy between  $\delta^{18}\text{O}$  and Mg/Ca  
191 temperatures on the order of  $\sim 10^\circ\text{C}$  [*Dunkley Jones et al.*, 2013]. The median disparity  
192 across all proxies is not affected by the inclusion of  $\delta^{18}\text{O}$  from these three sites.

193 Other than Sites 865 and 1051 (both at  $\sim 1500\text{m}$  depth) and the Bass River and Wil-  
194 son Lake cores (both  $< 150\text{m}$  deep [*Harris et al.*, 2010]), bottom water temperatures are  
195 underestimated in all simulations when compared to proxy estimates (Figure 3c). Cali-

196 bration uncertainty for benthics is shown as  $\pm 1^\circ\text{C}$  for both Mg/Ca [*Lear et al.*, 2002] and  
197  $\delta^{18}\text{O}$  [*Kim and O'Neil*, 1997].

### 3.2. The carbon pulse

198 Time series for atmospheric carbon anomalies show that the two weathering schemes  
199 yield very similar results, especially when initialized with the warmest climate (2520  
200 ppm, Figure 4, compare dotted lines with solid lines). Even with a colder initial climate  
201 (840 ppm) the two weathering schemes show less discrepancy in the recovery than when  
202 integrated under present day conditions [*Meissner et al.*, 2012]. As expected, temperature  
203 response decreases with increasing background  $\text{CO}_2$ , for example a 10 000 GtC pulse has  
204 a larger impact on temperature for the 840 ppm simulations than for the 2520 ppm  
205 simulations. Global mean ocean temperatures take over 5000 years to equilibrate. There  
206 is little to no recovery in atmospheric  $\text{CO}_2$  and temperatures during the 10 000 years of  
207 integration for simulations that started with present day global mean alkalinity, indicating  
208 that the climate system is so saturated in  $\text{CO}_2$  that the land and ocean can absorb little of  
209 the excess atmospheric carbon. Simulations with higher ocean alkalinity (dashed lines in  
210 Figure 4, middle panels) show a significantly faster recovery in atmospheric  $\text{CO}_2$ , especially  
211 for high emissions (7000 and 10 000 GtC). While the maximum increase in surface air  
212 temperature is similar for simulations that started with different alkalinities, deep ocean  
213 warming is slightly less for higher alkalinity simulations (Figure 4e and h).

214 All 36 simulations indicate an initial decrease in global mean oceanic oxygen concentra-  
215 tions followed by a recovery (last row of Figure 4). Figure 5 (first row) shows the values of  
216 vertical minimum oxygen concentrations in hypoxic regions during the 840, 1680, and 2520

217 ppm control simulations. While there is widespread hypoxia (defined here as regions with  
218 oxygen concentrations below  $90\mu\text{M}$ ) in the tropics in all simulations, the Arctic and parts  
219 of the Southern Ocean become hypoxic for higher atmospheric  $\text{CO}_2$  concentrations. Most  
220 of these hypoxic regions are located within the uppermost 1000m of the water column  
221 (Figure 5e). Only the Arctic Ocean as well as some continental shelves in the Atlantic  
222 Ocean experience bottom-water oxygen levels below  $30\mu\text{M}$  during the most oxygen de-  
223 pleted simulation (2520\_10000\_LB, Figure 5f). The bottom water of the Atlantic Ocean  
224 is also depleted but stays above  $60\mu\text{M}$  during this simulation.

225 Sediment chemistry timeseries are shown in Figure 6. The percentage of calcite in  
226 sediments increases during the first several thousand years in all simulations (Figure 6,  
227 third row). This is due to a temperature-driven increase in global mean photosynthesis and  
228 calcite production (see detailed discussion in *Meissner et al.* [2012]), which compensates  
229 for the initial increase in dissolution in sediments (Figure 6, first and second rows) and acts  
230 as a weak positive feedback on atmospheric  $\text{CO}_2$  concentrations. After several thousand  
231 years, the acidification signal of the carbon pulse reaches the deep ocean (note the steeper  
232 increase in dissolution: Figure 6, second row). Dissolution exceeds the downward flux of  
233 calcite and the total mass of calcite in the pore layer decreases. The last row in Figure 6  
234 shows the change in the global mean calcite compensation depth (diagnosed here as the  
235 mean depth of grid boxes with less than 10% dry weight  $\text{CaCO}_3$ ). A temporary shoaling  
236 of this metric can be seen for all simulations.

237 Figure 7 shows the simulated maximum SST anomalies at the eight sites compared  
238 to proxy reconstructions. While the reconstructions often exhibit considerable spread,

239 there is some overlap with the simulations at nearly all sites. Note that proxy data from  
240 Sites 865 [5] and 527 [6] likely underestimate the temperature anomaly for stratigraphic  
241 reasons (chemical erosion or “burndown” of the basal few thousand years of the PETM,  
242 *Dunkley Jones et al.* [2013]). Proxy reconstructions for all other sites are compatible with  
243 the high emission scenarios (7000-10000 Gt C) especially when started with a higher CO<sub>2</sub>  
244 background climate of 1680 or 2520 ppm.

245 Maximum proxy bottom water temperature anomalies are shown in Figure 8, with  
246 depth ranging from 80m (Wilson Lake, first panel) to 3400m (DSDP 527, last panel).  
247 Bottom temperature proxies from Site 1209 [4] show a small peak increase compared to  
248 most simulations and proxy data from all other sites, which might be due to chemical  
249 erosion and/or slow deposition rates [*Dunkley Jones et al.*, 2013]. Bottom temperature  
250 proxy reconstructions for other sites agree best with the higher emission scenarios.

251 The median model-proxy disparity for sea surface temperature anomalies is minimized  
252 for three distinctive scenarios: a low-carbon scenario (initial CO<sub>2</sub> of 840 ppm plus a carbon  
253 forcing of 4500 GtC; overall median disparity of 0.348°C); a medium scenario (1680 ppm  
254 + 7000 GtC, median disparity 0.133°C); and a high-carbon scenario (2520 ppm + 10000  
255 GtC, median disparity 0.20°C). One should bear in mind that the design of our model  
256 simulations (emission pulse over one year) tends to overestimate the surface temperature  
257 response. Furthermore, the proxy data in several deep sea sites are likely to have missed  
258 the peak temperature because of chemical erosion or low temporal resolution. We therefore  
259 conclude that based on SST anomalies, the carbon pulse was likely 7000-10000 Gt C or  
260 higher. When analyzing bottom temperatures, the minimum model-proxy disparities are

261 achieved for slightly more carbon intensive scenarios: 840 ppm + 7000 GtC (0.362°C);  
262 1680 ppm + 10 000 GtC (0.67°C); and 2520 ppm + 10 000 GtC (1.197°C). Overall, the  
263 amount of released carbon required to cause the reconstructed temperature anomalies  
264 depends heavily on the initial atmospheric carbon dioxide concentration: 7000 GtC for  
265 pre-PETM atmospheric CO<sub>2</sub> concentrations of 840 ppm; 7000-10 000 GtC for pre-event  
266 CO<sub>2</sub> concentrations of 1680 ppm and over 10 000 GtC for pre-PETM atmospheric CO<sub>2</sub> of  
267 2520 ppm.

#### 4. Discussion

268 Previous model-based estimates of the magnitude of the carbon perturbation required  
269 to trigger the PETM were constrained by the size of the carbon isotope excursion or the  
270 extent of deep ocean dissolution (e.g., *Panchuk et al.* [2008]). The results show consider-  
271 able variation due to the range of complexity of the models used as well as the unknown  
272 background chemistry of Paleocene ocean water. To refine our understanding of PETM  
273 atmospheric forcing, we apply a novel model-data combination: the UVic model which  
274 was built for long-term simulations with a special focus on ocean dynamics and feedbacks  
275 [*Weaver et al.*, 2001] and a recently published compilation of proxy surface and deep water  
276 temperature data [*Dunkley Jones et al.*, 2013]. Here we interpret the results and impli-  
277 cations of the model-data comparison beginning with a discussion of the uncertainties of  
278 the data and the sensitivity of model simulations, followed by a comparison of the results  
279 with those of previous investigations.

## 4.1. Uncertainties in proxy data reconstructions

Investigation of a climate event that took place 55 million years ago is fraught with challenges, both for modelling and proxy analysis. Deep ocean acidification resulting from the carbon release led to widespread dissolution of the carbonate microfossils which were deposited during and immediately before the PETM at deep sea sites [Zachos *et al.*, 2005; Colosimo *et al.*, 2006; Zachos *et al.*, 2007; Murphy *et al.*, 2010]; thus precise constraint of the peak warming signal is not possible at these locations. Oxygen isotope variations across the PETM are impacted by changes in salinity as well as temperature. In tropical sites, for example, an increase in evaporation is thought to have decreased the amplitude of the temperature signal [Zachos *et al.*, 2003]; conversely, fresh water input at high-latitude Site 690 may have increased the amplitude. The Mg/Ca values of seawater are known to change through time and temperature estimates based on them rely heavily on the calibration applied (e.g., Evans and Müller [2012]). Interpretation of Mg/Ca and particularly  $\delta^{18}\text{O}$  values is also confounded by possible alteration of carbonate microfossils during burial. Carbonate recrystallization decreases the magnitude of the PETM SST increase, especially at low-latitude deep-sea sites (e.g., Pearson *et al.* [2001]). GDGT lipid-based proxies, used in coastal and high-latitude PETM sections, circumvent such diagenetic issues but are subject to significant calibration uncertainty, especially during warm climate states (e.g., Kim *et al.* [2010]; Hollis *et al.* [2012]) and with changing productivity regimes [Taylor *et al.*, 2013]. For a more detailed discussion of the uncertainties within the pre-PETM and PETM proxy data set see Dunkley Jones *et al.* [2013].

## 4.2. Uncertainties in major ion seawater composition



300 Model simulations rely heavily on boundary conditions, which have significant uncer-  
301 tainties during this period. Topography and wind forcing, in particular, are major un-  
302 knowns which greatly influence the climate state. Little is known about the orbital pa-  
303 rameters or the background chemistry (pre-PETM ocean alkalinity [*Cui et al.*, 2011]). For  
304 example, *Lowenstein et al.* [2001] suggest that the  $\text{Mg}^{2+}/\text{Ca}^{2+}$  ratio increased from  $<2.3$   
305 in the Cretaceous to  $>2.5$  between 50 and 0 Ma. The  $\text{Ca}^{2+}$  concentration in seawater is  
306 believed to have reached maximum values two to three times greater than modern values  
307 in the Cretaceous and was also likely higher during the Eocene than today (*Horita et al.*  
308 [2002], their Figure 8). The major ion composition is crucial for calculating seawater  
309 chemistry, saturation and the capacity of carbon uptake by the ocean [*Tyrrell and Zeebe,*  
310 2004]. However, Ocean General Circulation Models (OGCMs) do not generally include  
311 sophisticated seawater chemistry models. In the group of climate models including full  
312 ocean GCMs, the UVic model has one of the most detailed biogeochemistry components.  
313 While we cannot take variations in the concentrations of any particular major seawater  
314 ion into account, we can vary the global mean ocean alkalinity as a measure of carbonate  
315 and bicarbonate ions in the ocean. Figure 1b and d show two extreme cases of background  
316 alkalinity and their impact on ocean sediments. The percent dry weight  $\text{CaCO}_3$  in the  
317 late Paleocene was probably between these two extremes [*Panchuk, 2007*], which gives  
318 us confidence that our simulations have spanned the parameter space of possible climate  
319 responses to a certain carbon pulse (including climate sensitivity) with regard to initial  
320 marine sediment cover [*Goodwin et al.*, 2009; *Goodwin and Ridgwell, 2010*]. While the  
321 long-term recovery of atmospheric  $\text{CO}_2$  is highly dependent on the initial alkalinity and

322 sediment cover (Figure 4, middle panels), the maximum temperature response acts on  
323 much shorter timescales (especially for surface temperatures) and is less influenced by the  
324 background ocean chemistry.

### 4.3. Climate sensitivity

325 The climate sensitivity of the UVic Earth System Climate Model is a key parameter  
326 for the analysis presented here. Climate sensitivity is often split into fast feedbacks (e.g.  
327 water vapor, snow albedo, sea ice albedo; also called the ‘Charney sensitivity’) and slow  
328 feedbacks (e.g. vegetation, ice sheets, ocean circulation). While the IPCC 2013 report  
329 states that the Charney sensitivity “is likely in the range 1.5°C to 4.5°C (high confidence),  
330 extremely unlikely less than 1°C (high confidence), and very unlikely greater than 6°C  
331 (medium confidence)” [Stocker *et al.*, 2013], these values have been challenged in the  
332 past, especially for warmer background climates [Pagani *et al.*, 2010; Lunt *et al.*, 2010].  
333 Based on a sensitivity study of the Pliocene Lunt *et al.* [2010] suggest a 30-50% higher  
334 climate sensitivity due to slow feedbacks not included in coupled GCMs. It should be  
335 noted, however, that their model does not include the sediment-alkalinity or weathering  
336 feedbacks, which are the main negative slow feedbacks in the climate system. In a more  
337 recent paper, Rohling *et al.* [2013] revisit climate sensitivity over the past 65 million years  
338 and find values which agree with the most recent IPCC report [Stocker *et al.*, 2013]. On  
339 the other hand, Schmittner *et al.* [2011] find that modern climate models are more likely to  
340 over- than to under-estimate climate sensitivity; a view that has been challenged recently  
341 [Fyke and Eby, 2012]. Cloud feedbacks, a significant source of uncertainty in future climate  
342 projections, are particularly poorly understood under CO<sub>2</sub> concentrations four to nine

343 times preindustrial values [*Abbot and Tziperman, 2009; Kiehl and Shields, 2013*]. *Lunt*  
344 *et al.* [2012] find that the main reasons for differences between early Eocene simulations  
345 by five different models include differences in surface albedo feedbacks, water vapor and  
346 lapse rate feedbacks, as well as prescribed aerosol loading, rather than differences in cloud  
347 feedbacks.

348 The simulations presented here show the reaction of the climate system to a carbon pulse  
349 within the first 10 000 years of the PETM. The UVic model includes the classic Charney  
350 feedbacks in addition to some of the slower feedbacks (e.g., vegetation, alkalinity, ocean  
351 circulation, weathering fluxes). With these feedbacks, the UVic ESCM has a climate  
352 sensitivity of 3.3°C under PETM boundary conditions, which is slightly lower than under  
353 preindustrial boundary conditions (3.5°C, *Weaver et al.* [2007]) and which falls within the  
354 range suggested by the IPCC and *Rohling et al.* [2013].

#### 4.4. High latitude temperatures and meridional temperature gradient

355 The latitudinal temperature gradient at the surface is slightly overestimated in our  
356 simulations compared to temperature reconstructions (Figure 2). This might be par-  
357 tially due to a potential underestimation of low-latitude SSTs by proxy data (e.g. *Huber*  
358 [2008]). However, the UVic model clearly faces the well-known problem of climate models  
359 simulating polar regions that are too cool in high-CO<sub>2</sub> climates (e.g. *Sloan and Barron*  
360 [1990]; *Heinemann et al.* [2009]; *Huber and Caballero* [2011]; *Valdes* [2011]; *Lunt et al.*  
361 [2012]; *Sagoo et al.* [2013]). All of our equilibrium model simulations fail to capture warm  
362 temperatures suggested by proxy data from two of the three high-latitude locations: the  
363 ACEX site [1] and Site 1172 [8] (Figure 3a and b), from the Arctic and Southern Oceans,

364 respectively. The extremely high temperatures reconstructed in ACEX are particularly  
365 puzzling. Our simulations show the Arctic Ocean to be largely isolated, with high river  
366 runoff and precipitation exceeding evaporation, leading to very fresh and stratified wa-  
367 ters. These conditions are in agreement with salinity proxies and fossil assemblages [*Sluijs*  
368 *et al.*, 2006; *Waddell and Moore*, 2008]. The surface water masses are therefore in close  
369 thermal equilibrium with the overlying atmosphere, with almost no heat exchange with  
370 deeper layers or other ocean basins. Consequently, the reconstructed high temperatures  
371 in the Arctic could have only been achieved by locally increased longwave radiation (e.g.  
372 polar stratospheric clouds [*Sloan and Pollard*, 1998], or changes in cloud condensation  
373 nuclei [*Kiehl and Shields*, 2013]), locally changed short wave radiation (e.g. obliquity, *Se-*  
374 *wall and Sloan* [2004]) or more efficient heat transport in the atmosphere (stronger winds  
375 and/or increase in latent heat transport).

376 Site 1172 [8] in the Pacific sector of the Southern Ocean records higher temperatures  
377 than Site 690 [7], which is at a similar latitude but in the Atlantic sector. A possible  
378 explanation for the warm proxy temperature estimates at Site 1172 given its location  
379 on the east coast of Australia, involves southward shifted westerlies concurrent with an  
380 intensified western boundary current, which would transport warm low-latitude waters  
381 further south than in our simulations. It is also possible that GDGT-based proxies are  
382 over-estimating SSTs at high-latitude locations, such as Site 1172 and the ACEX site  
383 (e.g., *Hollis et al.* [2012]; *Taylor et al.* [2013]), or that a seasonal bias of the proxies is  
384 causing additional model-data disagreement [*Lunt et al.*, 2012].

385 Bottom water temperatures in all simulations are underestimated in two-thirds of the  
386 sites considered (Figure 3c). Given that bottom temperatures reflect conditions at deep  
387 water formation sites, the model's underestimation of bottom water temperatures and  
388 overestimation of surface temperature gradient are likely connected. Although there is  
389 significant deep water formation in the Tethys Ocean, this water mass is not dense enough  
390 to significantly influence deep water circulation patterns. Therefore, our model does not  
391 support the long-since-refuted WSBW (warm saline bottom water) hypothesis (*Bice and*  
392 *Marotzke* [2001] and references therein).

#### 4.5. Temperature versus dissolution

393 Previous estimates of carbon input are based on simulations of deep ocean dissolution  
394 [*Panchuk et al.*, 2008; *Zeebe et al.*, 2009; *Cui et al.*, 2011]. Our simulations were not  
395 integrated long enough to capture the full dissolution event (Figure 6). While the UVic  
396 model is better skilled at simulating ocean dynamics and changes in three-dimensional  
397 temperature fields than most other climate models of intermediate complexity, it is compu-  
398 tationally too intensive to allow for long enough integrations to analyze maximum changes  
399 in the calcite compensation depth. Furthermore, there are two additional uncertainties  
400 to consider when analyzing modelled changes in the calcite compensation depth. First,  
401 deep ocean dissolution depends on background seawater ion concentration (Section 4.2,  
402 Figures 1 and 6) and would therefore require a range of long-term simulations spanning  
403 the parameter space of alkalinity [*Cui et al.*, 2011]. Second, it is still debated how calcite  
404 production and export react to increasing atmospheric CO<sub>2</sub>, with models showing both  
405 an increase and decrease in export [*Gehlen et al.*, 2007; *Ridgwell et al.*, 2007; *Schmittner*

406 *et al.*, 2008; *Meissner et al.*, 2012]. In addition, the stratigraphic record of the earliest  
407 part of the PETM at many of the study sites is compromised by chemical erosion. Given  
408 these model and stratigraphic complications, we do not interpret the apparent mismatch  
409 between the modelled CaCO<sub>3</sub> results and percentages in the study sections.

#### 4.6. How much CO<sub>2</sub> is enough?

410 The estimate of the amount of carbon required to generate PETM warming depends  
411 heavily on the pre-PETM carbon dioxide concentrations [*Pagani et al.*, 2006b]. Estimates  
412 of late Paleocene atmospheric CO<sub>2</sub> concentrations range widely from 200 ppm to 2800 ppm  
413 (*McInerney and Wing* [2011] and references therein). Here we find that simulated SSTs  
414 agree best with temperature reconstructions for atmospheric CO<sub>2</sub> concentrations between  
415 840 and 1680 ppm, while the best fit between model and data in terms of sediment cover  
416 is achieved for an atmospheric CO<sub>2</sub> concentration of 1680 ppm and global mean ocean  
417 alkalinity of 3.644 mol/m<sup>3</sup> (not shown). Further, proxy data and model simulations fit  
418 best with either a low-carbon scenario (pre-PETM atmospheric CO<sub>2</sub> of 840 ppm plus a  
419 carbon release of 4500-7000 GtC), a medium scenario (1680 ppm plus 7000-10 000 GtC)  
420 or a high-carbon scenario (2520 ppm plus > 10 000 GtC).

421 While a release of 4500 GtC agrees with earlier estimates based on the shoaling of  
422 the calcite compensation depth [*Zachos et al.*, 2005; *Zeebe et al.*, 2009], it underesti-  
423 mates bottom temperature anomalies in our study unless pre-PETM CO<sub>2</sub> concentrations  
424 were below 840 ppm. However, 840 ppm is the minimum CO<sub>2</sub> concentration required to  
425 achieve pre-event reconstructed temperatures. Given that the design of our simulations  
426 (pulse emission) entails an overestimation of maximum simulated temperature anoma-

427 lies, both pre-PETM atmospheric CO<sub>2</sub> concentrations and total release are likely to be  
428 conservative estimates. Furthermore, our simulations with modern background alkalinity  
429 likely overestimate the deep ocean temperature response and hence also underestimate the  
430 total release of carbon. Our study therefore agrees with *Pagani et al.* [2006a]’s climate  
431 sensitivity-based estimate (> 5400 GtC) and *Panchuk et al.* [2008]’s simulations based  
432 on the extent of seafloor CaCO<sub>3</sub> dissolution (> 6800 GtC). Our high-carbon scenario is  
433 also in line with *Cui et al.* [2011]’s *C<sub>org</sub>* scenario, who forced the GENIE model with a  
434 prescribed atmospheric δ<sup>13</sup>C (13 000 GtC).

#### 4.7. Deep-sea anoxia

435 All three of our best-fit simulations are consistent with a bottom water temperature  
436 increase of 4-5 °C (Figure 4) without a significant change in thermohaline circulation  
437 patterns, corroborating the study of *Thomas et al.* [2003]. While annual and global mean  
438 export production decreases by up to 20% in our simulations (not shown), overall the  
439 combination of warmer ocean temperatures and reduced ventilation leads to an expansion  
440 of hypoxic regions (Figure 5), a finding that has been observed in paleoredox proxies  
441 [*Chun et al.*, 2010]. The Arctic Ocean becomes almost entirely hypoxic in our high CO<sub>2</sub>  
442 simulations, due to a very stratified water column. Other than in the Arctic Ocean and  
443 the deep North Atlantic Ocean, hypoxic regions are situated within the first 1000-1500m  
444 of the water column. This is in contrast to *Winguth et al.* [2012] who found widespread  
445 dysoxia in bottom waters in a 2500 year long simulation with CCSM3 under 4480 ppm.  
446 While their simulations with the CCSM3 model were integrated under higher atmospheric  
447 CO<sub>2</sub> forcing than our simulations, their integration time of 2500 years was rather short

448 to equilibrate deep ocean temperatures and the associated vertical temperature gradient  
449 and stratification; the stratification in *Winguth et al.* [2012]’s simulations might therefore  
450 be overestimated. On the other hand, the UVic ESCM is missing wind-climate feedbacks  
451 which can influence upwelling and nutrient availability and therefore export production  
452 and deep-sea oxygen. The simulated deep-sea oxygen might also be overestimated because  
453 of the negative bias in simulated bottom water temperatures (Figure 3c). Observation of  
454 suboxia in the deep North Atlantic is consistent with *Pälike et al.* [2014] who found that  
455 Atlantic intermediate waters were suboxic during the PETM but those from the Pacific  
456 were not.

## 5. Conclusions

457 Estimates of late Paleocene atmospheric CO<sub>2</sub> concentrations and the magnitude of the  
458 PETM carbon release vary widely in the literature. Here we take advantage of a recently  
459 published compilation of recalculated paleotemperatures [*Dunkley Jones et al.*, 2013] to  
460 independently determine these variables using the UVic Earth System Climate Model.  
461 We integrated thirty-six 10 000-year long simulations under varying PETM boundary  
462 conditions. We find three scenarios that best align with proxy reconstructions of PETM  
463 temperature anomalies: a low-carbon scenario (late Paleocene atmospheric CO<sub>2</sub> concen-  
464 tration of 840 ppm and a PETM carbon pulse of 7000 GtC), a medium-carbon scenario  
465 (1680 ppm and 7000-10 000 GtC) and a high-carbon scenario (2520 ppm and > 10 000  
466 GtC). The low- and medium-carbon scenarios fit best with pre-PETM absolute temper-  
467 ature reconstructions. However, the number of locations for which we have reliable SST  
468 reconstructions is small, and the reconstructed temperatures at each of these locations



469 varies widely depending on the proxy and species used. Furthermore, several important  
470 boundary conditions for the modelling study are highly uncertain (alkaline run-off from  
471 rivers [Cui *et al.*, 2011], topography, and clouds). Ocean bottom waters remain well  
472 oxygenated in all our simulations other than for the Arctic and North Atlantic Oceans.

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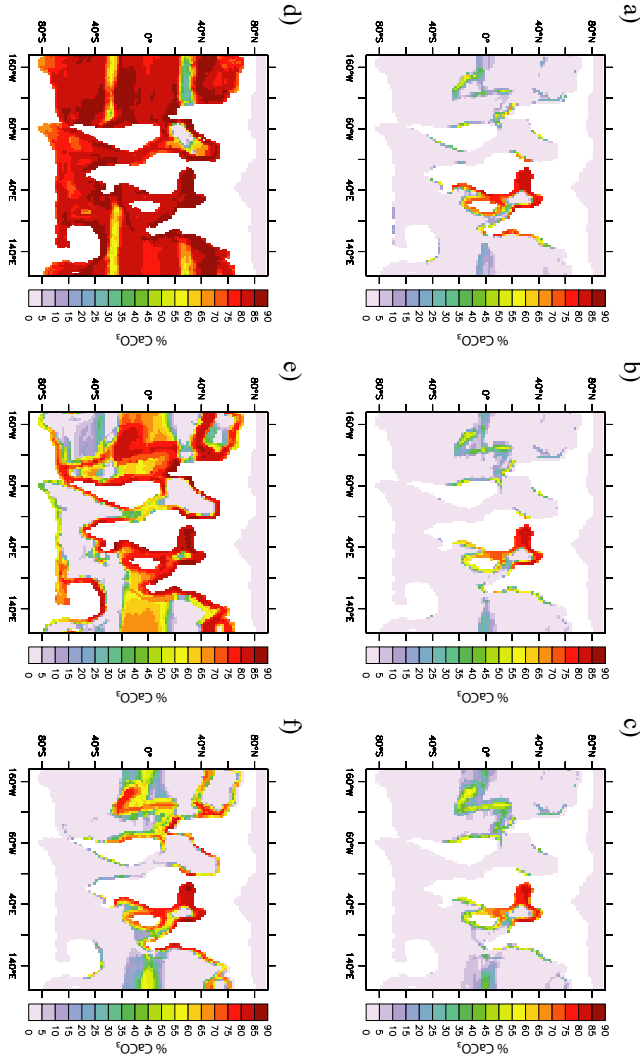
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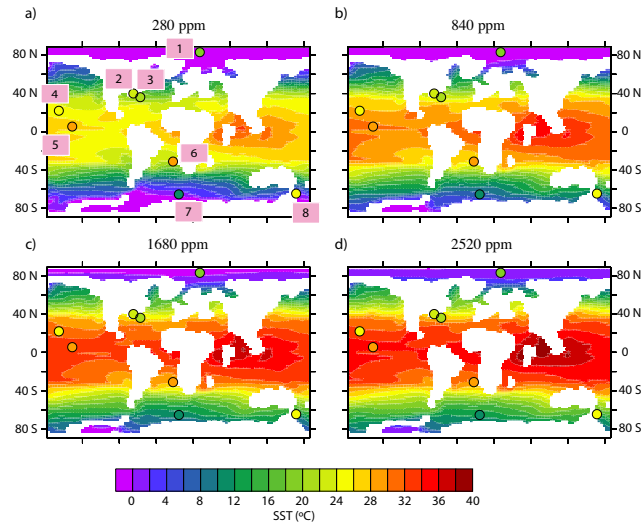
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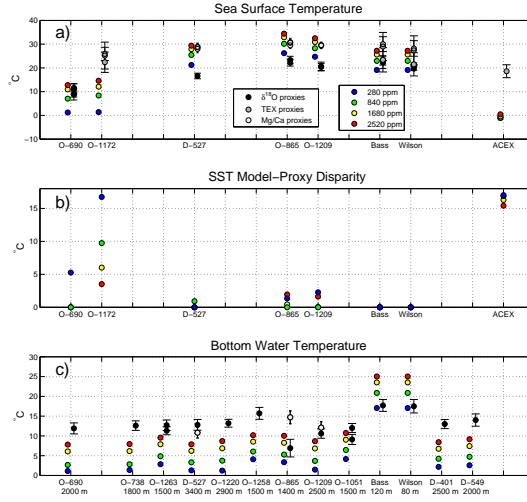
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**Figure 1.** Annual mean percent dry weight  $\text{CaCO}_3$  for the six warmer equilibrium simulations; (a) 840, (b) 1680, (c) 2520, (d) 1680\_Alk2, (e) 1680\_Alk15, (f) 1680\_Alk12.

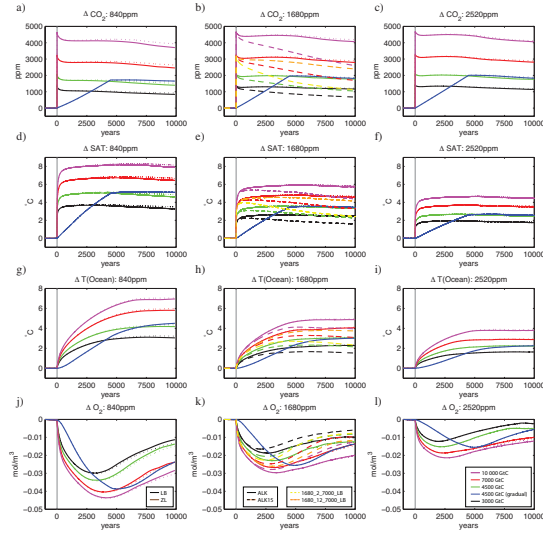


**Figure 2.** Annual mean sea surface temperatures in  $^{\circ}\text{C}$  for the four equilibrium simulations. Also shown are pre-PETM SST reconstructions from proxy data compiled by *Dunkley Jones et al.* [2013]; for sites with several reconstructions of pre-PETM SST, we have plotted the median value. Sites are labeled in (a) as follows: [1] Integrated Ocean Drilling Program (IODP) Leg 302 ACEX core, [2] Bass River, [3] Wilson Lake, [4] Ocean Drilling Program (ODP) Sites 1209, [5] 865, [7] 690, and [8] 1172, and [6] Deep Sea Drilling Project Site 527. Note that [2] and [3] are shown shifted slightly apart for ease of viewing.

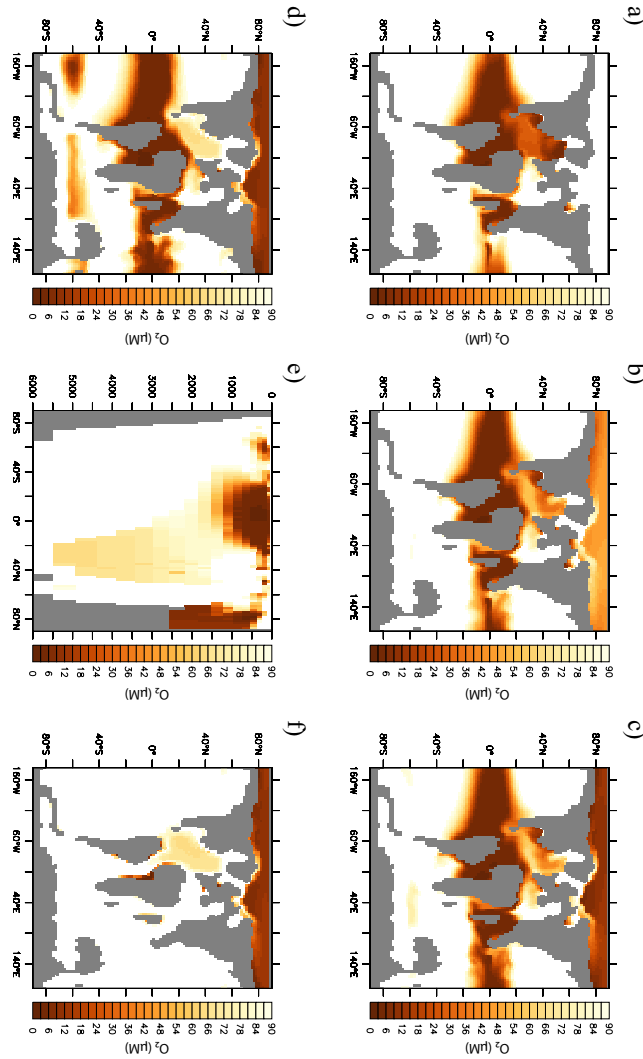


**Figure 3.** Annual mean sea surface ocean temperatures (a) and ocean bottom temperatures (c) in °C for the four equilibrium simulations compared to pre-PETM proxy data compiled by *Dunkley Jones et al.* [2013]. Model-data disparity (calculation described in the text) for each simulation and site is shown in (b). Note that in (c) the depth of ODP 1258 is taken at 1500 m (rather than 2500 m as in *Dunkley Jones et al.* [2013]), which is the maximum depth of the ocean model at that location.

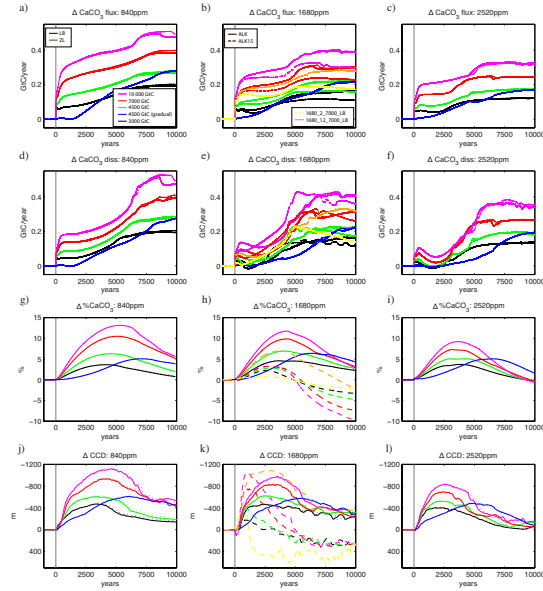




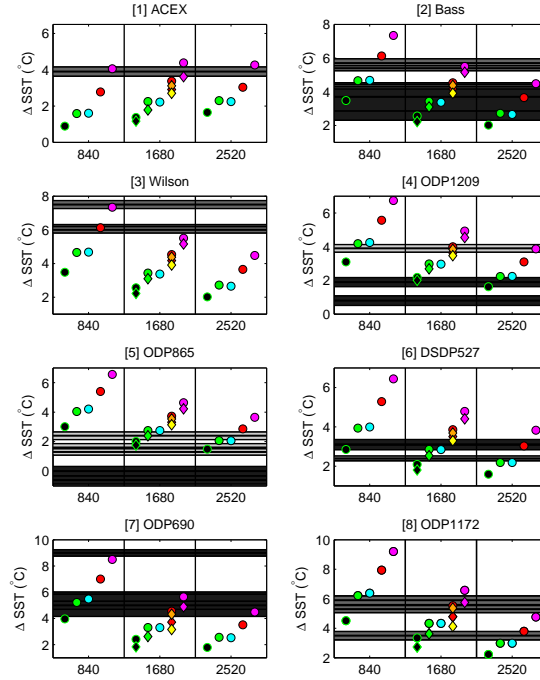
**Figure 4.** Annual mean model results for all transient simulations. Columns (left to right) show simulations initialised with 840, 1680, and 2520 ppm CO<sub>2</sub>; rows (top to bottom) show atmospheric CO<sub>2</sub> concentration anomalies in ppm, global mean surface atmospheric temperature (SAT) anomalies, global mean ocean temperature anomalies and global mean oceanic oxygen concentrations. Simulations with carbon pulses of 3000, 4500, 7000, and 10 000 GtC are plotted in black, green, red, and pink respectively; gradual release simulations (1 GtC/year for 4500 years) are shown in blue. Simulations with the LB and ZL weathering schemes are plotted with solid and dotted lines respectively. Simulations initialized with 1.5 times present day alkalinity are shown in dashed lines (colours corresponding to magnitude of carbon pulse). 1680\_12\_7000\_LB and 1680\_2\_7000\_LB are shown in orange and yellow dashed lines respectively. Light grey vertical lines show the time of carbon release.



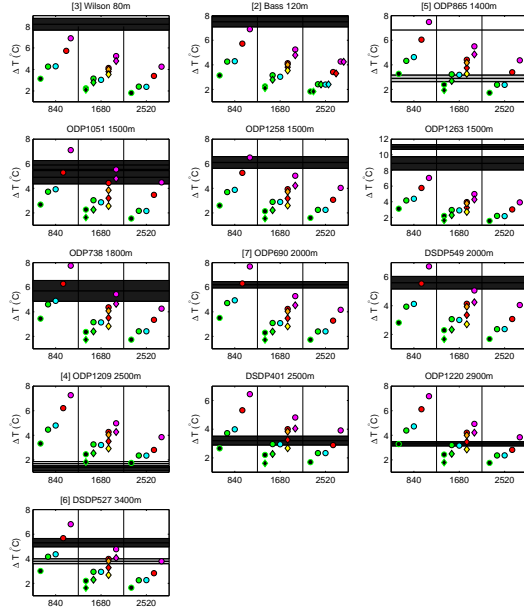
**Figure 5.** Annual mean minimum oxygen concentrations in the water column in  $\mu\text{M}$  ( $10^{-3}\text{mol/m}^3$ ). Only values below  $90 \mu\text{M}$  are shown. Upper panels show equilibrium simulations: (a) 840, (b) 1680, (c) 2520. Lower panels show simulation 2520\_10000\_LB at year 2600 after the pulse: (d) vertical minimum concentration, (e) zonal minimum concentration, (f) bottom water oxygen concentrations.



**Figure 6.** Annual and global mean sediment model results for all transient simulations. Columns (left to right) show simulations initialised with 840, 1680, and 2520 ppm  $\text{CO}_2$ ; rows (top to bottom) show anomalies of global mean downward flux of calcite into the sediments in GtC/year, dissolution of calcite in sediments (GtC/year), calcite pore layer portion (in %), and calcite compensation depth (defined here as the mean depth of grid boxes with less than 10% dry weight  $\text{CaCO}_3$ ; negative values designate a shoaling of this metric). Simulations with carbon pulses of 3000, 4500, 7000, and 10000 GtC are plotted in black, green, red, and pink respectively; gradual release simulations (1 GtC/year for 4500 years) are shown in blue. Simulations with the LB and ZL weathering schemes are plotted with solid and dotted lines respectively. Simulations initialized with 1.5 times present day alkalinity are shown in dashed lines (colours corresponding to magnitude of carbon pulse). 1680.12-7000\_LB and 1680.2-7000\_LB are shown in orange and yellow dashed lines respectively. Light grey vertical lines show the time of carbon release.



**Figure 7.** Maximum temperature anomaly for sea surface temperatures (ACEX, Bass, Wilson, Sites 1209, 865, 527, 690 and 1172). Each panel is split into 3 columns for simulations starting at 840 ppm (left), 1680 ppm (middle) and 2520 ppm (right). Carbon releases of 3000, 4500, 7000, and 10000 GtC are plotted in black, green, red, and pink respectively; gradual release simulations (1 GtC/year for 4500 years) are shown in blue. Simulations with present-day and 1.5 times present-day alkalinity are plotted with circles and diamonds respectively. Orange and yellow diamonds stand for simulations 1680\_12\_7000LB and 16800\_2\_7000\_LB respectively. Horizontal bars in light (Mg/Ca), medium (TEX), and dark ( $\delta O^{18}$ ) grey show proxy reconstructions with standard error.



**Figure 8.** Maximum temperature anomaly for bottom temperatures. Each panel is split into 3 columns for simulations starting at 840 ppm (left), 1680 ppm (middle) and 2520 ppm (right). Carbon releases of 3000, 4500, 7000, and 10 000 GtC are plotted in black, green, red, and pink respectively; gradual release simulations (1 GtC/year for 4500 years) are shown in blue. Simulations with present-day and 1.5 times present-day alkalinity are plotted with circles and diamonds respectively. Simulations 1680\_12\_7000\_LB and 16800\_2\_7000\_LB are plotted with orange and yellow diamonds respectively. Horizontal bars in light (Mg/Ca), medium (TEX), and dark ( $\delta O^{18}$ ) grey show proxy reconstructions with standard error. Note that the depth of ODP 1258 is taken at 1500 m (rather than 2500 m as in *Dunkley Jones et al.* [2013]), which is the maximum depth of the ocean model at that location.

**Table 1.** List of Simulations. LB and ZL denote the weathering scheme used (see text and*Meissner et al.* [2012])

Control Simulation	Transient Pulse		Transient Gradual Release (1 Pg C /year)	
	Emission (Pg C)	Name	Emission (Pg C)	Name
280	-	-	-	-
840	3000	840_3000_LB 840_3000_ZL	4500	840_4500G_LB 840_4500G_ZL
	4500	840_4500_LB 840_4500_ZL		
	7000	840_7000_LB 840_7000_ZL		
	10000	840_10000_LB 840_10000_ZL		
1680	3000	1680_3000_LB 1680_3000_ZL	4500	1680_4500G_LB 1680_4500G_ZL
	4500	1680_4500_LB 1680_4500_ZL		
	7000	1680_7000_LB 1680_7000_ZL		
	10000	1680_10000_LB 1680_10000_ZL		
1680_Alk12	7000	1680_12_7000_LB	-	-
1680_Alk15	3000	1680_15_3000_LB	-	-
	4500	1680_15_4500_LB		
	7000	1680_15_7000_LB		
	10000	1680_15_10000_LB		
1680_Alk2	7000	1680_2_7000_LB	-	-
2520	3000	2520_3000_LB 2520_3000_ZL	4500	2520_4500G_LB 2520_4500G_ZL
	4500	2520_4500_LB 2520_4500_ZL		
	7000	2520_7000_LB 2520_7000_ZL		
	10000	2520_10000_LB 2520_10000_ZL		