Warm arctic continents during the Palaeocene–Eocene thermal maximum

Johan W.H. Weijers a,1, Stefan Schouten a, Appy Sluijs b, Henk Brinkhuis b, Jaap S. Sinninghe Damsté a,c,*

a Royal Netherlands Institute for Sea Research (NIOZ), Department of Marine Biogeochemistry and Toxicology, P.O. Box 59, 1790 AB Den Burg — Texel, The Netherlands
b Palaeoecology, Institute of Environmental Biology, Utrecht University, Laboratory of Palaeobotany and Palynology, Budapestlaan 4, 3584 CD Utrecht, The Netherlands
c Department of Earth Sciences, Utrecht University, Budapestlaan 4, 3584 CD, Utrecht, The Netherlands

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Abstract

The Palaeocene–Eocene Thermal Maximum (PETM; ~55.5 Ma) is a geologically relatively brief episode of extreme warmth. Both deep and surface ocean temperatures increased by up to 5 °C in equatorial waters and up to 8 °C in mid and high latitude waters. From the continents, the annual mean air temperature response during the PETM is still largely unknown, mainly due to a lack of quantitative temperature proxies and sufficient suitable, continuous high resolution records. Recently, a new proxy for continental temperature reconstructions has been proposed, based on the distribution of membrane lipids of bacteria in present-day soils [J.W.H. Weijers, S. Schouten, J.C. van den Donker, E.C. Hopmans, J.S. Sinninghe Damsté (2007) Environmental controls on bacterial tetaether membrane lipid distribution in soils, Geochimica et Cosmochimica Acta 71, 703–713] and shown to reconstruct annual mean air temperature. In this study we applied this new proxy in an attempt to reconstruct the air temperature in high latitude continental areas during the PETM by analysis of a marine sedimentary sequence obtained from the Lomonosov Ridge in the central Arctic Ocean (Integrated Ocean Drilling Program Expedition 302, Site 004, Hole A). The results indicate a warming of ~8 °C above background values of ~17 °C. This warming is coincident with a similar rise in sea surface temperatures documented earlier. Our results thus further confirm the warm conditions in the Arctic, and point to a strongly reduced latitudinal temperature gradient during the PETM.

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* Corresponding author. Tel.: +31 222 369550; fax: +31 222 319674.
E-mail addresses: jweijers@nioz.nl, johan.weijers@bristol.ac.uk (J.W.H. Weijers), schouten@nioz.nl (S. Schouten), a.sluijs@uu.nl (A. Sluijs), h.brinkhuis@uu.nl (H. Brinkhuis), damste@nioz.nl (J.S. Sinninghe Damsté).
1 Currently at: Organic Geochemistry Unit, Bristol Biogeochemistry Research Centre, School of Chemistry, University of Bristol, Cantock’s Close, Bristol, BS8 1TS, United Kingdom.

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1. Introduction

Superimposed on the already warm climates of the early Cenozoic, the Palaeocene–Eocene Thermal Maximum (PETM) at ~55.5 million years ago represents a geologically relatively brief episode (~170 kyr; Sluijs et al., in press) of extreme global warming. Sediments deposited during the PETM are also characterized by a strong negative excursion (2.5 to 6‰) in the stable carbon isotope (δ13C) records of carbonates and organic carbon in both the marine and terrestrial realms (e.g., Kennett and Stott, 1991; Koch et al., 1992; Wing et al., 2005; Sluijs et al., 2006; Schouten et al., 2007). This excursion is considered to be associated with the release of massive amounts of 13C-depleted carbon into the ocean and atmosphere system (e.g., Dickens et al., 1995; Pagani et al., 2006a and references therein; Storey et al., 2007). Biotic responses to the climatic changes during the PETM include rapid floral range changes (Wing et al., 2005), large scale faunal dispersion (Koch et al., 1992; Smith et al., 2006), as well as massive benthic extinctions (Kennett and Stott, 1991) and proliferation of exotic foraminifera (Kelly et al., 1996) and the dinoflagellate taxon ‘Apectodinium’ (e.g., Bujak and Brinkhuis, 1998; Crouch et al., 2001).

During the PETM, the deep oceans warmed globally by 4 to 6 °C (e.g., Kennett and Stott, 1991; Bralower et al., 1995; Tripati and Elderfield, 2005). Sea surface temperatures (SST) in the tropics showed a warming of up to 5 °C (Bralower et al., 1995; Thomas et al., 1999; Zachos et al., 2003) while stronger warming of 5 to 8 °C has been reported for the mid and high latitude oceans (Kennett and Stott, 1991; Zachos et al., 2003; Sluijs et al., 2006; Zachos et al., 2006). Compared to the marine realm relatively little is known about changes in the annual mean air temperature (MAT) on the continents during the PETM, due to a lack of quantitative temperature proxies in combination with the relative short duration of the PETM. Several qualitative climate reconstructions indicate generally warmer conditions throughout on the continents around the PETM, notably at higher latitudes (e.g., Robert and Kennett, 1994; Markwick, 1998; Francis and Poole, 2002). Based, for example, on leaf margin analysis and oxygen isotope analysis of fossil teeth enamel, a temperature increase of about 5 to 7 °C has been inferred for the PETM in mid latitude North America (Fricke and Wing, 2004; Wing et al., 2005). Unfortunately, however, continuous high resolution records of MAT on the continents are not available for the PETM interval.

Recently, a new method for estimating terrestrial annual MAT has been developed based on the relative distribution of branched glycerol dialkyl glycerol tetraether (GDGT) membrane lipids derived from bacteria thriving in soils (Weijers et al., 2006, 2007b). The distribution of branched GDGTs in a globally distributed set of modern soils, expressed in the Methylation index of Branched Tetraethers (MBT) and the Cyclisation ratio of Branched Tetraethers (CBT), shows a significant linear correlation with modern annual MAT in the range of −6 to 27 °C (Weijers et al., 2007b). Since these membrane lipids are fluvially transported to the oceans (Hopmans et al., 2004), analysing marine sedimentary records in front of large river outflows potentially provides continuous high resolution records of river basin integrated continental temperature signals. Based on a sediment record from the Congo deep sea fan, for example, Weijers et al. (2007a) were able to reconstruct the change in annual MAT over the last deglaciation for the Congo River basin, covering almost the whole of tropical central Africa. Here, we apply this new method to a sedimentary sequence from the Lomonosov Ridge in the Arctic Ocean covering the PETM interval. During the PETM this site received considerable fluvial terrestrial organic matter input (Sluijs et al., 2006). By analysing the distribution of the soil-derived GDGT lipids in this sediment core we aimed to reconstruct annual MAT for the high latitude Arctic landmass during the PETM. In addition, palynomorph assemblages were analysed in order to constrain the geographic source of the temperature signal in this core.

2. Methods

2.1. Sample and site description

Marine sediment samples are derived from Integrated Ocean Drilling Program (IODP) Hole 302-4A (~87° 52.00′ N; 13° 10.64′ E; 1288 m water depth), positioned on the Lomonosov Ridge in the central Arctic Ocean (Fig. 1), recovered during IODP Expedition 302 (or Arctic Coring Expedition; ACEX) in August 2004 (Backman et al., 2006). This marine sedimentary sequence comprises a Palaeogene record containing the PETM interval between 391 and 378 m composite depth (mcd) (Fig. 3). Due to drilling disturbances and incomplete recovery the lower boundary of the PETM interval and the exact stratigraphic position of Core 31X are somewhat problematic (see Sluijs et al., 2006). In total 43 samples have been analysed (see data table in electronic Appendix) with a resolution of about 5 samples per meter, or, for the PETM interval, a time resolution of ca. 1 sample per 5 kyr using a ~170 kyr duration of the PETM (Sluijs et al., in press).
The Lomonosov Ridge represents a part of the Barentsz Shelf that rifted from the Svalbard margin and subsequently drifted away towards the pole where it submerged to its present depth during the Cenozoic (Jokat et al., 1992). During the late Palaeocene, this location was strongly influenced by fresh water input as evident from organic walled dinoflagellate cyst (dino-cyst) assemblages and high proportions of terrestrial palynomorphs (Sluijs et al., 2006). The PETM section has been recognized on basis of the concomitant occurrence of a \( \sim 6\% \) negative excursion in \( \delta^{13}C_{\text{org}} \) values and the occurrence of the dinocyst species \textit{Apectodinium augustum}, which is diagnostic of the PETM (Bujak and Brinkhuis, 1998; Sluijs et al., 2006).

2.2. Sample preparation and HPLC/MS analysis

Powdered and freeze dried sediments (1–3 g dry mass) were extracted with an accelerated solvent extractor ( Dionex2000), using a dichloromethane (DCM):MeOH 9:1 (v/v) solvent mixture, 3 times for 5 min at a pressure of ca 7.6 \times 10^6 \text{ Pa} and a temperature of 100 °C. The obtained total extracts were rotary evaporated and separated over an activated \( \text{Al}_2\text{O}_3 \) column using hexane:DCM 9:1 (v/v) and DCM:MeOH 1:1 (v/v) solvent mixtures into an apolar and polar fraction, respectively. The polar fraction, containing the branched GDGTs, was dried under a pure \( \text{N}_2 \) flow, dissolved ultrasonically in a hexane:propanol 99:1 (v/v) mixture at a concentration of 2 mg ml\(^{-1}\) and filtered over an 0.45 μm mesh PTFE filter (ø 4 mm) prior to HPLC/MS analysis.

High performance liquid chromatography–atmospheric pressure chemical ionization/mass spectrometry (HPLC–APCI/MS) analyses were performed on an Agilent 1100 series/Hewlett–Packard 1100 MSD series machine equipped with auto-injector and HP Chemstation software as described previously (Hopmans et al., 2000) with minor modifications. Injection volume was usually 10 μl and separation of the compounds was achieved in normal phase on a Prevail Cyano column (150 mm × 2.1 mm; 3 μm; Alltech). The flow rate of the hexane:propanol 99:1 (v/v) eluent was 0.2 ml min\(^{-1}\), isocratically for the first 5 min and thereafter with a linear gradient to 1.8% propanol in 45 min. Analyses were performed in selective ion monitoring mode in order to increase sensitivity and reproducibility. Branched GDGTs were quantified by integrating the area of the [M+H]\(^+\) peaks (protonated molecular ion).
and comparing these with an external standard curve prepared with known amounts of the isoprenoid GDGT crenarchaeol. The CBT ratio and MBT index were calculated as follows:

\[
\text{CBT} = -\log\left(\frac{[\text{Ib}] + [\text{IIb}]}{[\text{I}] + [\text{II}]}\right)
\]

\[
\text{MBT} = \frac{([\text{I}] + [\text{Ib}] + [\text{Ic}])}{([\text{I}] + [\text{Ib}] + [\text{Ic}]) + ([\text{II}] + [\text{IIb}] + [\text{IIc}]) + ([\text{III}] + [\text{IIIb}] + [\text{IIIC}]})
\]

Roman numerals correspond to the GDGT structures drawn in Fig. 2. Annual MAT was then calculated from the CBT and MBT values using the calibration equation given by Weijers et al. (2007b):

\[
\text{MBT} = 0.122 + 0.187\times\text{CBT} + 0.020\times\text{MAT} \quad (R^2 = 0.77).
\]

Based on duplicate measurement of the whole data series the average standard deviation of the absolute temperature estimates (analytical reproducibility) has been determined at 0.4 °C.

2.3. Palynological analysis

Sediments were oven-dried at 60 °C. Samples were treated with 30% HCl and twice with 30% HF for carbonate and silicate removal, respectively. After sieving over a 15-μm nylon mesh sieve, residues were analysed at 500× magnification.

3. Results and discussion

3.1. Temperature estimates based on soil derived branched GDGT lipids

Concentrations of the soil derived branched GDGT membrane lipids are ∼90 μg g⁻¹ dry weight sediment (dws) during the PETM interval which is higher than the values of 10 and 60 μg g⁻¹ dws obtained from the pre- and post-PETM sections of this core, respectively. Based on the rather constant and substantial TOC contents (∼2%) in combination with well preserved palynomorphs both within and outside the PETM section, this difference is unlikely to be the result of differences in preservation conditions (Sluijs et al., 2006). Instead, the higher concentrations of branched GDGTs during the PETM might suggest higher productivity of the terrestrial vegetation and/or enhanced river discharge during the PETM, which is consistent with the more humid conditions at the North Pole at that time (Pagani et al., 2006b). Since terrestrial organic matter input was high, it was possible to determine the degree of cyclisation (CBT) and methylation (MBT) of these terrestrial derived branched GDGTs in the sediments of this core and hence to reconstruct the continental annual MAT. The geographic source of the branched GDGTs in the core from the Lomonosov Ridge determines the continental area for which the MAT is reconstructed. Spreading along the Gakkel Ridge had already separated the Lomonosov Ridge from the Siberian Shelf by the latest Palaeocene (Jokat et al., 1992). Palaeoenvironmental and tectonic studies have indicated that the coring site itself was only shallowly submerged at the PETM (Backman et al., 2006; Sluijs et al., 2006). Hence, it could be argued that the bulk of the terrestrial material was derived from parts of the Lomonosov Ridge close to the coring site that were still above sea level. This hypothesis is supported by palynological data. In upper
Palaeocene and lower Eocene strata, spores produced by the fern family Osmundaceae are consistently present and locally abundant (Fig. 3). These spores are very large and cannot have been transported over long distances, and thus the site must have been very close to shore. The consistent occurrence of Osmundaceae spores implies that parts of the Lomonosov Ridge close to the core site were indeed still above sea level and were the source of these Osmundaceae spores and most likely the main portion of the branched GDGTs. The reconstructed MAT is thus thought to represent central Arctic polar conditions. The reconstructed annual MAT shows values of 16 to 18 °C for the latest Palaeocene and increases to values up to 25 °C during the PETM (Fig. 3). Towards the end of the PETM, temperatures drop again to values of around 18 to 20 °C. This transient temperature change parallels the negative δ¹³Corg excursion and the acme of the subtropical dinoflagellate *Apectodinium augustum* (Sluijs et al., 2006). Higher temperatures occur at the time of lower δ¹³Corg values and the subsequent cooling mirrors the recovery pattern of δ¹³Corg (Fig. 3).

Our MAT record for the central Arctic Lomonosov Ridge compares well with the SST reconstruction for the Arctic Ocean at this location by Sluijs et al. (2006). On the basis of the TEX₈₆ proxy, their reconstruction shows SSTs of ~18 °C for the latest Palaeocene, >23 °C for the PETM, and a cooling towards ~17 °C at the end of the PETM (Fig. 3). Thus, both proxies indicate a strong climatic warming in the Arctic region during the PETM, and a cooling towards the end of the PETM consistent with the recovery of the δ¹³Corg signal. Remarkably, the absolute temperatures estimated by both proxies are also similar, particularly so when taking into account uncertainties like the exact temporal origin of the temperature signals in this high latitude area, i.e. annual mean or seasonal temperatures. It has to be emphasised that both organic proxies are based on different compounds and source organisms. Where the TEX₈₆ SST proxy is based on the membrane lipids of marine Crenarchaeota which are ubiquitous in marine sediments (Schouten et al., 2002), the MBT MAT proxy is based on soil-derived branched GDGT membrane...
lipids which are only detected in trace amounts in open marine settings and thus indicative for a predominant terrestrial source (Hopmans et al., 2004). Both proxies are, moreover, differently calibrated and hence totally independent. The similarity between both records, therefore, emphasises once again that temperature is exerting the main control on GDGT lipid distributions.

In addition, the similar absolute estimates for both land and sea surface temperatures are a first and strong independent confirmation that temperatures in the Arctic during the PETM were indeed extremely high compared to the present-day situation.

Table 1
Overview of Palaeocene–Eocene continental temperature estimates

<table>
<thead>
<tr>
<th>Time interval</th>
<th>Location</th>
<th>Palaeo Lat.</th>
<th>Method</th>
<th>Temperature</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Early Eocene</td>
<td>Lomonosov Ridge</td>
<td>∼ 75° N</td>
<td>MBT/CBT</td>
<td>∼ 17 °C</td>
<td>This study</td>
</tr>
<tr>
<td></td>
<td>Ellesmere Island Canada</td>
<td>71° N</td>
<td>Crocodile remains</td>
<td>≥ 14 °C</td>
<td>Markwick (1998)</td>
</tr>
<tr>
<td></td>
<td>Bighorn Basin Wyoming, USA</td>
<td>45° N</td>
<td>Leaf margin analyses</td>
<td>∼ 18 °C</td>
<td>Wing et al. (2000)</td>
</tr>
<tr>
<td>PETM</td>
<td>Lomonosov Ridge</td>
<td>∼ 75° N</td>
<td>MBT/CBT</td>
<td>∼ 25 °C</td>
<td>This study</td>
</tr>
<tr>
<td></td>
<td>Coastal continent Antarctica</td>
<td>65° S</td>
<td>Clay mineral composition</td>
<td>≥ 15 °C</td>
<td>Robert and Kennett (1994)</td>
</tr>
<tr>
<td></td>
<td>Bighorn Basin Wyoming, USA</td>
<td>45° N</td>
<td>Leaf margin analyses</td>
<td>∼ 20 °C</td>
<td>Wing et al. (2005)</td>
</tr>
<tr>
<td>Late Palaeocene</td>
<td>Lomonosov Ridge</td>
<td>∼ 75° N</td>
<td>MBT/CBT</td>
<td>∼ 19 °C</td>
<td>This study</td>
</tr>
<tr>
<td></td>
<td>Bighorn Basin Wyoming, USA</td>
<td>45° N</td>
<td>Leaf margin analyses</td>
<td>∼ 15 °C</td>
<td>Wing et al. (2000)</td>
</tr>
</tbody>
</table>

a Minimum mean annual air temperature required for the presence of crocodiles.
b Underestimate; see text for discussion.
c This represents the latitude of the Ocean Drilling Program Site 690B at Maud Rise; Antarctic coastline is at ∼ 70° S.
d Formation temperature of kaolinite and thus minimum soil temperature; the overall clay mineral composition is indicative of sub-tropical conditions.

temperatures might be expected mainly because the season of maximum growth likely occurred during the polar day (summer) season, but also because the estimated MAT is similar to the TEX 86 SST for which a summer bias was already invoked (Sluijs et al., 2006). Branched GDGT membrane lipids, however, are produced by ostensibly non-phototrophic bacteria thriving in the anoxic parts of soils and peat bogs (Weijers et al., 2006). Therefore, these bacteria are expected not to be affected by polar night/day conditions. As, in addition, frost conditions are suggested to be absent during the PETM (Bice et al., 1996; Tripati et al., 2001; Jahren, 2007), especially on this ridge surrounded by warm Arctic waters (Sluijs et al., 2006), there is no a priori reason why the MBT proxy is recording summer temperatures alone. Other possibilities, like changing nutrient conditions in soils during polar night conditions cannot be excluded but remain purely speculative. Despite these uncertainties, the relatively smooth trend of our MAT record and the parallel evolution with the TEX 86 δ 13 C org records strongly suggest that air temperatures at the high latitude Arctic Lomonosov Ridge were high during the PETM and increased by up to 8 °C above background values.

3.2. Comparison to other Palaeocene–Eocene temperature estimates

As noted above, and in contrast to the marine realm, terrestrial temperature estimates for the PETM interval are very scarce. For high latitude regions only two semi-quantitative temperature estimates, i.e. minimum estimates, are available (Robert and Kennett, 1994; Markwick, 1998, see Table 1). In addition to the minimum temperature provided by the clay mineral assemblage derived from coastal Antarctica, its composition during
the PETM interval compares to that found in modern tropical to subtropical Atlantic sediments, which even suggests subtropical conditions on coastal Antarctica during the PETM (Robert and Kennett, 1994). Absolute estimates of annual MAT for the PETM are only derived from the Big Horn Basin area, Wyoming USA, which was located at a palaeolatitude of ∼45° N. Temperature estimates based on leaf margin analysis (Wing et al., 2000, 2005) are ∼6 °C lower than temperatures derived from oxygen isotope ratio analyses of phosphate in teeth from fossil animals from the same area (Fricke and Wing, 2004) (Table 1). This could be explained by the fact that florals of predominantly wet-soil environments, as used in the Big Horn Basin study, generally show a greater proportion of toothed leaves, resulting in possible temperature under-estimates of up to 10 °C (Kowalski and Dilcher, 2003). Considering this possible bias, this implies that subtropical to tropical temperature conditions might have occurred at the mid-latitude North American continent during the PETM. Remarkably, these mid-latitude PETM temperature estimates are, even if taking into account the uncertainties accompanying the different methods, at most similar to the maximum PETM temperatures reconstructed for the central Arctic Lomonosov Ridge and likely not substantially higher. One possible explanation might be that the peak PETM warmth was missed in the Bighorn Basin section due to the relatively poor stratigraphic resolution of the samples. On the other hand, due to the strong seasonal difference at the high latitudes, our Arctic air temperature estimates might indeed represent maximum (summer) estimates. Alternatively, our data are in support of the idea that the latitudinal thermal gradient between the high and mid latitudes was very small during the PETM, a feature which seems to be characteristic of greenhouse climates (e.g., Huber et al., 1995).

3.3. Implications

Our record of polar continental PETM air temperatures, in conjunction with the few other available high latitude temperature estimates (Kennett and Stott, 1991; Robert and Kennett, 1994; Sluijs et al., 2006), provide strong indications of subtropical conditions at the Earth’s polar regions during the PETM. This has considerable implications for our knowledge of the Earth’s climate state under elevated greenhouse gas levels in an ice free world. Several modelling studies have been conducted to simulate the response of Earth’s climate to high greenhouse gas levels for the PETM (e.g., Schmidt and Shindell, 2003; Shellito et al., 2003; Renssen et al., 2004; Shellito and Sloan, 2006). In agreement with proxy data (e.g., Kennett and Stott, 1991), these models show increased warming which is amplified at the high latitudes, i.e. ca. 5 to 8 °C (e.g., Renssen et al., 2004). However, the simulated absolute temperatures for the high latitudes are near freezing point (e.g., Shellito et al., 2003; Shellito and Sloan, 2006) and hence considerably lower than the indications from proxy data. Thus, proxy data indicate a latitudinal thermal gradient that is much flatter than simulated by climate models which generally yield gradients of ∼30 °C (e.g., Shellito et al., 2003; Shellito and Sloan, 2006).

Apparently, current climate models lack one or more physical mechanisms that are able to warm the polar regions without substantially increasing temperatures in the tropics during the early Cenozoic. Several reasons have been suggested explaining this model-data discrepancy. These include the presence of sea ice in models, a lack of proper boundary conditions like sea surface temperatures and topography, and an increased oceanic and/or atmospheric heat transport during the PETM. None of these, however, can explain the model-data discrepancy (Huber and Sloan, 1999, 2001; Sloan et al., 2001; Caballero and Langen, 2005). Considered to be more plausible feedback mechanisms in reducing the tropic-to-pole temperature gradient during the early Cenozoic in general and PETM in particular are a higher abundance of polar stratospheric clouds (Sloan and Pollard, 1998) and enhanced poleward oceanic heat transport induced by tropical cyclones (Sriver and Huber, 2007).

The seemingly sub-tropical conditions at the poles (Robert and Kennett, 1994; Sluijs et al., 2006 and this study) and conservative temperature estimates in excess of 33 °C for the mid latitudes (Zachos et al., 2006), might imply that temperatures at the equator were as high as 40 °C during the PETM. Published reconstructions of Cretaceous SSTs at low to mid latitudes using the TEX86, δ18O and Mg/Ca proxies (Schouten et al., 2003; Bice et al., 2006; Zachos et al., 2006; Forster et al., 2007) do suggest that tropical sea waters may have indeed been as high as 40 °C in the geological past. Thus, there may have been a latitudinal thermal gradient at the PETM. Nonetheless, it still must have been much smaller than today and, in addition, superimposed on temperatures that were globally substantially higher than currently assumed.

4. Conclusions

Using a new organic proxy for reconstructing continental air temperatures, the MBT index, we estimated temperatures for the Lomonosov Ridge in the Arctic
Ocean during the PETM. The reconstructed temperature signal is most likely derived from the Lomonosov Ridge itself, which was partially still above sea level. Air temperatures clearly increased by about 8 °C above already warm background temperatures, a similar amplitude as observed for high latitude sea surface waters. Although the exact temporal origin of the reconstructed temperature signal is somewhat uncertain, the estimated values of 17 °C prior to the PETM and 25 °C during the PETM match well with the reconstructed sea surface temperature for the Arctic Ocean at this time interval. These values confirm a much reduced latitudinal temperature gradient in the PETM greenhouse world compared to the present-day situation.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.epsl.2007.06.033.

References


