

Extreme warming of mid-latitude coastal ocean during the Paleocene-Eocene Thermal Maximum: Inferences from TEX₈₆ and isotope data

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ABSTRACT

Changes in sea surface temperature (SST) during the Paleocene-Eocene Thermal Maximum (PETM) have been estimated primarily from oxygen isotope and Mg/Ca records generated from deep-sea cores. Here we present a record of sea surface temperature change across the Paleocene-Eocene boundary for a nearshore, shallow marine section located on the eastern margin of North America. The SST record, as inferred from TEX₈₆ data, indicates a minimum of 8 °C of warming, with peak temperatures in excess of 33 °C. Similar SSTs are estimated from planktonic foraminifer oxygen isotope records, although the excursion is slightly larger. The slight offset in the oxygen isotope record may reflect on seasonally higher runoff and lower salinity.

Keywords: Paleocene, Eocene, isotopes, greenhouse.

INTRODUCTION

The Paleocene-Eocene Thermal Maximum (PETM) represents one of the more prominent and abrupt climate anomalies in Earth history with sea surface temperatures (SSTs) increasing by as much as 5 °C in the tropics and 8 °C in the high latitudes (Thomas et al., 2002; Zachos et al., 2003; Tripathi and Elderfield, 2004). The peak warmth was sustained for several tens of thousands of years before gradually returning to pre-event levels. Several lines of evidence indicate that a rise in greenhouse carbon levels (CH₄ and/or CO₂) was responsible for this global warming (e.g., Dickens et al., 1997; Bowen et al., 2004). The approximate mass of carbon released is still unknown, but has been estimated to be in excess of 2000 GtC (Dickens et al., 1997), and possibly as high as 4500 GtC (Zachos et al., 2005).

If the rise in SST documented in open ocean sites was a consequence of greenhouse warming, the SST in coastal oceans should have risen by as much, if not more. Moreover, coastal oceans would have been particularly sensitive to changes in runoff, and hence precipitation, though the response would have been highly variable both spatially and temporally. Indeed, previous investigations of shallow marine sequences have found evidence of significant environmental perturbation of the coastal oceans during the PETM,

including evidence of warming and changes in runoff (Bujak and Brinkhuis, 1998; Egger et al., 2003; Gibson et al., 1993). Much of the paleoclimatic information, however, has been derived from qualitative indexes such as fossil assemblages (Crouch et al., 2001, 2003), in part because traditional temperature proxies applied to deep-sea cores, such as oxygen isotopes, are not particularly well suited for application to shallow-marine, land-based sections. The general absence of planktonic foraminifera is one limitation. The effects of meteoric diagenesis, a process that can reset the primary oxygen isotopic composition of carbonates toward lower values, are another. Even where fossils are present and well preserved, deviations in local seawater salinity from the global mean increase the uncertainty in estimating temperature from δ¹⁸O, a problem that would have been compounded with rapid greenhouse warming and changes in precipitation and runoff.

In this investigation, we estimate coastal SST during the PETM in a shallow marine sequence using an organic-based proxy of SST, TEX₈₆, which is derived from the membrane lipids of marine crenarchaeota, a common component of picoplankton (Schouten et al., 2002, 2003). Studies of core top sediments have demonstrated a strong correlation between the number of cyclopentane rings in crenarchaeota lipids and mean annual SST (r²

= 0.92). Moreover, culture experiments show that changes in salinity and nutrients do not substantially affect the temperature signal recorded by TEX₈₆ (Wuchter et al., 2004), and it also seems to be unaffected by sedimentary redox conditions (Schouten et al., 2004). With the TEX₈₆-derived SST, we then use the oxygen isotopes to determine if this locality experienced substantial changes in salinity.

The section sampled for this study, Wilson Lake (Fig. 1), is located in New Jersey (39°39'N, 75°03'W) where the upper Paleocene–lower Eocene is accessible by coring. The Paleocene-Eocene boundary interval consists of unconsolidated siliciclastic sands and clays with low carbonate content (<15%) deposited during a sea-level transgression (Cramer et al., 1999; Gibson et al., 1993). Wilson Lake offers several advantages, one of which is high abundances of marine organic matter including dinoflagellates and crenar-

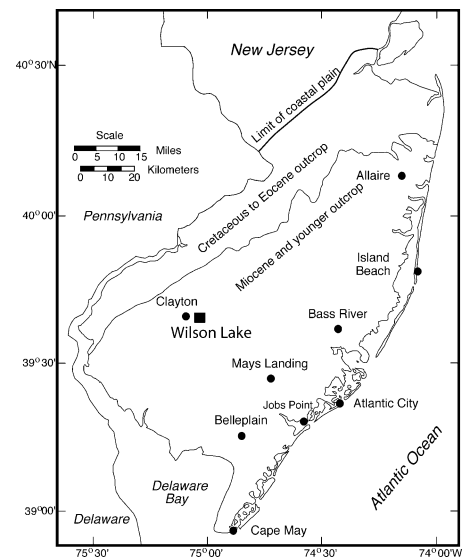


Figure 1. Location map showing the location of Wilson Lake (~39°39'N, 75°2'W) and other USGS and ODP cores (modified from Miller, 1997).

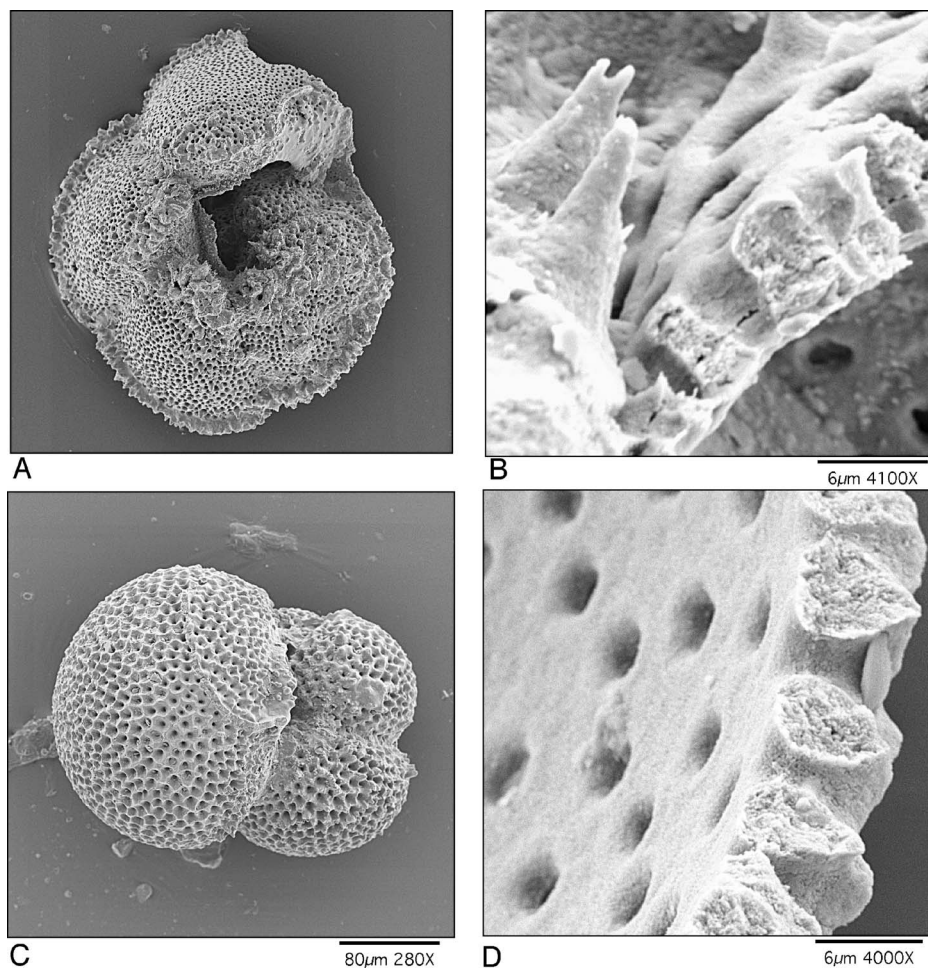


Figure 2. SEM photographs of planktonic foraminifera. **A:** *Morozovella* sp. (103.62 m). **B:** Cross section of shell wall of *Morozovella* sp. (103.62 m). **C:** *Subbotina* (104.58 m). **D:** Cross section of shell wall of *Subbotina* (104.58 m).

chaetotal lipids. Moreover, Wilson Lake samples yield well-preserved planktonic foraminifera with some shells exhibiting porcelain textures (Fig. 2), though poorly preserved specimens are present as well. The well-preserved shells should yield close to primary $\delta^{18}\text{O}$ values, which in combination with TEX_{86} , can be used to quantify changes in temperature as well as seawater $\delta^{18}\text{O}$.

FACIES DESCRIPTION AND METHODS

The Wilson Lake Paleocene-Eocene boundary section is marked by a distinct transition from glauconitic clayey sandstones to silty claystones. This, together with the absence of mollusks, suggests a middle shelf depositional setting, perhaps tens of kilometers offshore at a paleodepth between 25 and 100 m (Gibson et al., 2000). The uppermost Paleocene and lower Eocene were recovered near the bottom of the core between 92 and 112 m. Two unconformities are apparent in the lowermost Eocene (Gibbs et al., 2006), though the Paleocene-Eocene transition appears to be relatively complete. Flora representative of nanofossil zones NP9 and NP10 are present,

though the exact position of the boundary between these zones is uncertain.

Samples were collected every 20–40 cm over a 20 m interval, disaggregated and wet sieved to isolate the sand fraction from which foraminifera were collected. Stable-isotope analyses were carried out on planktonic and benthic foraminifera. The planktonic foraminifera included two taxa that resided in the mixed-layer, *Acarinina soldadoensis* and *Morozovella velascoensis (acuta)*, and a somewhat deeper dweller, *Subbotina triangularis*. Analyses were also carried out on benthic foraminifera *Cibicidoides*. Measurements were performed on an Autocarb coupled to a PRISM mass spectrometer at the University of California–Santa Cruz. Precision based on replicate analyses of in-house standard CM is better than ± 0.05 and $\pm 0.10\%$ for carbon and oxygen isotopes, respectively. All values are reported relative to VPDB.

For the TEX_{86} index, ~ 20 fine-fraction ($< 63 \mu\text{m}$) samples were selected and analyzed by high-performance liquid chromatography/atmospheric pressure positive ion chemical ionization–mass spectrometry

(Schouten et al., 2002). Briefly, the fine fractions were extracted with a Dionex Accelerated Solvent Extractor using a mixture of dichloromethane (DCM) and methanol (MeOH). The extract was fractionated into apolar and polar fractions, containing the crenarchaeotal lipids using a small column with activated alumina and using hexane/DCM (9:1 by volume) and DCM/MeOH (1:1 by volume) as effluents, respectively. Aliquots of polar fractions were dissolved in hexane/propanol (99:1 by volume) and filtered through $0.45 \mu\text{m}$ PTFE filters. The samples were analyzed with a Thermo Finnigan Quantum Ultra triple-quadrupole LC-MS, and separation was performed on an Econosphere NH_2 column, maintained at 30°C . The glycerol dialkyl glycerol tetraethers (GDGTs) were eluted using a changing mixture of hexane/propanol (99:1 for 5 min, then a linear gradient to 1.8 propanol in 45 min). Detection was achieved using atmospheric pressure chemical ionization–mass spectrometry of the eluent. Single-ion monitoring (SIM) was set to scan the five $[\text{M}^+]\text{H}$ ions of the GDGTs with a dwell time of 237 ms for each ion. All TEX_{86} analyses were performed at least in duplicate. The concentration of branched and isoprenoid tetraether lipids (BIT index) was measured on five samples to constrain the relative concentration of terrestrial organic matter (Hopmans et al., 2004).

RESULTS

The Wilson Lake foraminifera show distinct interspecies carbon isotope patterns not unlike those found in pelagic settings. For example, mixed-layer species, *M. velascoensis* and *A. soldadoensis*, yield the highest carbon isotope values, consistent with a near-surface habitat, while *S. triangularis* and benthic foraminifera yield the lowest carbon isotope values. The foraminiferal oxygen isotope values, on the other hand, exhibit weaker gradients, and in some intervals none at all.

The most prominent feature of the isotope records are large negative excursions in both carbon and oxygen isotopes across the Paleocene-Eocene boundary (110–109 m) (Fig. 3). The foraminifer $\delta^{13}\text{C}$ values decrease by 3% – 4% , while the $\delta^{18}\text{O}$ values decrease by 2.0% – 2.5% . Minimum $\delta^{13}\text{C}$ values of -3.5% are recorded by the benthic foraminifera, and $\delta^{18}\text{O}$ values of -4.3% by the mixed-layer planktonic foraminifera. These low $\delta^{13}\text{C}$ values are sustained over a 13 m interval to the base of the lower unconformity at ~ 96 m. After the initial $\delta^{18}\text{O}$ decrease in the mixed-layer foraminifer, the records deviate with the *A. soldadoensis* values increasing to levels similar to or lower than the benthics, while the *M. velascoensis* values remain low ($\sim -4.0\%$).

The TEX_{86} index shows a sharp increase across the boundary that is essentially coincident with the decrease in foraminiferal ox-

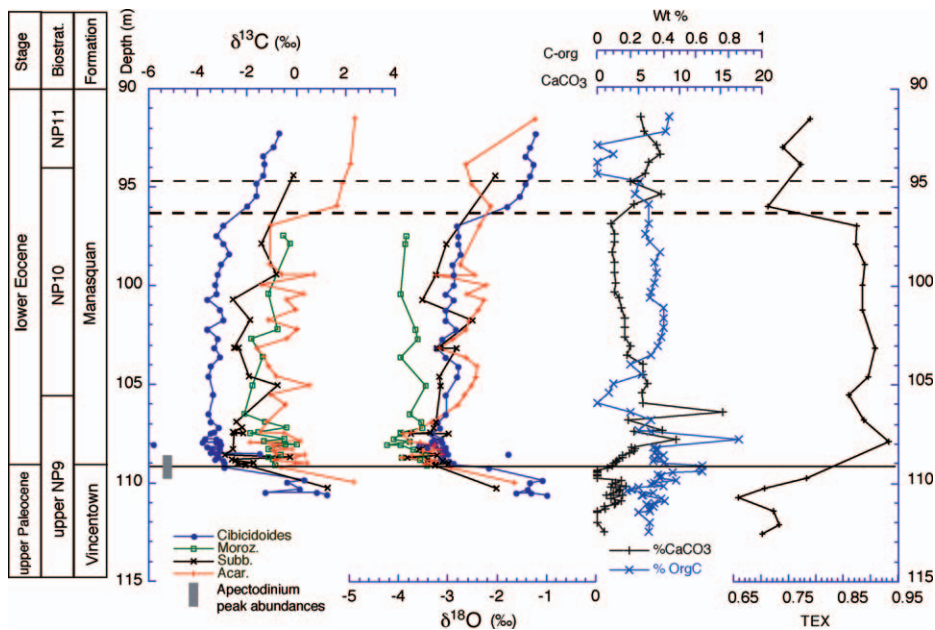


Figure 3. The columns to the far left show the lithology and nannofossil biostratigraphic zonation for Wilson Lake plotted versus subsurface depth (m). The biostratigraphic scheme follows the NP scheme of Martini (1971) where the NP9/NP10 boundary is defined as the first occurrence of *Rhombaster/Tribrachiatus bramlettei* and the NP10/NP11 boundary is approximated by the first occurrence of *T. orthostylus*. Stable-isotope, weight percent Corg and CaCO₃, and TEX₈₆ raw data are plotted versus subsurface depth. The stable-isotope data are from analyses of *Morozovella velascoensis (acuta)*, *Acarinina soldadoensis*, *Subbotina* spp., and *Cibicidoides* spp. The dashed lines at 94.79 and 96.32 m represent unconformities. The lower unconformity truncates the upper portion of the excursion layer. Gray bar in the left panel shows the level of the dinoflagellate *Apectodinium* abundance acme.

xygen isotope values. Application of the modern calibration to these values yields an increase in temperature from 31 to 40 °C at the height of the PETM, which are exceedingly high temperatures. However, the modern calibration is based on empirical core top data from 0 to 28 °C (Schouten et al., 2002, 2003). As a result, it was necessary to extrapolate out to higher TEX₈₆ values to interpret the SSTs. Therefore, we applied the more conservative calibration line based on core top data from 20 to 28 °C as proposed by Schouten et al. (2003) for SST >28 °C. This results in temperatures ranging from 25 °C prior to and after the PETM to 33 °C at the peak of the event. The BIT index for the five samples analyzed ranged between 0.03 and 0.11 (GSA Data Repository Table DR1¹), suggesting relatively low contributions of terrestrial organic matter (Hopmans et al., 2004).

Dinoflagellates and Palynomorphs

Palynological assemblages from Wilson Lake are characterized by the persistent dominance of dinocysts over other palynomorphs,

¹GSA Data Repository item 2006155, Table DR1, stable isotope, total inorganic and organic carbon, and dinoflagellate abundance data, is available online at www.geosociety.org/pubs/ft2006.htm, or on request from editing@geosociety.org or Documents Secretary, GSA, P.O. Box 9140, Boulder, CO 80301, USA.

including pollen. The dinocyst succession is marked by the successive dominance of typical late Paleocene–early Eocene taxa such as *Areoligera*, *Spiniferites*, *Cordosphaeridium*, *Senegalinium*, *Membranosphaera*, and, notably, *Apectodinium*. The global acme of the latter taxon is also recorded at Wilson Lake, peaking only at the onset of the PETM (e.g., Bujak and Brinkhuis, 1998; Crouch et al., 2001, 2003). The peak abundances of *Apectodinium* fall between 109.42 and 108.69 m, preceding slightly the maximum temperatures derived from $\delta^{18}\text{O}$ and TEX₈₆. An additional peak of *Apectodinium* is recorded in the upper part of the carbon isotope excursion. The sediments are nearly barren of terrestrial palynomorphs, an observation that is consistent with the BIT index, suggesting that either river discharge occurred far from the drill site, or vegetation was scarce in the hinterland of Wilson Lake.

DISCUSSION

Prior to this work, investigations that have attempted to constrain SST changes across the PETM have mostly focused on the magnitude of the anomalies rather than absolute temperatures (e.g., Thomas et al., 1999; Zachos et al., 2003; Tripathi and Elderfield, 2004), in part because of potential preservational artifacts (Schrag et al., 1995). The peak SST of 33 °C estimated from TEX₈₆ for this locality is high,

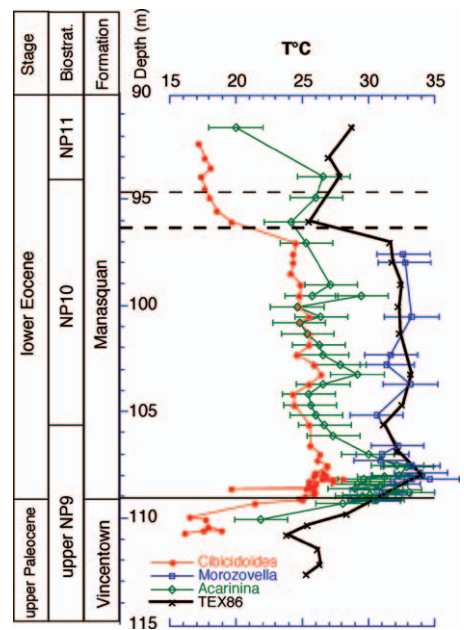


Figure 4. Sea surface temperatures as computed from (1) planktonic foraminifera $\delta^{18}\text{O}$ and (2) the TEX₈₆. The oxygen isotope-based curves were derived assuming seawater $\delta^{18}\text{O}_{\text{sw}}$ of -0.5‰ (SMOW) using standard paleotemperature equation (Erez and Luz, 1983). The errors bars on the planktonic foraminifera curves reflect the range of estimated temperatures associated with just a $\pm 0.5\text{‰}$ uncertainty in $\delta^{18}\text{O}_{\text{sw}}$.

especially if it is viewed as an annual mean, rather than summer maximum. In comparison, modern SST along this coast (over the shelf) ranges from 4 °C in winter to 28 °C in summer, with an annual mean of ~ 17 °C. Because coastal ocean temperatures often have a strong local/regional overprint, it is probably not valid to assume these paleotemperatures were representative of open Atlantic SST at this latitude. Nevertheless, based on GCM simulations, it appears a zonally averaged summer temperature of 33 °C for this paleolatitude ($\sim 35^{\circ}\text{--}37^{\circ}\text{N}$ at 55 Ma) would require a CO₂ concentration in excess of 2000 ppm (Shellito et al., 2003).

Modern calibration of TEX₈₆ is limited to temperatures below 28 °C, making the estimates of absolute temperatures above this value somewhat suspect. Yet, the absolute temperatures computed here are well within the range estimated from oxygen isotopes. In fact, if we use $\delta^{18}\text{O}_{\text{shell}}$ to estimate temperature assuming an ice-free world (mean ocean $\delta^{18}\text{O}$ of -1.0‰), but with a local $\delta^{18}\text{O}_{\text{sw}}$ of -0.5‰ due to evaporation (Zachos et al., 1994), the planktonic foraminiferal temperatures derived for the earliest Eocene are essentially identical to the TEX₈₆ temperatures, though the upper Paleocene temperatures are offset by 2 °C (Fig. 4). Alternatively, if we just consider the temperature anomaly interpreted from TEX₈₆ values (+8 °C), we can estimate relative changes in $\delta^{18}\text{O}_{\text{sw}}$ /salinity using the plankton-

ic foraminiferal oxygen isotope records. An 8 °C rise in temperature should lower $\delta^{18}\text{O}_{\text{shell}}$ by $\sim 1.70\%$. The benthic and *A. soldadoensis* excursions were roughly -1.85% and -2.2% , respectively, implying a possible $\delta^{18}\text{O}_{\text{sw}}$ change of -0.20% to -0.50% . The discrepancy could reflect a decrease in local sea surface salinity (SSS) (and $\delta^{18}\text{O}_{\text{sw}}$) due to higher runoff during the PETM. Assuming a $\Delta\delta^{18}\text{O}/\Delta\text{salinity}$ relationship of $0.15\%/ppt$ (Fairbanks, 1982), the -0.50% residual ($\Delta\delta^{18}\text{O}_{\text{sw}} = \Delta\delta^{18}\text{O}_{\text{shell}} - \Delta\delta^{18}\text{O}_{\text{TEX}}$) would require a modest salinity decrease of roughly 3–4 ppt.

Is a shift toward higher regional runoff and precipitation supported by the other lithologic and paleontologic data? The clay-rich excursion layer is relatively thick and dominated by kaolinite, patterns that have been observed elsewhere and attributed to higher humidity and more intense chemical weathering and runoff (e.g., Gibson et al., 2000; Egger et al., 2003). The *Apectodinium* acme is also associated with higher temperatures and enhanced runoff, stratification, and eutrophic conditions in coastal waters (Bujak and Brinkhuis, 1998; Crouch et al., 2003; Egger et al., 2003). This genus is morphologically very similar to modern cysts almost exclusively produced by heterotrophic dinoflagellates and thus would have required nutrient-rich conditions (Bujak and Brinkhuis, 1998). Nannofossil assemblages also indicate increased fertility during the PETM at Wilson Lake (Gibbs et al., 2006). Increased discharge by rivers likely supplied the necessary nutrients to fertilize the coastal ocean. On the other hand, there is very little terrestrial organic matter in this core. One possibility is that regional climate in this region became more seasonally extreme during the PETM, with a brief, intense wet season and prolonged dry season. Under this climate regime, the local landscape would have been sparsely vegetated and thus prone to excessive erosion during the wet season, which would explain both the increased flux of terrigenous sediment and scarcity of terrestrial organic matter.

Although the absolute SST and SSS values estimated for this location should be viewed with some caution until the uncertainties in the TEX_{86} temperature calibration are reduced, the estimated peak temperature of 33 °C is substantially higher than would be estimated from $\delta^{18}\text{O}$ of planktonic foraminifera (~ 25 °C) from tropical or subtropical deep-sea cores, consistent with the notion that the latter are biased toward heavier $\delta^{18}\text{O}$ values and colder temperatures (e.g., Schrag et al., 1995). As such, this coupled TEX_{86} /isotope approach shows promise for quantifying both absolute temperature and salinity change during the PETM, and thus should be applied to other clay-rich, shelf sections.

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