Subtropical sea-surface warming and increased salinity during Eocene Thermal Maximum 2

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ABSTRACT
Eocene Thermal Maximum 2 (ETM-2; ca. 54.2 Ma) represents the second largest of the major Eocene hyperthermals, yet comparatively little is known about the scale and rate of climatic change for key regions. Here we provide the first detailed records of subtropical sea-surface warming and salinization for ETM-2 at two subtropical locations, Ocean Drilling Program Sites 1209 (North Pacific) and 1265 (South Atlantic). Coupled planktic foraminiferal Mg/Ca and \( \delta^{18}O \) indicate 2–4 °C of rapid warming and local salinization of ~1–2 ppt at both sites. The increase in sea-surface temperature is equivalent to anomalies reported from higher latitude sites, and is consistent with theory on the expected pattern of spatial temperature response to greenhouse gas forcing in an ice-free world (i.e., no ice-albedo feedback). Similarly, the observed salinization is consistent with the hypothesis of enhanced meridional vapor transport and increased subtropical aridity in a warmer world.

INTRODUCTION
The early Eocene hyperthermals represent potential calibration points for establishing the response of global and regional warming and shifts in the hydrologic cycle to past changes in greenhouse gas (GHG) forcing because they represent transient and extreme warming events. The magnitude of warming (4–8 °C; e.g., Dunkley Jones et al., 2013) during the largest hyperthermal, the Paleocene–Eocene Thermal Maximum (PETM; ca. 56 Ma), and evidence for intensification of the hydrologic cycle (e.g., Schmitz and Pujalte, 2007) are roughly within projections given the estimates of GHG forcing and PETM climate sensitivity (PALEOSENS Project Members, 2012; Kiehl and Shields, 2013).

Subsequent but smaller hyperthermals represent additional climate sensitivity calibration points. The Eocene Thermal Maximum 2 (ETM-2; ca. 54 Ma) was characterized by surface- and deep-ocean warming (Lourens et al., 2005; Sluijs et al., 2009; Stap et al., 2009) that was approximately half that of the PETM. The magnitude of the accompanying carbon isotope excursion (CIE, −1.5‰ to −1.1‰ in benthic foraminifera; e.g., Stap et al., 2010a; Fig. DR1 in the GSA Data Repository1) and dissolution horizons (Lourens et al., 2005; Nicolo et al., 2007; Stap et al., 2009; Westerhold et al., 2011) indicates a global carbon cycle perturbation that is also approximately half that of the PETM, although this has yet to be adequately constrained by independent observations or modeling.

In addition to warming, the hyperthermals are also characterized by a mode shift, or intensification, of the hydrologic cycle. In the case of the PETM this includes widespread evidence of increased intensity and/or frequency of extreme precipitation events and drought in continental settings (Schmitz and Pujalte, 2007; Kraus and Riggins, 2007; Foreman et al., 2012), and changes in runoff and/or precipitation and evaporation (P-E) in marine settings (Zachos et al., 2003, 2006; Tripati and Elderfield, 2004; Nicolo et al., 2007; Tipple et al., 2011). In particular, open-ocean P-E, as inferred from changes in sea-surface salinity (ASSS), appears to have declined in the subtropics, and increased in high latitudes. Similarly, cursory evidence for intensification of the hydrologic cycle has been documented for ETM-2 in the Arctic and North America (Sluijs et al., 2009; Abels et al., 2012), but few other locations.

Previous attempts to quantify ETM-2 warming have been limited to a few planktic \( \delta^{18}O \) records that show modest (negative) anomalies (Lourens et al., 2005; Stap et al., 2010a), possibly due to the influence of competing environmental factors, such as higher SSS and acidification (e.g., Spero et al., 1997), both of which would be consistent with a global carbon cycle perturbation. Subtropical salinization would be expected due to increased subtropical evaporation and transport of \(^4\)O-depleted water vapor poleward on a global scale (e.g., Roberts et al., 2011; Tindall et al., 2010). To circumvent this, additional proxies such as Mg/Ca are required to constrain sea-surface temperature (SST), which, with adjustments for pH (Evans et al., 2016), could then allow for estimation of changes in local sea-surface \( \delta^{18}O_{\text{seawater}} \), and thus SSS (e.g., Zachos et al., 2003).

To assess whether the magnitude of ETM-2 warming was globally uniform in the absence of an ice-albedo feedback (e.g., Kiehl and Shields, 2013), and to test if regional P-E declined in the subtropics due to increased meridional vapor transport consistent with GHG warming (Carmichael et al., 2015), we applied coupled \( \delta^{18}O \) and Mg/Ca paleothermometry in planktic foraminifera from Ocean Drilling Program (ODP) Sites 1209 and 1265 to establish low- to mid-latitude SST and SSS anomalies.

METHODS
Specimens of Acarinina soldadoensis, a planktic foraminifera species that likely hosted photosymbionts and thus resided in the photic zone (e.g., Pearson et al., 1993), were collected from the 250–355 μm sieve size fraction at 3–5 k.y. resolution from ODP Sites 1209 and 1265, where ETM-2 is tightly constrained by carbon isotope and cycle stratigraphy (Lourens et al., 2005; Westerhold et al., 2007; Gibbs et al., 2012). Trace element ratios (e.g., Mg/Ca, Sr/Ca, Al/Ca, Ti/Ca) of oxidatively reductively cleaned specimens (10–20 shells) were measured by inductively coupled plasma–mass spectrometry following the methodology of Brown et al. (2011) with an interrun precision for Mg/Ca of <3% (2 standard deviation). Stable isotope analyses (\( \delta^{18}C \) and \( \delta^{18}O \), Vienna Pee Dee belemnite) were performed on shells from the same samples via isotopic ratio mass spectrometry with an interrun precision for \( \delta^{18}C \) and \( \delta^{18}O \) below 0.1% (2 relative standard deviation, RSD).

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1GSA Data Repository item 2018045, supplemental information including methodological details and figures that support the discussion, is available online at http://www.geosociety.org/datarepository/2018/ or on request from editing@geosociety.org

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and <0.16\% (2 RSD), respectively. A \%CaCO_3 record was also generated for ODP Site 1209 via coulometric analysis. In addition, we compiled previously published bulk CaCO_3 (% and δ^{13}C) and benthic foraminiferal δ^{18}O and δ^{18}H for Site 1265 (Stap et al., 2009, 2010a, 2010b) and bulk δ^{13}C and benthic foraminiferal δ^{18}O and δ^{18}H for Site 1209 (Gibbs et al., 2012; McCarren, 2009). Details of all analytical and numerical methods are provided in the Data Repository.

From the coupled planktic Mg/Ca and δ^{18}O data (Fig. 1), SST and SSS anomalies were computed (Fig. 2) following the method of Zachos et al. (2003), but accounting for additional recently discovered proxy sensitivities. Specifically, the exponential constant of the Mg/Ca proxy calibration (A value; see Equation 3 in the Data Repository) is potentially sensitive to Mg/Ca_{seawater} (Evans et al., 2016). In order to account for this sensitivity, we applied a range of constants (A values from 0.05 to 0.09) that encompass the error in Mg/Ca temperature regression fits for potential Mg/Ca_{seawater} values of the early Eocene (see the Data Repository; Evans et al., 2016) that contribute to the uncertainty in δSSST (i.e., ±1 °C). Furthermore, Evans et al. (2016) observed a pH effect on Mg/Ca that they quantified using a logistic fit to trends in field and laboratory data (Equation 1 in the Data Repository), although they could not statistically rule out a linear fit (Equation 2 in the Data Repository). Using a δ^{11}B-based estimate of ΔpH (Penman et al., 2014) for the PETM, Evans et al. (2016) applied a pH correction to planktic δ^{18}O and Mg/Ca data of Zachos et al. (2003) to highlight the significance of this effect. Because ΔpH has not yet been established for ETM-2, we apply a ΔpH of −0.05 pH units, estimated using Long-Term Ocean-Atmosphere-Sediment Carbon Cycle Reservoir (LOSCAR) cycle-coupled simulation of the event, as constrained by observations of the magnitude of the CIE, ASST, and changes in the carbonate compensation depth (CCD) (see the Data Repository). Applying both the logistic and linear pH adjustments, the pH effect on our Mg/Ca-δSSST is <−0.1 °C (Fig. DR3). Scaling the Site 1209 δ^{11}B-based PETM ΔpH (Penman et al., 2014) to ETM-2 using the planktic CIE recorded for each event (see the Data Repository) leads to a larger ΔpH (~0.11 pH units), although the added effect on our results is minimal (<0.1 °C decrease in δSSST and <0.2 ppt decrease in δSSS; Fig. DR3).

Planktic δ^{18}O values are also adjusted with LOSCAR-based ΔpH (~0.05 pH units) following the Globigerina bulloides relationship initially ascribed to a [CO_3] effect on δ^{18}O by Spero et al. (1997) (i.e., −2.51‰ per pH unit), before estimating and removing the Mg/Ca-based temperature influence to determine δ^{18}O_{seawater}. The residual shifts in δ^{18}O_{seawater} are then converted to δSSS using 0.25‰–0.50‰ per ppt from Zachos et al. (2003). This approach assumes (1) the complete absence of ice sheets to influence δ^{18}O_{seawater}, and (2) that the state of foraminifera preservation is uniform on short length scales, where lithology (i.e., \%CaCO_3) is relatively invariant (Fig. 1).

RESULTS

At both sites, Mg/Ca increases by ~25% during the CIE (Fig. 1). Given the long residence times of Mg and Ca in seawater (>1 m.y.),
Mg/Ca should have remained constant over the duration of the study interval (~200 k.y.); thus, the Mg/Ca anomalies must reflect ΔSSS. During the CIE of ETM-2, planktic δ18O values decrease by <0.5‰ during the CIE at South Atlantic ODP Site 1265, but at Pacific Site 1209 show little to no decrease beyond background variability. In contrast, benthic δ18O data show a similar 0.6‰–0.7‰ decrease at both locations (Fig. 1; Stap et al., 2010b), suggesting similar warming of the deep sea. Surface-ocean CIEs of ~1.5‰ are recorded in mixed-layer planktic foraminifera at both sites (Fig. 1). The %CaCO3 decreases by ~40% at Site 1265 (Stap et al., 2009) and by ~5% at Site 1209 (this study) during the CIE (Fig. 1).

**ΔSSS AND ΔSSS FROM THE SUBTROPICS DURING ETM-2**

We focus on establishing anomalies instead of absolute SST and SSS, because recrystallization of planktic foraminifera in carbonate-rich pelagic ooze can shift δ18O substantially (Pearson et al., 2001). Any overprint of δ18O, however, should be uniform over the short length scales studied here, and features such as anomalies (and cycles) are typically preserved (Kozdon et al., 2013). In contrast, Mg/Ca seems to be minimally affected by recrystallization in relatively closed systems such as the low-porosity pelagic muds at Sites 1209 and 1265 (Kozdon et al., 2013; Edgar et al., 2015). Dissolution, however, tends to decrease Mg/Ca and increase δ18O in foraminiferal calcite (Dekens et al., 2002). Combining the dissolution sensitivity of the modern foraminifera *G. ruber* (Dekens et al., 2002) with our simulated CCD response for ETM-2 (Fig. DR2) translates to a potential Mg/Ca decrease due to dissolution of 1%–3% and 2%–6% in the Pacific and Atlantic, respectively. In terms of SST, this would dampen ETM-2 warming by <0.5 °C in the Pacific, and <1 °C in the Atlantic. In contrast, δ18O dissolution sensitivity might suggest a potential increase in foraminiferal δ18O of <0.04‰ and <0.08‰ for the Pacific and Atlantic, respectively. This magnitude of δ18O enrichment would tend to amplify our calculated ΔSSS by only <0.25 ppt, for both sites. However, estimates of the effect of dissolution on our results are likely extremes, as the depth-dependent dissolution effect probably overcompensates for actual dissolution (e.g., Hönisch et al., 2013). Furthermore, the muted δ18O temperature signal is observed at both sites, which exhibit variable decreases in %CaCO3.

In general, Mg/Ca suggest a similar degree of warming in the equatorial Pacific and South Atlantic during ETM-2. The δ18O anomalies, in contrast, are small and unequal, indicating local salinity-related anomalies in δ18O seawater. Although the total rise in SST amounts to 2–4 °C during ETM-2 (Fig. 2) and residual shifts in δ18O seawater indicate an increase in SSS of +2 ± 1 ppt at Site 1209 and +1 ± 1 at 1265 (Fig. 2), suggesting increased subtropical aridity coincident with the warming.

**DISCUSSION**

The 2–4 °C warming in low to mid-latitudes established herein is similar in magnitude to bottom-water warming recorded by deep-sea benthic δ18O (~3 °C; Stap et al., 2010b), where negligible changes in pH and salinity are predicted, by inference, in areas of bottom-water formation in the Southern Ocean and the Arctic (3–5 °C ΔSST; Sluijs et al., 2009). The spatial pattern of warming for ETM-2 is similar (albeit smaller in magnitude) to the pattern recorded by Mg/Ca and GDGT (gycrole diakyl glycerol tetaethers) based temperature estimates during the PETM (e.g., Dunkley Jones et al., 2013) and expected pattern of warming in a largely ice-free world that lacks ice-albedo feedbacks (Kiehl and Shields, 2013).

Our finding of increased subtropical salinity during ETM-2 supports theoretical considerations of Eocene greenhouse warming effects on atmospheric vapor transport, ocean salinity, and δ18O seawater. Simulations with isocele-enabled climate models (Roberts et al., 2011; Tindall et al., 2010) suggest relatively enhanced evaporation in the subtropical and net transport of 18O-depleted moisture poleward, raising both the salinity and δ18O seawater of subropical surface waters. In one simulation involving the GISS (Goddard Institute for Space Studies) ModelE-R (Roberts et al., 2011), P-E decreases in the subtropical latitude bands from 15°S to 45°S, and from 10°N to 40°N, resulting in 0.5–2.0 ppt local increases in SSS, the largest increases occurring in the subtropical Pacific and the North Atlantic and the smallest in the equatorial Pacific. Similarly, the HadCM3 (Hadley Centre coupled atmosphere-ocean general circulation model) was used to simulate differences between preindustrial and Eocene δ18O seawater producing broadly similar results as the GISS ModelE-R, with some local differences in the Indian Ocean and North Atlantic Ocean basins (Tindall et al., 2010). It is interesting that near Site 1265, simulated changes in P-E appear to be small, and so the site would record a reduced δ18O seawater anomaly, consistent with our findings. While orbital forcing and ocean circulation might have contributed, the close covariation of SSS and SST with the CIE during the transient Eocene hyperthermals supports a primary feature of climate theory, that GHG warming drives increased meridional moisture transport, i.e., intensification of the hydrologic cycle (Carmichael et al., 2015).

Our estimate of surface salinization recorded during ETM-2 at Site 1209 is similar in magnitude to the original estimate of a 2–3 ppt increase for the PETM (Zachos et al., 2003). This original estimate was not adjusted for pH or for other effects on either δ18O and Mg/Ca. In order to compare the salinity anomalies for the PETM and ETM-2, we recalculated PETM ΔSSST and ΔSSS using planktic Mg/Ca and δ18O data from Zachos et al. (2003). Applying the same range in Mg/Ca temperature sensitivity (A values) as used for ETM-2 and adjusting for ΔpH using simulated LOSCAR Pacific surface PETM ΔpH (~0.17 units; Zeebe et al., 2009), we estimate a salinity anomaly of 3 ± 2 ppt for the PETM at Site 1209 (Fig. DR4). We used the LOSCAR PETM ΔpH over the δ18B-based PETM ΔpH to maintain consistency when comparing the SSS anomalies for the two events. The large uncertainty in the recalculated PETM ΔSSS reflects a broad range of possible Mg/Ca temperature sensitivities for the Eocene. When averages are compared, the PETM salinity anomaly at Site 1209 is slightly larger than the anomaly for ETM-2 (i.e., 3 ± 2 ppt for the PETM, and 2 ± 1 ppt for ETM-2; Fig. 2; Fig. DR4), as one would expect, due to the relatively larger perturbations in temperature and hydrological cycle during the PETM compared to ETM-2.

In summary, this study documents patterns of sea-surface warming and salinization of the subtropical (~28°N) Pacific and subtropical (~42°S) South Atlantic during ETM-2 that are consistent with theoretical predictions of the climatic response to GHG forcing, including globally uniform warming in the absence of ice-albedo feedback, a decrease in P-E, and a corresponding increase in SSS at low to mid-latitudes, suggesting enhanced meridional vapor transport.

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**REFERENCES CITED**


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