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Eocene-Oligocene southwest Pacific Ocean paleoceanography new insights from foraminifera chemistry (DSDP site 277, Campbell Plateau)

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Despite its major role in the Earth's climate regulation, the evolution of high-latitude ocean dynamics through geological time remains unclear. Around Antarctica, changes in the Southern Ocean (SO) circulation are inferred to be responsible for cooling from the late Eocene and glaciation in the early Oligocene. Here, we present a geochemical study of foraminifera from DSDP Site 277 (Campbell Plateau), to better constrain thermal and redox evolution of the high latitude southwest Pacific Ocean during this time interval. From 56 to 48 Ma, Mg/Ca- and $\delta^{18}\text{O}$ -paleothermometers indicate high surface and bottom water temperatures (24–26°C and 12–14°C, respectively), while weak negative Ce anomalies indicate poorly oxygenated bottom waters. This is followed by a cooling of ~4° between 48 and 42 Ma, possibly resulting from a weakening of a proto-EAC (East Australian Current) and concomitant strengthening of a proto-Ross gyre. This paleoceanographic change is associated with better ventilation at Site 277, recorded by an increasing negative Ce anomaly. Once this proto-Ross gyre was fully active, increasing biogenic sedimentation rates and decreasing *Subbotina* sp. $\delta^{13}\text{C}$ values indicate enhanced productivity. This resulted in a shoaling of the oxygen penetration in the sediment pile recorded by increasing the foraminiferal U/Ca ratio. The negative Ce anomaly sharply increased two times at ~35 and ~31 Ma, indicating enhanced seawater ventilation synchronously with the opening of the Tasmanian and Drake Passage gateways, respectively. The Oligocene glaciation is recorded by a major increase of bottom seawater $\delta^{18}\text{O}$ during the EOT (Eocene-Oligocene Transition) while Mg/Ca-temperatures remain rather constant. This indicates a significant ice control on the $\delta^{18}\text{O}$ record.

KEYWORDS

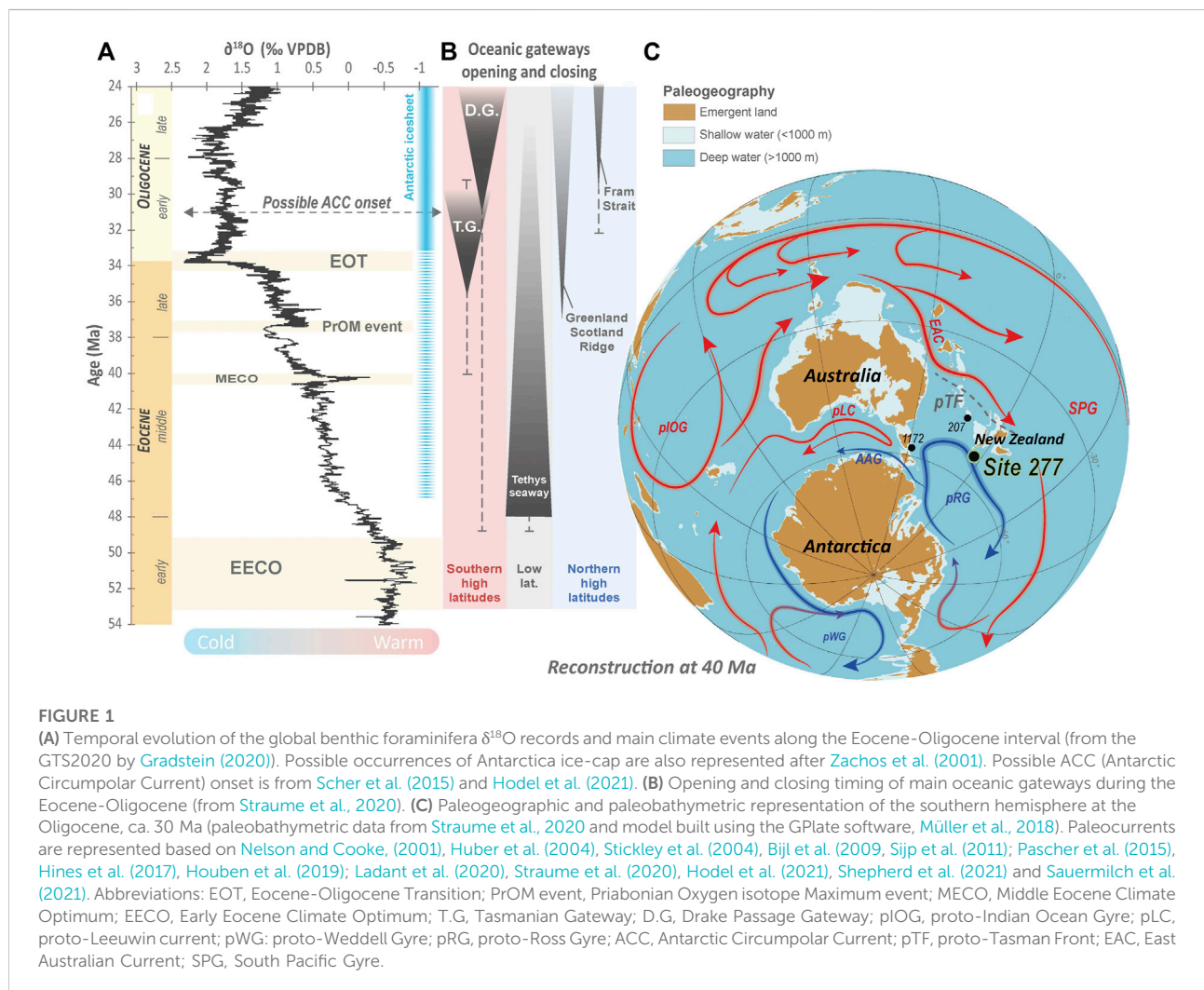
paleoceanography, climate, foraminifera, Eocene, Oligocene, southwest Pacific Ocean

1 Introduction

Facing ongoing global warming, understanding the feedbacks driving the Earth's climate is one of the greatest challenges of Earth sciences. As a pivotal period marking the transition from a greenhouse to the modern icehouse world, the Eocene-Oligocene interval has been studied for decades to identify and decipher the climate mechanisms driving this cooling (e.g., Zachos et al., 2001; Hutchinson et al., 2021) (Figure 1). Today, it is generally accepted that a reduction in atmospheric $p\text{CO}_2$ played a major role in this transition, resulting in the cooling and then glaciation (e.g., DeConto and Pollard, 2003; Pagani et al., 2011; Ladant et al., 2014; Kennedy-Asser et al., 2020; Hutchinson et al., 2021). However, ocean dynamics also play a key role in Earth's climate. Indeed, oceanic circulation controls global heat transfers while oceanic primary productivity and the associated biological pump is a major

atmospheric CO_2 sink (e.g., Jovane et al., 2009; Goldner et al., 2014; Elsworth et al., 2017; Toumoulin et al., 2020; Zhang et al., 2020; Sauermilch et al., 2021). Hence, constraining the evolution of oceanic circulation and the chemistry of different water masses during the Eocene-Oligocene transitions is fundamental for a better understanding of the Earth's climate changes.

Antarctica and the surrounding Southern Ocean have been a significant focus of the last decades given their high sensitivity to cooling and glaciation during the Eocene-Oligocene (e.g., Kennett et al., 1975; Nelson and Cooke, 2001; Zachos et al., 2001; Stickley et al., 2004; Bijl et al., 2009; , 2021; Bohaty et al., 2012; Hollis et al., 2012, 2015; Scher et al., 2014, 2015; Pascher et al., 2015; Hodel et al., 2021; Sauermilch et al., 2021; Shepherd et al., 2021). Located at the outlet of the Tasmanian gateway, the DSDP Site 277 on the Campbell Plateau (Figure 1) was drilled specifically to investigate the role played by the Antarctic Circumpolar Current (ACC) in the



thermal isolation of Antarctica from tropical heat fluxes (e.g., Kennett et al., 1975) (Figure 1). There is significant evidence that the development of the ACC played a major role in cooling and keeping Antarctica cool (e.g., Kennett, 1977; Kennett and Exon, 2004; Sauermilch et al., 2019, 2021). In this study, we investigate the role that other currents such as the proto-East Australian Current and the proto-Ross gyre (e.g., Nelson and Cooke, 2001; Bijl et al., 2009; Pascher et al., 2015; Sauermilch et al., 2021; Shepherd et al., 2021) played on the ocean heat fluxes, ocean redox state and primary productivity during the Eocene-Oligocene cooling.

Here, we investigate the physico-chemical properties of the southwest Pacific water masses recorded at Site 277 to better decipher the high latitude Southern Ocean dynamics during the Eocene-Oligocene cooling. Site 277 (52°13.43'S; 166°11.48'E) is located on the Campbell Plateau, South of New Zealand, between Auckland and Campbell Islands. The site was drilled to better understand the paleoceanographic history of the region related to long-term oceanic structural changes and Antarctic glaciation (Kennett et al., 1975). It is characterized by continuous sedimentation from the middle Paleocene to the late Oligocene (Kennett et al., 1975). The Eocene-Oligocene section of the Site 277, is composed of two main lithological units: Unit 2 (13–246 mbsf) and Unit 3 (246–454) mbsf. Unit 2 is a nannofossil ooze containing glauconite, radiolaria, foraminifera, minor amounts of diatoms and sponge spicules. Unit 3 only differs by the fact that siliceous fossils are poorly preserved and less abundant and by the occurrence of chert nodules (Hollis et al., 1997). Site 277 is thus characterized by a very uniform and mainly biogenic Eocene-Oligocene sedimentary record, with minor detrital materials (Kennett et al., 1975). It, therefore, constitutes an ideal target for paleoceanographic studies dealing with marine biocarbonates. The modern water depth at Site 277 is 1232 m below sea level. Its paleodepth during the Eocene-Oligocene interval is estimated to be 1,000–1,500 m, well above the lysocline at lower to middle bathyal water depths since the Paleocene (Hollis et al., 1997).

We applied several chemical proxies to planktic and benthic foraminifera from Site 277: (1) Mg/Ca and $\delta^{18}\text{O}$ ratios to infer seawater temperature and ice-volume evolution, (2) Ce anomaly (Ce/Ce*) as a seawater redox proxy, (3) U/Ca ratio as a sedimentary redox proxy, and (4) the $\delta^{13}\text{C}$ as a potential paleoproductivity proxy in combination with sedimentation rates (e.g., Katz et al., 2010 for a review, Chase et al., 2001; Boiteau et al., 2012; Chen et al., 2017; Remmelzwaal et al., 2019; Hodel et al., 2021). All these proxies were analyzed for both benthic (mixed *Gyroidinoides* sp. and *Cibicidoides* sp.) and planktic foraminifera (*Subbotina* sp.) from Site 277 to decipher physico-chemical features of bottom and surface water. These results provide new insights to better constrain southwest Pacific Ocean circulation, associated heat transfer, ocean and sedimentary redox evolution as well as paleoproductivity nearby Antarctica across the Eocene-Oligocene cooling.

2 Materials and methods

2.1 Foraminifera separation and cleaning

Based on the detailed core descriptions of Kennett et al. (1975), 27 sediment samples were selected for foraminifera analysis. Each sample was resuspended into Milli-Q water, ultrasonicated and sieved at 200 μm . Each fraction was dried for 48 h in an oven at 50°C. Then, planktonic (*Subbotina* sp., thermocline dwellers, e.g., Pearson et al., 2006; Hollis et al., 2012; Pascher et al., 2015) and benthic foraminifera (*Cibicidoides* sp. and *Gyroidinoides* sp.) were hand-picked separately under a stereo microscope. We only selected entire, clean and well-preserved specimens devoid of visible recrystallization, clay aggregates and oxyhydroxide patches (Figure 3). Each planktic and benthic foraminifera sample was then cleaned and digested in a clean laboratory following the procedure of Hodel et al. (2021). Foraminifera tests cleaning method consists of several ultrasonications in MQ water (Molina-Kescher et al., 2014; Osborne et al., 2017; Hodel et al., 2021) rather than the oxidizing-reducing cleaning of Barker et al. (2003). This allows the preservation of secondary coatings hosting the sedimentary redox signal (foraminiferal U/Ca ratio evolutions) that we investigate further in this contribution. For each sample, back-scattered electrons (BSE) images of foraminifera tests (and residues after chemical treatment) were obtained using a JEOL JSM-6360LV electron microscope operating at 20 kV at the *Observatoire Midi-Pyrénées* (PANGEE instrumental Platform, Toulouse, France).

2.2 Elemental chemistry

Major and trace element concentrations of each foraminiferal fraction were obtained through the all-in-one procedure developed in Hodel et al. (2021). Cleaned foraminifera fractions were dissolved into 1 ml of a 0.5 N acetic acid solution. Then, major and trace element concentrations were measured using a Thermo Scientific Element XR HR-ICP-MS at the *Observatoire Midi-Pyrénées* (Toulouse, France) (see Hodel et al., 2021 for more details). The oxide interferences (BaO+ on Nd+ and Eu+, LREE on MREE, and MREE on HREE, see table in [Supplementary Material S1](#)) were then corrected. Oxide production rate was determined based on the Cal-S standard material (Calcite, Yeghicheyan et al., 2003). The precision and accuracy of ICP-MS analyses were assessed by measuring the Cal-S standard. Element concentrations of all analyzed foraminiferal fractions are available in Supplementary data.

2.3 Oxygen and carbon isotope ratio analysis

Stable isotope analyses were performed at the ISTE^P Laboratory (Sorbonne Université, Paris, France) using a Kiel IV carbonate automatic device coupled with a DELTA V Advantage isotope ratio mass spectrometer. Foraminiferal oxygen and carbon isotope compositions were obtained by measuring the oxygen and carbon stable isotopes of carbon dioxide generated by the dissolution of planktonic and benthic foraminifera samples using anhydrous orthophosphoric acid at 70°C. Isotope values are reported in conventional delta (δ) notation as per mil (‰) deviations of the isotopic ratios ($^{18}\text{O}/^{16}\text{O}$) calculated to the VPDB scale. Accuracy and precision of 0.08‰ (1 σ) for both O and C isotopes were determined by repeated analyses of an internal Carrara marble standard, calibrated against the international standard NBS-19. Results of all the foraminiferal $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ analyses of this study are available in Supplementary data.

2.4 Paleotemperature calculation

2.4.1 Mg/Ca temperatures

Mg/Ca-derived paleotemperatures were calculated using the latest power law relationship established to describe foraminiferal Mg/Ca ratio and temperature covariations (Hollis et al., 2015; Hines et al., 2017 and reference therein, Eq. 1). The latter is an update of the exponential relationship of Lear et al. (2002) to better fit past seawater Mg/Ca ratios (Mg/Ca_{sw}).

$$T = \ln \left(\frac{[\text{Mg}/\text{Ca}_{\text{test}}] \times [\text{Mg}/\text{Ca}_{\text{sw}}^{t=0}]^H}{B \times [\text{Mg}/\text{Ca}_{\text{sw}}^{t=t}]^H} \right) \times \frac{1}{A}. \quad (1)$$

In this study, we used a value of 1.6 mol.mol⁻¹ for the Mg/Ca ratio of seawater at the time of test crystallization (i.e., Mg/Ca_{sw}^{t=t}; Stanley and Hardie, 1998; Evans and Müller, 2012; Hollis et al., 2015; Hines et al., 2017). Possible variations of this value along the studied interval are further discussed (see part 4.2.4). The Mg/Ca ratio of modern seawater (Mg/Ca_{sw}^{t=0}) has been fixed at 5.17 mol.mol⁻¹ (Stanley and Hardie, 1998; Evans and Müller, 2012). For surface seawater temperatures (SST), we used the general calibration of Anand et al. (2003), which is based on nine planktonic species (A = 0.09 and B = 0.38) and an H value of 0.29 (Creech et al., 2010; Hollis et al., 2012). For seafloor temperatures (SFTs), we used the calibration of Lear et al. (2002) based on three *Cibicidoides* species (A = 0.109 and B = 0.867) and an H value of 0.16 (Burgess et al., 2008). Errors on calculated temperatures were determined by considering the relative standard deviation (RSD) on Mg and Ca concentrations of 1.03% and 1.14% on average, respectively, as well as the calibration error

of $\pm 1.6^\circ\text{C}$ on the multi-species calibrations established by Lear et al. (2002) and Anand et al. (2003). The cumulative error calculated provides upper and lower uncertainties ($\pm 1.62^\circ\text{C}$ on average). All the paleotemperatures calculated based on Mg/Ca molar ratios are available in Supplementary data.

2.4.2 $\delta^{18}\text{O}$ temperatures

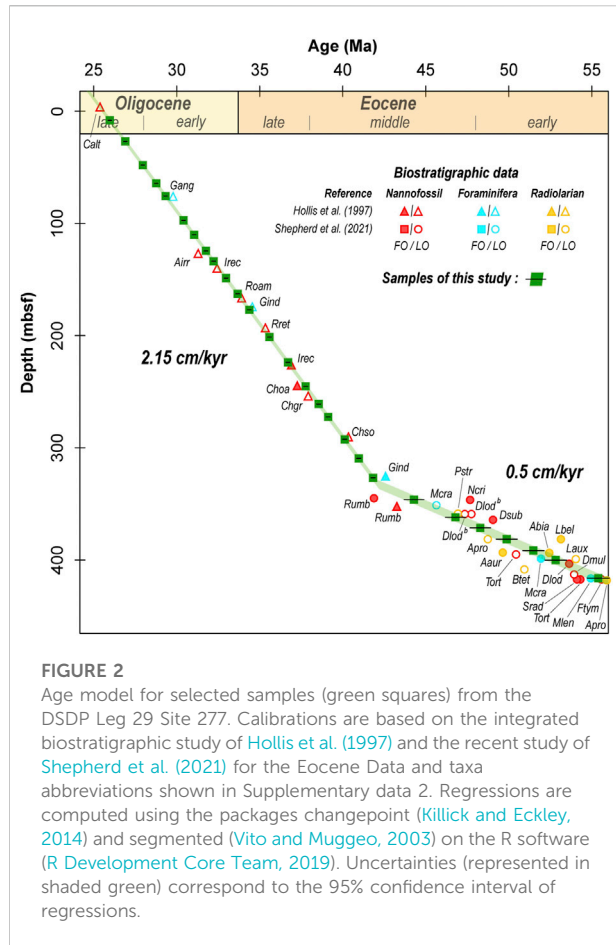
$\delta^{18}\text{O}$ -derived paleotemperatures were calculated using the equation of Kim and O'Neil (1997) for both benthic and planktonic foraminifera (Eq. 2):

$$T(^{\circ}\text{C}) = 16.1 + -4.64(\delta^{18}\text{O}_M - \delta^{18}\text{O}_{sw}) + 0.09(\delta^{18}\text{O}_M - \delta^{18}\text{O}_{sw})^2, \quad (2)$$

where $\delta^{18}\text{O}_M$ is the measured value and $\delta^{18}\text{O}_{sw}$ the seawater value. Here, $\delta^{18}\text{O}_{sw}$ is fixed to -1.23‰ , including an ice-volume component of -0.96‰ (Zachos et al., 1994) and a SMOW to VPDB correction of -0.27‰ (Kim and O'Neil, 1997). We fixed the ice-volume component because the evolution of global continental ice volume along the Eocene-Oligocene interval remains unconstrained. This likely induces a bias on calculated temperatures, notably across and after the Eocene-Oligocene Transition (EOT), which is debated further (see part 4.2.4.). Planktonic $\delta^{18}\text{O}$ values were corrected for paleolatitude (correction of -0.23‰ for ca. 65°S, Zachos et al., 1994). We also applied a correction of $+0.28\text{‰}$ for benthic species (Katz et al., 2003; Hollis et al., 2015). All the $\delta^{18}\text{O}$ -derived paleotemperatures are available in Supplementary data.

2.4 Age model and sedimentation rate calculations

Age estimations for each sample have been calculated using the biochronological data from Hollis et al. (1997) for Site 277 and updated with the recently published nannofossil data of Shepherd et al. (2021) for the Eocene interval. Data of Hollis et al. (1997) were recalibrated to the GTS 2012 (Gradstein et al., 2012) (see Supplementary Material S2). We used the *changeoint* (Killick and Eckley, 2014) and *segmented* (Vito and Muggeo, 2003) packages implemented on the R software (R Development Core Team, 2019) to identify statistically robust trends of sedimentation rate fitting with linear regressions and to calculate the absolute age of each selected sample (Figure 2) (see Hodel et al., 2021 for a similar approach). The 95% confidence interval of these regressions (*jack-knife* R script) defined uncertainties on obtained ages (from ± 0.10 to ± 0.6 Ma, Figure 2). The calculated ages with their associated uncertainties are reported in Supplementary Material S2.



3 Results

3.1 Sample ages

Calculated ages of our samples range from 55.4 ± 0.6 Ma to 26.0 ± 0.1 Ma. Our sampling frequency resolution is between 0.5 and 1.2 Ma from 26.0 to 41.8 Ma and between 1.3 and 2.6 Ma from 44.3 to 55.4 Ma. Our statistical approach allowed us to identify two main sedimentation regimes (in line with Hollis et al., 1997), namely a sedimentation rate of ~ 0.5 cm/kyr between 55.4 ± 0.6 Ma and 41.8 ± 0.1 Ma and of ~ 2.15 cm/kyr between 41.8 ± 0.1 Ma and 26.0 ± 0.1 Ma (Figure 2). Shepherd et al. (2021) recently mentioned two hiatuses at Site 277 in the early and middle Eocene that we do not observe with our statistical model. This likely relies on the fact that our linear regressions are based on several biostratigraphic data points. It results in a slightly less resolved but statistically more robust age model. However, it is worth noting that the calculated uncertainties of our ages take into account the poor distribution of the biostratigraphic markers for this time interval. Thus, the effect of these possible hiatuses on our ages is encompassed in the error bars.

3.2 Scanning electron microscopy imaging: Test preservation

Scanning electron microscopy (SEM) observations (Figure 3) have been used to assess the preservation state of benthic and planktonic foraminifera. As evidenced by previous studies (e.g., Hollis et al., 2015; Shepherd et al., 2021), planktonic foraminifera from Site 277 are characterized by a “frosty” texture (Sexton et al., 2006), resulting from minor authigenic calcite recrystallization after the deposition of the test on the seafloor and/or during their sinking through the water column (Figures 3A,B). Benthic foraminifera show a smoother texture without visible secondary calcite recrystallization. SEM observations of the residues after the foraminifera test acid dissolution (0.5 M CH_3COOH) indicate that only calcite has been dissolved, indicating minimal contamination by detrital material (e.g., clay mineral, quartz, mica and feldspar grains) or early-diagenetic mineral such as pyrite (Figures 3E,F).

3.3 Foraminiferal Rare Earth Element concentrations

All analyzed foraminifera show smooth seawater-like PAAS (Post Archean Australian Shales) normalized REE element patterns (Figures 4A,B) characterized by: 1) a depletion in LREE (Light Rare Earth Elements) compared to HREE (Heavy Rare Earth Elements, $(\text{Pr}/\text{Yb})_N$ ratios ranging from 0.27 to 0.57 and 0.29 to 0.60 for benthic and planktonic foraminifera, respectively), 2) a marked negative Ce anomaly ($\text{Ce}/\text{Ce}^* = \text{Ce}_N / (\text{Pr}_N \times (\text{Pr}_N/\text{Nd}_N))$, ranging from 0.31 to 0.90 and 0.29 to 0.92 for benthic and planktonic foraminifera, respectively) and 3) a pronounced positive La anomaly ($\text{La}/\text{La}^* = \text{La}_N / (\text{Pr}_N \times (\text{Pr}_N/\text{Nd}_N)^2)$ ranging from 1.58 to 2.48 and 1.60 to 2.23 for benthic and planktonic foraminifera, respectively). All these features are characteristic of modern seawater chemistry (e.g., Elderheld and Greaves, 1982; Elderfield, 1988; Grenier et al., 2013; Jeandel et al., 2013; Hathorne et al., 2014; Deng et al., 2017). Interestingly, temporal evolutions of the Ce/Ce^* values display clear stepwise tendencies (Figure 4C). From 55.4 Ma to 41.8 Ma, Ce/Ce^* values are high and fluctuate between 0.67 to 0.90 and 0.69 to 0.92 for benthic and planktonic foraminifera, respectively. From 41.8 Ma onward, Ce/Ce^* values decrease considerably in both benthic and planktonic species in a series of steps. First, Ce/Ce^* values decrease from 0.88 at 41.8 Ma to 0.62 at 39.1 Ma and stabilize *ca.* 0.62 from 39.1 to 35.6 Ma. A second marked drop occurs between 35.6 and 34.4 Ma down to *ca.* 0.47. Finally, a third major drop of the Ce/Ce^* values down to *ca.* 0.30 is noticeable between 31.8 and 31.0 Ma. Then the Ce anomaly remains constant until 26.0 Ma.

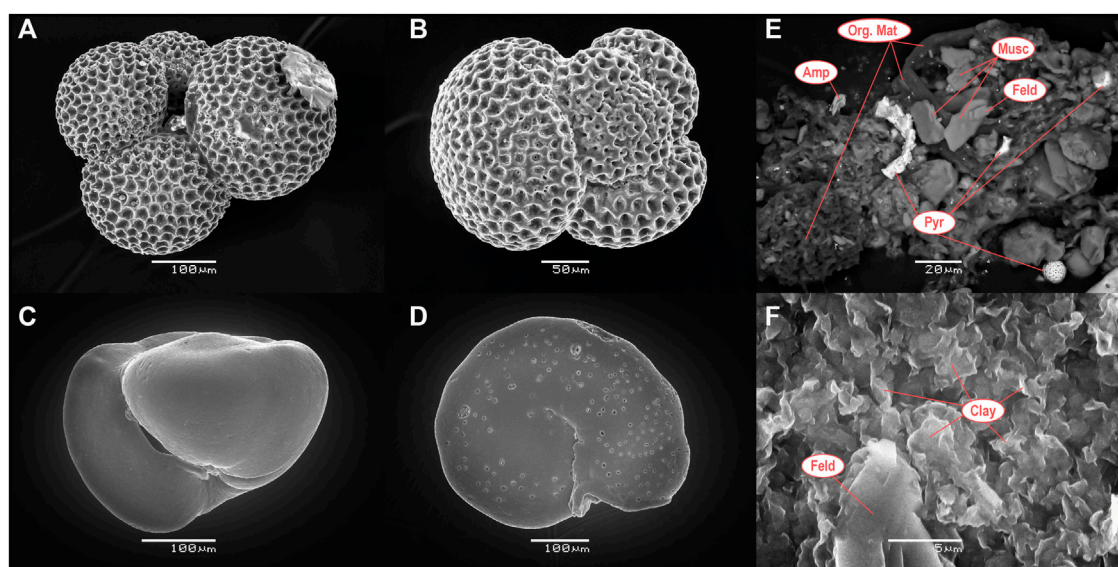


FIGURE 3

Back-scattered electron (BSE) SEM microphotographs of (A,B) planktonic foraminifera (*Subbotina* sp.) and (C,D) benthic foraminifera (*Gyroidinoides* sp. and *Cibicidoides* sp., respectively) from the DSDP Site 277. (E) and (F) are examples of residues remaining after 0.5 M CH_3COOH tests dissolution. Abbreviations stand for Pyr, pyrite; Org. Mat, organic matter; Musc, muscovite; Amp, amphibole; Feld, feldspar.

3.4 Foraminiferal major and traces element concentrations

3.4.1 Contamination evaluation

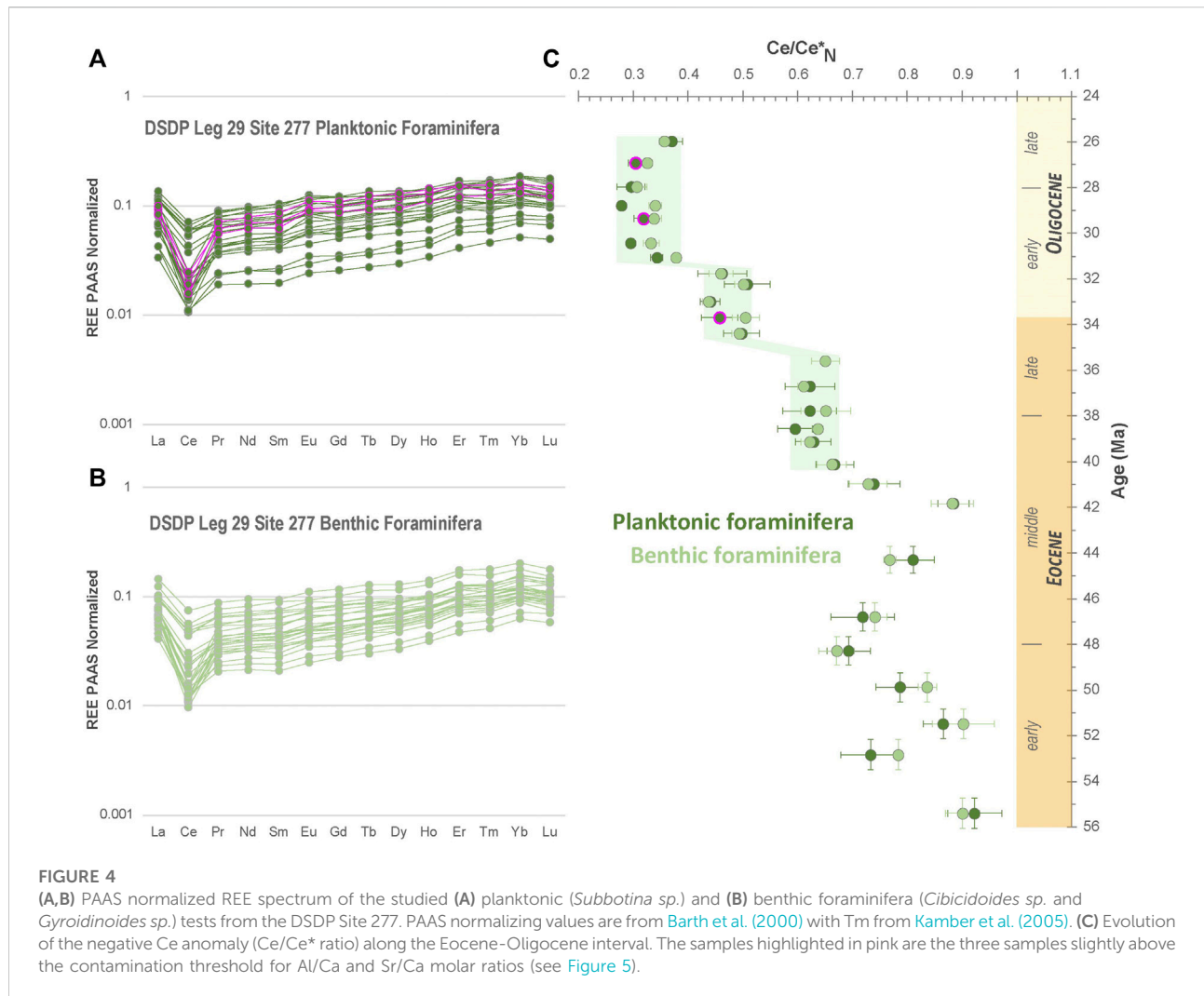
In addition to SEM observations and REE concentrations, Al/Ca vs. Mg/Ca and Sr/Ca vs. Mg/Ca diagrams can be used in chemical studies on foraminifera to monitor potential contaminations (e.g., Hollis et al., 2015; Osborne et al., 2017; Henehan et al., 2020; Hodel et al., 2021). In their laser ablation study, Hollis et al. (2015) calculated threshold values in such diagrams to discard samples with Mg/Ca-temperatures possibly biased by more than 1°C by detrital silicates, early-diagenetic (coatings) and/or recrystallization processes. In our study, all samples show low Al/Ca ratio with values ranging from 0.005 to 0.07 $\text{mmol}\cdot\text{mol}^{-1}$ and 0.008–0.166 $\text{mmol}\cdot\text{mol}^{-1}$ for benthic and planktonic foraminifera, respectively (Figure 5A). These values are well below the contamination threshold established by Hollis et al. (2015), namely Al/Ca > 2 $\text{mmol}\cdot\text{mol}^{-1}$ (Figure 5A). Only one sample (26.9 Ma) displays Al/Ca ratio above the threshold value of 0.140 fixed by Rae et al. (2011) and Henehan et al. (2020) for potential clay contamination (Figure 5A). With Sr/Ca ratio ranging from 0.92 to 1.38 and 0.90 to 1.67 for benthic and planktonic foraminifera, respectively, our data also lie within the range of 0.8–1.6 $\text{mmol}\cdot\text{mol}^{-1}$ established by Hollis et al. (2015) for acceptable values, except for two planktonic foraminifera samples showing ratios of 1.62 and 1.67. These two samples and the sample with Al/Ca ratio > 0.140 were discarded and not included in the discussion. They are circled in pink in the figures.

3.4.2 Mg/Ca ratio

Mg/Ca ratio ranges between 1.51 and 3.69 $\text{mmol}\cdot\text{mol}^{-1}$ and 1.58–3.06 $\text{mmol}\cdot\text{mol}^{-1}$ for benthic and planktonic foraminifera, respectively (Figure 6A). From 55.4 Ma to 46.8 Ma, both benthic and planktonic foraminifera show high Mg/Ca ratio, varying between 3.32 and 3.69 and 2.41 and 3.06, respectively. Mg/Ca ratio is rather stable during this time interval for benthic foraminifera, while it slightly decreases in the planktonic record. Then, the Mg/Ca ratio drops down to much lower values of 2.04 and 1.89 in the benthic and planktonic foraminiferal fractions, respectively at 44.3 Ma. From 44.3 Ma onwards, the Mg/Ca ratio of both benthic and planktonic foraminifera is more stable with values of 1.51–2.14 and 1.66 to 2.16 for benthic and planktonic foraminifera, respectively, except for some peaks in the benthic record at 41.0 Ma (Mg/Ca = 2.47), 39.1 Ma (Mg/Ca = 2.46) and 32.2 Ma (Mg/Ca = 2.44).

3.4.3 U/Ca ratio

Foraminiferal U/Ca ratio ranges from 0.007 to 0.078 $\mu\text{mol}\cdot\text{mol}^{-1}$ and 0.007–0.128 $\mu\text{mol}\cdot\text{mol}^{-1}$ for benthic and planktonic foraminifera, respectively (Figure 6C). U/Ca ratio is very low and stable from 55.4 to 41.8 Ma (from 0.007 to 0.020 $\mu\text{mol}\cdot\text{mol}^{-1}$ and 0.007–0.026 $\mu\text{mol}\cdot\text{mol}^{-1}$ for benthic and planktonic foraminifera, respectively). Then, it rapidly increases from 41.8 Ma onwards and reaches values of 0.058 and 0.068 $\mu\text{mol}\cdot\text{mol}^{-1}$ at 36.7 Ma. Further on, the U/Ca ratio decreases down to values of 0.017 and 0.020 $\mu\text{mol}\cdot\text{mol}^{-1}$ at



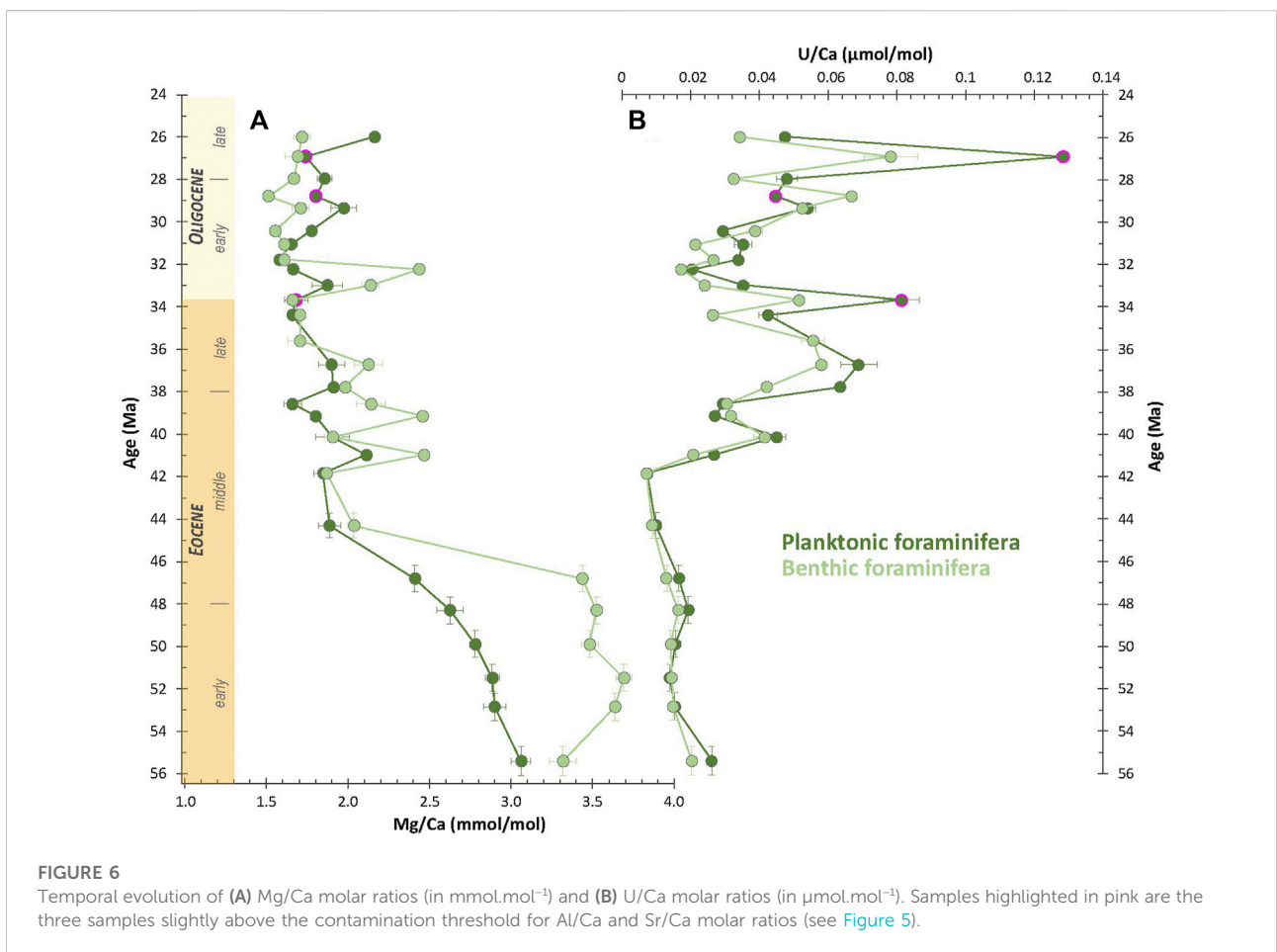
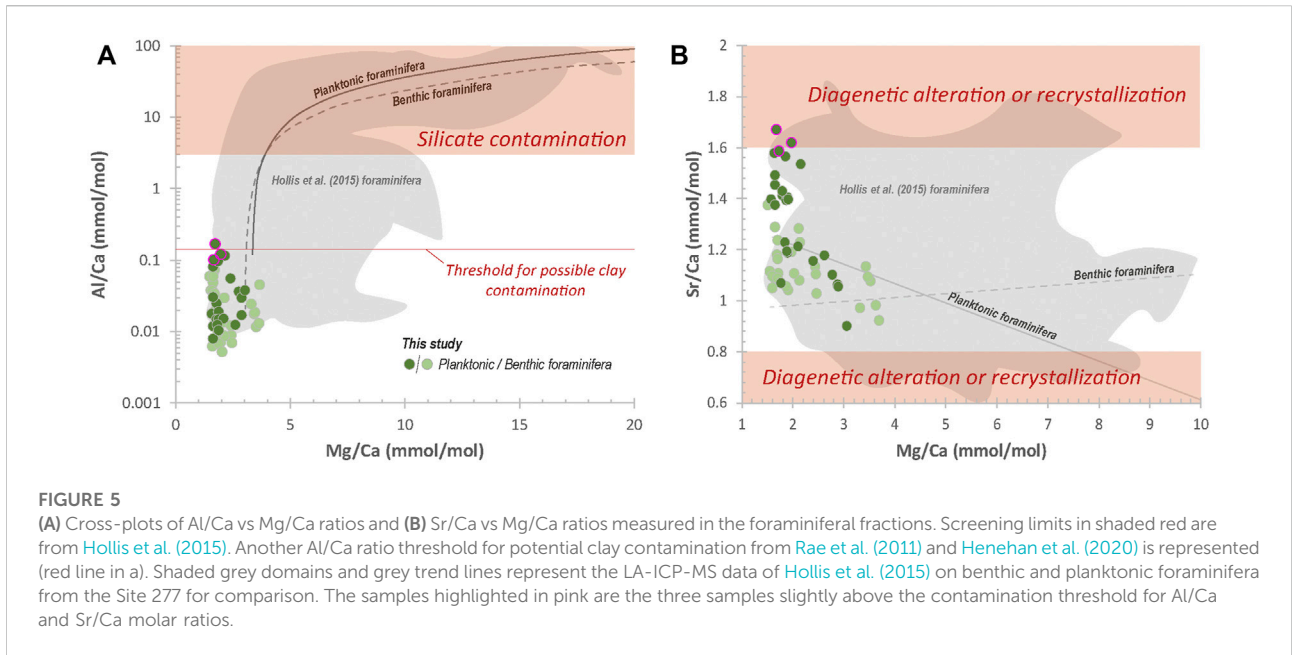
32.2 Ma in benthic and planktonic foraminifera, respectively. Finally, the U/Ca ratio increases again, more gradually, from 32.2 Ma to 26.0 Ma.

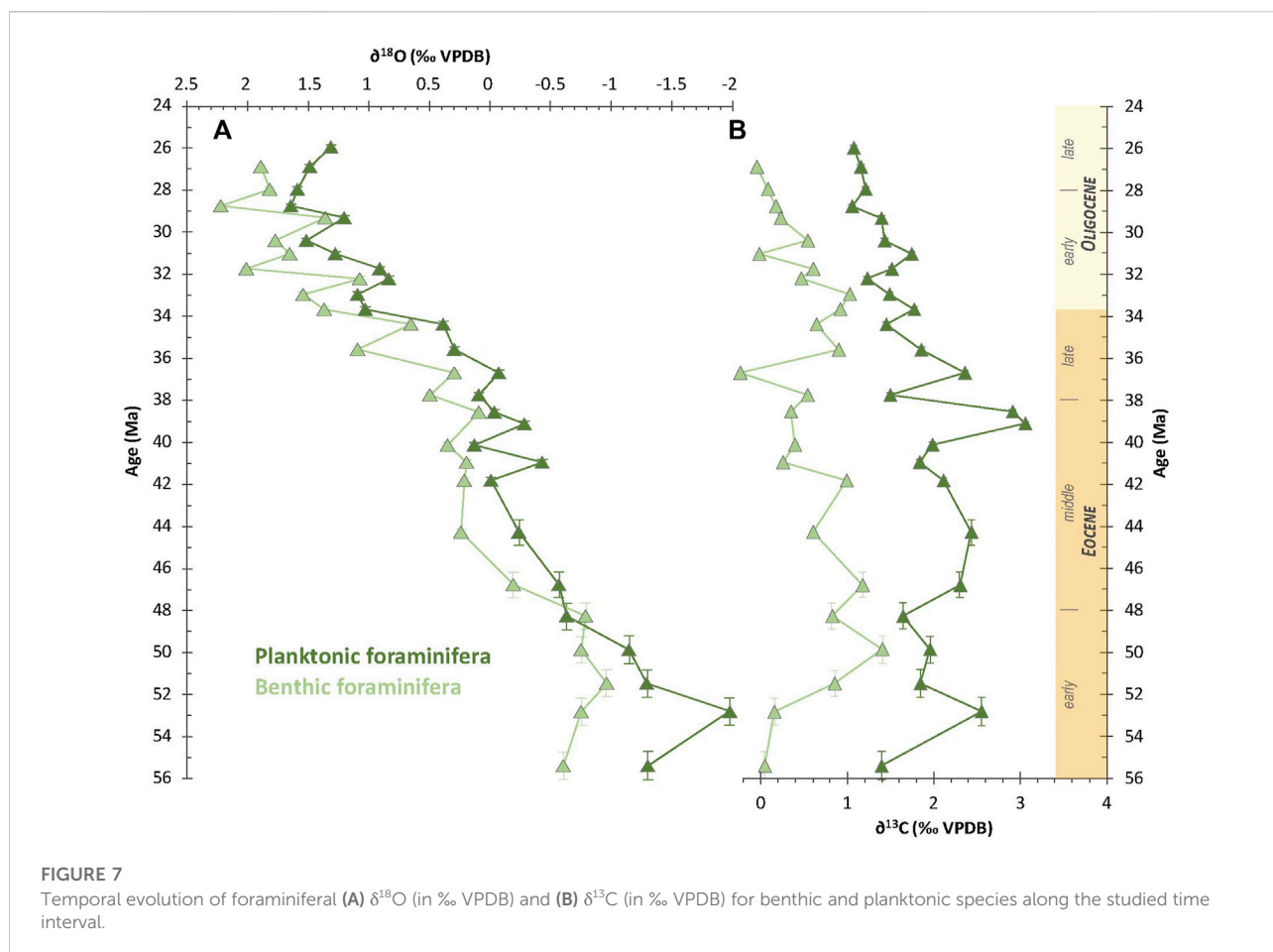
3.5 Oxygen and carbon isotope compositions

Foraminiferal $\delta^{18}\text{O}$ values range from -0.96 to $+2.22\text{‰}$ for benthic species and between -1.30 and $+1.64\text{‰}$ for planktonic species ([Figure 7A](#)). Overall, the benthic and planktonic $\delta^{18}\text{O}$ signals show a similar evolution, despite average values being lower by 0.5‰ for planktonic foraminifera. Values are both low during the early Eocene, ranging from -0.96 to -0.61‰ and -1.99 to -1.15‰ for benthic and planktonic foraminifera, respectively. From *ca.* 49 to 43 Ma, $\delta^{18}\text{O}$ increases up to -0.01‰ in planktonic foraminifera and $+0.23\text{‰}$ in benthic foraminifera, before stabilizing until *ca.* 36 Ma. From that time, the $\delta^{18}\text{O}$ of

benthic and planktonic foraminifera increases progressively up to $+1.89\text{‰}$ at 26.9 Ma and $+1.31\text{‰}$ at 26.0 Ma, respectively, although there are some second-order oscillations. Notably, $\delta^{18}\text{O}$ of both benthic and planktonic foraminifera increase from $+0.65$ to $+1.54\text{‰}$ and from $+0.39$ to $+1.09\text{‰}$, respectively between 34.4 Ma and 33.0 Ma, i.e. across the EOT.

Foraminiferal $\delta^{13}\text{C}$ values are between -0.24 and $+1.41\text{‰}$ and $+1.06$ to $+3.06\text{‰}$ for benthic and planktonic species respectively ([Figure 7B](#)). Benthic and planktonic signals do not show similar temporal trends. For benthic foraminifera, $\delta^{13}\text{C}$ increases from $+0.04$ to $+1.41\text{‰}$ during the 55.4 to 49.9 Ma interval and progressively decreases down to -0.24‰ at 36.7 Ma. Then it increases up to $+1.03\text{‰}$ at 33.0 Ma before decreasing down to -0.05‰ at 26.9 Ma. For planktonic foraminifera, $\delta^{13}\text{C}$ values range from $+1.39\text{‰}$ to $+2.54\text{‰}$ and do not show major evolution along the 55.4 to 40.1 Ma interval. Then planktonic foraminifera $\delta^{13}\text{C}$ values increase up to $+3.05\text{‰}$ at 39.1 and $+2.91\text{‰}$ at 38.5 Ma before decreasing down to $+1.07\text{‰}$ at 26.0 Ma.





4 Discussion

4.1 Pristine seawater chemical signal

SEM observations of planktonic foraminifera indicate minor recrystallizations as evidenced by “frosty” test texture (Sexton et al., 2006; Hollis et al., 2015; Shepherd et al., 2021) (Figure 3). Nonetheless, for both benthic and planktonic foraminifera, the LREE depletion compared to HREE as well as the marked Ce and La anomalies are typical of pristine seawater chemical signatures (e.g., Elderfield and Greaves, 1982; Elderfield, 1988; Bau and Dulski, 1996; Kamber and Webb, 2001; Bolhar et al., 2004; Deng et al., 2017; Hodel et al., 2021) (Figure 4). These features would undoubtedly have been deleted by early-diagenetic fluids and detrital contamination (e.g., Bolhar et al., 2004; Deng et al., 2017). This is in agreement with the Al/Ca and Sr/Ca ratios presented in Section 3.4.1, showing that diagenetic or recrystallization overprints can be excluded, except for three samples (Figure 5). Very low Al/Ca and Sr/Ca ratios in Al/Ca vs. Mg/Ca and Sr/Ca vs. Mg/Ca diagrams (Figure 5) also discard a significant effect of secondary coatings on Mg/

Ca ratios and therefore on Mg/Ca-derived temperatures. Indeed, all the samples that we further use in our discussion are below the calculated threshold values calculated by Hollis et al. (2015) to discard samples with Mg/Ca-temperatures possibly biased by more than 1°C by detrital silicates, early-diagenetic (coatings) and/or recrystallization processes. Hence, even if we did not use the conventional oxidizing-reducing cleaning of Barker et al. (2003), these chemical features attest of the validity of our Mg/Ca-derived temperatures. The latter is ultimately confirmed by the perfect consistency between 1) our Mg/Ca-SFT record and all the $\delta^{18}\text{O}$ -SFT records available for Site 277 (Hollis et al., 2012; Pascher et al., 2015; Shepherd et al., 2021; this study; Figure 8), and 2) our *Subbotina Sp.* Mg/Ca-derived thermocline temperatures and those of Hines et al. (2017) on the same site, whom applied a LA-ICP-MS method specially designed to remove the effects of potential post-depositional diagenetic contamination on Mg/Ca ratios (Figure 8). Therefore, we conclude that the elementary chemistry of the analysed benthic and planktonic foraminifera from Site 277 reflects pristine seawater composition for the investigated elements.

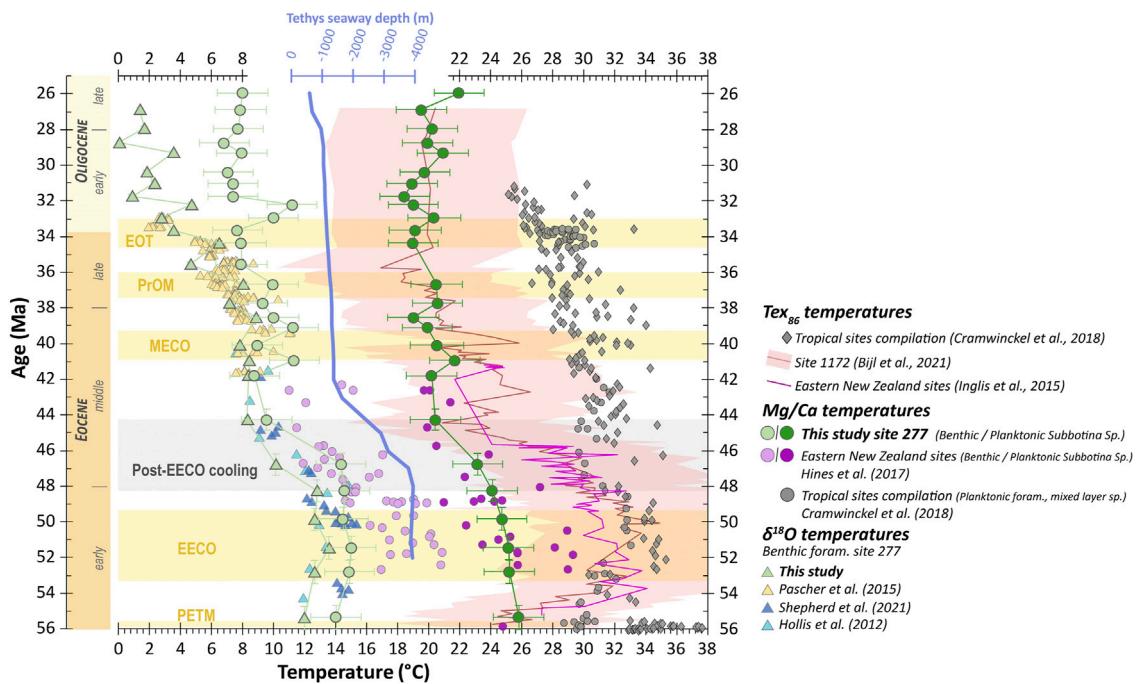


FIGURE 8

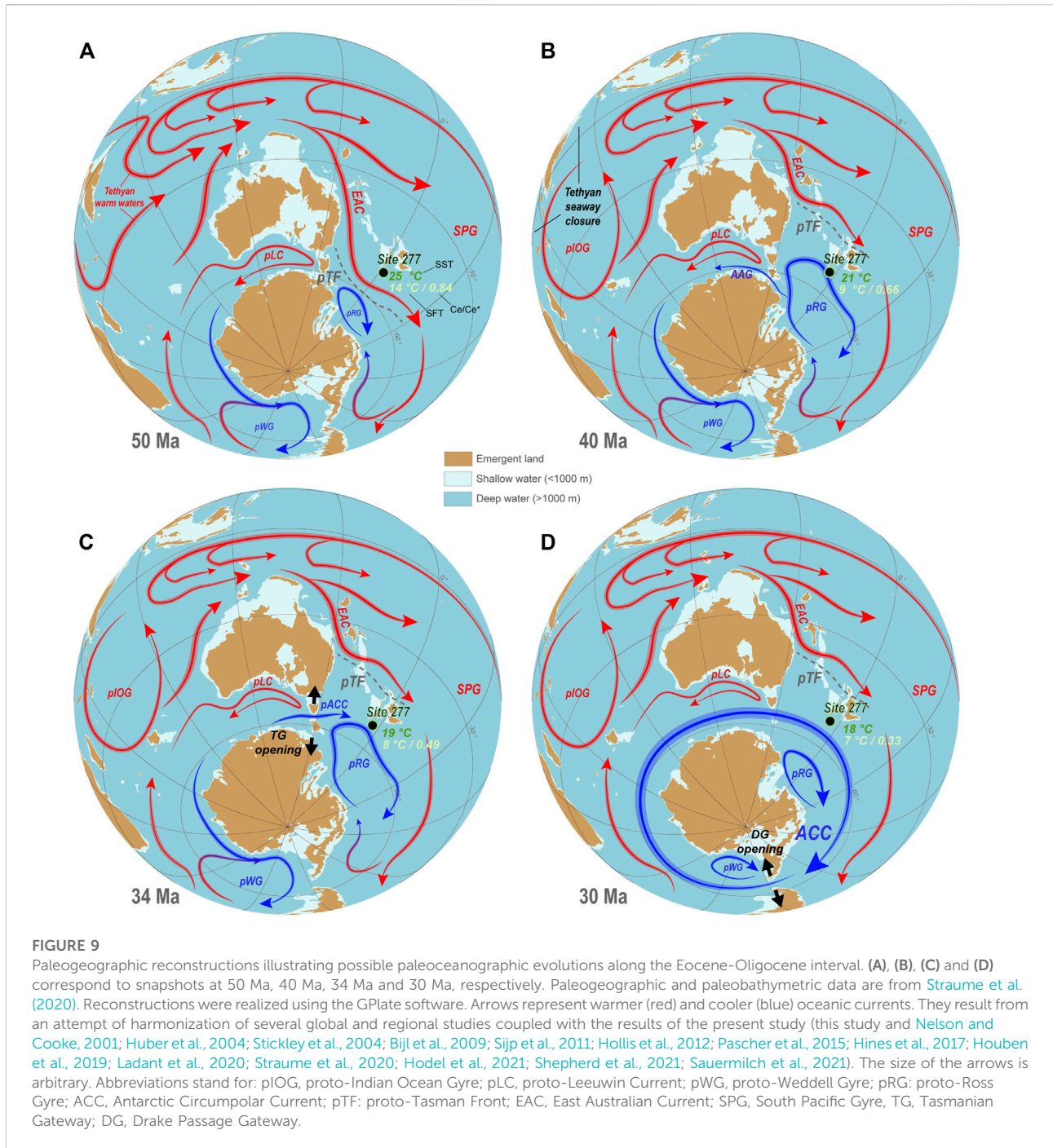
Temporal evolution of foraminiferal Mg/Ca- and $\delta^{18}\text{O}$ -derived SSTs (here thermocline temperatures obtained on *Subbotina sp.* foraminifera) and SFTs (seafloor temperatures obtained on *Cibicidoides sp.* and *Gyroidinoides sp.*) compared to available data from Hollis et al. (2012), Pascher et al. (2015), Hines et al. (2017), and Shepherd et al. (2021). TEX_{86} -derived SSTs (mixed-layer temperatures) for Site 1172 (Bijl et al., 2021) and Eastern New Zealand sites (Mid-Waipara and Hampden, Inglis et al., 2015) are also represented for comparison. Compilation of Mg/Ca- and TEX_{86} -derived SSTs (mixed-layer temperatures) from tropical sites worldwide by Cramwinckel et al. (2018) is also shown for comparison. It is worth noting that the post-EEOC cooling that we and other authors noticed in the southwest Pacific Ocean is not visible in that compilation. Tethys seaway depth evolution is from Straume et al. (2020).

Nevertheless, we cannot exclude potential effects of secondary processes on the O isotope compositions (e.g., Katz et al., 2010; Hollis et al., 2015). Indeed, $\delta^{18}\text{O}$ of foraminiferal calcite and derived temperatures are known to be more easily biased by such processes. *Post-mortem* recrystallization of planktonic tests once deposited on the seafloor can induce a cool bias on $\delta^{18}\text{O}$ -derived sea-surface temperatures (SSTs) by partially recording the bottom ocean temperature (Katz et al., 2010; Hollis et al., 2015). This bias is reflected by the relatively close $\delta^{18}\text{O}$ values of benthic and planktonic species throughout the studied interval, suggesting a temperature offset of only $\sim 2^\circ\text{C}$ between bottom and surface waters, much lower than the one estimated from Mg/Ca ratios ($\sim 12^\circ\text{C}$). For this reason, we focus our study on Mg/Ca-derived SSTs, which appear to be less affected by *post-mortem* recrystallization (Katz et al., 2010; Hollis et al., 2015) than $\delta^{18}\text{O}$ -derived SSTs. Finally, it is worth noting that the planktonic *Subbotina sp.* foraminifera picked for this study are thought to be thermocline-dwelling species (e.g., Pearson et al., 2006; Hollis et al., 2012; Pascher et al., 2015). Thus, calculated SSTs likely correspond to an integration of the thermocline paleotemperatures.

4.2 Southwest Pacific seawater temperatures

4.2.1 Early Eocene Climate Optimum

With a maximum temperature of 26°C in the early Eocene and a minimum temperature of 18°C in the late Oligocene, our Mg/Ca-derived SSTs are in line with TEX_{86} temperatures calculated and compiled by Bijl et al. (2009, 2021) for ODP Site 1,172 (Tasmania) located at an equivalent paleolatitude of $\sim 60^\circ\text{S}$ (see Figure 1 for Site 1172 location) (Figure 8). These high SSTs (from 23 to 26°C in the thermocline) are consistent with the multi-proxy SSTs of 26 – 28°C calculated by Hollis et al. (2012) for the Early Eocene Climate Optimum (EECO, Figure 1A) in the Canterbury Basin (New Zealand), as well as with numerous geochemical studies indicating warm subtropical-cool tropical conditions in the southwest Pacific Ocean in the early Eocene (Bijl et al., 2009; Hollis et al., 2009, 2015; Liu et al., 2009; Creech et al., 2010; Sluijs et al., 2011; Hines et al., 2017; Crouch et al., 2020) (Figure 8). Despite these numerous studies displaying consistently high temperatures, such high SSTs at high latitude (above 50°S) are still debated. This notably hinges on the fact that “proxy-based paleoclimate reconstructions from the



southwest and tropical Pacific imply little to no latitudinal temperature gradient during the early-Eocene, which is difficult to reconcile with the known climate dynamics and model studies” (in [Hines et al., 2017](#), see also [Cramwinckel et al., 2018](#) and [Hollis et al., 2019](#)). Recently, [Shepherd et al. \(2021\)](#) indicated that nannofossil assemblages from DSDP sites 207 and 277 (see [Figure 1](#) for sites locations) more likely attest to warm temperate rather than tropical conditions. Several

hypotheses have been proposed to explain this thermal overestimation, notably calibration issues, seasonal bias, and physiological processes ([Sluijs et al., 2006](#); [Hollis et al., 2012](#); [Taylor et al., 2013](#); [Hollis et al., 2015](#); [Inglis et al., 2015](#)). However, as raised by [Hollis et al. \(2015\)](#), given the consistency between the various proxies used in these studies ($\delta^{18}\text{O}$, Mg/Ca, and TEX_{86}), such overestimation must rely on common factor(s) affecting all proxies like a warm-season bias rather than calibration issues or

physiological factors (e.g., [Sluijs et al., 2011](#); [Hollis et al., 2012, 2015](#)). Benthic Mg/Ca and $\delta^{18}\text{O}$ data also testify to high and stable seafloor temperatures (SFTs) during the early Eocene, ranging from 14 to 15°C and 12–14°C, respectively, between 56 Ma and 48 Ma. These data are also consistent with those from [Hollis et al. \(2012\)](#) and [Shepherd et al. \(2021\)](#) showing similar SFTs for the Canterbury Basin and Site 277 ([Figure 8A](#)).

To explain such high seasonal SSTs and high SFTs during the early Eocene, [Hines et al. \(2017\)](#) proposed that an intensification of a proto-East Australian Current (EAC) during the PETM may have resulted in the distribution of tropical temperatures down into the high-latitude southwest Pacific Ocean. Similarly, based on the [Nelson and Cooke \(2001\)](#) study, [Shepherd et al. \(2021\)](#) proposed that New Zealand and the Tasman Sea were likely bathed by warm waters originating from the western limb of the South Pacific anticyclonic gyre and the southward extension of a proto-EAC ([Figure 9A](#)).

4.2.2 Middle Eocene cooling

Following the EECO, our Mg/Ca-derived SSTs show a progressive cooling during the early-mid-Eocene, which intensifies from 48.3 Ma to 44.3 Ma with SSTs decreasing from 24°C down to 20°C ([Figure 8A](#)). Synchronously, both Mg/Ca- and $\delta^{18}\text{O}$ -derived SFTs also attest to a marked cooling of *ca.* 4°C following the EECO ([Figure 8A](#)). In line with the emergence of cool-water nannofossil taxa at Site 277 from 49 Ma ([Shepherd et al., 2021](#)), this cooling likely marks the transition toward temperate conditions in the southwest Pacific Ocean (see also [Hollis et al., 2012, 2021](#)).

A first hypothesis to explain this southwest Pacific Ocean cooling could be a global climate event due to decreasing atmospheric $p\text{CO}_2$ (e.g., [Zachos et al., 2008](#)). However, other local mechanisms have been proposed given the apparent lack of (sub-) equatorial cooling at that time (e.g., [Bijl et al., 2013](#); [Cramwinckel et al., 2018](#); [Shepherd et al., 2021](#); [Figure 8](#)). Indeed, a comparison of SSTs between southwest Pacific and tropical sites shows that all southwest Pacific sites are affected by a significant cooling following the EECO, whereas no comparable drop is observed for tropical sites ([Figure 8](#) and references therein). To explain this, [Bijl et al. \(2013\)](#) proposed that local cooling could have been caused by a westbound flowing current across the Tasmanian Gateway producing a “cooling of Antarctic surface waters and coasts, which was conveyed to global intermediate waters through invigorated deep convection in southern high latitudes.” More recently, [Shepherd et al. \(2021\)](#) proposed that a migration of the proto-Ross gyre and the proto-Tasman front (cooler waters, e.g., [Nelson and Cooke, 2001](#); [Huber et al., 2004](#)) toward lower latitudes could also explain this post-EECO cooling.

As previously mentioned, the source of warm waters bathing the Campbell Plateau before this cooling was likely a proto-EAC fed by warm tropical-equatorial waters originating from both the tropical Indian Ocean and a South Pacific anticyclonic gyre (e.g.,

[Nelson and Cooke, 2001](#); [Hines et al., 2017](#); [Shepherd et al., 2021](#)) ([Figure 9A](#)). Therefore, we hypothesize that the northward migration proto-Ross gyre proposed by [Shepherd et al. \(2021\)](#) may have resulted from a retreat of this proto-EAC. Indeed, the Tasman gateway was not significantly opened at that time ([Stickley et al., 2004](#)) and the proto-EAC was thus likely to be the main control on the northward expansion of the proto-Ross gyre. Based on recent studies, we propose two mechanisms to explain such a proto-EAC weakening:

- 1) Paleomagnetic studies suggest a rifting between Southern Pacific and Indian oceans between 50 and 40 Ma (e.g., [Van Hinsbergen et al., 2015](#)). Meanwhile, these studies (see also [Bijl et al., 2021](#)) show a southward drift of the Pacific sector of Antarctica over this time interval, relative to the position of the Earth's spin axis. This possibly has relocated the proto-EAC, reduced its strength, and leads to an effective northward expansion of the proto-Ross gyre.
- 2) An alternate hypothesis is linked to the Tethys seaway. In the early Eocene, the Tethys seaway played a key role in global ocean circulation by driving an eastward TCC (Tethys Circum global Current), which provided warm saline water masses to the tropical Indian Ocean (e.g., [Jovane et al., 2009](#); [Ladant et al., 2020](#); [Straume et al., 2020](#)). [Hamon et al. \(2013\)](#) also modeled that the “final closure of the eastern Tethys seaway played a major role in the oceanic circulation reorganization during the middle Miocene”, notably by inducing “strong changes in the latitudinal density gradient and ultimately the reinforcement of the Antarctic Circumpolar Current”. This confirms the important influence of the Tethyan inputs on the Southern Ocean paleoceanography. Recently, [Straume et al. \(2020\)](#) suggested that a major step in the Tethys seaway closure occurred from 48 Ma to 42 Ma (depth loss of *ca.* 3,000 m), synchronously with the possible proto-EAC retreat and associated post-EECO cooling ([Figure 8](#)). Thus, we hypothesize that this Tethys seaway shoaling from 48 Ma to 42 Ma, possibly coupled with the partial obstruction of the Tethys-tropical Indian Ocean connection by the northward migration of the Indian plate (e.g., [Straume et al., 2020](#)), could also have resulted in a significant reduction of the warm eastward tropical-equatorial fluxes feeding the proto-EAC at that time ([Figure 9B](#)).

Further studies, notably modeling studies, are needed to validate or invalidate these hypotheses.

4.2.3 Middle to late Eocene

Following the post-EECO cooling, SSTs remained stable from 44 Ma to 26 Ma (20°C on average with a standard deviation of $\pm 1.8^\circ\text{C}$) ([Figure 8](#)), without noticeable shift at the Eocene-Oligocene Transition (33.7 Ma). This trend is

consistent with the TEX_{86} -derived SSTs of Bijl et al. (2021). In contrast, SFTs show more fluctuations during this time interval. After 44.3 Ma onward, SFTs calculated from both proxies remain stable until 36.7 Ma except for two isolated temperature rises of *ca.* $+2^{\circ}\text{C}$ noticeable in the Mg/Ca record at 41.0 Ma and 39.1 Ma. Then both proxies record a SFT drop of more than 3°C for $\delta^{18}\text{O}$ data and *ca.* 2°C for Mg/Ca data between 36.7 Ma and 35.6 Ma. Our sampling resolution is not sufficient to study in detail these potential short-lived events, which are not the purpose of this study.

4.2.4 Eocene-Oligocene transition: Major cooling, massive ice cap build-up, or both?

At the time of the EOT glaciation (from 34.4 Ma to 33.0 Ma), $\delta^{18}\text{O}$ -derived SFTs drop from 6°C to 3°C . Then $\delta^{18}\text{O}$ -derived SFTs progressively decrease during the Oligocene (with two positive peaks at 32.2 Ma and 29.3 Ma) down to a SFT of 1.37°C at 26.9 Ma. Interestingly, Mg/Ca-derived SFTs and SSTs do not follow this cooling trend and remain rather stable from 35.6 Ma to 26.0 Ma (Figure 8).

A possible explanation is that local phenomena could have influenced seawater temperature at Site 277 across the EOT. Indeed, this absence of cooling at the EOT in our Mg/Ca record is also noticeable in the TEX_{86} record of Bijl et al. (2021) from ODP Site 1172 (Figure 8). This could be due to a throughflow of the warm proto-Leeuwin current at the same time as EOT cooling as suggested by Houben et al. (2019) and/or to a subtropical front developing at these sites as the Tasmanian gateway opens (Sauermilch et al., 2021).

However, the dichotomy between ice-independent (Mg/Ca and TEX_{86}) and ice-dependent ($\delta^{18}\text{O}$) records remains puzzling. As mentioned above, southwest Pacific Mg/Ca and TEX_{86} -derived temperatures remain constant across the EOT, but $\delta^{18}\text{O}$ -derived temperatures drop (Figure 8). Several other sites show the same ambiguity worldwide: for instance, DSDP Site 511 at the Falkland Plateau (Houben et al., 2019), ODP sites 689 and 748 at Maud Rise and the Kerguelen Plateau (Bohaty et al., 2012). Lear et al. (2004) even suggest warming of *ca.* 2°C in the deep South Atlantic (DSDP Site 522) and Equatorial Pacific (ODP Site 1,218) oceans following the EOT. Therefore, either Mg/Ca and TEX_{86} temperatures are biased similarly or $\delta^{18}\text{O}$ temperatures are biased.

Based on the fact that the foraminiferal $\delta^{18}\text{O}$ is ice-dependent while the Mg/Ca is not, authors firstly concluded that this apparent absence of cooling in the benthic foraminifera Mg/Ca record implied that most of the $\delta^{18}\text{O}$ increase was due to an ice volume increase, modifying the seawater $\delta^{18}\text{O}$, rather than a major cooling (e.g., Lear et al., 2000; Billups and Schrag, 2003). Then, it has been shown that these results imply ice volume that cannot be accommodated by Antarctica alone (e.g., Coxall et al., 2005; DeConto et al., 2008) and therefore suggest a massive Arctic ice-cap formation. Based on a compilation of U_{37}^k and TEX_{86} -derived SSTs at low and high latitudes, Liu et al. (2009)

proposed that the EOT was accompanied by a heterogeneous cooling and modeled “that Northern Hemisphere glaciation was not required to accommodate the magnitude of continental ice growth during this time”. Today the timing of the Northern Hemisphere ice-cap onset is a subject of controversy (see Hutchinson et al., 2021 for a review). Several studies have found evidence of massive Arctic ice-cap formation (at least ephemeral) as early as the Eocene (e.g., Tripati et al., 2005, 2008; Moran et al., 2006; Eldrett et al., 2007; Stickley et al., 2009; Darby, 2014; Bernard et al., 2016; Tripati and Darby, 2018) while others argue for a Miocene (e.g., DeConto et al., 2008) or even a Pliocene (e.g., Zachos et al., 2001) Arctic glaciation. Given this controversy, several authors looked for alternative hypotheses to explain the lack of foraminiferal Mg/Ca-derived temperature cooling following the EOT. These hypotheses are the following:

- 1) Based on the sensitivity of benthic foraminiferal Mg/Ca to seawater saturation state (Dekens et al., 2002; Elderfield et al., 2006; Rosenthal et al., 2006; Yu and Elderfield, 2008), it has been proposed that the benthic foraminifera Mg/Ca signal might have been affected by an increase in bottom seawater saturation state associated with a deepening of the calcite compensation depth (CCD) (Lear et al., 2004, 2008, 2010; Coxall et al., 2005; Pusz et al., 2011; Bohaty et al., 2012).
- 2) Another possibility is that other forcings may have modified the Mg and Ca distribution in seawater and thus modified the Mg/Ca_{seawater} ratio itself (e.g., Bohaty et al., 2012). If so, our temperature calculation from foraminiferal Mg/Ca using a unique Mg/Ca seawater value for the whole Eocene-Oligocene ($\text{Mg/Ca}_{\text{sw}}^{\text{EOT}} = 1.6 \text{ mol.mol}^{-1}$ in Eq. 1, Hollis et al., 2015; Hines et al., 2017) is biased.

However, several observations call these hypotheses into question. Indeed, a possible role of carbonate saturation state change (CCD deepening) at Site 277 is improbable given: 1) its shallow depth (i.e. 1000-1500 m, well above the lysocline at lower to middle bathyal water depths since the Paleocene, Hollis et al., 1997); and 2) the Mg/Ca temperature data show that neither SFT nor SST cooled. Concerning a sudden Mg/Ca_{seawater} ratio change, our calculations of Mg/Ca-derived temperatures considering an unrealistic switch from an Eocene Mg/Ca_{seawater} ratio (1.6 mol.mol^{-1} , Hollis et al., 2015; Hines et al., 2017) to a modern Mg/Ca_{seawater} ratio ($5.17 \text{ mol.mol}^{-1}$, Stanley and Hardie, 1998; Evans and Müller, 2012) at the EOT does not result in a good agreement between Mg/Ca and $\delta^{18}\text{O}$ temperatures (see Supplementary Figure S1). Finally, as previously mentioned, the TEX_{86} -derived temperatures of Bijl et al. (2021) at Site 1172, which are not dependent on seawater carbonate nor to Mg chemistry, are also constant across the EOT (see Figure 8).

Hence, these findings tend to confirm that the $\delta^{18}\text{O}$ record has been affected by something other than seawater cooling across the EOT (e.g., Cramer et al., 2011). This is probably an

important ice control supported by Antarctica alone (e.g. Zachos et al., 2001; DeConto et al., 2008; Liu et al., 2009), or possibly by a bipolar glaciation as suggested by recent studies (Darby, 2014; Bernard et al., 2016; Tripathi and Darby, 2018). We do not have the key to resolve this enigma. However, we believe that it constitutes a critical obstacle in the understanding of this major climate transition and that further studies are needed. In particular, the depicted changes in seawater temperatures obtained from foraminiferal $\delta^{18}\text{O}$ values will remain biased as long as the evolution of the ice-volume component present in the $\delta^{18}\text{O}$ fractionation equation (see Eq. 2) is not further constrained.

4.3 Redox state of the southwest Pacific Ocean

The negative Ce anomaly ($\text{Ce}/\text{Ce}^* < 1$) is a common chemical feature of present-day seawater (e.g., Elderheld and Greaves, 1982; Elderfield, 1988; Grenier et al., 2013; Hathorne et al., 2014; Deng et al., 2017). This relies on the two valence states of Ce (III and IV) while other REE (apart Eu) are only trivalent. In modern oxic seawater, the oxidation of Ce^{3+} ions to Ce^{4+} ions leads to the precipitation of CeO_2 , hence removing Ce from seawater leading to a negative Ce anomaly in the REE pattern. Logically, the more oxygenated the seawater is, the more Ce is removed and the stronger the negative Ce anomaly is. Therefore, the amplitude of the Ce anomaly, quantified by the Ce/Ce^* ratio, constitutes a powerful proxy to reconstruct the past seawater redox state (e.g., Bau and Dulski, 1996; Deng et al., 2017). Because foraminifera accurately record the REE chemistry of the seawater in which they thrive in their calcitic test, the temporal evolution of the Ce/Ce^* values in well-preserved foraminifera allows to finely document the past seawater redox state (e.g., Kamber et al., 2014; Osborne et al., 2017; Remmelzwaal et al., 2019; Hodel et al., 2021). At Site 277, Ce/Ce^* values are similar for benthic and planktonic foraminifera throughout the studied interval. This is due to the fact that REE are at least partly incorporated in authigenic Fe-Mn coatings once on the seafloor and therefore benthic and planktonic foraminifera both record bottom water REE composition (e.g., Roberts et al., 2010; Molina-Kescher et al., 2014; Osborne et al., 2017).

4.3.1 Weakly oxygenated ocean along the early to middle Eocene at site 277

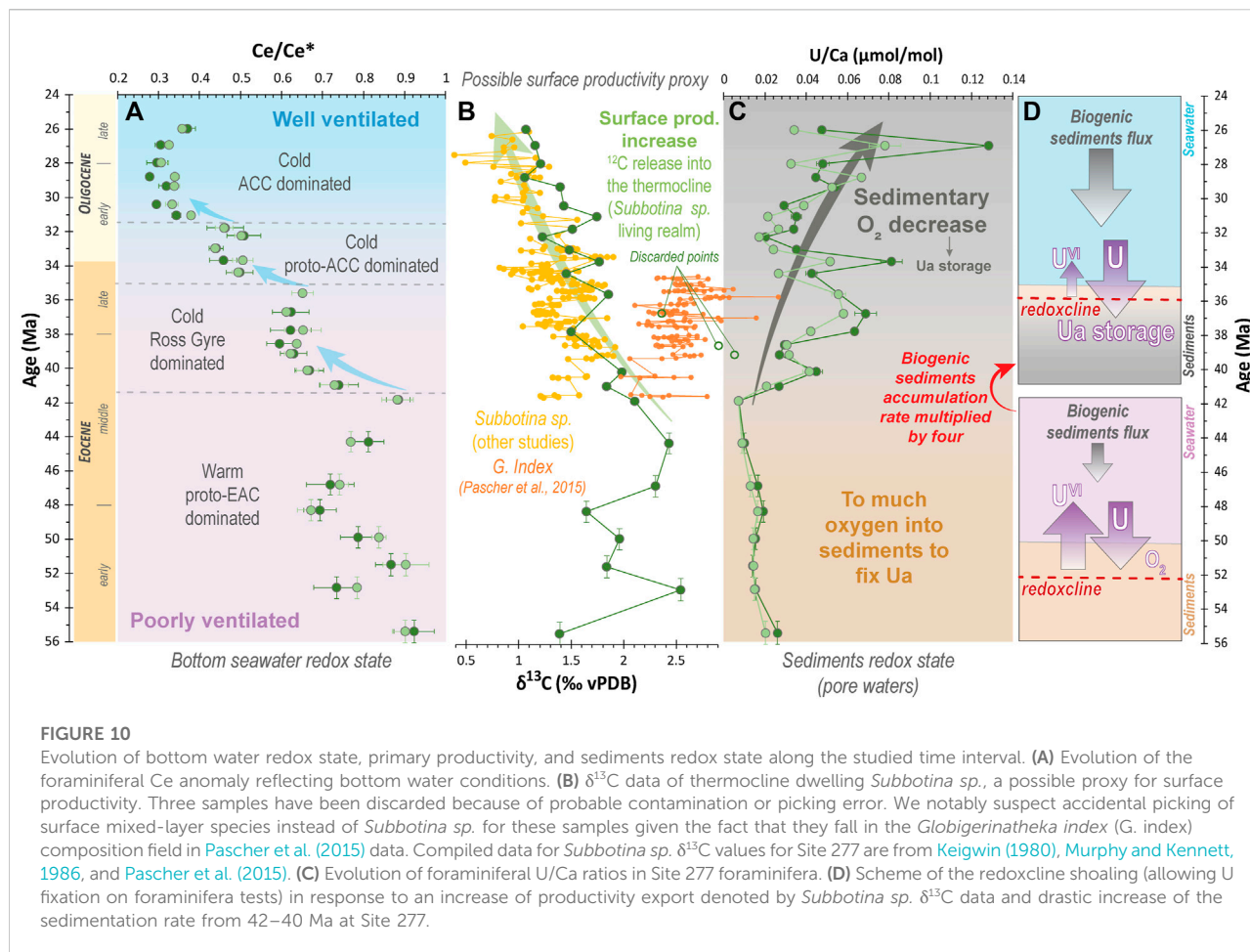
In the early to middle Eocene, Ce/Ce^* values are variable but remain high ($\text{Ce}/\text{Ce}^* > 0.67$) until 41.8 Ma (Figure 10), which indicates hypoxic bottom water conditions at the Campbell Plateau at that time (e.g., Remmelzwaal et al., 2019). Since this hypoxia is associated with very high SFTs and SSTs (above 8°C and 22°C , respectively, Figure 8), we propose that these low-oxygen conditions could be related to this particular warm climate. Indeed, previous studies of oceans' redox state evolution argued that the main warming events of the Earth's

history were often accompanied by open ocean hypoxic and shelf domain anoxic/euxinic episodes (e.g., Monteiro et al., 2012; Pälike et al., 2014; Chang et al., 2018; Remmelzwaal et al., 2019; Cornaggia et al., 2020; Clarkson et al., 2021). Among them, the PETM (*ca.* 56 Ma) and the ETM2 (*ca.* 54 Ma) are the last hyperthermal events for which widespread ocean deoxygenation has been proposed (e.g., Dickson and Cohen, 2012; Pälike et al., 2014; Chang et al., 2018; Yao et al., 2018; Remmelzwaal et al., 2019; Harper et al., 2020; Clarkson et al., 2021). Recently, Remmelzwaal et al. (2019) showed that temperature was likely the main driver of ocean deoxygenation during the PETM. Here, with similar temperatures (Figure 8), we suggest that the warm climate of the early to middle Eocene could have resulted in a poorly oxygenated ocean too. If so, the Campbell Plateau could have been particularly impacted by ocean deoxygenation given its rather shallow (*ca.* 1,000–1,500 mbsl) and isolated setting (e.g., Hollis et al., 1997).

From 41.8 Ma onwards, the Ce/Ce^* values decrease to *ca.* 0.6 and stabilize between 40.1 Ma and 35.6 Ma. This indicates progressive seawater oxygenation that further stabilized (Figure 10A). This possibly constitutes a response to the cooling of the surface and bottom waters between 48.3 and 44.3 Ma at Site 277 (retreat of the warm proto-EAC) and enhanced ventilation (vertical mixing through upwelling) at Site 277 once a proto-Ross gyre dominated system was established (Figure 8, Figure 9A,B, Figure 10A).

4.3.2 Enhanced ventilation after the Tasmanian and Drake Passage gateways opening

After the first oxygenation step, foraminiferal Ce/Ce^* values drop a second time between 35.6 Ma and 34.4 Ma (Figure 4B, Figure 10A) indicating a further oxygenation enhancement of the Campbell Plateau bottom waters. This drop is synchronous with the deep opening of the Tasmanian gateway from 35.5 Ma by Stickley et al. (2004). Finally, a last drop of the foraminiferal Ce/Ce^* values occurs between 31.8 Ma and 31.0 Ma. This third oxygenation step led to well-oxygenated bottom waters at the Campbell Plateau during the late Oligocene ($\text{Ce}/\text{Ce}^* < 0.4$) (Figure 9D, Figure 10A). Interestingly, as for the previous step, this ventilation enhancement is synchronous with a major gateway opening, namely the Drake Passage gateway, which likely opened to deep flow from *ca.* 31 Ma allowing the ACC onset (e.g., Scher et al., 2015; Hodel et al., 2021 and references therein). Hence, these results 1) tend to validate the timing of the Tasman and Drake Passage gateways deep openings *ca.* 35 Ma and *ca.* 31 Ma, respectively, and 2) testify that the opening of these gateways had major consequences on the Southern Ocean ventilation and water masses redox state Hodel et al. (2021) (Figure 9). Further work is needed with a higher spatial and temporal resolution to better constrain the involved mechanisms. In particular, it would be interesting to measure the evolution of the foraminiferal Ce anomaly for



several other drill sites in the southwest Pacific and southern Indian Ocean and compare it to that of Site 277 that we provide in this study.

4.4 Paleoproductivity and sedimentary redox state

Based on the fact that the U/Ca in cultured planktonic foraminifera decreases with increasing seawater $[\text{CO}_3^{2-}]$, the foraminiferal U/Ca ratio has been first considered as a promising proxy for seawater carbonate chemistry (e.g., Russell et al., 1996, 2004; Katz et al., 2010; Keul et al., 2013). However, Boiteau et al. (2012) pointed out that the U contents in downcore foraminifera were often much higher than the U incorporation threshold into biogenic calcite (lattice-bound U/Ca < 15 nmol.mol⁻¹). As a result, these authors suggested that most of the U measured in downcore foraminifera was hosted by reduced sedimentary coating (U⁴⁺ fixation, while U⁶⁺ remains soluble in oxygenated environments). From that point, they proposed that the foraminiferal U/Ca ratio was

directly related to authigenic U in sedimentary piles, providing “a simple means of evaluating redox changes in sediments where bulk sediment U data is unavailable or unreliable” (in Boiteau et al., 2012). Also, Boiteau et al. (2012), and later Chen et al. (2017), identified a clear correlation between glacial cycles and foraminiferal U/Ca in sediments of the last 400 kyrs, mirrored by high U/Ca ratios during glacial maxima and low U/Ca ratios during interglacial. They attributed this obvious correlation to either a “shoaling of the oxygen penetration depth (i.e. sedimentary redoxcline) due to increased export production or to reduced bottom water oxygen concentrations” (Boiteau et al., 2012). These processes both result in reducing conditions favoring authigenic U precipitation or adsorption onto the foraminifera tests during glacial intervals. Chase et al. (2001) also identified such glacial-interglacial U/Ca variations in foraminifera from a well-ventilated Sub-Antarctic setting during the last glacial maximum. Therefore, they argued that this tendency (high U/Ca ratios during glacial maxima and low U/Ca ratios during interglacials) more likely results from increased glacial productivity in the Sub-Antarctic region rather than from bottom water redox change.

Similarly, the temporal trends of the foraminiferal Ce anomaly, which recorded bottom seawater redox changes, attest to enhanced ventilation of the water masses at Site 277 from 41.8 Ma onwards (Figure 10A). Thus, the increase in foraminiferal U/Ca that we observe is unlikely to be related to a lower bottom seawater oxygenation and more likely relies on productivity/sedimentary processes (e.g., Chase et al., 2001; Boiteau et al., 2012). Thus, we hypothesize that a productivity enhancement and export may have reduced the oxygen penetration depth into the sediment, promoting more reductive conditions conducive to U fixation on foraminifera tests (e.g., Chase et al., 2001; Boiteau et al., 2012) (Figure 10D). This hypothesis is strengthened by 1) the sedimentation rate evolution, which is multiplied by four from 41.8 Ma without a major change in biogenic vs detrital contributions (the sedimentation remained mainly biogenic, Kennett et al., 1975; Hollis et al., 1997) (Figure 2) and 2) the *Subbotina* sp. $\delta^{13}\text{C}$ values, which start decreasing from that time (Figure 10B). Indeed, metabolic processes preferentially incorporate ^{12}C , thus phytoplankton blooms deplete the shallowest surface seawater in ^{12}C and enrich deeper surface water (where *Subbotina* sp. live) in ^{12}C after organic matter mineralization (e.g., Katz et al., 2010). Therefore, even if $\delta^{13}\text{C}$ is not only a productivity proxy and has to be interpreted carefully, *Subbotina* sp. $\delta^{13}\text{C}$ evolves as we would expect if there was a major phytoplanktonic bloom. This strengthens the idea of enhanced primary productivity from 42 Ma. If so, the rather shallow depth of Site 277, which remained well above the lysocline along the Eocene-Oligocene interval, is compatible with the fact that the organic matter did not have time to fully mineralize during its sinking through the water column, therefore allowing its deposition and burial. Then, the decomposition of this remaining organic matter into the sediment likely promoted reducing conditions and a shoaling of the sedimentary redoxcline, allowing the fixation of U on both benthic and planktonic foraminifera tests at Site 277 (Figure 10D). Increasing productivity has been already noticed in the Southern Ocean during the Eocene-Oligocene in response to seawater cooling and circulation intensification (e.g., Diester-Haass and Zahn, 1996; Plancq et al., 2014). Similarly, we can reasonably hypothesize that at Site 277, cooler seawater and more vigorous ocean circulation may have promoted an enhanced vertical mixing and nutrient supply through upwelling (Diester-Haass and Zahn, 1996; Miller et al., 2009; Plancq et al., 2014) once a proto-Ross gyre dominated system was established from ca. 42 Ma, then following the Tasmanian Gateway opening from ca. 35 Ma and finally with the establishment of the ACC following the Drake Passage Gateway opening from ca. 31 Ma.

5 Conclusion

Our integrated geochemical study of Site 277 foraminifera brings new insights into the paleoceanographic evolution of the southwest Pacific during the major climate transition of the Eocene-Oligocene.

Both Mg/Ca- and $\delta^{18}\text{O}$ -derived temperatures attest to very high SSTs (from ca. 24°C–26°C in the thermocline) and SFTs (from ca. 12°C to 14°C) at Site 277 during the EECO (from 55.4 Ma to 48.1 Ma). Such high temperatures are in line with other studies relating subtropical conditions in this area. This indicates that despite its high latitude location, the Campbell Plateau was bathed by warm waters, possibly originating from a proto-EAC fed by warm tropical-equatorial waters originating from both a South Pacific anticyclonic gyre and the tropical Indian Ocean (Figure 9A). Weak foraminiferal Ce anomaly (Ce/Ce* values close to 1) along this warm early to middle Eocene (from 55.4 Ma to 41.8 Ma) attests to poorly oxygenated bottom seawater at Site 277 at that time. Between 48.3 and 44.3 Ma, both Mg/Ca and $\delta^{18}\text{O}$ data recorded a marked bottom and surface cooling of ca. 4°C. Given the absence of an equivalent cooling at higher latitudes, we hypothesize that a proto-EAC weakening and a Ross-gyre enhancement pushed the proto-Tasman front (cooler waters) towards lower latitudes (Figure 9B). Then, bottom and surface temperatures remained rather constant until the EOT. Interestingly, our benthic Mg/Ca- and $\delta^{18}\text{O}$ -signals are significantly different from the EOT: $\delta^{18}\text{O}$ -temperatures decrease while Mg/Ca-temperatures remain constant. This dichotomy suggests that the hypothesis of a dominant ice control on the global EOT $\delta^{18}\text{O}$ decrease has to be reconsidered.

Major changes in redox and paleoproductivity took place from 41.8 Ma onwards, i.e., with the onset of a proto-Ross gyre dominated system (Figure 9B). We notably identified a major increase in productivity export from that time, which likely promoted more reductive sedimentary conditions. From 41.8 Ma as well, a drop of foraminiferal Ce/Ce* values attests to enhanced ventilation of bottom waters after the establishment of a proto-Ross gyre dominated system. Ce/Ce* values then drop a second time between 35.6 Ma and 34.4 Ma and a third time between 31.8 Ma and 31.0 Ma, synchronously with the Tasmanian and Drake Passage gateways opening. These results tend to confirm the timing of these major gateways opening and attest that their opening had major consequences on the Southern Ocean ventilation and water masses' redox state.

Thereby, altogether these results shed light on the role that major oceanic gateway closings and openings play on the Earth's climate by driving ocean currents and associated heat fluxes, water masses redox state and primary productivity fluxes.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

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A. Marquet on the ICP-MS and L. Emmanuel for O and C isotope analyses.

Conflict of interest

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Supplementary material

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/feart.2022.998237/full#supplementary-material>

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