



Changes in the Red Sea overturning circulation during Marine Isotope Stage 3

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Abstract. The oceanography of the Red Sea is controlled by the restricted exchange of water masses with the Indian Ocean and by high evaporation rates due to the arid climate of the surrounding land areas. In the northern Red Sea, the formation of oxygen-rich subsurface water ventilates the deeper parts of the basin, but little is known about the variability in this process in the past. The stable oxygen and carbon isotope records of epibenthic foraminifera from a sediment core of the central Red Sea and comparison with existing isotope records allow for the reconstruction of changes in the Red Sea overturning circulation (ROC) during Marine Isotope Stage 3. The isotope records imply millennial-scale variations in the ROC, in phase with the climate variability in the high northern latitudes. This suggests an immediate response of dense-water formation to the regional climate and hydrology of the northern Red Sea. Deep-water formation was intensified under the influence of cold and hyper-arid conditions during Heinrich stadials and was diminished during Dansgaard–Oeschger interstadials. While these changes are reflected in both stable oxygen and carbon isotope records, the latter data also exhibit changes in phase with the African–Indian monsoon system. The decoupling of the stable carbon and oxygen isotope records at the summer monsoon maximum centered around 55–60 ka may be associated with an increased inflow of nutrient-rich intermediate waters from the Arabian Sea to the central Red Sea. This process fueled local surface water productivity, resulting in enhanced remineralization of sinking organic matter and release of ¹²C at intermediate water depths.

1 Introduction

The landlocked basin of the Red Sea is bordered by the semi-arid to arid regions of the Arabian Peninsula and northern Africa (Fig. 1). The oceanography of the Red Sea is characterized by the restricted exchange with the Indian Ocean through the narrow and only 137 m deep Hanish Sill of the Bab al-Mandab Strait (Smeed, 1997, 2004). The exchange processes exhibit distinct seasonal contrasts, which are closely linked to the monsoonal wind system. During the NE monsoon in winter, a two-layer system prevails, with inflowing surface waters from the Gulf of Aden and the deeper water mass outflow of Red Sea water (RSW). During the summer SW monsoon, a three-layer system develops, with outflowing surface waters and diminished RSW outflow, but the intrusion of nutrient-rich Gulf of Aden Intermediate Water (GAIW) (Smeed, 1997, 2004). Enhanced evaporation rates result in high sea surface salinities (SSS), with maximum values above 40 psu in the northern Red Sea. As a result, warm (21.6–21.8 °C) and saline (40.5–40.6 psu) intermediate and deep-water masses form in winter as a result of the cooling of highly saline surface waters in the gulfs of Suez and Aqaba and occasionally in the northernmost open Red Sea (Cember, 1988; Eshel et al., 1994; Woelk and Quadfasel, 1996; Papadopoulos et al., 2015) (Fig. 2). As part of the Red Sea overturning circulation (ROC), the newly formed subsurface water masses extend southward and ventilate the deep-sea ecosystems, with estimated residence times of 40 to 90 years (Fig. 3) (Woelk and Quadfasel, 1996). Finally, RSW

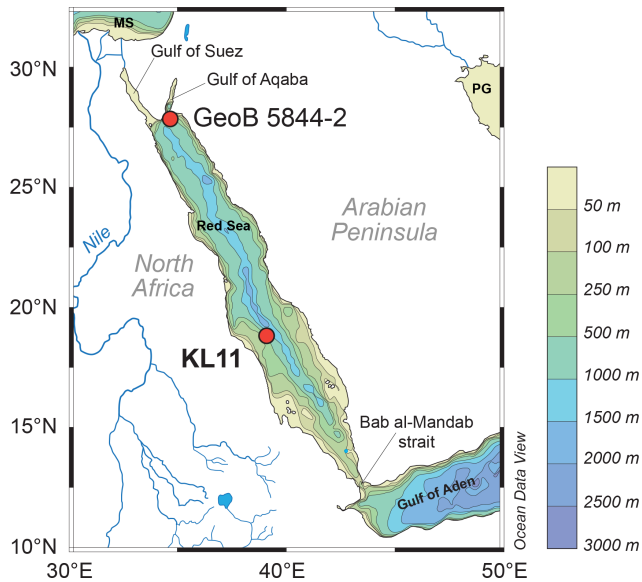


Figure 1. Bathymetric map of the Red Sea, with the locations of core GeoTü KL11 investigated in this study and GeoB5844-2 (Arz et al., 2007). Map generated with Ocean Data View (Schlitzer, 2015).

flows over the sill at Bab al-Mandab and spreads southward into the Indian Ocean along the African continental slope, where it can be traced back to the Agulhas Current (Roman and Lutjeharms, 2009).

An oxygen minimum zone is developed below the oxygen-saturated surface layer, between approximately 200 and 700 m water depth (Sofianos and Johns, 2007). The most extreme oxygen deficiencies are restricted to the central and southern Red Sea basin, with oxygen concentrations as low as $10 \mu\text{mol kg}^{-1}$ ($\sim 0.2 \text{ mL L}^{-1}$) (Fig. 3). The productivity in the surface water reveals regional and seasonal contrasts, with generally oligotrophic conditions in the central and northern Red Sea but meso- to eutrophic conditions south of 19°N (Raitsos et al., 2013). Maximum phytoplankton activity occurs during winter and is related to vertical mixing in the northern and central Red Sea and wind-induced horizontal intrusion of nutrient-rich water from the Gulf of Aden in the south (Naqvi et al., 1986; Eshel and Naik, 1997; Raitsos et al., 2013). Surface productivity is minimal during summer stratification in most areas of the Red Sea, but the intrusion of GAIW into the surface water results in phytoplankton blooms in the southern Red Sea (Dreano et al., 2016).

The paleoceanography of the Red Sea during the late Quaternary was closely linked to global sea level changes regulating the exchange of water masses between the Red Sea and the Indian Ocean. Reduced water exchange occurred during glacial sea level lowstands, leading to a drastic increase in Red Sea SSS (Thunell et al., 1988; Locke and Thunell, 1988). The glacial salinity increase resulted in a decrease in the abundance of pteropods and planktic foraminifera (Almogi-

Labin et al., 1998) and ultimately led to the development of “aplanktic” zones when the upper salinity limits for planktic foraminifera were exceeded (Fenton et al., 2000; Trommer et al., 2011). The strong salinity changes are reflected by high-amplitude fluctuations in the planktic $\delta^{18}\text{O}$, with glacial to interglacial differences of up to $\sim 5\text{‰}$ (Hemleben et al., 1996). The planktic $\delta^{18}\text{O}$ record facilitated the reconstruction of detailed eustatic sea level fluctuations for the past 500 kyr, allowing for the evaluation of glacial lowstands and interglacial highstands, short-term fluctuations in response to abrupt climate change, and polar ice sheet dynamics (Rohling et al., 1998, 2004, 2009; Siddall et al., 2003; Arz et al., 2007; Bouilloux et al., 2013; Grant et al., 2014).

The glacial–interglacial changes in SSS and alternating exposure and flooding of shallow shelf areas in the gulfs of Aqaba and Suez modulated the formation rate of intermediate and deep-water masses in the northern Red Sea and thus the ROC. Despite strongly increased SSS during glacials (Hemleben et al., 1996), the abolition of dense-water formation sites at low sea levels resulted in a reduction in the ROC, as indicated by increased proportions of low-oxygen-tolerant taxa and drops in the diversity of deep-sea benthic foraminifera (Badawi et al., 2005). The available benthic stable isotope records, which could provide further insights into deep-water circulation changes, are either intermittent (Hemleben et al., 1996) or combine epi- and infaunal taxa (Arz et al., 2007), which involve potential biases concerning the influence of strong porewater and metabolic effects (Theodor et al., 2016). To date, a detailed reconstruction of the changes in deep-water circulation and ventilation is still missing. Particularly, it remains unclear how the glacial ROC is influenced by the millennial-scale climate variability in the North Atlantic and if the stable carbon isotopic composition of the dissolved inorganic carbon (DIC) of the deep-water mass is also influenced by the African–Indian monsoon system.

These open questions will be addressed with our study. Especially, we present the first high-resolution composite epibenthic foraminiferal stable isotope record of Marine Isotope Stage (MIS) 3 from an intermediate water depth of the central Red Sea. The data are evaluated for (i) orbital and millennial-scale variability and (ii) the relations between the regional hydrological and biogeochemical processes with the climate variability in the high and low latitudes.

2 Material and methods

Sediment core GeoTü KL11 (in the following referred to as KL11; Figs. 1 and 3) was recovered in 1987 during RV *Me-teor* cruise M5/2 (Hemleben, 1996). It was retrieved from the central Red Sea ($18^\circ 44.5' \text{N}$, $39^\circ 20.6' \text{E}$) at 825 m water depth. Total core recovery was 21.0 m. For this study, we sampled the interval from 194 to 391 cm at 1 to 2 cm spacing, summing up to a total of 104 samples. We wet sieved 2–3 g freeze-dried sediment of each sample over a $63 \mu\text{m}$ mesh.

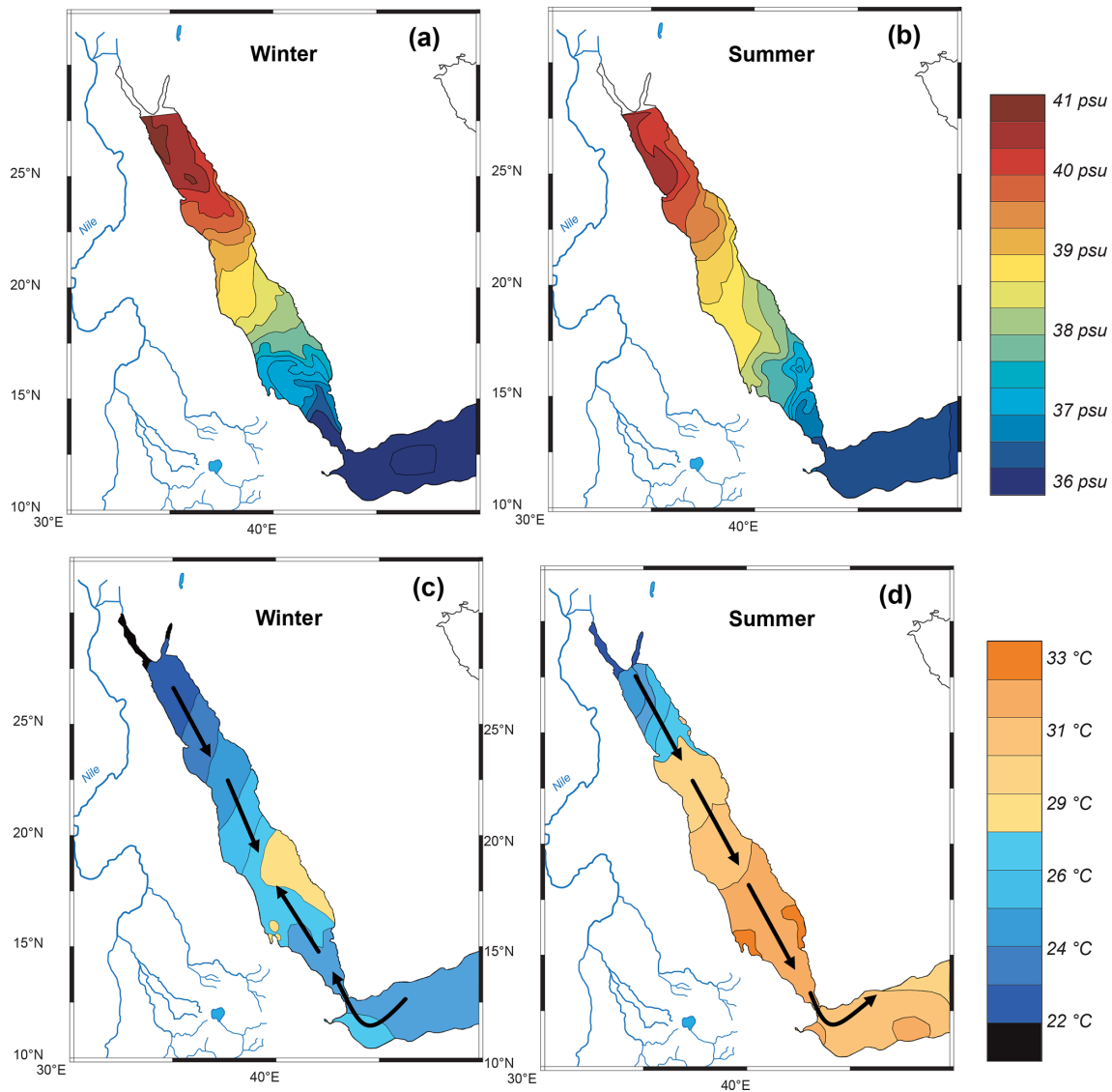


Figure 2. Seasonal sea surface salinity distribution during the (a) winter (November–March) and the (b) summer (June–September) (Sofianos et al., 2003). Seasonal sea surface temperature distribution and averaged net surface currents during the (c) winter (October–April) and the (d) summer (May–September) (Raitsois et al., 2013, 2015).

The residue was dried at 38 °C and subsequently dry-sieved over a 125 µm mesh. The fraction > 125 µm was investigated in our study.

For stable isotope analyses, we selected the benthic foraminiferal species *Cibicides mabahethi*, *Discorbinella bertheloti* s.l., and *Hanzawaia boueana* s.l., all inhabiting a preferentially epifaunal microhabitat (Rathburn and Corliss, 1994; Edelman-Furstenberg et al., 2001; Murray, 2006; Margreth et al., 2009). The species were not present in all samples but exhibited concurrent occurrences in some intervals, allowing for the generation of a composite stable isotope record with an average of the up to three epibenthic isotope measurements. For each species, at least five individuals were selected per sample, and their diameters and preser-

vation states were documented. The stable isotope analyses were performed on a Finnigan MAT 253 mass spectrometer in conjunction with an automatic Kiel IV carbonate preparation device at the Leibniz-Laboratory for Radiometric Dating and Stable Isotope Research, Kiel. Sample reaction was induced by the individual acid addition (99 % H_3PO_4 at 75 °C) under vacuum. The evolved carbon dioxide was analyzed 8 times for each sample. As documented by the performance of international carbonate standards (NBS19 +1.95 ‰ Vienna Pee Dee Belemnite (VPDB) (^{13}C), −2.20 ‰ VPDB (^{18}O); IAEA-603 +2.46 ‰ VPDB (^{13}C), −2.37 ‰ VPDB (^{18}O)) and the laboratory's internal carbonate standards (Hela +0.91 ‰ VPDB (^{13}C), +2.48 ‰ VPDB (^{18}O); HB1 −12.10 ‰ VPDB (^{13}C), −18.10 ‰

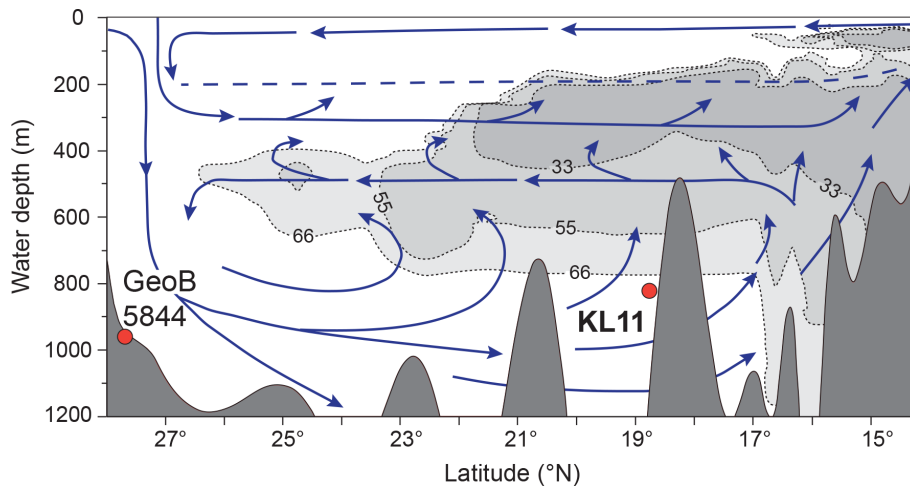


Figure 3. Generalized overturning circulation (blue arrows; Cember, 1988) and dissolved oxygen concentration (stippled lines and shadings; Sofianos and Johns, 2007) along a NNW–SSE transect of the Red Sea. Oxygen concentrations are given (in $\mu\text{mol kg}^{-1}$). The depth positions of sediment cores GeoTü KL11 and GeoB5844-2 are indicated. The blue stippled line represents the base of the pycnocline (Cember, 1988).

VPDB (^{18}O); SHK $+1.74\text{‰}$ VPDB (^{13}C), -4.85‰ VPDB (^{18}O)), the analytical precision of stable isotope analysis is better than $\pm 0.08\text{‰}$ for $\delta^{18}\text{O}$ and $\pm 0.05\text{‰}$ for $\delta^{13}\text{C}$. Values are given in δ notation versus VPDB.

Various age models have been published for core KL11, mainly based on graphical correlations of the planktic $\delta^{18}\text{O}$ record with the global standard $\delta^{18}\text{O}$ record (Hemleben et al., 1996), the Antarctic ice core record (Siddall et al., 2003), the Soreq speleothem record (Grant et al., 2012), and further refinement of the latter by including AMS ^{14}C ages (Hartman et al., 2020). The consideration of AMS ^{14}C ages in the Red Sea may be problematic because of the uncertainties in reservoir ages and potential early diagenetic effects (Rohling et al., 2008). Nevertheless, in order to avoid the tuning of our record to external time series, we established our age model for MIS 3 of KL11 by including two AMS ^{14}C dates (Schmelzer, 1998) and correlating the composite epibenthic $\delta^{18}\text{O}$ record with the benthic $\delta^{18}\text{O}$ record of the well-dated core GeoB5844-2 from 963 m water depth of the northernmost Red Sea (Arz et al., 2007) (Figs. 1 and 4; Table 1). The age model of the interval 25–65 ka of GeoB5844-2 is based on 5 AMS ^{14}C dates and on correlation of the magnetic paleointensity record to the North Atlantic paleointensity stack in the interval older than 40 kyr (Fig. 4). For consistency with the published radiocarbon ages and age model of Arz et al. (2007), we refrained from recalibrating the AMS ^{14}C dates using newer calibration curves. The synchronization of the two cores is justified because their benthic $\delta^{18}\text{O}$ records resemble each other due to the common main deep-water source situated in the northern Red Sea. The deep-water mass is relatively homogenous, with similar temperatures and salinities across the basin (Cember, 1988; Woelk and Quadfasel, 1996). A similar homogeneity of deep-water temperature and salinity obviously also persisted during the past

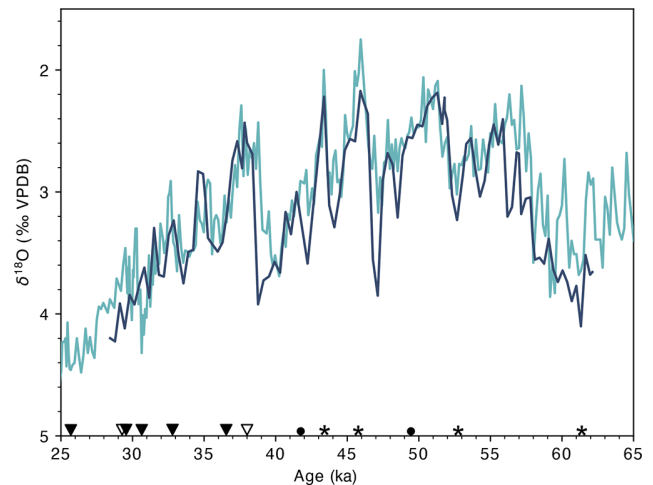


Figure 4. Alignment of benthic $\delta^{18}\text{O}$ records of cores KL11 from the central Red Sea (dark blue) and GeoB5844-2 from the northern Red Sea (light blue). Black triangles represent the radiocarbon dates and black dots the paleomagnetic data used for the establishment of the age model of GeoB5844-2 (Arz et al., 2007). The age model of KL11 is based on two radiocarbon dates of KL11 (white triangles; Schmelzer, 1998) and the graphical correlation of the $\delta^{18}\text{O}$ records. Black asterisks represent the graphical tie points.

glacial period, as illustrated by the close resemblance of the benthic $\delta^{18}\text{O}$ records from the northern and central Red Sea (Fig. 4). According to our age model, the studied core section covers the time interval from 62.1 to 28.4 ka, with an average sample resolution of 330 ± 90 years.

Table 1. Data used for establishing the age model for the studied section of KL11.

Core depth (cm)	Age (cal ka)	Datum	Reference
199.5	29.29 ± 0.35	AMS ¹⁴ C, KL11	Schmelzer (1998)
250.5	38.00 ± 0.71	AMS ¹⁴ C, KL11	Schmelzer (1998)
278.5	43.38	δ ¹⁸ O of GeoB5844-2	Arz et al. (2007)
292.5	45.92	δ ¹⁸ O of GeoB5844-2	Arz et al. (2007)
331.5	52.67	δ ¹⁸ O of GeoB5844-2	Arz et al. (2007)
385.5	61.32	δ ¹⁸ O of GeoB5844-2	Arz et al. (2007)

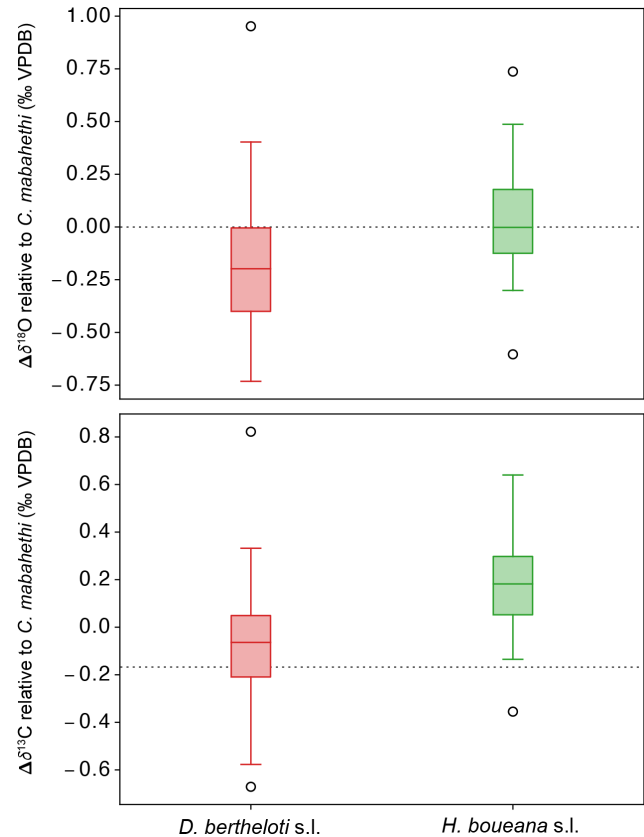
3 Results

The stable isotope values exhibit specific offsets between the different taxa in both δ¹⁸O and δ¹³C. The calculation of average offsets between the species is based on 60 paired analyses of *C. mabahethi* and *D. bertheloti* s.l. and 40 paired analyses of *C. mabahethi* and *H. boueana* s.l. Based on *C. mabahethi* as the reference value, the mean δ¹⁸O deviation of *D. bertheloti* s.l. is $-0.19‰ ± 0.30‰$ and that of *H. boueana* s.l. is $+0.02‰ ± 0.24‰$. The corresponding mean δ¹³C deviations are $-0.09‰ ± 0.30‰$ for *D. bertheloti* s.l. and $+0.17‰ ± 0.19‰$ for *H. boueana* s.l. (Fig. 5). The mean Δδ¹⁸O and Δδ¹³C values were used to adjust the stable isotope values of *D. bertheloti* s.l. and *H. boueana* s.l. to that of *C. mabahethi* and to generate the composite stable isotope records, which are described and evaluated below. The average standard deviation for the composite δ¹⁸O and δ¹³C records is $±0.16‰$ and $±0.13‰$, respectively. These values are considerably lower when compared to the observed amplitudes of temporal fluctuations (Figs. 6–8).

The composite δ¹⁸O record fluctuates between approximately 2.2‰ and 4.2‰ VPDB and shows rapid millennial-scale fluctuations, superimposed by a long-term trend from high to low values and back to high values. The short-term δ¹⁸O fluctuations are of the order of 0.5‰–1.5‰. Pronounced maxima are centered around 61, 47, 39, and 29 ka (Fig. 6a).

The values of the composite δ¹³C record fluctuate between approximately 0.6‰ and 1.9‰ VPDB. The record is also characterized by millennial-scale fluctuations, which are, however, of lower amplitude ($~0.5‰$ – $1‰$) when compared to those of the δ¹⁸O record. In addition, the short-term changes in the δ¹³C record are more pronounced in the interval between 48.5 and 28.4 ka, while a more gradual δ¹³C decrease is observed between 62.1 and 48.5 ka (Fig. 6b).

Accordingly, the δ¹⁸O and δ¹³C records reveal distinct differences in their correlation across the studied time interval. While the correlation of the entire composite isotope records delivers a coefficient of determination (R^2) of 0.43, it is considerably higher, with $R^2 = 0.64$ for the interval 48.5–28.4 ka but only $R^2 = 0.17$ for the interval 62.1–48.5 ka (Fig. 6c–e).

**Figure 5.** Box plots for the δ¹⁸O and δ¹³C deviations of *D. bertheloti* s.l. and *H. boueana* s.l., with reference to *C. mabahethi* in samples from MIS 3 of KL11.

4 Discussion

The epibenthic stable oxygen and carbon isotope records of KL11 display high-amplitude millennial-scale fluctuations (Fig. 6), suggesting rapid changes in deep-water temperature and/or salinity and shifts in the deep-water residence time and organic matter fluxes of the central Red Sea. Despite potential uncertainties in the age model, the pattern in the epibenthic δ¹⁸O record of KL11 closely resembles that of the high-latitude climate variability as documented in the North Greenland Ice Core Project (NGRIP) ice core record (Figs. 7 and 8) (North Greenland Ice Core Project members, 2004; Svensson et al., 2008; Wolf et al., 2010). Specifically, low δ¹⁸O values in KL11 coincide with Dansgaard–Oeschger interstadials and also exhibit the typical pattern of interstadial events, comprising a sharp and rapid increase phase, a plateau phase, and a slower decrease phase (e.g., 51, 46, and 38 ka). High δ¹⁸O values in KL11 coincide with stadials. Particularly high δ¹⁸O values are related to Heinrich stadials (Figs. 7 and 8). The close association of deep-water formation processes in the northern Red Sea and high northern latitude climate variability is nicely illustrated by the difference between the epibenthic (deep-water) δ¹⁸O record of KL11

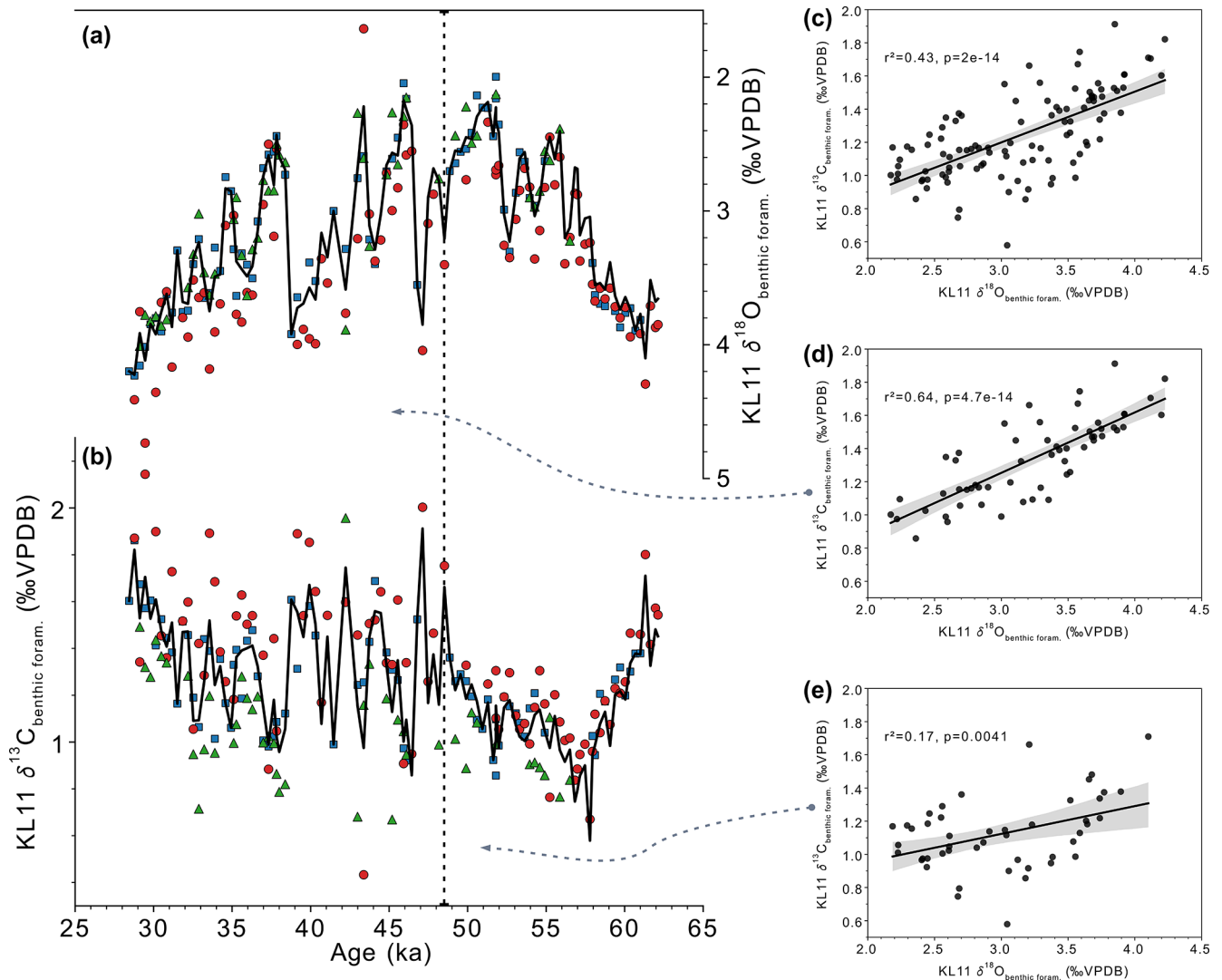


Figure 6. Composite stable (a) oxygen and (b) carbon isotope records for the interval ~ 62 – 28 ka of KL11 (black lines), representing the averages of up to three analyzed benthic foraminiferal species. Before calculating the averages, the isotope values of *Discorbinella bertheloti* s.l. and *Hanzawaia boueana* s.l. were corrected by their mean species-specific offsets from the signal of *Cibicides mabahethi*, as shown in Fig. 5. The mean standard deviation of the calculated averages in the composites is ± 0.16 ‰ for $\delta^{18}\text{O}$ and ± 0.13 ‰ for $\delta^{13}\text{C}$. The markers indicate isotopic measurements of *C. mabahethi* (blue squares), *D. bertheloti* s.l. (red dots), and *H. boueana* s.l. (green triangles). The correlation between composite epibenthic $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ data are shown in panel (c) for the entire time interval and separated for the intervals of (d) 48.5 to 28.4 ka and (e) 62.1 to 48.5 ka. The shaded area around the calibration indicates the 95 % confidence interval of the regressions.

and the planktic (surface water in the vicinity of deep-water formation sites) $\delta^{18}\text{O}$ record of GeoB5844-2 (Fig. 7b). The $\Delta\delta^{18}\text{O}$ record reveals maximum values for stadials and Heinrich events, highlighting the formation of significantly colder and probably also more saline water masses at deep-water formation sites during Northern Hemisphere cold events. This provides evidence for a dominant Northern Hemisphere climate control of Red Sea deep-water formation and, thus, on the ROC during MIS 3. This interpretation is generally corroborated by the close resemblance to the $\Delta\delta^{18}\text{O}$ record based on the endobenthic (*Bulimina marginata*) and planktic

(*Globigerinoides ruber*) $\delta^{18}\text{O}$ data of GeoB5844-2 (Fig. 7c). However, the consideration of *B. marginata* is less reliable, since this species prefers an endobenthic microhabitat (Jorissen and Wittling, 1999), and therefore, its $\delta^{18}\text{O}$ signal may additionally reflect changes in the alkalinity gradient of the porewater (Schmiedl and Mackensen, 2006).

Heinrich stadials are associated with a strong reduction or even cessation of North Atlantic deep-water formation (McManus et al., 2004; Denton et al., 2010). Proxy data and model results demonstrated that the slow-down of the Atlantic meridional overturning circulation also influenced

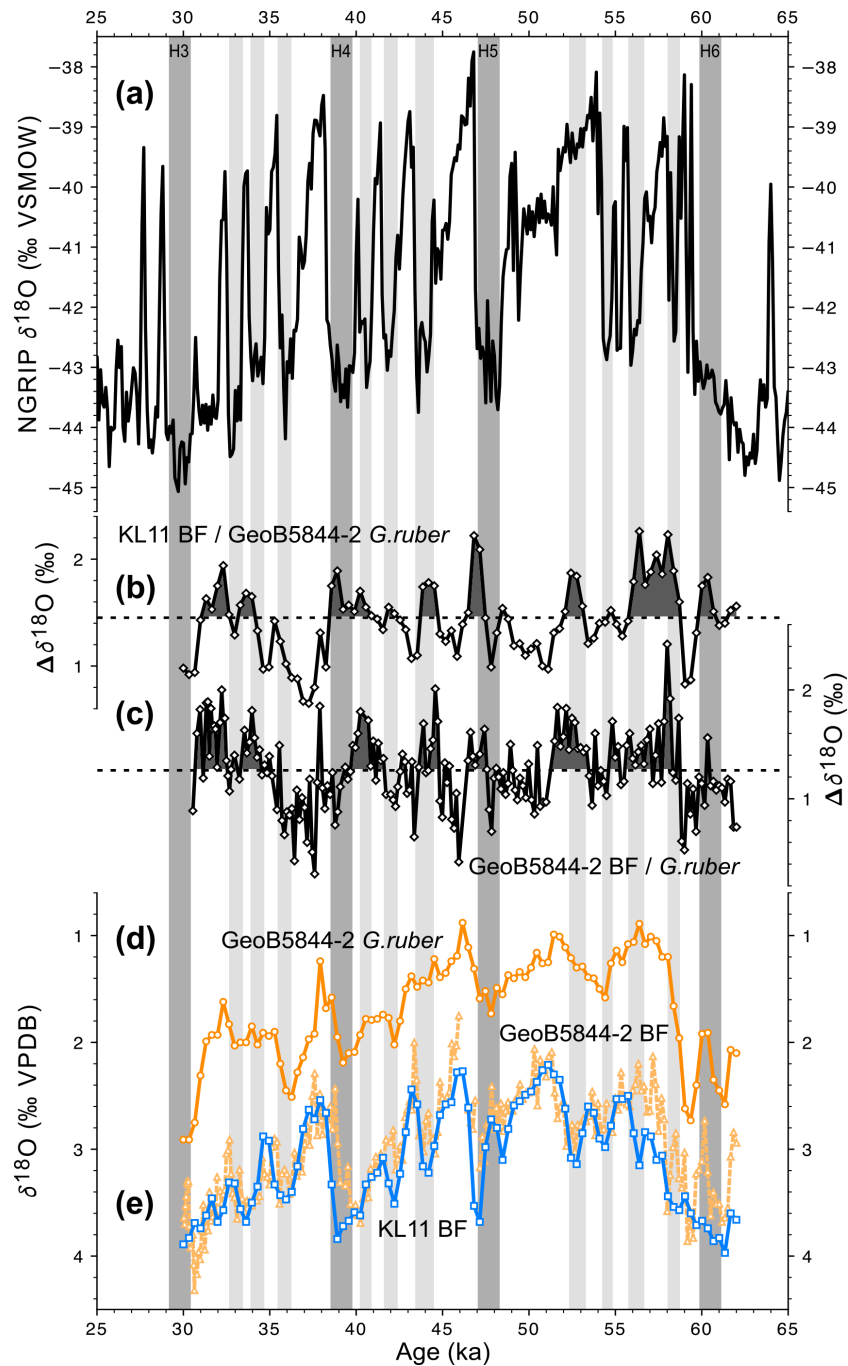


Figure 7. Comparison of epibenthic and planktic stable oxygen isotope records from the Red Sea, their resulting difference, and the Greenland stable oxygen isotope record. **(a)** $\delta^{18}\text{O}$ record of the North Greenland Ice Core Project (NGRIP members, 2004) against the extended GICC05 age scale (Svensson et al., 2008; Wolf et al., 2010). **(b)** Difference in the stable oxygen isotope records ($\Delta\delta^{18}\text{O}$) of the composite epibenthic $\delta^{18}\text{O}$ of KL11 from the central Red Sea and the planktic (*Globigerinoides ruber*) $\delta^{18}\text{O}$ of GeoB5844-2 from the northern Red Sea (Arz et al., 2007). For the calculation of the $\Delta\delta^{18}\text{O}$ values, the single records were resampled at a spacing of 330 years. **(c)** Difference in the stable oxygen isotope records ($\Delta\delta^{18}\text{O}$) of the endobenthic (*Bulimina marginata*) and planktic (*G. ruber*) $\delta^{18}\text{O}$ of GeoB5844-2 from the northern Red Sea (Arz et al., 2007). **(d)** Planktic (*G. ruber*) $\delta^{18}\text{O}$ of GeoB5844-2 from the northern Red Sea (Arz et al., 2007). **(e)** Composite epibenthic $\delta^{18}\text{O}$ of KL11 from the central Red Sea (blue) and endobenthic (*B. marginata*) $\delta^{18}\text{O}$ of GeoB5844-2 from the northern Red Sea (stippled orange; Arz et al., 2007). The horizontal stippled lines in the $\Delta\delta^{18}\text{O}$ records mark the mean values. Gray bars represent Northern Hemisphere stadials and Heinrich events.

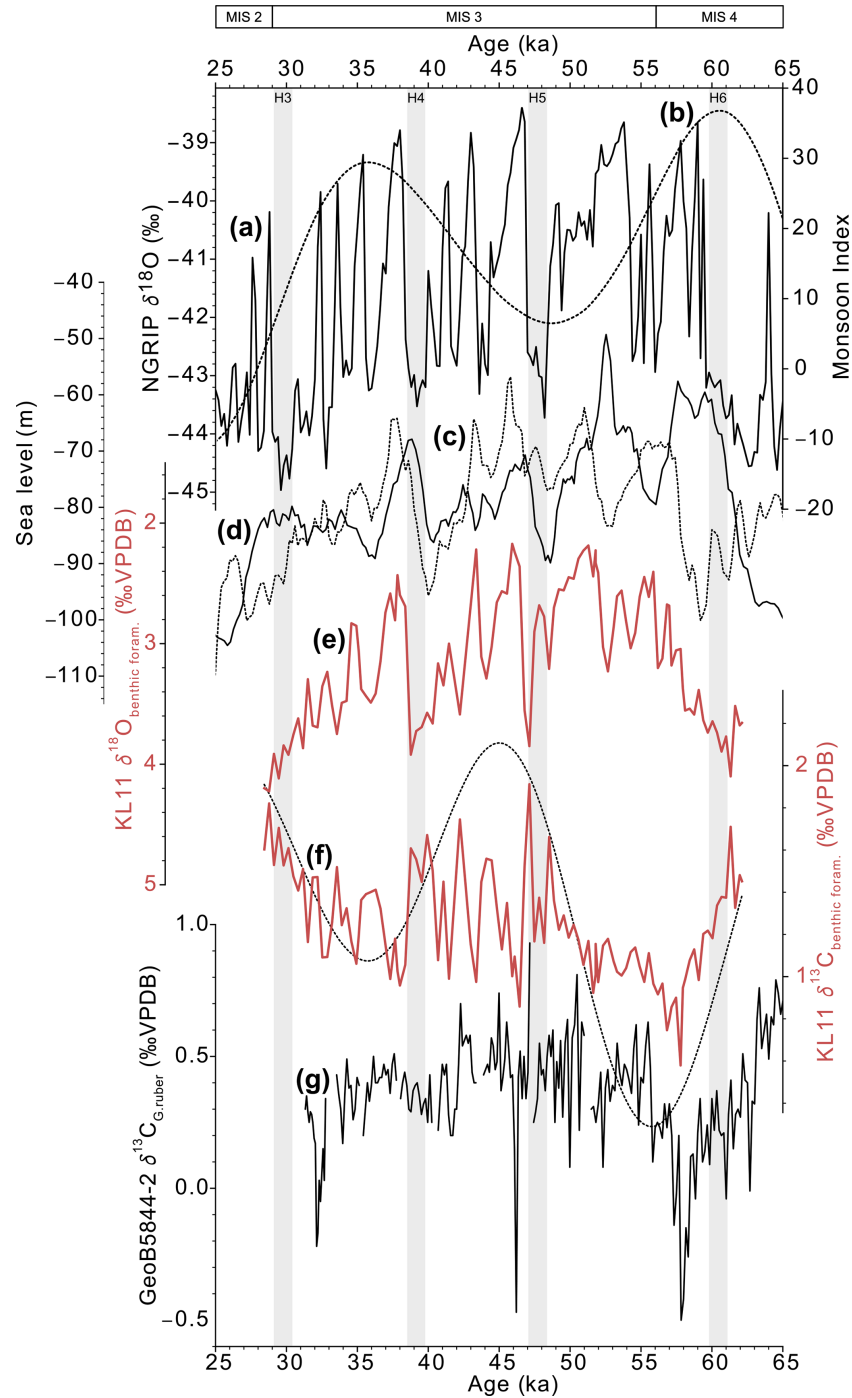


Figure 8. Composite epibenthic stable oxygen and carbon isotope records of the central Red Sea compared with the planktic stable carbon isotope record of the northern Red Sea, sea level reconstructions for the Red Sea, and climate changes in the high northern latitudes for Marine Isotope Stages (MISs) 2 to 4. **(a)** $\delta^{18}\text{O}$ record of the North Greenland Ice Core Project (NGRIP members, 2004) against the extended GICC05 age scale (Svensson et al., 2008; Wolf et al., 2010). **(b)** Monsoon index calculated according to Rossignol-Strick (1983), based on the 21 June insolation at 23.45°N and at the Equator (Laskar et al., 2004). **(c)** Sea level reconstructions for the northern Red Sea (dashed line, Arz et al., 2007) and **(d)** central Red Sea (black line, Siddall et al., 2003; Rohling et al., 2008). **(e)** Composite epibenthic $\delta^{18}\text{O}$ record of KL11 from the central Red Sea. **(f)** Composite epibenthic $\delta^{13}\text{C}$ record of KL11 from the central Red Sea. The stippled line represents the band-pass-filtered precession (23 kyr) component of the epibenthic $\delta^{13}\text{C}$ record. Note that minima in the $\delta^{13}\text{C}$ precession component correspond to maxima in the monsoon index. **(g)** $\delta^{13}\text{C}$ record of the planktic foraminifer *Globigerinoides ruber* white of GeoB5844-2 from the northern Red Sea (Arz et al., 2007). Note that the insolation maximum centered around 58–62 ka is associated with a transient decrease in the planktic and epibenthic $\delta^{13}\text{C}$ values. The positions of Heinrich stadials H3–H6 are indicated by light gray bars.

the hydrology of mid- to low-latitude regions in the Northern Hemisphere, leading to megadroughts in northern Africa and the Mediterranean region (Mulitza et al., 2008; Hamann et al., 2008; Ehrmann et al., 2017; Allard et al., 2021) and weakening of the Indian summer monsoon (Schulz et al., 1998). It is likely that during these droughts, enhanced evaporation also resulted in increased sea surface salinity in the northern Red Sea, fostering the formation of dense waters in this region. Previous studies have confirmed a strong influence of changes in the Mediterranean and North Atlantic climate on the environmental changes in the northern Red Sea (Arz et al., 2003a, b, 2007; Lamy et al., 2006). Alkenone temperature reconstructions for the northern Red Sea display a general cooling trend during MIS 3, punctuated by short-term sea surface temperature (SST) drops of the order of 1 to 4 °C during Heinrich stadials. We speculate that the strong cooling could be associated with the inflow of cold-air masses from the north responding to phases of intensified Siberian High. The strength of the Siberian High is linked to the North Atlantic climate variability. It is generally stronger during glacials due to the enhanced surface cooling of the Eurasian continent (Vandenberghé et al., 2006). In the eastern Mediterranean region, northeasterly outbreaks of cold air are linked to the strength of the Siberian High (Rohling et al., 2002; Casford et al., 2003). The close correspondence of grain size in Asian loess deposits and the Greenland ice core record suggests a stronger Siberian High during stadials of the last glacial period (Vandenberghé et al., 2006; Cheng et al., 2022), possibly also explaining the strong SST drops in the northern Red Sea during Heinrich events. Pollen data from the Aegean region suggest extremely severe climate deterioration during Heinrich stadial H5 (Müller et al., 2011), which is also reflected in the deep-sea $\delta^{18}\text{O}$ record of the Red Sea (Figs. 6–8).

Similarly, the reconstructed sea level fluctuations in the northern Red Sea also occurred in phase with Northern Hemisphere climate variability, with transient sea level rises of up to 25 m during Greenland interstadials (Arz et al., 2007). These results contrast with earlier findings from the central Red Sea, where estimated short-term sea level changes are of the order of 35 m and occurred in phase with Antarctic climate changes (Siddall et al., 2003), suggesting a strong southern meltwater component. Subsequent model simulations and comparison with stable oxygen isotope data suggest similar meltwater contributions from both Antarctic and northern ice sheets during the last glacial period (Rohling et al., 2004) (Fig. 8d). To date, the establishment of accurately dated sea level records is limited by the various age scales used in Greenland and Antarctic ice core records (review in Siddall et al., 2008). Some of the discrepancies in the available sea level reconstructions from the Red Sea may be attributed to the different age model strategies, involving potential dating uncertainties and diagenetic biases (Rohling et al., 2008). In addition, the extraction of the sea level component from the benthic $\delta^{18}\text{O}$ signal strongly depends on the

correct reconstruction of temperature and salinity effects at deep-water formation sites, which appears challenging. Despite these biases and conflicting reconstructions millennial-scale sea level changes during the last glacial period seem to essentially follow an Antarctic climate pattern (Siddall et al., 2008). However, the subsequent study of Grant et al. (2012) from the eastern Mediterranean Sea suggested that the large-scale sea level variability reflects a global signature of climate changes recorded in both Antarctic and Greenland ice cores.

The close resemblance of the epibenthic $\delta^{18}\text{O}$ record with that of the Greenland ice core record suggest a direct atmospheric control on short-term changes in deep-water circulation during the last glacial period. Since maximum deep-water formation is observed during Heinrich stadials, when sea level dropped, the moderate further exposure of shelf areas in the northern Red Sea obviously did not play a dominant role on deep-water formation. Instead, the more restricted exchange between the Red Sea and the Indian Ocean and the associated SSS rise in the Red Sea likely facilitated dense-water formation (Arz et al., 2007) (Figs. 7 and 8). However, the succession of Dansgaard–Oeschger and Heinrich events is largely superimposed by salinity effects in the planktic $\delta^{18}\text{O}$ records of the Red Sea (Siddall et al., 2003; Arz et al., 2007) (Fig. 7). According to these findings, cooling and enhanced evaporation rates prevailed in the northern Red Sea region during Heinrich stadials and fostered the formation of dense waters in the deep-water formation sites. In the time interval ~ 49 –28 ka, epibenthic $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ records of KL11 show a positive correlation, with high $\delta^{18}\text{O}$ values during Heinrich stadials being associated with high epibenthic $\delta^{13}\text{C}$ values. The lack of differences between the $\delta^{13}\text{C}$ records of KL11 and GeoB5844-2 affirm that the epibenthic stable carbon isotope can be used to reflect the residence time. This confirms our conclusion of more vigorous ventilation of deep-water masses with low residence times during Heinrich stadials and reduced ventilation with the presence of older and thus ^{12}C -enriched deep-water masses during Dansgaard–Oeschger interstadials.

For the correct interpretation of the deep-sea $\delta^{13}\text{C}$ record, the potential contribution of $\delta^{13}\text{C}$ changes in the dissolved inorganic carbon (DIC) in the source areas of deep-water formation in the northern Red Sea must be considered. Therefore, we compared the epibenthic $\delta^{13}\text{C}$ record of KL11 with the $\delta^{13}\text{C}$ record of the planktic foraminifer *Globigerinoides ruber* (white) of GeoB5844-2 from the northernmost Red Sea (Arz et al., 2007) (Fig. 8). The epibenthic carbon isotopic composition reflects the addition of ^{12}C to the intermediate and deep water through decomposition of sinking organic matter. Despite the preferentially epifaunal lifestyle of the measured taxa, additional porewater effects of around 0.2‰–0.5‰ cannot be ruled out, as suggested by the analyses on morphologically similar species of the genera *Cibicides* and *Cibicidoides* (McCorkle et al., 1990; Schmiedl et al., 2004; Theodor et al., 2016). Various studies have shown

that the $\delta^{13}\text{C}$ of *G. ruber* reflects not only the $\delta^{13}\text{C}$ of ambient DIC but is also influenced by biological fractionation effects, mainly by algal photosymbiosis and the metabolism of the foraminifer (Rohling and Cooke, 1999; Schiebel and Hemleben, 2017, and references therein). Accordingly, the $\delta^{13}\text{C}$ of *G. ruber* commonly deviates significantly from $\delta^{13}\text{C}_{\text{DIC}}$. Depending on the magnitude of the different effects, the deviations of $\delta^{13}\text{C}_{G,\text{ruber}}$ from $\delta^{13}\text{C}_{\text{DIC}}$ range from +1‰, due to symbiotic enrichment (Bemis et al., 1998), to −0.5‰ to −1‰, due to metabolic depletion (Niebler, 1995; Katz et al., 2010; Birch et al., 2013). Despite these offsets, the epibenthic and planktic $\delta^{13}\text{C}$ records of KL11 reflect a generally similar long-term trend, particularly in the older part of the record. However, the planktic $\delta^{13}\text{C}$ record lacks systematic Heinrich-stadial-associated fluctuations. Thus, the alteration of surface water $\delta^{13}\text{C}_{\text{DIC}}$ by millennial-scale changes in productivity and addition of ^{12}C from land can be largely ruled out. These processes play a more prominent role on orbital timescales. Specifically, last glacial $\delta^{13}\text{C}$ values of the Red Sea surface waters are approximately 0.5‰–1.0‰ lower than Holocene values (Schmelzer, 1998; Bouilloux et al., 2013), suggesting the redistribution of carbon from exposed shelf areas into the Red Sea basin during glacial sea level low stands.

The decoupling of the $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ records in the interval ~62–49 ka suggests the additional influence of orbital-scale regional biogeochemical processes on the $\delta^{13}\text{C}$ signal in the central Red Sea (Fig. 8). The core position of KL11 is situated close to the border between the more oligotrophic central and northern Red Sea and the mesotrophic to eutrophic area in the southern Red Sea (Raitso et al., 2013). According to the modern situation, changes in surface water productivity at site KL11 depend on vertical mixing and formation of eddies during winter (Eshel and Naik, 1997; Raitso et al., 2013) but may also be influenced by the intrusion of nutrient-rich intermediate waters from the Gulf of Aden during summer (Trommer et al., 2010; Dreano et al., 2016).

The past exchange of water masses between the Red Sea and Indian Ocean and the advection of nutrient-rich waters from the Gulf of Aden through the strait of Bab al-Mandab depend on orbital and suborbital glacioeustatic changes and shifts in monsoon wind intensity (Siddall et al., 2004; Trommer et al., 2011; Bouilloux et al., 2013). During the time interval ~60–50 ka, the global sea level was approximately 60 m lower than at present but higher than during the sea level low stands of MIS 4 (approximately −100 m) and the last glacial maximum (approximately −120 m) (Siddall et al., 2003; Rohling et al., 2004; Grant et al., 2014) (Fig. 8). Within MIS 2–4, sea level was at a maximum between ~60 and 50 ka. Accordingly, the exchange of intermediate waters likely persisted during this period, including the inflow of nutrient-rich GAIW during summer, particularly during phases of enhanced summer monsoon winds (Fig. 8).

The time interval 58–62 ka coincides with the stronger of two maxima in the Northern Hemisphere monsoon in-

dex during MIS 3 (Fig. 8b). The monsoon index has been calculated according to Rossignol-Strick (1983), based on the 21 June insolation gradient between 23.45° N and at the Equator (Laskar et al., 2004). Generally, maxima in the African-Asian summer monsoon are coherent with insolation maxima (Cheng et al., 2016). However, upwelling proxies from marine sediment cores of the Arabian Sea exhibit time lags of ~8 kyr between insolation and monsoon maxima, which has been attributed to the impact of ice volume of the Northern Hemisphere and the transport of latent heat from the southern subtropical Indian Ocean to the Tibetan Plateau (Clemens and Prell, 2003; Clemens et al., 2010). A new model study demonstrated that the observed time lag in the Arabian Sea did not necessarily document changes in Indian summer monsoon intensity. It may rather be caused by shifts in the upwelling area from coastal regions during insolation maxima to more open-ocean areas during insolation minima (Jalihal et al., 2022). The decrease in the epibenthic $\delta^{13}\text{C}$ in KL11 lags the monsoon index maximum by only ~3.5 kyr which is comparable to the observed lags of Mediterranean sapropels (Lourens et al., 1996) and also lags in Asian speleothem records (Clemens et al., 2010). The close relation of the monsoon index and epibenthic $\delta^{13}\text{C}$ suggests a more or less immediate response of surface water productivity and related organic matter fluxes at site KL11 to the strength of the summer monsoon. Our results support previous evidence from planktic foraminifera, which exhibit a close correspondence between the high-productivity indicator *Globigerinita glutinata* and summer insolation at site KL11 during the last interglacial period (Trommer et al., 2011). The development of summer phytoplankton blooms in the southern part of the modern Red Sea is related to the intrusion of nutrient-rich intermediate waters from the Gulf of Aden, which can be traced as far north as 19° N (Trommer et al., 2010; Dreano et al., 2016). The different proxy data suggest that this process was intensified in the past during phases of increased Indian summer monsoon and that nutrient-rich waters reached the position of KL11 temporarily during MIS 3. The timing suggests that the nutrient-rich waters were derived from areas along the southern Arabian Peninsula, where maximum upwelling occurred during phases of an intensified Indian summer monsoon (Jalihal et al., 2022).

5 Conclusions

We established a high-resolution composite epibenthic stable oxygen and carbon isotope record from the central Red Sea for the last glacial period. The records show high-amplitude variations during MIS 3, suggesting millennial-scale changes in the thermohaline circulation of the Red Sea. Despite generally reduced ROC during glacial sea level lowstands, deep-water formation increased during cold and hyper-arid conditions in the northern Red Sea borderlands. Inversely, the

formation of dense waters was reduced during warmer and more humid conditions resulting in a diminished ROC.

The millennial-scale changes in aridity in the northern Red Sea region are in phase with the abrupt climate variability in the high northern latitudes, as documented in the Greenland ice core record, which could suggest a link to the strength of the Siberian High and related inflow of cold-air masses to the sites of Red Sea deep-water formation. Our results suggest that the regional hydroclimatic changes modulate the thermohaline circulation of the Red Sea during glacial boundary conditions. Instead, millennial-scale sea level changes and related changes in the exchange between the Red Sea and the Indian Ocean play a subordinate role in the preconditioning of dense-water formation in the northern Red Sea. This is also shown by the lack of coherent millennial-scale changes in the planktic stable isotope records from the northernmost Red Sea representing the general source signal in the area of deep-water formation.

The KL11 stable oxygen and carbon isotope records are decoupled in the early part of MIS 3, caused by a transient decrease in the $\delta^{13}\text{C}$ values in phase with an increase in the Northern Hemisphere summer insolation around 55–60 ka. This suggests a connection between the biogeochemical processes of the central Red Sea to the African–Indian monsoon dynamics. During the summer monsoon maximum and concomitant moderate sea level rise, nutrient-rich intermediate waters intruded from the Gulf of Aden and delivered nutrients as far north as 19° N. The influx of nutrients fueled surface water productivity at site KL11 in the central Red Sea. The associated remineralization of sinking organic matter led to the observed transient decrease in epibenthic $\delta^{13}\text{C}$ values.

Data availability. The new data are available in the Supplement of this paper.

Supplement. The supplement related to this article is available online at: <https://doi.org/10.5194/cp-20-267-2024-supplement>.

Author contributions. GS and WE initiated the initial project (REVENT). RHH was in charge of the sample processing and species selection. NA performed the stable isotope analysis. RHH and GS wrote the first draft of the paper. HWA contributed planktic isotope data. All authors contributed to the interpretation and discussion of the data, and to the writing of the submitted paper.

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