

Reconstructing the thermal structure of the upper ocean: Insights from planktic foraminifera shell chemistry and alkenones in modern sediments of the tropical eastern Indian Ocean

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[1] Shell chemistry of planktic foraminifera and the alkenone unsaturation index in 69 surface sediment samples in the tropical eastern Indian Ocean off West and South Indonesia were studied. Results were compared to modern hydrographic data in order to assess how modern environmental conditions are preserved in sedimentary record, and to determine the best possible proxies to reconstruct seasonality, thermal gradient and upper water column characteristics in this part of the world ocean. Our results imply that alkenone-derived temperatures record annual mean temperatures in the study area. However, this finding might be an artifact due to the temperature limitation of this proxy above 28°C. Combined study of shell stable oxygen isotope and Mg/Ca ratio of planktic foraminifera suggests that *Globigerinoides ruber* sensu stricto (s.s.), *G. ruber* sensu lato (s.l.), and *G. sacculifer* calcify within the mixed-layer between 20 m and 50 m, whereas *Globigerina bulloides* records mixed-layer conditions at ~50 m depth during boreal summer. Mean calcifications of *Pulleniatina obliquiloculata*, *Neogloboquadrina dutertrei*, and *Globorotalia tumida* occur at the top of the thermocline during boreal summer, at ~75 m, 75–100 m, and 100 m, respectively. Shell Mg/Ca ratios of all species show a significant correlation with temperature at their apparent calcification depths and validate the application of previously published temperature calibrations, except for *G. tumida* that requires a regional Mg/Ca-temperature calibration ($Mg/Ca = 0.41 \exp(0.068 \cdot T)$). We show that the difference in Mg/Ca-temperatures of the mixed-layer species and the thermocline species, particularly between *G. ruber* s.s. (or s.l.) and *P. obliquiloculata*, can be applied to track changes in the upper water column stratification. Our results provide critical tools for reconstructing past changes in the hydrography of the study area and their relation to monsoon, El Niño-Southern Oscillation, and the Indian Ocean Dipole Mode.

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1. Introduction

[2] In the tropical Indo-Pacific, semi- and inter-annual changes in the wind strength and precipitation mainly caused

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by monsoon, El Niño-Southern Oscillation (ENSO), and the Indian Ocean Dipole Mode (IOD), significantly affect the thickness and evolution of the mixed-layer and the thermocline as well as the intensity and direction of the surface and subsurface circulation [e.g., Susanto *et al.*, 2001; Rao *et al.*, 2002; Gordon, 2005; Qu and Meyers, 2005]. Reconstruction of the upper water column structure in this region therefore can provide critical information for understanding past behavior of these climate systems and the mechanisms of past, and future, regional climate change.

[3] The vertical structure of the water column can be best reconstructed by using differences in a geochemical proxy for temperature such as shell stable oxygen isotopes ($\delta^{18}O$) or Mg/Ca ratios of various planktic foraminifera species that thrive at different water depths due to their preferred range of temperature, salinity, chlorophyll, light, etc. [e.g., Fairbanks *et al.*, 1980; 1982; Hemleben *et al.*, 1989; Sautter and Thunell, 1991; Ortiz *et al.*, 1995]. As habitat depths of planktic foraminifera

minifera vary in different regions due to different hydrographic characteristics, it is crucial to determine the regional habitat and calcification depths of various species, their seasonality and their environmental control [e.g., *Elderfield and Ganssen*, 2000; *Anand et al.*, 2003; *Mulitza et al.*, 2003; *McConnell and Thunell*, 2005; *Farmer et al.*, 2007; *Cléroux et al.*, 2008; *Mohtadi et al.*, 2009; *Regenberg et al.*, 2009; *Steph et al.*, 2009] in order to minimize errors in paleoceanographic reconstructions based on $\delta^{18}\text{O}$ or Mg/Ca.

[4] Here we present paired Mg/Ca and $\delta^{18}\text{O}$ measurements on seven planktic foraminifera species in 69 sediment surface samples from the eastern equatorial Indian Ocean, W and S off Indonesia. We use shell $\delta^{18}\text{O}$ to infer calcification depths for different species, and compare the shell Mg/Ca ratio with the modern temperature data that correspond to the inferred calcification depths. In addition, we evaluate published Mg/Ca-temperature calibrations in order to assess the best possible modern calibration of this temperature proxy for the eastern equatorial Indian Ocean. We further analyze alkenone-derived sea surface temperatures (SST) in the same surface samples to compare different SST proxies and to assess how modern environmental conditions are preserved in surface sediments. Finally, we introduce different proxies to reconstruct thermal gradients and upper water column characteristics that help to assess past behavior of ENSO, monsoon, and IOD from sedimentary records in the tropical eastern Indian Ocean.

2. Study Area

[5] The study area is part of the Indo-Pacific Warm Pool with mean SST generally exceeding 28°C. The seasonal climate in this region is affected by the Australian-Indonesian monsoon (AIM) winds [e.g., *Tapper*, 2002; *Qu and Meyers*, 2005; *Kida and Richards*, 2009]. The northern and southern portions of the study area are characterized by different seasonality. During the SE monsoon from June–September (boreal summer), alongshore winds induce Ekman pumping and coastal upwelling off S and SW Indonesia that decrease SST by 1–2°C (Figure S1 in the auxiliary material) and increase chlorophyll *a* concentrations south of 4°S (Figure 1a).¹ The NW monsoon season from December to March is characterized by the southward progression of the Intertropical Convergence Zone (ITCZ), increased precipitation, a rather uniform SST distribution, and low chlorophyll *a* concentrations in the entire tropical eastern Indian Ocean (Figure 1b). In contrast to off S and SW Indonesia, a relatively thick mixed-layer of about 70–80 m persists throughout the year off W and NW Indonesia (Figures 1a, 1b, and S1). Surface salinity varies between 33.5 and 34.2 psu in the study area (Figure S2). Seasonal salinity changes are rather small in both regions, with maximum variability of about 0.6 psu in the upwelling areas off southern Indonesia (Figure S2).

[6] On inter-annual timescales, ENSO and the IOD strongly affect the hydrography of the equatorial Indian Ocean [e.g., *Susanto et al.*, 2001; *Qu and Meyers*, 2005; *Susanto and Marra*, 2005; *Zhong et al.*, 2005; *Halkides et al.*, 2006; *Du et al.*, 2008; *Horii et al.*, 2008]. Briefly, El Niño and positive IOD years are associated with anomalously strong

SE winds that reinforce coastal upwelling and induce up to 5°C decrease in SST and higher primary production off S and SW Indonesia during boreal summer (Figures 1c–1e). In contrast, La Niña and negative IOD years amplify the NW monsoon climatic features, i.e., enhanced westerly winds, positive precipitation anomalies, and a uniformly high SST in the study area (Figure 1f).

3. Material and Methods

3.1. Sampling

[7] Surface sediment samples were collected during the RV SONNE cruises 184 (34 samples) [*Hebbeln et al.*, 2005] and 189 (35 samples) [*Wiedicke-Hombach et al.*, 2007] from off W and S Indonesia (Figure 2 and Table 1). The top 1 cm of multicore samples were freeze-dried, washed and sieved for the planktic foraminifera study, and freeze-dried and ground for alkenone analyses, respectively. In order to address different environmental conditions in the study area both on the seasonal and interannual timescales, we grouped the sample material in seven different basins (Figure 2). The Simeulue Basin (SB) off NW Sumatra, the Nias Basin (NB) south of the SB, and the Northern Mentawai Basin (NMB) off W Sumatra are characterized by oligotrophic to mesotrophic conditions and insignificant seasonal and interannual variations in marine productivity, SST, and thermocline depth. In contrast, the Southern Mentawai Basin (SMB), the Java Basin (JB), the Lombok Basin (LB), and the Savu Sea (SS) show a pronounced seasonality of the upper water column structure, primary productivity, and SST due to the occurrence of coastal upwelling in boreal summer.

[8] A study by *Mohtadi et al.* [2007] on the same surface samples showed that carbonate dissolution can be neglected in this set of samples, as either the basins lack the aragonite lysocline (SB, NB, NMB and SMB), or the samples are retrieved from water depths well above the calcite lysocline (JB, LB, SS) and do not show any evidence of selective calcite dissolution.

3.2. Radiocarbon Analyses

[9] Radiocarbon dating was performed on 16 selected surface sediments from all of the different basins in the study area (Table 2). Samples were measured at the Keck Carbon Cycle Accelerator Mass Spectrometry Laboratory, University of California, Irvine (UCI), and at the National Ocean Sciences Accelerator Mass Spectrometry Facility, Woods Hole (OS). UCI samples contained mono-species tests of *Globigerinoides sacculifer* (without sac-like final chamber), while OS samples included multispecies tests of planktic foraminifera. Radiocarbon ages were converted to calendar ages using Marine 09 calibration [*Hughen et al.*, 2004] of CALIB 6.0 software assuming no deviation from the global ocean carbon reservoir.

3.3. Planktic Foraminifera

[10] For all analyses presented in this study, we used whole tests of *Globigerinoides ruber sensu stricto* (s.s.), *G. ruber sensu lato* (s.l.), and *Globigerina bulloides* from the 250–355 μm size fraction, and of *G. sacculifer* (without sac-like final chamber), *Neogloboquadrina dutertrei*, *Pulleniatina obliquiloculata*, and *Globorotalia tumida* from the 355–500 μm size fraction, respectively. Determination of

¹Auxiliary materials are available in the HTML. doi:10.1029/2011PA002132.

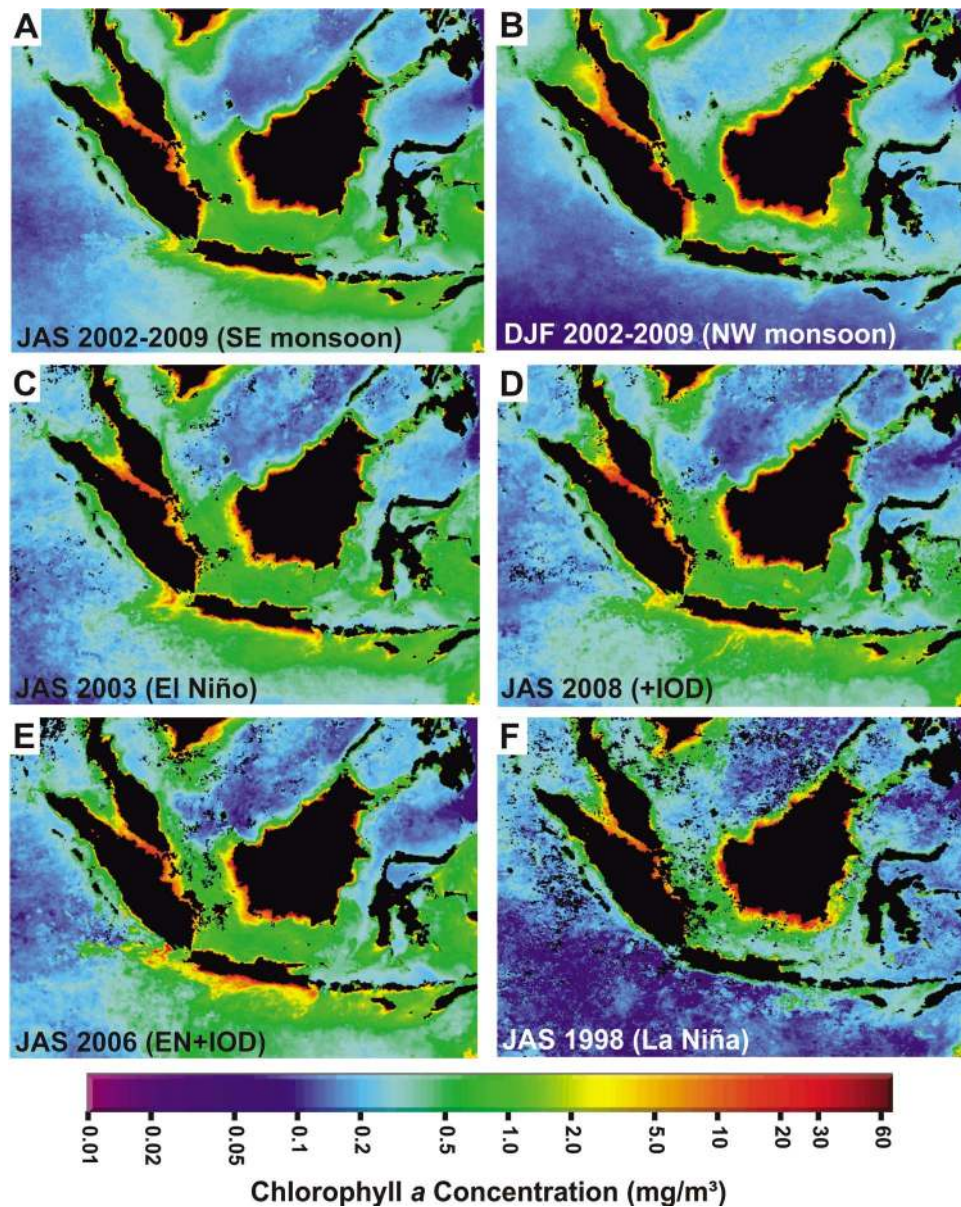


Figure 1. Remote sensing images of the mean seasonal chlorophyll *a* concentration around Indonesia (<http://oceancolor.gsfc.nasa.gov>). Mean values between 2002 and 2009 during (a) the SE monsoon (July–September) and the (b) NW monsoon (December–February) seasons. Here, the effect of monsoon-related seasonal upwelling off S and SW Indonesia is apparent. Chlorophyll *a* concentrations (July–September) during (c) a moderate El Niño year (2003) and (d) a positive IOD year (2008). Chlorophyll *a* concentrations (July–September) during (e) a positive IOD/El Niño year (2006), and (f) a strong La Niña year (1998). Note the enhanced (reduced) chlorophyll *a* concentrations off S and SW Indonesia during the positive IOD/El Niño (La Niña) years indicative of intensified (weakened) upwelling.

G. ruber s.s. and s.l. follows the concept of Wang [2000], in which *G. ruber* s.l. corresponds to the more compact and higher trochospiral forms previously described as *G. elongatus* [d'Orbigny, 1826], *G. pyramidalis* [Van den Broeck, 1876], and *G. cyclostomus* [Galloway and Wissler, 1927].

3.3.1. Oxygen Isotope and Mg/Ca Analyses

[11] A Finnigan MAT 251 mass spectrometer was used to measure the $\delta^{18}\text{O}$ composition of the planktic foraminifera (Table 1). Approximately 5–20 individual tests were picked for each measurement. The isotopic composition of the carbonate sample was measured on the CO_2 gas evolved by

treatment with phosphoric acid at a constant temperature of 75°C. For all stable isotope measurements a standard gas (Burgbrohl CO_2 gas) was used. Samples were calibrated against PDB by using the NBS 19 and an internal laboratory standard (Solnhofen Limestone). All isotopic data given here are relative to the PDB standard. Analytical standard deviation is about $\pm 0.07\%$ (Isotope Laboratory, Faculty of Geosciences, University of Bremen).

[12] For Mg/Ca analyses, planktic foraminifera tests were utilized from the same size fraction as for the $\delta^{18}\text{O}$ analyses. Samples contained about 30–40 intact tests of *G. ruber* s.s.,

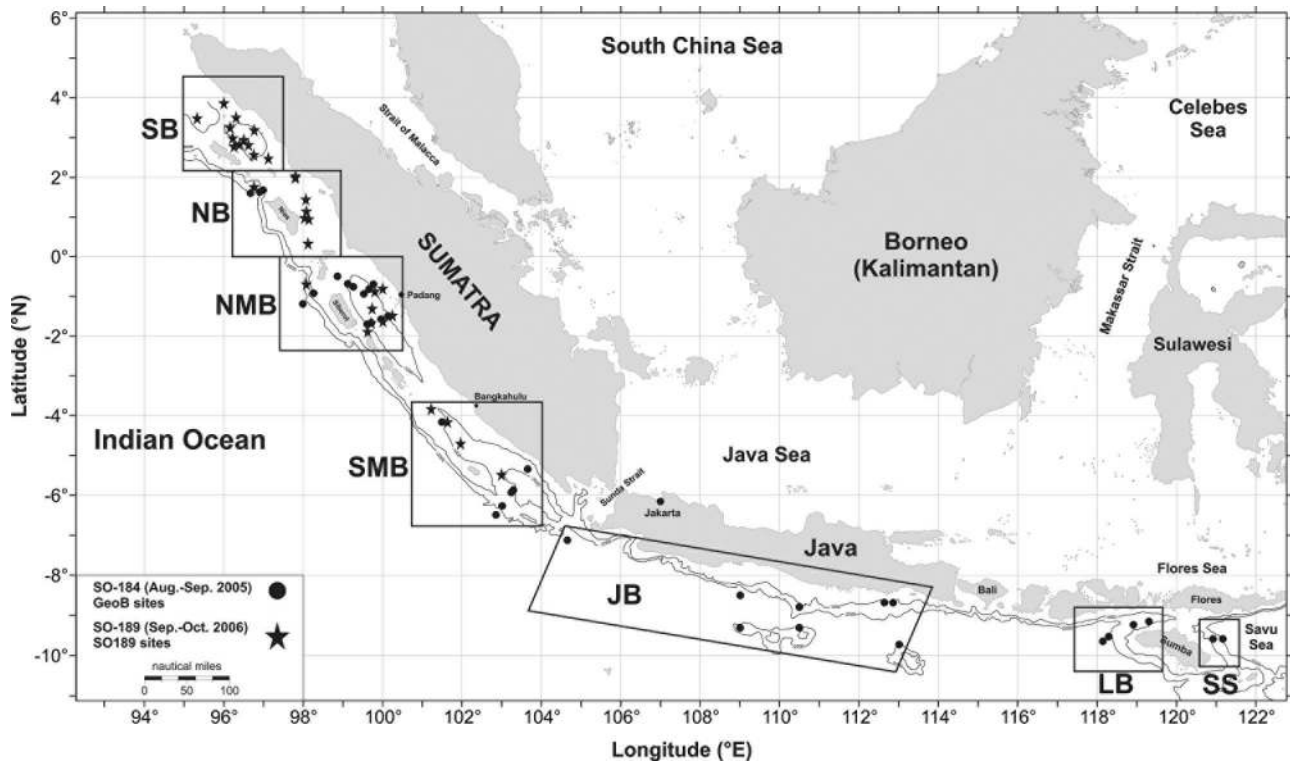


Figure 2. Schematic map of the study area showing the position of the investigated surface sediments collected during the SO-184 (dots) and SO-189 (stars) cruises. Rectangles indicate different fore-arc basins. SB: Simeulue Basin; NB: Nias Basin; NMB: Northern Mentawai Basin; SMB: Southern Mentawai Basin; JB: Java Basin; LB: Lombok Basin; SS: Savu Sea. The four southern basins are characterized by seasonal upwelling during boreal summer.

G. ruber s.l., *G. sacculifer*, *P. obliquiloculata*, and *N. dutertrei*, about 40–50 tests of *G. bulloides*, and about 10–15 tests of *G. tumida*. Samples were cleaned applying a modified method originally proposed by *Barker et al.* [2003] consisting of five water washes and two methanol washes followed by two oxidation steps with 1% NaOH-buffered H_2O_2 , and a weak acid leach with 0.001M QD HNO_3 . Samples were then dissolved into 0.075M QD HNO_3 and centrifuged for 10 min. at 6000 rpm, transferred into test tubes and diluted. Mg/Ca ratios were measured using a Perkin Elmer Optima 3300 R Inductively Coupled Plasma Optical Emission Spectrophotometer (ICP-OES) equipped with an auto sampler and an ultrasonic nebulizer U-5000 AT (Cetac Technologies Inc.) housed at the Faculty of Geosciences, University of Bremen. The Mg/Ca values are reported as $mmol\ mol^{-1}$. Instrumental precision was determined using an external, in-house standard ($Mg/Ca = 2.92\ mmol\ mol^{-1}$), which was run after every fifth sample. Relative standard deviation was 0.01 $mmol\ mol^{-1}$ (0.34%) for the external standard and $\sim 0.48\%$ for the ECRM 752–1 standard [*Greaves et al.*, 2008]. Replicate measurements on 36 samples revealed an average standard deviation of 0.11 $mmol\ mol^{-1}$. Fe/Ca, Mn/Ca, and Al/Ca ratios were determined in conjunction with Mg/Ca as clay contamination and post-depositional Mn-rich carbonate coatings can exert a significant control on Mg/Ca ratios [*Rosenthal et al.*, 2000]. We exclude this possibility as these ratios were small in all samples ($< 0.1\ mmol\ mol^{-1}$ for Mn/Ca and Fe/Ca, and not detectable, i.e., $< 0\ mmol\ mol^{-1}$, for Al/Ca).

3.3.2. Calcification Depths

[13] To estimate the calcification depths for all species, the measured $\delta^{18}O$ values of planktic foraminifera were compared to the expected equilibrium $\delta^{18}O$ of calcite at different water depths. Seawater $\delta^{18}O$ ($\delta^{18}O_{SW}$) was calculated for six different water depths at 0 m, 20 m, 50 m, 75 m, 100 m, and 150 m from the WOA 05 salinity data [*Antonov et al.*, 2006] using the salinity- $\delta^{18}O_{SW}$ relationship of *Morimoto et al.* [2002] for the Western Pacific Warm Pool:

$$\delta^{18}O_{SW}(SMOW) = -14.3 + 0.42\ \text{salinity} \quad (1)$$

[14] The Standard Mean Ocean Water (SMOW) values were converted to the PDB scale by subtracting 0.27‰. In a next step, species-specific $\delta^{18}O$ -temperature equations were used to calculate the equilibrium $\delta^{18}O$ of calcite (Table 3). *Bemis et al.* [1998] demonstrated that linear and quadratic equations provide equally good fits at warm ocean temperatures. Therefore, we only used $\delta^{18}O$ -temperature equations with a linear relationship (Table 3). The water depth in which the expected equilibrium $\delta^{18}O$ of calcite matches the measured $\delta^{18}O$ values of planktic foraminifera is assumed to approximate the mean calcification depth of each species.

[15] Several studies have shown that physiological (vital) effects influence the shell isotopic composition of planktic foraminifera and result in shell $\delta^{18}O$ disequilibrium with the ambient water, ranging between -1% and 0.5% [see e.g., *Niebler et al.*, 1999, and references therein]. Correcting the

Table 2. Radiocarbon Dating on Selected Surface Sediments From the Eastern Tropical Indian Ocean^a

Surface Sample [0–1 cm]	Lab-ID	Species	¹⁴ C Age [years]	±Error [years]	1Sigma (68%) Cal. Age ± Error [years BP, BP = 1950 AD] CALIB 6.0 (Marine09)
SO189-11MC (SMB)	UCI-78814	<i>G. sacculifer</i>	1280	15	828 ± 40
SO189-60MC (NB)	UCI-78815	<i>G. sacculifer</i>	625	15	273 ± 15
SO189-64MC (NB)	UCI-78816	<i>G. sacculifer</i>	1185	15	715 ± 25
SO189-87MC (SB)	UCI-78817	<i>G. sacculifer</i>	435	15	>1950 AD
SO189-97MC (SB)	UCI-78818	<i>G. sacculifer</i>	-55	15	>1950 AD
GeoB 10008-4 (NMB)	UCI-78819	<i>G. sacculifer</i>	215	15	>1950 AD
GeoB 10010-1 (NMB)	UCI-78820	<i>G. sacculifer</i>	-295	50	>1950 AD
GeoB 10016-2 (NB)	UCI-78821	<i>G. sacculifer</i>	640	20	281 ± 20
GeoB 10022-6 (NMB)	UCI-78822	<i>G. sacculifer</i>	-335	15	>1950 AD
GeoB 10026-2 (NMB)	UCI-78823	<i>G. sacculifer</i>	-180	15	>1950 AD
GeoB 10041-3 (JB)	UCI-78824	<i>G. sacculifer</i>	-325	15	>1950 AD
GeoB 10049-5 (JB)	UCI-78825	<i>G. sacculifer</i>	-230	15	>1950 AD
GeoB 10058-1 (JB)	UCI-78826	<i>G. sacculifer</i>	-295	15	>1950 AD
GeoB 10063-5 (LB)	UCI-78827	<i>G. sacculifer</i>	-350	15	>1950 AD
GeoB 10065-9 (LB)	OS-65991	mixed planktonic	>modern		>1950 AD
GeoB 10069-4 (SS)	OS-65992	mixed planktonic	>modern		>1950 AD

^aRadiocarbon ages are converted to calendar ages using Marine 09 calibration [Hughen et al., 2004] of CALIB 6.0 software. Samples were measured at the Keck Carbon Cycle Accelerator Mass Spectrometry Laboratory, University of California, Irvine (UCI), and at the National Ocean Sciences Accelerator Mass Spectrometry Facility, Woods Hole (OS), respectively. SB: Simeulue Basin; NB: Nias Basin; NMB: Northern Mentawai Basin; SMB: Southern Mentawai Basin; JB: Java Basin; LB: Lombok Basin; SS: Savu Sea. Note that radiocarbon ages are converted to calendar ages assuming no deviation from the global ocean carbon reservoir of 400 years.

$\delta^{18}\text{O}$ values for the species-specific disequilibrium effects would accordingly result in shallower or deeper apparent calcification depths of each species. However, most of the existing $\delta^{18}\text{O}$ -temperature equations have not considered the shell $\delta^{18}\text{O}$ disequilibrium. Moreover, results from field and culture studies on the vital effect of planktic foraminifera differ significantly, or even contradict in terms of the sign (positive or negative disequilibrium) of the vital effect [see Niebler et al., 1999, and references therein]. For instance, the proposed $\delta^{18}\text{O}$ disequilibrium for *G. bulloides* ranges between -0.5‰ [Spero and Lea, 1996] and $+0.5\text{‰}$ [Ganssen, 1983]. Since no data on seawater $\delta^{18}\text{O}$ exist from the study area, we use in the following the measured shell $\delta^{18}\text{O}$ of planktic foraminifera for calculating their apparent calcification depth, and later discuss possible vital effects on the inferred habitat depths.

3.4. Alkenone Analyses

[16] For alkenone analysis, samples were extracted with a Dionex Accelerated Solvent Extractor 200 in three cycles using dichloromethane as eluent. Extracts were dried under a stream of nitrogen, saponified with 0.5 ml 1-propanolic KOH (5%) for 24h at 20°C followed by a solid phase clean-up using silica gel columns to remove the KOH. These purified extracts were analyzed by gas chromatography with a HP-6890 instrument equipped with a HP PTV Inlet on a DB-1 capillary column (30 m*0.25 mm i.d.; film thickness 0.25 μm) coupled to a flame ionization detector. Samples were injected splitless in dichloromethane using a cool injection program with solvent venting. Hydrogen was the carrier gas at a flow rate of 0.9 ml min⁻¹. A temperature program of 2 min. isothermal at 56°C, 56–150°C at 24°C min⁻¹, 150–320°C at 4.7°C min⁻¹ and 10 min isothermal was used, giving a good separation of all major compounds. Alkenones were identified by retention times. Quantification was performed relative to external calibration with n-C₃₆ alkane. The reproducibility of SST was better than 0.006

U₃₇^{K'} units or 0.2°C. The ketone unsaturation index U₃₇^{K'} was converted to temperature according to Conte et al. [2006]:

$$T(^{\circ}\text{C}) = -0.957 + 54.3(U_{37}^{K'}) - 52.9(U_{37}^{K'})^2 + 28.3(U_{37}^{K'})^3 \quad (2)$$

3.5. Auxiliary Data

[17] The 1° by 1° grid data of the World Ocean Atlas 2005 (WOA 05, <http://www.nodc.noaa.gov>) were used to estimate annual mean and boreal summer temperatures [Locarnini et al., 2006] and salinity [Antonov et al., 2006], and to calculate seawater $\delta^{18}\text{O}$ at the studied sites (section 3.3.2). Annual mean and boreal summer vertical distributions of nitrate [Garcia et al., 2006] have been used as indicators for

Table 3. Comparison of Commonly Used Species-Specific Foraminiferal Temperature: The $\delta^{18}\text{O}$ Relationship Using a Linear Approximation

Reference	Source	T (°C) = a + b ($\delta^{18}\text{O}_C - \delta^{18}\text{O}_{\text{sw}}$)	
		a	b
Shackleton [1974]	<i>Uvigerina sp.</i>	16.9	-4.0
Bouvier-Soumagnac and Duplessy [1985]	<i>N. dutertrei</i>	10.5	-6.58
Bemis et al. [1998]	<i>G. bulloides</i>	12.6	-5.07
	<i>O. universa</i>	14.9	-4.80
Peeters [2000]	<i>G. bulloides</i>	14.2	-4.81
Multiza et al. [2003]	<i>G. ruber</i>	14.2	-4.44
	<i>G. sacculifer</i>	14.91	-4.35
	<i>G. bulloides</i>	14.62	-4.70
Spero et al. [2003]	<i>G. sacculifer</i>	12.0	-5.67
Farmer et al. [2007]	<i>G. ruber</i>	15.4	-4.78
	<i>G. sacculifer</i>	16.2	-4.94
	<i>N. dutertrei</i>	14.6	-5.09
	<i>P. obliquiloculata</i>	16.8	-5.22
	<i>G. tumida</i>	13.1	-4.95

nutrient availability. However, most of the 1° by 1° grids of the WOA 05 were from the open ocean further offshore and thus, did not capture the nearshore upper water column dynamics at the sampling sites. Another limitation of using WOA 05 data is that several surface sample sites, particularly samples from the SB, NB, and NMB, are situated within only one WOA 05 grid box. These restrictions hamper a detailed, site-specific study, especially in the case of seawater $\delta^{18}\text{O}$ calculation and the habitat depth estimation of planktic foraminifera. On the other hand, local effects such as bioturbation and different ages of the surface samples might bias the geochemical proxies and their relation to the modern hydrographic data. We therefore calculated also mean values for each basin in order to reduce local effects and deviation of the geochemical proxies from the WOA 05 data. In summary, the insufficient coverage of the study area by the WOA 05 data sets remains the main limitation for a precise calibration of the surface data to the modern hydrographic data.

4. Results

4.1. Radiocarbon Ages

[18] Assuming that surface samples with an intact fluffy layer represent modern conditions, only multicore top samples without a surface fluffy layer were selected for radiocarbon measurements on *G. sacculifer* in order to estimate the oldest possible ages of the surface samples in the study area. Nonetheless, the results reveal modern ages for the selected surface samples (Table 2). Notable are the three surface samples from the Nias Basin (NB) that show older ages than in the other basins. We attribute this to lower sedimentation rates on the outer slope (GeoB 10016-2), stronger bioturbation on the shelf (SO189-64MC, 69 m water depth), and possibly a combination of both processes (SO189-60MC). Therefore, we infer that other surface samples from the NB still represent present-day conditions. Low sedimentation rates and stronger bioturbation might also be responsible for the oldest measured age of ~800 years reported from the southern Mentawai Basin (SO189-11MC).

4.2. Shell $\delta^{18}\text{O}$ -Derived Calcification Depths

4.2.1. *G. ruber* Sensu Stricto and Sensu Lato

[19] Average shell $\delta^{18}\text{O}$ values of *G. ruber* s.s. are indistinguishable in the SB, NB, and the NMB (around -3.15‰), and in the JB, LB, and the SS (around -2.9‰, Table 1), respectively. Average value in the SMB (~-3.0‰) lies between those for the other basins. Average shell $\delta^{18}\text{O}$ values of *G. ruber* s.l. are very similar to those of *G. ruber* s.s. in the SB, NB, NMB, and the SMB, but on average 0.14‰ lighter in the JB, LB, and the SS (Table 1).

[20] Calcification depths inferred from *G. ruber* s.s. and *G. ruber* s.l. shell $\delta^{18}\text{O}$ suggest comparable habitat depths for both species regardless of the equation used (Figure 3). Average calcification depths of *G. ruber* s.s. are slightly deeper in the SB and NB, where a thick mixed-layer exists throughout the year, and slightly shallower in the upwelling areas of JB, LB and SS (Figures 3a–3d). In contrast, average calcification depth of *G. ruber* s.l. does not change in different basins (Figures 3e–3h). However, the range of the estimated calcification depths from the individual measurements is larger than that of *G. ruber* s.s., with three samples appearing as outliers, showing values above -2.4‰ (GeoB 10050-1) and below -3.7‰ (SO189-89MC and 97MC).

4.2.2. *G. sacculifer* and *G. bulloides*

[21] Average shell $\delta^{18}\text{O}$ values of *G. sacculifer* are similar in the SB, NB, NMB, and the SMB (around -2.75‰), and between -2.5‰ and -2.6‰ in the JB, LB, and the SS (Table 1). Average $\delta^{18}\text{O}$ values of *G. bulloides* are lighter in the non-upwelling environments of the SB, NB, and the NMB (between -3.05‰ and -3.09‰), and heavier in the upwelling areas of the SMB, JB, LB, and the SS (between -2.5‰ and -2.8‰, Table 1). Shell $\delta^{18}\text{O}$ -derived average calcification depth of *G. sacculifer* varies between 20 m and 75 m, depending on the $\delta^{18}\text{O}$:temperature equation applied (Figures 4a–4d). Shallowest calcification depths between 20 m and 50 m are obtained when using the *Farmer et al.* [2007] equation (Figure 4b), while the *Mulitza et al.* [2003] equation suggests deepest calcification depths of about 75 m for *G. sacculifer* (Figure 4c). The species-specific equation proposed by *Spero et al.* [2003] suggests calcification depths between 50 m and 75 m (Figure 4a), while the equation proposed by *Shackleton* [1974] implies slightly shallower calcification depths closer to the 50 m isoline (Figure 4d). In general, calcification depths of *G. sacculifer* do not change significantly in different basins, except for *Farmer et al.* [2007] and *Shackleton* [1974] equations that suggest a slightly deeper calcification depth in the Simeulue and Nias Basins, similar to the inferred calcification depths for *G. ruber* s.s. (Figures 3a–3d).

[22] For *G. bulloides*, estimation of the expected equilibrium $\delta^{18}\text{O}$ of calcite is based on boreal summer temperature and salinity data of WOA 05 (solid lines in Figure 4e–4h). This inference is based on the analyses of a sediment trap time series beneath the upwelling area off S Java by *Mohtadi et al.* [2009] showing that ~90% of the total flux of *G. bulloides* larger than 250 μm occurs during the boreal summer season. Hence, the strong seasonality of *G. bulloides* needs to be considered when estimating its calcification depth in the upwelling areas of SMB, JB, LB, and the SS. In the other basins, characterized by non-upwelling environments

Figure 3. Shell $\delta^{18}\text{O}$ -based calcification temperatures and habitat depth estimates for (a–d) *G. ruber* s.s. and (e–h) *G. ruber* s.l. in the tropical eastern Indian Ocean. Solid lines indicate the expected upper water column $\delta^{18}\text{O}$ of calcite in the study area, averaged for different depths (0–75 m) using the WOA 05 annual mean temperature [*Locarnini et al.*, 2006] and salinity [*Antonov et al.*, 2006] data, salinity: $\delta^{18}\text{O}_{\text{SW}}$ relationship of *Morimoto et al.* [2002], and species-specific $\delta^{18}\text{O}$:temperature equations for each panel. Gray dots are measured shell $\delta^{18}\text{O}$ values of the entire surface samples, black dots are average values for each basin. Average values represent the following basins from left to right: Simeulue Basin, Nias Basin, Northern Mentawai Basin, Southern Mentawai Basin, Java Basin, Lombok Basin, and the Savu Sea. Shaded envelopes indicate the error range calculated from the combined analytical error of the measured and the expected $\delta^{18}\text{O}$ of calcite, and the standard deviation of their difference.

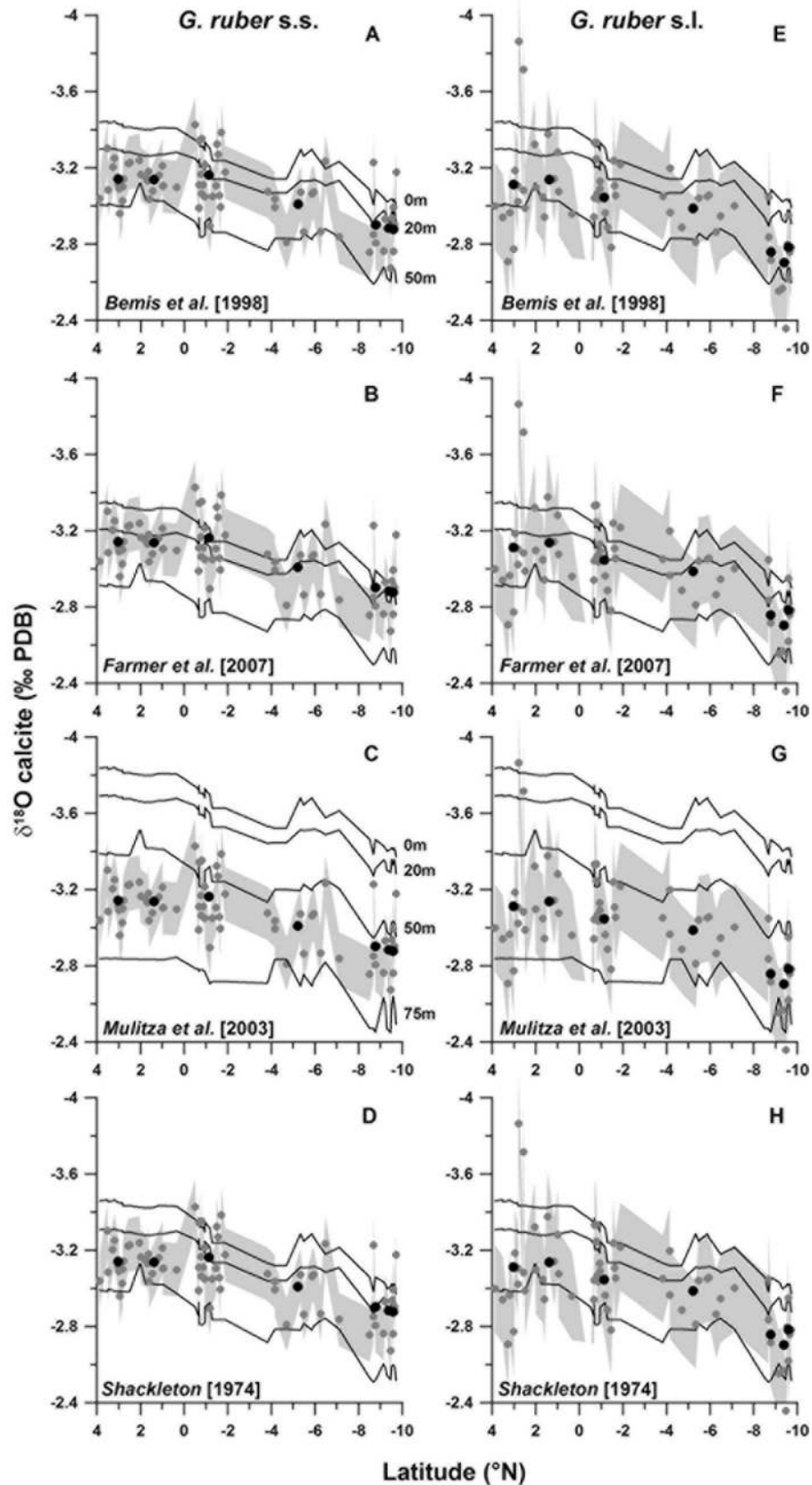


Figure 3

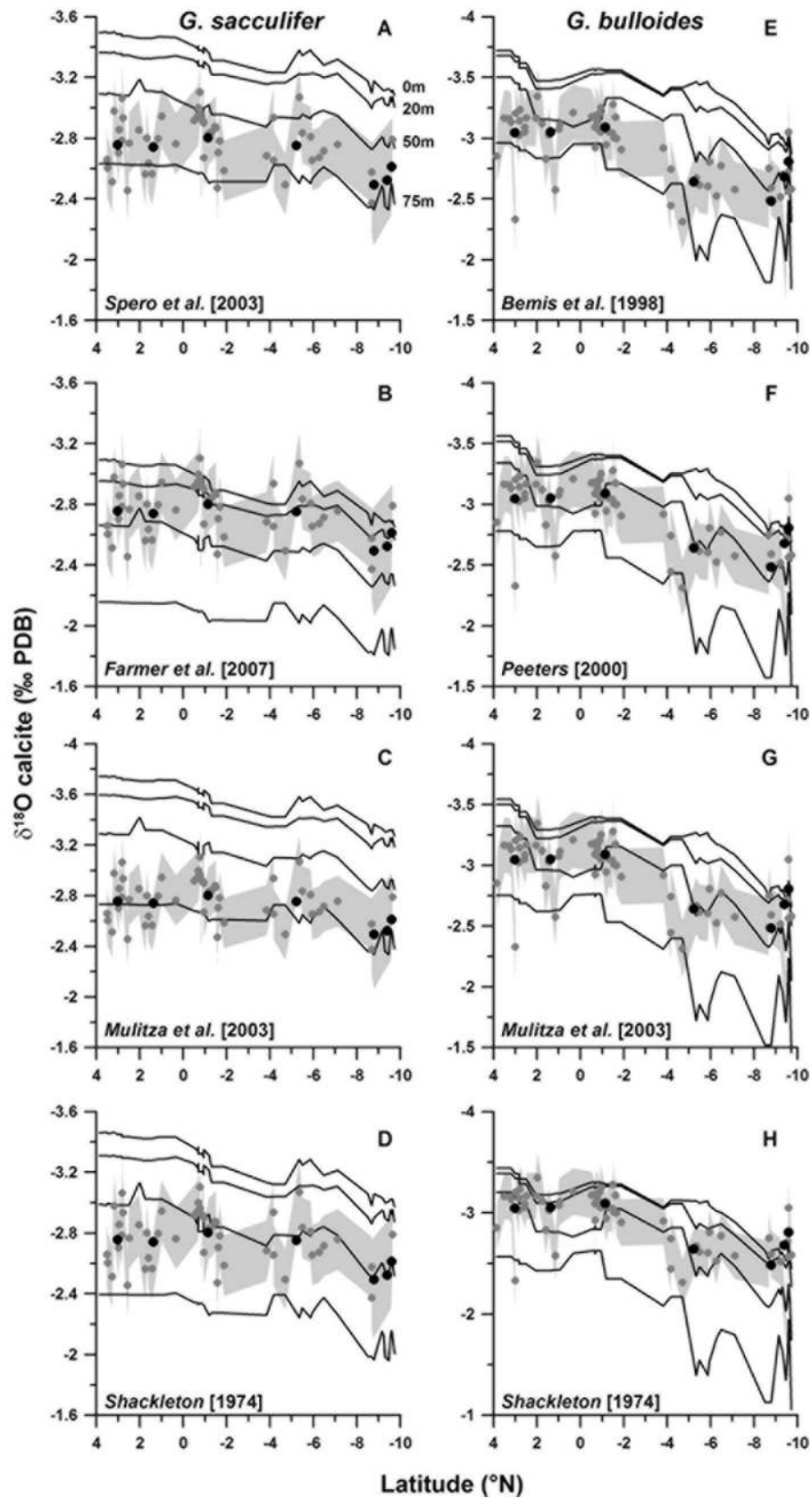


Figure 4. Shell $\delta^{18}\text{O}$ -based calcification temperatures and habitat depth estimates for (a–d) *G. sacculifer* and (e–h) *G. bulloides* in the tropical eastern Indian Ocean. Shaded envelopes, solid lines and dots are as in Figure 3, except the solid lines in *G. bulloides* panels (Figures 4e–4h) representing boreal summer values. Note the different scales for each panel.

(SB, NB, NMB), annual mean and summer temperatures and salinities do not differ significantly (Figures S1 and S2), enabling us to apply summer values for *G. bulloides* in the entire study area.

[23] Shell $\delta^{18}\text{O}$ -derived average calcification depths of *G. bulloides* range around the 50 m isoline and tend to decrease in the JB, LB and the SS, regardless of the equation used (Figures 4e–4h). Calcification depths are slightly deeper when applying the Bemis *et al.* [1998] equation (Figure 4e), and shallower when using the Shackleton [1974] equation (Figure 4h). The two equations proposed by Mulitza *et al.* [2003] and Peeters [2000] are almost identical and therefore, the resulting calcification depths for *G. bulloides* are similar for different basins (Figures 4f and 4g).

4.2.3. *N. dutertrei*, *P. obliquiloculata*, and *G. tumida*

[24] Average shell $\delta^{18}\text{O}$ values of *N. dutertrei* range between about -2.0‰ and -2.1‰ in the SB, NB, NMB, and the SMB, and between -1.6‰ and -1.8‰ in the JB, LB, and the SS, respectively (Table 1). Average shell $\delta^{18}\text{O}$ values of *P. obliquiloculata* range between -2.3‰ and -2.4‰ in the SB, NB, and the NMB, and between about -1.4‰ and -1.7‰ in the JB, LB, and the SS, respectively (Table 1). Average value for the SMB lies in between, at about -1.9‰ . Average shell $\delta^{18}\text{O}$ values of *G. tumida* differ significantly for different basins, ranging between about -0.4‰ and 0.5‰ , which might be due to the small amount of data on this species, or different amounts of secondary calcite. *G. tumida* is absent in the surface samples from the SB, and only present in one (two) surface sample(s) from the NB (NMB) suggesting the preference of this species for the nutrient-rich subsurface waters beneath the upwelling areas in the southern part of the study area (Table 1).

[25] Estimation of the expected equilibrium $\delta^{18}\text{O}$ of calcite for *P. obliquiloculata*, *N. dutertrei*, and *G. tumida* is based on boreal summer temperature and salinity data of WOA 05, due to their preferred seasonality similar to *G. bulloides* [Mohtadi *et al.*, 2009]. Shell $\delta^{18}\text{O}$ -derived average calcification depths of *N. dutertrei* show insignificant changes by applying different equations and vary between 75 m and 100 m in the study area (Figures 5a–5c). Calcification depths are slightly shallower in the SMB (~ 75 m) and somewhat deeper in the SB. Estimated average calcification depths for *P. obliquiloculata* are similar to those of *N. dutertrei* in the SB, NB, NMB, and the SMB, and slightly deeper in the JB, LB, and the SS when applying the Shackleton [1974] and Bouvier-Soumagnac and Duplessy [1985] equations (Figures 5d and 5f). In general, the Farmer *et al.* [2007] equation implies shallower average calcification depths between 50 m and 75 m in the SB, NB, NMB, and the SMB, and deeper calcification depths below 75 m in the JB, LB, and the SS (Figure 5e). Shell $\delta^{18}\text{O}$ -derived average calcification depths of *G. tumida* lie between 100 m and 150 m when applying the $\delta^{18}\text{O}$:temperature equation of Shackleton [1974], and at ~ 150 m when using the species-specific equation of Farmer *et al.* [2007] (Figures 5g and 5h).

4.3. Shell Mg/Ca Values and Alkenone-Derived SST

[26] Of all the planktic foraminifera measurements, shell Mg/Ca values were unrealistically high in one sample from the SB (SO189-112MC), and two samples from the NB (SO189-64MC and SO189-65MC, see Table 1). These

samples were retrieved from shallow waters at or shallower than 80 m close to the shore, offshore two (small) river mouths. Since there are no anomalous values in the $\delta^{18}\text{O}$, Fe/Ca or Al/Ca of these samples, the high Mg/Ca values cannot easily be explained by increased freshwater input, inorganic calcite precipitation, dissolution or insufficient cleaning of the samples. Likewise, examination of the planktic foraminiferal tests does not show any significant diagenetic overprint. To this end, we do not have any sound explanation of these anomalous values and exclude these samples from calculation of average Mg/Ca in these basins.

[27] Average shell Mg/Ca of *G. ruber* s.s. and *G. ruber* s.l. shows higher values in the SB, NB, NMB, and the SMB, and decreases significantly in the JB, LB, and the SS (Table 1). Highest average Mg/Ca value occurs in the NB (~ 6 mmol mol $^{-1}$). Average Mg/Ca values of *G. ruber* s.s. and *G. ruber* s.l. are similar in different basins, except for the LB, where *G. ruber* s.l. value is on average 0.45 mmol mol $^{-1}$ lower than the *G. ruber* s.s. value.

[28] Likewise, average shell Mg/Ca of *G. sacculifer* shows higher values in the SB, NB, NMB, and the SMB, and decreases significantly in the JB, LB, and the SS (Table 1). Highest average Mg/Ca value occurs in the NB (~ 4.7 mmol mol $^{-1}$). Average shell Mg/Ca of *G. bulloides* is high in the SB, NB, and the NMB (>7.6 mmol mol $^{-1}$) and decreases considerably to values < 6.4 mmol mol $^{-1}$ in the upwelling areas of the SMB, JB, and the LB (Table 1). The SS is represented by only one sample with a relatively high Mg/Ca value of ~ 7 mmol mol $^{-1}$.

[29] Shell Mg/Ca values of *P. obliquiloculata* and *N. dutertrei* show a decrease from higher values in the non-upwelling areas of the SB, NB, and the NMB (≥ 3 mmol mol $^{-1}$) to lower values in the upwelling areas of the SMB, JB, LB, and the SS (≤ 2.75 mmol mol $^{-1}$, Table 1). This pattern can also be observed for *G. tumida*, although only three samples exist from the non-upwelling basins.

[30] Average alkenone-based SST estimates for different basins do not vary considerably ($28^\circ\text{C} \pm 0.4^\circ\text{C}$, Table 1), with lowest (highest) average SST of $\sim 27.8^\circ\text{C}$ (28.4°C) recorded in the JB (NB). Moreover, SST estimates are remarkably similar within the different basins and do not deviate by more than 0.5°C from the observed average SST for each basin, except for one sample (SO189-31MC) recording the lowest SST (27°C) in the study area.

5. Discussion

5.1. Mg/Ca Versus $\delta^{18}\text{O}$ -Derived Calcification Depths/Temperatures

[31] In the following, calcification depths are considered to be between 20 m and 50 m for *G. ruber* s.s., at 50 m for *G. ruber* s.l., *G. sacculifer*, and *G. bulloides*, at 75 m for *N. dutertrei* and *P. obliquiloculata*, and at 100 m for *G. tumida*. The deeper habitat inferred for various species by applying the plankton tow and culture based equations of Mulitza *et al.* [2003, Figures 3c, 3g, and 4c] and Spero *et al.* [2003, Figure 4a] might be a result of generally higher $\delta^{18}\text{O}$ values recorded by surface samples. Our results agree with previous studies suggesting habitat depths of *G. ruber* s.s., *G. ruber* s.l., *G. sacculifer* and *G. bulloides* within the mixed-layer, and of *N. dutertrei*, *P. obliquiloculata*, and *G. tumida* within the thermocline [e.g., Fairbanks and

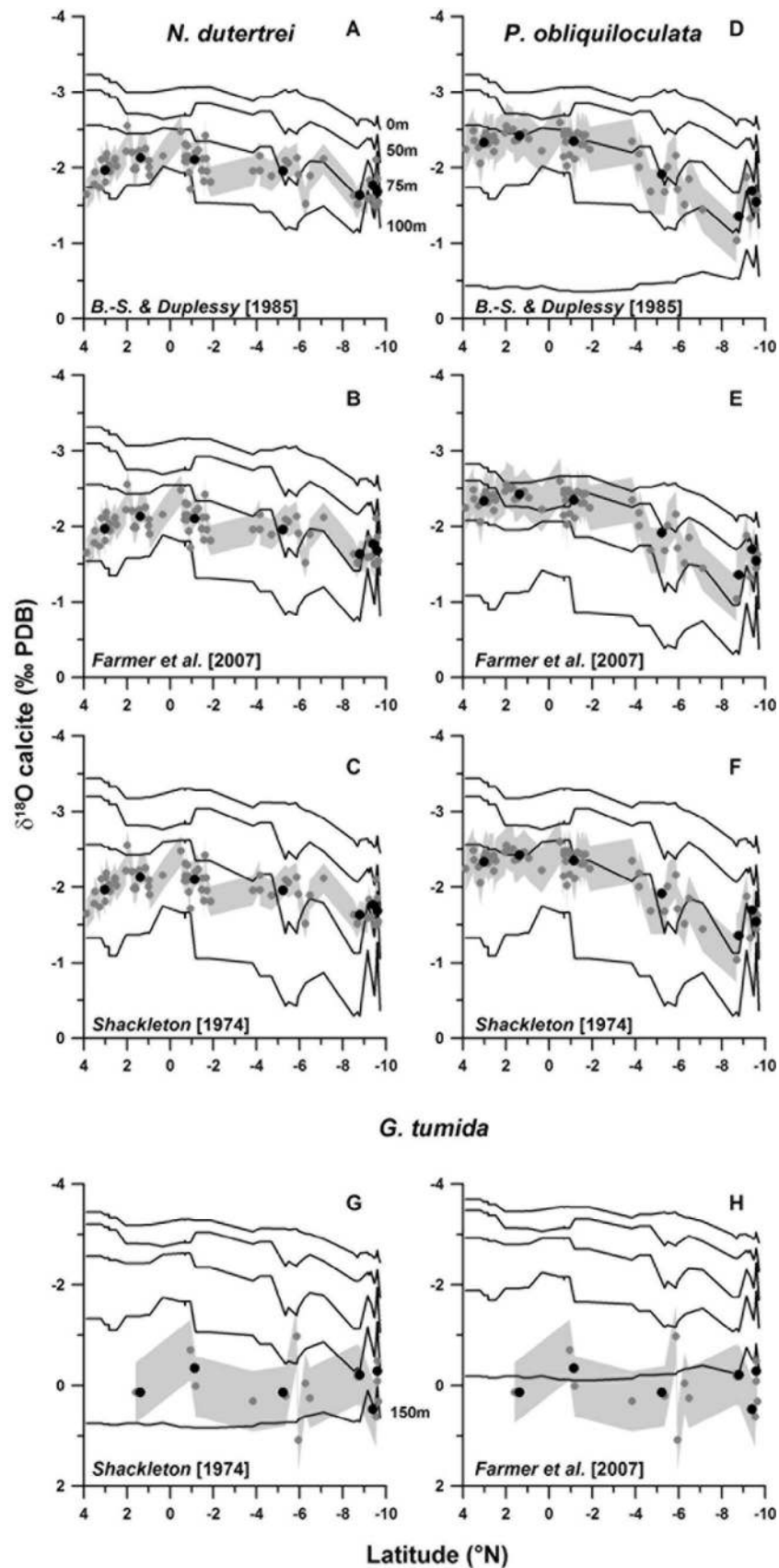


Figure 5. Shell $\delta^{18}\text{O}$ -based calcification temperatures and habitat depth estimates for (a–c) *N. dutertrei*, (d–f) *P. obliquiloculata*, and (g and h) *G. tumida* in the tropical eastern Indian Ocean. Shaded envelopes, solid lines and dots are as in Figure 3, except that the solid lines represent boreal summer values.

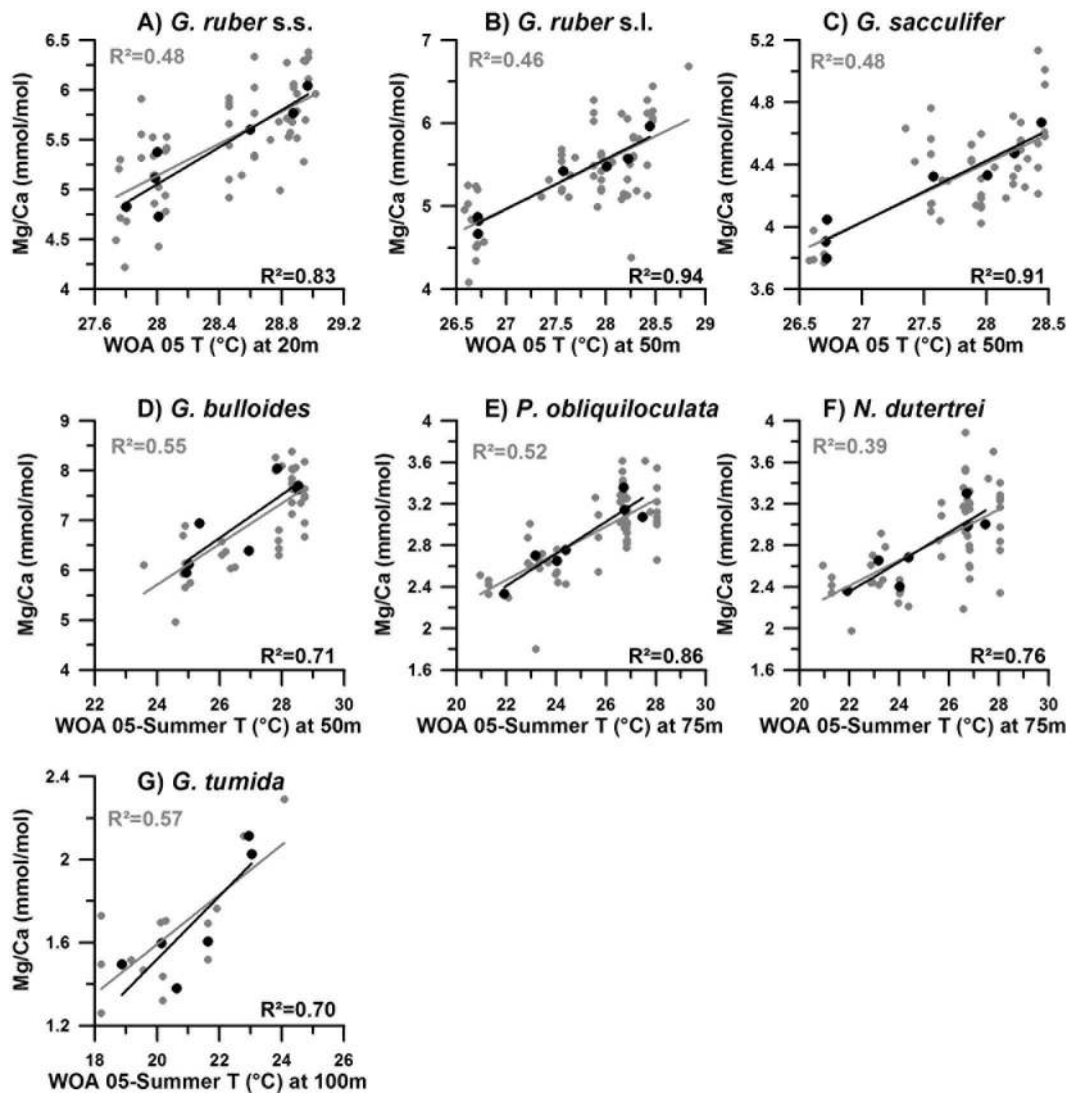


Figure 6. Relationship between shell Mg/Ca ratio and the calcification temperatures of various planktic foraminiferal species in the study area. Gray: the entire data sets; black: average values for each basin. Temperatures are taken from the WOA 05 data set, at depths that correspond to the estimated calcification depth of each species (see Figures 3–5). Mg/Ca values are plotted versus annual mean temperatures at (a) 20 m and (b and c) 50 m, and versus summer temperatures at (d) 50 m, (e and f) 75 m, and (g) 100 m, respectively.

Wiebe, 1980; Fairbanks *et al.*, 1982; Peeters, 2000; Field, 2004; Kuroyanagi and Kawahata, 2004; Farmer *et al.*, 2007].

[32] Shell Mg/Ca ratios of all species show a significant, linear relationship to temperatures from the WOA 05 data at depths that correspond to the estimated calcification depth of each species (Figure 6). Since salinities in the upper water column of the study area are lower than 35 psu (Figure S2), a positive salinity effect on shell Mg/Ca ratios [Arbuszewski *et al.*, 2010] can be neglected. The large scatter in the data (gray dots in Figure 6) might be due to the (a) relatively narrow temperature range for each species, (b) different amount of encrustation [e.g., Schmidt and Mulitza, 2002], (c) different genetic types [e.g., Darling *et al.*, 2003], (d) different ages of the surface samples, (e) a limited coverage of local hydrography by the WOA 05 data, both spatially and

vertically, and (f) interannual variability in local hydrography caused by ENSO or IOD that cannot be resolved in our core top study. Therefore, it is not surprising that average values for each basin (black dots in Figure 6) show improved correlation coefficients (R^2) at or greater than 0.7 for all species by reducing the above mentioned uncertainties, although a larger R^2 is also a result of fewer data points. It should be noted that for *G. ruber* s.s., *G. bulloides*, and *G. tumida*, correlation coefficients between Mg/Ca and WOA 05 temperatures are also significant at adjacent depths due to the fact that calcification of planktic foraminifera and hence, incorporation of Mg in their calcite shell, occur within a depth range, rather than at a fixed depth (see Figure S3 and discussion below).

[33] In face of a small temperature range for each species in the study area, Mg/Ca-temperature relationship could be

expressed as a linear function (Figure 6). However, previous studies showed that shell Mg/Ca in planktic foraminifera increases exponentially with temperature [e.g., *Elderfield and Ganssen*, 2000; *Dekens et al.*, 2002; *Anand et al.*, 2003; *McConnell and Thunell*, 2005; *Cléroux et al.*, 2008; *Regenberg et al.*, 2009]. The species-specific Mg/Ca-temperature exponential relationships from this study (Figure 7, black dots and lines) fit within the range of previously published equations, although they occur at the warm end of the existing calibrations (Figure 7, gray lines). The only exception is the Mg/Ca-temperature relationship of *G. tumida* that shows a comparable slope with the *Rickaby and Halloran* [2005] equation, but a different intercept (Figure 7g), which might be due to the much shallower inferred habitat of this species in the study area, or to the different size-fractions used in the other studies (300–355 μm [*Rickaby and Halloran*, 2005] and 355–400 μm [*Regenberg et al.*, 2009]). We therefore calculated a regional Mg/Ca-temperature calibration for *G. tumida*:

$$\text{Mg/Ca} = 0.41 \exp(0.068 * T) \quad (3)$$

[34] For all the other species, the existing Mg/Ca-temperature calibrations can be used for the eastern equatorial Indian Ocean. For *G. ruber* s.s., *G. ruber* s.l., and *G. sacculifer* the regional calibrations fit best to the equations proposed by *Anand et al.* [2003] for the same size-fractions used in this study ($\text{Mg/Ca} = 0.34 \exp(0.102 * T)$ and $\text{Mg/Ca} = 0.38 \exp(0.09 * T)$ for *G. ruber*, Figures 7a and 7b, $\text{Mg/Ca} = 0.347 \exp(0.09 * T)$ for *G. sacculifer* without sac, Figure 7c). Regional Mg/Ca-temperature relationship of *G. bulloides* is best explained by the equation of *Elderfield and Ganssen* [2000] ($\text{Mg/Ca} = 0.81 \exp(0.081 * T)$). For *N. dutertrei*, the regional calibration fits best to the equation proposed by *Regenberg et al.* [2009] ($\text{Mg/Ca} = 0.65 \exp(0.065 * T)$, Figure 7e). Finally, regional Mg/Ca-temperature relationship of *P. obliquiloculata* matches best to the equation for deep-dwelling foraminifera proposed by *Cléroux et al.* [2008] ($\text{Mg/Ca} = 0.78 \exp(0.052 * T)$, Figure 7f).

[35] As mentioned above, $\delta^{18}\text{O}$ -disequilibrium effect on the calculation of apparent calcification depths of planktic foraminifera might also impact their Mg/Ca-temperature relationships. Correcting the measured $\delta^{18}\text{O}$ values for a range of disequilibrium values suggested for *G. sacculifer*, *N. dutertrei*, and *P. obliquiloculata* [see *Niebler et al.*, 1999; *Regenberg et al.*, 2009, and references therein] would increase their calcification depths, but decrease significantly (0.1–0.3) the correlation coefficient of their Mg/Ca-temperature calibration. This finding, in accordance with results of *Regenberg et al.* [2009], justifies the application of the disequilibrium-uncorrected calibration for these species. There is no disequilibrium effect reported for *G. ruber* s.l. and

G. tumida. Correcting the measured $\delta^{18}\text{O}$ values for a range of disequilibrium values suggested for *G. ruber* s.s. (*G. bulloides*) would increase (increase or decrease) its calcification depth, while the correlation coefficient of the Mg/Ca-temperature calibration remains high (Figure S3). This might be a result of a bimodal vertical distribution of these species in the water column [e.g., *Peeters et al.*, 2002; *Kuroyanagi and Kawahata*, 2004] due to different hydrologic conditions in the study area (upwelling and non-upwelling areas), or different genotypes of these species in sediment samples [e.g., *Darling et al.*, 2003]. In spite of these possible effects, the water depths with a significant Mg/Ca-temperature correlation coefficient for these species match the range of their $\delta^{18}\text{O}$ -derived calcification depths (20–50 m for *G. ruber* and 20–75 m for *G. bulloides*, Figures 3a–3d and 4e–4h), justifying the application of the disequilibrium-uncorrected calibration for these species. Nonetheless, future field studies on both planktic foraminifera and local hydrography are imperative in order to assess the $\delta^{18}\text{O}$ -disequilibrium effect of planktic foraminifera, and to better calculate their apparent calcification depth.

5.2. Use of Temperature Proxies for Reconstructing Upper Water Column Hydrology

[36] Conversion of the Mg/Ca values to temperatures using the species-specific calibrations that match the apparent calcification depths of planktic foraminifera reveals a relatively consistent picture in the study area (Figure 8). In the non-upwelling basins of SB, NB, and the NMB, Mg/Ca temperatures of the surface-dwelling species *G. ruber* s.s., *G. ruber* s.l., *G. sacculifer*, and *G. bulloides* reflect mixed-layer temperatures between 0 m and 50 m at or above 28°C, whereby the succession of these species reflects decreasing (increasing) temperatures (habitat depths). Alkenone-based temperatures appear to match Mg/Ca temperatures of *G. sacculifer*, also reflecting mixed-layer temperatures in these basins. Since there are no significant seasonal changes in the mixed-layer conditions of these basins, our sediment surface data suggest that these proxies can be used to reconstruct past mixed-layer conditions of the eastern equatorial Indian Ocean, from ~4° N to ~4° S. Mg/Ca temperatures of *P. obliquiloculata* and *N. dutertrei* reflect upper thermocline temperatures at ~70 m (26°C–28°C) and ~100 m (23°C–25°C), respectively. Despite the sparse data of *G. tumida* from these basins, it appears that this species records nearly the same temperatures as *N. dutertrei*, at ~100 m water depth (23°C–24°C). Accordingly, these three species can be used to reconstruct the upper thermocline conditions in the eastern Indian Ocean, from ~4° N to ~4° S.

[37] In the southern basins of SMB, JB, LB, and the SS, calibrated temperatures based on shell Mg/Ca- and calcification depth of planktic foraminifera suggest that *G. ruber* s.s., *G. ruber* s.l., and *G. sacculifer* reflect annual mean mixed-layer temperatures of 27°–28°C between 0 m and

Figure 7. (a–g) Shell Mg/Ca ratio to $\delta^{18}\text{O}$ -derived calcification temperature of various planktic foraminiferal species from this study (black dots and lines) compared to other species-specific temperature calibrations (gray lines). Note that the exponential fits in Figures 7a–7d (Figures 7e–7g) are extrapolated to lower (higher) temperatures than in the original publications, and that all Mg/Ca values from this study are at the warm end of the published calibrations. 1a: calibration with calculated exponential value; 1b: calibration with assumed exponential value; 8a: species-specific calibration; 8b: deep-dwelling (multispecies) calibration.

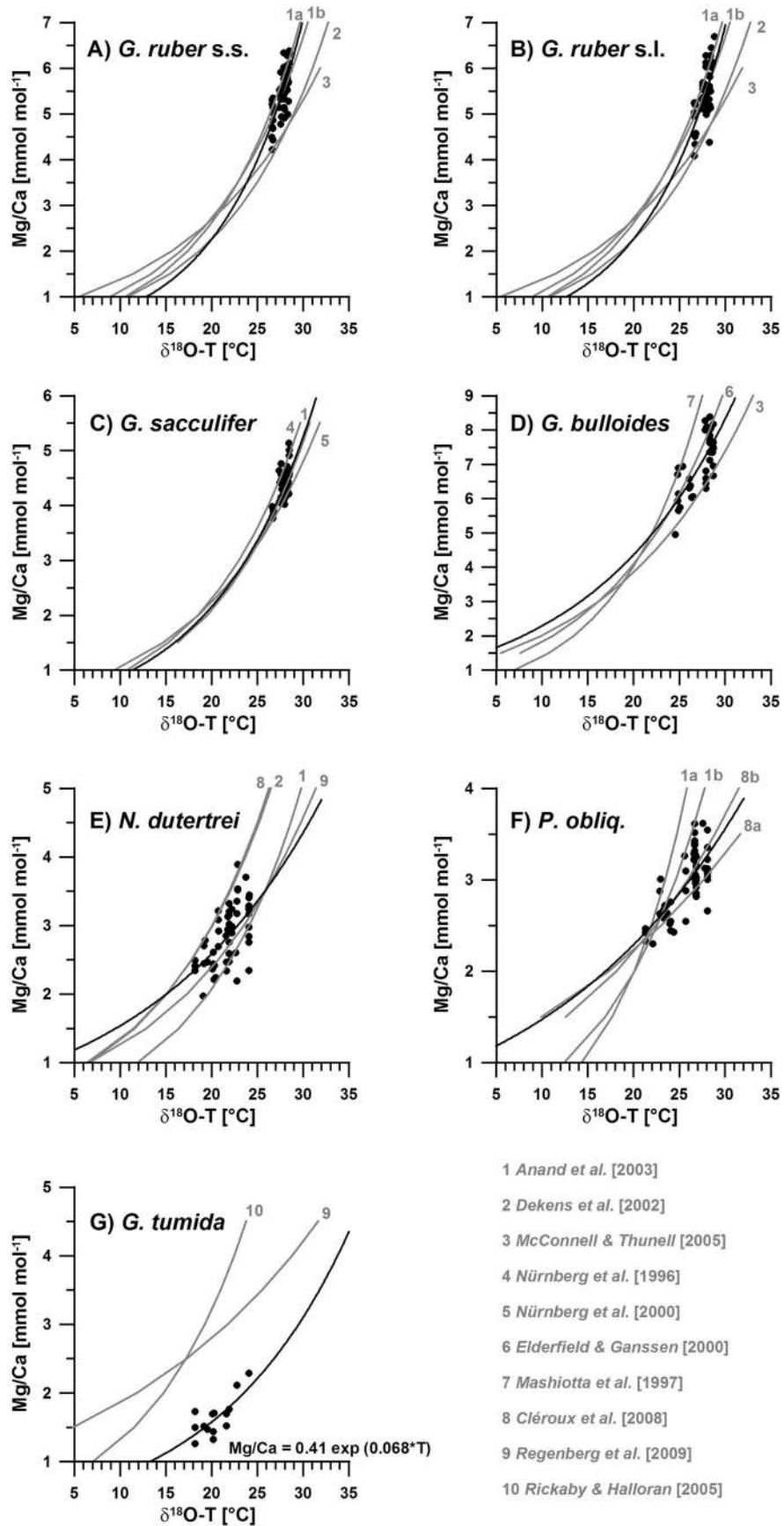


Figure 7

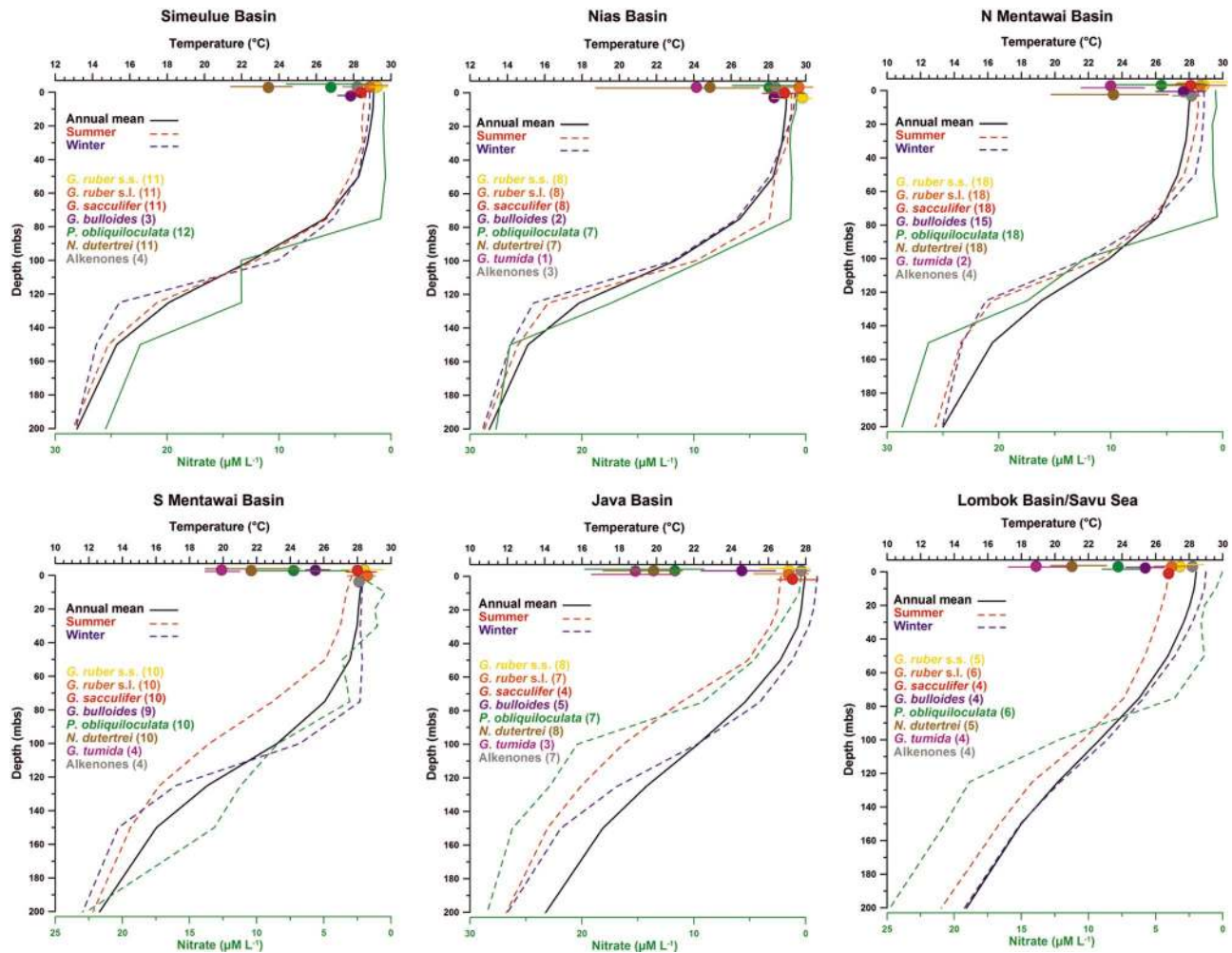


Figure 8. Temperature profiles and nitrate concentration of the upper 200 m in the different basins of the study area (WOA 05, [Garcia *et al.*, 2006; Locarnini *et al.*, 2006]). Solid black lines: annual mean temperatures; dashed red lines: summer temperatures; dashed blue lines: winter temperatures; solid green lines: annual mean nitrate concentrations in micromole per liter ($\mu\text{M L}^{-1}$); dashed green lines: boreal summer nitrate concentrations; mbs: meter below surface. Color dots indicate average temperatures, calculated from shell Mg/Ca ratio of planktic foraminiferal species, and alkenone unsaturation index. Alkenone temperatures are calculated after Conte *et al.* [2006]. Mg/Ca-temperatures are calculated using species-specific equations of Anand *et al.* [2003] for *G. ruber* s.s., *G. ruber* s.l., and *G. sacculifer*, Cléroux *et al.* [2008] for *P. obliquiloculata*, Regenberg *et al.* [2009] for *N. dutertrei*, Elderfield and Ganssen [2000] for *G. bulloides*, and own equation for *G. tumida*. Color lines denote the range of the calculated temperatures in different samples; values in brackets refer to the number of the observed surface sediment samples. The bottom (top) three panels refer to basins that are (not) affected by seasonal upwelling.

50 m (Figure 8). Alkenone-based temperatures remain at $\sim 28^\circ\text{C}$, suggesting that this proxy can only be used to reconstruct annual mean SST south of 4°S . It should be noted that alkenone temperatures in the study are at the upper limit of this proxy [e.g., Conte *et al.*, 1998] and might underestimate the growth temperature of the alkenone-producing coccolithophores, particularly during boreal winter when SST is highest. Downcore studies from the study area show a strong seasonal signal in the alkenone-based SST estimates during glacial periods when SSTs lie within the temperature response of alkenone unsaturation [Lückge *et al.*, 2009; Mohtadi *et al.*, 2010]. Hence, it is well possible that alke-

nes in surface sediments simply record the limit of their temperature response, which corresponds to modern annual mean SST in the study area but masks their seasonality.

[38] Shell Mg/Ca of *G. bulloides* seems to record summer temperatures ranging between 20 m and 75 m, with average values reflecting temperatures of $24^\circ\text{--}26^\circ\text{C}$ at the base of the mixed-layer during boreal summer, at ~ 50 m. Mg/Ca temperatures of *P. obliquiloculata* and *N. dutertrei* reflect upper thermocline temperatures at ~ 60 m ($21\text{--}24^\circ\text{C}$) and ~ 80 m ($20\text{--}22^\circ\text{C}$) during boreal summer, respectively, thus recording a slightly shallower depth than in the non-upwelling basins. *G. tumida* appears to record temperatures between

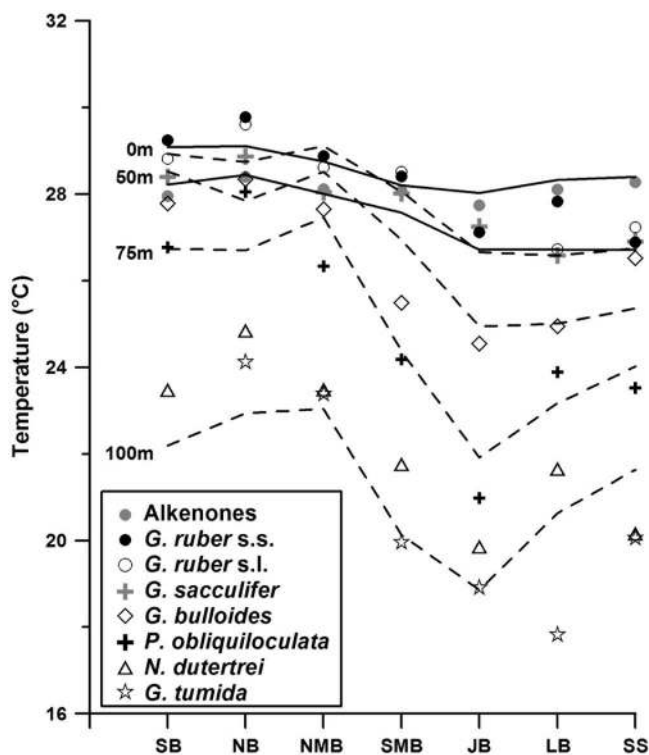


Figure 9. Alkenones- and Shell Mg/Ca-derived temperatures averaged for each basin. Calibrations are as in Figure 8. Solid lines (dashed lines) indicate annual mean (summer) upper water column temperatures in the different basins at 0 m and 50 m (0–100 m); averaged for each depth at each basin using the WOA 05 database [Locarnini et al., 2006]. SB: Simeulue Basin; NB: Nias Basin; NMB: Northern Mentawai Basin; SMB: Southern Mentawai Basin; JB: Java Basin; LB: Lombok Basin; SS: Savu Sea. The latter four basins are characterized by monsoon-induced seasonal upwelling.

19°C and 20°C at ~100 m depth during boreal summer, at a similar depth as in the northern, non-upwelling basins. In general, these results are in good agreement with the inferred habitat depths from a sediment trap time series in the upwelling area off southern Java [Mohtadi et al., 2009] suggesting habitat depths of 0–30 m for *G. ruber*, 60–80 m for *P. obliquiloculata*, and 60–90 m for *N. dutertrei*.

[39] Comparison of the Mg/Ca data with the nitrate concentrations in the upper water column further supports previous findings that *G. bulloides* and the thermocline-dwelling species are also bound to the nutricline (Figure 8) [e.g., Fairbanks and Wiebe, 1980]. In particular, *N. dutertrei* and *G. tumida* are apparently independent of the thermocline temperature, but dwell at depths characterized by a nitrate concentration of 10–15 $\mu\text{M L}^{-1}$. In general, it appears that temperature changes at the upper thermocline in different basins do not significantly affect the water depth that defines the shell Mg/Ca ratio. Rather, the thermocline species appear to track the top of the nutricline that mostly corresponds to the same water depths (70–100 m), regardless of the absolute temperature values at these depths (Figure 9). For instance, Mg/Ca-derived temperatures of *P. obliquiloculata* track the summer 75 m isotherm that

represents the upper thermocline (nutricline), although temperatures at this depth vary between ~22°C and ~28°C in the study area (Figure 9). To this end, our data do not allow a conclusion on whether temperature or nutrient is decisive for the planktic foraminiferal calcification depths. This enterprise requires extensive regional field studies in the future.

[40] In search of a robust proxy for reconstructing past changes in the thermocline depth and/or changes in the upper water column stratification/mixing, our data show that both the pattern and the magnitude of the temperature difference between *G. ruber* s.s. (or *G. ruber* s.l.) and *P. obliquiloculata* in surface sediments of the study area track the difference between temperatures at 20 m and at 75 m from the WOA 05 data (Figure 10a). Likewise, temperature difference between *G. ruber* s.s. and *N. dutertrei* effectively follows the difference between 20 m and 100 m, suggesting that these differences can be used to track past changes in the water column structure in the eastern equatorial Indian Ocean (Figure 10a). Temperature difference between *G. bulloides* and *P. obliquiloculata* (*N. dutertrei*) shows the same pattern as observed for the difference between summer temperatures at 50 m and at 75 m (100 m, Figure 10b). However, the calculated difference between *G. bulloides* and *N. dutertrei* temperatures lies between the WOA 05 difference of 50–75 m and 50–100 m, which might be due to the Mg/Ca temperatures of *N. dutertrei* that reflect temperatures between 75 m and 100 m water depth (Figure 9).

[41] Temperature difference between *G. sacculifer* and thermocline-dwelling species follows the same pattern as observed between *G. bulloides* (*G. ruber* s.s.) and thermocline species in the study area (Figure 10c). However, the temperature difference slightly overshoots the expected difference in the JB, and is less than expected in the NB. Finally, the difference between alkenone and *G. bulloides* temperatures appears to be an appropriate measure of the seasonality of the mixed-layer, as it depicts the difference between annual mean SST and summer temperatures at 50 m (Figure 10c). However, our results do not resolve the applicability of this proxy for reconstructing seasonality in geological past (see discussion above).

[42] Our study cannot resolve the interannual effect of ENSO and IOD on different proxies at different sites. Since interpretation of past changes in the thermal structure of the water column or seasonality might vary at different sites, a careful understanding of various processes contributing to variance in these proxies is essential. For this purpose, long-duration field studies (sediment trap and plankton tow) are required that record the gradient and the entire range of climate variability, and their effect on the here introduced proxies.

6. Conclusions

[43] Evaluation of geochemical proxies in modern surface sediments from the eastern equatorial Indian Ocean allows the following conclusions:

[44] Shell $\delta^{18}\text{O}$ and Mg/Ca ratio of planktic foraminifera reveal that *G. ruber* s.s. (~20 m), *G. ruber* s.l. (20–50 m), and *G. sacculifer* (~50 m) reflect annual mean conditions within the mixed-layer. Calcification depths of these species vary slightly in the different basins of the study area that differ with regard to seasonal changes in temperature and

productivity, yet still remain within the mixed-layer. *G. bulloides* (~50 m), *P. obliquiloculata* (~75 m), *N. dutertrei* (75–100 m), and *G. tumida* (~100 m) record boreal summer conditions in nutrient-rich environments at the base of the mixed-layer, and at the upper thermocline, respectively. Our findings from the coastal tropical eastern Indian Ocean are in good agreement with core top results from open ocean

environments [e.g., Elderfield and Ganssen, 2000; Cléroux et al., 2008; Regenberg et al., 2009] and sediment-trap data [e.g., Anand et al., 2003; Mohtadi et al., 2009]. Alkenone-based temperatures record annual mean SST, possibly due to the temperature limitation of this proxy at ~28°C that corresponds to annual mean SST in the study area.

[45] Shell Mg/Ca ratios of planktic foraminifera significantly correlate to the temperature at their $\delta^{18}\text{O}$ -derived calcification depths. Mg/Ca-temperatures occur at the warm end of the existing species-specific calibrations for planktic foraminifera, except *G. tumida* that apparently inhabits warmer waters in the study area and therefore, requires a regional Mg/Ca-temperature calibration ($\text{Mg/Ca} = 0.41 \exp(0.068 * T)$). Conversion of shell Mg/Ca to temperature can be best assessed by applying the equations proposed by Anand et al. [2003] for *G. ruber* s.s., *G. ruber* s.l., and *G. sacculifer*, by Elderfield and Ganssen [2000] for *G. bulloides*, by Cléroux et al. [2008] for *P. obliquiloculata*, and by Regenberg et al. [2009] for *N. dutertrei*.

[46] Temperature difference between the mixed-layer species (*G. ruber* s.s., *G. ruber* s.l., and *G. bulloides*) and the thermocline-species (*P. obliquiloculata*, *N. dutertrei*) in the study area depicts the temperature difference between the mixed-layer (20–50 m) and the upper thermocline (75–100 m) during summer, and can be used to reconstruct past changes in the stratification of the upper water column. Temperature difference between alkenones and *G. bulloides* tracks the present-day difference between annual mean and boreal summer temperatures of the mixed-layer.

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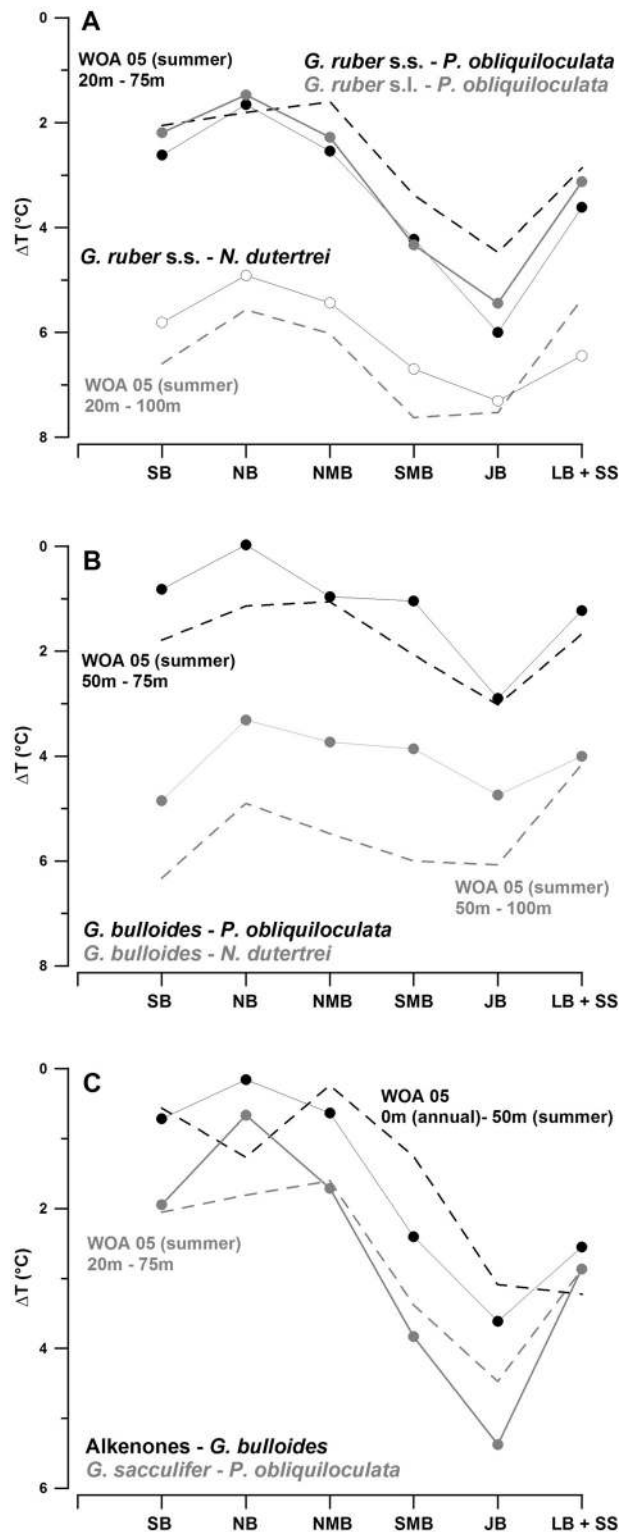


Figure 10. (a) Mg/Ca-based average temperature difference (ΔT) between *G. ruber* s.s. and *P. obliquiloculata* (black dots), *G. ruber* s.l. and *P. obliquiloculata* (gray dots), and between *G. ruber* s.s. and *N. dutertrei* (black circles) in different basins. Dashed lines represent temperature difference between 20 m and 75 m (black), and 20 m and 100 m (gray) from the WOA 05 database for boreal summer. (b) Mg/Ca-based average temperature difference (ΔT) between *G. bulloides* and *P. obliquiloculata* (black dots), and between *G. bulloides* and *N. dutertrei* (gray dots) in different basins. Dashed lines represent temperature difference between 50 m and 75 m (black), and 50 m and 100 m (gray) from the WOA 05 database for boreal summer. (c) Difference between average alkenone and *G. bulloides* temperatures (black dots), and between *G. sacculifer* and *P. obliquiloculata* (gray dots) temperatures in different basins. Dashed lines represent temperature difference between annual mean SST (0 m) and summer temperatures at 50 m (black), and between 20 m and 75 m during boreal summer (gray) from the WOA 05 database. Abbreviations for the different basins are as in Figure 9.

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