

# Mc/Ca, Sr/Ca and stable isotope from planktonic froaminifera T. sacculifer: testing a multi-proxy approach for inferring paleo-temperature and paleo-salinity [preprint]

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#### 2 Mg/Ca, Sr/Ca AND STABLE ISOTOPE FROM PLANKTONIC FORAMINIFERA T. SACCULIFER: TESTING A MULTI-PROXY APPROACH FOR INFERRING 3 4 PALEO-TEMPERATURE AND PALEO-SALINITY

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#### ABSTRACT

24 Over the last decades, sea surface temperature (SST) reconstructions based on the Mg/Ca of 25 for aminiferal calcite have frequently been used in combination with the  $\delta_{18}O$  signal from the 26 same material, to provide estimates of  $\delta_{18}$ O of the water ( $\delta_{18}$ O<sub>w</sub>), a proxy for global ice volume 27 and sea surface salinity (SSS). However, because of error propagation from one step to the next, 28 better calibrations are required to increase accuracy and robustness of existing isotope and 29 element to temperature proxy-relationships. Towards that goal, we determined Mg/Ca, Sr/Ca 30 and the oxygen isotopic composition of Trilobatus sacculifer (previously referenced as 31 Globigerinoides sacculifer), collected from surface waters (0-10m), along a North-South 32 transect in the eastern basin of the tropical/subtropical Atlantic Ocean. We established a new paleo-temperature calibration based on Mg/Ca, and on the combination of Mg/Ca and Sr/Ca. 33 Subsequently, a sensitivity analysis was performed in which, one, two, or three different 34 35 equations were considered. Results indicate that foraminiferal Mg/Ca allow for an accurate 36 reconstruction of surface water temperature. Combining equations,  $\delta_{18}$ Ow can be reconstructed 37 with a precision of about  $\pm 0.5\%$ . However, the best possible salinity reconstruction based on 38 locally calibrated equations, only allowed reconstruction with an uncertainty of  $\pm 2.49$ . This was 39 confirmed by a Monte Carlo simulation, applied to test successive reconstructions in an 'ideal 40 case', where explanatory variables are known. This simulation shows that from a pure statistical 41 point of view, successive reconstructions involving Mg/Ca and  $\delta_{18}$ Oc preclude salinity





42 reconstruction with a precision better than  $\pm 1.69$  and hardly better than  $\pm 2.65$ , due to error 43 propagation. Nevertheless, a direct linear fit to reconstruct salinity based on the same measured 44 variables (Mg/Ca and  $\delta^{18}Oc$ ) was established. This direct reconstruction of salinity lead to a 45 much better estimation of salinity ( $\pm 0.26$ ) than the successive reconstructions.

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#### I. INTRODUCTION

50 Since Emiliani's pioneering work (1954), oxygen isotope compositions recorded in fossil 51 foraminiferal shells became a major tool to reconstruct past sea surface temperature. After 52 Shackleton's seminal studies (1967, 1968 and 1974), it became clear that part of the signal 53 reflected glacial-interglacial changes in continental ice volume and hence sea level variations. 54 The oxygen isotope composition of foraminiferal calcite ( $\delta_{18}O_c$ ) is thus controlled by the 55 temperature of calcification (Urey, 1947; Epstein et al., 1953) but also by the oxygen isotope composition of seawater ( $\delta_{18}O_w$ )). The relative contribution of these two factors cannot be 56 57 deconvolved without an independent measure of the temperature at the time of calcification 58 such as e.g. Mg/Ca (e.g. Nürnberg et al., 1996; Rosenthal et al., 1997; Rathburn and DeDeckker, 59 1997; Hastings et al., 1998; Lea et al., 1999; Lear et al., 2002; Toyofuku et al., 2000; Anand et 60 al., 2003, al., Kisakurek et al., 2008; Duenas-Bohorquez et al., 2009, 2011; Honisch et al., 2013; 61 Kontakiotis et al., 2016; Jentzen et al., 2018). The sea surface temperature (SST) reconstructed from Mg/Ca of foraminiferal calcite has, therefore, increasingly been used in combination with 62 63 the  $\delta_{18}$  or signal measured on the same material, to estimate  $\delta_{18}$  ow, global ice volume and to 64 infer past sea surface salinity (SSS) (e.g. Rohling 2000, Elderfield and Ganssen, 2000; Schmidt 65 et al., 2004; Weldeab et al., 2005; 2007). These studies also showed that, because of error propagation, inaccuracies in the different proxies combined for the reconstruction of past sea 66 water  $\delta_{18}O$  and salinity obstruct meaningful interpretations. Hence, while there is an 67 68 understandable desire to apply empirical proxy-relationships down-core, additional calibrations 69 appear necessary to make reconstructions more robust. Calibrations using foraminifera sampled 70 from surface seawater (0-10m deep), provide the best possibility to avoid most of the artefacts 71 usually seen when using specimen from core tops or culture experiments for calibration 72 purposes. Here, we report a calibration based on *Globigerinoides sacculifer*, which should now 73 and will be referenced in this manuscript as Trilobatus sacculifer (Spezzaferri et al., 2015), 74 from the Atlantic Ocean. Mg and Sr concentrations were measured on individual specimens 75 with Laser Ablation-Inductively Coupled Plasma-Mass Spectrometry (LA-ICP-MS), while the





76 oxygen isotope composition of the same tests as used for the elemental analyses was 77 subsequently measured by Isotope ratio Mass Spectrometry (IRMS). Environmental parameters 78 (temperature: T, salinity: S, dissolved inorganic carbon: DIC and alkalinity: ALK) but also the 79 isotopic composition (O18w) of the seawater the foraminifers were growing in, were measured. 80 The primary objectives of this study are to test and improve the calibration of (1) both the 81 Mg/Ca and oxygen isotope paleothermometer for T. sacculifer, a paleoceanographically 82 relevant species; (2) evaluate the agreement between observed and predicted  $\delta_{18}O_w$  and (3) test 83 potential for SSS reconstructions of the Atlantic Ocean. Our results indicate that the best 84 possible salinity reconstruction based on locally calibrated equations from the present study, 85 only allowed reconstruction with an uncertainty of  $\pm 2.49$ . Such an uncertainty does not allow for viable (paleo)salinity data. This is subsequently confirmed by a Monte Carlo simulation, 86 87 applied to test successive reconstructions in an 'ideal case', where explanatory variables are 88 known. This simulation shows that from a pure statistical point of view, successive 89 reconstructions involving Mg/Ca and  $\delta_{18}$ Oc preclude salinity reconstruction with a precision 90 better than  $\pm 1.69$  and hardly better than  $\pm 2.65$ , due to error propagation. Nevertheless, a direct linear fit based on the same measured variables (Mg/Ca and  $\delta^{18}Oc$ ), and leading to much better 91 92 estimation of salinity  $(\pm 0.26)$ , could be established.

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- 94 95

#### 2. MATERIAL AND METHODS

#### 96 2.1. Collection procedure

97 Foraminifera were collected between October and November 2005, on board of the research 98 vessel Polarstern (ANT XXIII/1) during a meridional transect of the Atlantic Ocean 99 (Bremerhaven/Germany - Cape Town/South of Africa; Fig. 1a). Foraminifera were 100 continuously collected from a depth of ca. 10 m using the ship's membrane pump (3 m<sub>3</sub>/h). The 101 water flowed into a plankton net (125 µm) that was fixed in a 1000 L plastic tank with an 102 overflow (Fig 1b). Every eight hours, the plankton accumulated in the net was collected. 103 Temperature and salinity of surface seawaters were continuously recorded by the ship's 104 systems, and discrete water samples were collected for later analyses of total ALK, DIC and 105  $\delta_{18}O_w$  (see Tab. 1). Plankton and water samples were poisoned with buffered formaldehyde 106 solution (20%) and HgCl2 (1.5 ml with 70gL-1 HgCl2 for 1 L samples), respectively. In total, 107 more than seventy plankton samples were collected during the transect, covering a large range 108 in both temperature and salinity. Specimens of T. sacculifer from thirteen selected stations, 109 selected as to maximize temperature and salinity ranges, were picked and prepared for analyses.





110 Salinity, temperature, DIC, ALK and  $\delta_{18}$ Ow data reported in this paper represent 111 October/November values for the selected stations.

112

#### 113 **2.2. Description of species**

114 Trilobatus sacculifer is a spinose species with endosymbiotic dinoflagellates inhabiting the 115 shallow (0-80 m deep) tropical and subtropical regions of the world oceans. This species 116 displays a large tolerance to temperature (14-32°C) and salinity (24-47) (Hemleben et al., 1989; 117 Bijma et al., 1990). Based on differences in the shape of the last chamber of adult specimens, 118 various morphotypes can be distinguished. Among others the last chamber can be smaller than 119 the penultimate chamber, in which case it is called kummerform (kf). This species shows an 120 ontogenetic depth migration and predominantly reproduces at depth around full moon (Bijma 121 and Hemleben, 1993). Just prior to reproduction a secondary calcite layer, called gametogenic 122 (GAM) calcite is added (Bé et al., 1982; Bijma and Hemleben, 1993; Bijma et al., 1994). 123 Juveniles (<100µm) ascend in the water column and reach the surface after less than approximately 2 weeks. Pre-adult stages then slowly descend within 9-10 days to the 124 125 reproductive depth. In our samples (collected between 0 and 10 m depth), T. sacculifer 126 specimens have not yet added the Mg-enriched gametogenic calcite, which generally occurs 127 deeper in the water column just prior to reproduction. Therefore, only the trilobus morphotype 128 without GAM calcite is considered (230µm to 500µm) in this study, which limits the 129 environmental, ontogenetic and physiological variability between samples and should be taken 130 into account when compared to other calibrations based on core top and/or sediment trap 131 collected specimens.

132

#### 133 2.3. Seawater analysis

134 The DIC and ALK analyses of the sea water were carried out at the Leibniz Institute of Marine 135 Sciences at the Christian-Albrechts University of Kiel, (IFM-GEOMAR), Germany. Analyses 136 were performed by extraction and subsequent coulometric titration of evolved CO<sub>2</sub> for DIC 137 (Johnson et al., 1993), and by open-cell potentiometric seawater titration for ALK (Mintrop et 138 al., 2000). Precision / accuracy of DIC and ALK measurements are 1 µmol kg-1 / 2 µmol kg-1 139 and 1.5 µmol kg-1 / 3 µmol kg-1, respectively. Accuracy of both DIC and ALK was assured by 140 the analyses of certified reference material (CRM) provided by Andrew Dickson from Scripps 141 Institution of Oceanography, La Jolla, USA. Measurements of  $\delta_{18}O_w$  were carried out at the 142 Faculty of Geosciences, Utrecht University, Netherlands. Samples were measured using a 143 GasBench II - Delta plus XP combination. Results were corrected for drift with an in-house





- 144 standard (RMW) and are reported on V-SMOW scale, with a precision of 0.1‰ and accuracy
- 145 verified against NBS 19 of 0.2‰ respectively. For reconstruction calculations δ18Ow data were
- 146 corrected to the PDB scale by subtracting 0.27‰ (*Hut*, 1987).
- 147

#### 148 2.4. Carbonate analysis

149 2.4.1. Foraminiferal sample preparation

150 Under a binocular microscope, maximum test diameter of each specimen was measured and 151 individual tests were weighed on a microbalance (METTLER TOLEDO, precision  $\pm 0.1 \mu g$ ). 152 Since the foraminifera were never in contact with sediments, the rigorous cleaning procedure 153 required for specimens collected from sediment cores, was not necessary. Prior to analysis the tests were cleaned following a simplified cleaning procedure: All specimens were soaked for 154 155 30 min in a 3-7% NaOCl solution (Gaffey and Brönniman, 1993). A stereomicroscope was used 156 during cleaning and specimens were removed from the reagent directly after complete 157 bleaching. The samples were immediately and thoroughly rinsed with deionised water to ensure complete removal of the reagent. After cleaning, specimens were inspected with scanning 158 159 electron microscopy and showed no visible signs of dissolution. This cleaning procedure 160 preserves original shell thickness and thus maximises data acquisition during laser ablation. 161 Foraminifera were fixed on a double-sided adhesive tape and mounted on plastic stubs for LA-162 ICP-MS analyses.

163

#### 164 2.4.2. Elemental composition analysis

165 For each station, 5-8 specimens were analysed. Their last chambers were ablated using an 166 Excimer 193 nm deep ultraviolet laser (Lambda Physik) with GeoLas 200Q optics (Reichart et 167 al, 2003) creating 80 µm diameter craters. Pulse repetition rate was set at 6 Hz, with an energy 168 density at the sample surface of 1 J/cm2. The ablated material was transported on a continuous 169 helium flow into the argon plasma of a quadrupole ICP-MS instrument (Micromass Platform) 170 and analysed with respect to time. Ablation of calcite requires ultraviolet wavelengths as an 171 uncontrolled disruption would result from higher wavelengths. By using a collision and reaction 172 cell spectral interferences on the minor isotopes of Ca (42Ca, 43Ca and 44Ca) were reduced and 173 interferences of clusters like 12C16O16O were prevented. Analyses were calibrated against NIST 174 (U.S. National Institute of Standards and Technology) 610 glass using the concentration data 175 of Jochum et al. (2011) with Ca as internal standard. For Ca quantification, mass 44 was used 176 while monitoring masses 42 and 43 as internal check. In the calcite, the Ca concentration was 177 set at 40%, allowing direct comparison to trace metal/Ca from traditional wet-chemical studies.





178 Mg concentrations were calculated using masses 24 and 26; Sr concentrations were calculated 179 with mass 88. One big advantage in using LA-ICP-MS measurements is that single laser pulses 180 remove only a few nanometers of material, which allows high resolution trace elements profiles 181 to be acquired (e.g. Reichart et al., 2003; Regenberg et al., 2006; Dueñas-Bohórquez et al., 182 2009, 2010, Hathorne et al., 2009; Munsel et al., 2010; Dissard et al., 2009; 2010a and b; Evans et al., 2013; 2015; Steinhardt 2014, 2015; Fehrenbacher et al., 2015; Langer et al., 2016; Koho 183 184 et al., 2015; 2017; Fontanier et al., 2017; De Nooijer et al., 2007, 2014, 2017a and b; Jentzen et 185 al., 2018, Schmitt et al., 2019; Levi et al., 2019). Element concentrations were calculated for 186 the individual ablation profiles integrating the different isotopes (glitter software). Even though 187 the use of a single or very few specimens, can be criticised when determining foraminifera 188 Mg/Ca and  $\delta_{18}$ O in order to perform paleoclimate reconstructions instead of more traditional 189 measurements, Groeneveld et al., (2019) recently demonstrated that for both proxies, single 190 specimen variability is dominated by seawater temperatures during calcification, even if the 191 presence of an ecological effect leading to site-specific seasonal and depth habitat changes is 192 also noticeable.

193

#### 194 **2.5. Stable isotope analysis**

195 The specimens used for elemental composition analyses using LA-ICP-MS were subsequently 196 carefully removed from the plastic stubs and rinsed with deionised water before measuring their 197 stable isotope composition. Depending on shell weight, 2 to 3 foraminifera were necessary to 198 obtain a minimum of 20µg of material, required for each analysis. Analyses were carried out in 199 duplicate for each station. The results, compiled in table 2, represent average measurements. 200 The analyses were carried out at the Department of Earth Sciences of Utrecht University (The 201 Netherlands), using a Kiel-III -Finnigan MAT-253 mass spectrometer combination. The  $\delta_{18}O_c$ 202 results are reported in ‰ PDB. Calibration was made with NBS-19 (precision of 0.06-0.08 ‰ 203 for sample size  $20-100 \mu g$ , accuracy better than 0.2%).

204

#### 205 2.6. Statistical analysis

Within this manuscript, all statistical analyses with regards to elemental and isotopic data, were
 carried out using the program R with default values (R Development Core Team (2019).
 3. RESULTS

211 **3.1. Elemental composition** 





212	Overall values of the Mg/Ca and Sr/Ca ratios in the tests of T. sacculifer varied from 1.78 to
213	5.86 mmol/mol (Fig. 2a) and 1.41 to 1.52 mmol/mol (fig. 2b), respectively (Tab. 2). These
214	Mg/Ca concentrations compare well with results found in literature for this species from either
215	culture experiments, plankton tow, or surface sediment, growing at the same temperatures (e.g.
216	Nürnberg et al., 1996; Anand et al. 2003, Regenberg et al., 2009, Fig. 3). Similarly, the overall
217	variation in Sr/Ca-values reported in this study is comparable to that observed in core top and
218	cultured G. ruber and T. sacculifer combined, for comparable salinity and temperature
219	conditions, (varying between 1.27 to 1.51mmol/mol; e.g. Cleroux et al., 2008; Kisakürek et al.,
220	2008; Dueñas-Bohórquez et al., 2009).
221	
222	
223	
224	The relationship between measured temperatures and both Mg/Ca and Sr/Ca ratios were
225	calculated using least square differences. Both show a good correlation with surface water
226	temperature (Fig. 2, Tab. 3). The Mg/Ca ratio increases exponentially by 8.3%/°C (best fit)
227	(Mg/Ca and Sr/Ca ratios given in mmol/mol):
228	$Mg/Ca=(0.42\pm0.13) \exp((0.083\pm0.001)*T[^{\circ}C]), R^{2}=0.86 \text{ pvalue}=2,9e-06 \text{ (equation 1)}$
229	
230	whereas Sr/Ca ratio increases linearly by 0.6%/°C (Fig. 2a and b), best fit:
231	
232 233	$Sr/Ca=(0.009\pm0.002)*T+(1.24\pm0.05), R^2=0.67 \text{ pvalue}=5.e-04$ (equation 2)
233	Combining Mg and Sr data for a non-linear multivariate regression allows improvement of the
234	correlation with temperature, best fit:
235	correlation with temperature, best fit.
230	
238	$T = -(27\pm15) + (8\pm1) \ln(Mg/Ca) + (28\pm11) Sr/Ca, pvalue Mg/Ca: 2.10^{-4}$ (equation 3)
238	$R^{2}=0.92 \text{ pvalue}= 2.e-04 \text{ (equation 3)}$
239 240	R <sup>2</sup> -0.92 pvalue= 2.e-04
240 241	Regression for the relationship between salinity and Mg/Ca ratios does not show any clear
241	correlation ( $R_2=0.09$ , p-value=0.32). This is in good agreement with previous culture
243 244	experiments studies which only report a minor sensitivity of Mg/Ca to salinity in planktonic
244	foraminifera (e.g. Dueñas-Bohórquez et al., 2009; Hönisch et al., 2013; Kisakürek et al., 2008;
245	Nürnberg et al., 1996). The correlation observed between Sr/Ca ratios and salinity (R2=0.29, p-





246	value=0.053) is better compared to that between Mg/Ca and salinity, but remains relatively
247	weak. Nevertheless, recalculated regressions of Mg/Ca, incorporating salinity, show an
248	improvement of the correlation with temperature, best fit:
249	
250	$ln(Mg/Ca[mmol/mol]) = (-5.02\pm2) + (0.09\pm0.009)*T + (0.11\pm0.05)*S,$
251	$R_2=0.91 \text{ pvalue} = 5e-06$

252

This result is in good agreement with the recent study of Gray and Evans (2019), who reported the minor Mg/Ca sensitivity of *Trilobatus sacculifer* to salinity ( $3.6 \pm 0.01\%$  increase per salinity unit) and described, based on previously published culture experiments' data (Dueñas-Bohórquez et al., 2009; Hönisch et al., 2013; Kisakürek et al., 2008; Lea et al., 1999; Nürnberg et al., 1996), a similar fit allowing to assess the sensitivity of foraminiferal Mg/Ca of *T. sacculifer* to temperature and salinity combined.

259 
$$Mg/Ca = exp(0.054(S-35) + 0.062T-0.24)$$
 RSE: 0.51 Gray and Evans (2019)

In order to compare both equations, Mg/Ca values from our study were used to reconstruct temperature and salinity using the fit established per Gray and Evans (2019), versus reconstructed temperature and salinity using our fit. The observed R<sub>2</sub> are then 0.99 and 0.48 for temperature and salinity, respectively. We can conclude, that if the equation of Evans and Gray is in perfect agreement with our equation with regards to the temperature parameter, this is not the case for salinity, which shows a strong difference between the two equations, most probably explained by the weak correlation of Mg/Ca to salinity.

#### 267 **3.2. Stable isotopes concentration**

The  $\delta_{18}O$  (PDB) values of the tests ( $\delta_{18}Oc$ ) and of the seawater ( $\delta_{18}Ow$ ) vary from -0.70 to -2.98‰ and from 0.74 to 1.25‰, respectively (Tab. 1 and 2). The relationship between temperature and the foraminiferal  $\delta_{18}O$  (expressed as a difference to the  $\delta_{18}Ow$  of the ambient seawater) was estimated with a linear least squares regression:

273 
$$T = (11.82 \pm 1.3) - (4.82 \pm 0.45)^* (\delta_{18}Oc - \delta_{18}Ow) [\%]; R^2=0.90$$
 (equation 4)

274

275 The oxygen isotope fractionation ( $\delta_{18}$ Oc -  $\delta_{18}$ Ow) shows a strong correlation with *in situ* surface 276 water temperature (linear increase of 0.17‰/°C).





# 278 **3.3.** Comparison with previously established *T. sacculifer* temperature reconstruction

#### 279 equations

As mentioned above, average juvenile and pre-adult *T. sacculifer* specimen only spend between 9 to 10 days in surface waters. Therefore, measured *in situ* temperature is representative of the calcification temperatures. This is supported by the strong correlation between measured temperature and  $\delta_{18}$ O analyses (R2=0.90, equation 4), and measured temperature vs. Mg/Ca, (R2=0.87, equation 1). Nevertheless, diurnal variations in temperatures cannot be discarded and may induce a slight offset between measured average temperature and mean calcification temperature.

287

For comparison, three Mg/Ca temperature calibrations for *T. sacculifer* were considered in this manuscript. The equation of Nürnberg et al. (1996) based on laboratory cultures, (2) the equation established by Anand et al. (2003) based on sediment trap samples and (3) the equation derived by Regenberg et al. (2009) based on surface sediment samples of the Tropical Atlantic Ocean. In each of these studies only *T. sacculifer* without SAC chamber were considered, (Tab. 3).

Similarly, in addition to equation 4 established in this study, three δ18O based paleo-temperature
equations for *T. sacculifer* were used for comparison with our data set: (1) Erez and Luz, (1983)
and, (2) Spero et al. (2003), both based on cultured specimens, and (3) Mulitza et al. (2003)
based on surface water samples (Fig. 4; Tab. 3).

298

#### 299 **3.4.** Correlation between measured δ<sub>18</sub>O/Salinity

Salinity and the oxygen isotope composition of surface seawater were measured for 23 stations located between 33°N and 27°S of the Eastern Atlantic Ocean (Tab. 4), including the thirteen stations represented in figure 1, where foraminifera were sampled. The  $\delta_{18}O_w$ -salinity relationship (equation 5) is plotted in figure 5.

304 305

$$\delta_{18}Ow = (0.194 \pm 0.04) * S - (5.8 \pm 1.5), R^2 = 0.53$$
 (equation 5)

306

For comparison, the  $\delta_{18}$ Ow-salinity relationship for the tropical Atlantic Ocean calculated by Paul et al. (1999) (from 25°S to 25°N) based on GEOSECS data, and by Regenberg et al. (2009), based on data from Schmidt 1999 (30°N–30°S), are plotted in the same figure. Temporal, geographical and depth differences in sampling, as well as analytical noise, are most probably responsible for the observed variations.





### 312

313

#### 4. DISCUSSION

#### 314 4.1. Intra-test variability

The Mg/Ca and Sr/Ca composition of foraminiferal calcium carbonate was determined using 315 316 laser ablation ICP-MS of the final (F) chamber of size-selected specimen. Eggins et al., (2003) 317 report that the Mg/Ca composition of sequentially precipitated chambers of different species 318 (including T. sacculifer) are consistent with temperature changes following habitat migration 319 towards adult life-cycle stages. As described for T. sacculifer in the Red Sea (Bijma and 320 Hemleben, 1994), juvenile specimens (<100µm) migrate to the surface, where they stay about 321 9-10 days, before descending to the reproductive depth (80m). The addition of GAM calcite 322 proceeds immediately prior to gamete release (Hamilton et al., 2008). The specimens 323 considered in this study were collected between 0 and 10 meters depth, and GAM calcite was 324 not detected. This limits the impact of variability due to migration, reduces potential ontogenic 325 vital effects responsible for inter-chamber elemental variations (Dueñas-Bohórquez, 2010) and, 326 avoids variability due to variable amounts of GAM calcite precipitated (Nürnberg et al., 1996). 327 This is confirmed by the strong correlation (R<sup>2</sup>=0.87) observed between our Mg/Ca-328 reconstructed temperature vs. measured surface temperature. In agreement with measurements 329 on specimens from culture experiments (Dueñas-Bohórquez, 2009), Mg-rich external surfaces 330 (GAM calcite) were not observed in our samples.

Because the diameter of the laser beam used in this study was 80µm, it represents a reliable mean value of elemental concentration of the last chamber wall, for every analysis of a single shell a full ablation of the wall chamber was performed (until perforation was completed). For comparison, results from traditional ICP-OES Mg/Ca analyses (Regenberg et al., 2009), electron microprobe (Nurnberg et al., 1996) and laser ablation ICP-MS (this study) are plotted in figure 3a and suggest comparable foraminiferal Mg/Ca ratios for *T. sacculifer* at similar temperatures.

338

#### **4.2. Incorporation of Sr into Mg/Ca-Temperature calibrations**

Combining Mg and Sr data to compute temperature was first suggested by Reichart et al. (2003) for the aragonitic species *Hoeglundina elegans*. It has been demonstrated that variables other than temperature, such as salinity and carbonate chemistry (possibly via their impact on growth rate) are factors influencing Sr incorporation into calcite (e.g. Lea et al., 1999, Dueñas-Bohórquez et al., 2009; Dissard et al., 2010a; Dissard et al., 2010b). The good correlation of Sr/Ca with temperature in our results (R2=0.67, p value= 5.e-04, Fig 2b), also suggests that





temperature exerts a major control on the amount of Sr incorporated into T. sacculifer' tests. 346 347 However, Sr/Ca concentration also shows a correlation with salinity (R<sub>2</sub>=0.29, p-value=0.053), 348 which is not observed for Mg ( $R_2=0.09$ , p-value=0.32). Therefore, the incorporation of Sr into 349 the Mg-T reconstruction equation might improve temperature reconstruction by accounting for 350 the impact of salinity. It has recently been suggested that the Sr incorporation in benthic 351 foraminiferal tests is affected by their Mg contents (Mewes et al., 2015; Langer et al.; 2016). 352 However, as pointed out in Mewes et al., (2015), calcite's Mg/Ca needs to be over 30-50mmol 353 in order to noticeably affect Sr partitioning. There is no obvious reason to assume that 354 planktonic foraminifera should have a different Mg/Ca threshold. Therefore, with a 355 concentration between 2 to 6 mmol/mol (Sadekov et al., 2009), the observed variation in Sr concentration in T. sacculifer' tests can be safely considered to be independent of the Mg/Ca 356 357 concentrations. Hence, other environmental parameters such as temperature, salinity and/or 358 carbonate chemistry, potentially via an impact on calcification rates, must control Sr/Ca values. 359

360 The standard deviation of measured temperatures versus reconstructed temperature was 361 calculated for each of the three Mg-temperature equations established in this study. For 362 equation (1), based on Mg/Ca only, SD= 1.37, for equation (3), based on both Mg/Ca and Sr/Ca, 363 SD=0.98, and for equation (4), based on Mg/Ca ratio and salinity, SD=1.03. Incorporation of 364 Sr into the Mg-Temperature reconstruction equation resulted in the standard deviation the closest to 1 (SD=0.98), indicating that this statistically improved reconstructions possibly by 365 366 attenuating the salinity effect as well as potentially other environmental parameters such as 367 variations in carbonate chemistry or the effect of temperature itself. Therefore, the combination 368 of Mg/Ca and Sr/Ca should be considered to improve temperature reconstructions (Tab. 3). For 369 the remainder of this discussion, and in order to compare our data with previously established 370 calibrations for T. sacculifer, the equation based on Mg/Ca alone (equation 1) will be 371 considered.

372

#### 373 **4.3** Comparison with previous *T. sacculifer* Mg/Ca-Temperature calibrations.

Mg/Ca ratios measured on *T. sacculifer* from our study show a strong correlation with measured surface water temperature (R<sub>2</sub>=0.86, p value=2.9e-06) (Fig. 2a), increasing exponentially by 8.3% per °C. The relation with temperature (equation 1) is comparable to the one published by Nürnberg et al., (1996) and within the standard error of the calibration (Fig. 3a). This implies that the temperature controlled-Mg incorporation into *T. sacculifer* tests is similar under culture conditions as it is in natural surface waters. The equation established by Duenas-Bohorquez et





al., (2010) based on *T. sacculifer* specimen from culture experiments integrates ontogenetic
(chamber stage) effects. Even though incorporating the ontogenetic impact may improve
temperature reconstructions based on Mg/Ca ratios, this is not routinely done for paleotemperature reconstruction using *T. sacculifer*. Therefore, the equation of Nürnberg et al.,
(1996) is used in our study for comparison of various reconstruction scenarios.

385 A comparable regression (similar slope) has been established for T. sacculifer from tropical 386 Atlantic and Caribbean surface sediment samples by Regenberg et al. (2009) (Fig 3a). This 387 regression predicts Mg concentrations that are about 0.15 mmol/mol higher compared to our 388 study. Because the Mg-T calibration from Regenberg et al. (2009) is based on sediment-surface 389 samples, Mg concentrations were correlated with reconstructed mean annual temperatures. This 390 potentially leads to an over or under-estimation of temperatures depending on the seasonality 391 of the growth period and might explain the observed difference between the two regressions. 392 Due to sample limitation, we analysed foraminifera from a wider size fraction (230µm to 393 500µm), compared to Regenberg et al. (2009) (355-400µm), introducing an additional bias 394 between the two datasets (Duenas-Bohorquez et al., 2010; Friedrich et al., 2012). Finally, 395 Regenberg et al. (2009), compiled data of samples from the tropical Atlantic and Caribbean 396 Ocean, while we collected samples from the Eastern tropical Atlantic. All of these potential 397 biases can easily explain the small discrepancy observed between our regression and the one 398 from Regenberg et al., (2009). Interestingly, Jentzen et al., (2018), were able to compare Mg/Ca 399 ratios measured on T. sacculifer from both surface sediment samples of the Caribbean sea and 400 specimen sampled with a plankton net nearby. They observed a similar systematic increased 401 Mg/Ca ratio in fossils tests of T. sacculifer (+0.7 mmol/mol-1) compared to living specimens, 402 arguing that different seasonal signals were responsible for the observed difference. However, 403 it is interesting to note that the Mg/Ca differences observed between living T. sacculifer (e.g. 404 this study and Jentzen et al., 2018) and fossils specimens (e.g. Regenberg et al., 2009 and 405 Jentzen et al., 2018) could also be explained by the presence of GAM calcite on T. sacculifer 406 from sediment samples, as GAM calcite is enriched with Mg compared to pre-gametogenetic 407 calcite precipitated at the same temperature (Nurnberg et al., 1996). If Jentzen et al., (2018) and Regenberg et al. (2009) do not describe the presence or absence of GAM calcite on T. sacculifer 408 409 specimens analysed in their studies, a study on the population dynamics of T. sacculifer from 410 the central Red Sea Bijma and Hemleben (1990) concluded that the rate of gametogesis 411 increased exponentially between 300 and 400µm to reach a maximum of more than 80% at 412  $355\mu$ m (sieve size =500 $\mu$ m real test length). It can therefore safely be assumed that the Mg/Ca





- 413 difference between living specimens from the plankton and empty shells from the sediment is
- 414 due to GAM calcite.
- 415 The Mg-Temp data obtained by Jentzen et al., (2018) is however, in good agreement with the 416 equation established by Regenberg et al., (2009), and will therefore not be considered separately 417 in this study. The overall strong similarity observed between our regression and the one from 418 Regenberg et al. (2009), indicates nevertheless that Mg-temp calibrations established on T. 419 sacculifer specimen from plankton tow, can be applied to T. sacculifer (without Sac) from the 420 surface-sediment, even if these applications have to be considered with care and only on 421 sediment samples showing no sign of dissolution. 422 In contrast, the equation of Anand et al., (2003) based on sediment trap samples, is appreciably
- 423 different (Fig. 3b). This may be due to: (1) difference in cleaning and analytical procedures, (2) 424 uncertainty in estimated temperature (3) addition of GAM calcite at greater depth. Anand et al., 425 (2003) fixed the intercept of the exponential regression for T. sacculifer to the value obtained 426 for a multispecies regression and subsequently recalculated for each species the pre-exponential 427 coefficients. Using this approach their new equation for T. sacculifer is: Mg/Ca= 0.35 exp 428 (0.09\*T), which is identical to Nürnberg et al., (1996) and equation 1 from our study. Still, this 429 implicitly assumes a common temperature dependence exists for all species, which is not 430 realistic. To avoid a priori assumptions only the primary equation of Anand et al., (2003) (see 431 Tab. 3) is considered in this study.
- 432

#### 433 **4.4.** Comparison with previous δ<sub>18</sub>O-Temperature calibrations.

434 As for Mg/Ca, the oxygen isotope composition also shows a strong correlation with measured 435 surface water temperature (R<sub>2</sub>=0.90). The *T. sacculifer*  $\delta_{18}$ O-temperature equation of Spero et 436 al., (2003), based on a culture experiment, is very similar to equation 4 in our study. However, 437 sensitivity (slope) differs within the uncertainties calculated for equation 4. As no uncertainties are given for the Spero et al., (2003) equation, it is difficult to determine whether these 438 439 equations are statistically different or not. In contrast, the equation of Mulitza et al., (2003), has 440 a similar slope (within uncertainties) but a higher intercept (Fig. 4a). The equation of Erez and 441 Luz, (1983) differs considerably from equation 4, for both slope and intercept parameters. 442 Bemis et al., (1998) suggested a bias in the calibration due to uncontrolled carbonate chemistry 443 during the experiments of Erez and Luz (1983) (a decrease in pH, e.g. due to bacterial growth 444 in the culture medium or to a higher CO<sub>2</sub> concentration in the lab (air conditioners, numerous people working in the same room etc), would quickly lead to an increase in  $\delta_{18}O$  of culture-445 grown foraminifera). This could explain the observed effect between our study (equation 4) and 446





447 the calibration from Erez and Luz (1983). Although the equation of Mulitza et al., (2003) is 448 also based on T. sacculifer collected from surface waters, their equation is significantly different 449 from equation (4). This deviation could possibly be due to a difference in size fractions considered in the two studies (230 to 500  $\mu$ m, and 150 to 700  $\mu$ m for this study and Mulitza et 450 al., (2003), respectively). Berger et al. (1979), already reported that large T. sacculifer tests are 451 enriched in  $\delta_{18}O$  relative to smaller ones (variation of 0.5‰ between 177 and 590µm). 452 453 Similarly, in culture experiments, larger shells of *Globigerina bulloides* are isotopically heavier 454 relative to smaller specimens (variation of approximatively 0.3‰ between 300 to 415µm, Bemis et al., 1998). Jentzen et al., (2018) reported that: 'Enrichment of the heavier <sup>18</sup>O isotope 455 in living specimens below the mixed layer and in fossil tests is clearly related to lowered in situ 456 457 temperatures and gametogenic calcification'. Gametogenic calcite has been shown to enrich 458  $\delta_{18}$ O signatures by about 1.0-1.4‰ relative to pregametogenic T. sacculifer (Wyceh et al., 459 2018). Finally, variation in light intensity (e.g. due to different sampling period and/or sampling 460 location), may have influenced the  $\delta_{18}O$  composition via an impact on symbiont activity (Spero 461 and DeNiro, 1987). Bemis et al. (1998) demonstrated that in seawater with ambient [CO<sub>32-]</sub>, 462 Orbulina universa shells grown under high light level (> 380 µEinst m-2 s-1) are depleted in O18 by on average 0.33% relative to specimens grown under low light levels (20-30 µEinst m-2 s-463 464 1). The different correlation between  $\delta_{18}O$  and temperature reported by Mulitza et al., (2003) 465 may be caused by size fraction differences, different sampling time, light intensity, differences 466 in calcification depth or hydrography, or a combination of factors. These are all potential biases that could explain the steeper intercept observed by Mulitza et al., (2003) relative to our study. 467 468

#### 469 **5. Reconstructions**

470 A few scenarios are considered in the following section, in which one, two or three proxy471 equations are combined to solve for salinity.

472

473 Three Mg/Ca-paleo-temperature equations (Nürnberg et al., 1996; Regenberg et al., 2009; and Anand et al., 2003) were used to compare "reconstructed" temperatures to the known in situ 474 475 surface waters temperatures. The mean foraminiferal Mg/Ca ratio measured at each of our 476 stations was inserted into each of the three equation and solved for temperature (Fig. 3b.). The 477 linear regression of reconstructed temperatures based on Nürnberg et al. (1996) overlaps almost 478 perfectly with the theoretical best fit. This confirms that calibrations based on culture 479 experiments (the primary geochemical signal recorded in the tests) are very well-suited for reconstructing surface water temperature. The regression from Regenberg et al., (2009) 480





481 reconstructed surface temperature that are too warm. This is in agreement with the fact that the 482 Mg/Ca ratio from surface sediment foraminifera are slightly higher than for living specimen 483 (Jentzen et al. 2018). The offset increases with decreasing temperature (0.5°C and 1.5°C 484 respectively at 30°C and 16°C). Finally, the reconstructed temperature using the equation from Anand et al. (2003), shows a strong systematic offset. Because the equation of Nürnberg et al., 485 486 (1996) matched our measured temperatures almost perfectly, their equation will be used to 487 analyse further reconstruction. Still, we acknowledge that downcore reconstructions will 488 inevitably also involve GAM calcite and hence other calibrations established using specimens 489 collected deeper in the water column or in the sediment should be better suitable. Similarly, 490 three  $\delta_{18}$ O-paleo temperature equations (Erez and Luz, 1983; Mulitza et al., 2003; Spero et al., 2003) were tested to reconstruct  $\delta_{18}$ Oc- $\delta_{18}$ Ow. The equation of Erez and Luz, (1983), shows a 491 492 significant systematic overestimation of  $\delta_{18}$ Oc- $\delta_{18}$ Ow, and will therefore not be considered any 493 further. Measured surface water temperatures at our 13 stations were inserted into the equations 494 of Mulitza et al., (2003) and Spero et al., (2003) to derive  $\delta_{18}$ Oc- $\delta_{18}$ Ow (Fig. 4). The  $\delta_{18}$ Oc-495  $\delta_{18}$ Ow reconstructions based on the equation of Mulitza et al. (2003) and Spero et al. (2003), 496 are both slightly more positive, than the theoretical best fit. In order to test the robustness of 497 δ18Ow reconstructions from paleoceanographic literature (e.g. Nürnberg and Groeneveld, 2006; 498 Bahr et al., 2011), we use the reconstructed temperatures based on the Mg/Ca-paleo-499 temperature equation from Nürnberg et al., (1996) to predict  $\delta_{18}$ Ow using measured  $\delta_{18}$ Oc and 500 the equations from Mulitza et al., (2003) and Spero et al. (2003). The reconstructed  $\delta_{18}$ Oc-501  $\delta_{18}$ Ow from inserting the Mg/Ca temperature into these equations is slightly overestimated 502 (0.5‰), but the offsets remain small enough to consider these as reasonable reconstructions.

503

Nevertheless, when reconstructing  $\delta_{18}$ Ow by inserting the Mg/Ca temperature and measured  $\delta_{18}$ Oc in both equations, the correlation coefficients of the linear regressions are weak (R<sup>2</sup> = 0.19 and 0.13 for Spero et al., 2003 and Mulitza et al., 2003, respectively) demonstrating that the reconstructed  $\delta_{18}$ Ow is not very reliable, therefore no reconstruction of salinity using these equations will be further tested in this manuscript.

509

510 Nevertheless, to test the robustness of theoretical and empirical salinity reconstructions, we 511 have the perfect data set at hand, as every parameter is known from *in situ* measurement or 512 sampling. We will use the equations 1, 4 and 5 established in this study and presented in table 513 3, for demonstration purposes.





515	$Mg/Ca = ae^{bT}$	Eq. 1
516	with $a=0.42(\pm 0.13)$ and $b=0.083(\pm 0.001)$	
517		
518	$T=c+d(\delta^{18}0c-\delta^{18}0w)$	Eq. 4
519	with $c=12.08(\pm 1.46)$ and $d=-4.73(\pm 0.51)$	
520		
521	$\delta^{18}Ow = eS + f$	Eq. 5
522	with $e=0.171(\pm 0.04)$ and $f=-4.93(\pm 1.66)$	
523	Classically, from those equations it is possible to extract var	iables estimated from the
524	observation $Mg/Ca$ and $\delta^{18}Oc$ through the equations:	
525	$\hat{T} = \frac{1}{b}(\log (Mg/Ca) - \log(a))$	Eq.1'
526	$\delta^{\widehat{18}Ow} = \delta^{18}Oc - \frac{1}{d}(\hat{T} - c)$	Eq. 4'
527	$\hat{S} = \frac{1}{e} \left( \delta^{\widehat{180w}} - f \right)$	Eq. 5'
528		

529

Given that  $\hat{T}$  is estimated from the fit from Eq. 1' (fig. 3a) and  $\delta^{\widehat{18}Ow}$  is estimated from Eq. 4', 530  $\hat{S}$  is finally calculated from Eq. 5' (figure 5). Hence, the error in  $\hat{S}$  is an accumulation of errors 531 from successive fits. In this study the standard deviation of the fit between  $\hat{S}$  and the measured 532 533 salinity for the 13 stations is  $\pm 2.49$  and the R<sup>2</sup> is 0.33 (p-value 0.04) (Fig. 6a and b). In conclusion, even the best possible salinity reconstruction based on locally calibrated equations 534 1, 4 and 5 from the present study only allows salinity reconstructions with a precision of  $\pm 2.49$ . 535 536 In the modern Atlantic Ocean, and based on recent sea surface salinity estimation (Vinogradova 537 et al., 2019), such a variability would not allow to distinguish water masses between 60°N to 538 60°S. Similarly, on a temporal timescale, given the regional salinity variations expected in most of the ocean over glacial-interglacial cycles is less than  $\pm 1, 2\sigma$  (Gray and Evans, 2019), such an 539 540 incertitude on salinity reconstruction would not even allow to distinguish modern versus last 541 glacial maximum water masses.

542

543 In the following steps, we quantify the error propagation more precisely. In simple cases, error 544 accumulation in an equation can be assessed by calculating the partial derivatives and by 545 propagating the uncertainties of the equation with respect to the predictors (Clifford, 1973). 546 However, for complex functions the calculation of partial derivatives can be tedious. Here, error





547 propagation related to  $\hat{S}$  was computed by a Monte Carlo simulation, which is simple to implement (Anderson, 1976). It is important to note that the propagated error with a 548 549 reconstructed salinity is a combination of fitting errors and errors associated with measurement inaccuracies (Mg/Ca and  $\delta^{18}$ Oc). First, we will only consider the error related to the fitting 550 procedure, (Eq. 1',4' and 5', assuming that variables (i.e. the data) are perfectly known without 551 uncertainties). For example, the fitting error related to Eq. 4' is computed by fitting  $\delta^{18}$ Ow 552 from measured  $\delta^{18}$ Oc and measured Temperature, i.e. the data are known and not 553 approximated. This is done by adding random Gaussian noise, with standard deviation 554 corresponding to the RMSE (Root Mean Square error) of each fit (respectively 1.32°C for 555 Eq.1', 0.15‰ for Eq. 4' and 0.55 for Eq. 5'). The resulting standard deviation error for the 556 557 reconstructed Salinity based on 10000 fits following the Monte-Carlo approach amounted to 558  $\pm 1.69$  (each fit using sampling from random distributions defined above). Hence,  $\pm 1.69$  is the 559 smallest possible error for salinity reconstructions, using the three steps above, only due to its mathematics. We can also estimate the error propagation at each step:  $\hat{T}\pm 1.32^{\circ}C$  (Eq.1'), 560  $\delta^{\widehat{18}Ow\pm 0.45\%}$  (Eq.4') and  $\hat{S}_{\pm 1.69}$  (Eq.5'). Now we will include the uncertainties related to 561 estimating the variables using proxy data. Hereto, some Gaussian noises simulating the 562 uncertainties of measured variables (Mg/Ca and  $\delta^{18}$ Oc) were introduced with standard 563 deviations taken from Table 2. The resulting standard deviation error increased to  $\pm 2.65$ . 564 Therefore, it can be concluded that statistically speaking,  $\delta^{\widehat{18}Ow}$  cannot be reconstructed to a 565 precision better than  $\pm 0.45\%$ , while salinity cannot be reconstructed to a precision better than 566  $\pm 1.69$  (fitting errors only) and, in reality hardly better than  $\pm 2.65$  (full to error propagation). 567 568

569 Finally, to complete this analysis, a direct linear fit to estimate salinity using exp  $(-\delta^{18}Oc)$ 570 and Mg/Ca was performed and led to an error of ±0.26 and a R<sup>2</sup> = 0.82 (p-value 2.10-4):

571

572 
$$\hat{S} = -0.16(\pm 0.02) e^{-\delta^{18}Oc} + 0.28(\pm 0.1) \frac{Mg}{ca} + 35.80(\pm 0.33) (\text{R}_2=0.81, \text{ p-value} \approx 2.10.4)$$
 Eq. 6  
573

574 This demonstrates that the direct reconstruction using the exact same variables as those initially 575 measured (Mg/Ca and  $\delta^{18}Oc$ ), led to a much better estimation of salinity that the successive 576 reconstruction.

- 577
- 578
- 579





580	Finally, to complete this analysis, a direct linear fit to estimate salinity using exp $(-\delta^{18}Oc)$
581	and Mg/Ca was performed and led to an error of $\pm 0.26$ and a R <sup>2</sup> = 0.82 (p-value 2.10-4):
582	
583	$\hat{S} = -0.16(\pm 0.02) e^{-\delta^{18}OC} + 0.28(\pm 0.1) \frac{Mg}{Ca} + 35.80(\pm 0.33) \text{ (R}_2 = 0.81, \text{ p-value} \approx 2.104) \qquad \text{Eq. 6}$
584	
585	This demonstrates that the direct reconstruction using the exact same variables as those initially
586	measured (Mg/Ca and $\delta^{18}Oc$ ), led to a much better estimation of salinity that the successive
587	reconstruction.
588	
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593	6. Implications
594	We analyzed shell Mg/Ca and Sr/Ca ratios, and $\delta_{18}O$ in <i>T. sacculifer</i> collected from surface
595	water along a North-South transect of the Eastern Tropical Atlantic Ocean. We find a strong
596	correlation between Mg/Ca ratios and surface water temperature, confirming the robustness of
597	surface water temperature reconstructions based on T. sacculifer Mg/Ca.
598	Insertion of the Sr/Ca ratio into the paleo-temperature equation improves the temperature
599	reconstruction. We established a new calibration for a paleo-temperature equation based on
600	Mg/Ca and Sr/Ca ratios for live T. sacculifer collected from surface water:
601	
602	
603	
604	
605	$T = (-27 \pm 15) + (8 \pm 1) \cdot \ln(Mg/Ca) + (28 \pm 11) \cdot Sr/Ca$
606	Scenarios were tested using previously published reconstructions. Results were compared to
607	reconstructions performed using local calibrations established in this study and therefore
608	supposed to represent the best possible calibration for this data set:
609	(1) Mg/Ca ratios measured in T. sacculifer specimens collected in surface water allow accurate
610	reconstruction of surface water temperature.





- 611 (2)  $\delta_{18}$ Ow can be reconstructed with an uncertainty of ±0.45‰. Such  $\delta_{18}$ Ow reconstructions
- 612 remain a helpful tool for paleo-reconstructions considering the global range of variation of
- 613 surface  $\delta_{18}$ Ow (from about -7 to 2‰, LeGrande and Schmidt 2006; ).
- 614

(3) In contrast, the best possible salinity reconstruction based on locally calibrated equations 1,

- 4 and 5 from the present study, only allowed reconstruction with an uncertainty of  $\pm 2.49$ . Such an uncertainty renders these reconstructions meaningless and does not allow for viable (paleo)salinity data.
- This is confirmed by a Monte Carlo simulation, applied to test successive reconstructions in an 'ideal case', where explanatory variables are known. This simulation shows that from a pure statistical point of view, successive reconstructions involving Mg/Ca and  $\delta_{18}$ Oc preclude salinity reconstruction with a precision better than  $\pm 1.69$  and hardly better than  $\pm 2.65$ , due to error propagation.
- Nevertheless, a direct linear fit to reconstruct salinity based on the same measured variables (Mg/Ca and  $\delta^{18}Oc$ ) was established (Eq. 6) and presented in table 3. This direct reconstruction of salinity should lead to a much better estimation of salinity (±0.26) than the successive reconstructions.

628

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**Table 1.** Measured temperature, salinity, DIC, ALK, and  $\delta_{18}$ Ow of the stations selected for this 880

study (October/November 2005). Mean annual temperature per station (World Ocean Atlas, 2005). 881

					Salinity	DIC	Alkalinity	
Stations	Latitude	Longitude	Measured	Mean annual	(±0.05)	(µmol/kg)	(µmol/kg)	δı8Ow (PDB)
				Tag		precision	precision 1.5	precision
			T°C (±0.05)	T°C		1µm/Kg	μm/Kg	0.1 ‰
						accuracy 2	accuracy 4	accuracy
			Oct/Nov.			µm/Kg	µm/Kg	0.2 ‰
25	22°38.640'N	20°23.578'W	24.91	21.8222	36.63	2069	2391	1.1
29	18°8.088'N	20°55.851'W	26.09	22.6964	36.24	2037	2369	0.9
31	14°32.128'N	20°57.251'W	28.24	25.0457	35.78	2009	2330	0.8
35	10°23.424'N	20°4.869'W	29.73	26.7731	35.63	1982	2304	1.2
38	7°2.114'N	17°27.818'W	29.43	27.764	34.67	1929	2257	0.7
40	4°22.323'N	15°16.911'W	28.47	27.7331	34.35	1915	2214	0.8
42	2°15.702'N	13°33.854'W	27.56	26.9215	35.72	2002	2332	1.1
46	1°35.741'S	10°33.846'W	25.91	25.6623	36.13	2053	2346	1.0
49	4°44.752'S	8°6.641'W	24.59	25.6229	36.07	2057	2369	0.9
52	8°6.086'S	5°29.077'W	23.80	24.864	35.99	2062	2360	0.7
56	11°51.783'S	2°30.743'W	22.18	23.6074	36.38	2071	2387	1.0
62	17°59.620'S	2°25.321'E	19.11	21.2856	35.99	2100	2369	1.1
66	22°26.998'S	6°6.922'E	18.71	20.1148	35.68	2070	2349	1.0





**Table 2.** Mean elemental (Mg/Ca and Sr/Ca) and isotopic ( $\delta_{18}$ Oc) composition per station, measured in foraminiferal calcite in mmol/mol and ‰ PDB, respectively. Elemental and isotopic compositions were determined on the same material (n varying from 5 to 9 specimens per station); isotopic analyses were done in duplicate for each station. Mean  $\delta_{18}$ Oc- $\delta_{18}$ Ow measured per stations in ‰ PDB.

Stations	Measured	Measured	Measured	Measured	Recons.	Recons.	Recons.
	Mg/Ca	Sr/Ca	δ18Oc ‰ (V-PDB)	δ18Oc- δ18Ow	δ18Ow (Mulitza)	δ18Ow (Spero)	δ18Ow (this study)
	mmol/mol	mmol/mol	precision 0.08‰	‰ (V-PDB)	‰ (V-PDB)	‰ (V-PDB)	‰ (V-PDB
25	$3.22\pm0.51$	$1.53\pm0.08$	-1.76	-2.82	0.38	0.40	0.88
29	$4.01\pm0.24$	$1.52\pm0.06$	-1.75	-2.63	1.00	0.87	1.44
31	$4.78\pm0.37$	$1.56\pm0.18$	-2.51	-3.33	0.73	0.49	1.11
35	$5.46\pm0.38$	$1.59\pm0.08$	-2.35	-3.59	1.27	0.94	1.62
38	$4.31 \pm 1.14$	$1.58\pm0.14$	-2.89	-3.59	0.07	-0.10	0.49
40	$4.07\pm0.64$	$1.57\pm0.07$	-2.98	-3.78	-0.18	-0.32	0.25
42	$3.79\pm0.49$	$1.53\pm0.08$	-2.38	-3.44	0.21	0.12	0.67
46	$3.92 \pm 1.24$	$1.47\pm0.07$	-1.67	-2.66	1.02	0.91	1.46
49	$2.99 \pm 0.39$	$1.55\pm0.11$	-1.83	-2.74	0.10	0.16	0.62
52	$2.97\pm0.30$	$1.50\pm0.03$	-1.34	-2.08	0.57	0.64	1.09
56	$3.31\pm0.53$	$1.50\pm0.03$	-1.06	-2.10	1.15	1.15	1.65
62	$2.20\pm0.24$	$1.47\pm0.07$	-0.70	-1.76	0.38	0.64	0.99
66	$1.66\pm0.17$	$1.48\pm0.09$	-0.74	-1.75	-0.46	-0.02	0.23





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**Table 3.** Calibration equations for *T. sacculifer*.

Mg/Ca Relationship with Temperature         Mg/Ca=0.42( $\pm 0.13$ )e^(T*0.083( $\pm 0.001$ ))         Eq. 1         0.86         2.9e-00           Nümberg et al., 1996         Mg/Ca=0.37( $\pm 0.065$ )e^(T*0.091( $\pm 0.007$ ))         0.93         0.93         0.93         0.93           Anand et al., 2003         Mg/Ca=0.6( $\pm 0.021$ )e^(T*0.048( $\pm 0.012$ ))         Regenberg et al., 2009         Mg/Ca=0.6( $\pm 0.16$ )e^(T*0.048( $\pm 0.012$ ))         0.93         0.93           Sr/Ca Relationship with Temperature         Sr/Ca Relationship with Temperature and Salinity         Sr/Ca=(0.0094 $\pm 0.002$ )*T+(1.29 $\pm 0.05$ )         Eq. 2         0.67         5.e-04           Mg/Ca and Sr/Ca Relationship with Temperature and Salinity         T=(-27\pm15)+(8\pm1)*In(Mg/Ca)+(28\pm11)*Sr/Ca         Eq. 3         0.93         2.e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27\pm15)+(8\pm1)*In(Mg/Ca)+(28\pm11)*Sr/Ca         Eq. 3         0.93         2.e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27\pm15)+(8\pm1)*In(Mg/Ca)+(28\pm11)*Sr/Ca         Eq. 3         0.93         2.e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27\pm15)+(8\pm1)*In(Mg/Ca)+(28\pm11)*Sr/Ca         Eq. 4         0.91         5.e-06           This study         T=12.08(\pm1.46)-4.73(\pm0.51)*(6ix0c-6ix0w)         T=0.01 \pm 0.001*5         0.91         5.e-06           Matrix study         T=12.08(\pm1.46)-4.	Source		R <sup>2</sup>	p-values
Temperature           Titis study         Mg/Ca=0.42(±0.13)e^{(T*0.083(±0.001))}         Eq. 1         0.86         2.9e-0           Nimberg et al., 2003         Mg/Ca=0.37(±0.065)e^{(T*0.091(±0.007))}         Eq. 1         0.86         2.9e-0           Ngenberg et al., 2003         Mg/Ca=0.6(±0.16)e^{(T*0.075(±0.006))}         Sr/Ca Relationship with           Temperature           This study         Sr/Ca=(0.0094±0.002)*T+(1.29 ± 0.05)         Eq. 2         0.67         5.e-04           Mg/Ca and StorCa Relationship with         Temperature         Tel:2.272:15)+(8±1)*In(Mg/Ca)+(28±11)*Sr/Ca         Eq. 2         0.67         5.e-04           Mg/Ca and StorCa Relationship with         Temperature           This study Mg/Ca)         In(Mg/Ca)=(-5.10±2)+(0.09±0.009)*T+(0.11±0.05)*S         0.91         5.e-06           This study Mg/Ca)         In(Mg/Ca)=(-5.10±2)+(0.09±0.009)*T+(0.11±0.05)*S         0.91         5.e-06           This study Mg/Ca)         IT=12.08(±1.46)-4.73(±0.51)*(6 inOc ~6 inOc)         Eq. 40 <th></th> <th></th> <th></th> <th>P . uides</th>				P . uides
Nümberg et al., 1996       Mg/Ca=0.37(±0.065)e^{T}0(.091(±0.007))       0,93         Anand et al., 2003       Mg/Ca=1.06(±0.021)e^{T}0.048(±0.012))       0,93         Regenberg et al., 2009       Mg/Ca=0.6(±0.16)e^{T}0.048(±0.012))       0,93         Sr/Ca Relationship with Temperature       This study       Sr/Ca=(0.0094±0.002)*T+(1.29 ± 0.05)       Eq. 2       0,67       5.e-04         Mg/Ca and Sr/Ca Relationship with Temperature       This study       T=(-27±15)+(8±1)*Bn/Ca       Eq. 3       0,93       2.e-04         Mc/Ca Relationship with Temperature and Salinity       T=(-27±15)+(8±1)*Bn/Ca Eq. 3       0,91       5.e-06         This study (Mg/Ca)       In(Mg/Ca)=(-5.10±2)+(0.09±0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Mg/Ca)       Sr/Ca = (1.81±0.5) + (0.008±0.002) T - (0.01±0.01)*S       0,71       0.002         ðis OR elationship with Temperature       T= 12.08(±1.46)-4.73(±0.51)*(ôISOc - ôISOw)       Eq. 4       0,88       1,6 e-00         Muitz at et al., (2003)       T= 15.35(±0.71).4.22(±0.25)*(ôISOc - ôISOw)       Eq. 4       0,88       1,6 e-00         Muitz at et al., (2003)       T= 12.5.67*(ôISOc - ôISOw)       Eq. 4       0,88       1,6 e-00         Serie et al., (2003)       T= 12.5.67*(ôISOc - ôISOw)       Eq. 5       0.38       1,2 e-01         Salinity				
Nümberg et al., 1996       Mg/Ca=-0.37( $\pm$ 0.065)e^(T*0.091( $\pm$ 0.007))       0,93         Anand et al., 2003       Mg/Ca=-0.6( $\pm$ 0.012)e^(T*0.048( $\pm$ 0.012))       0,93         Regenberg et al., 2009       Mg/Ca=-0.6( $\pm$ 0.16)e^(T*0.048( $\pm$ 0.012))       0,93         Sr/Ca Relationship with Temperature       5.e-04       0,67       5.e-04         Mg/Ca and Sr/Ca Relationship with Temperature       7.e-(27±15)+(8±1)*In(Mg/Ca)+(28±11)*Sr/Ca       Eq. 2       0,67       5.e-04         Mg/Ca and Sr/Ca Relationship with Temperature       7.e-(27±15)+(8±1)*In(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0,93       2.e-04         Mc/Ca Relationship with Temperature and Salinity       7.e-(27±15)+(8±1)*In(Mg/Ca)+(0.09±0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Mg/Ca)       In(Mg/Ca)=(-5.10±2)+(0.09\pm0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Mg/Ca)       In(Mg/Ca)=(-5.10±2)+(0.09\pm0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Mg/Ca)       Sr/Ca = (1.81±0.5) + (0.008\pm0.002) T - (0.01\pm0.01)*S       0,71       0.002         õis O e faitonship with Temperature       T= 12.08(±1.46)-4.73(±0.51)*(õis O e -õis Ow) Eq. 4       0,88       1,6 e-0t         This study       T = 15.35(±0.71)-4.22(±0.25)*(õi 80C e -õi 80Cw)       0.88       1,6 e-0t         Sere ot al., (2003)       T = 12.5.67*(õi 80C e -õi 80Cw)	This study	Mg/Ca=0.42(±0.13)e^(T*0.083(±0.001)) Eq. 1	0,86	2.9e-06
Regenberg et al., 2009       Mg/Ca=0.6(±0.16)e^A(T*0.075(±0.006))         Sr/Ca Relationship with Temperature       Sr/Ca=(0.0094±0.002)*T+(1.29 ± 0.05)       Eq. 2       0.67       5.e-04         Mg/Ca and Sr/Ca Relationship with Temperature       This study       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0.93       2.e-04         Me/Ca Relationship with Temperature and Salinity       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0.93       2.e-04         Me/Ca Relationship with Temperature and Salinity       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0.93       2.e-04         Me/Ca Relationship with Temperature and Salinity       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0.93       2.e-04         ðiso Relationship with Temperature       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±10.05)*(5       0.91       5.e-06       0.71       0.002         ðiso Relationship with Temperature       T=12.08(±1.46)-4.73(±0.51)*(6)isOc -6)isOw)       Eq. 4       0,88       1,6 e-0         Erez and Luz, (1983)       T= 16.06(±0.549)-5.08(±0.52)*(6)isOc -6)isOw)       Eq. 4       0,88       1,6 e-0         Muitza et al., (2003)       T=12.5.67*(6)isOc -6)isOw)       Eq. 5       0.38       1,2 e-0         Salinity (this study) $\delta_{15}O_{1} = (0.16(\pm 0.02)) e^{(-6^{n}_{18}O_{1}) + 0.28(\pm 0.1) Mg/Ca+35.80(\pm 0.33) Eq. 6       0.8$			0,93	
Sr/Ca Relationship with Temperature           This study         Sr/Ca=(0.0094±0.002)*T+(1.29 ± 0.05)         Eq. 2         0.67         5.e-04           Mg/Ca and Sr/Ca Relationship with Temperature         Temperature         903         2e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27±15)+(8±1)*In(Mg/Ca)+(28±11)*Sr/Ca         Eq. 3         0.93         2e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27±15)+(8±1)*In(Mg/Ca)+(28±11)*Sr/Ca         Eq. 3         0.93         2e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27±15)+(8±1)*(0.09±0.009)*T+(0.11±0.05)*S         0.91         5.e-06           This study (Sr/Ca)         Sr/Ca = (1.81±0.5) + (0.008±0.002) T - (0.01±0.01)*S         0.71         0.002           ðisO Relationship with Temperature         T=12.08(±1.46)-4.73(±0.51)*(6isOc - \deltaisOw)         Eq. 4         0.88         1.6 e-00           Erez and Luz, (1983)         T= 16.06(±0.549)-5.08(±0.32)*(618Oc - \delta18Ow)         T= 12-5.67*(818Oc - \delta18Ow)         T= 12-5.67*(818Oc - \delta18Ow)         T= 12-5.67*(818Oc - \delta18Ow)         T= 12-2.67*(818Oc - \delta18Ow)         <	Anand et al., 2003	Mg/Ca=1.06(±0.021)e^(T*0.048(±0.012))		
Sr/Ca Relationship with Temperature           This study         Sr/Ca=(0.0094±0.002)*T+(1.29 ± 0.05)         Eq. 2         0.67         5.e-04           Mg/Ca and Sr/Ca Relationship with Temperature         Temperature         903         2e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27±15)+(8±1)*In(Mg/Ca)+(28±11)*Sr/Ca         Eq. 3         0.93         2e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27±15)+(8±1)*In(Mg/Ca)+(28±11)*Sr/Ca         Eq. 3         0.93         2e-04           Me/Ca Relationship with Temperature and Salinity         T=(-27±15)+(8±1)*(0.09±0.009)*T+(0.11±0.05)*S         0.91         5.e-06           This study (Sr/Ca)         Sr/Ca = (1.81±0.5) + (0.008±0.002) T - (0.01±0.01)*S         0.71         0.002           ðisO Relationship with Temperature         T=12.08(±1.46)-4.73(±0.51)*(6isOc - \deltaisOw)         Eq. 4         0.88         1.6 e-00           Erez and Luz, (1983)         T= 16.06(±0.549)-5.08(±0.32)*(618Oc - \delta18Ow)         T= 12-5.67*(818Oc - \delta18Ow)         T= 12-5.67*(818Oc - \delta18Ow)         T= 12-5.67*(818Oc - \delta18Ow)         T= 12-2.67*(818Oc - \delta18Ow)         <	Regenberg et al., 2009	Mg/Ca=0.6(±0.16)e^(T*0.075(±0.006))		
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $				
Mg/Ca and St/Ca Relationship with Temperature       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0,93       2 e-04         Me/Ca Relationship with Temperature and Salinity       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0,93       2 e-04         Me/Ca Relationship with Temperature and Salinity       In(Mg/Ca)=(-5.10±2)+(0.09±0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Sr/Ca)       Sr/Ca = (1.81±0.5) + (0.008±0.002) T - (0.01±0.01)*S       0,71       0.002         õisO Relationship with Temperature       T=12.08(±1.46)-4.73(±0.51)*(õisOc -õisOw) Eq. 4       0,88       1,6 e-00         This study       T=12.08(±1.46)-4.73(±0.51)*(õisOc -õisOw) Eq. 4       0,88       1,6 e-00         Erez and Luz, (1983)       T=16.06(±0.549)-5.08(±0.32)*(õ18Oc -õ18Ow)       1,6 e-00         Mulitza et al., (2003)       T=12.5.67*(õ18Oc -õ18Ow)       1,6 e-00         Spero et al., (2003)       T=12.5.67*(õ18Oc -õ18Ow)       1,2 e-01         Mecta linear fit to reconstruct salinity       S= -0.16 (±0.02) e ^(- õ^*_{18}Oc) + 0.28 (±0.1) Mg/Ca+35.80 (±0.33) Eq. 6       0.82       < 2e-04	<u>*</u>			
Temperature         This study       T=(-27±15)+(8±1)*ln(Mg/Ca)+(28±11)*Sr/Ca       Eq. 3       0,93       2 e-04         Me/Ca Relationship with         Temperature and Salinity         This study (Mg/Ca)       In(Mg/Ca)=(-5.10±2)+(0.09±0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Sr/Ca)       Sr/Ca = (1.81±0.5) + (0.09±0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Sr/Ca)       Sr/Ca = (1.81±0.5) + (0.09±0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (Mg/Ca)       Sr/Ca = (1.81±0.5) + (0.09±0.009)*T+(0.11±0.01)*S       0,71       0.002         This study (Mg/Ca)       Sr/Ca = (1.81±0.5) + (0.09±0.009)*T+(0.11±0.01)*S       0,71       0.002         This study       T= 12.08(±1.46)-4.73(±0.51)*(6180c - 6180w)       Tent colspan="2">Tent colspan="2">Tis study (Tent colspan="2">Tent colspan="2">State colspan="2">State colspan="2">State colspan="2">State colspan="2">State colspan="2">State colspan="2">State colspan="2">Colspan= 20         Me/Ca Relationship with       Tent colspan="2">Tent colspan="2">Tent colspan="2">Tent colspan= 20       State colspan="2"Tent co		Sr/Ca=(0.0094±0.002)*T+(1.29 ± 0.05) Eq. 2	0,67	5.e-04
$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$				
Me/Ca Relationship with Temperature and Salinity       In (Mg/Ca)=(-5.10±2)+(0.09±0.009)*T+(0.11±0.05)*S       0.91       5.e-06         This study (Sr/Ca)       Sr/Ca = (1.81±0.5) + (0.008±0.002) T - (0.01±0.01)*S       0.71       0.002         õis0 Relationship with Temperature       T= 12.08(±1.46)-4.73(±0.51)*(õis0c -õis0ow) Eq. 4       0.88       1,6 e-0i         This study       T= 12.08(±1.46)-4.73(±0.51)*(õis0c -õis0ow)       Eq. 4       0.88       1,6 e-0i         Ferz and Luz, (1983)       T= 16.06(±0.549)-5.08(±0.32)*(õi80c -õi80w)       Mulitza et al., (2003)       T= 15.35(±0.71)-4.22(±0.25)*(õi80c -õi80w)       88       1,6 e-0i         Meiza et al., (2003)       T= 12-5.67*(õi80c -ði80w)       T= 12-5.67*(õi80c -ði80w)       88       1,2 e-0i         Salinity (this study)       Õis0w = (0.171±0.04)*S - (4.93 ±1.66)       Eq. 5       0.38       1,2 e-0i         Mainity       S = -0.16 (±0.02) e ^(- õ^1is0c)+ 0.28 (±0.1) Mg/Ca+35.80 (±0.33) Eq. 6       0.82       < 2e-0i				
Temperature and Salinity         This study (Mg/Ca) $\ln(Mg/Ca)=(-5.10\pm 2)+(0.09\pm 0.009)*T+(0.11\pm 0.05)*S$ 0.91       5.e-06         This study (Sr/Ca)       Sr/Ca = (1.81\pm 0.5) + (0.008\pm 0.002) T - (0.01\pm 0.01)*S       0.71       0.002         õis 0 Relationship with       Temperature       0.71       0.002         This study       T= 12.08(±1.46)-4.73(±0.51)*(õis0- ôis0w) Eq. 4       0.88       1,6 e-06         Erez and Luz, (1983)       T= 16.06(±0.549)-5.08(±0.32)*(õl80c -ôl80w)       Eq. 4       0.88       1,6 e-06         Mulitza et al., (2003)       T= 15.35(±0.71)-4.22(±0.25)*(õl80c -ôl80w)       Eq. 5       0.38       1,2 e-02         Sero et al., (2003)       T= 12-5.67*(õl80c -ôl80w)       Eq. 5       0.38       1,2 e-02         Mulitza et al. (2003)       T= 10.06(±0.02) e ^(- ô^180c) + 0.28 (±0.1) Mg/Ca+35.80 (±0.33) Eq. 6       0.82       < 2e-04		$T=(-27\pm15)+(8\pm1)*\ln(Mg/Ca)+(28\pm11)*Sr/Ca$ Eq. 3	0,93	2 e-04
This study (Mg/Ca)       ln(Mg/Ca)=(-5.10±2)+(0.09±0.009)*T+(0.11±0.05)*S       0,91       5.e-06         This study (St/Ca)       St/Ca = (1.81±0.5) + (0.008±0.002) T - (0.01±0.01)*S       0,71       0.002 $\delta_{18}O$ Relationship with       Temperature       0 <th0< th="">       0       <th0< th=""></th0<></th0<>				
This study (Sr/Ca)       Sr/Ca = (1.81±0.5) + (0.008±0.002) T - (0.01±0.01)*S       0,71       0.002 $\delta_{180}$ Relationship with       Temperature       0 <th0< th="">       0       0       0       <t< td=""><td></td><td></td><td></td><td></td></t<></th0<>				
$\delta_{18}O$ Relationship with Temperature       T = 12.08(±1.46)-4.73(±0.51)*( $\delta_{18}Oc - \delta_{18}Ov$ ) Eq. 4       0.88       1.6 e-0t         This study       T = 16.06(±0.549)-5.08(±0.32)*( $\delta_{18}Oc - \delta_{18}Ow$ )       Eq. 4       0.88       1.6 e-0t         Multiza et al. (2003)       T = 16.06(±0.549)-5.08(±0.32)*( $\delta_{18}Oc - \delta_{18}Ow$ )       Eq. 5       0.38       1.6 e-0t         Spero et al. (2003)       T = 12-5.67*( $\delta_{18}Oc - \delta_{18}Ow$ )       Eq. 5       0.38       1.2 e-0t         measured $\delta_{18}O$ vs. measured $\delta_{18}Ow = (0.171\pm0.04)*S - (4.93\pm1.66)$ Eq. 5       0.38       1.2 e-0t         direct linear fit to reconstruct salinity       S = -0.16 (±0.02) e^(-\delta^{18}Oc) + 0.28 (±0.1) Mg/Ca+35.80 (±0.33) Eq. 6       0.82       < 2e-0t         921       922       923       924       925       924       925       924				5.e-06
Temperature           This study         T= 12.08(±1.46)-4.73(±0.51)*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ ) Eq. 4         0.88         1,6 e-0           Erez and Luz, (1983)         T= 16.06(±0.549)-5.08(±0.32)*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )         0.88         1,6 e-0           Mulitza et al., (2003)         T= 15.35(±0.71)-4.22(±0.25)*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )         0.88         1,6 e-0           Spero et al., (2003)         T= 12-5.67*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )         0.88         1,2 e-0           measured $\delta_{18}O$ vs. measured $\delta_{18}O_{w} = (0.171\pm0.04)*S - (4.93\pm1.66)$ Eq. 5         0.38         1,2 e-0           direct linear fit to reconstruct salinity         S = -0.16 (±0.02) e^{(-\delta^{18}O_{c})+ 0.28 (±0.1)} Mg/Ca+35.80 (±0.33) Eq. 6         0.82         < 2e-04		$Sr/Ca = (1.81\pm0.5) + (0.008\pm0.002) T - (0.01\pm0.01)*S$	0,71	0.002
This study         T= 12.08(±1.46)-4.73(±0.51)*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )         Eq. 4         0,88         1,6 e-0t           Erez and Luz, (1983)         T= 16.06(±0.549)-5.08(±0.32)*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )         T= 15.35(±0.71)-4.22(±0.25)*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )         Nulitza et al., (2003)         T= 15.35(±0.71)-4.22(±0.25)*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )         Nulitza et al., (2003)         T= 12-5.67*( $\delta_{18}O_{c} - \delta_{18}O_{w}$ )           measured $\delta_{18}O$ vs. measured Salinity (this study) $\delta_{18}O_{w} = (0.171\pm0.04)*S - (4.93\pm1.66)$ Eq. 5         0.38         1,2 e-013           direct linear fit to reconstruct salinity $S = -0.16 (\pm 0.02) e^{-(-\delta^{-}18}O_{c}) + 0.28 (\pm 0.1) Mg/Ca+35.80 (\pm 0.33) Eq. 6$ $0.82 < 2e-04$ $2e-04$ 921         922         923         924         925         925         925         925         925         925         925         925         925         925         926         926         925         925         926         925         926         925         926         925         926         926         927         926         926         926         926         926         926         927         927         927         926         926         926         926         926         926         927         926         926         926				
Erez and Luz, (1983)       T = 16.06(±0.549)-5.08(±0.32)*( $\delta$ 18Oc - $\delta$ 18Ow)         Mulitza et al., (2003)       T = 15.35(±0.71)-4.22(±0.25)*( $\delta$ 18Oc - $\delta$ 18Ow)         Spero et al., (2003)       T = 12-5.67*( $\delta$ 18Oc - $\delta$ 18Ow)         measured $\delta$ 1sO vs. measured $\delta$ 1sOw = (0.171±0.04)*S - (4.93 ±1.66)       Eq. 5       0.38       1,2 e-0         direct linear fit to reconstruct       S = -0.16 (±0.02) e ^(- $\delta^{18}$ Oc)+ 0.28 (±0.1) Mg/Ca+35.80 (±0.33) Eq. 6       0.82       < 2e-04				
Mulitza et al., (2003)       T= $15.35(\pm 0.71) \cdot 4.22(\pm 0.25)*(\delta 180c - \delta 180w)$ Spero et al., (2003)       T= $12 \cdot 5.67*(\delta 180c - \delta 180w)$ measured $\delta_{18}O$ vs. measured Salinity (this study) $\delta_{18}O_w = (0.171\pm 0.04)*S - (4.93 \pm 1.66)$ Eq. 5 $0.38$ $1,2 e \cdot 0.5$ direct linear fit to reconstruct salinity based on measured variables (Mg/Ca and $\delta_{18}O_c$ ) $S = -0.16 (\pm 0.02) e^{(-\delta_{18}O_c) + 0.28 (\pm 0.1)} Mg/Ca + 35.80 (\pm 0.33) Eq. 6$ $0.82 < 2e \cdot 0.46 (\pm 0.22) e^{-0.28 (\pm 0.1)} Mg/Ca + 35.80 (\pm 0.33) Eq. 6$ 921       922       923       924       925	•		0,88	1,6 e-06
Spero et al., $(2003)$ T = 12-5.67*( $\delta 180c - \delta 180w$ )         measured $\delta 180$ vs. measured Salinity (this study) $\delta 180w = (0.171\pm0.04)*S - (4.93\pm1.66)$ Eq. 5 $0.38$ $1,2 e-0.5$ direct linear fit to reconstruct salinity based on measured variables (Mg/Ca and $\delta 180c$ )       S = -0.16 ( $\pm 0.02$ ) e ^(- $\delta ^{18}Oc) + 0.28$ ( $\pm 0.1$ ) Mg/Ca+35.80 ( $\pm 0.33$ ) Eq. 6 $0.82$ < 2e-04         921       922       923       924       925       925				
measured $\delta_{18}Ovs.$ measured Salinity (this study) $\delta_{18}Ove = (0.171 \pm 0.04)*S - (4.93 \pm 1.66)$ Eq. 5 $0.38$ $1,2 e-0.38$ direct linear fit to reconstruct salinity based on measured variables (Mg/Ca and $\delta_{18}Ove)$ $S = -0.16 (\pm 0.02) e^{(-\delta_{18}Ove) + 0.28 (\pm 0.1)} Mg/Ca + 35.80 (\pm 0.33) Eq. 6       0.82 < 2e-04         921       922       923       924       925       925   $				
Salinity (this study)direct linear fit to reconstruct salinity based on measured variables (Mg/Ca and $\delta_{18}O_c$ )S = -0.16 (±0.02) e ^(- $\delta^{18}O_c$ )+ 0.28 (±0.1) Mg/Ca+35.80 (±0.33) Eq. 6 (±0.33) Eq. 6 (±0.34) Eq. 6 (±0.34) Eq. 6<	Spero et al., (2003)	$T = 12-5.67*(\delta 180c - \delta 180w)$		
salinity based on measured variables (Mg/Ca and õisOc) 921 922 923 924 925		$\delta_{18}O_w = (0.171 \pm 0.04)^*S - (4.93 \pm 1.66) \qquad \qquad Eq. \ 5$	0.38	1,2 e-03
922 923 924 925	salinity based on measured variables	S = -0.16 (±0.02) e ^(- $\delta^{18}O_c$ )+ 0.28 (±0.1) Mg/Ca+35.80 (±0.33) Eq. 6	0.82	< 2e-04
923 924 925	921			
924 925	922			
925	923			
	924			
026	925			
	926			

- 928
- 929
- 930





- **Table 4.** Temperature, salinity and  $\delta_{18}O_w$  of the stations used to determine the salinity/  $\delta_{18}O_w$
- 935 relationship (equation 5)

Stations	Latitude	Longitude	T°C(±0.05)	Salinity(±0.05)	δ18Ow (SMOW) precision 0.1% accuracy 0.2%
19	33°20.14'N	14°38.45'W	22,09	36,83	1,3
21	30°23.42'N	16°24.99'W	23,01	36,91	1,4
23	25°20.68'N	18°4.17'W	24,87	37,01	1,8
25	22°38.64'N	20°23.58'W	24,91	36,63	1,3
29	18°8.09'N	20°55.85'W	26,09	36,24	1,1
31	14°32.13'N	20°57.25'W	28,24	35,78	1,1
35	10°23.424'N	20°4.869'W	29.73	35.63	1,5
36	9°5.71'N	19°14.21'W	29,29	35,63	1,1
37	7°43.88'N	18°5.42'W	29,25	34,92	1,0
38	7°2.11'N	17°27.82'W	29,43	34,67	1,0
39	5°49.51'N	16°29.68'W	29,34	34,34	1,0
40	4°22.32'N	15°16.91'W	28,47	34,35	1,1
42	2°15.70'N	13°33.85'W	27,56	35,72	1,3
43	0°57.53'N	12°33.06'W	26,48	36,05	1,3
46	1°35.74'S	10°33.85'W	25,91	36,13	1,3
47	2°17.53'S	10°1.35'W	26,16	36,2	1,2
49	4°44.75'S	8°6.64'W	24,59	36,07	1,2
51	6°55.67'S	6°24.31'W	24,28	36,01	1,1
52	8°6.09'S	5°29.08'W	23,8	35,99	1,0
56	11°51.79'S	2°30.74'W	22,18	36,38	1,3
62	17°59.62'S	2°25.32'E	19,11	35,99	1,3
66	22°26.99'S	6°6.92'E	18,71	35,68	1,3
69	25°0.20'S	8°17.16'E	18,19	35,64	0,9
72	27°2.39'S	10°35.53'E	18,5	35,64	1,0





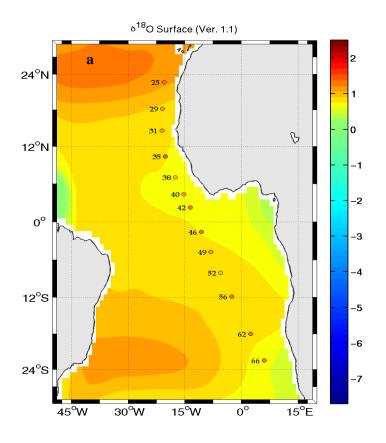
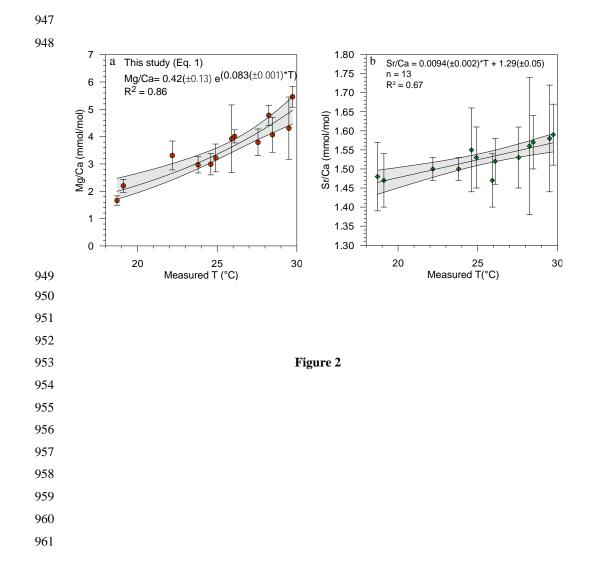




Figure 1

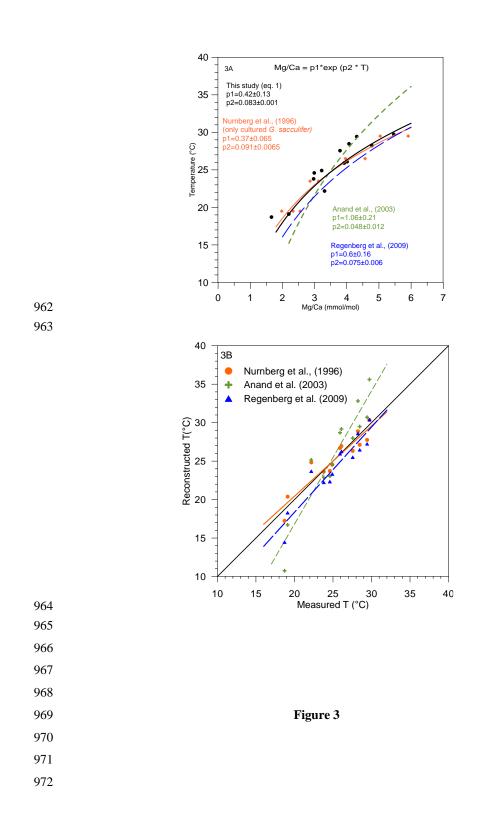






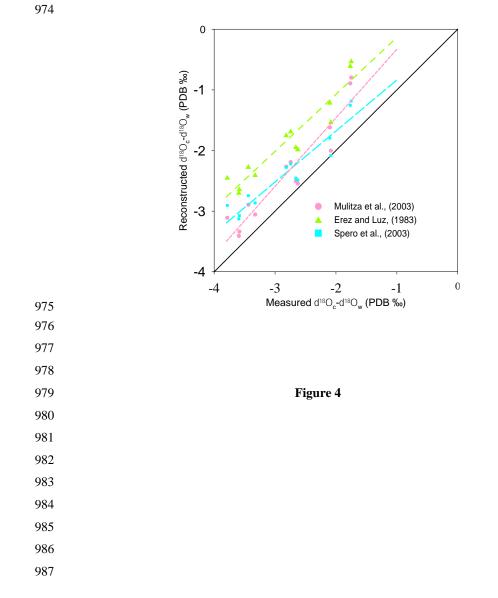






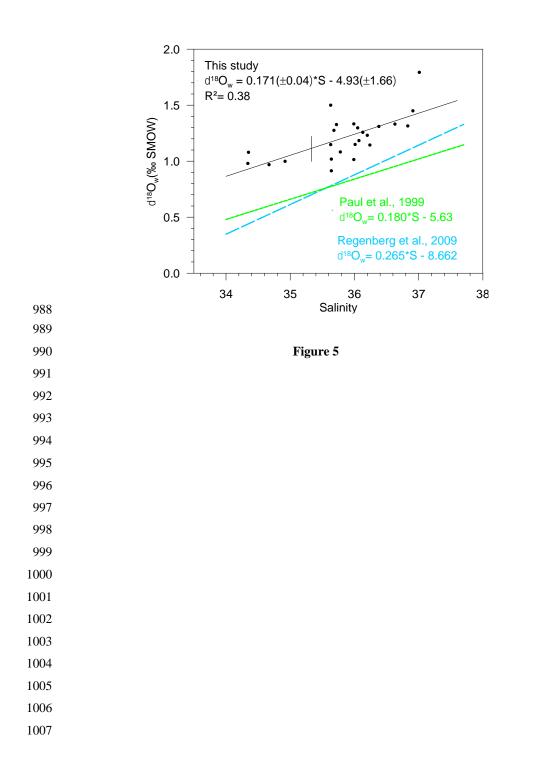






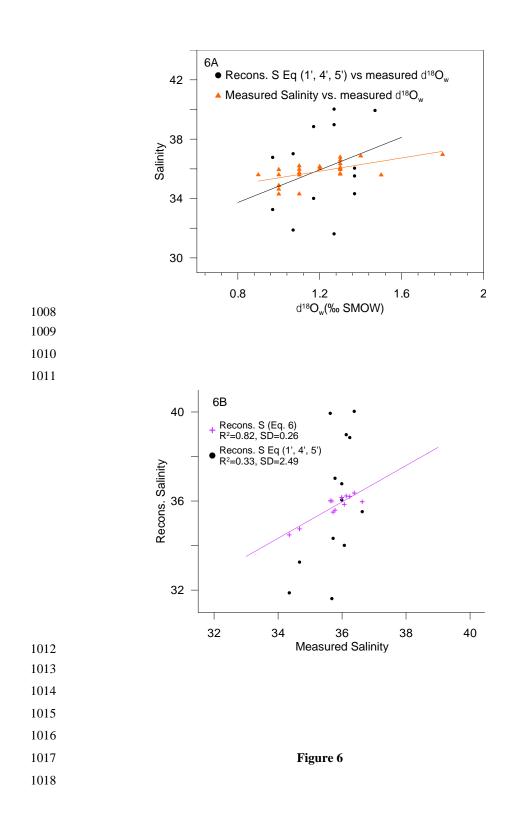
















1019	FIGURE LEGENDS
1020	
1021	Fig. 1: Stations used in this study, plotted on gridded data set described in LeGrande and
1022	Schmidt (2006) (a). (http://data.giss.nasa.gov/o18data/grid.html). Set up for planktonic
1023	foraminifers collections (b).
1024	
1025	Fig. 2: (a) Mg/Ca and (b) Sr/Ca (mmol/mol) and 95% confidence intervals plotted versus
1026	measured surface temperature (°C). Each point represents an average of the Mg/Ca and Sr/Ca
1027	per station.
1028	
1029	Fig. 3 a) Mg/Paleo-temperature equations established in this study (equation 1) (black dots, and
1030	full lines), based on the data of Nürnberg et al., (1996) (Orange diamond and large full orange
1031	line); Anand et al., (2003) (small green dotted line) and Regenberg et al., (2009) (large blue
1032	dotted line) and 3b) Reconstructed Mg-temperatures (Oct/Nov. 2005) plotted versus measured
1033	temperatures (°C) presented in Table 1. For each station mean measured Mg/Ca was inserted
1034	into the equation of Nürnberg et al., (1996) (only cultured specimens of T. sacculifer) (orange
1035	dots, full line), the equation of Anand et al., (2003) (green crosses, small dashed line), and the
1036	equation of Regenberg et al., (2009) (blue triangles, large dashed lines).
1037	
1038	Fig. 4: Reconstruction of $\delta_{18}$ Oc- $\delta_{18}$ Ow by inserting the measured temperature into three $\delta_{18}$ O
1039	based paleo-T-equation: The equation of Spero et al., (2003) (light blue squares, large light blue
1040	dashed line), the equation of Mulitza et al., (2003) (pink dots, small pink dashed line), the
1041	equation sorted by Erez and Luz (1983) (green triangles, green dashed line) plotted versus
1042	measured $\delta_{18}$ Oc- $\delta_{18}$ Ow (% PDB). The diagonal line represents the 1:1 regression.
1043	
1044	Fig. 5: Measured surface $\delta_{18}$ Ow ( $\infty$ SMOW) plotted versus measured surface salinity (stations
1045	listed in Tab. 4) (black dots and full line). Regression lines of the $\delta_{18}$ Ow-salinity relationship
1046	calculated by Paul et al., (1999) for the tropical Atlantic Ocean (from $25^{\circ}S$ to $25^{\circ}N$ ) based on
1047	GEOSECS data (green line), and by Regenberg et al., (2009) (blue dashed line) based on
1048	Schmidt (1999) data for the Atlantic Ocean for the water depth interval of 0–100 m.
1049	
1050	Fig. 6: a) Measured salinity (orange triangles) and reconstructed salinity based on equations 1,
1051	4 and 5 from the present study (black dots), plotted versus measured $\delta_{18}$ Ow.





- b) Reconstructed salinity based on 1) successive reconstructions using equations 1, 4 and 5
- 1053 from the present study (black dots) and 2) direct linear fit (Eq. 6) based on the same measured
- 1054 variables (Mg/Ca and  $\delta^{18}Oc$ ) (purple crosses), plotted versus measured salinity.

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