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# Contribution of changes in opal productivity and nutrient distribution in the coastal upwelling systems to late Pliocene/early Pleistocene climate cooling

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## Abstract

The global late Pliocene/early Pleistocene cooling (~3.0–2.0 million years ago, Ma) concurred with extremely high diatom and biogenic opal production in most of the major coastal upwelling regions. This phenomenon was particularly pronounced in the Benguela Upwelling System (BUS), off Namibia, where it is known as the Matuyama Diatom Maximum (MDM). Our study focuses on a new diatom silicon isotope ( $\delta^{30}\text{Si}$ ) record covering the MDM in the BUS. Unexpectedly, the variations in  $\delta^{30}\text{Si}$  signal follow biogenic opal content, whereby the highest  $\delta^{30}\text{Si}$  values correspond to the highest biogenic opal content. We interpret the higher  $\delta^{30}\text{Si}$  values during the MDM as a result of a stronger degree of silicate utilization in the surface waters caused by high productivity of mat-forming diatom species. This was most likely promoted by weak upwelling intensity dominating the BUS during the Plio/Pleistocene cooling combined with a large silicate supply derived from a strong Southern Ocean nutrient leakage responding to the expansion of Antarctic ice cover and the resulting stratification of the polar ocean 3.0–2.7 Ma ago. A similar scenario is hypothesized for other major coastal upwelling systems (e.g. off California) during this time interval, suggesting that the efficiency of the biological carbon pump was probably sufficiently enhanced in these regions during the MDM to have significantly increased the transport of atmospheric  $\text{CO}_2$  to the deep ocean. In addition, the coeval extension of the area of surface water stratification in both the Southern Ocean and the North Pacific, which decreased  $\text{CO}_2$  release to the atmosphere, led to further enhanced atmospheric  $\text{CO}_2$  drawn-down and thus contributed significantly to late Pliocene/early Pleistocene cooling.

## 1 Introduction

The causes and consequences of global late Pliocene/early Pleistocene cooling (ca. 3.0–2.0 Ma) have been subject to numerous investigations in order to understand mechanisms driving global climate from a state warmer than today towards colder

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1991), at which hydrological conditions were comparable (Dekens et al., 2007; Liu et al., 2008; Herbert et al., 1998). The most pressing questions regarding the cause of the MDM are therefore: (i) how to explain the co-occurrence of the largest amount of Antarctic diatom mats concomitant with warm and mainly stratified surface water conditions in the coastal upwelling regions during the late Pliocene/early Pleistocene? Why did the decline of the Antarctic diatom flora accompany the decrease in opal production when upwelling intensified at the beginning of the early Pleistocene, despite the fact that the development of the upwelling system should have enhanced nutrient supply and sustained higher diatom productivity? (ii) Given the importance of the changes in global opal productivity for the carbon cycle, to what extent did these major shifts of the centers of opal production promote late Pliocene/early Pleistocene climate cooling?

To answer these fundamental questions, we first investigate the causes of the onset and the end of the MDM in the BUS, one of the major areas where this high biogenic opal production was most pronounced. Using a comparison of proxies, we reconstruct regional biogeochemical cycling between 3.0 and 1.5 Ma in the BUS. We then compared our results from this region to other upwelling systems and regarding related changes in the polar oceans during this time interval we propose a scenario illustrating a possible significant impact of the MDM on the Plio-Pleistocene global climate transition.

## 2 Material and methods

### 2.1 IODP Site 1082

To conduct this study, we generated a new  $\delta^{30}\text{Si}$  record of diatom opal for the well-dated Benguela ODP (Ocean Drilling Program) Site 1082, Hole A (21°06' S, 11°49' E, 1279 m water depth) located on the Walvis Ridge in the northern Cape basin, which is compared with nitrogen isotopes ( $\delta^{15}\text{N}$ ) and biogenic opal content (BSi) and mass accumulation rates (MAR). Stratigraphy of Site 1082 has previously been described

extensively by Etourneau et al. (2009). In the center of the BUS, at Site 1084 (Fig. 1), the upwelling is at present active year-round while in the north, where the IODP Site 1082 is located, the upwelling cells develops only seasonally (Shannon, 1985), making the study site at a suitable location for recording past changes in upwelling intensity. Silicic acid ( $\text{Si}(\text{OH})_4$ ) and nitrate ( $\text{NO}_3^-$ ) are today mainly supplied from the Southern Ocean (Fig. 1) through the resurgence of Antarctic Intermediate Water (AAIW) and Subantarctic Mode Waters (SAMW) that form in the polar frontal system zone of the Atlantic Sector of the Southern Ocean (Shannon, 1985). When the wind-driven coastal upwelling off Namibia intensifies, nutrients are carried to the surface and support high productivity levels.

## 2.2 Silicon isotope compositions ( $\delta^{30}\text{Si}$ ) of diatoms

The downcore  $\delta^{30}\text{Si}$  record was obtained from diatom biogenic opal. The extraction of diatoms from the sediment followed the procedure of Morley et al. (2004). In the fraction 11–23  $\mu\text{m}$ , diatoms are by far the most abundant opaline organisms. After each extraction and purification step, the diatom opal samples were visually controlled under the microscope, and were then, according to the result, treated again with heavy liquid separation (at a slightly different density than before) or were wet-sieved again to purify this size-fraction as much as possible. Sponge spicules could not be observed in the samples, whereas radiolarian fragments were still present. By applying the separation steps carefully, the “contaminations” originating from other organisms could be reduced to a very small amount (below approximately 5%), so that a significant influence on the Si isotope data is highly unlikely. Also, abundances of radiolarian fragments were compared to the  $\delta^{30}\text{Si}$  data and no evidence was found for any correlation.

The diatom opal was then dissolved in 1 ml 0.1 N NaOH at 130 °C for several hours (usually overnight), and then centrifuged. The supernatant solution was thereafter transferred into new teflon vials. 200  $\mu\text{l}$  of  $\text{H}_2\text{O}_2$  were added to remove any remaining organic matter. After most of the reaction had ended, the samples were dried down and re-dissolved in 1 ml 0.1 N NaOH at 130 °C overnight. The still warm resulting samples

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were then diluted with 4 ml MQ water, and were neutralised with 0.1 ml 1 N HCl. Sample aliquots were used to determine Si concentrations using a colorimetric method on a photospectrometer (Hansen and Koroleff, 1999). The chromatographic separation and purification (Georg et al., 2006; Reynolds et al., 2008) was achieved with a precleaned 1 ml BioRad ion exchange column filled with AG50W-X8 (200–400 mesh) cation exchange resin.

All  $\delta^{30}\text{Si}$  ratios were measured on a Nu instruments MC-ICPMS at GEOMAR in Kiel, which is equipped with an adjustable entrance slit for medium resolution to ensure peak separation of the  $^{30}\text{Si}$  peak and the molecular interference of  $^{14}\text{N}^{16}\text{O}$ . The sample and standard solutions were introduced into the plasma via a Cetac Aridus II desolvator equipped with a PFA nebulizer at a 60 to 80  $\mu\text{l min}^{-1}$  uptake rate. Measurements were achieved with a standard-sample-standard bracketing method (Albarède et al., 2004). Samples and standards were measured with a concentration of 0.4 to 0.6 ppm depending on the performance of the instrument on the measurement day. All silicon isotope results in this study are presented in the  $\delta^{30}\text{Si}$  notation as follows:

$$\delta^{30}\text{Si} = \left( \left( \frac{{}^{30}\text{Si}/{}^{28}\text{Si}_{\text{sample}}}{{}^{30}\text{Si}/{}^{28}\text{Si}_{\text{standard}}} \right) - 1 \right) \times 1000$$

representing deviations of the measured  $^{30}\text{Si}/^{28}\text{Si}$  from the international Si standard NBS28 in parts per thousand (‰).

Measurements of the inter-laboratory standards IRMM018 and Big Batch gave average values of  $\delta^{30}\text{Si} = -1.62 \pm 0.26\text{‰}$  ( $2\sigma$ ) and  $-10.73 \pm 0.27\text{‰}$  ( $2\sigma$ ), respectively. These values are in good agreement with values obtained by other laboratories (Reynolds et al., 2007). Samples were measured three to five times within a day session, which resulted in external errors between 0.06 and 0.29‰ in  $\delta^{30}\text{Si}$  ( $2\sigma$ ). Duplicate measurements over a longer period time ( $n = 20$  sessions/days within one year) of an in-house matrix standard gave a reproducibility of  $\pm 0.25\text{‰}$  in  $\delta^{30}\text{Si}$  ( $2\sigma$ ).

In order to test if the isotopic ratios are consistent with mass dependent fractionation (mean for kinetic and equilibrium fractionation), as occurs during biological processes, we plotted the  $\delta^{30}\text{Si}$  versus the  $\delta^{29}\text{Si}$  (Fig. 2). The theoretical mass dependent

fractionation between both isotopes is 0.5135 (Reynolds et al., 2006). The calculated fractionation for the data of our study is 0.5006. Both are, within the uncertainties of stable silicon isotope measurements on a MC-ICPMS, indistinguishable from each other and suggest that any molecular interferences were fully eliminated.

### 3 Results and discussion

#### 3.1 Diatom- $\delta^{30}\text{Si}$ evidence for local changes in Si cycling during the MDM

During the MDM interval, the diatom  $\delta^{30}\text{Si}$  values at Site 1082 were in the range expected for an upwelling area but experienced large amplitude changes ranging between values of 0.2 and 1.7‰ (Fig. 3). The lowest values around 0.3–0.4‰ generally coincide with minima in BSi content (<20%) and BSi accumulation rates ( $1\text{ g cm}^{-2}\text{ ka}^{-1}$ ), whereas the highest values (up to 1.4–1.7‰) correspond to the maxima of opal accumulation rates and diatom productivity (respectively, up to 60% and  $4\text{ g cm}^{-2}\text{ ka}^{-1}$ ). Given that diatoms fractionate the dissolved seawater Si by  $-1.1\text{‰}$  during incorporation into their opal frustules (De La Rocha et al., 1997),  $\delta^{30}\text{Si}$  values of up to 1.7‰ suggest that the surface waters in which the diatoms grew had isotopic values in the order of 2.8‰, whereas surface water values of only 1.5‰ are reconstructed for minimum diatom signatures of 0.4‰. For comparison, the bulk sediment and diatom-bound  $\delta^{15}\text{N}$  was at a minimum during the MDM and revealed very light values around 0–2‰, while after 2.4 Ma, the values increased to 4–5 during the mid-Pleistocene (Fig. 3) (Etourneau et al., 2009; Robinson et al., 2002) which is consistent with the distribution found in the modern situation (Pichevins et al., 2005).

Similar to nitrogen isotopes ( $\delta^{15}\text{N}$ ), silicon isotopes ( $\delta^{30}\text{Si}$ ) are fractionated during utilization in the surface waters in a way that the lighter Si isotopes are preferentially incorporated into diatom frustules (De La Rocha et al., 1998, 2000; Brzezinski et al., 2002; Varela et al., 2004), which leaves the ambient seawater enriched in the heavier isotopes. The degree of depletion thus determines the magnitude of isotopic

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fractionation and together with water mass mixing controls the dissolved signature in ambient seawater (e.g. Reynolds et al., 2006).

In this study, on the basis of our current knowledge, we assume that the  $\delta^{30}\text{Si}$  signature of the water masses during the MDM was similar to nowadays (De La Rocha and Bickle, 2005). We also consider the effects of differences in Si isotope fractionation between diatom species and their respective ecological niches negligible. As previously documented, the composition of diatom species changed at  $\sim 3.0$  Ma, from mixed diatom assemblages to a dominance of *T. antarctica*, a mat-forming diatom species characterizing the MDM, replaced again thereafter at  $\sim 2.4$ – $2.0$  Ma by dominating *Chaetoceros* spores and cetae (Berger et al., 2002; Lange et al., 1999; Pérez et al., 2001). Changes in diatom assemblages may therefore potentially have affected the isotopic signal at IODP Site 1082. However, no drastic fluctuations in the  $\delta^{30}\text{Si}$  – and  $\delta^{15}\text{N}$  – values accompanied these faunistic changes (Fig. 3). Instead, the  $\delta^{30}\text{Si}$  record followed the BSi MAR variations since 3.0 Ma which implies that the  $\delta^{30}\text{Si}$  signature was unlikely affected by changes in diatom assemblages but rather reflects past nutrient utilization and supply conditions. This assumption is supported by findings of De La Rocha et al. (1997) who reported results from culture experiments showing that three different diatom species had strictly the same fractionation factor. There is therefore no proven evidence that a change in diatom assemblage impacts the  $\delta^{30}\text{Si}$  values reconstructed in the BUS during the MDM.

Besides, *Chaetoceros* is a diatom species living in the surface waters whereas *T. antarctica* can migrate through the water column to feed in the nutricline and realize photosynthesis at the surface. Few studies focused on the effects of differences in depth, in which the diatoms grew and the possible differences of the isotopic signature of seawater on the  $\delta^{30}\text{Si}$  values of the diatoms. Cardinal et al. (2007) and Fripiat et al. (2011) demonstrated that in the Southern Ocean, the  $\delta^{30}\text{Si}$  values of the diatoms living in the mixed layer were nearly the same as those measured on diatoms growing in deeper layers. Some variations were found but remained overall low and are much smaller than those reconstructed by our  $\delta^{30}\text{Si}$  record. We therefore conclude that the

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effects of different diatom assemblages and their respective environment of growth are not significant enough to influence the  $\delta^{30}\text{Si}$  variations.

Dissolution of diatom frustules during the downward transport in the water column to the sediment may also affect the isotopic signature of Si. According to Demarest et al. (2009), if the preservation efficiency between different diatom assemblages is >20 % different from each other, then the  $\delta^{30}\text{Si}$  might change by  $\pm 0.1\%$ , whereby this process increases linearly as dissolution progresses. As previously demonstrated, *Chaetoceros* shows a high resistance to dissolution (Koning et al., 2001), and *T. antarctica* builds mats which usually are also well protected against dissolution (Ragueneau et al., 2000). This feature explains why well-preserved diatoms are found in the BUS sediment during the Plio-Pleistocene climate transition (Lange et al., 1999). We are therefore confident that the  $\delta^{30}\text{Si}$  measured on the biogenic opal is not significantly influenced by dissolution and its effect is most likely minor regarding the large amplitude variations in our  $\delta^{30}\text{Si}$  record.

Today, the BUS is mainly supplied with Si from the Southern Ocean waters through advection of SAMW and AAIW (Fig. 1). During the MDM, the delivery of Si to the BUS was probably even more pronounced in response to the extension of the Antarctic ice cap and the progressive stratification of the Southern Ocean during global cooling at  $\sim 3.0\text{--}2.7\text{ Ma}$  (Haug et al., 1999). This most likely allowed the transfer of a large amount of unused Si to the low-latitude upwelling systems (Cortese et al., 2004). The presence of diatom species typical for Southern Ocean waters in the BUS (Berger et al., 2002; Lange et al., 1999; Pérez et al., 2001) confirms the close connection between the two regions during the MDM. According to sedimentary archives, productivity during the MDM was mostly dominated by carbonate producers in the Southern Ocean (Gersonde et al., 1999) from where waters supplying the BUS originated. Very few diatoms or other opal producers were found throughout the Southern Ocean at that time (Cortese et al., 2004), which supports the conclusion that the original silica pool was poorly utilized. This observation therefore minimizes potential effects of diatom dissolution in the Southern Ocean water column potentially affecting the  $\delta^{30}\text{Si}$  values advected

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waters to the BUS because this process, as previously observed in culture experiments (Demarest et al., 2009), should have led to low  $\delta^{30}\text{Si}$  signatures in the BUS and not the high ones observed.

Moreover, supposing a similar reorganization of the nutrient cycles between the North Pacific and the Southern Ocean during the extension of the polar ice cover and the stratification of the polar oceans  $\sim 3.0\text{--}2.7$  Ma ago, the decline in biogenic opal producers in both regions may have been accompanied by a reduced silicate utilisation in the Southern Ocean and thereby have led to lower  $\delta^{30}\text{Si}$  values comparable to those reconstructed in the North Pacific (from  $\sim 1.6$  to  $\sim 1.2\text{‰}$  around  $\sim 2.73$  Ma) (Reynolds et al., 2008). This implies that the isotopic signature of waters entering the BUS was most likely low or at least much lower than the values calculated for the surface waters in the BUS for the subsequent period between 3.0 and 2.0 Ma. We therefore argue that the extracted isotopic signal of the diatoms in the Benguela upwelling system most likely reflects a local signature and recorded the utilization of Si by locally produced diatoms rather than changes in the advected dissolved  $\delta^{30}\text{Si}$  signature.

### 3.2 Evidence for a weak BUS during the MDM

According to our results, a relatively high degree of silicic acid utilisation should correspond to heavy diatom  $\delta^{30}\text{Si}$  values, i.e. close to  $1.8\text{‰}$ , and vice versa. Consequently, the MDM was characterized by a high rate silicate utilisation by locally-produced diatoms, mostly *T. antarctica* mats. However, these high isotopic values occurred during the interval of highest BSi concentrations. In addition, the highest BSi values also appeared during a period where the nitrate was largely unutilised as illustrated by the low  $\delta^{15}\text{N}$  values (Fig. 3) (Etourneau et al., 2009). Although the bulk  $\delta^{15}\text{N}$  values recorded an isotopic signature derived from a mixture of organic matter, its good coherency with the diatom-bound  $\delta^{15}\text{N}$  values determined at the same site (Robinson et al., 2002) demonstrates that the sedimentary  $\delta^{15}\text{N}$  signatures mostly reflected

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changes in surface water conditions and nitrate uptake by locally produced diatom species. Taken together, the high BSi MAR and  $\delta^{15}\text{N}$  records would therefore suggest a strong upwelling of nutrient-rich subsurface waters. The MDM in the BUS thus should also have corresponded to a lower and not a higher degree of silicate utilisation because of the silica-rich subsurface waters supply, whereas the opposite is observed.

An anticorrelation between the  $\delta^{15}\text{N}$  and  $\delta^{30}\text{Si}$  records has also been described in the North Pacific (Reynolds et al., 2008) during the onset of significant Northern Hemisphere glaciation and the associated establishment of stratification of the surface waters at 2.73 Ma. A similar pattern has been found in the Southern Ocean during the last glacial-interglacial cycles (De La Rocha et al., 1998; Brzezinski et al., 2002). In the latter study, the authors ascribed the high  $\delta^{15}\text{N}$  and the low  $\delta^{30}\text{Si}$  and BSi during the last glacial periods to enhanced iron (Fe) fertilization that would have generated preferential utilization and uptake of N by diatoms instead of Si and therefore, to a higher degree of fractionation of  $\delta^{15}\text{N}$  compared with Si isotopes (Brzezinski et al., 2002). Comparatively, in the North Pacific, the  $\delta^{15}\text{N}$  increased with the stratification of the surface waters at 2.73 Ma, while the  $\delta^{30}\text{Si}$  and BSi MARs declined. Reynolds et al. (2008) argued that the inverse relationship between the proxies could not have been caused by an increase in Fe fertilization as the opal productivity decreased instead of increased. They rather ascribed the anticorrelation to the combined sea ice and surface water stratification effects that limited the light and largely inhibited the upwelling of nutrients, thus causing a more complete N utilization in this region compared with that of Si.

Contrary to the North Pacific or the Southern Ocean, the BUS has been located along the coast and is supplied with large amounts of Fe through aeolian dust. It is therefore unlikely that Fe has played a major role for the observed  $\delta^{15}\text{N}$ ,  $\delta^{30}\text{Si}$ , and BSi variations, despite possible changes in Fe supply tied to varying hinterland climate conditions (Dupont, 2006). In upwelling areas such as the Benguela system, for which currently no water column or sedimentary diatom Si isotope data exist, it is expected that during periods of most intense upwelling activity the continuous supply of relatively



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characterized by long periods of stratified conditions of surface waters and weak upwelling activity in the BUS due to the presence of *T. antarctica*, a mat-forming diatom species which requires extended periods of stable and stratified conditions for growing and consuming an important amount of silicate to build their mats. In addition, the appearance of a smaller dominant diatom species (*Chaetoceros*), typical of modern upwelling conditions, only occurred when the MDM declined after 2.4–2.0 Ma (Fig. 2). Warm sea surface temperatures (SST) and low north-south SST gradients reported through the Benguela region between 3.0 and 2.4 Ma also indicated weak upwelling conditions (Etourneau et al., 2009; Marlow et al., 2000) during the period of increased diatom and opal productivity. This assumption is furthermore consistent with pollen data obtained at Site 1082 demonstrating that wetter conditions dominated the first part of the MDM due to weak upwelling activity and enhanced moisture transport to the continent (Dupont et al., 2005; Dupont, 2006). Thus, compelling evidence from different markers strongly supports weak coastal upwelling systems off Namibia as a feature of the late Pliocene/early Pleistocene cooling.

This weak BUS activity during the MDM was probably closely linked to the system of atmospheric pressure cells that engender strong trade winds and cause the upwelling of deep waters along the Namibian coast. The ocean high pressure cells were likely situated at a more southern position during the MDM owing to warmer atmospheric temperatures and a reduced ice cap over Antarctica, as well as warmer SSTs in the Southern Ocean (Martinez-Garcia et al., 2010; Pollard and Deconto, 2009). This would probably have maintained the South Atlantic high pressure cells far from the African continental low and prevented the formation of strong trade winds along the shore, especially in the northern part of the BUS, thus resulting in a weaker upwelling intensity. The transfer of advected nitrate and silicate-rich waters masses supplying the BUS during the MDM associated with long periods of stratified surface water conditions therefore offered favourable and stable conditions for the growth of mat-forming diatoms, the most significant contributor of biogenic opal production. Silicate was nearly completely consumed by *T. antarctica* for growing and building their extensive mats

and progressively became the limiting factor of growth as revealed by their high  $\delta^{30}\text{Si}$  signatures. By comparison, the utilization of nitrate remained much lower owing to its important supply from the Southern Ocean to the BUS, as indicated by the low  $\delta^{15}\text{N}$  signatures.

In contrast, between 2.4 and 2.0 Ma the MDM decline is marked by a pronounced reduction of the biogenic opal production and a shift from Antarctic species (*T. antarctica*) to upwelling diatom species (*Chaetoceros* spores and cetae) in concert with the intensification of the upwelling activity (Etourneau et al., 2009; Marlow et al., 2000) and the establishment of a strong meridional atmospheric circulation (Etourneau et al., 2010). In parallel, the polar frontal system in the Southern Ocean developed (Liu et al., 2008), which was accompanied by an increase in Southern Ocean siliceous productivity (Cortese et al., 2004), likely stimulated by increasingly iron supply via aeolian dust (Martinez-Garcia et al., 2011). This probably restricted the Si and N export to low-latitude upwelling regions and the nutrient access to upwelling productivity. Concomitantly, the  $\delta^{30}\text{Si}$  values and biogenic opal content in the BUS decreased whereas the  $\delta^{15}\text{N}$  increased (Fig. 1). Any significant effects of the changing diatom community on Si isotope fractionation are unlikely because no abrupt variation in  $\delta^{30}\text{Si}$  was observed during the shift from Antarctic to upwelling species. We also suppose that the  $\delta^{30}\text{Si}$  was essentially only measured on biogenic opal derived from the same diatom species as those producing the organic matter on which bulk  $\delta^{15}\text{N}$  was measured. Hence, taken together these arguments strongly suggest that the decreasing  $\delta^{30}\text{Si}$  and increasing  $\delta^{15}\text{N}$  values have been the consequence of a diminished utilisation of silica and a higher utilisation of nitrate, respectively, although the  $\delta^{15}\text{N}$  may be also overprinted by heavier isotope signatures circulating through the nutricline (Etourneau et al., 2009). The development of the upwelling system likely triggered a more continuous supply of both bioavailable Si and N. However, the higher utilisation of nitrate likely drove the latter to become a more limiting factor for phytoplankton growth, as illustrated by the higher  $\delta^{15}\text{N}$  values, so that Si utilization by local diatoms was reduced and led to lighter  $\delta^{30}\text{Si}$  values. In addition, the production of *Chaetoceros* instead of *T. antarctica*

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probably drove to reduced silicate utilization and biogenic opal production as the former species generates less biogenic opal owing to their smaller size than the Antarctic species and its lesser competitive capacity against the other micro-organisms.

#### 4 Implications for late Pliocene/early Pleistocene atmospheric CO<sub>2</sub>

5 The shift of the centres of opal deposition away from the polar oceans and low latitudes most likely had fundamental consequences for the carbon cycle and as such may also have had dramatic consequences for global climate. Our new results suggest a possible scenario linking the MDM, nutrient distribution and opal productivity changes in the BUS, as well as in other coastal upwelling systems (Fig. 4), to the late Pliocene/early  
10 Pleistocene global climate shift (Fig. 5).

Here, we propose that during the warm Pliocene (Fig. 5a) vertical mixing probably dominated the surface waters of the polar oceans (Haug et al., 1999; Sigman et al., 2004) whereas conditions of warm surface waters and weak upwelling activity governed the low-latitude eastern boundary current regions (e.g. Etourneau et al., 2009; this study). Siliceous productivity was mainly focused in the Southern Ocean and North  
15 Pacific while low phytoplankton productivity, limited by nutrient (Si and N) supply, developed along the upwelling areas (Cortese et al., 2004). Contrary to the low latitude coastal upwelling systems, the ocean/atmosphere CO<sub>2</sub> exchange was probably intense and acted as a source in polar regions, which thus contributed to maintain high atmospheric pCO<sub>2</sub> levels (350–400 ppm).  
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During the late Pliocene/early Pleistocene cooling (Fig. 5b), stratification of the Southern Ocean and North Pacific due to the extension of the polar ice caps from 3.0–2.7 Ma (Sigman et al., 2004) likely led to limitation of the ocean-atmosphere exchange in these regions, and therefore progressively reduced the capacity of the polar ocean  
25 to release CO<sub>2</sub> to the atmosphere, as was the case in the low latitude coastal upwelling regions during the preceding warm Pliocene. This stratification was accompanied by an efficient nutrient leakage towards the low latitudes, stimulating diatom productivity

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to 250–300 ppm). In addition, the strengthening of the global upwelling activity from 2.4–2.0 Ma likely increased the exposure of deeper water CO<sub>2</sub> to the atmosphere, thus counterbalancing the effects of the phytoplankton production.

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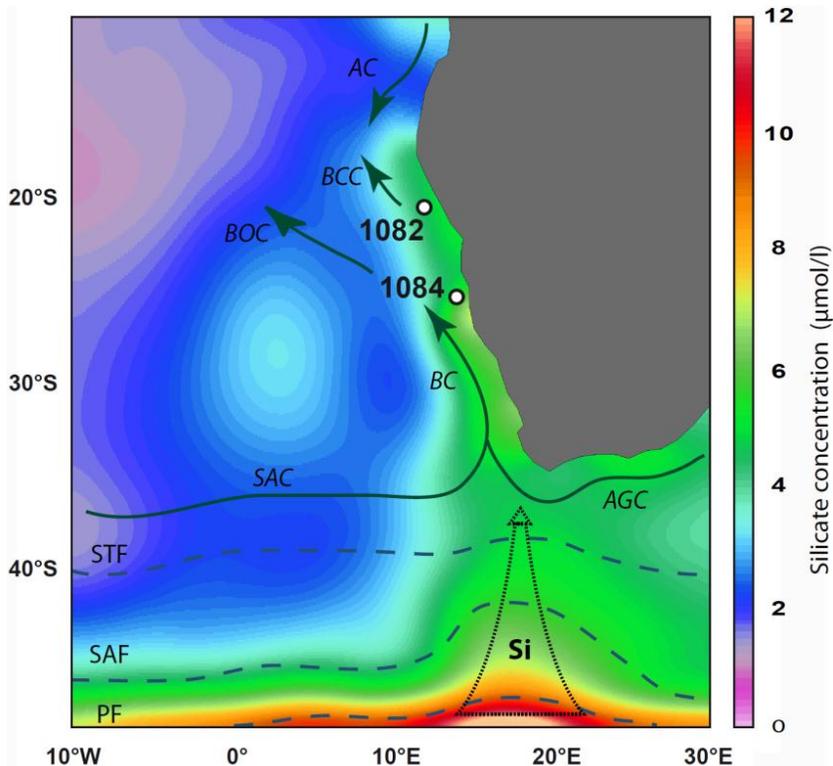
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**Fig. 1.** Silicate concentration ( $\mu\text{mol l}^{-1}$ ) at 10 m water depth and location of the Benguela ODP Sites 1082 and 1084. As illustrated here, Si is supplied in the BUS by the upwelling of SAMW and AAIW formed in the Southern Ocean, in the polar front system zone. AC, Angola Current; AGC, Aghulas Current; BC, Benguela Current; BCC, Benguela Coastal Current; BOC, Benguela Oceanic Current; SAC, South Atlantic Current. STF, Subtropical Front; SAF, Subantarctic Front, PF; Polar Front.

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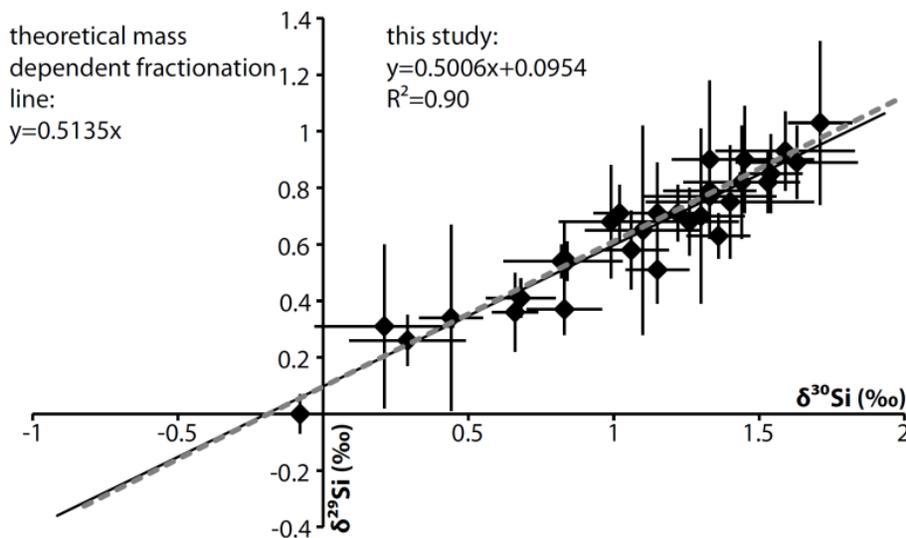
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**Fig. 2.**  $\delta^{30}\text{Si}$  (‰) versus  $\delta^{29}\text{Si}$  (‰).

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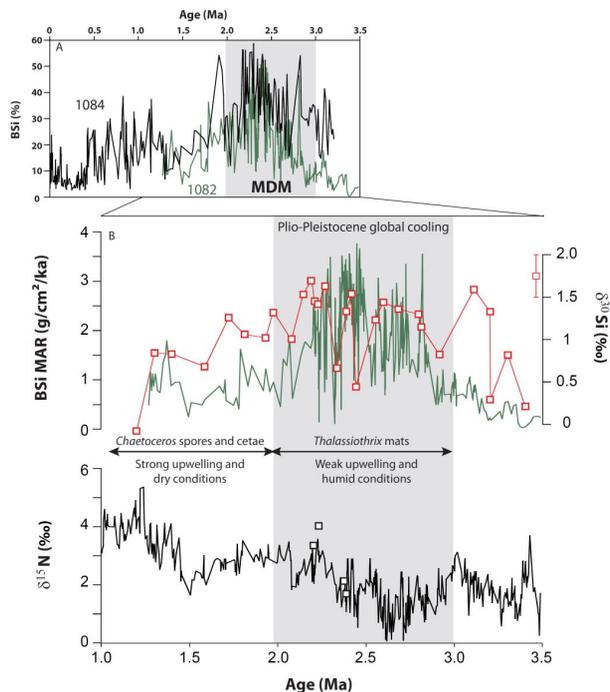
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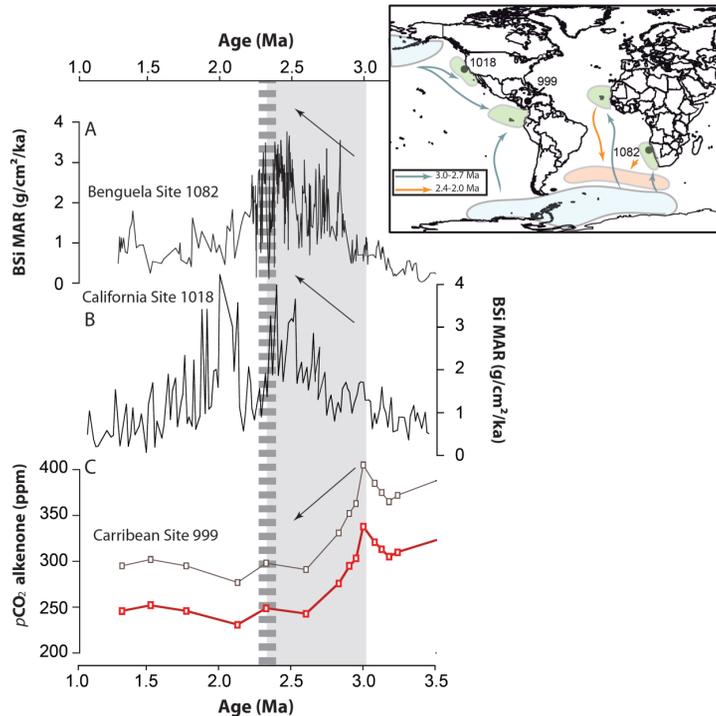
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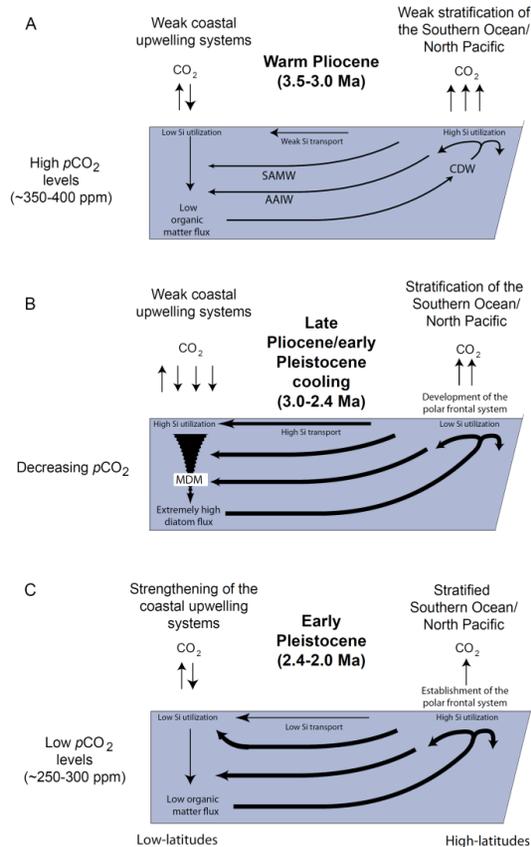
**Fig. 3.** **(A)** BSi content (%) at the Benguela Sites 1082 (green) and 1084 (black) (Etourneau et al., 2009; Lange et al., 1999; Pérez et al., 1999; Robinson et al., 2002). **(B)** BSi mass accumulation rate (MAR) ( $\text{g cm}^{-2} \text{ka}^{-1}$ ) (green) (Etourneau et al., 2009),  $\delta^{30}\text{Si}$  (red) (this study) and bulk  $\delta^{15}\text{N}$  (black) (Etourneau et al., 2009) records at Site 1082. Age models of Sites 1082 and 1084 are described in Etourneau et al. (2009). Black horizontal arrows indicate diatom species dominance with *Thalassiothrix antarctica* mats, characteristic of weak upwelling conditions, during the MDM and *Chaetoceros* spores and cetae, typical of strong upwelling conditions, during the mid-to-late Pleistocene (Marlow et al., 2000). In **(B)**, the typical  $2\sigma$  external reproducibility of the  $\delta^{30}\text{Si}$  measurements is indicated by the error bar.



**Fig. 4.** (A) BSi MAR at the Benguela Site 1082 and (B) at the California Site 1018 (Janecek, 2000). (C)  $p\text{CO}_2$  derived from alkenones at the Caribbean Site 999 (Seki et al., 2010). The grey zone corresponds to the period of increasing siliceous productivity and decreasing  $p\text{CO}_2$  as illustrated by the black arrows. The dashed line indicates the maximum biogenic opal production during the MDM. The associated map shows the respective locations of the different sites and the major opal centres shifts, from the Southern Ocean and the North Pacific (blue) to the low-latitudes upwelling regions (green) around 3.0–2.7 Ma, and from the latter to the Atlantic sector of the Southern Ocean (orange) around 2.4–2.0 Ma (modified from Cortese et al., 2004).

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**Fig. 5.** Proposed scenario representing the Si transport from the Southern Ocean/North Pacific towards upwelling systems and its utilization by phytoplankton productivity during **(A)** the warm Pliocene, **(B)** the late Pliocene/early Pleistocene cooling and **(C)** the early Pleistocene. AAIW, Antarctic Intermediate Waters, CDW, Circumpolar Deep Water, and SAMW, Subantarctic Mode Waters.

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