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## Deglacial biogenic opal peaks revealing enhanced Southern Ocean upwelling during the last 513 ka

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### ABSTRACT

Strength of Southern Ocean upwelling controls the exchange of carbon dioxide (CO<sub>2</sub>) between deep ocean reservoirs and atmosphere, as well as the communication of dissolved silicon with the euphotic zone of the Southern Ocean. The silicon supply could limit diatom opal productivity in the high-latitudes of Southern Ocean and the subsequent burial of biogenic opal in underlying sediments. Here we report a record of biogenic opal export off the Prydz Bay south of the polar front of the Southern Ocean, indicating strengthened upwelling during the past five glacial terminations. In all five terminations (I–V), opal peaks occur in line with Northern Hemisphere summer insolation intensity as well as the existing IRDs, indicating that freshwater injection associated with retreat of the Northern Hemisphere ice sheets could be the cause of enhanced upwelling in the Southern Ocean during terminations. This could in turn promote CO<sub>2</sub> outgassing, finally accelerating the completion of the terminations. In addition, the enhanced upwelling could export the Si-rich deep water to low latitudes via Antarctic Intermediate Water (AAIW) and Subantarctic Mode Water (SAMW), potentially leading to deglacial opal peaks in subtropical North Atlantic.

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### 1. Introduction

To improve the understanding of future climate change under the increasing anthropogenic CO<sub>2</sub> levels, a number of attempts have been made to interpret the close association between atmospheric CO<sub>2</sub> and global temperature in the late Quaternary (Siegenthaler et al., 2005). It is known that no single source can account for the full amplitude of glacial-interglacial variability of atmospheric CO<sub>2</sub> (Sigman and Boyle, 2000; Archer et al., 2000). Although multiple carbon sources may be involved (Köhler et al., 2005; Peacock et al., 2006), there is general agreement that deep ocean is a large carbon

reservoir accounting for the major amplitude of CO<sub>2</sub> glacial-interglacial variability in the past (Adkins et al., 2002; Sigman and Boyle, 2000; 2007; Watson and NaveiraGarabato, 2006; Toggweiler et al., 2006; Sikes, 2012).

It is proposed that changes in Southern Ocean upwelling is of crucial importance on regulating atmospheric CO<sub>2</sub> levels during Terminations. That is, the enhanced upwelling, perhaps associated with changes in Southern Westerlies, will stimulate the ventilation of carbon-rich deep water mass to the surface, promoting the CO<sub>2</sub> outgassing and finally triggering the global warming and Terminations (Skinner et al., 2010; Anderson et al., 2009). This hypothesis is supported by various paleoceanographic records broadly (e.g. <sup>13</sup>C-depleted carbon during deglaciation (Köhler et al., 2005; Smith et al., 1999; Spero and Lea, 2002), Nd isotope records in the Tobago Basin and East Equatorial Pacific (Pahnke et al., 2008; Pena et al., 2013), acute drop in deglacial <sup>14</sup>C activity of dissolved inorganic carbon (DIC) in North Pacific intermediate waters (Marchitto et al.,

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2007), etc), indicating a vital role played by upwelling branches of the global meridional overturning circulation (MOC) during the last deglaciation (Anderson et al., 2009; Siani et al., 2013).

From physical oceanographic perspective, upwelling regions in the SO consists of two branches associated with the upper and lower cell of global MOC, respectively in north and south of Antarctic Divergence Zone (Lumpkin and Speer, 2007; Marshall and Speer, 2012; Fig. 1B). The  $27.6 \text{ kg m}^{-3}$  density surface, outcropping south of the polar front all the way around Antarctica (see the red line in Fig. 1B), represents the average boundary between the upper and lower MOC cells in the SO (Marshall and Speer, 2012). Plenty of studies already demonstrated the variability of the upwelling strength associated with the upper cell during the last deglaciation (e.g. Anderson et al., 2009; Skinner, 2012), however, no reports exist discussing the upwelling in the lower cell. The latter is thus highly desirable to fully understand the outgassing process associated with the SO upwelling variability.

Consumption of dissolved silicic acid ( $\text{H}_4\text{SiO}_4$ ) by primary producers (diatoms) and cycling pathways represents primary prerequisites for effective carbon sequestration (Ragueneau et al., 2000). Diatoms live in the euphotic zone where they utilize the silicic acid to form opal tests. Although iron could limit the physiological status and growth rate of individual diatom cells (Boyd et al., 2000), biogenic opal productivity is ultimately limited by the supply of dissolved Si acid (Pondaven et al., 2000; Nelson et al., 2002). The paucity of silicate in the modern high-latitude Southern Ocean surface waters (Fig. 1B) has inspired the use of biogenic opal export as a measure of the silicate-rich deep waters upwelling to surface (Gnanadesikan and Toggweiler, 1999; Anderson et al., 2009; Meckler et al., 2013).

## 2. Materials and methods

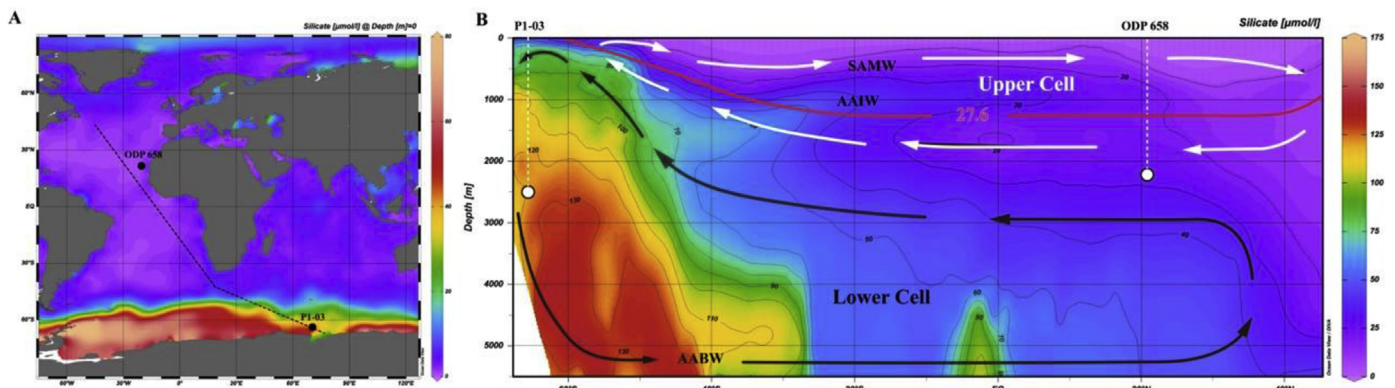
The gravity core P1-03 was drilled from the front edge of the continental slope deposition fan off the Prydz Bay ( $73^\circ 00' 56''\text{E}$ ,  $65^\circ 59' 22''\text{S}$ ) with a water depth of 2,542 m during the “30th Chinese National Antarctic Research Expedition” cruise in 2013–2014, (Fig. 1). P1-03 lies in the south of the  $27.6 \text{ kg m}^{-3}$  density surface outcropping location, having the potential to trace the surface Silicic Acid situation in the lower cell of the global MOC. The core is 5.64 m long, and the sediments mostly consist of gray clay silt without turbidite. The core was subsampled at an interval of 2 cm and totally 282 samples were prepared for the next analysis.

The AMS<sup>14</sup>C datum was determined from 4 sediment layers for organic carbon and carbonate carbon at Beta Analytic Lab, Miami, USA (Table 1). Organic matter is measured on acid insoluble organic carbon component of bulk samples and carbonate carbon is manually picked up from planktonic foraminifera *Neogloboquadrina pachyderma* (sin) (150–250  $\mu\text{m}$ ). We converted the AMS<sup>14</sup>C age into the calibrated calendar age using the online program Calib 7.04 (Stuiver and Reimer, 1993) and  $\Delta R$  value 830a according to Domack et al. (2001) in Palmer Deep west of Antarctic Peninsula.

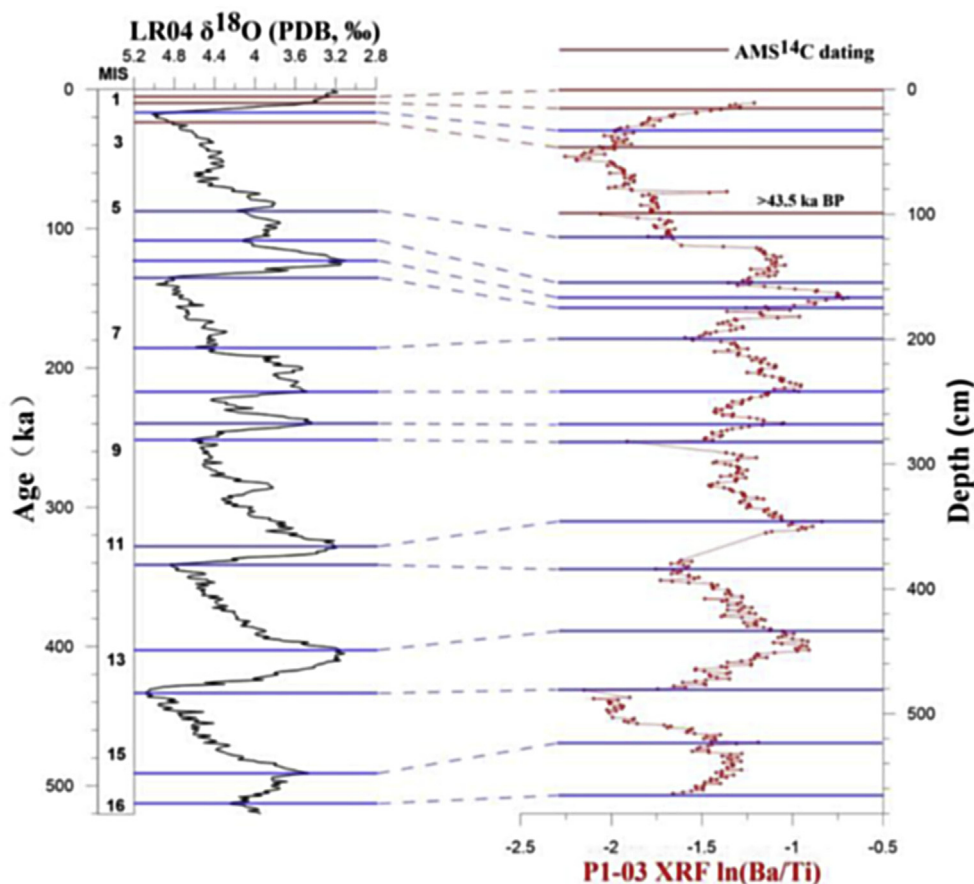
The measurements of X-ray fluorescence (XRF) scanning for P1-03 were performed with an Avaatech XRF Core Scanner at Tongji University. Data were obtained at a resolution of 1 cm over an area of  $1.2 \text{ cm}^2$  directly at the split core surface of the archive half, and we got the relative contents of the 29 elements from Al to Ba. The core surface was covered with 4 mm thick SPEXCerti Prep Ultralene1 foil to avoid contamination of the XRF measurement unit. Although the XRF data is unable to provide the percentage of elements, it has a positive correlation with the ICP-MS measured element contents (Tjallingii et al., 2007), and they have a linear logarithmic relationship (Weltje and Tjallingii, 2008). The calculations of normative Ba/Ti, Si/Al, Ca/Ti are based on the assumption that sedimentary Ti and Al are of detrital origin, and the composition of the Ti and Al bearing phases of the terrigenous material (detrital Ba/Ti, Si/Al, Ca/Ti) remained constant in space and time (Jaccard et al., 2009). In order to evaluate biogenic Ba, Si and Ca components, eliminate the influence of terrigenous material and deposition “dilution effect”, the normative calculations to Al and Ti have been widely applied in recent studies (Murray and Leinen, 1996; 2000; Wei et al., 2003).

Biogenic opal was determined by alkaline extraction of silica (Mortlock and Froelich, 1989), and average errors for this method range from 0.2 to 1.0%. Biogenic opal percentage was measured by molybdate-blue spectrophotometry at key laboratory of Marine Sedimentology and Environmental Geology.

Both sieve analysis and a laser particle sizer (Malvern Mastersizer 2000) were used for grain size measurement. First,  $\text{H}_2\text{O}_2$  was used to remove organic material, and HCl ( $0.25 \text{ mol l}^{-1}$ ) to remove  $\text{CaCO}_3$ . Samples were then placed in a water bath at  $85^\circ\text{C}$  for 4 h and  $\text{Na}_2\text{CO}_3$  ( $1 \text{ mol l}^{-1}$ ) was added to remove biogenic silica (opal). Samples were washed using de-ionized water and dried. The coarser fraction was retained on a mesh ( $>2 \text{ mm}$ , gravel), weighed and its abundance calculated in percent weight. The finer fractions



**Fig. 1.** Silicate concentration and location of study sites. Circles mean the sediment cores, the black dashed line indicates the section location. **A.** Map of annual average surface silicate concentration, indicating the scarcity of this nutrient in modern ocean (Levitus, 2009). **B.** Meridional depth section of silicate concentration in major present-day water masses (arrows) and the locations of P1-03 and North Atlantic core ODP 658. The white and black arrows indicate the water flows of the Upper Cell and Lower Cell of the global MOC. The red line is the  $27.6 \text{ kg m}^{-3}$  density surface. The maps were made using the Ocean Data View software (Schlitzer, 2000). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)



**Fig. 2.** Comparison between the XRF ln(Ba/Ti) curve of Core P1-03 and the stacked global benthic LR04  $\delta^{18}\text{O}$  (Lisiecki and Raymo, 2005). The solid red lines indicate Accelerator Mass Spectrometry (AMS) $^{14}\text{C}$  dating layers, and the solid blue lines indicate chronologic tie points. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

**Table 1**  
AMS $^{14}\text{C}$  age and calibrated age of core P1-03.

Depth (cm)	AMS $^{14}\text{C}$ age (a.BP)	Calibrated age (a.BP(1 $\sigma$ ))	Dating material
0–2	5560 $\pm$ 30	4961 $\pm$ 59	Organic Carbon
16–18	11,890 $\pm$ 40	12,607 $\pm$ 40	Organic Carbon
44–46	18,360 $\pm$ 60	22,265 $\pm$ 106	Organic Carbon
98–100	>43,500		planktonic foraminifera ( <i>N. pachyderma</i> )

(<2 mm) were immersed in de-ionized water, scattered by sodium metaphosphate and analysed with a laser particle sizer.

### 3. Results

#### 3.1. Chronological framework

Our sediment core spans the last four glacial-interglacial cycles starting from MIS13 (Figs. 3 and 4). Due to the absence of biogenic carbonate associated with development of continuous foraminiferal oxygen isotope stratigraphy and carbonate-based AMS $^{14}\text{C}$  data series in the studied core P1-03, we established the age model by organic carbon AMS $^{14}\text{C}$  dates for the last 30ka BP and by the correlation of Ba/Ti with LR04 stacked stable oxygen isotope for the older record (Lisiecki and Raymo, 2005). In a recent study (Wu et al., 2015), based on the close coupling between bioBa and LR04 curve, biogenic Ba has been chosen to reconstruct the age model of

sediment core at Prydz trough during the late quaternary. BioBa is a primary productivity proxy in paleoceanography study (Bonn et al., 1998; Dymond, 1992; Paytan and Griffith, 2007; Schenau et al., 2001; Tribouillard et al., 2006). Given the interglacial high productivity and glacial low productivity in the south of the Polar Front (Jaccard et al., 2013; Lamy et al., 2014; Anderson et al., 2014), the variation of primary productivity thus is in close relationship with climate changes during glacial interglacial cycles. Assuming that sedimentary Ti is of detrital origin, Ba abundance normalized to Ti yields an estimate of the sedimentary concentration of biogenic Ba (Bonn et al., 1998; Nurberg et al., 1997; Schroeder et al., 1997), which serves as a tool to reconstruct the chronological framework. Eighteen stratigraphic tie points for the study core P1-03 (including the organic carbon AMS $^{14}\text{C}$  dates, Table 1) was chosen for detailed comparison with the standard LR04  $\delta^{18}\text{O}$  stack (Fig. 2). During MIS 2 and MIS 1 with 3 AMS $^{14}\text{C}$  dates, the closely correlation between Ba/Ti and LR04 (Fig. 2) could verify the reasonableness of the age

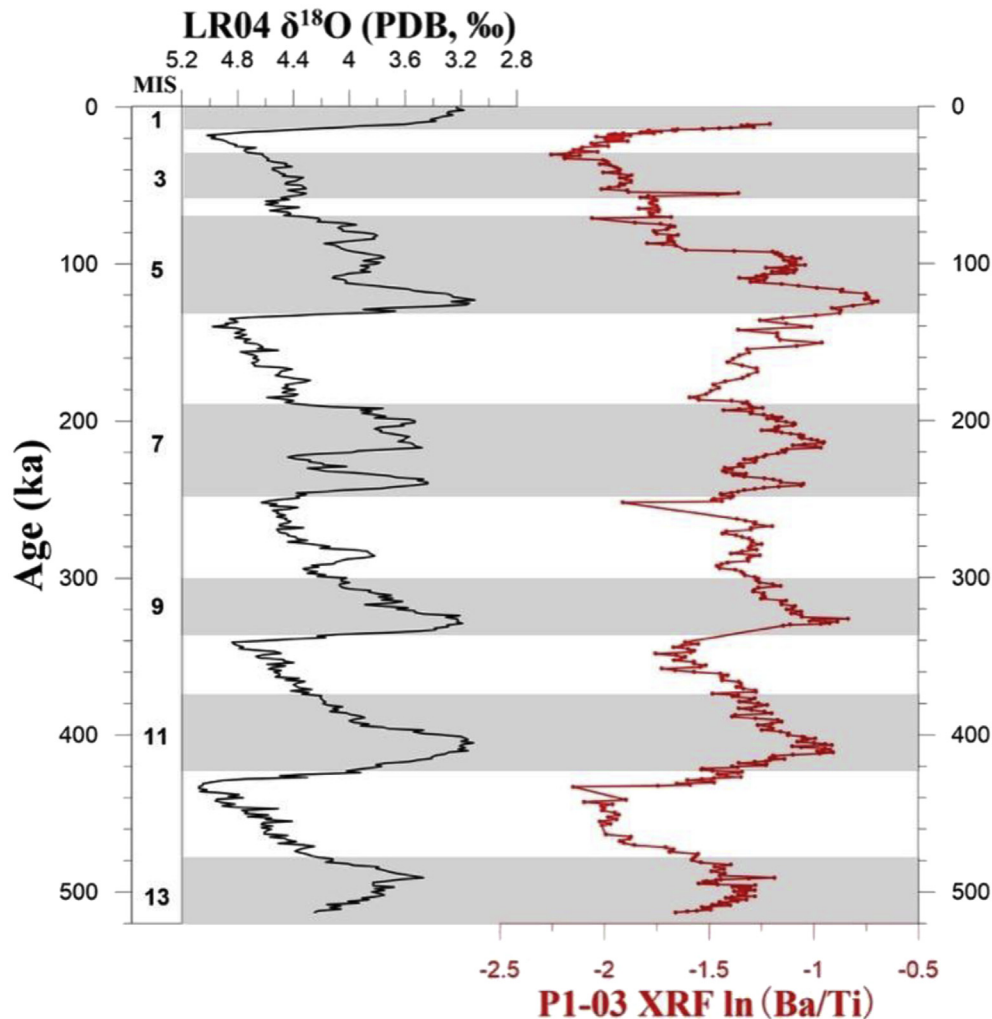


Fig. 3. Chronological framework and comparison between the XRF  $\ln(\text{Ba}/\text{Ti})$  curve of Core P1-03 and the stacked global benthic LR04  $\delta^{18}\text{O}$  (Lisiecki and Raymo, 2005).

framework reconstruction. By graphical comparing the P1-03 XRF  $\text{Ba}/\text{Ti}$  curve with the LR04  $\delta^{18}\text{O}$  stack, Core P1-03 spans the last 513 ka covering the Marine Isotope Stage (MIS) 1–13 (Fig. 3).

### 3.2. Biogenic opal content and XRF $\ln(\text{Si}/\text{Al})$

Both the biogenic opal contents (%) and the XRF  $\ln(\text{Si}/\text{Al})$  (Fig. 4) are characterized by pronounced peaks during each glacial termination. They are very similar to the deglacial opal peaks in the record from the subtropical North Atlantic, at Ocean Drilling Program (ODP) Site 658 (20°44'57"N, 18°34'51"W, 2263 m water depth) (Tiedemann et al., 1989; Meckler et al., 2013). The biogenic opal contents are characterized by a relatively wide range of parameter values (3.67–21.66), with average values of 9.17, similar with the core ODP-658 average values of 10.61. Besides the deglacial opal peaks, the biogenic opal and  $\ln(\text{Si}/\text{Al})$  generally display clear glacial/interglacial oscillation with higher values during interglacials.

### 3.3. XRF $\ln(\text{Ca}/\text{Ti})$

The XRF  $\ln(\text{Ca}/\text{Ti})$  is used as a proxy for carbonate preservation, and the values span from 0.24 to 1.82 with an average 0.57 (Fig. 4J). The low values of  $\ln(\text{Ca}/\text{Ti})$  during glacial terminations are supposed to correspond to the dramatic increasing of Epica Dome C

$\text{CO}_2$  concentration in Fig. 4 (Siegenthaler et al., 2005).

### 3.4. Sand fraction of sediment

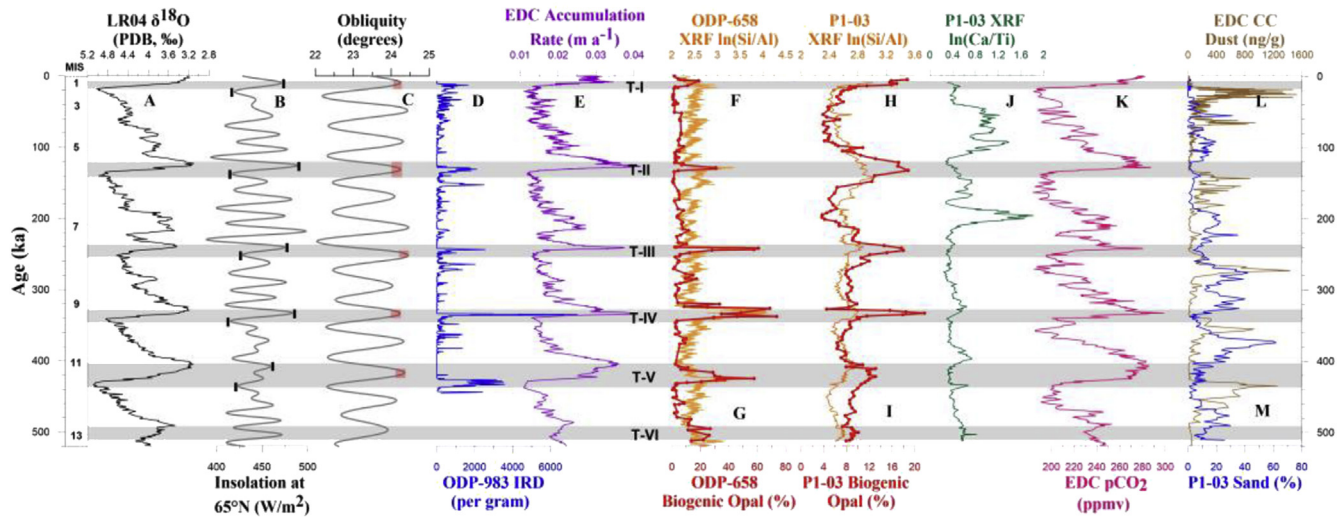
We use the sand fraction data to evaluate the terrestrial influence during glacial terminations (Fig. 4M). This data is characterized by a wide range of values (0–61.74%), with average values of 12.08%. There is no significant correlation between sand fraction and biogenic opal contents, implying the terrestrial influence on biogenic opal could be ignored. Especially the relative low values in glacial terminations, indicating that the deglacial opal peaks are not terrestrial sourced.

## 4. Discussion

### 4.1. Deglacial opal peaks and enhanced Southern Ocean upwelling

The biogenic opal concentration of core P1-03 is characterized by pronounced peaks during each glacial termination (Fig. 4I), indicating a maxima in opal production. Meanwhile, it is noted that the  $\ln(\text{Si}/\text{Al})$  record varies in line with the deglacial opal peaks (Fig. 4H). On one hand, opal preservation efficiency in the SO is not significantly different from the global average (2–6%; DeMaster, 2002; Nelson et al., 2002; Pondaven et al., 2000); on the other





**Fig. 4.** Records of (A) The stacked global benthic LR04  $\delta^{18}\text{O}$  (Lisiecki and Raymo, 2005). (B) 21 July insolation at  $65^\circ\text{N}$  (Berger, 1978). Black bars high light the highest and lowest insolation value bounding each major termination. (C) Obliquity (Berger, 1978). Red shading indicates the timing of opal peaks from P1-03. (D) Ice-rafted debris (IRD) record from ODP 983 in the northeast Atlantic (Barker et al., 2015). (E) Accumulation Rate at Epica Dome C (Bazin et al., 2013). (F)  $\text{In}(\text{Si}/\text{Al})$  record from ODP 658 (Meckler et al., 2013). (G) Biogenic opal record from ODP 658 (Tiedemann et al., 1989). (H)  $\text{In}(\text{Si}/\text{Al})$  record from P1-03. (I) Biogenic opal concentration from P1-03. (J)  $\text{In}(\text{Ca}/\text{Ti})$  record, a proxy for carbonate preservation, from P1-03. (K)  $\text{pCO}_2$  record from EDC (Siegenthaler et al., 2005). (L) Coulter Counter dust mass concentration from EDC (Lambert et al., 2008). (M) Sand fraction record from P1-03. Grey shading indicates glacial terminations I to VI, MIS 1 to MIS 13 are interglacial periods. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

hand, the glacial-interglacial variation of biogenical opal coincides with primary productivity as inferred from XRF  $\text{In}(\text{Ba}/\text{Ti})$  (Fig. 3). All together, this confirms that the sedimentary opal variation represents production changes rather than preservation efficiency. In addition, it is worthy noting that there is no significant correlation between Sand/EDC CC Dust and our biogenic opal record (Lambert et al., 2008), especially during glacial terminations (Fig. 4 L and M). This indicates that the influence of terrigenous and airborne dust on the opal record is trivial.

We attribute the deglacial opal peaks to the enhanced supply of dissolved Si to thermocline and surface water in upwelling regions of the MOC lower cell, where upwelled silicate rich deep-water masses were at its maximum during deglaciation. The enhanced ventilation associated with increased upwelling in the SO, would lead to an increase in both Si and oceanic Fe supply to surface waters and promote opal production (Ayers and Strutton, 2013). Our results, consistent with other opal flux records in Southern Ocean (Chase et al., 2003; Anderson et al., 2009), indicate the key role of the upwelling branch on nutrient supply to the surface as emphasized in the latest MOC diagram (Marshall and Speer, 2012).

#### 4.2. Impact on low-latitude surface and thermocline waters

Enhanced upwelling in the SO during deglaciation would have introduced the Si-rich signature during the formation of Antarctic Intermediate Water (AAIW) and Subantarctic Mode Water (SAMW), which are known to feed the lower thermocline in the low latitude (Sarmiento et al., 2004). If the deglacial opal peaks in the lower cell were truly a chemical signature of enhanced MOC upwelling branch of the SO (Fig. 4 H and I), an attendant maximum supply of dissolved Si and the growth of diatoms in low-latitude regions would be expected. We use the opal production at a site in subtropical North Atlantic to test this prediction (Fig. 1). A clear deglacial correlation between the high-latitude SO and low-latitude are found in the similar time range (Fig. 4; Meckler et al., 2013).

Coincidentally, enhanced supply of southern sourced Si-rich water drove the major ecological changes observed in the other

low-latitude area during glacial terminations (Kienast et al., 2006; Bradtmiller et al., 2006, 2009; Anderson et al., 2009; Pahnke et al., 2008; Calvo et al., 2011; Hendry and Robinson, 2012), indicating a pervasive opal peaks during deglaciation. Other evidence also indicates the increased Southern Ocean upwelling and deep-water injection into low-latitude thermocline during glacial terminations, such as  $\Delta^{14}\text{C}$  record from Southeast Pacific (Siani et al., 2013), Nd isotope record from the Tobago Basin and East Equatorial Pacific (Pahnke et al., 2008; Pena et al., 2013), widespread minimum in  $\delta^{13}\text{C}$  of planktonic foraminifera (Spero and Lea, 2002),  $\delta^{13}\text{C}$  record of atmospheric  $\text{CO}_2$  derived from the Taylor Dome ice core (Smith et al., 1999).

However, it has also been speculated that the Si-rich water feeding the low-latitude opal peaks originated from the upward mixing of abyssal water (Meckler et al., 2013). But the similarity of the deglacial opal peaks in the low-latitude Atlantic and Pacific Oceans would require a process that could operate in both basins despite the different deep-water mass configurations (Bradtmiller et al., 2006, 2007). Moreover, the deglacial  $\text{CO}_2$  outgassing that spread via AAIW pathway (partly) (Siani et al., 2013), could be used to argue against the alternative deep mixing mechanism. So we suggest that increase in the silicate concentration or formation rate of SAMW or AAIW could play a more important role than this coexist mechanism.

#### 4.3. Link to Northern Hemisphere summer insolation and atmospheric $\text{CO}_2$

Since Termination V, each glacial termination occurs when boreal summer insolation intensity begins to rise at a low value, with much of the remainder occurring during the rising limb of the insolation curve (Fig. 4B). This means not only the termination is strongly correlated with the rising boreal summer insolation, but also rising insolation plays a key role in driving the termination to completion. This hypothesis is supported by empirical evidence from the stalagmite (Cheng et al., 2009) as well as climate modeling (He et al., 2013). The amount and rate of insolation rising may also be important controls on ice sheets (Cheng et al., 2009), then

influence the Southern Ocean upwelling. Supporting this idea is the observation that the two terminations with low opal peaks (T-III and T-V) are associated with relatively low insolation shifts. In contrast, the terminations with high opal peaks (T-I, T-II and T-IV) are associated with high magnitudes. The Termination VI, with the insolation shift and decline during this period, has the least significant opal peak value (Fig. 4 B,I). Because of the resolution of our age model and the graphical comparing with the orbital tuned LR04  $\delta^{18}\text{O}$  stack, it's a pity that we don't have a chance to get the precise evaluation on "which is the driving force", especially for the completely orbital forcing driven insolation. However, we could suggest that there is a close relationship between the boreal summer insolation and SO upwelling, by the good correlation between these parameters.

Increased upwelling in the Southern Ocean, inferred from the opal records, coincided with the deglacial rise in atmospheric  $\text{CO}_2$  (Fig. 4K). During deglaciation, the weakest carbonate preservation in the deep Southern Ocean derived from XRF In(Ca/Ti), was consistent with  $\text{CO}_2$  increasing, and the maximum in the rate of  $\text{CO}_2$  rise was synchronous with the opal peaks (Fig. 4I). These lead us to conclude that enhanced upwelling of the Southern Ocean was a primary contributor to the deglacial rise in atmospheric  $\text{CO}_2$ . Our results also support the interpretation with respect to the  $\text{SO}^{14}\text{C}$ -depleted deep water during LGM and its injection into intermediate waters during deglaciation, in line with rising atmospheric  $\text{CO}_2$  (Marchitto et al., 2007; Sikes, 2012).

#### 4.4. Termination mechanisms

Along with the growing realization of the key role played by the upwelling branch of the MOC on climate system, the Southern Ocean becomes a focus to understand processes associated with modern and ancient climate variability (Huybers and Wunsch, 2010; Marshall and Speer, 2012). A series of mechanisms have been proposed to explain the central role played by the upwelling branch of the Southern Ocean on regulating the glacial-interglacial variation of atmospheric  $\text{CO}_2$  changes. First, Northern Hemisphere cooling during glacials could cause a reorganization of global atmospheric circulation, leading to a southward movement in the Southern Hemisphere westerlies (Law et al., 2008; Toggweiler and Russell, 2008; Govin et al., 2009; Anderson et al., 2009; Abarzúa, 2012; Lamy, 2012). This facilitates the upwelling branch of the MOC and enhances the ventilation between the abyss and the surface, promoting atmospheric  $\text{CO}_2$  increase, and resulting in observed productivity peaks (Anderson et al., 2009). This mechanism is further supported by the coincident southward shift/intensification of the Southern Hemisphere westerlies inferred from EDC accumulation rate record (Bazin et al., 2013; WAIS, 2013; Fig. 4E) during terminations. Second, the bipolar seesaw mechanism (Stocker and Johnsen, 2003). A cessation of the MOC in the Northern Hemisphere induced by freshwater injection owing to the North Atlantic/Arctic ice sheet retreat (Zhang et al., 2013), supported by IRD record in the North Atlantic (Barker et al., 2015; Fig. 4D), could warm the southern hemisphere. As a consequence, this will cause sea-ice retreat in the Southern Ocean, promoting  $\text{CO}_2$  outgassing (Keeling and Stephens, 2001). Third, sea-ice cover (Stephens and Keeling, 2000). Retreated sea ice cover in deglacial times could have allowed winds to drive air-sea exchange more efficiently between deep and surface waters, thereby increased the concentration of atmospheric  $\text{CO}_2$  (Cheng et al., 2009; Marshall and Speer, 2012). This effect would be amplified by sea-ice/albedo feedback. Fourth, Antarctic local orbital forcing might act together with the Northern Hemisphere during the terminations, by caused the sea-ice decline and the increased interhemispheric temperature gradient (Laepple et al., 2011; WAIS, 2013).

Our results show that opal peaks occur in line with Northern Hemisphere summer insolation and IRD peaks during terminations. We agree with the scenario in which a set of mechanisms is ultimately linked to the rise in boreal summer insolation that eventually causes  $\text{CO}_2$  rise. The rising insolation may trigger the initial disintegration of the massive ice sheet, which in turn stimulates a weakening of MOC as well as a cold anomaly in the Northern Hemisphere. The resulting warming in the Southern Ocean, associated with bipolar thermal seesaw, causes the sea ice retreat and changes in Southern Westerlies, promoting  $\text{CO}_2$  outgassing and glacial terminations (Broecker et al., 1985; Broecker, 1998; Crowley, 1992; Stocker et al., 1992; Stocker and Johnsen, 2003; Blunier and Brook, 2001; Wang et al., 2007; Toggweiler and Russell, 2008; Anderson et al., 2009).

## 5. Conclusions

In this study we report a record covering the last ~500ka from Prydz Bay in south of Antarctic Divergence Zone (i.e. upwelling regions of the lower cell of the global MOC). Within the age uncertainty, the opal peaks in our records are identified to be accompanied with  $\text{CO}_2$  rise during the last 6 terminations. Our results thus provide additional evidence to support the role played by the Southern Ocean upwelling on  $\text{CO}_2$  outgassing.

The chronology of our records cannot provide precise phase comparison with the core from subtropical North Atlantic, and thus can not support/role out the role of enhanced vertical diffusion in the tropics on  $\text{CO}_2$  outgassing. High resolution records with more reliable age model are thus still required to evaluate their roles on  $\text{CO}_2$  outgassing in the future. Due to the limitation of paleomagnetic analysis in the high latitude, we give up building age model via comparison of relative paleointensity (RPI) between P1-03 and global reference stack in this study. Fortunately, the relative abundance of diatom *Eucampia antarctica* — a sea surface temperature proxy, has the potential to build a reliable chronological framework by comparison to the Antarctic ice core local temperature reconstruction (Jaccard et al., 2016). We are planning to apply this approach in subsequent research, and compare the new age framework to this study.

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