Photic zone changes in the north-west Pacific Ocean from MIS 4–5e

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Abstract. In comparison to other sectors of the marine system, the palaeoceanography of the subarctic North Pacific Ocean is poorly constrained. New diatom isotope records of \(\delta^{13}C\), \(\delta^{18}O\), \(\delta^{30}Si\) (\(\delta^{13}C_{\text{diatom}}, \delta^{18}O_{\text{diatom}}, \text{and } \delta^{30}Si_{\text{diatom}}\)) are presented alongside existing geochemical and isotope records to document changes in photic zone conditions, including nutrient supply and the efficiency of the soft-tissue biological pump, between Marine Isotope Stage (MIS) 4 and MIS 5e. Peaks in opal productivity in MIS 5b/c and MIS 5e are both associated with the breakdown of the regional halocline stratification and increased nutrient supply to the photic zone. Whereas the MIS 5e peak is associated with low rates of nutrient utilisation, the MIS 5b/c peak is associated with significantly higher rates of nutrient utilisation. Both peaks, together with other smaller increases in productivity in MIS 4 and 5a, culminate with a significant increase in freshwater input which strengthens/re-establishes the halocline and limits further upwelling of sub-surface waters to the photic zone. Whereas the MIS 5e peak is associated with low rates of nutrient utilisation, the MIS 5b/c peak is associated with significantly higher rates of nutrient utilisation. Both peaks, together with other smaller increases in productivity in MIS 4 and 5a, culminate with a significant increase in freshwater input which strengthens/re-establishes the halocline and limits further upwelling of sub-surface waters to the photic zone. Whilst \(\delta^{30}Si_{\text{diatom}}\) and previously published records of diatom \(\delta^{15}N\) (\(\delta^{15}N_{\text{diatom}}\)) (Brunelle et al., 2007, 2010) show similar trends until the latter half of MIS 5a, the records become anti-correlated after this juncture and into MIS 4, suggesting a possible change in photic zone state such as may occur with a shift to iron or silicon limitation.

1 Introduction

The modern-day subarctic north-west Pacific Ocean represents a major component of the global oceanic system acting as the one of the terminuses of the deep water thermohaline circulation. Today high precipitation and low evaporation in the region maintain a year-round halocline in the water column (water depth = 100–150 m), reinforced in the summer/early autumn months by the presence of a seasonal thermocline (water depth = 50 m) (Emile-Geay et al., 2003; Antonov et al., 2010; Locarnini et al., 2010). This stratification exerts a major impact on the regional ocean by limiting the mixing of surface waters with underlying nutrient-rich and carbon-rich deep water and by preventing convection and formation of North Pacific Deep Water (Emile-Geay et al., 2003; Menviel et al., 2012).

The initial development of the halocline and stratified water column has been attributed to the onset of major Northern Hemisphere glaciation (NHG) at 2.73 Ma, which increased the flux of freshwater to the region, via increased monsoonal rainfall and/or glacial meltwater, and sea surface temperatures (SSTs) (Sigman et al., 2004; Haug et al., 2005; Swann et al., 2006; Nie et al., 2008). The decrease of abyssal water upwelling associated with this may have contributed to the establishment of globally cooler conditions and the expansion of glaciers across the Northern Hemisphere from 2.73 Ma (Haug et al., 2005). Whilst the halocline appears to have prevailed through the late Pliocene and early Quaternary glacial–interglacial cycles (Swann, 2010), other studies have shown that the stratification boundary may have broken down in the late Quaternary at glacial terminations and during the early part of interglacials (Sarnthein et al., 2004; Jaccard et al., 2005, 2009, 2010; Galbraith et al., 2007, 2008; Gebhardt et al., 2008; Brunelle et al., 2010; Kohfeld and Chase, 2011).

Developing a complete understanding of the nature of regional stratification in the subarctic North Pacific Ocean is important for a number of reasons. Firstly, the palaeoceanographic history of the region remains poorly constrained relative to other sectors of the global ocean. Secondly, with evidence of a pervasive link between the subarctic Pacific and Southern oceans (Haug et al., 2005; Jaccard et al., 2005, 2009, 2010; Galbraith et al., 2007, 2008; Gebhardt et al., 2008; Brunelle et al., 2010; Kohfeld and Chase, 2011).
2010) records from the former can be used to further investigate teleconnections between these regions (Haug and Sigman, 2009; Sigman et al., 2010). Thirdly, with subsurface waters in the ocean interior rich in carbon and nutrients (Gallbraith et al., 2007; Gebhardt et al., 2008; Menven et al., 2012), any weakening/removal of the halocline has potential implications for the regional soft-tissue biological pump and ocean–atmospheric exchanges of CO$_2$.

To further understand the subarctic north-west Pacific Ocean, diatom isotope measurements of $\delta^{13}$C, $\delta^{18}$O, $\delta^{30}$Si ($\delta^{13}$C$_{diatom}$, $\delta^{18}$O$_{diatom}$, and $\delta^{30}$Si$_{diatom}$) are presented here from the open waters of ODP Site 882 between Marine Isotope Stage (MIS) 4 and MIS 5e (Fig. 1). Existing research from the region has revealed two periods of elevated opal concentration in this interval alongside large changes in proxies relating to nutrient supply and utilisation (Jaccard et al., 2005, 2009; Brunelle et al., 2007, 2010). The new diatom isotope data presented here will allow the changes in photic zone conditions and the response of the soft-tissue biological pump to be further constrained. Diatoms, unicellular siliceous algae, are ideally suited for this purpose as they (1) occupy the uppermost sections of the water column above the halocline, (2) dominate export production in high-latitude and upwelling zones (Nelson et al., 1995), and (3) represent the halocline, (2) dominate export production in high-latitude (1) occupy the uppermost sections of the water column above the halocline, (2) dominate export production in high-latitude and upwelling zones (Nelson et al., 1995), and (3) represent a key component of the soft-tissue biological pump to be further constrained. Diatoms, unicellular siliceous algae, are ideally suited for this purpose as they

2 Methods

ODP Site 882 is located on the western section of the Detroit Seamounts at a water depth of 3244 m (50°22’N, 167°36’E) (Fig. 1). The age model used in this study is derived from the astronomical calibration of high-resolution GRAPE density and magnetic susceptibility measurements with linear interpolation between selected tie points (Jaccard et al., 2009). Ages are constrained by two radiocarbon dates and verified by correlating magnetic susceptibility and benthic foraminifera $\delta^{18}$O records from ODP Sites 882 and 883. Samples were prepared for diatom isotope analysis using techniques previously employed at this site (Swann et al., 2006, 2008) with the 75–150 µm fraction analysed. Diatom biovolumes, calculated following Hillebrand et al. (1999) and Swann et al. (2008), show that samples in this fraction are dominated by a single taxa Coscinodiscus radiatus (Ehrenb.) (Fig. 2) which blooms throughout the year with elevated fluxes often occurring in autumn/early winter (Takahashi, 1986; Takahashi et al., 1996; Onodera et al., 2005). Consequently, the diatom isotope measurements obtained here are interpreted as primarily reflecting annually averaged conditions with a slight bias towards autumn/early winter months. Smaller size fractions which contain a greater diversity of taxa were not analysed due to the potential for vital effects in $\delta^{18}$O$_{diatom}$ (Swann et al., 2008). Sample purity was assessed for all samples using light microscopy and SEM with unclean samples disregarded for isotope analysis. Both techniques show the excellent preservation of diatoms in the sediment record and suggest that issues of dissolution/diagenesis are not relevant to this study.

$\delta^{18}$O$_{diatom}$ and $\delta^{30}$Si$_{diatom}$ were analysed following a combined step-wise fluorination procedure at the NERC Isotope Geoscience Laboratory (UK) (Leng and Sloane, 2008) with measurements made on a Finnigan MAT 253 and values converted to the VSMOW and NBS28 scale respectively using the NIGL within-run laboratory diatom standard BFCmod which has been calibrated against NBS28. A small subset of the $\delta^{18}$O$_{diatom}$ data was previously published as part of an investigation in $\delta^{18}$O$_{diatom}$ vital effects in Swann et al. (2008) (see Table S1 in the Supplement). Where sufficient material remained following $\delta^{18}$O$_{diatom}$ and $\delta^{30}$Si$_{diatom}$ analysis, samples were analysed for $\delta^{13}$C$_{diatom}$ using a Costech elemental analyser linked to an Optima mass spectrometer via cold trapping (Hurrell et al., 2011). Replicate analyses of sample material across the analysed interval indicate an analytical reproducibility (1σ) of 0.4‰, 0.06‰ and 0.3‰ for $\delta^{18}$O$_{diatom}$, $\delta^{30}$Si$_{diatom}$ and $\delta^{13}$C$_{diatom}$ respectively.

3 Results

Through the analysed interval, $\delta^{13}$C$_{diatom}$ largely follows previously published siliceous productivity (opal) records from the region in indicating two intervals of higher productivity from 130 to 114 ka BP (MIS 5e) and from 101 to 86 ka BP (MIS 5b/c) (Jaccard et al., 2005, 2009) (Fig. 3). Before/after each of these intervals $\delta^{13}$C$_{diatom}$ is lower at $< -18$‰. These trends are also largely mirrored by the $\delta^{30}$Si$_{diatom}$ and $\delta^{13}$N$_{diatom}$ records of diatom-bound nitrogen ($\delta^{13}$N$_{diatom}$) (Brunelle et al., 2007, 2010) records of nutrient supply/utilisation, except during MIS 5e when values for both remain low and comparable to those in MIS 5d (Fig. 3).
4 Discussion

4.1 Environmental controls on diatom isotopes

Given the limited number of published diatom isotope records in palaeoceanography, the section below summarises the main controls on δ18O<sub>diatom</sub>, δ30Si<sub>diatom</sub> and δ13C<sub>diatom</sub>. Diatom isotopes act as an alternative proxy to records from planktonic foraminifera at sites, such as ODP Site 882, depleted in carbonates. Measurements of δ18O<sub>diatom</sub> can be interpreted in the same way as those of planktonic foraminifera (δ18O<sub>foram</sub>) (Swann and Leng, 2009) with variations linked to changes in temperature (−0.2‰°C<sup>−1</sup>) (Brandriss et al., 1998; Moschen et al., 2005; Dodd and Sharp, 2010; Crespin et al., 2010) and surface water δ18O (δ18O<sub>water</sub>). During biomineralisation diatoms uptake silicon, in the form of silicic acid (H<sub>4</sub>SiO<sub>4</sub>), with the lighter 28Si preferentially used over 29Si and 30Si. With an enrichment factor independent of temperature, the concentrations of CO<sub>2</sub> in the water (pCO<sub>2aq</sub>) and other vital effects (De La Rocha et al., 1997; Milligan et al., 2004), δ30Si<sub>diatom</sub> reflects changes in photic zone silicic acid utilisation which is regulated by the biological demand for silicic acid, the rate at which nutrients are supplied to the photic zone and the δ30Si composition of the silicic acid substrate (δ30Si<sub>dSi</sub>) (De La Rocha, 2006; Reynolds et al., 2006).

A number of studies have examined the controls on δ13C<sub>diatom</sub> on carbon from bulk cellular diatom organic material including the cytoplasm. Whilst palaeoenvironmental reconstructions solely analyse the cell wall, which is preserved in the sediment and protected from dissolution by the diatom frustule (Abramson et al., 2009), it is assumed that the controls on cell wall δ13C<sub>diatom</sub> are similar to those for bulk δ13C<sub>diatom</sub> as the cell-wall organic matter forms a key template for diatom biomineralisation (Hecky et al., 1973; Swift and Wheeler, 1992; Kröger et al., 1999; Sumper et al., 2004). During photosynthesis, organic carbon matter is formed from HCO<sub>3</sub>− and CO<sub>2aq</sub> (Tortell et al., 1997) using both active and indirect transportation mechanisms (Sültemeyer et al., 1993) and C<sub>3</sub> and C<sub>4</sub> photosynthetic pathways (Reinfelder et al., 2000). Marine studies including those from the Bering Sea and North Pacific Ocean have demonstrated that the majority of diatom carbon originates from HCO<sub>3</sub>− via direct transportation (Tortell and Morel, 2002; Cassar et al., 2004; Martin and Tortell, 2006; Tortell et al., 2006, 2008). Although HCO<sub>3</sub>− : CO<sub>2aq</sub> uptake ratios may alter with inter-species variations in cell morphologies (Martin and Tortell, 2008), no link exists with changes in pCO<sub>2aq</sub>Fe availability, growth rates, primary productivity or frustule area : volume ratios (Cassar et al., 2004; Martin and Tortell, 2006; Tortell et al., 2006, 2008).

With 12C preferentially fractionated over 13C (Laws et al., 1995), δ13C<sub>diatom</sub> predominantly reflects changes in photosynthetic carbon demand driven by variations in biological productivity or carbon cellular concentrations. Smaller magnitude variations in δ13C<sub>diatom</sub> may then arise with changes in the composition of the dissolved inorganic carbon substrate (δ13C<sub>DIC</sub>) and through the intracellular and extra-cellular balance of CO<sub>2</sub> with an increase in photic zone pCO<sub>2aq</sub> reducing δ13C<sub>diatom</sub> (Laws et al., 1995; Rau et al., 1996, 1997). Whilst questions remain over the potential for δ13C<sub>diatom</sub> to be impacted by changes in HCO<sub>3</sub>− : CO<sub>2</sub> uptake, growth rates, amino acid composition, cell morphology as well as the diffusion of carbon into the cell by the enzyme RuBisCO (Laws et al., 1995, 1997, 2002; Rau et al., 1996,
4.2 Changes in the regional biological pump (MIS 5e to MIS 5b)

Previously published opal concentration data (Jaccard et al., 2009) together with δ\textsuperscript{13}C\textsubscript{diatom} data from this study indicates two intervals of high siliceous productivity at ODP Site 882 through the analysed interval: the first from 130 to 114 ka BP corresponding to the last interglacial (MIS 5e: 130–116 kaBP), the second from 101 to 86 kaBP covering the latter half of MIS 5c (105–93 kaBP) and most of MIS 5b (93–86 kaBP) (Jaccard et al., 2009) (Fig. 3, green shading). Whilst records of biogenic barium (BioBa) capture the MIS 5e peak (Jaccard et al., 2009), they fail to do so with the second flux event. Modern day calibrations have noted the lack of a relationship between BioBa and export production in the region (Serno et al., 2014) and speculated that the mismatch can be attributed to early diagenetic remobilisation of barium following a change in redox state (Gebhardt et al., 2008). On the other hand, all evidence points against an actual preservation/dissolution issue in this BioBa record (see Jaccard et al., 2009) and so, in line with Jaccard et al. (2009), we interpret BioBa as a measure of organic carbon export rather than siliceous productivity. With the isotope records reported here derived from diatoms and the siliceous fraction of the sediment record, we focus our discussion on the opal siliceous productivity record and only used BioBa as a proxy of organic carbon export.

Similar to the Southern Ocean, the modern-day subarctic north-west Pacific Ocean photic zone is largely limited by iron availability (Harrison et al., 2009; Tsuda et al., 2003). Accordingly, increases in bioavailable iron represent a plausible mechanism for explaining the two main (opal inferred) productivity peaks during MIS 5. Today iron supply is thought to primarily occur via aeolian dust deposition originating from East Asia and the Badain Juran Desert.
(Yuan and Zhang, 2006) and other global regions (Hsu et al., 2012). Additional iron is then derived from volcanic activity (Banse and English, 1999), continental margins (Lam and Bishop, 2008), advection of waters from the Okhotsk Sea (Nishioka et al., 2007) and winter mixing of surface/sub-surface water (Shigemitsu et al., 2012). Both productivity peaks occur without a corresponding increase in aeolian dust at “Station 3” (close to ODP Site 882 at 50°00' N, 164°59'E) (Shigemitsu et al., 2007) or in East Asian winter monsoon records from the Chinese Loess Plateau and other marine sites (Sun et al., 2006; Zhang et al., 2009). Whilst a doubling in aeolian dust does occur at “Station 3” during the early stages of MIS 5e, this ceases before any increase in $\delta^{13}C_{\text{diatom}}$, opal or other proxy at ODP Site 882 (Shigemitsu et al., 2007). The absence of a significant increase in bioavailable iron would appear to rule out a major role for iron in driving the two productivity peaks in MIS 5e and MIS 5b/c. This would be in line with evidence indicating that productivity peaks during the last deglaciation across the North Pacific Ocean also occur without a corresponding increase in aeolian dust or other iron source input (Kohfeld and Chase, 2011). Others have also argued that iron only exerts a secondary or minor control on regional water column productivity in the palaeo-record (Kienast et al., 2004; Lam et al., 2013) whilst we are unable to account for possible changes in the flux of bioavailable iron from the Okhotsk Sea, winter mixing and other sources identified above.

4.2.1 Nutrient utilisation and supply

The deep and intermediate waters of the subarctic North Pacific Ocean contain some of the highest nutrient levels in the world (Whitney et al., 2013). Accordingly productivity peaks over glacial–interglacial cycles, including those covered in this study, have been linked to changes in the regional halocline and water column stratification which would alter the advection of nutrient- and carbon-rich sub-surface waters into the photic zone (Jaccard et al., 2005; Gebhardt et al., 2008). A key difference between the two productivity events in MIS 5e and MIS 5b/c is the response of the biological community to raised photic zone nutrient availability. Although productivity is high during MIS 5e, values are low for $\delta^{30}Si_{\text{diatom}}$ at ODP Site 882 (< 1.0 ‰, n = 3) and for $\delta^{15}N_{\text{diatom}}$ (< 6 ‰) at a nearby site (49°72'N, 168°30'E) (Brunelle et al., 2010) (Fig. 3). In contrast during MIS 5b/c the productivity peak is concordant with an increase in $\delta^{30}Si_{\text{diatom}}$ and $\delta^{15}N_{\text{diatom}}$ to ca. 1.2–1.3 ‰ and > 6 ‰ respectively (Fig. 3).

Changes in $\delta^{30}Si_{\text{diatom}}$ may reflect either increased biological uptake of silicic acid (consumption) and/or changes in the supply of silicic acid to the photic zone. The modern-day regional stratified water column is best represented by a closed system model in which a finite amount of silicic acid exists for biomineralisation (Reynolds et al., 2006). In contrast an unstratified water column would be reflected by an open system model with continual supply of silicic acid. By assuming that the two productivity peaks reflect a weakening in the stratification, an open system model can be used to investigate the controls on $\delta^{30}Si_{\text{diatom}}$:

$$\delta^{30}Si_{\text{diatom}} = \delta^{30}Si(OH)_4 + \epsilon \cdot f.$$  

(1)

where $\delta^{30}Si(OH)_4$ is the isotopic composition of dissolved silicic acid supplied to the photic zone, $\epsilon$ is the enrichment factor between diatoms and dissolved silicic acid and $f$ is the fraction of utilised Si(OH)$_4$ remaining in the water. Existing work from the North Pacific Ocean has estimated $\delta^{30}Si(OH)_4$ at 1.23 ‰ and $\epsilon$ as 1.0 (Reynolds et al., 2006). Using changes in Si(OH)$_4$ consumption (Eq. 1) and silicic productivity (opal), the supply of Si(OH)$_4$ into the photic zone can be constrained relative to mean conditions during MIS 5e as

$$\text{Si(OH)}_4_{\text{supply}} = \frac{\text{Opal}_{\text{sample}} / \text{Opal}_{\text{MIS 5e}}}{(1 - f_{\text{sample}} / f_{\text{MIS 5e}})} \times (1 - f)_{\text{MIS 5e}}.$$  

(2)

Estimates of Si(OH)$_4$ consumption and supply from Eqs. (1) and (2) are only applicable for intervals when the water column represents an open system (e.g. the productivity peaks in MIS 5e and MIS 5b/c) and are dependant on modern-day estimates of $\delta^{30}Si(OH)_4$ being representative of past conditions. This assumption is based on evidence that $\delta^{30}Si(OH)_4$ is relatively resilient to change, outside of seasonal biological fluxes, over timescales similar to this study except in extreme circumstances linked to major reductions in the flux of riverine silicon into the ocean (Rocha and Bickle, 2005). The results show that the productivity peaks in MIS 5e and MIS 5b/c are both closely correlated with elevated levels of Si(OH)$_4$ being supplied to the photic zone (Fig. 3), supporting the suggestion that these intervals are linked to a reduction in water column stratification and an increase in the vertical flux of nutrients bearing sub-surface waters into the photic zone (Jaccard et al., 2005; Gebhardt et al., 2008). However, whilst the increase in Si(OH)$_4$ supply in MIS 5b/c is matched by a corresponding increase in biological consumption of Si(OH)$_4$, the ratio of regenerated to preformed nutrients in the ocean interior, the opposite occurs during MIS 5e when the rates of Si(OH)$_4$ consumption are at their lowest over the analysed interval. (Fig. 3). Whilst reduced Si(OH)$_4$ consumption during MIS 5e could be linked to iron limitation, records indicate that aeolian dust deposition was equally low during both the MIS 5e and the MIS 5b/c productivity peaks. However, as before we are unable to account for changes in iron supply from non-aeolian sources.

4.2.2 Implications for $pCO_2$

Understanding the mechanisms that regulate changes in atmospheric concentrations of CO$_2$ ($pCO_2$) remains a key objective in palaeoclimatology. Previous research has demonstrated that the Southern Ocean and low-latitude oceans
act as the dominant source/sink of atmospheric CO$_2$ over glacial–interglacial cycles (Pichevin et al., 2009; Fischer et al., 2010; Sigman et al., 2010). Whilst the North Pacific Ocean does not need to be invoked to explain the full amplitude of glacial–interglacial changes, recent work has advocated a potential role for the region in regulating atmospheric pCO$_2$ over the last termination (Rae et al., 2014).

Today the net annual ocean–atmosphere exchange of CO$_2$ in the subarctic north-west Pacific Ocean is close to zero, but alters from being a sink of atmospheric CO$_2$ in spring to a source in winter (Takahashi et al., 2006; Ayers and Lozier, 2012) (Fig. 4a). This seasonal variability can be attributed to changes in the biological pump and in SST which affects the solubility of CO$_2$ (Honda et al., 2002; Chierici et al., 2006; Ayers and Lozier, 2012). A weakening of the halocline stratification in MIS 5e and MIS 5b/c would have increased the advection of nutrient- and carbon-rich waters from the ocean interior, raising photic zone pCO$_2$ (aq) and the potential for CO$_2$ to be ventilated into the atmosphere due to an air–sea disequilibrium in CO$_2$. This, however, is dependant on the response and relative efficiency of the biological pump in taking advantage of the increased nutrient supply and altering the ratio of regenerated : performed nutrients to re-export carbon into the deep ocean (Sigman et al., 2004, 2010; Marinov et al., 2008). Whilst changes in the temperature/thermocline may also have been important, the only SST record for the region (Martínez-Garcia et al., 2010) does not contain the temporal resolution to investigate this further and does not provide a surface–subsurface depth temperature transect.

During MIS 5e a scenario of both higher pCO$_2$ (aq) and incomplete/low rates of nutrient utilisation suggests the regional ocean could have ventilated CO$_2$ into the atmosphere faster than the soft-tissue biological pump reabsorbed and sequestered CO$_2$ into the deep ocean (Fig. 4b) despite evidence for higher organic carbon export (BioBa) in this period (Fig. 3). The culmination of this interval at the end of MIS 5e would have resulted in the system returning to a stratified state, perhaps similar to the modern-day water column (Fig. 4a) with minimal air–sea fluxes of CO$_2$. In contrast the opal productivity peak in MIS 5b/c is marked by similar levels of photic zone Si(OH)$_4$ supply as in MIS 5e but with ca. 20 % higher rates of Si(OH)$_4$ consumption (Figs. 3 and 4c). The combination of high siliceous productivity (opal/δ$^{13}$C$_{diatom}$) and a highly efficient biological pump ($\delta^{30}$Si$_{diatom}$/δ$^{15}$N$_{diatom}$) during this interval suggests that the net flux of ocean–atmosphere CO$_2$ exchanges arising from the sea–air disequilibrium could have remained close to zero if photosynthetic carbon demand were similar to the rate of sub-surface carbon flux to the photic zone. Whilst a highly efficient soft-tissue biological pump raises the possibility for the region to have acted as a net sink of atmospheric CO$_2$, the potential and significance for this is limited by the relatively low proportion of surface waters which reach the deep ocean interior (Gebbie and Huybers, 2011) and low BioBa in this interval (Fig. 3).

Although the data suggest that changes in the regional photic zone may have contributed to variations in atmospheric pCO$_2$ during MIS 5, both via the soft-tissue biological pump and associated changes in ocean alkalinity, it is not possible to quantify the magnitude of any fluxes or access whether they were accompanied by a change in diatom silicification and cellular Si:C ratios. Firstly, insufficient purified diatom material remains to measure diatom silicon concentrations. Secondly, although diatom elemental carbon measurements obtained during the analysis of $\delta^{13}$C$_{diatom}$ increase from < 0.3 wt% in MIS 5e to ca. 0.4 wt% in MIS 5b–d (see Supplement Table S1), the analytical reproducibility for C$_{diatom}$ is relatively high at 0.1 % (1σ) (Hurrell et al., 2011) and measurements are derived from the cell wall material and not the bulk cellular matter formed.
during photosynthesis. Furthermore, whilst other cores from the region show a similar double peak in opal productivity during MIS 5 (Narita et al., 2002) records at other sites suggest that the second peak is restricted to MIS 5b with no increase in MIS 5c (Shigemitsu et al., 2007). Such discrepancies either suggest poor stratigraphic controls on the age model for either core, or the potential for significant spatial variability across the region and reiterates that the magnitude of any ocean–atmosphere fluxes of CO$_2$ would be low compared to those occurring elsewhere in the marine system such as the Southern Ocean and low-latitude oceans.

4.2.3 Freshwater controls on siliceous productivity

Records show that the decline in siliceous productivity for both intervals culminates with large decreases in $\delta^{18}$O$_{diatom}$ of ca. 3–5‰ from ca. 113 kaBP and 85 kaBP (Fig. 3, red shading). The magnitude of change is too large to be driven by reductions in deep water upwelling or shifts in ocean water masses from both higher and lower latitudes, which would only alter $\delta^{18}$O$_{water}$ by ca. 1‰ (LeGrande and Schmidt, 2006). Instead the drop in $\delta^{18}$O$_{diatom}$ suggests an input of isotopically depleted freshwater that may be similar in origin to events documented at the same site during the late Pliocene/early Quaternary (Swann, 2010).

Although the modern-day regional halocline is maintained by high precipitation and low evaporation (Emile-Geay et al., 2003), it is difficult to envisage a sufficient increase in precipitation to initiate a 3–5% decrease in $\delta^{18}$O$_{diatom}$. This is reiterated by evidence that monsoonal activity was largely stable during MIS 5b–e (Sun et al., 2006; Zhang et al., 2009). At the same time the potential for a glacial source is questioned by evidence indicating a restricted glaciation in north-east Russia, closest to ODP Site 882, at the Last Glacial Maximum (LGM) (Barr and Clark, 2011, 2012), although other work suggests these ice sheets may have been considerably larger prior to the LGM (Bigg et al., 2008; Barr and Solomon, 2014). Recent work has shown that both of the major decreases in $\delta^{18}$O$_{diatom}$ coincide with increases in IRD accumulation in some, but not all, cores from the Okhotsk Sea (Nürnberg et al., 2011). It has also been argued that the regional water column was regulated by significant inputs of meltwater from the North American ice sheets during the last deglaciation (Lam et al., 2013). In either case, the decrease in $\delta^{18}$O$_{diatom}$ at the end of each siliceous productivity peak suggests that inputs of freshwater helped re-establish/strengthen the halocline, limiting the upwelling of nutrient-/CO$_2$-rich sub-surface waters and biological activity. However, with the decrease in $\delta^{18}$O$_{diatom}$ only occurring after the initial decline in productivity, freshwater can only be acting as a secondary control in re-establishing the halocline.

Previous work has suggested a link between changes in the subarctic north-west Pacific Ocean and the Southern Ocean (Jaccard et al., 2005, 2010; Brunelle et al., 2007; Shigemitsu et al., 2007; Galbraith et al., 2008; Sigman et al., 2010). The most viable mechanisms for synchronous changes between polar regions are temperature- and salinity-driven variations in water column density (Brunelle et al., 2007). For example a cooling of polar SST would reduce the rate of sub-surface upwelling into the photic zone (de Boer et al., 2007), lowering nutrient availability and potentially triggering the initial decline in siliceous productivity. At the same time, a decrease in SST would increase the sensitivity of the water column to subsequent changes in salinity, making the region highly vulnerable to inputs of freshwater which would strengthen the water column and inhibit productivity (Sigman et al., 2004). Additional reductions in siliceous productivity may then arise from lower North Pacific overturning and associated deep-water incursions and upwelling in the North Pacific (Schmittner, 2005). Support for a series of events similar to this at ODP Site 882 lies with the concordant decreases at ODP Site 882 between supplied Si(OH)$_4$/$\delta^{30}$Si$_{diatom}$/opal and Antarctic ($\delta$D)/NGRIP ($\delta^{18}$O$_{ice}$) ice-core records (NGRIP; 2004; Jouzel et al., 2007) at the start of each productivity decline from ca. 118 and 89 kaBP respectively (Fig. 3). The final switch to a low productivity system then coincides with the later decreases in $\delta^{18}$O$_{diatom}$ at 113 and 85 kaBP, suggesting that the climatic deterioration associated with lower $\delta$D/$\delta^{18}$O$_{ice}$ may have fuelled the increase in precipitation and/or an advancement of regional glaciers around the North Pacific Basin that triggered the increase in freshwater input. Whilst it remains unclear what initiated either siliceous productivity peak, it can be speculated that reductions in freshwater after 100 kaBP could have weakened the halocline and created the conditions for the second productivity bloom to eventually develop later in MIS 5b/c.

4.3 Photic zone changes from MIS 4-5a

Previous research has documented reduced levels of productivity in the north-west Pacific Ocean during the last glacial in response to surface water stratification (Narita et al., 2002; Jaccard et al., 2005, 2010; Brunelle et al., 2007, 2010; Shigemitsu et al., 2007; Galbraith et al., 2008; Gebhardt et al., 2008). From the latter half of MIS 5a onwards records of $\delta^{15}$N$_{diatom}$ and $\delta^{30}$Si$_{diatom}$/Si(OH)$_4$ consumption become anti-correlated (Fig. 3). Combined with a long-term shift to lower rates of Si(OH)$_4$ supply and higher rates of Si(OH)$_4$ consumption, this supports suggestions that changes in dust/iron inputs in the last glacial may have helped regulate the biological pump by altering the biological demand for individual nutrients (Brunelle et al., 2007, 2010; Galbraith et al., 2008; Shigemitsu et al., 2008), in this case by increasing biological uptake of silicon over nitrogen to the extent that Si(OH)$_4$ consumption in MIS 4 was up to 40% higher than during MIS 5e. Elevated Si(OH)$_4$ consumption may also indicate that the availability of Si(OH)$_4$ rather than iron may have ultimately limited siliceous productivity over...
this interval, in line with a previous suggestion by Kienast et al. (2004).

Superimposed on a trend of low siliceous productivity during MIS 5a and MIS 4 are two small–moderate increases in opal at ca. 76–74 ka BP and ca. 70 ka BP (Fig. 3, green shading). The increase at 70 ka BP does not coincide with any samples analysed in this study, but the increase at 76–74 ka BP coincides with higher $\delta^{13}$C$_{\text{diatom}}$, $\delta^{30}$Si$_{\text{diatom}}$ and Si(OH)$_4$ supply/consumption (open model). Similar to before, both opal peaks culminate with a 2–3 ‰ reduction in $\delta^{18}$O$_{\text{diatom}}$ (Fig. 3, red shading), reiterating the role of freshwater in controlling photic zone dynamics in an era that coincides with increased monsoonal and thus precipitation variability (Sun et al., 2006; Shigemitsu et al., 2007; Zhang et al., 2009). However, whereas the earlier declines in siliceous productivity during MIS 5e and MIS 5b/c are accompanied by reductions in both Si(OH)$_4$ supply and consumption, here the declines initially occur with reduced Si(OH)$_4$ supply and higher rates of Si(OH)$_4$ consumption. This advocates the aforementioned suggestion that the photic zone shifted to a new state from the end of MIS 5a, highlighted by further large changes in Si(OH)$_4$ consumption in MIS 4 that do not coincide with a changes in siliceous productivity or $\delta^{18}$O$_{\text{diatom}}$ (Fig. 3).

5 Conclusions

Results here provide evidence for significant temporal changes in the strength and efficiency of the regional soft-tissue biological pump from MIS 4–5e, altering the ratio of regenerated to preformed nutrients in the water column. In particular the results show evidence of an inefficient soft-tissue biological pump from 124 to 114 ka BP, creating the potential for the region to have played a role in maintaining the warm climate of the last interglacial through the ventilation of oceanic CO$_2$ to the atmosphere. In addition to highlighting temporal changes in the biological pump, the data also reveal that the end of both these and other siliceous productivity fluxes over the analysed interval are linked to significant increases in freshwater input to the region, re-establishing/strengthening the halocline and limiting the subsurface supply of nutrient- and carbon-rich waters to the photic zone. However, further work is needed to resolve the source of these freshwater inputs and the mechanisms responsible for initiating the increase in siliceous productivity and Si(OH)$_4$ supply to the photic zone. Finally, whilst these findings reiterate earlier work in indicating a highly dynamic and changing water column in the subarctic North Pacific Ocean during the last glacial–interglacial cycle, further work is needed to assess the spatial representativeness of these results in other sectors of the subarctic North Pacific Ocean.

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References


Kienast, S. S., Hendy, I. L., Crusius, J., Pedersen, T. F., and Calvert, S. E.: Export production in the subarctic North Pa


