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26th November 2014

Dear Erin,

Please find attached a revised version of our manuscript "Photic zone changes in the North West Pacific Ocean from MIS 4-5e" (ref: cp-2014-107). We thank you for your editorial comments and apologise for the typos and for not integrating some of your comments into our previous version. A summary of the most recent changes are listed below together with a marked-up copy of the manuscript highlighting the changes made within it.

Best wishes,

George

Editor comments (responses in bold)

1) There are some small typos in the text. **We apologise for this and have check the lastest version of the manuscript carefully.**

2) ...I can only see a note to 'assuming modern day estimates of d30Si(OH)4 are valid for the past'... the authors should include the confirmation given in their response document, that this assumption is valid given the processes that we expect ought to change values of $\delta^{30}Si(OH)_4$. **We have included this text in the revised manuscript.**

3) ...the authors posted a 2nd interactive comment which notes the presence of a low resolution SST data set for ODP site 882. This data is not cited in the revised manuscript... **We have included the citation in the revised manuscript and mention the availability of this data and its limitations.**

4) Reviewer 2 sought for clarification that the SST cooling with ONHG was in annual SSTs given that alkenones show a warming. This hasn't been corrected on the revised ms (line 44 p.1) **We apologise for this typo. Reduced SST should have read increased SST and we have amended the text accordingly.**

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 $\delta^{13}C$, $\delta^{18}O$ and $\delta^{30}Si$ ($\delta^{13}C_{diatom}$, $\delta^{18}O_{diatom}$, $\delta^{30}Si_{diatom}$), 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. Whilst $\delta^{30}Si_{diatom}$ and previously published records of diatom $\delta^{15}N$ ($\delta^{15}N_{diatom}$) (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.

Keywords: diatom, freshwater; MIS 5e; nutrient; productivity; stratification

1 Introduction

The modern day subarctic North West Pacific Ocean represent 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 maintains 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., 2009; Locarnini et al., 2009). This stratification exerts a major impact on the regional ocean by limiting the mixing of surface waters with underlying nutrient 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~~ reduced Sea Surface Temperatures (SST) and~~ increased the flux of freshwater to the region, via increased monsoonal rainfall and/or glacial meltwater, and Sea Surface Temperatures (SST) (Sigman et al., 2004; Haug et al., 2005; Swann et al., 2006; Nie et al., 2008). The decrease ~~cessation~~ 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, 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 (Galbraith et al., 2007; Gebhardt et al., 2008; Menviel 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 $\delta^{13}C$, $\delta^{18}O$ and $\delta^{30}Si$ ($\delta^{13}C_{diatom}$, $\delta^{18}O_{diatom}$ and $\delta^{30}Si_{diatom}$) are presented here from the open waters of ODP Site 882 between 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); 3) represent a key component of the soft-tissue biological pump in transferring carbon into the ocean interior by incorporating c. 23.5% of all carbon produced by net primary production into their cellular organic matter (Mann, 1999).

2 Methods

ODP Site 882 is located on the western section of the Detroit Seamounts at a water depth of 3,244 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 BFC_{mod} 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 Supplementary Table 1). 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 analysis 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 130-114 ka BP (MIS 5e) and 101-86 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 $\delta^{15}N$ records of diatom bound nitrogen ($\delta^{15}N_{diatom}$) (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). Following a return to lower values in MIS 5a, all productivity/nutrient proxies show a series of abrupt oscillations that continue into MIS 4 with values in this interval equivalent to the peaks and minima documented in MIS 5b-e.

Measurements of $\delta^{18}O_{diatom}$ can be classified into three stages: 1) periods of relative stability in MIS 5e and MIS 5b-c (124-114 and 102-87 ka BP); 2) periods of significant decreases (≥4‰) in MIS 5d and MIS 5a (113-100 and 85-76 ka BP); 3) periods of increase variability in MIS 5a-4 (75-57 ka BP) (Fig. 3). Intervals of high and stable $\delta^{18}O_{diatom}$ values in MIS 5e and MIS 5b-c coincide with peaks in $\delta^{13}C_{diatom}$ and opal concentrations. The termination of both productivity phases, as indicated by changes in $\delta^{13}C_{diatom}$, $\delta^{15}N_{diatom}$, $\delta^{30}Si_{diatom}$ and opal concentrations, are then concordant with the large reductions in $\delta^{18}O_{diatom}$ during MIS 5d and MIS 5a, suggesting a link between the processes controlling $\delta^{18}O_{diatom}$ and photic zone productivity/nutrient utilisation. This is reinforced by the often synchronous changes between $\delta^{18}O_{diatom}$, $\delta^{30}Si_{diatom}$ and opal concentrations during MIS 5a and into MIS 4.

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 $\delta^{18}O_{diatom}$, $\delta^{30}Si_{diatom}$ and $\delta^{13}C_{diatom}$. Diatom isotopes act as an alternative proxy to records from planktonic foraminifera at sites, such as ODP Site 882, depleted in carbonates. Measurements of $\delta^{18}O_{diatom}$ can be interpreted in the same way as those of planktonic foraminifera ($\delta^{18}O_{foram}$) (Swann and Leng, 2009) with variations linked to changes in temperature (−0.2‰/°C) (Brandriss et al., 1998; Moschen et al., 2005; Dodd and Sharp, 2009; Crespin et al. 2010) and surface water $\delta^{18}O$ ($\delta^{18}O_{water}$). During biomineralisation diatoms uptake silicon, in the form of silicic acid (H_4SiO_4), with the lighter ^{28}Si preferentially used over ^{29}Si and ^{30}Si. With an enrichment factor independent of temperature, the concentrations of CO_2 in the water ($pCO_{2(aq)}$) and other vital effects (De La Rocha et al., 1997; Milligan et al., 2004), $\delta^{30}Si_{diatom}$ 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 $\delta^{30}Si$ composition of the silicic acid substrate ($\delta^{30}Si_{DSi}$) (De La Rocha, 2006; Reynolds et al., 2006).

A number of studies have examined the controls on $\delta^{13}C_{diatom}$ 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 $\delta^{13}C_{diatom}$ are similar to those for bulk $\delta^{13}C_{diatom}$ 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 and Kröger, 2004). During photosynthesis, organic carbon matter is formed from both HCO_3^- and $CO_{2(aq)}$ (Tortell et al., 1997) using both active and indirect transportation mechanisms (Sültemeyer et al., 1993) and C_3 and C_4 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_3^- via direct transportation (Tortell and Morel, 2002;

Cassar et al., 2004; Martin and Tortell, 2006, Tortell et al., 2006, 2008). Although $HCO_3^-:CO_{2(aq)}$ uptake ratios may alter with inter-species variations in cell morphologies (Martin and Tortell, 2008), no link exists with changes in $pCO_{2(aq)}$, 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 ^{12}C preferentially fractionated over ^{13}C (Laws et al., 1995), $\delta^{13}C_{diatom}$ predominantly reflect changes in photosynthetic carbon demand driven by variations in biological productivity or carbon cellular concentrations. Smaller magnitude variations in $\delta^{13}C_{diatom}$ may then arise with changes in the composition of the dissolved inorganic carbon substrate ($\delta^{13}C_{DIC}$) and through the intracellular and extra-cellular balance of CO_2 with an increase in photic zone $pCO_{2(aq)}$ reducing $\delta^{13}C_{diatom}$ (Laws et al., 1995; Rau et al., 1996, 1997). Whilst questions remain over the potential for $\delta^{13}C_{diatom}$ to be impacted by changes in $HCO_3^-:CO_2$ 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, 1997, 2001; Popp et al., 1998; Cassar et al., 2006), many of these physiological processes as well as the impact of inter-species vital effects (Jacot Des Combes et al., 2008) can be partially circumvented by analysing samples comprised of a single taxa. Consequently, ~~W~~with samples in this studies overwhelmingly dominated by *C. radiatus*, changes in $\delta^{13}C_{diatom}$ are primarily interpreted as reflecting changes in photic zone productivity (Fig. 2). We argue that the impact of a changes in $\delta^{13}C_{DIC}$ ~~arising from variations in $pCO_{2(aq)}$ or $\delta^{13}C_{DIC}$ with negligible/minimal influence~~ is negligible due to the aforementioned evidence that $\delta^{13}C_{DIC}$ exerts only a minimal impact on $\delta^{13}C_{diatom}$, likely within analytical error, although the lack of carbonates in the sediments prevents an independent $\delta^{13}C$ record being established to prove this beyond doubt . ~~(Fig. 2)~~. Similarly we argue that higher $\delta^{13}C_{diatom}$ values in MIS 5e, when higher $pCO_{2(aq)}$ should have acted to reduce $\delta^{13}C_{diatom}$, points towards $pCO_{2(aq)}$ not exerting a significant control on $\delta^{13}C_{diatom}$, although we are aware of the circular reasoning with this argument.

4.2 Changes in the regional biological pump (MIS 5e to MIS 5b)

Previously published opal concentration data (Jaccard et al. 2009) together with $\delta^{13}C_{diatom}$ data from this study indicates two intervals of high siliceous productivity at ODP Site 882 through the analysed interval, the first from 130-114 ka BP corresponding to the last interglacial (MIS 5e: 130-116 ka BP), the second from 101-86 ka BP covering the latter half of MIS 5c (105-93 ka BP) and most of MIS 5b (93-86 ka BP) (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. ~~Assessing the accuracy of BioBa as a measure of export production is beyond the scope of this study. However~~ Modern day calibrations ~~others~~ 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 remobilization 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 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., 1999; 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 5c, this ceases before any increase in $\delta^{13}C_{diatom}$, 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 Norrh 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 $\delta^{30}Si_{diatom}$ at ODP Site 882 (<1.0‰, n=3) and for $\delta^{15}N_{diatom}$ (<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 $\delta^{30}Si_{diatom}$ and $\delta^{15}N_{diatom}$ to c. 1.2-1.3‰ and >6‰ respectively (Fig. 3).

Changes in $\delta^{30}Si_{diatom}$ 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 $\delta^{30}Si_{diatom}$:

$$\delta^{30}\,Si_{diatom} = \delta^{30}\,Si\left(OH\right)_4 + \varepsilon * f \tag{Eq. 1}$$

where $\delta^{30}Si(OH)_4$ is the isotopic composition of dissolved silicic acid supplied to the photic zone, ε 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 ε as 1.0 (Reynolds et al. 2006). Using changes in $Si(OH)_4$ consumption (Equation 1) and siliceous productivity (opal), the supply of $Si(OH)_4$ into the photic zone can be constrained relative to mean conditions during MIS 5e as:

$$Si\left(OH\right)_{4\ supply} = \frac{Opal_{sample} / Opal_{MIS\,5e}}{\left(1 - f^{sample}_{consumed}\right) / \left(1 - f^{MIS5e}_{consumed}\right)} \tag{Eq. 2}$$

Estimates of $Si(OH)_4$ consumption and supply from Equations 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. 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 suggesting 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 $Si(OH)_4$, increasing 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 *p*CO_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 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 atmosphere 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 c. 20% higher rates of $Si(OH)_4$ consumption (Fig. 3, 4c). The combination of high siliceous productivity (opal/$\delta^{13}C_{diatom}$) and a highly efficient biological pump ($\delta^{30}Si_{diatom}$/$\delta^{15}N_{diatom}$) 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 was 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).

~~Whilst~~Although the data suggests 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 c. 0.4 wt.% in MIS 5b-d (see Supplementary Information) 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 suggests 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 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 c. 3-5‰ from c. 113 ka BP and 85 ka BP (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 c. 1‰ (LeGrande et al., 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 in the region (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 Solomina, 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 northern 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 mechanism 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 Atlantic 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 (δD)/NGRIP ($\delta^{18}O_{ice}$) ice-core records (NGRIP, 2004; Jouzel et al., 2007) at the start of each productivity decline from c. 118 and 89 ka BP 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 ka BP, suggesting that the climatic deterioration associated with lower δD/$\delta^{18}O_{ice}$ 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 ka BP 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 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 increase in opal at c. 76-74 ka BP and c. 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 $\delta^{13}C_{diatom}$, $\delta^{30}Si_{diatom}$ and $Si(OH)_4$ supply/consumption (open model). Similar to before, both opal peaks culminate with a 2-3‰ reduction in $\delta^{18}O_{diatom}$ (Fig. 3, red shading), reiterating the role of freshwater in controlling photc 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 $\delta^{18}O_{diatom}$ (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 124-114 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 reveals 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 sub-surface 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 ~~initial~~ 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.

Acknowledgements

Thanks are owed to Eric Galbraith and Sam Jaccard for providing information on the ODP Site 882 age model and previously published BioBa and Opal data. Funding for GEAS was provided in part by a Natural Environment Research Council (NERC) postdoctoral fellowship award (NE/F012969/1). Finally we thank two anonymous reviewers who comments helped to improve this manuscript.

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Supplementary material

Supplementary Table 1: ODP Site 882 $\delta^{13}C_{diatom}$, $\delta^{18}O_{diatom}$ and $\delta^{30}Si_{diatom}$ data together with C_{diatom} and relative species biovolume composition in the analysed samples.

Figure legends

Figure 1: Location of ODP Site 882 in the subarctic North West Pacific Ocean.

Figure 2: Relative diatom species biovolumes in samples analysed for $\delta^{18}O_{diatom}$.

Figure 3: Data from ODP Site 882 showing changes in: 1) productivity [$\delta^{13}C_{diatom}$, BioBa (Jaccard et al., 2005) and opal concentrations (Jaccard et al., 2009)]; 2) nutrient dynamics [$\delta^{30}Si_{diatom}$ and $\delta^{15}N_{diatom}$ (Brunelle et al., 2010)]; 3: modelled $Si(OH)_4$ supply/consumption in an open system model; and 4) freshwater input ($\delta^{18}O_{diatom}$) together with EPICA Antarctic δD (Jouzel et al., 2007) and NGRIP Greenland $\delta^{18}O_{ice}$ (NGRIP, 2004). Changes in the supply/consumption of $Si(OH)_4$ are relative to mean conditions in MIS 5e. Green/red shading indicate the increases in productivity and decreases in $\delta^{18}O_{diatom}$ respectively that are discussed in the text.

Figure 4: Schematic models showing subarctic North West Pacific Ocean conditions for: A) Modern day: halocline water column with nutrient poor surface waters limiting biological export; B) MIS 5e: no halocline and enhanced upwelling of nutrient and carbon rich sub-surface waters leading to increased productivity. Low rates of nutrient utilisation suggest a possible increase in $p\mathrm{CO}_2$ and release of CO_2 to the atmosphere; C) MIS 5b-c: conditions similar to MIS 5e but with higher rates of nutrient consumption and a more efficient

soft-tissue biological pump limiting/preventing ventilation of CO_2.