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1	1 Coupled climate and subarctic Pacific nutrient upwelling over the last 850, 000 year		
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16	Highlights: "Coupled climate and subarctic Pacific nutrient upwelling over the last 850, 000		
17	years"		
18 19	• Increased Bering Sea sea ice caused expansion of intermediate water in glacials		
20	• Intermediate water expansion inhibited deep water nutrient supply to surface waters		
21	• Region contributed to global ocean-air CO ₂ exchange on glacial timescales		
22	• Tight correlation between subarctic nutrient upwelling and global CO2		
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24 25	Abstract		
26	High latitude deep water upwelling has the potential to control global climate over glacial		
27	timescales through the biological pump and ocean-atmosphere CO ₂ exchange. However, there		
28	is currently a lack of continuous long nutrient upwelling records with which to assess this		

mechanism. Here we present geochemical proxy records for nutrient upwelling and glacial 29 North Pacific Intermediate Water (GNPIW) formation in the Bering Sea over the past 850 kyr, 30 which demonstrates that glacial periods were characterised by reduced nutrient upwelling, 31 when global atmospheric CO₂ and temperature were also lowered. We suggest that glacial 32 expansion of sea ice in the Bering Sea, and the simultaneous expansion of low nutrient GNPIW, 33 inhibited vertical mixing and nutrient supply across the subarctic Pacific Ocean. Our findings 34 35 lend support to the suggestion that high latitude sea ice and the resultant intermediate water formation, modulated deep water upwelling and ocean-atmosphere CO₂ exchange on glacial-36 37 interglacial timescales.

38 Keywords: Upwelling; Sea Ice; CO₂; subarctic; Bering Sea; Glacial

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40 **1. Introduction**

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42 Recent studies have shown a sharp increase in both surface ocean subarctic Pacific productivity and CO₂ ventilation during the last deglaciation (Galbraith et al., 2007; Gray et al., 2018), and 43 suggested that the equivalent of \sim 30 ppm atmospheric CO₂ was released into the atmosphere 44 due to increased overturning (Rae et al., 2014). This was proposed to coincide with a collapse 45 of glacial North Pacific Intermediate Water (GNPIW), allowing overturning to bring nutrient 46 and CO₂-rich deep water to the surface (Gray et al., 2018). Although a pervasive link between 47 subarctic Pacific export productivity and Pleistocene glacial-interglacial cycles has been 48 demonstrated (Jaccard et al., 2010, 2005), it is not yet known if this was primarily controlled 49 by changes in iron supply (Galbraith et al., 2008; Praetorius et al., 2015), water column 50 stratification and/or light limitation (Jaccard et al., 2010; Lam et al., 2013) and/or the extent of 51 GNPIW (Knudson and Ravelo, 2015a; Li et al., 2017) and nutrient upwelling (Gray et al., 52 53 2018). If the upwelling of nutrient and CO₂-rich water was the predominant control on subarctic

Pacific productivity variations (Kender et al., 2018), this would imply the region has been an active source of pCO_2 throughout the Pleistocene and not just the last deglacial, because they are coupled with ice core temperature records (Jaccard et al., 2010, 2005). One way to test for GNPIW and its effect on upwelling nutrients is from the Bering Sea, which today contains an active upwelling zone.

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Surface waters in the Bering Sea are fed by the subarctic Pacific Alaskan Stream, which enters 60 61 through passes and straits in the Aleutian arc and circulates northward around the basin, forming the shelf-adjacent Bering Slope Current (BSC) (Stabeno et al., 1999) (Fig. 1). 62 Turbulent eddies in the BSC extend below the surface waters at ~300 m water depth, causing 63 64 upwelling of North Pacific Deep Water (NPDW), which contains some of the highest nutrient concentrations in the marine system (Stabeno et al., 1999). This seasonally raises photic zone 65 CO₂ and fuels a high productivity region at the Bering shelf break, known as the "green belt" 66 (Springer et al., 1996) (Fig. 1). Due to its terminal position in the ocean conveyor and deep 67 bathymetric eddies, any glacial reduction in NPDW upwelling across the subarctic Pacific 68 69 should also be apparent as reduced nutrient availability in the green belt.

Modern North Pacific Intermediate Water (NPIW) is sourced from the Okhotsk Sea and is 70 widely distributed across the North Pacific ocean at a water depth between ~300 - 800 m 71 characterized by salinity minima with density centred at 26.8 $\sigma \theta$ (Talley, 1993; Yasuda, 1997). 72 During the last glacial maximum, intensified NPIW (termed GNPIW) expanded down to 2000 73 74 m (Matsumoto et al., 2002), with previous studies suggesting that the Bering Sea was a key source of GNPIW (Horikawa et al., 2010; Ohkushi et al., 2003) as a result of enhanced brine 75 rejection on the Beringian shelf, following increased sea ice growth since ~900 kyr (Jang et al., 76 77 2017; Kender et al., 2018; Knudson and Ravelo, 2015b). Given its density, the formation of GNPIW from the Bering Sea has the potential to suppress upwelling by preventing eddies in
the BSC from reaching NPDW. Expansion of this low nutrient GNPIW from the Bering Sea
would crucially act to enhance stratification across the subarctic region – not just the smaller
Bering Sea – causing region-wide impact on vertical mixing, nutrient supply and the biological
pump.

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Here, we present a new high resolution long sedimentary nitrogen isotope record ($\delta^{15}N$), in 84 combination with previously published lower resolution results (Kim et al., 2017), from IODP 85 Site U1343 in the Bering Sea green belt (Fig. 1) to assess variation in nitrate utilisation between 86 Marine Isotope Stages (MIS) 1 and 21, where we assume that the sedimentary bulk δ^{15} N is a 87 88 representation of total macronutrient utilisation. To account for changes in export production, we normalise the opal mass accumulation rate (MAR) and produce the first high latitude record 89 of nutrient delivery from deep to surface water (termed the "upwelling index"; see Methods). 90 By semi-quantitatively assessing the offset between the opal MAR and $\delta^{15}N$ in this way, we 91 constrain the impact of local growth rate and iron supply, to reveal long-term variability in 92 93 macronutrient supply from deep water upwelling. We also present a new high-resolution multispecies benthic foraminiferal oxygen isotope record for an improved age model and produce 94 the first assessment of the influence of GNPIW at 2 km water depth in the Bering Sea. From 95 this, we discuss the extent to which oceanographic changes in Bering Sea upwelling and 96 GNPIW production may have contributed towards NPDW upwelling and glacial-interglacial 97 *p*CO₂ changes in the subarctic Pacific over the last 850 kyr. 98

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100 2. Materials and Methods

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IODP Site U1343 (54°33.4'N, 176°49.0'E, water depth 1950 m) is situated on a topographic
high in the Bering Sea green belt adjacent to the northern continental shelf, proximal to the
modern winter sea ice edge and at a depth just below the oxygen minimum zone (Fig. 1).
Marine sediments are composed primarily of fine clays and biogenic material, and are
characteristically distinct from shelf-transported materials (Takahashi et al., 2011).

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108 **2.2.** Foraminiferal isotopes

Foraminiferal preservation is notably good at this site given the shallow North Pacific-wide 109 lysocline which limits carbonate preservation at other Bering Sea sites (Takahashi et al., 2011), 110 providing an excellent opportunity to create a robust high resolution $\delta^{18}O_{U1343}$ age model 111 (Asahi et al., 2016; Kender et al., 2018). In the absence of a single species consistently 112 occurring in all samples, 262 samples of ~100 µg of foraminiferal calcite were measured for 113 δ^{18} O using ten different species of benthic foraminifera (*Cassidulina parkerianus, Cassidulina* 114 teretis, Elphidium batialis, Globobulimina auriculata, Globobulimina pacifica, Islandiella 115 norcrossi, Nonionella labradorica, Uvigerina bifurcata, Uvigerina senticosa and Valvulineria 116 *spp.*) at an average sampling resolution of 0.84 m from 2 - 222 m CCSF-A. Measurements 117 were made using an IsoPrime[®] mass spectrometer with a Multicarb device at the British 118 Geological Survey, where the analytical precision of duplicates was <0.05‰. The stable 119 isotope data were calibrated to the VPDB scale through the international NBS standards. 120 Species-specific offsets from Kender et al. (2018) were applied to fit the data to the most 121 commonly occurring species, E. batialis, and are presented in Figure 2 and Supplementary 122 Table 2. Given that benthic foraminiferal δ^{13} C is sensitive to diagenetic alteration, as calcite 123 dissolution, re-precipitation and overgrowth can alter the isotopic composition of their tests 124

125 (Cook et al., 2016), poor linear regressions between foraminiferal δ^{13} C and both raw δ^{18} O (R² 126 = 0.09, p < 0.001) and δ^{18} O adjusted to *E. batialis* (R² = 0.00, p > 0.05) confirms that diagenetic 127 alteration of foraminiferal shells cannot explain the glacial-interglacial variability in the δ^{18} O 128 (Fig. S1).

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130 **2.3. Age Model, Sedimentation rates and Opal MAR calculations**

Benthic foraminiferal isotope data presented here were combined with previously published 131 data from Site U1343 (Asahi et al., 2016), to create a total of 751 $\delta^{18}O_{U1343}$ data points with an 132 average time step of 1.1 kyr at an average resolution of 0.30 m from 0 - 222 m CCSF-A. The 133 age model, was defined by correlating to the LR04 global composite stack (Lisiecki and 134 135 Raymo, 2005), and constructed by selecting 17 age-depth tie points during periods of rapid isotopic change primarily at deglacial intervals (Table S2). Although there is a possible time 136 lag, where Pacific benthic δ^{18} O records may lag the Atlantic Ocean (Asahi et al., 2016; Lisiecki 137 and Raymo, 2005), the high sample resolution and strong correlation between $\delta^{18}O_{U1343}$ and 138 LR04 (r = 0.82, $p = \langle 0.001 \rangle$) suggests that these are within the expected uncertainty of the age 139 140 model.

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The high sedimentation rate at IODP Site U1343 and a lack of hiatuses in core material provides confidence in the continuity of its record (Asahi et al., 2016; Takahashi et al., 2011). Based on the new composite age model, sedimentation rates were calculated between each age-depth tie point. The average sedimentation rate was 29.15 cm kyr⁻¹, with higher rates during interglacials and a notable sedimentation rate crash at ~630 kyr (likely due to low biogenic opal; Fig S2). These new sedimentation rates were applied to existing opal data (Kim et al., 2014) to calculate an opal MAR record using linear interpolation of the ship dry bulk density data and a ten-point
smoothing average (Takahashi et al., 2011).

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151 **2.4. Bulk sedimentary** δ^{15} N

Bulk samples at an average resolution of 0.81 m CCSF-A, totalling 276 samples between 0 -152 222 m CCSF-A were analysed for δ^{15} N with 500 mg of raw material run on a Carlo Erba 1108 153 elemental analyzer, interfaced to a Thermo Finnigan Delta Plus XP IRMS at the University of 154 California, Santa Cruz, with a precision of 0.15‰ based on both sediment standards and 155 replicates. Results, relative to atmospheric nitrogen (AIR), were combined with previously 156 published data from Site U1343 (Kim et al., 2017), to produce a composite record of 373 data 157 points ($\delta^{15}N_{U1343}$), with an average resolution of 0.60 m between 0 and 222 m CCSF-A; average 158 resolution of 2.3 kyr on the new age model (Fig. S2). 159

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- 161 **2.5. Oceanographic indices**
- 162 **2.5.1.** Δδ¹⁸**O**849 U1343

To assess regional variation and discrepancies between Site U1343 and the deep North Pacific 163 Ocean, a Pacific end-member record from ODP Site 849 (Mix et al., 1995) (0°11'N, 110°31'W; 164 3851 m water depth) was used to remove high frequency and glacial-interglacial variability. A 165 consistent offset of ~0.5 ‰ was found between $\delta^{18}O_{U1343}$ and $\delta^{18}O_{849}$, and so this was added to 166 δ^{18} O_{U1343} first, to ensure offset values reflected only differences in δ^{18} O amplitude. After being 167 linearly interpolated at an interval of 2 kyr (Fig. 2), a three-point smoothing spline was applied 168 to the $\delta^{18}O_{U1343}$, and was subtracted from the similarly processed $\delta^{18}O_{849}$ record ($\Delta\delta^{18}O_{849}$ -169 U1343), following a similar methodology to Knudson & Ravelo (2015a). Lower $\delta^{18}O_{U1343}$ 170

become more similar to those of shallower water Bering Sea Site U1342 at ~800 m water depth (Fig. 1, 3a), which is suggested to be influenced by GNPIW (Knudson and Ravelo, 2015a). Therefore, where $\Delta\delta^{18}O_{849-U1343}$ is approximately 0, Site U1343 is assumed to reflect the same water mass as the deep North Pacific Ocean, which upwells along the Bering Sea shelf edge today (Stabeno et al., 1999). Whereas more positive $\Delta\delta^{18}O_{849-U1343}$ values occur when $\delta^{18}O_{U1343}$ is lower than $\delta^{18}O_{849}$, suggesting a colder and/or more saline intermediate water mass is present at Site U1343, similar to the GNPIW signature at Site U1342.

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9 **2.5.2.** Upwelling Index

To account for the changes in the isotope values of deep North Pacific source water ($\delta^{15}N_{source}$) 180 and gain insights into regional nitrate utilisation, we follow previous work (Galbraith et al., 181 2008; Knudson and Ravelo, 2015b) by subtracting the record of ODP Site 1012 in the eastern 182 tropical North Pacific (Liu et al., 2005) (32°17'N, 118°24'W; 1172 m water depth) which is 183 influenced by waters from the Eastern Tropical North Pacific denitrification zone, and is 184 thought to be a site of near complete nitrate utilisation throughout the Pleistocene (Galbraith et 185 al., 2008). $\delta^{15}N_{1012}$ was linearly interpolated at a time interval of 2 kyr (the average resolution 186 of $\delta^{15}N_{1012}$) and subtracted from the similarly interpolated $\delta^{15}N_{U1343}$, to produce a record driven 187 by nutrient utilisation for Site U1343 ($\Delta \delta^{15} N_{U1343-1012}$) (Fig S3). 188

Given that high $\Delta \delta^{15} N_{U1343-1012}$ values can either reflect higher export production (more rapid removal of nitrate from the euphotic zone) or a slower nutrient resupply (slower replenishment of the surface nitrate pool), and as diatoms are the dominant contributors to export production, the opal MAR can be used to constrain the impact of variable productivity on $\Delta \delta^{15} N_{U1343-1012}$. Comparison of the opal MAR with the opal (%) confirms that the MAR record is not an artefact of the age model and calculated sedimentation rates (Fig S2). The opal MAR record was also linearly interpolated at an interval of 2 kyr and then normalised so that all values ranged between 0 and 1. Then the similarly normalised $\Delta\delta^{15}N_{U1343-1012}$ was subtracted to create a semiquantitative proxy termed the 'upwelling index', with low (higher) values indicating decreased (increased) upwelling nutrient supply on the Bering shelf slope (Equation 1; Fig. 4b).

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$$Upwelling \ Index = Normalised \ Opal \ MAR - Normalised \ \Delta\delta^{15}N_{U1343-1012}$$
(1)

In doing this we assume that the rate of export production is a reflection on total primary 200 productivity, and that the majority of macronutrient delivery to the photic zone at IODP Site 201 U1343 was from upwelling of NPDW and that rates of nutrient utilisation were controlled by 202 both nutrient upwelling and delivery of iron (Fe) to the photic zone (predominantly derived 203 from sea ice and shelf-derived material (Aguilar-Islas et al., 2008)). During times of low Fe 204 205 input and incomplete nutrient utilisation, when increased upwelling will decrease nutrient utilisation (as the nutrient pool is larger) while productivity will show a smaller increase (from 206 the small amount of upwelled Fe which will be scaled proportionally with macronutrient 207 increase), causing the upwelling index to increase at a lower rate than for times of higher Fe 208delivery. The result of the upwelling index can be classified by three potential states which 209 210 reflect interglacial, glacial and deglacial changes in biogeochemical cycling (Fig. 5).

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212 **3. Results**

213 **3.1.** Δδ¹⁸O₈₄₉ - U1343: GNPIW formation

Over the last 850 kyr benthic foraminiferal $\delta^{18}O_{U1343}$ varied between +2.47 and +5.02 ‰ (mean = +3.58 ‰), with an amplitude of change similar to LR04 and ODP Site 849 in the eastern equatorial Pacific (range from +3.08 to +5.10 ‰; mean = +4.12 ‰; Fig. 3a, 4a). The highest $\Delta\delta^{18}O_{849-U1343}$ values occur during glacial maxima of MIS 4, 6, 8, 12, 16 and 20 (Fig. 3b). As

the two δ^{18} O records are used to create the age models, with rapid deglacial changes in δ^{18} O as 218 the tie points, rapid $\Delta \delta^{18}O_{849-U1343}$ shifts at deglacials may be artificially amplified. However, 219 $\Delta \delta^{18}O_{849-U1343}$ values are significantly higher in glacial periods (mean = 0.05) than interglacial 220 221 periods (mean = 0.00; p < 0.05), with values gradually increasing as glacial periods developed and became more intense, suggesting an increasing influence of GNPIW in the Bering Sea 222 slope region. During deglacial periods GNPIW collapses, and the $\Delta \delta^{18}O_{849-U1343}$ becomes more 223 negative during interglacials, as $\delta^{18}O_{U1343}$ is more similar to the deep water record from the 224 eastern equatorial Pacific Site 849, suggesting NPDW influence at the 2 km water depth at 225 226 Bering Sea slope Site U1343.

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228 **3.2.** δ^{15} N and Opal MAR

Today Bering Sea green belt primary productivity is dominated by diatoms which primarily 229 230 bloom in spring/summer following seasonal sea ice melt, which provides a source of dissolved iron and promotes water column stability, allowing diatoms to remain in the photic zone and 231 utilise available nutrients (Aguilar-Islas et al., 2008). Estimates of annual primary productivity 232 in the green belt are >170 g C m⁻² yr⁻¹ (Sambrotto et al., 2008) with blooms sustained by tidal 233 mixing and lateral supply of iron rich waters from the shelf (Aguilar-Islas et al., 2008). 234 Conditions, however, are reversed during winter months (November to April) when sea ice 235 expansion and associated reductions in light availability reduces primary productivity, leading 236 to incomplete nutrient utilisation and the emergence of high-nutrient low chlorophyll (HNLC) 237 238 conditions similar to the lower subarctic Pacific and central Bering Sea (Galbraith et al., 2008; Knudson and Ravelo, 2015b; Sambrotto et al., 2008). With diatoms as the primary contributors 239 to biogenic material within the IODP Site U1343 sediment record (Takahashi et al., 2011), the 240 241 opal mass accumulation (MAR) record, recalculated using sedimentation rates from our new

age model, is thought to approximate changes in primary productivity (Kim et al., 2014), where 242 productivity was significantly higher in interglacials (mean = $3.30 \text{ g cm}^{-2} \text{ kyr}^{-1}$) than in glacials 243 $(\text{mean} = 2.66 \text{ g cm}^{-2} \text{ kyr}^{-1}; \text{ p} < 0.001; \text{ Fig. 3e})$ (Kim et al., 2014). Although the phytoplankton 244 community structure will likely have varied in response to changing seasonality, diatoms have 245 remained the dominant contributors to export production in the Bering Sea (Takahashi et al., 246 2011). As changes in the composition of the phytoplankton community will be minor in 247 comparison to glacial-interglacial variation in export production, we assume these changes are 248 unlikely to have impacted our geochemical signals on millennial timescales. Furthermore, 249 silica supply is not considered a limiting nutrient in the Bering Sea (Tsunogai et al., 1979), and 250 hence the opal MAR likely reflects first order changes in productivity (Kim et al., 2014). 251

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Changes in $\delta^{15}N_{U1343}$ are assumed to primarily reflect temporal variations in nitrate utilisation 253 which is influenced by the amount of productivity and nutrient availability, which is in turn 254 dependent on the rate of upwelling of macronutrient rich NPDW, driven by eddies in the BSC 255 along the shelf break. Terrestrial input with low δ^{15} N is not considered to have driven the record 256 due to the lack of covariation of $\delta^{15}N$ and sedimentary C/N ratio (Fig. S2), and the broad 257 similarity with ODP Site 882 and MD2416 in the North Pacific which are far from land (Fig. 258 3c) (Galbraith et al., 2008; Jaccard et al., 2010). Similarly, variable inorganic contribution is 259 not considered to bias the bulk δ^{15} N record, as the intercepts on %TOC vs. %TN do not vary 260 significantly between glacial and interglacials (Fig. S4). In addition, diatom-bound $\delta^{15}N$ 261 records from the Bering Sea and subarctic Pacific show similar trends to their bulk $\delta^{15}N$ 262 counterparts (Brunelle et al., 2007; Galbraith et al., 2008; Studer et al., 2012), indicating bulk 263 sediment nitrogen is a good proxy for nutrient utilisation in surface water. $\delta^{15}N_{U1343}$ values 264

range from +2.72 to +8.89 ‰ (mean = +5.59 ‰) with peak values occurring in glacials, particularly during MIS 2 at ~14 kyr and MIS 8 at ~245 kyr (Fig. 2c).

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Prior to ~ MIS 13, there appears to be less distinctive glacial-interglacial variability in δ^{15} N-U1343, which could reflect sedimentary preservation, however diagenetic alteration of nitrogen is considered less of an influence in this high sedimentation rate setting (Brunelle et al., 2007; Robinson et al., 2012). Alternatively, the greater glacial-interglacial variability in nutrient utilisation over the last ~600 kyr may be expected, given that 100 kyr cycles in the late Quaternary are characterised by colder and more severe glacial periods, and warmer interglacial periods, than the preceding 41 kyr cycles (McClymont et al., 2013).

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Low $\delta^{15}N_{U1343}$ excursions (< 4.5‰) may be a result of inorganic or terrestrial input, as the 276 modern deep ocean average δ^{15} N is ~4.5‰ (Sigman et al., 2009). However, the δ^{15} N record is 277 not a direct recorder of deep ocean δ^{15} N and more likely reflects surface water δ^{15} N, which will 278 be a combination of the deep water signal, as well as surface water processes such as 279 productivity. Confidence in the fidelity of low $\delta^{15}N_{U1343}$ values (<5.0‰) is provided by other 280 subarctic Pacific records from open ocean sites great distances from the continent, such as from 281 Site MD2416 (Galbraith et al., 2008), Site 882 (Jaccard et al., 2010) (Fig. 3c) and Site 887 282 (Studer et al., 2012). Furthermore, diatom-bound $\delta^{15}N$ results also show values less than 5‰ 283 in the subarctic (Brunelle et al., 2007; Studer et al., 2012). Although nitrate uptake can only 284 account for increases in $\delta^{15}N_{U1343}$, terrestrial/inorganic nitrogen input is unlikely to impose a 285 first-order control on the glacial-interglacial variability at U1343. 286

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3.3. Glacial-interglacial variability in the upwelling index

The upwelling index is highly correlated with the LR04 deep ocean δ^{18} O record (r = -0.72, p < 289 290 0.001) – a combination of global ice volume and deep ocean temperature – with a significantly lower upwelling index in glacials (mean = -0.40) than in interglacials (mean = -0.10; p < 0.001) 291 (Fig. 4). Minimal upwelling index values during glacial maxima (due to increased rates of 292 nitrate utilisation ($\delta^{15}N_{U1343}$) despite lowered primary productivity (opal MAR)), suggest there 293 was a smaller pool of nutrients in the photic zone, concomitant with expanded GNPIW (Fig. 294 3b-c, e). We propose that presence of GNPIW was a pivotal mechanism in reducing nutrient 295 supply from deep water upwelling (see Discussion). A rapid deglaciation phase occurred at all 296 major glacial terminations (dashed lines in Fig. 4) when nutrient upwelling peaked, indicating 297 ample nutrient supply from resumed NPDW upwelling and the establishment of a high 298 productivity/low nitrate utilisation state. 299

Although upwelling is also correlated with relative sea level (Elderfield et al., 2012) (r = -0.55, 300 p < 0.001) and Antarctic air temperature (Jouzel et al., 2007) (r = 0.56, p < 0.001) (Fig. 4), the 301 high correlation with global surface ocean temperature (Snyder, 2016) (r = 0.60, p < 0.001) and 302 303 ice core atmospheric CO₂ records over the last 800 kyr are of particular interest (r = 0.60, p < 0.600.001) (Lüthi et al., 2008). These tight correlations are despite the limitations to our 304 productivity and nutrient utilisation proxies and possible inter-site age model discrepancies, 305 306 therefore suggesting a common link between upwelling, atmospheric temperature and CO₂. Although correlation does not imply causation, the consistency in the relationship between the 307 upwelling index and the ice-core derived CO₂ provides support for a shared underlying 308 309 mechanism. Exceptions to this correlation between the upwelling index and CO₂ occurs during MIS 13, when the upwelling index was high but pCO_2 was lower than subsequent interglacials, 310 and MIS 11, when the upwelling index was lower than might be expected given the higher 311

 pCO_2 for this interglacial (Fig. 4). While the former discrepancy may be a result of the MAR 312 calculations, where sedimentation is relatively high during MIS 13, this also correspond with 313 the Mid-Brunhes Event (MBE) where interglacial conditions became warmer (Jouzel et al., 314 2007). A productivity peak at MIS 13 in response to increased sea ice melt may explain the 315 high upwelling index result at this time. Alternatively, notably warm conditions during "super 316 interglacial" MIS 11 (Melles et al., 2012) likely caused earlier and rapid sea ice melt, which 317 318 would have reduced iron availability to the green belt, reducing the duration of the spring bloom and limiting summer productivity (Aguilar-Islas et al., 2008). After post-MBE warming, from 319 320 MIS 11 onward, the relationship between upwelling index and Antarctic CO₂ is particularly strong (r = 0.70, p < 0.001). 321

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323 **4. Discussion**

324 Changes in nutrient upwelling are likely to have been controlled by a combination of sea level change, sea ice presence and GNPIW extent (Kender et al., 2018). Geochemical and biological 325 proxy evidence indicate that an expansion of sea ice occurred during glacials throughout the 326 last 850 kyr (Detlef et al., 2018; Stroynowski et al., 2017; Teraishi et al., 2016), causing lower 327 rates of primary productivity (Kim et al., 2014) and actively restricting air-sea gas exchange. 328 Coincident with glacials and maximal sea ice extent, our $\Delta\delta^{18}O_{849-U1343}$ results provide 329 evidence for intermediate water presence to at least 2 km water depth, supporting data from 330 IODP Site U1342 which indicates GNPIW presence to at least 800 m water depth (Knudson 331 332 and Ravelo, 2015a) (Fig. 3a-b). Brine rejection on the Bering Sea shelf during enhanced sea ice formation in glacials is the proposed mechanism for formation of a more saline water body, 333 which would sink to intermediate depths at the shelf slope and propagate southward as GNPIW 334 335 (Knudson and Ravelo, 2015a), as observed for NPIW in the modern Okhotsk Sea.

Although direct sea ice reconstructions from the Bering Sea are of lower resolution, they 337 338 broadly support GNPIW expansion, as glacial periods during this time interval were characterised by extended thick pack ice which peaked during the mid-to-late glacials, when 339 we see the largest offsets in $\Delta \delta^{18}O_{849-U1343}$ (Detlef et al., 2018; Stroynowski et al., 2017). 340 Changes in sea-ice would have been in combination with sea level falls of ~50-150 m during 341 glacials (Elderfield et al., 2012) (Fig. 4d), causing closure of the Bering Strait (modern depth 342 of ~50 m) and retention of both cold/fresh surface waters and denser modified shelf-water 343 344 (following brine rejection) within the Bering Sea, as the northward flow into the Arctic Ocean was prevented (Kender et al., 2018; Knudson and Ravelo, 2015a). The resultant expanded 345 GNPIW at the shelf edge likely prevented eddies in the BSC from reaching NPDW, isolating 346 nutrients in the photic zone without significant resupply (low upwelling index) and causing the 347 observed high rates of nitrate utilisation but low productivity, whilst also trapping CO₂ in deep 348 349 waters during glacials over the last 850 kyr (Fig. 5b).

350

In addition to sea ice and GNPIW expansion during glacials triggering a reduction in 351 productivity and nutrient upwelling, lowered sea level would have hindered Alaskan Stream 352 water from flowing into the basin by shoaling the passes along the eastern Aleutian arc (Fig. 353 1b). Although we cannot quantify this effect, it would have caused some degree of decrease in 354 surface water flow around the Bering Sea basin. This is important because the strength of BSC 355 flow controls eddy field fluctuation, where increased flow maintains higher primary 356 productivity by promoting vertical nutrient supply, and vice versa (Mizobata and Saitoh, 2004). 357 There is evidence for glacial reductions in the inflow of the Alaskan Stream from reduced 358 359 proportions of the diatom species Neodenticula seminae near the inflow sites at the Aleutian

arc in the southern Bering Sea, as well as at the slope (Stroynowski et al., 2017; Teraishi et al., 360 2016). The resultant decrease in BSC flow in glacials would have therefore reduced large eddy 361 formation, further lowering NPDW upwelling of nutrients, productivity and photic zone CO₂ 362 (Mizobata and Saitoh, 2004). Simultaneously, sea ice expansion over a shallowed Beringian 363 shelf during glacials may have increased the mobilisation of sediments and transport of shelf-364 derived iron to the slope (Aguilar-Islas et al., 2008). Reduced vertical macronutrient supply but 365 366 sustained iron supply from sea ice, which is a key source of bioavailable iron in the green belt today (Aguilar-Islas et al., 2008), could help to explain the co-occurrence of low productivity 367 and high nutrient utilisation in glacials (Fig. 2, 5b). Overall, our results support previous 368 proposals that enhanced sea ice and GNPIW formation during periods of low glacial sea level 369 were a key mechanism for controlling deep water upwelling and green belt nutrient supply over 370 371 the last 850 kyr (Kender et al., 2018). Reduced upwelling of nutrient-rich NPDW due to GNPIW presence and lowered BSC strength and physical overturning in glacials could then 372 have caused the green belt region to act as a larger net carbon sink. 373

374

375 During deglaciation, conditions would have reversed with the break up and retreat of sea ice (Kim et al., 2014; Stroynowski et al., 2017) and sea level rise, causing re-strengthening of the 376 BSC and both local and regional collapse of GNPIW (Fig. 5b). Coupled with high shelf-derived 377 iron from deglacial flooding of the Beringian shelf and potentially increased deglacial Yukon 378 River flow (VanLaningham et al., 2009), this rapid restoration of regional upwelling nutrients 379 explains the observed productivity peaks during deglaciations (Kim et al., 2014) from at least 380 381 MIS 20-21 onwards. Modelling has shown that late in the last deglacial, wind strength across the Bering Sea and subarctic remained greatly increased due to the remanence of large North 382 America ice sheets (Gray et al., 2018). This would have aided eddy formation in the Bering 383

Sea and upwelling of NPDW across the subarctic region through promoting vertical mixing.
Once ice sheets had declined and sea ice diminished, seasonally variable upwelling and vertical
mixing across the subarctic Pacific would have become re-established in interglacials, similar
to modern conditions (Fig. 5c).

388

We have demonstrated for the first time that the glacial reductions in Bering Sea slope primary 389 productivity were driven at least in part by reduced upwelling of nutrients (Fig. 4b), and 390 391 propose that this was largely controlled by sea ice and expanded low- nutrient GNPIW (Fig. 3b). Whilst we do not suggest that this mechanism solely explains the observed proxy evidence 392 from across the North West Pacific Ocean during the Pleistocene, similar productivity and 393 nutrient utilisation δ^{15} N records are seen at ODP Site 882 and MD2416 (Fig. 3c, d, f) (Galbraith 394 et al., 2008; Jaccard et al., 2010), whilst an upwelling index derived for ODP Site 882, using a 395 $\Delta \delta^{15} N_{882-1012}$ record for regional nutrient utilisation and opal (%) as a productivity record 396 (Jaccard et al., 2010) (Fig. 3d, f), shows remarkably similar trends to the Bering Sea upwelling 397 index. Although the upwelling index for Site 882 only covers a short time period, the high 398 correlation between the upwelling index for Sites U1343 and 882 (r = 0.77, p < 0.001) provides 399 support for a common controlling mechanism of GNPIW between the Bering Sea and wider 400 subarctic Pacific region. 401

402

We propose that reduced upwelling of nutrients in the subarctic Pacific Ocean, as recorded at ODP Site 882 and MD2416, was the cause of reduced glacial export production, forced by the expansion of low nutrient GNPIW, similar to the Bering Sea slope (at U1343). The formation of dense GNPIW in the Bering Sea likely propagated south through the deep western Kamchatka Strait (Fig. 1), although this does not discount additional contributions of NPIW

from the Okhotsk Sea (Max et al., 2014). This mechanism was proposed for the last glacial 408 cycle in the subarctic Pacific Ocean, where model and proxy evidence from the Bering and 409 Okhotsk Seas indicated that enhanced GNPIW during Heinrich Stadial 1 increased 410 intermediate-to-deep ocean stratification, isolating abyssal carbon in NPDW (Gong et al., 411 2019). Furthermore, the disappearance of GNPIW during the latter part of last deglaciation 412 allowed renewed upwelling of deeper water rich in nutrients and is considered responsible for 413 increases in surface water proxies for CO₂ (Gray et al., 2018). Our new data support GNPIW 414 production as the underlying mechanism driving the strength of nutrient upwelling both along 415 416 the Bering Sea slope and across the open subarctic Pacific Ocean over the past 850 kyr.

417

418 Changes in the Southern Ocean are considered to be a dominant component in driving glacialinterglacial changes in pCO₂ (Gottschalk et al., 2016; Jaccard et al., 2013). For example, glacial 419 reduction of atmospheric pCO_2 has been linked to an increased rate of biological sequestration 420 in the Southern Ocean and/or reduced supply of nutrients and CO₂ upwelled from the deep 421 ocean as a result of intermediate water formation and expanded sea ice cover (Sigman et al., 422 423 2010). Although net CO₂ flux in the Southern Ocean is regarded as the largest single control on glacial-interglacial variability in pCO_2 (Sigman et al., 2010), we demonstrate here that 424 similar processes occurred in the subarctic Pacific region. With modelling studies suggesting 425 426 that ~ 30 ppm atmospheric CO₂ was released into the atmosphere due to increased overturning in the subarctic region at the last deglaciation (Rae et al., 2014), we surmise that similar 427 magnitude ocean-atmospheric CO₂ fluxes may have occurred over the last 850 kyr in response 428 429 to changes in deep water nutrient upwelling across the Bering Sea and subarctic Pacific Ocean. This aids ongoing efforts to fully explaining observed trends in atmospheric pCO_2 over this 430 interval (Gray et al., 2018; Rae et al., 2014). 431

432 **5.** Conclusion

Overall, we constructed a new high resolution benthic oxygen isotope stratigraphy for IODP 433 Site U1343, to aid the development of a new nutrient upwelling index which uses $\delta^{15}N_{\text{bulk}}$ and 434 opal MAR to assess glacial-interglacial variability in the supply of nutrients from NPDW 435 upwelled at the Bering shelf edge. We find a tight correlation between the upwelling index 436 results and global temperature, sea level and atmospheric pCO_2 , as well as the upwelling index 437 calculated for the short term record at ODP Site 882 in the western subarctic Pacific Ocean. 438 We posit that increased sea ice volumes in the Bering Sea and the resultant expansion of 439 GNPIW caused suppression of both nutrient and CO₂ upwelling during periods of Bering Strait 440 441 closure, both in the Bering Sea and across the subarctic Pacific Ocean region.

Results here suggests that the regional significance of sea ice and deep water ventilation in the 442 subarctic Pacific Ocean has been generally underestimated when modelling Quaternary climate 443 change, where GNPIW growth and propagation from the Bering Sea influences both local and, 444 importantly, regional upwelling of NPDW and export production. The strong correlation 445 446 between high latitude upwelling and glacial cycles in atmospheric CO₂ (Fig. 4b-c), and the bipolar nature of Quaternary glaciations, also supports suggestions that sea ice was a primary 447 control on the past global climate state (Sigman et al., 2010). Overall, improved understanding 448 449 of this mechanistic relationship between the Bering Sea and the subarctic Pacific will aid future efforts to constrain the role of glacial-interglacial oceanographic variability in global climate, 450 in complement to Southern Ocean-driven dynamics. 451

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453 **References**

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Fig. 1: Oceanographic setting of the Bering Sea and subactic Pacific Ocean. a) Location of Site U1343 (57 33'N, 175 49'W; 1954 m; this study) on the Bering slope and Site U1342 (54 83'N, 176 92'E; 818m) in the southern Bering Sea from IODP Expedition 323. The location of North West Pacific ODP Sites 882 (50.35, 167.58; 3244 m) and MD2416 (51.27 N, 167.73 E; 2317m) are also shown, together with surface (red arrows) and deep water (blue arrows) circulation. The high productivity 'green belt' region (Springer et al., 1996), stimulated by shelf-adjacent Bering Slope Current (BSC) is also marked (green patterned band). b) Cross sections of the Aleutian passes and volumes of transport through which Alaskan Stream water exchanges with the Bering basin (Stabeno et al., 1999; Takahashi et al., 2011).



Fig. 2 : Benthic foraminiferal δ^{18} O results from U1343. a) Raw δ^{18} O values are categorised by species/genus and plotted against *E. batialis* (black) to which species-specific offsets are applied (Kender et al., 2018). b) Raw δ^{18} O data (red dots) are plotted against δ^{18} O_{U1343} (black line) which has been tuned to *E. batialis*. c) Comparison of δ^{18} O_{U1343} from this study (black line) against previous data from IODP Site U1343 (Asahi et al., 2016) (blue dots), which were combined in d) to create the updated age model for IODP Site U1343 (orange dots), which has been smoothed using linear interpolation to a 2 kyr resolution (black line).



Fig. 3: Geochemical proxy results from IODP Site U1343 from MIS 2 to 20. a) Benthic foraminiferal δ^{18} O results from IODP Site U1343 compared to those from the shallower IODP Site U1342 and deep Pacific ODP Site 849. The data from Site U1342 has been recalculated here to match benthic foraminiferal offset for *E. batialis*, used for Site U1343 (Kender et al., 2018). b) The offset in δ^{18} O between IODP Site U1343 and ODP Site 849 ($\Delta\delta^{18}O_{849-U1343}$). c) Bulk $\delta^{15}N$ data from IODP Site U1343 compared with deep North Pacific ODP Site 1012 (a site of complete nutrient utilisation) together with bulk $\delta^{15}N$ records from ODP Site 882 and MD2416 in the subarctic North Pacific Ocean. d) The $\delta^{15}N$ offset between IODP Site U1343 and ODP Site 1012 ($\Delta\delta^{15}N_{U1343-1012}$) which constrains changes in source water $\delta^{15}N$ to reflect changes in Bering Sea nutrient utilisation. e) Opal mass accumulation rate (MAR) from IODP Site U1343 [adapted from Kim et al. (2014), see Methods]. f) The opal (%) and Ba/Al productivity records from ODP Site 882 (Jaccard et al., 2010). Blue bars represent periods of Bering Strait closure (>50 m sea level drop) (Elderfield et al., 2012) with vertical dashed lines indicating deglaciations.



Fig. 4: Comparison of the upwelling indexes against other global and regional paleoenvironmental records. a) Composite LR04 benthic foraminiferal δ 180 stack (Lisiecki and Raymo, 2005) over the last 850 kyr. b) The upwelling index for Site U1343 (red) and Site 882 (grey), which show suppressed rates of upwelling during glacial periods following Bering Strait closure (blue bars). The upwelling index shows strong correlation with the composite EPICA Dome C Ice Core atmospheric CO2 record over the last 800 kyr (c) (Lüthi et al., 2008), changes in relative sea level (m) (d) (Elderfield et al., 2012), where the blue lines at -50 m represent modern Bering Strait depth, and global temperature records (temperature deviation from present day) (e) (Jouzel et al., 2007; Snyder 2016). Vertical dashed lines indicate deglaciations.





b) Deglacial: high upwelling index



c) Interglacial: intermediate upwelling index



Fig. 5: Schematic models representing glacial, deglacial and interglacial regimes of biogeochemical cycling in the Bering Sea. a) A low upwelling index indicates reduced upwelling caused by an expansion of NPIW which forms in glacials following the closure of the Bering Strait, leading to an isolated pool of nutrients in the photic zone. This results in reduced primary productivity (opal MAR) but higher rates of nutrient utilisation due to the reduced supply of deep water nutrients into the photic zone. b) A high upwelling index indicates ample nutrient supply from resumed NPDW upwelling which occurs during deglaciation, leading to a high productivity/low nutrient utilisation state. c) An intermediate upwelling index results indicates upwelling typical of an interglacial, similar to the modern regime.



- Cassidulina spp.
- Elphidium batialis
- Globobulimina spp.
- Islandiella norcrossi
- Nonionella spp.
- Uvigerina spp.
- Valvulineria spp.

Fig. S1: Benthic foraminiferal comparison of δ^{13} C and δ^{18} O. a) δ^{13} C data categorised by species/genus. The poor relationship for δ^{13} C against both raw δ^{18} O (b) and δ^{18} O adjusted to *E. batialis* (c) suggests that diagenetic alteration of foraminiferal shells cannot explain the glacial-interglacial variability in the δ^{18} O.



Fig. S2: Bulk δ^{15} N and Biogenic Opal data for IODP Site U1343. a) Bulk δ^{15} N data (red dots) from this study and previously published lower resolution data (Kim et al., 2017), smoothed using a linearly interpolation to create a 2 kyr resolution dataset (black line). b) C/N ratios, which are low and show a lack of covariation to δ^{15} N, suggesting that terrestrial input of nitrogen was low and did not drive the δ^{15} N record. c) New sedimentation rates calculated for IODP Site U1343 using the updated age model presented in this study, which were then applied to d) the biogenic opal record (Kim et al. 2014) to calculate e) the opal mass accumulation rate (MAR) data.



Fig. S3: Assessing the relative contribution of $\delta^{15}N_{1012}$ to $\Delta\delta^{15}N_{U1343-1012}$. Results demonstrate that there is no significant difference between the upwelling index record calculated using $\Delta\delta^{15}N_{U1343-1012}$ compared to $\delta^{15}N_{U1343}$ (where the latter does not subtract the ODP 1012 record) (t = -1.93, p = 0.05).



Fig. S4: %TOC vs %TN cross plots for IODP Site U1343 from this study and previously published lower resolution data (Kim et al., 2017). a) Shows the cross plot for the whole dataset, with an intercept of 0.0193, suggesting minimal influence of inorganic nitrogen. Similarity between b) glacial intercept of 0.0181 and c) interglacial intercept of 0.0192, suggests that variable contribution of terrestrial/inorganic nitrogen does not drive the bulk δ^{15} N record presented here.

Table S1: Benthic foraminiferal δ^{18} O offsets applied to species/genus, from *Elphidium batialis* (Kender et al., 2018). The total number of samples for each group is also noted.

Species	Offset from Elphidium batialis (‰)	Number of samples
Elphidium batialis	0	117
Valvulineria spp.	+ 0.25	6
Islandiella norcrossi	+ 0.48	9
Cassidulina spp.	+ 0.56	11
Nonionella spp.	+ 0.65	2
Globobulimina spp.	+ 0.85	61
Uvigerina spp.	+ 0.87	56

Table S2: New age-depth tie points for Site U1343, building upon work of Asahi et al. (2016). Age model constructed by tuning to LR04 global benthic stack (Lisiecki and Raymo, 2005) (see Methods).

	Age (kyr)
Depth (CCSF – A) (m)	
0.96	10.15
14.23	57.38
36.72	131.41
48.03	181.46
59.73	219.06
68.72	242.57
79.52	279.96
96.24	335.14
114.56	396.17
119.35	424.39
129.90	480.93
145.02	512.78
152.01	545.32
161.63	580.48
173.39	621.39
174.57	641.03
184.09	700.03
188.27	725.84
191.58	753.63
203.57	790.75
209.04	812.86