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1 **Reduced Arctic sea ice extent during the mid-Pliocene Warm Period**  
2 **concurrent with increased Atlantic-climate regime**

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23 **Revised submission to Earth and Planetary Science Letter (EPSL)**

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## 26 **Abstract**

27 Quantifying the contribution of poleward oceanic heat transport to the Arctic Ocean is  
28 important for making future sea ice and climate predictions. To highlight its potential  
29 importance in a warmer world, we present a new record of water-mass exchange between  
30 the Atlantic and the Arctic Oceans using the authigenic neodymium isotopic composition  
31 of marine sediments from the Fram Strait during the past ~3.4 to 2.6 Ma. In this study, we  
32 target the mid-Pliocene Warm Period (mPWP: 3.264–3.025 Ma) of the Pliocene epoch,  
33 the most recent geological analogue for future climate change. We complement our semi-  
34 quantitative water mass exchange reconstruction with estimates of spring sea ice  
35 concentration based on source-specific biomarkers. Our estimates of volume transport of  
36 warm waters into the Arctic Ocean suggest long-term secular changes from the lowest  
37 during the Marine Isotope Stage M2 “glacial” (3.312–3.264 Ma), to near complete  
38 “Atlantification” of the Eurasian sector of the Arctic Ocean during the mPWP. Orbital  
39 forcing is found to be the dominant controlling factor for modulating northward volume  
40 transport of Atlantic-derived water masses, with consequential reduction in Arctic spring  
41 sea ice concentration by ~30-35%. Current generation models often produce diverging  
42 results, however, and have not yet been validated against proxy data in northern high  
43 latitude settings during the mPWP. Our new results of northward volume transport and  
44 sea ice extent therefore provide much needed input for validation of current generation  
45 models aimed at improving the robustness of future climate modelling in the Arctic.

46 **Keywords: mid-Pliocene, North Atlantic Current, Arctic, Sea ice, Atlantification**

## 47 **1. Introduction**

48 The most dramatic changes observed in the Arctic Ocean during the recent past are the  
49 unprecedented reductions in sea ice extent and thickness (Kinnard et al., 2011). Although  
50 coupled ice-ocean model simulations suggest that the recent warming in the Northern  
51 Hemisphere is responsible for this decline (Petrie et al., 2015), there is disagreement  
52 between data and models over the impact of atmospheric warming versus oceanic heat  
53 transport on sea ice decline (Ding et al., 2018; Dowsett et al., 2012; Haywood et al., 2013).  
54 Studies based on proxy reconstructions of heat and volume transport through the Fram  
55 Strait (Spielhagen et al., 2011), and in-situ observations in the eastern Arctic Ocean  
56 (Polyakov et al., 2017), suggest that enhanced oceanic heat transport by the North  
57 Atlantic Current (NAC) over the past few decades likely explains the weakened  
58 stratification, increased vertical mixing and reduced sea ice in the Atlantic sector of the  
59 Arctic, collectively termed “Atlantification” (Polyakov et al., 2017; Spielhagen et al., 2011).  
60 In order to improve our understanding about Arctic sea ice variability, particularly within  
61 the current context of rapid global warming, it is imperative to reconstruct sea ice  
62 conditions during previous warm climate states, and decipher the underlying mechanisms  
63 that control its distribution. One such period in Earth’s history is the Pliocene (5.33–2.58  
64 Ma), which experienced higher global temperatures than pre-industrial (Dowsett et al.,  
65 2009), and was characterized by a gradual transition from relatively warm climates during  
66 the Early Pliocene towards cooler conditions in the Late Pliocene. Some previous organic  
67 geochemical-based proxy climate reconstructions for the Pliocene have been conducted  
68 for the North Atlantic and Fram Strait (Clotten et al., 2018; Knies et al., 2002), and similar  
69 studies have been carried out for other warm interglacials such as the Eemian (Marine

70 Isotope Stage (MIS) 5e) (Stein et al., 2017). However, the roles of atmospheric warming  
71 versus northward heat transport were not assessed as part of these studies.

72 Here we aimed to identify the potential impact of future changes in oceanic heat  
73 transport into the Arctic Ocean and the effects of “Atlantification” in a warmer than modern  
74 climate. To achieve this, we conducted a semi-quantitative assessment of northward  
75 volume transport of Atlantic water through the Fram Strait during a geological period when  
76 (1) climatic conditions in terms of temperature and atmospheric CO<sub>2</sub> level were analogous  
77 to modern/or future projected scenarios and (2) global oceanographic and tectonic  
78 settings were nearly identical to today. The mid-Pliocene Warm Period (mPWP: 3.264–  
79 3.025 Ma) is known to be warmer (globally) than today (Dowsett et al., 1992; Haywood et  
80 al., 2016), with atmospheric CO<sub>2</sub> concentrations estimated to be in the range 350-450  
81 ppmv (Berends et al., 2019; Foster et al., 2017). Hence, the mPWP has been proposed  
82 as a possible reference for future warm climate states (IPCC, 2013). Confirmation of  
83 increased polar ocean heat transport and reduced sea ice in the Arctic Ocean during the  
84 mPWP (Raymo et al., 1996) would therefore be of clear benefit for the assessment of  
85 coupled ocean-ice-atmosphere model simulations of the mPWP (Haywood et al., 2016).

86 To achieve this objective, we first reconstructed an orbital-resolution record of  
87 watermass mixing between the NAC and Arctic-derived polar waters (PW) in the Fram  
88 Strait (Fig. 1), based on authigenic neodymium (Nd) isotopes ( $\epsilon_{Nd}$ ). The radiogenic  
89 isotope composition of Nd in seawater reveals changes in watermass mixing and  
90 circulation patterns due to its quasi-conservative behavior (Martin, 2002) and lower  
91 average oceanic residence time (360–2000 years) compared to the global ocean mixing  
92 time ~1500 years (Tachikawa et al., 1999) . Critically, in contrast to stable oxygen ( $\delta^{18}O$ )

93 or carbon isotope ( $\delta^{13}\text{C}$ ) measurements, the Nd isotope ratios are not affected by isotopic  
94 fractionation resulting from any biological or other low-temperature processes, so  
95 represent a robust proxy for paleo-water mass circulation (Martin, 2002). In a modern  
96 context, the majority of Atlantic-derived water masses are transported northward into the  
97 Arctic Ocean along the Svalbard continental margin, which is the northernmost extension  
98 of the NAC (Fig. 1). This warm water submerges into the Arctic Ocean or is deflected  
99 westward and submerged southward below cold and less saline waters of the East  
100 Greenland Current (EGC). All of these modern water masses possess characteristic Nd  
101 isotope signatures (Fig. 1) (Laukert et al., 2017; Werner et al., 2014). Less radiogenic  
102 values are indicative of a stronger influence of NAC flowing into the Nordic Seas (present-  
103 day  $\epsilon_{\text{Nd}} = -13.2$  to  $-13.0$ ) (Teschner et al., 2016) while more radiogenic Nd isotope  
104 signatures reflect enhanced contribution from Arctic-derived polar waters (PW) (Laukert  
105 et al., 2017) (e.g.  $\epsilon_{\text{Nd}} = -9.9$ ). For this study, an orbital-resolution ( $\sim 5$  ka) authigenic  $\epsilon_{\text{Nd}}$   
106 record was obtained through analysis of bulk sediments from Ocean Drilling Program  
107 (ODP) Hole 910C (hereafter referred to 910C) on the Yermak Plateau, eastern Fram  
108 Strait, ( $80^{\circ}15.894'\text{N}$ ,  $6^{\circ}35.430'\text{E}$ , water depth: 556.4 m) covering the interval between 3.4  
109 Ma and 2.6 Ma. This new record is supplemented by a previously published low-resolution  
110 (60-70 ka) record of authigenic  $\epsilon_{\text{Nd}}$  from ODP Hole 911A ( $80^{\circ} 28.466' \text{ N}$ ,  $8^{\circ} 13.640' \text{ E}$ ,  
111 water depth: 902 m) (hereafter referred to 911A) at the eastern flank of the Yermak  
112 Plateau (Teschner et al., 2016).

113 To identify the corresponding changes in sea ice coverage and carbonate  
114 chemistry, the sea ice biomarker proxy  $\text{IP}_{25}$ , a related open-water highly branched  
115 isoprenoid (HBI) lipid (HBI III), and calcium carbonate ( $\text{CaCO}_3$ ) abundance related to

116 carbonate chemistry and productivity/or preservation, were also analyzed. Over the last  
117 decade, source-specific highly branched isoprenoid (HBI) lipid biomarkers have emerged  
118 as reliable proxies for reconstructing past sea ice extent in the polar oceans (Belt, 2018  
119 and references therein). The most frequently studied biomarker is the mono-unsaturated  
120 HBI IP<sub>25</sub>, first identified in Arctic sea ice and sediments by Belt et al. (2007), and has since  
121 been used as a binary measure of seasonal Arctic sea ice in the past for time scales  
122 ranging from recent decades to several millions of years. Further, by combining  
123 sedimentary IP<sub>25</sub> concentrations with those of various phytoplankton biomarkers in the  
124 form of the IP<sub>25</sub>-phytoplankton (PIP<sub>25</sub>) index, semi-quantitative estimates of sea ice extent  
125 can be achieved (Muller et al., 2011; Belt, 2018 for a review). Finally, when a further tri-  
126 unsaturated HBI (often referred to as HBI III; Belt et al., 2015) is used as the open water  
127 counterpart to IP<sub>25</sub>, the resulting PIP<sub>25</sub> index (i.e. P<sub>III</sub>IP<sub>25</sub>) exhibits a reasonably good linear  
128 relationship to spring sea ice concentration (%SpSIC) for the Barents Sea and  
129 neighboring regions (Smik et al., 2016).

130         North Atlantic and Arctic waters are characterized by distinct carbonate (e.g.  
131 alkalinity and pH) characteristics, so carbonate abundance in sediments from the Fram  
132 Strait (mixing zone) can be used to infer changes in carbonate productivity, preservation  
133 and dissolution resulting from variable paleo-oceanographic changes. For example, warm  
134 and carbonate-rich North Atlantic waters lead to better preservation compared to cold  
135 carbonate depleted Arctic waters. In some previous studies, therefore, high carbonate  
136 preservation in sediments from the Fram Strait has been attributed to increased influence  
137 of Atlantic water masses (e.g. Zamelczyk et al., 2014). In the Norwegian–Greenland Sea,  
138 high carbonate content has also been interpreted to reflect the influence of warm Atlantic

139 water masses, while low carbonate content were attributed to cold surface waters and  
140 increasing dilution by terrigenous material (Huber et al., 2000). Therefore, a combined  
141 authigenic Nd isotope and carbonate record from the Fram-Strait were employed in the  
142 present study to reconstruct northward volume and heat transport by the NAC.

143

## 144 **2. Material and methods**

145 Sediments of ODP Hole 910C (80°15.894' N, 6°35.430' E; water depth: 556.4 m)  
146 have been analyzed in this study. The deep-water Nd isotope signal was extracted from  
147 the Fe-Mn oxyhydroxide fraction of bulk sediment following the leaching procedure  
148 described below. Further details of Nd isotopes, highly branched isoprenoid biomarkers  
149 and calcium carbonate abundance measurements are given in the following sections.

### 150 **2.1 Neodymium isotope analysis in authigenic fractions**

151 We measured the neodymium (Nd) isotope composition in authigenic phases extracted  
152 from the bulk sediments of 910C. This new record of authigenic  $\epsilon_{Nd}$  is supplemented by  
153 an earlier published low-resolution (60-70 ka) record from ODP Hole 911A at the eastern  
154 flank of the Yermak Plateau (Teschner et al., 2016). Therefore, for comparison, and to  
155 avoid discrepancies related to the analytical methods for the extraction of authigenic Nd  
156 from sediments and its isotope measurements, we adopted the same method of Teschner  
157 et al. (2016). The procedure thus began with extracting the past seawater signal  
158 contained in the diagenetic coatings from ~2 g of sample material with a 0.05 M  
159 hydroxylamine hydrochloride and 15% acetic acid solution (HH leach), buffered to a pH  
160 of ~3.5 to 4.0, without rinsing before the HH leach. The rare earth elements (REEs) in the  
161 solution were separated using cation exchange columns filled with AG50WX8 resin (mesh



162 200–400). Nd was separated from the other REEs using columns filled with Ln-Spec resin  
163 (50–100 mesh). Nd isotopes were analyzed using a multi-collector inductively coupled  
164 plasma mass spectrometer (MC-ICP-MS, Thermo Scientific Neptune Plus) at the National  
165 Centre for Polar & Ocean Research (NCPOR), Goa, India. All Nd isotope ratios  
166 ( $^{143}\text{Nd}/^{144}\text{Nd}$ ) presented here were corrected for mass bias following an exponential law  
167 using the known value of  $^{146}\text{Nd}/^{144}\text{Nd}$  of 0.7219. The instrument bias was normalized to  
168 the accepted  $^{143}\text{Nd}/^{144}\text{Nd}$  value of the JNdi-1 standard of 0.512115 (Tanaka et al., 2000).  
169 Repeat measurements of the JNdi-1 standard yielded a long-term average reproducibility  
170 of  $\pm 0.3 \text{ } \epsilon_{\text{Nd}}$  ( $2\sigma$ ;  $n = 103$ ) over a period of nine months. Average procedural blank  
171 ascertained for Nd ( $n = 4$ ) was 170 pg which is less than 1% of the total Nd analyzed in  
172 samples, so blank correction was not applied. All Nd isotope ratios are reported in epsilon  
173 notation according to Equation 1.

$$174 \quad \epsilon_{\text{Nd}} = \left[ \frac{{}^{143}\text{Nd}/{}^{144}\text{Nd}_{\text{sample}}}{{}^{143}\text{Nd}/{}^{144}\text{Nd}_{\text{CHUR}}} - 1 \right] \times 10^4 \quad \text{Eq (1)}$$

175 In order to check the quality of the authigenic Nd isotope analyses, which includes  
176 chemical extractions of the authigenic Nd and its isotopic measurements, we analyzed a  
177 total of 16 replicates. Data from these replicates (with a variable range of  $\epsilon_{\text{Nd}}$ ) are highly  
178 consistent (Supplementary Fig. S1); majority are falling on the equiline (1:1 line) within  
179 their uncertainty.

## 180 **2.2 HBI biomarkers**

181 The HBI biomarkers IP<sub>25</sub> and HBI III were extracted from freeze-dried subsamples  
182 (~2–4 g) from 910C. Samples were saponified in a methanolic KOH solution (~5 mL H<sub>2</sub>O:  
183 MeOH (1:9); 5% KOH) for 60 min (70 °C). Hexane (3×2 mL) was added to the saponified  
184 content, with supernatant solutions, containing non-saponifiable lipids (NSLs), transferred

185 with glass pipettes to clean vials and dried over a gentle stream of N<sub>2</sub> to remove traces  
186 of H<sub>2</sub>O/MeOH. NSLs were then re-suspended in hexane (0.5 mL) and fractionated using  
187 column chromatography (SiO<sub>2</sub>; 0.5 g). Non-polar lipids, including IP<sub>25</sub> and HBI III, were  
188 eluted with hexane (6 mL). Each non-polar fraction was further purified to remove  
189 saturated components using silver-ion chromatography (Belt et al., 2015) with saturated  
190 compounds eluted with hexane (2 mL) and unsaturated compounds, including HBIs,  
191 collected in a subsequent acetone fraction (3 mL). Prior to extraction, samples were  
192 spiked with an internal standard (9-octylheptadec-8-ene, 9-OHD, 10 µL; 10 µg mL<sup>-1</sup>) to  
193 permit quantification of HBIs. Analysis of fractions containing IP<sub>25</sub> and HBI III was carried  
194 out using gas chromatography–mass spectrometry (GC–MS) following the methods and  
195 operating conditions described elsewhere (Belt et al., 2012). Mass spectrometric analysis  
196 was carried out in total ion current (TIC) and selected ion monitoring (SIM) modes. The  
197 identification of IP<sub>25</sub> and other HBIs was based on their characteristic GC retention indices  
198 (e.g. RI<sub>HP5MS</sub> = 2081 and 2044 for IP<sub>25</sub> and HBI III, respectively) and mass spectra (Belt,  
199 2018). Quantification of all HBIs was achieved by comparison of mass spectral responses  
200 of selected ions (e.g. IP<sub>25</sub>, *m/z* 350; HBI III, *m/z* 346) in SIM mode with those of the internal  
201 standard (9-OHD, *m/z* 350) and normalized according to their respective instrumental  
202 response factors, derived from solutions of known biomarker concentration, and sediment  
203 masses (Belt et al., 2012).

204 Concentrations of IP<sub>25</sub> and HBI III were combined in the form of the P<sub>III</sub>IP<sub>25</sub> index  
205 (Eq. 4), with the latter then used to provide semi-quantitative estimates of spring sea ice  
206 concentration (SpSIC (%), Eq. 5) according to a recent regional calibration (Smik et al.,

207 2016). A root mean-square error of 11% associated with SpSIC estimates, was also  
208 calculated using regional calibration data (Köseoğlu et al., 2018; Smik et al., 2016)

$$209 \quad P_{III}P_{25} = \frac{IP_{25}}{(IP_{25} + (0.63 * HBI_{III}))} \quad Eq (4)$$

$$210 \quad SpSIC (\%) = \frac{(P_{III}IP_{25} - 0.0692)}{0.0107} \quad Eq (5)$$

211 Finally, we used the non-parametric CP3O algorithm from the R package ECP (R Core  
212 Team, 2018) to carry out change-point analysis on SpSIC estimates to identify significant  
213 shifts in the time series profile (Supplementary Fig. S2). All biomarker and %SpSIC data  
214 are provided in Supplementary Data 2.

### 215 **2.3 Analysis of carbon**

216 Analyses of total carbon (TC) and organic carbon ( $C_{org}$ ) were performed with a LECO SC-  
217 632 at the Geological Survey of Norway, Trondheim. For TC determination, subsamples  
218 of 300-400 mg were combusted at 1350°C and the release of  $CO_2$  was measured. For  
219  $C_{org}$  analysis, sub-samples of 400-450 mg were placed in carbon-free pervious ceramic  
220 combustion boats. These were placed on a heating plate at 50 °C ( $\pm 5^\circ C$ ) and treated  
221 with 10 vol.% hydrochloric acid (HCl) to remove inorganic carbon (carbonate) and  
222 subsequently rinsed with distilled water and dried in the drying oven prior to analysis.  
223 Carbonate content was calculated as  $CaCO_3 = (TC - C_{org}) \times 8.33$  on the  
224 assumption that calcite is the dominant form in the carbonate fraction (Vogt et al., 2001).  
225 Results are provided in weight percentage (wt. %) and the standard deviation of the TC  
226 and  $C_{org}$  measurements based on the repeated measurements of a standard was  $\pm 0.026$   
227 wt% ( $1\sigma$ , n=8) and  $\pm 0.028$  wt. % ( $1\sigma$ , n=11), respectively.

### 228 **2.4 Age control for sediments deposited at ODP Hole 910C**

229 The age constraints for 910C is based on correlation of bio-stratigraphic and  
230 magneto-stratigraphic datums with Hole 911A together with additional benthic stable  
231 isotope data from 910C for the Pliocene (2.44 – 5.76 Ma). The age model based on the  
232 tie points and associated uncertainties have already been discussed in previous studies  
233 (Grøsfjeld et al., 2014; Knies et al., 2014b; Mattingsdal et al., 2014). Briefly, five tie points  
234 formed the basis of the age model for our target interval between ~3.4 and 2.6 Ma in  
235 910C (see Supplementary Table S1). Two tie-points at 190 mbsf and 305 mbsf inferred  
236 from seismic correlation between 910C and 911A mark the magneto-stratigraphic  
237 boundaries at 2.58 Ma (Matuyama/Gauss) and 3.6 Ma (Gauss/Gilbert) (Mattingsdal et al.,  
238 2014). Support for this age model is provided by the biostratigraphic “Datum A” (~2.78  
239 Ma) at ~223 mbsf in 910C (Sato and Kameo, 1996) and the glacial-to-interglacial  
240 oscillations of the benthic  $\delta^{18}\text{O}$  record of 910C (Knies et al., 2014a). Between “Datum A”  
241 (2.78 Ma) and the inferred Gauss/Gilbert boundary (3.6 Ma), we have originally applied  
242 an age model based on linear sedimentation rates between these two fix-points (Knies et  
243 al., 2014a). One major climate transition (i.e. MIS M2 glaciation) expressed by a sharp  
244 increase in the global  $\delta^{18}\text{O}$  stack (Lisiecki and Raymo, 2005) (Supplementary Fig. S4)  
245 falls within our targeted time interval of 3.4 to 2.6 Ma. We used the more radiogenic  $\epsilon_{\text{Nd}}$   
246 peak at 260.4 mbsf in 910C as an additional tie point to define the MIS M2 glaciation  
247 (Supplementary Table S1), corresponding to a pronounced IRD pulse in Hole 911A  
248 (Supplementary Fig. S6). The calculated sedimentation rates between fix points either  
249 side of this new tie point are within the same order of magnitude (8 to 15 cm/ka) thus  
250 justifying this additional age fix point. The age of the “Datum A” corresponding to the depth  
251 223 mbsf was constrained based on the occurrence of calcareous nanofossils in 910C

252 and 911A (Sato and Kameo, 1996) and is slightly shifted from the original age of 2.78 Ma  
253 (Knies et al., 2014b) to 2.83 Ma in the revised age model (Supplementary Table S1).  
254 Together with the new tie points for biostratigraphic “Datum A” and shifted radiogenic  $\epsilon_{Nd}$   
255 peak to MIS M2, we used the linear sedimentation rates between all tie points to establish  
256 the age model for 910C between 3.4 - 2.6 Ma (Supplementary Table S1). Based on the  
257 revised chronology, the most negative excursion in the authigenic  $\epsilon_{Nd}$  profile is now shifted  
258 from 2.981 to 3.081 Ma, while the most positive excursion defines the MIS M2 glaciation  
259 (Supplementary Fig. S4). Considering the uncertainty in our age model, it might be  
260 challenging to resolve all individual peaks and troughs corresponding to glacial-  
261 interglacial stages in our proxy records of authigenic  $\epsilon_{Nd}$ , biomarkers and  $CaCO_3$ ;  
262 however, the most prominent excursions in our proxy records during the mPWP can be  
263 resolved with confidence, which is the primary target interval of the present study. All  
264 information on previously published and new tie points are provided in Supplementary  
265 Table S1.

266

### 267 **3. Results**

#### 268 **3.1 Authigenic $\epsilon_{Nd}$ record from the Yermak Plateau.**

269 The new  $\epsilon_{Nd}$  record from the Yermak Plateau allows identification of the maximum limit of  
270 water mass exchange between the NAC and Arctic derived Polar waters (PW),  
271 particularly during the major climatic transitions of the Pliocene to the earliest Pleistocene  
272 (3.4–2.6 Ma). These include the MIS M2 glaciation (3.312–3.264 Ma), the mPWP (3.264–  
273 3.025 Ma) and the intensification of Northern Hemispheric glaciation (iNHG) at ~2.7 Ma  
274 ago. The authigenic  $\epsilon_{Nd}$  record shows long-term secular changes from -9.2 during the MIS

275 M2 glacial period to -14.4 (5.2  $\epsilon_{Nd}$  unit) during the mPWP; the modern value of -11.7  
276 reported (Lambelet et al., 2016) from the core site falling within this range. Thereafter, an  
277 increasing trend up to -7.8 at ~2.6 Ma is clearly discernable, with several prominent  
278 positive excursions associated with iNHG cold stages (Fig. 2c). Our  $\epsilon_{Nd}$  record for 910C  
279 exhibits a larger range (6.6  $\epsilon_{Nd}$  unit) compared to that of 911A (3.4  $\epsilon_{Nd}$  unit) (Teschner et  
280 al., 2016) within the time period 2.5–3.5 Ma (Fig. 2c), most likely due to the higher  
281 temporal resolution.

### 282 **3.2 Biomarker and CaCO<sub>3</sub> records.**

283 The occurrence of seasonal sea ice throughout the record is confirmed by the near  
284 continuous presence of the biomarkers IP<sub>25</sub> and HBI III (Fig. 2d). Although the  
285 concentration of HBI III is mainly lower than that of IP<sub>25</sub>, it is the more abundant biomarker  
286 during the mPWP (ca. 3.150–2.970 Ma), consistent with more productive open-water  
287 conditions, as also shown by the carbonate record, which reaches its highest values  
288 during the MIS KM1-K2 (~3.150-3.050 Ma) (Fig. 2f). The CaCO<sub>3</sub> abundance measured in  
289 the bulk sediments ranges from 0.5 to 6%; however, a sharp three fold increase from the  
290 mean value ~2% to 6% is evident during the mPWP, which coincides with the prominent  
291 negative excursion in the  $\epsilon_{Nd}$  record (Fig. 2f).

292

### 293 **4. Discussion**

294 The authigenic ferromanganese oxyhydroxide fraction extracted from the bulk sediments  
295 has been demonstrated to record the  $\epsilon_{Nd}$  signal of bottom waters of the Yermak Plateau  
296 (Teschner et al., 2016; Werner et al., 2014). Hence, temporal variations in authigenic  $\epsilon_{Nd}$   
297 during glacial-interglacial periods have been primarily attributed to watermass exchange

298 between the NAC and PW, changes in sediment provenance, and variable weathering  
299 input due to glacial erosion (Teschner et al., 2016). However, other factors/mechanisms  
300 that contributed to the past authigenic  $\epsilon_{Nd}$  variability are discussed in the following section.

#### 301 **4.1 Factors contributing to past authigenic $\epsilon_{Nd}$ variability**

302 ODP Hole 910C is placed in the mixing zone between Atlantic- and Arctic-derived  
303 waters (Fig. 1) and is therefore well suited to monitor the relative influence of two water  
304 masses: (i) relatively warmer, high salinity water (i.e. the NAC characterized by a less  
305 radiogenic Nd isotope signature and (ii) relatively cold and less saline water (i.e. Arctic-  
306 derived PW) characterized by more radiogenic Nd isotopes. In the open ocean away from  
307 ocean margins and regions of deep-water formation, Nd appears to behave quasi-  
308 conservatively (Rempfer et al., 2011). Therefore, the variability in authigenic  $\epsilon_{Nd}$  record  
309 from the open ocean is mainly explained by the mixing of water masses with distinct  $\epsilon_{Nd}$   
310 signatures (Lang et al., 2016). However, contributions from other sources of dissolved Nd  
311 can substantially influence the authigenic  $\epsilon_{Nd}$  record. Assuming that the modern  
312 geological and tectonic setting of the study region have largely remained stable over the  
313 past ~4.6 Ma (Knies et al., 2014a), we discuss the following potential mechanisms and  
314 factors that may have contributed to the variability and changes in the authigenic  $\epsilon_{Nd}$   
315 record of 910C: (i) changes in weathering regimes and sediment sources; (ii) boundary  
316 exchange processes; and (iii) volumetric exchange of the NAC and PW.

317 Dissolved radiogenic isotope signatures in seawater originate from weathering  
318 processes of the continental crust (Frank, 2002) and, therefore, the glacial-interglacial  
319 changes in chemical weathering could influence the  $\epsilon_{Nd}$  record. Teschner et al. (2016)  
320 reconstructed past water mass mixing and erosional inputs prior and post intensification

321 of Northern Hemisphere glaciation (iNHG, ~2.7 million years ago) based on records of  
322 radiogenic isotopes of Sr, Nd and Pb at ODP Hole 911A. Changes in the authigenic  $\epsilon_{Nd}$   
323 record were highlighted for two different scenarios; (i) prior to the iNHG, the Pb and Nd  
324 isotopes composition was characterized by unradiogenic values and low variability due  
325 to the limited extent of ice sheets. These observations are consistent with earlier  
326 inferences from the Arctic Ocean (Haley et al., 2007) and suggest constant erosional  
327 supply of material to the Yermak Plateau, most likely from local sources (i.e. Svalbard).  
328 (ii) After the iNHG, conditions changed dramatically with higher-amplitude  $\epsilon_{Nd}$  variability  
329 in both deep waters and detrital sediments inputs due to changes in weathering inputs  
330 associated with the waxing and waning of the Eurasian ice sheets, water mass exchange  
331 and increased supply of ice-rafted debris (IRD). Comparison of the IRD record (Knies et  
332 al., 2014b) with our  $\epsilon_{Nd}$  record, shows higher IRD flux during the periods of M2 and iNHG  
333 (~2.7 Ma), and low and stable IRD fluxes during the mPWP (Supplementary Fig. S6). The  
334 latter corresponds to the IRD record from Site U1307 on Eirik Drift where coarse IRD is  
335 largely absent during the mPWP, and IRD is only present in small abundances during  
336 (de)glacials between ~3 Ma and 2.75 Ma. Therefore, higher variability in IRD supply and  
337 change of its sources could influence the authigenic  $\epsilon_{Nd}$  record in 910C; however, this can  
338 be excluded for the interglacial periods prior to the iNHG, particularly during the mPWP.  
339 It is also important to note that the timing of the iNHGs was further shifted to post MIS G2  
340 (2.64 Ma) based on the Pb isotope and geochemical studies of the IRD on the lower  
341 eastern flank of the Reykjanes Ridge (Bailey et al., 2013). Therefore, we suggest that the  
342 observed variability and changes in the radiogenic Nd isotope record in 910C is affected  
343 by glacial weathering input probably during the MIS M2 glaciation and after the iNHG. In



344 contrast, it is unlikely to be significantly affected by the changes in chemical weathering  
345 inputs and/sediment transport from distant sources during our targeted time interval of  
346 mPWP due to the stability of the climatic conditions and glacial erosion was rather limited.

347         The chemical weathering of Iceland-derived basaltic material can influence the Nd  
348 isotope composition of the NAC resulting in a shift towards more radiogenic values.  
349 However, in an earlier study, it has been suggested that present day exchange with  
350 Iceland derived basaltic material does not affect the deep water  $\epsilon_{Nd}$  signature of the main  
351 path of North Atlantic inflow, although it can influence the signature of southward flowing  
352 currents such as the East Greenland Current (Chen et al., 2012; Lacan and Jeandel,  
353 2004).

354         Seawater interactions with the continental margins (boundary exchange) could be  
355 a potential source for radiogenic isotope signatures of seawater, particularly in the Nordic  
356 Seas where basaltic formations are highly susceptible to dissolution and exchange with  
357 seawater (Chen et al., 2012; Lacan and Jeandel, 2004). The effects of boundary  
358 exchange have been reported from different continental margins in the subpolar regions  
359 including the Nordic Seas, and model results confirmed the importance of this input  
360 mechanism (Rempfer et al., 2011). Due to the large shelf areas of the Arctic Ocean,  
361 boundary exchange might be expected to be significant, although the water column data  
362 available so far do not provide clear evidence for this process (Andersson et al., 2008).  
363 Further, Laupkert et al (2017) suggested recently that  $\epsilon_{Nd}$  values around -10 are present  
364 in the eastern and western Fram Strait below ~500 m, implying that there is no evidence  
365 for boundary exchange processes influencing the  $\epsilon_{Nd}$  record to a significant extent on the  
366 Yermak Plateau.

367 In summary, with the absence of any significant ice-rafting prior to ~2.7 Ma (except  
368 MIS M2) in the Nordic Seas (Fig. 2e), increased sea surface temperatures (SST) by 3–  
369 7°C (Lawrence et al., 2009) between 3.4 and 2.6 Ma compared to the Holocene mean  
370 annual SST (Fig. 2g; dashed line) (Calvo et al., 2002), and thus no widespread Northern  
371 Hemisphere glacial advances, we attribute the large range in  $\epsilon_{Nd}$  (-14.8 to -9.0) in 910C  
372 prior to the iNHG to changes in watermass circulation rather than to variable glacial  
373 weathering input. As such, the prominent negative excursion in the  $\epsilon_{Nd}$  record during the  
374 mPWP (i.e. -14.4  $\epsilon_{Nd}$  units; Fig. 2c) is most likely due to an increase in volume transport  
375 of the NAC, resulting in an Atlantic-dominated climate regime of the Eurasian sector of  
376 the Arctic Ocean. Further, the prominent negative excursion in  $\epsilon_{Nd}$  record coincides with  
377 a sharp three-fold increase in  $CaCO_3$  abundance during the mPWP (Fig. 2f). This  
378 suggests an increased flow of warm NAC with higher pH resulted in better preservation  
379 of carbonate and/or increase in productivity during interglacial periods in the eastern Fram  
380 Strait (Supplementary Figs. S7b, d, e), consistent with earlier reports from modern and  
381 Quaternary sediments (Huber et al., 2000).

382 To test the hypothesis of increased “Atlantification” and its concurrent sea ice  
383 decline further, we quantified the volumetric changes of the AW-derived water masses  
384 and sea ice concentration at 910C using (1) a simplified binary mixing model by  
385 constraining the end member values of  $\epsilon_{Nd}$  for two water masses and (2) semi-quantitative  
386 estimates of spring sea ice concentration (%SpSIC) based on a regional calibration of  
387 biomarker distributions in modern sediments (Smik et al., 2016).

#### 388 **4.2 Quantifying water mass exchange based on authigenic $\epsilon_{Nd}$ record**

389           Compilation and reassessment of seawater Nd data from the literature shows that  
390 the characteristic NAC  $\epsilon_{\text{Nd}}$  signature near its origin in the inter-gyre region (north of 46°  
391 N) displays  $\epsilon_{\text{Nd}}$  values between  $-14.0 \pm 0.3$  and  $-15.1 \pm 0.3$  (Dubois-Dauphin et al., 2017),  
392 which changes gradually during transport across the Arctic Mediterranean due to mixing  
393 of more radiogenic signatures of PW ( $\epsilon_{\text{Nd}} = -9.9 \pm 0.7$ , 1 SD (standard deviation) and [Nd]  
394  $=27.1$ ) (Laukert et al., 2017). We have assigned  $\epsilon_{\text{Nd}}$  and [Nd] values for the NAC ( $-15 \pm 1$   
395 (1 SD) and  $16 \pm 1$  pmol/kg) and PW ( $-9.9 \pm 1$  (1 SD) and  $27 \pm 1$  pmol/kg) end-members,  
396 respectively, which are clearly distinct from the modern value in the Fram Strait (mean  
397  $\epsilon_{\text{Nd}} = -11.7 \pm 0.8$  (2SD)) (Laukert et al., 2017). With this identification of suitable end-  
398 member values for  $\epsilon_{\text{Nd}}$ , we therefore adopt a simple binary mixing approach for the  
399 determination of the percentage Atlantic water component (%AWC) on the assumption  
400 that Nd behaves quasi-conservatively and end-member compositions were invariant  
401 during the studied time interval. Such assumptions are discussed in more detail in the  
402 Supplementary Note 2. Meanwhile, we note that this method was successfully employed  
403 in a previous study (Lang et al., 2016) using a Nd isotope record from the late Pliocene  
404 (3.3–2.4 Ma ago) to quantify the mixing proportion of southern source water and north  
405 Atlantic deep water (NADW) in the North Atlantic.

#### 406 **4.2.1 Binary estimates of Atlantic water mass mixing using authigenic $\epsilon_{\text{Nd}}$ record**

407 We have used the  $\epsilon_{\text{Nd}}$  record from 910C to generate the semi-quantitative estimate of  
408 water-mass mixing between NAC and PW during the Late Pliocene to early Pleistocene  
409 (~3.4 - 2.6 Ma). The underlying assumptions of this approach are: (i) Nd isotopes exhibit  
410 quasi-conservative behaviour, (ii) mixing of Atlantic- and Arctic-derived waters at 910C is  
411 binary, and (iii) modern day end-members have been invariant between 3.4 and 2.6 Ma.

412 We used the following binary mixing equation constrained by our current understanding  
 413 of end-member compositions:

$$414 \quad \varepsilon_{Nd_{910C}} = \frac{\varepsilon_{Nd_{AW}} * C_{AW} * f_{AW} + \varepsilon_{Nd_{PW}} * C_{PW} * f_{PW}}{C_{AW} * f_{AW} + C_{PW} * f_{PW}} \quad Eq (2)$$

$$415 \quad f_{AW} + f_{PW} = 1 \quad Eq (3)$$

416 where  $\%AWC_{\varepsilon_{Nd}} = f_{AW} * 100$  is the relative contribution of Atlantic water component to  
 417 910C ( $\%PWC_{\varepsilon_{Nd}} = 100 - \%AWC_{\varepsilon_{Nd}}$ ),  $C_{PW}$  and  $C_{AW}$  represent the concentration of Nd in  
 418 the Arctic (PW) and the Atlantic (AW),  $\varepsilon_{Nd910C}$  is the value of Nd isotope compositions of  
 419 sediment leach from 910C, and  $\varepsilon_{AW}$  and  $\varepsilon_{PW}$  are the end-members of isotope composition  
 420 of Atlantic and Arctic water masses, respectively.  $f_{AW}$  and  $f_{PW}$  represent the fractions of  
 421 Nd coming from the Atlantic (AW) and Arctic (PW) waters.

422 In order to validate the use of this binary mixing model to 910C, we also calibrated  
 423 our approach by comparison of semi-quantitative estimates of modern day volume  
 424 transport with in situ observations. We thus estimated the modern day volume transport  
 425 of NAC using a contemporary  $\varepsilon_{Nd}$  value at the borehole site of 910C and compared that  
 426 with a mooring-based observation (Beszczynska-Moeller et al., 2012). Our estimate of  
 427 %NAC based on  $\varepsilon_{Nd}$  ( $47 \pm 9\%$ ) (Supplementary Fig. S9) compares well with a value of  $45$   
 428  $\pm 5\%$  (Supplementary Fig. S9b) measured from an array of moorings in Fram Strait ( $78^\circ$   
 429  $50' N$ ) over the period 1997–2010 (Beszczynska-Moeller et al., 2012).

430 We have determined the uncertainty associated fractions of NAC volume  
 431 estimates using a Monte-Carlo error propagation method with 10,000 iterations in  
 432 MATLAB, which is represented as an error envelop (at 95% confidence) (Fig. 3a).

433 However, we offer some caution that our %NAC estimates may be subject to changes in  
434 the future when more suitable archives allow generation of orbital resolution records of  
435 NAC and PW end-member behaviour. For now, the uncertainties reported here for our  
436  $\epsilon_{Nd}$ -based estimates of %NAC may be underestimated due to limited knowledge of end-  
437 member  $\epsilon_{Nd}$  values for Atlantic and Arctic waters. On the other hand, our main conclusions  
438 over our targeted time interval (3.4 – 2.6 Ma) are not influenced by such uncertainties.

439 Our estimates of %AWC in 910C indicate three distinct peaks with values close to  
440 100%, indicating the presence of a dominant Atlantic watermass in the water column  
441 during the three interglacial events (Haywood et al., 2013) (i.e. MIS KM3, K1, and G17)  
442 within or close to the mPWP; albeit within the limitation of the age constraints of 910C  
443 (Fig. 3 a). For MIS KM5c, with near-modern orbital configuration, the %AWC ( $51 \pm 11\%$ )  
444 was similar to today ( $45 \pm 5\%$ ) (Beszczynska-Moeller et al., 2012; Zhang et al., 2013) but  
445 was close to  $\sim 0\%$  during the preceding MIS M2 glaciation (3.305–3.285 Ma) (Fig. 3a),  
446 consistent with previous observations of a weaker NAC and concurrent cooling in the  
447 circum-Arctic (De Schepper et al., 2015). Importantly, although %AWC estimates for the  
448 glacial periods (M2 and iNHGs) might potentially suffer higher uncertainty due to  
449 enhanced IRD flux and weathering inputs associated with higher glacial activity, such  
450 effects during the mPWP are likely insignificant, in practice, due to the relatively stable  
451 climate and lower IRD fluxes (Blake-Mizen et al., 2019; Knies et al., 2014a) (Fig. 2e and  
452 Supplementary Fig. S6). Pertinent to our reconstructed reduced flux of %AWC during the  
453 M2 glaciation, we note that a similar situation has been reported for MIS6 based on  
454 authigenic coupled isotope records of Nd and Hf from the central Arctic Ocean (Chen et  
455 al., 2012).

## 456 **4.2 Sea ice reconstruction**

457 Extensive sea ice cover (>60% SpSIC) prevailed from 3.36–3.18 Ma, including maximum  
458 extent during MIS M2 (Fig. 3b). Thereafter, %SpSIC reduced substantially. According to  
459 Smik et al. (2016), biomarker-based %SpSIC estimates above ca. 68% also imply the  
460 occurrence of some summer sea ice (>5% SuSIC) (Supplementary Fig. S3). Similarly,  
461 while the occurrence of some summer sea ice was a common feature up to ca. 3.18 Ma  
462 (Fig. 3b), coincident with consistently low %AWC (i.e. below the modern value of 45%;  
463 Fig. 3a), ice-free summers were likely a common feature at the Yermak Plateau  
464 thereafter, especially during the mPWP. Change-point analysis carried out on our  
465 %SpSIC estimates shows a statistically significant decrease of ca. 30–35% starting at ca.  
466 3.15 Ma before increasing again at ca. 2.97 Ma (Supplementary Fig. S2). Prior to this,  
467 extensive sea ice cover similar to the modern (spring) maximum prevailed, including  
468 maximum extent during MIS M2 when the %AWC was at a minimum (Fig. 3a, b). The  
469 reduction in SpSIC during the mPWP likely reflects a response to increased %AWC,  
470 analogous to observations made for eastern Fram Strait (Spielhagen et al., 2011) and the  
471 Barents Sea spanning recent decades/centuries (Cabedo-Sanz and Belt, 2016). Similar  
472 observations have been reported for the Early Holocene and the Last Interglacial (MIS  
473 5e/Eemian), implying that increased Atlantic Water inflow is one important factor  
474 controlling sea ice conditions in an area covering northern Svalbard/Yermak Plateau and  
475 the northern Barents Sea continental margin (Müller et al., 2012; Belt et al., 2015; Stein  
476 et al., 2017). According to our SpSIC estimates, maximum sea ice extent during the  
477 mPWP exhibited a closer resemblance to that of modern-day late summer (i.e. minimum)  
478 conditions (Fig. 3b). These new data support the boundary conditions used in the

479 Pliocene Research, Interpretation and Synoptic Mapping (PRISM) project, which  
480 assumes a conservative sea ice extent, an ice-free Arctic Ocean in summer, and winter  
481 sea ice conditions approximately equivalent to modern summer ice extent (Dowsett et al.,  
482 2010).

### 483 **4.3 Forcing factors modulating North Atlantic volume transport and its impact**

484 Our new reconstructions of watermass mixing and carbonate abundances follow  
485 the periodicities of eccentricity (~100 ka), obliquity (~40) and precessional cycles (~20  
486 ka) (Figs. 4a, b). Further, the %AWC and %SpSIC records show good alignment with the  
487 eccentricity (Fig. 4c) and summer insolation in the northern hemisphere (Fig. 5b), implying  
488 orbitally-paced control over changes to oceanic heat flow into the Arctic Ocean. Cross  
489 wavelet analysis highlights the common highest power between these two time series in  
490 colour bands (Fig. 4c). The vector arrows indicate an in-phase relation (pointing  
491 rightward) during 3.2 – 2.9 Ma at the eccentricity band (64 – 128 ka), implying that  
492 orbitally-controlled, enhanced NAC resulted in an increase in marine productivity and  
493 reduction in sea ice coverage during the mPWP (Supplementary Fig. S7c, d). In  
494 particular, during the three interglacials with high eccentricity (i.e. KM3, K1, and G17; Fig.  
495 5a, d), increased seasonality combined with warmer summers (higher solar insolation) in  
496 the Northern Hemisphere (Fig. 5a, b) may have resulted in an increased oceanic heat  
497 transport with consequential decline in Arctic sea ice extent and polar amplification of this  
498 warming. Alternatively, an orbitally-forced reduction in Arctic sea ice coverage may have  
499 changed buoyancy and salinity in the Atlantic, and thus been responsible for increased  
500 northward ocean heat transport during mPWP interglacials leading to a strongly positive  
501 ice-albedo feedback. Our proxy data do not reveal any correspondence with variable

502 atmospheric CO<sub>2</sub> estimates (Figure 5c, d and e) implying only a minor influence of  
503 greenhouse gas-derived radiative forcing in modulating NAC heat transport and reduction  
504 in Arctic sea ice. Further, tectonic changes could have driven circulation changes as has  
505 been reported for the Bering Strait and Nordic Sea related to reconfiguration of oceanic  
506 gateways (De Schepper et al., 2015; Horikawa et al., 2015). However, the strong signal  
507 of orbital cycles in our proxy records clearly indicate that the orbital forcing played the  
508 dominant role over all other controlling factors.

509         Based on multi-proxy records, it has been inferred that the Atlantic Meridional  
510 Overturning Circulation (AMOC) was significantly stronger in the mPWP compared to  
511 today (Raymo et al., 1996; Frank et al., 2002; Dowsett et al., 2009), which could have  
512 contributed to enhanced northward heat transport during the mPWP interglacials  
513 (Dowsett et al., 2009; Lawrence et al., 2010; Naafs et al., 2012), consistent with our  
514 findings. However, the exact mechanism(s) responsible for changes in northward heat  
515 transport remain a topic of debate (Haywood et al., 2016; Haywood et al., 2013; Zhang  
516 et al., 2013), but could potentially be resolved through further ocean modelling studies  
517 that integrate the new proxy data presented herein.

518         Regardless of the ultimate driver(s), our estimates of %AWC show a clear  
519 dominance of a warm and well-mixed Atlantic-dominated climate regime in the Eurasian  
520 Arctic during MIS KM3, K1, and (within the given age uncertainties) G17, with lower than  
521 modern sea ice extent (including ice-free summers) and higher marine productivity,  
522 consistent with modeled and reconstructed amplification of Arctic surface temperatures  
523 (Ballantyne et al., 2013), and a rise in annual mean surface air temperatures between  
524 4°C to 5°C ( $\Delta t = \text{Plio-KM5c} - \text{pre-industrial}$ ) (Prescott et al., 2018). This implies that the



525 increase in %AWC with concurrent reduction in %SpSIC during these mPWP interglacials  
526 resembles modern observations of an advanced “Atlantification” of the study region  
527 (Cabedo-Sanz and Belt, 2016; Naafs et al., 2010; Spielhagen et al., 2011).

528 The conclusion of increased “Atlantification” of the Arctic during the mPWP from  
529 our new proxy records from the Atlantic-Arctic gateway confirms previous studies from  
530 lower latitudes a (Naafs et al., 2010; Raymo et al., 1996). Since current generation models  
531 have not yet been validated against any proxy-based observations of “Atlantification” in  
532 the Eurasian sector of the Arctic during the mPWP, our new Nd isotope, biomarker and  
533 CaCO<sub>3</sub> records thus provide important input for testing the robustness of future climate  
534 modelling for northern high latitude settings.

## 535 **5. Conclusions**

536 Our new Nd isotope record of past water mass exchange in the Atlantic-Arctic  
537 gateway relative to the modern-day setting suggests a near doubling of NAC volume  
538 transport during mPWP interglacials KM3, K1, and G17 with different orbital  
539 configurations and thus stronger seasonality than today. This resulted in a warm and well-  
540 mixed Atlantic-dominated climate regime (“Atlantification”) of the Eurasian sector of the  
541 Arctic Ocean, reduced spring sea ice concentration, and the possibility of ice-free  
542 conditions during summers. In contrast, the mPWP interglacial with near-modern orbits  
543 (MIS KM5c) does not show significant deviation from today’s NAC volume transport or  
544 sea ice extent. This study demonstrates a dominant role of orbital forcing in modulating  
545 northward ocean heat transport and Arctic sea ice coverage during the mPWP. It also  
546 highlights the importance of improving data-model comparison studies for the Arctic  
547 Ocean that integrate reconstructions of water mass flow and ocean circulation, as well as

548 temperature and sea ice, for climate states of the past that may be analogous to the  
549 future.

#### 550 **Data availability**

551 All the data are provided in the supplementary and also will be archived in PANGEA upon acceptance of  
552 the manuscript.

#### 553 **Code availability**

554 The MATLAB codes for uncertainty estimates on the volumetric water fractions of Atlantic water are  
555 available from the corresponding author W. Rahaman on request.

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562 codes for Monte-Carlo error propagation on the water fractions estimates. Finally, we thank two  
563 anonymous reviewers who provided supportive and constructive feedback on the initial version  
564 of this manuscript.

#### 565 **Author contributions**

566 W.R., J.K. and M.T. designed the study. W.R., J.K., S.T.B and A.H wrote most of the text. Analysis  
567 of model results was completed by A.H and J.T. Mo.T. and L.N. analyzed authigenic Nd isotope  
568 compositions in bulk sediment. L.S, D.K and S.T.B. measured the concentrations of the IP<sub>25</sub> and  
569 HBI III biomarkers in bulk sediments and interpreted outcomes. All authors contributed to  
570 interpreting results, discussion and improvement of this paper.

571 **Competing interests**

572 The authors declare no competing interests.

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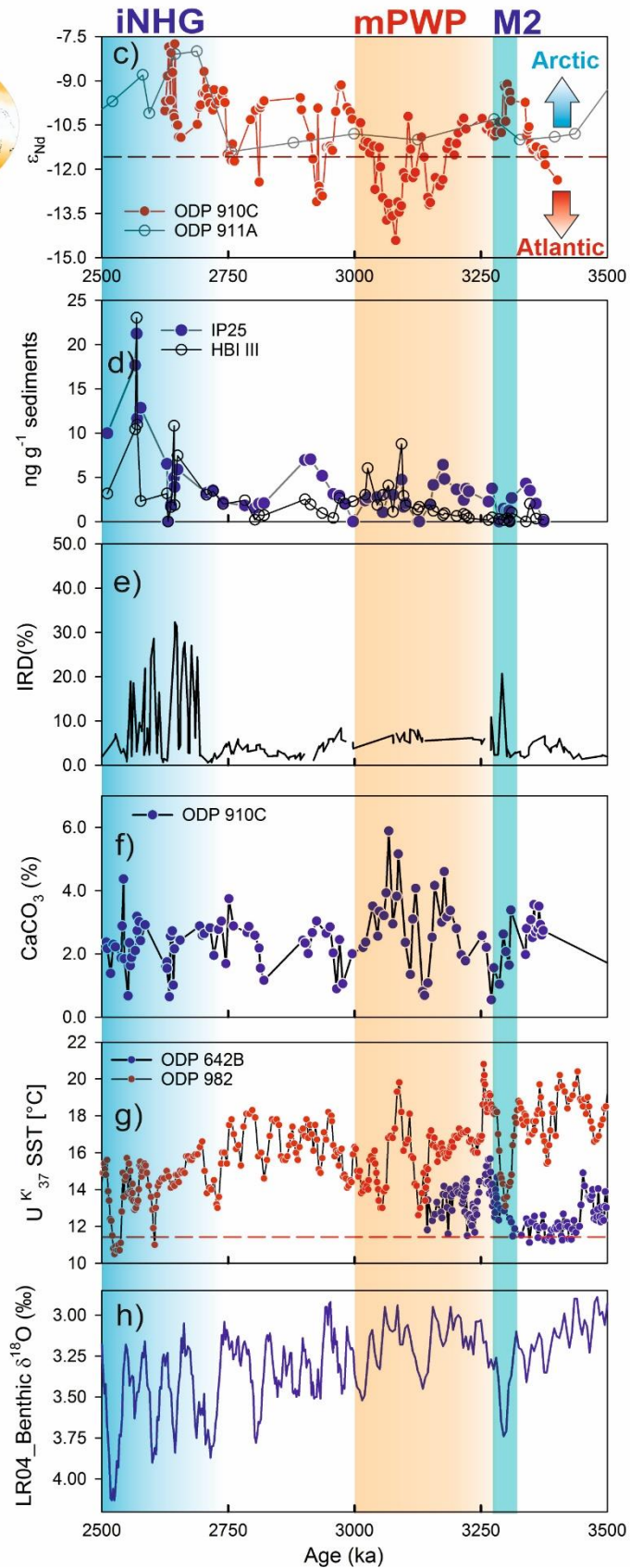
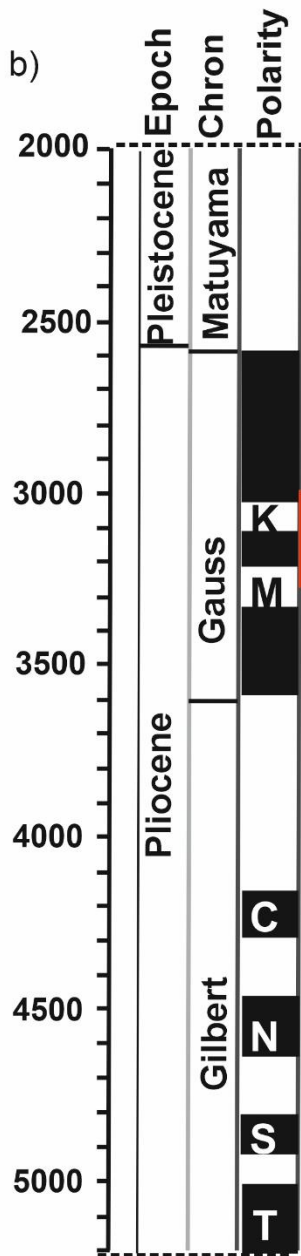
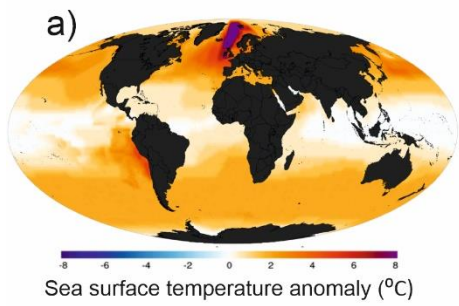


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575 **Fig. 1. Water mass circulation and their characteristic Nd isotope compositions.** Locations of ODP

576 Sites 910 (red star) and 911 (filled yellow circle) with schematic flow paths of the main water masses in the

577 northern North Atlantic and Nordic Seas and their present-day  $\epsilon_{Nd}$  signatures(Teschner et al., 2016). Dark  
578 red arrows mark the warm inflowing Atlantic water; dark blue arrows represent the cold deep and surface  
579 water masses flowing out of the Arctic Ocean(Andersson et al., 2008; François and Catherine, 2004; Lacan  
580 and Jeandel, 2004). White numbers mark the average  $\epsilon_{Nd}$  values of the bedrocks of Svalbard(Tütken et al.,  
581 2002), the Norwegian Caledonian Margin and Iceland(Laskar et al., 2004), the Putorana basalts in Russia  
582 (Sharma et al., 1992), and Greenland (François and Catherine, 2004). Positions of ODP site 982 (58° N,  
583 16° W) and ODP Hole 642B (67° 20' N, 2° 90' E) are shown.



585 **Fig. 2. Water mass exchange and associated changes in the Fram Strait during the Late-Pliocene**  
586 **and Pleistocene** (a) Sea surface temperature (SST) anomalies during the mPWP (~3.3 – 3.0 Ma)  
587 compared to today (Dowsett et al., 2009). b) Pliocene-Pleistocene time scale with paleo-magnetic reversals.  
588 Red block represents the time slice of mPWP. c) Authigenic  $\epsilon_{Nd}$  record from 910C (this study) and 911A  
589 (Teschner et al., 2016). d) Record of sea ice and open water biomarkers IP<sub>25</sub> and HBI III. (e) Record of IRD  
590 (%) from ODP site 911A (Knies et al., 2014b). f) Record of CaCO<sub>3</sub> abundance (wt. %). g) Record of  
591 alkenon UK<sub>37</sub> derived SST at ODP Sites 982(Lawrence et al., 2009) (58° N, 16° W) and ODP Hole  
592 642B(Bachem et al., 2017) (67° 20' N, 2° 90' E). Dashed lines indicate Holocene average SSTs for the  
593 Norwegian Sea(Calvo et al., 2002) at 11.6 °C. h) Benthic  $\delta^{18}O$  (LR04) stack(Lisiecki and Raymo, 2005).  
594 The shaded bands represent the major climatic transitions: M2 glaciation (blue shade, 3.312–3.264 Ma),  
595 mid-Pliocene Warm Period (mPWP) (brown shade, 3.3–3.0 Ma) and intensification Northern Hemisphere  
596 glaciation (iNHG, ~2.7 Ma).

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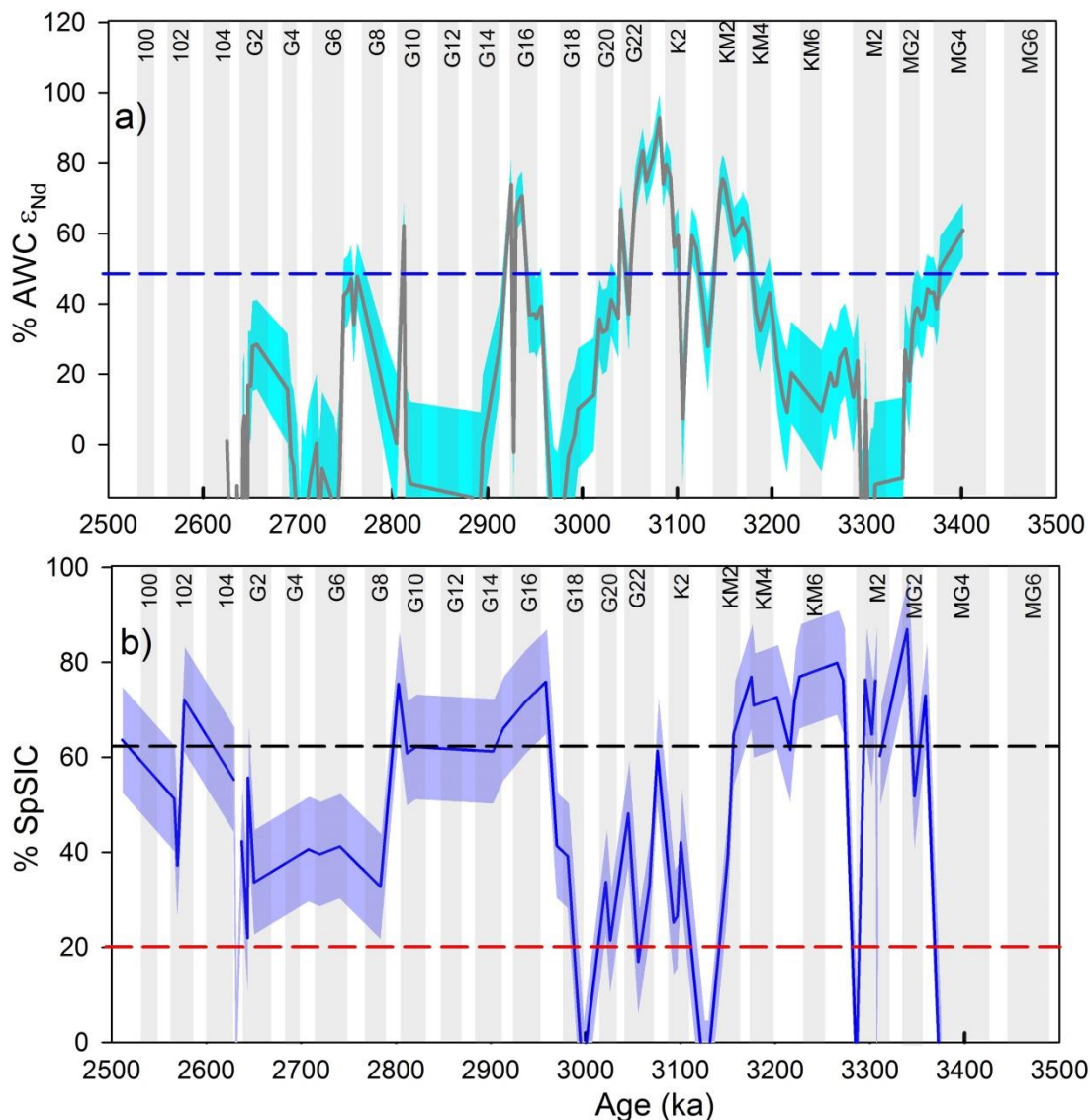
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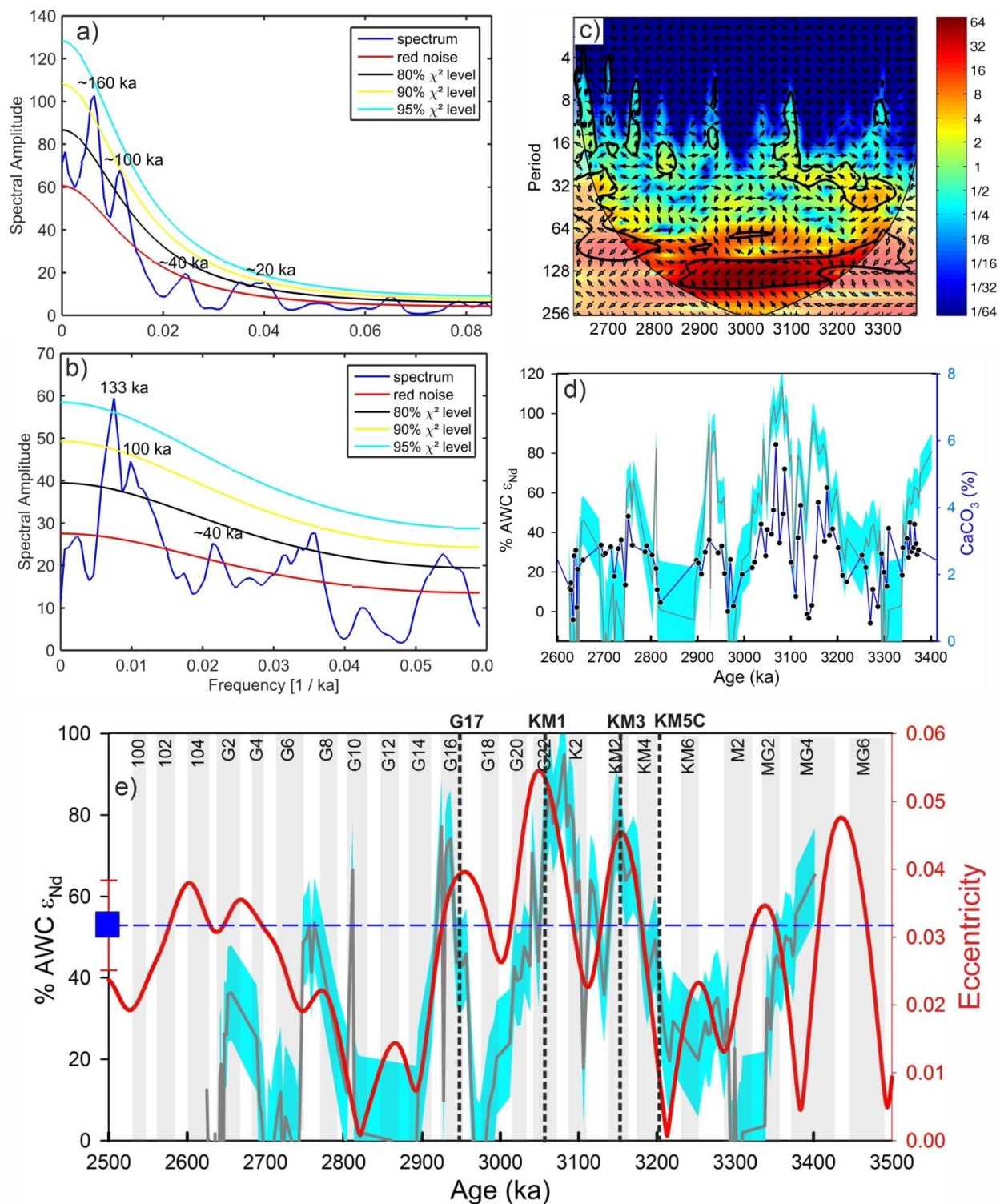
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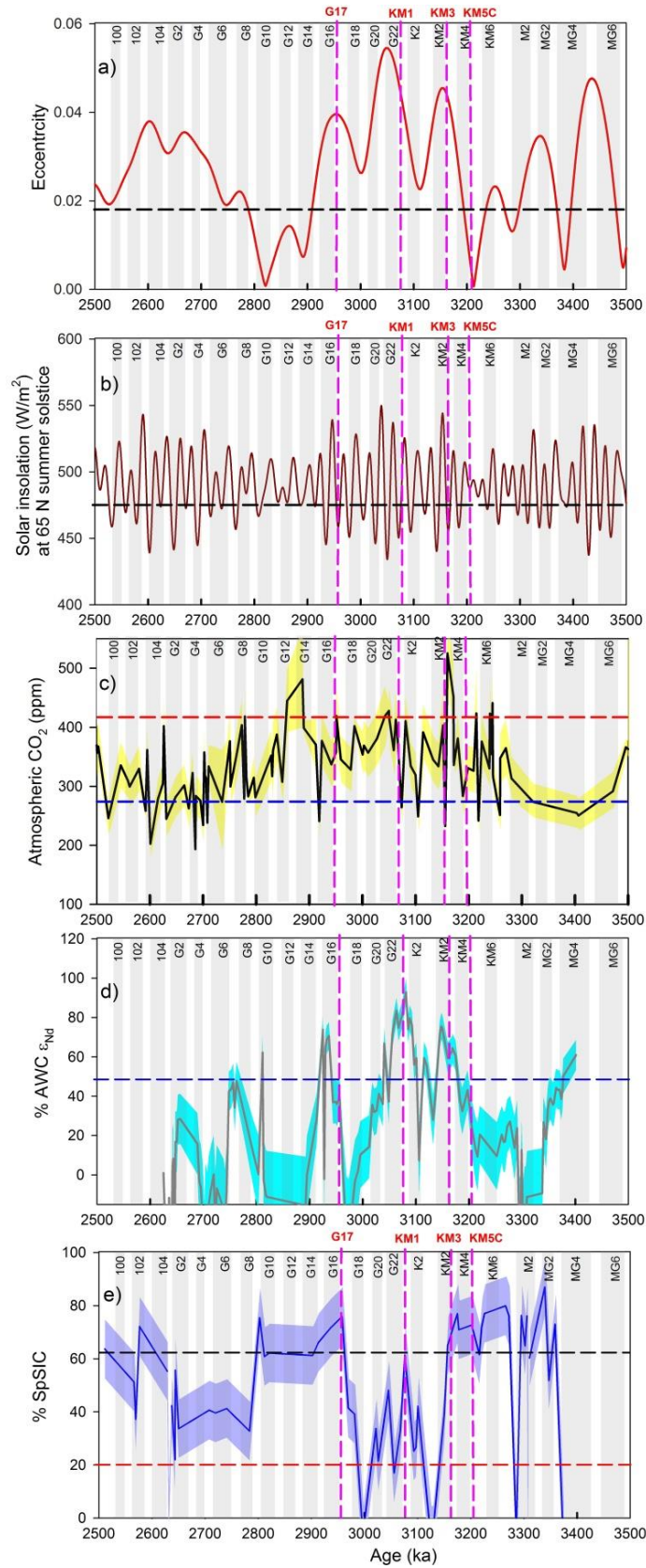
609 **Fig. 3. North Atlantic (NAC) volume transport and corresponding Arctic Sea Ice changes.** a) Fraction  
 610 of Atlantic Water Component (%AWC $\epsilon_{Nd}$ ). Dark gray line: Best estimate. Shading: 95% confidence interval.  
 611 Blue dashed line indicates modern Atlantic flow based on mooring estimate (Beszczynska-Moeller et al.,  
 612 2012). (b) Spring sea ice (%). Solid blue line represents mean value. Blue shade represents root-mean-  
 613 square error (RMSE) on the mean value. Blue and red dashed horizontal lines represent the modern mean  
 614 (1988-2017, NSIDC) sea ice maximum (62%, Apr-June; spring) and minimum (20%, September; late  
 615 summer) concentrations at the core site.



618 **Fig. 4. Identification of orbital cycles in proxy records.** Power spectrum analysis of (a) NAC volume  
 619 transport and (b) CaCO<sub>3</sub> abundance (%) records from the Yermak Plateau. They show periodicities of orbital



620 cycles at different significance level. c) Cross wavelet analysis of two time series highlights the common  
621 highest power between these two time series which is highlighted in color code. Vector arrow indicates  
622 phase relation between the time series. The 5% significance level against red noise is shown as a thick  
623 contour. The thin solid line indicates cone of influence. The relative phase relationship is shown as arrows  
624 (with in-phase pointing right, anti-phase pointing left, and  $\epsilon_{Nd}$  leading  $CaCO_3$  by  $90^\circ$  pointing straight down  
625 and vice versa). d) Comparison of the record of NAC with  $CaCO_3$  percentage, an indicator of marine  
626 productivity. Both the curves overall follow the same pattern; the highest abundance in calcium carbonate  
627 and thus the highest productivity was observed during mPWP when NAC flow was maximum.



629 **Fig. 5 Role of orbital forcing in modulating watermass exchange and spring sea ice extent.** (a)  
630 Record of eccentricity (Laskar et al., 2004). Dashed horizontal line represents modern eccentricity. Vertical  
631 dashed lines (pink) indicate four interglacial periods KM5C, KM3, KM1 and G17. Among these four  
632 interglacial periods, KM5C is most similar to that of the modern orbital forcing(Haywood and Valdes, 2004).  
633 (b) Record of solar insolation at 60° N summer solstice (Laskar et al., 2004). Dashed line represents modern  
634 value of summer insolation. c) Record of atmospheric pCO<sub>2</sub> derived from boron isotopes ( $\delta^{11}\text{B}$ ) (Foster et  
635 al., 2017). Yellow band represents error envelopes ( $1\sigma$  SD). Black and red colour dashed lines represent  
636 pre-industrial CO<sub>2</sub> (280 ppm) and present CO<sub>2</sub> (~410 ppm) level. These forcing parameters are compared  
637 with fraction of d) Atlantic Water (%AWC) and (e) Spring sea ice (%).

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