Deep Ocean storage of heat and CO2 in the Fram Strait, Arctic Ocean during the last glacial period

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Abstract

The Fram Strait is the only deep gateway between the Arctic Ocean and the Nordic Seas and thus is a key area to study past changes in ocean circulation and the marine carbon cycle. Here, we study deep ocean temperature, δ 180, carbonate chemistry (i.e., carbonate ion concentration, [CO32-]), and nutrient content in the Fram Strait during the late glacial (35,000–19,000 years BP) and the Holocene based on benthic foraminiferal geochemistry and carbon cycle modelling. Our results indicate a thickening of Atlantic water penetrating into the northern Nordic Seas, forming a subsurface Atlantic intermediate water layer reaching to at least ~2600 m water depth during most of the late glacial period. The recirculating Atlantic layer was characterized by relatively high [CO32-] and low δ 13C during the late glacial, and provides evidence for a Nordic Seas source to the glacial North Atlantic intermediate water flowing at 2000–3000 m water depth, most likely via the Denmark Strait. In addition, we discuss evidence for enhanced terrestrial carbon input to the Nordic Seas at ~23.5 ka. Comparing our δ 13C and qualitative [CO32-] records with results of carbon cycle box modelling suggests that the total terrestrial CO2 release during this carbon input event was low, slow, or directly to the atmosphere.

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27 Key points

- 28 Ambiguous interspecies differences in benthic foraminiferal δ^{18} O from the Nordic
- 29 Seas for the last glacial maximum are due to bioturbation.
- 30 Deep ocean temperature and δ^{18} O in central Fram Strait were ~1.5 °C and 1‰ higher
- 31 during the last glacial maximum.
- 32 Intermittent Nordic Seas outflow through Denmark Strait could have been the source
- of glacial North Atlantic water at 2000–3000 m water depth.

34

36 Abstract

The Fram Strait is the only deep gateway between the Arctic Ocean and the Nordic 37 Seas and thus is a key area to study past changes in ocean circulation and the marine carbon 38 cycle. Here, we study deep ocean temperature, δ^{18} O, carbonate chemistry (i.e., carbonate ion 39 concentration, $[CO_3^{2-1}]$, and nutrient content in the Fram Strait during the late glacial (35,000– 40 19,000 years BP) and the Holocene based on benthic foraminiferal geochemistry and carbon 41 42 cycle modelling. Our results indicate a thickening of Atlantic water penetrating into the northern Nordic Seas, forming a subsurface Atlantic intermediate water layer reaching to at 43 least ~2600 m water depth during most of the late glacial period. The recirculating Atlantic 44 layer was characterized by relatively high $[CO_3^{2-}]$ and low $\delta^{13}C$ during the late glacial, and 45 provides evidence for a Nordic Seas source to the glacial North Atlantic intermediate water 46 flowing at 2000–3000 m water depth, most likely via the Denmark Strait. In addition, we 47 discuss evidence for enhanced terrestrial carbon input to the Nordic Seas at ~23.5 ka. 48 Comparing our δ^{13} C and qualitative [CO₃²⁻] records with results of carbon cycle box 49 modelling suggests that the total terrestrial CO₂ release during this carbon input event was 50 low, slow, or directly to the atmosphere. 51

52

53 **1 Introduction**

Late Pleistocene glaciations were characterized by millennial- and decadal- scale variations in global/regional climate and atmospheric carbon dioxide levels (i.e., pCO₂), superimposed on gradual longer-term trends of increasing ice volume and decreasing atmospheric pCO₂ (e.g., Bard et al., 1990; Barker et al., 2011; Marcott et al., 2014; Bereiter et al., 2015). For example, Greenland ice core records show that the high latitude North Atlantic region underwent ~25 millennial-scale climate oscillations during the last glacial period

(~110,000–19,000 years ago; 110–19 ka), referred to as Dansgaard-Oeschger (DO) events 60 61 (Dansgaard et al., 1993). These DO events, consisting of warm interstadials and cold stadials, are characterized by an abrupt atmospheric warming over Greenland of 8–16 °C from cold 62 stadials to warm interstadials followed by a gradual cooling and eventually a sudden cooling 63 back to stadial conditions (Huber et al., 2006). Associated with DO events, high-resolution 64 records of atmospheric pCO₂ from Antarctic ice cores reveal millennial-scale (up to 25 ppm) 65 66 and centennial-scale (up to 10 ppm) variations (e.g., Marcott et al., 2014; Bereiter et al., 2015; Bauska et al., 2018). 67

The Atlantic Meridional Overturning Circulation (AMOC), which is a crucial 68 69 regulator of the earth's climate variability, is thought to have played a leading role in these (sub)millennial-scale variations in climate and atmospheric pCO₂ through its control on heat 70 redistribution and ocean-atmosphere gas exchange (e.g., Broecker, 1998; Fischer et al., 2010). 71 Modern exchange of surface and deep water between the Arctic Ocean, the Nordic Seas and 72 the North Atlantic Ocean represents the northern limb of the AMOC (Hansen and Østerhus, 73 74 2000). Northward inflow of warm Atlantic water across the Greenland-Scotland ridge advects 75 heat, salt and carbon to the Nordic Seas and the Arctic Ocean, where the major part of the inflow water densifies and returns as deep overflows to the North Atlantic (Hansen and 76 77 Østerhus, 2000). The evolution of this circulation pattern during the last glacial period and its contribution to glacial climate and atmospheric pCO₂ variations is still elusive. For example, 78 deep ocean exchanges between the Arctic Ocean and the Nordic Seas are particularly poorly 79 constrained during the Last Glacial Maximum (LGM; 25-19 ka), when global ice sheets 80 81 reached their maximum integrated volume (e.g., Clark et al., 2010). Moreover, carbon 82 exchanges between the deep ocean, the surface ocean and the atmosphere are even less constrained in the Arctic Ocean and Nordic Seas for the last glacial period. This is partly due 83 to limited availability of deep-sea sediments dating from the LGM from the central Arctic 84

Ocean, because of extensive sea ice cover resulting in very low or no sedimentation (e.g., Hanslik et al., 2010). In addition, δ^{18} O values measured in benthic foraminifera from the deep northern and central Nordic Seas show large and ambiguous interspecies differences (Bauch et al., 2001; Ezat et al., 2019).

The Fram Strait is an important gateway between the Arctic Ocean and the Nordic 89 Seas and thus is a key area to investigate the heat and carbon exchanges between the Arctic 90 91 Ocean, the Nordic Seas and the Atlantic Ocean. In addition, marine records from the deep 92 Fram Strait document enhanced flux of terrestrial carbon to the northern Nordic Seas at ~24 ka (Hebbeln et al., 1994). Changes in terrestrial carbon storage have also been suggested to 93 94 have played a role in some abrupt centennial-scale changes in atmospheric pCO₂ (e.g., Bauska et al., 2016). Given the vast and dynamic permafrost and subglacial carbon reservoirs in the 95 Arctic region (e.g., Vonk et al., 2012; Tarnocai et al., 2009; Köhler et al., 2014; Wadham et 96 al., 2019; Meyer et al., 2019), and the increased melting and release of huge icebergs from 97 surrounding ice sheets (e.g., Luckman et al., 2006; Bjørk et al., 2012; Mankoff et al., 2020), it 98 99 is imperative to study the impact of a possible 'terrestrial carbon mobilization' event at ~24 ka 100 on deep ocean carbonate chemistry and atmospheric pCO₂.

Here, we assess the interspecies differences in benthic foraminiferal δ^{18} O and study 101 the glacial evolution of deep-ocean temperatures, δ^{18} O and carbonate chemistry in the deep, 102 central Fram Strait (~2600 m water depth) for the 35–19 ka period using benthic foraminiferal 103 element/Ca, radiocarbon and isotope measurements from two deep ocean sediment cores 104 (~2600 m water depth). In addition, we use the CYCLOPS carbon cycle box model to assess 105 the effects of a possible 'terrestrial carbon mobilization' event at ~ 24 ka on atmospheric pCO₂ 106 107 and the carbonate chemistry of the deep 'Northern Component Water', and compare the model results with our proxy reconstructions. The overall purpose is to study the exchange of 108 deep water through the Fram Strait and explore the contribution of ocean circulation and 109

biogeochemical changes in the northern Nordic Seas to glacial climate and carbon cyclevariations.

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113 2 Material and methods

114This study is based on new and published data from two sediment cores HH12-946MC115(78°53' N; 01°45' W; 2637 m water depth) and HH12-948MC (78°52' N; 00°21' E; 2542 m

water depth) from the central Fram Strait (Ezat et al., 2019; Figure 1). Published results from

sediment cores HLY0503-18TC (88°45' N; 146°68' E; 2654 m water depth; Cronin et al.,

118 2012), PS1243 and MD99-2276 (69°37' N, 06°55' W, 2711 m water depth; Bauch et al., 2001;

119 Thornalley et al., 2015), 1294-4 (77° 59.9' N; 5°22.3' E; 2668 m water depth; Hebbeln et al.,

120 1994), 1295-5 (77°59.2' N; 2°24.8' E; 3112 m water depth; Hebbeln et al., 1994), and HM52-

43 (64° 25' N, 0.73° E, 2781 m water depth; Veum et al., 1992) are also closely compared

and discussed (Figure 1).

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124 **2.1 Stable carbon and oxygen isotopes**

125 We use published foraminiferal δ^{18} O and δ^{13} C records of the planktic species 126 *Neogloboquadrina pachyderma*, the shallow infaunal benthic species *Oridorsalis umbonatus* 127 and the epifaunal benthic species *Cibicidoides wuellerstorfi* from cores HH12-946MC and 128 HH12-948MC (Ezat et al., 2019). In order to increase the temporal resolution and extend the 129 investigated time interval, new measurements were obtained on *O. umbonatus* following the 130 methods described in Ezat et al. (2019).

131 In addition, new stable isotope measurements on *C. wuellerstorfi* from core HH12-

132 948MC were made after acid leaching. About 6 specimens of *C. wuellerstorfi* were crushed

and leached by 0.01M HCl for 15 minutes, including ultrasonication for 5 minutes.

134 Thereafter, the samples were rinsed immediately by milliQ water. The δ^{18} O and δ^{13} C were

then measured at the Godwin Laboratory for Paleoclimate Research, University of Cambridge, UK. These new measurements were performed to investigate the possibility that the low δ^{18} O previously recorded in *C. wuellerstorfi* (Bauch et al., 2001; Thornalley et al., 2015; Ezat et al., 2019) could be caused by diagenetic coating with extremely low δ^{18} O. It was not possible to perform this 'acid leaching' test on core HH12-946MC, because all specimens of *C. wuellerstorfi* in this core were already used for stable isotope analyses and radiocarbon dating (see below).

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143 2.2 Benthic element/Ca analyses and seawater temperature and $\delta^{18}O_{water}$ calculations

144 For element/Ca analyses, we used the most persistent and abundant benthic foraminiferal species in our study region, O. umbonatus (Bauch et al., 2001; Ezat et al., 2019). 145 A total of 25 to 30 pristine specimens of *O. umbonatus* were carefully picked from the 150-146 147 250 µm size fraction from sediment cores HH12-948MC and HH12-946MC. After crushing, the samples were cleaned following the reductive-oxidative method described in Pena et al. 148 149 (2005) and originally published in Boyle and Keigwin (1985). At the day of analyses, the samples were dissolved using 330 µl 0.1M HNO₃. After centrifugation for 5 minutes (7000 150 rpc), 300 µl of the samples were transferred to new acid-cleaned vials. For blank samples 151 152 (which have been treated with cleaning reagents as actual samples), 50 μ l of a high purity 2000 ppm Ca^{2+} solution was added before adding the acid, to enable determination of the 153 blanks as element/Ca. An aliquot of 50 µl of each sample was added to 200 µl 0.1M HNO₃ 154 and analyzed on an ICP-OES to determine their $[Ca^{2+}]$. Accordingly, the samples were re-155 diluted to 10 ppm [Ca²⁺] using '0.3M HF - 0.1M HNO₃' mixture and were analyzed by HR-156 ICP-MS at the University of Cambridge following Misra et al. (2014). Based on repeated 157 measurements of in-house standard solutions (n=11), the analytical precision (2 sd) for both 158 B/Ca and Mg/Ca is 0.9%. Blank samples (n=9) show average B/Ca and Mg/Ca of 2.5 159

µmol/mol and 0.027 mmol/mol, respectively, suggesting insignificant contamination from
vials or cleaning/dissolution reagents. Mn/Ca, Fe/Ca (except two samples) and Al/Ca (except
four samples) are < 25 µmol/mol, 50 µmol/mol and 50 µmol/mol respectively, indicating
negligible contamination by terrigenous materials or diagenetic coatings (e.g., Barker et al.,
2005).

Mg/Ca was converted to bottom water temperature (BWT; °C) using the equation 165 from Barrientos et al. (2018): (O. umbonatus Mg/Ca = A * $exp(0.102 \pm 0.01*BWT)$). The pre-166 exponential constant (A) is calibrated to our core-top samples from core HH12-948MC 167 168 (which are <2000 years BP) yielding a value of 1.45. Elderfield et al. (2006) has shown that Mg/Ca data of *O. umbonatus* from sites with BWT <2 °C do not follow a simple calibration 169 curve. This could be related to the 'carbonate ion' effect (e.g., Elderfield et al., 2006), partial 170 dissolution, and/or use of a wide size range of foraminiferal specimens (e.g., Tisserand et al., 171 2013). Nevertheless, the consistency between published BWT reconstructions based on 172 Mg/Ca measurements of O. umbonatus and independent BWT reconstructions from clumped 173 isotopes (Thornalley et al., 2015) and Mg/Ca measured on ostracods (Cronin et al., 2012) 174 from the central Nordic Seas (see Thornalley et al., 2015) provides confidence in the use of 175 this species in BWT reconstructions in our study region. 176 Seawater δ^{18} O was calculated by removing the temperature component of the δ^{18} O-O. 177

177 Seawater δ^{10} O was calculated by removing the temperature component of the δ^{10} -O 178 *umbonatus* using the equation from Shackleton (1974). To convert from the Pee Dee 179 Belemnite scale for carbonate δ^{18} O to Standard Mean Ocean Water for δ^{18} O-water, 0.2 was 180 added (Shackleton, 1974). Using other available equations (Bemis et al., 1998; Marchitto et 181 al., 2014) would change the amplitude of glacial-interglacial change in seawater δ^{18} O by 182 <0.1‰.

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184 **2.3 Radiocarbon analyses**

185	Four ¹⁴ C dates from core HH12-946MC on specimens of C. wuellerstorfi (~ 250–
186	$600 \ \mu g$ carbonate per sample) were measured following Gottschalk et al. (2018) using the
187	Mini Carbon Dating System (MICADAS) at the University of Bern (Szidat et al., 2014). In
188	brief, foraminiferal samples were loaded into septum-sealed glass vials. The air in the vials
189	was removed and replaced by He using two concentric needles inserted through the septum by
190	the automated carbonate handling system. The foraminiferal samples were then weakly
191	leached by adding 200 μl 0.01 M HCl for 3 min at room temperature. After a second flush of
192	the vials with He (to remove CO_2 produced during leaching), 0.5 mL 85% (~ 15 M)
193	orthophosphoric acid (H ₃ PO ₄) was added with a gas-tight syringe for carbonate dissolution.
194	The reaction was run to completion over night at a temperature of 70°C. To evaluate possible
195	methodological bias when comparing between 'MICADAS-based' C. wuellerstorfi and
196	published 'graphitization-based' O. umbonatus and N. pachyderma ¹⁴ C dates measured at the
197	¹⁴ Chrono Centre, Queen's University Belfast, Northern Ireland (Ezat et al., 2019), one N.
198	pachyderma ¹⁴ C date was measured by MICADAS following the methods outlined above.
199	The MICADAS-based N. pachyderma date is consistent with 'graphitization-based' N.
200	pachyderma dates (Table 1; Supplemental Figure 1), indicating negligible methodological
201	bias.

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203 **2.4 Age models**

The age models of cores HH12-946MC and HH12-948MC have been published previously (Ezat et al., 2019) and are based on calibrated planktic ¹⁴C dates (Supplemental Tables 2, 3) using the Marine13 dataset in Calib 7.04 (Reimer et al., 2013). Here we update the age models using the recently published Marine20 dataset (Heaton et al., 2020) in Calib 8.2 (Stuiver et al., 2020) (Tables S1, S2). We used ΔR (the regional difference from the average global marine reservoir correction) = 1000 years (i.e., reservoir age = ~1500 years)

for the glacial parts of the records and $\Delta R=0$ for the Holocene. Although changes in the 210 211 reservoir age in the Fram Strait during the last glacial period remain unconstrained, the assigned value of 1500 ¹⁴C years are consistent with reservoir age estimates from the southern 212 and central Norwegian Sea (Thornalley et al., 2015; Ezat et al., 2017) as well as with 213 modelling results (e.g., Butzin et al., 2020). The resulting sedimentation rates range from 0.6 214 to 9 cm/kyr in core HH12-946MC, with an average of 2 cm/kyr for both the Holocene and 215 216 glacial parts (Tables S1, S2). These relatively low sedimentation rates can potentially induce large smoothing effects from bioturbation. However, because the ¹⁴C dates of *N. pachyderma* 217 and *O. umbonatus* are in chronological for the glacial part of the record, and because ¹⁴C dates 218 219 of O. umbonatus are older than those of N. pachyderma (except for one sample at 28.25 cm, but the difference is less than the analytical uncertainty; Table 1), we conclude that the impact 220 of burrowing activity is minor. Furthermore, our study mainly target the relatively long-term 221 222 changes from late MIS 3 to LGM. According to the age model, the transition from late MIS3 to the LGM is characterized by a pronounced increase in δ^{18} O values measured in N. 223 pachyderma and O. umbonatus (Ezat et al., 2019; see also 'Results and Discussion' section), 224 which provides confidence in our age model at such timescales. 225

The original age models of sediment cores 1294, 1295 from the eastern Fram Strait 226 227 were based on uncorrected planktic radiocarbon dates (Hebbeln et al., 1994). We updated the original age models of cores 1294, 1295 by the same approach as for cores HH12-946MC and 228 HH12-948MC (Tables S3, S4). The age models of sediment cores PS1243 and MD99-2276 229 from the central Nordic Seas were based on correlation to the well-dated sediment cores from 230 the southern Norwegian Sea (see Rasmussen and Thomsen, 2004; Ezat et al., 2017) using the 231 planktic δ^{18} O (Thornalley et al., 2015). Unfortunately, because of missing data from the 232 deglaciation, the planktic δ^{18} O records from the Fram Strait cannot be correlated with the 233

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- 237
- 238 2.5 Carbon cycle modelling

Given the evidence of an abrupt +10 ppm jump in atmospheric CO₂ (e.g., Ahn and 239 Brook, 2014; Bereiter et al., 2015) and an enhanced influx of terrestrial carbon to the Nordic 240 Seas (Hebbeln et al., 1994) at the Heinrich Stadial 2/Interstadial 2 transition (~23.5 ka), we 241 are interested if and how the 'mobilization of terrestrial carbon' could be recorded in 242 243 'Northern Component Water'. For this, we used the glacial configuration of the CYCLOPS carbon cycle box model (Hain et al., 2010, 2014) as a point of comparison to our 244 observational records and in order to assess the effects of possible mobilization of terrestrial 245 carbon at this time. The baseline model scenario is spun-up to the reference LGM conditions, 246 including Southern Ocean changes that dominated simulated glacial CO₂ drawdown (Hain et 247 248 al., 2010; Sigman et al., 2010), as well as a 'Glacial North Atlantic Intermediate Water' (GNAIW; e.g., Lynch-Stieglitz et al., 2007) representation of the AMOC that is needed to 249 account for the gross pattern of LGM and deglacial carbon isotope changes (Sigman et al., 250 251 2003; Hain et al., 2014). The GNAIW configuration in CYCLOPS has the same 21.5 Sv strength (1 Sv = $10^6 \text{ m}^3/\text{s}$) as the default NADW configuration, but the lower limb of the 252 southward return flow is restricted to mid-depth as opposed to present-day NADW flowing in 253 the deep Atlantic basin. This initial steady state of the model is perturbed with a sequence of 254 Atlantic circulation changes that is designed to mimic the Heinrich stadial 2/interstadial 2 255 256 transition changes; starting with GNAIW, switching to a 'Heinrich Stadial' collapsed state of the AMOC for the duration of Heinrich stadial 2 (4000 years), followed by the default 257 NADW circulation to represent vigorous and deep overturning during interstadial 2 (1300 258

records from the southern Norwegian Sea. Finally, in this study, we define a Heinrich stadial

as the whole stadial period during which a Heinrich event occurred (e.g., Barker et al., 2011).

years), and finally a return to the initial GNAIW circulation. This treatment is equivalent to 259 260 the representation of the deglacial AMOC changes described in Hain et al. (2014). Assuming these simplistic 'ad hoc' changes adequately reflect Atlantic changes over the Heinrich stadial 261 2/interstadial 2 interval, the simulated biogeochemical changes in the "Northern Component 262 Water" model reservoir, which represents the subsurface high-latitude North Atlantic 263 including the Nordic Seas, serve as a comparison baseline for our observational data. To 264 265 simulate the spectrum of hypotheses of how terrestrial carbon may have been mobilized from the largely glaciated high-northern continents we consider both (1) oxidation on land with 266 CO₂ release directly to the atmosphere, and (2) erosion and transfer of terrestrial carbon to the 267 268 deep Nordic Seas/Arctic Ocean (as represented by the 'Northern Component Water'). In each of these two pathway cases, we simulate 100-year mobilization events incrementing the total 269 amount of carbon released from zero to 88 Pg C (1 Pg $C = 10^{15}$ gram carbon), similar to the 270 271 experiments of Köhler et al. (2011, 2014) where 125 Pg C from land was released to the atmosphere at 14.6 ka within a time window of 50 to 200 years. In this paper, we only discuss 272 273 the results from the carbon addition experiments. For a summary of the results from idealized circulation scenarios, see Text S1 (Supporting Information). 274

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276 **3 Results and Discussion**

277 **3.1 Interspecies benthic for aminiferal** δ^{18} O differences and implications

One significant obstacle to reconstruct deep ocean conditions in the northern and central Nordic Seas for the last glacial period is the ambiguous large δ^{18} O differences between the epifaunal benthic foraminiferal species *C. wuellerstorfi* and the shallow infaunal species *O. umbonatus* (Bauch et al., 2001; Thornalley et al., 2015; Ezat et al., 2019). Epifaunal δ^{18} O is ~1‰ lower than coeval infaunal δ^{18} O from sediments dated to the LGM (Figure 2). In

addition, the epifaunal δ^{18} O from the LGM sediments is up to 1‰ lower compared to the 283 Holocene, which is in contrast to all other ocean basins and the SE Norwegian Sea (Figure 2). 284 It has been suggested that this low epifaunal δ^{18} O from the LGM sediments indicates short-285 term, but regional ventilation events by brine formation (Bauch et al., 2001). However, it 286 remains possible that C. wuellerstorfi specimens were transported from younger/shallower 287 sediments or have a diagenetic carbonate coating with very low δ^{18} O. Our results show that 288 intensive acid leaching before δ^{18} O measurements (see Methods) did not change the δ^{18} O 289 values (Figure 2d). Thus, we can here eliminate the possibility that the low C. wuellerstorfi 290 δ^{18} O values are due inorganic carbonate coatings. 291

292 Radiocarbon dating of C. wuellerstorfi could reveal whether these C. wuellerstorfi specimens are autochthonous. The abundance of C. wuellerstorfi in the glacial sediments from 293 the central and northern Nordic Seas is very low (~<5% of the benthic foraminiferal 294 295 assemblages compared to 40% during the Holocene) with an average of 7 specimens per sample (~300 µg carbonate) (Bauch et al., 2001; Ezat et al., 2019). Our MICADAS ¹⁴C dates, 296 297 obtained on small samples of C. wuellerstorfi from the LGM sediments, range from 14,000 to 15,400 ¹⁴C years BP. Furthermore, they date ~ 4500 ¹⁴C years younger than the ages obtained 298 on coeval O. umbonatus and N. pachyderma (Figure 2c, Table 1). This clearly indicates that 299 these C. wuellerstorfi specimens are not of LGM age and that the specimens probably have 300 been reworked from younger sediments. These results confirm that caution is needed when 301 using low abundant 'proxy carriers' from low sedimentation rate areas (c.f., Peng and 302 Broecker, 1984). However we also note that down-mixing of Holocene specimens cannot 303 fully explain the young ages of *C. wuellerstorfi* because δ^{18} O values in *C. wuellerstorfi* from 304 the LGM sediments are even lower than those from the Holocene. The deglacial sediments in 305 our two records from the central Fram strait are barren of calcareous foraminifera, which have 306 been attributed to either post-depositional dissolution or unfavourable conditions (Zamelczyk 307

et al., 2014; Ezat et al., 2019). If post-depositional dissolution was the reason for the absence of foraminifera in the deglacial sections of our Fram Strait cores, an earlier downcore mixing of deglacial foraminifera with very low δ^{18} O (typical of HS1 in this region) into the LGM sediments may have occurred. However, we emphasize that the reworking process of *C*. *wuellerstorfi* into the LGM sediments remains not fully explained.

Regardless of the transport mechanism of younger C. wuellerstorfi into the LGM 313 sediments in the deep Fram Strait, it is clear that the geochemical signals in C. wuellerstorfi 314 (e.g., δ^{18} O and δ^{13} C) cannot be used in this context to infer hydrographic and ventilation 315 changes in the region during the LGM. Given the striking similarity of δ^{18} O measured in C. 316 317 wuellerstorfi from LGM sediments in the Fram Strait (Ezat et al., 2019; this study) and the central Nordic Seas (Bauch et al., 2001) (Figure 2), our results suggest that inferences of the 318 ocean circulation during the LGM based on δ^{18} O and δ^{13} C in *C. wuellerstorfi* from the central 319 320 Nordic Seas (e.g., Bauch et al., 2001; Thornalley et al., 2015; Knies et al., 2018; Mackensen and Schmiedl, 2019) also should be treated with caution. Radiocarbon dating of the C. 321 322 wuellerstorfi from the central Nordic Seas could settle the issue about reworking. We thus base our reconstructions of ocean circulation and carbonate chemistry in the northern Nordic 323 Seas on shell geochemistry of O. umbonatus. 324

It is also notable that δ^{18} O of *C. wuellerstorfi* from the deep southern Norwegian Seas do not show the same low values for the LGM and that δ^{18} O values of both *C. wuellerstorfi* and *O. umbonatus* are consistently high (Veum et al., 1992) (Figure 2b) similar to *O. umbonatus* δ^{18} O from the central and northern Nordic Seas (Figure 2). This provides confidence in using *C. wuellerstorfi* geochemistry from LGM sediments from the southern Norwegian Sea records, in contrast to the northern (and likely central) Nordic Seas deep records.

332

3.2 Deep temperature and $\delta^{18}O$ evolution

334	Our BWT reconstructions from ~2600 m water depth from the central Fram Strait
335	based on Mg/Ca in O. umbonatus varies between -0.2 and 1° C during MIS 3 (~40–26 ka)
336	with an average of 0.6±0.5 °C (Figure 3b). This is ~1.5±0.5 °C warmer than the modern and
337	average Holocene temperatures, which are ~-0.8 $^{\circ}$ C (Figure 3b). These BWT changes are in
338	agreement with those from the central Arctic Ocean at the same water depth based on Mg/Ca
339	in ostracods (Cronin et al., 2012) (Figure 3). Our BWT record further shows an increasing
340	trend along the transition from MIS 3 to the LGM with the highest BWT (~1.2 $^{\circ}$ C) recorded
341	at ~26–24 ka (Figure 3b). At ~23.5 ka, close to the timing of interstadial 2, the BWT
342	decreased to ~0 °C and thereafter it fluctuated between ~0 and 0.5 °C at 22–20 ka (Figure 3b).
343	Deep benthic foraminiferal δ^{18} O records from the Nordic Seas (excluding epifaunal
344	records from the central and northern Nordic Seas; see section 3.1) reveal a glacial-
345	interglacial change of only 0.7‰ (Figure 2; 3a), which is smaller than records from other
346	ocean basins. Our deep-water δ^{18} O estimates based on δ^{18} O and temperature correction using
347	Mg/Ca in O. umbonatus shows a 1‰ glacial-interglacial change, which is similar to the
348	estimated global average ocean δ^{18} O change (Figure 3d). Seawater δ^{18} O in the deep Nordic
349	Seas increased from 0.7‰ during late MIS 3 to 1‰ during the LGM (Figure 3d). This pattern
350	mimics the global ocean changes in deep-water δ^{18} O (Figure 3d). The consistent amplitude
351	and evolution of seawater $\delta^{18}O$ variations in the Nordic Seas (and possibly the central Arctic
352	Ocean) and the mean global ocean change are in agreement with persistent exchange of deep
353	water with other ocean basins (though likely with different modes and rates) and argue against
354	extreme isolation of the Arctic Ocean. This is consistent with the new ¹⁴ C-based evidence
355	from the Fram Strait (Ezat et al., 2019). Based on the previously inferred 'stagnation' of the
356	deep Arctic Ocean and the Nordic Seas (ventilation ages up to 10,000 years), geothermal
357	heating was suggested as a significant contributor to the glacial deep ocean warming in the

region (Thornalley et al., 2015). In contrast, recent ¹⁴C evidence from the Fram Strait (Ezat et 358 al., 2019) and our deep-water δ^{18} O evolution (Figure 3d) suggest smaller ventilation ages and 359 more exchange with other ocean basins, which would have limited the contribution of 360 geothermal heating to the deep-water warming. We propose that these observations can be 361 reconciled if the glacial warming observed in the Nordic Seas is attributed instead to 362 thickening and/or deepening of the Atlantic intermediate water, allowing it to influence 363 364 deeper areas than at present. This also suggests a significant reduction and/or shoaling of cold deep water formation by open ocean convection. Planktic δ^{18} O records from the Nordic Seas 365 display an increase from the southern Norwegian Sea (64° 25' N), central Nordic Seas (69°37' 366 367 N) to the Fram Strait (78°53' N) during the Holocene (Figure 3e), which likely reflects the cooling of the Atlantic water as it flows northward. This gradient is largely diminished during 368 the glacial period in particular at 22–19 ka (Figure 3) probably suggesting overall colder sea 369 370 surface temperatures in the region and a dominant subsurface flow (and thus negligible heat loss) instead of a surface flow subject to air-sea exchange. 371

372

373 **3.3** Carbonate chemistry and δ^{13} C records

Epifaunal benthic foraminiferal geochemistry is typically used to reconstruct deep 374 ocean chemistry changes (e.g., Duplessy et al., 1988; Yu and Elderfield, 2007; Mackensen 375 and Schmiedl, 2019). Given the reworking problem that we identify with the available 376 epifaunal species (C. wullerstorfi) from the glacial sediments in our study area (see section 377 3.1), we instead use δ^{13} C and B/Ca measured in the shallow infaunal benthic foraminiferal 378 species O. umbonatus for the reconstruction of carbonate chemistry changes. In the North 379 380 Atlantic Ocean, O. umbonatus lives in the upper 1 cm of the sediments (Corliss, 1985), which may indicate influence from pore water chemistry. However, the overall low productivity and 381 supply of carbon to the deep sea as well as low sediment accumulation rates in the Fram Strait 382

likely suggest insignificant decoupling between bottom water and shallow (< 2 cm deep in the 383 sediments) pore water chemistry. In addition, the central Arctic Ocean equivalent of O. 384 umbonatus (O. tener) is considered to have an epifaunal microhabitat (Murray, 2006; 385 Barrientos et al. 2018). In general, B/Ca in benthic foraminifera, including infaunal species, 386 show a positive correlation with the deep-water carbonate saturation ($\Delta[CO_3^{2-}]$), providing a 387 method for the reconstruction of deep-ocean carbonate ion concentration ($[CO_3^{2-}]$) (Yu and 388 Elderfield, 2007; Yu et al., 2008; Rae et al., 2011). Carbonate ion concentration broadly 389 records the difference or ratio between the two master variables of the carbonate system, 390 Alkalinity and Dissolved Inorganic Carbon (DIC), allowing $[CO_3^{2-}]$ reconstructions to inform 391 on past changes in carbon storage in the ocean (Broecker and Peng, 1982; Yu et al., 2008, 392 2016). Thus, we use O. umbonatus B/Ca and δ^{13} C as qualitative indicators of [CO₃²⁻] and 393 nutrients, respectively. Given the shallow infaunal habitat of O. umbonatus, this should 394 395 nonetheless be taken with caution.

The B/Ca in O. umbonatus increased from ~52 µmol/mol during late MIS 3 to ~ 62 396 µmol/mol during the LGM and subsequently decreased to ~43 µmol/mol during the Holocene 397 (Figure 4b). This suggests higher glacial $[CO_3^{2-}]$, with highest values during the LGM as seen 398 in many studies for mid-depth records from the North Atlantic Ocean (Yu et al., 2008, 2010, 399 2020). The δ^{13} C in the infaunal species O. umbonatus is on average ~1% lower during the last 400 glacial (~40–19 ka) compared to the Holocene (Figure 4e). The glacial-interglacial changes in 401 δ^{13} C and B/Ca of *O*. *umbonatus* are generally consistent with δ^{13} C and B/Ca in *C*. 402 wuellerstorfi from the subpolar North Atlantic Ocean at water depths between 2000 and 3000 403 m (Yu et al., 2008) (Figure 4). This supports the inference of Yu et al. (2008) that the Nordic 404 Seas have contributed to the lower glacial North Atlantic intermediate water (LGNAIW) 405 water mass. Idealized numerical modelling shows that under extensive sea-ice cover and 406 subsurface inflow of Atlantic water into the Nordic Seas, Atlantic water recirculates as a 407

western boundary current out of the Nordic seas (Jensen et al., 2018). The Denmark Strait
may therefore have provided an overflow pathway of the subsurface Atlantic water to the
North Atlantic Ocean.

411

412 **3.4** Changes in the terrestrial carbon input and carbon cycle modelling

During the later part of Heinrich stadial 2 and the transition to interstadial 2, δ^{13} C of 413 414 organic material and C/N from three nearby records show a significant and relatively abrupt decrease suggesting an increase in the input of terrestrial carbon (Hebbeln et al., 1994; see 415 Figure 4d). This is associated with a decrease in %CaCO₃ and an increase in Ice Rafted 416 Debris (IRD) (Hebbeln et al., 1994). Although a limited number of records of $\delta^{13}C$ measured 417 418 in organic material are available from the region (Hebbeln et al., 1994), the decrease in %CaCO₃ and increase in IRD close to the end of Heinrich stadial 2 are widely recorded from 419 420 the northern Nordic Seas (e.g., Jessen et al., 2010; Zamelczyk et al., 2014) and suggest that the increase in the input of terrestrial carbon was probably a geographically wide-spread 421 422 feature. This possibly hints to a mobilization event of Arctic permafrost and/or carbon release from ice sheets. Arctic permafrost, which can be divided into terrestrial (~1,000 Pg C), ice 423 complex (~400 Pg C) and subsea (~1,400 Pg C) permafrost, is a major active component of 424 425 the global carbon cycle (e.g., Tarnocai et al., 2009; Vonk et al., 2012). Also, ice sheets represent an interactive player in the global carbon cycle (e.g., Wadham et al., 2019). Within 426 the limitation of our chronology due to uncertainty about past changes in R (section 2.4), this 427 event may have occurred sometime between 24.5 ka and 23 ka. We do not have constraints on 428 the source, magnitude or transport pathways of this possible 'terrestrial carbon' mobilization 429 event. However, some of this mobilized carbon likely entered the atmosphere directly, while a 430 smaller part entered the Arctic Ocean system. 431

Although ocean biological and physical processes have likely been leading 432 433 mechanisms for the glacial/interglacial and millennial-scale variations in atmospheric pCO₂, and may contribute to centennial-scale changes (e.g., Ezat et al., 2017; Rae et al., 2018), 434 oxidation of terrestrial organic carbon may also have played a role in abrupt centennial-scale 435 events (e.g., Köhler et al., 2014; Bauska et al., 2016; Meyer et al., 2009). Given the evidence 436 of an abrupt increase of 10 ppm in atmospheric CO_2 (e.g., Bereiter et al., 2015) and an 437 438 enhanced influx of terrestrial carbon to the Nordic Seas (Hebbeln et al., 1994) at the Heinrich stadial 2/interstadial 2 transition, we are interested in finding out if and how the mobilization 439 of terrestrial carbon in the northern high latitudes is recorded in the deep 'Northern 440 441 Component Water'. To test the relative sensitivity of atmospheric pCO_2 and 'Northern Component Water' carbonate chemistry, we superimpose two sets of carbon mobilization 442 scenarios onto the CYCLOPS model idealized circulation scenario, with one set adding 443 444 respired carbon directly to the atmosphere (Figure 5, left panel) or alternatively directly to 'Northern Component Water' (Figure 5, right panel). In both cases, carbon is added for 100 445 446 years with cumulative mobilization ranging 0-88 Pg C for individual realizations. If ~45 Pg C are directly released to the atmosphere, simulated atmospheric pCO₂ experiences an abrupt 447 and short-lived 10 ppm increase, similar to reconstructions of interstadial 2 (Figure 4g). In all 448 449 cases where the carbon is added to the atmosphere the global surface ocean immediately absorbs about half of the added carbon and the associated ocean acidification and Suess effect 450 are dispersed and not concentrated near the boreal source and 'Northern Component Water'. 451 The carbonate chemistry and δ^{13} C of DIC remain principally controlled by large-scale changes 452 in ocean circulation, and not by the carbon addition. In contrast, if the carbon is released to 453 the 'Northern Component Water', the model predicts abrupt and severe decline of $[CO_3^{2-}]$ and 454 δ^{13} C of DIC in the 'Northern Component Water' because the carbon becomes concentrated in 455 the interior Atlantic rather than dispersed globally. For that reason carbon release to the 456

'Northern Component Water' is unlikely to result in a transient atmospheric pCO₂ spike. It is 457 expected that the anomalies in $[CO_3^{2-}]$ and $\delta^{13}C$ of DIC in the 'Northern Component Water' 458 depend on the rate of carbon release because with NADW circulation during interstadial 2, the 459 water residence time in the 'Northern Component Water' is short and advection and 460 dissipation of the added carbon to the global deep ocean is facilitated by the vigorous AMOC. 461 From these idealized carbon mobilization scenarios, we conclude that carbonate chemistry 462 and δ^{13} C from the Nordic Seas should be relatively sensitive to respiration of terrestrial 463 organic matter at depth, where carbon remains relatively concentrated until ocean circulation 464 dissipates the added carbon. This pathway of carbon mobilization has only a minimal 465 466 transient effect on atmospheric pCO₂ because the carbon is initially concentrated and then dissipated within the ocean interior. The alternative pathway, direct release to the atmosphere, 467 causes an immediate CO₂ transient that is moderated by the global surface ocean Revelle 468 469 buffer factor and its total buffer capacity, with the global dispersion of the added carbon precluding the generation of significant regionally focused carbonate chemistry anomalies. 470 We only observe modest changes in our B/Ca and δ^{13} C records at the Heinrich stadial 471 2/interstadial 2 transition (Figure 4b,c). Thus, if the terrestrial organic matter influx at 472 Heinrich stadial 2/interstadial 2 (Hebbeln et al., 1994; Figure 4) recorded a mobilization event 473 474 of subglacial or permafrost carbon, the total carbon release was likely low or slow, or released directly to the atmosphere. 475

476

477 **4.** Conclusions

478 In this study, we reconstructed deep ocean hydrographic changes in the northern 479 Nordic Seas during the late glacial (35–19 ka). Furthermore, we investigated the ambiguously 480 large differences of δ^{18} O between two benthic foraminiferal species *Oridorsalis umbonatus* (shallow infaunal) and *Cibicidoides wuellerstorfi* (epifaunal) that have been recorded from sediments dating from the last glacial maximum (LGM) in the central and northern Nordic Seas. Using species-specific foraminiferal ¹⁴C dating, we showed that the specimens of *C*. *wuellerstorfi* have been reworked. We therefore based our reconstructions on the most persistent and abundant benthic species *O. umbonatus*. We propose that the reconstruction of glacial ocean circulation in low-resolution deep-sea sediments from this region based on δ^{18} O and δ^{13} C measured in *C. wuellerstorfi* should be treated with caution.

Our bottom water temperatures (BWT) from Mg/Ca show, in agreement with previous 488 studies from the Arctic Ocean and central Nordic Seas, that the deep northern Nordic Seas 489 490 were on average 1.5 °C warmer during the late glacial compared to the Holocene. Furthermore, local (and potentially regional) seawater δ^{18} O was 0.7 and 1‰ higher during 491 late MIS 3 and the LGM, respectively, relative to the Holocene. The similarity in the 492 magnitude and evolution between our seawater δ^{18} O record and global average ocean δ^{18} O 493 supports the recent ¹⁴C evidence of persistent exchange between the Arctic Ocean and the 494 Nordic Seas and other ocean basins during the late glacial. In addition, the similar signatures 495 of elevated $[CO_3^{2-}]$ and lower $\delta^{13}C$ during the late glacial in the subpolar North Atlantic 496 between 2000–3000 m water depth and the northern Nordic Seas is a strong indication that the 497 498 Nordic Seas may have contributed significantly to the lower glacial North Atlantic intermediate water. Altogether, our results indicate that during the late glacial the Atlantic 499 water thickened and deepened to at least 2600 m water depth and flowed as a subsurface 500 intermediate layer in the Nordic Seas and the Arctic Ocean with active overflows to the North 501 Atlantic Ocean likely via the Denmark Strait. Open ocean convection in the Nordic seas 502 503 probably ceased or became intermittent.

Finally, evidence from previous studies suggests enhanced terrestrial carbon flux to
the northern Nordic Seas at the Heinrich stadial/interstadial 2 transition (~23.5 ka).

506 Comparing our $[CO_3^{2-}]$ and $\delta^{13}C$ records with carbon cycle model results, we conclude that 507 either the total subglacial or permafrost carbon release either was low or slow, or released 508 directly to the atmosphere.

509

510 Figure & Table Captions

511	Figure 1. Ma	p showing major	ocean currents in t	the Nordic	Seas and Arcti	c Ocean and
0	I IGAL C IT ITIG		occum currents m	the rorate		e occan an

512 the location of sediment cores. Yellow dots refer to the location of sediment cores from this

study (HH12-946MC and HH12-948MC, Ezat et al., 2019) and puslished records discussed in

this study (HLY0503-18TC, e.g., Cronin et al., 2012; 1294-4 and 1295-5, Hebbeln et al.,

515 1994; PS1243 and MD99-2276, e.g., Thornalley et al., 2015; HM52-43, Veum et al., 1992).

516 Atlantic water inflow is indicated by solid red arrows (surface inflow) and dashed red arrows

517 (submerged inflow). Freshwater circulation are shown by white arrows. Nordic Seas

518 overflows are indicated by black arrows.

519

- 520 Figure 2. Benthic δ^{18} O records from the Nordic Seas. Southern Norwegian Sea (red
- 521 curve; Veum et al., 1992), central Nordic Seas (blue curves; Bauch et al., 2001) and

522 Fram Strait records HH12-946MC (purple curves), HH12-948MC (black curves) (Ezat

et al., 2019; this study). Numbers in plot (c) are un-calibrated 14 C dates obtained on C.

wuellerstorfi (italic, underlined font) and *O. umbonatus* (normal font). Triangles in (d) shows

525 δ^{18} O in *C. wuellerstorfi* after extensive acid leaching. Grey and orange highlighting refer to

the Last Glacial Maximum (LGM) and the Holocene, respectively.

528	Figure 3. Deep Fram Strait, central Nordic Seas and Arctic Ocean temperature and
529	δ^{18} O during the last glacial period. (a) δ^{18} O in <i>O. umbonatus</i> from Fram Strait records
530	HH12-946MC (circles; Ezat et al., 2019; this study), HH12-948MC (triangles; Ezat et al.,
531	2019) and Central Nordic Seas core MD99-2276 (squares; Thornalley et al., 2015). (b) Deep
532	ocean temperature based on Mg/Ca in O. umbonatus from cores core MD99-2276 (squares;
533	Thornalley et al., 2015), HH12-946MC (circles) and HH12-948MC (triangles). Error bars
534	represent the combined 'analytical' (based on 2 s.d. of replicate measurements of in-house
535	standard) and 'calibration' uncertainties. (c) Deep ocean temperature based on Mg/Ca in
536	ostracods from the central Arctic Ocean, core HLY0503-18TC (Cronin et al., 2012). (d)
537	Seawater δ^{18} O based on Mg/Ca and δ^{18} O in <i>O. umbonatus</i> from cores HH12-946MC (circles,
538	this study), HH12-948MC (triangles, this study) and core MD99-2276 (squares, Thornalley et
539	al., 2015). Dashed line represents average global ocean δ^{18} O following Spratt and Lisiecki
540	(2016). (e) Planktic δ^{18} O records from the southern Norwegian Sea (red, core HM52-43,
541	Veum et al., 1992), central Nordic Seas (dark green, core PS1243, Bauch et al., 2001; green,
542	core MD99-2276, Thornalley et al., 2015), and Fram Strait (blue, HH12-946MC; light blue,
543	HH12-948MC; Ezat et al., 2019 and this study). Red arrows above the x-axis refer to the
544	location of the calibrated planktic ¹⁴ C dates that have been used in the age model of core
545	HH12-946MC.

546

Figure 4. Evolution of deep ocean chemistry in Fram Strait and subpolar North Atlantic Ocean. (a) Deep ocean temperature from Fram Strait records HH12-946MC (circles), HH12948MC (triangles). (b) B/Ca in *O. umbonatus* from cores HH12-946MC (circles) and HH12-

550 948MC (triangles). Errors bars are based on 2 s.e. of replicate measurements of in-house

standard (see Methods). (c) δ^{13} C in *O. umbonatus* from cores HH12-946MC (circles) and

552 HH12-948MC (triangles). (d) δ^{13} C in organic matter from the Fram Strait records 1295-5

(dashed line and open circles) and 1294-4 (solid line and filled circles) (Hebbeln et al., 1994). (e) Carbonate ion concentration based on B/Ca measured in from core NEAP 8K (Yu et al., 2008). (f) δ^{13} C measured in *C. wuellerstorfi* from core NEAP 8K (Yu et al., 2008). (g) Atmospheric pCO₂ from Antarctic ice cores (Bereiter et al., 2015 and references therein).

557

Figure 5. Modeled changes in deep 'Northern Component Water' from idealized
circulation changes (blue line) and carbon addition (0–88 Pg C; 1 Pg C= 10¹⁵ gram
carbon) scenarios (black lines). The model results from the 'direct carbon addition to the
atmosphere' and the 'carbon addition to the deep North Atlantic' experiments are shown
(black lines) at the left and right panels, respectively. Abbreviations: GNAIW, Glacial North
Atlantic Intermediate Water; HS, Heinrich Stadial; NADW, North Atlantic Deep Water.

564

Table 1. Radiocarbon dates on *Neogloboquadrina pachyderma*, *Cibicidoides wuellerstorfi*and *Oridorsalis umbonatus* from core HH12-946MC. Grey-highlighting indicates samples
measured by 'gas source-based techniques' (this study; see Methods), while other samples
were measured by 'graphitization-based' methodology (Ezat et al., 2019).

569

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580 Data availability statement

- 581 The new data presented in this article are available in the 'Supporting Information' and
- will be available at Zenodo database upon acceptance of this manuscript, DOI:
- 583 https://doi.org/10.5281/zenodo.4436094.
- 584

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Figure 1.



0°

Figure 2.



Figure 3.



Figure 4.



Figure 5.



1.0

0.5

Sediment	Calendar	¹⁴ C Age (years BP)			
Depth (cm)	Age (ka)	N. pachyderma	C. wuellerstorfi	O. umbonatus	
25.25	19.7	18075 ±90		18580 ±88	
25.75	20.1	18399 ±75		18511 ±82	
27.25	20.6	18765 ±73	14460 ±190	18787 ±95	
28.25	21.2	19335 ±77	14206 ±205	19232 ± 148	
29.25	22.6	19833 ±284	15175 ±169		
29.75	23.3		15372 ±205		
30.75	24.7	22372 ±108			
32.75	25.5	23076 ±131		25327 ±161	
33.75	27.1	24741 ±134		26118 ± 170	
35.75	30.7	28333 ±225		29715 ±340	