

**Title: A Diminished North Atlantic Nutrient Stream during Younger Dryas  
Climate Reversal**

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**Abstract:** The high rate of biological productivity in the North Atlantic is stimulated by the advective supply of nutrients into the region via the Gulf Stream (nutrient stream). It has been proposed that the projected future decline in the Atlantic Meridional Overturning Circulation (AMOC) will cause a reduction in nutrient supply and resulting productivity. Here we examine how the nutrient stream changed over the Younger Dryas climate reversal that marked the transition out of the last ice age. Gulf Stream nutrient content decreased, and oxygen content increased at the Florida Straits during this time of weakened AMOC. The decreased nutrient stream was accompanied by a reduction in biological productivity at higher latitudes in the North Atlantic, supporting the link postulated in theoretical and modelling studies.

**One-Sentence Summary:** Paleooceanographic data supports hypothesis linking delivery of nutrients by Gulf Stream to high latitude productivity.

## **Main Text:**

The Gulf Stream and North Atlantic Drift serve as a northward flowing conduit of nutrient-rich intermediate waters (1, 2). In the subpolar North Atlantic the density surfaces bearing these nutrients shoal and the nutrients are incorporated into the deep winter mixed layer (3). During the spring bloom these nutrients drive primary productivity in the sunlit surface ocean. The high nutrient waters of the Gulf Stream have been termed the “nutrient stream” and this advective pathway is thought to be the dominant source of nutrients supporting phytoplankton productivity in the subpolar North Atlantic (4, 5). The Gulf Stream originates in the Florida Straits and as it moves northward along the western boundary of the subtropical gyre, both the mass and nutrient transport of the current is augmented by recirculating water within the North Atlantic basin. However, the ultimate source of the nutrients to the incipient Gulf Stream in the Florida Straits is the import of high nutrient intermediate-depth waters from the tropics, which are replenished by the high-nutrient Southern Hemisphere intermediate and mode waters as part of the upper (northward) limb of the Atlantic Meridional Overturning Circulation (AMOC) (2, 6). In addition to supplying the nutrient stream, the Southern Hemisphere intermediate and mode waters supply the nutrients that drive much of the primary production at low and mid latitudes (7, 8).

The AMOC is projected to weaken over the coming century with the increase in anthropogenic greenhouse gasses (9). Consequently, the nutrient stream is projected to weaken, leading to basin scale changes in biogeochemistry and decreased primary production. Declines in primary production would impact the important North Atlantic fisheries, and may also impact the uptake of anthropogenic CO<sub>2</sub>. While a future decline in North Atlantic export production under future warming scenarios has long been projected (10), it has only more recently been recognized that the decline in the nutrient stream is the dominant mechanism driving this change (11). Whitt (5) found that, when driven with the high emissions RCP8.5 scenario, the Community Earth System Model (CESM) shows a 35% decline in Gulf Stream nitrate transport at 30°N in 2080 in response to a 39% decline in AMOC and find that the decreased nutrient stream is the primary driver of the associated decline in North Atlantic export productivity. Tagklis et al. (12) showed that the weakening of AMOC over the next century in the multi-model mean of seven CMIP5 models results in reduced upper-ocean phosphate concentration, upper-ocean apparent oxygen utilization, and primary productivity in the North Atlantic, and they attribute these changes to the decreased nutrient stream. Using biogeochemical data from an ice core, Osman et al. (13) reconstructed a decline in North Atlantic primary productivity over the industrial era, which they attributed to a weakening AMOC. There are multiple lines of evidence that the AMOC weakened during the Younger Dryas cold interval that punctuated the transition out of the last ice age (14). In this paper we assess the link between the AMOC and nutrient stream for the Younger Dryas event through the reconstruction of nutrient and oxygen concentrations in the deepest waters of the Florida Straits and explore the consequences for North Atlantic productivity. While the details of the background climate state and time scale of change differ from the present day, this past climate event provides an opportunity to test the mechanisms that have been identified in the climate models.

## **Reconstruction of Seawater Biogeochemistry in the Florida Straits**

Sediment core KNR166-2-26JPC was recovered from 24°19.61'N, 83°15.14'W in the Florida Straits at 546 m water depth. Due to the high sedimentation rates, this core provides exceptional time resolution over the Younger Dryas climate reversal. The coring site along the Florida margin is currently bathed by Antarctic Intermediate Water (AAIW) (potential density = 27.3 kg m<sup>-3</sup>), which at this location is characterized by high nutrient and low oxygen concentrations and supplies new nutrients to the North Atlantic (2, 6) (**Fig. 1**). In this paper we reconstruct seawater dissolved oxygen concentration at this location using carbon isotope measurements in paired epifaunal and deep infaunal benthic foraminifera, following an approach pioneered by McCorkle and Emerson (15, 16). *Planulina ariminensis* occupies a habitat on top of the sediments and records seawater  $\delta^{13}\text{C}$  values, and *Globobulimina* has a habitat deep within the sediments where the dissolved carbon has much lower  $\delta^{13}\text{C}$  values and oxygen is reduced to near zero values from the respiration of organic matter within the sediments. The difference between the  $\delta^{13}\text{C}$  values in the tests of these two benthic foraminifera is used to quantitatively reconstruct past oxygen concentrations (17). We also present abundance data for three thermocline dwelling species of planktonic foraminifera which are currently abundant in areas with subsurface low oxygen layers, *Globorotalia menardii*, *Globorotalia tumida*, and *Pullentina obliquiloculata* (18). At this location, we have previously published Cd/Ca measurements in benthic foraminifera which were used to reconstruct changes in seawater Cd<sub>w</sub>, which mimics the distribution of the major nutrient PO<sub>4</sub> in the ocean (19). We have also published benthic foraminiferal Mg/Li (20) to reconstruct temperature at this site over the deglaciation. We also use a previously published age model for this core (21). We estimate past PO<sub>4</sub> concentrations, using the Cd<sub>w</sub> reconstructions, and Apparent Oxygen Utilization (AOU), a measure of oxygen consumption in the subsurface ocean that is determined from the difference between in-situ and saturation oxygen concentration, using the quantitative temperature and oxygen reconstructions. We then use the PO<sub>4</sub> and AOU to estimate separately changes in the PO<sub>4</sub> concentration inherited from the source regions (preformed PO<sub>4</sub>, P<sub>pre</sub>) and the PO<sub>4</sub> that has been contributed to the subsurface from the remineralization of organic matter. Because intermediate waters leave the surface slightly undersaturated in oxygen, AOU overestimates true subsurface oxygen utilization, and P<sub>pre</sub> can be underestimated by up to 0.2  $\mu\text{M}$ , depending on the climate state. Detailed methods can be found in the Supplementary Information.

### **Weakened Florida Straits Nutrient Stream during the Younger Dryas**

While the intermediate waters that supply the nutrient stream have high PO<sub>4</sub> and oxygen concentrations (high P<sub>pre</sub>, low AOU) when they leave the surface in the Southern Hemisphere, these properties change as the waters transit northward through the tropics. Export productivity from overlying surface waters adds PO<sub>4</sub> and consumes O<sub>2</sub> via the oxidation and remineralization of sinking organic matter. Since the productivity is itself fed by a continuous supply of preformed nutrients, the accumulation of remineralized PO<sub>4</sub> and depletion of oxygen both depend on the northward flow of nutrient-charged intermediate waters (22). As the AMOC weakened during the Younger Dryas (14), this contribution of nutrient-rich, low oxygen intermediate water from the south would have diminished, replaced by lower PO<sub>4</sub> and higher oxygen intermediate waters formed locally in the North Atlantic. We therefore would expect an increase in the oxygen concentrations in the deep Florida Straits. Indeed, we reconstruct high oxygen values during the Younger Dryas from the dual carbon isotope approach, and increased oxygenation of the waters in the Florida Straits is also supported by the disappearance of the planktonic foraminifera *G. menardii*, *G. tumida*, and *P. obliquiloculata* (**Fig. 2**). The small

warming of the waters transiting the site of about 2°C during the Younger Dryas interval (20) would imply a decrease in oxygen concentration at saturation with the atmosphere, and the decrease in AOU is consequently slightly larger than the increase in oxygen. This is, again, what is expected if the oxygen increase is driven by the diminished contribution of the high AOU intermediate waters from the tropics.

A more direct measure of the nutrient stream comes from estimates of the concentration of the major nutrient PO<sub>4</sub> in the deep Florida Straits (**Fig. 3**). This estimate is based on the Cd/Ca in *H. elegans*, a different species of benthic foraminifera than was used to reconstruct oxygen via the carbon isotope measurements. The PO<sub>4</sub> at this site reflects contributions of intermediate waters formed in both hemispheres, including the regenerated PO<sub>4</sub> from the remineralization of organic matter in the tropics. We find a strong Younger Dryas decrease in PO<sub>4</sub> and, given that the intermediate water Northern and Southern end member values did not change by much over this time period (**Fig. S2b**), the lower PO<sub>4</sub> in the Florida Straits is consistent with the weakened AMOC. To probe this further, we use the AOU to separate the PO<sub>4</sub> concentration into its preformed (P<sub>pre</sub>) and regenerated (PO<sub>4</sub> – P<sub>pre</sub>) components. The PO<sub>4</sub> reduction primarily reflects a small decrease in P<sub>pre</sub>, and a much larger decrease in the regenerated PO<sub>4</sub>, as inferred from the large AOU decrease (**Fig. 3**). Today the intermediate waters formed in the Southern Hemisphere have a higher P<sub>pre</sub> than those that form in the North Atlantic, so a lower P<sub>pre</sub> is expected during a time of weakened AMOC when a larger proportion of Northern Hemisphere sourced intermediate waters would have bathed the site. Similarly, with a weakened AMOC we expect a lesser contribution of intermediate waters that have transited through the tropics, where they gain a high regenerated PO<sub>4</sub> content. There may have also been a positive feedback whereby the reduced northward transport of nutrients led to less productivity and remineralization in the tropics. The reduction of both pre-formed and regenerated PO<sub>4</sub> throughout the intermediate-depth North Atlantic, coupled with a decreased contribution of intermediate waters from the Southern Hemisphere, is also inferred for the Last Glacial Maximum, based on an inversion of the spatially more extensive data-set for that time period (23).

## **Biogeochemical Consequences of a Weakened vs. Collapsed AMOC**

While it has been established that the surface branch of the AMOC replenishes the modern nutrient stream, and that a future century-scale weakening of the AMOC will result in a weaker nutrient stream and lower North Atlantic productivity, it is not immediately apparent that these concepts and expectations translate directly to the Younger Dryas, a millennial scale event that occurred on the deglaciation. Schmittner (24) simulated a millennial scale AMOC collapse and recovery using freshwater forcing in an intermediate complexity model. While a productivity decline is observed, the mechanism behind this decline under a collapsed AMOC differs from the simulations for a future weakening of the AMOC. The decline in productivity in this case is attributed to strong near-surface density stratification, and the intermediate water nutrient content in the upper 1 km of the North Atlantic is higher over the duration of the shutdown rather than lower as in the future scenarios. Another set of equilibrium experiments under Last Glacial Maximum boundary conditions in a different intermediate complexity model confirmed this behavior for a simulation where the large scale AMOC is completely collapsed due to strong freshwater forcing (25). However, when AMOC is weakened, but not shut down completely, the PO<sub>4</sub> in the upper 2 km of the North Atlantic is lower than the control state, consistent with the pattern seen for the future transient (12) and in our reconstruction for the Younger Dryas. The transient response of a long paleoclimate simulation using CESM1 (26) can be used to explore

the relationship between AMOC, biogeochemical properties in the upper North Atlantic, and productivity. In this simulation, as the AMOC and nutrient stream weaken, the nutrient content of the upper North Atlantic and primary productivity both decline (**Fig. S2a**). However, when the AMOC is almost absent, a state with low productivity and *high* open ocean North Atlantic intermediate water nutrient content is seen, as in the AMOC collapse scenarios of the intermediate complexity models. The intermediate water  $Cd_w$  records in the Florida Straits and in the open ocean North Atlantic converge at low values during the Younger Dryas, consistent with what the model shows for a very weak AMOC and diminished nutrient stream, but not a fresh-water induced collapse (**Fig. S2b**).

## Impact of Weakened Younger Dryas Nutrient Stream on North Atlantic Productivity

Finally, we examine paleoclimate records that have been interpreted as indicators of export productivity. These include records of fluxes of organic components to the sediments as well as microfossil assemblages associated with different productivity regimes. Most of the records that resolve the Younger Dryas climate oscillation show decreased productivity relative to the millennia immediately before and after the event (**Fig. 4, Table S1**). While changes in oceanic conditions influencing productivity over the Younger Dryas may have influenced each site differently, the overall reduction of productivity in the region is consistent with the diminished supply of nutrients. This provides observational support of the causal chain inferred from global climate model simulations, that AMOC weakening leads to reduced advective supply of nutrients and declining North Atlantic primary productivity. We also highlight that while North Atlantic primary productivity is expected to decline in the case of either a weakening or collapse in the AMOC, the sign of biogeochemical changes in the intermediate waters of the open ocean North Atlantic will be different. The data for the Younger Dryas are consistent with a weakened rather than a collapsed AMOC. While most climate models suggest a weakening of the AMOC over the coming century, a collapse has not been ruled out (27, 28), and it worth noting the non-linear response of some biogeochemical impacts between these two scenarios.

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References 32-63 are only called out in the Supplementary Materials.

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**Supplementary Materials**

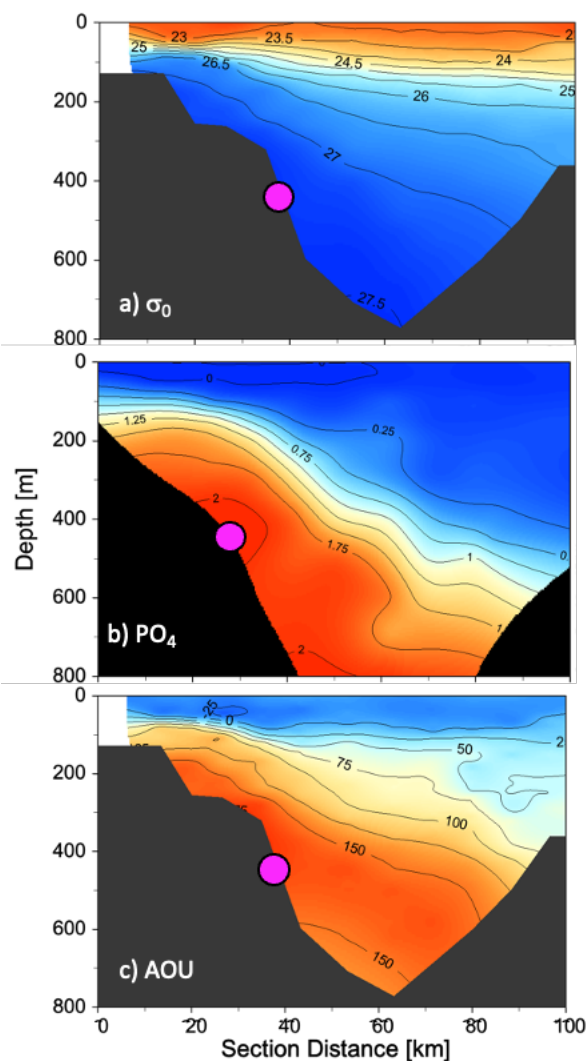
Materials and Methods

Figs. S1 to S4

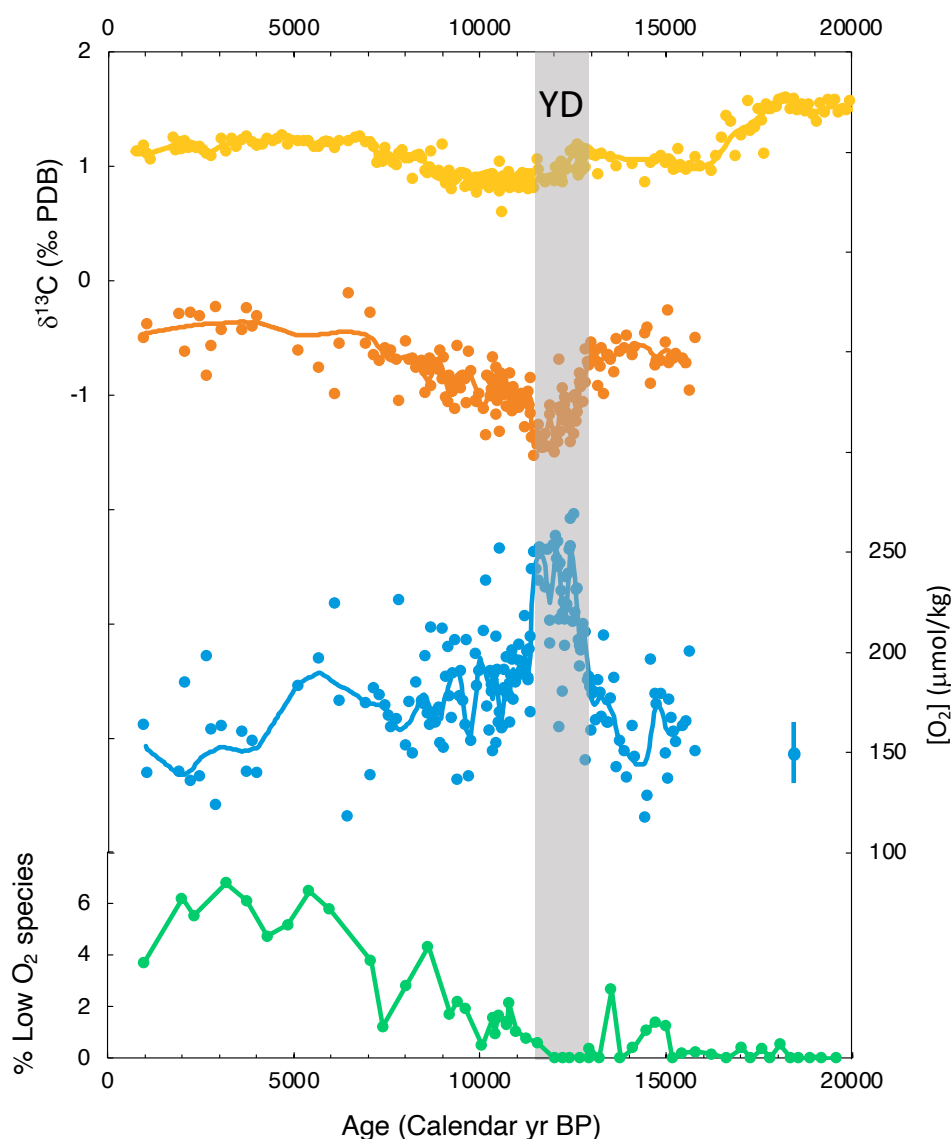
Tables S1 to S2

References (28–50)

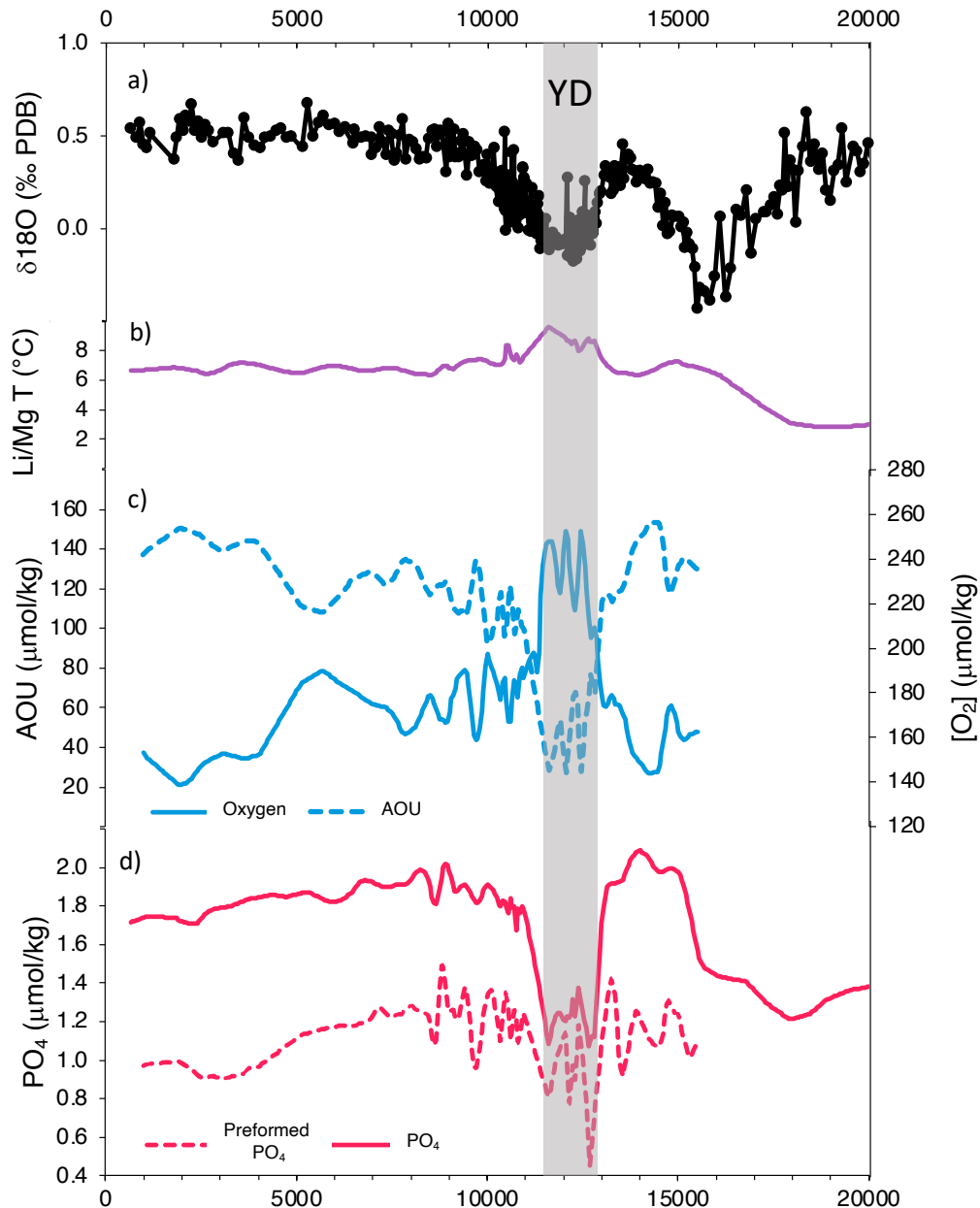
Data S1



**Fig. 1. The nutrient stream in the Florida Straits.** The WOCE A5 section across the Florida Current at 26°N from Florida (left) to the Bahamas (right), just downstream of the location of sediment core KNR166-2-26JPC is shown (29, 30). a) Potential density ( $\sigma_0$ )  $\text{kg m}^{-3}$ , b)  $\text{PO}_4$  ( $\mu\text{mol kg}^{-1}$ ), c) AOU ( $\mu\text{mol kg}^{-1}$ ). The depth of core KNR166-2-26JPC on the Florida Margin is indicated with a magenta circle for all three panels.

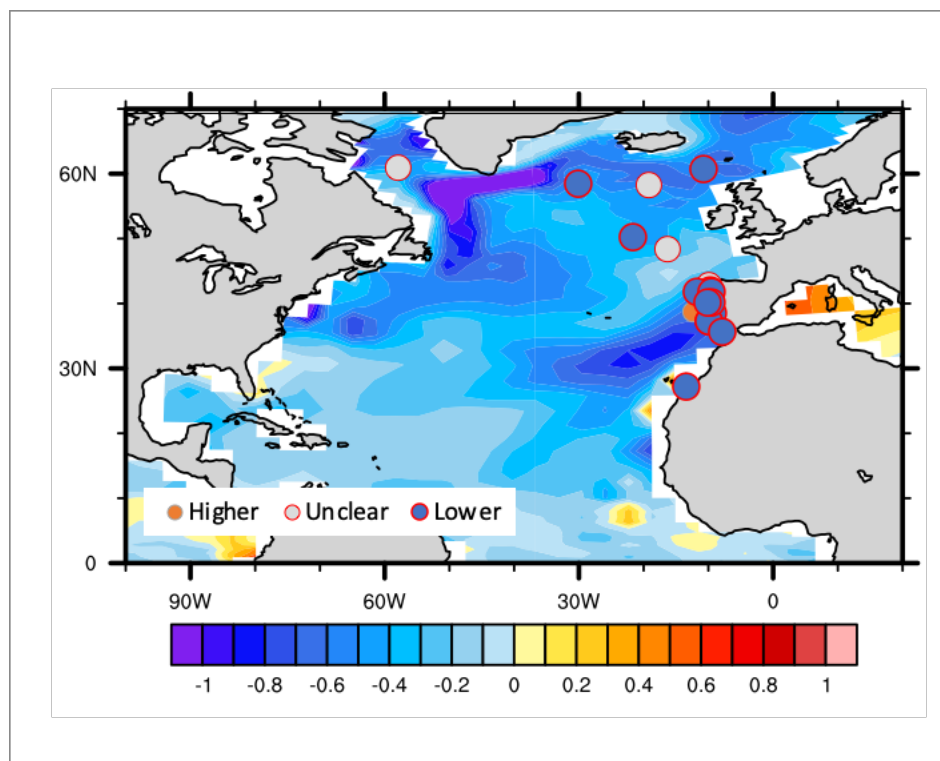


**Fig. 2. History of oxygenation in the Florida Straits.** The average carbon isotopic composition of tests of the foraminifera *P. ariminensis* (yellow) and *Globulimina* (orange) from KNR166-2-26JPC, and the difference in these values converted to oxygen (blue). The 1-sigma calibration error for the oxygen proxy is indicated in blue. The solid line is a loess smooth through the data. Also shown in green are the proportion of planktonic foraminiferal specimens that are associated with low-oxygen values in the subsurface. The grey bar indicates the Younger Dryas time period.



**Fig. 3. Florida Straits circulation and biogeochemistry over the deglaciation.** a) The ice-volume-corrected oxygen isotope ratio of benthic foraminifera on the Florida Margin at KNR166-2-26PC, which reflects changes in the density contrast across the Florida Straits and the strength of the upper branch of the Atlantic meridional overturning circulation (AMOC) (14, 26, 31). b) T estimate from Li/Mg (20) c) Smoothed reconstructed oxygen time series as in Figure 3 (solid line), and the Apparent Oxygen Utilization (AOU) based on Li/Mg T reconstruction

(dashed line). d)  $\text{PO}_4$  estimated from Cd/Ca ratio (19), and preformed  $\text{PO}_4$  estimated using AOU. The grey bar indicates the Younger Dryas time period.



**Fig. 4. Younger Dryas productivity anomaly.** Circles are sediment core locations where reconstructed productivity is clearly lower than time periods before and after the Younger Dryas (blue), higher than the time periods before or after (orange) or where there is no clear anomaly relative to the time before and after (grey) (Table S1). The background is anomaly in the flux of particulate organic carbon ( $\text{mol m}^{-2} \text{yr}^{-1}$ ) between a time in the C-iTRACE simulation (26) with a weak AMOC (17.5ka) and a strong AMOC (19.5 ka).



## Supplementary Materials for

### **A Diminished North Atlantic Nutrient Stream during Younger Dryas Climate Reversal**

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#### **The PDF file includes:**

Materials and Methods  
Figs. S1 to S4  
Tables S1 to S2  
References

#### **Other Supplementary Materials for this manuscript include the following:**

Data S1

## Materials and Methods

### Previously Published Age Model and Measurements

We use the age model published by Waelbroeck et al. 2019 (21) for JPC26. Stable isotope measurements on *Planulina ariminensis* were published in Lynch-Stieglitz et al. 2014 (31) and Lynch-Stieglitz et al. 2019 (32). Cd/Ca and Mg/Li measurements on *Hoeglundina elegans* are published in Valley et al. 2017 (19) and Valley et al. 2019 (20). The calculation of bottom water temperature from Mg/Li and seawater Cd ( $Cd_w$ ) from Cd/Ca is described in these publications.

### *Globobulimina* Stable Isotope Measurements

The sediment core KNR166-2-26JPC was sampled at 2 cm intervals. Specimens of *Globobulimina* were picked from the  $>250\text{ }\mu\text{m}$  sieve size fraction. Where available, 5 specimens were analyzed for each sample depth (average number of specimens analyzed per depth was 2.5). Analyzed individuals had a median mass of 43  $\mu\text{g}$  (range 20-219  $\mu\text{g}$ , 96%  $< 100\text{ }\mu\text{g}$ ). Each specimen was photographed before analysis (**Figure S3**) and analyzed individually using a Thermo Kiel IV carbonate preparation device coupled to a MAT253 and calibrated using NBS-19 and NBS-18. The standard deviation (1 sigma) of replicate analyses of an in-house carbonate standard in the 20-100  $\mu\text{g}$  size range analyzed at the same time as the samples was .06 for  $\delta^{18}\text{O}$  and .03 for  $\delta^{13}\text{C}$ . Measurements with oxygen isotope values that were greater than 2 s.d. away from a robust Loess smoothed version of the record were flagged (4% of the data) and not included in the  $\delta^{13}\text{C}$  averages calculated for each depth (**Figure S4**)

### Determination of Biogeochemical Variables from Proxy Measurements

#### *PO<sub>4</sub>*

$\text{PO}_4$  was estimated from  $Cd_w$  using the relationship described in Elderfield and Rickaby (33) with  $\alpha = 2.5$  for the Atlantic Ocean (34).

#### *O<sub>2</sub>*

In situ oxygen concentration is calculated from the difference between the  $\delta^{13}\text{C}$  of *Globobulimina* and *P. ariminensis* using the relation in Hoogakker et al. (17). While this relation was developed using *C. wuellerstorfi*, in this region *C. wuellerstorfi* are not found at the shallow water depths of this core site, but *P. ariminensis*, another elevated epibenthic species (35) is abundant and reliably records the  $\delta^{13}\text{C}$  of seawater (36). Where *P. ariminensis*  $\delta^{13}\text{C}$  was not available at the same depth as the *Globobulimina*  $\delta^{13}\text{C}$  data, the average  $\delta^{13}\text{C}$  of *P. ariminensis* from the two depths above and below the sample depth was used. This proxy for past oxygen concentration is based on the assumption that *Globobulimina* records the  $\delta^{13}\text{C}$  at the depth in the sediments where pore water oxygen goes to zero, that *P. ariminensis* records bottom water  $\delta^{13}\text{C}$ , and that the pore water  $\delta^{13}\text{C}$  differs from surface water due to the remineralization of organic matter using oxygen supplied by diffusion from above. There is evidence that these assumptions can be violated to some extent in some locations, with observations of deeper depth habitats for *Globobulimina* and evidence for sulfate reduction contributing to the low pore water  $\delta^{13}\text{C}$ , and locations where the proxy doesn't seem to give reasonable values (37-40). Despite these complications,  $\delta^{13}\text{C}$  difference between these species at an increasing number of locations is well correlated to overlying oxygen concentration. At our core site, the reconstructed Holocene values are similar to the modern measurements (150  $\mu\text{M}$ ) near this site. The oxygen

reconstruction matches well with the completely independent Cd-based PO<sub>4</sub> reconstruction, also giving us confidence in the application of the proxy at this site.

### *AOU*

Apparent Oxygen Utilization (AOU) is defined to be the difference between O<sub>2</sub> concentration at saturation and the in-situ oxygen concentration. Oxygen at saturation (TEOS-10) is calculated using the Mg/Li derived temperature. Where Mg/Li temperature is not available at the same sample depth as *Globobulimina*, the average Mg/Li T value for the depths above and below is used. While the oxygen saturation changes are dominated by temperature, we adjust the modern salinity value by scaling the glacial-interglacial difference for the North Atlantic measured at Feni Drift of 1.115 (41) to global ice volume changes with time (42). While we do not adjust for any local changes in salinity, a 1 psu difference in salinity corresponds to only a 2 μmol kg<sup>-1</sup> difference in oxygen saturation. Note that AOU may not be the same as the True Oxygen Utilization (TOU), if the seawater water was not saturated with respect to oxygen when it left the surface. Intermediate waters in today's North Atlantic are estimated to leave the surface with an undersaturation of up to 15-20 μM (43), and as a result AOU overestimates TOU by a similar amount. The initial disequilibrium in intermediate waters in the North Atlantic may increase to up to 35 μM for cold climates (44).

### *PO<sub>4pre</sub>*

For the depths where both AOU and Cd-based PO<sub>4</sub> estimates are available, preformed PO<sub>4</sub> is calculated using AOU:  $PO_{4pre} = PO_4 - (1/170) \cdot AOU$ , where 1:170 is the average P:-O<sub>2</sub> ratio of remineralized carbon (45). When preformed PO<sub>4</sub> is calculated using AOU, the true preformed PO<sub>4</sub> will be underestimated by a proportionate amount, (AOU-TOU)/170. If the disequilibrium for the deglaciation was in the range of 15-35 μM, this would correspond to an underestimate of true PO<sub>4pre</sub> by .1-.2 μM. If the degree of disequilibrium changed over the Younger Dryas climate oscillation, this could change the partitioning of the remineralized and preformed changes by a similar amount. The value of P:-O<sub>2</sub> ratio 1:170 from Anderson and Sarmiento (45) is widely used in models and data studies, and matches the values for the intermediate Atlantic in more recent studies (46). However, these more recent studies suggest a global average ratio of about 1:150. Using this value would reduce P<sub>pre</sub> for the portion of our record with highest AOU values by about .1 μM, and the qualitative inferences we draw here would not change.

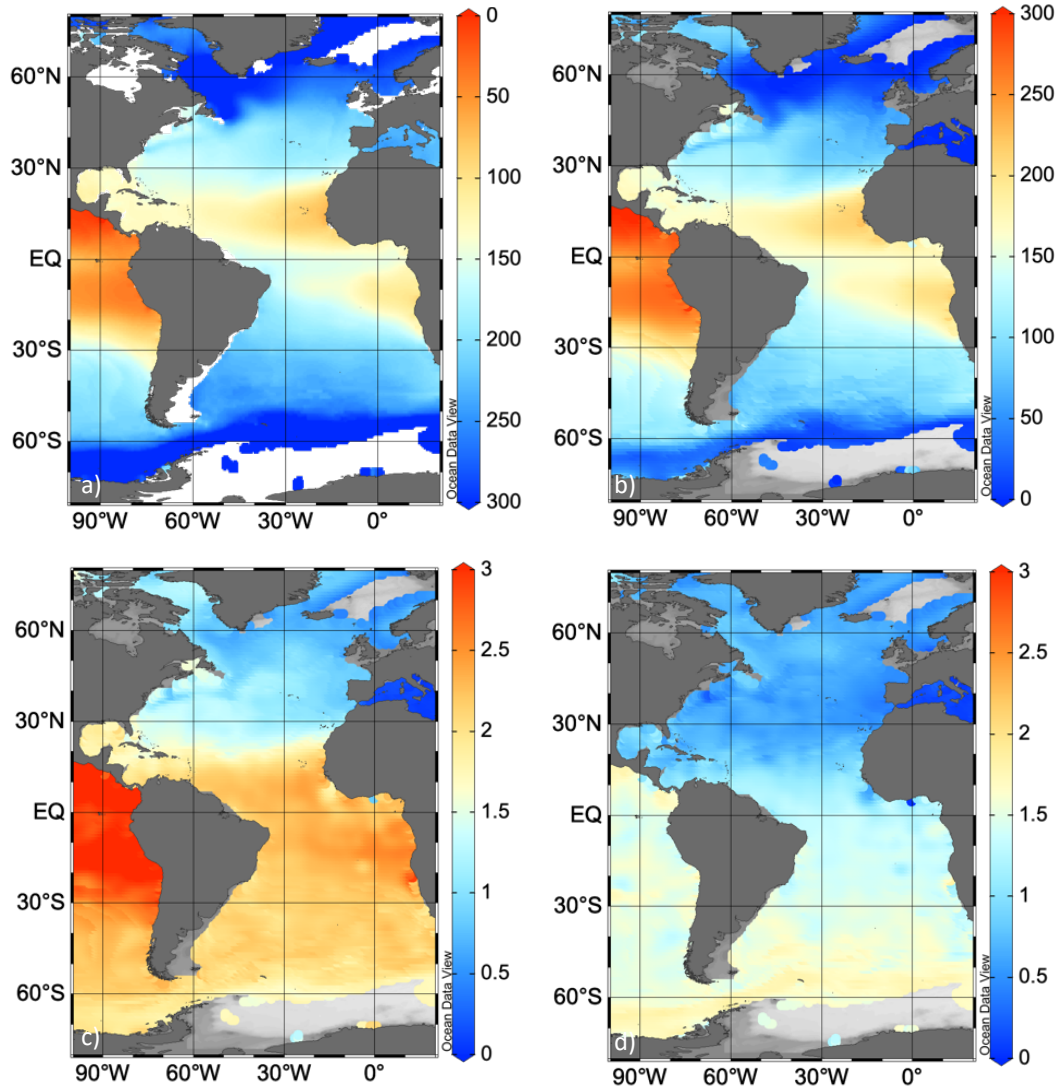
### Planktonic Species Counts

Samples were selected for counting to be approximately 300 years apart based on the age model. We isolated portions of each sample from greater than 150 micrometers sieve fraction. We split the samples, and examined a portion which contained a minimum of 300 total planktonic foraminifera. The total number of planktonic foraminifera, *Globorotalia menardii* and *Globorotalia tumida* (*G. menardii* complex) and *Pullentinia obliquiloculata* were counted. Lastly, prior to this work with the core we had removed *Globogerinoides ruber* from the samples for isotopic analysis and counts of total foraminifera were adjusted to reflect this.

### N. Atlantic Productivity Compilation

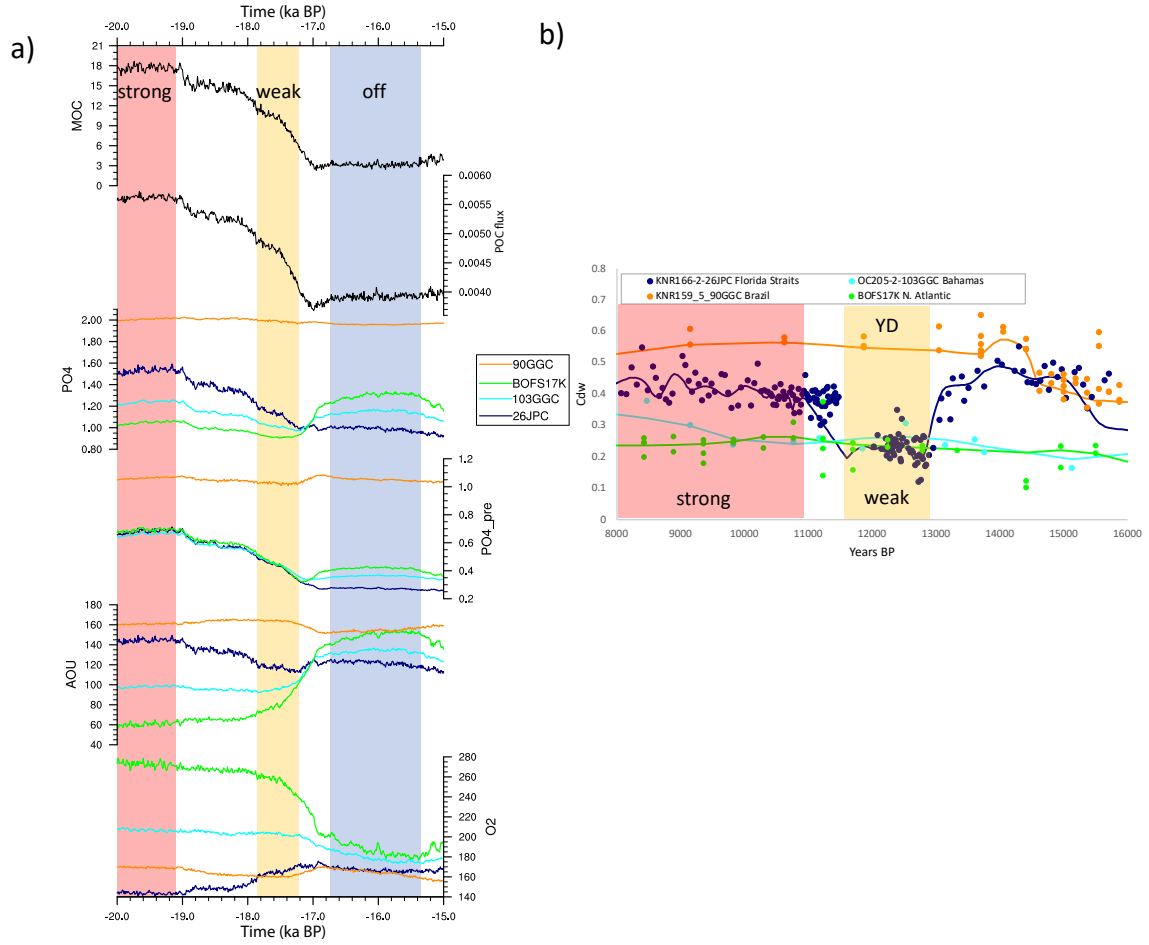
We examined each productivity record that had at least one Younger Dryas aged data point to see if the Younger Dryas productivity was consistently lower or higher than for the data points immediately before and after the Younger Dryas (**Table S1**). If there was no difference or there

was a change across the Younger Dryas (e.g. Younger Dryas productivity was higher than before the Younger Dryas and lower than after the Younger Dryas), the Younger Dryas productivity anomaly was marked “unclear”. Interpretations of productivity proxies and identification of the Younger Dryas interval were taken as written from the original papers.

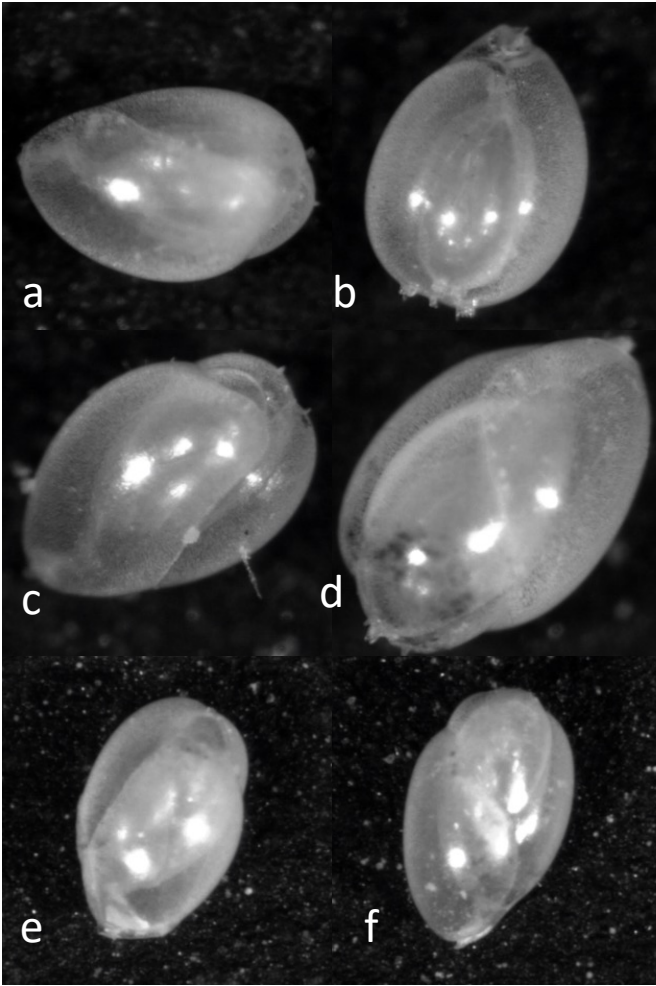


**Fig. S1.**

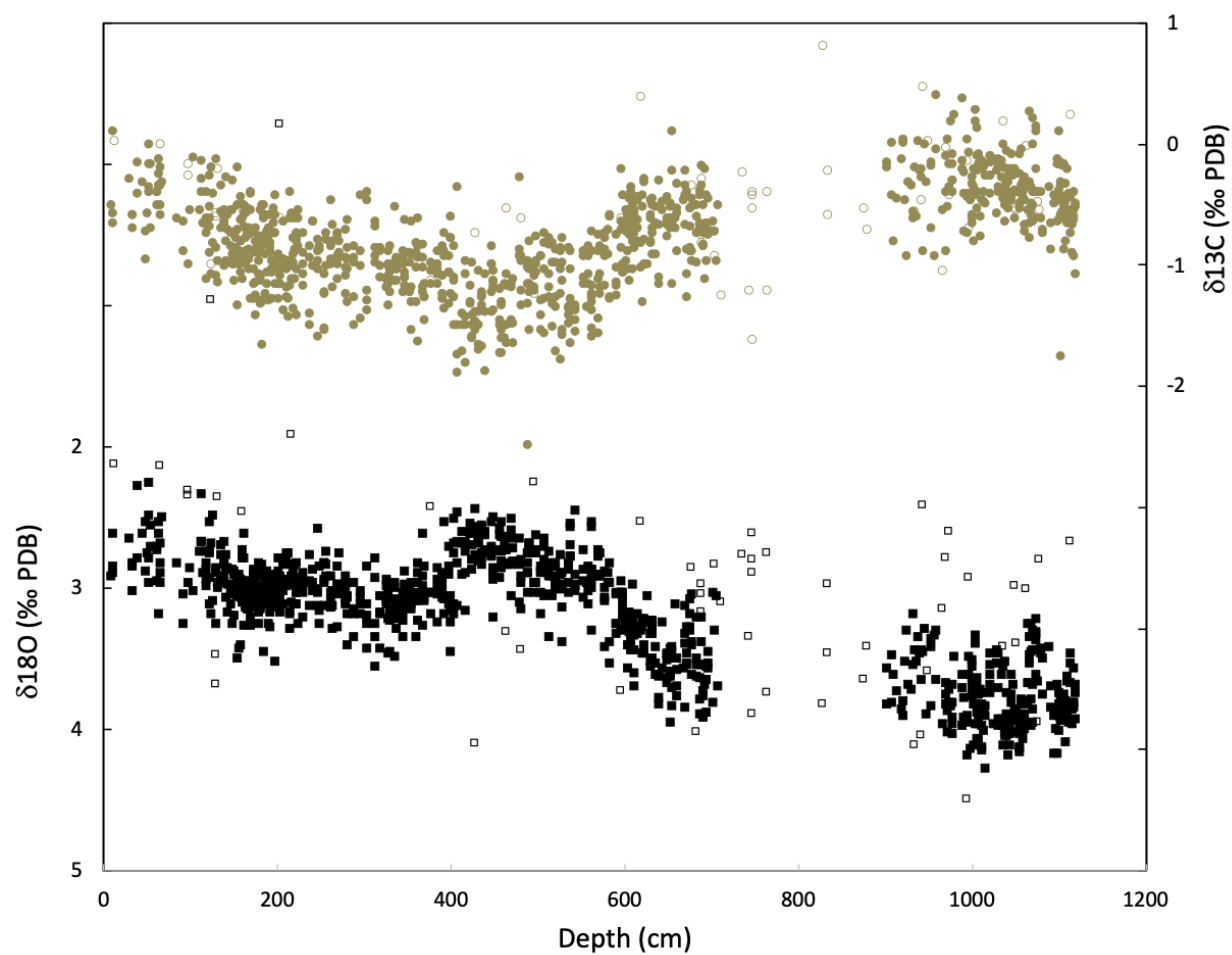
Properties along the 27.3 isopycnal from modern climatology (30, 47, 48). a) Oxygen ( $\mu\text{mol kg}^{-1}$ ) b) AOU ( $\mu\text{mol kg}^{-1}$ ), c)  $\text{PO}_4$  ( $\mu\text{mol kg}^{-1}$ ), d) Preformed  $\text{PO}_4$  ( $\mu\text{mol kg}^{-1}$ ).



**Fig. S2. AMOC strength and regional nutrient patterns.** (a) Time series of AMOC, North Atlantic productivity (averaged over domain shown in Figure 4), and biogeochemical properties from the C-iTRACE simulations between 20 and 15ka (26). Properties are shown at core sites where nutrient reconstructions are available over the Younger Dryas.  $P_{pre}$  is calculated using AOU, in the same way as for the proxy reconstructions. In the model, a weaker AMOC ("weak") is accompanied by lower PO<sub>4</sub>,  $P_{pre}$ , AOU, and higher O<sub>2</sub> at the Florida Straits (dark blue) than for the time when AMOC is strong ("strong"). However, for the open ocean N. Atlantic intermediate water sites (green and light blue), when the AMOC collapses ("off"), nutrient concentrations and AOU increase to levels higher than seen in the intermediate waters in the Florida Straits. (b) Reconstructions of seawater Cd<sub>w</sub>, a proxy for PO<sub>4</sub> over the Younger Dryas. Intermediate water Cd<sub>w</sub> remain high in the South Atlantic, and low in the North Atlantic over the entire time period. Florida Straits values converge towards N. Atlantic values during the Younger Dryas as is seen during times of moderate AMOC weakening in the model ("weak"). Locations, depths, and modern density for the core sites shown are in **Table S2**.



**Fig. S3. Example specimens of benthic foraminifera.** *Globobulimina* from KNR166-2-26JPC, depth in core and mass of test indicated for each: a) 206 cm 36  $\mu\text{g}$  b) 206 cm 44  $\mu\text{g}$ , c) 404 cm 44  $\mu\text{g}$  d) 404 cm 89  $\mu\text{g}$ , e) 686 cm 25  $\mu\text{g}$ , f) 686 cm 25  $\mu\text{g}$



**Fig. S4. Individual *Globobulimina* carbon and oxygen isotope measurements vs. core depth.** Light symbols indicate individuals that were not included in the average  $\delta^{13}\text{C}$  value for each depth.

| Core                    | Latitude<br>(°N) | Longitude<br>(°E) | Younger<br>Dryas<br>Productivity | Productivity<br>Proxy         | Reference |
|-------------------------|------------------|-------------------|----------------------------------|-------------------------------|-----------|
| GeoB5546-2              | 27.5             | -13.7             | low                              | organic carbon                | (49)      |
| DS97-2P                 | 58.9             | -30.4             | low                              | foram accum rates             | (50)      |
| ENAM33                  | 61.3             | -11.1             | low                              | foram accum rates             | (50)      |
| BOFS 5K                 | 50.7             | -21.9             | low                              | benthic foram assemblage      | (51)      |
| SO75-26KL               | 38.8             | -9.5              | low                              | organic carbon                | (52)      |
| U1385                   | 37.6             | -10.1             | low                              | nannofossil abundance         | (53)      |
| SHAK06-5K               | 37.6             | -10.2             | low                              | nannofossil abundance         | (54)      |
| OMEXII-9K               | 42.3             | -10.1             | low                              | planktonic assemblage         | (55)      |
| N3KF24                  | 42.1             | -12.0             | low                              | planktonic assemblage         | (55)      |
| MD03-<br>2697/MD99-2331 | 42.2             | -9.7              | low                              | planktonic assemblage         | (55)      |
| MD95-2040               | 40.6             | -9.9              | low                              | planktonic assemblage         | (55)      |
| MD95-2039               | 40.6             | -10.4             | low                              | planktonic assemblage         | (55)      |
| M39029-7                | 36.1             | -8.2              | low                              | planktonic assemblage         | (55)      |
| BOFS 14K                | 58.6             | -19.4             | unclear                          | benthic foraminifera          | (51)      |
| MD952042                | 37.8             | -10.2             | unclear                          | C37 mass accumulation<br>rate | (56)      |
| PO200-10-26-2           | 41.5             | -9.7              | unclear                          | organic carbon                | (52)      |
| HU2008-029-004          | 61.5             | -58.0             | unclear                          | multiproxy biogenic<br>fluxes | (57)      |
| SU92-03                 | 43.2             | -10.1             | unclear                          | planktonic assemblage         | (55)      |
| OMEXII-5K               | 41.8             | -10.0             | unclear                          | planktonic assemblage         | (55)      |
| MD95-2042               | 37.8             | -10.2             | unclear                          | planktonic assemblage         | (55)      |
| BENGAL<br>13078#16      | 48.8             | -16.5             | unclear                          | benthic foraminifera          | (58)      |
| MD01-2446               | 39.1             | -12.6             | high                             | planktonic assemblage         | (55)      |
| MD99-2339               | 35.9             | -7.5              | high                             | planktonic assemblage         | (55)      |

**Table S1. Productivity records shown in Fig. 4**

| Core            | Latitude<br>(°N) | Longitude<br>(°E) | Depth<br>(m) | Modern $\sigma_t$ | Reference |
|-----------------|------------------|-------------------|--------------|-------------------|-----------|
| KNR159-5-90GGC  | -27.4            | -46.6             | 1108         | 27.3              | (59, 60)  |
| KNR166-2-26JPC  | 24.3             | -82.3             | 546          | 27.3              | (19)      |
| OCE205-2-103GGC | 26.1             | -78.1             | 965          | 27.6              | (61)      |
| BOFS17K         | 58.0             | -16.5             | 1150         | 27.6              | (62, 63)  |

**Table S2. Cd Records shown in Fig. S2**

**Data S1. (separate file)**

An excel file containing measured and derived variables. Data are also be archived at the World Data Center for Paleoclimatology.