2016-06-17

The 8.2-ka BP event in north-eastern North America: first combined oxygen and hydrogen isotopic data from peat in Newfoundland

Daley, Tim

http://hdl.handle.net/10026.1/9780

10.1002/jqs.2870
Journal of Quaternary Science
Wiley

All content in PEARL is protected by copyright law. Author manuscripts are made available in accordance with publisher policies. Please cite only the published version using the details provided on the item record or document. In the absence of an open licence (e.g. Creative Commons), permissions for further reuse of content should be sought from the publisher or author.
The 8.2-ka BP event in north-eastern North America: first combined oxygen and hydrogen isotopic data from peat in Newfoundland

T. J. DALEY, *1 K. E. BARBER, 2 P. D. M. HUGHES, 2 N. J. LOADER, 3 M. LEUENBERGER 4 and F. A. STREET-PERROTT 3

1 School of Geography, Earth and Environmental Sciences, Plymouth University, Plymouth PL4 8AA, UK
2 Palaeoenvironmental Laboratory, Geography and Environment, University of Southampton, Highfield, Southampton SO17 1BJ, UK
3 Department of Geography, College of Science, Swansea University, Singleton Park, Swansea SA2 8PP, UK
4 Climate and Environmental Physics, Physics Institute, and Oeschger Centre for Climate Change Research, University of Bern, CP-J3012 Bern, Switzerland

Received 8 January 2016; Revised 27 April 2016; Accepted 6 May 2016

ABSTRACT: Finding direct evidence for atmospheric circulation change in terrestrial records of Holocene climate variability remains a fundamental challenge. Here we present the first combined stable oxygen and hydrogen isotopic palaeorecord from a peatland core in Newfoundland, Canada. Sphagnum cellulose samples were isolated from a core from Nordan’s Pond Bog, Newfoundland, and analysed for δD values. Combined with existing Δ18O data, the resulting δD/Δ18O bi-plot correlates directly with existing measurements of the modern (late 20th century) isotopic composition of precipitation from GNIP stations in Nova Scotia and Labrador, implying a close relationship between the estimated isotopic composition of source water used by the mosses and that of the source precipitation. We use the relative variations between the two isotope records to test the hypothesis that atmospheric circulation changed in the millennium following the 8.2-ka BP climate event. The data reveal a secondary complex isotopic response ~200 years (8250–8050 b.p.) after a primary oxygen isotopic event that is widespread in the North Atlantic region. This secondary event is characterized by a divergence in oxygen and hydrogen isotope records that can most plausibly be explained by the augmentation of precipitation moisture from a more distant and more continental vapour source.

Copyright © 2016 The Authors. Journal of Quaternary Science Published by John Wiley & Sons Ltd.

KEYWORDS: 8.2 ka; hydrogen; isotopes; Newfoundland; North Atlantic; Sphagnum.

Introduction

Ensemble model-simulated projections for 21st-century climate in the North Atlantic region suggest future warming may be mitigated in part by a decrease in the heat transfer from low to higher latitudes associated with slow down in Atlantic Meridional Overturning Circulation (AMOC) (Hofmann and Rahmstorf, 2009; Collins et al., 2013). Changes in the surface oceanic heat gradient between low and high latitudes will impact upon the gradient in atmospheric temperature, thereby influencing the strength and position of the northern mid-latitude atmospheric jetstream circulation (hereafter jet) (Tang et al., 2014; Francis and Vavrus, 2015). In this context, quantification of the impact of past changes in the northern mid-latitude atmospheric jet in response to oceanic circulation changes is valuable for constraining model simulations of similar scenarios in future climate projections (LeGrande and Schmidt, 2009; Tindall and Valdes, 2011; Holmes et al., 2016).

The most compelling evidence for past changes in the northern atmospheric jet has been interpreted from the Greenland ice core records, where the multitude of independent indicators is sufficient to provide a convincing record (Dawson et al., 2003; Vinther et al., 2003; Alley and Ágústsdóttir, 2005; Jouzel et al., 2000; Daley et al., 2000; Daley et al., 2014). Past changes in d relate most strongly to variations in atmospheric conditions over the vapour source region (Dansgaard, 1964; Merlivat and Jouzel, 1979; Gunthe et al., 2000; Sonntag et al., 1983; Rozanski et al., 1993). Variations in d may therefore indicate changes in the source area for local precipitation. Traditionally, relatively high positive values of d (~10) have been used to diagnose moisture originating from surface evaporation in an environment with relatively low humidity and/or high wind speeds and/or relatively high ocean temperatures. Values <10 have indicated a moisture source that was humid and/or with relatively low surface water temperatures. Previous research estimated that a 10% increase in relative humidity over the ocean decreased d by ~6‰ (Rozanski, 1985). Recently, however, it has been demonstrated that the combined record of oxygen and hydrogen data is significant, not least for the ability to detect changes in the source of moisture (Dansgaard, 1964; Merlivat and Jouzel, 1979; Sonntag et al., 1983; Rozanski et al., 1993; Jouzel et al., 2007; Pfahl and Sodemann, 2014; Steen-Larsen et al., 2014). This insight is afforded by variations in the rate of kinetic fractionation of the isotopomers of hydrogen and oxygen in water and which are expressed as variations in the deuterium excess (d), given by the formula d = −8δD − 1008Ω. Deuterium excess data have already demonstrated their interpretative value in modern instrumental studies (Pfahl and Wernli, 2008; Daley et al., 2012; Steen-Larsen et al., 2014) and through analyses of ice cores (Jouzel et al., 2007; Steffensen et al., 2008), but they also now have a potential role in disentangling the causes of what can often be large variations in mid-latitude isotopic records that are rarely related to a single driver such as atmospheric temperature or the amount of precipitation alone (Araguás-Araguás et al., 2000; Daley et al., 2012; Young et al., 2015; Holmes et al., 2016).
effect of wind speed over the sites of oceanic evaporation is a relatively minor contributory factor compared with relative humidity in the labelling of δ in meteoric waters (Pfahl and Sodemann, 2014; Steen-Larsen et al., 2014). Furthermore, Pfahl and Sodemann (2014) have demonstrated that relative humidity dominates any signal from the surface ocean temperatures. Indeed, they suggest that palaeorecords of variations in δ should now be interpreted as variations driven primarily by relative humidity changes at the site of moisture source. Ice core records are the only source of combined oxygen and hydrogen isotopic measurements of palaeo-precipitation (e.g. Jouzel et al., 2007) but they are limited to high-latitude and high-altitude locations and so there remains a gap in evidence from the mid-latitudes with which to test ideas of atmospheric circulation change.

The 8.2-ka BP climate event provides an ideal test case for a first comparison between hydrogen and oxygen records from a mid-latitude Sphagnum moss archive given that it was the most severe climatic event in the Holocene and appears to have had most pronounced impact in the northern mid- to high latitudes (Alley and Agüestdöttir, 2005; Rohling and Pälike, 2005; Thomas et al., 2007; Daley et al., 2011; Morrill et al., 2013). The 8.2-ka BP event was a circum-North Atlantic region cold event lasting ~150 years approximately 8200 years ago and occurred in response to changes in the salinity balance of the North Atlantic Ocean (Barber et al., 1999). An injection of meltwater from the catastrophic flood of Glacial Lake Agassiz-Ojibway provided the trigger that initiated an oceanic-atmospheric response, the spatial parameters of which are already well reproduced in numerical model experiments (LeGrande et al., 2006; LeGrande and Schmidt, 2008; Tindall and Valdes, 2011; Holmes et al., 2016). The most pronounced isotopic excursion reported thus far was found in a peat core from Newfoundland (Daley et al., 2009). Progressively reduced values for the isotopic excursion were observed with increasing distance from the Labrador Sea (Marshall et al., 2007; Dominguez-Villar et al., 2009; Daley et al., 2011; Holmes et al., 2016). While the spatial pattern of the climate event has been reproduced well by model experiments, the duration of the event has remained a challenge for those models to simulate (Daley et al., 2011; Tindall and Valdes, 2011). Recent work has demonstrated that data model comparisons offer the best signal over the longer-term records implies that there remains a limited to an alternative, more distant source. This process would be evidenced by a change in the deuterium excess and by a relative decrease in δ18O values if the distance from vapour source to site of precipitation were greater than that which is observed in the modern day. In this study, therefore, we used combined hydrogen and oxygen isotopic data from the same sub-samples of Sphagnum cellulose from a peatland in Newfoundland to test directly the hypothesis that deuterium excess (and therefore atmospheric circulation) changed in the millennium following the 8.2-ka BP event.

Understanding the hydrogen isotopic signal of the source water in Sphagnum cellulose

Sphagnum cellulose isotopic values are modified from the source water (precipitation) isotopic composition by local environmental processes and by the biochemical processes occurring to produce cellulose (Ménot-Combes et al., 2002; Daley et al., 2010). First, isotopic values may be modified by any evaporation of precipitation waters on the surface of the bog before use of the water by the plant. Secondly, isotopic values are modified by biochemical fractionation during cellulose synthesis. The processes leading to the oxygen isotopic labelling of Sphagnum cellulose are relatively well understood. Sphagnum δ18O values have been shown to more closely approximate that of the source precipitation than bog surface waters based on long-term surface monitoring of several sites in Europe (Daley et al., 2010). The close linkage between the oxygen isotopic composition of Sphagnum cellulose and its meteoric source water has been applied in subsequent palaeoclimatic studies (Kaislahti Tillman et al., 2010; Roland et al., 2015). Several studies have detected differences in Sphagnum isotopic values associated with microtopographical variations and distance from the water table (Brenninkmeijer et al., 1982; Daley et al., 2010; Loader et al., 2016). The study by Daley et al. (2010) alongside earlier pioneers (Brenninkmeijer et al., 1982; Aravena and Warner, 1992) also observed similar scales of variation between surface samples during individual sampling experiments. While evaporation undoubtedly influences surface variability, extended monitoring over 2 years demonstrated that any topographic control on variation in δ18O (Sphagnum) values was not consistent through time, partly because this difference was enhanced in summer bog waters (when most sampling in these studies has taken place) relative to spring or autumn waters in a way that was not consistently captured by the mosses. This longer-term pattern is probably due to temporary suspension of photosynthesis under dry conditions, when the evaporative effect would be greatest (Williams and Flanagan, 1996). The limited effect of this local spatial fractionation signal over the longer-term records implies that there remains reliability in the decadal-synthesized signal in peatland
palaeorecords, while recognizing that there still remains a challenge to characterize and quantify the evaporative influence (Loader et al., 2016).

Biochemical fractionation affects oxygen and hydrogen to different extents due to the different pathways followed during cellulose synthesis (Estep and Hoering, 1981; Sternberg and DeNiro, 1983; Sternberg et al., 1986; Yakir and DeNiro, 1990; Sternberg and Ellsworth, 2011). For both elements, the cellulose synthesis process involves reactions that take place in light conditions (photosynthesis) and those that occur during biosynthesis later on in the pathway (Yakir and DeNiro, 1990). Cellulose synthesis of oxygen atoms exhibits a nearly constant enrichment of ~27‰ relative to source water in plants from all photosynthetic modes and under a range of environmental conditions (DeNiro and Epstein, 1979, 1981; Sternberg et al., 1986; Yakir and DeNiro, 1990; Waterhouse et al., 2013). Enrichment results from isotopic fractionation during hydration of carbonyl groups of the intermediates of cellulose synthesis (DeNiro and Epstein, 1981). Any temperature-dependent effect is likely to be limited although may be present (Sternberg and Ellsworth, 2011). Plants of all photosynthetic modes have a stage in which oxygen passes through a carbonyl stage, which explains the occurrence of 27‰ in all plant types (Sternberg et al., 1984a,b; Yakir and DeNiro, 1990).

Hydrogen fixation in cellulose, by contrast, is not the same for all plants. This difference results partly from the variable extent of biochemical fractionation occurring in different biosynthetic pathways (Estep and Hoering, 1980, 1981; Yakir and DeNiro, 1990). Hydrogen in cellulose is derived entirely from hydrogen in leaf water (Yapp and Epstein, 1982). Fractionation is regulated by two sets of processes: those occurring in the leaf during photosynthesis (leading to strong depletion in the cellulose) and subsequent processes of cellulose synthesis from the intermediates (leading to enrichment) (Estep and Hoering, 1981; Yakir and DeNiro, 1990; Luo and Sternberg, 1991). The photosynthetic depletion relates to preferential hydration of nicotinamide adenine dinucleotide phosphate (NADP) with protium rather than deuterium to form NADPH (Yakir and DeNiro, 1990). Subsequent enrichment occurs during exchange with hydrogen in heterotrophic cellulose synthesis (Estep and Hoering, 1981; Yakir and DeNiro, 1990; Luo and Sternberg, 1991). The fractionation factor that describes the isotopic discrimination in the formation of cellulose from water is defined as:

\[
\alpha = \left(\frac{\delta_{\text{cellulose}}}{1000}\right) + 1 = \left(\frac{\delta_{\text{source water}}}{1000}\right) + 1
\]

In modern Patagonian Sphagnum, αD has been determined as 0.922 and reflects a greater net photosynthetic depletion of Sphagnum with respect to source waters (Pendall et al., 2001). Strong depletion of Sphagnum δD relative to vascular plants has also been reported from sites in Europe (Brenninkmeijer et al., 1992). Investigation of the fractionation of hydrogen in Sphagnum cellulose has been limited, although the studies above indicate that a strong depletion signal may be present in the genus (Loader et al., 2016). It is possible that this may reflect the simple physiology of Sphagnum reflecting limited post-photosynthetic exchange of intermediates used in cellulose synthesis.

Methods

A 9-m-long peat core (NDN02/1) was collected from Nordan’s Pond Bog in 2002 (Fig. 1) using a large-bore Russian corer (Hughes et al., 2006). Hydrogen isotope analyses were undertaken on aliquots of the same Sphagnum

![Figure 1. Map of the North Atlantic region showing modern ocean surface currents, Glacial Lake Agassiz outflow, NDN02/1 site location (open triangle), Global Network of Isotopes in Precipitation (GNIP) monitoring stations at Goose Bay, Labrador and Truro, Nova Scotia (open circles), and published reference site locations for existing high-resolution records. Solid circles = lake sediment cores, solid triangles = ice cores, open squares = speleothem sites. Dark arrows indicate warm surface ocean currents: GS = Gulf Stream, NAD = North Atlantic Drift, IC = Irminger Current. Grey arrows indicate cool surface ocean currents: EGC = East Greenland Current, LC = Labrador Current. Thick open arrow indicates routing of final drainage of glacial Lake Agassiz.](image)
cellulose sub-samples previously reported for oxygen isotope analyses (Daley et al., 2009). The cellulose sub-samples had previously been prepared from Sphagnum leaves that were isolated from the peat matrix using a stacked sieve system and density-separation procedure (Daley et al., 2009, 2010). Four adjacent cellulose samples at 8-cm separation were selected at ~1000-year intervals based on the published chronology (Daley et al., 2011) to provide a broad stratigraphic framework of δD_{Sphagnum} values throughout the Holocene. Sample resolution was increased to one sample every 4 cm from ~8400 to 8000 a BP and contiguous 1-cm samples were analysed across the primary isotopic event observed in the oxygen isotope stratigraphy, providing sub-centennial and decadal resolution, respectively, such that hydrogen sample resolution equalled that of the oxygen record through 576–708 cm (7594–8812 a BP).

We used an online equilibration method for the measurements of δD_{Sphagnum} values because of the benefits offered in terms of the rapidity of the process, the small sample sizes necessary, the limited fractionation effects and the avoidance of dangerous chemical procedures (Filot et al., 2006; Loader et al., 2015). The procedure used is described in detail in Filot et al. (2006) and is summarized here. Samples of 0.5–0.7 mg of dry alpha cellulose were weighed into tin capsules and wrapped loosely to allow water vapour to enter while remaining closed enough to prevent sample loss. Samples were then placed into an equilibration chamber and subjected to a continuous flow of a standard ‘Meerwasser’ water vapour of known isotopic composition (+15‰ vs. Vienna Standard Mean Ocean Water (VSMOW)) at 110°C and a flow rate of 7.8 μL min⁻¹ delivered by a helium carrier gas stream at 40 mL min⁻¹. After 540 s the water supply was stopped and the sample transferred into an AS128 autosampler and then the pyrolysis reactor after only a few seconds. Pyrolytic conversion took place at 1450°C over a glassy carbon granulate on a 2-mm layer of silver wool and 20-mm bed of quartz wool in a thermo-chemical elemental analyser. Pyrolysis products were separated using a 1-m GC column then carried by a ConFlow II open split unit to a Thermo Finnigan Delta plus XL isotope ratio mass spectrometer. Results are given in standard delta notation relative to VSMOW. Repeat measurements of three separate cellulose standards (Merck; IAEA; Sigma) indicated measurement error was limited to ~5‰. The D/H ratio of non-exchangeable carbon-bound hydrogen was estimated from the bulk hydrogen isotope data using a mass balance relationship (Filot et al., 2006).

Results and discussion

δD_{Sphagnum} values for core NDN02/1 from Nordan’s Pond Bog are presented as a time series alongside δ¹⁸O_{Sphagnum} values (Daley et al., 2009) using the age–depth model from Daley et al. (2011) in Fig. 2. The grey bars in the figure represent the 2σ range on repeat sample measurements. Age error estimates are reported here based on the maximum and minimum ages from the published CLAM age depth model (Daley et al., 2011). With the exception of the early Holocene event, δD_{Sphagnum} values show variation centred on ~−120 ± 10‰ with no general increasing or decreasing trend. δD_{Sphagnum} values through the early Holocene event exceeded the range of variability found throughout the rest of the record, albeit with a lower sampling resolution from ~7950 (± 50) a BP to present.

At ~8450 (± 50) a BP, δD_{Sphagnum} values fell ~15‰ in ≤55 years stabilized for ~40 years then dropped a further ~22‰ in approximately 10 years to an isotopic minimum ~8350 (± 40) a BP. The subsequent recovery was very rapid (~30‰ in ~20 years). δD_{Sphagnum} values oscillated for the following ~100 years then recovered to ~130 ± 5‰ for the next ~200 years.

A good general correspondence between the hydrogen and oxygen isotopic records is observed. Both datasets show relatively high values at the base of the record, a strong isotopic depletion with a minimum at ~8350 (± 40) a BP, higher and generally rising values between ~7800 (± 60) a BP and ~4200 (± 90) a BP, followed by a fall to lower values centred on ~3500 (± 55) a BP rising to higher values again ~2200 (± 40) a BP. Lower correspondence between the two records is observed in the upper section of the core. Between ~10000 (± 55) and ~600 (± 50) a BP, δ¹⁸O_{Sphagnum} values exhibit a relatively low-magnitude isotopic decline. This corresponds to a large decline in δD_{Sphagnum} values. In the top four samples, δD_{Sphagnum} values again show relatively higher variability than the equivalent δ¹⁸O_{Sphagnum} values for these levels.

The duration of the primary early Holocene isotopic event for both records was similar if the primary event is considered to have been initiated from the point of inflexion in both records at 656 cm (~8450 ± 50 a BP). The most notable difference between the two records is in the presence of a second isotopic minimum that succeeds the primary event. In the δ¹⁸O_{Sphagnum} data, a second minimum was centred on ~8150 (± 40) a BP, was of roughly half the magnitude of the primary event and had a duration of ~200 years. In the δD_{Sphagnum} data, we find no evidence for a contemporaneous secondary minimum. Rather, the secondary oxygen minimum is associated with δD_{Sphagnum} values within 1σ of the mean for the record. During the period in which δ¹⁸O_{Sphagnum} values are observed to fall, δD_{Sphagnum} values show a general increasing trend. Comparison of the two isotopic tracers therefore suggests that a high-magnitude isotopic event occurred ~8450 (± 50) a BP with a minimum centred on ~8350 (± 40) a BP and had a total duration of ~150 years, consistent with the broader north Atlantic region isotopic anomaly. These followed a divergence in δD_{Sphagnum} and δ¹⁸O_{Sphagnum} values. This divergence lasted approximately 200 years and probably represented a sustained change in δ (Fig. 2).

To estimate the isotopic composition of the cellulose source water (δw), we applied the published cellulose–water enrichment factors of δD = 0.922 (Pendall et al., 2001) and α¹⁸O = 1.274 (Daley et al., 2010) to the δSphagnum cellulose values. The resulting estimates overlap modern GNIP measurements of the isotopic composition of precipitation from Goose Bay, Labrador and Truro, Nova Scotia (IAEA/WMO, 2004) in the bi-plot in Fig. 3. Average seasonal values for modern GNIP data are presented in Table 1. The linear regression for both the measured meteoric waters (or regional meteoric water line, RMWL; δD = 7.47δ¹⁸O + 5.38; R² = 0.97) and the NDN02/1-inferred palaeo-estimates (δD = 8.26δ¹⁸O + 5.77; R² = 0.61) are remarkably similar (Fig. 3). The 95% confidence and prediction limits are given as dotted and dashed lines, respectively (Fig. 3). The close correspondence between δw estimates and the RMWL suggests that δw was equivalent to δprecipitation. The mean of δ for the length of the record is 4.05 (n = 51). The minor offset of the palaeo-slope to the right of the RMWL may have two explanations. First, the offset may represent the relatively consistent time-integrated local evaporation effect on δSphagnum values from which the estimates are derived. Or, secondly, it may represent consistently lower deuterium excess in cellulose relative to measured annual weighted δprecipitation values because of the seasonality of Sphagnum moss growth. Annually, deuterium excess values vary through
the year such that during summer months relatively low values of $d$ are observed, reflecting warmer ocean water and higher relative humidity in the sites of moisture sourcing. In the winter, higher values of $d$ are observed. In global meteoric waters, this is observed as an inter-hemispheric variation over the year (Clark and Fritz, 1997; Gibson et al., 2005; Pfahl and Sodemann, 2014; Steen-Larsen et al., 2014). Given that Sphagnum would suspend growth in the absence

Figure 2. Combined records of $\delta^18$O Sphagnum values (black) and $\delta^13$O Sphagnum values (dark grey; Daley et al., 2009) from NDN02/1. All values are given in delta notation relative to VSMOW. Chronology is that published in Daley et al. (2011). Dating control points are highlighted by black triangles. Values represent the best estimate based on the published CLAM age depth model (Daley et al., 2011).
of liquid water, either during periods of aridity or when unavailable (frozen) and that average temperatures in Newfoundland in the modern day are significantly below 0 °C for at least 3 months of the year, there would be a natural bias in the record towards summer season cellulose synthesis.

Modern climate data for 1971–2000 from Musgrave Harbour (49.450°N, 53.967°W; altitude 3 m AOD; 43 km north-west of Nordan’s Pond Bog) demonstrate the range of temperature seasonality. Mean monthly temperatures vary from a January maximum of 2.3 °C and minimum of −9.8 °C to a July maximum of 19.9 °C and minimum of 10.8 °C. Mean monthly maximum temperatures between 1971 and 2000 were below

![Graphical representation of seasonal temperature data](image)

**Figure 3.** Comparison of Holocene Sphagnum source water reconstruction from NDN02/1 with modern GNIP precipitation data (IAEA/WMO, 2004) using $\delta ^{18}O = 1.0274$ (Daley et al., 2010) and $\delta D = 0.922$ (Pendall et al., 2001) for cellulose–water enrichment. GNIP variation in the isotopic composition of meteoric waters is given by season. Closed symbols = Truro; open symbols = Goose Bay. Cellulose palaeo-source water estimates = grey hexagons. Black hexagons in expanded insert = samples from the period 8350–8050 a BP with relatively high values of $\delta D$. 


<table>
<thead>
<tr>
<th></th>
<th>Autumn</th>
<th>Winter</th>
<th>Spring</th>
<th>Summer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Goose Bay $\delta ^{18}O$</td>
<td>−14.6</td>
<td>−19.8</td>
<td>−15.4</td>
<td>−12.4</td>
</tr>
<tr>
<td>$\sigma$</td>
<td>2.1</td>
<td>4.0</td>
<td>3.0</td>
<td>1.4</td>
</tr>
<tr>
<td>$\delta D$</td>
<td>−102.0</td>
<td>−142.0</td>
<td>−113.9</td>
<td>−89.2</td>
</tr>
<tr>
<td>$\sigma$</td>
<td>15.6</td>
<td>30.5</td>
<td>22.5</td>
<td>12.0</td>
</tr>
<tr>
<td>Truro $\delta ^{18}O$</td>
<td>−8.5</td>
<td>−12.6</td>
<td>−8.8</td>
<td>−7.0</td>
</tr>
<tr>
<td>$\sigma$</td>
<td>1.9</td>
<td>3.3</td>
<td>2.4</td>
<td>1.3</td>
</tr>
<tr>
<td>$\delta D$</td>
<td>−56.7</td>
<td>−87.5</td>
<td>−62.2</td>
<td>−48.9</td>
</tr>
<tr>
<td>$\sigma$</td>
<td>15.3</td>
<td>24.4</td>
<td>18.3</td>
<td>11.2</td>
</tr>
</tbody>
</table>
0 °C for the winter months (December, January and February). This would, correspondingly, explain a plot with a slope consistent with meteoric waters, but a lower intercept. On balance, it is more likely that the small persistent offset, associated with a mean value for $d$ slightly lower than RMWL intercept (4.05; $n=51$) and similarity in slope results from the seasonal growth bias. An evaporative effect would be expected to become more effective under higher surface air temperatures and be reflected in a local evaporative line with a lower slope angle.

Figure 4. $\delta^{18}$O$_{\text{precipitation}}$ values (black), $\delta^{18}$O$_{\text{precipitation}}$ values (dark grey) and deuterium excess values ($d$; dashed line) from NDN02/1 Sphagnum cellulose (top) compared with decadal-scale terrestrial isotopic time series for 9200–7400 a BP (BP = 1950 AD) for the North Atlantic region synchronized by start date ($T=0$; adapted from Daley et al., 2011). The time from start of primary event progresses from left to right. (A) $\delta^{18}$O$_{\text{precipitation}}$ reconstruction (black line) with $\delta^{18}$O$_{\text{precipitation}}$ reconstruction (grey line) and $d$ values (dashed line) from Sphagnum peat core NDN02/1, Newfoundland (Daley et al., 2009). Grey shaded box indicates period of higher $d$ values. (B) Greenland composite $\delta^{18}$O ice record (Thomas et al., 2007). (C) Variation in lake-sediment $\delta^{18}$O$_{\text{calcite}}$ from Lake Rouge, Estonia (Veski et al., 2004). (D) Sub-decadal variation in $\delta^{18}$O$_{\text{calcite}}$ from speleothem YD01, from Pippikin Pot Cave (Daley et al., 2011). (E) Oxygen isotopic variation in lake-sediment calcite from Hawes Water, northern England (Marshall et al., 2007). (F) Variation in $\delta^{18}$O$_{\text{calcite}}$ from annually banded speleothem WSC97-10-5, from White Scar Cave (Daley et al., 2011). (G) Oxygen isotopic variation in lake-sediment calcite from Lough Avolla, western Ireland (Holmes et al., 2016). (H) $\delta^{18}$O$_{\text{precipitation}}$ reconstruction from lake-sediment calcite from the Ammersee, Germany (von Grafenstien et al., 1998). (I) Variation in $\delta^{18}$O$_{\text{calcite}}$ from speleothem LV5, northern Spain (Domínguez-Villar et al., 2009).
In a recent review of the 8.2-ka BP event, Daley et al. (2011) compiled published isotopic records from sites around the North Atlantic region. The records were presented in calendar years since published start date for the event, given as T = 0. Comparison of the new $^{18}D_{\text{Sphagnum}}$ record with these existing reconstructions (Fig. 4) demonstrates that the secondary anomaly in $^{18}O$ values in NDN02/1 (Daley et al., 2009) is replicated neither in $^{18}D_{\text{Sphagnum}}$ values nor more broadly and is associated with relatively high $d$ values (Fig. 4). Only three mechanisms could explain this divergence. The first is that there was strong localized evaporation of the source water before cellulose synthesis. The second is that Nordan’s Pond Bog, in effect, became more continental via increased distance between moisture source and the site. A third is that moisture from an evaporated, non-oceanic source contributed significantly to the source of the precipitation at Nordan’s Pond Bog. A conclusion based on the first possibility would seem inconsistent with the broader evidence for a consistent slope to the reconstructed source water line and the integrated offset based on a predominance of Sphagnum growth in the summer months. Furthermore, the period 8250–8050 BP was a relatively cool (Leuenberger et al., 1999; Hughes et al., 2006; Thomas et al., 2007). While it is possible that vapour pressure deficits may have increased in cool, dry air conditions (Alley and Ágústdóttir, 2005), it seems unlikely that this was so sustained and so effective as to have produced the observed divergence in $^{18}O$ and $D$ values. Further still, palaeoecological evidence from the core indicates that relatively wet bog surface conditions prevailed, suggesting minimal evaporation from the site (Hughes et al., 2006; Charman et al., 2009). Finally, localized evaporation cannot explain the decrease in $^{18}O$ values. Instead, the second explanation would appear consistent with the observed extension of perennial sea ice in the North Atlantic and the decrease in $^{18}O$ values (Müller et al., 2009). The development of a longer pathway from water source to the peatland site is likely to have involved variation in the mean position or intensity of the northern atmospheric jet. Model simulations have demonstrated enhanced Atlantic anticyclonic activity during the 8.2-ka BP event (LeGrande and Schmidt, 2008; Clarke et al., 2009; Tindall and Valdes, 2011), but none of which we are aware has run through the following millennium with similar temporal resolution. The enhanced temperature gradient between low and high latitudes induced by the slowdown in AMOC (Kleiven et al., 2008) and extension of sea ice (Ellison et al., 2006; Müller et al., 2009) probably influenced the atmospheric jet such as to increase the intensity of zonal flow. However, it seems unlikely that a more intense and zonal atmospheric jet would enable the sourcing of oceanic moisture for Newfoundland precipitation from a more distant Atlantic source. A third possibility, then, is that the remaining bodies of continental and microenvironmental water relations. Hydrological Processes 14: 1341–1355.


