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the reconstruction of past $\delta^{18}\text{O}_{\text{lw}}$ composition provides valuable insights into the SHW evolution at high southern latitudes.

Over recent decades, it has been widely recognized that the oxygen isotope composition of aquatic cellulose ($\delta^{18}\text{O}_{\text{cell}}$) is a reliable recorder of host water $\delta^{18}\text{O}$ values (e.g. Epstein et al., 1977; DeNiro and Epstein, 1981; Sternberg, 1989, 2009). Furthermore, results from laboratory (Sauer et al., 2001) and field studies (Mayr et al., 2013) demonstrate convincingly that $\delta^{18}\text{O}$ of cellulose extracted from submerged aquatic mosses are highly correlated to their host waters owing to the absence of uncertainties related to evapotranspiration. However, achieving a high-resolution $\delta^{18}\text{O}_{\text{cell}}$ record could be impeded, because oxygen isotope analysis of moss cellulose requires large quantities of moss remains for cellulose extraction. An approach to tackle this problem is the isotope analyses of purified bulk organic matter (OM) of preserved aquatic moss shoots, which needs much less material, and can potentially improve the temporal resolution of paleoclimatic reconstructions based on the moss cellulose alone without losing paleoclimatic information (Zhu et al., 2014).

In a previous study, $\delta^{18}\text{O}_{\text{cell}}$ values of aquatic moss debris were used to infer $\delta^{18}\text{O}_{\text{lw}}$ of Laguna Potrok Aike over the last deglaciation (Mayr et al., 2013). In the present study, we used handpicked subfossil shoots of a single aquatic moss species from sediment sections covering the last glacial-interglacial transition period to generate a composite record of the $\delta^{18}\text{O}_{\text{lw}}$ inferred from purified bulk moss OM and extracted cellulose fractions. The aims of the study are: (1) to present a high-resolution $\delta^{18}\text{O}_{\text{lw}}$ record of Laguna Potrok Aike for the period containing large global climatic shifts by employing isotope proxies of aquatic mosses, (2) to highlight climatic changes on the southern South American continent during the last glacial-interglacial transition and (3) to evaluate the SHW impact on factors determining the $\delta^{18}\text{O}_{\text{lw}}$ of Laguna Potrok Aike.

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surface area and type, suggesting that isotope composition of the main source waters (precipitation and groundwater) are similar for all water bodies and evaporation is a main driver of the $\delta^{18}\text{O}_{\text{lw}}$ (Mayr et al., 2007).

Long-term isotopic data from the next GNIP (Global Network of Isotopes in Precipitation) station located on the leeward side of the southern Andes (south of 50°S) is available from Punta Arenas, located about 140 km southwest of Laguna Potrok Aike. For an observation period from 1990 to 2009, weighted monthly mean $\delta^{18}\text{O}$ values of precipitation at Punta Arenas station are positively correlated with monthly mean air temperatures ($R^2 = 0.89$) and exhibit an isotopic range of about 5‰ (IAEA/WMO, 2014; Fig. 3).

3 Material and methods

3.1 Material

In 2008 sediment cores were retrieved from two drilling sites in Laguna Potrok Aike within the framework of the PASADO project (Ohlendorf et al., 2011; Zolitschka et al., 2013; Fig. 1b). Sediment samples used in this study are from the composite profile 5022-2CP of site 2 which has a composite depth (cd) of 106 m, consisting of undisturbed pelagic sediments, volcanic tephra layers and mass movement sediments that resulted from lake internal sediment redistribution. The composite profile is divided into five lithological units based on the prevailing sedimentary structures and frequency of deposits of mass movement (Kliem et al., 2013b). Mass movement deposits and tephra layers were removed from the composite profile resulting in an event-corrected composite depth profile (cd-ec) of 45.8 m (Kliem et al., 2013b). The sediment section investigated in this study ranges between ca. 10 and 30 m (cd) or between 9.6 and 21.4 m (cd-ec) and consist of lithological unit B and C-1 (Fig. 3). Both unit B and C-1 mainly comprise pelagic laminated silts intercalated with thin fine sand and coarse silt layers originating from mass movement deposits. Pelagic silts are poorly laminated in

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There is a strong scatter between calibrated ages and event-corrected composite depth, particularly between 10–13.5 m cd-ec (Fig. 4). In order to obtain a reliable age-depth model, only the youngest ages were included in the age-depth model under the assumption that older than expected ^{14}C ages are the result of admixture of reworked old organic matter to the young counterparts. The main difference between the present and the previous age-depth model by Kliem et al. (2013b) is in the depth range between 12 and 15 m (cd-ec) where the calibrated ages derived from the present age-depth model are up to 1700 years younger than in the previous one (Fig. 4). We argue that the present age-model based on more AMS ^{14}C dates tends to be more reliable in this depth range based on a 1.5 m thick sediment section consisting of a multi-layered volcanic tephra bed at the depth from 16.8–18.2 m (cd) corresponding to the event-corrected composite depth of 14.7 m (cd-ec) (Wastegård et al., 2013). Chemical analyses indicated that this volcanic tephra is the R_1 tephra derived from volcano Reclús (Wastegård et al., 2013). Based on high-resolution dating of 1 mm peat layers immediately beneath the Reclús tephra layer at two sites at the Strait of Magellan, McCulloch et al. (2005) have provided a weighted pooled mean age of 12638 ± 60 ^{14}C years BP for the Reclús R_1 tephra. We recalibrated this ^{14}C age with CALIB 7.0 using SHCal13 (Hogg et al., 2013) and obtained a 2-sigma range of calibrated ages between 14 559 and 15 210 cal BP (Fig. 4). Stern (2008) and Sagredo et al. (2011) reported nearly the same ages. According to the new age-depth model, the depth of 14.7 m (cd-ec) has an age of 15 102 cal BP well within the reported age range of the Reclús R_1 tephra, whereas the previous model gives a considerably older age of 16 034 cal BP (Kliem et al., 2013b). The higher reliability of the present age-depth model is validated by the consistency with the independently dated tephra ages.

According to the new age-depth model the investigated sediment section covers the last glacial-interglacial transition from 26 000 to 8500 cal BP and ranges from the LGM to the early Holocene. The temporal boundary between lithological units B and C-1 is around 17 600 cal BP.

3.3 Laboratory methods

3.3.1 Isolation of moss remains

To acquire as much moss remains as possible, ca. 10 cm³ of sediments from every sample was screened. Each freeze-dried sediment sample was moistened with deionized water, placed on a magnetic stirrer and stirred for 2 h to disaggregate the material. Subsequently, the sample was carefully screened through a 200 μm sieve to obtain the coarse plant-debris fraction. The sieve fraction (> 200 μm) consists mainly of subfossil plant fragments such as shoots and leaves of mosses and remains of vascular plants, which were usually well preserved. Moss shoots were handpicked from the coarse sieve fraction under a binocular. To gain species-specific moss samples, we tried to pick only shoots of *D. perplicatus*. However, due to the similarity of the fragments of *D. perplicatus* and *V. pachyloma* and some not easily identifiable branches without leaves, an admixture of such moss fragments to the *D. perplicatus* samples cannot be ruled out. The remaining plant material in the coarse sieve fraction (> 200 μm) could contain fragments of *B. inundata*, *V. pachyloma* and other unidentifiable mosses and individual leaves of *D. perplicatus* as well as remains of aquatic and possibly terrestrial vascular plants and is termed as “residue” hereafter.

Each moss sample was first treated with a mixture of HCl and HF (10 % respectively) and left for 16 h at room temperature to completely remove attached carbonates and minerogenic components. Samples were then rinsed with deionized water three times to remove reagents and remaining clastic matter and freeze-dried. The cleaned moss samples were weighed and homogenized by cutting the moss branches into fine segments with scissors to avoid loss of fine moss material compared to milling. Bulk OM of moss branches was first analysed for $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values, before cellulose extraction was conducted. HCl-HF treatment of moss tissue prior to cellulose extraction has no effect on the $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values of cellulose (Zhu et al., 2014).

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3.3.2 Cellulose extraction

Cellulose was extracted from moss shoots and the residue fraction using the Cuprammonium solution (CUAM) method that has shown high reliability in yielding clean and pure cellulose from freshwater sediments, peat mosses and aquatic plants (Wissel et al., 2008; Moschen et al., 2009; Zhu et al., 2014). This method produces pure cellulose by dissolving and re-precipitating cellulose from whole plant material. Samples were first bleached with NaClO_2 (7 %) acidified with concentrated acetic acid (96 %) in a water bath for 10 h at 60 °C. The residual material was washed two times with hot deionized water (~ 70 °C) to remove the reagents and freeze-dried. The dry sample was mixed with ca. 30 mL CUAM solution (15 g L^{-1}) while placed in a dark room and stirred on a magnetic stirrer for 6 h and left for further 10 h at room temperature to completely dissolve the cellulose. After separation of not dissolved non-cellulose material, the cellulose solution was carefully decanted into a centrifuge tube and treated with 3 mL H_2SO_4 (20 %) for cellulose precipitation. The white precipitated cellulose was then rinsed three times with deionized water and freeze-dried.

3.3.3 Stable isotope measurements

For carbon isotope analyses, an amount of moss OM or cellulose equivalent to 100 μg of carbon was weighed into tin capsules. Samples were combusted at 1020 °C using an elemental analyser (Thermo Scientific Flash, 2000) interfaced on-line with an isotope ratio mass spectrometer (Thermo Scientific Delta V Advantage). Carbon content was determined by peak integration of mass-to-charge ratio (m/z) 44, 45 and 46, and calibrated against elemental standards. For oxygen isotope analyses, an amount of moss OM or cellulose providing 125 μg of oxygen was weighed into silver capsules. Immediately prior to oxygen isotope analysis, samples were placed overnight (16 h) in a vacuum drier at 100 °C to avoid analytical bias by adsorbed air moisture. Vacuum-dried samples were then pyrolysed at 1450 °C in a high temperature pyrolysis analyzer (HTO, HEKAtech) and measured on-line with a coupled isotope ratio mass

spectrometer (Micromass IsoPrime). Oxygen content was determined by peak integration of m/z 28, 29 and 30, and calibrated against elemental standards. Each sample was measured at least 2 times for both carbon and oxygen isotopes. Isotope ratios are expressed as δ -values in per mil (‰), where

$$\delta = (R_{\text{sample}}/R_{\text{standard}} - 1) \times 1000$$

with R_{sample} and R_{standard} as isotope ratios ($^{13}\text{C}/^{12}\text{C}$, $^{18}\text{O}/^{16}\text{O}$) of samples and standards, respectively. Isotope values are reported on the VPDB scale for carbon and the VSMOW scale for oxygen. Laboratory standards were inserted between samples to monitor the performance of the instrument and for calibration purposes. The standards USGS24 (-16.05‰), IAEA-CH-6 (-10.45‰) and IAEA-CH-7 (-32.15‰) were used for calibration of carbon isotope ratios of laboratory standards and samples, respectively (Coplen et al., 2006). The benzoic acid standards IAEA-601 ($23.14 \pm 0.19\text{‰}$) and IAEA-602 ($71.28 \pm 0.36\text{‰}$) (Brand et al., 2009) were used for calibration of oxygen isotope ratios of laboratory standards and samples, respectively. The overall precision of replicate analyses was better than $\pm 0.1\text{‰}$ for carbon and $\pm 0.3\text{‰}$ for oxygen isotope ratios. Ratios of carbon and oxygen content (C/O) of moss OM and cellulose were calculated on a weight base.

3.4 Reconstructed lake water $\delta^{18}\text{O}$ values

Modern field calibration datasets published in Zhu et al. (2014) were used to reconstruct the $\delta^{18}\text{O}_{\text{lw}}$ values from both bulk OM ($\delta^{18}\text{O}_{\text{OM}}$) and cellulose ($\delta^{18}\text{O}_{\text{cell}}$) of submerged aquatic mosses as well as the residue fraction applying the equations:

$$\delta^{18}\text{O}_{\text{lw}} = 1.156(\pm 0.036) \delta^{18}\text{O}_{\text{OM}} - 32.2(\pm 0.8); \quad (1)$$

$$\delta^{18}\text{O}_{\text{lw}} = 1.028(\pm 0.021) \delta^{18}\text{O}_{\text{cell}} - 30.4(\pm 0.5). \quad (2)$$

The uncertainty of the prediction (standard error of the regression) is 0.4‰ for Eq. (1) and 0.3‰ for Eq. (2), respectively (Zhu et al., 2014). The application of both equations

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2014). Nevertheless, some samples from lithological unit C-1 show around 1 ‰ more positive $\delta^{18}\text{O}_{\text{lw-corr}}$ values inferred from bulk moss OM compared to the moss cellulose reference line (1 : 1), while more positive $\delta^{18}\text{O}_{\text{lw-corr}}$ values inferred from residue cellulose are found for a couple of samples from lithological unit B. In terms of $\delta^{13}\text{C}$ values, bulk moss OM generally follows the moss cellulose reference line with an almost constant depletion. However, a marked bias towards more ^{13}C enriched values is observed for the samples of residue cellulose from lithological unit B, which indicates the presence of the remains of aquatic vascular plants in the residue fraction. According to Fig. 8 and Zhu et al. (2014), the observed positive $\delta^{13}\text{C}$ bias range of 2–4 ‰ suggests a 10–20 % contribution of aquatic vascular plants to the residue fraction, which results, however, only in a positive $\delta^{18}\text{O}$ bias of less than 0.2 ‰ which is well within the analytical uncertainty. Therefore, it is reliable to use the residue cellulose for an auxiliary $\delta^{18}\text{O}_{\text{lw}}$ reconstruction in the present study.

A composite $\delta^{18}\text{O}_{\text{lw-corr}}$ record based on aquatic moss shoots is constructed by the combination of bulk moss OM and moss cellulose applying moving average smoothing with a 500 years window (Fig. 9). The $\delta^{18}\text{O}_{\text{lw-corr}}$ record documents a mean $\delta^{18}\text{O}_{\text{lw-corr}}$ value of ca. -6.5‰ between 26 000 and 21 000 cal BP (Fig. 9). Subsequently, a $\delta^{18}\text{O}_{\text{lw-corr}}$ decrease of ca. 1 ‰ occurred between 21 000 and 17 600 cal BP and the minimum of the complete record of -7.5‰ was reached. From 17 600 till 12 800 cal BP, $\delta^{18}\text{O}_{\text{lw-corr}}$ strongly increased by an amplitude of nearly 3 ‰ interrupted by a millennial period with declining values of up to ca. 0.7 ‰ beginning at around 15 600 cal BP. Afterwards, the $\delta^{18}\text{O}_{\text{lw-corr}}$ values appeared to be subjected to millennial fluctuations and reached ultimately close to -3‰ in the early Holocene, similar with the present-day values (Fig. 9).

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5 Discussion

5.1 Factors controlling lake water $\delta^{18}\text{O}$ of Laguna Potrok Aike

Variations in $\delta^{18}\text{O}_{\text{lw}}$ are controlled by changes in the isotope composition of input waters (precipitation, surface inflow and groundwater inflow) and changes in the magnitude of subsequent evaporative ^{18}O enrichment (Edwards et al., 2004). It has been found that long-term temporal isotopic variation in precipitation at middle and high latitudes closely follows long-term changes in mean annual air temperature (Rozanski et al., 1992; Teranes and McKenzie, 2001; Darling et al., 2005). Under modern comparably stable conditions, seasonal or short-term variations of meteorological parameters and changes in $\delta^{18}\text{O}$ of precipitation ($\delta^{18}\text{O}_{\text{p}}$) do not have pronounced impacts on $\delta^{18}\text{O}_{\text{lw}}$ of Laguna Potrok Aike (Mayr et al., 2007). However, the highly significant positive correlation between monthly mean air temperature and weighted mean $\delta^{18}\text{O}_{\text{p}}$ at Punta Arenas, as shown in Fig. 3, indicates a potential influence of long-term local temperature changes on $\delta^{18}\text{O}_{\text{p}}$. Other than temperature change, $\delta^{18}\text{O}$ variations of precipitation can, however, also arise from changes in the direction of air masses bringing moisture to southern Patagonia. Precipitation brought from easterly directions is more enriched in heavy isotopes than those brought by westerly winds. The mean $\delta^{18}\text{O}_{\text{p}}$ of the former and the latter is -8 and -15‰ , respectively (Mayr et al., 2007). Thus, within a longer period with increasing air temperature and more frequent easterlies, the $\delta^{18}\text{O}_{\text{p}}$ and, in turn, $\delta^{18}\text{O}$ of inflow and $\delta^{18}\text{O}_{\text{lw}}$ could shift to more positive values. In addition, $\delta^{18}\text{O}_{\text{lw}}$ of lakes in semi-arid southern Patagonia are subjected to strong modification by evaporation, based on the fact that dry, extremely windy and highly evaporative conditions dominate the leeward side of the southern Andes (Garreaud et al., 2013). Today, the mean $\delta^{18}\text{O}$ of inflow (precipitation and groundwater) of Laguna Potrok Aike are around -13‰ , while $\delta^{18}\text{O}_{\text{lw}}$ values have a range between -3‰ and -4‰ (Mayr et al., 2007) indicating high evaporative ^{18}O enrichment of more than 9‰ relative to meteoric waters.

(i) $\delta^{18}\text{O}$ of meteoric water and groundwater markedly lower than present

All estimates of regional temperatures in southern Patagonia indicate a pronounced decrease during the last Glacial. Alkenone derived sea-surface temperatures (SST) from marine sediment cores off the Chilean coast (Fig. 10f) indicate lower SSTs by ca. 6°C for the last Glacial relative to the present (Lamy et al., 2007; Caniupan et al., 2011). For the South American continent, lower air temperatures by $8\text{--}10^\circ\text{C}$ during the LGM than today have been inferred from coupled ocean–atmosphere simulations (Rojas et al., 2009). Furthermore, Trombotto (2002) has suggested a lowering of the mean annual air temperature of at least 14°C in southern Patagonia during the LGM based on the presence of ice-wedge casts.

As discussed in Sect. 5.1, $\delta^{18}\text{O}_p$ is positively correlated with surface air temperature. A mean spatial gradient of $\delta^{18}\text{O}_p$ with surface air temperature of $0.53\text{‰}\cdot^\circ\text{C}^{-1}$ (Gourcy et al., 2005) or $0.58\text{‰}\cdot^\circ\text{C}^{-1}$ (Rozanski et al., 1993) has been reported. On the temporal scale, an average $\delta^{18}\text{O}_p$ -temperature coefficient of about $0.6\text{‰}\cdot^\circ\text{C}^{-1}$ is observed at mid- and high-latitudes (Rozanski et al., 1992). According to this relation, distinctly lowered temperatures would cause a strong ^{18}O depletion of precipitation in the order of 6‰ in southern Patagonia during the full Glacial compared to the present. If the groundwater flowing into Laguna Potrok Aike is mainly recharged by regional precipitation, the strong ^{18}O depletion of glacial precipitation would also have a direct impact on the $\delta^{18}\text{O}$ of inflow. Under these circumstances, $\delta^{18}\text{O}$ of surface and subsurface inflow into Laguna Potrok Aike during the full Glacial would be about -19‰ (present value: -13‰) assuming that the modern balance of precipitation from the Pacific and Atlantic was retained. This large ^{18}O depletion of inflow would result in an ^{18}O enrichment of about 12‰ between $\delta^{18}\text{O}_{\text{IW-corr}}$ (-6.5‰) recorded and $\delta^{18}\text{O}$ of inflow (-19‰) during the full Glacial compared to the modern magnitude of ^{18}O enrichment of about 9‰ (Mayr et al., 2007).

Today, climate in the south-eastern Patagonian steppe is characterized by strong westerly winds which are adiabatically warmed and dried while passing the Andes,

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leading to semi-arid and highly evaporative conditions in eastern Patagonia (Garreaud et al., 2013) that can explain the modern ^{18}O enrichment of lake water. Enrichment during the full Glacial might also have been caused by evaporation induced by a similar foehn-wind effect. It might have been strengthened by the thick Patagonian Ice Sheet covering the southern Andes which might have increased adiabatic warming and drying of subsiding air masses coming from westerly directions. This föhn-wind effect could be very pronounced in a cold and dry environment during the Glacial, which is corroborated by palynological studies of Laguna Potrok Aike sediments (Recasens et al., 2012). At Lake Hoare in the modern McMurdo Dry Valley of Antarctica strong and dry regional föhn-winds heat adiabatically by about 20°C (from -30 to -10°C) upon their descent from the surrounding ice plateau, even in sunless austral winters (Clow et al., 1988). At a mean annual temperature of less than -15°C in the Dry Valley region, relative humidity averages to only 0.54 and the annual sublimation (ablation) rate of surface ice of lakes reaches about 300 mm (Clow et al., 1988; Chinn, 1993). In a similar way, strong and extremely dry downslope föhn-winds passing the ice-covered southern Andes could have resulted in higher-than-expected evaporation and sublimation rates during the Glacial. Thus, isotopic enrichment of lake water during the full Glacial could have been stronger than expected.

This interpretation is largely based on the predominance of the SHW at the latitude of Laguna Potrok Aike (52°S) during the Glacial. The Patagonian Ice Sheet covering the southern Andes from 38 to 56°S during the LGM (Glasser et al., 2008) implies the existence of westerly winds within this latitudinal belt, because a positive mass balance of modern glaciers in the southern Andes is favored by low summer temperature and high precipitation and the latter is, in turn, largely related to the westerly winds from the Pacific (Schneider et al., 2003). In fact, paleoclimate studies from sites between 30 and 45°S in southwestern South America have implied much higher precipitation during the Glacial compared to the present (e.g. Heusser, 1989; Lamy et al., 1999; Moreno et al., 1999; Valero-Garcés et al., 2005).

(ii) Moderate change in $\delta^{18}\text{O}$ of source water compared to the present

If the SHW is located in a more equatorward position (Williams and Bryan, 2006), the balance between westerly and easterly winds would shift towards more easterly winds which could consequently dominate in southern Patagonia during the Glacial. Assuming almost 100 % precipitation moisture from the Atlantic, $\delta^{18}\text{O}_p$ could be roughly estimated for about -14‰ (cf. discussion above). In this case, $\delta^{18}\text{O}$ of inflow into Laguna Potrok Aike would be more positive than the estimation in scenario (i) and the magnitude of ^{18}O enrichment would be smaller accordingly.

If the glacial temperature in southern Patagonia was lowered by more than 10°C (cf. discussion in scenario (i)), the local mean annual temperature at Laguna Potrok Aike would be lower than -3°C during the Glacial and the formation of permafrost would be fostered. The occurrence of a relict sand wedge dated to 35 ± 3 ka in the Laguna Potrok Aike catchment area (Kliem et al., 2013a) indeed suggests permafrost conditions during the Glacial around the lake. Deep permafrost during the Glacial would have major impacts on the hydrological and isotopic water balance of Laguna Potrok Aike. Groundwater recharge from meteoric water may then have been precluded due to impervious permafrost layers. Hence, any isotopic change in the precipitation may not have been transmitted into the groundwater. Decreased precipitation and then limited surface and subsurface inflows during the full Glacial would also generally make a smaller contribution to lake water budget than today. Thus, the expected large negative shift in $\delta^{18}\text{O}$ of inflow, as discussed in scenario (i), may not have occurred. In addition, deep permafrost could have largely prohibited the exchange between the groundwater and the lake water body (subsurface in- and outflow), thus converting Laguna Potrok Aike into a closed lake system with extremely prolonged lake water residence time under non-overflow conditions. Under these circumstances, even small evaporative isotopic enrichment effects could sum up to considerable cumulative ^{18}O enrichment of lake water.

lowering lake level that reached a depth of -33 m below the modern one shortly before 6790 cal BP (Haberzettl et al., 2008; Anselmetti et al., 2009; Zolitschka et al., 2013).

Like the patterns of changes in the SHW during the LGM, for the period of the last deglaciation towards the early Holocene, the reconstructed development of the SHW based on the paleoclimatic sites in southern Patagonia remains controversial (cf. Kilian and Lamy, 2012; Villa-Martínez et al., 2012).

6 Conclusions

This study presents a high-resolution $\delta^{18}\text{O}_{\text{lw}}$ reconstruction for Laguna Potrok Aike located in semi-arid southern Patagonia throughout the last glacial-interglacial transition by using purified bulk OM and extracted cellulose of subfossil submerged aquatic mosses. These data provide a unique continental proxy record of the environmental development in the high southern latitudes during this period of fundamental climatic shifts. The temporal evolution of $\delta^{18}\text{O}_{\text{lw-corr}}$ of Laguna Potrok Aike is largely controlled by changes in $\delta^{18}\text{O}$ of the source water of lake, surface air temperature and evaporative ^{18}O enrichment.

Considering the occurrence of permafrost during the Glacial, the $\delta^{18}\text{O}_{\text{lw-corr}}$ record between 26 000 and 21 000 cal BP is best explained by a main water supply from isolated groundwater whose $\delta^{18}\text{O}$ may not be as depleted as that of glacial meteoric water. Moreover, probably reduced interchange between in- and outflows and generally decreased inflows would have prolonged the lake water residence time. Under these circumstances, the higher than expected $\delta^{18}\text{O}_{\text{lw-corr}}$ during this period could be achieved despite possibly weakened evaporation under glacial conditions. Between 21 000 and 17 600 cal BP, coinciding with the timing of a reconstructed lake level overflow of Laguna Potrok Aike, large ^{18}O depletion in $\delta^{18}\text{O}_{\text{lw-corr}}$ record is observed. Low $\delta^{18}\text{O}_{\text{lw-corr}}$ together with the overflow situation could be linked to an increased

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Table 1. AMS ^{14}C ages for the modeled event-corrected sediment depth in the range of 9.37–26.48 m (cd-ec) for 5022–2CP of Laguna Potrok Aike. All ^{14}C ages derive from samples collected in pelagic sediment sections. Ranges of calibrated ages (at 95 % confidence intervals, 2 s) are the output of age-modeling software *clam 2.2* (Blaauw, 2010) applying the SHCal13 calibration curve (Hogg et al., 2013) and smoothed spline with a smoothing level of 0.5. Accepted ^{14}C ages are shown in bold.

Lab. No. ^d	Sediment depth (m cd)	Event corrected sediment depth (m cd-ec)	^{14}C Age (BP)	Error (±)	$\delta^{13}\text{C}$ (‰)	C-mass (mg)	Sample description	Range of calibrated ages (2 s)	Median probability
Poz-8392 ^a	9.69	9.37	7580	50	-28.3	2.56	Stems of aquatic moss	8203–8421	8355
Poz-48915	10.81	10.35	9390	90	-28.7	1.74	Bulk aquatic moss tissues	10254–10784	10557
AA93659	10.95	10.49	11379	57	-26.6	–	Bulk sediment	13079–13292	13185
AA93660	12.22	11.52	12200	200	-29.4	–	Wood, plant fragments	13574–14902	14115
AA93661	12.99	12.18	14042	70	-25.3	–	Bulk sediment	16674–17276	17000
Poz-5985^a	13.04	12.22	8930	50	-18.9	2.28	Bone of Tuco Tuco	9780–10188	10016
AA93662	14.06	12.34	16101	84	-25.5	–	Wood, transparent shell fragments	19122–19612	19378
Poz-48917	14.08	12.36	16360	90	-29.3	1.18	Bulk aquatic moss tissues	19476–19980	19704
Poz-49760	14.37	12.66	19380	100	-27.3	1.77	Bulk aquatic moss tissues	22983–23581	23284
Poz-8548 ^a	14.78	13.00	10240	60	8.4	3.61	Calcite fraction of bulk sample	11611–12067	11872
Poz-48918	15.07	13.30	17460	100	-28.5	3.14	Bulk aquatic moss tissues	20717–21355	21026
Poz-49761	15.37	13.60	11490	60	-25.1	1.50	Bulk aquatic moss tissues	13147–13428	13291
Poz-8396 ^a	15.55	13.78	11200	60	-30.0	1.69	Stems of aquatic moss	12831–13130	13023
Poz-48919	15.73	13.95	12050	70	-31.6	1.41	Bulk aquatic moss tissues	13712–14089	13868
Poz-49763	15.87	14.08	10840	60	-28.8	2.03	Bulk aquatic moss tissues	12654–12790	12711
Poz-48920	15.95	14.18	11120	70	-33.1	2.97	Bulk aquatic moss tissues	12771–13081	12935
AA93664	16.07	14.30	10980	140	-29.7	–	Wood, seeds, plant fragments	12663–13082	12847
Poz-49764	16.15	14.38	12040	60	-27.4	1.70	Bulk aquatic moss tissues	13719–14054	13856
Poz-8397^a	16.40	14.81	12490	70	-31.2	1.60	Stems of aquatic moss	14198–15001	14583
Poz-49765	16.42	14.65	12590	60	-27.4	1.52	Bulk aquatic moss tissues	14409–15139	14844
Poz-5072^a	16.48	14.70	12850	70	-25.8	2.64	Stems of aquatic moss	15068–15575	15267
Poz-49022	18.28	14.73	12720	70	-29.2	2.43	Bulk aquatic moss tissues	14753–15304	15082
AA93666	18.28	14.73	12783	64	-27.3	–	Plant fragments (large in quality)	14906–15415	15179
Poz-48922	18.40	14.85	13530	70	-27.6	1.82	Bulk aquatic moss tissues	14409–15139	16235
Poz-5073^a	18.51	14.96	13450	70	-28.7	2.69	Stems of aquatic moss	15881–16354	16132
Poz-48923	18.67	15.11	14540	80	-29.3	1.58	Bulk aquatic moss tissues	17450–17913	17671
Poz-37017^b	18.69	15.13	14540	70	-27.6	1.56	Stems of aquatic moss	17465–17900	17672
Poz-48925	21.13	15.94	16150	80	-24.0	1.30	Bulk aquatic moss tissues	19190–19662	19435
Poz-37022^b	22.09	16.54	17460	80	-29.2	1.63	Stems of aquatic moss	20760–21317	21023
AA93669	22.57	17.23	18850	170	-26.3	–	Bulk sediment	22363–23080	22686
Poz-37007^b	23.25	17.70	18700	120	-39.9	0.91	Stems of aquatic moss	22304–22852	22527
AA93670	25.01	18.93	20600	270	-26.0	–	Bulk sediment	24118–25442	24762
Poz-37020^b	27.20	19.50	20490	120	-28.0	1.11	Stems of aquatic moss	24242–25045	24604
Poz-34236^{b,c}	36.38	26.48	25110	180	-25.0	1.45	Stems of aquatic moss	28713–29531	29108

^a Haberzettl et al. (2007).

^b Kliem et al. (2013b).

^c Not shown in Fig. 4, but serving as connection point with previous age-depth by Kliem et al. (2013b).

^d Poz: Poznan Radiocarbon Laboratory; AA: NSF-Arizona AMS Laboratory.

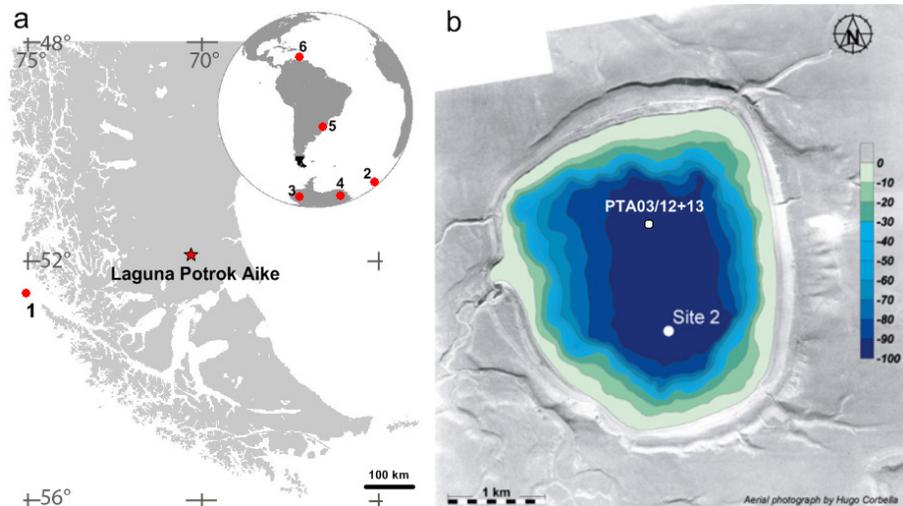


Figure 1. (a) Location of Laguna Potrok Aike (red star) in southern Patagonia indicated as a black area on the inserted map. Location of sites presented in Fig. 10: 1, MD07-3128 at the Chilean offshore (Caniupán et al., 2011); 2, TN057-13-4PC in the Southern Atlantic (Anderson et al., 2009); 3, WDC in West Antarctica (WAIS Divide Project Members, 2013); 4, EDML of East Antarctica (EPICA Community Members, 2006); 5, Botuverá Cave in Southern Brazil (Wang et al., 2007); 6, Cariaco Basin (Deplazes et al., 2013). (b) Sediment samples investigated in the present study derive from the drilling site 2 shown on the bathymetric map of Laguna Potrok Aike inserted into an aerial photograph (provided by Hugo Corbella). At site 2, hydraulic piston cores were taken in 2008 within the framework of PASADO. The piston core PTA03/12+13 taken in 2003 has been used for the reconstruction of oxygen isotope composition of lake water in Mayr et al. (2013).

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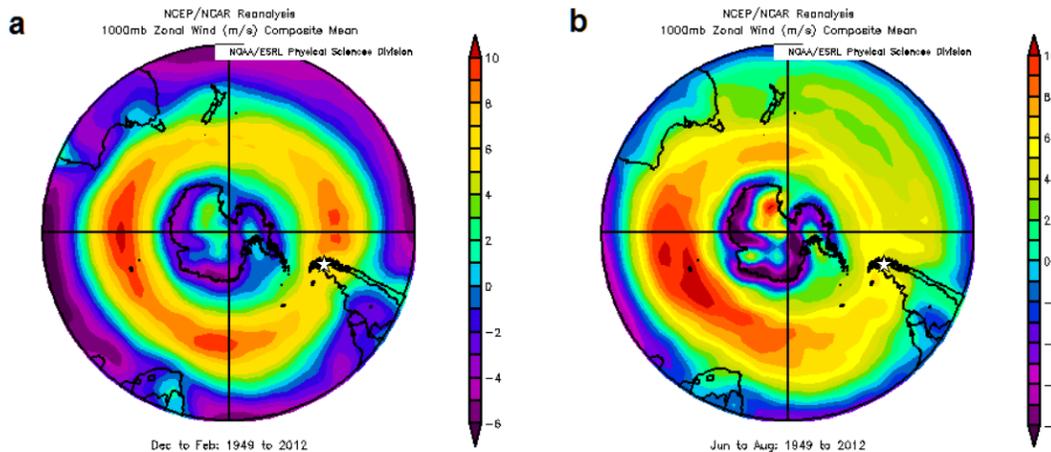


Figure 2. Mean near-surface (1000 mb) zonal wind (m s^{-1}) in Southern Hemisphere for austral summer (a) and winter months (b) based on NCEP/NCAR Reanalysis. Location of Laguna Potrok Aike is indicated by white stars. Data source: <http://www.esrl.noaa.gov/psd/cgi-bin/data/composites/printpage.pl>, accessed on 6 February 2014.

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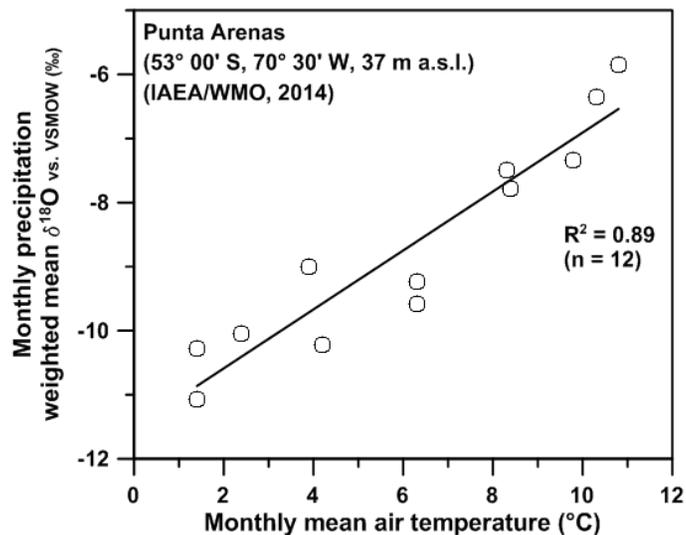


Figure 3. Relationship between monthly mean air temperature and weighted mean $\delta^{18}\text{O}$ of precipitation at GNIP station Punta Arenas (53°00' S, 70°30' W, 37 m a.s.l.) for the period from 1990 to 2009 (IAEA/WMO, 2014).

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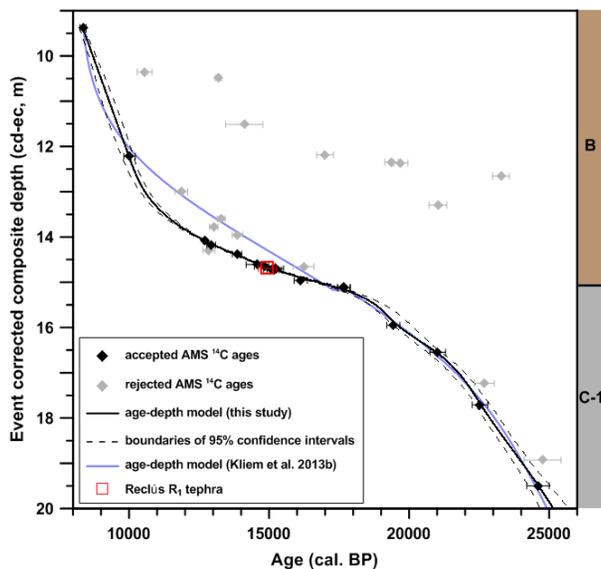


Figure 4. Age-depth model for the sediment section between 9 and 20 m event-corrected composite depth (cd-ec) from the composite profile 5022-2CP of Laguna Potrok Aike (cf. Table 1 for details). The age-depth model used in the present study is shown as a black line which is constructed by clam 2.2 applying a smooth spline with a smoothing level of 0.5 (Blaauw, 2010). Dashed lines represent the upper and lower boundary of 95 % confidence intervals. The accepted AMS ^{14}C ages are shown as black diamonds and the rejected ones in grey. Error bars represent the range of calibrated ages at 95 % confidence intervals. The previous age-depth model by Kliem et al. (2013b) is given as a blue line. The red open square represents the depth of the Reclús R_1 tephra. Its ^{14}C age from McCulloch et al. (2005) is recalibrated by CALIB 7.0 using SHCal13 (Hogg et al., 2013). Error bars are given for 2 sigma age range. Two lithological units (cf. text for detail) occurring in the investigating depth range are shown on the right.

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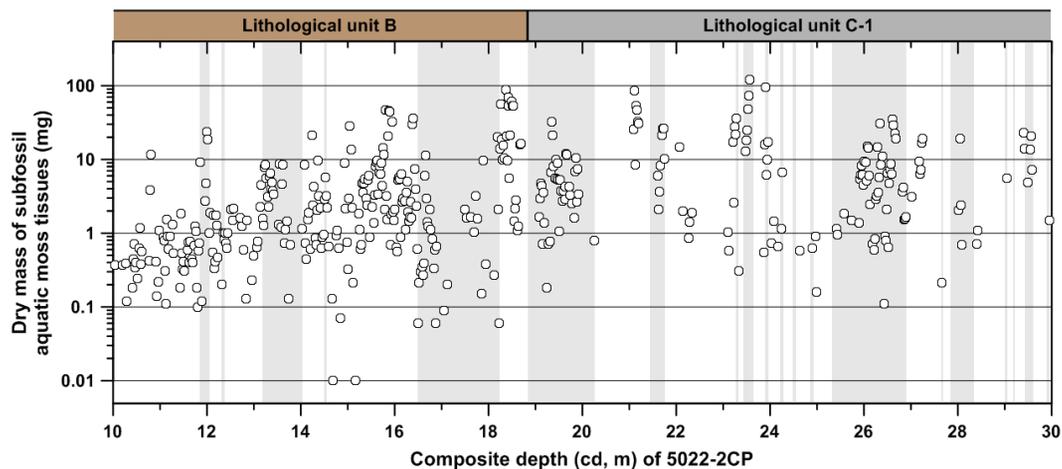


Figure 5. Weighed dry mass of subfossil aquatic moss remains handpicked from sediment samples within the investigated composite depth range of 5022-2CP of Laguna Potrok Aike. Vertical grey bars represent mass movement deposits and volcanic ash layers. Two lithological units (cf. text for detail) occurring in the investigating depth range are shown at the top of the figure. Note that the y axis is in log-scale.

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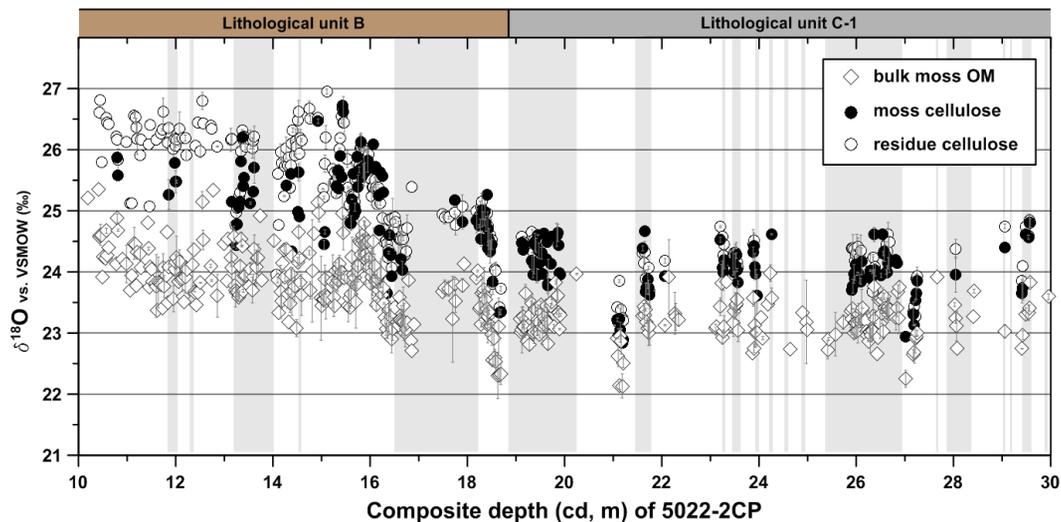


Figure 6. $\delta^{18}\text{O}$ values of all measured samples within the investigated composite depth range of 5022-2CP of Laguna Potrok Aike. Bulk aquatic moss organic matter (OM) is represented by open diamonds, aquatic moss cellulose by closed circles and residue cellulose by open circles. Standard deviations are shown as bars. Vertical grey bars represent mass movement sediment sections and volcanic ash layers. Two lithological units (cf. text for detail) occurring in the investigating depth range are shown at the top of the figure.

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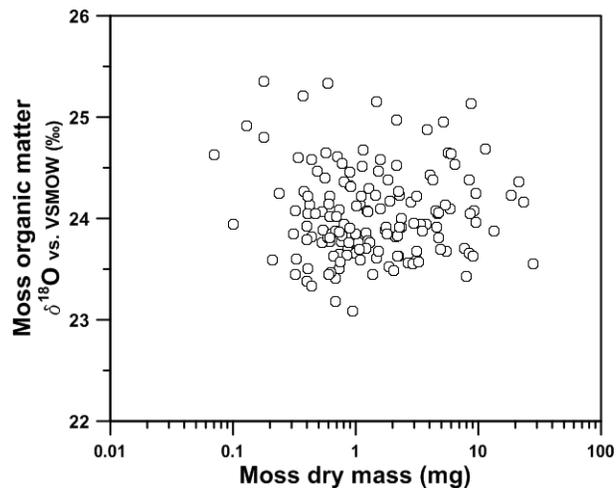


Figure 7. Relationship between dry mass of handpicked *Drepanocladus perplicatus* and $\delta^{18}\text{O}$ values of moss organic matter for samples within the composite depth between 10 and 16 m. Note that the x axis is in log-scale.

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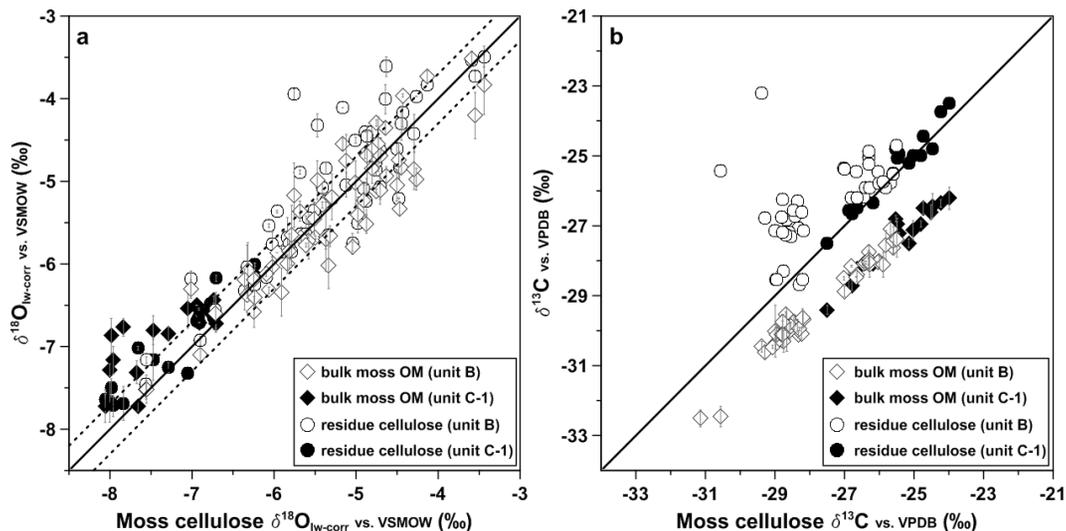


Figure 8. (a) Reconstructed lake-water $\delta^{18}\text{O}$ ($\delta^{18}\text{O}_{\text{lw-corr}}$) from bulk aquatic moss organic matter (OM) (diamonds) and residue cellulose (circles) in relation to $\delta^{18}\text{O}_{\text{lw-corr}}$ values reconstructed from aquatic moss cellulose. Samples from the sediment sections of mass movement deposits and tephra layers are excluded. Samples from lithological unit B and C-1 are shown in open and closed symbols, respectively. One-to-one line of $\delta^{18}\text{O}_{\text{lw}}$ values reconstructed from aquatic moss cellulose is presented as a black line. (b) Same as (a), but for $\delta^{13}\text{C}$ values. Lower and upper limit of one-to-one line in (a) are presented in dashed lines according to the standard error of regression for modern calibration data set (Zhu et al., 2014). Standard deviations of individual values are given as bars in (a) and (b).

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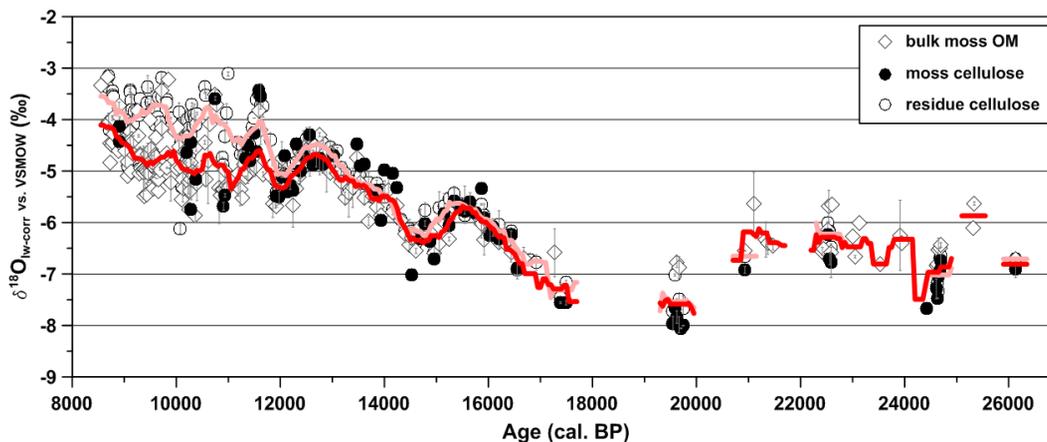


Figure 9. Isotopic record of reconstructed lake-water $\delta^{18}\text{O}$ ($\delta^{18}\text{O}_{\text{lw-corr}}$) from bulk aquatic moss organic matter (OM), aquatic moss cellulose and residue cellulose during the last Glacial-Interglacial transition period. Samples from the sediment sections of mass movement deposits and tephra layers are excluded. Color lines represent the moving average smoothing using a 500 years window (red: smoothing of average $\delta^{18}\text{O}_{\text{lw-corr}}$ of composite aquatic moss record combining moss OM with moss cellulose; pale red: smoothing of $\delta^{18}\text{O}_{\text{lw-corr}}$ of residue cellulose). Discontinuity prior to 17 600 cal BP is caused by insufficient moss material.

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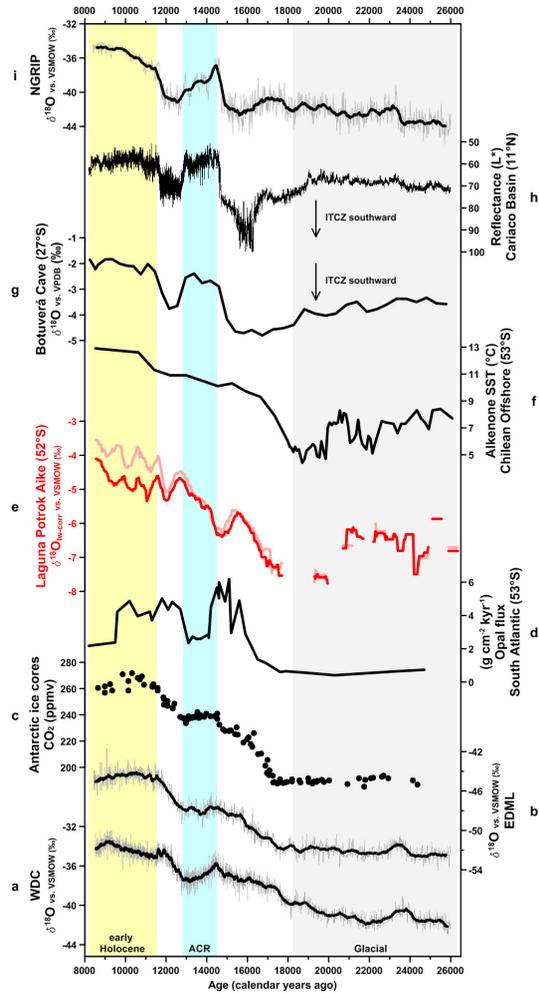
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Figure 10. Reconstructed lake-water $\delta^{18}\text{O}$ of Laguna Potrok Aike in comparison to global proxy records. **(a)** $\delta^{18}\text{O}$ record from the WDC in West Antarctica (WAIS Divide Project Members, 2013). **(b)** $\delta^{18}\text{O}$ record from EDML of East Antarctica (EPICA Community Members, 2006). **(c)** CO_2 concentration from Antarctic ice cores (Schmitt et al., 2012). **(d)** Opal flux of TN057-13-4PC (53° S) in the Southern Atlantic (Anderson et al., 2009). **(e)** Reconstructed lake-water $\delta^{18}\text{O}$ ($\delta^{18}\text{O}_{\text{lw-corr}}$) of Laguna Potrok Aike in this study, smoothed by a 500 years window (red: smoothing of average $\delta^{18}\text{O}_{\text{lw-corr}}$ of composite aquatic moss record combining moss cellulose with moss OM; pale red: smoothing of $\delta^{18}\text{O}_{\text{lw-corr}}$ of residue cellulose). Ocean water effect during the LGM and deglaciation has been corrected according to Lea et al. (2002). **(f)** Alkenone derived SST record from the offshore core MD07-3128 (53° S) (Caniupán et al., 2011). **(g)** $\delta^{18}\text{O}$ record of Botuverá Cave (27° S) in Southern Brazil (indicator of the ITCZ position) (Wang et al., 2007). **(h)** Sediment total reflectance from Cariaco Basin (indicator of the ITCZ position) (Deplazes et al., 2013). **(i)** $\delta^{18}\text{O}$ record of NGRIP on GICC05 time scale (Anderson et al., 2006; Rasmussen et al., 2006). ACR: Antarctic Cold Reversal.

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