





Stable Southern Hemisphere westerly winds throughout the Holocene until intensification in the last two millennia

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The Southern Hemisphere westerly winds sustain the Southern Ocean's role as one of Earth's main carbon sinks, and have helped sequester nearly half the anthropogenic CO₂ stored in the ocean. Observations show shifts in the vigor of this climate regulator, but models disagree how future change impacts carbon storage due to scarce baseline data. Here, we use the hydrogen isotope ratios of sedimentary lipids to resolve Holocene changes in Southern Hemisphere westerly wind strength. Our reconstruction reveals stable values until ~2150 years ago when aquatic compounds became more ²H-enriched. We attribute this isotope excursion to wind-driven lake water evaporation, and regional paleoclimate evidence shows it marks a trend towards a negative Southern Annular Mode – the Southern Ocean's main mode of atmospheric variability. Because this shift is unmatched in the past 7000 years, our findings suggest that previously published millennium-long Southern Annular Mode indices used to benchmark future change may not capture the full range of natural variability.

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The Southern Ocean covers 20% of the Earth's surface and has stored more than 40% of all anthropogenic carbon emissions sequestered by the oceans since the industrial revolution¹, while absorbing around 75% of the excess heat generated during this period². This outsized role as a global climate regulator is attributed to the Southern Hemisphere westerly winds (SHW), which draw up deep water and mix it while encircling the Antarctic³. The vigor of the Southern Ocean's role as a carbon pump is modulated by the opposing effects of CO₂ uptake and outgassing at the sea surface⁴. This balance is strongly influenced by variations in the strength and position of the SHW⁵. These, in turn, are largely driven by the Southern annular mode (SAM)⁶: the main pattern of atmospheric climate variability over the Southern Ocean, which is typically described as the pressure difference between mid and high austral latitudes⁷. The SHW belt shifts poleward in the positive SAM polarity and expands equatorward during a negative SAM phasing⁸.

Instrumental data reveal an increasingly positive SAM phasing and associated poleward SHW shift over the past decades^{8,9}, which has already intensified regional hydroclimate extremes¹⁰. While this recent (last ~50 years) trend is primarily attributed to a combination of anthropogenic ozone depletion and greenhouse gas emissions¹¹, recent studies indicate that natural SAM variability is substantial and cannot be explained by direct radiative forcing mechanisms alone^{12,13}. These findings have drawn further attention to observational evidence for SAM modulation by other internal modes of climate change, most notably the El Niño-Southern Oscillation (ENSO)^{14,15}. However, while such teleconnections can modify the impact of anthropogenic forcing on coupled SHW-SAM change¹⁶, their dynamics remain poorly documented and incompletely understood¹⁷. This knowledge gap mainly stems from the scarcity and brevity of baseline data⁸: our knowledge of past SAM variability does not extend beyond the last millennium, and only two records span that entire period^{12,13}. Longer paleoclimate timeseries are imperative to resolve the full range of natural variability and better represent future SHW-SAM behavior and its impact on atmospheric CO₂ concentrations^{5,18}.

Here, we present a record of centennial-scale variations in SHW strength over the past 7 ka based on lake sediments. To do so, we investigated a lacustrine sediment sequence from South Georgia (54.4°S and 36.6°W), a sub-Antarctic Island in the SHW core belt (Fig. 1a). To avoid the uncertainties (for example, lags or non-linearity) of wind-attributed feedbacks like changes in precipitation or temperature and their impact on vegetation¹⁹, we rely on a less indirect measure of wind strength: sedimentary leaf wax hydrogen isotope ratios ($\delta^2\text{H}$). Modern observations help constrain the climatic controls on the $\delta^2\text{H}$ values of source water used by plants in South Georgia. Compound-specific analysis reveals that the isotopic composition of lake water (recorded by aquatic plants) became ²H-enriched compared to its meteoric source water (recorded by terrestrial plants). The direction and magnitude of this isotopic offset is indicative of evaporative water loss²⁰, a process that is typically wind-driven in similar closed (endorheic) basins in the SHW core belt²¹.

Setting and approach

South Georgia (54.4°S and 36.6°W) is one of the few landmasses that is located year-round in the SHW core belt and, therefore, well-situated to detect the fingerprints of past variations in this globally relevant circulation system (see introduction). As can be seen in Fig. 1a, this belt contracts in summer and expands during winter. On longer timescales, SAM variations can cause similar latitudinal SHW shifts in response to changes in the meridional pressure gradient between 45°S and 60°S⁷. While we should note

that the relation between SAM and regional climate depends on the choice of index and data due to a lack of observations²², the station-based Marshall SAM index reveals a negative correlation with re-analyzed wind stress on South Georgia for the period prior to 1979²³—when SAM variability is attributed to natural variations captured by proxy records like ours²⁴. Numerous studies have also linked SAM-related shifts in temperature and hydroclimate to island-wide changes in glacier extent or ecosystem dynamics^{25–32}. Local weather observations from Grytviken (Fig. 1a) reveal a maritime subpolar climate with an average annual temperature of 2 °C and 1450 mm of evenly distributed yearly precipitation²³ (Fig. 1c).

Our study site—informally named Lake Diamond²⁹—is located in the north-central part of South Georgia, in the Olsen Valley that abuts the Cumberland West Bay (Fig. 1b). The basin covers an area of ~0.1 km², sits at ~15 m a.s.l and has a maximum depth of 3.5 m. Vegetation is restricted to areas below 100 m a.s.l (Fig. 1b) and is dominated by tussock grass (*Festuca contracta*) and greater burnet (*Acaena magellanica*). Based on reconstructed glacial sediment fluxes, previous studies showed that meltwater from the up-valley Diamond glacier incidentally spilled into the lake during flood events over the past 7 ka^{29,32}. Timeseries analysis of Sentinel-2 satellite imagery reveals that this has not happened since image acquisition commenced in 2015³³. In the absence of in- or outlets, we consider the basin closed (endorheic): water is thus primarily sourced from precipitation falling in the ~1 km² Lake Diamond catchment (Fig. 1b). Satellite imagery also shows that our study site is typically snow- and ice-covered for 6 months and that ice-off occurs between late October and early November.

We present lipid biomarker analyses from Lake Diamond sediments that were previously studied by²⁹. The investigated sequences consist of 46 cm long surface core LDS2 and 246 cm piston core LDP2: based on a robust published radiocarbon chronology ($n = 24$), this record covers the past 10 ka. We restricted our analyses to the past 7.2 ka, when the Diamond basin was occupied by a lake. This interval is contained in the uppermost 221.5 cm of the composite record and has previously been described as units D and F by ref. 29.

To reconstruct sub-Antarctic surface climate change during this period, and assess its links to SHW core belt shifts, hydrogen isotope ratios ($\delta^2\text{H}$) were measured on plant leaf wax samples ($n = 56$) from Lake Diamond sediments (see methods). For this purpose, we extracted, separated, quantified, and analyzed *n*-alkanes: aliphatic hydrocarbons produced by plants and preserved in sedimentary archives around the world³⁴. $\delta^2\text{H}$ values of these lipid biomarkers track the $\delta^2\text{H}$ values of the source water used during biosynthesis^{35–37}.

Isotope values of precipitation are influenced by a range of parameters, including condensation temperature, evaporation, precipitation amount, and the source and pathway of moisture³⁸: near South Georgia, all of these variables co-vary with SHW change^{12,39–41}. Weather station data from nearby Grytviken (~10 km away) and interpolated precipitation isotopes reveal a strong positive ($R^2 = 0.92$, $p < 0$) relationship between modern (1905–2019 CE) surface temperatures and meteoric $\delta^2\text{H}$ (Figs. 1c, 2b)^{23,42}. Based on observations and ice core data from James Ross Island (Fig. 1a)^{43,44}, this relationship could have existed throughout the past 7 ka. The isotope composition ($\mu = -60.3\text{‰}$, $2\sigma = 1.1\text{‰}$) of two modern samples furthermore indicates that lake water $\delta^2\text{H}$ values approximate the weighted local annual precipitation mean (-67.5‰), despite collection during the austral summer (February 2015). Figure 2a shows that evaporative loss of lighter ¹⁶O and ¹H has a minor impact on modern lake water composition as our water samples plot just below the Local Meteoric Water Line (LMWL), constructed with data from

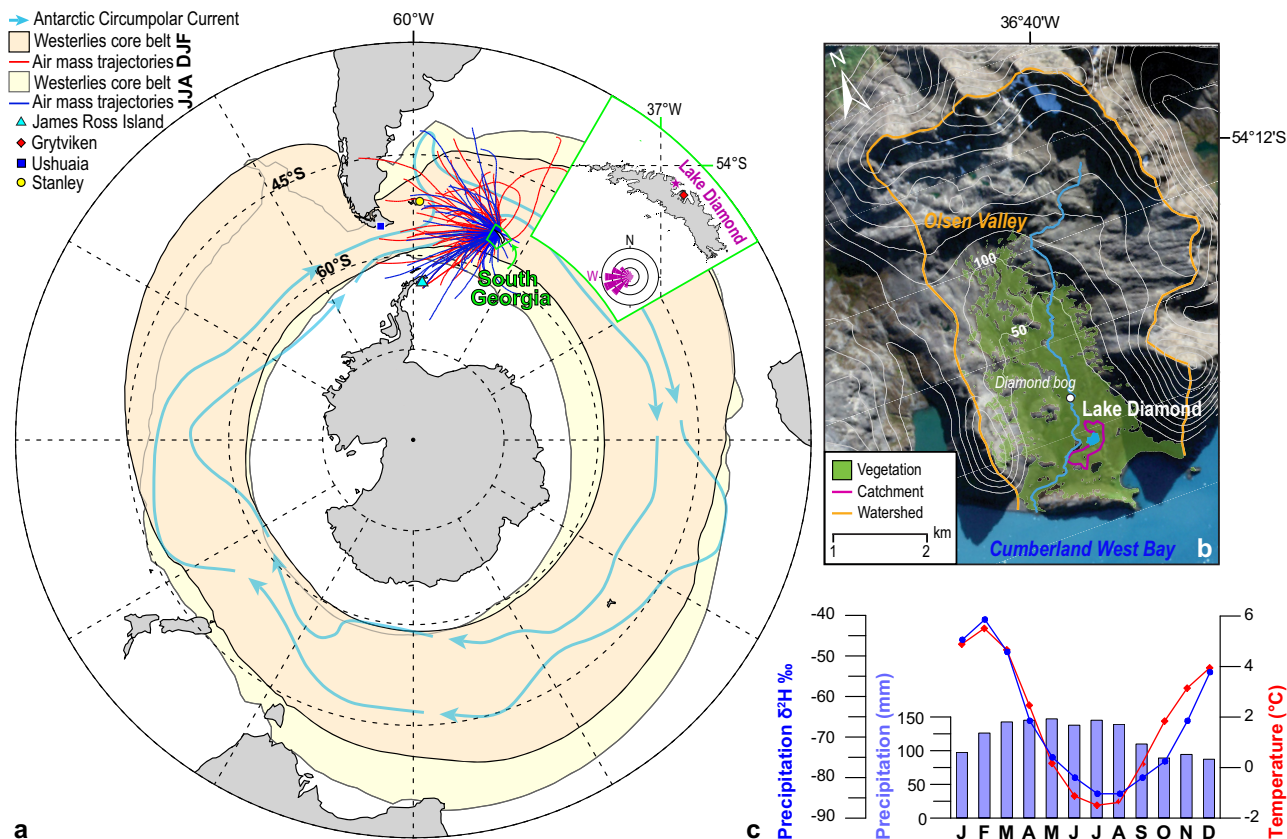


Fig. 1 Key localities and observations. **a** Overview map of the Southern Ocean with an inset of South Georgia, highlighting Lake Diamond as well as the nearest weather stations of Grytviken, Ushuaia, and Stanley in color coding that corresponds with Fig. 2. The average position and extent of the Southern Westerlies core belt, defined as areas where mean zonal wind speeds exceed 5 m/s^{-1} between 1981–2010 CE¹²², is shown for the Austral summer (DJF) and winter (JJA). The purple wind rose shows the prevalence of this dominant wind system. Forty-eight hours daily backward air mass trajectories from 2019 are shown for summer (red) and winter (blue)¹²³. The long-term position of the Antarctic Circumpolar Current (ACC) is delimited in cyan by the Antarctic convergence to the South and the Sub-Antarctic front to the North after¹²⁴. **b** Close-up of the Lake Diamond catchment and the wider Olsen valley as modified after²⁹. Vegetated areas, defined as surfaces with normalized difference vegetation index (NDVI) values >0.3 during the growing season (January 2020)³³, are highlighted in green. Sentinel-2 satellite image courtesy of the ESA Copernicus Open Access Hub. **c** Long-term monthly local climatology based on 1905–2012 CE station data from nearby Grytviken²³ and spatial $\delta^2\text{H}$ interpolation⁴².

the nearest GNIP stations at Ushuaia ($\sim 2000 \text{ km}$ away) and Stanley ($\sim 1500 \text{ km}$ away)⁴⁵ (Fig. 1a).

Results and discussion

Holocene lipid $\delta^2\text{H}$ records lake water evaporation. Because modern meteoric $\delta^2\text{H}$ correlates with seasonal temperature change in South Georgia, and Lake Diamond water $\delta^2\text{H}$ integrates a somewhat enriched annual precipitation average (see Figs. 1c, 2), we expect reconstructed values to capture changes in temperature and evaporative ^2H -enrichment. To better constrain this paleo signal, we disentangle the different plant sources producing our sedimentary n -alkanes and the environmental signals they record in the text below.

While aquatic photoautotrophs directly use lake water for biosynthesis³⁵, terrestrial plants use soil moisture³⁵, which is often isotopically skewed towards the growing season when biosynthesis occurs⁴⁶ and differently impacted by evaporative enrichment than lake water^{47,48}. This is particularly true for closed (endorheic) basins like Lake Diamond due to their long water residence time²⁰. To constrain these different $\delta^2\text{H}$ source signatures, we differentiate between aquatic and terrestrial n -alkane producers. It has previously been argued that mid-chain (C_{21} – C_{25}) n -alkane homologs are produced by aquatic plants, whereas long-chain (C_{27} – C_{33}) compounds typically derive from

terrestrial vegetation^{49,50}. Recent studies have called this generalization into question^{51–53}. Here, we rely on reported n -alkane distributions from representative terrestrial vegetation at nearby Grytviken to understand the sources of n -alkanes in Lake Diamond (Fig. 1a)⁵⁴. Critically, this selection comprises two species that dominate the Lake Diamond catchment today: tussock grass and greater burnet (see Fig. 3a). As can be seen in Fig. 4, mid-chain homologs are negligible in these terrestrial plants but abundant in sediment samples. This is in line with evidence from upwind southern Patagonia, where these compounds are attributed to aquatic macrophytes⁵⁵. We thus argue that mid-chain n -alkanes are primarily synthesized by aquatic producers in Lake Diamond. In this respect, we note that moss patches in the catchment only contain traces of *Sphagnum*³²—the only peat-forming species synthesizing considerable amounts of mid-chain n -alkanes^{56,57}. The inference that mid- and long-chain homologs derive from different sources is supported by Principal Component Analysis (PCA: see methods). The ordination diagram of Fig. 5c separates these homologous groups on PC2. Based on these lines of reasoning, we infer that mid- and long-chain n -alkanes derive from aquatic and terrestrial plants, respectively.

Down-core n -alkane $\delta^2\text{H}$ values can also be affected by vegetation shifts because of species-specific differences in apparent fractionation between source water and leaf

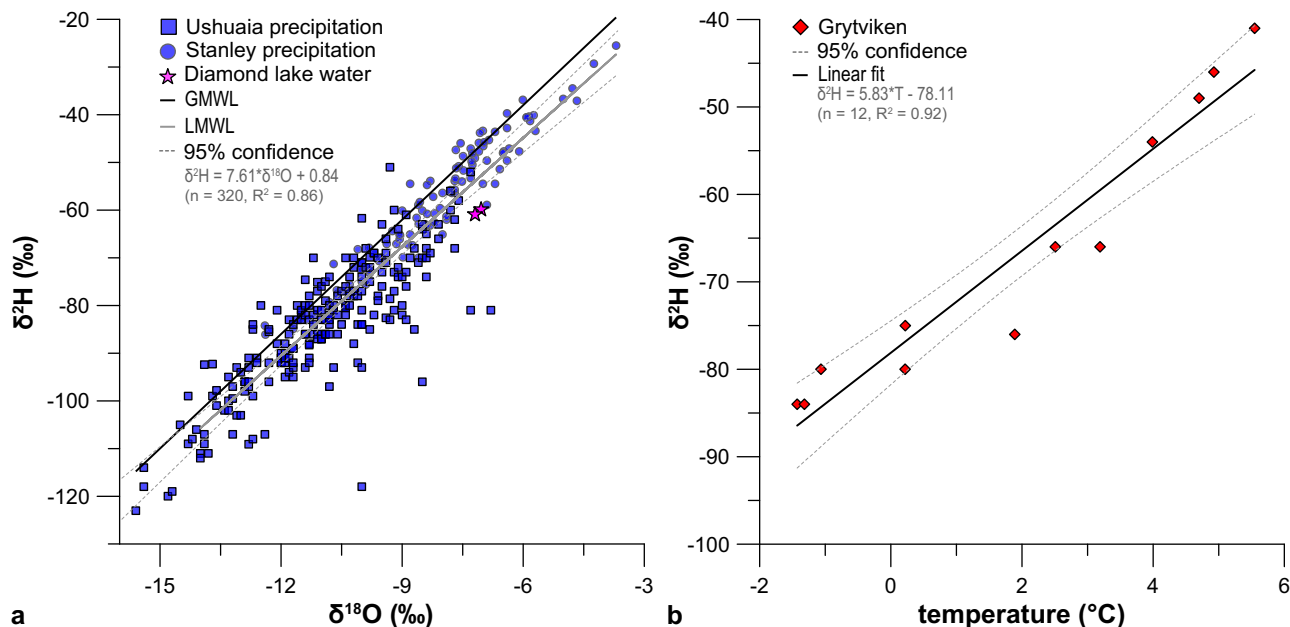


Fig. 2 Isotope characteristics of meteoric and lake water. **a** Relationships between source water precipitation hydrogen isotope values ($\delta^2\text{H}$) and oxygen isotopes ($\delta^{18}\text{O}$) in samples collected between 1965–2016 CE from the most proximal weather stations in Stanley and Ushuaia along the global (GMWL: black) and local (LMWL: gray) waterlines⁴⁵. **b** Monthly temperatures at the nearby (~10 km away) Grytviken weather station (see Fig. 1a)^{23,42}. Regression fits with summary statistics between plotted variables are shown in gray. The pink star ($n = 2$) in panel **a** highlights the isotope value of Lake Diamond water samples taken in February 2015.

waxes^{35,58,59}. To constrain vegetation changes in the catchment area over the past 7 ka, we use palynological evidence and n -alkane Average Chain Length (ACL: see methods). The pollen data seen in Fig. 3a show that tussock grasses (*Festuca contracta*) and greater brunet (*Acaena magellanica*) have dominated terrestrial plant cover around Lake Diamond over the past 7 ka. The stable abundance of *Callitriche antarctica* and *Montia fontana*, the only mid-chain n -alkane-synthesizing macrophytes preserved in the Diamond pollen record, suggest aquatic vegetation also changed little⁶⁰. Supporting this evidence, both ACL values and concentrations of mid-chain ($\text{C}_{21}\text{--}\text{C}_{25}$) and long-chain ($\text{C}_{27}\text{--}\text{C}_{33}$) homologs fluctuate around a stable mean throughout our study period (Fig. 3a–c). Finally, we find no relationship between n -alkane abundance and $\delta^2\text{H}$ values of various homologs, indicating that the inferred minor vegetation changes do not explain changes in isotope values (Fig. 5a, b).

Hydrogen isotope values of all analyzed n -alkanes are relatively stable on centennial timescales from the beginning of the Lake Diamond record until ~2150 cal. yr. BP (Fig. 6). Variability then visibly deviates from this long-term mean, as also indicated by change-point analysis (see methods). Deviation from the long-term mean is primarily captured by hydrogen isotope values of mid-chain aquatic homologs ($\delta^2\text{H}_{\text{aq}}$), which become progressively more positive. In contrast, hydrogen isotope values of long-chain terrestrially derived homologs ($\delta^2\text{H}_{\text{terr}}$) exhibit a different and more subdued response that does not significantly deviate from the long-term mean (Fig. 6b). As noted, modern water samples from Lake Diamond are indistinguishable from amount-weighted annual precipitation $\delta^2\text{H}$ values, which in turn track surface temperatures on South Georgia at seasonal timescales (Fig. 1c). Based on the calculated $\delta^2\text{H}$ -temperature dependence of $5.83\text{‰}\text{°C}^{-1}$ (Fig. 2b), the observed ~100‰ $\delta^2\text{H}_{\text{aq}}$ shift to more positive values during the past two millennia would require ~17 degrees of warming. As local and regional temperature reconstructions reveal a shift towards moderately cooler annual (~1 °C) and growing season (~2 °C) conditions during this period

(Fig. 6a)^{27,44,61}, this is highly implausible. Furthermore, the dissimilarity between changes in $\delta^2\text{H}_{\text{aq}}$ and $\delta^2\text{H}_{\text{terr}}$ values suggests that at least one of these records does not simply reflect $\delta^2\text{H}$ values of precipitation. We also exclude large variations in precipitation seasonality: as the isotopic composition of soil moisture used by land plants is typically skewed towards the growing season when biosynthesis occurs⁶², the observed isotopic enrichment of $\delta^2\text{H}_{\text{aq}}$ would also be captured by $\delta^2\text{H}_{\text{terr}}$ if caused by an increase in less negative (relatively ^2H -enriched) spring-summer precipitation (see Fig. 1c).

We interpret the shift to more positive $\delta^2\text{H}_{\text{aq}}$ values after ~2150 cal. yr BP to reflect an increase in evaporative enrichment of the lake water during this time. The observed positive shift in $\delta^2\text{H}_{\text{aq}}$ values, as well as their ^2H -enrichment relative to $\delta^2\text{H}_{\text{terr}}$, are both characteristic of basins with a negative water balance^{20,47}. As noted (see setting and approach), evaporation is often an important source of water loss in closed (endorheic) basin lakes like Lake Diamond^{63,64}. Because evaporative isotope enrichment can outweigh other environmental controls on $\delta^2\text{H}$ values in leaf waxes, and because the positive hydrogen isotope shift during the past two millennia (the Common Era) is only recorded in Diamond Lake by lipids derived from aquatic plants that track lake water³⁵, our interpretation provides an explanation for all of the observations without invoking unreasonable changes in temperature or the seasonality of precipitation. It also allays concerns that the incidental spills of glacial meltwater inferred by²⁹ left a detectable imprint: as this water source is ^2H -depleted⁶⁵, substantial input would result in isotope changes in the opposite direction of what we observe in our record.

Based on the strong seasonal link between meteoric $\delta^2\text{H}$ and temperature in South Georgia (Figs. 1c, 2b), and the tendency of terrestrial compounds to record precipitation $\delta^2\text{H}$ in small catchments like that of Lake Diamond⁶⁶, we infer that down-core $\delta^2\text{H}_{\text{terr}}$ values reflect temperature change. As seen in Fig. 6, this interpretation is supported by the fact that the most prominent feature of our $\delta^2\text{H}_{\text{terr}}$ record—a ~20‰ oscillation towards heavier

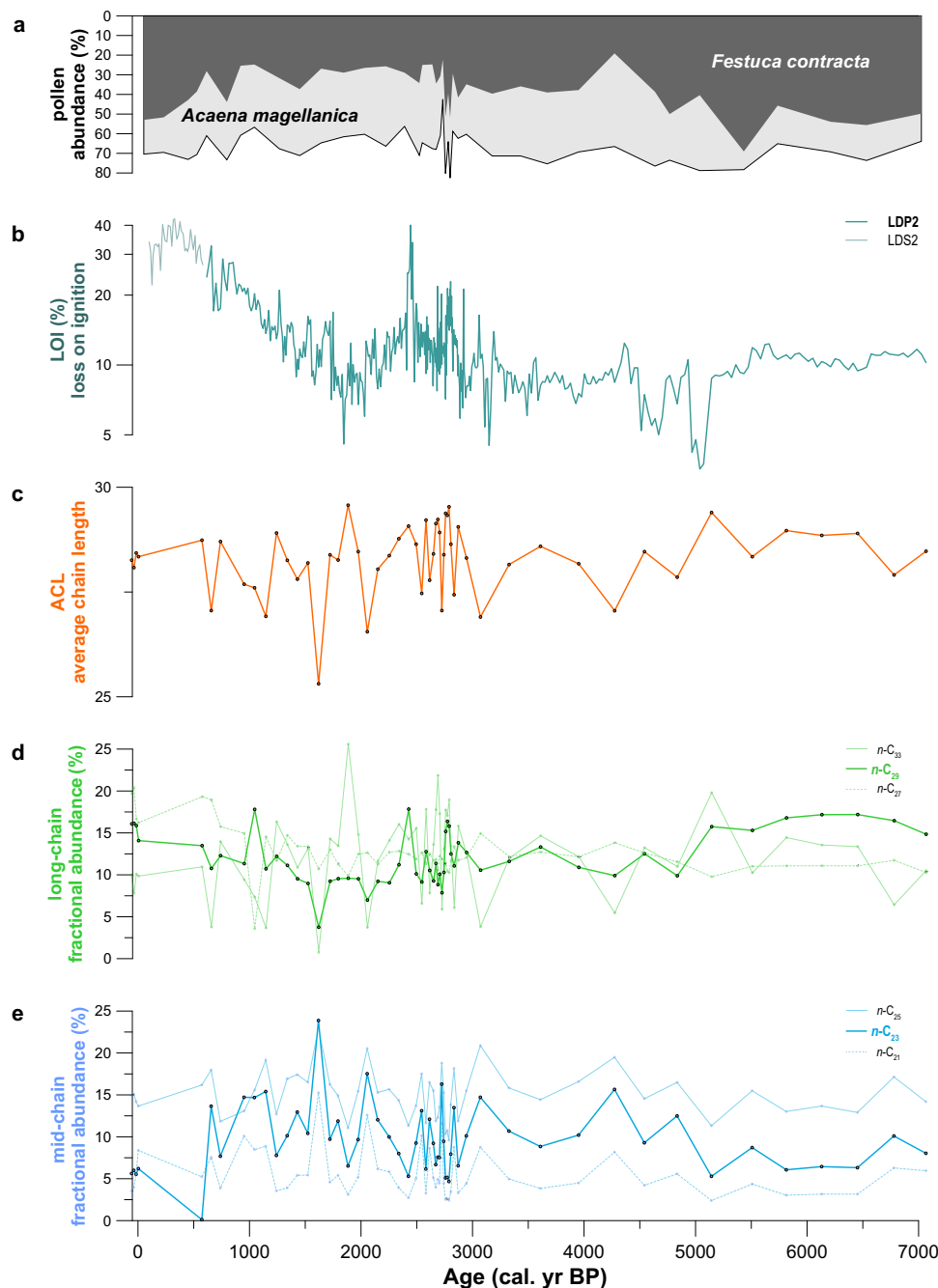


Fig. 3 Mid- to Late Holocene leaf wax distributions, pollen abundances, and organic content (LOI) in Lake Diamond. **a** Fractional abundances of mid-chain sedimentary *n*-alkanes, highlighting the predominantly aquatic C₂₃ homolog. **b** Similar, but for long-chain compounds, highlighting terrestrially-sourced C₂₉. **c** The average chain length (ACL) distribution of down-core sedimentary *n*-alkane samples. **d** Variations in organic content as reflected by LOI measurements in both LDP2 and LDS2 cores. **e** A cumulative pollen diagram for core LDP2 after⁶⁰, highlighting the abundance of two taxa that have dominated local vegetation throughout the studied period: *Festuca* and *Acaena*.

isotopic values between 1200–600 cal. yr BP—coincides with warming phase South Georgia^{27,30,60}.

To quantify the impact of lake water evaporation through time, we calculate the isotopic difference between $\delta^2\text{H}_{\text{terr}}$ (taken to reflect $\delta^2\text{H}$ of precipitation) and $\delta^2\text{H}_{\text{aq}}$ (taken to reflect $\delta^2\text{H}$ of precipitation and evaporative enrichment of lake water) after²⁰, subtracting $\delta^2\text{H}_{\text{terr}}$ from $\delta^2\text{H}_{\text{aq}}$, and defining this parameter as $\epsilon_{\text{terr-aq}}$. We chose the C₂₉ and C₂₃ *n*-alkanes to represent terrestrial and aquatic lipids, respectively, based on their maximum separation along PC2 (Fig. 5c), and use their $\delta^2\text{H}$ values to calculate $\epsilon_{\text{terr-aq}}$ change through time.

Due to core top sediment loss in LDP2 and extensive subsampling of LDS2²⁹, we have no measurements of $\delta^2\text{H}_{\text{terr}}$ and $\delta^2\text{H}_{\text{aq}}$ for the past 500 years (see Fig. 6). To extend our inferred evaporative enrichment reconstruction into this period, which overlaps with notable regional climate transitions including negative and positive phases of the Southern Annular Mode (SAM) and the culmination of the Little Ice Age (LIA)^{12,67}, we rely on proxy-to-proxy calibration⁶⁸. Specifically, we use sediment organic content (measured by loss on ignition or LOI) as a substitute for measured $\epsilon_{\text{terr-aq}}$ values during this interval (see methods), based on (1) statistically significant

($\rho = 0.66$, $p < 0.001$, $n = 14$) correspondence between these measurements (Fig. 7 and methods section) and (2) available LOI data from surface core LDS2 (Fig. 3b). We cannot be certain why this strong link between LOI and $\epsilon_{\text{terr-aq}}$ exists, but widespread evidence for contemporaneous climate deterioration argues against an increase in lake productivity (Fig. 6a, c)^{27,69}. Vegetation change also seems an unlikely culprit, as the previously discussed pollen evidence from Lake Diamond reveals that Holocene variability was minimal and unrelated to measured δD values (see section 2.1, and Figs. 3a, 5b). Finally, the non-significant anti-correlation

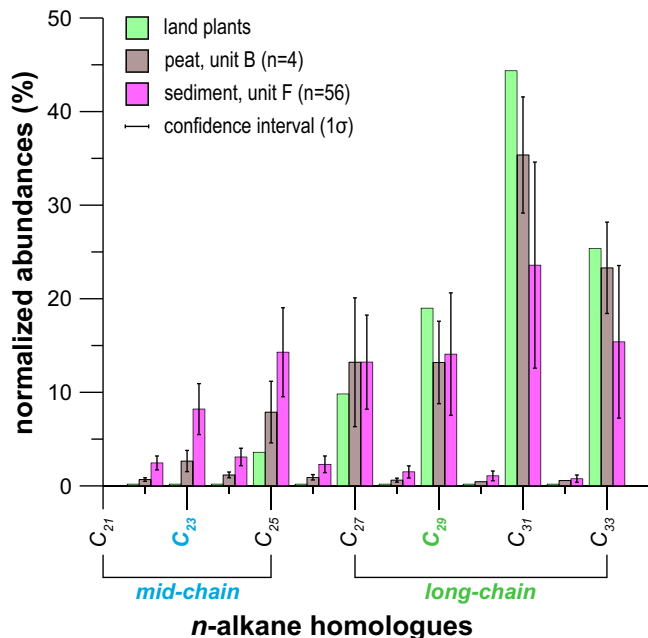


Fig. 4 Chain-length distributions. The abundance of n -alkane homologues in terrestrial plants collected in South Georgia by ref. ⁵⁴, and sediment and peat samples from the investigated archive from Lake Diamond (LDP2). We distinguish between mid- and long-chain compounds after the definition by ref. ⁴⁹ in color coding that corresponds with Figs. 3, 5, 6. Confidence intervals are based on down-core variations of LDP2 samples (see Fig. 6).

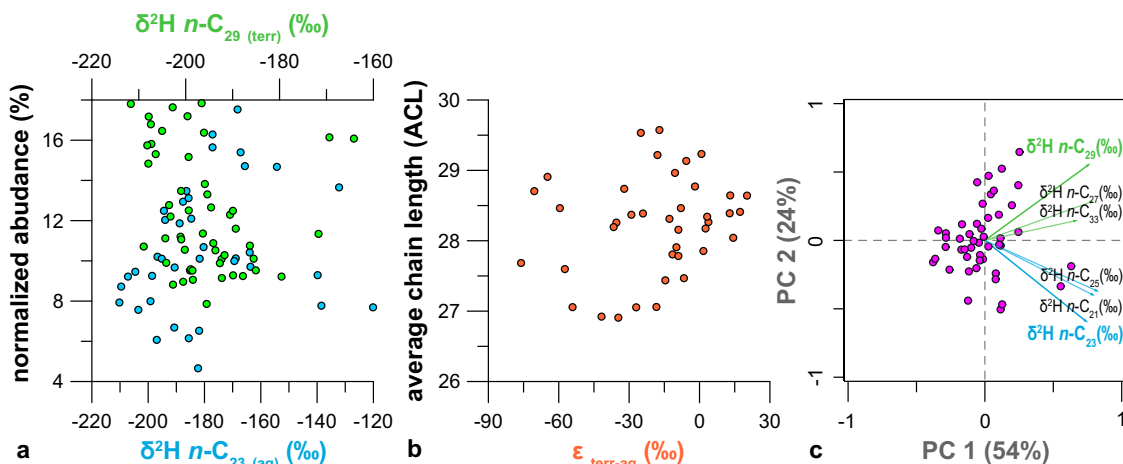


Fig. 5 Relations between n -alkane chain length and isotope values. Bi-plots showing **a** The hydrogen isotope ($\delta^2\text{H}$) composition of representative mid- (C_{23}) and long-chain (C_{29}) sedimentary n -alkane sample combinations versus their normalized abundance, **b** The isotopic offset between compounds produced by representative aquatic (C_{23}) and terrestrial (C_{29}) plants ($\epsilon_{\text{terr-aq}}$) versus the Average Chain Length (ACL) of all alkanes measured in these samples ($n = 39$), and **c** Sample (pink dots) and variable (vectors) scores for the main ordination axes (PC 1–2: labeled with their explanatory power) of principal component analysis (PCA) carried out on measured $\delta^2\text{H}$ values of analyzed n -alkanes ($n\text{-C}_{21}$ to $n\text{-C}_{33}$ without $n\text{-C}_{31}$: see methods section).

($R^2 = -0.03$) between LOI and the mass-specific magnetic susceptibility (χ_{bulk})-derived glacier reconstruction by²⁹ allays concerns about the imprint of dilution with glacial clastic input during the investigated period. We hypothesize that the correlation between LOI and $\delta^2\text{H}_{\text{aq}}$ values can be attributed to the impact of inferred lake evaporation on water levels: particularly in a shallow and conically-shaped basin like Lake Diamond²⁹, low stands focus organic deposition into the center where our cores were extracted⁷⁰. In support of this hypothesis, we also observe a strong ($\rho = -0.52$, $p < 0.05$, $n = 13$) correspondence in LDP2 between $\epsilon_{\text{terr-aq}}$ and *Pediastrum* (Fig. 7)⁶⁰. The abundance of this green algae genus is widely used as an indicator of lake level shallowing⁷¹, also in upwind southern Patagonia⁷². As *Pediastrum* synthesizes some mid-chain n -alkanes⁷³, we should re-iterate that there is no relation between the abundance and isotopic value of these homologues in Lake Diamond (Fig. 5a).

Wind-driven evaporation during the past two millennia. We propose the inferred increase in evaporative water loss from Lake Diamond after ~ 2150 cal. yr BP was driven by strengthening winds. The previously discussed evidence for local and regional cooling at this time rules out direct temperature effects (Fig. 6a)^{27,44,61}. We also reject decreasing relative humidity as an explanation because such a change would be reflected by an increase in $\delta^2\text{H}_{\text{ter}}$ values due to leaf water enrichment⁴⁸, which we do not observe (Fig. 6b). Our interpretation is supported by macrofossil evidence from peatlands, sensitive archives of sub-Antarctic surface water balance^{74,75}, which reveal a sharp transition ~ 2200 cal. yr BP towards colder wetter (humid) conditions in South Georgia (Fig. 6c).

Variations in wind strength can leave a measurable isotopic imprint on closed (endorheic) basins in the SWH core belt like Lake Diamond⁷⁶. Nearby paleoclimate records indicate that wind strength indeed increased after ~ 2150 cal. yr BP, when $\epsilon_{\text{terr-aq}}$ began to deviate. A record from neighboring Annenkov Island shows an increase in the accumulation rates of long-distance transported (LDT) pollen—an established indicator of wind strength^{77,78}. Further upwind (52.5°S and 72.6°W), close to the Andean crest, where strengthening of the westerlies is coupled to an increase in orographic precipitation⁷⁹, the amount of hygrophytic pollen increased throughout the last two millennia⁸⁰. A similar trend is observed in upwind Lago Guanaco (51.5°S and

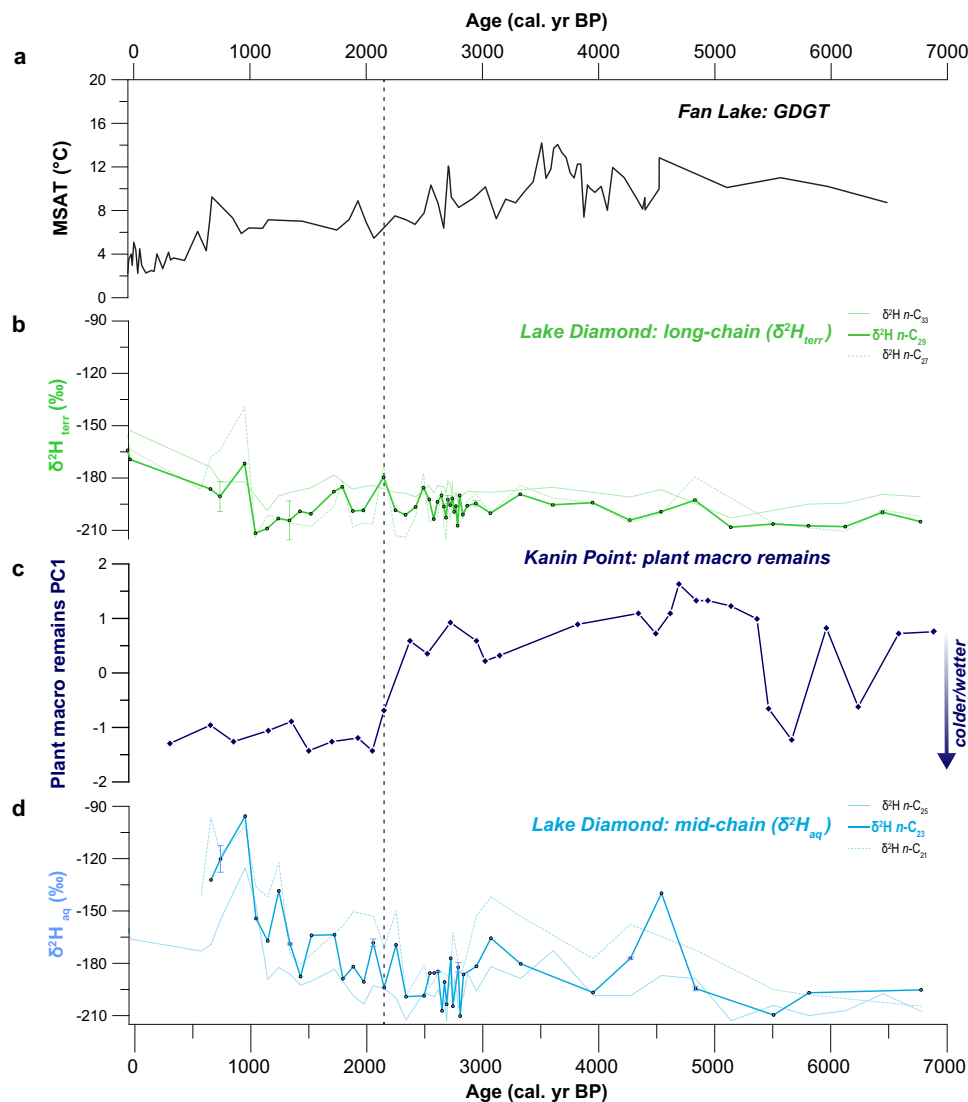


Fig. 6 Comparing compound-specific hydrogen isotope values ($\delta^2\text{H}$) from Lake Diamond with local temperature and hydroclimate proxies. a GDGT-derived mean summer air temperatures (MSAT) from neighboring Annenkov Island²⁷. **b** Terrestrially produced long-chain *n*-alkane $\delta^2\text{H}$ values from Lake Diamond (this study), highlighting representative C_{29} homolog. **c** Principal component analysis (PCA) of plant macrofossil data from ref. ⁶⁹. **d** Aqueatically sourced short-chain *n*-alkane $\delta^2\text{H}$ values from Lake Diamond (this study), highlighting representative C_{23} homolog. The ~2150 cal. yr BP age where change-point analysis detects a significant deviation from the long-term mean is shown by a stippled line. All color coding matches that of Figs. 3–6.

72.5 °W), where (1) a carbonate-derived $\delta^{18}\text{O}$ -based reconstruction of the balance between wind-driven precipitation and surface evaporation from Lago Guanaco is similar ($\rho = 0.53$, $p < 0.001$, $n = 35$) to our inferred wind-driven evaporation record from Lake Diamond during the overlapping ~100–950 cal. yr BP period (Fig. 8)²¹, and (2) gradual wind-driven precipitation increases during this period are reflected by the dominance of forest over grassland⁴⁰ (Fig. 9d). We attribute the delayed response of the latter transition to the well-documented nonlinear relation between regional changes in vegetation and climate¹⁹.

Links to the southern annular mode (SAM). The Lake Diamond $\epsilon_{\text{terr-aq}}$ wind-driven lake evaporation reconstruction suggests that the past two millennia were characterized by major regional shifts in core belt SHW strength that appears unmatched in the past 7 ka. As seen in Figs. 8, 9, the transition towards windier conditions seems to mark a major reorganization of the regional climate system as the pattern of change observed in Lake Diamond broadly coincides with (1) pronounced annual cooling towards a

post-glacial temperature minimum off the northeastern Antarctic Peninsula (Fig. 9b)—recorded by ice core $\delta^2\text{H}$ values from James Ross Island (Fig. 1a) and (2) extension of the sea ice season towards a Holocene maximum along the western Antarctic Peninsula (Fig. 9c) as indicated by a rise in sea ice diatom biomarker concentrations⁸¹. A concomitant increase in seabird-derived phosphorus (P) from a rookery on the Falklands also attests to the broad ecological impacts of these climatic changes (Fig. 9e)⁸². Sea ice cover and wind stress have a major impact on the sub-Antarctic food web through impacts on the transport of nutrient waters as well as productivity and feeding^{83,84}. This reconstructed pattern of coupled cooling, mid-latitude westerly intensification, and the Bellingshausen Sea ice expansion is diagnostic for a negative phase of the Southern Annular Mode or SAM^{12,85,86}. The SAM is the main pattern of Southern Ocean climate variability⁹ and affects the strength and position of the SHW by modulating the atmospheric pressure gradient between 40–65°S (see introduction). There is a strong correlation (Spearman’s correlation coefficient $\rho = 0.75$, $p < 0.0001$, $n = 35$) between our inferred evaporation record and the sole annual

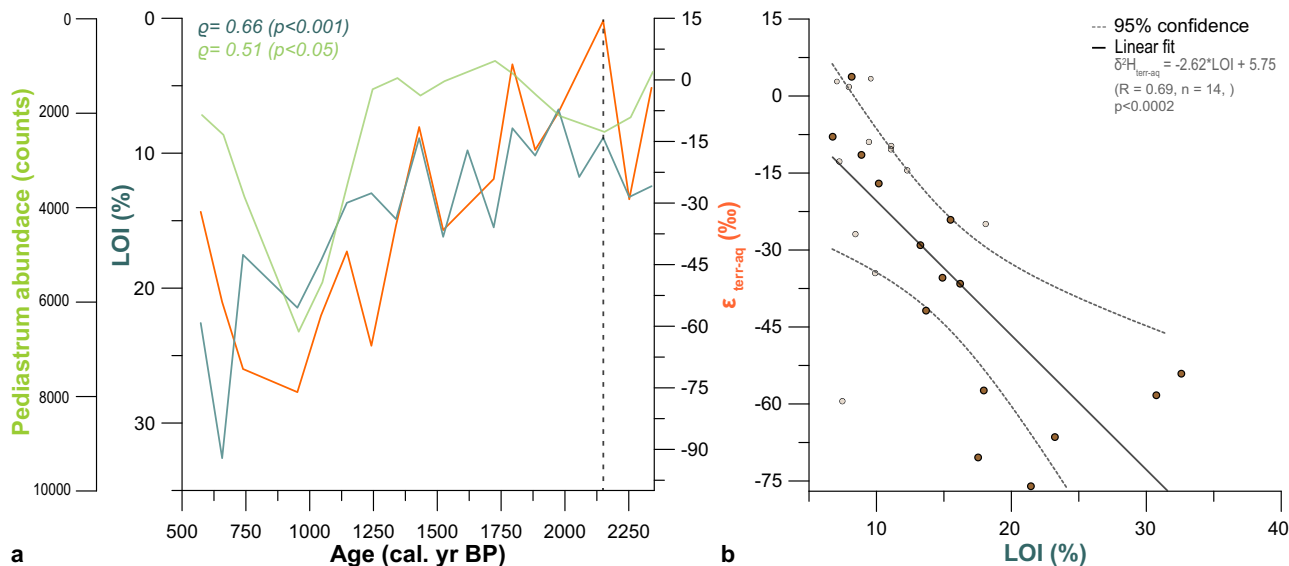


Fig. 7 Proxy-to-proxy calibration. **a** $\epsilon_{\text{terr-aq}}$ and LOI values, as well as *Pediastrum* counts from ref. ⁶⁰ in core LDP2 over the past two millennia (unit F3: see methods) and **b** A bi-plot for this period that captures the association between these parameters with a linear regression fit (opaque dots) and shows how data from the preceding 2.4–7 ka BP period covered by LDP2 (transparent dots) relate to this model (also see methods). For the latter purpose, we excluded values ($n = 7$) from the ~2.4–2.7 ka BP to avoid closed-sum effects from dilution with alluvial clastic input during this time, as reported by ref. ²⁹.

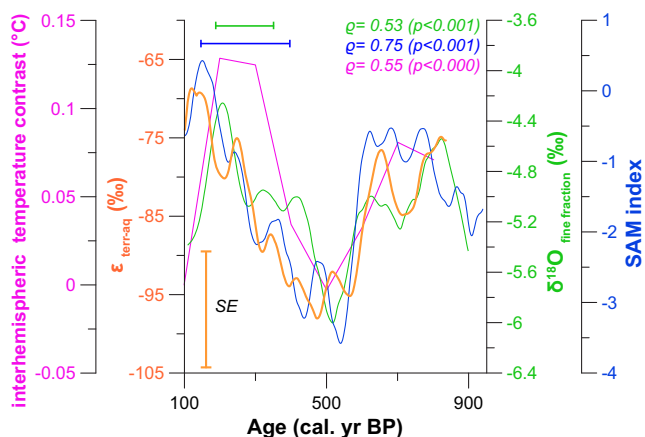


Fig. 8 Comparing inferred $\epsilon_{\text{terr-aq}}$ values from Lake Diamond (orange) with other regional proxies of wind-driven change. In blue, the Southern Annular Mode (SAM) index by ref. ¹². And in green, the $\delta^{18}\text{O}$ fine fraction SHW record from Lago Guanaco by ref. ²¹. Shown in purple is the area-weighted inter-hemispheric temperature contrast between the extratropics (30–90°) of both hemispheres (North–South) using 30° averages of ensemble means presented by ref. ¹¹⁰. All reconstructions have been smoothed using a 70-year moving average. In the upper part of the figure, we show chronological uncertainties (horizontal bars) and correlation coefficients (italics) in matching colors. The vertical bar at the base marks the reported standard error (SE) of our PTP-derived model (see methods).

SAM reconstruction that extends back to 1000 CE, demonstrating that Lake Diamond sediments indeed capture shifts in the behavior of this dominant regional climate mode (Fig. 8)¹². These also include the positive SAM trend—characterized by less windy conditions over Lake Diamond—during the past 500 years. We thus argue that our reconstruction of wind-driven evaporation is linked to SAM variability over the past 7 millennia. However, to acknowledge that links with SHW strength (1) depend on the choice of index and reanalysis data²², and (2) could be non-stationary through time¹³, we refer to SAM-like changes in the following paragraphs after⁸⁵. Concretely, we define these as

changes in wind-driven evaporation that were likely driven by shifts in the meridional pressure gradient.

Positive modern SAM phasing was not unprecedented during Holocene. The Lake Diamond record provides valuable information concerning long-term SAM behavior. Notably, the period from 7 ka BP until ~2150 cal. yr BP was comparatively stable and mostly characterized by positive SAM-like conditions (Fig. 9). Based on baseline data that do not extend beyond the past millennium¹³, it has been argued that today's positive SAM phasing—is attributed to a combination of anthropogenic greenhouse forcing and ozone depletion^{11,87}—falls outside the range of natural variability⁸. Our findings indicate that this may not be the case and that the negative SAM polarity of the past two millennia was exceptional rather than today's positive phasing. While acknowledging the possibility that the relation between wind-driven evaporation in Lake Diamond and the SAM is non-stationary beyond the past millennium¹³, we use the relationship ($R^2 = 0.65$, $p = 0.000$) between the Lake Diamond data and the SAM index by¹² to approximate modern limits. Based on this basic calculation, we find that $\epsilon_{\text{terr-aq}}$ values greater than -45‰ exceed the observed 1999 CE SAM maximum (Fig. 9a). Notwithstanding the uncertainties of this simple approach, it highlights that SAM values have been near or exceeded observational limits during most of the Mid- to Late Holocene, and underscores the potential of this period to better understand the impacts of a more positive SAM, which is predicted for the future⁸⁷. Our findings corroborate previous proxy-derived evidence that strongly positive SAM phases do not require anthropogenic radiative forcing^{13,85}. Taken together, this body of work hints at the possibility that internal variations, which will be superimposed on future anthropogenic change but remain poorly resolved in climate models⁸⁷, play a critical but undervalued role in driving SAM shifts.

Was SAM-like change driven by Northern Hemisphere cooling? In the following discussion, we examine whether the multi-centennial Common Era climate variations captured by the Lake Diamond record and other reconstructions of SAM-like change in

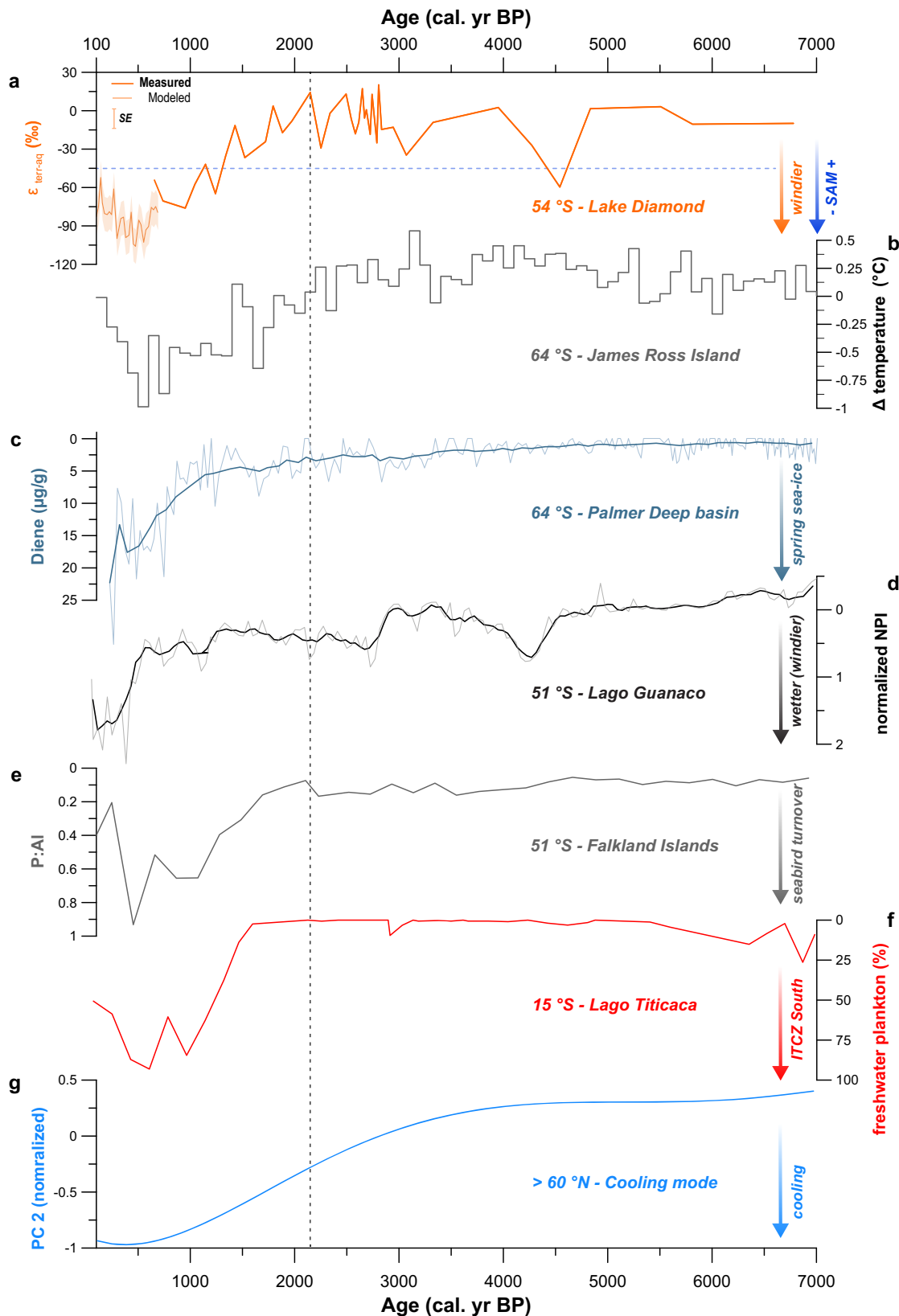


Fig. 9 could have been caused by modes of internal variability. Specifically, the El Niño-southern oscillation or ENSO¹⁴. El Niño events are associated with negative SAM phases because positive temperature anomalies in the Tropical Pacific trigger Rossby wave trains that propagate positive pressure differences poleward⁸⁸. Conversely, La Niña events coincide with positive SAM phases as

ensuing Tropical Pacific cooling leads to low sea-level pressure anomalies in the sub-Antarctic. Recently, it has been shown that these connections also operate on centennial timescales¹⁷. However, while the reconstructed SAM minimum ~500 cal. yr BP (Fig. 8) coincides with a phase of El Niño-dominated conditions⁸⁹, no consistent link exists between records of SAM-like variability

Fig. 9 Comparing evaporation in Lake Diamond with regional proxies of sea ice, wind, seabird, and temperature shifts. **a** Lake Diamond $\epsilon_{\text{terr-aq}}$ values. Please note that we distinguish between reconstructed and PTP-inferred values (see methods), while also displaying the SE of this model. **b** James Ross Island (JRI) ice core temperature reconstruction after⁴⁴, showing 100-year averaged anomalies compared to the CE 1961–1990 mean. **c** Diatom-derived diunsaturated (diene) highly branched isoprenoid (HBI) biomarker concentrations from⁸¹. **d** Raw and smoothed standardized *Nothofagus* (southern beech tree)/Poaceae (grass) Index (NPI) values from⁴⁰. **e** Seabird-derived ratios of bio-element Phosphorus (P) over lithogenic Aluminum (Al) after⁸². **f** Freshwater plankton abundances (%) in Lake Titicaca from¹⁰⁴. **g** Normalized smoothed principal component 2 (PC2) values of the global cooling mode inferred by ref. ¹⁰⁹.

and ENSO reconstructions⁹⁰. Indeed, there is growing consensus that ENSO-SAM links are non-stationary^{13,91}, and also depend on the strength of the latter⁹². In light of this evidence, we note that the Common Era excursion captured in the Lake Diamond record and the other records of SAM-like change in Fig. 9 coincides with the onset of ENSO strengthening to a Holocene maximum^{93,94}. Such long-term trends are, however, often characterized as ENSO-like and attributed to other mechanisms that likewise affect tropical Pacific surface temperature change⁸⁸.

There is, for example, mounting evidence for a global atmospheric teleconnection that is (1) considered analogous to ENSO as it also impacts equatorial convection, but (2) driven by the temperature gradient between Earth's hemispheres, and (3) operating at the multi-centennial timescales resolved by our record⁹⁵. The emerging observational, model-based, and proxy-derived evidence supporting this framework shows that differential extratropical temperature change increases poleward heat export to the cooler hemisphere^{96–98}. To facilitate this, the inter-tropical convergence zone (ITCZ) moves into the warmer hemisphere so that more energy is transported across the equator^{99,100}. A similar response is observed during ENSO variations as the ITCZ migrates south into a warming South Pacific during El Niño events and vice versa during La Niña¹⁰¹. Critically, model experiments reveal that this feedback also alters the strength and position of the core SHW belt through poleward Rossby wave train propagation¹⁰²—a key characteristic of modern ENSO-SAM interactions⁸⁸.

To assess whether this global atmospheric teleconnection may help explain the Common Era excursion observed in our isotope record and the other reconstructions of SAM-like change shown in Fig. 9, we focus on anticipated changes in ITCZ position and meridional temperature gradient⁹⁵. In light of the presented shift towards anomalously negative SAM values (Figs. 8, 9a), and the association of this state with El Niño-like (negative ENSO) conditions⁸⁸, these include evidence for a southward ITCZ migration in response to Northern Hemisphere cooling or Southern Hemisphere warming. Proxy evidence from the sensitive southern limit (15°S) of the present-day South American ITCZ reveals major contemporaneous precipitation increases—the main expression of such a shift¹⁰³. As seen in Fig. 9f, this is recorded by a sharp increase in freshwater plankton abundances in oligohaline Lake Titicaca from 2 ka BP onwards¹⁰⁴. A transition from dry savanna to a wetter rain forest, reflected by a sharp rise in *Moraceae* (mulberry) pollen ~2160 cal. yr BP in southern Amazonia (13–14°S)¹⁰⁵ also supports this interpretation. Both observations are in broad agreement with global evidence for a southward ITCZ migration during the Late Holocene in response to orbitally-driven Northern Hemisphere cooling^{96,106}. This trend was amplified during the last two millennia by the negative radiative effects of Arctic sea ice expansion¹⁰⁷ and is identified as one of the major temperature shifts of the Holocene¹⁰⁸. Recent proxy-validated simulations even identify this feedback as the main driver of global temperature change during the Common Era¹⁰⁹.

As seen in Fig. 9g, the evolution of this cooling mode tracks the millennial-scale trend of mid to late Holocene SAM-like

variability in our study region and shows that cooling rates intensify around ~2100 cal. yr BP. To further investigate the correspondence between the Common Era SAM-like changes discussed here and (relative) Northern Hemisphere cooling, we compare our Lake Diamond reconstruction and the millennium-long SAM index of¹² to the temperature contrast between the extratropics (30–90°) of both hemispheres (North minus South)—a measure of Earth's meridional temperature gradient. To do so, we rely on the area-weighted ensemble means of the 30° latitude average presented by¹¹⁰. Figure 8 underlines that the Northern Hemisphere cooled differentially during this period and reveals a notable degree of agreement ($\rho = 0.55$, $p < 0.000$, $n = 22$) with coeval SAM-like changes.

Following the above, we argue that the inferred shift towards a negative SAM-like state after ~2150 cal. yr BP is consistent with a change in Earth's thermal gradient in response to amplified Arctic cooling. Our findings thus provide additional evidence for the existence of a previously hypothesized atmospheric teleconnection that links the high latitudes in both hemispheres and impacts SHW strength⁸⁸. This raises the possibility that ongoing amplified Arctic warming may (1) reinforce the predicted trend towards a more positive phase of SAM⁸⁷ and (2) impact the ability of the SHW to help regulate the global climate in the future.

Methods

Biomarker analysis. In total, 56 samples were extracted from the lacustrine sequence of piston core LDP2 ($n = 52$) and surface core LDS2 ($n = 4$) at 2–4 cm intervals. In addition, we analyzed five samples from the underlying peat deposit in LDP2. Prior to analysis, these sediments were homogenized and freeze-dried. We extracted lipids from each sample with a 9:1 v/v solution of dichloromethane (DCM) and methanol using a Dionex 350 Accelerated Solvent Extractor. The Total Lipid Extract (TLE) that was obtained was then dried under a stream of nitrogen gas in a Biotage TurboVap. Next, TLEs were separated into aliphatic, ketone, and polar fractions using silica gel flash column chromatography. For this purpose, we sequentially used four bed volumes of hexane, DCM, and methanol as eluents. Aliphatic compounds were then treated with copper to remove elemental sulfur and eluted through silver nitrate-impregnated silica gel to remove unsaturated hydrocarbons. We subsequently identified and quantified C_{21} – C_{33} *n*-alkanes in the saturated fraction following the protocol outlined by ref. ¹¹¹ and by comparing sample mass spectra to a drift standard containing homologous *n*-alkanes (C_8 – C_{40}), phytane, pristane, and 5 α -androstandane. Measurements were made using the protocol by ref. ¹¹¹ on an Agilent 7890 gas chromatograph (GC) equipped with both mass selective (MSD) and flame ionization (FID) detectors, as well as an HP-5ms column. Average Chain Lengths (ACL) were determined¹¹²:

$$ACL = \frac{\sum(C_i)(X_i)}{\sum(C_i)}, \quad (1)$$

where C is the concentration and X is the chain length of *n*-alkane i .

Hydrogen isotope measurements of these homologs were performed on a Thermo Trace GC coupled to a Thermo Delta V Isotope Ratio Mass Spectrometer (IRMS) via an Isolink Conflo IV interface using the same column and protocol by ref. ¹¹¹ as for GC-MSD/FID analysis. Based on chromatographic evidence of co-elution with a hopanoid compound, we excluded *n*- C_{31} isotope ratios from our record. Most compounds were measured in triplicate, and the precision of replicate measurements was typically better than 2.5‰ (1 σ). Measurements with intensities lower than 1000 mV on mass two were excluded. To monitor analytical uncertainty, an external transesterified *n*-alkanoic acid mixture calibrated against authenticated A4 and F8 lipid mixtures—provided by A. Schimmelmann from the University of Indiana—was measured after every sixth sample. Throughout the analysis, precision ranged from 0.95 to 2.4‰ among four homologs. To convert compound-specific δ^2H values to the VSMOW scale after¹¹³, we regularly measured the authenticated A5 lipid mixture—provided by A. Schimmelmann from the University of Indiana. We also report the δ^2H offset between terrestrially-sourced *n*- C_{29} and aquatically produced *n*- C_{23} ($\epsilon_{\text{terr-aq}}$) after¹¹⁴.

Core stratigraphy. The stratigraphy of the presented Lake Diamond record is described in detail by ref. 29. As stated before, our analysis prioritizes sediments deposited under lacustrine conditions. These encompass the upper 221.5 cm of piston core LDP2 that cover the past 7.2 ka²⁹, subdivide this sequence into three lithological units: the compacted silt-dominated organic-rich sediments of unit D (194–221.5 cm), the poorly sorted sandy mass wasting deposit of unit E (185.5–194 cm) and the silty-sandy dark brown organic-rich deposits of unit F (0–185.5 cm). Here, we complement the multi-proxy dataset from²⁹ with palynological and loss-on-ignition (LOI) analyzes. To allow proxy-to-proxy calibration for the past 500 years—which are not covered by core LDP2²⁹—we measured LOI at ~1 cm increments on sub-sectioned pollen samples from surface core LDS2 ($n = 35$). For this purpose, we extracted 0.5 ml of sediment with a syringe and dried it overnight at 105 °C before burning samples at 550 °C for 4 h after¹¹⁵. These data do not overlap with the observational period as the uppermost 6.5 cm (150 years) has been used for ²¹⁰Pb dating⁶⁰ and biomarker analysis. Pollen samples were analyzed at 4–10 cm intervals from the composite lake Diamond record ($n = 41$) as detailed by ref. 60. Slide preparation was carried out using standard techniques after¹¹⁶. Eucalyptus tablets were added to calculate concentrations, while identification was aided by¹¹⁷ and reference material collected during fieldwork in 2011 and 2015.

Geostatistics. We used basic predictive models to construct waterlines (Fig. 2a), constrain the controls on meteoric $\delta^2\text{H}$ (Fig. 2b), and substitute LOI for $\epsilon_{\text{terr-aq}}$ (see Section 3.1 and Fig. 7b). For this purpose, the following linear ordinary least squares (OLS) regression curves were fitted with the help of Stata 16:

$$\delta^2\text{H} = 7.61 * \delta^{18}\text{O} + 0.84 \quad (2)$$

$$\delta^2\text{H} = 5.83 * T - 78.11 \quad (3)$$

$$\epsilon_{\text{terr-aq}} = -2.62 * \text{LOI} + 5.75 \quad (4)$$

To avoid artifacts stemming from autocorrelation⁶⁸, we relied on the Durbin–Watson d statistic using 1.5 and 2.5 as cut-off values following the recommendations of ref. 118. Proxy-to-proxy calibration of $\epsilon_{\text{terr-aq}}$ against LOI focused on the upper part of LDP2, where both datasets significantly deviate from a stable long-term mean (see section 2.1 and Fig. 7a). To pinpoint this transition, we relied on change-point analysis using the `cpt.mean` function and AMOC method in version 2.2.2 of the change-point R package¹¹⁹. The detected change-point at 60 cm (~2150 cal. yr BP) core depth corresponds to the transition into unit F4²⁹. We did not incorporate include data from underlying unit F3 to avoid closed-sum effects from dilution with alluvial clastic input deposited in this interval as reported by²⁹. We do, however, show how $\epsilon_{\text{terr-aq}}$ –LOI combinations from this preceding 2.4–7 ka BP period covered by LDP2 relate to equation 5 in Fig. 7b. Finally, we carried out a principal component analysis (PCA) on samples ($n = 49$) where $\delta^2\text{H}$ was measured at least once on each homolog (n -C21 to n -C33 without n -C31; see biomarker analysis) to evaluate the degree of co-variance between compounds (see section 2.1 and Fig. 4c). To do so, we used version 5 of the Canoco software¹²⁰. Following the recommendations of ref. 121, all n -alkane homologs were centered and standardized prior to analysis.

Data availability

The authors declare that all data that we generated for this study and are presented in the figures of this manuscript are available in Supplementary Data 1 and have been uploaded to the DataverseNO repository. Here, the files can be accessed using the following DOI - <https://doi.org/10.18710/IYNKEZ>.

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Author contributions

W.G.M.v.d.B. and W.J.D.A. designed the study. W.G.M.v.d.B. and L.T.O. wrote the paper. W.J.D.A., L.T.O., M.Z., and W.G.M.v.d.B. performed the analyses. J.B. and A.E.B. led fieldwork. All authors contributed to the main text during multiple rounds of comments.

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The authors declare no competing interests

Additional information

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