

Holocene water balance variations in Great Salt Lake, Utah: application of GDGT indices and the ACE salinity proxy

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Key Points:

- Glycerol dialkyl glycerol tetraethers produced by bacteria and archaea detect Holocene limnological changes at Great Salt Lake.
- The Archaeol and Caldarchaeol Ecometric salinity index remains near its upper limit in Great Salt Lake from 7.2 ka to present.
- A 5.5 ka step shift in microbial lipids suggests limnological change coeval with the end of a regional dry spell in the mid-Holocene.

Abstract

Great Salt Lake, Utah, is a hypersaline terminal lake in the Great Basin, and the remnant of the late glacial Lake Bonneville. Holocene hydroclimate variations cannot be interpreted from the shoreline record, but instead can be investigated by proxies archived in the sediments. GLAD1-GSL00-1B was cored in 2000 and recently dated by radiocarbon for the Holocene section with the top 11 m representing ~7 ka to present. Sediment samples every 30 cm (~220 years) were studied for the full suite of microbial membrane lipids, including those responsive to temperature and salinity. The Archaeol and Caldarchaeol Ecometric (ACE) index detects the increase in lipids of halophilic archaea, relative to generalists, as salinity increases. We find Holocene ACE values ranged from 81-98, which suggests persistent hypersalinity with <50 g/L variability across 7.2 ka. The temperature proxy, MBT'_{5Me}, yields values similar to modern mean annual air temperature for months above freezing (MAF = 15.7°C) over the last 5.5 ka. Several GDGT metrics show a step shift in microbial communities and limnology at 5.5 ka. Extended archaeol detects elevated salinity during the regional mid-Holocene drought, not readily detected in the ACE record that is often near the upper limit of the index. We infer that the mid-Holocene GSL was shallower and saltier than the late Holocene. The current drying may be returning the lake to conditions not seen since the mid-Holocene.

Keywords: GDGT, ACE, Holocene, Great Salt Lake

Plain Language Summary

Great Salt Lake in Utah is the remnant of a once much larger lake and is currently at a historically low level. We study a lake sediment core, collected in 2000 from the floor of Great Salt Lake, and recently dated. We take new samples from the core and measure them for molecules made by microbes, whether living in the lake or washed in from the surrounding soils. We reconstruct lake conditions during the last 7,200 years and assess whether lake level fluctuated during that time. Over the past 7,200 years, we find evidence that the lake was shallower from 7,200 to 5,500 years ago but has been relatively stable until the modifications of the lake in the 20th century and the current drying trend.

1. Introduction

Paleoclimate reconstructions of the Holocene provide insights into the stability or sensitivity of our present climate state. While it is a time of relative climatic stability compared to the glacial-interglacial cycles that preceded it, the changes may in some cases be profound especially in zones prone to drought-stress. In southwestern North America, lake sediments and speleothems record a generally wet early Holocene (11 to 8 ka), and a dry middle Holocene (8 to 4 ka) with a return to relatively moist late Holocene (Lachniet et al., 2020 and references therein). Although many Holocene climate modeling studies have struggled to simulate the mid-Holocene warmth, recent modeling has found this emerges with the addition of vegetation feedbacks (Thompson et al., 2022). As the mid-Holocene warming may be an analog for the current warming, it is studied here as context for the current drought in the region.

Great Salt Lake (GSL), UT, USA, at 41.1°N is a remnant of a larger lake, Lake Bonneville. Lake Bonneville reached its maximum size during the termination of the Last Glacial Maximum (LGM) at ~18 ka then shrank to the areal extent of modern GSL by ~13 ka (Oviatt et al., 2021). The drying between the LGM and Holocene greatly increased salinity in the residual lake and left behind the Bonneville salt flats. This drastic shrinking and lake level drop was recorded in preserved shorelines (Oviatt et al., 2021). In contrast, any recessional shorelines of the mid-Holocene arid interval have been lost to subsequent late Holocene transgressions. Instead, continuous geochemical records from sediment cores have potential to reconstruct Holocene climate. As its name implies, Great Salt Lake is salty with current salinity of 120-180 g/L (data for 2010-2021; Rupke and McDonald, 2012), making it hypersaline (>50 g/L salinity). In a closed-basin lake like GSL, salinity is controlled by the hydrologic balance between input from rivers, springs/groundwater, and precipitation, and output as evaporation, with increased evaporation resulting in lower lake levels and higher salinity. Using organic geochemical evidence, we seek to reconstruct GSL salinity and limnological conditions across the Holocene to provide longer context for present-day drying trends and declining lake levels (Wurtsbaugh et al., 2017).

Archaeol is a diphytanyl glycerol diether (DGD), a membrane lipid made by a diverse range of archaea (e.g. Vandier et al., 2021 and references therein) including abundant production by halophilic *Euryarchaeota* (Dawson et al., 2012; Macalady et al., 2004; Teixidor et al., 1993). Proportional abundances of archaeol have been used to track the contrasting salinity of

hypersaline, marine, estuarine, and freshwater environments (Turich and Freeman, 2011). Application of this method to lake sediment cores has revealed large salinity fluctuations in the late Pleistocene in Lake Elsinore (CA) and Searles Lake (CA) (Feakins et al., 2019; Peaple et al., 2022), but the response to more subtle variations in the Holocene has not yet been tested. Another DGD, extended archaeol, has been identified as a more diagnostic biomarker for halophilic *Euryarchaeota* (Bale et al., 2019; Vandier et al., 2021), but not yet reported from southwestern North American hypersaline lakes.

Aside from DGDs, archaea also produce a number of isoprenoid glycerol dialkyl glycerol tetraethers (isoGDGTs) and these have been used as microbial and environmental indicators in aquatic environments. GDGT-0 (previously referred to as caldarchaeol) is produced by *Crenarchaeota*, *Thaumarchaeota*, and some *Euryarchaeota* (Schouten et al., 2013). High proportions of this compound, in the absence of abundant crenarchaeol, have been used to denote production by methanogenic *Euryarchaeota*, which can be indicative of stratified waters (Baxter et al., 2021; Blaga et al., 2009; Naeher et al., 2014). In stably stratified hypersaline lakes, mixing may occur during lake freshening events (Peaple et al., 2022) or in lakes shallow enough for wind mixing (Stefanescu et al., 2021).

Branched GDGTs (brGDGTs) are bacterial membrane lipids that have also been widely applied to reconstruct paleoenvironments including applications to soils and lacustrine sediments for temperature reconstructions (e.g. Martinez-Sosa et al., 2021; Peterse et al., 2012; Weijers et al., 2007). Complications with lake depth have been noted with this paleothermometry technique in nearby alpine lakes with temperature stratification (Stefanescu et al., 2021). As GSL is located at mid-elevation (1,280 masl) and has density primarily controlled by salinity (rather than temperature), those alpine lake depth complications are not expected to apply. However, the hypersaline environment of GSL may carry additional complications, as salinity outliers have been noted in existing calibrations (Martinez-Sosa et al., 2021). Here, using DGDs, isoGDGTs, and brGDGTs, we generate a 7.2 ka hydroclimate history of GSL to track climate change and limnological conditions from the middle to late Holocene, paying particular attention to the limitations of GDGT proxies in this extreme environment.

1.1. Study location

GSL is a hypersaline terminal lake located in the eastern part of the United States' Great Basin (**Figure 1**). At the lake's historic mean elevation of 1,280 masl (m above sea level), the lake is more than 10 m deep across the central part of the basin and averages 5.5 m deep (Belovsky et al., 2011; Shope and Angeroth, 2015). In 2000, the Global Lakes Drilling (GLAD) initiative drilled four sites to a total of 120 m below lake floor reaching back to the boundary of oxygen isotope stages 6 and 7 (~175 ka) (Dinter et al., 2000). The original study reported density, magnetic susceptibility, minerology, inorganic carbon, and organic carbon from the recovered cores (Dinter et al., 2000). Recently the Holocene sedimentary section in the upper 11 m of GLAD1-GSL00-1B (41.094°N, 112.365°W, 8.4 m water depth) was dated using radiocarbon (Bowen et al., 2019). The Holocene sedimentary sequence is the target for the biomarker reconstructions here.

The modern lake water balance of Great Salt Lake includes inflow from rivers (66%), precipitation (31%), and groundwater (3%) (Jones et al., 2009). The water leaves by evaporation from the closed basin (Jones et al., 2009), although recent work revises groundwater inflow up to 10-12% (Bunce et al., 2022). Instrumental records from 1966 to present indicate 7 m of fluctuation in lake levels (Rupke and McDonald, 2012; United States Geological Survey, 2022b) due to the changing water balance. During the high stand of the last glacial maximum/Heinrich Stadial 1 (HS1), GSL occupied a much greater areal extent, up to ~52,000 km², forming freshwater Lake Bonneville (Baxter et al., 2005; Oviatt et al., 2021). Continued downcutting at the outflow point eventually caused rapid overflow during the Bonneville flood, dropping lake levels by 100 m (Oviatt et al., 2021). Since then, the drier climate caused continued lake shrinkage to modern levels by the start of the Holocene, leading to the formation of the Bonneville Salt Flats (west of the present-day GSL) and a reduction of lake size to <10% of its former extent.

Riverine inflow to GSL primarily comes from the Bear (32.5 ± 34.4 m³/s), Jordan (3.5 ± 1.2 m³/s), and Weber Rivers (8.1 ± 15.9 m³/s; all reported as 2000-2020 annual mean discharge). The Bear River records low discharge (2-5 m³/s) in the dry summer months from July to September, with flow rising consistently during the fall to a plateau of 20-50 m³/s in January and February. Peak flows occur in March to April, reaching >50 m³/s, and then drops back down to 2-5 m³/s by June (United States Geological Survey, 2022a).

The modern GSL is separated into a northern and southern section by a railroad causeway built in 1959. As rivers exclusively flow into the South Arm, salinities are 120-180 g/L, whereas the North Arm has salinities of 290-340 g/L and a permanent halite bottom crust (data for 2010-2021; Jagmiecki et al., 2021; Rupke and McDonald, 2012). Salinity is heterogeneous around the shallow lake margins with freshwater influence near river inflows during peak discharge associated with times of snowmelt and evaporative enrichment along other margins, especially during summer and drought. At times in the instrumental record, a halocline has developed in the south arm, with higher salinities at depth and fresher conditions at the surface, but it is currently well mixed (Rupke and McDonald, 2012). While the North Arm is a restricted evaporation pond, the South Arm has major river inflow and is more representative of the pre-causeway GSL (Belovsky et al., 2011).

GSL has fluctuated between 7.6 m and 13.7 m deep over the historical record (1847-present) (Belovsky et al., 2011) with fluctuations during the seasonal cycle of summer evaporation and spring snow melt, interannual variability, and recent declining trends. In August 2022, GSL lake levels reached a historic low at just 6.8 m deep, with a surface elevation of 1,277 m, 3 m below the historic average (Ramirez, 2022). Current drought and diversion of inflowing rivers primarily for agriculture are projected to lead to further declines in lake levels in the coming decades (Wurtsbaugh et al., 2017). The decreasing areal extent of GSL is leaving infrastructure at abandoned shorelines and there are growing concerns over health effects associated with exposed lake bed dust (Null and Wurtsbaugh, 2020; Perry et al., 2019). The increase in salinity is projected to negatively impact the lake ecosystem, which will be detrimental to migratory shore and water birds (Jewell, 2021; Sorensen et al., 2020). Changing salinities and lake access will also affect existing resource extraction, including the brine shrimp cyst and mineral extraction industries (Naftz, 2017; Wurtsbaugh et al., 2017). This study seeks to generate Holocene paleolimnological reconstructions that extend beyond the instrumental record to provide longer baselines against which current trends can be compared.

2. Age model

Bowen et al. (2019) measured the radiocarbon of various materials (including brine shrimp cysts, algal mats, *n*-alkanes, carbonate, total organic carbon, and terrestrial macrofossils) from the GSL

sediment core GLAD1-GSL00-1B and formulated a radiocarbon age model based upon the cysts only ($n = 15$ dates). While some components are older (carbonates) or younger (charcoal), we are interested in all the organics that formed in the lake waters as age control for the lake biomarker-based record. In addition to cysts, the short chain *n*-alkanes, algal mats, and total organic carbon (TOC) all appear to form in lake waters. Thus, these additional age constraints are used to inform a revised age model here ($n = 27$ dates, **Figure 2a**), which is modestly different from the prior age model (**Figure 2b**). The revised age model is retained, as it has the most constraints. Despite producing older ages (≤ 300 years) between 5.5 and 3 ka (**Figure 2c**), this is trivial for the 220-year resolution of this study but represents uncertainty on the timing of the 5.5 ka transition identified in this study.

3. Materials and methods

3.1. Lipid extraction

We subsampled sediment core GLAD1-GSL00-1B at 30 cm intervals (approximately every 220 yrs) with each sample spanning ~ 1 cm (integrating ~ 7 yr) and samples were stored frozen at -20°C until analysis. Sediment samples (~ 10 g) were freeze-dried then homogenized with a mortar and pestle. Total lipid extracts (TLE) were obtained for each sample by immersion in 9:1 dichloromethane (DCM):methanol (MeOH) at 100°C and 1.034×10^4 kPa for 15 min using a Dionex ASE 350 Accelerated Solvent Extraction at the University of Southern California.

3.2. GDGT quantification

TLEs were dissolved in 40 μL 99:1 hexane:isopropanol and filtered through 0.45 μm PTFE (polytetrafluoroethylene) filters. 10 μL (or $\frac{1}{4}$ of the sample) was injected and archaeol, isoGDGTs, and brGDGTs were separated using an Agilent 1260 Infinity high-performance liquid chromatograph (HPLC) coupled to an Agilent 6120 single quadrupole mass spectrometer with two Ethylene Bridged Hybrid (BEH) Hydrophilic Interaction (HILIC) silica columns (2.1 mm \times 150 mm, 1.7 μm ; Waters) in series at 30°C following the methods of Hopmans et al. (2016). The mobile phase is hexane:isopropanol with a linear gradient transition between 98.2:1.8 for 25 mins, 96.5:3.5 for 25 mins, 90:10 for 50 mins, and reverting to the initial mixture for 20 mins, for a total of 120 minutes at a flow rate of 0.20 mL min^{-1} . Ionization was via positive ion atmospheric pressure chemical ionization (APCI) and detection was in selective ion

monitoring (SIM) mode, targeting the m/z for archaeol (653.8), isoGDGTs (1302.3, 1300.3, 1298.3, 1296.3, 1292.3), and brGDGTs (1050.0, 1048.0, 1046.0, 1036.0, 1034.0, 1032.0, 1022.0, 1020.0, 1018.0) at a mass tolerance of 1 dalton. Integration of peak areas was done using the MATLAB package ORIGAmI and each compound was quantified by comparison to the peak area for 500 ng of a C₄₆ internal standard (m/z 743.8) from the University of Arizona (Huguet et al., 2006). Six samples were rerun in SIM mode with an additional target to detect extended archaeol (m/z 723.8), injecting 10 μ L out of 200 μ L as it is an abundant compound, but otherwise using the same conditions. We compared the raw concentrations of summed branched and summed isoprenoidal compounds (Σ isoGDGT and Σ brGDGT) in ng/g sediment, to assess the productivity of each compound class. Here, we also include 7-methyl isomers (isomers with methyl groups bonded in the C₇ position) in the Σ brGDGT calculations due to their high abundance in this hypersaline environment (Wang et al., 2021). An example chromatogram is shown in **Figure S1**.

The ACE index, a paleosalinity proxy, was calculated using the following equation from Turich and Freeman (2011):

$$ACE = \frac{archaeol}{archaeol + GDGT-0} \times 100 \quad (1)$$

ACE values were converted to salinity according to the calibration of Turich and Freeman (2011), where:

$$ACE = 0.35 \times salinity - 5.4 \quad (2)$$

We also monitored for extended archaeol, biomarker specific to halophilic *Euryarchaeota*, in a subset of samples (Bale et al., 2019; Vandier et al., 2021) and calculate the ratio R_{EA} following Vandier et al. (2021):

$$R_{EA} = \frac{extended\ archaeol}{archaeol + extended\ archaeol} \quad (3)$$

The MBT'_{5Me} index, a brGDGT temperature proxy, was calculated according to De Jonge et al. (2014a):

$$219 \quad MBT'_{5Me} = \frac{Ia+Ib+Ic}{Ia+Ib+Ic+IIa+IIb+IIc+IIIa} \quad (4)$$

220 where I-III indicates the number of methyl groups (4-6), and a-c the number of cyclopentane
 221 rings (0-2). While MBT'_{5Me} is most frequently used as an indicator for temperature, recent work
 222 by Martinez-Sosa et al. (2021) and Wang et al. (2021) found evidence that salinity, particularly
 223 hypersalinity, may bias temperature estimates. We convert the MBT'_{5Me} values to mean annual
 224 temperature for months above freezing (MAF) using BayMBT₀, the Bayesian calibration of
 225 Dearing Crampton-Flood et al. (2020) available for both global surface soils and lake sediments.
 226 Martinez-Sosa et al. (2021) suggested that alkaline, saline lakes may receive greater brGDGT
 227 input from surrounding soils than from in lake production. Thus, we test both the soil and lake
 228 calibrations here.

229 The CBT' index is used as a pH indicator and was also calculated following the equation of De
 230 Jonge et al. (2014a):

$$231 \quad CBT' = \log_{10} \left(\frac{Ic+IIa'+IIb'+IIc'+IIIa'+IIIb'+IIIc'}{Ia+IIa+IIIa} \right) \quad (5)$$

232 where ' denotes structural isomers with methyl groups bonded in the C₆ rather than C₅ position.
 233 We calculated pH values using the global soils and lakes calibrations of Raberg et al. (2022):

$$234 \quad CBT' = 0.50 \times \text{soil pH} - 3.65 \quad (6)$$

$$235 \quad CBT' = 0.23 \times \text{lake pH} - 1.98 \quad (7)$$

236 As the CBT' formulation in Raberg et al. (2022) included a negative sign, both calibration
 237 equations were adjusted to maintain the positive formulation of CBT' (equation 5) as originally
 238 defined by De Jonge et al. (2014a).

239 The IR_{6Me} index is the ratio of 6-methyl brGDGT isomers to the sum of 5- and 6-methyl
 240 brGDGT isomers, calculated according to De Jonge et al. (2014b):

$$241 \quad IR_{6Me} = \frac{IIa'+IIb'+IIc'+IIIa'+IIIb'+IIIc'}{IIa+IIb+IIc+IIIa+IIIb+IIIc+IIa'+IIb'+IIc'+IIIa'+IIIb'+IIIc'} \quad (8)$$

242 The fractional cyclization (fC) index shows the proportion of brGDGTs with cyclopentane rings
 243 and is calculated according to Martinez-Sosa and Tierney (2019):

$$244 \quad fC = \frac{\Sigma(b)+2\Sigma(c)+\Sigma(b')+2\Sigma(c')}{\Sigma(a)+\Sigma(b)+\Sigma(c)+\Sigma(b')+\Sigma(c')} \times 0.5 \quad (9)$$

245 The branched and isoprenoidal tetraether (BIT) index measures the proportion of branched to
 246 isoprenoidal inputs, with the isoprenoidal inputs represented only by crenarchaeol. BIT is
 247 calculated using the equation of Hopmans et al. (2004):

$$248 \quad BIT = \frac{Ia+IIa+IIa'+IIIa+IIIa'}{Ia+IIa+IIa'+IIIa+IIIa'+cren} \quad (10)$$

249 However, BIT is limited in environments where crenarchaeol is a minor constituent; in such
 250 cases it may be more useful to compare the summed branched and isoprenoidal compounds using
 251 the ratio $R_{i/b}$:

$$252 \quad R_{i/b} = \frac{\Sigma isoGDGT}{\Sigma brGDGT} \quad (11)$$

253 BIT was originally developed as a measure of terrestrial GDGT input to marine environments,
 254 but recent work by Xiao et al. (2016) introduced the $\Sigma IIIa/\Sigma IIa$ ratio as a measure solely of
 255 brGDGT sourcing without including crenarchaeol:

$$256 \quad \Sigma IIIa/\Sigma IIa = \frac{IIIa+IIIa'}{IIa+IIa'} \quad (12)$$

257 Martin et al. (2020) and Ramos-Roman et al. (2022) later extended this to lacustrine
 258 environments and included the 7-methyl isomers (denoted by '''):

$$259 \quad \Sigma IIIa/\Sigma IIa = \frac{IIIa+IIIa'+IIIa'''}{IIa+IIa'+IIa'''} \quad (13)$$

260 4. Results

261 We measured the individual isoGDGT and brGDGT abundances along with archaeol abundance
 262 (data available at NOAA; So et al., 2022). IsoGDGTs are dominated by GDGT-0 (**Figure 3a**)

and we show the distribution of the remaining isoGDGTs (**Figure 3b**). Σ isoGDGT and Σ brGDGT were similar in magnitude (**Figure 3c**). Σ isoGDGT values had a mean of 168 ng/g ($1\sigma = 56$ ng/g) and Σ brGDGT values had a mean of 117 ng/g ($1\sigma = 39$ ng/g). Only 5 out of the 35 samples had Σ brGDGT exceeding Σ isoGDGT (by <30 ng/g).

The salinity proxy, the Archaeol and Caldarchaeol Ecometric (ACE), remained relatively invariant throughout the record with a mean of 90.5 ($1\sigma = 3.7$) from 7.2 ka to present; Z-scores range from -2.5 to 2.0 (**Figure 4a**). Statistical analysis using the R Package EnvCpt (Beaulieu and Killick, 2018) for environmental time series changepoint detection confirms a single changepoint in the mean and variance for the MBT'_{5Me} index between a mean of 0.49 ($1\sigma = 0.05$) before 5.5 ka to a mean of 0.67 ($1\sigma = 0.05$) from 5.5 ka to present (**Figure 4b**, **Figure S2b**). The same changepoint analysis identified shifts at the same time in other indices as follows. IR_{6Me} values showed a small increase from 0.61 ($1\sigma = 0.02$) before 5.5 ka to 0.69 ($1\sigma = 0.03$) after (**Figure 4c**, **Figure S2c**). The fC index shifted from 0.23 ($1\sigma = 0.02$) before 5.5 ka to 0.15 ($1\sigma = 0.03$) after (**Figure 4d**, **Figure S2d**). The Σ IIIa/ Σ IIa ratio including 7-methyl isomers remained constant over 7.2 ka (**Figure 4e**), averaging 0.84 ($1\sigma = 0.13$). However, when 7-methyl isomers are excluded, a step shift at 5.5 ka is seen, with Σ IIIa/ Σ IIa averaging 0.64 ($1\sigma = 0.08$) pre-shift and 0.38 ($1\sigma = 0.06$) post-shift (**Figure S2e**). Additional indices were relatively invariant. CBT' values averaged -0.01 ($1\sigma = 0.08$) across the record (**Figure S3c**). The BIT index was high and invariant across 7.2 ka (**Figure S3f**), with a mean of 0.95 ($1\sigma = 0.02$). $R_{i/b}$ averaged 1.51 ($1\sigma = 0.05$) across 7.2 ka, indicating archaeal isoGDGTs are generally more abundant than bacterial brGDGTs (**Figure S3g**). As ACE was relatively insensitive, we measured extended archaeol in six samples. We found R_{EA} was as high as 40% prior to 5.5ka (mean 20%, $1\sigma = 12\%$) and consistently low after 5.5 ka (mean 6%, $1\sigma = 3\%$) (**Figure 4a**).

5. Discussion

5.1. GSL Holocene paleosalinity

Measured ACE values for the Holocene GSL interpreted with the calibration of Turich and Freeman (2011) indicate a mean salinity of 274 g/L ($1\sigma = 11$ g/L) with <50 g/L of variability for the whole record. Thus, from 7.2 ka to present, we find GSL to be hypersaline and its salinity to be relatively invariant (**Figure 5**). The low variability (<50 g/L) likely results from applying the

ACE index at its upper limits. With all samples having ACE values >80 (almost all archaeol with little GDGT-0), the index would have limited sensitivity to variations at such high salinities. At deeper depths in the same GLAD core that extends through Pleistocene pluvials, the ACE proxy could be applied to study a larger range of salinities. However, in the Holocene hypersaline terminal lake system, we approach the upper limit of the index.

Modern lake measurements indicate the salinity of the North Arm brine is over 300 g/L, and the South Arm brine is close to 140 g/L (the latter being more reflective of GSL prior to causeway separation). Historical measurements (data for 1966-2020, Rupke and McDonald, 2012) indicate the salinity range of surface samples and depth profiles in the North Arm is 250-340 g/L and the South Arm is 100-220 g/L. Although the available ACE calibration (Turich and Freeman, 2011) suggests salinities in the range of 248-295 g/L during the mid to late Holocene, these estimates appear too high, as this is higher than the modern and historical measurements for the South Arm, with such values found only in the North Arm today. Overestimates of salinity using ACE (values ranging 14-67) have previously been reported in the Piedmont Basin (Italy), which was deposited at the onset of the Miocene Messinian salinity crisis (Natalicchio et al., 2017). This was attributed to archaeol and GDGT-0 production by methanotrophic *Euryarchaeota* and *Thaumarchaeota* in addition to halophilic archaea. In the Holocene GSL, the *Euryarchaeota* are likely producers based on high GDGT-0 and low crenarchaeol (**Figure 3a**).

Absolute salinities are therefore approximations given the limited datasets for calibration and the lack of laboratory quantification standards for each of the analytes. Turich and Freeman (2011) noted that HPLC-MS measurements made in different laboratories have differential ionization of archaeol and GDGT-0, which Natalicchio et al. (2017) also noted. Given the present lack of robust quantification and comparability, we interpret the ACE index as a qualitative indicator of salinity. In addition to relative response factors, the different ecologies of different aquatic communities and their limited calibration each affect the quantitative interpretation of the proxy at present. We can rule out low GDGT-0 relative response factors based on analyses with the same instrument conditions in other sedimentary archives across the full range of the ACE index (Feakins et al., 2019; Peuple et al., 2021). In the Holocene GSL, we may expect some archaeol and GDGT-0 production by halophilic methanogens but not *Thaumarchaeota* given the high GDGT-0/crenarchaeol ratios (based on the inferences of Natalicchio et al. (2017)). Instead, the

dominant producers are likely methanogenic *Euryarchaeota*, which can be indicative of stratified waters (Baxter et al., 2021; Blaga et al., 2009; Naeher et al., 2014). The narrow range and consistently high calculated ACE values of 80-100 suggest limited microbial ecology variability or a loss of sensitivity to salinity variations near the index maximum.

We can also test the ACE salinity proxy through comparison to other evidence for hypersalinity. Bowen et al. (2019) found brine shrimp cysts throughout the core section. In modern GSL, brine shrimp are abundant in the south arm and previous studies have identified the optimum salinity range for these shrimp to be 120-160 g/L, below which brine shrimp are limited by predation and above which by physiological stress (Great Salt Lake Salinity Advisory Committee, 2021). The presence of cysts seems to indicate ACE overestimates salinity. However, there are likely spatial and temporal differences between ACE and cysts as salinity recorders as we will explore. Brine shrimp live in shallow, marginal shore environments while their cysts float near the surface and can be transported/deposited in deeper parts of the lake. In comparison, halophilic archaea can be found throughout the water column and may record salinities at different depths including the saltier deep water at density stratified times. Seasonally, brine shrimp become very productive in late spring when temperatures reach 18-20°C while archaea may be most productive during the warm summer. Thus, high ACE values may not indicate a calibration problem for this system and instead the mismatch between high ACE and cyst may reflect seasonally varying or depth stratified salinity.

A biomarker more specific to halophilic archaea, but that has not yet been developed into a quantitative proxy, is extended archaeol (Bale et al., 2019; Vandier et al., 2021). We measured extended archaeol for six samples (three pre-5.5 ka, three post-5.5 ka) and detect this compound in all six. We calculate R_{EA} , defined by Vandier et al. (2021) as the ratio of extended archaeol to the sum of extended archaeol and archaeol, and find R_{EA} to range from 3-30%, with the pre-5.5 ka average being 20% ($1\sigma = 12\%$) and the post-5.5 ka being 6% ($1\sigma = 3\%$) (**Figure 4a, 5b**). The presence of this compound confirms the presence of abundant halophilic *Euryarchaeota*. Vandier et al. (2021) broadly identifies an increase in R_{EA} with an increase in salinity and this, together with generally higher R_{EA} pre-5.5 ka, may suggest higher salinities during the arid mid-Holocene. However, as extended archaeol is a relatively novel compound and its abundance

cannot currently be directly converted to salinity values, further studies are needed to validate this interpretation.

Overall, both high ACE values, constant brine shrimp cyst presence, and high quantities of extended archaeol indicate GSL hypersalinity over the past 7.2 ka. Given that GSL ACE values extend beyond the upper limits of the calibration of Turich and Freeman (2011) and the lack of an applicable local calibration, the calculated salinities may be overestimates and we consider reconstructions to be semi-quantitative using existing ACE calibrations. The limited variability detected by ACE suggests a role for an additional hypersalinity marker, extended archaeol to discern variations in extreme salinity in this setting.

5.2. MBT'_{5Me} temperature record

Using the BayMBT₀ lake calibration, the mean annual temperature of months above freezing (MAF) averaged 18°C (1σ = 3°C) throughout the record, with a mean of 15°C (1σ = 2 °C) before 5.5 ka and 20°C (1σ = 2 °C) after (**Figure 6**). Lower MAF estimates resulted from the soil calibration, averaging 14°C (1σ = 3°C) throughout the record, 10°C (1σ = 2°C) before 5.5 ka, and 15°C (1σ = 2°C) after 5.5 ka. After the step change at 5.5 ka, the soil-based calibration yields MAF estimates that closely match those of present-day Salt Lake City (15.7°C; National Centers for Environmental Information, 2022), whereas the lake-based calibration yielded temperatures that were on average 4°C too warm. A warm bias for saline lakes had previously been reported from the global lakes calibration dataset including nearby Mono Lake, California (Martinez-Sosa et al., 2021).

Reconstructed temperatures apparently increased by 5°C at 5.5 ka (**Figure 6**). Temperatures reconstructed using the soil calibration before 5.5 ka were much cooler than expected when compared against modern temperatures, whereas using the lake calibration temperatures were similar to modern. Previous climate records suggest the mid-Holocene to be a warm, arid period relative to the late Holocene and no known climatic processes would support a step change in MAF of that magnitude at that time (see **Section 5.4**). As we also detect a step shift at 5.5 ka in several GDGT indices (see **Section 5.3** and **Figure S2**), it is likely that non-thermal changes in GSL at 5.5 ka affected the MBT'_{5Me} proxy, whether through a change in the microbial community between a saltier, shallower lake and a deeper lake with increased overturning and

bottom water oxygenation, and/or a shift in the productivity within the lake, and/or changes in the influx of microbial lipids from surrounding soils. We do find evidence from extended archaeol relative to archaeol that the pre-5.5 ka GSL was saltier (**Figure 5b**). We also find an increase in IR_{6Me} , decrease in fC , and decrease in $\Sigma IIIa/\Sigma IIa$ excluding 7-methyl isomers, which also point to a shift in the microbial community and their lipids within the lake. Whereas exploration of soil versus lake provenance is, to some extent, inconclusive (see supplementary), some of the questions remain unclear and signal a need for more regional calibration of microbial communities in saline lake systems. While we are not certain about the nature of the microbial community change, we see signs of non-thermal factors and we do not interpret the temperatures for 7.2-5.5 ka (**Figure 6**, yellow shading).

5.3. Limnology shift at 5.5 ka

Although the ACE index may be relatively insensitive to salinity change in the last 7.2 ka given the hypersalinity of the lake, other biomarker indices show evident changes. Changepoint analysis revealed a shift in several GDGT indices that indicate shifts in limnological conditions at 5.5 ka (MBT'_{5Me} , IR_{6Me} , fC , and $\Sigma IIIa/\Sigma IIa$) (**Figure 4**, **Figure S2**). We see an increase in MBT'_{5Me} values at 5.5 ka (**Figure 4b**). As MBT'_{5Me} is a paleotemperature proxy, these values alone would indicate increased temperatures. The increase in IR_{6Me} at this time means a proportional increase in 6-methyl brGDGT isomers (**Figure 4c**). Greater proportions of 6-methyl isomers have been linked to higher pH (De Jonge et al., 2014a; Raberg et al., 2021; Raberg et al., 2022), comparatively more aquatic production than soil production (De Jonge et al., 2014b; Kirkels et al., 2020), lower soil water content (Dang et al., 2016), and/or higher conductivity (Raberg et al., 2021). A decrease in fC is a decrease in the proportions of cyclized brGDGTs (**Figure 4d**). Less cyclization of brGDGTs has been linked to lower pH/conductivity (Raberg et al., 2021), and/or comparatively more soil production (Kirkels et al., 2020). We also see a step decrease at 5.5 ka in $\Sigma IIIa/\Sigma IIa$ values when 7-methyl isomers are excluded (**Figure 4e**). Lower $\Sigma IIIa/\Sigma IIa$ values indicates a lower proportion of brGDGTs with six methyl groups compared to those with five methyl groups and has been linked to increased soil inputs to lacustrine environments (Martin et al., 2020; Ramos-Roman et al., 2022). However, when 7-methyl isomers are accounted for, the $\Sigma IIIa/\Sigma IIa$ ratio remains relatively constant. Principal component

analysis (PCA) of GSL samples further supports a shift in lake conditions at 5.5 ka with samples pre- and post-5.5 ka plotting in separate clusters (**Figure S4a**).

MBT'_{5Me} values are low prior to this shift and, using the soil calibration, yield unrealistically cold temperatures (**Figure 6**). Lake production of brGDGTs has a 'colder' distribution than soils (Martinez-Sosa et al., 2021), thus a shift to *in situ* aquatic production might explain the observed pattern. The values pre-5.5 ka may instead represent more *in situ* lake production of brGDGTs during the middle Holocene, making it more appropriate to use the lake calibration and this seems to be supported by lake calibration temperatures matching more closely to modern Salt Lake City temperatures during this time. One possibility is that lower lake levels prior to 5.5 ka combined with arid conditions restricted river inflows that would have brought soil-produced brGDGTs to GSL. Mixed brGDGT sourcing is possible for Holocene GSL based on its distributions of tetra-, penta-, and hexamethylated brGDGTs (Ramos-Roman et al., 2022; Russell et al., 2018) falling between those of global lakes and soils (**Figure S5, S6**), although neither pre- nor post-5.5 ka distributions can be unambiguously ascribed to a purely lake-like or soil-like signal compared to large global datasets (**Figure S4b, S6**). Thus, we do not find strong support for a shift from lake to soil-derived compounds.

Part of the ambiguity may be due to the uniqueness of GSL. PCA analysis shows that GSL has an extreme brGDGT distribution compared to global distributions, especially prior to 5.5 ka (**Figure S4b**). Hypersaline lakes differ in their chemistry and are isolated bodies of water which may have fundamentally different microbial communities and/or lipid responses to environmental change than their freshwater counterparts or other saline lakes. Wang et al. (2021) noted warm-biased MBT'_{5Me} values with increasing salinity. In contrast to Martinez-Sosa et al. (2021) who had attributed this warm bias to soil inputs, Wang et al. (2021) attributed this to increased production of 7-methyl penta- and hexamethylated isomers in place of 5-methyl ones by lake microbes. This would explain the temperature overestimates we observe when using the BayMBT₀ lake calibration post-5.5 ka (**Figure 6**). Overall, both studies indicate warm biases in MBT'_{5Me} in saline lakes, although the variations need additional study.

Across the 5.5 ka transition, IR_{6Me} and fC increase and decrease respectively (**Figure 4c, 4d**). pH estimated from CBT' using the soil calibration of Raberg et al. (2022) averaged 7.2 (1 σ = 0.02)

and using the lake calibration averaged 8.6 ($1\sigma = 0.03$). Both calibrations yield near-constant pH values throughout the whole record meaning shifts in IR_{6Me} and fC are likely not due to changes in pH. Pre-5.5 ka, higher fC could indicate higher conductivity which may reflect higher temperature and/or higher salinity, both of which align with middle Holocene aridity (see **Section 5.4**). Lower IR_{6Me} values are harder to reconcile as they would suggest lower conductivity, opposite that of fC. Aside from conductivity, low IR_{6Me} may otherwise indicate higher soil water content and/or a greater influx of organic material produced in soils. However, these factors seem secondary to pH and conductivity as controls of IR_{6Me} . Additionally, an interpretation of high soil water content or soil production would contrast with both regional climate records, which highlight mid-Holocene aridity, and brGDGT provenance studies, which do not suggest shifts from lake to soil sourcing. Due to the large number of factors with potential to influence IR_{6Me} at this site, we do not interpret the exact environmental implications of the IR_{6Me} step change.

The decrease in $\Sigma IIIa/\Sigma IIa$ values with the exclusion of 7-methyl isomers would suggest greater soil input after 5.5 ka, but constant values when these isomers are included indicate otherwise (**Figure 4e**). Throughout the 7.2 ka record, the quantity of 7-methyl isomers is comparable to or exceeds their 5- and 6-methyl counterparts. Wang et al. (2021) reported high abundances of 7-methyl isomers in hypersaline lakes and low amounts in freshwater lakes. The high abundance of 7-methyl isomers at hypersaline GSL may explain the trends in the different formulations of $\Sigma IIIa/\Sigma IIa$ (**Figure 4e**). It is likely that 7-methyl brGDGTs are produced within the hypersaline lake and their high abundance results in differing $\Sigma IIIa/\Sigma IIa$ trends when these isomers are included or excluded. The coincident changepoint in four GDGT indices (MBT'_{5Me} , IR_{6Me} , fC, and $\Sigma IIIa/\Sigma IIa$) (**Figure 4, Figure S2**) suggests a change in both the bacterial and archaeal communities suggesting that the microbial and limnological conditions of GSL pre-5.5 ka are distinct from post-5.5 ka.

Our biomarker reconstruction detects a changepoint in GSL limnology at 5.5 ka in several metrics and those non-thermal effects appear to disrupt the temperature proxy. Though ACE shows that GSL is hypersaline throughout the 7.2 ka record, both high ACE and extended archaeol show that the mid-Holocene lake was saltier and shallower than the subsequent 5.5 ka.

We infer that the shallower, saltier mid-Holocene lake hampered paleothermometry in this time, while the late Holocene temperature reconstructions appear reasonable.

5.4. Comparison to regional climate

The reconstruction of shallower, saltier (**Figure 7a**) GSL in the mid-Holocene is consistent with evidence for regional aridity. In Lehman Caves (NV) located 260 km southwest of GSL, a decrease in speleothem Mg/Ca shows increased aridity at ~8 ka (Steponaitis et al., 2015) (**Figure 7b**). In nearby Leviathan Cave (NV) 180 km southwest of GSL, a decrease in speleothem $\delta^{13}\text{C}$ values at ~4 ka is interpreted as an increase in effective moisture (Lachniet et al., 2020) (**Figure 7c**). Collectively, these cave records bracket a mid-Holocene arid period from around 8–4 ka. A proxy synthesis and reanalysis effort across western North America (Routson et al., 2022) identifies a mid-latitude moisture increase in this region after 6 ka (**Figure 7d**). Preceding this moisture increase, we observe low lake levels in many Great Basin lakes (Steponaitis et al., 2015) (**Figure 7e**), including Owens Lake (CA) (Bacon et al., 2020), Lake Tahoe (CA/NV) (Benson et al., 2002), and Pyramid Lake (NV) (Benson et al., 2002), as well as at small lakes in the Midwest (Shuman et al., 2002) and in the sub-alpine region of the Rocky Mountains (Shuman and Serravezza, 2017). A compilation of southwestern US climate records by Lachniet et al. (2020) also found peak aridity at 7.5 ka (**Figure 7f**). Thus, it seems likely that widespread mid-Holocene aridity in mid-latitude western North America contributed to a shallower GSL recorded from the start of this record at 7.2 ka to 5.5 ka, with a shift to wetter conditions represented locally at 5.5 ka.

Most of the precipitation in the Great Basin falls during the winter with moisture delivered by westerlies from the Pacific, and a much smaller fraction falls during the summer as part of the North American Monsoon (NAM) with moisture from the Gulf of California and eastern Pacific (Lachniet et al., 2020). Moisture from the Gulf of Mexico can also contribute to the NAM, but this rarely reaches the Great Basin (Jana et al., 2018). Changes in the incursion of the NAM and the position of westerlies over western North America have been inferred from the reanalysis of a large network of proxy data (Routson et al., 2022). They identified a strong NAM that peaked at 7 ka, coincident with weakened/northward-deflected westerlies. The latter could explain the dry mid-Holocene in the mid-latitudes, including at GSL from 7.2 to 5.5 ka, as it would have

decreased delivery of Pacific moisture. Following this, in the late Holocene, a reduction in the NAM and more zonal westerly flow, resulted in wetter conditions in western North America including at GSL.

Routson et al. (2022) suggested that these changes in circulation were linked to meridional temperature gradients. Changes in broad scale atmospheric circulation and Pacific SSTs due to high summer insolation and Laurentide ice sheet (LIS) collapse have previously been connected to Holocene shifts in the precipitation regime over the western US. Summer northern hemisphere insolation peaked in the Early Holocene ~11-10 ka and has steadily decreased up to the present (Lachniet et al., 2020). In the same time period, the LIS rapidly retreated from 10-8 ka (Shuman et al., 2002), mostly having disappeared by ~7 ka (Peltier et al., 2015). Steponaitis et al. (2015) attributed the onset of mid-Holocene aridity in the Great Basin at ~8 ka to a northward displacement of winter storm tracks after the collapse of the remnant LIS. In contrast, Lachniet et al. (2020) suggested summer insolation to be the primary driver of regional climate variation. They find a connection between southwestern US drought and increased SSTs in the western tropical Pacific (WTP), warmth in the Arctic, and low Arctic sea ice, these phenomena being caused by high summer insolation. Both Steponaitis et al. (2015) and Lachniet et al. (2020) also suggest that a La Niña-like configuration of the tropical Pacific (a response to orbital forcing (Brierley et al., 2020) may have contributed to regional aridity.

Changes in summer insolation and the LIS have also been linked to hydroclimate changes in other regions of the US. Lake and pollen records indicate a moist to dry transition in the Midwest, and a dry to moist transition in the northeast and southeast between 9 and 8 ka, attributed to the collapse of the LIS Hudson Bay Dome and the subsequent proportional increase in the influence of summer insolation (Shuman et al., 2002). In the northeast, plant wax isotopes show an increased ratio of summer to winter precipitation after 8 ka, as the LIS glacial anticyclone previously prevented northward transport of subtropical moisture during the summer (Shuman et al., 2006). From ~9-7 ka, lakes in California, the Great Basin, and the Rocky Mountains declined while those in western Canada rose. These trends reversed by ~5.7 ka, possibly due to the decline in summer insolation after the LIS melted (Shuman and Serravezza, 2017). Similarly, between 5.7-5.2 ka, lakes across central and eastern mid-latitude North America show an increase in effective moisture, which terminated mid-Holocene aridity and

coincided with the end of the Holocene temperature maximum in this region (Shuman and Marsicek, 2016). A global compilation of Holocene temperature proxies (“Temperature 12K”) including their age model uncertainties, places the central estimate for mid-Holocene warmth around 6.45 ka within a range of 7.65–4.95 ka (Kaufman et al., 2020).

Model studies also suggest similar patterns of mid-Holocene aridity driven by large scale climatic processes. Diffenbaugh et al. (2006) found insolation to be the primary driver of North American midcontinent aridity and model efforts by Shin et al. (2006) suggested persistent La Niña-like Pacific SST patterns (i.e. warm western tropical Pacific, cool eastern tropical Pacific). Bhattacharya et al. (2018) found both models and proxy records from the Gulf of California region indicate a weakened NAM during the last glacial and NAM strengthening as the LIS retreated and westerly storm tracks, which previously mixed in cold, dry air that weakened the NAM, were displaced/weakened. While the LIS had largely disappeared by the middle Holocene, PMIP4-CMIP6 simulations still show enhanced seasonality and stronger monsoons in the northern hemisphere compared to preindustrial times (Brierley et al., 2020). The model dynamics support the proxy evidence for a strong NAM and weakened westerlies in the middle Holocene, leaving dry conditions in mid-latitude western North America (Routson et al., 2022). In the late Holocene, once NAM weakened, the more zonal westerlies (Routson et al., 2022) may have increased the moisture flux to the GSL region, resulting in the more mesic conditions experienced until recently.

The late Holocene was comparably wetter than the mid-Holocene, with lake high stands from ~4.5 to ~3 ka recorded at Owens Lake (CA) (Bacon et al., 2020; Benson et al., 2002), Walker Lake (NV) (Benson et al., 1991), Pyramid Lake (NV) (Briggs et al., 2005), Winnemucca Lake (NV) (Briggs et al., 2005), Mono Lake (CA) (Stine, 1990), and Silver Lake (CA) (Enzel et al., 1989). Although the biomarker record was stable without change points within the late Holocene, we do observe some variability within the GSL record. We note an ACE spike at 2.8 ka (**Figure 7a**) so this may have been a short lived drying in GSL. This may correspond to the onset of the Late Holocene Dry Period (2.8–1.85 ka) identified by Mensing et al. (2013) with lake low stands at Mono Lake (CA) (Stine, 1990), Walker Lake (CA) (Adams, 2007; Benson et al., 1991), Zaca Lake (CA) (Kirby et al., 2014), and Bear Lake (UT/ID) (Moser and Kimball, 2009). Tree ring studies are optimal for shorter timescales with limber pine assemblages in the western Great

Basin spanning the last 4000 years (Millar et al., 2019). Within the Late Holocene Dry Period, they found wet interludes occurred at the Neoglacial and Little Ice Age, whereas drought conditions stunted tree growth in the last 20 years, the second driest interval in the last 4000 years, surpassed only during 20 BC to 150 AD (Millar et al., 2019). The 220-year resolution of sampling in this study of GSL is not sufficient to contribute to finer records of hydroclimate (20-200 yr droughts). Although finer sampling is possible (samples span 1 cm, integrating ~7 yrs), there are limits to the resolution in lake sediments in a large lake with a large catchment, for example terrestrial plant wax, may experience considerable pre-aging on the landscape (Bowen et al., 2019) that may preclude detailed Holocene reconstructions. Longer records are the focus of future study as the GLAD drilling extended to 175 ka, capturing Pleistocene pluvials that may be resolved by the ACE salinity proxy which had approached its upper limit for the Holocene hypersaline terminal lake system.

6. Conclusions

We reconstructed a Holocene paleoclimate record for GSL using GDGT biomarker analyses of lacustrine sediments from the GLAD1-GSL00-1B sediment core spanning 7.2 ka to present. Although absolute salinity interpretations of the ACE index in GSL are uncertain, the index is near its upper limit ($>80/100$) suggesting that hypersaline conditions existed throughout the record. Applying the calibration of Turich and Freeman (2011), ACE values in these Holocene GSL sediments suggest salinities >250 g/L, though comparison to the present-day salinity range along with brine shrimp cyst evidence suggests this may be an overestimate, or represent spatial or temporal differences between the two recorders.

For the brGDGT thermometer, the BayMBT₀ soil calibration yielded MAF estimates consistent with modern Salt Lake City MAF from 5.5 ka to the present, whereas the lake calibration was warm-biased, consistent with reports of saline lake outliers in prior calibration studies (Martinez-Sosa et al., 2021). In contrast, the soil calibration yields unreasonably cold temperatures during the mid-Holocene while the lake calibration yields temperatures similar to modern MAF. Using either calibration alone, at 5.5 ka we find an apparent step shift in temperature of 5°C. However, such a large, abrupt warming is unlikely as the Holocene is a generally stable time. As we do not find provenance evidence to support a switch in calibration for the MBT'_{5Me} index, we maintain

interpretations of the late Holocene temperature records that are consistent with modern MAF, but we do not find temperature estimates to be secure for the mid-Holocene.

Changepoint analysis reveals that several other GDGT indices (IR_{6Me} , fC , $\Sigma IIIa/\Sigma IIa$ without 7-methyl isomers) also show a step shift at 5.5 ka strongly suggesting non-thermal effects on MBT'_{5Me} , likely a change in the lake depth influencing microbial communities before 5.5 ka. We find some support from extended archaeol measured on a subset of samples that found greater salinity before 5.5 ka. We interpret these data to indicate a shallow, saltier lake from 7.2 to 5.5 ka, in line with regional evidence for dry conditions during the mid-Holocene (Lachniet et al., 2020).

This paleolimnological study provides context for recent changes in the instrumental record, with a decline in lake level of 7 m from 1986 to 2022, which has reduced the surface area from 8550 km² to 2460 km² (Utah Department of Natural Resources, 2022) due to unsustainable water use and climate change (Wurtsbaugh et al., 2017). Our biomarker record extends the instrumental baseline against which the current drying can be compared, finding relatively stable conditions at GSL over the last 5.5 ka, with modern conditions approaching that of the mid-Holocene aridity. The mid-Holocene salty and shallow GSL is a prelude to the current changes associated with water extraction and drying.

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Supporting Information

Supporting Information may be found in the online version of this article.

Conflict of Interest

The authors declare no financial conflicts of interests for any author or their affiliations.

Open Research

Data files are archived at the NOAA paleoclimatology database (So et al., 2022).

Figure Captions

Figure 1. Maps showing a) the Great Salt Lake (GSL) core site of GLAD1-GSL00-1B and the Lake Bonneville shoreline (top right), and b) the locations of regional comparison records.

Figure 2. Age model based on all lake organic carbon radiocarbon dates (Bowen et al., 2019) from the GLAD core using the Bayesian R package BACON (Blaauw and Christen, 2011). a) Calibrated radiocarbon age probability distribution functions (blue), age model mean (red line) and uncertainty (gray shading), b) comparison between the age model mean (red line) for all lake organic radiocarbon dates (used in this study) and the cyst-only model (blue line) presented by Bowen et al. (2019), and c) their difference. The biomarker transition at 5.5 ka in the all-lake OC model is ~200 years older than in the cyst-only model.

Figure 3. Change in the distribution and concentration of isoGDGTs in the GSL over the 7.2 ka record, showing a) isoGDGT proportional abundances, b) isoGDGT proportional abundances without GDGT-0, and c) isoGDGT (black) and brGDGT (gray) concentration through time.

Figure 4. Temporal variations in GDGT indices showing a) ACE Z-score (blue) and R_{EA} (black cross), b) MBT'_{5Me} , c) IR_{6Me} , d) fC , and e) $\Sigma IIIa/\Sigma IIa$. We use statistical methods to identify a changepoint in several indices at 5.5 ka (**Figure S2**). The portion of the record prior to the step change is highlighted (yellow shading).

Figure 5. Salinity indicators in a 7.2 ka record from GLAD1-GSL00-1B core. a) ACE variations near the upper limit of the index. b) Calculated ACE Z-scores (blue line) and R_{EA} (black cross). Arrow indicates direction of salinity increase.

Figure 6. GSL mean annual temperature for months above freezing (MAF) calculated using the BayMBT₀ lake (blue) and soil (red) calibrations of Dearing Crampton-Flood et al. (2020) are shown along with one standard deviation uncertainty (light red envelope for soil calibration

temperatures, dashed blue lines for lake calibration temperatures). The black line indicates the Salt Lake City mean annual air temperature for months above freezing (MAF). Yellow shading indicates the period when the MBT'_{5Me} proxy was likely affected by different limnological conditions and microbial production, rather than a change in temperature.

Figure 7. Compilation of regional paleoclimate reconstructions, showing a) GSL ACE Z-score (this study), b) Lehman Caves (NV) Mg/Ca record (Steponaitis et al., 2015), c) Leviathan Cave (NV) $\delta^{13}\text{C}$ record (Lachniet et al., 2020), d) precipitation index for mid-latitude western North America (Routson et al., 2022), e) Great Basin lakes with low lake levels (Steponaitis et al., 2015); and f) aridity index for the southwestern US (Lachniet et al., 2020).

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Figure 1.

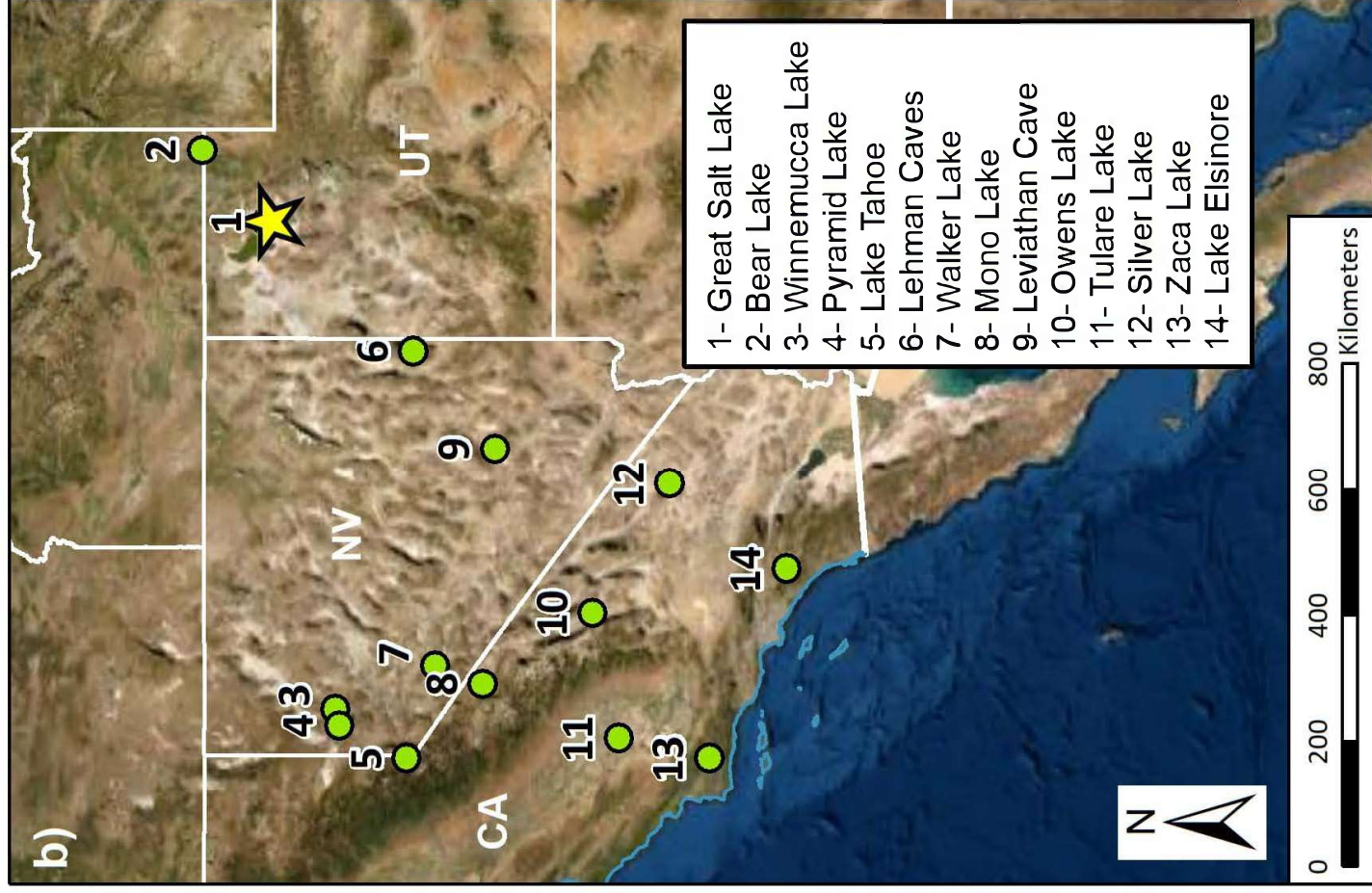
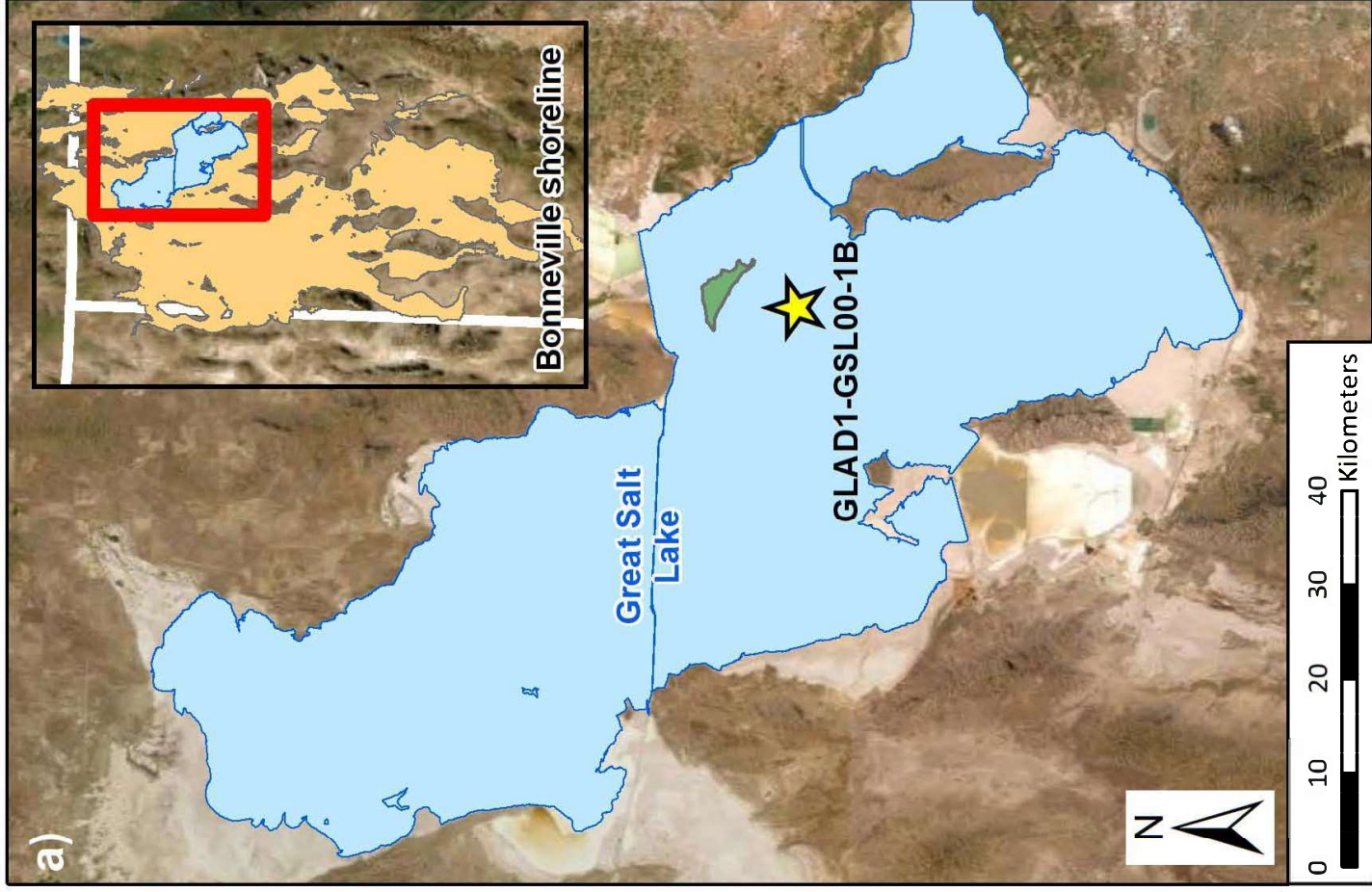


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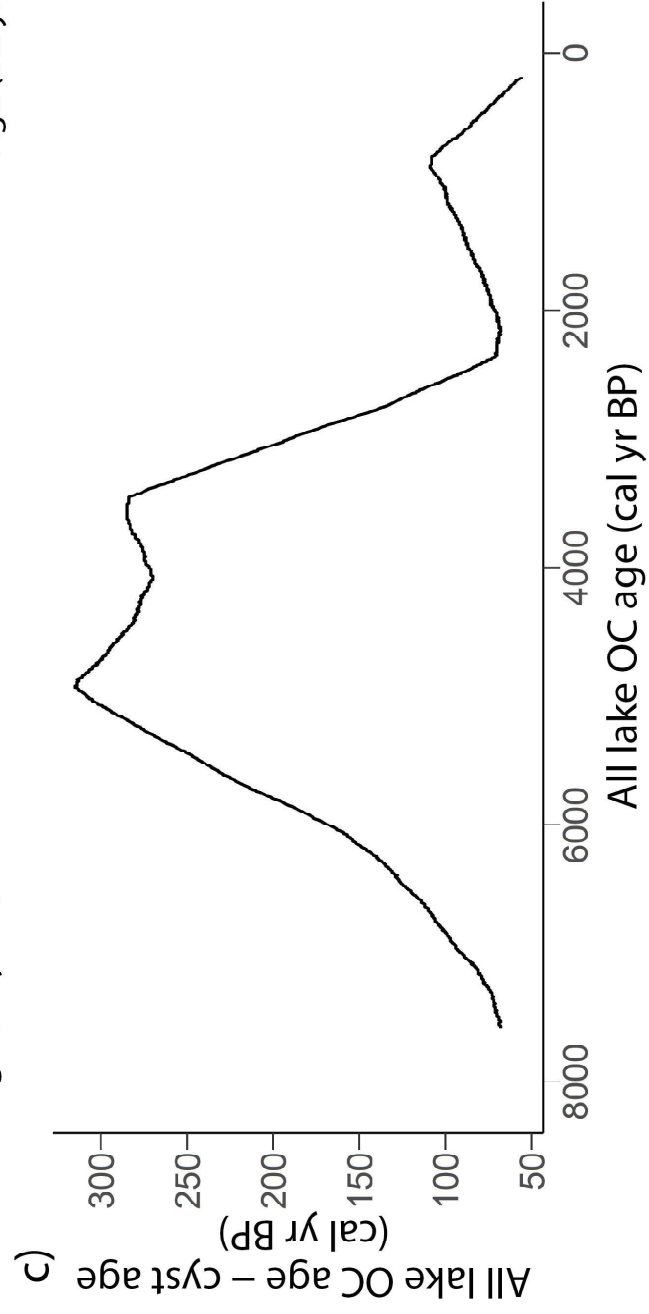
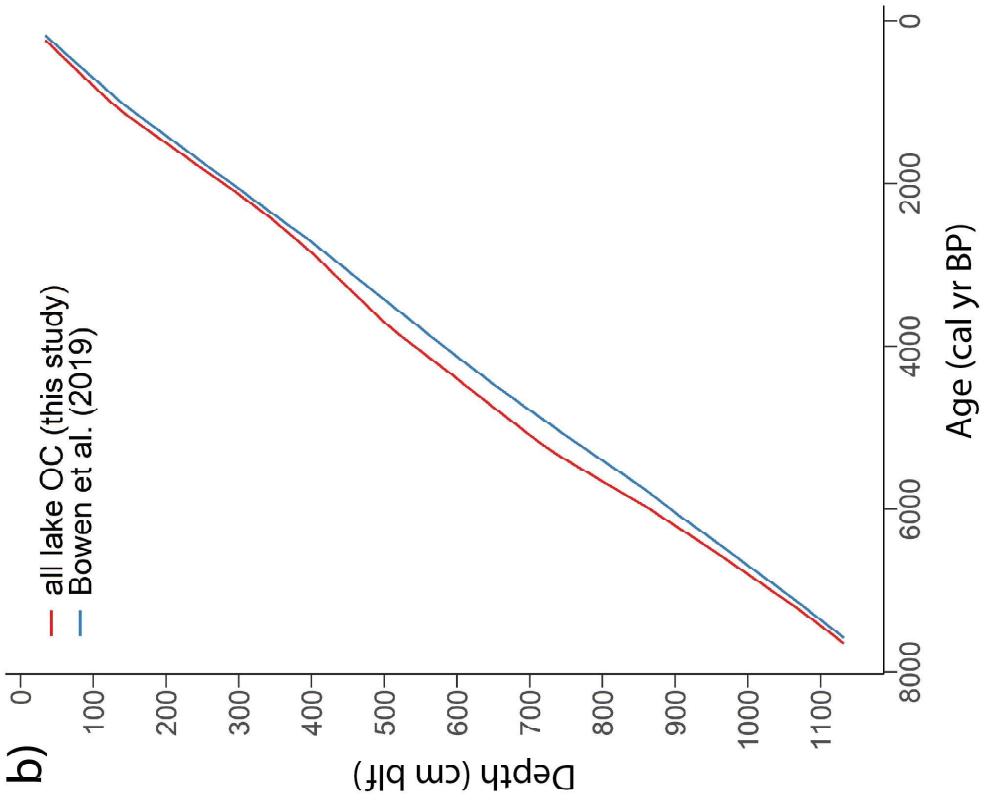
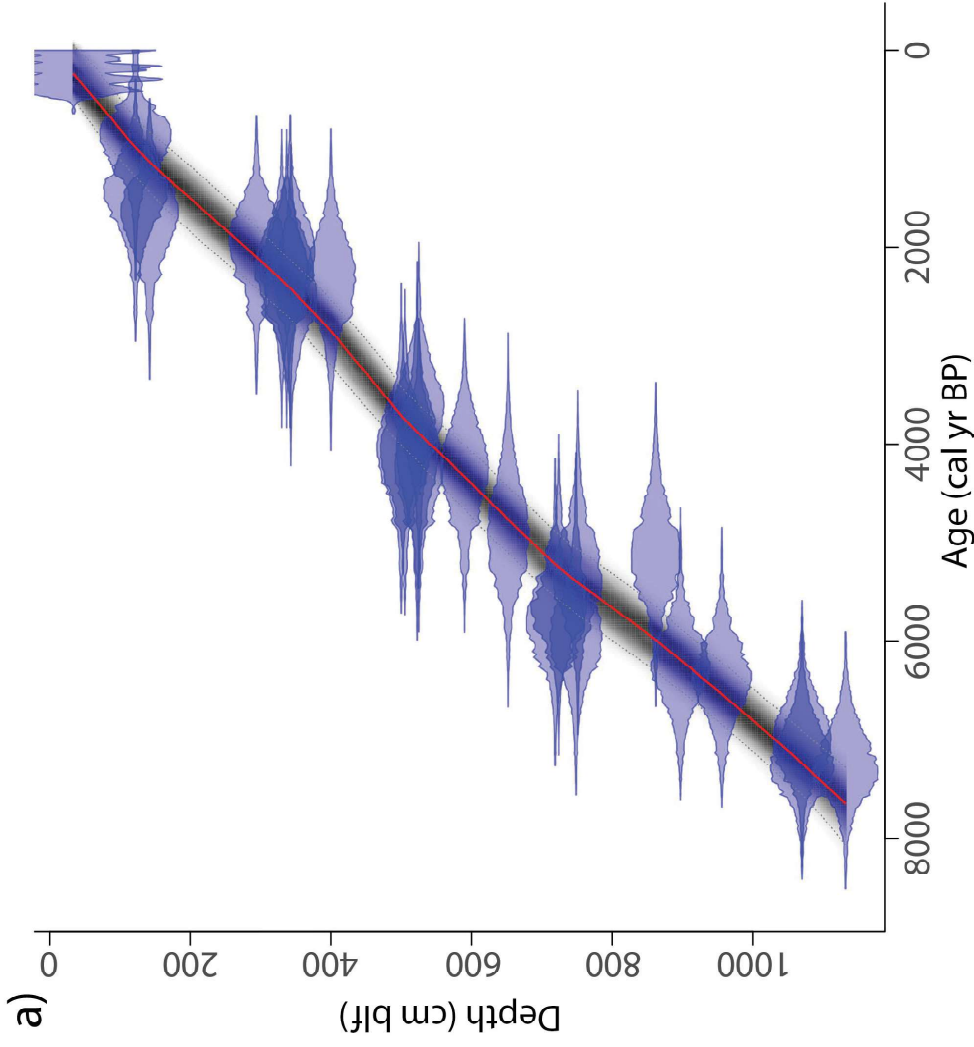


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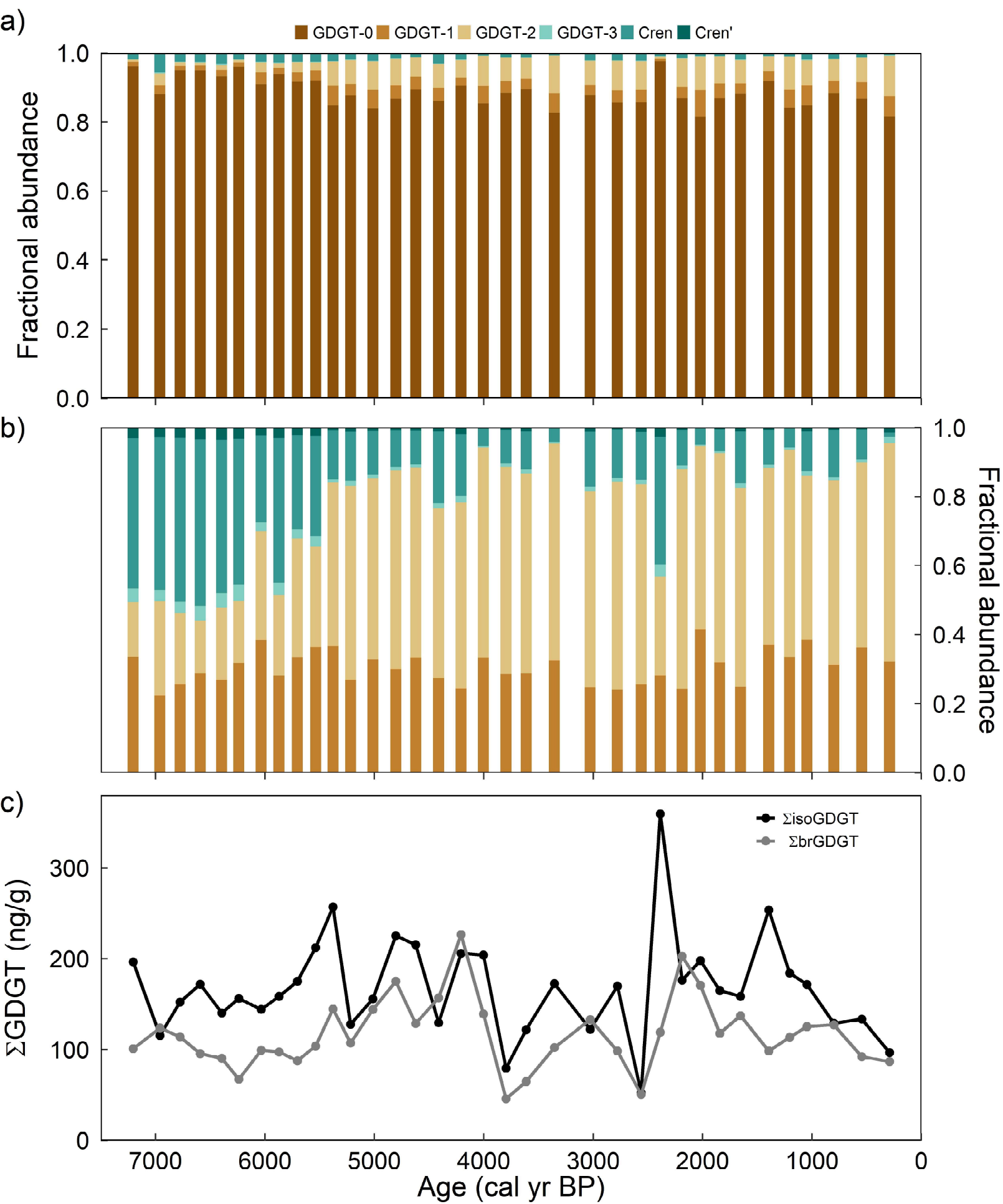


Figure 4.

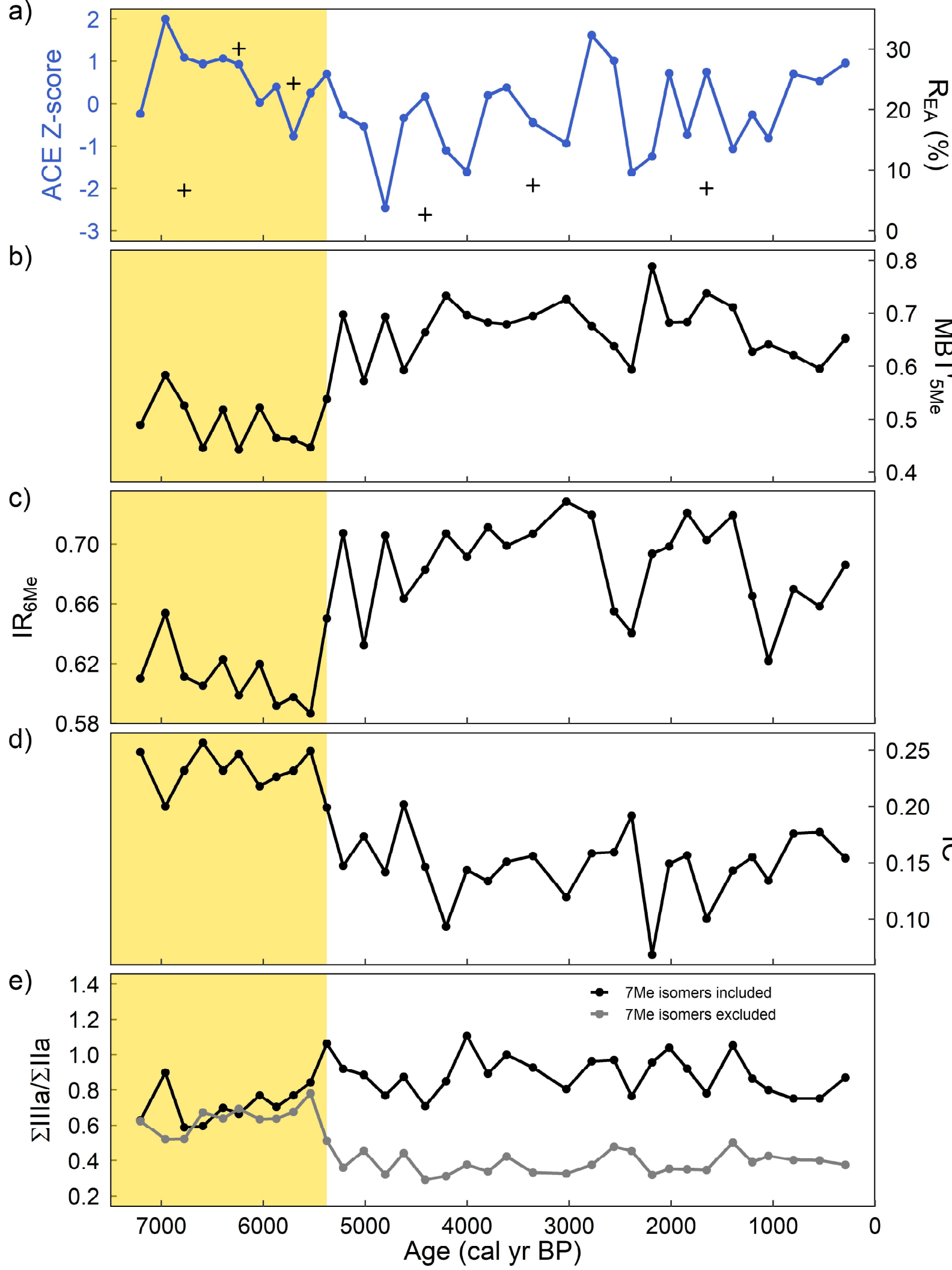


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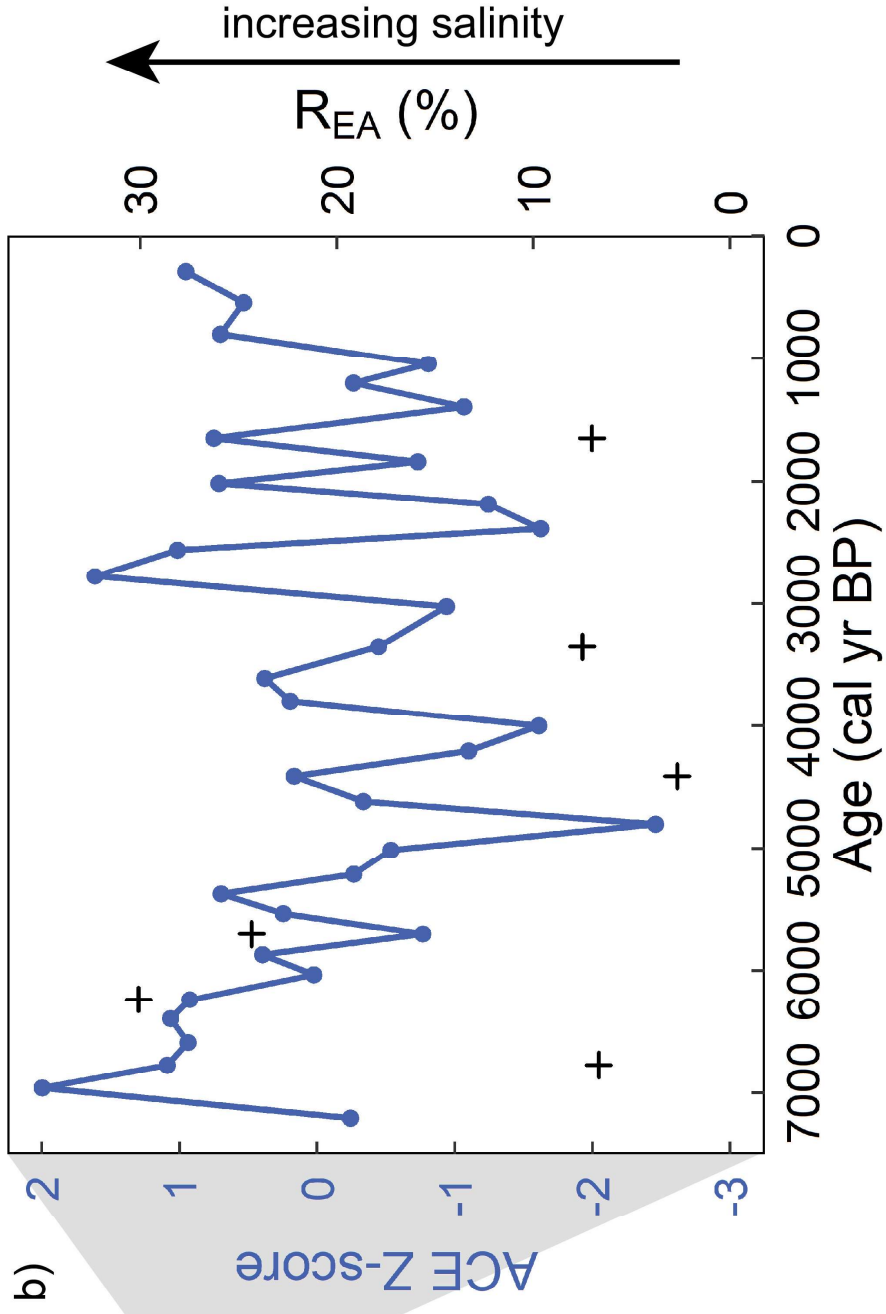
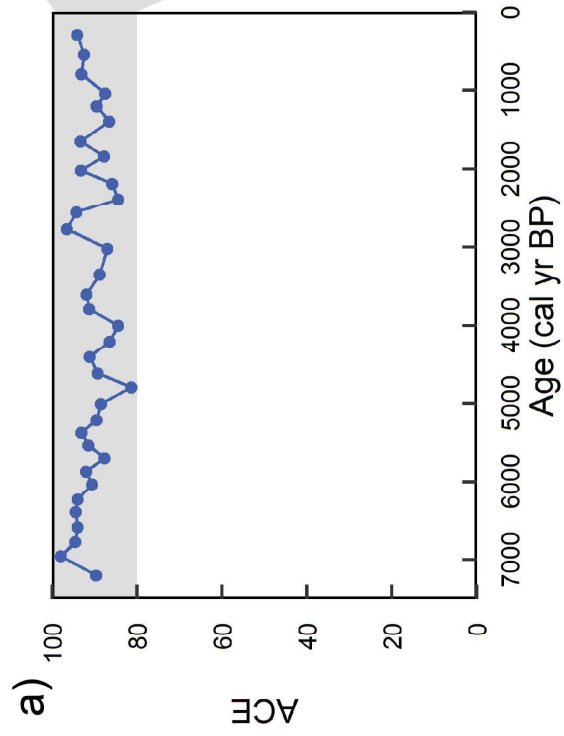


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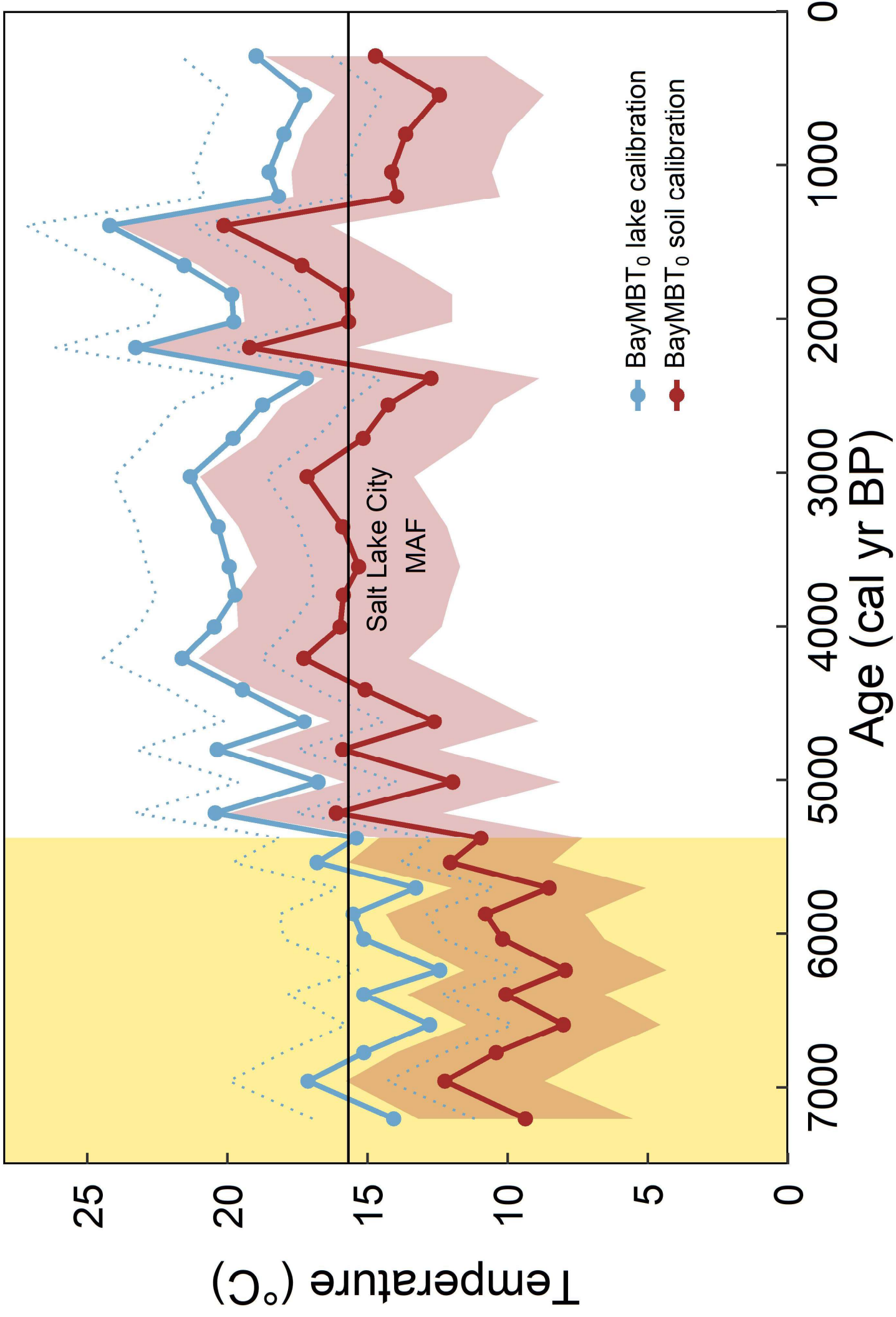
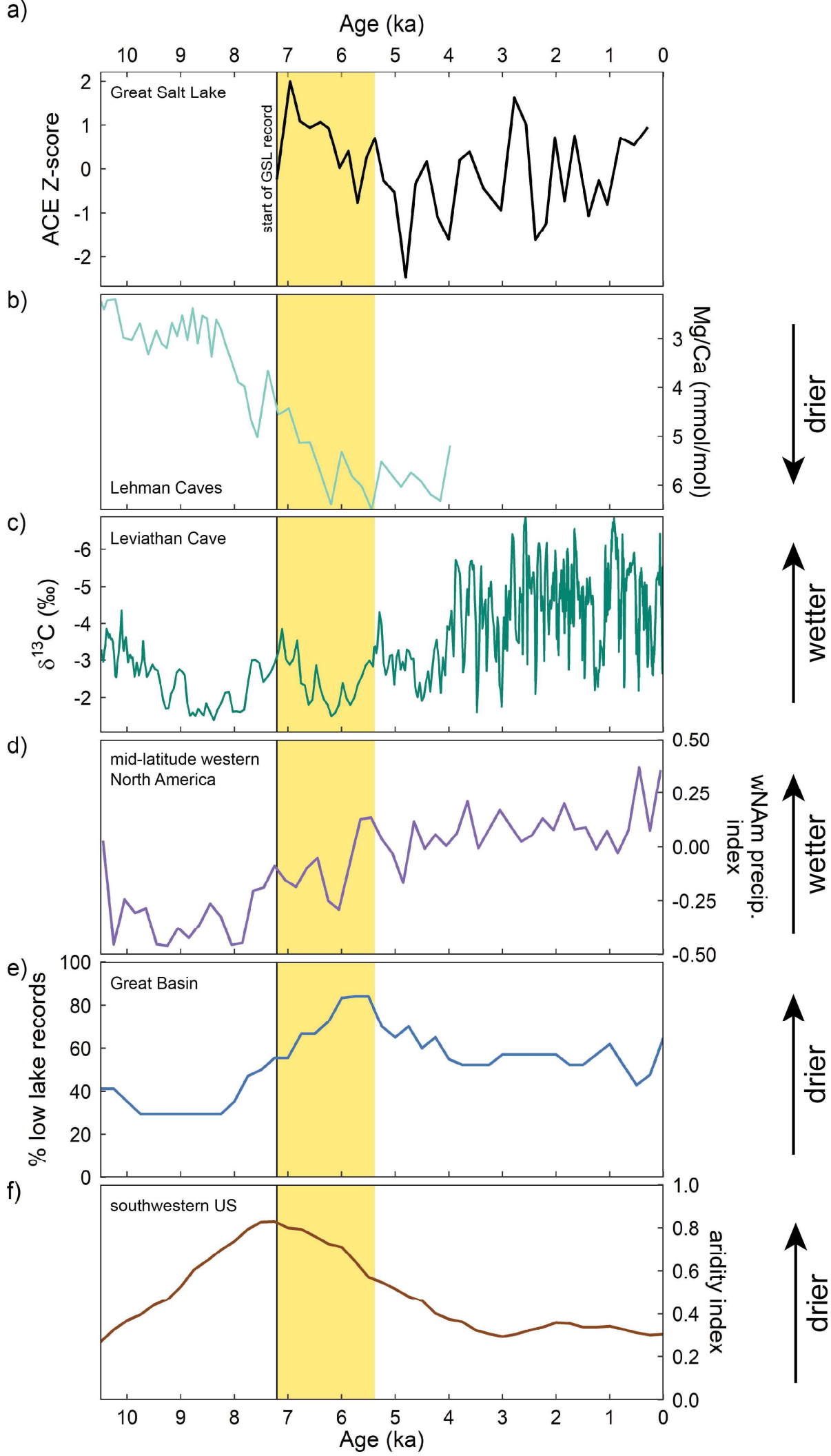


Figure 7.



Holocene water balance variations in Great Salt Lake, Utah: application of GDGT indices and the ACE salinity proxy

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Figures S1 to S6

Introduction

The supporting information associated with the manuscript referenced above includes six supplementary figures. Figure S1 is an example chromatogram of a GSL sample from this study. Figure S2 shows changepoint analysis for detecting changes in the mean and variance of select GDGT indices calculated for this study's GSL samples. Figure S3 is a compilation of all GDGT indices calculated for this study's GSL samples. Figure S4 shows principal component analyses using brGDGT distributions of GSL samples, compared to global lakes and soils. Figure S5 shows the brGDGT distributions of lake sediment and soils compared against GSL sediments. Figure S6 is a ternary diagram of tetra-, penta-, and hexamethyl brGDGT proportions in GSL samples compared to global lakes and soils.

Supplementary Figures

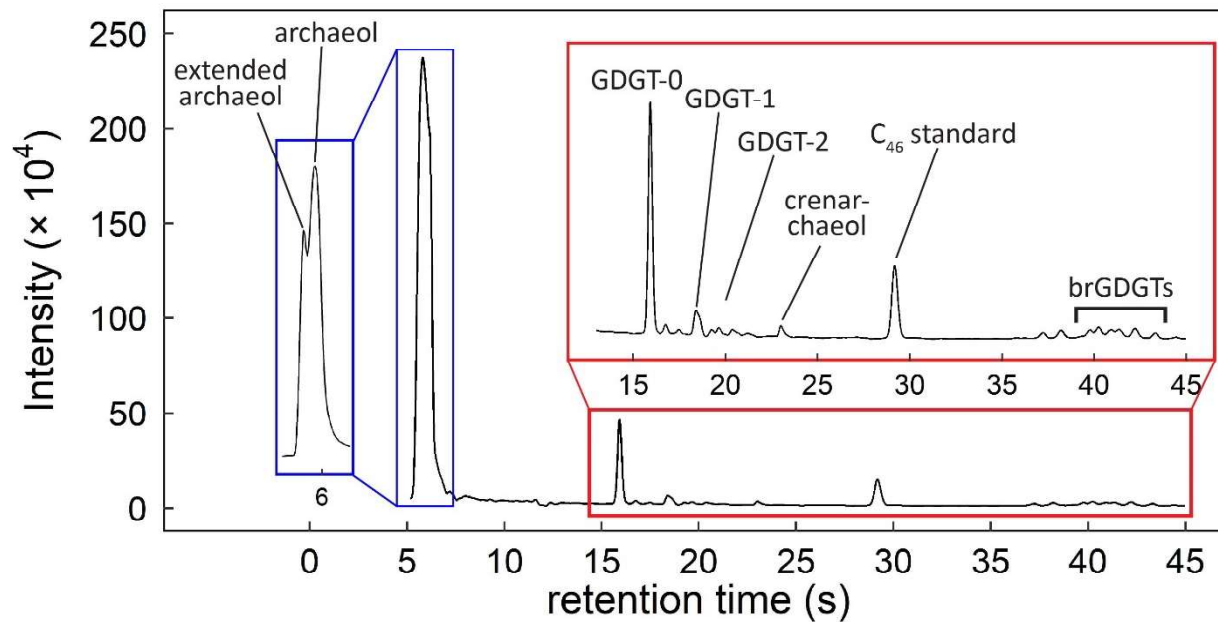


Figure S1. An example chromatogram representative of GSL samples from this study (GLAD1-GSL00-1B, 906-907cm). Labeled in the chromatogram are the individual peaks of archaeol, extended archaeol, GDGT-0, GDGT-1, GDGT-2, crenarchaeol, and the C_{46} standard along with the region containing brGDGTs.

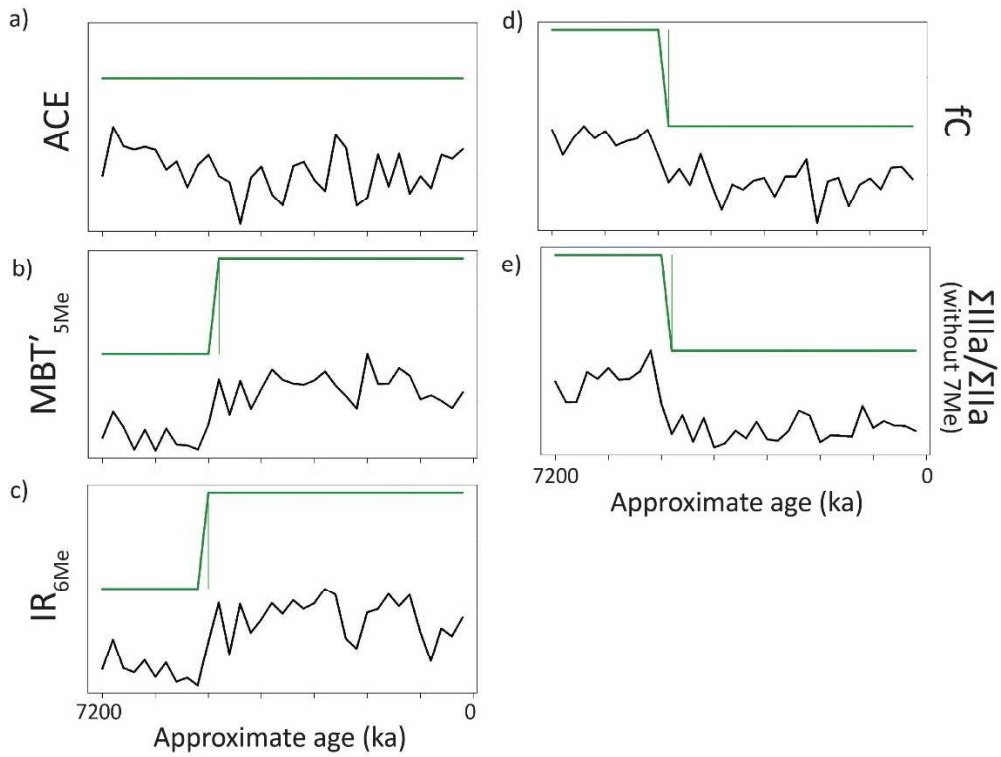


Figure S2. Environmental time series changepoint detection using the R package EnvCpt applied to a) ACE, b) MBT'_{5Me} , c) IR_{6Me} , d) fC , and e) $\Sigma IIIa/\Sigma IIa$ without 7-methyl isomers. The method used identifies changepoints in the series mean and variance.

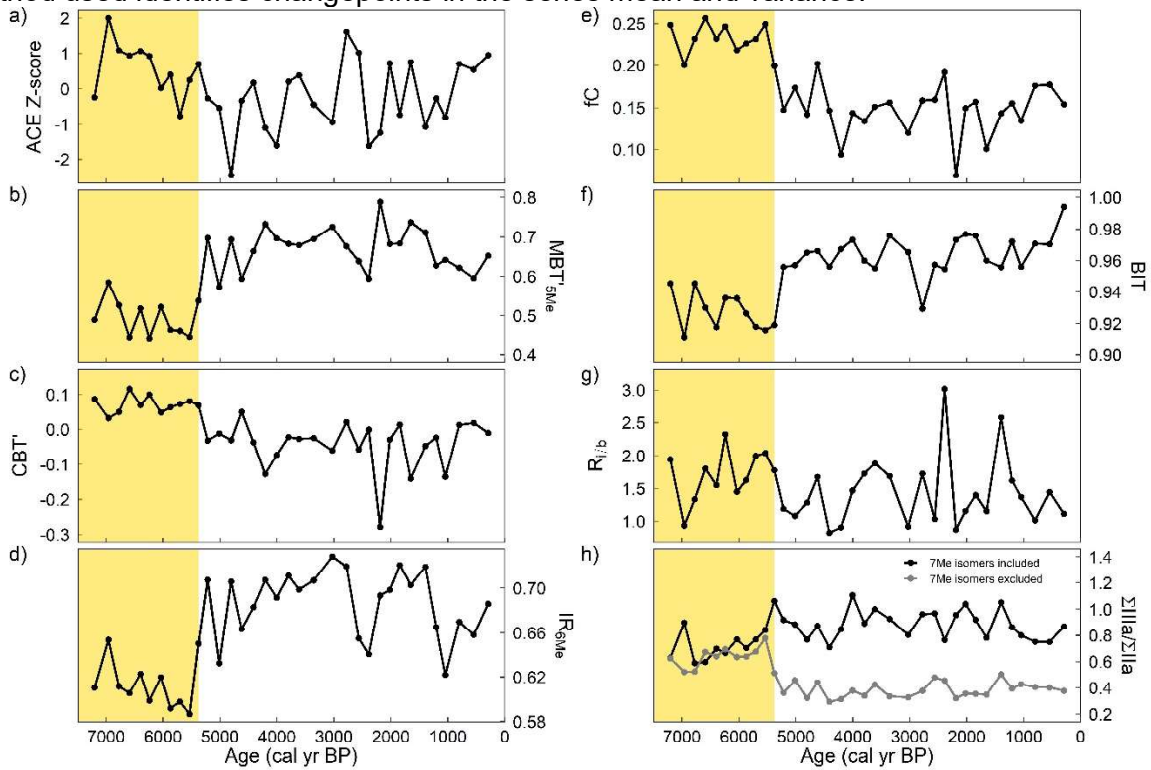


Figure S3. Temporal variations in GDGT indices showing a) ACE Z-scores, b) MBT'_{5Me} , c) CBT' , d) IR_{6Me} , e) fC , f) BIT , g) $R_{i/b}$, and h) $\Sigma IIIa/\Sigma IIa$. Yellow shading indicates the portion of the record prior to the step change seen in four indices (MBT'_{5Me} , IR_{6Me} , fC , and $\Sigma IIIa/\Sigma IIa$).

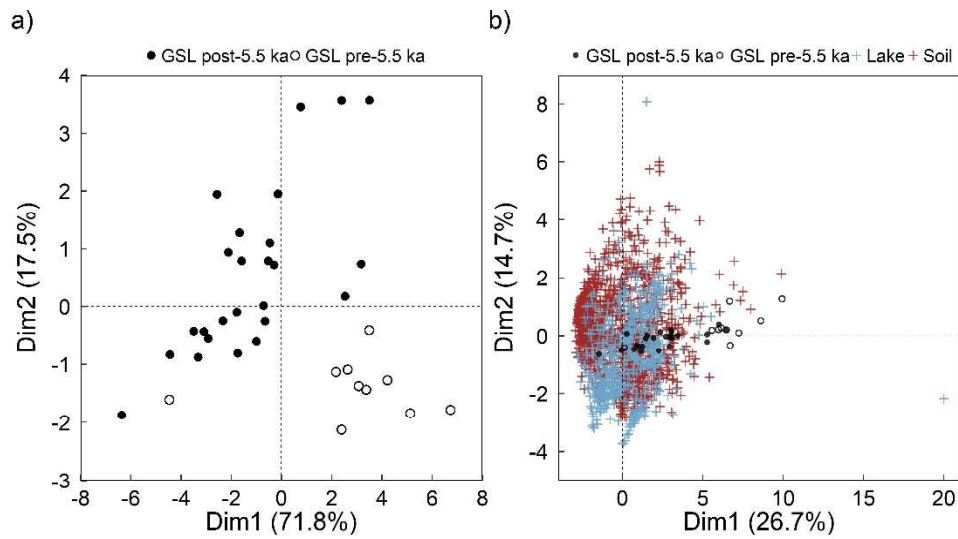


Figure S4. Principal component analyses (PCA) using brGDGT distributions of a) pre- (open circle) and post-5.5 ka (filled circle) GSL sediments, and b) compared to global lakes (blue cross) and soils (red cross) compilation of Raberg et al. (2022).

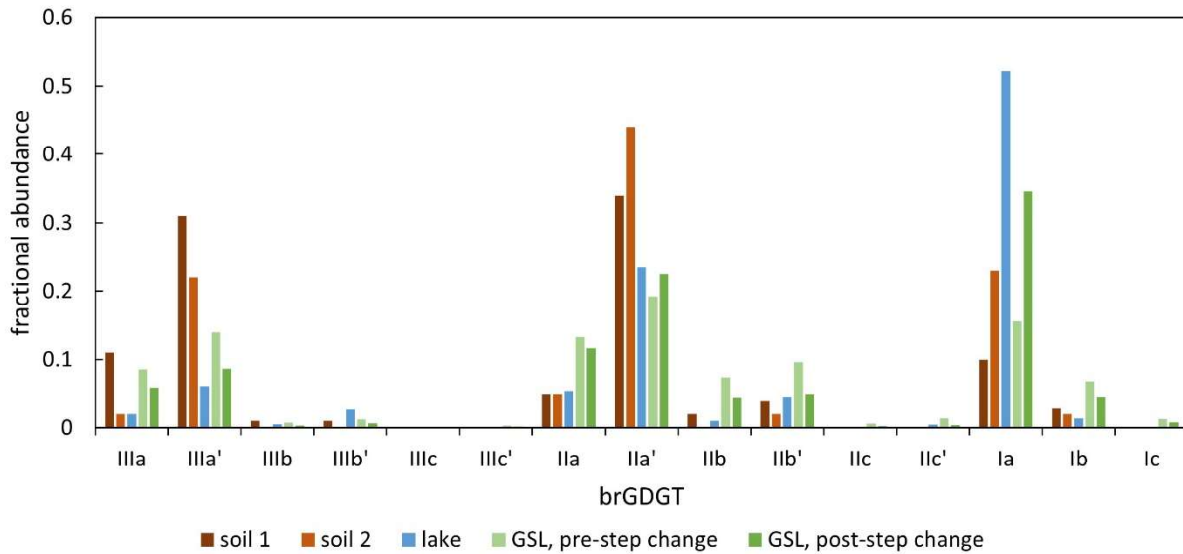


Figure S5. BrGDGT distribution of GSL samples from before and after the 5.5 ka step shift, and comparison to soils and lake distributions from southwestern US sites. Comparison brGDGT distributions were obtained from the soil compilation of Dearing Crampton-Flood et al. (2020), and lake compilation of Martinez-Sosa et al. (2021), showing Mono Lake (CA).

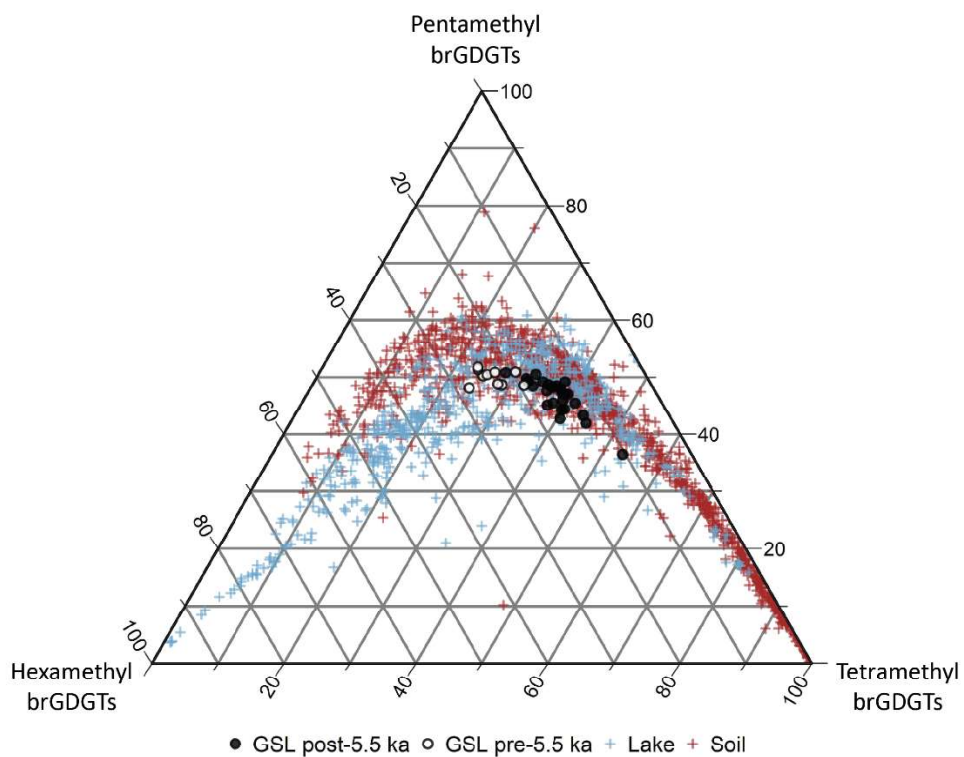


Figure S6. Ternary diagram showing the proportions of tetra-, penta-, and hexamethyl brGDGTs (I, II, and III respectively) in GSL sediments, differentiated into pre-5.5 ka (white circle) and post 5.5 ka (black circle), and the global lakes (blue cross) and soils (red cross) compilation of Raberg et al. (2022).