

Multi-proxy evidence for millennial-scale changes in North Pacific Holocene hydroclimate from the Kenai Peninsula lowlands, south- central Alaska

1 **Ellie Broadman^{1*}, Darrell S. Kaufman¹, Andrew C. G. Henderson², Edward E. Berg³, R. Scott**
2 **Anderson¹, Melanie J. Leng^{4,5}, Sean A. Stahnke¹, Samuel E. Muñoz⁶**

3 ¹School of Earth and Sustainability, Northern Arizona University, Flagstaff, Arizona, USA

4 ²School of Geography, Politics & Sociology, Newcastle University, Newcastle upon Tyne, UK

5 ³Kenai National Wildlife Refuge, U.S. Fish and Wildlife Service, Soldotna, Alaska, USA

6 ⁴National Environmental Isotope Facility, Isotope Geosciences Facility, British Geological Survey,
7 Keyworth, Nottingham, UK

8 ⁵Centre for Environmental Geochemistry, School of Biosciences, University of Nottingham,
9 Nottingham, UK

10 ⁶Marine Science Center, Department of Marine & Environmental Science, Northeastern University,
11 Nahant, Massachusetts, USA

12 *** Correspondence:**

13 Ellie Broadman

14 ebb42@nau.edu

15 **Keywords:** diatom oxygen isotopes, biogenic silica, Kenai lowlands, Aleutian Low, hydroclimate,
16 south-central Alaska, Holocene

17 **Abstract**

18 The Holocene hydroclimate of south-central Alaska has been studied extensively, but conflicting

19 interpretations between oxygen isotope paleoclimate datasets are seemingly as common as

20 converging reconstructions, in part due to the challenges of interpreting oxygen isotope ratios in

21 terms of climate. Here, we present a new Holocene record of biogenic silica abundance (BSi), diatom

22 flora, and diatom oxygen isotopes ($\delta^{18}\text{O}_{\text{BSi}}$) analyzed in sediments from Sunken Island Lake (SIL) in
23 the Kenai Peninsula lowlands, which we interpret in the context of previously published paleoclimate
24 records, and use to understand regional changes in hydroclimate. Changes in lake level documented
25 by aerial photography coupled with a survey of regional lake water isotopes indicate SIL is sensitive
26 to changes in the balance of precipitation and evaporation (P-E). However, an analysis of SIL $\delta^{18}\text{O}_{\text{BSi}}$
27 over the instrumental period indicates that $\delta^{18}\text{O}_{\text{BSi}}$ is sensitive to both P-E and the isotope
28 composition of precipitation ($\delta^{18}\text{O}_{\text{precip}}$), which is driven by changes in the Aleutian Low atmospheric
29 pressure cell (AL). We attribute a $\sim 2\text{‰}$ increase in $\delta^{18}\text{O}_{\text{BSi}}$ from 5.5 – 4.5 ka cal BP to a stronger AL,
30 which resulted in the delivery of isotopically heavier precipitation to the Kenai lowlands, and wetter
31 conditions during the late Holocene. These interpretations are supported by late Holocene increases
32 in the relative abundance of planktonic diatoms and BSi-inferred storminess, and by evidence for
33 higher-than-present lake levels on the paleo-shorelines above SIL at $\sim 1.5 - 0.5$ ka cal BP. Our dataset
34 demonstrates that this region was characterized by relatively low lake levels and dry climate in the
35 early Holocene, a strengthening of the AL in the late Holocene, and wetter climate during the late
36 Holocene until recent decades.

37 **1 Introduction**

38 The hydroclimate of the northeastern Pacific continental margin is of interest to both modern and
39 paleo-climatologists due to the importance of the synoptic-scale ocean-atmospheric dynamics that
40 influence the region. The terrestrial hydroclimate of southeastern Beringia is particularly sensitive to
41 these dynamics, in part due to the pronounced influence of changes in the position and intensity of
42 the Aleutian Low (AL), an atmospheric low pressure cell that affects the strength and trajectory of
43 winter storms (Rodionov et al., 2007). AL variability can therefore cause increased or decreased
44 winter season precipitation, as well as changes to the oxygen isotope composition of precipitation
45 ($\delta^{18}\text{O}_{\text{precip}}$), which result from different source waters and storm trajectories. These changes in

46 $\delta^{18}\text{O}_{\text{precip}}$ are often inferred from proxy data in paleoclimate archives and used to reconstruct AL
47 variability. Comparisons between $\delta^{18}\text{O}$ records and the North Pacific Index (NPI – an index of area-
48 weighted sea level pressure over the North Pacific), which is strongly inversely related to the strength
49 of the AL (Trenberth and Hurrell, 1994), are often used to confirm the relationship between $\delta^{18}\text{O}$ and
50 the AL (e.g. Bailey et al., 2015). In recent decades, many studies on the Holocene hydroclimate of
51 southern Alaska and southwestern Yukon have used fluctuations in $\delta^{18}\text{O}$ to reconstruct changes in the
52 AL, contributing substantially to our knowledge of the pressure cell’s Holocene evolution (Anderson
53 et al., 2005; Fisher et al., 2008; Schiff et al., 2009; Jones et al., 2014, 2019; Bailey et al., 2015, 2018).
54 Despite this wealth of literature, our confidence in Holocene AL variability remains limited because
55 of contradictory interpretations of paleo-AL behavior among paleoclimate datasets (e.g. Schiff et al.,
56 2009; Jones et al., 2014; Kaufman et al., 2016) and confounding factors associated with the variety of
57 materials analyzed for $\delta^{18}\text{O}$ (bulk organic matter, endogenic calcite, biogenic silica, glacier ice, leaf
58 waxes, and *Pisidium* shells). Additionally, a recent study of $\delta^{18}\text{O}_{\text{precip}}$ in Anchorage, Alaska revealed
59 that, while $\delta^{18}\text{O}_{\text{precip}}$ is influenced by synoptic-scale atmospheric circulation patterns related to AL
60 conditions, there is no statistically significant relation between $\delta^{18}\text{O}_{\text{precip}}$ and the NPI, calling into
61 question the efficacy of using $\delta^{18}\text{O}$ records to reconstruct shifts in AL strength and position in this
62 region (Bailey et al., 2019).

63 While the details of the Holocene history of the AL remain unclear (Kaufman et al., 2016), some
64 features of the hydroclimate of southern Alaska are relatively consistent among existing paleoclimate
65 datasets. For example, Barron and Anderson (2011) identified a shift in terrestrial hydroclimate
66 associated with a strengthened AL at ~4 ka cal BP throughout the northeastern Pacific Rim, which
67 would have increased moisture flow to southern Alaska. This late Holocene hydroclimatic shift has
68 been corroborated by more recent $\delta^{18}\text{O}$ reconstructions that also identify a strengthened AL at this
69 time (Jones et al., 2014; Bailey et al., 2018). This inferred AL shift also corresponds roughly to a

70 period characterized by wetter conditions throughout southeastern Beringia in the late Holocene
71 (Kaufman et al., 2016).

72 To further examine Holocene hydroclimate and related environmental changes in south-central
73 Alaska, we present a new biogenic silica (BSi), diatom flora, and diatom oxygen isotope ($\delta^{18}\text{O}_{\text{BSi}}$)
74 record from Sunken Island Lake in the Kenai Peninsula lowlands. Diatoms are abundant in the
75 sedimentary record of most non-alkaline lakes and have an unambiguous source of $\delta^{18}\text{O}$ from the
76 lake water in which they make their silica-based frustules, a process thought to occur in isotopic
77 equilibrium with lake water (Leng and Barker, 2006). Therefore, $\delta^{18}\text{O}_{\text{BSi}}$ is a proxy well suited to
78 reconstructing paleoclimate conditions in this region. Our results provide evidence for a late
79 Holocene increase in both effective moisture and southerly-derived precipitation, and highlight the
80 complexity of interpreting paleo- $\delta^{18}\text{O}$ data in the dynamic and multi-faceted North Pacific
81 hydroclimate setting.

82 **1.1 Study area**

83 The Kenai Peninsula is located in south-central Alaska, bordered by Cook Inlet and the Gulf of
84 Alaska (GoA) (Fig. 1A). The Kenai Mountains occupy the southeastern portion of the peninsula, and
85 act as a barrier to storms traveling from the GoA, creating a rain shadow effect in the Kenai lowlands
86 to the northwest. Mean annual temperature at the Kenai Moose Pens SNOTEL site was 2.7°C, mean
87 annual rainfall was 47.2 cm, and mean annual snow-water equivalent was 11.4 cm between 1995 and
88 2019 (National Water and Climate Center, <https://www.wcc.nrcs.usda.gov/index.html>). The majority
89 of precipitation in the Kenai Peninsula occurs from November to March, when the AL strengthens
90 and moves eastward, as opposed to spring and summer months when it weakens and moves westward
91 (Overland et al., 1999) (Fig. 2). These seasonal changes in the AL also have a strong influence on the
92 average path of winter storms: a strong AL directs storms from the southwest to the GoA and south-

93 central region of the state, whereas a weak AL sends storms west and northwest to travel across
94 Alaska before reaching the same region (Rodionov et al., 2007) (Fig. 3). These seasonal changes in
95 precipitation are accompanied by seasonal changes in temperature and $\delta^{18}\text{O}_{\text{precip}}$ (Fig. 2). The relative
96 strength and position of the AL also vary interannually, resulting in stormier and less stormy years in
97 south-central Alaska. Both seasonal and interannual shifts in average storm trajectories associated
98 with the AL also influence $\delta^{18}\text{O}_{\text{precip}}$ (Fig. 3), where interannual fluctuations superimpose an isotope
99 imprint on the average seasonal change.

100 Sunken Island Lake (SIL; 60.592°, -150.883°; 76 m a.s.l.) is one of many kettles holes that formed in
101 the Kenai lowlands during the recession of the Moosehorn stade of the Naptowne glaciation, which
102 commenced ~23 – 19 ka cal BP following ice advances from both the Alaska Range and the Kenai
103 Mountains (Reger et al., 2007). SIL comprises a large, deep sub-basin in its southern portion (16.8 m
104 maximum water depth), and two smaller sub-basins in the north and east (7.4 m and 7.0 m maximum
105 water depth respectively) (Fig. 1B). The lake is topographically closed, lacking a permanent inflow
106 and outflow, and is therefore primarily fed by precipitation and groundwater. SIL is thermally
107 stratified in the summer months: in June 2017, water surface temperature was 16.2°C, and bottom
108 water temperatures were 10.6°C at 7.0 m water depth in the eastern sub-basin and 6.4°C at 16.8 m
109 water depth in the southern sub-basin. Conductivity of SIL water ranged from 35 to 45 $\mu\text{S cm}^{-1}$.
110 Aerial photographs show that lake surface elevation was ~78.5 m a.s.l in 1950 CE, and lowered to
111 ~75.5 m a.s.l by 2011 CE, indicating that substantial lake-level fluctuations occur at this site (Fig.
112 1C-D).

113 2 Methods

114 2.1 Regional water sampling and isotope analyses

115 To determine whether evaporation influences the isotope composition of lake water ($\delta^{18}\text{O}_{\text{lake}}$) at SIL
116 and to characterize the $\delta^{18}\text{O}$ of the components of the local hydrologic system, a water sampling
117 campaign was undertaken in the Kenai lowlands in June 2017, June 2018, and August 2018. Water
118 samples were collected in 125 mL polyethylene bottles from 27 lakes (including SIL), 2 rivers, and a
119 groundwater well ($n = 84$). Following each field season, water samples were processed using
120 Wavelength-Scanned Cavity Ringdown Spectroscopy at the Colorado Plateau Analytical Laboratory
121 at Northern Arizona University. Analytical precision was $\leq 0.3\text{‰}$ for $\delta^{18}\text{O}$ and $\leq 0.6\text{‰}$ for δD (1σ).

122 2.2 Sediment core recovery, sedimentological analyses, and geochronology

123 Sediment core SIL-04-01 (821 cm long) was taken from the northern sub-basin of SIL (60.5962° , –
124 150.8779° ; 7.4 m water depth) (Fig. 1B) in June 2004 using a 5-cm-diameter Livingstone piston
125 corer. Upon recovery, core segments were split, and the sediment stratigraphy described. Magnetic
126 susceptibility (MS) was analyzed at 5 mm spacing using a Bartington MS2E surface probe. Loss on
127 ignition (LOI) was completed for each contiguous 1 cm increment, where each 1 cm^3 sample was
128 weighed and dried overnight in an oven, then re-weighed and burned at 550°C for 2 hours, and
129 finally weighed again to calculate LOI (Dean, 1974). SIL-04-01 has been previously analyzed for
130 pollen, macrofossils, and charcoal (Anderson et al., 2019).

131 Plant and insect macrofossils from SIL-04-01 provide an existing geochronological framework
132 (Anderson et al., 2019), but the core has been heavily sub-sampled for previous analyses. Additional
133 material was needed for our sediment-intensive diatom oxygen isotope sampling. Therefore, in June
134 2017 sediment cores were taken from four sites across SIL (Fig. S1) using a percussion piston coring
135 system. Upon recovery, cores were split, described, and analyzed for bulk density and MS using

136 Geotek MSCL-S and MSCL-XYZ automated core loggers at the National Lacustrine Core Facility
137 (LacCore) at the University of Minnesota, Minneapolis. Sediment cores from the deepest, southern
138 basin of the lake (16.8 m water depth; SIL17-2 and SIL17-3) were not analyzed in this study because
139 the stratigraphy of these sequences indicates they may contain one or more slump deposits (Fig. S1).
140 Of the other two sediment sequences, the core from the eastern sub-basin (SIL17-1; 60.5916°, –
141 150.8854°; 7.0 m water depth; Fig. 1B) was > 1 m longer (401 cm) than the core from the northern
142 basin (SIL17-4). SIL17-1 was therefore used in this study due to the large sample sizes needed for
143 analyzing $\delta^{18}\text{O}_{\text{BSi}}$. Though SIL17-1 was the longest sediment sequence collected in 2017, it did not
144 capture the entire Holocene record, and therefore the lower portion of SIL-04-01 was still needed to
145 complete the $\delta^{18}\text{O}_{\text{BSi}}$ dataset. MS measurements and prominent tephra layers were used to provide tie
146 points between cores SIL17-1 and SIL-04-01 in order to establish a geochronology for the upper ~4
147 m of SIL17-1 using existing ages from SIL-04-01 (Fig. S2). This transfer of previously acquired ages
148 gave rise to a composite ~750 cm “master core” (SIL-MC), which comprises ~4 m of SIL17-1 and
149 the lower ~3.5 m of SIL-04-01 (Supplementary Data). SIL-MC was used for all sampling and
150 original analyses presented in this study. To compare existing analyses from SIL-04-01 (Anderson et
151 al., 2019) to the BSi, diatom assemblage, and $\delta^{18}\text{O}_{\text{BSi}}$ data, the depths and values from SIL-04-01
152 were adjusted according to regression relationships derived from the cores’ shared tie points from the
153 MS and tephtras (Supplementary Data).

154 Percent biogenic silica by mass (BSi) was determined for SIL-MC using wet-alkaline extraction
155 (10% Na_2CO_3), molybdate-blue reduction, and spectrophotometry (Mortlock and Froelich, 1989).
156 Sampling resolution was every 0.5 cm for the upper 15 cm, and every 8 cm downcore ($n = 130$). A
157 duplicate was analyzed for every 4 or 5 samples ($n = 30$) to determine the reproducibility of the
158 procedure. Duplicates were processed in separate batches, and samples in each batch were selected
159 randomly from all sample depths.

160 ^{210}Pb and ^{137}Cs profiles were used to constrain the chronology for the last ~100 years of surface core
161 SIL17-1B (Fig. S3; Table S1). The previously published ^{210}Pb -based chronology from SIL-04-01
162 (Anderson et al., 2019) was not used because SIL17-1 includes sediments deposited between 2004
163 and 2017, necessitating a new surface sediment chronology that incorporates these youngest
164 sediments that were analyzed for BSi and $\delta^{18}\text{O}_{\text{BSi}}$. ^{210}Pb , ^{214}Pb , and ^{137}Cs γ -activities were measured
165 on 14 oven-dried and powdered samples from the upper 19 cm of SIL17-1B using a Canberra Broad
166 Energy Germanium Detector (BEGe; model no. BE3830P-DET), with a count time of 24 hours. The
167 Constant Rate of Supply (CRS) model (Appleby, 2001) was used to estimate ages and confidence
168 intervals of the samples with excess ^{210}Pb activities above equilibrium with ^{214}Pb . Samples were
169 analyzed at the Earth Surface Systems Laboratory at the Marine Science Center at Northeastern
170 University.

171 The down-core chronology for SIL-MC was established using 16 AMS ^{14}C dates on plant and insect
172 macrofossils from SIL17-1 ($n = 1$) and SIL-04-01 ($n = 15$). Ages from SIL-04-01 were “mapped”
173 onto SIL17-1 using least-squares regressions derived from prominent marker layers and MS peaks
174 shared between the sediment sequences (Fig. S2). For all ^{14}C ages, sediment samples 1-2 cm thick
175 were sieved at 150 μm , and macrofossils were then picked, dried in an oven at 50°C, and identified
176 under a Zeiss light microscope. Macrofossils were prepared and converted to graphite at Northern
177 Arizona University, and ^{14}C content was measured at the Keck-Carbon Cycle AMS facility at UC
178 Irvine.

179 Uncalibrated ^{14}C dates and the ages from the ^{210}Pb model output were incorporated into an age model
180 for the entire sediment sequence using the R package Bacon (v2.2) (Blaauw and Christen, 2011),
181 which calibrates ^{14}C years to calendar years using IntCal13 (Reimer et al., 2013). Chronological
182 uncertainties for the BSi and $\delta^{18}\text{O}_{\text{BSi}}$ time series were generated in GeoChronR (McKay et al., 2018)
183 using the ensemble output from the Bacon age model.

184 To constrain fluctuations in Holocene lake level, sediments from satellite fens and ice-shoved
185 ramparts surrounding SIL were acquired and dated. Cores were collected from five satellite fens near
186 SIL using a 2.5-cm-diameter piston corer designed to capture the lowermost 45 cm of peat above
187 mineral soil. AMS ¹⁴C dates were determined for macrofossils in the lowermost peat horizons. A
188 prominent 1400-m long ice-shoved rampart was also excavated at two sites. Thrust events in soil
189 profiles in the ice-shoved rampart were identified, and AMS ¹⁴C dating was completed on
190 macrofossils below the thrust events in order to chronologically constrain the thrust event.

191 **2.3 Diatom assemblage analysis**

192 Twenty 1-cm-thick samples from SIL-MC were selected at 30-50 cm increments for diatom species
193 analysis. These samples were treated with 30% H₂O₂ and 70% HNO₃ to remove organic matter
194 before creating slides for counting using Naphrax© mounting medium. A Zeiss light microscope was
195 used to count three hundred valves per slide along transects at 1000X magnification, and taxonomic
196 identifications were made according to Foged (1971), Foged (1981), Krammer and Lange-Bertalot
197 (1986-1991), Mann et al. (2004), and McGlaughlin and Stone (1986). Diatom nomenclature was
198 updated following Spaulding et al. (2019). Following identification, diatom species counts were
199 converted to percent relative abundance and graphed with Tilia (v.2.1.1) (Grimm, 2015). An
200 incremental sum-of-squares cluster analysis (CONISS) was applied to dominant taxa with a relative
201 abundance > 5% in at least one sample (Grimm, 1987) to determine zones in the sequence. A
202 principal component analysis (PCA) (ter Braak and Prentice, 1988) was completed on the correlation
203 matrix of untransformed percentage data for all dominant taxa.

204 **2.4 Diatom oxygen isotope analysis**

205 Samples for diatom oxygen isotope analysis (1 cm thick) were taken from SIL-MC every ~10 cm
206 downcore, and every 1 cm of the upper 20 cm (n = 106). Because the oldest sediments were taken

207 from heavily-sampled SIL-04-01, sampling intervals are less consistent over the deepest ~3.5 m of
208 SIL-MC. Sampling avoided visible tephra layers. Samples were purified using a series of chemical
209 digestions, sieving, and heavy liquid separations (*cf.* Morley et al., 2004). All samples were visually
210 inspected for contamination under a Zeiss light microscope, and 30 samples were inspected further
211 using a Zeiss Supra 40VP variable pressure field emission scanning electron microscope (SEM) (e.g.
212 Fig. 4) and energy dispersive X-ray spectroscopy (EDS). $\delta^{18}\text{O}_{\text{BSi}}$ values were measured by using the
213 stepwise fluorination method (Leng and Sloane, 2008) in the stable isotope facility at the British
214 Geological Survey in Keyworth, UK. $\delta^{18}\text{O}$ are reported as per mil (‰) deviations of the isotopic ratio
215 ($^{18}\text{O}/^{16}\text{O}$) calculated to the VSMOW scale using a within-run laboratory standard calibrated against
216 NBS-28. The measured $\delta^{18}\text{O}$ of the standard silica (BFC) was 28.9‰ ($n = 16$) with an analytical
217 reproducibility of 0.2‰. Analytical reproducibility of the sample material was $< 0.3\text{‰}$ for $\delta^{18}\text{O}$ (1σ).

218 **3 Results**

219 **3.1 Regional water isotopes**

220 Water isotope values from the 75 lake water samples (Table S2) have a $\delta^{18}\text{O}$ range of 8.3‰ (−15.1 to
221 −6.8‰ VSMOW) and a δD range of 38.9‰ (−121.9 to −83.0‰ VSMOW), yielding a Local
222 Evaporation Line (LEL) with a slope of 4.65 (Fig. 5). Water from SIL plots on the far right of the
223 LEL ($\delta^{18}\text{O}$ range: −9.3 to −8.4‰ VSMOW; δD range: −94.0 to −87.5‰ VSMOW; Fig. 5). $\delta^{18}\text{O}$ and
224 δD values are lower for samples from groundwater and the Kenai and Moose Rivers, some of which
225 are situated close to the Global Meteoric Water Line (GMWL).

226 **3.2 Sediment stratigraphy and geochronology**

227 The ^{210}Pb activities exhibit a gradual decline over the upper 11 cm of core SIL17-1B towards an
228 equilibrium of ~8.3 Bq/kg (Fig. 6; Fig. S3; Table S1). Four samples were excluded from the CRS
229 calculations because they contain low levels of excess ^{210}Pb , likely due to the presence of old

230 material reintroduced to the lake bottom. The ^{137}Cs activities were not used as an independent
231 chronological control because ^{137}Cs activities were generally low, and the peak is poorly defined. The
232 offset between the ^{137}Cs peak and the ^{210}Pb -inferred age model (Fig. 6), as well as low excess ^{210}Pb in
233 the 4 samples described above, might be due to slumping associated with the Prince William Sound
234 earthquake of 1964 CE, which has been previously linked to turbidite deposits (Praet et al., 2017;
235 Boes et al., 2018) and subsidence (Hamilton and Shennan, 2005) in the Kenai lowlands. The
236 sedimentation rate since 1950 CE inferred from the ^{210}Pb model (~ 1.2 mm/year) is comparable to that
237 reported by Anderson et al. (2019) (~ 1.3 mm/year).

238 The oldest reliable age in the SIL age-depth model is from 705.5 cm and is calibrated to 12261 ± 280
239 a BP (Fig. 6; Table 1). Two older ages were included in the age-depth model, but they are
240 stratigraphically reversed and imply an exceedingly low sedimentation rate, or an unconformity. This
241 implies the age model is not reliable in the inorganic mud below 705.5 cm depth, but diatoms are
242 essentially absent in these sediments ($\text{BSi} \leq 1.3\%$ below 712 cm) so this has no influence on our BSi-
243 and isotope-based conclusions. Three ^{14}C ages between 8 – 6 ka cal BP were excluded from the age
244 model, all of which were small macrofossil samples (0.03 – 0.06 mg graphitized carbon) with
245 relatively high analytical uncertainties (214 – 563 years). The ~ 770 cm sediment sequence captures
246 at least the entire Holocene and the late Younger Dryas, with average 95% confidence intervals of \pm
247 312 years (min = 1; max = 1268 years).

248 SIL sediments are composed primarily of gyttja, with 26 visible tephra (≥ 1 mm thick) that
249 correspond to pronounced peaks in MS (Fig. 7; Table S3). The basal ~ 50 cm is interbedded, gray,
250 inorganic clays, silts, and sands likely sourced from sediments associated with the unstable, recently
251 deglaciated landscape that dominated the Kenai lowlands after ~ 19 ka cal BP (Reger et al., 2007).
252 Organic matter content, as measured by LOI, fluctuates between 20 – 30% throughout the sediments
253 above the basal inorganic unit (Anderson et al., 2019). BSi ranges from 0.2 – 24.2% over the

254 Holocene (mean = 9.8%), with the increase in BSi content starting in the early Holocene and peaking
255 at its highest values in the mid Holocene between ~6.3 – 4.6 ka cal BP, and decreasing thereafter
256 (Supplementary Data). Analytical reproducibility as indicated by differences between duplicate
257 samples averages $0.97 \pm 0.86\%$ (1σ) across the core.

258 3.3 Diatom assemblages

259 The diatom assemblages from SIL are diverse, composed of 141 identified taxa, from which 15
260 dominant species were identified. Dominant species were grouped into one of three habitat types as
261 specified by Spaulding et al. (2019): planktonic diatoms, which are non-motile and occupy the water
262 column; facultatively planktonic diatoms, which may be motile, often dwelling in the lake's benthos,
263 but can elect to live in the water column when it is ecologically advantageous; and benthic diatoms,
264 which can be motile, or live attached to substrates on the lake floor or in shallow sediments.

265 Dominant genera at SIL include planktonic (*Aulacoseira*, *Discostella*), facultatively planktonic
266 (*Pseudostaurosira*, *Staurosira*, *Staurosirella*), and benthic (*Nitzschia*, *Pinnularia*, *Planothidium*,
267 *Sellaphora*) diatoms. Based on the CONISS dendrogram, changes in the relative abundances of these
268 taxa were divided into four zones (Fig. 8).

269 Zone 1 (12 – 8.5 ka cal BP; 695 – 570 cm) is characterized by a high abundance of *Staurosirella*
270 *pinnata*, and the highest recorded abundances of *Discostella stelligera*, *Pinnularia saprophila*,
271 *Nitzschia palea*, *Sellaphora disjuncta*, and *Sellaphora saugerresii*. Approaching 9 ka cal BP, other
272 benthic taxa increase in abundance as the three dominant species in the zone diminish.

273 Zone 2 (8.5 – 4.6 ka cal BP; 570 – 300 cm) is dominated by facultatively planktonic taxa such as *S.*
274 *pinnata*, *Staurosirella leptostauron* var. *dubia*, *Pseudostaurosira brevistriata*, *Pseudostaurosira*
275 *pseudoconstruens*, and *Pseudostaurosira parasitica*. At ~6 ka cal BP there is an increase in the
276 relative abundance of planktonic taxa such as *Aulacoseira subarctica*, *Aulacoseira valida*, and *D.*

277 *stelligera*, which then diminish. *Planothidium joursacense* becomes the dominant benthic species in
278 this zone.

279 Zone 3 (4.6 – 1 ka cal BP; 300 – 80 cm) features an increase in planktonic species, particularly *A.*
280 *valida* and *D. stelligera*. Relatively high abundances of *S. pinnata* are present throughout this zone,
281 but facultatively planktonic taxa generally become less abundant with time. *Planothidium*
282 *frequentissimum* and *P. joursacense* are the dominant benthic taxa in this zone.

283 Zone 4 (1 ka cal BP – present; 80 – 0 cm) is dominated by the highest recorded abundances of the
284 facultatively planktonic species *Staurosira construens* and *S. pinnata*, and by a decrease in the
285 relative abundance of both planktonic and benthic taxa.

286 The first two principle components (PCs) of the stratigraphic diatom assemblage data account for
287 65.4% of the overall variance in the record ($\lambda_1 = 0.419$; $\lambda_2 = 0.235$), and largely track changes in *S.*
288 *pinnata* (λ_1) and the opposing relation between *D. stelligera* and several fragilarioid species including
289 *S. construens*, *P. brevistriata*, and *P. parasitica* (λ_2) (Fig. 9).

290 **3.4 Diatom oxygen isotopes**

291 The $\delta^{18}\text{O}_{\text{BSi}}$ data show a Holocene range of 5.7‰ (+26.5 to +32.2‰ VSMOW, $n = 98$) with a mean
292 of +29.2 ‰ ($n = 55$) prior to 4.5 ka cal BP, which shifts to a mean of +30.7‰ ($n = 24$) between 4.5
293 and ~1 ka cal BP, followed by a decrease in the mean to +28.3‰ ($n = 27$) over the past ~1 ka cal BP
294 (Fig. 7; Supplementary Data). SEM images indicate that contamination by clay minerals and tephra
295 is insignificant (e.g. Fig. 4). EDS data reveal that the percent of Al_2O_3 , commonly used as an
296 indicator of clay contamination in purified biogenic silica (Brewer et al., 2008), is < 1% in all
297 samples. Because sedimentary diatom frustules may contain up to 1% Al_2O_3 incorporated into the

298 silica matrix (Koning et al., 2007), this result further indicates that samples analyzed for $\delta^{18}\text{O}_{\text{BSi}}$
299 comprise pure biogenic silica.

300 **4 Discussion**

301 **4.1 Climate controls on proxy datasets**

302 To determine controls on $\delta^{18}\text{O}_{\text{BSi}}$ and BSi in SIL, the data covering the instrumental period were
303 analyzed for their statistical relations with climate variables of interest. Daily precipitation data
304 collected at Kenai airport (http://climate.gi.alaska.edu/acis_data) were summed into annual totals,
305 and then re-sampled to average over the intervals of the $\delta^{18}\text{O}_{\text{BSi}}$ and BSi sediment samples from SIL
306 using the software package Analyseries (Paillard et al., 1996). Significance calculations were
307 adjusted to correct for auto-correlated time series using a lag-one autocorrelation model (Bretherton
308 et al., 1998) prior to calculating correlation coefficients. Additionally, to relate the $\delta^{18}\text{O}_{\text{BSi}}$ data to
309 North Pacific ocean-atmosphere circulation, the measured $\delta^{18}\text{O}_{\text{BSi}}$ data over the instrumental period
310 and the NPI and PDO index values were binned and averaged within three intervals corresponding to
311 recognized shifts in North Pacific sea-surface temperatures (SSTs) in 1944 CE (to negative PDO
312 conditions) and 1975 CE (to positive PDO conditions) (Khapalova et al., 2018). As the AL and PDO
313 are synoptic-scale patterns primarily impacting climate on multi-decadal timescales (Trenberth and
314 Hurrell, 1994; Mantua et al., 1997), binning $\delta^{18}\text{O}_{\text{BSi}}$ data to reflect known regime shifts is appropriate
315 for comparing the paleo-data to these modes of variability.

316 **4.1.1 Controls on $\delta^{18}\text{O}_{\text{BSi}}$**

317 Assuming a fractionation factor of $-0.2\text{‰}/\text{°C}$ (Brandriss et al., 1998; Moschen et al., 2005; Dodd and
318 Sharp, 2010) between the bottom water of the core site for SIL17-1 (10.6°C) and the youngest
319 sedimentary diatoms at SIL ($\delta^{18}\text{O}_{\text{BSi}} = 28.3\text{‰}$), $\delta^{18}\text{O}_{\text{lake}}$ was calculated to within 2.1‰ of the
320 measured value (-7.2‰ calculated versus -9.3‰ measured). The offset between measured and

321 calculated $\delta^{18}\text{O}_{\text{lake}}$ is unsurprising, in part due to uncertainty in the fractionation factor between water
322 and diatom silica, for which published values range from $-0.16\text{‰}/^{\circ}\text{C}$ to $-0.49\text{‰}/^{\circ}\text{C}$ (Juillet-Leclerc
323 and Labeyrie, 1987; Brandriss et al., 1998; Moschen et al., 2005; Shemesh et al., 1992; Crespin et al.,
324 2010; Dodd and Sharp, 2010). Furthermore, a growing body of evidence suggests diagenetic
325 alteration may overprint the $\delta^{18}\text{O}$ of diatom silica in the decades following sedimentary deposition
326 (Dodd et al., 2017; Tyler et al., 2017; Menicucci et al., 2017). There is no clear solution to account
327 for this possible diagenetic alteration in paleoenvironmental reconstructions, but we acknowledge
328 that it adds uncertainty to any sedimentary $\delta^{18}\text{O}_{\text{BSi}}$ reconstruction, including at SIL. In one study
329 using marine sedimentary diatoms, a universal correction was applied to each $\delta^{18}\text{O}_{\text{BSi}}$ value
330 (Menicucci et al., 2019), but this approach does not account for the uncertainty in the duration of
331 diagenetic alteration, which may occur over decades (Dodd et al., 2017) to millions of years
332 (Menicucci et al., 2017). In light of the uncertainties regarding the fractionation between lake water
333 and diatom silica, as well as the potential influence of diagenetic alteration of sedimentary diatom
334 $\delta^{18}\text{O}$, the offset between calculated and measured $\delta^{18}\text{O}_{\text{lake}}$ at SIL is expected, and does not reduce our
335 confidence that the $\delta^{18}\text{O}_{\text{BSi}}$ dataset is broadly indicative of changes in $\delta^{18}\text{O}_{\text{lake}}$.

336 As described above, the temperature-dependent fractionation between diatom silica and lake water is
337 small ($\sim -0.2\text{‰}/^{\circ}\text{C}$), meaning it is often damped by larger fluctuations in $\delta^{18}\text{O}_{\text{lake}}$ that can occur due
338 to changes in both precipitation/evaporation balance (P-E) and $\delta^{18}\text{O}_{\text{precip}}$ (Leng and Barker, 2006).
339 Therefore, the $\delta^{18}\text{O}_{\text{BSi}}$ dataset from SIL is interpreted primarily in terms of these hydroclimatic
340 variables, as has been done for other $\delta^{18}\text{O}_{\text{BSi}}$ records from southern Alaska (Schiff et al., 2009; Bailey
341 et al., 2015; Bailey et al., 2018), rather than in terms of Holocene temperature changes. The $\delta^{18}\text{O}_{\text{lake}}$
342 composition of SIL demonstrates it is heavily influenced by evaporative enrichment relative to local
343 meteoric water, where the average $\delta^{18}\text{O}_{\text{lake}}$ is -8.8‰ ($n = 6$) and the average $\delta^{18}\text{O}$ of ground and river
344 waters, which reflect an average of local meteoric waters, is -15.9‰ ($n = 8$) (Fig. 5; Table S2).

345 Combined with aerial photographic evidence for rapid lake-level fluctuations, which also indicates
346 the influence of evaporation (Fig. 1C-D), these data suggest that P-E is likely a key driver of
347 sedimentary $\delta^{18}\text{O}_{\text{BSi}}$ at SIL. Assuming this condition applies in the past, then sedimentary $\delta^{18}\text{O}_{\text{BSi}}$
348 would be sensitive to P-E. The influence of P-E on $\delta^{18}\text{O}_{\text{BSi}}$ is revealed through shifts during the
349 instrumental period that are negatively correlated with changes in the amount of annual precipitation
350 at Kenai airport ($r = -0.81$, $p = 0.02$) (Fig. 10B). This result indicates that during wetter years (higher
351 P-E), less evaporative enrichment results in lower $\delta^{18}\text{O}_{\text{lake}}$, and subsequent lower $\delta^{18}\text{O}_{\text{BSi}}$; during drier
352 years (lower P-E), more evaporative enrichment increases $\delta^{18}\text{O}_{\text{lake}}$ and $\delta^{18}\text{O}_{\text{BSi}}$.

353 One major driver of changes in P-E in southern Alaska is the variability in the position and intensity
354 of the AL (Jones et al., 2014; Bailey et al., 2018). Because a stronger AL tends to increase the
355 amount of winter (November – March) precipitation arriving in the Kenai lowlands, wetter winter
356 conditions accompany a strong AL. Increased winter precipitation resulting from a strong AL might
357 result in lower $\delta^{18}\text{O}_{\text{lake}}$ values at SIL due to higher P-E (indicating an overall wetter climate), but also
358 by increasing the relative annual contribution of isotopically light winter precipitation (compared to
359 summer; Fig. 2).

360 The AL also modulates the dominant path of storms arriving in the Kenai lowlands, which in turn can
361 alter average $\delta^{18}\text{O}_{\text{precip}}$. Because a strong AL encourages south-to-north (meridional) transport from
362 the GoA to the Kenai lowlands (Cayan and Peterson, 1989; Mock et al., 1998; Rodionov et al., 2007;
363 Berkelhammer et al., 2012), these storms tend to travel less distance and cross fewer continental
364 barriers before arriving in the Kenai lowlands. Conversely, a weak AL encourages west-to-east
365 (zonal) transport from further west in the North Pacific Ocean. Storms during a weak AL tend to
366 travel greater distances and cross more continental barriers, and therefore are likely to experience
367 more rain-out of ^{18}O , depleting $\delta^{18}\text{O}_{\text{precip}}$ arriving in the Kenai lowlands. This relation between the
368 AL and $\delta^{18}\text{O}_{\text{precip}}$ has been corroborated by several isotope-enabled model experiments

369 (Berkelhammer et al., 2012; Porter et al., 2014). The rain-out effect of ^{18}O due to shifting average
370 storm tracks has an opposing influence on $\delta^{18}\text{O}_{\text{lake}}$ to that of P-E: a strong AL leads to heavier
371 $\delta^{18}\text{O}_{\text{precip}}$, and therefore higher $\delta^{18}\text{O}_{\text{lake}}$, compared to a weak AL (Fig. 3). While there is little
372 instrumental data to support this relation between $\delta^{18}\text{O}_{\text{precip}}$ and the AL on the Kenai Peninsula,
373 $\delta^{18}\text{O}_{\text{precip}}$ recorded in Anchorage during the three strongest (2019, 2015, 2016) and weakest (2009,
374 2011, 2018) AL years from 2005–2018 reveal average $\delta^{18}\text{O}_{\text{precip}}$ was heavier during years with an
375 anomalously strong AL as indicated by the NPI. During the winter months (November – March),
376 average $\delta^{18}\text{O}_{\text{precip}}$ was 1.7‰ heavier during the strongest (–18.5‰) versus the weakest (–20.2‰) AL
377 years, and average $\delta^{18}\text{O}_{\text{precip}}$ was 3.0‰ heavier during the strongest (–15.7‰) versus the weakest (–
378 18.7‰) AL years annually (Bailey et al., 2019).

379 $\delta^{18}\text{O}_{\text{BSi}}$ values during the instrumental period reveal the relationship between AL strength and
380 $\delta^{18}\text{O}_{\text{precip}}$ may apply at SIL: $\delta^{18}\text{O}_{\text{BSi}}$ is broadly consistent with shifts in the PDO and NPI indices (Fig.
381 10A), where negative NPI (strong AL) and positive PDO index (associated with strong AL) values
382 correspond to higher $\delta^{18}\text{O}_{\text{BSi}}$. This relation is further corroborated by the difference in $\delta^{18}\text{O}_{\text{precip}}$ in
383 Anchorage measured in the strongest and weakest AL years described above (Bailey et al., 2019).
384 This suggests $\delta^{18}\text{O}_{\text{BSi}}$ reflects changes in storm-track-driven $\delta^{18}\text{O}_{\text{precip}}$ in addition to P-E, which have
385 opposing influences on the $\delta^{18}\text{O}_{\text{lake}}$ of SIL and other water bodies in the study area (Fig. 3), making it
386 challenging to constrain the relative importance of these two related hydroclimatic influences in the
387 paleo-data. These opposing influences might also explain the relative stability of $\delta^{18}\text{O}_{\text{BSi}}$ at SIL,
388 because the decrease in $\delta^{18}\text{O}_{\text{BSi}}$ resulting from wetter conditions would be balanced by changes in
389 $\delta^{18}\text{O}_{\text{precip}}$. The offset between $\delta^{18}\text{O}_{\text{lake}}$ at SIL at local river and ground waters (~7‰; Table S2) is
390 larger than the difference between $\delta^{18}\text{O}_{\text{precip}}$ in the strongest and weakest AL years (up to 3‰; Bailey
391 et al., 2019). However, the overall range of variability in measured $\delta^{18}\text{O}_{\text{precip}}$ at Anchorage from
392 2005–2018 is 31.1‰ (average = –16.5‰; min = –36.8‰; max = –5.7‰; Bailey et al., 2019),

393 indicating that changes in $\delta^{18}\text{O}_{\text{precip}}$ might be capable of driving larger fluctuations in $\delta^{18}\text{O}_{\text{lake}}$ if
394 sustained on longer timescales. The relationship between binned $\delta^{18}\text{O}_{\text{BSi}}$ values with known shifts in
395 the PDO and AL indices demonstrate these modes of variability likely influence lake sediment
396 sequences on multi-decadal timescales, despite the lack of significant correlation on an event-specific
397 basis (Bailey et al., 2019) where storm-to-storm variability and local topography might be more
398 likely to confound the fingerprint of these synoptic-scale patterns.

399 **4.1.2 Controls on BSi**

400 BSi in high altitude/latitude lakes might respond to a number of factors, including length of the ice-
401 free season (McKay et al., 2008), nutrient availability (Perren et al., 2017), dilution by minerogenic
402 material associated with storminess (Krawiec and Kaufman, 2014), or changing water chemistry that
403 can influence preservation (Bradbury et al., 1989). SEM imaging of diatoms in SIL sediments
404 indicates that dissolution is not a factor affecting BSi in this sequence (Fig. 4). At SIL, BSi over the
405 instrumental period is negatively correlated with annual precipitation at Kenai airport ($r = -0.66$, $p =$
406 0.01) (Fig. 10C), with periods of higher precipitation corresponding with lower BSi. This result is
407 consistent with a dilution effect whereby increased storminess in the Kenai lowlands causes increased
408 transport of minerogenic material to the lake, and subsequently dilutes BSi, as has been reported
409 elsewhere in southern Alaska (Krawiec and Kaufman, 2014).

410 While apparent over the instrumental period, dilution by clastic material related to storminess is
411 unlikely to be the only control on BSi at SIL. For example, BSi and organic matter content inferred
412 from LOI (Anderson et al., 2019) are moderately correlated ($r = 0.42$, $p < 0.01$) (Fig. S4). Organic
413 matter (OM) fluctuations typically represent combined autochthonous and allochthonous OM and
414 can therefore be tied to catchment-scale sediment composition and productivity (e.g. Shuman, 2003),
415 changes in lake level (e.g. Digerfeldt et al., 1992), or dilution by minerogenic material (e.g. Nesje and

416 Dahl, 2001). If OM and BSi differ, then BSi might reflect processes that impact diatoms directly,
417 such as changes in seasonality (Buczko et al., 2018) or taxon-specific responses of diatoms to
418 changing climate or environmental conditions (Lotter and Hofmann, 2003). The correlation between
419 OM and BSi suggests organic content and diatom abundance in SIL sediments are driven by
420 catchment-scale processes to some extent. However, the highest OM in the lake sediments occurs in
421 the early Holocene (~11.5 – 9.5 ka cal BP) and late Holocene (~2.5 – 0.5 ka cal BP), while the period
422 of highest BSi content occurs from ~6.3 – 4.6 ka cal BP, suggesting some independent controls on
423 OM and BSi have been responsible for fluctuations in these metrics as recorded at SIL on century- to
424 multi-millennial timescales (Fig. 7; Fig. S4), and these controls may not have been stable over the
425 course of the sediment sequence. For example, relatively high OM from ~11.5 – 9.5 ka cal BP might
426 reflect heightened organic input from the *Populus-Alnus-Salix* hardwood forest that occupied the
427 region surrounding SIL prior to the arrival of *Picea* (Anderson et al., 2019). Increasing BSi from ~9 –
428 6 ka cal BP might be related to increasing summer temperature, reflected by the chironomid-based
429 reconstruction from nearby Rainbow Lake (Clegg et al., 2011).

430 **4.2 Holocene hydroclimate and environmental change**

431 **4.2.1 Pleistocene-Holocene transition (~12.3 – 11 ka cal BP)**

432 Age-model uncertainties in the basal sediments of the SIL cores make it difficult to extrapolate
433 results older than ~12.3 ka cal BP (Fig. 6). The continuous deposition of organic sediments after
434 ~12.3 ka cal BP is roughly coincident with the inferred lake-level rise during the middle of the
435 Younger Dryas (12.2 ka cal BP) at Discovery Pond, 22 km to the north (Kaufman et al., 2010). Lake
436 level likely started rising prior to this at SIL, as the top of the basal inorganic unit is ~20 cm lower
437 than the sediments dated to ~12.3 ka cal BP, but age model uncertainties preclude an exact
438 assessment of when this occurred.

439 The oldest diatomaceous sediments at SIL (~11.9 ka cal BP) contain a high relative abundance of *S.*
440 *pinnata* (PC1; Fig. 8; Fig. 11A), a small, pioneering, fragiliarioid species typical of lakes in recently
441 stabilized landscapes (Smol et al., 2005; Hausmann and Pienitz, 2009). The dominance of *P.*
442 *saprophila*, a large benthic diatom, might also indicate that lake levels were low enough to yield a
443 proportionately extensive littoral zone, allowing a generally rare benthic diatom to succeed in this
444 environment (Wolin and Duthie, 1999). BSi content remains low (< 5%) throughout this interval
445 (Fig. 11C), most likely due to cold temperatures and turbid waters in the recently deglaciated
446 landscape prohibiting large diatom blooms.

447 The $\delta^{18}\text{O}_{\text{BSi}}$ values show large excursions during the Pleistocene-Holocene transition (Fig. 11D):
448 $\delta^{18}\text{O}_{\text{BSi}}$ decreases from +29.2‰ at ~12 ka cal BP to +26.2‰ (the lowest recorded $\delta^{18}\text{O}_{\text{BSi}}$) at ~11.5 ka
449 cal BP, then increases to +30.7‰ at ~11 ka cal BP. These changes in $\delta^{18}\text{O}_{\text{BSi}}$ occur during a major
450 vegetation transition to hardwood forest inferred from the SIL pollen record (Anderson et al., 2019).
451 Large climate shifts inferred from proxy data during the late YD and earliest Holocene have been
452 observed in other records from this region, including at nearby Horse Trail Fen, where inferred $\delta^{18}\text{O}$
453 of environmental water was substantially lower from 11.7 – 10.8 ka cal BP than during the following
454 ~2 ka cal BP (Jones et al., 2019) (Fig. 11E). At Hundred Mile Lake in the Matanuska Valley, Yu et
455 al. (2008) also found positive excursions in $\delta^{18}\text{O}_{\text{carbonate}}$ from *Pisidium* shells, ostracods, and *Chara*
456 encrustations in the earliest Holocene (~11.7 – 11.4 ka cal BP), followed by deviations in $\delta^{18}\text{O}_{\text{carbonate}}$,
457 organic matter, carbonate, and silicate content from ~11.2 – 10.9 ka cal BP during a period of
458 inferred earliest Holocene warmth. At Lone Spruce Pond in southwestern Alaska, shifts in BSi as
459 well as $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ of organic matter occur between ~12 – 11.5 ka cal BP (Kaufman et al., 2012),
460 and are associated with a warming climate. The mechanism for these excursions in the late YD and
461 earliest Holocene is unclear, though Kaufman et al. (2010) suggest that a strengthened AL in the
462 latter half of the YD promoted wetter and warmer climate conditions across southern Alaska, a

463 possibility that is echoed by Jones et al. (2019). While the precise cause of the changes in $\delta^{18}\text{O}_{\text{BSi}}$ at
464 SIL is not certain, it seems that a rapidly changing hydroclimate and environment are important
465 features of the glacial-interglacial transition, and that atmospheric circulation patterns may be one of
466 the drivers.

467 **4.2.2 Early and middle Holocene (~11 – 4.5 ka cal BP)**

468 Dated transitions from minerogenic sediments to terrestrial peat in satellite fens surrounding SIL
469 from ~9.5 – 7.7 ka cal BP (Fig. 11F) indicate that the water level at SIL declined during the early
470 Holocene to several meters lower compared to the late Holocene water level, as revealed by the ice-
471 shoved rampart analyses. The ~2 m fen profiles of wet sedge-dominated peat indicate a generally
472 rising water level throughout the mid and into the late Holocene, where the rising zone of peat
473 accumulation tracked the rising lake level. One lake sediment core that captures the upper ~6 ka cal
474 BP from SIL (SIL17-1) shows a basal beach sand (Fig. S1), indicating lake level was rising following
475 an early Holocene low stand.

476 In the earliest Holocene, the SIL diatom assemblage is dominated by facultatively planktonic and
477 large benthic taxa. By ~10.4 ka cal BP, the relative abundance of these habitat types diminishes,
478 giving way to a high relative abundance of *D. stelligera* (PC2; Fig. 8; Fig. 11A), a small planktonic
479 diatom that has been observed to be related to increases in lake nutrient content at other sites (Law et
480 al., 2015). The increase in *D. stelligera* at SIL coincides with the highest relative abundances of
481 *Alnus* pollen (Anderson et al., 2019), a genus known to increase local nitrogen availability (Shaftel et
482 al., 2011). Perren et al. (2017) documented the same relationship between *D. stelligera* and *Alnus* at
483 Lone Spruce Pond in southwestern Alaska, where they peak later in the Holocene. Given the
484 evidence for lowered lake levels at SIL (described above) and warm, dry conditions elsewhere in the
485 Kenai lowlands (Anderson et al., 2006; Jones et al., 2009; Anderson et al., 2019) at this time, the

486 increase in *D. stelligera* is likely related to the aforementioned landscape and soil processes rather
487 than to a high lake level *per se*, as the higher relative abundance of planktonic diatoms might suggest
488 (Wolin and Duthie, 1999). Following these high abundances of *D. stelligera*, the diatom assemblage
489 is dominated by facultatively planktonic taxa, making interpretations of lake-level changes difficult,
490 but indicating continued nitrogen enrichment (McGowan et al., 2005; Scheffer and van Nes, 2007).
491 At ~6 ka cal BP there is an increase in the percentage of *D. stelligera* and *Aulacoseira* spp.,
492 potentially due to rising lake levels increasing the water column depth and providing more habitat for
493 colonization by these planktonic diatom taxa.

494 Low and progressively increasing BSi from ~11.5 – 8 ka cal BP likely reflects the prevalent and
495 increasing presence of *Alnus* on the landscape (Anderson et al., 2019) (Fig. 11B), which fixes
496 nitrogen and encourages diatom blooms (Perren et al., 2017) (Fig. 11C). The highest BSi in the
497 dataset then occur between ~6.3 and ~4.6 ka cal BP, approximately concurrent with the highest
498 pollen concentrations (Anderson et al., 2019), though these high BSi values are interrupted by a
499 minimum at ~5.5 ka cal BP. These fluctuations might represent the influence of changes in runoff,
500 which controls mineral delivery to the lake, coupled with increased land cover and productivity as a
501 result of a shift to a wetter climate. Changes in runoff might also impact the delivery of other limiting
502 nutrients for diatom growth, such as silicon and phosphorus, which would cause changes in diatom
503 abundance. The arrival and establishment of *Picea mariana* at ~4.5 ka cal BP (Anderson et al., 2019)
504 is thought to be associated with a shift to wetter climate (Hu et al., 1996; Lynch et al., 2002),
505 providing an additional line of evidence for a shift in hydroclimate at this time.

506 Persistent fluctuations in $\delta^{18}\text{O}_{\text{BSi}}$ during most of the early and mid Holocene (Fig. 11D) are difficult
507 to attribute to any one hydroclimate variable. Jones et al. (2014, 2019) interpreted elevated $\delta^{18}\text{O}$ of
508 both total organic matter (TOM) and cellulose of peat and rapid peat accumulation in nearby Horse
509 Trail Fen to represent a weakened AL, an increased contribution of summer precipitation, and overall

510 wet conditions in the early Holocene. The inverse relation between the Horse Trail Fen and SIL $\delta^{18}\text{O}$
511 values (Fig. 11D-E) might be explained by the different seasonal influences on $\delta^{18}\text{O}_{\text{TOM}}$ in peat and
512 $\delta^{18}\text{O}_{\text{BSi}}$ in lake sediment. Peat accumulation is heavily dependent on summer moisture, and therefore
513 likely reflects a disproportionate influence of summer precipitation compared to climate proxies
514 found in lake sediments, which instead are more likely to reflect annual average or winter-dominated
515 conditions as represented by $\delta^{18}\text{O}_{\text{lake}}$ (Jones et al., 2014). LaBrecque and Kaufman (2016) also found
516 evidence for the advance of an outlet glacier from ~10.8 – 9.8 ka cal BP at Emerald Lake in the
517 Kenai Mountains, which they interpret as indicative of either lower summer temperature or higher
518 winter snowfall. In contrast, several studies (Anderson et al., 2006; Jones et al., 2009; Anderson et
519 al., 2019) have interpreted the early Holocene as a dry period in the Kenai lowlands based on pollen
520 and plant macrofossil evidence. The early-mid Holocene SIL $\delta^{18}\text{O}_{\text{BSi}}$ record does little to corroborate
521 or reject any of these hypotheses regarding hydroclimate, though the diatom assemblage and fen peat
522 data support the notion that lake levels were low and progressively rose following the early
523 Holocene.

524 $\delta^{18}\text{O}_{\text{BSi}}$ values remained relatively low (mean = +29.1‰, $n = 41$) until ~5.7 ka cal BP, when they
525 increased steadily until reaching sustained higher values (~ +31.2‰) by ~4.5 ka cal BP,
526 encompassing the largest step-wise shift in the Holocene (Fig. 11D). This observed shift to
527 higher $\delta^{18}\text{O}_{\text{BSi}}$ at ~4.5 ka cal BP could be interpreted to reflect a decrease in P-E, or drier conditions.
528 However, periodic late Holocene increases in the relative abundance of planktonic diatoms at SIL,
529 the presence of peats in satellite fens following the early Holocene low stand, and regional evidence
530 for wetter conditions at this time (Anderson et al., 2006) are all lines of evidence that challenge this
531 interpretation. If increased evaporation (decreased P-E) is not responsible for increasing $\delta^{18}\text{O}_{\text{lake}}$ and
532 $\delta^{18}\text{O}_{\text{BSi}}$ at ~4.5 ka cal BP, another possible driver of the shift would be an increase in $\delta^{18}\text{O}_{\text{precip}}$ values.
533 Isotopically heavier precipitation could have occurred due to a change in average storm track

534 trajectories associated with a shift to a stronger AL (Fig. 3), which would favor meridional
535 atmospheric flow and therefore relatively less rain-out of ^{18}O (as described in Section 4.1.1).
536 Numerous recent studies (Barron and Anderson, 2011; Jones et al., 2014; Bailey et al., 2018) have
537 suggested that the AL strengthened and increased in variability between 5 – 4 ka cal BP, including at
538 nearby Horse Trail Fen (Jones et al., 2014; 2019) (Fig. 11E), which is the only other full Holocene
539 $\delta^{18}\text{O}$ record from the Kenai lowlands. While we cannot unilaterally attribute the increase in $\delta^{18}\text{O}_{\text{BSi}}$
540 from ~5.5 – 4.5 ka cal BP at SIL to an increase in $\delta^{18}\text{O}_{\text{precip}}$ rather than to decreased P-E based on the
541 datasets presented in this study, several lines of evidence indicate the climate became wetter and the
542 AL strengthened at ~4.5 ka cal BP, suggesting that a stronger AL is the more likely driver of this
543 shift. The aforementioned datasets that have previously identified a shift in AL activity at this time
544 span from southwestern Yukon to Adak Island (Fig. 1A), and together with SIL $\delta^{18}\text{O}_{\text{BSi}}$ indicate a
545 synoptic-scale shift in North Pacific ocean-atmosphere circulation and terrestrial climate conditions
546 around the boundary between the mid and late Holocene.

547 **4.2.3 Following the increase in $\delta^{18}\text{O}_{\text{BSi}}$ at ~4.5 ka cal BP (~4.5 – 1 ka cal BP)**

548 Diatom assemblages following the ~4.5 ka cal BP transition remain diverse, though the relative
549 abundance of planktonic taxa increases slightly from ~4 to 1 ka cal BP, suggesting a shift to a deeper
550 water column (Wolin and Duthie, 1999) (Fig. 11A). Intermittent increases in *Aulacoseira* spp., which
551 form heavy colonies requiring turbulence for suspension in the water column to remain in the photic
552 zone (Rühland et al., 2008; Lotter et al., 2010), could indicate the presence of persistent stronger
553 winds associated with storminess throughout the late Holocene (Wang et al., 2008; Andrén et al.,
554 2015; Solovieva et al., 2015).

555 Following the highest Holocene $\delta^{18}\text{O}_{\text{BSi}}$ values between ~4.7 – 2.7 ka cal BP (mean = +31.2‰, $n =$
556 12), $\delta^{18}\text{O}_{\text{BSi}}$ decreased from ~2.7 – 1 ka cal BP (mean = +30.2‰, $n = 13$), but still remained elevated

557 relative to the early and mid-Holocene (mean = +29.2‰, $n = 54$). The sustained enriched $\delta^{18}\text{O}$ values
558 likely indicate the continued contribution of ^{18}O enriched precipitation due to the shift to dominantly
559 stronger AL conditions (Fig. 11D). The slight decreasing trend in $\delta^{18}\text{O}$ values could be interpreted as
560 representative of a progressively weakening AL over the late Holocene (Fig. 3). However, BSi
561 decreases over this interval, potentially indicating an increase in storminess and subsequent BSi
562 dilution by mineral matter transported to the lake during storms (Fig. 11C). Elevated percentages of
563 planktonic taxa in the late Holocene might also indicate higher lake levels associated with increased
564 precipitation. Charcoal accumulation rates from nearby Paradox Lake (Anderson et al., 2006) also
565 reveal fire frequency declined in the late Holocene, implying regional climate was wetter at this time.
566 The neoglacial advance of glaciers in the Kenai Mountains in the last ~4 ka cal BP also indicates
567 regionally wetter conditions that promoted the accumulation of snow and ice (Barclay et al., 2009;
568 Kaufman et al., 2016). Additionally, ice-shoved ramparts above the modern and inferred paleo-
569 shorelines of SIL indicate lake levels were higher than either the early Holocene or the present by
570 ~1.5 ka cal BP (Fig. 11F). Therefore, it is possible that following the initial change in precipitation
571 source water at ~4.5 ka cal BP, the influence of P-E on $\delta^{18}\text{O}_{\text{lake}}$ became more prominent. Though the
572 opposing effects of changes in $\delta^{18}\text{O}_{\text{precip}}$ and P-E cannot be definitively teased apart, the evidence for
573 wetter conditions at SIL, in the Kenai lowlands, and across eastern Beringia (Kaufman et al., 2016)
574 supports the interpretation of an increase in precipitation during the late Holocene. We therefore
575 interpret the overall decrease in $\delta^{18}\text{O}_{\text{BSi}}$ over the late Holocene, combined with decreasing BSi and
576 periodic, intermittent increases in the abundance of planktonic diatoms, to indicate an overall wetter
577 climate during this interval, though these changes in climate conditions cannot be unequivocally
578 attributed to AL strength.

579 **4.2.4 Last millennium (~1 ka cal BP – present)**

580 $\delta^{18}\text{O}_{\text{BSi}}$ values decreased during the last millennium (mean = +28.3‰, $n = 27$), reaching the lowest
581 value since 11.5 ka cal BP (+26.5‰ at 1996 CE) (Fig. 11D). The initial decrease starting at ~1 ka cal
582 BP might have been related to the documented advance of glaciers in the First Millennium CE in
583 south-central Alaska (Barclay et al., 2009) and throughout the northeastern Pacific Cordilleran
584 (Reyes et al., 2006), suggesting colder and wetter conditions throughout this region during the last
585 millennium, consistent with lower $\delta^{18}\text{O}_{\text{BSi}}$ associated with both a P-E isotope imprint, and possibly
586 cooler air temperatures that could lower $\delta^{18}\text{O}_{\text{precip}}$ (Dansgaard, 1964). Superimposed on the
587 decreasing trend are excursions to lower $\delta^{18}\text{O}_{\text{BSi}}$ at ~1250 – 1400 CE and ~1700 – 1750 CE (Fig. 12).
588 These shifts correspond to documented glacial advances (Fig. 12; Wiles and Calkin, 1993; Wiles et
589 al., 1999; Daigle and Kaufman, 2009; LaBrecque and Kaufman, 2016) throughout the western Prince
590 William Sound region that have been attributed to early and late stages of the Little Ice Age (LIA),
591 and are therefore likely associated with changing regional hydroclimate conditions, which impacted
592 both glaciers and lakes. Given the negative correlation between $\delta^{18}\text{O}_{\text{BSi}}$ and annual precipitation at
593 Kenai airport over the instrumental record (Fig. 10B), it follows that these negative $\delta^{18}\text{O}_{\text{BSi}}$
594 excursions might represent wet intervals during the LIA that resulted in glacier expansion.
595 Additionally, ice-shoved rampart evidence for higher-than-modern lake levels persists from ~1090
596 CE until ~1550 CE (Fig. 11F). In light of the agreement between SIL $\delta^{18}\text{O}_{\text{BSi}}$ and evidence for
597 regional glacial advances, we interpret the continued and enhanced overall decreasing trend in
598 $\delta^{18}\text{O}_{\text{BSi}}$, coupled with decreasing BSi, as increasingly wet conditions in the last millennium.
599 Since 2000 CE, both $\delta^{18}\text{O}_{\text{BSi}}$ values and BSi increased, likely as a result of lowered lake levels,
600 reduced P-E (for $\delta^{18}\text{O}_{\text{BSi}}$), as well as increased temperatures and longer ice-free seasons (for BSi).
601 The percentage of planktonic diatoms also decreased, while facultatively planktonic taxa such as *S.*
602 *pinnata* (PC1; Fig. 8) increased to their highest abundances in the Holocene, particularly in surface

603 sediments. An increased abundance of *S. pinnata* has been previously associated with nutrient
604 enrichment (McGowan et al., 2005; Scheffer and van Nes, 2007) and disturbance (Anderson, 2000),
605 which is consistent with warming, drying, fire, and road construction on this landscape in recent
606 decades. The elevations of dated terrestrial peats from perched satellite fens also indicate that current
607 conditions are the driest in the Holocene, with modern lake levels well below both dated late
608 Holocene ice-shoved ramparts and inferred early Holocene lake level minima (Fig. 11F). These
609 dramatic recent changes have occurred at a time of unprecedented rates of climate warming (Alaska
610 Climate Research Center, 2009), suggesting the landscape and lakes in the Kenai lowlands will likely
611 be subjected to continued alteration.

612 **5 Conclusions**

613 The multi-proxy record of environmental change from Sunken Island Lake in the Kenai lowlands
614 reflects regional hydroclimate conditions over the Holocene. In the Kenai lowlands, a stronger AL is
615 associated with an increase in storms that track from the south and deliver more precipitation that is
616 relatively enriched in ^{18}O . An increase in $\delta^{18}\text{O}_{\text{BSi}}$ at ~ 4.5 ka cal BP implies an increase in $\delta^{18}\text{O}_{\text{precip}}$
617 associated with AL strength, supporting prior work showing enhanced AL strength around 4 ka cal
618 BP (Barron and Anderson, 2011; Jones et al., 2014; Bailey et al., 2018). Multiple lines of evidence
619 support the interpretation of an increasingly wet late Holocene (beginning at ~ 4.5 ka cal BP)
620 compared to the early and mid Holocene, including high shorelines and ice-shoved ramparts,
621 decreased BSi, and increased planktonic diatom abundances between 4 and 1 ka cal BP. This trend
622 towards wetter conditions during the late Holocene has been reversed in the 21st Century, when
623 multiple lines of evidence indicate rapidly falling lake levels and decreased P-E. Despite the
624 uncertainty and challenges associated with reconciling multiple, competing influences on this oxygen
625 isotope record, our study demonstrates how the use of multiple hydroclimate indicators (BSi, diatom
626 flora, and $\delta^{18}\text{O}_{\text{BSi}}$) can help evaluate the relative contributions of these influences, rather than

627 attributing changes in $\delta^{18}\text{O}_{\text{BSi}}$ to only one primary influence (i.e. AL variability). Specifically,
628 intervals of lower BSi and/or higher planktonic diatom relative abundances clarify interpretations of
629 the mid to late Holocene $\delta^{18}\text{O}_{\text{BSi}}$ data where it might otherwise have been more difficult to discern
630 changes in $\delta^{18}\text{O}_{\text{precip}}$ from changes in P-E. Given the complexity of relating shifts in synoptic-scale
631 patterns to $\delta^{18}\text{O}_{\text{precip}}$ (Bailey et al., 2019), this approach of contemplating a more complex explanation
632 of $\delta^{18}\text{O}$ paleo-data may lead to fewer conflicting $\delta^{18}\text{O}$ interpretations among paleoclimate studies.

633 **6 Conflict of Interest**

634 The authors declare that the research was conducted in the absence of any commercial or financial
635 relationships that could be construed as a potential conflict of interest.

636 **7 Author Contributions**

637 EB led the study with guidance from DSK, co-led fieldwork, purified diatom oxygen isotope
638 samples, prepared and analyzed diatom assemblage samples, analyzed water isotope samples,
639 prepared samples for geochronological analyses, and wrote the original manuscript. DSK co-led field
640 work and reviewed early versions of the manuscript. ACGH provided expert assistance with the
641 interpretations of the diatom isotope and flora analyses. EEB participated in fieldwork and conducted
642 analyses on the SIL fen cores and ice-shoved rampart samples. RSA participated in fieldwork,
643 provided previously acquired sediments and data, and identified plant macrofossils for dating. MJL
644 oversaw the analysis of the diatom oxygen isotope samples at the British Geological Survey. SAS
645 produced and interpreted the BSi data. SEM analysed and interpreted $^{210}\text{Pb}/^{137}\text{Cs}$ gamma data. All
646 co-authors read and edited the manuscript.

647 **8 Funding**

648 This project was funded by National Science Foundation award 1602106 to DSK, and by grants and
649 scholarships to EB from the Geological Society of America (GSA), GSA Limnogeology Division,
650 the Phycological Society of America, LacCore/CSDCO, and the School of Earth and Sustainability at
651 Northern Arizona University.

652 **9 Acknowledgments**

653 We wish to thank the Kenai Natives Association for permitting our sediment and water sampling at
654 Sunken Island Lake; the U.S. Fish and Wildlife Service, Kenai National Wildlife Refuge for their
655 interest in this research and for assisting with field work; Eric Sandberg and Molly McCormick for
656 housing our field teams and assisting with water and sediment sampling; Nicholas McKay for
657 assistance with GeoChronR and for valuable insights that improved this manuscript; David Fortin
658 and Ann Jade Wong for assistance with field work; Dick Reger for describing the ice-shoved rampart
659 soil profiles; Katherine Whitacre, Kathryn Geyer, and Mackenzie Sanchez for assistance with
660 laboratory work; Chris Ebert for assisting with the preparation of the ¹⁴C samples; UC Irvine Keck
661 Carbon Cycle Laboratory for analyzing the ¹⁴C samples; Jack Lacey and Hilary Sloane for analysis of
662 the diatom oxygen isotopes at the British Geological Survey; Jamie Brown of the Colorado Plateau
663 Isotope Lab for analyzing the water samples; LacCore/CSDCO for assisting with Initial Core
664 Description; and Polar Field Services/CH2MHill for outfitting our field teams. We are grateful for
665 suggestions from Jonathan Tyler and one anonymous reviewer that greatly improved this manuscript.

666 **10 References**

- 667 Alaska Climate Research Center (2009). Temperature changes in Alaska.
668 <http://climate.gi.alaska.edu/ClimTrends/Change/TempChange.html> [accessed 11 Dec 2019]
- 669 Anderson, N.J. (2000). Diatoms, temperature and climatic change. *Eur. J. Phycol.* 35, 307-314.
670 doi:10.1080/09670260010001735911

- 671 Anderson, L., Abbott, M.B., Finney, B.P., and Burns, S.J. (2005). Regional atmospheric circulation
672 change in the North Pacific during the Holocene inferred from lacustrine carbonate oxygen isotopes,
673 Yukon Territory, Canada. *Quat. Res.* 64, 21-35. doi:10.1016/j.yqres.2005.03.005
- 674 Anderson, R.S., Hallett, D.J., Berg, E., Jass, R.B., Toney, J.L., de Fontaine, C.S., and DeVolder, A.
675 (2006). Holocene development of Boreal forests and fire regimes on the Kenai Lowlands of Alaska.
676 *Holocene* 16(6), 791-803. doi:10.1191/0959683606hol966rp
- 677 Anderson, R.S., Berg, E., Williams, C., and Clark, T. (2019). Postglacial vegetation community
678 change over an elevational gradient on the western Kenai Peninsula, Alaska: pollen records from
679 Sunken Island and Choquette Lakes. *J. Quat. Sci.* 34, 309-322. doi:10.1002/jqs.3102
- 680 Andrén, E., Klimaschewski, A., Self, A.E., Amour, N.S., Andreev, A.A., Bennett, K.D., Conley,
681 D.J., Edwards, T.W.D., Solovieva, N., and Hammarlund, D. (2015). Holocene climate and
682 environmental change in north-eastern Kamchatka (Russian Far East), inferred from a multi-proxy
683 study of lake sediments. *Global Planet. Change* 134, 41-54. doi:10.1016/j.gloplacha.2015.02.013
- 684 Appleby, P. (2001). "Chronostratigraphic techniques in recent sediments," in Tracking
685 Environmental Change Using Lake Sediments, Vol. 1. eds. W. Last and J. Smol. (Dordrecht, The
686 Netherlands: Kluwer Academic Publishers), 171-203
- 687 Bailey, H.L., Kaufman, D.S., Henderson, A.C.G., and Leng, M.J. (2015). Synoptic scale controls on
688 the $\delta^{18}\text{O}$ in precipitation across Beringia. *Geophys. Res. Lett.* 42, 4608-4616.
689 doi:10.1002/2015GL063983
- 690 Bailey, H.L., Kaufman, D.S., Sloane, H.J., Hubbard, A.L., Henderson, A.C.G., Leng, M.J., Meyer,
691 H., and Welker, J.M. (2018). Holocene atmospheric circulation in the central North Pacific: A new
692 terrestrial diatom and $\delta^{18}\text{O}_{\text{diatom}}$ dataset from the Aleutian Islands. *Quat. Sci. Rev.* 194, 27-38.
693 doi:10.1016/j.quascirev.2018.06.027
- 694 Bailey, H.L., Klein, E.S., and Welker, J.M. (2019). Synoptic and mesoscale mechanisms driver
695 winter precipitation $\delta^{18}\text{O}$ / $\delta^2\text{H}$ in south-central Alaska. *J. Geophys. Res.-Atmos.* 124, 4252-4266.
696 doi:10.1029/2018JD030050
- 697 Barclay, D.J., Wiles, G.C., and Calkin, P.E. (2009). Tree-ring crossdates for a First Millennium AD
698 advance of Tebenkof Glacier, southern Alaska. *Quat. Res.* 71, 22-26.
699 doi:10.1016/j.yqres.2008.09.005
- 700 Barron, J. A., and Anderson, L. (2011). Enhanced Late Holocene ENSO/PDO expression along the
701 margins of the eastern North Pacific. *Quat. Int.* 235, 3-12. doi:10.1016/j.quaint.2010.02.026
- 702 Berkelhammer, M.B., Stott, L.D., Yoshimura, K., Johnson, K., and Sinha, A. (2012). Synoptic and
703 mesoscale controls on the isotopic composition of precipitation in the western United States. *Clim.*
704 *Dynam.* 38, 433-454. doi:10.1007/s00382-011-1262-3
- 705 Blaauw, M., and Christen, J.A. (2011). Flexible paleoclimate age-depth models using an
706 autoregressive gamma process. *Bayesian Anal.* 6(3), 457-474. doi:10.1214/11-BA618
- 707 Boes, E., Van Daele, M., Moeraut, J., Schmidt, S., Jensen, B.J.L, Praet, N., Kaufman, D., Haessler,
708 P., Loso, M.G., and De Batist, M. (2018). Varve formation during the past three centuries in three

- 709 large proglacial lakes in south-central Alaska. *Geol. Soc. Am. Bull.* 130, 757-774.
710 doi:10.1130/B31792.1
- 711 Bradbury, J.P., Forester, R.M., and Thompson, R.S. (1989). Late Quaternary paleolimnology of
712 Walker Lake, Nevada. *J. Paleolimnol.* 1(4), 249-267. doi:10.1007/BF00184000
- 713 Brandriss, M.E., O'Neil, J.R., Edlund, M.B., and Stoermer, E.F. (1998). Oxygen isotope fractionation
714 between diatomaceous silica and water. *Geochim. Cosmochim. Acta* 62, 1119–1125.
715 doi:10.1016/S0016-7037(98)00054-4
- 716 Bretherton, C.S., Widmann, M., Dymnikov, V.P., Wallace, J.M., and Bladé, I. (1998). The effective
717 number of spatial degrees of freedom of a time-varying field. *J. Climate* 12, 1990-2009.
718 doi:10.1175/1520-0442(1999)012<1990:TENOSD>2.0.CO;2
- 719 Brewer, T.S., Leng, M.J., Mackay, A.W., Lamb, A.L., Tyler, J.T., Marsh, N.G., 2008. Unraveling
720 contamination signals in biogenic silica oxygen isotope composition: the role of major and trace
721 element geochemistry. *J. Quat. Sci.* 23(4), 321-330. doi:10.1002/jqs.1171
- 722 Buczkó, K., Szurdoki, E., Braun, M., and Magyari, E. (2018). Reconciling diverse diatom-based lake
723 responses to climate change in four mountain lakes in the South-Carpathian Mountains during the
724 last 17 kyrs. *Quat. Int.* 477, 117-137. doi:10.1016/j.quaint.2017.03.013
- 725 Cayan, D.R., and Peterson, D.H. (1989). “The influence of North Pacific circulation on streamflow in
726 the West,” in *Aspects of Climate Variability in the Pacific and Western Americas*, Vol. 55, ed. D.H.
727 Peterson (Washington, DC: American Geophysical Union), 375–398
- 728 Clegg, B.F., Kelly, R., Clarke, G.H., Walker, I.R., and Hu, F.S. (2011). Nonlinear response of
729 summer temperature to Holocene insolation forcing in Alaska. *P. Natl. Acad. Sci. USA.* 108(48),
730 19299-19304. doi:10.1073/pnas.1110913108
- 731 Crespin, J., Sylvestre, F., Alexandre, A., Sonzogni, C., Pailles, C., and Perga, M-E. (2010). Re-
732 examination of the temperature-dependent relationship between $\delta^{18}\text{O}_{\text{diatoms}}$ and $\delta^{18}\text{O}_{\text{lake water}}$ and
733 implications for paleoclimate inferences. *J. Paleolimnol.* 44, 547-557. doi:10.1007/s10933-010-9436-
734 2
- 735 Daigle, T.A., and Kaufman, D.S. (2009). Holocene climate inferred from glacier extent, lake
736 sediment and tree rings at Goat Lake, Kenai Mountains, Alaska, USA. *J. Quat. Sci.* 24, 33-45.
737 doi:10.1002/jqs.1166
- 738 Dansgaard, W. (1964). Stable isotopes in precipitation. *Tellus*, 16(4), 436–468. doi:10.1111/j.2153-
739 3490.1964.tb00181.x
- 740 Dean, W.E. (1974). Determination of carbonate and organic matter in calcareous sediments and
741 sedimentary rocks by loss on ignition; comparison with other methods. *J. Sed. Res.* 44(1), 242-248.
742 doi:10.1306/74D729D2-2B21-11D7-8648000102C1865D
- 743 Digerfeldt, G., Almendinger, J.E., and Björck, S. (1992). Reconstruction of past lake levels and their
744 relation to groundwater hydrology in the Parkers Prairie sandplain, west-central Minnesota.
745 *Palaeogeogr. Paleocl.* 94, 99-118. doi:10.1016/0031-0182(92)90115-L

- 746 Dodd, J.P., and Sharp, Z.D. (2010). A laser fluorination method for oxygen isotope analysis of
747 biogenic silica and a new oxygen isotope calibration of modern diatoms in freshwater environments.
748 *Geochim. Cosmochim. Acta* 74(4), 1381-1390. doi:10.1016/j.gca.2009.11.023
- 749 Dodd, J.P., Sharp, Z.D., Fawcett, P.J., Brearley, A.J., McCubbin, F.M., 2012. Rapid post-mortem
750 maturation of diatom silica oxygen isotope values. *Geochem. Geophys.* 13(9), Q09014.
751 doi:10.1029/2011GC004019
- 752 Fisher, D., Osterberg, E., Dyke, A., Dahl-Jensen, D., Demuth, M., Zdanowicz, C., Bourgeois, J.,
753 Koerner, R.M., Mayewski, P., Wake, C., Kreutz, K., Steig, E., Zheng, J., Yalcin, K., Goto-Azuma,
754 K., Luckman, B., and Rupper, S. (2008). The Mt Logan Holocene - late Wisconsinan isotope record:
755 tropical Pacific - Yukon connections. *Holocene* 18, 667-677. doi:10.1177/0959683608092236
- 756 Foged N. (1971). Diatoms found in a bottom sediment sample from a small deep lake on the
757 Northern Slope, Alaska. *Nova Hedwigia* 21(1-4): 923-1034
- 758 Foged, N. (1981). Diatoms in Alaska. *Bibliotheca Phycologica* 53, 1-317
- 759 Grimm, E.C. (1987). Coniss - a Fortran-77 program for stratigraphically constrained cluster-analysis
760 by the method of incremental sum of squares. *Computat. Geosci.* 13, 13-35. doi:10.1016/0098-
761 3004(87)90022-7
- 762 Grimm, E.C. (2015). TILIA Software. Version 2.1.1. <https://www.tiliait.com/download/>
- 763 Hamilton, S., and Shennan, I. (2005). Late Holocene great earthquakes and relative sea-level change
764 at Kenai, southern Alaska. *J. Quat. Sci.* 20, 95-111. doi:10.1002/jqs.903
- 765 Hausmann, S., and Pienitz, R. (2009). Seasonal water chemistry and diatom changes in six boreal
766 lakes of the Laurentian Mountains (Québec, Canada): impacts of climate and timber harvesting.
767 *Hydrobiologia* 635(1), 1-14. doi:10.1007/s10750-009-9855-0
- 768 Hu, F.S., Brubaker, L.B., and Anderson, P.M. (1996). Boreal ecosystem development in the
769 northwestern Alaska Range since 11,000 yr B.P. *Quat. Res.* 45(2), 188-201.
770 doi:10.1006/qres.1996.0019
- 771 Jones M.C., Peteet D.M., Kurdyla D., and Guilderson, T. (2009). Climate and vegetation history
772 from a 14,000-year peatland record, Kenai Peninsula, Alaska. *Quat. Res.* 72, 207-217.
773 doi:10.1016/j.yqres.2009.04.002
- 774 Jones, M.C., Wooller, M., and Peteet, D.M. (2014). A deglacial and Holocene record of climate
775 variability in south-central Alaska from stable oxygen isotopes and plant macrofossils in peat. *Quat.*
776 *Sci. Rev.* 87, 1-11. doi:10.1016/j.quascirev.2013.12.025
- 777 Jones, M.C., Anderson, L., Keller, K., Nash, B., Littell, V., Wooller, M., and Jolley, C.A. (2019). An
778 assessment of plant species differences on cellulose oxygen isotopes from two Kenai Peninsula
779 Alaska peatlands: implications for hydroclimatic reconstructions. *Front. Earth Sci.* 7(25), 1-25.
780 doi:10.3389/feart.2019.00025
- 781 Juillet-Leclerc, A., Labeyrie, L. (1987). Temperature dependence of the oxygen isotopic fractionation
782 between diatom silica and water. *Earth Planet. Sci. Lett.* 84, 69-74. doi:10.1016/0012-

- 783 821X(87)90177-4
- 784 Kaufman D.S., Anderson R.S., Hu F.S., Berg, E., and Werner, A. (2010). Evidence for a variable and
785 wet Younger Dryas in southern Alaska. *Quat. Sci. Rev.* 29: 1445–1452.
786 doi:10.1016/j.quascirev.2010.02.025
- 787 Kaufman, D.S., Axford, Y., Anderson, R.S., Lamoureux, S.F., Schindler, D.E., Walker, I.R., and
788 Werner, A. (2012). A multi-proxy record of the Last Glacial Maximum and last 14,500 years of
789 paleoenvironmental change at Lone Spruce Pond, southwestern Alaska. *J. Paleolimnol.* 48, 9-26.
790 doi:10.1007/s10933-012-9607-4
- 791 Kaufman, D.S., Axford, Y.L., Henderson, A.C.G., McKay, N.P., Oswald, W.W., Saenger, C.,
792 Anderson, R.S., Bailey, H.L., Clegg, B., Gajewski, K., Hu, F.S., Jones, M.C., Massa, C., Routson,
793 C.C., Werner, A., Wooller, M.J., and Yu, Z. (2016). Holocene climate changes in eastern Beringia
794 (NW North America) — A systematic review of multi-proxy evidence. *Quat. Sci. Rev.* 147, 312-339.
795 doi:10.1016/j.quascirev.2015.10.021
- 796 Khapalova, E.A., Jandhyala, V.K., Fotopoulos, S.B., and Overland, J.E. (2018). Assessing change-
797 points in surface air temperature over Alaska. *Front. Enviro. Sci.* 6(121), 1-17.
798 doi:10.3389/fenvs.2018.00121
- 799 Koning, E., Gehlen, M., Flank, A.M., Calas, G., Epping, E., 2007. Rapid post-mortem incorporation
800 of aluminum in diatom frustules: evidence from chemical and structural analyses. *Marine Chem.* 106,
801 208-222. doi:10.1016/j.marchem.2006.06.009
- 802 Krammer, K., and Lange-Bertalot, H. (1986-1991). Bacillariophyceae Band 1-4. Stuttgart: Gustav
803 Fischer Verlag.
- 804 Krawiec, A.C.L., and Kaufman, D.S. (2014). Holocene storminess inferred from sediments of two
805 lakes on Adak Island, Alaska. *Quat. Res.* 82, 73-84. doi:10.1016/j.yqres.2014.02.007
- 806 LaBrecque, T.S., and Kaufman, D.S. (2016). Holocene glacier fluctuations inferred from lacustrine
807 sediment, Emerald Lake, Kenai Peninsula, Alaska. *Quat. Res.* 85, 34-43.
808 doi:10.1016/j.yqres.2015.11.004
- 809 Law, A.C., Anderson, N.J., and McGowan, S. (2015). Spatial and temporal variability of lake
810 ontogeny in south-western Greenland. *Quat. Sci. Rev.* 126, 1-16.
811 doi:10.1016/j.quascirev.2015.08.005
- 812 Leng, M.J., and Barker, P.A. (2006). A review of the oxygen isotope composition of lacustrine
813 diatom silica for palaeoclimate reconstruction. *Earth-Sci. Rev.* 75, 5-27.
814 doi:10.1016/j.earscirev.2005.10.001
- 815 Leng, M.J., and Sloane, H.J. (2008). Combined oxygen and silicon isotope analysis of biogenic
816 silica. *J. Quat. Sci.* 23, 313-319. doi:10.1002/jqs.1177
- 817 Lotter, A.F., and Hofmann, G. (2003). “The development of the late-glacial and Holocene diatom
818 flora in Lake Sedmo Rilsko (Rila Mountains, Bulgaria),” in *Aspects of Palynology and*
819 *Palaeoecology*, ed. S. Tonkov (Moscow, Russia: Pensoft Publishers), 171-183

- 820 Lotter, A.F., Pienitz, R., and Schmidt, R. (2010). "Diatoms as indicators of environmental change in
821 subarctic and alpine regions," in *The Diatoms: Application for the Environmental and Earth*
822 *Sciences*, eds. J.P. Smol and E.F. Stoermer (Cambridge: United Kingdom: Cambridge University
823 Press), 97-115
- 824 Lynch, J.A., Clark, J.S., Bigelow, N.H., Edwards, M.E., and Finney, B.P. (2002). Geographic and
825 temporal variations in fire history in boreal ecosystems of Alaska. *J. Geophys. Res.-Atmos.* 108, NO.
826 D1, 8152. doi:10.1029/2001JD000332
- 827 Mann, D.G., McDonald, S.M., Mayer, M.M., Droop, S.J.M., Chepurnov, V.A., Loke, R.E., Ciobanu,
828 A., and Du Buf, J.M.H. (2004). The *Sellaphora pupula* species complex (Bacillariophyceae):
829 Morphometric analysis, ultrastructure and mating data provide evidence for five new species.
830 *Phycologia* 43, 459-482. doi:10.2216/i0031-8884-43-4-459.1
- 831 Mantua, N.J., Hare, S.R., Zhang, Y., Wallace, J.M., and Francis, R.C. (1997). A Pacific interdecadal
832 climate oscillation with impacts on salmon production. *B. Am. Meteorol. Soc.* 78, 1069-1079.
833 doi:10.1175/1520-0477(1997)078<1069:APICOW>2.0.CO;2
- 834 Menicucci, A.J., Spero, H.J., Matthews, J., and Parikh, S.J. (2017). Influence of exchangeable
835 oxygen on biogenic silica oxygen isotope data. *Chem. Geol.* 466, 710-721.
836 doi:10.1016/j.chemgeo.2017.07.020
- 837 Menicucci, A.J., Thunell, R.C., and Spero, H.J. (2019). 220 year diatom $\delta^{18}\text{O}$ reconstruction of the
838 Guaymas Basin thermocline using microfluorination. *Palaeogeogr. Paleocl.* 35(2), e2019PA003749.
839 doi:10.1029/2019PA003749
- 840 McGowan, S., Patoine, A., Graham, M.D., and Leavitt, P. (2005). Intrinsic and extrinsic controls on
841 lake phytoplankton synchrony as illustrated by algal pigments. *SIL Proceedings 1922-*
842 *2010*, 29(2), 794-798. doi:10.1080/03680770.2005.11902787
- 843 McKay, N.P., Kaufman, D.S., and Michelutti, N. (2008). Biogenic silica concentration as a high
844 resolution, quantitative temperature proxy at Hallet Lake, south-central Alaska. *Geophys. Res. Lett.*
845 35, L05709, doi:10.1029/2007GL032876
- 846 McKay, N.P., Emile-Geay, J., Heiser, C., and Khinder, D. (2018). GeoChronR development
847 repository. Retrieved from <http://dx.doi.org/10.5281/zenodo.60812>. doi:10.5281/zenodo.60812
- 848 McLaughlin, R.B., and Stone, J.L. (1986). Some late Pleistocene diatoms of the Kenai Peninsula,
849 Alaska. *Nova Hedwigia* 82, 1-148
- 850 Mock, C.J., Bartlein, P.J., and Anderson, P.M. (1998). Atmospheric circulation patterns and spatial
851 climatic variations in Beringia. *Int. J. Climatol.* 10, 1085-1104. doi:10.1002/(SICI)1097-
852 0088(199808)18:10<1085::AID-JOC305>3.0.CO;2-K
- 853 Mortlock, R.A., and Froelich, P.N. (1989). A simple method for the rapid determination of biogenic
854 opal in pelagic marine sediments. *Deep-Sea Res.* 36, 1415-1426. doi:10.1016/0198-0149(89)90092-7
- 855 Morley, D.W., Leng, M.J., Mackay, A.W., Sloane, H.J., Rioual, P., and Battarbee, R.W. (2004).
856 Cleaning of lake sediment samples for diatom oxygen isotope analysis. *J. Palaeolimnol.* 31, 391-
857 401. doi:10.1023/B: JOPL.0000021854.70714.6b

- 858 Moschen, R., Lücke, A., and Schleser, G.H. (2005). Sensitivity of biogenic silica oxygen isotopes to
859 changes in surface water temperature and paleoclimatology. *Geophys. Res. Lett.* 32, L07708.
860 doi:10.1029/2004GL22167
- 861 Nesje, A., and Dahl, S.O. (2001). The Greenland 8200 cal. yr BP event detected in loss-on-ignition
862 profiles in Norwegian lacustrine sediment sequences. *J. Quat. Sci.* 16(2), 155-166.
863 doi:10.1002/jqs.567
- 864 Overland, J.E., Adams, J.M., and Bond, N.A. (1999). Decadal variability of the Aleutian Low and its
865 relation to high-latitude circulation. *J. Clim.* 12, 1542-1548. doi:10.1175/1520-
866 0442(1999)012<1542:DVOTAL>2.0.CO;2
- 867 Paillard, D., Labeyrie, L., and Yiou, P. (1996). Macintosh program performs time-series
868 analysis. *Eos* 77(39), 379. doi:10.1029/96EO00259
- 869 Perren, B.B., Axford, Y., and Kaufman, D.S. (2017). Alder, nitrogen, and lake ecology: terrestrial-
870 aquatic linkages in the postglacial history of Lone Spruce Pond, southwestern Alaska. *PLoS ONE*
871 12(1), e0169106. doi:10.1371/journal.pone.0169106
- 872 Praet, N., Moernaut, J., Van Daele, M., Boes, E., Haeussler, P., Strupler, M., Schmidt, S., Loso, M.,
873 and Batist, M. (2017). Paleoseismic potential of sublacustrine landslide records in a high-seismicity
874 setting (south-central Alaska). *Mar. Geol.* 384, 103-119. doi:10.1016/j.margeo.2016.05.004
- 875 Porter, T.J., Pisaric, M.F.J., Field, R.D., Kokelj, T.W.D., deMontigny, P., Healy, R., and LeGrande,
876 A.N. (2014). Spring-summer temperatures since AD 1780 reconstructed from stable oxygen isotope
877 ratios in white spruce tree-rings from the Mackenzie Delta, northwestern Canada. *Clim. Dynam.* 42,
878 771-785. doi:10.1007/s00382-013-1674-3
- 879 Reimer, P.J., Bard, E., Bayliss, A., Beck, J.W., Blackwell, P.G., Ramsey, C.B., Buck, C.E., Cheng,
880 H., Edwards, R.L., Friedrich, M., Grootes, P.M., Guilderson, T.P., Haflidason, H., Hajdas, I., Hatté,
881 C., Heaton, T.J., Hoffmann, D.L., Hogg, A.G., Hughen, K.A., Kaiser, K.F., Kromer, B., Manning,
882 S.W., Niu, M., Reimer, R.W., Richards, D.A., Scott, E.M., Southon, J.R., Staff, R.A., Turney,
883 C.S.M., van der Plicht, J. (2013). IntCal13 and Marine13 radiocarbon age calibration curves, 0-
884 50,000 years cal BP. *Radiocarbon* 55(4), 1869-1887. doi:10.2458/azu_js_rc.55.16947
- 885 Reger, R.D., Sturmman, A.G., Berg, E.E., and Burns, P.A.C. (2007). A guide to the late Quaternary
886 history of northern and western Kenai Peninsula, Alaska: Guidebook 8. Anchorage, Alaska: State of
887 Alaska Department of Natural Resources, Division of Geological and Geophysical Surveys.
- 888 Reyes, A.V., Wiles, G.C., Smith, D.J., Barclay, D.J., Allen, S., Jackson, S., Larocque, S., Laxton, S.,
889 Lewis, D., Calkin, P.E., and Clague, J.J. (2006). Expansion of alpine glaciers in Pacific North
890 America in the first millennium A.D. *Geology* 34(1), 57-60. doi:10.1130/G21902.1
- 891 Rodionov, S.N., Overland, J.E., and Bond, N.A. (2005). Spatial and temporal variability of the
892 Aleutian climate. *Fish. Oceanogr.* 14, 3-21. doi:10.1111/j.1365-2419.2005.00363.x
- 893 Rodionov, S.N., Bond, N.A., and Overland, J.E. (2007). The Aleutian Low, storm tracks, and winter
894 climate variability. *Deep-Sea Res. II* 54, 2560-2577. doi:10.1016/j.dsr2.2007.08.002
- 895 Rühland, K., Paterson, A.M., and Smol, J.P. (2008). Hemispheric-scale patterns of climate-related

- 896 shifts in planktonic diatoms from North American and European lakes. *Glob. Change Biol.* 14(11),
897 2740e2754. doi:10.1111/j.1365-2486.2008.01670.x
- 898 Scheffer, M., and van Nes, E.H. (2007). Shallow lakes theory revisited: various alternative regimes
899 driven by climate, nutrients, depth and lake size. *Hydrobiologia* 584, 455-466. doi:10.1007/s10750-
900 007-0616-7
- 901 Schiff, C., Kaufman, D.S., Wolfe, A.P., Dodd, J., and Sharp, Z. (2009). Late Holocene storm-
902 trajectory changes inferred from the oxygen isotope composition of lake diatoms, south Alaska. *J.*
903 *Paleolimnol.* 41, 189-208. doi:10.1007/s10933-008-9261-z
- 904 Shaftel, R.S., King, R.S., and Back, J.A. (2011). Alder cover drives nitrogen availability in Kenai
905 lowland headwater streams, Alaska. *Biogeochemistry* 107, 135-148. doi:10.1007/s10533-010-9541-3
- 906 Shemesh, A., Charles, C.D., and Fairbanks, R.G. (1992). Oxygen isotopes in biogenic silica: global
907 changes in ocean temperature and isotopic composition. *Science* 256, 1434-1436.
908 doi:10.1126/science.256.5062.1434
- 909 Shuman, B. (2003). Controls on loss-on-ignition variation in cores from two shallow lakes in the
910 northeastern United States. *J. Paleolimnol.* 30, 371-385. doi:10.1023/B:JOPL.0000007226.68831e3
- 911 Smol, J.P., Wolfe, A.P., Birks, H.J.B., Douglas, M.S., Jones, V.J., Korhola, A., Pienitz, R., Rühland,
912 K., Sorvari, S., Antoniades, D., and Brooks, S.J. (2005). Climate-driven regime shifts in the
913 biological communities of arctic lakes. *P. Natl. Acad. Sci. USA.* 102(12), 4397e4402.
914 doi:10.1073/pnas.0500245102
- 915 Solovieva, N., Klimaschewski, A., Self, A.E., Jones, V.J., Andren, E., Andreev, A.A., Hammarlund,
916 D., Lepskaya, E.V., and Nazarova, L. (2015). The Holocene environmental history of a small coastal
917 lake on the north-eastern Kamchatka Peninsula. *Global Planet. Change* 134, 55e66.
918 doi:10.1016/j.gloplacha.2015.06.010
- 919 Spaulding, S.A., Bishop, I.W., Edlund, M.B., Lee, S. and Potapova, M. (2019). Diatoms of North
920 America. <https://diatoms.org/> [accessed 11 Dec 2019]
921
- 922 ter Braak, C.J.F., and Prentice, I.C. (1988). A theory of gradient analysis. *Adv. Ecol. Res.* 18,
923 271e317. doi:10.1016/S0065-2504(03)34003-6
- 924 Trenberth, K.E., and Hurrell, J.W. (1994). Decadal atmosphere-ocean variations in the Pacific. *Clim.*
925 *Dynam.* 9, 303-319. doi:10.1007/BF00204745
- 926 Tyler, J.J., Sloane, H.J., Rickaby, R.E.M., Cox, E.J., and Leng, M.J. (2017). Post-mortem oxygen
927 isotope exchange within cultured diatom silica. *Rapid. Commun. Mass Sp.* 31(20), 1749-1760.
928 doi:10.1002/rcm.7954
- 929 Wang, L., Lu, H., Liu, J., Gu, Z., Mingram, J., Chu, G., Li, J., Rioual, P., Negendank, J.F., Han, J.,
930 and Liu, T. (2008). Diatom-based inference of variations in the strength of Asian winter monsoon
931 winds between 17,500 and 6000 calendar years BP. *J. Geophys. Res.-Atmos.* 113, D2101.
932 doi:10.1029/2008JD010145

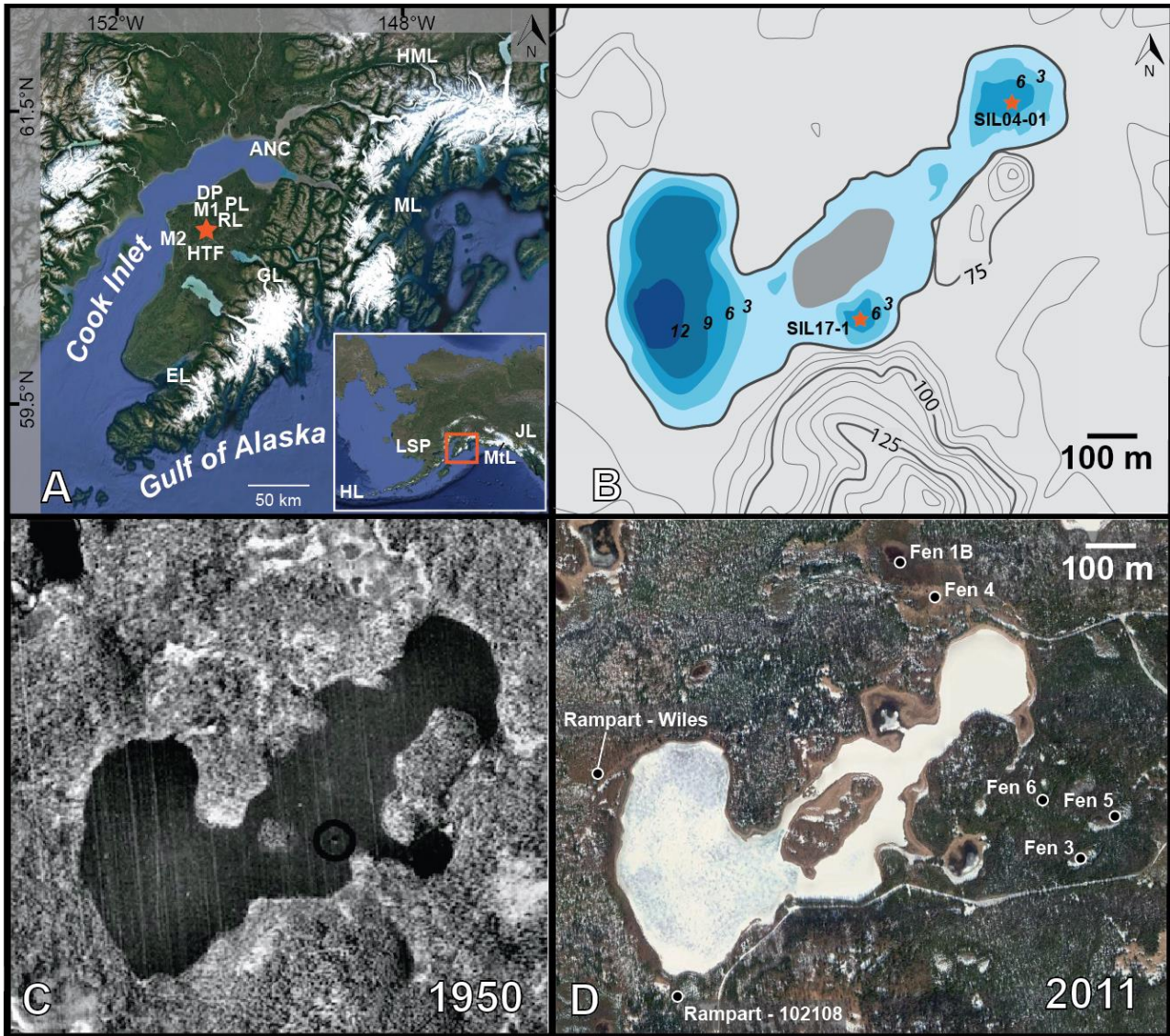
- 933 Wiles, G.C., and Calkin P.E. (1993). Neoglacial fluctuations and sedimentation of an iceberg-calving
934 glacier resolved with tree rings (Kenai Fjords National Park, Alaska). *Quat. Int.* 18, 35-42.
935 doi:10.1016/1040-6182(93)90051-G
- 936 Wiles, G.C., Barclay, D.J., and Calkin, P.E. (1999). Tree-ring-dated ‘Little Ice Age’ histories of
937 maritime glaciers from western Prince William Sound, Alaska. *Holocene* 9(2), 163-173.
938 doi:10.1191/095968399671927145
- 939 Wolin, J.A., and Duthie, H.C. (1999). “Diatoms as indicators of water level change in freshwater
940 lakes,” in *The Diatoms: Application for the Environmental and Earth Sciences*, eds. J.P. Smol and
941 E.F. Stoermer (Cambridge: United Kingdom: Cambridge University Press), 183–202
- 942 Yu, Z.C., Walker, K.N., Evenson, E.B., and Hadjas, I. (2008). Late glacial and early Holocene
943 climate oscillations in the Matanuska Valley, south-central Alaska. *Quat. Sci. Rev.* 27, 148-161.
944 doi:10.1016/j.quascirev.2007.02.020

945 **11 Supplementary Material**

946 See attached Supplementary Figures and Tables as well as Supplementary Data (Excel sheet).

947 **12 Data Availability Statement**

948 The depths and ages for all sediment cores used in this study can be found in the attached data
949 supplement. The BSi and $\delta^{18}\text{O}_{\text{BSi}}$ data can be found in the attached data supplement, and at the [World](#)
950 [Centre for Paleoclimatology](#). The diatom assemblage data can be found at the [Neotoma Paleoecology](#)
951 [Database](#). *Note to editor: Data will be transferred when the manuscript is accepted for publication.*



953

954 **Figure 1:** (A) Location of Sunken Island Lake in the Kenai lowlands indicated by the orange star.
 955 Locations of the following sites are marked: Anchorage (ANC), Kenai Moose Pens SNOTEL
 956 meteorological station (M1), Kenai airport meteorological station (M2), Heart Lake (HL; Bailey et
 957 al., 2018), Lone Spruce Pond (LSP; Kaufman et al., 2012), Mount Logan (MtL; Fisher et al., 2008),
 958 Jellybean Lake (JL; Anderson et al., 2005), Emerald Lake (EL; LaBrecque and Kaufman, 2016),
 959 Horse Trail Fen (HTF; Jones et al., 2014, 2019), Discovery Pond (DP; Kaufman et al., 2010),
 960 Paradox Lake (PL; Anderson et al., 2006), Rainbow Lake (RL; Clegg et al., 2011), Goat Lake (GL;

961 Daigle and Kaufman, 2009), Mica Lake (ML; Schiff et al., 2009); and Hundred Mile Lake (HML; Yu
962 et al., 2008). Images from Google Earth. (B) Bathymetric map of Sunken Island Lake, with water
963 depths shown in increments of 3 m, and contour lines in increments of 5 m. Orange stars indicate
964 sites of sediment cores analyzed for this study. (C-D) Aerial photographs of Sunken Island Lake in
965 1950 CE (DeVolder, *pers. comm.*) (C) and in 2011 CE (Google Earth) (D). Locations of sampled
966 satellite fens and ice-shoved ramparts are shown in (D).

967

968

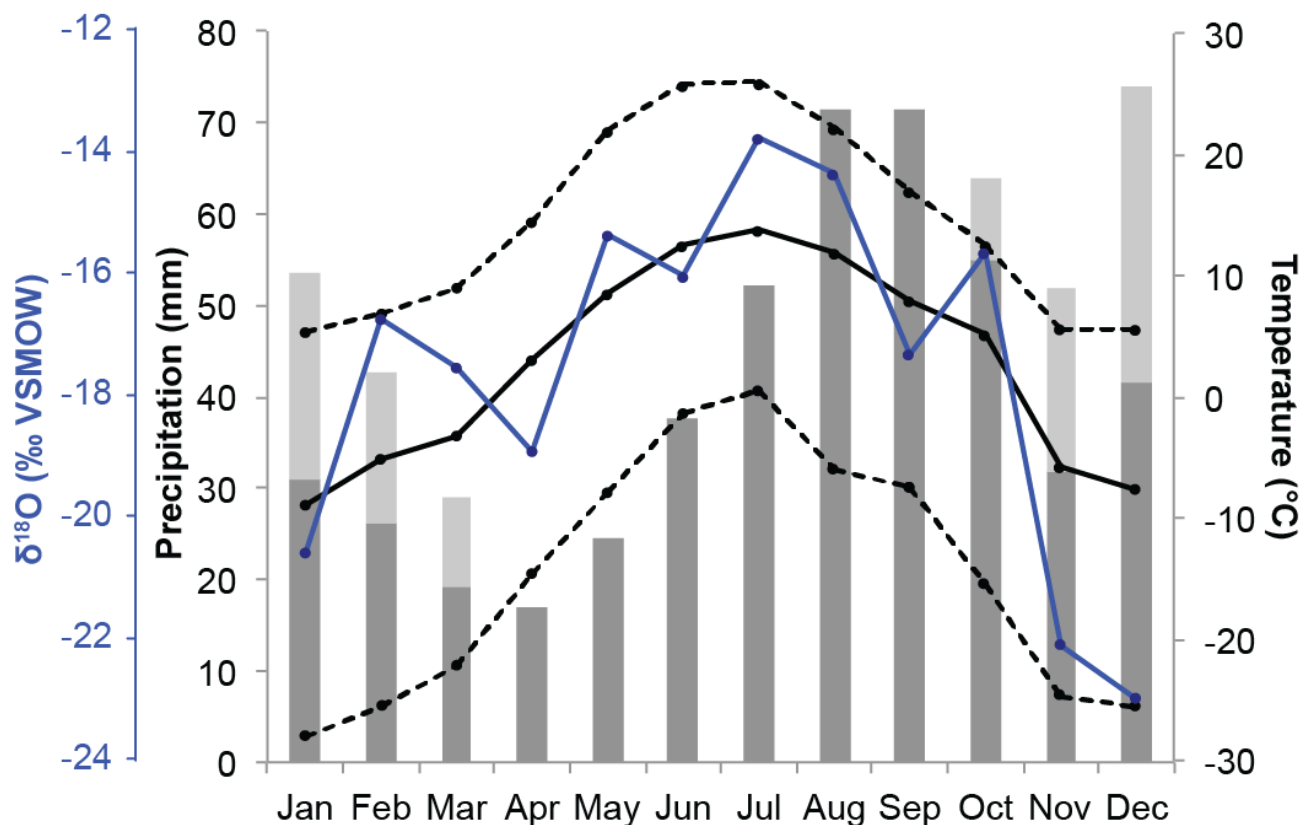
969

970

971

972

973

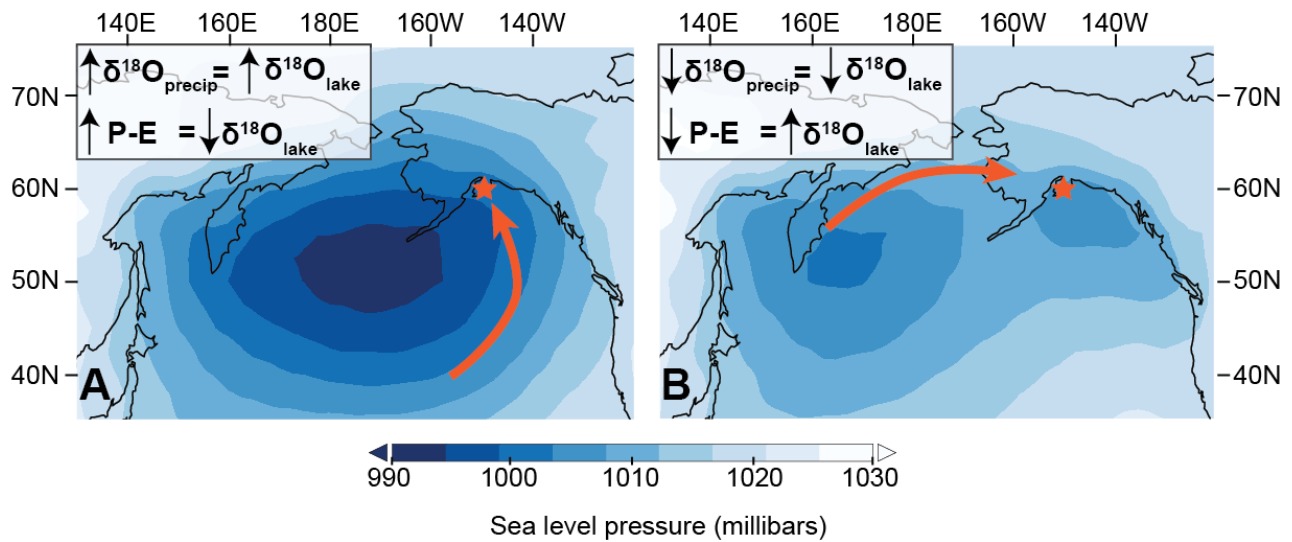


974

975 **Figure 2:** (A) Meteorological data from the Kenai Moose Pens SNOTEL site in Soldotna, AK
 976 (<https://www.wcc.nrcs.usda.gov/index.html>), showing the average (solid line), high (upper dashed
 977 line), and low (lower dashed line) temperatures alongside average monthly precipitation (dark gray
 978 bars) and average monthly snow-water equivalent (SWE; light gray bars) for 1995 – 2019 CE. (B)
 979 The same precipitation and SWE data as in (A) shown alongside monthly $\delta^{18}\text{O}_{\text{precipitation}}$ from the
 980 Tideview Station in Anchorage for 2005 – 2017 CE (Bailey et al., 2019).

981

982



983

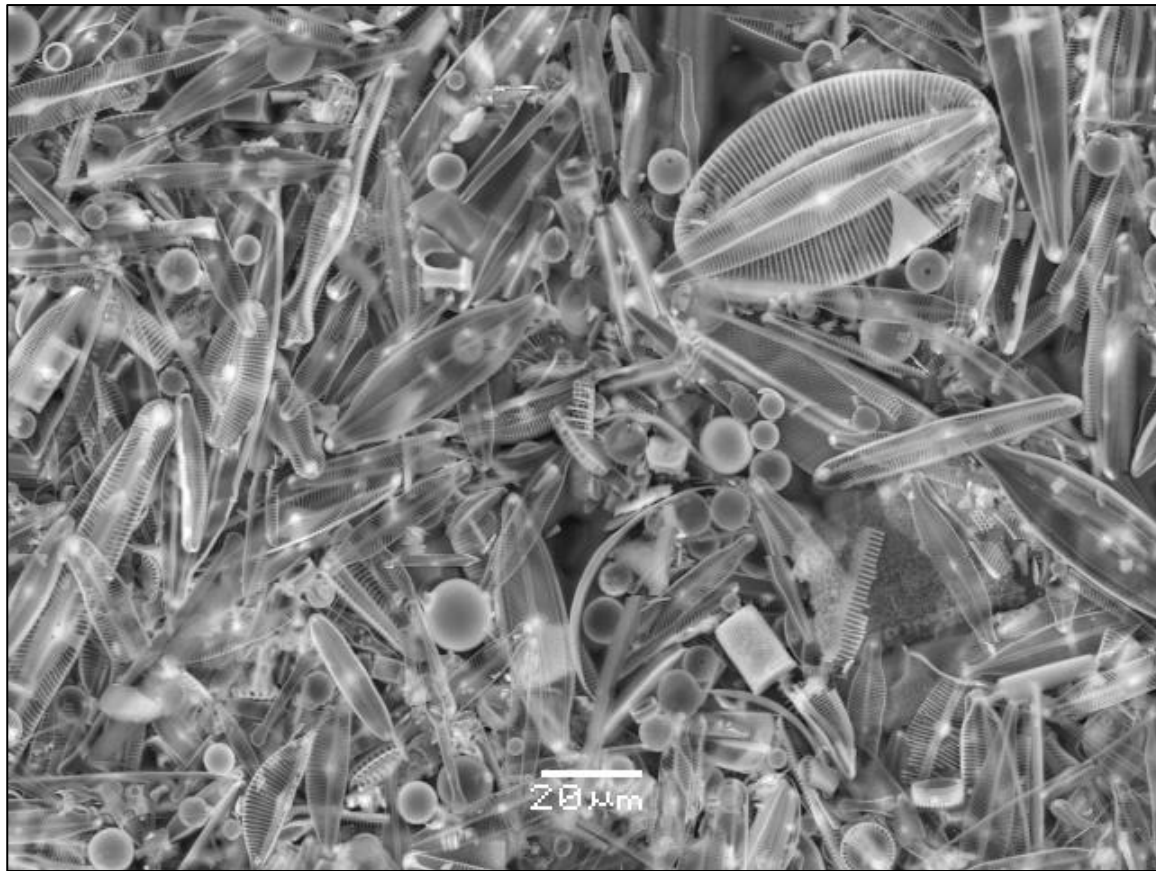
984 **Figure 3:** Study region showing the relation between Aleutian Low strength and associated effect on
 985 precipitation/evaporation balance P-E, $\delta^{18}\text{O}_{\text{precipitation}}$ and $\delta^{18}\text{O}_{\text{lake}}$. Average sea level pressure for the
 986 ten strongest (A) and weakest (B) Aleutian Low years from 1951 – 2000 CE is shown, as indicated
 987 by the North Pacific Index (after Rodionov et al., 2007). Orange arrows represent the tendency of
 988 storm tracks in the ten strongest and weakest Aleutian Low years for the same ten strongest (A) and
 989 weakest (B) AL years (Rodionov et al., 2007). The competing influences of $\delta^{18}\text{O}_{\text{precip}}$ and P-E on
 990 $\delta^{18}\text{O}_{\text{lake}}$ at Sunken Island Lake are described in white text.

991

992

993

994



995

996 **Figure 4:** Scanning electron microprobe image of purified biogenic silica from 383.5 cm below lake
997 floor (5662 ± 86 a cal BP) at Sunken Island Lake. Spherical features are chrysophyte cysts.

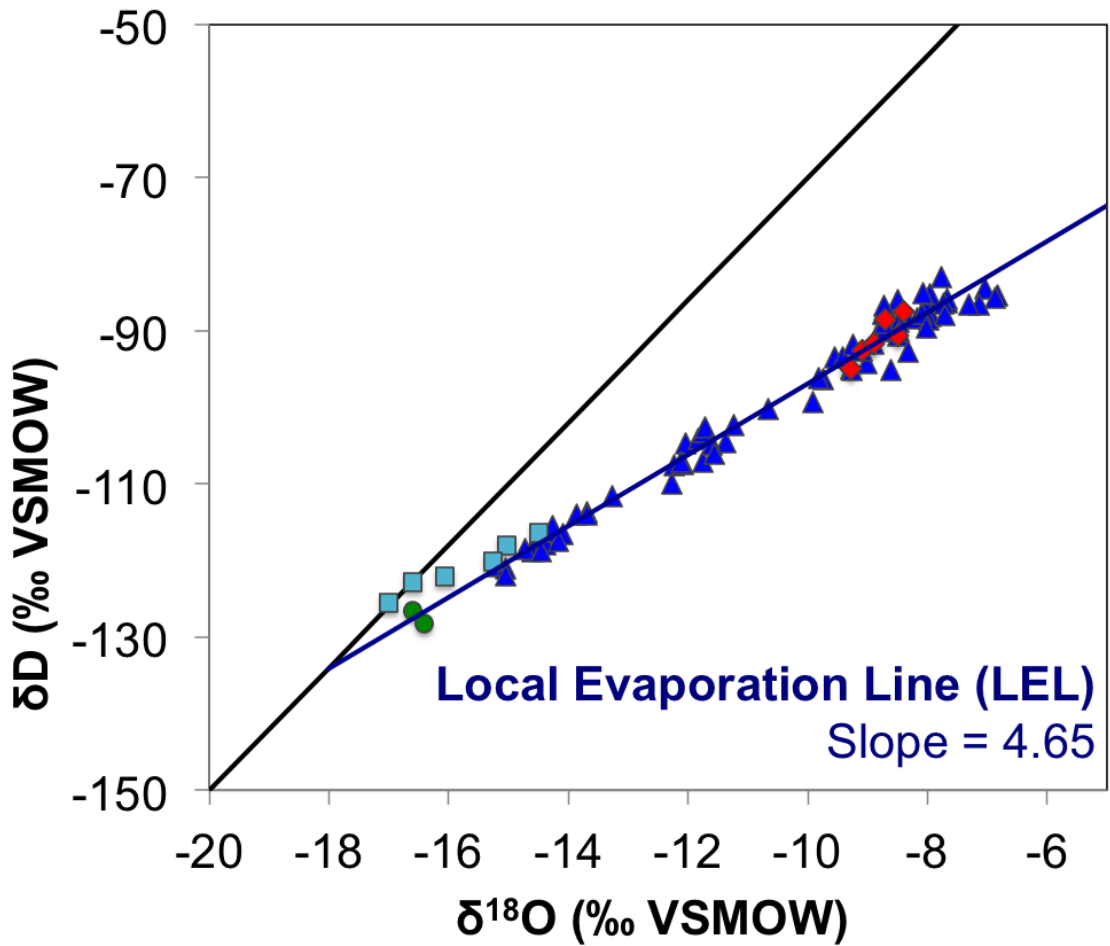
998

999

1000

1001

1002

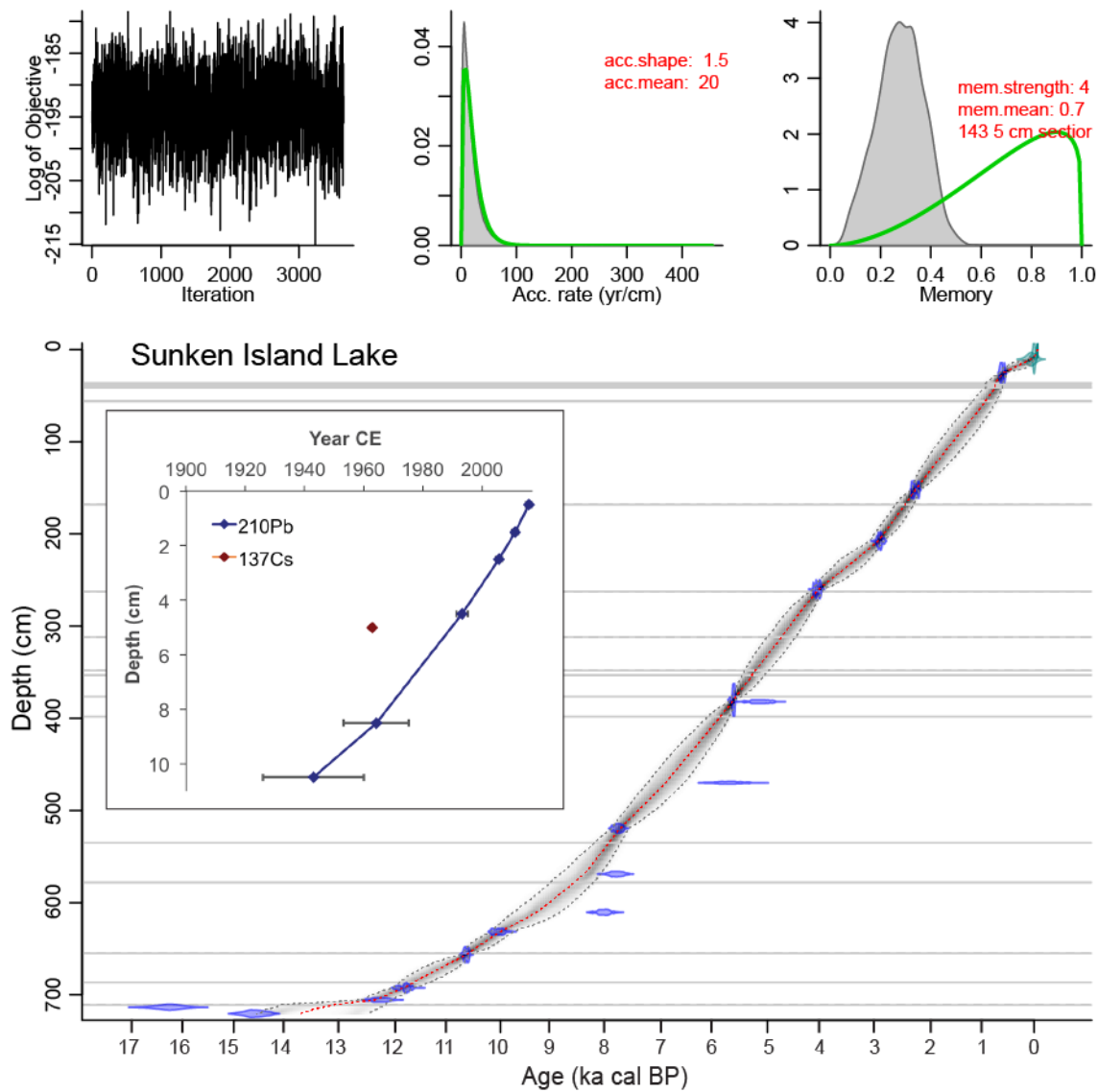


▲ Lakes ◆ Sunken Island ■ Rivers ● Groundwater

1003

1004 **Figure 5:** Water isotope data for lakes, rivers, and groundwater in the Kenai lowlands collected
 1005 between 2017 and 2018 CE, with Local Evaporation Line (LEL; dark blue) plotted for all lake water
 1006 samples, and the Global Meteoric Water Line (GMWL; black). Data are in Table S2.

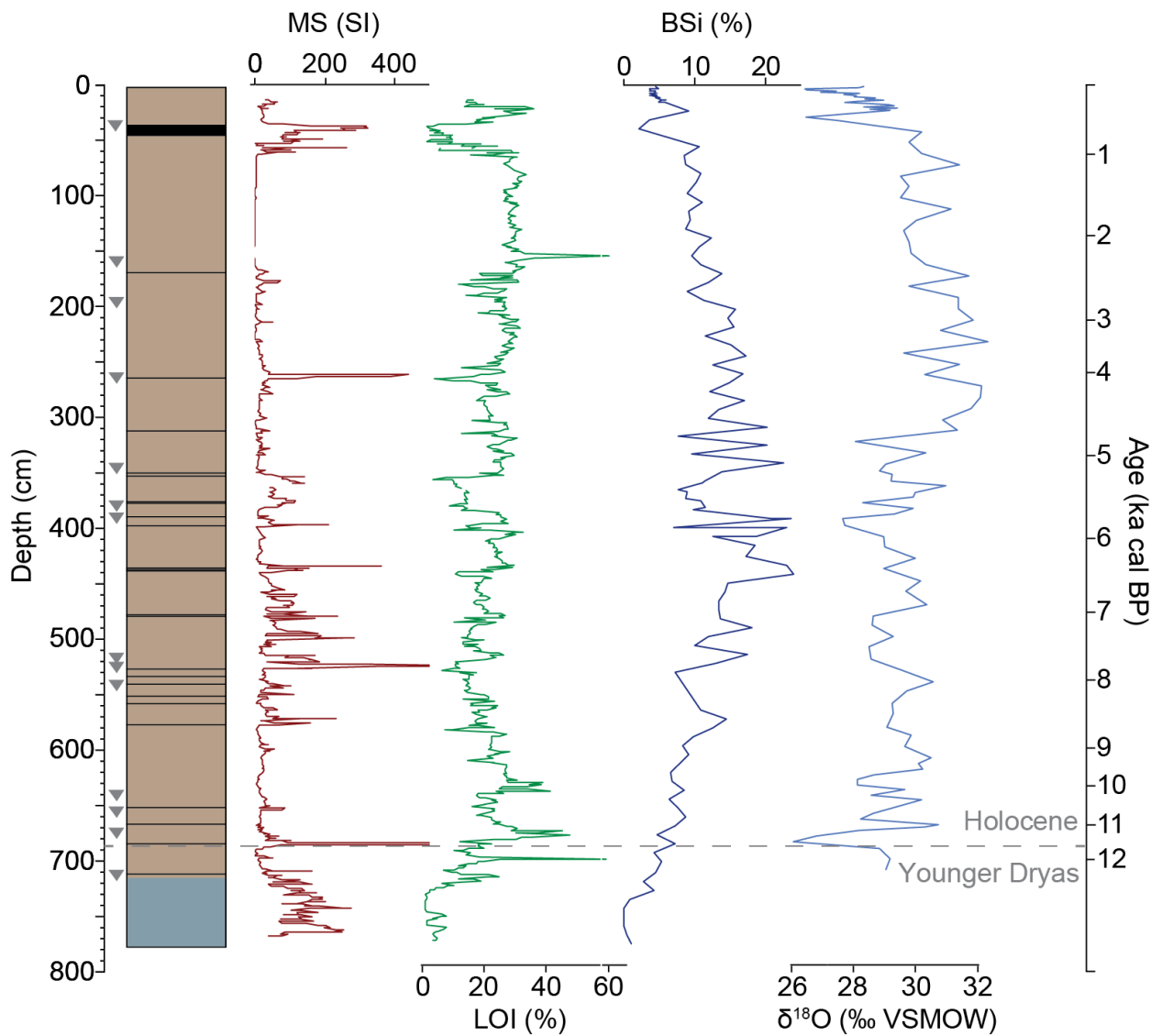
1007



1008

1009 **Figure 6:** Age-depth model for Sunken Island Lake master core (SIL-MC), created using Bacon
 1010 (v2.2; Blaauw and Christen, 2011). Gray horizontal lines mark visible tephra deposits that were
 1011 assumed to have been deposited instantaneously. The depths and basal ages of these layers are in
 1012 Table S3. Inset shows ^{210}Pb and ^{137}Cs profiles of the near surface (data in Fig. S3 and Table S1).

1013

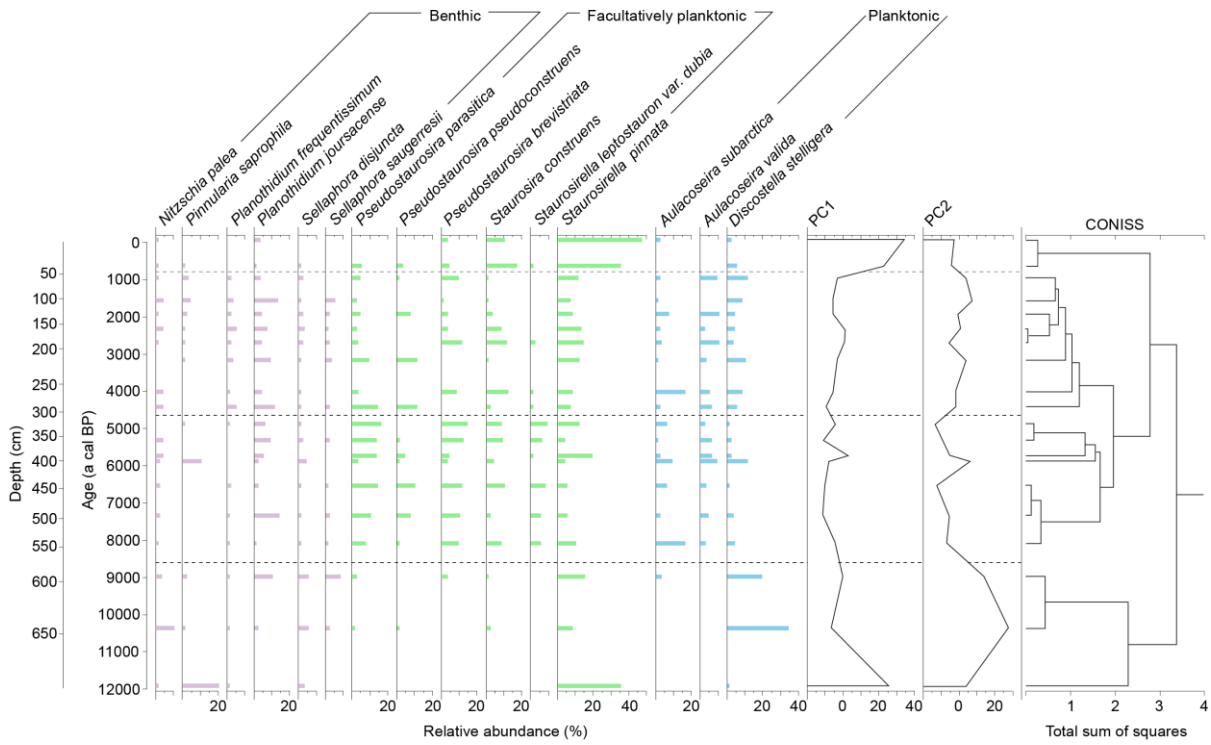


1014

1015 **Figure 7:** Stratigraphy of Sunken Island Lake master core (SIL-MC) with magnetic susceptibility
 1016 (MS), loss on ignition (LOI), and biogenic silica (BSi). The ambient lake sediment (brown) is
 1017 dominantly gyttja; macroscopically visible tephras are shown as black bands, and basal gray clay and
 1018 sand is indicated in blue. Age scale is based on the age model shown in Fig. 6. Gray triangles show
 1019 calibrated ^{14}C ages prior to 13 ka cal BP shown in Table 1. Dashed gray line shows boundary
 1020 between the Younger Dryas and the Holocene (11.7 ka cal BP).

1021

1022

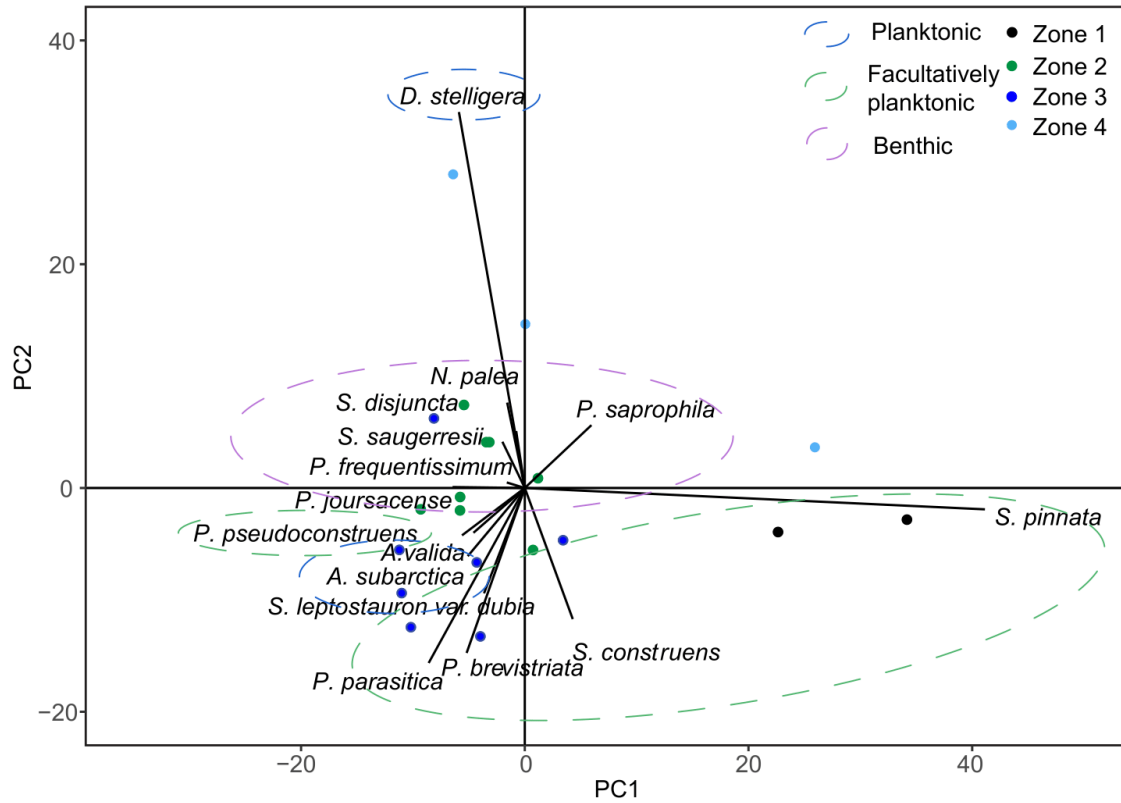


1023

1024 **Figure 8:** Relative abundance of 15 dominant diatom taxa in Sunken Island Lake core SIL-MC.

1025 CONISS-designated zones are indicated by dashed lines. Analyzed and plotted using Tilia (v.2.1.1)

1026 (Grimm, 2015).



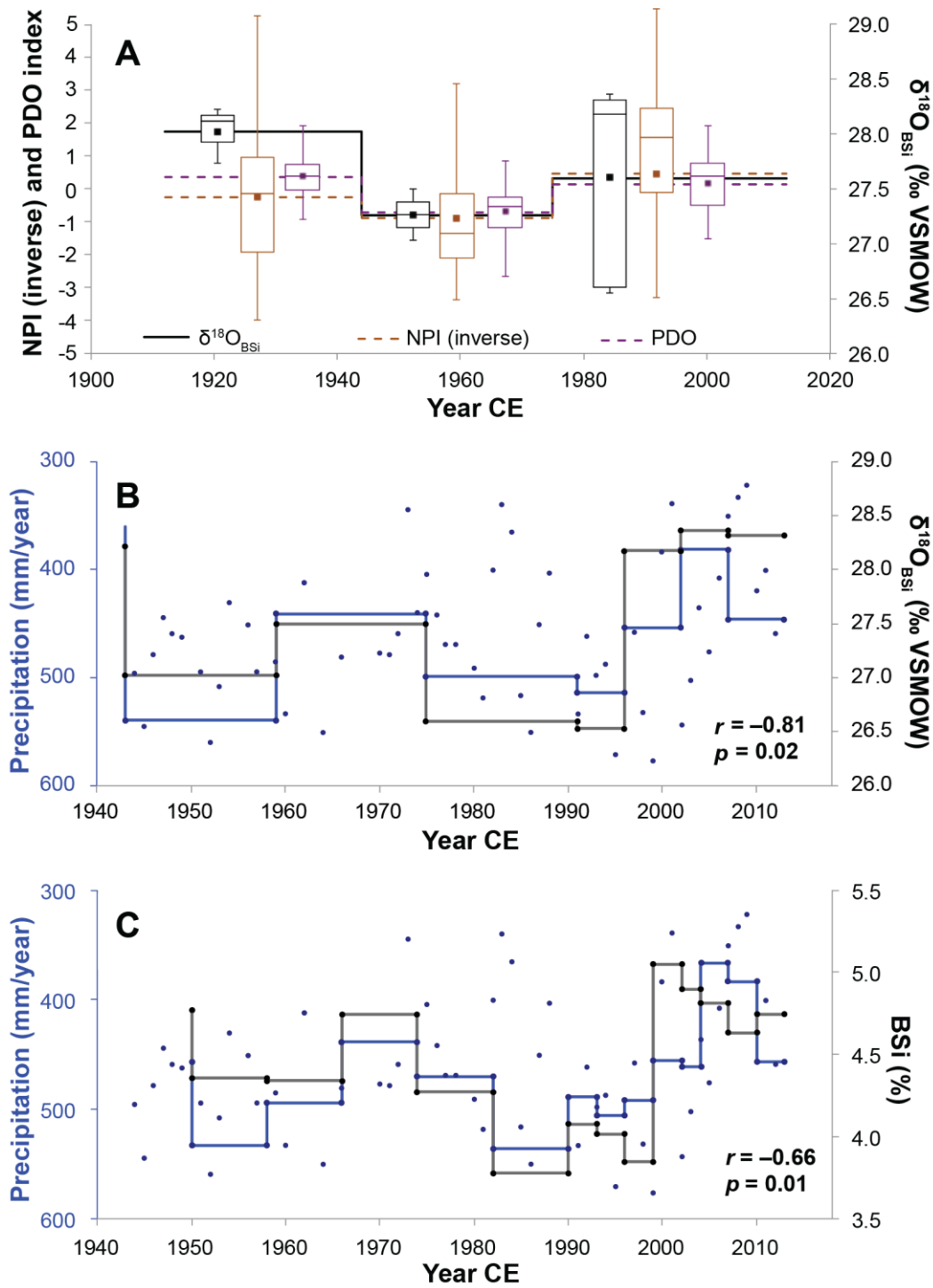
1027

1028 **Figure 9:** Principal component analysis (PCA) of diatom assemblages by sample. CONISS zone is
 1029 indicated by dot color, and habitat types (after Spaulding et al., 2019) are enclosed in dashed circles.
 1030 PC1 explains 41.9% of the variance in the dataset, and PC2 explains 23.5%.

1031

1032

1033



1034

1035 **Figure 10:** Comparison between instrumental climate data and proxy data from Sunken Island Lake
 1036 for the period of 1900-2015 CE. (A) $\delta^{18}\text{O}_{\text{BSi}}$ compared with the North Pacific Index (NPI) and the
 1037 Pacific Decadal Oscillation (PDO) index (data from <https://climatedataguide.ucar.edu> and
 1038 <http://research.jisao.washington.edu/pdo/> respectively). The three intervals represent known PDO

1039 regime shifts (Khapalova et al., 2018); index values are averaged over each interval. (B) $\delta^{18}\text{O}_{\text{BSi}}$ and
1040 (C) BSi compared with total annual precipitation (mm/year) at Kenai airport (data from
1041 http://climate.gi.alaska.edu/acis_data). Data were binned in panels B and C using the Analyseries
1042 software package (Paillard et al., 1996). Binned data are shown by thick lines, while dots are
1043 individual data points. The highest and lowest precipitation data points are not shown in the range of
1044 data displayed in this figure. Note change in x-axis scale following panel A, and inverse y-axes in
1045 panels B and C.

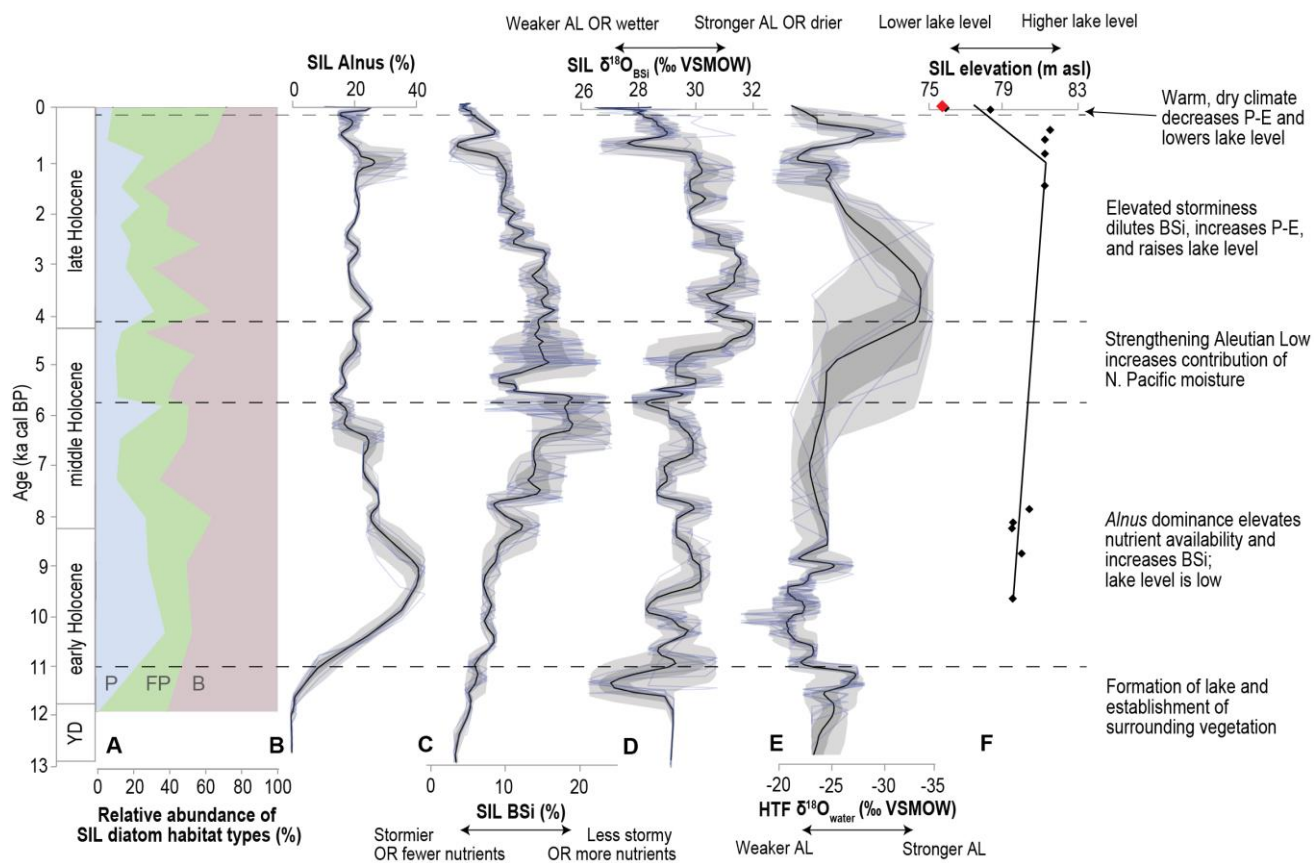
1046

1047

1048

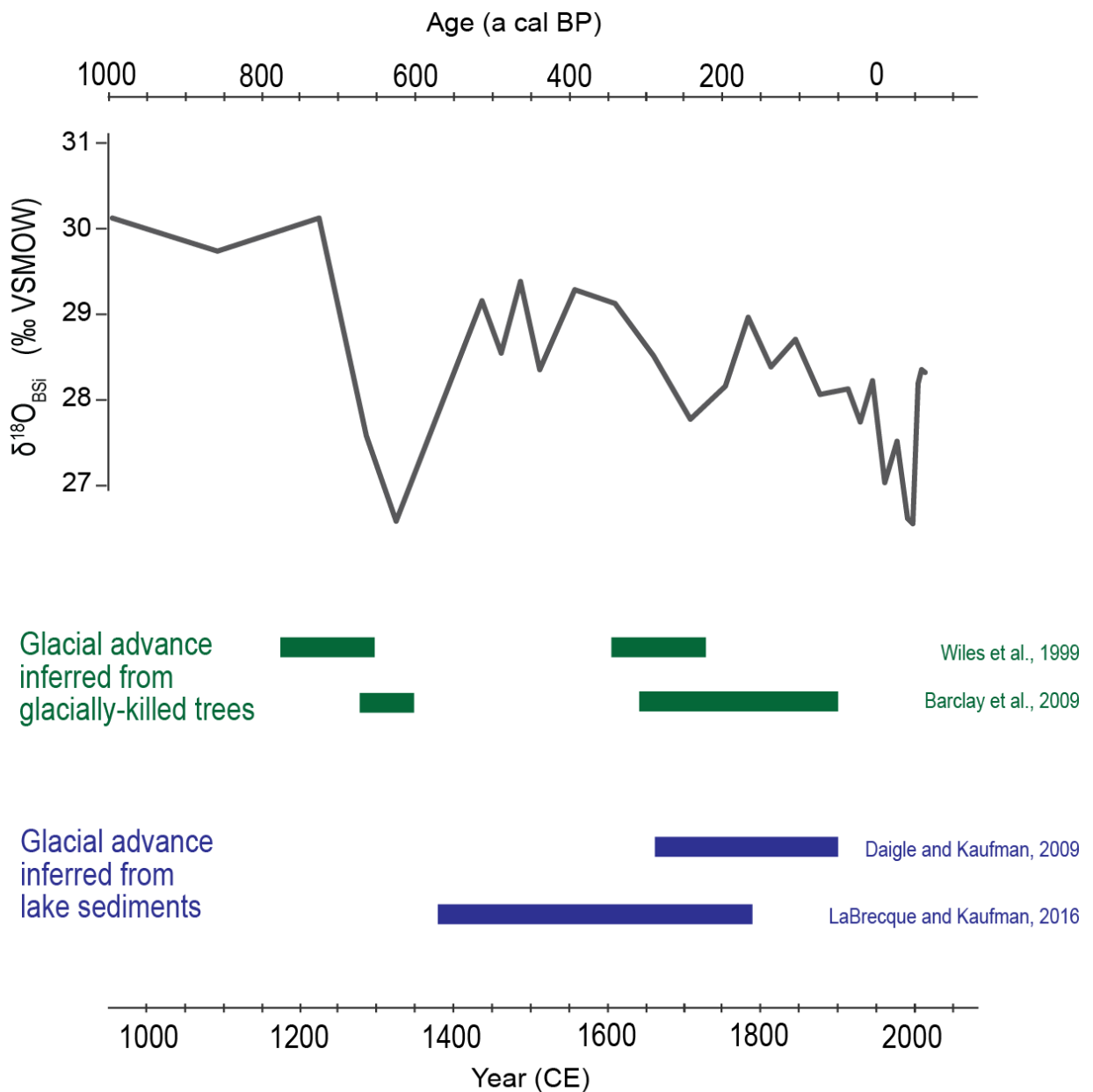
1049

1050



1051

1052 **Figure 11:** Sunken Island Lake (A) proportions of diatom assemblage habitat types: planktonic (P;
 1053 blue), facultatively planktonic (FP; green), and benthic (B; purple), (B) relative abundance of *Alnus*
 1054 (Anderson et al., 2019), (C) biogenic silica (BSi), and (D) $\delta^{18}\text{O}_{\text{BSi}}$ shown alongside (E) $\delta^{18}\text{O}_{\text{water}}$
 1055 inferred from $\delta^{18}\text{O}_{\text{TOM}}$ at Horse Trail Fen (note flipped x-axis values) (Jones et al., 2019), and (F)
 1056 Sunken Island Lake level inferred from dated basal peats in satellite fens and ice shoved ramparts.
 1057 Black lines in panels B-E are the mean of the age-model ensembles, and dark and light gray shading
 1058 encompass 68% and 95% of the ensemble members, respectively; blue lines show 5 representative
 1059 members of the 101-member (SIL) and 55-member (HTF) ensemble. Data plotted using GeoChronR
 1060 (McKay et al., 2018). Black dots in panel F are radiocarbon ages from satellite fen basal peats and
 1061 ice-shoved ramparts, and red dot indicates current lake elevation (Table 1). Dashed horizontal lines
 1062 correspond to important paleoenvironmental events/features, as annotated on the far right.



1063

1064 **Figure 12:** The most recent millennium of the $\delta^{18}\text{O}_{\text{BSi}}$ from Sunken Island Lake shown alongside
 1065 inferred glacial advances from sites in the Kenai Mountains and the western Prince William Sound
 1066 region. Periods of glacial advance are shown as horizontal bars in green for tree-ring records from
 1067 glacially-killed trees (Barclay et al., 2009; Wiles et al., 1999) and in blue for sediment records from
 1068 Goat Lake (Daigle and Kaufman, 2009) and Emerald Lake (LaBrecque and Kaufman, 2016).

1069

Core/Site ID	Midpoint depth blf (cm)	¹⁴ C age (year BP)	Calibrated age (a cal BP)*	Dated material	Lab ID
SIL-04-01	25.1	593 ± 35	603 ± 65	<i>Alnus</i> fruit, bryophytes, plant fibers	DAMS009385
SIL-04-01	152.1	2195 ± 15	2249 ± 57	algal copropel	UCIAMS69752
SIL-04-01	207.7	2788 ± 28	2888 ± 69	wood fragments	DAMS009386
SIL-04-01	260.0	3720 ± 20	4042 ± 57	algal copropel, <i>Betula</i> fruit	UCIAMS69753
SIL17-1A	382.0	4490 ± 120	5137 ± 290	insect chitin, bryophytes, plant fragments	UCIAMS202769
SIL-04-01	380.0	4915 ± 20	5631 ± 34	plant fragments	UCIAMS34293
SIL-04-01	470.0	4970 ± 230	5718 ± 563	insect chitin, bryophytes	UCIAMS17428
SIL-04-01	519.5	6960 ± 60	7794 ± 114	wood and plant fragments	UCIAMS17579
SIL-04-01	569.0	7000 ± 130	7831 ± 248	insect chitin, bryophytes	UCIAMS17429
SIL-04-01	610.5	7230 ± 110	8060 ± 214	bryophytes, plant fragments	UCIAMS34294
SIL-04-01	631.5	8915 ± 50	10039 ± 172	cladocera chitin	UCIAMS17580
SIL-04-01	656.5	9415 ± 25	10644 ± 68	bryophytes, plant fragments, seeds	UCIAMS69745
SIL-04-01	692.5	10125 ± 20	11771 ± 83	bryophytes	UCIAMS34295
SIL-04-01	705.5	10390 ± 80	12261 ± 280	bryophytes	UCIAMS34296
SIL-04-01	713.5	13460 ± 160	16208 ± 499	insect chitin, aquatic plants	UCIAMS17581
SIL-04-01	720.5	12455 ± 35	14599 ± 335	algal copropel	UCIAMS69755
FEN1B	210.0	7215 ± 25	8014 ± 95	graminoid peat	UCIAMS134786
FEN3	150.0	7860 ± 20	8625 ± 56	graminoid peat	UCIAMS134359
FEN4	180.0	7320 ± 20	8110 ± 69	graminoid- <i>Drepanocladus</i> peat	UCIAMS134787
FEN5	210.0	6925 ± 20	7746 ± 67	<i>Sphagnum</i> -graminoid peat	UCIAMS134788
FEN6	210.0	8445 ± 20	9479 ± 38	graminoid peat	UCIAMS134789
Wiles	Ice-shoved rampart	350 ± 40	399 ± 91	twig fragments	Beta-245726
102108-1	Ice-shoved rampart	1570 ± 15	1471 ± 55	plant detritus, bark flakes	UCIAMS54689
102108-2	Ice-shoved rampart	630 ± 15	593 ± 50	plant detritus	UCIAMS54690
102108-3	Ice-shoved rampart	920 ± 15	860 ± 60	plant detritus	UCIAMS54691

1070 **Table 1:** ¹⁴C ages for Sunken Island Lake master core, basal ages for satellite fen cores, and ages for
1071 ice-shoved ramparts.

1072 *Calibrated age is the median of the calibrated age probability density function. Uncertainty is one
1073 half of the two sigma range