

Timing and causes of mid-Holocene mammoth extinction on St. Paul Island, Alaska

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Relict woolly mammoth (*Mammuthus primigenius*) populations survived on several small Beringian islands for thousands of years after mainland populations went extinct. Here we present multiproxy paleoenvironmental records to investigate the timing, causes, and consequences of mammoth disappearance from St. Paul Island, Alaska. Five independent indicators of extinction show that mammoths survived on St. Paul until $5,600 \pm 100$ y ago. Vegetation composition remained stable during the extinction window, and there is no evidence of human presence on the island before 1787 CE, suggesting that these factors were not extinction drivers. Instead, the extinction coincided with declining freshwater resources and drier climates between 7,850 and 5,600 y ago, as inferred from sedimentary magnetic susceptibility, oxygen isotopes, and diatom and cladoceran assemblages in a sediment core from a freshwater lake on the island, and stable nitrogen isotopes from mammoth remains. Contrary to other extinction models for the St. Paul mammoth population, this evidence indicates that this mammoth population died out because of the synergistic effects of shrinking island area and freshwater scarcity caused by rising sea levels and regional climate change. Degradation of water quality by intensified mammoth activity around the lake likely exacerbated the situation. The St. Paul mammoth demise is now one of the best-dated prehistoric extinctions, highlighting freshwater limitation as an overlooked extinction driver and underscoring the vulnerability of small island populations to environmental change, even in the absence of human influence.

mammoth | extinction | Holocene | St. Paul Island | ancient DNA

During the wave of late Quaternary extinctions, nearly two-thirds of megafaunal genera disappeared worldwide (1, 2). Proposed extinction drivers include environmental change, human impacts, or some combination, with the relative contributions of these varying by species and region (2). Woolly mammoths, an iconic Ice Age species, vanished from mainland Asia and North America between 14,000 and 13,200 y ago, with some mainland populations perhaps persisting until approximately 10,500 y ago (1, 3, 4); however, relict populations survived on two newly formed Beringian islands into the middle Holocene (4–7) (Fig. 1A).

Wrangel Island is relatively large (7,600 km²), and the late survival of mammoths until 4,020 y ago (5) is consistent with a retreat to refugia in northeastern Siberia as temperatures rose (6). Conversely, the remarkable persistence of mammoths on the much smaller St. Paul Island (110 km²) until the mid-Holocene suggests that isolated megafaunal populations can survive thousands of years after habitat fragmentation. Because there is no evidence of humans on St. Paul Island before the arrival of Russian whalers in 1787 CE (7), the island affords a unique opportunity to study the

processes governing the persistence and extinction of small megafaunal populations in the absence of humans.

St. Paul Island is a remnant of the Bering Land Bridge that became isolated between 14,700 and 13,500 y ago due to sea level rise during the last deglaciation. The island rapidly shrank in area until 9,000 y ago, and then slowly shrank until 6,000 y ago (Fig. 1B). Today, St. Paul is highly isolated (>450 km from Alaska and Aleutians) and is characterized by low relief (maximum elevation 203 m above sea level), a few freshwater lakes, no springs or streams, no permafrost, and moderately productive moss-herbaceous tundra vegetation (8, 9). Apart from a small

Significance

St. Paul Island, Alaska, is famous for its late-surviving population of woolly mammoth. The puzzle of mid-Holocene extinction is solved via multiple independent paleoenvironmental proxies that tightly constrain the timing of extinction to $5,600 \pm 100$ y ago and strongly point to the effects of sea-level rise and drier climates on freshwater scarcity as the primary extinction driver. Likely ecosystem effects of the mega-herbivore extinction include reduced rates of watershed erosion by elimination of crowding around water holes and a vegetation shift toward increased abundances of herbaceous taxa. Freshwater availability may be an underappreciated driver of island extinction. This study reinforces 21st-century concerns about the vulnerability of island populations, including humans, to future warming, freshwater availability, and sea level rise.

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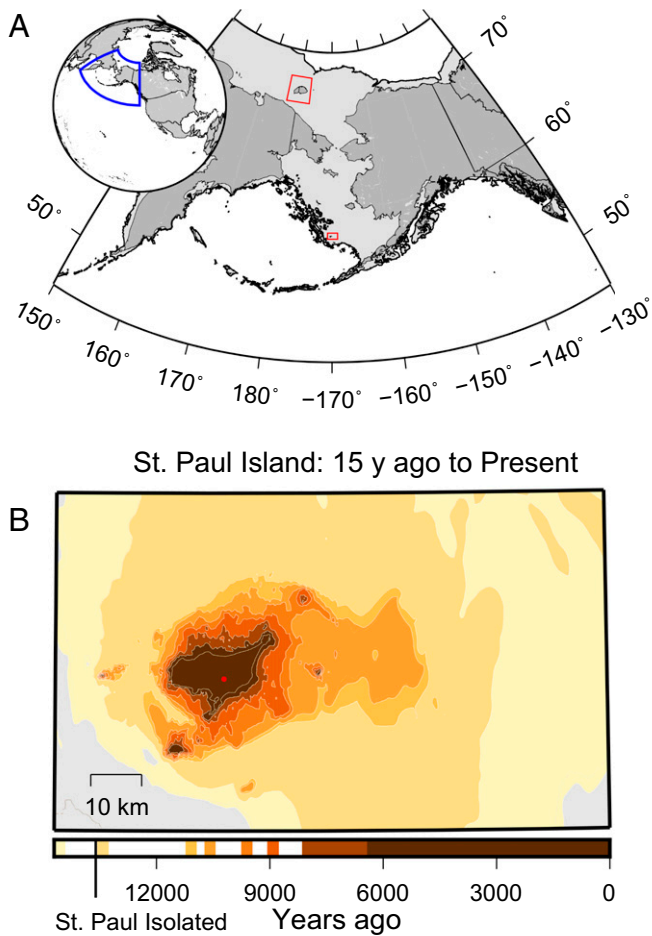


Fig. 1. Location of St. Paul Island, Alaska and areal losses due to rising sea level. (A) Map of current continents (dark gray) and the past position of the Bering Land Bridge (light gray) with red boxes indicating Wrangel Island (Upper) and St. Paul Island (Lower). (B) The shrinking area of St. Paul Island from 15,000 y ago to the present due to rising sea level during the late Pleistocene through the middle Holocene. The red dot indicates the location of Lake Hill.

population of reindeer (*Rangifer tarandus*) introduced to the island in 1911 (10), the largest terrestrial mammal on St. Paul today is the Arctic fox (*Alopex lagopus*). Previous radiocarbon dates on five mammoth remains from St. Paul indicated a last appearance date (LAD) of 6,480 y ago (11); however, LADs rarely represent the last surviving individual, creating uncertainty in the actual extinction timing. Previously proposed hypotheses for the cause of extinction of the St. Paul population include changes in vegetation (8) or increasing snowpack (12), but to date there is little direct evidence to support any proposed hypothesis.

In this study, we seek to better understand the timing, causes, and consequences of the mid-Holocene extinction of this late-surviving mammoth population. To achieve this goal, we collected in 2013 a series of sediment cores from Lake Hill, a small freshwater lake near the center of St. Paul (Fig. 1C and *SI Appendix, Fig. S1*). Lake Hill was cored previously in the 1960s to test hypotheses about whether the coastal Bering Land Bridge was a glacial refugium for *Picea* and other plant species (9). From our new sediment cores, we analyzed four independent proxies known to correlate with megafaunal presence (13–16): sedimentary ancient DNA (sedaDNA) and three coprophilous fungal spore types—*Sporormiella*, *Sordaria*, and *Podospora* (Fig. 2 and *SI Appendix, SI Text*) (16). We also reconstructed past environments using multiple

proxies from the cores, including cladocerans, diatoms, pollen, plant macroremains, and stable isotopes, and dated the cores using radiocarbon dates from terrestrial plant macroremains and tephrochronology (*SI Appendix, SI Text*). In addition, we collected and radiocarbon-dated 14 newly identified St. Paul mammoth remains using ultrafiltered collagen extracts and measured their stable isotopic composition ($\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values) (*SI Appendix, SI Text and Table S1*).

Results and Discussion

Timing of Extinction. The five independent indicators of extinction timing are in close agreement and together indicate that the St. Paul mammoths went extinct $5,600 \pm 100$ y ago (Fig. 2), which is 900 y later than the previous LAD value for St. Paul mammoths (11) and 1,580 y before the final extinction of mammoths on Wrangel Island (4, 7). The youngest of the newly dated mammoth remains from St. Paul provided a median calibrated radiocarbon date of 5,530 y ago (95% confidence interval, 5,585–5,330 y ago),

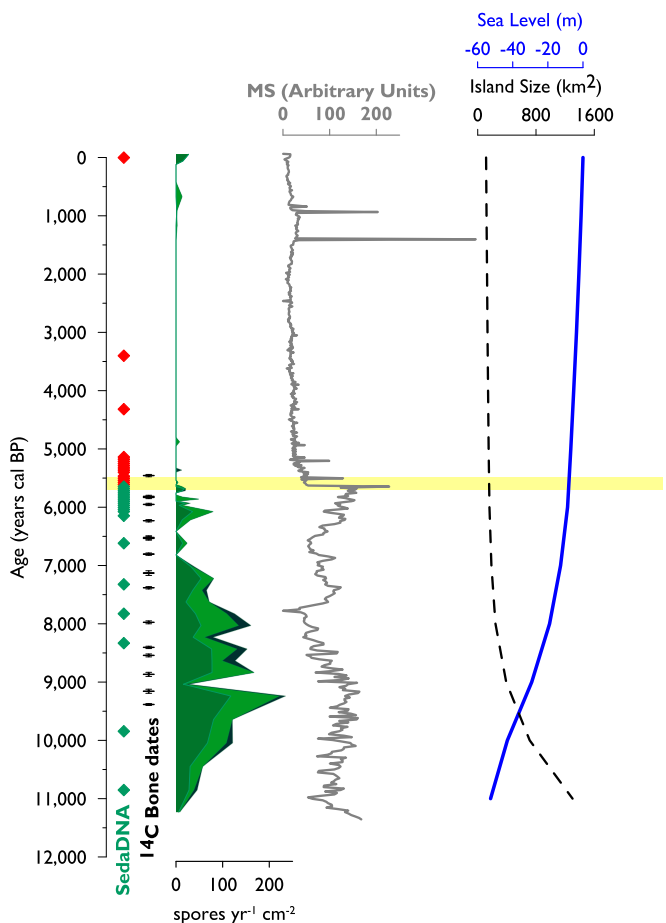


Fig. 2. Timing of woolly mammoth extinction on St. Paul. Five indicators are plotted against time (calendar years before 1950 CE): sedimentary ancient DNA (sedaDNA; green and red circles indicate inferred presence and absence, respectively), direct AMS radiocarbon dates on mammoth remains (black crosses), and spore accumulation rates for the coprophilous fungi *Sporormiella* (medium green, bottom of stack), *Sordaria* (light green, middle), and *Podospora* (dark green, top) (spores yr⁻¹ cm⁻²). Magnetic susceptibility, in arbitrary units (AU) of Lake Hill sediments, sea level (m), and St. Paul Island area (km²) are shown for comparison. Note the truncation by plot of highest peak of magnetic susceptibility at 1,500 AU. The yellow band indicates inferred timing of extinction at $5,600 \pm 100$ y ago. Magnetic susceptibility is a measure of sediment characteristics; its rapid decline indicates a rapid reduction in watershed erosion at the same time as the extinction.

which overlaps the extinction timing inferred from sedaDNA and fungal spores of $5,650 \pm 80$ y ago (SI Appendix, Fig. S3). Mammoth sedaDNA is present in all tested sediment samples between $10,850 \pm 150$ y ago (the oldest sample tested) and $5,650 \pm 80$ y ago (Fig. 2 and SI Appendix, Fig. S5), and absent in samples younger than $5,610 \pm 80$ y ago (Fig. 2). *Sporormiella* and *Sordaria* terminate at $5,680 \pm 80$ y ago and $5,650 \pm 80$ y ago, respectively, the latter exactly coinciding with the last appearance of mammoth sedaDNA. Declining *Sporormiella* and *Sordaria* abundances from 9,000–5,650 y ago (Fig. 2) might indicate declining mammoth population size, but coprophilous spore abundances may be influenced by sedimentary processes (14). *Sporormiella* abundances rise again at 1890 ± 50 CE, presumably reflecting the introduction of caribou (*Rangifer tarandus*) in 1911 CE (10). *Podospira* spores are less abundant than *Sporormiella* and *Sordaria* and disappear earlier, $7,020 \pm 170$ y ago. Overall, the close agreement among these five independent proxies makes $5,600 \pm 100$ y ago one of the most robust and precise estimates of timing ever recorded for a prehistoric species extinction.

Drivers of Extinction. The disappearance of mammoths from St. Paul Island was accompanied by pronounced lake shallowing and increased water turbidity between 7,850 and 5,600 y ago, indicated by significant shifts from pelagic to littoral cladocerans, decreases in planktonic diatoms, and increases in tycho planktonic diatoms

(Fig. 3 and SI Appendix, SI Text). A concurrent rise in conductivity is indicated by an increased abundance of diatoms and cladocerans that are tolerant of high electrolyte concentrations, notably the cladoceran *Alona circumfibrata* (Fig. 3), which is resilient to fluctuating salinity and can be an indicator of enhanced salinity in lakes (17) (SI Appendix, SI Text).

Bayesian change-point analysis indicates that the most probable change-point for *A. circumfibrata* occurs at 6,430 y ago, with its rapid rise, and a secondary change point occurs at 5,430, when it declines. Lake shallowing and enhanced evaporative loss is also supported by a trend of increasing stable oxygen isotope values (expressed as $\delta^{18}\text{O}$ values) in Lake Hill between 8,500 and 5,300 y ago (Fig. 3), with $\delta^{18}\text{O}$ values significantly higher after 5,600 y ago than before ($P = 9.4 \times 10^{-15}$, two-sample t test). Stable nitrogen isotope values (expressed as $\delta^{15}\text{N}$ values) from mammoth remains seem to increase between 9,500 and 5,500 y ago (Fig. 3), consistent with decreasing precipitation and soil moisture (18, 19). However, the slope of a linear regression fitted to the $\delta^{15}\text{N}$ time series does not significantly differ from 0 at a two-sigma threshold ($P = 0.07$) because of the single enriched value at 9,500 y ago, and thus the $\delta^{15}\text{N}$ data are suggestive rather than definitive. Finally, an abrupt decrease in magnetic susceptibility at $5,650 \pm 80$ y ago coincides exactly with the *Sordaria* and sedaDNA disappearances and indicates a rapid shift in depositional regime to one characterized by

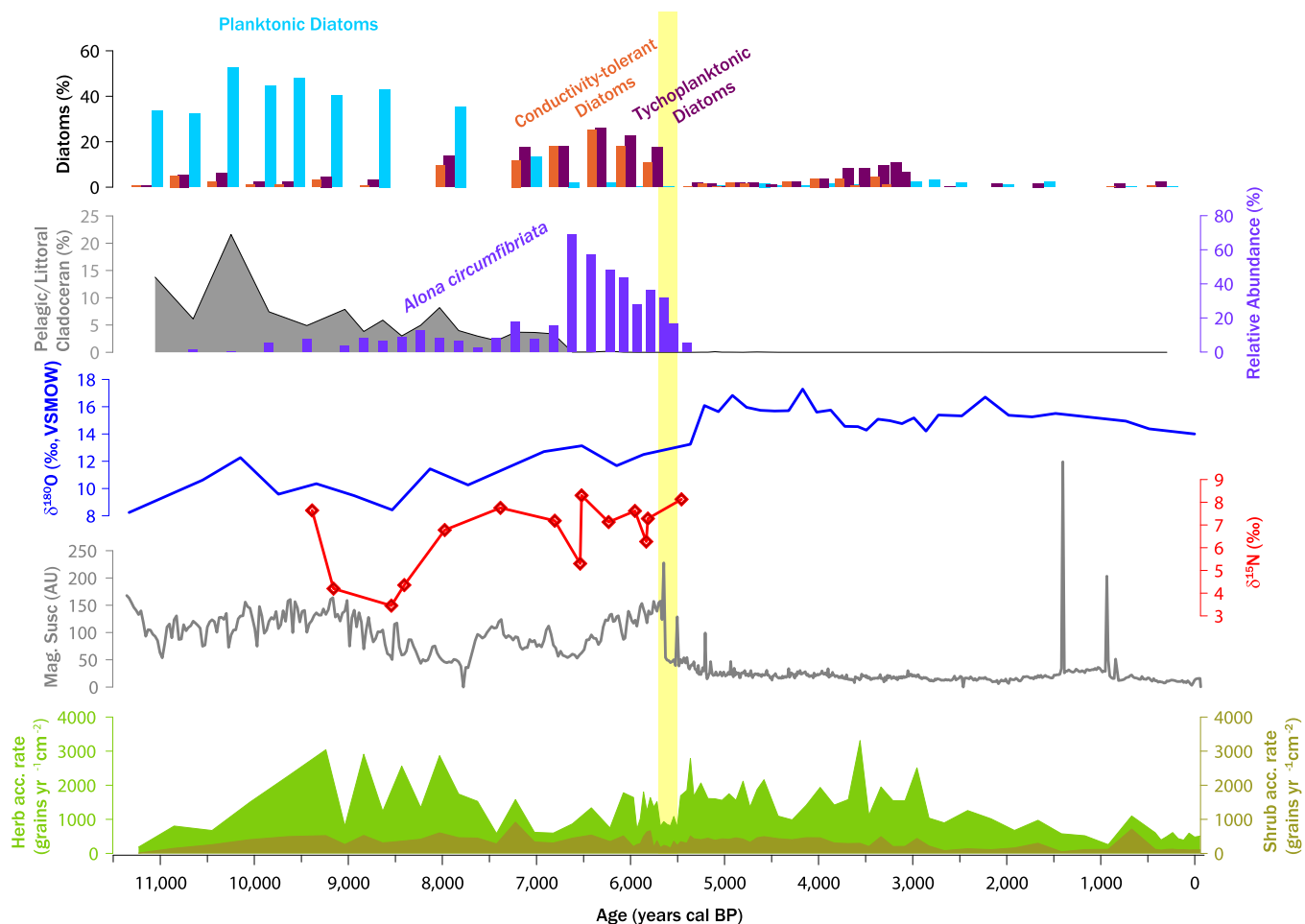


Fig. 3. Paleoenvironmental proxies at Lake Hill, plotted against time. Shown are relative abundances of planktonic (blue bars), tycho planktonic (maroon bars), and high-conductivity-tolerant diatoms (orange bars); ratio of pelagic and littoral cladocerans (gray curve); relative abundance (%) of cladoceran *Alona circumfibrata*, which is tolerant of high lake water conductivity (purple bars); $\delta^{18}\text{O}$ values from aquatic invertebrate chitin (blue curve); $\delta^{15}\text{N}$ values from mammoth collagen (red curve); sediment magnetic susceptibility (in AU); and accumulation rates for herb and shrub pollen types (light green and tan curves, respectively). The estimated timing of extinction (yellow vertical bar) and the magnetic susceptibility curve are included to help establish the stratigraphic correlation between Figs. 2 and 3.

decreased inputs of terrestrial sediments, likely due to reduced rates of watershed erosion after the extinction. Taken together, these proxies strongly indicate lake shallowing, enhanced evaporation, increased turbidity, and increased solute concentrations at Lake Hill during the mid-Holocene extinction window of mammoth on St. Paul Island.

Freshwater availability on oceanic islands is an underappreciated, but important, driver of vertebrate mortality. On Mauritius, a mass mortality event 4,150 y ago coincided with a severe drought that concentrated animals near increasingly toxic and saline lakes (20). Modern elephants (*Loxodonta* and *Elephas*), the closest living equivalents to mammoths, consume 70–200 L/day (21), and their survival depends on water availability (22). Elephants prefer clear water and will dig “wells” near sediment-choked water sources to filter and clarify drinking water (23). St. Paul mammoths may have needed even more water than modern elephants for evaporative cooling during the Holocene interglacial (24), because many of the morphological and physiological adaptations for *M. primigenius* were for retaining heat (25, 26). As island size decreased, freshwater resources on St. Paul became more limited. Surface availability of fresh water is presently—and was probably at the time of extinction—restricted to shallow groundwater perched on a seawater base and several freshwater coastal lagoons and three closed-basin crater lakes (27). Of these, the freshwater lake at Lake Hill is currently the deepest (1.3 m) (9) and thus was the primary (or only) freshwater source for mammoths during dry periods (20). Thus, freshwater availability on St. Paul is highly limited, and diminished freshwater availability is a plausible agent of extinction on St. Paul.

Several previously hypothesized extinction drivers for St. Paul can be discarded. Human colonization during the mid-Holocene is unlikely, given the extreme isolation of St. Paul in the Bering Sea and the lack of any archaeological record (7). Polar bear predation is unlikely, because all St. Paul polar bear remains postdate the mammoth extinction by >1,000 y (11). Volcanic activity (11, 28) is unlikely, given the absence of visible tephra in Lake Hill sediments during the extinction window. Extinction due to increasing winter snowpack (12, 29) is inconsistent with the increase in $\delta^{18}\text{O}$ values at Lake Hill during the Holocene (Fig. 3). Extinction due to vegetation shifts (8) is unlikely, given the stable vegetation composition on St. Paul during the Holocene (Fig. 3), although declining pollen accumulation rates in the early Holocene might indicate declining tundra productivity.

Reduction in island size per se (8) is not a direct driver, because the modern size of the island (110 km²) was reached by 9,000 y ago, more than 3,000 y before extinction. However, decreasing island size may have indirectly caused extinction, by reducing mammoth population sizes and reducing the number and volume of freshwater resources. If so, the St. Paul mammoth extinction may be a natural example of extinction debt (30), and suggests that the delay between habitat loss and final extinction can last for millennia.

Causes of Freshwater Scarcity. The causes of reduced freshwater availability on St. Paul are less certain, with likely factors including sea level rise, climate change, and resource overuse. Rising sea levels gradually decreased land area and availability of freshwater resources (Fig. 1), but a freshwater crisis might have been triggered by mid-Holocene regional climate change and variability. The Lake Hill findings and $\delta^{15}\text{N}$ values from mammoth bones indicate increased aridity between 8,000 and 5,300 y ago on St. Paul (Fig. 3). Other lake records from mainland Alaska show decreased effective moisture and increased hydrological variability between 6,800 and 5,000 y ago (31–34), suggesting that water scarcity of St. Paul might have been caused by regionally enhanced hydrological variability during the mid-Holocene.

Mammoth activity around Lake Hill probably contributed to the degradation of freshwater quality. As ecosystem engineers,

elephants strongly modify their local environments, especially at water holes and other areas of heavy use (35). As plant cover is destroyed, environmental deterioration rapidly radiates outward (12). The abrupt decrease in magnetic susceptibility, shift from terrigenous to lacustrine sediments (Fig. 3), disappearance of pelagic cladocerans, near-disappearance of planktonic diatoms, and presence of tychoplanktonic diatoms all coincide with the time of mammoth extinction. Thus, mammoth activity likely denuded the lake margin, accelerated rates of erosion, and contributed to lake infilling and decreased water quality. The increase in magnetic susceptibility before extinction (Fig. 3) is consistent with intensified water use and rates of watershed erosion, perhaps as other water sources were depleted.

Conclusions and Implications. Multiple reinforcing lines of evidence indicate that reduced freshwater availability triggered the extinction of St. Paul mammoths at $5,600 \pm 100$ y ago. Sea level rise and declining island area set the stage for extinction (8) by reducing island carrying capacity, decreasing population size, and increasing mammoth dependency on diminishing freshwater resources, leaving mammoths vulnerable to environmental fluctuations. Regional climate variability and aridification then may have applied the coup de grâce, possibly exacerbated by resource degradation by the mammoths. Thus, the St. Paul history demonstrates that in the absence of human pressures, small megafaunal populations can persist for thousands of years, while highlighting the susceptibility of island populations to environmental fluctuations, including freshwater availability, particularly during periods of rising sea level. Island biogeographic theory predicts that rates of extinction should be elevated on small islands with limited resources; these findings support that theory while pointing to freshwater scarcity as an underappreciated driver. Our results suggest that as sea levels rise and hydroclimatic variability increases over the coming centuries due to anthropogenic warming (36), freshwater scarcity may become an increasingly critical limiting resource for island vertebrate populations.

Methods

Specific details of all methods are provided in *SI Appendix, SI Text*. Island size through time was reconstructed using a GIS bathymetric map of the St. Paul Island area and the sea level rise curve from the Bering Strait (37). Three overlapping cores were collected at the primary Lake Hill core in 2013 CE (57.17809N, 170.24828W) and used to construct a composite core using clearly identifiable marker layers, such as visible tephra and lithologic transitions, in the overlapping sections (*SI Appendix, Fig. S2*). Samples for all paleoecological and paleoenvironmental proxies were taken at National Lacustrine Core Facility (LacCore) mostly at identical depths, to eliminate problems with proxy correlation. A Bayesian age model for the upper 740 cm of the composite core (*SI Appendix, Fig. S3*) was constructed using OxCal from accelerator mass spectrometry (AMS) radiocarbon (¹⁴C) dating of six terrestrial macrobotanical remains and a tephra layer representing the Aniakhak CFE II tephra, with a known age of $3,595 \pm 4$ cal yr BP (*SI Appendix, Table S3*). Tephra extraction was done following standard methods (38). Radiocarbon dating and isotopic analyses of mammoth teeth were derived from collagen extracted and purified following the modified Longin method with ultrafiltration (39). All AMS ¹⁴C results were corrected for isotopic fractionation according to standard conventions (40). Conventional ¹⁴C ages were calibrated with OxCal 4.2.3 (41) using the IntCal13 curve (42).

SedaDNA from 38 samples throughout the upper 740 cm of the core was extracted, sequenced, and bioinformatically processed at the Paleogenomics Laboratory, University of California Santa Cruz (*SI Appendix, Fig. S5 and Table S5*). Sequencing data are available from the National Center for Biotechnology Information's Short Read Archive (BioProject accession no. PRJNA320875). Extraction and processing of spores and pollen followed a modified version of the LacCore protocol (43). Plant macrofossil samples were taken every 10 cm, but shifted to finer resolution (sampling every 2–5 cm) for the lower portions of the core. All macrofossil samples were disaggregated in a glass Petri dish with distilled water and were manually collected. Samples were not sieved.

Cladocera, diatom, and stable isotopic analyses were all conducted at the Alaska Stable Isotope Facility, University of Alaska Fairbanks. Preparation of

diatom samples was done following a modified previously published protocol (44). Cladocera were processed with an adapted version of a method described previously (45). The $\delta^{18}\text{O}$ values of chironomid head capsules and other aquatic insect chitin were processed and analyzed following previously published protocols (46, 47). Bulk sediment samples were collected from the core at 16-cm intervals for isotopic and other geochemical analyses and processed following previously published protocols (48).

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- Guthrie RD (2006) New carbon dates link climatic change with human colonization and Pleistocene extinctions. *Nature* 441(7090):207–209.
- Barnosky AD, Koch PL, Feranec RS, Wing SL, Shabel AB (2004) Assessing the causes of late Pleistocene extinctions on the continents. *Science* 306(5693):70–75.
- Haile J, et al. (2009) Ancient DNA reveals late survival of mammoth and horse in interior Alaska. *Proc Natl Acad Sci USA* 106(52):22352–22357.
- Nikoskiy PA, Sulerzhitsky LD, Pitulko VV (2011) Last straw versus Blitzkrieg overkill: Climate-driven changes in the Arctic Siberian mammoth population and the Late Pleistocene extinction problem. *Quat Sci Rev* 30:2309–2328.
- Vartanyan SL, Arslanov KA, Karhu JA, Possnert G, Sulerzhitsky LD (2008) Collection of radiocarbon dates on the mammoths (*Mammuthus primigenius*) and other genera of Wrangel Island, northeast Siberia, Russia. *Quat Res* 70(1):51–59.
- Lorenzen ED, et al. (2011) Species-specific responses of Late Quaternary megafauna to climate and humans. *Nature* 479(7373):359–364.
- Veltre DW, Veltre MJ (1981) A preliminary baseline study of subsistence resource utilization in the Pribilof Islands. Alaska Department of Fish and Game, Division of Subsistence, Technical Paper no. 57. Available at: www.subsistence.adfg.state.ak.us/TechPap/tp057.pdf. Accessed June 22, 2016.
- Guthrie RD (2004) Radiocarbon evidence of mid-Holocene mammoths stranded on an Alaskan Bering Sea island. *Nature* 429(6993):746–749.
- Colinvaux P (1981) Historical ecology in Beringia: The south land bridge coast at St. Paul Island. *Quat Res* 16(1):18–36.
- Hanna GD (1922) The reindeer herds of the Pribilof Islands. *Sci Mon* 15:181–186.
- Veltre DW, Yesner DR, Crossen KJ, Graham RW, Coltrain JB (2008) Patterns of faunal extinction and paleoclimatic change from mid-Holocene mammoth and polar bear remains, Pribilof Islands, Alaska. *Quat Res* 70(1):40–50.
- Bryson RA, Agenbroad LD, McEnaney DeWall K (2010) Paleoclimate modeling and paleoenvironmental interpretations for three instances of island dwelling mammoths. *Quat Int* 217(1–2):6–9.
- Willerslev E, et al. (2003) Diverse plant and animal genetic records from Holocene and Pleistocene sediments. *Science* 300(5620):791–795.
- Baker AG, Bhagwat SA, Willis KJ (2013) Do dung fungal spores make a good proxy for past distribution of large herbivores? *Quat Sci Rev* 62:21–31.
- Rawlence NJ, et al. (2014) Using palaeoenvironmental DNA to reconstruct past environments: Progress and prospects. *J Quat Sci* 29(7):610–626.
- van Geel B, Zazula GD, Schweger CE (2007) Spores of coprophilous fungi from under the Dawson tephra (25,300 ^{14}C years BP), Yukon Territory, northwestern Canada. *Palaeogeogr Palaeoclimatol Palaeoecol* 252:481–485.
- Thienpont JR, et al. (2013) Exploratory hydrocarbon drilling impacts to Arctic lake ecosystems. *PLoS One* 8(11):e78875.
- Stevens RE, Hedges REM (2004) Carbon and nitrogen stable isotope analysis of northwest European horse bone and tooth collagen, 40,000 BP–present: Palaeoclimatic interpretations. *Quat Sci Rev* 23(7–8):977–991.
- Newsome SD, Miller GH, Magee JW, Fogel ML (2011) Quaternary record of aridity and mean annual precipitation based on $\delta^{15}\text{N}$ in ratite and dromornithid eggshells from Lake Eyre, Australia. *Oecologia* 167(4):1151–1162.
- de Boer EJ, et al. (2015) A deadly cocktail: How a drought around 4200 cal yr BP caused mass mortality events at the infamous “dodo swamp” in Mauritius. *Holocene* 25(5):758–771.
- Benedict FG (1936) *The Physiology of the Elephant* (The Carnegie Institution of Washington, Washington, DC).
- Western D (1975) Water availability and its influence on the structure and dynamics of a savanna large mammal community. *E Afr Wildl J* 13:265–286.
- Ramey EM, Ramey RR, Brown LM, Kelley ST (2013) Desert-dwelling African elephants (*Loxodonta africana*) in Namibia dig wells to purify drinking water. *Pachyderm* 53:66–72.
- Dunkin RC, Wilson D, Way N, Johnson K, Williams TM (2013) Climate influences thermal balance and water use in African and Asian elephants: Physiology can predict drivers of elephant distribution. *J Exp Biol* 216(Pt 15):2939–2952.
- Kubiak H (1982) Morphological characters of the mammoth: an adaptation to the arctic-steppe environment. *Paleoecology of Beringia*, eds Hopkins DM, Matthews JV, Schweger CE (Academic, New York), pp 281–289.
- Tridico SR, Rigby P, Kirkbride KP, Haile J, Bunce M (2014) Megafaunal split ends: Microscopical characterisation of hair structure and function in extinct woolly mammoth and woolly rhino. *Quat Sci Rev* 83:68–75.
- Feulner AJ (1980) Water resources reconnaissance of the southeastern part of St. Paul Island. USGS Water-Resources Investigations Report 80-61 (US Geological Survey, Reston, VA).
- Clegg BF, Hu FS (2010) An oxygen-isotope record of Holocene climate change in the south-central Brooks Range, Alaska. *Quat Sci Rev* 29(7–8):928–939.
- Finney BP, Bigelow NH, Barber VA, Edwards ME (2012) Holocene climate change and carbon cycling in a groundwater-fed, boreal forest lake: Dune Lake, Alaska. *J Paleolimnol* 48:43–54.
- Kuusaaari M, et al. (2009) Extinction debt: A challenge for biodiversity conservation. *Trends Ecol Evol* 24(10):564–571.
- Jones MC, Wooller M, Peteet DM (2014) A deglacial and Holocene record of climate variability in south-central Alaska from stable oxygen isotopes and plant macrofossils in peat. *Quat Sci Rev* 87:1–11.
- Laws RM (1970) Elephants as agents of habitat and landscape change in East Africa. *Oikos* 21:1–15.
- Van Wyk P, Fairall N (1969) The influence of the African elephant on the vegetation of the Kruger National Park. *Koedoe* 12:57–89.
- Mayewski PA, et al. (2004) Holocene climate variability. *Quat Res* 62(3):243–255.
- Feeley TC, Winer GS (2009) Volcano hazards and potential risks on St. Paul Island, Pribilof Islands, Bering Sea, Alaska. *J Volcanol Geotherm Res* 182(1–2):57–66.
- Clark PU, et al. (2016) Consequences of twenty-first-century policy for multi-millennial climate and sea-level change. *Nat Clim Change* 6:360–369.
- Hu A, et al. (2010) Influence of Bering Strait flow and North Atlantic circulation on glacial sea-level changes. *Nat Geosci* 3(2):118–121.
- Blockley SPE, et al. (2005) A new and less destructive laboratory procedure for the physical separation of distal glass tephra shards from sediments. *Quat Sci Rev* 24(16–17):1952–1960.
- Brown TA, Nelson DE, Vogel JS, Southon JR (1988) Improved collagen extraction by modified Longin method. *Radiocarbon* 30(2):171–177.
- Stuiver M, Polach HA (1977) Discussion: Reporting of ^{14}C data. *Radiocarbon* 19(3):355–363.
- Bronk RC (2009) Bayesian analysis of radiocarbon dates. *Radiocarbon* 51:337–360.
- Reimer PJ, et al. (2013) Intcal13 and Marine13 radiocarbon age calibration curves 0–50,000 years cal BP. *Radiocarbon* 55(4):1869–1887.
- Faegri K, Kaland PE, Krzywinski K (1989) *Textbook of Pollen Analysis* (Wiley, New York), 4th Ed.
- Battarbee RW, et al. (2001) Diatoms. Tracking Environmental Change Using Lake Sediments, Vol. 3: Terrestrial, Algal and Siliceous Indicators, eds Smol JP, Birks HJB, Last WM (Kluwer, Dordrecht), pp 155–202.
- Korhola A, Rautio M (2001) Cladocera and other branchiopod crustaceans. Tracking Environmental Change Using Lake Sediments, Vol. 4: Zoological Indicators, eds Smol JP, Birks HJB, Last WM (Kluwer, Dordrecht), pp 5–41.
- Wooller MJ, et al. (2004) Quantitative paleotemperature estimates from $\delta^{18}\text{O}$ of chironomid head capsules preserved in arctic lake sediments. *J Paleolimnol* 31(3):267–274.
- Wooller MJ, et al. (2012) Reconstruction of past methane availability in an Arctic Alaska wetland indicates climate influenced methane release during the past similar to 12,000 years. *J Paleolimnol* 48(1):27–42.
- Wooller MJ, et al. (2012) An ~11,200-year paleolimnological perspective for emerging archaeological findings at Quartz Lake, Alaska. *J Paleolimnol* 48(1):83–99.