



Title	Impact of Climate Change on Hunter-Fisher-Gatherer Cultures in Northern Japan Over the Past 4,400 Years
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Impact of climate change on hunter-fisher-gatherer cultures in northern Japan over the past 4400 years

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22 **Text S1**

23

24 **1.1. History of Hokkaido**

25 Human culture on inland Hokkaido developed through the succession from the Jomon
26 (~8,000–300 BCE), Zoku-Jomon (~300 BCE–700 CE), Satsumon (~700–1200 CE), and
27 Ainu cultures (after ~1200 CE) (*e.g.*, Fujimoto, 1965; Fujio, 2021). Along the northeastern
28 coast of Hokkaido, the Okhotsk culture (marine culture) continued from ~500 to 900 CE.
29 The Jomon culture, distributed through the entire Japanese Archipelago, was gradually
30 replaced by the Yayoi culture with rice farming during the first millennia BCE. Until ~300
31 BCE, the Yayoi culture spread to Honshu Island except for its northernmost area. The
32 people in Hokkaido and northern Honshu continued to hunt, fish, and gather in the
33 Zoku-Jomon period (Nishimoto, 1985; Minagawa, 2014). The Zoku-Jomon culture
34 expanded to South Sakhalin and the Kuril Islands from ~500 BCE to 1 BCE. After ~200
35 CE, the people of the Okhotsk culture migrated from Sakhalin Island, settled along the
36 coast of the Okhotsk Sea, and engaged in intense hunting of marine mammals, fishing,
37 terrestrial hunting, gathering, and primitive farming (Nishimoto, 1985; Takahashi, 2002;
38 Minagawa, 2014; Leipe et al., 2017). This culture was modified to the Tobinitai culture
39 characterized by intense salmon fishing and disappeared until ~1200 CE (Yamaura, 1983).
40 The people of the Satsumon culture fished for salmon, hunted animals, gathered food, and
41 engaged in primitive farming (Fujimoto, 1981; Minagawa, 2014). The Ainu culture started
42 around 1200 CE, expanded northward to Sakhalin Island, and then later eastward to the
43 Kuril Islands at 1500 CE.

44

45 **Text S2**

46

47 **2.1. Radiocarbon dating**

48 The remains of *Sphagnum* in the peats were prepared using the acid–alkali–acid (AAA)
49 treatment (Okuno et al., 2001). The samples were combusted with CuO at 850°C for 3 h in
50 a sealed quartz glass tube to produce CO₂, and the CO₂ was purified in liquid N₂ and
51 EtOH–liquid N₂ traps (Kitagawa et al., 1993). The purified CO₂ was reduced to graphite
52 with an iron powder catalyst. The graphite was then pressed into targets and analyzed at the
53 accelerator mass spectrometry facility at the Museum of the University of Tokyo. Stable
54 carbon isotope compositions were measured using an elemental analyzer mass spectrometer
55 equipped with a continuous flow system (Elementar EA1110 and Thermo Fisher Scientific
56 Delta Plus Advantage). Conventional ages were converted to calendar ages using the OxCal
57 program (ver. 4.4; Bronk Ramsey, 2021) and the IntCal20 dataset (Reimer et al., 2020).
58 Median values were used for the creation of the age–depth model. The sample at 51 cm
59 depth in MHWL-3 shows two different ages (Table S1). We chose the older age for the
60 age–depth model because younger material can be incorporated during coring.

61

62 **2.2. Purification of cellulose**

63 Cellulose was separated and purified from the tissue fraction according to the following
64 procedure. The sample was ultrasonicated three times in a mixture of methanol and toluene
65 (1:1) solution for 30 min (first) and 5 min (second and third), and the supernatant was
66 discarded each time to remove lipids. The same three-round extraction procedure was
67 repeated with acetone solution. After drying, the sample was heated four times in a 28 g/L
68 sodium chlorite solution in a water–acetone mixture (35: 1) at 70°C for 1 h. The
69 supernatant was discarded each time to remove lignin. The residue was rinsed three times
70 with Milli-Q water at 70–80°C and then rinsed three times at room temperature. The
71 residue was further ultrasonicated four times, with gentle shaking in a 17% sodium
72 hydroxide solution at 80°C for 45 min. The supernatant was discarded to remove the
73 hemicellulose. The residue (purified cellulose) was dried at 60°C.

74

75 **2.3. Cellulose isotopes**

76 Analysis of cellulose oxygen isotopes was conducted using a continuous flow
77 pyrolysis elemental analyzer/isotope mass spectrometer (TCEA/Delta plus XL) (Sharp et
78 al., 2001). The cellulose (150 µg) was wrapped in a silver foil and decomposed in a furnace
79 at 1375°C. The cellulose yielded CO from the degradation of its unexchangeable oxygen
80 and carbon, and the $\delta^{18}\text{O}$ of this CO was analyzed by mass spectrometry. A cellulose
81 standard (Merck), with known $\delta^{18}\text{O}$ (27.4 ‰) relative to VSMOW, was simultaneously
82 measured and used for calibration of $\delta^{18}\text{O}$. The accuracy of the analysis of $\delta^{18}\text{O}$ was
83 ± 0.3 ‰. The average value of duplicate analyses was used for discussion.

84

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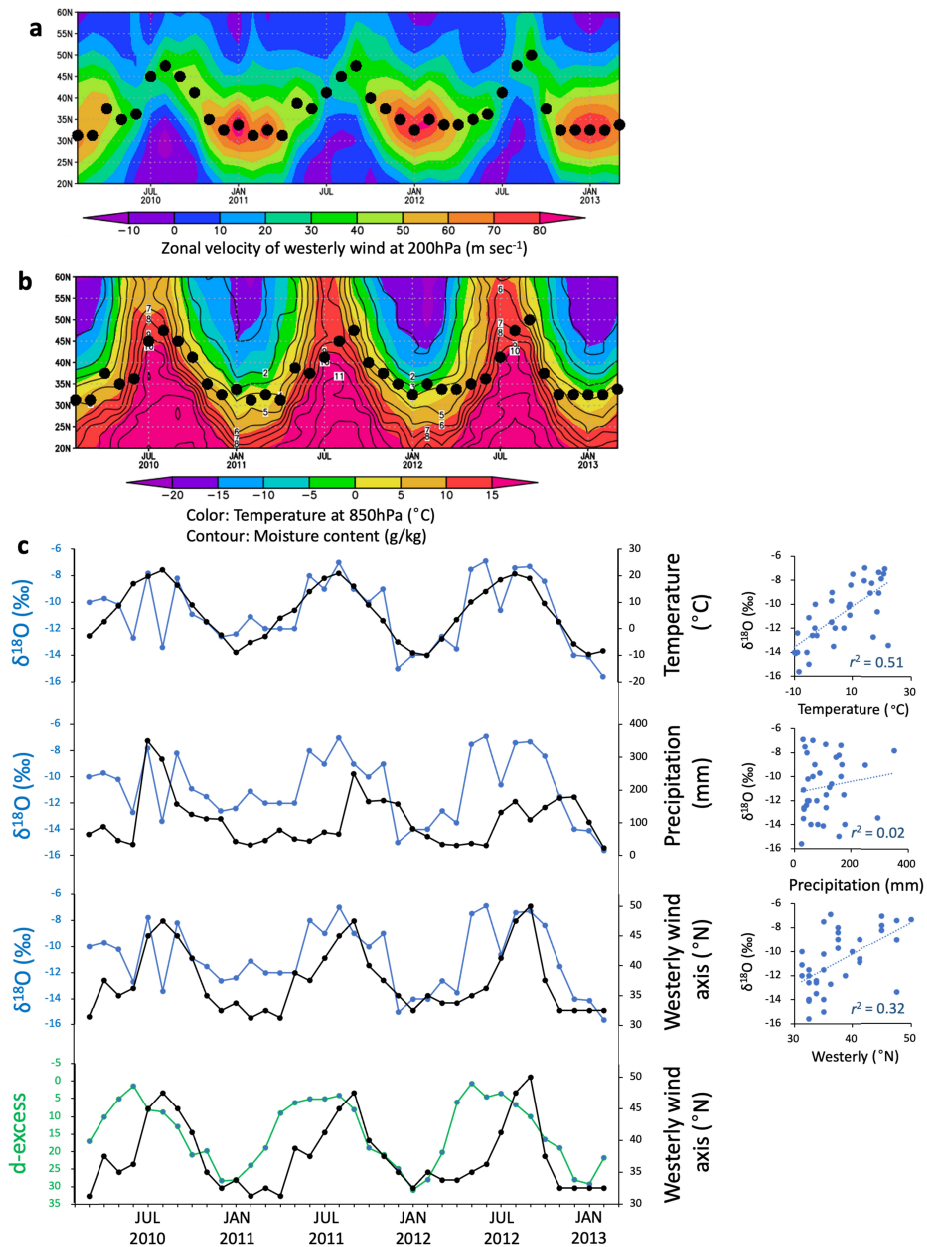
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- 121

123 **Table S1.** Raw radiocarbon dates of *Sphagnum* in cores MHWL-1 and MHWL-3

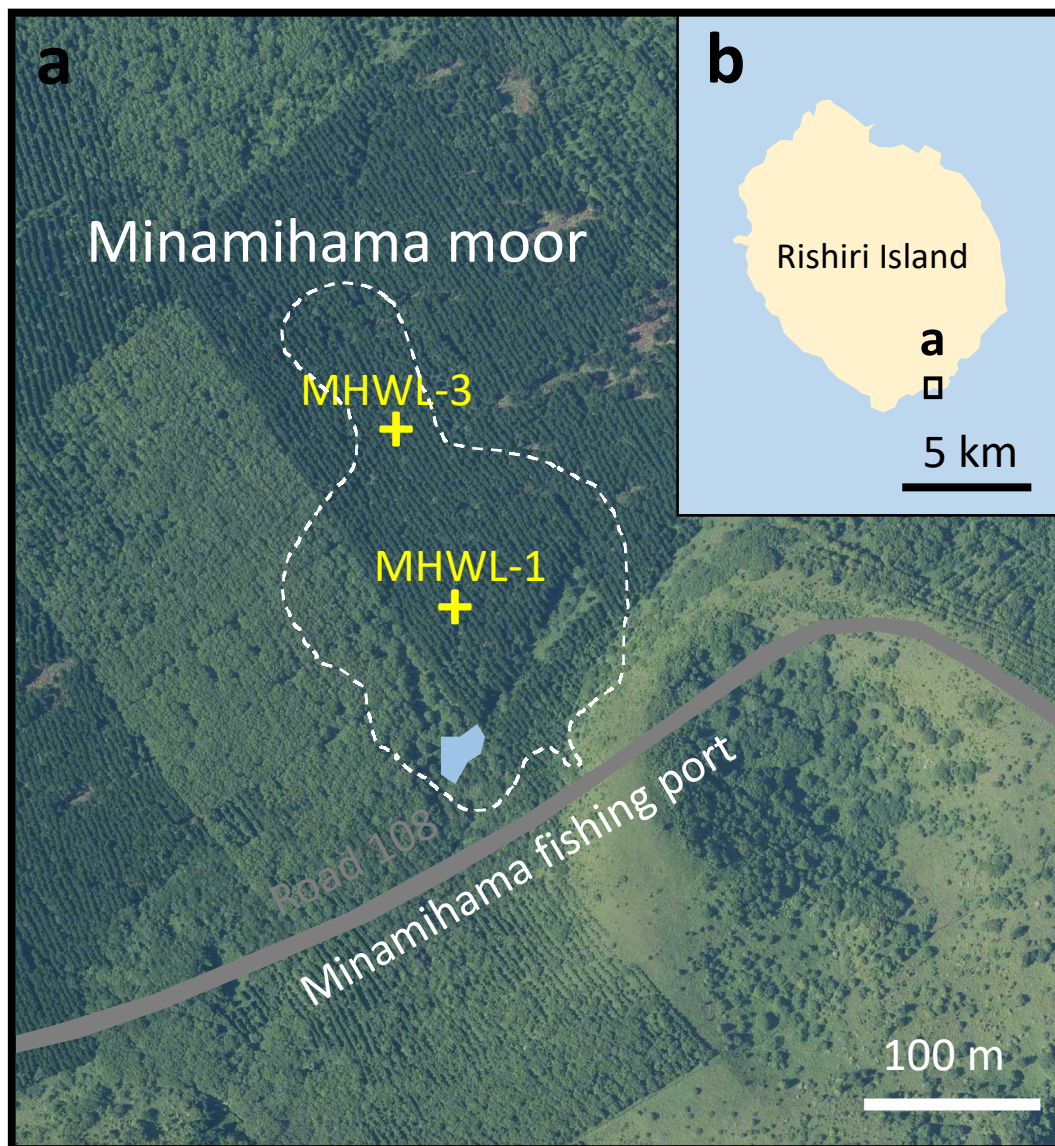
Core	Depth (cm)	Age (years BP) $\pm 1\sigma$			Note
MHWL-1	91	1137	\pm	20	
MHWL-1	221	2178	\pm	21	
MHWL-1	461	3908	\pm	23	
MHWL-3	51	212	\pm	32	not used
MHWL-3	51	290	\pm	21	
MHWL-3	131	1200	\pm	19	
MHWL-3	181	1625	\pm	21	
MHWL-3	271	1971	\pm	21	
MHWL-3	351	2430	\pm	27	



125

130 **Figure S1. (a)** Zonal wind velocity at 200 hPa (positive westward) along 145 °E, **(b)**
 131 temperature and moisture content at 850 hPa along 145°E (Sakurai et al., 2021), **(c)**
 132 precipitated water $\delta^{18}\text{O}$, d-excess, temperature and precipitation at the town of Teshio
 133 (2010–2013; Li et al., 2017) ~40 km southeast of the study site, and the position of the
 134 westerly jet. Black solid dots indicate the positions of the westerly jet.

131



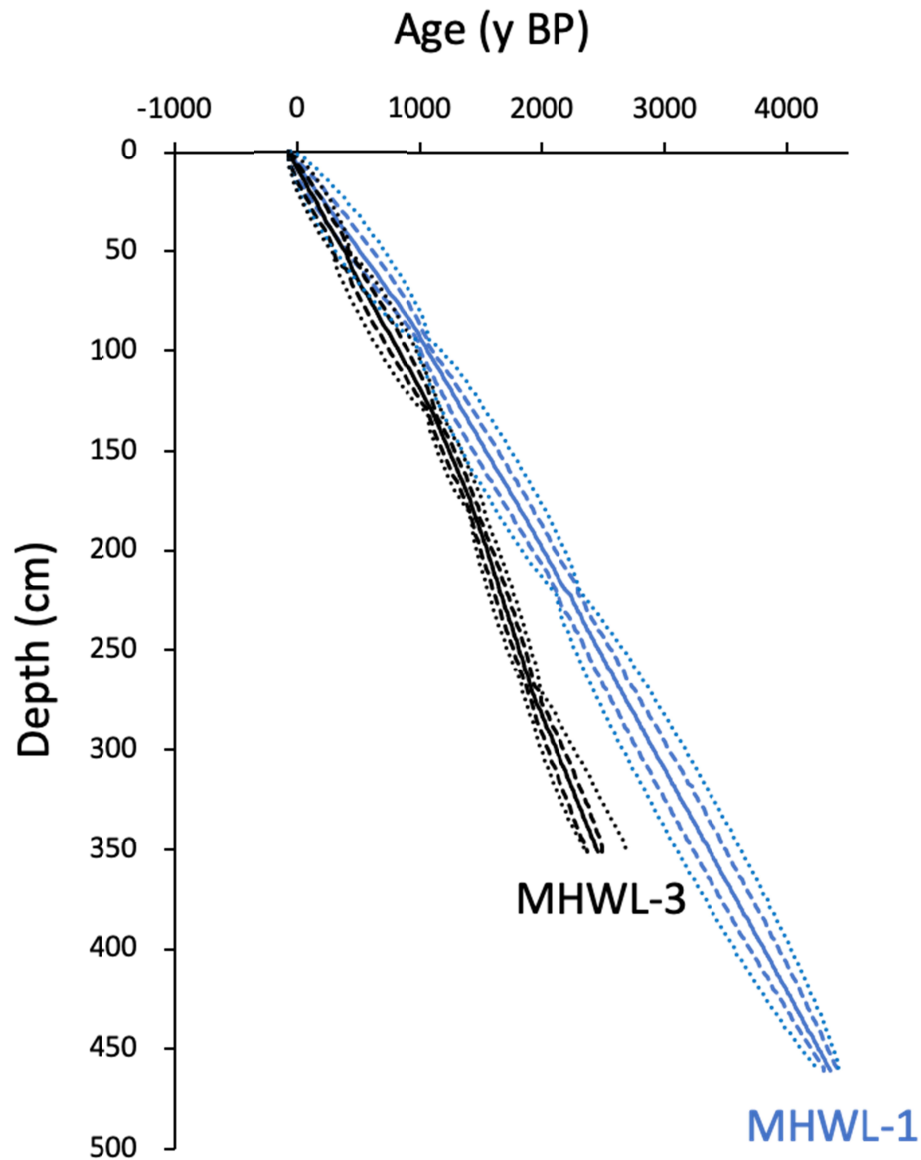
132

133 **Figure S2.** Locations of coring sites in (a) the Minamihama moor and at (b) Rishiri Island.

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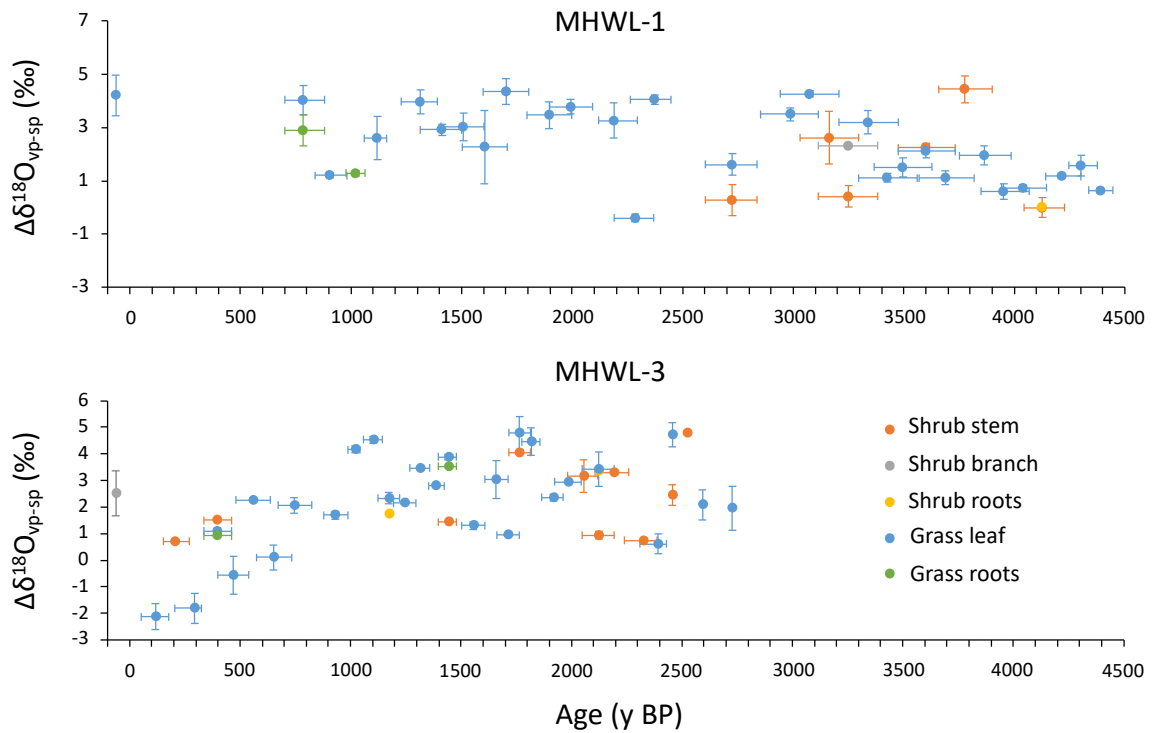
141 **Figure S3.** The age-depth model for cores MHWL-1 and MHWL-2. Solid, broken and
142 dotted lines in the age–depth plot indicate the median and the 1σ and 2σ range of the
143 modeled age, respectively.

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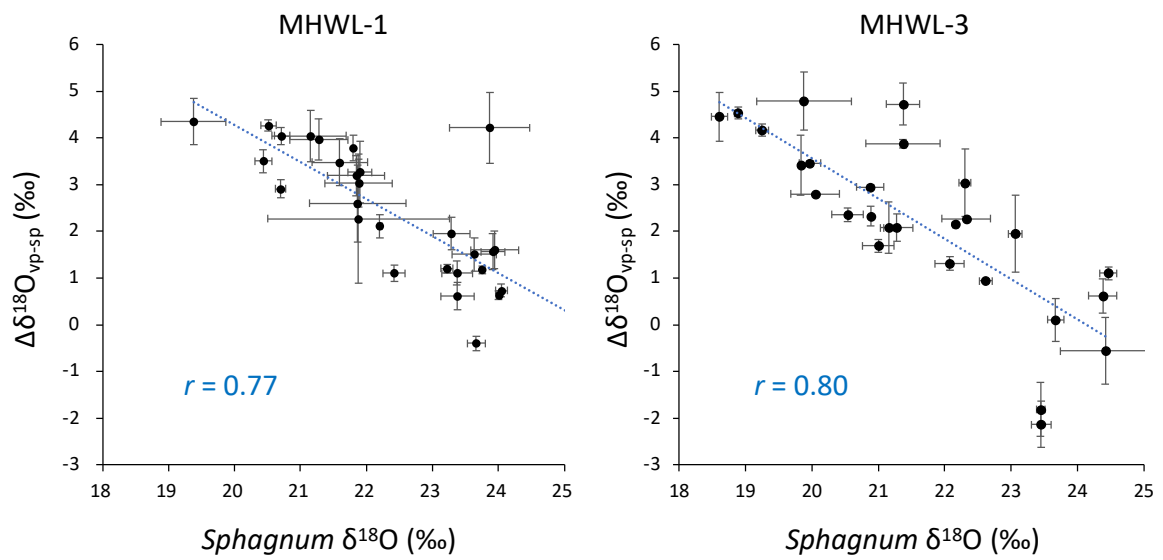
145

146 **Figure S4.** Changes in the difference of $\delta^{18}\text{O}$ between vascular plants and *Sphagnum*

147 ($\Delta\delta^{18}\text{O}_{\text{vp-sp}}$) in cores MHWL-1 and MHWL-3. The vertical and horizontal bars of the

148 sample value indicate the 1σ interval ranges.

149



150

151 **Figure S5.** Cellulose $\delta^{18}\text{O}$ values of *Sphagnum* and the difference of $\delta^{18}\text{O}$ between grass
152 leaf and *Sphagnum* in cores MHWL-1 and MHWL-3. Bars indicate 1 σ intervals.

153