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1 **Vegetation succession, carbon accumulation and hydrological change in sub-Arctic**
2 **peatlands (Abisko, N. Sweden)**

3

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13

14

15 **Keywords**

16 Climate change; permafrost; tephra; testate amoebae; carbon accumulation; *Sphagnum*
17 *lindbergii*.

18

19 **Abstract**

20 We present a multiproxy study of the hydrological, ecological and carbon accumulation
21 dynamics of two sub-Arctic peatlands in Abisko, Sweden. High-resolution analyses of plant
22 macrofossils, testate amoebae, pollen, mineral content, bulk density, and carbon and nitrogen
23 were undertaken. The peat records were dated using tephrochronology, ¹⁴C and ²¹⁰Pb. Local
24 plant succession and hydrological changes in peatlands were synchronous with climatic shifts,
25 although autogenous plant succession towards ombrotrophic status during peatland
26 development was also apparent. The Marooned peatland experienced a shift ca. 2250 cal yr
27 BP from rich to poor-fen, as indicated by the appearance of *Sphagnum fuscum*. At Stordalen,

28 a major shift to wetter conditions occurred between 500-250 cal yr BP, which is most
29 probably associated with climate change during the Little Ice Age. During the last few
30 decades we observe a deepening of water table and an increase in shrub pollen, coinciding
31 with recent climate warming and the associated expansion of shrub communities across the
32 Arctic zone. Rates of carbon accumulation vary greatly between the sites, illustrating the
33 importance of local vegetation communities, hydrology and permafrost dynamics. We also
34 use multiproxy data to elucidate the palaeoecology of *Sphagnum lindbergii* and find it is
35 indicative of wet conditions in peatlands.

36

37 **Introduction**

38 Peatlands located in the sub-Arctic and Arctic zones are important archives for
39 palaeoenvironmental reconstruction owing to their sensitivity to climate and hydrological
40 change (Turetsky *et al.*, 2002; Lamarre *et al.*, 2012; Swindles *et al.*, 2015a). The recent rapid
41 increase of global air temperatures at high latitude (IPCC, 2013) has led to partial or complete
42 thawing of permafrost in many regions (Turetsky *et al.*, 2002; Johansson *et al.*, 2006;
43 Callaghan *et al.*, 2010). Numerous peatland studies have been conducted in areas of
44 permafrost degradation that focus on different aspects of ecosystem response, including
45 changes in carbon accumulation rates (Loisel *et al.*, 2014; Swindles *et al.*, 2015a), organic
46 matter decomposition rates (Scanlon and Moore, 2000), plant succession and peatland
47 development pathways (Zoltai, 1993; Kuhry, 2008; Lemus-Lauzon *et al.*, 2016; Teltewskoi *et*
48 *al.*, 2016; Treat *et al.*, 2016) and hydrology (Swindles *et al.*, 2015a,b). Peatland carbon
49 accumulation rates depend on complex relationships between soil temperature and moisture,
50 litter composition and production, and degree of decomposition (Vardy *et al.*, 1998; Malmer
51 and Wallén, 2004; Sannel and Kuhry, 2009; Treat *et al.*, 2014). The importance of the length

52 of the growing season and incoming radiation has also been highlighted for carbon
53 accumulation in northern peatlands (Charman *et al.*, 2013).

54 Here we present a palaeoecological study from the Abisko region of northern Sweden
55 to examine the long term dynamics of permafrost peatlands. The Abisko region has
56 experienced rapid warming during the twentieth century (Callaghan *et al.* 2010; Swindles *et*
57 *al.*, 2015a), and documented degradation of permafrost peat and deepening of active layer
58 during recent decades (Zuidhoff and Kolstrup, 2000; Łkerman and Johansson, 2008; Kokfelt
59 *et al.*, 2009). Furthermore, climate model projections suggest continued marked temperature
60 increases in the near future in this region (Swindles *et al.*, 2015a, b). Detailed long-term
61 palaeoecological studies of these degrading permafrost peatlands are important for
62 understanding relationships between climate, vegetation, hydrology and carbon accumulation,
63 which has implications for understanding the response of high latitude peatlands to climate
64 warming (Gao and Couwenberg, 2015). We focus on the late Holocene, because this period
65 contains several phases of climate change (e.g. Roman Warm Period, cold Migration Period,
66 Medieval Warm Period, Little Ice Age). Although several palaeoecological studies have been
67 conducted in Abisko(e.g. Zuidhoff and Kolstrup, 2000; Malmer and Wallén, 2004; Kokfelt *et*
68 *al.*, 2009), this study represents the first multiproxy late Holocene data from peatlands in this
69 region.

70

71 The aims of this study are to: i) determine the drivers of vegetation change over the
72 late Holocene (last 2250 years); ii) reconstruct the late Holocene hydrological conditions of
73 two peatlands to evaluate possible links with climate and autogenic peatland processes; and
74 iii) assess the impact of climatic and ecological changes on carbon accumulation rates; and iv)
75 elucidate the palaeoecology of *Sphagnum lindbergii*. We also present new tephra data from

76 our study sites in northern Scandinavia – extending the current Northern European
77 tephrostratigraphy.

78

79 **Study sites**

80 The study sites are located in the Abisko region of northern Sweden's sub-Arctic zone (Fig.
81 1). The Marooned (7°57'24.0" N, 19°59'11.4" E) and Stordalen (68°21'24.3" N, 19°02'53.5"
82 E) peatlands were selected due to their high-latitude location near the tree line and at a
83 sensitive climatic boundary subject to alternating influences of air masses of Atlantic and
84 Arctic origin (Shemesh *et al.*, 2001). Because of this, we expect these peatlands to be
85 sensitive to relatively small changes in climate. They also contrast in terms of permafrost;
86 Stordalen is in an advanced state of permafrost degradation, whereas Marooned has no
87 remaining permafrost (Swindles *et al.*, 2015a), (Swindles *et al.*, 2015b). It has been suggested
88 that the Marooned site thawed in the mid-late twentieth century (Swindles *et al.*, 2015a). Both
89 peatlands are part of large peat complexes in this area, and typical of the peatlands of northern
90 Scandinavia in terms of microform characteristics and vegetation communities. The
91 vegetation communities of both peatlands are characterized by *Sphagnum fuscum*, *Rubus*
92 *chamaemorus*, *Eriophorum vaginatum*, *Eriophorum angustifolium* and *Betula nana*.
93 *Sphagnum balticum*, *Drepanocladus* sp. and *Carex rostrata* are also present in each site. Peat
94 depth was highly variable in both sites and varied from <1m to >3m in places. However, peat
95 probing proved difficult in places owing to the presence of permafrost. The Stordalen
96 monolith was extracted from the top of a Palsa-Peat Plateau complex containing permafrost
97 whereas the monolith at Marooned was in a wet hollow at the edge of a thaw lake. The
98 Marooned peatland no longer contains permafrost.

99 Peat initiation in Marooned took place (as determined from the core) before deposition
100 of the Hekla 4 tephra (~4287 cal yr BP), but our reconstructions here focus on the last 2650

101 cal yr BP (i.e. the large-capacity monolith required for our detailed multiproxy analysis).
102 While in Stordalen peat accumulation commenced ca. 4700 cal BP as a result of
103 terrestrialisation (Kokfelt *et al.*, 2010), our reconstructions cover ca. 900 cal yr BP. It has
104 been reported that the ombrotrophic *Sphagnum* peat (depth <1 m) above the fen peat in
105 Stordalen began not later than 800 cal yr BP (Malmer and Wallén, 2004). However, both
106 monoliths captured the marked transition from Cyperaceae- to *Sphagnum* dominated peat and
107 the vegetation succession which is our primary interest in this study.

108

109

110 **Materials and Methods**

111 *Overview*

112 Our palaeoecological investigation employs plant macrofossils, pollen, testate
113 amoebae, as well as analysis of the physical and geochemical properties of peat. Chronologies
114 are based on radiocarbon and lead-210 dating and the presence of microscopic tephra layers.
115 The recent progress in tephrochronology (Lowe, 2011) allows for the detection of even tiny
116 amounts of dispersed ash (cryptotephra) and provides precise chronological control and
117 synchronisation markers for paleoecological reconstructions (e.g. Wulf *et al.*, 2013; Davies,
118 2015).

119 The analysis of plant macrofossil remains reveals a record of local plant communities
120 and hydrological conditions (Barber *et al.*, 2004; Mauquoy *et al.*, 2008), while pollen analysis
121 provides information on vegetation composition and abundance at a wider regional scale
122 (Berglund *et al.*, 1996). The testate amoebae record is used to reconstruct past changes in
123 hydrological conditions; in this study we used the transfer function of Swindles *et al.* (2015b)
124 to reconstruct palaeo water-table depths. We used micro-charcoal as an indicator of fire
125 occurrence and frequency (Mooney and Tinner, 2011). Finally, we determined carbon

126 accumulation rates to assess the effect of climatic and ecological changes on peatland carbon
127 sequestration (e.g. Charman *et al.*, 2013).

128

129 *Coring and subsampling, chronology of the core*

130 Two short peat monoliths, Marooned (38 cm) and Stordalen (30 cm), were taken from
131 each site following excavation of a small trench following De Vleeschouwer *et al.* (2010) .
132 The monoliths were large in volume to allow this detailed multiproxy study, and the accurate
133 determination of bulk density for carbon accumulation estimates.

134 A narrow-chamber core of 1m was also taken from Marooned using a Russian corer
135 immediately adjacent to the monolith for analysis of tephra. Not enough material was
136 available in the core for a detailed multiproxy study. To determine the age of the peat
137 samples, radiocarbon measurements were performed on handpicked terrestrial plant
138 macrofossils at Direct AMS, Seattle, USA. High-resolution ^{210}Pb -dating was applied to the
139 uppermost part of the monolith samples. Samples were acid digested with concentrated HCl,
140 HNO_3 , and H_2O_2 , and a spike of ^{209}Po added as a yield tracer. Material was plated onto silver
141 disks and the activity measured by alpha spectrometry. The date of each sample was derived
142 from the ^{210}Pb inventory using a constant rate of supply model (Appleby, 2001). The
143 calibrated dates are expressed as cal yr BP. Simple age models based on linear-interpolation
144 were generated using Clam v 2.2 (Blaauw, 2010).

145

146 *Plant macrofossil analysis*

147 Plant macrofossil remains were analysed contiguously at 1-cm intervals resulting in 68
148 samples. Samples of 5 cm^3 were washed and sieved under a warm-water spray using a 0.20-
149 mm mesh sieve. Initially, the entire sample was examined with a stereomicroscope to obtain
150 volume percentage of individual subfossils of vascular plants and mosses. The subfossil

151 carpological remains and vegetative fragments (leaves, rootlets, epidermis) were identified
152 using identification keys (Smith, 2004; Mauquoy and van Geel, 2007). Refer to Gałka et al.
153 (2016) for detailed information on the methods used for plant macrofossil analysis of peats.
154 We use the nomenclature of Mirek *et al.* (2002) for vascular plants and of Smith (2004) for
155 bryophytes.

156

157 *Palynological and charcoal analyses*

158 Samples of 1 cm³ were prepared using the standard procedure of Erdtman's acetolysis
159 (Berglund and Ralska-Jasiewiczowa, 1986). *Lycopodium* tablets were added to each sample to
160 enable quantitative analysis of microfossil concentration. Approximately 500 terrestrial pollen
161 grains per sample were counted. For taxonomical identification pollen keys (Beug, 2004) and
162 a reference collection of modern pollen were used. Non-pollen palynomorphs (NPPs) were
163 counted along with the pollen. NPPs were identified according to van Geel (1978) and
164 Shumilovskikh *et al.* (2015). Microcharcoal particles were counted along with the pollen and
165 were grouped into four size classes: 10-30 µm, 30-70 µm, 70-100 µm and >100 µm (Rull,
166 2009). The arboreal pollen (AP) + non-arboreal pollen (NAP) sum was used for percentage
167 calculations. The pollen diagram was stratigraphically ordered and zoned with constrained
168 cluster analysis (CONISS) and divided into local pollen assemblage zones.

169

170 *Testate amoebae*

171 Testate amoebae were extracted using a modified version of Booth et al. (2010). Peat
172 samples (2cm³) were placed in boiling water for 15 minutes and shaken. Extracts were passed
173 through a 300 µm sieve, back-sieved at 15 µm and allowed to settle before sub-samples were
174 used to make slides for microscopy. 100 to 200 amoebae were counted and identified to
175 species level or 'type' in each sample using high-power transmitted light microscopy at 200 to

176 400x magnification. For further detailed information on taxonomy and sources used see
177 Swindles *et al.* (2015b). The Abisko transfer function (Swindles *et al.*, 2015b) was applied to
178 the data to generate water-table depth reconstructions.

179

180 *Physical properties and geochemical analysis*

181 The difference in sample mass before and after freeze drying was used to calculate
182 moisture content, with bulk density calculated by dividing dry sample mass by volume
183 (Chambers *et al.*, 2011). A volume of 0.5×2.5×2.5 cm (3.125 cm³) was used for bulk density
184 determination. C and N content was measured on a Thermo Scientific Flash 2000 Series
185 CHNS/O Analyser. Carbon density (g C cm⁻³) was calculated by multiplying bulk density
186 with percentage carbon content, and carbon accumulation rates were calculated using the age
187 depth model described above (Charman *et al.*, 2013). We note that these are apparent
188 accumulation rates and must be treated with some degree of caution as they are from
189 uppermost peats that not have undergone full decomposition.

190

191

192 *Tephra*

193 Contiguous 5 cm samples were ashed at 550 °C and treated with 10% HCl before
194 mounting onto slides (Swindles *et al.*, 2010). Where tephra shards were identified under 400 x
195 magnification, cores were subsampled at 1 cm intervals. Extraction for geochemical analysis
196 followed the density separation method (cleaning float 2.25 g cm⁻³, retaining float 2.50 g cm⁻³)
197 (Blockley *et al.*, 2005). Shards were mounted onto slides (Dugmore and Newton, 1992) and
198 major element geochemistry was determined using an electron probe microanalyser (EPMA).
199 Glass shards from Marooned peatland were analysed on the JEOL JXA8230 EPMA
200 (University of Leeds) and samples from Stordalen peatland were analysed on the Cameca

201 SX100 EPMA (Tephra Analytical Unit, University of Edinburgh). Assignments to specific
202 eruptions were based on stratigraphy and visual comparison of tephra geochemistry with the
203 Tephabase database (Newton *et al.*, 2007) and other published literature. The full analytical
204 setup is presented in Appendix 1.

205

206 **Results**

207 *Chronology, peat accumulation rate, and lithology*

208 The Marooned core has a much older basal date (38 cm = ca. 2650 cal yr BP) when
209 compared with the Stordalen profile (25 cm = ca. 900 cal yr BP), resulting in a slower mean
210 peat accumulation rate at Marooned (~0.14 mm/year) compared with Stordalen (~0.31
211 mm/year) over the respective periods. Full radiocarbon and ²¹⁰Pb data for both profiles are
212 presented in Appendix 2 and age-depth model on Fig. 2. Detailed lithostratigraphic
213 information is presented in Table 1 and Fig. 3.

214

215 *Plant macrofossils*

216 Four zones in the local vegetation development were visually delimited at the
217 Marooned peatland (Fig. 3A). In the first zone MA-ma-1 (2650 to 2250 cal yr BP) *Carex* sp.
218 with brown mosses is dominant. The second zone MA-ma-2 (2250 to 600 cal yr BP) contains
219 small amounts of *Betula nana*, *Empetrum nigrum* and *Andromeda polifolia*, but is primarily
220 characterised by the dominance of *Sphagnum fuscum* and *S. magellanicum*. The third zone
221 (MA-ma-3, 600 to -5 cal yr BP) and the fourth zone MA-ma-4 (-5 to -62 cal yr BP) are
222 dominated by *S. fuscum*. *Dicranum elongatum* appears for the first time at ca. -10 cal yr BP.
223 An expansion of shrubs is represented by the presence of *B. nana* and *E. nigrum*.

224 Three phases in the local vegetation development were delimited at Stordalen (Fig.
225 3B). The first zone, ST-ma-1 (ca. 900-280 cal yr BP), is characterised by the continuous

226 presence of vegetative Cyperaceae macrofossils and *Carex* sp. fruits. In the bottom part of the
227 second stage ST-ma-2 (ca. 280-150 cal yr BP) *Drepanocladus* sp. was the dominant species.
228 *Sphagnum lindbergii* played a dominant role at ca. 220 cal yr BP. In the third zone ST-ma-3
229 (ca. 150-62 cal yr BP) *Sphagnum balticum* was dominant. Since ca. 100 cal yr BP
230 *Polytrichum commune* appeared and *A. polifolia* macrofossils re-appeared.

231

232 *Pollen*

233 Four phases in the regional and local vegetation development were statistically
234 delimited at Marooned peatland (Fig. 4). The first phase MA-po-1 (ca. 2650-2300 cal yr BP)
235 is characterised by the highest presence of Cyperaceae pollen (up to 10%) and by the
236 occurrence of *Menyanthes trifoliata* and *Aster* type. Among non-pollen palynomorphs
237 *Botryococcus*, and copepod spermatophores were present. In the second phase (ca. 2300-1000
238 cal yr BP) there is a peak in *Salix* (up to 15%) and *Rubus chamaemorus* (up to 10%). In the
239 third zone MA-po-3 (ca. 1000-25 cal yr BP) there is a peak of Ericaceae (up to 30%) ca. 450
240 cal. BP, and at ca. 5 cal yr BP *Picea* (up to 2%) and *R. chamaemorus* (up to 5%) increased.
241 *Thalictrum*, *Calluna*, *Equisetum*, and palynomorphs such as the shells of *Habrotrocha*
242 *angusticollis* and ascospores of Xylariaceae were present. The middle part of this zone also
243 contains abundant microcharcoal. In the fourth zone MA-po-4 (ca. -25- -62 cal yr BP) *Picea*
244 increased up to 3%, and *Pinus* up to 87% together with an increase in the role of Filicales
245 monoletae, Ericaceae, and *Arctostaphylos*.

246

247 *Testate amoebae*

248 Two phases in the testate amoebae sequence were identified at Marooned (Fig. 5A). In
249 first zone MA-ta-1 (ca. 2650-30 cal yr BP) *Diffflugia pristis* type (up to 30%), *Diffflugia pulex*
250 (up to 40%) and *Nebela militaris* (up to 25%) were the dominant taxa. The reconstructed

251 water-table depth (WTD) ranged between 22 and 28 cm below the peatland surface. During
252 the second zone MA-ta-2 (30-62 cal yr BP) *Longinebla militaris* increased to 40%. At ca. 5
253 cal yr BP there are peaks of *Archerella flavum* (up to 17%), and at ca. -40 cal yr BP *Euglypha*
254 *tuberculata* (up to 12%). The WTD in this zone ranged between 21 and 28 cm below the
255 peatland surface. Overall, the WTD reconstruction suggests relatively dry conditions
256 throughout.

257 Three phases in the testate amoebae succession were delimited at Stordalen (Fig. 5B).
258 The first phase ST-ta-1 (ca. 900-480 cal yr BP) is characterised by *Trigonopyxis arcuata* type
259 (up to 65%) and deep, stable WTD ca. 35 cm. In the phase ST-ta-2 (ca. 480-320 cal yr BP)
260 there is an increase in the abundance of *Archerella flavum*, *Assulina scandinavica*,
261 *Hyalosphenia papilio*, and *Placocista spinosa* type. In the upper part of phase ST-ta-2 (ca.
262 480-320 cal yr BP) *Physochila griseola* appears for the first time (up 20%) and reconstructed
263 WTD indicates a fluctuating water table from 34 to 8 cm. In the third phase St-ta-3 (ca. 320-
264 62 cal yr BP) *Archerella flavum* increases up to 65% and *P. spinosa* type up to 27%. The
265 WTD ranged between 8 and 18 cm. The Stordalen profile, in contrast to Marooned, contains a
266 clear 'wet-shift' in hydrology.

267

268 *Geochemical analysis*

269 The bulk density values for Marooned and Stordalen are typical of northern peatlands
270 (Loisel *et al.*, 2014) but vary noticeably during the different phases of peatland development.
271 In general, values are higher for the fen phase when peat is mainly composed of *Cyperaceae*,
272 while the values decrease for the more ombrotrophic phase, when the main peat forming
273 plants are mosses, which tend to form a less dense peat (Fig. 6 and Fig. 7). This pattern may
274 also be due partly to compaction. C/N ratio increases in both sites through time and is higher
275 in the later ombrotrophic phases before decreasing in the near-surface zone.

276

277 *Carbon accumulation rates*

278 Our study sites have high variability of carbon accumulation (CA) rates during the
279 studied time periods. In Marooned the CA rates range from ~0 to 128 gC m⁻² yr⁻¹ and are
280 lowest in zone B (ca. 500 yr BP – 2250 yr BP) and highest in the boundary between zones C
281 and D (ca. 50 yrs BP) (Fig. 6). The CA rates at Stordalen vary from ~0 to 27 gC m⁻² yr⁻¹ and
282 are generally much slower than at Marooned. The CA at Stordalen is highest in the boundary
283 between zones B and C (ca. 300 BP) and lowest at the boundary between A and B (ca. 500
284 BP) (Fig. 7).

285

286 *Tephra*

287 All tephra data are displayed in Appendix 3. Two tephra layers (MR-1 = 70 cm, MR-2
288 = 85 cm) were identified in the 1 m core at Marooned. MR-1 contains glass shards matching
289 the geochemistry of glass from the Hekla-S/Kebister eruption (~3720 cal yr BP) (Wastegård
290 *et al.*, 2008). MR-2 contains glass shards matching the geochemistry of glass produced during
291 the eruption of Hekla 4 (~4287 cal yr BP; Pilcher *et al.*, 1995).

292 The peat at the Stordalen peatland contains one tephra layer (SD-1). SD-1 is at 23 cm
293 depth and contains tephra shards matching the major element geochemistry of glass from the
294 eruption of Hekla in AD 1158 (Pilcher *et al.*, 2005). This tephra layer is historically dated and
295 with a peak within a one-centimeter horizon provides a precise chronological marker for the
296 Stordalen profile.

297

298 **Discussion**

299

300 1. Ecohydrological development

301 In the bottom part of both peat sequences the main peat-forming plants included
302 Cyperaceae (Fig. 6, phase A, Fig. 7, phase A and B). The presence of *Carex* fruits and
303 rootlets, as well brown mosses in both sites indicate that during this period the peatlands were
304 operating as fens, a widespread wetland type in the permafrost area of the northern
305 hemisphere (Vardy *et al.*, 1998; Kuhry, 2008; Teltewskoi *et al.*, 2016; Treat *et al.*, 2016).
306 Pollen data indicate that between 2650 and 2300 cal yr BP the non-peatland community at
307 Marooned was dominated by *Betula*. Previous research indicates that the proportion of *Betula*
308 in regional plant communities gradually increased since ca. 4100 cal yr BP (Bigler *et al.*,
309 2002), as a result of climate cooling in northern Fennoscandia (Berglung *et al.*, 1996; Korhola
310 *et al.*, 2000).

311 Around 2,250 cal yr BP, a new plant community dominated by *Sphagnum fuscum* and
312 *S. magellanicum* developed at Marooned. These species indicate a transition from
313 groundwater fed fen to rainwater-fed bog (Hughes, 2000), due most likely to autogenic plant
314 succession toward hummock development (Malmer, 2014). *S. fuscum* is a typical
315 ombrotrophic species and usually occurs on dry areas of the bog (Hölzer, 2010; Laine *et al.*,
316 2011), but can also build high hummocks in fens (Laine *et al.*, 2011). Autogenic plant
317 community changes from fen to dry bog may be supported by appearance of *Rubus*
318 *chamaemorus* and *Ledum/Empetrum* pollen. These plants usually grow on the top of
319 hummocks in ombrotrophic peatlands in the sub-Arctic zone of Scandinavia (Zuidhoff and
320 Kolstrup, 2005). Further evidence of drying is provided by testate amoebae communities
321 which indicate dry hydrological conditions with a reconstructed WTD of between 24 and 30
322 cm below the peat surface in the first ombrotrophic stage (Fig. 6, zone B). The development
323 of poor fen and then bog communities ca. 2250 cal yr BP may also be caused by permafrost
324 aggradation and physical separation of this part of peatland from the groundwater (Vardy *et*

325 *al.*, 1998). According to Kokfelt *et al.* (2010) a period of notable permafrost aggradation
326 occurred ca. 2650–2100 cal yr BP in the Abisko region.

327

328 2. Little Ice Age (LIA) conditions

329 Stable hydrological conditions with DWT at ca. 23 cm were inferred between ca. 500
330 and 50 cal yr BP at Marooned (Fig. 6, zone B and C). In this period *S. fuscum* and
331 *Polytrichum strictum* were the dominant species, with a persistent presence of *Dicranum*
332 *undulatum*. *D. undulatum* and *P. strictum* which usually occur in dry areas of peatlands, often
333 on hummocks (Hedenäs and Bisang, 2004; Smith, 2004).

334 During the LIA we noted changes in local plant communities in Marooned, mainly
335 shrubs that coincide with the increase in micro-charcoal (Figs. 4 and 6). The appearance of *P.*
336 *strictum*, and increasing abundances of *Ericaceae*, *Calluna vulgaris* pollen in Marooned
337 peatland may indicate the deepening of water table which favoured the development of shrub
338 communities (Zuidhoff and Kolstrup, 2005). The lack of macro-charcoal (e.g. burned
339 branches of shrubs or *Sphagnum* leaves) in the *S. fuscum* peat layer indicates that fire was
340 most likely not present at the site, but possible elsewhere in the surrounding environment.
341 Furthermore, it has been suggested that fires are generally not an important driver in
342 modifying *Sphagnum* communities (Kuhry, 1994; Magnan *et al.*, 2012). Fire may be favoured
343 by deep water tables in some peatlands which may lead to widespread thaw and subsidence of
344 permafrost (Jones *et al.*, 2015). Additionally, fire is known to lower water reserves in adjacent
345 unburned wetland areas (Watts *et al.*, 2015).

346 A very different hydrological pattern is apparent at Stordalen, where a marked shift
347 toward wetter conditions occurred from ca. 450 to 400 cal yr BP (WTD from 35 to 16 cm),
348 before a shift to drier conditions from ca. 400 to 330 cal yr BP and reverting to wet conditions
349 from ca. 330 cal yr BP to present day (Fig. 7, zone B). In this time period of variable

350 hydrological conditions (Zone B) plant assemblages dominated by *Carex* sp. are observed,
351 indicative of poor fen conditions. Hydrological changes between ca. 500 and 300 cal yr BP in
352 Stordalen may coincide with hydrological shifts caused by regional scale climatic changes
353 during the Little Ice Age (LIA) which have been documented in various parts of Europe
354 (Mauquoy *et al.*, 2008; Swindles *et al.*, 2007; Gałka *et al.*, 2014; Feurdean *et al.*, 2015).
355 During the LIA moisture patterns differed between North-East Europe and Central-East
356 Europe. North-West Europe and Scandinavia incurred shifts toward wetter climate (Charman
357 *et al.*, 2006; Väiliranta *et al.*, 2007; Swindles *et al.*, 2007, 2010; Mauquoy *et al.*, 2008) whilst
358 Central-East Europe tended toward drier conditions (Gałka *et al.*, 2014, 2017; Feurdean *et al.*,
359 2015). The different palaeohydrological pathways in peatland development in North-West
360 Europe, Scandinavia and Central-East Europe could be connected with the fluctuations in
361 largescale atmospheric circulation (Feurdean *et al.*, 2015; Gałka *et al.*, 2017). However,
362 hydrological shifts associated with the LIA were not observed at Marooned. Differences in
363 the hydrological pattern in Marooned and Stordalen may partly result from autogenic drivers
364 of peatland development. Plant autogenic succession is also an important factor to be
365 considered when examining peatland development (Swindles *et al.*, 2012, Loisel and Yu,
366 2013; Gałka *et al.*, 2016). The lack of a clear hydrological signal at Marooned peatland during
367 the LIA could be explained by the dominance of *S. fuscum* which has a wide ecological
368 tolerance across a range of hydrological conditions (Rydin and McDonald, 1985).
369 Furthermore, plant assemblages with dominant *S. fuscum* are most usually characterised by
370 testate amoebae indicative of dry habitats, especially when hummocks have developed.
371 Therefore, the peatland may have been climatically insensitive at this time owing to the
372 dominance of *S. fuscum*. Another reason for the lack of visible hydrological changes during
373 LIA may be the very low rates of peat accumulation at this time in Marooned.
374

375 3. Recent warming

376 The hydrological similarity between the two sites is clear in the context of the last few
377 decades. At both sites there is a trend toward drier conditions (lowering water table) (Fig. 6,
378 upper part of zone E and Fig. 7, upper part of zone D). At Marooned this deepening of the
379 water table coincides with the development of shrub communities including *Betula nana* and
380 *C. vulgaris*, and in the case of Stordalen, this drying corresponds with the presence of
381 *Polytrichum commune* and a significant increase of *A. polifolia*. The trend to dry hydrological
382 conditions marked by the appearance of *P. strictum* and *D. elongatum* has been documented
383 in previous studies at Stordalen (Kokfelt *et al.*, 2010). The increasing role of shrub
384 communities in the Abisko region may be also have been caused by modern climate warming,
385 which has facilitated the expansion of shrubs in the Arctic zone (Myers-Smith *et al.*, 2011;
386 Loranty and Goetz, 2012).

387

388 4. Carbon accumulation and C/N ratio

389 The CA rates here need to be interpreted with some caution, as no correction for incomplete
390 decomposition was applied. However, the data suggest that Stordalen has the highest rates of
391 apparent CA in zones B and C (500-150 cal yr BP, Fig. 7). CA accumulation rate increased
392 from ca. 500-300 cal yr BP in Stordalen (Zone B). In Marooned CA is lowest in zones B and
393 C (2300-100 cal yr BP) and higher in the other parts of the peat profile. The Marooned site
394 switched from a period of very slow CA at 200 cal yr BP which also mirrors an increase in
395 organic content (Fig. 6). In contrast, Stordalen has been closer to steady state despite marked
396 vegetation changes over time (Fig. 7). However, the shift from a sedge to *Sphagnum*
397 dominated system at ~140 cal yr BP led to a moderate decline in CA.

398 In deeper layers of peat profile in Marooned and Stordalen C/N ratios are low (Fig. 6
399 and 7), indicating strong microbial transformation of the herbaceous peat (Kuhry and Vitt,

400 1996; Krüger *et al.*, 2015). An increase of C/N ratios in upper part of both sites correlated
401 with the spread of *Sphagnum* species. The range of values are considered typical for
402 ombrotrophic peatlands (Kuhry and Vitt, 1996; Malmer and Wallén, 2004) and indicate
403 relatively low levels of decomposition. In Marooned a decrease of C/N ratio in the last two
404 decades took place (Fig. 6), what is associated with increase of peat decomposition and loss of
405 C over N during microbial decomposition (e.g. Krüger *et al.*, 2015).

406

407 5. Palaeoecology of *Sphagnum lindbergii*

408 *Sphagnum lindbergii* is a circumpolar species which occurs in oligotrophic to
409 mesotrophic habitats from pools to lawns in bogs and poor fens, often bordering nutrient-poor
410 flushes and lakes located in Arctic, sub-Arctic and boreal zones (Laine *et al.*, 2011) and in
411 high-altitude peatlands further south in mountain areas (Lemly *et al.*, 2007; Hölzer, 2010;
412 Wojtuń *et al.*, 2013). It grows together with *S. riparium*, *S. balticum*, *S. flexuosus*, and *S.*
413 *majus* (Hölzer, 2010; Laine *et al.*, 2011). In Stordalen, *S. lindbergii* grew together with
414 *Drepanocladus* sp., *Andromeda polifolia* and *Carex* ssp. for ca. 50 years and disappeared with
415 the appearance and spread of *S. balticum*. The depth of the water table during the presence of
416 *S. lindbergii* was ca. 10 cm (Fig. 7), indicative of a wet poor fen environment. In the *S.*
417 *lindbergii* population dominant species of testate amoebae were *Archerella flavum*, *Placocista*
418 *spinosa* type, *Hyalosphenia papilio*, and *Nebela tinctoria*, what indicates mostly usually wet
419 habitats and a relatively wide range of pH (e.g. 3.0-6.5 in Swindles *et al.*, 2015b).

420 In previous palaeoecological studies conducted in the Stordalen peatland (Kokfelt *et*
421 *al.*, 2010), *S. lindbergii* was found alongside *Drepanocladus exannulatus* and sedges and from
422 ca. 700 to 120 cal yr BP it was the dominant species in the plant assemblage. In others sites in
423 northern hemisphere, e.g. the Hudson Bay Lowlands, Alaska, and in the Sudety Mts. *S.*
424 *lindbergii* macrofossils during a wet poor-fen stage with *S. jensseni/balticum*, *S.*

425 *magellanicum*, *S. riparium*, *S. balticum*, *S. russowii*, *Eriophorum angustifolium*, and *P.*
426 *strictum* was found (Kuhry, 2008; Jones *et al.*, 2013; Lemus-Lauzon *et al.*, 2016; Gałka and
427 Fiałkiewicz–Kozieł, unpubl. data).

428 Based on the fossil data outlined here, it may be concluded that the ecological
429 requirements and accompanying plants associated with *S. lindbergii* has not changed
430 significantly over time when compared with the habitat identified from present data. The
431 presence of *S. lindbergii* macrofossils in a relatively thin (usually, several cm thick) peat layer
432 suggest that this species is not very competitive and mainly plays a role as a transitional
433 species during fen to bog transition. Therefore, in palaeoecological reconstructions the
434 presence of *S. lindbergii* is likely to indicate wet phases in peatland development, which most
435 likely coincide with wet climatic conditions. However, the role of autogenic processes during
436 *Sphagnum* succession should also be considered during reconstructions, with presence of *S.*
437 *lindbergii* in the transition between fen and bog.

438

439 6. Improvement of tephra dispersal maps in Scandinavia

440 The peatlands included in this study contain a total of three tephra layers, all of Icelandic
441 origin and produced by different eruptions of the same volcano (Hekla). They have all been
442 previously detected in northern Europe. The only tephra layer identified in the top 25 cm of
443 peat at Stordalen was SD-1, correlated to the eruption of Hekla 1158. Until recently, Hekla
444 1158 had been reported at only one site in northern Europe (Pilcher *et al.*, 2005). However,
445 this discovery and recent identifications of the Hekla 1158 tephra at two other sites in
446 northern Sweden (Watson *et al.*, 2016) indicate that the Hekla 1158 tephra could provide an
447 important dating isochron in northern Scandinavia.

448 Despite the close proximity of the Marooned and Stordalen peatlands, tephra shards
449 from the eruption of Hekla 1158 were not detected in the Marooned peatland. However, two

450 tephra layers correlated to the eruptions of Hekla-S (=MR-1) and Hekla 4 (=MR-2) were
451 identified. The identification of MR-1 represents the most northerly occurrence of the Hekla-S
452 tephra which has been identified at ~20 sites in northern Europe (Lawson *et al.*, 2012). MR-2
453 contains glass shards matching the geochemistry of glass from the most widespread tephra
454 deposit in northern Europe, Hekla 4 (4287 cal yr BP).

455

456 **Summary**

457 We present a multi-proxy palaeoecological dataset from two peat profiles from Abisko, Arctic
458 Sweden and conclude the following:

- 459 1. Local plant succession and hydrological changes in peatlands were synchronous with
460 climatic shifts, although autogeneous plant succession during peatland development towards
461 ombrotrophic status were also influential.
- 462 2. We observe a fall in water tables and an increase in shrub pollen during the last few
463 decades, coinciding with recent climate warming and associated with expansion of shrub
464 communities noted in various parts of the Arctic zone.
- 465 3. Rates of carbon accumulation vary greatly between the sites and in time, illustrating the
466 importance of local vegetation communities, hydrology and permafrost dynamics.
- 467 4. Our paleoecological data of *Sphagnum lindbergii* supported by its recent ecological
468 requirements and its other fossils finding in various part of world prove that *S. lindbergii* can
469 be indicative of wet conditions at the peatland's surface.
- 470 5. Three tephra layers (Hekla 1158, Hekla-S, and Hekla-4) of Icelandic origin, produced by
471 different eruptions of Hekla were recorded.

472

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490

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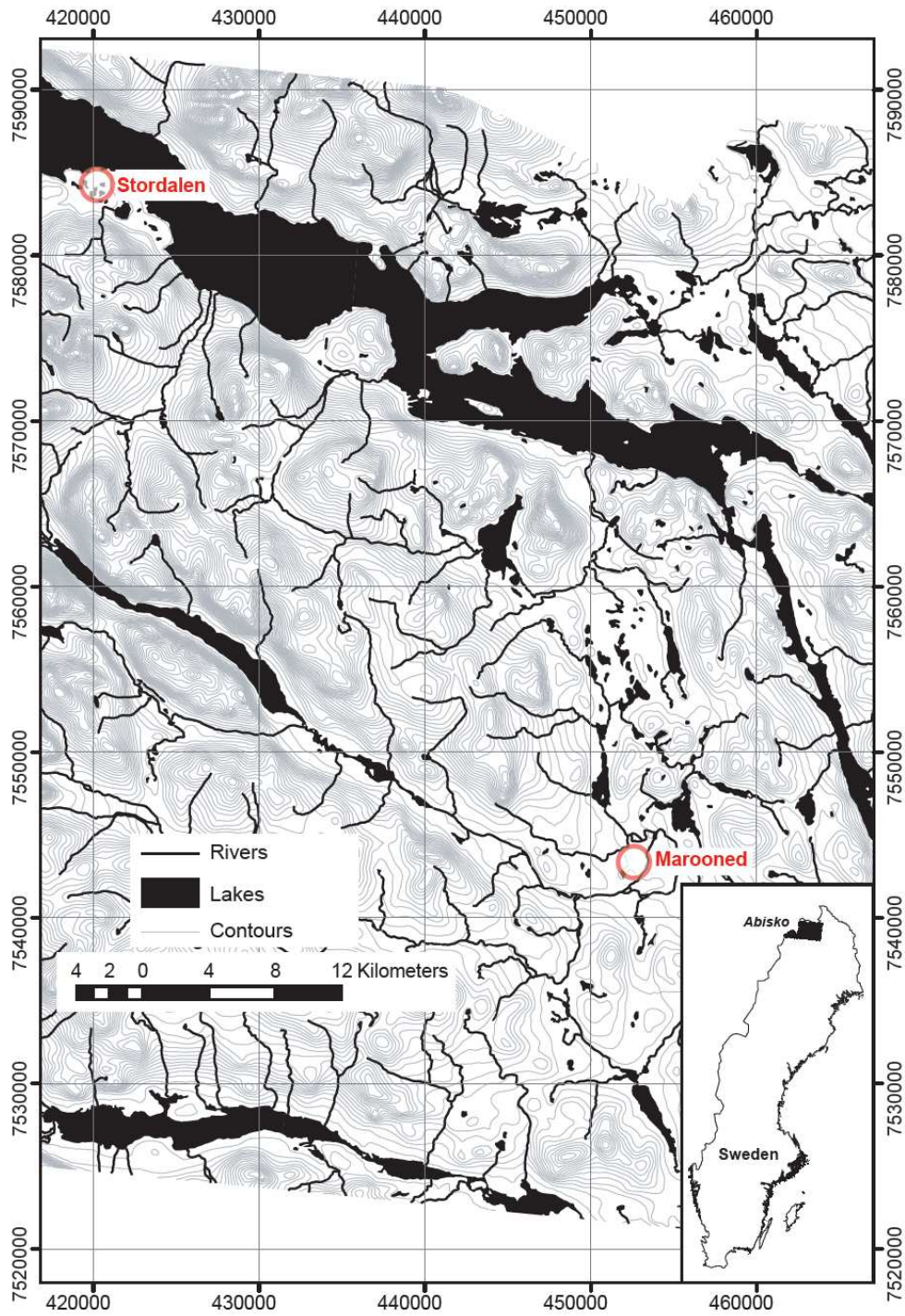
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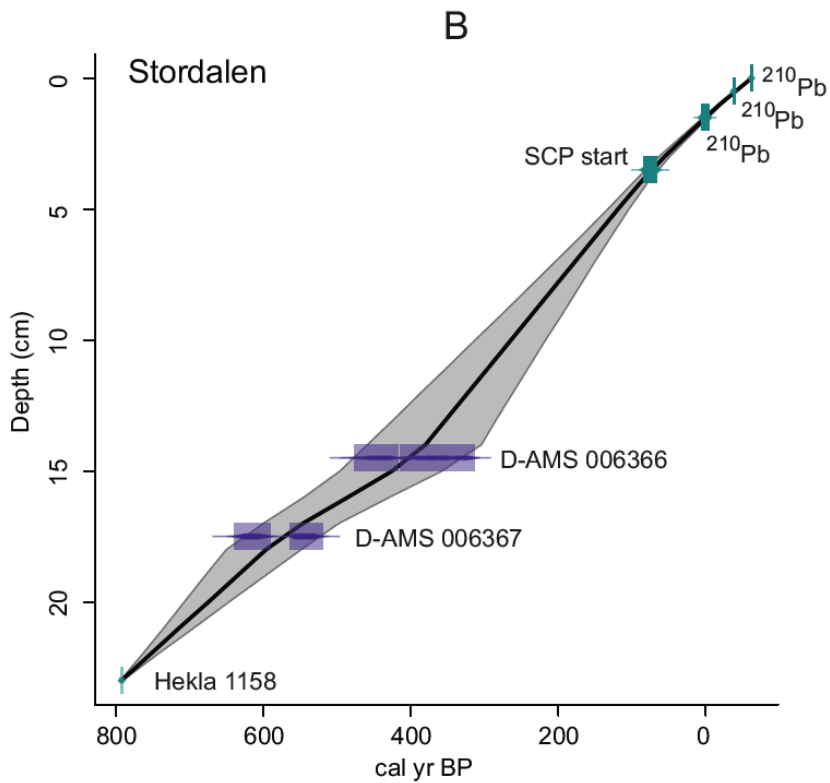
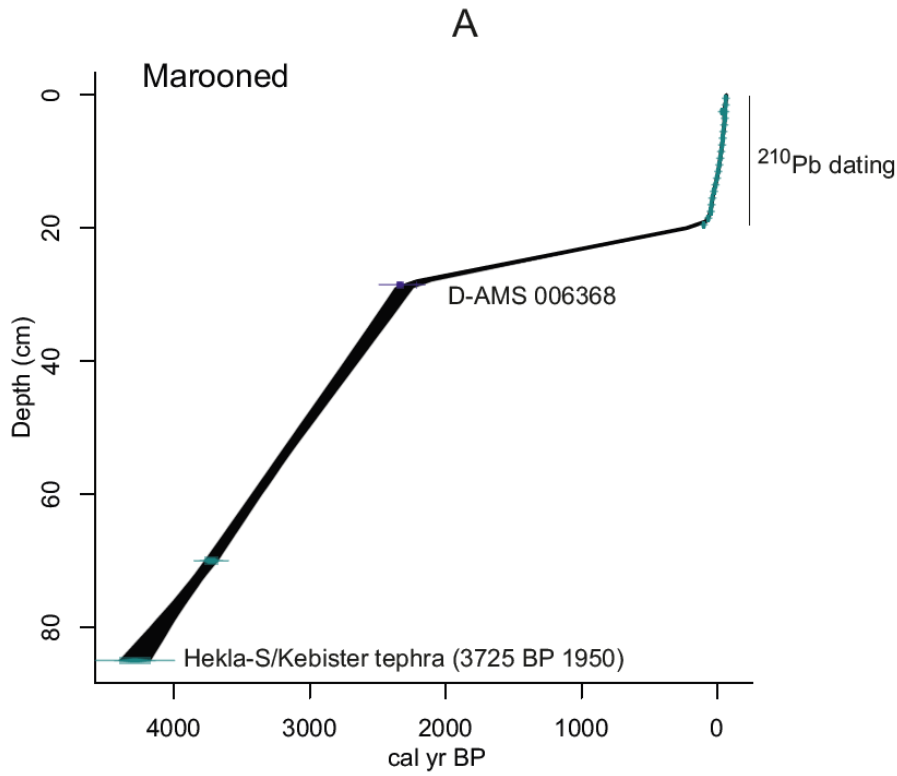
742 **Figure caption**



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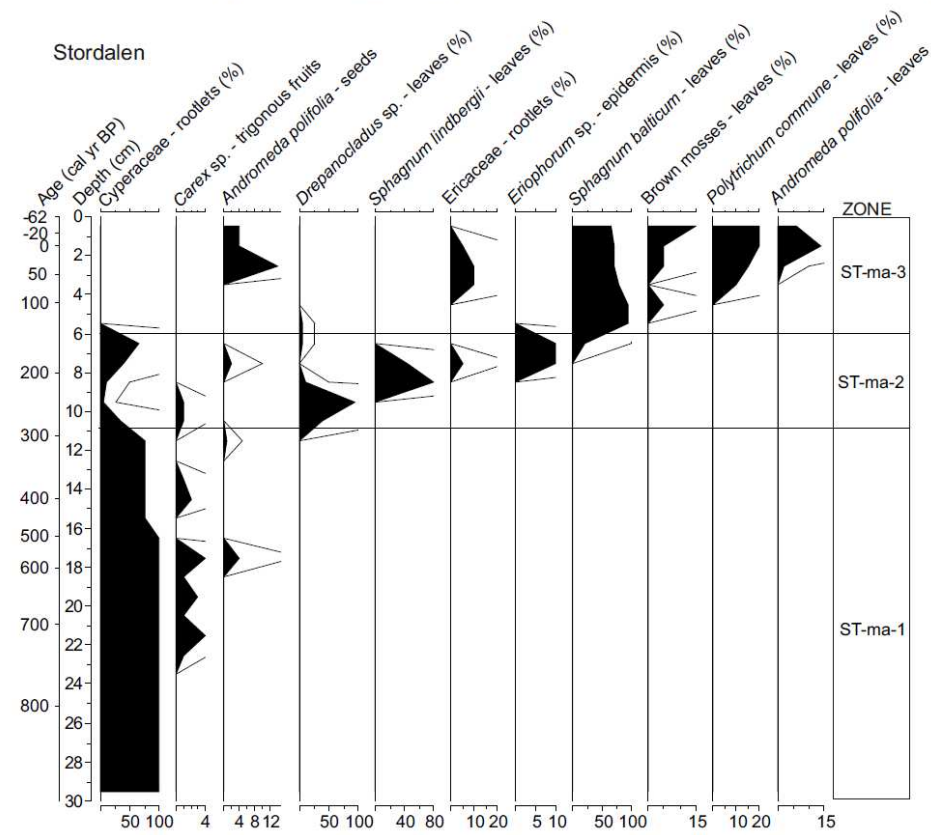
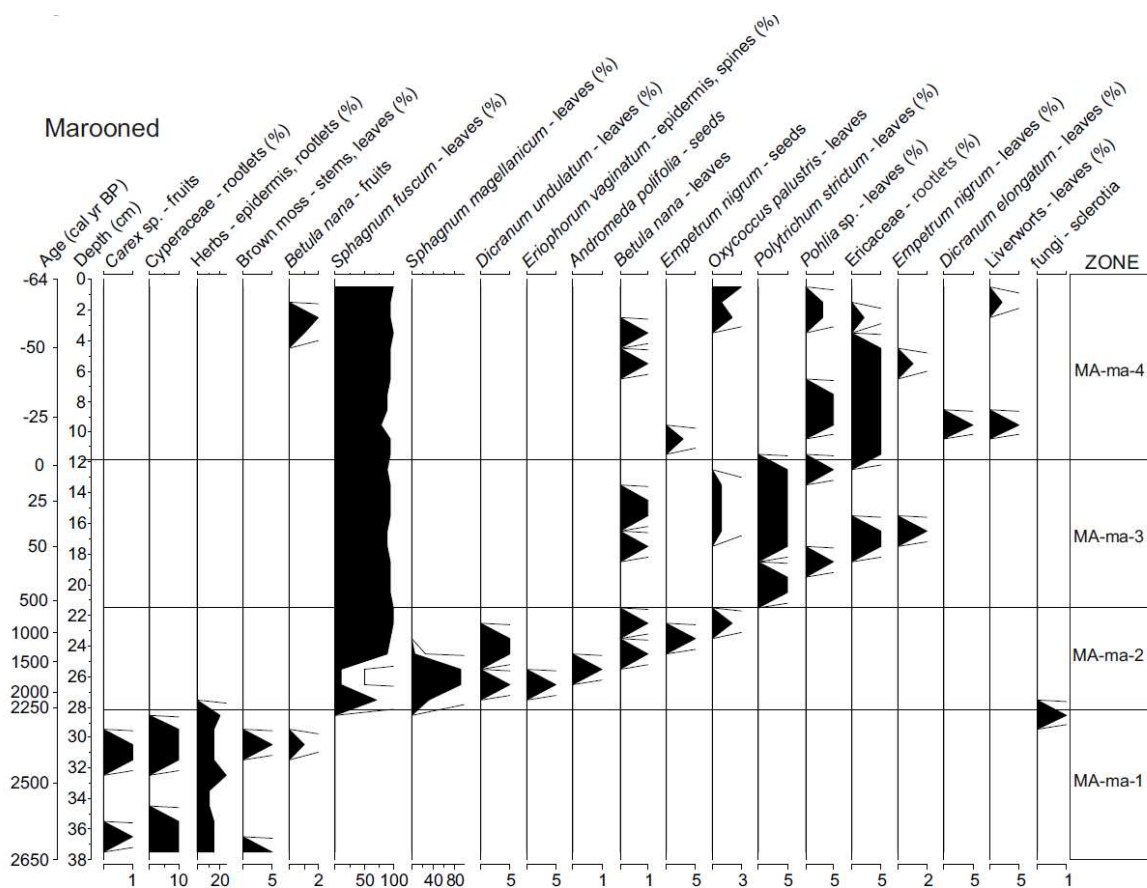
744 Figure 1. Study site.

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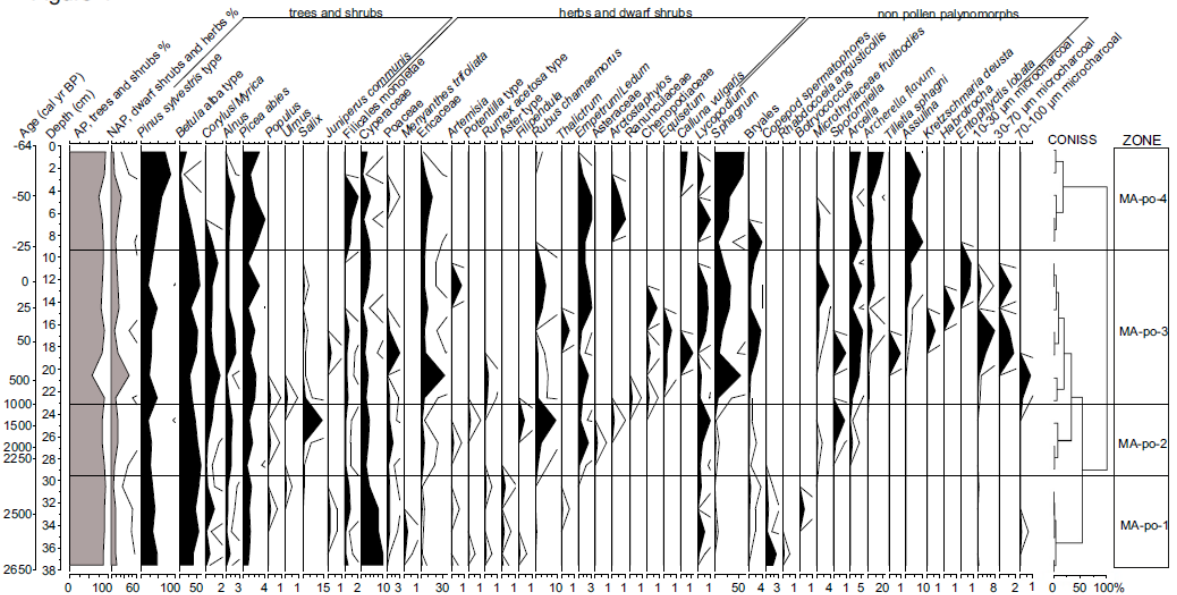
746

747 Figure 2. Age-depth model of the peat profile from Marooned and Stordalen peatlands.



748
 749 Figure 3. Percentage plant macrofossils diagram presenting local vegetation development in
 750 Marooned and Stordalen peatlands.

Figure 4



751

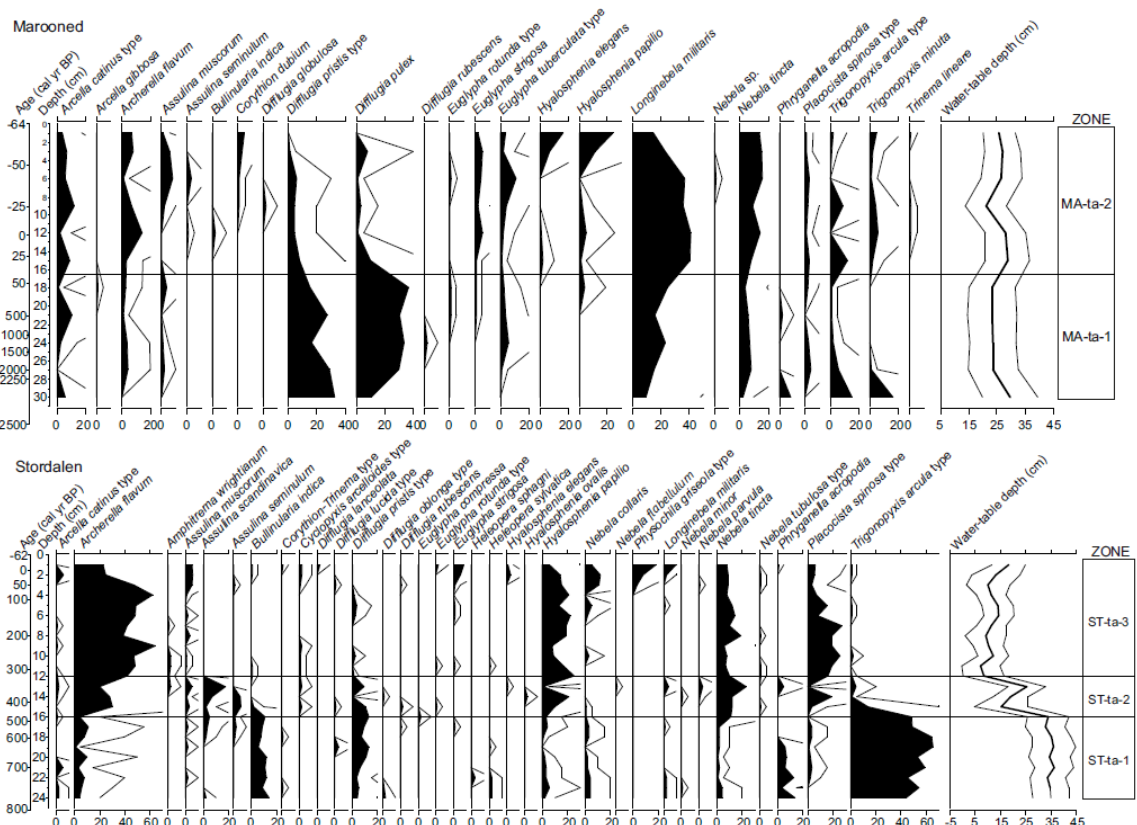
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Figure 4. Percentage pollen diagram presenting local and regional vegetation changes in the

753

Marooned peatland.

Figure 5



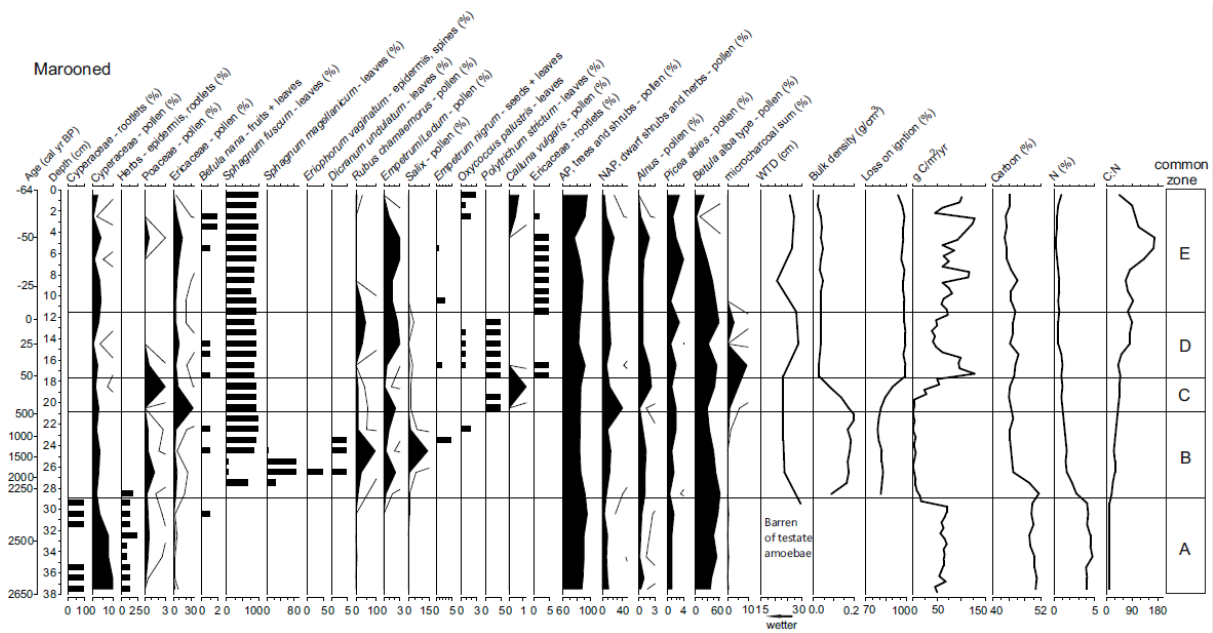
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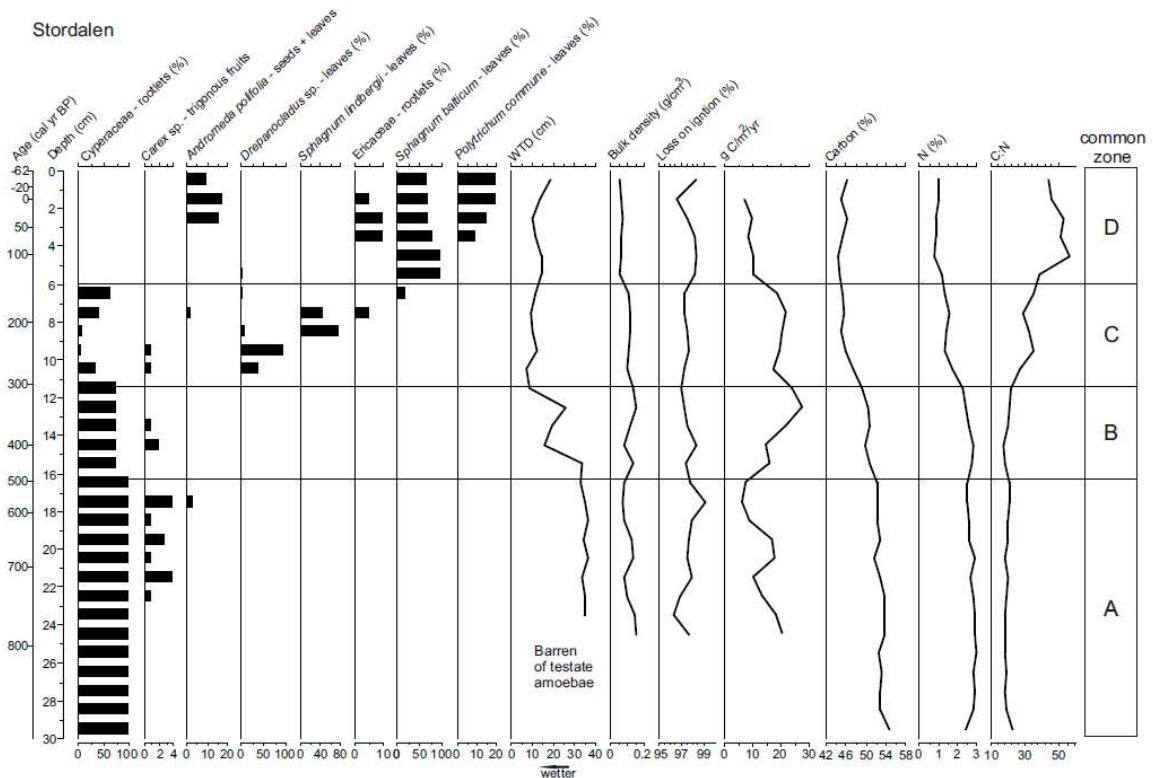
Figure 5. Percentage testate amoebae diagram presenting changes in testate amoebae

756

communities and water table changes in Marooned and Stordalen peatlands.



757
 758 Figure 6. Comparison of chosen taxa from three: plant macrofossils, pollen, and testate
 759 amoebae with the quantitative reconstruction of water-table depth (DWT) and results of
 760 geochemical analysis in Marooned peatland.



761
 762 Figure 7. Comparison of chosen taxa from two data set: plant macrofossils and testate
 763 amoebae with the quantitative reconstruction of water-table depth (DWT) and results of
 764 geochemical analysis in Stordalen peatland.

765 **Table**

766 Table 1. Lithostratigraphic description of the sediment sequence

Site	Description of peats
Marooned	38-28 cm herbaceous-moss peat 28-0 cm <i>Sphagnum</i> – brown moss peat
Stordalen	30-11 cm herbaceous-moss peat 11-6 cm <i>Sphagnum</i> - brown moss-herbaceous peat 6-0 cm fresh <i>Sphagnum</i> peat

767

768

769

770 **Appendix caption**

771 1. Analytical setup of tephra EPMA analysis.

772 Appendix 1

773 Analytical setup of tephra geochemical analysis.

774

775 EPMA: Cameca SX100 EPMA housed at the University of Edinburgh

776 Geochemical analysis was conducted using a Cameca SX100 electron probe microanalyser at

777 Tephra Analytical Unit (TAU), School of Geosciences, University of Edinburgh. All analyses

778 were conducted with a beam diameter of 5 µm, 15kV and beam currents of 2 nA (Na, Mg, Al,

779 Si, K, Ca, Fe) and 80 nA (P, Ti, Mn) (Hayward, 2012). Secondary glass standards (basalt

780 (BCR-2G) and rhyolite (Lipari)), were analysed before and after unknown tephra samples.

781 Marooned

782 EPMA: JEOL JXA8230 EPMA housed at the University of Leeds

783 Geochemical analysis was conducted using a beam diameter to 10 µm, 15 kV and a beam

784 current of and a beam current of 10 nA. Secondary glass standards (Lipari) were analysed

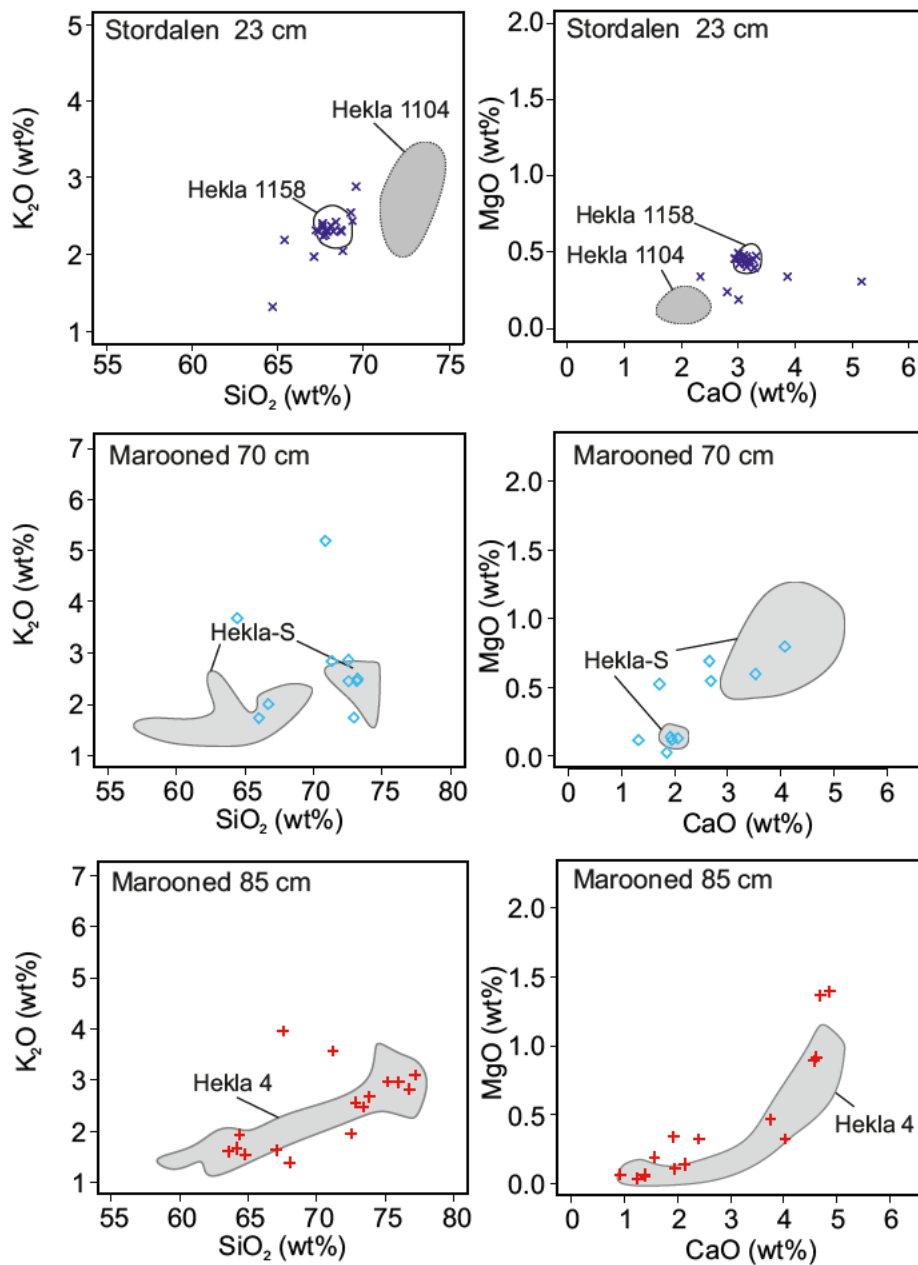
785 before and after unknown tephra samples.

786

787 2. Radiocarbon dates and ²¹⁰Pb measurements: S – Stordalen, M – Marooned

788 Appendix 2. Radiocarbon dates and ²¹⁰Pb measurements: S – Stordalen, M - Marooned

Site/Depth (cm)	Lab. No.	Age ¹⁴ C date	Calibrated range 95.4%	Dated material
S14-15	D-AMS 006366	340±24	477-314	<i>Sphagnum</i>
S17-18	D-AMS 006367	553±31	640-518	<i>Sphagnum</i>
M28-29	D-AMS 006368	2317±26	2360-2211	<i>Sphagnum</i> , herb epidermis
210Pb measurements				
S,1	²¹⁰ Pb	1988	1	Bulk peat
S,3	²¹⁰ Pb	1949	3	Bulk peat
S,5	SCP start	1875	5	Bulk peat
M0,5	²¹⁰ Pb	2012,61	0,01	Bulk peat
M1,5	²¹⁰ Pb	2011,12	0,06	Bulk peat
M2,5	²¹⁰ Pb	2008,60	0,09	Bulk peat
M3,5	²¹⁰ Pb	2005,03	0,18	Bulk peat
M4,5	²¹⁰ Pb	2000,70	0,28	Bulk peat
M5,5	²¹⁰ Pb	1994,52	0,41	Bulk peat
M6,5	²¹⁰ Pb	1989,59	0,69	Bulk peat
M7,5	²¹⁰ Pb	1985,03	0,48	Bulk peat
M8,5	²¹⁰ Pb	1978,01	0,53	Bulk peat
M9,5	²¹⁰ Pb	1971,48	0,67	Bulk peat
M10,5	²¹⁰ Pb	1964,21	0,69	Bulk peat
M11,5	²¹⁰ Pb	1956,38	0,90	Bulk peat
M12,5	²¹⁰ Pb	1946,70	0,96	Bulk peat
M13,5	²¹⁰ Pb	1936,20	0,87	Bulk peat
M14,5	²¹⁰ Pb	1925,00	0,90	Bulk peat
M15,5	²¹⁰ Pb	1913,56	0,97	Bulk peat
M16,5	²¹⁰ Pb	1907,17	0,81	Bulk peat
M17,5	²¹⁰ Pb	1901,37	0,99	Bulk peat
M18,5	²¹⁰ Pb	1883,16	0,70	Bulk peat
M19,5	²¹⁰ Pb	1849,26	0,73	Bulk peat



789

790 Appendix 3. Geochemical bi-plots of major elements of tephra shards (glass) from Stordalen
 791 and Marooned plotted against envelopes for the glass geochemistry of known tephras based
 792 on type data from the TephraBase database. All data have been normalised.

793