



Hässeldala – a key site for Last Termination climate events in northern Europe

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2 northern Europe

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13 The Last Termination (19 000-11 000 a BP) with its rapid and distinct climate
14 shifts provides a perfect laboratory to study the nature and regional impact of
15 climate variability. The sedimentary succession from the ancient lake at
16 Hässeldala Port in southern Sweden with its distinct Lateglacial/early Holocene
17 stratigraphy (>14.1-9.5 cal. ka BP) is one of the few chronologically well-
18 constrained, multi-proxy sites in Europe that capture a variety of local and
19 regional climatic and environmental signals. Here we present Hässeldala's multi-
20 proxy records (lithology, geochemistry, pollen, diatoms, chironomids,
21 biomarkers, hydrogen isotopes) in a refined age model and place the observed
22 changes in lake status, catchment vegetation, summer temperatures and
23 hydroclimate in a wider regional context. Reconstructed mean July temperatures
24 increased between ~14.1 and ~13.1 cal. ka BP and subsequently declined. This
25 latter cooling coincided with drier hydroclimatic conditions that were likely
26 associated with a freshening of the Nordic Seas and started a few hundred years
27 before the onset of Greenland Stadial 1 (~12.9 cal. ka BP). Our proxies suggest a
28 further shift towards colder and drier conditions as late as ~12.7 cal. ka BP,
29 which was followed by the establishment of a stadial climate regime (~12.5-11.8
30 cal. ka BP). The onset of warmer and wetter conditions led the Holocene
31 warming over Greenland by ~200 years. Hässeldala's proxies thus highlight the
32 complexity of environmental and hydrological responses across abrupt climate
33 transitions in northern Europe.

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46 The transition period from the Last Glacial Maximum into the present
47 Interglacial (the Holocene) occurred between 19 and 11 thousand years (ka)
48 before present (BP) and is informally known as the 'Last Deglaciation' or the
49 'Last Termination'. This interval is characterised by a series of abrupt climate
50 shifts of colder (stadial) and warmer (interstadial) states, which are most
51 prominently expressed in the North Atlantic region (Björck *et al.* 1996; Lowe *et*
52 *al.* 2008; Steffensen *et al.* 2008). These stadials and interstadials are often
53 referred to as 'Oldest Dryas' (>14.7 ka BP), 'Bølling' (14.7-14.1 ka BP), 'Older
54 Dryas' (14.1-13.9 ka BP), 'Allerød' (14.1-12.9 ka BP) and 'Younger Dryas' (12.9-
55 11.7 ka BP) (Rasmussen *et al.* 2014) in reference to earlier pollen-stratigraphic
56 work in Scandinavia (see e.g. Mangerud *et al.* 1974; Wohlfarth 1996).

57 Greenland ice core records have played a pivotal role in understanding
58 the underlying causes of abrupt deglacial climate variability, because they (i)
59 record climate shifts in a multitude of atmospheric proxies (Steffensen *et al.*
60 2008); (ii) provide a precise chronological template for past climatic events
61 (Rasmussen *et al.* 2014); and (iii) can be synchronized to Antarctic ice cores
62 using methane measurements (Blunier & Brook 2001). This synchronization
63 allows for a direct correlation of Northern and Southern Hemisphere climatic
64 changes and supports the hypothesis of the Atlantic bipolar seesaw concept
65 (Broecker 1998), which attributes a large role to the Atlantic Meridional
66 Overturning Circulation in triggering abrupt climate shifts (Stocker & Johnsen
67 2003; Knutti *et al.* 2004; EPICA 2006). The INTIMATE working group thus
68 proposed to use the Greenland ice core isotope stratigraphy as a template

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3 69 against which marine and terrestrial records could be compared to decipher
4 70 leads and lags between different regions (Björck *et al.* 1998; Walker *et al.* 1999).
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6 71 Unfortunately, rather than using independent chronological information to
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8 72 enable a critical appraisal of event sequences, many simply wiggle-matched their
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10 73 respective proxy stratigraphies to the Greenland ice core oxygen isotope curve.
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12 74 Lowe *et al.* (2008) and later Blockley *et al.* (2012) and Rasmussen *et al.* (2014)
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14 75 therefore refined the INTIMATE event stratigraphy by including cryptotephra
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16 76 layers that can be found in marine, ice core and terrestrial records and used as
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18 77 time-synchronous marker horizons. This latter approach has clear advantages as
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20 78 compared with simple wiggle-matching, but it soon became clear that a reliance
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22 79 only on cryptotephra layers to correlate between sites can be misleading as the
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24 80 ash may be reworked and/or originate from multiple eruptions of the same
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26 81 volcano with similar geochemical signatures (see e.g. Davies *et al.* 2004; Pyne-
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28 82 O'Donnell *et al.* 2008, 2010).

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30 83 Although considerable advances have been made with respect to
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32 84 cryptotephra studies (Lowe *et al.* 2015) and radiocarbon calibration (Reimer *et*
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34 85 *al.* 2013), the lack of high-resolution, independent chronologies for terrestrial
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36 86 and marine records still makes it impossible to compare the exact regional and
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38 87 hemispheric phasing of Last Termination climatic events, albeit with a few
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40 88 exceptions (Lane *et al.* 2013; Lohne *et al.* 2013, 2014; Muschitiello & Wohlfarth
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42 89 2015; Rach *et al.* 2014; Muschitiello *et al.* 2015b). Moreover, the detection of
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44 90 spatiotemporal differences of up to several hundred years in respect to
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46 91 temperature, precipitation, and vegetation changes both at the start of (Rach *et*
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48 92 *al.* 2014; Muschitiello & Wohlfarth 2015) and during Younger Dryas (Lane *et al.*
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50 93 2013), suggests that the propagation of climatic signals on regional to sub-
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52 94 continental scales is still not fully understood. The assumption of more or less
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54 95 synchronous hydroclimatic and ecosystem responses even on regional scales is
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56 96 therefore clearly an oversimplification.

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58 97 While chronological uncertainties surely play a major role when
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60 98 attempting to correlate between different archives (Blaauw 2012), the detection
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100 99 of past environmental and climatic shifts in various terrestrial archives also very
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102 100 much depends on (i) the choice of biological, chemical and physical proxies to be
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104 101 studied and their respective environmental and climatic information (e.g.,

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3 102 seasonal temperature, precipitation, windiness, run-off, aquatic and terrestrial
4 103 vegetation, lake status changes); (ii) understanding the complicated ecological
5 104 processes that drive biotic and abiotic changes (e.g. Jackson & Overpeck 2000;
6 105 Ammann *et al.* 2013; Birks & Birks 2014) and lake development (Engstrom *et al.*
7 106 2000); and (iii) the transfer functions and other methods used to derive
8 107 temperature estimates from micro- and macrofossil data sets (e.g. Juggins 2013;
9 108 Brooks & Langdon 2014).

109 The Last Termination, with its rapid and distinct climate shifts, is an
110 excellent time period to study the nature and regional impact of climate
111 variability. Few lake sedimentary successions in Europe however fulfil the
112 requirement of being chronologically well-constrained, multi-proxy sites that are
113 able to capture the variety of climatic and environmental signals preserved
114 within the fossil record and enable unequivocal comparisons to the Greenland
115 ice core template chronology (Ammann *et al.* 2013; Birks & Birks 2014; Rach *et*
116 *al.* 2014; Muschitiello & Wohlfarth 2015). One of these is the sedimentary
117 succession from the ancient lake at Hässeldala Port in southern Sweden (Fig 1A-
118 C), which displays a distinct Lateglacial stratigraphy (Davies *et al.* 2004). Since
119 the discovery of several cryptotephra layers in Hässeldala Port's sediments
120 (Davies *et al.* 2003), the site has been the focus of a variety of proxy studies, most
121 of which were performed on various parallel sediment cores (Davies *et al.* 2004;
122 Wohlfarth *et al.* 2006; Watson 2008; Kylander *et al.* 2013; Steinhorsdottir *et al.*
123 2013, 2014; Ampel *et al.* 2014; Muschitiello *et al.* 2015a, b; Karlatou-
124 Charalampopoulou 2016) (Fig. 1C, Table 1). Except for the most recent studies
125 (Steinhorsdottir *et al.* 2013; Ampel *et al.* 2014; Muschitiello *et al.* 2015b;
126 Karlatou-Charalampopoulou 2016), the chronology of core #2 and visual core-
127 to-core correlations were used (Davies *et al.* 2004; Wohlfarth *et al.* 2006; Watson
128 2008; Kylander *et al.* 2013). The latter approach was justified because total
129 organic carbon (TOC) and/or loss-on-ignition (LOI) measurements for each
130 analysed profile show the same characteristic pattern and allowed for a direct
131 correlation between parallel sediment cores and their lithostratigraphic units
132 (Wohlfarth *et al.* 2006).

133 Alignment of the various parallel profiles using a Monte Carlo based
134 approach and the most recent and updated age model for core #5 (Muschitiello

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3 135 *et al.* 2015b) now offer the possibility to present the suite of proxy records for
4 136 Hässeldala Port on one common time scale and to discuss local and regional
5 137 responses to Last Termination climate events. The different proxies analysed in
6 138 Hässeldala Port's sedimentary profiles have been described in detail in various
7 139 publications (see Table 1 for references), but except for the chironomid-based
8 140 temperature reconstruction and the biomarker data set, which have been
9 141 presented on an updated chronology (Muschitiello *et al.* 2015a, b), none of the
10 142 other proxies has been discussed within the context of the new chronology for
11 143 core #5. Here we combine Hässeldala Port's various sediment cores and proxies
12 144 and present these on the chronology for core #5 to discuss the timing and impact
13 145 of Lateglacial climatic shifts in southern Sweden and to address leads and lags
14 146 between changes in hydroclimate and vegetation across Europe.
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25 148 Hässeldala Port - background

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28 149 The ancient lake that existed during the Last Termination at Hässeldala Port
29 150 (56°16' N, 15°01' E; 63 m a.s.l.) in Blekinge province, southern Sweden (Fig. 1A-
30 151 C), filled in during the early Holocene. Today the site is a peat bog that is partly
31 152 covered by a mixed pine-birch forest, shrubs (*Juniperus*, *Ericales*), grass, mosses
32 153 and ferns. Peat has been excavated extensively during the first half of the 20th
33 154 century and Lateglacial sediments can therefore be found in a depth of less than
34 155 three meters. This, and the easy access to the site, has led to several coring
35 156 campaigns during the past fifteen years. Coring was always performed using a
36 157 Russian corer of 1 m length and 10 cm diameter.
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44 158 Since Hässeldala Port's Lateglacial and early Holocene sediment
45 159 succession is contained within one meter of sediment, it precisely fits the length
46 160 of the coring chamber and can be recovered without the need of overlaps. The
47 161 deepest part of the basin was found to be in the south-western corner of the bog.
48 162 It is from this restricted area, where all analysed sediment cores have been
49 163 obtained (Fig. 1C, Table 1). The exact GPS location of each coring point has
50 164 however not been recorded and lithostratigraphic transects combining all
51 165 studied profiles are therefore not possible, other than those shown in Wohlfarth
52 166 *et al.* (2006).
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3 167 The first to acknowledge the potential of Hässeldala Port's Lateglacial
4 168 sediments were Davies *et al.* (2003), who described five different cryptotephra
5 169 horizons from core #1 (Fig. 1C, Tables 1, 2). Of these three were correlated to
6 170 known eruptions based on their geochemical signatures: the Borrobol tephra
7 171 (BT), the newly defined Hässeldala tephra (HDT) and the Askja-S tephra (AsT)
8 172 (Table 2). Two tephras could not be correlated to any known eruption. Using the
9 173 pattern of the loss-on-ignition (LOI) curve of core #1, Davies *et al.* (2003)
10 174 tentatively assigned climatic/pollen stratigraphic periods and suggested that the
11 175 BT would date to the Bølling-Allerød Interstadial and the HDT and the AsT to the
12 176 early Holocene. However, the finding of two superimposed cryptotephras in
13 177 Lateglacial sediments in Scotland, which both seem to have been derived from
14 178 the same volcanic source, led Matthews *et al.* (2011) to suggest that Hässeldala
15 179 Port's BT is younger than the BT found in Scotland and likely an equivalent to the
16 180 Penifiler tephra.

17 181 Subsequent analyses were made on cores #2 (LOI, ^{14}C chronology) and #3
18 182 (TOC, cryptotephra, pollen stratigraphy) (Fig. 1C, Tables 1, 2) (Davies *et al.* 2004;
19 183 Andersson 2004; Wohlfarth *et al.* 2006). Both profiles showed an identical
20 184 stratigraphy and a good match between the LOI/TOC curves, which facilitated
21 185 visual inter-core correlations. The different age models established for core #2
22 186 were therefore transferred to core #3 and provided age estimates for the three
23 187 geochemically identified cryptotephra layers of 14.3-13.7 cal. ka BP ('BT'), 11.6-
24 188 11.2 cal. ka BP (HDT) and 11.1-10.6 cal. ka BP (AsT) (Wohlfarth *et al.* 2006). The
25 189 reconstructed vegetation development showed that arctic and sub-arctic plant
26 190 species dominated during the Lateglacial and that *Pinus sylvestris* and *Betula*
27 191 *pubescens* were present at the transition between the Younger Dryas and
28 192 Preboreal pollen zones (Andersson 2004; Wohlfarth *et al.* 2006). Moreover,
29 193 approximate ages for the regional Lateglacial pollen zone boundaries were
30 194 established using a suite of different age models. These suggested ages of 14.2-
31 195 13.7 cal. ka BP for the Older Dryas/Allerød pollen zone transition, of 13.1-12.6
32 196 cal. ka BP for the Allerød/Younger Dryas transition and of 11.9-11.3 cal. ka BP
33 197 for the Younger Dryas/Preboreal transition (Wohlfarth *et al.* 2006).

34 198 A set of twelve new sediment cores were later obtained by Watson
35 199 (2008), analysed for LOI, cryptotephra, Coleoptera and chironomids (Fig. 1C,

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3 200 Tables 1, 2) and correlated to cores #2 and #3. Six of these profiles were
4 201 subsampled and aggregated for Coleoptera analysis; LOI analyses were
5 202 undertaken on four cores to aid cross-core correlation (#HP4, #HP6, #HP9,
6 203 #HP12) and chironomids were analysed from core #HP4 (Fig. 1C, Table 1).
7 204 Cryptotephra was found in core #HP4 and was tentatively correlated to core #1
8 205 using the pattern of the LOI curve (Watson 2008) (Table 2). The Coleoptera
9 206 assemblages, which represent mean summer air temperatures, suggested mild
10 207 and cool summers until about 13.4 cal. ka BP with reconstructed mean summer
11 208 temperatures of about 14 °C (Watson 2008). At the Younger Dryas/Holocene
12 209 transition, Coleoptera, which are typical of acid bog environments and dry shrub
13 210 heathland such as *Amara alpina* and *Olophrum boreale*, appear suggesting a
14 211 change in lake status. The chironomid assemblages also show clear shifts in
15 212 species composition and point to a shift from oligotrophic to increasingly
16 213 eutrophic lake water conditions (Watson 2008). Moreover, the chironomid
17 214 assemblages allowed reconstructing mean summer surface water temperatures
18 215 between c. 14.2 and 11.8 cal. ka BP in great detail (Watson 2008; Muschitiello *et*
19 216 *al.* 2015b).

20 217 Kylander *et al.* (2013) analysed the geochemistry in core #4 (Fig. 1C,
21 218 Table 1) and, using lithostratigraphic markers and the TOC curve, matched this
22 219 profile to cores #2 and #3 to obtain a general chronostratigraphic framework.
23 220 Higher sediment accumulation rates were reconstructed for the pre-Allerød
24 221 period, while a relative constant sediment supply seems to have characterised
25 222 the Allerød and Younger Dryas pollen zones.

26 223 The most recent investigations focussed on core #5 (Fig. 1C) and
27 224 provided a new chronology and age model, an atmospheric CO₂ reconstruction
28 225 using leaf stomata (Steinhorsdottir *et al.* 2013, 2014), lake status changes based
29 226 on diatom assemblages (Ampel *et al.* 2014) and geochemistry (Muschitiello *et al.*
30 227 2015a) and a hydroclimate reconstruction using lipid biomarkers (Muschitiello
31 228 *et al.* 2015b). These latter analyses were based on hydrogen isotopes derived
32 229 from aquatic (*n*-C₂₁) and terrestrial (*n*-C₂₇₋₂₉₋₃₁) alkanes and can be used as
33 230 proxies for moisture source composition and terrestrial evaporation
34 231 (Muschitiello *et al.* 2015b) (Table 1). All sub-samples were taken in 1 cm
35 232 increments along core #5 and split to accommodate various proxy analyses.

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3 233 However, to provide enough leaf material for stomatal analyses, several parallel
4 234 cores had to be sub-sampled after precise lithostratigraphic correlation to the
5 235 master core #5 (Steinthorsdottir *et al.* 2013, 2014). Lastly, a new high-resolution
6 236 pollen stratigraphy, combined with LOI measurements, was established on parts
7 237 of core #6 (Muschitiello *et al.* 2015b; Karlatou-Charalampopoulou 2016) (Fig. 1C,
8 238 Table 1).

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13 239 Although the lithostratigraphy and the LOI/TOC curves for the parallel
14 240 profiles facilitated visual core-to-core correlations (Wohlfarth *et al.* 2006), this
15 241 eyeballing exercise did not allow for a precise transfer of proxy data from one
16 242 core to another. Muschitiello *et al.* (2015b) therefore statistically aligned cores
17 243 #2, #HP4, #5 and #6 with each other. This made it possible to update the
18 244 existing chronology for core #5 and provided a possibility to discuss climate
19 245 responses across the Allerød/Younger Dryas pollen zone transition in greater
20 246 detail (Muschitiello *et al.* 2015b).

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28 29 30 248 Lithostratigraphy and core-to-core correlations

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32 249 Hässeldala Port's sediment cores are all derived from the southwestern part of
33 250 the peatbog (Fig. 1C) and display an almost identical lithostratigraphic
34 251 succession, which is also reflected in the respective TOC/LOI curves (Davies *et al.*
35 252 2003, 2004; Wohlfarth *et al.* 2006; Kylander *et al.* 2013; Steinthorsdottir *et al.*
36 253 2013, 2014; Karlatou-Charalampopoulou 2016). Since the bottom topography of
37 254 the bog is not even, the depth of the Lateglacial/early Holocene profiles varies
38 255 slightly, as does the thickness of individual layers (Wohlfarth *et al.* 2006) (Figs 2,
39 256 3A). Despite this, the recovered Lateglacial/early Holocene successions show a
40 257 distinct and repetitive lithostratigraphic pattern (Figs 2, 3A): the bottommost
41 258 sediments in all cores are composed of grey silty sand and clayey silt/silty clay
42 259 (units 1 and 2) with low organic matter content; higher, but variable LOI/TOC
43 260 values characterise the silty clay gyttja and/or clay gyttja layers of units 3 to 6
44 261 (Fig. 2); and the dark brown silty clayey gyttja/clayey gyttja of unit 7 with its
45 262 high LOI/TOC values is distinct in all profiles, as is the overlying greenish-
46 263 brown/brown clayey gyttja of unit 8 with its markedly lower LOI/TOC values
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3 264 (Wohlfarth *et al.* 2006) (Fig. 2). The sediments in the uppermost part are made
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5 265 up of clay gyttja, gyttja and peaty gyttja (units 9-12) with high LOI/TOC values.

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7 266 Lithostratigraphic core-to-core correlations are thus straightforward and
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9 267 are also supported by the very similar pattern of the accompanying LOI/TOC
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11 268 curves (Figs 2, 3A). These display the same characteristic features in all six
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13 269 sequences with low organic contents in the bottommost part, gradually rising,
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15 270 but variable TOC/LOI values, a distinct peak in organic content, a decrease in
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17 271 LOI/TOC values and subsequent high values (Fig. 3A). However visual core-to-
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19 272 core correlations or visual curve matching do not allow transferring proxies
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21 273 from one sediment core to another or projecting the various proxies on the new
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23 274 chronology of core #5. We therefore follow the approach presented in
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25 275 Muschitiello *et al.* (2015a, b), who used a Monte Carlo Markov Chain algorithm
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27 276 for core-to-core alignment. This method provides a suite of possible alignments
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29 277 between stratigraphic series and the target series and estimates the optimal
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31 278 correlation (see Muschitiello *et al.* 2015b for further details). For Hässeldala
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33 279 Port, we use the high-resolution LOI/TOC curves of cores #1-3, HP4 and 6 as
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35 280 stratigraphic series and the LOI curve of core #5 as target series (Fig. 3A). Prior
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37 281 to the statistical analyses, the TOC values of cores #1, 2 and 3 were recalculated
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39 282 to LOI values to facilitate the statistical correlation. The alignment of the LOI
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41 283 curves of cores #1-3, HP4 and 6 to the LOI curve of core #5 then provided a
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43 284 means to transfer the plant macrofossil record and the ¹⁴C dates of core #2, the
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45 285 pollen stratigraphy of core #3 (Wohlfarth *et al.* 2006), the chironomid-based
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47 286 summer temperature reconstruction of core #HP4 (Watson 2008) and the high-
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49 287 resolution pollen stratigraphy of core #6 (Karlou-Chalampopoulou 2016) to
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51 288 the stratigraphy of core #5 (Muschitiello *et al.* 2015b). However, since the
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53 289 chironomid data set of core #HP4 had a higher temporal sampling resolution
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55 290 than samples taken from core #5, the data set had to be interpolated to the
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57 291 sampling resolution of master core #5 (Muschitiello *et al.* 2015b).

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60 293 Age-depth model and chronology

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295 Using the same core-to-core alignment approach as outlined above, Muschitiello

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3 296 *et al.* (2015b) updated the earlier age model for core #5 (Steinthorsdottir *et al.*
4 297 2013, 2014) by transferring the ¹⁴C dates from core #2 to the stratigraphy of
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6 298 core #5 (Table 3, Fig. 4).
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8 299 Age-depth curves were constructed (Muschitiello *et al.* 2015b) using
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10 300 Bacon2.2. (Blaauw & Christen 2011) and OxCal4.2 (Bronk Ramsey 2010) after
11 301 calibration with IntCal13 (Reimer *et al.* 2013). Since both age models provided
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13 302 very similar results, we here use OxCal4.2 modelled weighed mean ages for the
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15 303 respective age-depth assignments (Table 4). Parts of the uppermost and/or
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17 304 lowermost layers in some profiles do not overlap with the length of core #5. The
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19 305 chronology for these parts had therefore to be estimated using the modelled
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21 306 sedimentation rate of the respective lithostratigraphic units (Fig. 3B, Table 4).
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23 307 The new composite age model for core #5 (Muschitiello *et al.* 2015b) compares
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25 308 well with the previous age model (Steinthorsdottir *et al.* 2013, 2014) in terms of
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27 309 shape of the age-depth relationship and age output values. It however provides a
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29 310 higher number of time constraints, especially between 13 and 11 cal. ka BP
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31 311 (Muschitiello *et al.* 2015b).

312 When plotted on the new time scale of core #5, the main features of the
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314 six LOI/TOC curves fall within the same time interval, such as the decrease in
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316 LOI/TOC values at 13 cal. ka BP, their peak between 12.8 and 12.7 cal. ka BP, as
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318 well as the mid-point of their gradual rise at 11.5 cal. ka BP (Fig. 3B). Moreover,
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320 modelled ages differ by less than 150 years for both the 'Borrobol' and Askja-S
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322 tephra and by less than 50 years for the Hässeldala tephra between cores #1 and
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324 3 (Fig. 3B). The good temporal correspondence of the main features in the six
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326 sediment cores gives confidence in the statistical curve alignments and provides
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328 a new chronological basis for discussing the Lateglacial environmental and
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330 climatic history of southern Sweden.
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333 Local and regional environmental and climatic development

334 >14.1 cal. ka BP

335 Glacial varve chronologies, glacial stratigraphy and geomorphology, and pollen
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337 stratigraphic studies in Blekinge province, southernmost Sweden, reconstruct a
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339 rapid decay of the Scandinavian Ice Sheet margin in response to the temperature
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3 328 rise at the start of the Bølling-Allerød interstadial at 14.7 cal. ka BP (Björck &
4 329 Möller 1987; Ringberg 1991; Wohlfarth *et al.* 1994; Lundqvist & Wohlfarth 2001;
5 330 Anjar *et al.* 2013). The combination of ice sheet load, ice sheet melt and isostasy
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7 331 created a large ice lake in the Baltic Sea basin (Björck 1995; Andrén *et al.* 2011),
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9 332 whose highest shoreline reached up to ~65-67 m a.s.l. in Blekinge (Björck 1981;
10 333 Ringberg 1991).

13 334 The local glacial varve chronology shows that coastal areas in southern
14 335 Blekinge had become free of active ice within 100 years (Fig. 5A) and that the
15 336 formation of the highest coastline at 65-67 m a.s.l. likely corresponds to an ice
16 337 marginal position at around local varve year 0 (Ringberg 1991). Rapid land uplift
17 338 followed coastal deglaciation and led to an apparent lowering of the Baltic Ice
18 339 Lake shoreline (Björck 1981; Wohlfarth *et al.* 1994). The deposition of
19 340 exceptionally thick varves, which can be seen in many clay-varve diagrams from
20 341 Blekinge (corresponding to local varve years +88 to +220) (Ringberg 1991;
21 342 Wohlfarth *et al.* 1994), signals accelerated ice sheet melt and likely correlates to
22 343 fluvio-glacial delta surfaces at around 55 m a.s.l. (Björck 1981). More or less
23 344 coinciding with the appearance of the thick varves, the area around Hässeldala
24 345 Port became free of active ice, as seen in the deglaciation ages for sites such as
25 346 Trehörnan, Hemsjön, Kroksjön II and Skälgylet (Fig. 5B, Table 5). Glacial varves
26 347 did however not form at Hässeldala Port since the site was located at much
27 348 shallower depths and close to the shoreline of the Baltic Ice Lake (Fig. 5C). Pollen
28 349 stratigraphic studies of the varved-clay succession of Farslycke (Ising, 1998) and
29 350 Kroksjön II (Björck 1981) (Fig. 5B) and plant macrofossil finds of *Salix polaris*, *S.*
30 351 *reticulata*, *S. herbacea*, *Dryas octopetala*, and *Betula nana* in the thick varves
31 352 (Wohlfarth *et al.* 1994) suggest that herb and shrub vegetation had become
32 353 established on ice-free land areas. Pollen assemblages (Farslycke, Kroksjön) and
33 354 ¹⁴C ages on terrestrial plant remains (Farslycke, Skälgylet) assign the thick
34 355 varves to the Bølling pollen zone (Björck 1981, 1984; Björck & Möller 1987; Ising
35 356 1998; Wohlfarth & Possnert 2000; Lundqvist & Wohlfarth 2001). Varves
36 357 assigned to local varve years +220 to +360 become gradually thinner displaying
37 358 a greater distance to the rapidly melting ice sheet margin, which was now
38 359 located some 50-100 km to the north of Blekinge's present coastline, and drained
39 360 by large overland rivers (Björck & Möller 1987; Möller 1987). Lake isolation

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3 361 studies, combined with pollen stratigraphies, show that land uplift was still rapid
4 362 and that the shoreline of the Baltic Ice Lake was successively lowered to around
5 363 50-55 m a.s.l. at the end of the regional Bølling pollen zone (Figs 5B, C) and to
6 364 around 45-50 m a.s.l. (Fig. 5D) at the end of the regional Older Dryas pollen zone
7 365 (Björck 1981, 1984).

8 366 The gradual emergence of new land areas meant that Hässeldala Port's
9 367 location changed from an archipelago setting during the Bølling pollen zone to
10 368 further inland during the Older Dryas pollen zone (Figs 5C, D). The bottommost
11 369 silty fine sand (unit 1) in Hässeldala Port's successions (Figs 2, 6A) could derive
12 370 from melting of stagnant ice or run-off from poorly vegetated slopes. Coleoptera
13 371 assemblages suggest mild and cool summers during this early period with mean
14 372 summer air temperatures of about 14 °C (Watson 2008) and the presence of
15 373 *Hippophaë rhamnoides* and *Nymphaea alba* in the aquatic pollen record of
16 374 Hässeldala Port (Andersson 2004) indicates minimum mean July temperatures
17 375 of >10 °C and possibly as much as 11-12 °C (Iversen, 1954; Kolstrup 1980). In
18 376 contrast, mean July surface water temperatures reconstructed from chironomid
19 377 assemblages are around 7-8 °C (Fig. 6E) and may represent a lake water
20 378 temperature that was strongly influenced by run-off from stagnant ice. The herb,
21 379 grass, shrub and dwarf-shrub dominated pollen assemblages (HÄP-1)
22 380 (Andersson 2004; Wohlfarth *et al.* 2006) suggest that these lowermost
23 381 sediments were deposited during the regional Older Dryas pollen zone (Figs 6H,
24 382 J).

25 383 Plant macrofossil finds and pollen stratigraphic records in Blekinge
26 384 dating to the regional Older Dryas pollen zone (Berglund 1966; Björck 1981;
27 385 Wohlfarth *et al.* 1994, 2006) reveal a shrub-grass-herb vegetation, that
28 386 comprised *Artemisia*, *Betula nana*, *Calluna vulgaris*, *Caltha palustris*, *Dryas*
29 387 *octopetala*, *Empetrum nigrum*, *Ephedra*, *Helianthemum*, *Hippophaë rhamnoides*,
30 388 *Juniperus communis*, *Plantago*, *Polygonum*, *Rumex*, *Saxifraga*, *Salix polaris*, and
31 389 *Urtica* (Table 6). Pollen of *Populus tremula* (Berglund 1966) and plant
32 390 macrofossil finds of *Betula pubescens* (Wohlfarth *et al.* 1994) (Table 6) moreover
33 391 point to the regional presence of shrub-like trees. Regional minimum mean July
34 392 temperatures of >10 °C for *Betula pubescens* (Iversen 1954) compare well to
35 393 those inferred from Coleoptera species and pollen in Hässeldala's sediments.

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5 395 *14.1 – 12.7 cal. ka BP*

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7 396 Hässeldala's silty clay, gyttja clay and clayey gyttja (units 2 – 7) and the
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9 397 accompanying LOI curve demonstrate a gradually higher organic matter content
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11 398 of the sediments, but also that minerogenic inputs still dominate (Figs 6A, B). LOI
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13 399 displays a first marked increase at 13.7 cal ka BP, coinciding with the start of unit
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15 400 3, and thereafter fluctuates between lower (13.5-13.4 cal. ka BP; 13.05-12.95 cal.
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17 401 ka BP) and higher values (13.7-13.55 cal. ka BP; 13.3-13.1 cal. ka BP; 12.9-12.7
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19 402 cal. ka BP) (Fig. 6B). Accumulation rates seem to have been rather stable
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21 403 throughout this time interval, except for around 14.1-13.7 and 13.0-12.9 cal. ka
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23 404 BP, when sediment accumulation was slightly higher (Kylander *et al.* 2013).

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25 405 The tephra layer that had been assigned to the Borrobol tephra (Davies *et*
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27 406 *al.* 2003, 2004; Wohlfarth *et al.* 2006) (Fig. 6F), but has been discussed as being
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29 407 an equivalent of the Penifiler tephra (Mathews *et al.* 2011) falls within the
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31 408 interval of low LOI values. The core-to-core alignment and the new chronology
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33 409 for core #5 suggest that this tephra layer dates to around 14.1-13.7 cal. ka BP.

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35 410 In conjunction with the increase in LOI values at 13.7 cal. ka BP (Fig. 6B),
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37 411 diatom-inferred lake water pH shifts to more acid values (Fig. 6C). This distinct
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39 412 shift could be the result of increased surface flow that transported organic acids
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41 413 from surrounding soils into the shallow lake (Ampel *et al.* 2014) and could
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43 414 indicate that the catchment vegetation had become denser. Lake waters become
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45 415 neutral around 13.5 cal. ka BP and remain, apart from a few minor fluctuations,
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47 416 around a pH of 7 until 12.7 cal. ka BP (Fig. 6C). The diatom assemblages and the
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49 417 crysophyte:diatom ratios (C:D) in the lowermost diatom zone (DAZ-I; 14.1-13.2
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51 418 cal. ka BP) (Fig. 6C) suggest cold lake water conditions, low light and nutrient
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53 419 availability, but also a gradual shortening of the ice-covered season (Ampel *et al.*
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55 420 2014). Leaf-wax derived hydrogen isotopes, $\delta D_{n-C_{21}}$ and $\Delta\delta D_{terr-aq}$, which reflect
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57 421 the isotopic composition of the source moisture and terrestrial evaporation,
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59 422 respectively indicate a saline moisture source and minor shifts between drier
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423 and wetter conditions (Fig. 6D). However the measurement errors are too large
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425 to render these values significant. Chironomid-derived mean July surface water
temperatures show a step-wise increase from 8 °C at 14.1 cal. ka BP to maximum

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3 426 values of 12-13 °C at 13.2-13.1 cal. ka BP (Fig. 6E). Coleoptera assemblages are
4 427 still composed of species typical of cool and mild climates, but reconstructed
5 428 summer temperatures now reach >14 °C (Watson 2008) and seem to be
6 429 comparable to chironomid-inferred temperatures.

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10 430 The Hässeldala Port proxies suggest marked changes between 13.2 and
11 431 12.7 cal. ka BP. In diatom zone DAZ-II (13.2-12.6 cal. ka BP) (Fig. 6C) the habitat
12 432 structure in the shallow lake changes, open water taxa decline, but the diatom
13 433 flora becomes more diverse and attached-growing species become more
14 434 abundant. This points to denser aquatic vegetation, possibly in response to a lake
15 435 level lowering and/or higher nutrient availability (Ampel *et al.* 2014). The C:D
16 436 ratio shows a decline in the number of crysophyte cysts relative to diatoms,
17 437 which points to a shorter ice-covered season (Fig. 6C). $\delta D_{n-C_{21}}$ values start to
18 438 decrease at 13.2 cal. ka BP and reach minimum values between 13.0-12.8 cal. ka
19 439 BP, coincident with higher $\Delta\delta D_{terr-aq}$ values (Fig. 6G). This shift in hydrogen
20 440 isotope values suggests a freshening of the marine moisture source region and
21 441 distinctly drier conditions at Hässeldala Port (Muschitiello *et al.* 2015b). The
22 442 trend of decreasing $\delta D_{n-C_{21}}$ values and increasing $\Delta\delta D_{terr-aq}$ values was shortly
23 443 interrupted at 12.7 cal. ka BP, when more saline moisture reached the site and
24 444 conditions became wetter. The observed two-step decrease in δD_{aq} values (13.0-
25 445 12.8 and after 12.7 cal. ka BP) has been interpreted as a progressive freshening
26 446 of the Atlantic precipitation water source for Hässeldala Port caused by the
27 447 melting of the Fennoscandian Ice Sheet (Muschitiello *et al.* 2015b). The distinct
28 448 shift in hydrogen isotope values and diatom assemblages around 13.2 cal. ka BP
29 449 coincides with a decline in mean July surface water temperatures to around 10-
30 450 11 °C (Fig. 6C-E).

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46 451 The main features in local pollen assemblage zones HÄP-2 (14.1-13.2 cal.
47 452 ka BP) (rise in *Juniperus communis* pollen percentages) and HÄP-3 (13.2-12.7 cal.
48 453 ka BP) (increase in *Empetrum* and Ericales pollen values) (Andersson 2004;
49 454 Wohlfarth *et al.* 2006) allow for a correlation to the regional Allerød I and II
50 455 pollen zones (Björck & Möller 1987), respectively (Fig. 6H, J). *Nymphaea alba*
51 456 pollen are continuously present in low numbers in HÄP-2 and HÄP-3 (Andersson
52 457 2004) and thus suggest that minimum mean July temperatures were >10 °C
53 458 (Kolstrup 1980). HÄP-3 correlates to local pollen zone Hä-1 and to parts of local
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3 459 pollen zone Hä-2, which were recently established on core #6 (Karlou-
4 460 Charampopoulou 2016) (Fig. 6I) and detail the vegetation changes that occurred
5 461 at the regional Allerød/Younger Dryas transition (see below).

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8 462 Pollen stratigraphic data sets for Blekinge (Berglund 1966; Björck &
9 463 Möller 1987) show that herbs and shrubs still dominate, but that the regional
10 464 vegetation had become more diverse. Plant species now also include *Astragalus*
11 465 *alpinus*, *Cerastium alpinum*, *Dryopteris*, *Isoetes*, *Jasione montana*, *Lycopodium*,
12 466 *Lastrea*, *Myriophyllum alterniflorum*, *Populus tremula*, *Potentilla palustris*, *Prunus*
13 467 *padus*, *Pimpinella*, *Polypodium*, *Ranunculus*, *Rubus*, *Sanguisorba*, *Sorbus*, *Typha*
14 468 *latifolia* and *Valeriana officinalis* (Berglund 1966) (Table 6). Moreover, plant
15 469 macro-remains of *Betula pubescens* have been reported (Berglund 1966).

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18 470 Minimum mean summer temperatures based on indicator species, such as
19 471 *Betula pubescens* (Iversen 1954), *Hippophaë rhamnoides*, *Typha latifolia*
20 472 (Kolstrup 1980), and *Jasione montana* (Kolstrup 1979) (Table 6), suggest that
21 473 minimum mean July temperatures were around 12-15 °C. However these values
22 474 relate to the whole of the regional Allerød pollen zone and do not allow any
23 475 differentiation of changes through time. The chironomid-based summer surface
24 476 water temperature reconstruction and the age model for Hässeldala Port now
25 477 provide a more detailed view of the temperature evolution between 14.1 and
26 478 12.7 cal. ka BP and indicate that warm mean July temperatures were only
27 479 attained between 13.4-13.1 cal. ka BP (Fig. 6E).

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30 480 The increase in organic matter content seen in Hässeldala Port's
31 481 sediments between 12.9-12.7 cal. ka BP (Figs 3A, 6B) is a feature that is typical
32 482 for many Lateglacial sequences in southern Sweden (Björck & Möller 1987;
33 483 Berglund *et al.* 1994). It has been long argued that this increase represents the
34 484 in-wash of topsoil material that was released during the final melting of remnant
35 485 ice or could indicate a lake level lowering due to drier climatic conditions
36 486 (Digerfeldt 1971). A C/N ratio of 13 and bulk $\delta^{13}\text{C}$ values of around -18‰ for
37 487 Hässeldala Port do not support increased in-wash of terrestrial organic material,
38 488 but rather point to a dominance of aquatic sourced organic matter (Kylander *et*
39 489 *al.* 2013). Moreover, higher $\Delta\text{SD}_{\text{terr-aq}}$ values and lower mean July temperatures
40 490 that coincide with the peak in LOI indicate drier and slightly colder conditions
41 491 (Figs 6D, E), which could have led to a lake level lowering. This scenario

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3 492 compares well with the findings of Ampel *et al.* (2014), who had suggested a
4 493 lowering of the lake level as a result of drier conditions. A lower lake level and
5 494 expansion of the aquatic vegetation could have raised the nutrient level in the
6 495 already shallow lake as reflected by high LOI values.
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11 497 *12.7-11.8 cal. ka BP*

12 498 The lithological boundary between the dark brown clayey gyttja (unit 7) and the
13 499 greenish-brown/brown clayey gyttja (unit 8) is reflected by a decrease in LOI
14 500 values (Figs 6A, B). LOI values remain low until *c.* 11.8 cal. ka BP and gradually
15 501 increase thereafter (Fig. 6B). This increase is coupled to a change in sediment
16 502 colour and lithology (unit 9). The initial decrease and later increase in lake
17 503 organic productivity is mirrored by geochemical proxies (Kylander *et al.* 2013),
18 504 which suggest relatively stable conditions between 12.7 and 12.0 cal. ka BP, but
19 505 changes in source material and/or hydrology and in aquatic productivity
20 506 thereafter.
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29 507 Opportunistic diatom taxa (e.g. *Staurosirella pinnata*), which compete
30 508 well under challenging environmental conditions, increase in abundance in
31 509 diatom zone DAZ-III (12.6-11.8 cal. ka BP), as do the relative numbers of
32 510 chrysophyte cysts (12.6-12.3 cal. ka BP) (Ampel *et al.* 2014) (Fig. 6C). The
33 511 increase in chrysophyte cysts suggests that nutrient levels had decreased, which
34 512 would be in line with lower lake organic productivity as inferred by lower LOI
35 513 values (Fig. 6B). This change in lake status (low nutrient, low productivity)
36 514 together with colder temperatures and longer ice-cover seasons gave certain
37 515 diatom species an advantage (Ampel *et al.* 2014).
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45 516 The decline in mean July surface water temperatures, which starts at 12.7
46 517 cal. ka BP, continues until 12.5-12.4 cal. ka BP, when the lowest reconstructed
47 518 temperatures of 8 °C are reached (Fig. 6E). Coincident with the decline in
48 519 temperatures, $\delta D_{n-C_{21}}$ values decrease and $\Delta\delta D_{terr-aq}$ values increase, suggesting a
49 520 fresher moisture source and progressively drier conditions (Fig. 6D). Mean July
50 521 surface water temperatures fluctuate and stay around 9-10 °C between 12.4-11.9
51 522 cal. ka BP, but subsequently start rising again from 9 °C to 12 °C (Fig. 6E). This
52 523 steady rise more or less coincides with distinct changes in $\delta D_{n-C_{21}}$ and $\Delta\delta D_{terr-aq}$
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3 524 values (Fig. 6D), which show a shift towards a more saline marine moisture
4 525 source of precipitation and wetter conditions around 11.8 cal. ka BP
5 526 (Muschitiello *et al.* 2015b).

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8 527 The decrease in *Empetrum* pollen values and the increase in herb and
9 528 grass pollen percentages, especially for *Artemisia*, defines local pollen zone HÄP-
10 529 4 (Andersson 2004; Wohlfarth *et al.* 2006) (12.7-11.7 cal. ka BP). This local
11 530 pollen zone, which partly overlaps with local pollen zones Hä-3, Hä-4 and Hä-5
12 531 on core #6 (Karlathou-Charalampopoulou 2016), was correlated to the regional
13 532 Younger Dryas pollen zone (Figs 6H-J) of Berglund (1966) and Björck & Möller
14 533 (1987) (Andersson 2004; Wohlfarth *et al.* 2006) (see below). *Nymphaea alba*
15 534 pollen, which are present during HÄP-4 (Andersson 2004) suggest minimum
16 535 mean July temperatures of >10 °C (Kolstrup 1980), which compares to those
17 536 reconstructed from chironomid assemblages (Fig. 6E).

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20 537 Pollen stratigraphies for Blekinge (Berglund 1966; Björck & Möller 1987)
21 538 show that the expanding herb and shrub vegetation comprised *Artemisia*,
22 539 *Astragalus alpinus*, *Betula nana*, *Calluna vulgaris*, *Cerastium alpinum*,
23 540 *Chamaenerion alpinum*, *Dryas octopetala*, *Dryopteris*, *Empetrum nigrum*, *Ephedra*,
24 541 *Helianthemum*, *Hippophaë*, *Jasione montana*, *Juniperus communis*, *Lycopus*
25 542 *europaeus*, *Minuartia*, *Myriophyllum alterniflorum*, *M. spicatum*, *M. verticillatum*,
26 543 *Oxytropis capestris*, *Oxyria digyna*, *Plantago*, *Polygonum*, *Potentilla palustris*,
27 544 *Populus tremula*, *Ranunculus*, *Rumex acetosella*, *Rubus*, *Sanguisorba*, *Sorbus*,
28 545 *Saxifraga*, *Selaginella selaginoides*, and *Urtica* (Table 6). Interestingly, also *Pinus*
29 546 *sylvestris* macrofossil remains have been reported for this time interval
30 547 (Berglund 1966). This is surprising given that these had not been found during
31 548 the Allerød pollen zone. On the other hand, especially high pollen values for
32 549 *Pinus* in Allerød lake sediments in Blekinge could be an indication for the
33 550 presence of these trees prior to the regional Younger Dryas pollen zone (Björck
34 551 1979). *Pinus* may have thus survived in micro-climatically favourable habitats or
35 552 the macrofossil finds of *Pinus sylvestris* relate to the very end of the regional
36 553 Younger Dryas pollen zone.

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39 554 Plant indicator species (Table 6), such as *Hippophaë rhamnoides*,
40 555 *Myriophyllum alterniflorum*, *M. spicatum*, *M. verticillatum* and *Jasione montana*,
41 556 would suggest that minimum mean summer temperatures reached 10-12 °C
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3 557 (Kolstrup 1979; 1980). These values however relate to the regional Younger
4 558 Dryas pollen zone in general and do not allow differentiation of changes through
5 559 time. The chironomid-based temperature reconstruction for Hässeldala Port
6 560 shows that mean July surface water temperatures fluctuated between 8-10 °C
7 561 during the Younger Dryas and rose to >10 °C only at 11.9 cal. ka BP (Fig. 6E).
8 562 Since *Pinus sylvestris* requires minimum mean summer temperatures of >12 °C
9 563 (Iversen 1954), we speculate that the macrofossil finds reported by Berglund
10 564 (1966) stem from the latest part of the regional Younger Dryas pollen zone,
11 565 when summer temperatures had already increased.
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21 567 *In and out of the Younger Dryas pollen zone*

22 568 It is interesting to look closer into the sequence of events around the start and
23 569 end of the regional Younger Dryas pollen zone since several of Hässeldala Port's
24 570 proxies (LOI, hydrogen isotopes and diatom assemblages) have been analysed on
25 571 the same split samples of core #5. The chironomid-derived temperature
26 572 reconstruction and the pollen stratigraphies were however established on
27 573 parallel cores that were aligned to the chronology of core #5 using the approach
28 574 of Muschitiello *et al.* (2015b). The former therefore allow for a direct
29 575 comparison, independent of the age assignment, while minor time lags detected
30 576 for the latter could be an artefact because age model errors are not taken into
31 577 account here.
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40 578 The most recent pollen stratigraphy for Hässeldala defines local pollen
41 579 zone Hä-2 (12.77-12.54 cal. ka BP) as a transition zone between the regional
42 580 Allerød and Younger Dryas pollen zones (Figs 6I, J) (Karlou-Charalampopoulou
43 581 2016). This pollen zone is characterised by a first increase in *Artemisia* pollen
44 582 values and a decrease in pollen concentrations. These changes seem to have
45 583 occurred more or less coincident with several other proxies. Sediment organic
46 584 matter content declines, the number of cryptophyte cysts over diatoms starts to
47 585 increase (Fig. 6C), hydrogen isotope proxies show a brief return to wetter
48 586 conditions at *c.* 12.7 cal. ka BP that was followed by progressively drier
49 587 conditions (Fig. 6D) and chironomid-inferred summer surface lake water
50 588 temperatures decline (Fig. 6E). Slight differences between the lower boundaries
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3 589 of HÄP-4 and Hä-3 are due to how pollen zones transitions were originally
4 590 defined, as the increase in *Artemisia* pollen values in core #3, which is
5 591 comparable to the lower boundary of Hä-3, actually already occurs at 12.7 cal. ka
6 592 BP (Andersson 2004).

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10 593 Hä-5 (11.85-11.58 cal. ka BP) reflects the transition zone between the
11 594 regional Younger Dryas and Preboreal pollen zones (Figs 6I, J) and is defined by
12 595 decreasing *Juniperus* and *Artemisia* pollen percentages and rising *Empetrum*
13 596 pollen values (Karlathou-Charalampopoulou 2016). The start of Hä-5 predates the
14 597 transition between pollen zones HÄP-4 and HÄP-5 (11.7 cal. ka BP), but
15 598 coincides with rising sediment organic matter content, changes in diatom
16 599 assemblages, marked shifts in hydrogen isotope values and a rise in chironomid-
17 600 inferred summer surface water temperatures (Fig. 6B-I).

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23 601 Hässeldala Port's proxies thus provide no clear evidence for distinct local
24 602 lags in response to regional climatic changes at the start and end of the Younger
25 603 Dryas. Minor divergences, as seen in the comparison between the two pollen
26 604 stratigraphic records (Fig. 6H, I), are an artefact of where and how pollen zone
27 605 boundaries are drawn and defined.

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34 607 *11.8-10 cal. ka BP*

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36 608 Hässeldala Port's sediments shift from clayey gyttja (unit 8) to gyttja (units 9-11)
37 609 around 11.8 cal. ka BP and to peaty gyttja (unit 12) at 10.9 cal. ka BP (Fig. 6A).
38 610 The steady increase in LOI values and subsequent values of >80% depict the
39 611 increase in sediment organic matter and show that the small lake gradually filled
40 612 in (Fig. 6B). This is also reflected in the diatom assemblages of DAZ-IV, which
41 613 suggest a transition towards a less alkaline mire environment that gradually
42 614 turned into a rather dry mire environment (Ampel *et al.* 2014). Coleoptera,
43 615 which are typical of acid bog environments and dry shrub heathland, such as
44 616 *Amara alpina* and *Olophrum boreale* now appear (Watson 2008). The trend to
45 617 overall wetter conditions as seen in decreasing $\Delta\delta D_{\text{terr-aq}}$ values was briefly
46 618 interrupted around 11.5 cal. ka BP, when hydrogen isotope values suggest both a
47 619 freshening of the marine sources of precipitation and a return to slightly drier
48 620 conditions (Fig. 6D). The timing of this shift compares to the Preboreal
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3 621 Oscillation, a short cooling phase identified in North Atlantic paleo-records
4 622 (Björck *et al.* 1996, 1997) and predates the oscillation discussed earlier by
5 623 Wohlfarth *et al.* (2006) based on the LOI curve of core #3. The Hässeldala tephra,
6 624 identified in core #1 and 3 (Davies *et al.* 2003, 2004; Wohlfarth *et al.* 2006),
7 625 occurs around 11.3 cal. ka BP and thus after the oscillation seen in the hydrogen
8 626 isotope proxies (Fig. 6D, F). Cryptotephra, which has been geochemically
9 627 assigned to the Askja-S tephra was also found in cores #1 and 3 and can now be
10 628 dated to around 10.9-10.7 cal. ka BP (Fig. 6F).

11 629 Local pollen zone HÄP-5 (11.7-11.3 cal. ka BP) (Andersson 2006)
12 630 overlaps with local pollen zones Hä-5 and Hä-6 in core #6 (11.85-11.3 cal. ka BP)
13 631 (Karlou-Charalampopoulou 2016) (Fig. 6H, I). The distinct peak in *Empetrum*
14 632 pollen percentages, a decrease in *Artemisia* pollen values and overall a slight
15 633 increase in tree pollen, mainly of *Betula*, allows correlations to the YD-PB
16 634 transition zone of Berglund (1966) and Björck & Möller (1987). The YD-PB
17 635 transition zone has been assigned to the early part of the regional Preboreal
18 636 pollen zone (Björck *et al.* 1996), thus marking the start of the Holocene.
19 637 Macroscopic charcoal appears in large quantities around 11.8-11.6 cal. ka BP and
20 638 the first macroscopic finds of *Betula pubescens* occur at 11.6-11.4 cal. ka BP
21 639 (Wohlfarth *et al.* 2006) (Fig. 6G). Tree pollen percentages increase further in
22 640 HÄP-6 (11.3-10.9 cal. ka BP) and HÄP-7 (10.9-10.3 cal. ka BP), herb and shrub
23 641 pollen decline (Andersson 2006), and the presence of *Pinus sylvestris* is attested
24 642 by finds of needles at 11.0 cal. ka BP (Wohlfarth *et al.* 2006) (Fig. 6G, H). Aquatic
25 643 pollen of *Typha angustifolia* and *Nymphaea alba* in HÄP-6 and of *Myriophyllum*
26 644 *alterniflorum/spicatum* in HÄP-7 (Andersson 2004) suggest that minimum July
27 645 temperatures had increased distinctly and may have reached >14 °C (Kolstrup
28 646 1980). Wetter conditions, as suggested by decreasing $\Delta\delta D_{\text{terr-aq}}$ values (Fig. 6D),
29 647 and warmer summers likely favoured the further establishment and expansion
30 648 of trees in Hässeldala Port's catchment. HÄP-6 and HÄP-7 are correlated to the
31 649 regional Preboreal pollen zones of Berglund (1966).

32 650 The vegetation during the regional early Preboreal pollen zone in
33 651 Blekinge now also included *Botanococcus umbellatus*, *Caltha palustris*, *Isoetes*
34 652 *lacustris*, *Littorella uniflora*, *Menyanthes trifoliata*, *Rubus fruticosus*, and *Typha*
35 653 *latifolia*, and *Pinus sylvestris* (Berglund, 1966) (Table 6). This latter species, as

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3 654 well as the presence of *Typha latifolia* and *Betula pubescens*, in addition to other
4 655 indicator species, suggests that minimum mean July temperatures now reached
5 656 >12 °C (Iversen 1954; Kolstrup 1980). This assumption compares nicely to
6 657 higher summer temperatures inferred from chironomid assemblages (Fig. 6E)
7 658 and aquatic pollen.

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14 660 Wider implications

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18 662 Hässeldala Port's pollen-stratigraphic records of cores #3 (Andersson 2004;
19 663 Wohlfarth *et al.* 2006) and #6 (Karlatau-Charampopoulou 2016) and the refined
20 664 age model now provide a new chronological template for Lateglacial climatic and
21 665 environmental changes in southern Sweden (Table 7). Andersson's (2004) HÄP-
22 666 1, which is correlated to the regional Older Dryas pollen zone, is dated to >14.06
23 667 cal. ka BP. HÄP-2 compares to the regional Allerød I pollen zone and ranges
24 668 between 14.06 and 13.24 cal. ka BP and HÄP-3 or the regional Allerød II pollen
25 669 zone dates to between 13.24 and 12.67 cal. ka BP. Wohlfarth *et al.* (2006) had
26 670 earlier correlated local pollen zone HÄP-4 (12.67-11.68 cal. ka BP) with the
27 671 regional Younger Dryas pollen zone, but this correlation now needs to be
28 672 revised, following the high-resolution pollen stratigraphy of core #6 (Karlatau-
29 673 Charampopoulou 2016). This latter work shows that a Late Allerød/early
30 674 Younger Dryas transition zone (Hä-2: 12.77-12.54 cal. ka BP) overlaps with parts
31 675 of Andersson's (2004) pollen zones HÄP-3 and HÄP-4 (Fig. 6I, J) and suggests
32 676 that changes in vegetation composition already commenced at 12.77 cal. ka BP
33 677 (Table 7). Karlatau-Charampopoulou's (2016) local pollen zones Hä-3 and Hä-4
34 678 correlate to the regional Younger Dryas pollen zone as defined by Björck &
35 679 Möller (1987) and extend between 12.54 and 11.85 cal. ka BP (Table 7). The
36 680 transition between the regional Younger Dryas and Preboreal pollen zones
37 681 (Berglund 1966; Björck & Möller 1987), which Björck *et al.* (1996) had assigned
38 682 to the early Holocene, is also clearly seen in Hässeldala Port's pollen
39 683 stratigraphy. It is represented by pollen zone Hä-5 (Karlatau-Charampopoulou
40 684 2016) (Fig. 6 I, J) and dated to 11.85-11.58 cal. ka BP (Table 7).

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3 685 Muschitiello & Wohlfarth (2015) earlier discussed the time-transgressive
4 686 nature of vegetation changes in northern Europe around the Allerød – Younger
5 687 Dryas transition using well-dated pollen stratigraphic records. They showed that
6 688 the early response seen in Sluggan Bog, Northern Ireland (Walker *et al.* 2012)
7 689 (13.0 cal. ka BP) predates vegetation shifts in sites located further to the east by
8 690 *c.* 300-350 years (Fig. 7) and hypothesize that this may have been caused by
9 691 regional cooling around the Nordic Seas. Hässeldala Port now adds an important
10 692 data point to this network and supports Muschitiello & Wohlfarth's (2015)
11 693 conclusions that vegetation shifts reconstructed for sites in Norway, Sweden and
12 694 western Germany postdate those in Northern Ireland and also occurred 100-250
13 695 years later than the onset of Greenland Stadial 1 (Fig. 7). The change in
14 696 vegetation composition seen in Kråkenes (Norway) at 12.75-12.7 cal. ka BP
15 697 (Birks *et al.* 2000), Madtjärn (Swedish west coast) at 12.7-12.6 cal. ka BP (Björck
16 698 *et al.* 1996), Hässeldala Port between 12.77-12.54 cal. ka BP and in Meerfelder
17 699 Maar (Germany) at 12.7-12.65 cal. ka BP (Rach *et al.* 2014) thus only took place
18 700 at a time when critical threshold temperatures had been reached and when
19 701 Greenland ice cores reflect the establishment of full stadial climate conditions
20 702 and a stadial atmospheric circulation regime (Steffensen *et al.* 2008; Rach *et al.*
21 703 2014; Muschitiello & Wohlfarth 2015).

22 704 This hypothesis can be tested by comparing the timing of hydroclimate
23 705 shifts and vegetation records using the two available detailed δD stratigraphies
24 706 of Hässeldala Port (Muschitiello *et al.* 2015b) and Meerfelder Maar (Rach *et al.*
25 707 2014). The δD_{aq} records from Hässeldala Port and Meerfelder Maar show a
26 708 consistent trend towards more negative values at the start of the Younger Dryas
27 709 pollen zone (Fig. 8). This suggests large-scale cooling and more negative δD
28 710 values of the marine moisture source of precipitation – the North Atlantic Ocean
29 711 – associated with a southward diversion of the storm track trajectories and sea-
30 712 ice expansion (Rach *et al.* 2014). More interestingly, the two records display an
31 713 opposite hydro-climate behaviour between southern Sweden and western
32 714 Germany during the Late Allerød pollen zone/Greenland Interstadial 1a (*c.* 13.1-
33 715 12.9 cal. ka BP), with relatively lower δD_{aq} values at Hässeldala Port and
34 716 relatively higher δD_{aq} values at Meerfelder Maar. This pattern is also evident in
35 717 the $\Delta\delta D_{terr-aq}$ profiles, which suggest drier conditions at Hässeldala Port as

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3 718 opposed to more humid conditions at Meerfelder Maar (Fig. 8). Muschitiello *et al.*
4 719 (2015b) suggested that such opposite hydro-climate patterns prior to the start of
5 720 the Younger Dryas are an expression of increased freshwater forcing from the
6 721 Fennoscandian Ice Sheet to the Nordic Seas, which affected the hydro-climate
7 722 system at synoptic scale. Climate model simulations indeed show that freshwater
8 723 forcing from the Fennoscandian Ice Sheet results in relatively lower atmospheric
9 724 pressure over the western sector of Europe and Greenland, whereas Northern
10 725 Europe is affected by relatively higher-pressure conditions (Muschitiello *et al.*
11 726 2015b). This scenario could explain the spatially complex differences in moisture
12 727 availability across the North Atlantic (Muschitiello *et al.* 2015b). The hydro-
13 728 climate records from Hässeldala Port and Meerfelder Maar also show that
14 729 southern Sweden likely experienced drier conditions as compared to Western
15 730 Europe during the Younger Dryas (Fig. 8). This is in line with evidence of
16 731 increasing continentality and more severe aridity in Europe from south to north
17 732 (e.g. Birks *et al.* 2014).

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28 733 The transition out of the Younger Dryas is equally interesting. At *c.* 11.9
29 734 cal. ka BP, δD_{aq} values from Hässeldala Port and Meerfelder Maar start to rise
30 735 and increase by $\sim 40\text{‰}$ and 30‰ , respectively (Fig. 8). The shifts in δD_{aq} are
31 736 paralleled by changes in $\Delta\delta D_{terr-aq}$, which indicate progressively wetter
32 737 conditions both in Western Germany and southern Sweden (Fig. 8). Moreover,
33 738 these hydro-climate shifts were initiated *c.* 300 years earlier in Europe than the
34 739 transition into the Holocene in Greenland (Fig. 8). This sequence of events is in
35 740 agreement with the hypothesis of a gradual northward recession of the North
36 741 Atlantic sea-ice margin during the second half of the Younger Dryas, which paved
37 742 the way for a strengthening of the Atlantic Meridional Overturning Circulation
38 743 associated with the start of the warm Holocene interglacial (e.g. Lane *et al.* 2013;
39 744 Pearce *et al.* 2013; Bartolomé *et al.* 2015).

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49 745 Concluding, we find distinct and in-phase gradual perturbations of the
50 746 hydro-climate system across the North Atlantic leading the transitions into the
51 747 Younger Dryas stadial and the Holocene interglacial. These features imply that
52 748 complex atmosphere-ocean feedbacks are at work prior to an abrupt climate
53 749 change.
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8 754 The Lateglacial sedimentary successions from the ancient lake at Hässeldala Port
9 755 in southern Sweden spans the period 14.5-9.5 cal. ka BP and forms one of the
10 756 best chronologically-constrained sites in Europe. Hässeldala Port's sediments
11 757 were analysed using a variety of proxies that allow capturing local and regional
12 758 climatic and environmental signals. The refined age model for the multi-proxy
13 759 records (lithology, geochemistry, pollen, diatoms, chironomids, biomarkers,
14 760 hydrogen isotopes) now enables discussing environmental and climatic changes
15 761 in a regional context and provides a new chronological framework for the
16 762 vegetation development in southern Sweden. Hässeldala Port's multi proxy
17 763 records imply that regional cooling and hydroclimate changes in the Nordic Sea
18 764 region led to more or less synchronous local responses in and around the ancient
19 765 lake. These changes were however offset by several hundred years as compared
20 766 to the start and end of Greenland stadial GS-1. Hässeldala thus provides further
21 767 insights into the spatial and temporal complexity of environmental and
22 768 hydroclimatic responses to North Atlantic rapid climate shifts.

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6 1022 *Figure 1.* A. Location of the study area in southern Sweden. B. Topography
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8 1023 around Hässeldala Port's peatbog. C. Close-up of Hässeldala Port's peatbog
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10 1024 showing the location of the cored successions (see Table 1 for details on the
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12 1025 different sequences). All studied profiles were derived from the deepest part of
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14 1026 the bog. Present-day lakes are shown in blue and peat bogs in a light brown
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16 1027 colour. Elevations around Hässeldala Port reach between 35 and 80 m above
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18 1028 present day sea level (a.s.l.).
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21 1030 *Figure 2.* Lithostratigraphy, total organic carbon (TOC) and loss-on-ignition (LOI)
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23 1031 for Hässeldala Port's core #2 (Wohlfarth *et al.* 2006) and #5 (Steinthorsdottir *et*
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25 1032 *al.* 2013). The brown and grey shaded rectangles mark the distinct patterns that
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27 1033 can be recognised in all profiles. See Fig. 1C for the location of the cores.

28 1034 *Figure 3.* A. TOC/LOI (%) curves for cores #1, #2, #3, #HP4, #5, and #6 according
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30 1035 to depth. The brown and grey shaded rectangles mark the distinct patterns that
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32 1036 can be recognised in all profiles. See Table 1 for references for each analysed
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34 1037 sediment core. B. TOC/LOI (%) curves for cores #1, #2, #3, #HP4, #5, and #6
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36 1038 shown on the revised chronology for core #5 (Muschitiello *et al.* 2015b) (see text
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38 1039 and Fig. 4). The grey shaded area shows those parts of the profiles, where the age
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40 1040 was interpolated using known sedimentation rates. Tephra layers that have been
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42 1041 geochemically identified are the Askja-S tephra (AsT), the Hässeldala tephra
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44 1042 (HDT) and the Borrobol tephra (BT) (Davies *et al.* 2003, 2004; Wohlfarth *et al.*
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46 1043 2006), while other tephtras in core #1 and #HP4 have not been further analysed.
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48 1044 None of these tephtras has been used in the construction of the age model.

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50 1046 *Figure 4.* Age model of Hässeldala Port's composite radiocarbon-dated
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52 1047 succession using the IntCal13 calibration curve (Reimer *et al.* 2013). Circles
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54 1048 indicate calibrated ¹⁴C dates used to construct the age model. Blue colored circles
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56 1049 refer to dates that were transferred from core #2 onto core #5's depth scale. The
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58 1050 orange dotted lines show the 95% confidence bounds of the model, and the red

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3 1051 dotted line the modelled weighted mean age. See Table 3 for details on all ^{14}C
4 1052 dates.

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7 1053 *Figure 5.* Shoreline displacement and relative timing of the deglaciation in
8 1054 Blekinge province, southernmost Sweden during the regional Bølling and Older
9 1055 Dryas pollen zones. See Table 5 for details of all sites shown here. A. Ice
10 1056 recession lines (white dashed line) according to the stacked local clay-varve
11 1057 chronology (Ringberg 1991). The formation of the highest coastline of the Baltic
12 1058 Ice Lake estimated at 65-67 m a.s.l. (blue line) corresponds approximately to
13 1059 local varve years -10 to +10, i.e. to a position of the active ice margin around
14 1060 Farslycke and sites 171 and 172. The white shaded part shows the extension of
15 1061 the former ice sheet. B. Ice recession lines (white dashed line) according to the
16 1062 stacked local clay-varve chronology (Wohlfarth *et al.* 1994). The shoreline at 50-
17 1063 55 m a.s.l. (orange line) formed coincident with the deposition of the thick varves
18 1064 (local varve years +88 to +220). By this time, the active ice margin was located to
19 1065 the north of the study region, but stagnant ice remained above the highest
20 1066 coastline (Björck & Möller 1987). C. Land uplift around Hässeldala Port during
21 1067 the last deglaciation and location of the site in respect to the former coastlines of
22 1068 the Baltic Ice Lake. A. The coastline at 50-55 m a.s.l. (orange line) corresponds to
23 1069 the end of the regional Bølling pollen zone. D. The coastline at 45-50 m a.s.l.
24 1070 (dark green line) corresponds to the regional Older Dryas pollen zone. During
25 1071 the early part of the regional deglaciation Hässeldala Port was situated in an
26 1072 archipelago in the Baltic Ice Lake, but soon became part of a more continuous
27 1073 land area.

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34 1075 *Figure 6.* Hässeldala Port's lithostratigraphy and multi-proxy records shown on
35 1076 the revised chronology of core #5. A. Simplified lithostratigraphy of core #5. The
36 1077 lowermost unit 1 was only present in cores #1-3 and HP4 (see Figs 2 and 3A, B
37 1078 for details) and has been interpolated to core #5. B. Loss-on-ignition (LOI) curve
38 1079 for core #5. C. Reconstructed pH, diatom assemblage zones (DAZ-I to DAZ-IV)
39 1080 and crysophyte:diatom (C:D) ratios for core #5 (Ampel *et al.* 2014). D. Leaf-wax
40 1081 derived hydrogen isotope parameters for aquatic ($\delta\text{D}_n\text{-C}_{21}$) and terrestrial plants
41 1082 ($\delta\text{D}_n\text{-C}_{27-29-31}$) and calculated terrestrial evaporation ($\Delta\delta\text{D}_{\text{terr-aq}}$) for core #5

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3 1083 (Muschitiello *et al.* 2015b). All proxies in B-D were analysed on the same split
4 1084 samples. E. Chironomid-derived mean summer surface water temperatures for
5 1085 core #HP4 (Watson 2008) are shown on the revised core #5 chronology. F.
6 1086 Geochemically determined cryptotephra layers (AsT = Askja-S tephra; HDT =
7 1087 Hässeldala tephra; BT = Borrobol tephra) in cores #1-3 (Davies *et al.* 2003; 2004;
8 1088 Wohlfarth *et al.* 2006) are shown on the timescale of core #5 using the core-to-
9 1089 core alignment described in the text. G. The presence/absence of charcoal and of
10 1090 *Betula pubescens* and *Pinus sylvestris* macrofossils in core #2 is here shown on
11 1091 the timescale of core #5. H. Local pollen assemblage zones (HÄP-1 to HÄP-7) of
12 1092 core #3 (Andersson 2004; Wohlfarth *et al.* 2006) plotted on the revised core #5
13 1093 chronology. I. Local pollen assemblage zones of core #6 (Karatou-
14 1094 Charalampopoulou 2016) plotted on the revised core #5 chronology. J. Regional
15 1095 pollen zones (Björck & Möller 1987) and inferred Lateglacial and early Holocene
16 1096 vegetation in southern Sweden.
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18 1098 *Figure 7.* Age estimates for Allerød-Younger Dryas pollen zone transitions (one
19 1099 sigma) inferred from well-dated North European pollen stratigraphic sites
20 1100 (green bars): Kråkenes (Birks *et al.* 2000); Madtjärn (Björck *et al.* 1996); Sluggan
21 1101 Bog (Walker *et al.* 2012) and Meerfelder Maar (Rach *et al.* 2014). The age of the
22 1102 local pollen zone Hä-3 (brown bar) in Hässeldala Port (Karatou-
23 1103 Charampopoulou 2016), which is a transitional zone between the regional
24 1104 Allerød and Younger Dryas pollen zones, is shown for comparison. Age estimates
25 1105 for the onset of Greenland Interstadial 1a (GI-1a) and Greenland Stadial 1 (GS-1)
26 1106 are expressed on the IntCal13 time scale after synchronization with the GICC05
27 1107 scale (Muscheler *et al.* 2014).

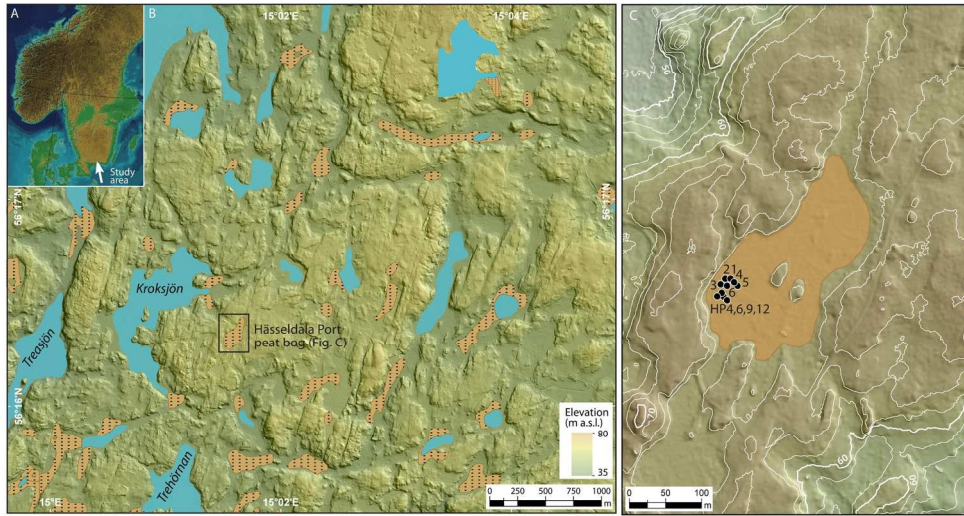
28 1108 *Figure 8.* Comparison of Lateglacial and early Holocene vegetation and
29 1109 hydroclimate changes as reconstructed from Hässeldala Port (HÄ) (Muschitiello
30 1110 *et al.* 2015b) and Meerfelder Maar (MFM) (Rach *et al.* 2014) compared to the
31 1111 Greenland event stratigraphy (displayed and plotted on the IntCal13 time scale -
32 1112 Muscheler *et al.* 2014). The red star marks the position of the Vedde Ash in
33 1113 Meerfelder Maar and in the NGRIP ice core.
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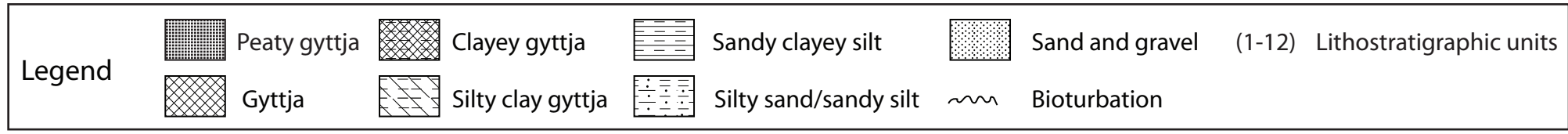
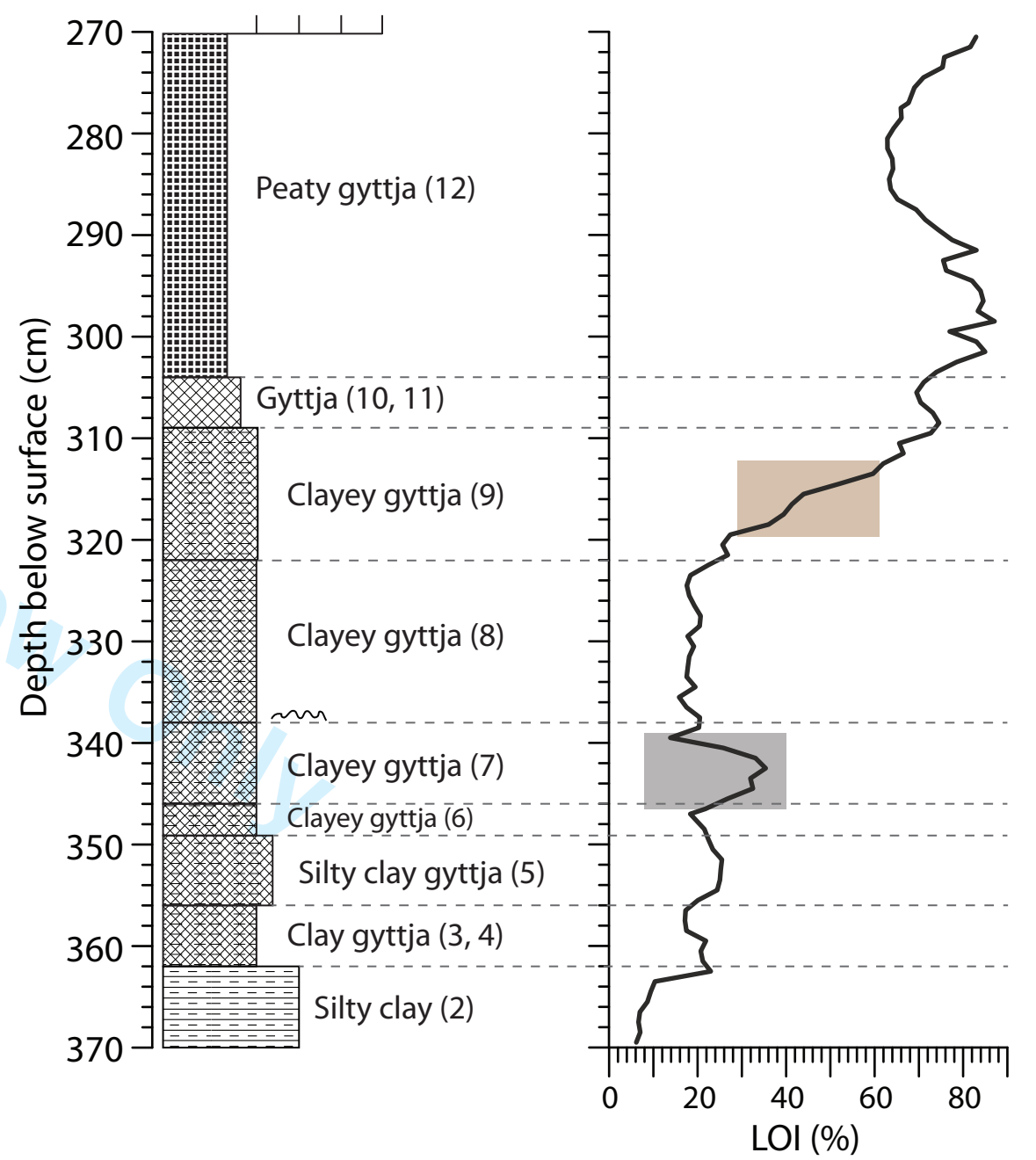
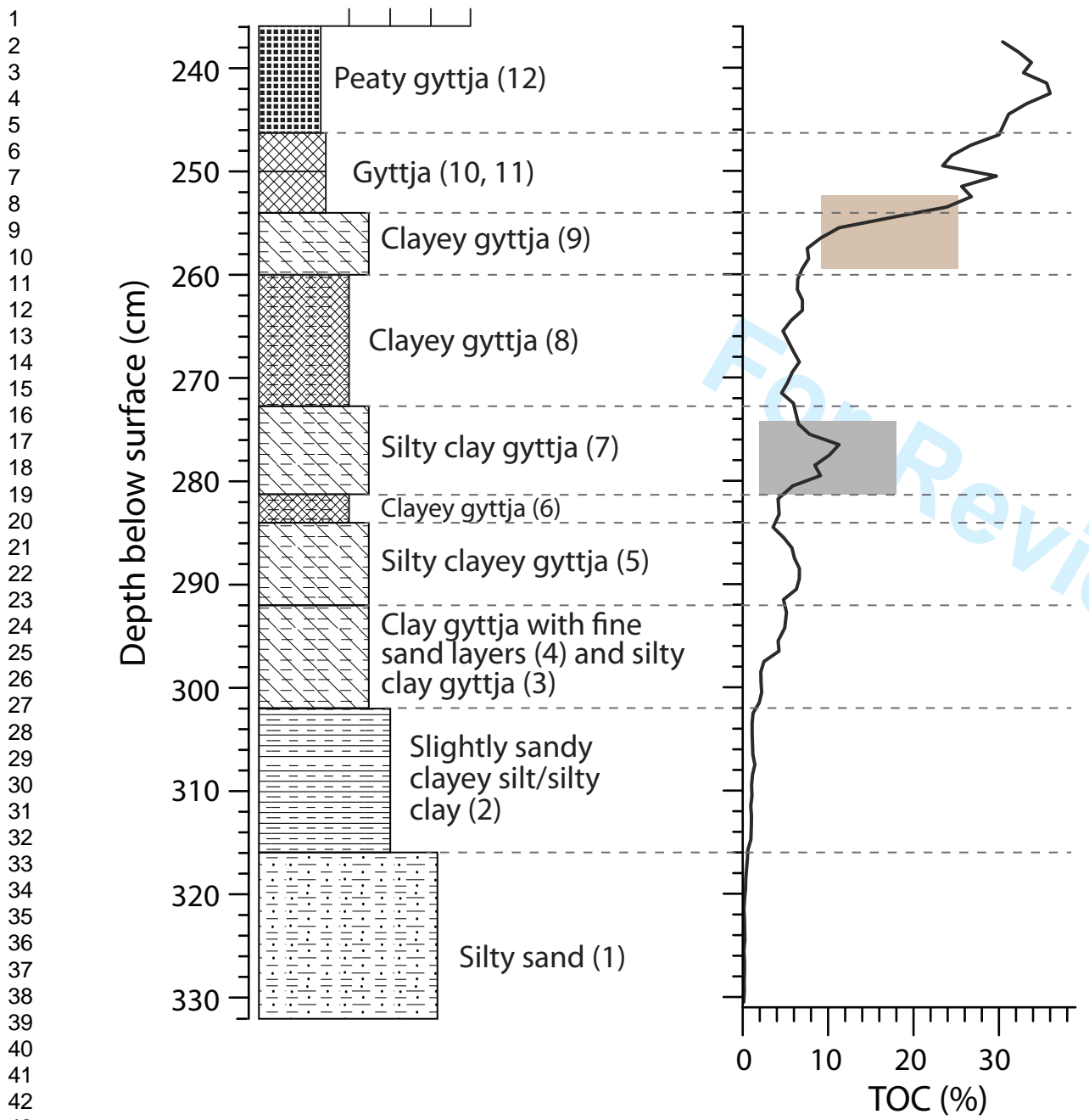
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Core #2

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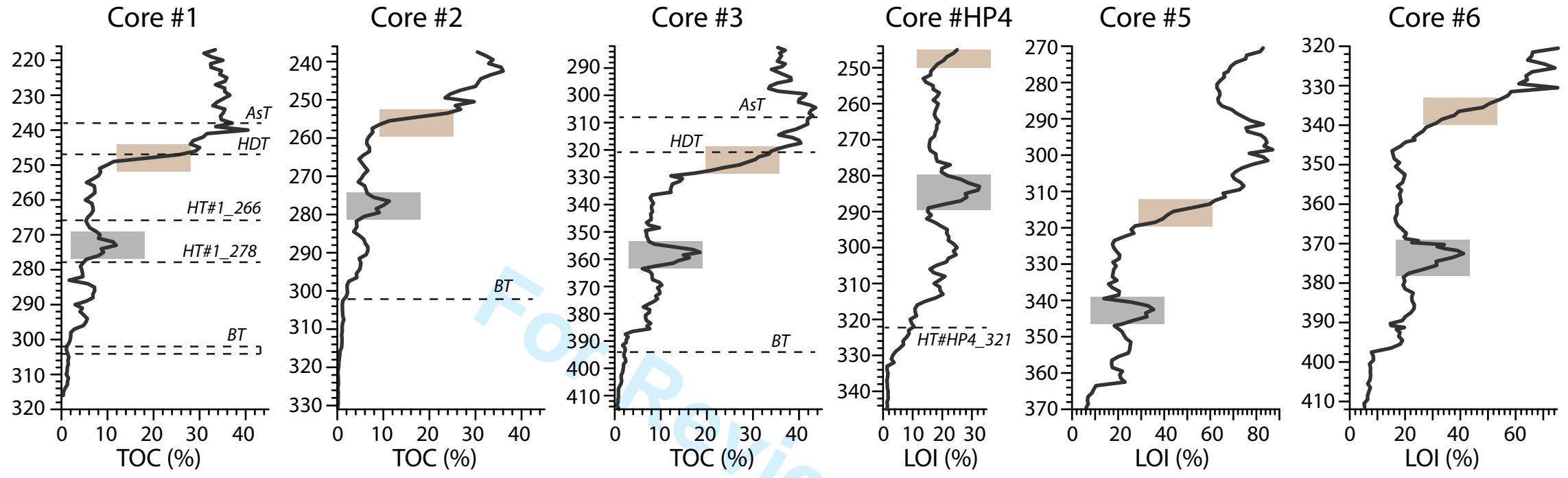
Core #5



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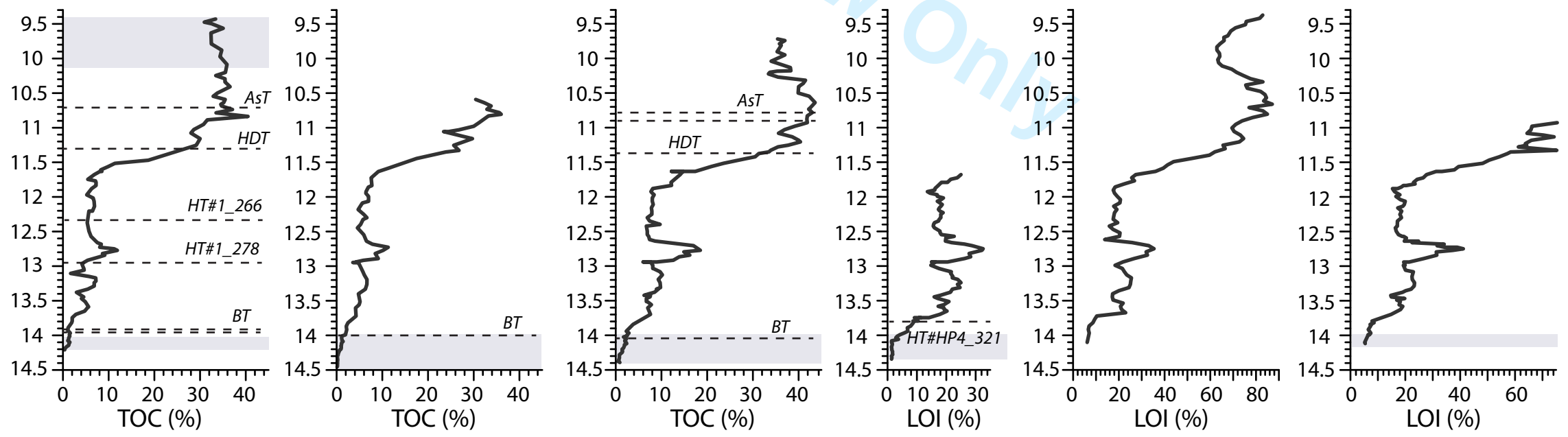
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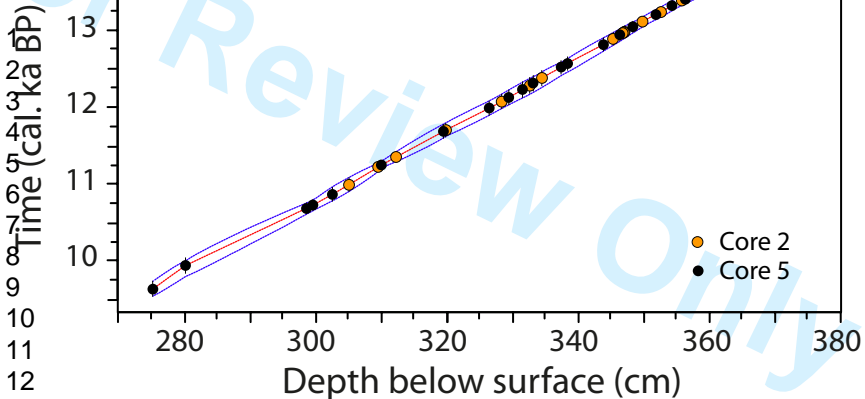
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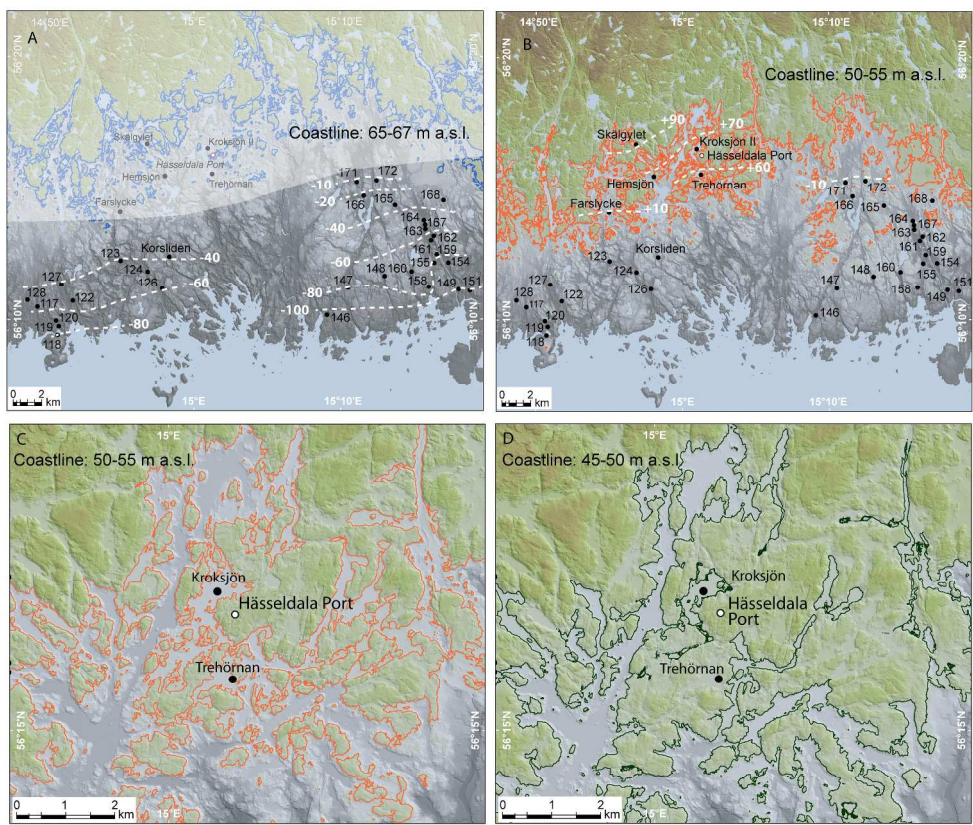
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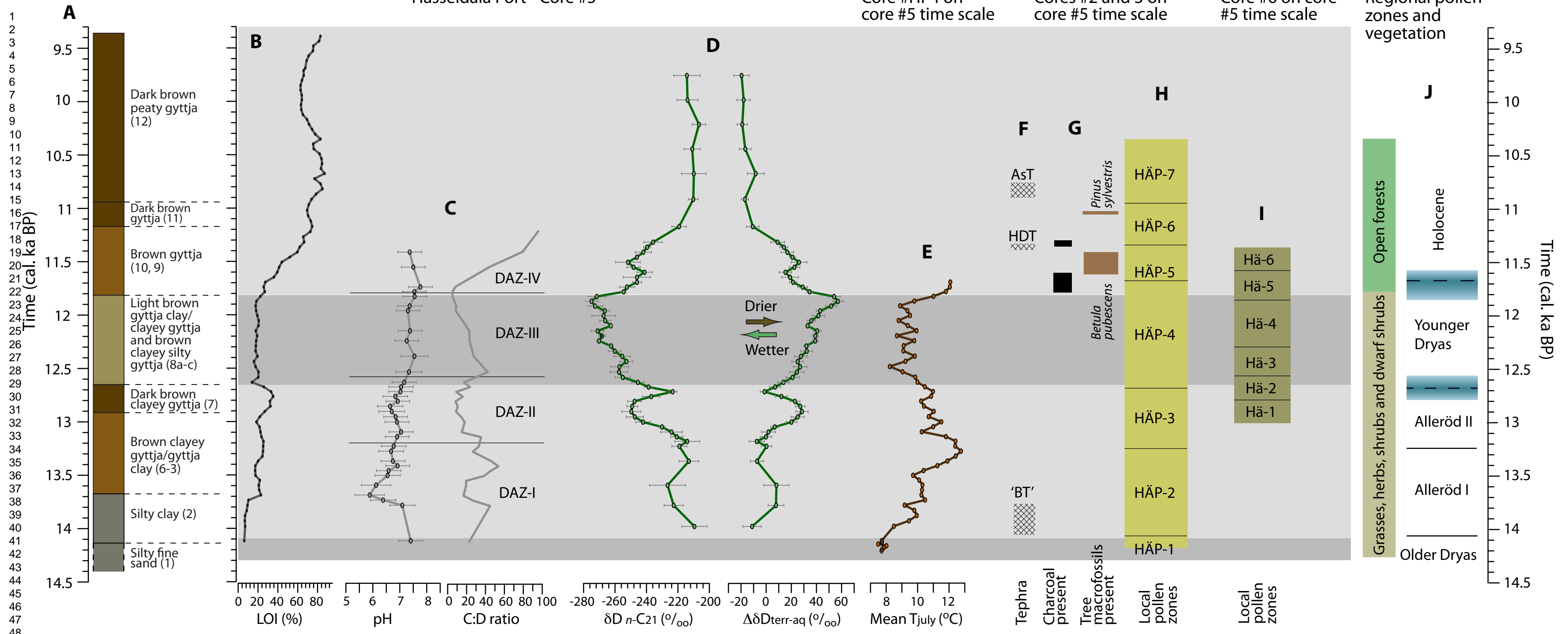
Hässeldala Port - Core #5

Core #HP4 on core #5 time scale

Cores #2 and 3 on core #5 time scale

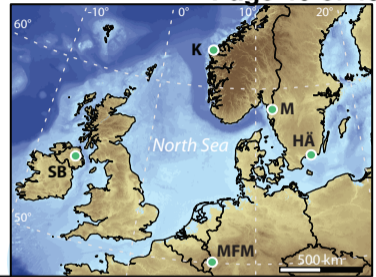
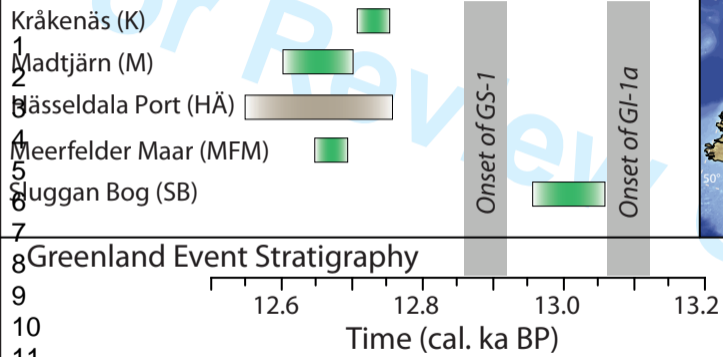
Core #6 on core #5 time scale

Regional pollen zones and vegetation

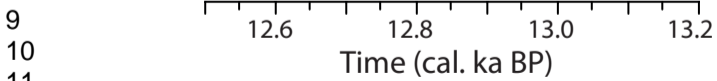


Boreas

Allerød - Younger Dryas pollen zone transition



Greenland Event Stratigraphy



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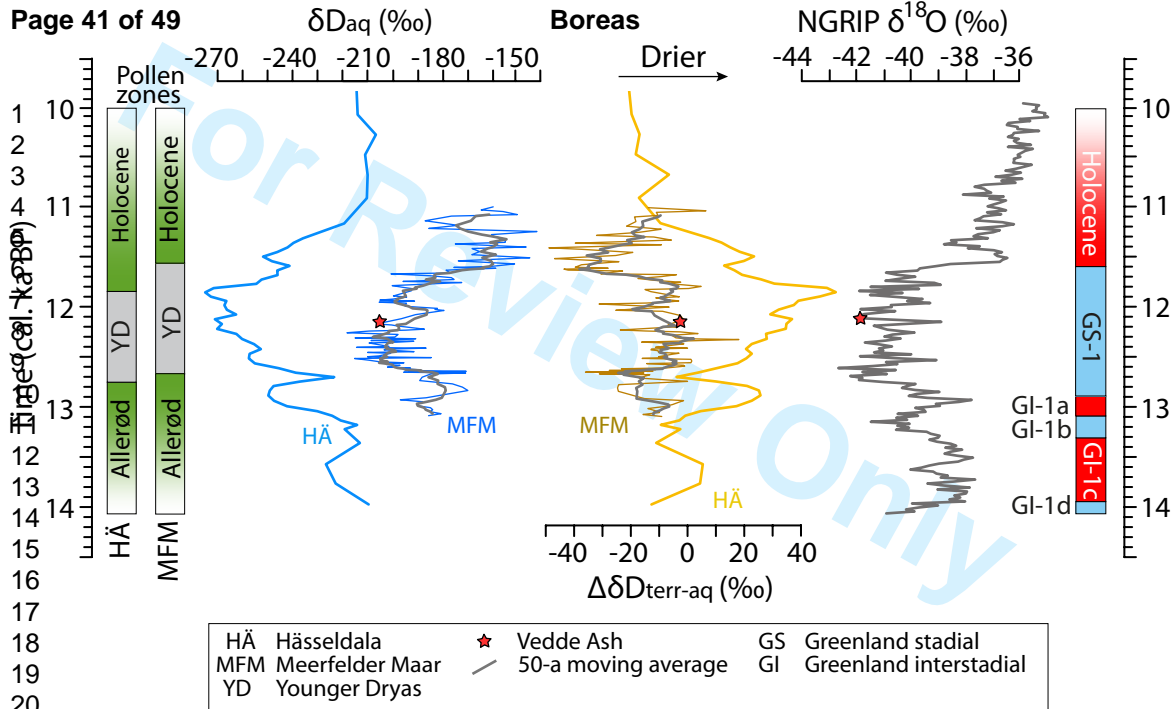


Table 1. Hässeldala Port sedimentary successions and analysed proxies. LOI = loss-on-ignition; TOC = total organic carbon; C/N = carbon:nitrogen ratio; XRF = major elements. Watson (2008) obtained a total of 14 parallel sediment cores from Hässeldala and sampled six of these for coleoptera analysis, two for tephra, four for LOI, and one for chironomids (#HP4). Correlation between these cores was based on stratigraphic correlations and on the pattern of the respective LOI curves (Watson 2008). See Fig. 1C for the location of the various cores.

Core #	Proxy	Reference
1	LOI, TOC, tephra	Davies <i>et al.</i> (2003)
2	TOC, tephra, ^{14}C	Davies <i>et al.</i> (2004)
3	TOC, tephra, pollen stratigraphy	Davies <i>et al.</i> (2004); Andersson (2004), Wohlfarth <i>et al.</i> (2006)
4	TOC, C/N, $\delta^{13}\text{C}_{\text{bulk}}$, $\delta^{15}\text{N}_{\text{bulk}}$, XRF	Kylander <i>et al.</i> (2013)
HP4	LOI, tephra, chironomids	Watson (2008), Muschitiello <i>et al.</i> (2015b)
HP6	LOI	Watson (2008)
HP9	LOI, tephra	Watson (2008)
HP12	LOI	Watson (2008)
5	LOI, ^{14}C , leaf stomata	Steinhorsdottir <i>et al.</i> (2013, 2014)
	Diatoms	Ampel <i>et al.</i> (2014)
	XRF	Unpublished
	Biomarkers	Muschitiello <i>et al.</i> (2015a, b)
	$\delta\text{D}_{\text{wax}}$, $\delta^{13}\text{C}$	Muschitiello <i>et al.</i> (2015b)
6	LOI, pollen stratigraphy	Muschitiello <i>et al.</i> (2015b); Karlatou-Charalampopoulou (2016)

Table 2. Cryptotephra found in Hässeldala Port's sediments (Davies *et al.* 2003, 2004) and Watson (2008). The Askja-S tephra (AsT), the Hässeldala tephra (HDT) and the Borrobol tephra (BT) have been geochemically identified in cores #1, 2 and 3 (Davies *et al.* 2003, 2004). Other cryptotephra layers found in cores #1 and HP4 could not be geochemically identified. See Fig. 1C for the location of the cores and Fig. 3A for details on the position of the individual tephra horizons. Tephra shard concentrations peak in core #1 at 304-303 cm depth, but remain high as far as 301 cm depth. Note that Matthews *et al.* (2011) suggested that Hässeldala Port's BT is younger than the BT found in Scotland and likely an equivalent to the Penifiler tephra.

Cryptotephra	Core #1 depth (cm)	Core #2 depth (cm)	Core #3 depth (cm)	Core #HP4 depth (cm)
Askja-S (AsT)	HT#1_238		HT#3_308-310	
Hässeldala (HDT)	HT#1_247		HT#3_321-322	
NN	HT#1_266			
NN	HT#_278			
NN				HT#HP4_321
Borrobol (BT)	HT#1_304-303 (302-301)	HT#2_302	HT#3_394	

Table 3. AMS ^{14}C dates for Hässeldala Port based on selected terrestrial plant remains and used to construct the composite age-depth model (Wohlfarth *et al.* 2006; Muschitiello *et al.* 2015b). Sample ID's shown in italics refer to dates that were transferred from core #2 to core #5 (see Fig. 4). However not all of these were used to construct the age model, since some of the ^{14}C dates were clear outliers. B = *Betula nana* – leaves, leaf fragments, seeds, buds, twigs; Bi = *Betula* indet. – twigs, leaf fragments, seeds, bark; Bp = *Betula pubescens* – seeds; C = charcoal; D = *Dryas octopetala* – leaves, leaf fragments, seeds, buds; M = moss; P = *Pinus sylvestris* – needles; T = undetermined terrestrial material. The sample depth refers to core #5 and the depth shown in brackets refers to the original depth in core #2 (Wohlfarth *et al.* 2006).

Sample depth (cm)	Thickness error (cm)	Sample ID	Material analysed	^{14}C age (a)	^{14}C error (1 sigma)	Used in the age model
369.5	0.5	UBA-20296	B, D	12 211	57	Yes
367.5	0.5	UBA-20297	D, S	11 972	62	Yes
366.6 (298.45)	1.15	<i>Ua-20516</i>	B, D, T	12 355	190	No
365.25	0.75	UBA-20298	B, D, S	11 673	56	No
363	0.5	UBA-20299	B, D	11 819	57	Yes
362.65 (296.2)	1.4	<i>Ua-20517</i>	B, D	11 920	90	Yes
360.5	0.5	UBA-20300	B, D	11 735	60	Yes
360.15 (294.35)	0.75	<i>Ua-20518</i>	B, D	11 990	110	No
358.5	0.5	UBA-20301	B, D	11 614	55	Yes
358.4 (292.85)	0.95	<i>Ua-20519</i>	B	11 805	240	Yes
356.5	0.5	UBA-20302	B, D	11 565	56	Yes
356 (291)	1.1	<i>Ua-20520</i>	B	11 525	85	Yes
354.5	0.5	UBA-20303	B	11 515	65	Yes
352.85 (289)	1.2	<i>Ua-20521</i>	T	11 490	85	Yes
352	1	UBA-20304	B	11 225	60	Yes
350 (287)	1.25	<i>Ua-20522</i>	B	11 455	125	Yes
348.5	0.5	UBA-20305	B	11 118	60	Yes
347.25 (285)	1.2	<i>Ua-20523</i>	B, T	11 245	95	Yes
347.1 (283.25)	0.9	<i>Ua-20524</i>	B, T	11 275	95	Yes
346.95 (281.5)	1.05	<i>Ua-20525</i>	B	11 200	165	Yes
346.5	0.5	UBA-20306	B	10 998	63	Yes
345.5 (279.5)	1	<i>Ua-20526</i>	B, T	10 935	80	Yes
344	1	UBA-20307	B	10 894	63	Yes
339.9 (274.75)	1.8	<i>Ua-20527</i>	B, T	11 070	135	No
338.5	0.5	UBA-20308	B	10 644	58	Yes
337.5	0.5	UBA-20309	B	10 484	50	Yes
337.4 (272.1)	0.95	<i>Ua-20528</i>	B, T	10 935	80	No

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	334.55 (270.3)	0.95	<i>Ua-20529</i>	B, T	10 515	75	Yes
	333.25	1.75	UBA-20310	B	10 331	51	Yes
	332.7 (268.45)	1	<i>Ua-16740</i>	B	10 165	95	Yes
	331.6	1.5	UBA-20311	B	10 301	57	Yes
	329.5	0.5	UBA-23306	Bi	10 331	53	Yes
	328.5	0.5	UBA-20312	B	10 639	51	No
	328.4 (265.5)	1.05	<i>Ua-16745</i>	B, D	10 285	95	Yes
	326.5	0.5	UBA-20313	B	10 130	69	Yes
	323.35 (259.1)	1.15	<i>Ua-16747</i>	B	9860	85	No
	319.85 (257)	1.15	<i>Ua-16750</i>	B, C	10 205	85	Yes
	319.5	0.5	UBA-21574	B	10 102	67	Yes
	315.9 (255)	1.1	<i>Ua-16752</i>	B, Bp, P	9720	90	No
	312.25 (253)	1.1	<i>Ua-16761</i>	B, T	9955	90	Yes
	310	1	UBA-22308	Bi, M	9864	47	Yes
	309.5 (251)	1.15	<i>Ua-16766</i>	P	9765	85	Yes
	305.05 (246.75)	1.15	<i>Ua-16768</i>	P	9625	70	Yes
	302.5	0.5	UBA-21575	B	9532	59	Yes
	299.5	0.5	UBA-21577	B	9442	52	Yes
	298.5	0.5	UBA-21578	B	9416	47	Yes
	290	1	UBA-21576	B, C	7496	55	No
	280	1	UBA-23309	Bi	9129	50	Yes
	275	0.5	UBA-21579	T	8605	46	Yes

Table 4. Core ID, core depth and modelled and estimated ages for each core from Hässeldala Port (see Fig. 1C for the location of the cores and Fig. 3A for their TOC/LOI profiles). The statistical alignment of the TOC/LOI profiles of cores #1-HP4 and #6 to core #5 using a Monte Carlo algorithm (Muschitiello *et al.* 2015a, b) allowed transferring the chronology of core #5 to each of the profiles (Fig. 3B). The sedimentation rate in the uppermost/lowermost layers in some of the profiles had to be estimated (see main text), because these parts did not overlap with core #5: ¹ = sedimentation rate 23 a cm⁻¹; ² = sedimentation rate 22 a cm⁻¹; ³ = sedimentation rate 16 a cm⁻¹.

Core ID	Core depth (cm)	Depth (cm) interval for which the age could be modelled by correlation to core #5	Modelled median age (cal. ka BP)	Depth (cm) interval for which the age was interpolated using sedimentation rates	Estimated ages (cal. ka BP)
#1	217-316			217-225 ¹	9.4-10.1
			10.2-14.0	306-316 ²	14.0-14.2
#2	237-331	237.5-301.5	10.6-14.0	302.5-330.5 ³	14.0-14.5
				290.5-415 ³	14.0-14.4
#3	282-415	282.5-289.5	9.7-14.0		
#4	336-436	—	—	—	—
#HP4	245-345	245-329	11.7-14.0		
				330-345 ²	14.0-14.3
#5	270-370	278.5-369.5	9.8-14.1		
#6	320-412	320.5-410.5	10.9-14.1	410.5-411.5	14.1

Table 5. Clay-varve localities investigated in Blekinge province, southernmost Sweden (Björck 1981; Ringberg 1991; Wohlfarth *et al.* 1994; Ising 1998). The local varve year corresponds to the year when the respective site had become ice free and refers to the local varve chronology for Blekinge. The numbering of the sites follows Ringberg (1991). See Fig. 5A and B for the location of the sites.

Site #	Site name	Local varve year	References
146	Vieryd 1	-103	Ringberg (1991)
149	R Fornanäs 4	-102	Ringberg (1991)
151	R Angelskog 6	-100	Ringberg (1991)
154	R Folkets Hus 9	-86	Ringberg (1991)
155	R Persborg 10	-85	Ringberg (1991)
118	Sternö	-80	Ringberg (1991)
147	Hjälmseryd 2	-80	Ringberg (1991)
158	R Långkärra 13	-79	Ringberg (1991)
160	Torneryd 15	-79	Ringberg (1991)
148	Saxemara 3	-73	Ringberg (1991)
159	R Snäckebacken 14	-73	Ringberg (1991)
119	Munkahus	-72	Ringberg (1991)
120	Drösebro	-67	Ringberg (1991)
161	R Herstorp I 16	-65	Ringberg (1991)
162	R Silverforsen 17	-63	Ringberg (1991)
126	Skyekärr	-60	Ringberg (1991)
117	Stilleryd	-59	Ringberg (1991)
163	R Herstorp II 18	-52	Ringberg (1991)
128	Horsaryd	-49	Ringberg (1991)
167	R Sörby 22	-49	Ringberg (1991)
122	Gustafsborg	-48	Ringberg (1991)
124	Trensum	-48	Ringberg (1991)
164	R Herstorp III 19	-48	Ringberg (1991)
127	Karlshamn	-41	Ringberg (1991)
123	Häggarp	-41	Ringberg (1991)
	Korsliden	-39	Wohlfarth <i>et al.</i> (1994)
165	Kjettorp I 20	-37	Ringberg (1991)
168	Kallinge Järnvägen 23	-31	Ringberg (1991)
166	Trofta	-22	Ringberg (1991)
171	Härsjölund 26	-11	Ringberg (1991)
172	Tubbarp I 27	-11	Ringberg (1991)
	Farslycke	+10	Ising (1998)
	Trehörnan	+65	Wohlfarth <i>et al.</i> (1994)
	Hemsjön	+72	Wohlfarth <i>et al.</i> (1994)
	Kroksjön II	+77	Björck (1981), Wohlfarth <i>et al.</i> (1994)
	Skälgylet	+89	Wohlfarth <i>et al.</i> (1994)

Table 6. Plant species present in Blekinge, southernmost Sweden during the last glacial/interglacial transition according to pollen stratigraphic and plant macrofossil analyses (Björck 1981; Berglund 1966; Wohlfarth *et al.* 1994, 2006; Steinthorsdottir *et al.* 2013). Asterisk indicates finds of plant macrofossil remains.

Time interval	Plant species
>14.1 ka BP	<i>Artemisia</i> , * <i>Betula nana</i> , * <i>Betula pubescens</i> , <i>Calluna vulgaris</i> , <i>Caltha palustris</i> , * <i>Dryas octopetala</i> , <i>Empetrum nigrum</i> , <i>Ephedra</i> , <i>Helianthemum</i> , <i>Hippophaë rhamnoides</i> , <i>Juniperus</i> , <i>Plantago</i> , <i>Polygonum</i> , <i>Populus tremula</i> , <i>Rumex</i> , * <i>Salix polaris</i> , * <i>S. reticulata</i> , * <i>S. herbacea</i> , <i>Saxifraga</i> , <i>Urtica</i>
14.1 – 12.7 ka BP	<i>Artemisia</i> , <i>Astragalus alpinus</i> , * <i>Betula nana</i> , * <i>Betula pubescens</i> , <i>Calluna vulgaris</i> , <i>Cerastium alpinum</i> , * <i>Dryas octopetala</i> , <i>Dryopteris</i> , * <i>Empetrum nigrum</i> , <i>Ephedra</i> , <i>Helianthemum</i> , <i>Hippophaë</i> , <i>Isoëtes</i> , <i>Jasione montana</i> , * <i>Juniperus communis</i> , <i>Lycopodium</i> , <i>Lastrea</i> , <i>Myriophyllum alterniflorum</i> , <i>Plantago</i> , <i>Polygonum</i> , <i>Populus tremula</i> , <i>Potentilla palustris</i> , <i>Prunus padus</i> , <i>Pimpinella</i> , <i>Polypodium</i> , <i>Ranunculus</i> , <i>Rubus</i> , * <i>Salix polaris</i> , * <i>S. reticulata</i> , * <i>S. herbacea</i> , <i>Sanguisorba</i> , <i>Sorbus</i> , <i>Saxifraga</i> , <i>Typha latifolia</i> , <i>Urtica</i> , <i>Valeriana officinalis</i>
12.7 – 11.7 ka BP	<i>Artemisia</i> , <i>Astragalus alpinus</i> , * <i>Betula nana</i> , <i>Calluna vulgaris</i> , <i>Cerastium alpinum</i> , <i>Chamaenerion alpinum</i> , * <i>Dryas octopetala</i> , <i>Dryopteris</i> , <i>Empetrum nigrum</i> , <i>Ephedra</i> , <i>Helianthemum</i> , <i>Hippophaë</i> , <i>Jasione montana</i> , * <i>Juniperus communis</i> , <i>Lycopus europaeus</i> , <i>Lycopodium</i> , <i>Lastrea</i> , <i>Minuartia</i> , <i>Myriophyllum alterniflorum</i> , <i>M. spicatum</i> , <i>M. verticillatum</i> , <i>Oxytropis capestris</i> , <i>Oxyria digyna</i> , * <i>Pinus sylvestris</i> , <i>Plantago</i> , <i>Polygonum</i> , <i>Potentilla palustris</i> , <i>Populus tremula</i> , <i>Ranunculus</i> , <i>Rumex acetosella</i> , <i>Rubus</i> , * <i>Salix polaris</i> , * <i>S. reticulata</i> , * <i>S. herbacea</i> , <i>Sanguisorba</i> , <i>Sorbus</i> , <i>Saxifraga</i> , <i>Selaginella selaginoides</i> , <i>Urtica</i>
11.7 – 10 ka BP	<i>Artemisia</i> , <i>Astragalus alpinus</i> , * <i>Betula nana</i> , <i>Botanococcus umbellatus</i> , <i>Calluna vulgaris</i> , <i>Cerastium alpinum</i> , <i>Chamaenerion augustifolium</i> , <i>Caltha palustris</i> , * <i>Dryas octopetala</i> , <i>Dryopteris</i> , <i>Empetrum nigrum</i> , <i>Ephedra</i> , <i>Helianthemum</i> , * <i>Hippophaë</i> , <i>Isoëtes lacustris</i> , * <i>Juniperus communis</i> , <i>Littorella uniflora</i> , <i>Lycopodium</i> , <i>Lastrea</i> , <i>Menyanthes trifoliata</i> , <i>Myriophyllum alterniflorum</i> , <i>M. spicatum</i> , <i>M. verticillatum</i> , * <i>Pinus sylvestris</i> , <i>Polygonum</i> , <i>Populus tremula</i> , <i>Ranunculus</i> , <i>Rumex acetosella</i> , <i>Rubus fruticosus</i> , <i>Sanguisorba</i> , <i>Sorbus</i> , * <i>Salix polaris</i> , * <i>S. reticulata</i> , <i>Saxifraga</i> , <i>Typha latifolia</i> , <i>Urtica</i>

Table 7. Revised time scale for southern Sweden's regional Lateglacial pollen zones (Björck & Möller 1987) based on Hässeldala Port's pollen stratigraphies (Andersson 2004; Karlatou-Charampopoulou 2016) and chronology.

Regional pollen zone	Age assignment (cal. ka BP)
Younger Dryas/Preboreal transition zone	11.85-11.58
Younger Dryas	12.54-11.85
Alleröd/Younger Dryas transition zone	12.77-12.54
Alleröd II	13.24-12.77
Alleröd I	14.06-13.24
Older Dryas	>14.06

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