## 1 Last ice sheet recession and landscape emergence above sea level in east-central Sweden, evaluated

- 2 using in situ cosmogenic <sup>14</sup>C from quartz
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## 24 Abstract

25 In situ cosmogenic <sup>14</sup>C (*in situ* <sup>14</sup>C) in quartz provides a recently developed tool to date exposure of bedrock surfaces up to ~25,-000 years. From outcrops located in east-central Sweden, we test the 26 accuracy of in situ <sup>14</sup>C dating against (i) a relative sea level (RSL) curve constructed from radiocarbon 27 dating of organic material in isolation basins, and (ii) the timing of local deglaciation constructed from 28 a clay varve chronology complemented with traditional radiocarbon dating. Five samples of granitoid 29 30 bedrock were taken along an elevation transect extending southwestwards from the Baltic Sea coast near Forsmark. Because these samples derive from bedrock outcrops positioned below the highest 31 postglacial shoreline, they target the timing of progressive landscape emergence above sea level. In 32 33 contrast, in situ <sup>14</sup>C concentrations in an additional five samples taken from granitoid outcrops above the highest postglacial shoreline, located 100 km west of Forsmark, should reflect local deglaciation 34

35 ages. The ten in situ <sup>14</sup>C measurements provide robust age constraints that, within uncertainties,

- 36 compare favorably with the RSL curve and with the local deglaciation chronology. These data
- 37 demonstrate the utility of *in situ* <sup>14</sup>C to accurately date ice sheet deglaciation, and durations of
- 38 postglacial exposure, in regions where cosmogenic <sup>10</sup>Be and <sup>26</sup>Al routinely return complex exposure
- 39 results.

## 40 1. Introduction

- 41 The pacing of retreat of ice sheets in North America and Eurasia since their maximum expansion
- 42 during the last glaciation remains an active research field (e.g., Hughes et al., 2016; Stroeven et al.,
- 43 2016; Patton et al., 2017; Dalton et al., 2020, 2023). Understanding the triggers and processes causing
- 44 the demise of these ephemeral ice sheets yields the best blueprint for understanding the future
- 45 behavior of the Greenland and Antarctic ice sheets in a warming climate. Coupling the behavior of
- 46 deglaciating ice sheets over the course of the Late Glacial and early Holocene to increasingly precise
- 47 climate reconstructions, including and climatic events, requires increased precision in ice sheet
- 48 reconstructions (e.g., Bradwell et al., 2021). Increased Perecision can be achieved enhanced through a
- 49 coupling of geomorphological mapping of ice sheet margins (such as moraines, grounding zone
- 50 wedges, lateral meltwater channels, and ice-dammed lake shorelines and spillways) with numerical
- 51 field constraints from a diverse array of dating techniques (e.g., Stroeven et al., 2016; Bradwell et al.,
  2021; Regnéll et al., 2023).

53 Ice sheet reconstructions, especially in North America, have attained a high level of detailbecome

54 highly detailed through radiocarbon dating (Dyke et al., 2002; Dalton et al., 2020). With the advance of

- offshore imaging of glacial geomorphology (Greenwood et al., 2017, 2021; Bradwell et al., 2021),
- 56 radiocarbon dating has received a renewed upswing in recent years (e.g., Dalton et al., 2020; Bradwell
- 57 et al., 2021). However, large tracts of landscape\_areas lack radiocarbon age constraints on ice sheet
- 58 retreat simply due to because of an absence -lack of datable organic material. Fortunately, optically\_-
- 59 stimulated luminescence ages on buried sand layers (e.g., Alexanderson et al., 2022) and cosmogenic
- 60 nuclide apparent exposure ages on exposed bedrock and erratics have narrowed some of the gaps
- 61 (e.g., Hughes et al., 2016; Stroeven et al., 2016; Dalton et al., 2023). In studies using cosmogenic
- 62 nuclides, an 'apparent' exposure age is derived from a simple calculation from the nuclide
- 63 concentration under consideration (Lal, 1991; Gosse and Phillips, 2001). However, <u>C</u>eorrectly
- 64 interpreting the exposure age relies on modelling that considers geological factors that can reduce the
- 65 nuclide concentration relative to the time since initial subaerial exposure (such as erosion and burial by
- 66 glacial ice, water, snow, and/or soil; Gosse and Phillips, 2001; Schildgen et al., 2005; Ivy-Ochs and
- 67 Kober, 2008). Exposure dating is the only technique available in regions where ice sheet erosion has
- 68 left the surface bare or covered by a thin drape of till. Kleman et al. (2008) show that for Fennoscandia,

69 these conditions are widespread in coastal regions where ice accelerated towards its streaming sectors

- 70 and where wave wash during glacial rebound further thinned or removed pre-existing sediment
- 71 covers.

72 Coastal sectors in formerly glaciated regions provide sites important to the study of paleoglaciology. 73 They offer an abundance of bedrock exposures from which patterns and processes of subglacial 74 erosion can be studied through cosmogenic nuclide exposure dating (e.g., Hall et al., 2020). Also, 75 because of the interplay with postglacial sea level, coastal areas yield data on glacioisostatic rebound 76 that are critical to geodynamic modelling of Earth rheology and thicknesses of former ice sheets (e.g., 77 Lambeck et al. (1998, 2010) and Patton et al. (2017), for Fennoscandian examples). Geodynamic models require validation against measurements of vertical crustal motion (Steffen and Wu. 2011). 78 such as those provided by recent global positioning system (GPS) measurements (e.g., Lidberg et al., 79 2010) and postglacial records of crustal rebound afforded by relative sea level (RSL) curves (e.g., Påsse 80 and Andersson, 2005). The construction of RSL curves, detailing the history of land surface emergence 81 82 from sea level, is traditionally done using either sediments accumulated in isolation basins at different elevations above sea level or by dating uplifted gravel beach ridges. Typically, isolation basins, and their 83 sediments, show a progression from marine, to brackish, and finally to freshwater environments as 84 85 their bedrock sills they are uplifted through tidal levels (Long et al., 2011). Histories of land uplift above 86 sea level are documented using micro- and macrofossil analyses of isolation basin sediments and 87 radiocarbon dating on macrofossils (Romundset et al., 2011). Uplifted beach ridges can be radiocarbon 88 dated from a variety of materials (Blake, 1993) but most confidently from driftwood, whalebone, and shells (e.g., Dyke et al., 1992). Gravel beach ridges have also been investigated using OSL and <sup>10</sup>Be 89 90 exposure dating even though, other than the highest beach ridge, they may be prone to clast reworking (Briner et al., 2006: Simkins et al., 2013: Bierman et al., 2018). A distinct advantage of 91 92 constructing RSL curves using cosmogenic nuclides is that land surface emergence above sea level may 93 be additionally dated from boulders (Briner et al., 2006) or bedrock (Bierman et al., 2018). 94 The potential for cosmogenic surface exposure dating of last ice sheet retreat in recently glaciated low-95 relief cratonic landscapes would seemingly be high because of the frequent outcropping of glacially sculptured quartz-bearing crystalline bedrock. However, the ice sheet may have been either non-96 97 erosive or erosion was insufficiently deep to remove all the cosmogenic nuclide inventory from previous exposure periods. Apparent ages are therefore often older than indicated by radiocarbon 98 dating (Heyman et al., 2011; Stroeven et al., 2016) because they include a component of nuclide 99 100 inheritance. Apparent ages younger than indicated by radiocarbon dating can also occur if sampled 101 rock surfaces have been shielded, for example by sediments, following deglaciation. Concentrations of 102 <sup>10</sup>Be and <sup>26</sup>Al, in either bedrock or erratic boulders, therefore often reflect complex exposure histories 103 rather than simple deglacial exposure durations (Heyman et al., 2011; Stroeven et al., 2016).

In this study we use <sup>14</sup>C produced *in situ* in quartz-bearing bedrock (*in situ* <sup>14</sup>C) because it potentially
circumvents an overt reliance on the need for deep erosion (>-3 m) to remove the inherited signal from
previous exposure periods (Gosse and Phillips, 2001). The reason for this is that, <u>B</u>because of its short
half-life of 5700 ± 30 years, <u>nuclide inherited in situ</u> <sup>14</sup>Cance will have largely decayed away if ice sheet
burial at investigated sites during the last glacial phase (marine isotope stage 2; MIS2) exceeded 25-30
ka, that is, ca. 5 half-lives (Briner et al., 2014).
Some studies assessing changes in glacier and ice sheet extents over Late Glacial to Holocene

- 111 timescales have used *in situ* <sup>14</sup>C (Miller et al., 2006; Fogwill et al., 2014; Hippe et al., 2014;
- 112 Schweinsberg et al., 2018; Pendleton et al., 2019; Young et al., 2021; Schimmelpfennig et al., 2022). In
- 113 such-these studies, in situ <sup>14</sup>C has been applied with other nuclides with longer half-lives, in particular
- <sup>10</sup>Be, to unravel complex histories of glacier advance and retreat (e.g., Goehring et al., 2011) and
- 115 spatial patterns in glacial erosion in mountainous terrain (e.g., Steinemann et al., 2021). However,
- 116 <u>E</u>extensive regions formerly covered by ice sheets are characterized by low relief, and low elevation
- 117 terrain.<sub>5</sub> <u>Tand-the effectiveness of *in situ* <sup>14</sup>C in dating ice sheet retreat in these non-alpine settings and</u>
- 118 in quantifying shoreline displacement from bedrock samples has not been previously assessed. The
- 119 aim of this study is therefore to validate the use of <sup>14</sup>C formed *in situ* in bedrock as a reliable
- 120 chronometer by evaluating its performance in duplicating (i) a previously-established Holocene RSL
- 121 curve based on radiocarbon dating (Hedenström and Risberg, 2003; SKB, 2020) and (ii) the timing of
- 122 deglaciation above the highest (post-glacial) shoreline in nearby east-central Sweden according to
- 123 reconstructions of deglaciation of the last ice sheet (Hughes et al., 2016; Stroeven et al., 2016).
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## 125 2. Study Area

- Our study is focused on a region that includes low elevation, low relief, Forsmark-Uppland and 126 adjoining higher elevation and relief Dalarna-Gävleborg in east-central Sweden (Fig. 1). This region was 127 selected because Forsmark is the location of a planned geological repository for spent nuclear fuel 128 129 (e.g., SKB 2022). As such, this region has been intensively studied and has a wealth of and therefore also has abundant geologic data relevant to our study. This includes in-depth analyses of bedrock and 130 131 environmental properties, including influences of glacial and postglacial processes (e.g., Lönnqvist and Hökmark, 2013; Hall et al., 2019; Moon et al., 2020; SKB, 2020). 132 133 From spatio-temporal ice sheet reconstructions by Kleman et al. (2008), the study area was glaciated
- 134 16-20 times for a total duration of c. 330 kyr over the past 1 Ma. The last deglaciation of the study area
- 135 is well-constrained by two recent reconstructions that differ in their approach (Hughes et al., 2016;
- 136 Stroeven et al., 2016). The Hughes et al. (2016) reconstruction is explicitly based relies primarily upon
- 137 on-chronological constraints supplied <sub>3</sub>from radiocarbon, thermal luminescence, optically stimulated

138 luminescence (OSL), infrared stimulated luminescence, electron spin resonance, terrestrial cosmogenic 139 nuclide (TCN), and U-series dating. Published landform data, mostly with respect to end moraines and 140 generally accepted correlations of ice-margin positions between individual moraines, provide 141 complementary evidence. but In contrast, the Stroeven et al. (2016) reconstruction combines 142 geomorphological constraints for ice sheet margin outlines, including ice-marginal depositional 143 landforms and meltwater channels, ice-dammed lakes, eskers, lineations, and striae, -with chronological constraints supplied by radiocarbon, varve, OSL, and TCN dating. Whereas Hughes et al. 144 145 (2016) reconstruct ice sheet retreat every 1 ka, and for every ice margin plot its position as "most 146 credible", "min", and "max", Stroeven et al. (2016) present ice margin positions for every 100 years 147 inside the Younger Dryas standstill position (Stroeven et al., 2015). These marginal positions are 148 temporally and spatially defined by the "Swedish Time Scale" clay varve record along the Swedish east coast (De Geer, 1935, 1940; Strömberg, 1989, 1994; Brunnberg, 1995; Wohlfarth et al., 1995). From 149 150 Stroeven et al. (2016), the last deglaciation of the study area occurred 10.8  $\pm$  0.3 ka BP, which overlaps 151 the timing of deglaciation of the study area from Hughes et al. (2016), within uncertainty (Fig. 1). The 152 highest postglacial shoreline in east-central Sweden is located at a present elevation of ~200 m a.s.l. in Dalarna-Gävleborg, ~100 km west of Forsmark (SGU, 2015). The exposure duration of bedrock above 153 154 the highest postglacial shoreline therefore represents the time since local deglaciation. Hence, in situ 155 <sup>14</sup>C ages from bedrock above the highest postglacial shoreline should conform to the reconstructed 156 deglaciation age of  $10.8 \pm 0.3$  ka from Stroeven et al. (2016). 157 Below the highest postglacial shoreline, in the Forsmark-Uppland region, the last deglaciation occurred in a marine environment and the landscape has progressively emerged above sea level 158 159 through postglacial isostatic uplift. A RSL curve constructed from radiocarbon dating of basal organic 160 sediments trapped in isolation basins along elevation transects describes the progressive emergence 161 of the Forsmark-Uppland landscape above sea level (Robertsson and Persson, 1989; Risberg, 1999; 162 Bergström, 2001; Hedenström and Risberg, 2003; Berglund, 2005; SKB, 2020). Ages calculated from in situ <sup>14</sup>C from bedrock outcrops along an elevation transect would then mirror the Forsmark RSL curve 163 164 for their corresponding elevations (but be slightly older because of nuclide production through shallow water before emergence). 165 166 A potential complication to the accurate exposure age dating of bedrock surfaces using in situ <sup>14</sup>C in 167 east-central Sweden is that the most recent period of ice sheet burial may not have been sufficiently 168 long to decay the any in situ <sup>14</sup>C inventory inherited from preceding prior exposure. Here, the extent of the Fennoscandian Ice Sheet during interstadial MIS3 and the timing of ice advance across the 169 170 Forsmark region during late MIS3 are crucially important. Kleman et al. (2020) have identified ice-free conditions around Idre (330 km NW, up-ice, of our study area; Fig. 1) between 55 ka and 35 ka, which 171

172 implies inundation of our study area by ice after 35 ka. Combined with a well-constrained final

173 deglaciation age of 10.8±0.3 ka (Stroeven et al. 2016), it appears that our study area has most recently

174 (during MIS2) been inundated by glacial ice for at most 24 ka. This inference is in line with results from

ice sheet modelling indicating a 22 kyr duration of ice-cover at Forsmark during MIS2 (SKB, 2020).

176 Consequently, it is possible that in situ <sup>14</sup>C concentrations may reflect subaerial exposure of bedrock in

177 our study area during MIS3 in addition to Holocene exposure, resulting in an offset towards older ages

178 relative to the RSL curve for Forsmark (Hedenström and Risberg, 2003; SKB, 2020) and the deglaciation

179 chronologies of Hughes et al. (2016) and Stroeven et al. (2016).

180

# 181 3. Methods

# 182 3.1. Sampling of bedrock outcrops for *in situ* <sup>14</sup>C measurement

We used the following sampling strategy to evaluate the accuracy of bedrock exposure ages derived 183 184 from in situ <sup>14</sup>C against the Forsmark RSL curve and the deglaciation of the last ice sheet in east-central Sweden. A rigorous scheme was applied to ensure that we avoided sampling quartz altered through 185 hydrothermal processes that is likely to occur in major pegmatite intrusions, outcrops located in major 186 deformation zones, and outcrop-scale veins, fractures, and adjacent rock volumes. Consequently, 187 sampling was done on outcrops of metagranitoid from the early-Svecokarelian GDG-GSDG suite that 188 dominates the Bergslagen lithotectonic unit (Stephens and Jansson, 2020). A petrological examination 189 190 using transmitted light polarization microscopy was applied to thin sections to ascertain that the quartz was unlikely to contain multi-fluid phase, vapour phase, or solid-phase inclusions. All samples were 191 192 collected using an angle grinder, which permits sampling of hard crystalline bedrock isolated from 193 outcrop edges, fractures, and quartz veins, and consistently limits sample thicknesses to 3 cm.

194 We collected a total of ten samples for in situ <sup>14</sup>C analyses. Five of these were collected along a SW-NE 195 transect near Forsmark (Fig. 1b). These outcrops were chosen because they span an elevation gradient of 9.4-56.0 m a.s.l. and exposure ages derived from in situ <sup>14</sup>C can therefore be evaluated against the 196 Forsmark RSL curve. We collected a further five samples from locations above the highest shoreline (Fig. 197 1a) to determine the age of local deglaciation for comparison with published deglaciation chronologies 198 199 (Hughes et al., 2016; Stroeven et al., 2016). Sample locations were logged on a 2 m-resolution LiDAR 200 digital elevation model (DEM) displayed in ArcGIS 10 on a tablet computer. A GPS add-in tool in ArcGIS 201 10 was used to record positional data, within a horizontal precision of 2 m. The elevation of each sample 202 location was extracted from the DEM and has a precision of tens of centimetres. The influence of these 203 minor positional uncertainties on our <sup>14</sup>C calculations is trivial and none of the sample sites is influenced

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204 by topographic shielding that could reduce the accumulation of <sup>14</sup>C in bedrock.

205 Each sampled bedrock outcrop formed a local topographic high, which minimizes the risk of burial by

soil and snow (Supplement 1). Moss mats were present on all sampled outcrops. Although we avoided

207 sampling bedrock that was moss-covered, we cannot be certain that moss mats did not formerly cover

208 the sample sites. Given a compressed thickness of 0.5 cm and an estimated density of 0.7 g f cm<sup>-3</sup>, this

may have contributed to a shielding of the sampled rock surfaces of 0.35 g f cm<sup>-2</sup>, which is negligible and

210 is therefore excluded from our age inferences.

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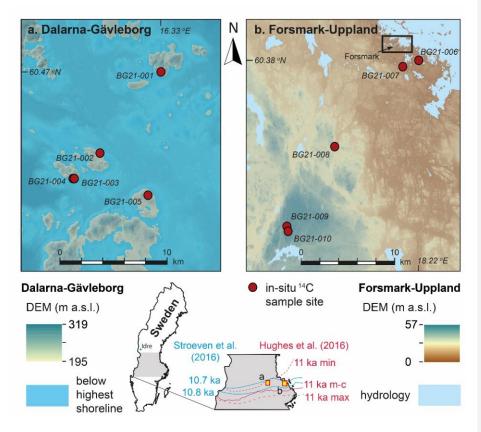


Figure 1. Sample locations for in situ <sup>14</sup>C dating in (a) Dalarna-Gävleborg and (b) Forsmark-Uppland. The 212 five Dalarna-Gävleborg sample sites are located on what were islands above the highest postglacial 213 shoreline (shown), whereas the five sample sites from Forsmark-Uppland are located below the highest 214 215 shoreline (not shown because the entire area was submerged). See inset maps for locations of panels a and b and for the 10.7 ka BP and 10.8 ka BP retreat isochrones (blue) from Stroeven et al. (2016) and 11 216 217 ka BP (most-credible, minimum, and maximum) retreat isochrones (red) from Hughes et al. (2016). The 218 rectangle in panel b approximately indicates the site selected for the planned geological repository for spent nuclear fuel at Forsmark. DEM with 2 m resolution, from LiDAR data, Lantmäteriet. 219

## 220 3.2. Laboratory preparation for accelerator mass spectrometry (AMS)

221 Samples were physically and chemically processed at the Purdue Rare Isotope Measurement Laboratory (PRIME Lab) at Purdue University, U.S.A. Concentrations of in situ <sup>14</sup>C were determined from purified 222 223 quartz separates through automated procedures (Lifton et al., 2023). Approximately 5 g of quartz from 224 each sample was added to a degassed LiBO<sub>2</sub> flux in a re-usable 90% Pt/10% Rh sample boat and heated 225 to 500 °C for one hour in ca. 6.7 kPa of Research Purity O<sub>2</sub> to remove atmospheric contaminants, which 226 were discarded. The sample was then heated to 1100 °C for three hours to dissolve the quartz and release the in situ <sup>14</sup>C, again in an atmosphere of ca. 6.7 kPa of Research Purity O<sub>2</sub> to oxidize any evolved 227 228 carbon species to CO2. The CO2 from the 1100 °C step was then purified, measured quantitatively, and converted to graphite for <sup>14</sup>C AMS measurement at PRIME Lab (Lifton et al., 2023). To test for data 229 reproducibility, sample BG21-002 was randomly selected to undergo laboratory preparation and AMS a 230 second time. Measured concentrations of in situ<sup>14</sup>C are calculated from the measured isotope ratios via 231 AMS following Hippe and Lifton (2014) (Table 1). 232

233 **3.3. Exposure age calculations** 

234 The expage calculator version 202403312 (http://expage.github.io/calculator) is used to calculate apparent exposure ages. It is based on the original CRONUS calculator v. 2 (Balco et al., 2008), the LSDn 235 236 production rate scaling (Lifton et al., 2014), and the CRONUScalc calculator (Marrero et al., 2016), using the geomagnetic framework of Lifton (2016) with the SHA.DIF.14k model for the last 14 kyr. Exposure 237 ages are calculated using resulting time-varying <sup>14</sup>C production rates accounting for decay and 238 interpolated to match the measured <sup>14</sup>C concentration. The production rate from muons is calibrated 239 against the Leymon High core <sup>14</sup>C data of Lupker et al. (2015) and the production rate from spallation is 240 calibrated against updated global <sup>14</sup>C production rate calibration data (Schimmelpfennig et al., 2012; 241 Young et al., 2014; Lifton et al., 2015; Borchers et al., 2016; Phillips et al., 2016; Koester and Lifton, 2023. 242 corrigendum in prep). This calibration is done iteratively for spallation and muons to reach convergence, 243 using the expage production rate calibration methods (Fig. 2). 244

Exposure age calculations along the Forsmark-Uppland transect account for <sup>14</sup>C production during emergence through shallow water. However, Bburial of sampled surfaces by snow is excluded from the age calculations for all sample sites because we neither know how snow burial depths and durations vary between sites nor vary through time. The effect of snow burial would be to slightly decrease cosmogenic nuclide production in the underlying rock surface (Schildgen et al., 2005) and we have minimized this effect through our sampling strategy.

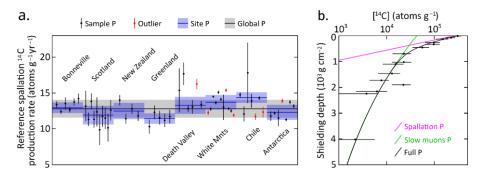


Figure 2. Production rate calibration of <sup>14</sup>C in quartz. (a) Reference spallation <sup>14</sup>C production rate 251 calibration based on data from Schimmelpfennig et al. (2012), Young et al. (2014), Lifton et al. (2015), 252 Borchers et al. (2016), and Phillips et al. (2016), corrected per Hippe and Lifton (2014) and compiled in 253 Koester and Lifton (2023). An uncertainty-weighted production rate is calculated for each of the eight 254 sites. Outliers, which are not included in the uncertainty-weighted production rates, are determined 255 256 based on the requirement that there should be at least three samples yielding a reduced chi-square statistic  $(X_R^2)$  with a p-value of at least 0.05 for the assumption that the individual production rates from 257 a site are derived from one normal distribution. For  $X_R^2$ , but not the uncertainty-weighting, we use the 258 259 largest of the sample-specific production rate uncertainty based on the <sup>14</sup>C concentration uncertainties 260 and 5% of the sample production rate. This procedure does not punish samples with low measurement 261 uncertainties, which otherwise risk exclusion as outliers. We adopt a global reference spallation <sup>14</sup>C 262 production rate of 13.3512.81 ± 1.131.25 atoms g<sup>-1</sup> yr<sup>-1</sup>, calculated as the arithmetic mean of the eight 263 site production rates with the uncertainty being based on an uncertainty-weighted deviation of all 264 included single sample production rates, excluding outliers. (b) Calibration of <sup>14</sup>C production rate from muons based on the data of Lupker et al. (2015). The calibration is based on the method used in the 265 266 CRONUScalc calculator (Marrero et al., 2016; Phillips et al., 2016). The figure shows the best fit <sup>14</sup>C concentration profiles produced from spallation, slow muons, and full production. The best fit yields 267 268 near zero production from fast muons (cf. Lupker et al., 2015). The production rate calibration has been 269 carried out using the expage-202403306 calculator in an iterative way to make the global reference 270 spallation <sup>14</sup>C production rate converge with the production rate from muons.

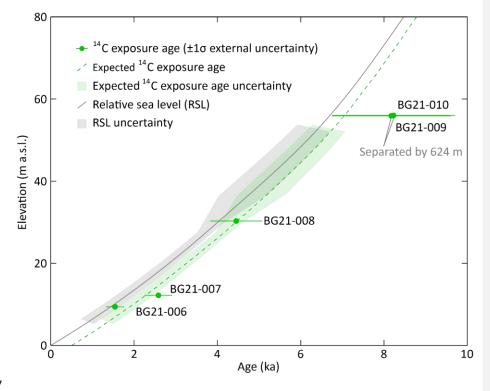
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## 272 4. Results

| 273 | Analytical results for in situ <sup>14</sup> C samples and procedural blanks are presented in Table 1. The mean and      |
|-----|--|
| 274 | standard deviation are used to correct measured <sup>14</sup> C sample inventories (Table 1) because procedural          |
| 275 | blanks are well-constrained during the analytical time frame. Inferred ages for the five in situ <sup>14</sup> C samples |

276 from the Forsmark-Uppland transect (i.e., below the highest postglacial shoreline) are shown relative to

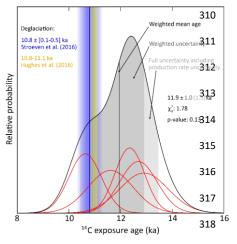
the Holocene RSL curve for Forsmark and the expected in situ <sup>14</sup>C exposure age curve considering 277 278 subaqueous cosmogenic nuclide production (Figure 3; Tables 1 and 2). Exposure age uncertainties are 279 large with internal uncertainties (measurement uncertainties; Balco et al., 2008) of 5-9% and external uncertainties of <u>12-2013-25</u>% (also including production rate uncertainties, which are high relative to 280 <sup>10</sup>Be (Borchers et al., 2016; Phillips et al., 2016). Apparent exposure ages increase consistently with 281 elevation and match expected ages within uncertainty. The two highest samples have near-identical 282 283 apparent exposure ages and elevations. However, these samples provide independent ages because 284 they are horizontally separated by 624 m (Figure 1b). There is good agreement between ages inferred 285 from these in situ <sup>14</sup>C data and the RSL curve constructed from organic radiocarbon dating of isolation 286 events (Hedenström and Risberg, 2003; SKB, 2020).



**Figure 3**. Apparent <sup>14</sup>C exposure ages for five Forsmark samples from below the highest shoreline (Fig. 1b; Table 2) with 1 $\sigma$  external uncertainties. The expected exposure ages are calculated assuming the RSL curve is correct, the <sup>14</sup>C spallation production rate is correct, partial exposure as the sample approaches the water surface, and full post-glacial exposure for the duration above sea level. Hence, the expected exposure age curve is a few hundred years older than the RSL curve. The RSL curve is from SKB (2020) and uncertainties for the 1–6 ka interval are calculated from the original radiocarbon data in Hedenström

and Risberg (2003). The RSL uncertainty envelope is also transposed onto the expected exposure age curve.

Apparent exposure ages for the five in situ <sup>14</sup>C samples located above the highest shoreline in Dalarna 296 297 and Gävleborg (Fig. 1a) are shown in Figure 4 and Table 2. The weighted mean age from all five samples 298 is  $11.92 \pm 1.53$  ka. These data display a  $X_p^2$  of 1.78 and a p-value of 0.13 based on 1 $\sigma$  internal uncertainties (Fig. 4a), which does not support a rejection of the hypothesis that the apparent exposure ages represent 299 300 the same population. In addition to the samples being from the same population, the exposure ages are consistent, within uncertainty, with the expected deglaciation age of  $10.8 \pm 0.3$  ka (Stroeven et al. 2016). 301 302 Replicate measurements on sample BG21-002 closely agree and an age based on a weighted mean <sup>14</sup>C 303 concentration is shown in Figure 4. Sample BG21 001 provides the youngest apparent age but, because 304 this sample was from a low profile outcrop (Supplement 1), this age may reflect partial shielding of the sampled bedrock surface by a past shallow soil cover or perhaps a deeper snow cover than the other 305 sites. We therefore consider this sample as least likely to provide a reliable age. Removing this sample 306 307 from consideration indicates that the remaining four sample sites are more clustered, with an older weighted mean age of  $11.612.4 \pm 1.31$  ka, which displays a  $X_{\pm}^{\pm}$  of 0.423 and a p-value of 0.743 based on 308 309 1σ internal uncertainties (Fig. 4b).



**Figure 4.** <u>Summed Pprobability density plotsdistributionsNormalized kernel density estimates</u> of the exposure ages from samples above the highest shoreline (Fig. 1a; Table 2). The individual samples (red curves) display 1 $\sigma$  internal uncertainty (measurement uncertainty). For the repeat sample BG21-002, the exposure age is calculated with a weighted mean <sup>14</sup>C concentration using a 2% uncertainty. <del>(a)</del> The probability density and data for all five samples. For the full set of samples, <u>T</u>the cosmogenic nuclide ages yield a reduced chi-square ( $X_R^2$ ) of 1.78 and a p-value of 0.13 based on internal uncertainties, which indicates that they are from the same population. <del>(b)</del> The probability density and data with sample

- 326 BG21-001 excluded as an outlier. These cosmogenic nuclide ages yield a  $X_{\pm}^2$  of 0.423 and a p-value of
- 327 0.743 based on internal uncertainties, which again indicate that they are from the same population. All
- 328 ages are referenced to the sampling year 2021. The weighted ages of  $11.92 \pm 1.53$  ka and  $11.612.4 \pm 1.532$  ka and 11.612.532 ka and 11.612.532 ka and 11.612.532 ka and 11.612.532
- 329 <u>1.31 ka both overlap with the deglaciation age from Stroeven et al. (2016).</u>

|                 | 5 Measure   |                |              |   | Notes |                           |   | PB2-06022022       |                                   | PB2-04212022                          | PB2-03222022            | Procedural Blanks | <u>BG21-010</u>     | BG21-009            | BG21-008            | BG21-007            | BG21-006            | BG21-005            | BG21-004            | BG21-003            | BG21-002R           | BG21-002            | BG21-001            | Sample  | Table 1. <i>In situ</i> <sup>1</sup>       | <b>BG21-006</b> 4 | <b>BG21-005</b> 4                             | <b>BG21-004</b> 4                             | <b>BG21-002R</b> 4  | <b>BG21-003</b> 4          | <b>BG21-002</b> 4   | <b>BG21-001</b> 4   | Sample ID PCE                                    | Xtra                             |  |
|-----------------|---|----------------|--------------|---|-------|---------------------------|---|--------------------|-----------------------------------|---------------------------------------|-------------------------|-------------------|---------------------|---------------------|---------------------|---------------------|---------------------|---------------------|---------------------|---------------------|---------------------|---------------------|---------------------|---|--|-------------------|---|---|---------------------|----------------------------|---------------------|---------------------|--|----------------------------------|--|
| alues calculat  | d relative to O   | Vienna Peede   | <u>b ID.</u> | Purdue Carbon Extraction and Graphitization System. |       |                           |   | PCEGS-169          |                                   |                                       | PCEGS-135               |                   | PCEGS-161           | PCEGS-160           | PCEGS-158           | PCEGS-157           | PCEGS-155           | PCEGS-153           | PCEGS-152           | PCEGS-148           | PCEGS-150           | PCEGS-147           | PCEGS-146           | PCEGS <sup>1</sup> #  | <sup>14</sup> C sample measurement details | 155 202101965     | <del>153</del> <u>20210196</u> 4              | <u>452</u> <u>20210196</u> 3                  | 150 202201473       | 148 202101962              | 147 202101961       | 146 202101960       | PCEGS #" PLID"                                   | on and meas                      |  |
| ed using Dilu   | X-2 standard  | e Belemnite.   |              | ion and Grap  |       |                           |   | 202201459          | 202201434                         | 202201452                             | 202201450               |                   | 202101969           | 202101968           | 202101967           | 202101966           | 202101965           | 202101964           | 202101963           | 202101962           | 202201473           | 202101961           | 202101960           |   | leasuremen                                 | 5.0               | <u>5</u>                                      | 963<br>51<br>natt                             | <u>5</u>            | <del>5</del> .             | 5                   | 5.                  | Q  | uremer                           |  |
| Ited Mass C a   | -<br>hitization bla   |                |              | hitization Sy                                       |       |                           |   | l)                 | l:                                | 1                                     | l)                      |                   | 4.99961             | 5.01906             | 5.07653             | 5.03589             | 5.06572             | 5.07578             | 5.05927             | 5.01070             | 5.04116             | 5.02383             | 5.02378             | Mass<br>Quartz<br>(g)                                       | it details                                 | 2 5.5 ± 0.1       | 3 4.6 ± 0.1                                   | 7 <u>11.9 ± 0.2</u>                           | } 7.7 ± 0.1         | ) <u>17.6 ± 0.3</u>        | 3 7.8 ± 0.1         | 3 <u>5.0 ± 0.</u> 1 | g) (µg)  | in situ <sup>14</sup> C at PRIME |  |
| and corrected   | nk (hased on  | quot (ca. a p  |              | stem.   |       |                           |   | $2.3 \pm 0.1$      | $\frac{2.3 \pm 0.1}{2.3 \pm 0.1}$ | $\frac{1.8 \pm 0.1}{2.2 \pm 0.4}$     | $\frac{1.4 \pm 0.1}{2}$ |                   | 42.2 ± 0.6          | $55.3 \pm 0.7$      | $4.0 \pm 0.1$       | $6.9 \pm 0.1$       | $5.5 \pm 0.1$       | $4.6 \pm 0.1$       | $11.9 \pm 0.2$      | $17.6 \pm 0.3$      | $7.7 \pm 0.1$       | $7.8 \pm 0.1$       | $5.0 \pm 0.1$       | <u>C yield</u><br>( <u>µg)</u>                              | 4  | 306.8 ± 3.7       | 304.5 ± 3                                     | <del>305.7 ± 3</del>                          | 305.3 ± 3           | 303.4 ± 3                  | 303.3 ± 3           | 393.8±4             | ( <i>ju</i> ted<br>( <i>ju</i> ted)<br>(line: 0) | d                                |  |
| for mean prid   | AMS Split M   |                |              |   |       |                           |   | 307.3 ± 3.7        | <u>307.4 ± 3.7</u>                | $\frac{307.0 \pm 3.7}{207.4 \pm 3.7}$ | $305.2 \pm 3.7$         |                   | 306.0 ± 3.7         | $305.6 \pm 3.7$     | 308.9 ± 3.8         | 309.2 ± 3.8         | 306.8 ± 3.7         | $304.5 \pm 3.7$     | $305.7 \pm 3.7$     | 303.4 ± 3.7         | $305.3 \pm 3.7$     | 303.3 ± 3.7         | 393.8 ± 4.8         | Diluted<br>Mass C<br>(µg)                                   |  | I INC             |   |   |                     |                            |                     |                     |  | 25 cm                            |  |
| ocedural blan   | ace (C) and st  |                | )<br>)       |   |       |                           |   | <u>298.3 ± 3.6</u> | 290.4 ± 3.0                       | 298.0 ± 3.0                           | <u>296.2 ± 3.6</u>      |                   | 297.0 ± 3.6         | $296.6 \pm 3.6$     | 299.9 ± 3.6         | <u>300.1 ± 3.7</u>  | 297.8 ± 3.6         | 295.6 ± 3.6         | 296.8 ± 3.6         | 294.5 ± 3.6         | 296.4 ± 3.6         | 294.4 ± 3.6         | 382.3 ± 4.6         | AMS Split<br>Mass C <sup>3</sup><br>(µg)                    |  | -45.2 ± 0.2       | -45.4 ± 0.2                                   | -44.6 ± 0.2                                   | $-45.2 \pm 0.2$     | <del>-43.9 ± 0.2</del>     | $-44.8 \pm 0.2$     | $-45.9 \pm 0.2$     | <b>8</b> 13 <b>С</b> с<br>(‰vрав)                |                                  |  |
| k (All blanks). | Measured relative to OX-2 standard.<br>Corrected for mass-dependent graphitization blank (based on AMS Split Mass C) and stable C composition | iupic analysis |              |   |       |                           |   | -40.3 ± 0.2        |                                   |                                       |                         |                   | $-40.1 \pm 0.2$     | -38.0 ± 0.2         | 6 -45.4 ± 0.2       | -45.0 ± 0.2         | -45.2 ± 0.2         | -45.4 ± 0.2         | -44.6 ± 0.2         | -43.9 ± 0.2         | -45.2 ± 0.2         | -44.8 ± 0.2         | -45.9 ± 0.2         | VPDB <sup>4</sup> )   |  | 0.1277±0.0056     | 0.4600 ± 0.0127                               | 0.4618 ± 0.0079                               | 0.4558 ± 0.0135     | 0.4633 ± 0.0108            | 0.4555 ± 0.0096     | $0.3399 \pm 0.0075$ | ""C/10C (10-11)                                  |                                  |  |
|                 | sition  |                |              |   |       | Mean                      | M | 0.4920 ± 0.0291    | 0.3364 ± 0.0315                   | 0.5364 ± 0.0215                       | $0.4853 \pm 0.0298$     |                   | 3.3197 ± 0.0680     | $3.3393 \pm 0.0946$ | $2.3565 \pm 0.0634$ | $1.6838 \pm 0.0507$ | $1.2766 \pm 0.0562$ | $4.5997 \pm 0.1272$ | $4.6181 \pm 0.0789$ | $4.6325 \pm 0.1075$ | $4.5575 \pm 0.1350$ | $4.5548 \pm 0.0964$ | $3.3992 \pm 0.0745$ | 14C/ <sup>13</sup> C <sup>6</sup>                           |  | 0.1172 ± 0.0059   | 0.4667 ± 0.0134                               | 0.4691 ± 0.0083                               | 0.4624 ± 0.0142     | 0.4709 ± 0.0113            | 0.4623 ± 0.0102     | $0.3412 \pm 0.0079$ | " <b>"C/C</b> iotal (70"')                       |                                  |  |
|                 |   |                |              |   |       | viean ± 1σ (145,163 only) |   | 0.3486 ± 0.0312    | 0.0922 ± 0.0000                   | 0.3037 ± 0.0292                       | $0.3413 \pm 0.0320$     |                   | 3.3399 ± 0.0721     | 3.3681 ± 0.1005     | $2.3076 \pm 0.0669$ | $1.6007 \pm 0.0536$ | $1.1715 \pm 0.0594$ | 4.6668 ± 0.1339     | $4.6905 \pm 0.0832$ | $4.7091 \pm 0.1134$ | $4.6239 \pm 0.1422$ | 4.6226 ± 0.1016     | 3.4118 ± 0.0785     | <sup>14</sup> C/C <sub>totaf</sub><br>(10 <sup>-14</sup> )  |  | 0.1243 ± 0.0101   | $0.6566 \pm 0.0225$                           | $0.6630 \pm 0.0159$                           | $0.6519 \pm 0.0237$ | <del>0.6604 ± 0.0197</del> | $0.6470 \pm 0.0181$ | $0.6177 \pm 0.0179$ | ""C (10° at)                                     |                                  |  |
|                 |   |                |              |   |       | <u>1 U.5894 ± U.UZ14</u>  |   |                    |                                   |                                       |                         |                   | 4.5648 ± 0.1321     | 4.6013 ± 0.1703     | $3.0145 \pm 0.1185$ | $1.9221 \pm 0.0960$ | $1.2426 \pm 0.1010$ | 6.5656 ± 0.2251     | $6.6300 \pm 0.1588$ | $6.6042 \pm 0.1969$ | 6.5186 ± 0.2368     | 6.4703 ± 0.1806     | $6.1771 \pm 0.1793$ | <sup>14</sup> ℃7<br>(1 <mark>0<sup>5</sup> at</mark> )      |  | 0.2453 ± 0.0199   | $\frac{1.2935 \pm 0.0444}{1.2935 \pm 0.0444}$ | $\frac{1.3105 \pm 0.0314}{1.3105 \pm 0.0314}$ | $1.2931 \pm 0.0470$ | <del>1.3180 ± 0.0393</del> | $1.2879 \pm 0.0360$ | 1.2296 ± 0.0357     | [""C] ( <i>10º at g-)</i> "                      |                                  |  |
|                 |   |                |              |   |       |                           |   | h                  | l;                                | i li                                  | h                       |                   | $0.9130 \pm 0.0264$ | $0.9168 \pm 0.0339$ | 0.5938 ± 0.0234     | $0.3817 \pm 0.0191$ | $0.2453 \pm 0.0199$ | $1.2935 \pm 0.0444$ | $1.3105 \pm 0.0314$ | $1.3180 \pm 0.0393$ | $1.2931 \pm 0.0470$ | $1.2879 \pm 0.0360$ | $1.2296 \pm 0.0357$ | [ <sup>14</sup> C]<br>(10 <sup>5</sup> at g <sup>-1</sup> ) |  |                   |   |   |                     |                            |                     |                     |  |                                  |  |

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Table 2-2. Apparent i-4n situ <sup>14</sup>C ages from quartz, Dalarna-Gävleborg and Forsmark-Uppland.

| Sample <sup>11D</sup> | Lat<br>(°) | Long<br>(°) | Elevation<br>(m a.s.l.) | <sup>14</sup> C <u>aAge ± Unc.<sup>Ext.</sup></u><br>(± Unc. <sup>Int.</sup> ) <sup>2a</sup> (ka) |
|-----------------------|------------|-------------|-------------------------|---|
| BG21-001              | 60.47432   | 16.33134    | 236.5                   | 10.6 ± 2.2 (± 0.6)  |
| BG21-002              | 60.40615   | 16.22197    | 212.6                   | 12.3 ± 2.9 (± 0.8)  |
| BG21-002R             | 60.40615   | 16.22197    | 212.6                   | 12.4 ± 3.0 (± 1.1)  |
| BG21-003              | 60.38459   | 16.17649    | 216.3                   | 12.9 ± 3.2 (± 0.9)  |
| BG21-004              | 60.38451   | 16.17440    | 217.8                   | 12.7 ± 3.0 (± 0.7)  |
| BG21-005              | 60.36888   | 16.30526    | 248.1                   | 11.6 ± 2.6 (± 0.9)  |
| BG21-006              | 60.38490   | 18.22308    | 9.4                     | 1.5 ± 0.2 (± 0.1)   |
| BG21-007              | 60.37892   | 18.19129    | 12.2                    | 2.6 ± 0.3 (± 0.2)   |
| BG21-008              | 60.30504   | 18.04993    | 30.3                    | 4.5 ± 0.6 (± 0.2)   |
| BG21-009              | 60.22988   | 17.94989    | 56.0                    | 8.2 ± 1.5 (± 0.5)   |
| BG21-010              | 60.22431   | 17.95051    | 55.9                    | 8.2 ± 1.4 (± 0.4)   |

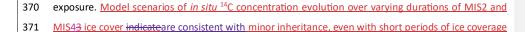
## Notes

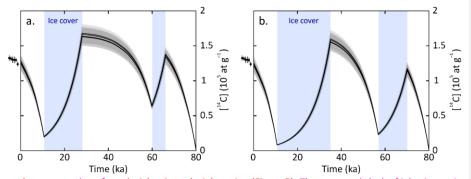
All samples have a thickness of 3 cm, a density of 2.7 g cm<sup>-3</sup>, and a shielding factor of 1. Zero erosion is assumed. 2 <sup>e-14</sup>C aAge- and 1σ external uncertainty (1σ internal uncertainty).<del>and Unc.<sup>Ent</sup> is external uncertainty and Unc.<sup>Int</sup> is (finternal uncertainty). Both are 1σ.</del>

#### 336 5. Discussion

The in situ <sup>14</sup>C bedrock exposure ages from the Forsmark-Uppland transect (i.e., below the highest 337 338 postglacial shoreline) consistently increase with elevation and overlap the expected exposure age curve, within uncertainty (Fig. 3). Because-the-apparent exposure ages accurately reflect the timing of 339 landscape emergence, in situ-<sup>14</sup>C is indicated as having high potential as a chronometer over Late 340 341 Glacial Holocene timescales in low relief, low elevation settings. This study adds to precious few demonstrations of the ability of applications of cosmogenic nuclides isotopes to defininge postglacial 342 343 landscape emergence above sea level (Briner et al., 2006; Bierman et al., 2018). Briner et al. (2006) 344 present good (visual) congruence with a record of shoreline emergence built from radiocarbon-dated driftwood and fauna by Dyke et al. (1992) using <sup>10</sup>Be measurements on boulders in beaches derived 345 from wave-washed till. Their study also mentions that building a relative sea level curve from pebbles, 346 cobbles and plucked bedrock suffered from inheritance problems, an experience shared by Matmon et 347 al. (2003) while attempting the dating of chert on beach ridges in southern Israel and heeded by 348 349 Bierman et al. (2018). Bierman et al. (2018) successfully dated landscape emergence on Greenland using <sup>10</sup>Be across a range of settings, including bedrock below the highest shoreline, cobbles from beach 350 351 ridges at the highest shoreline, and boulders and bedrock above the highest shoreline. They note that 352 success hinges on the requirement of warm-based ice and deep glacial erosion in exposing bedrock devoid of an inherited cosmogenic nuclide inventory. In many regions, however, including east-central 353 Sweden and more widely in Fennoscandia, these requirements are not met either because of cold-354 based conditions (Patton et al., 2016; Stroeven et al., 2016) or weakly erosive warm-based ice such as 355 at Forsmark (Hall et al., 2019; SKB, 2020), during all or much of glacial time. Cosmogenic nuclide 356 inheritance is therefore a part of the landscape fabric. Bierman et al. (2018) advocate the use of in situ 357 358 <sup>14</sup>C as a methodology to circumvent inheritance problems. Our study is the first to follow-up on that 359 suggestion, and shows, convincingly, that using in situ <sup>14</sup>C can extend the study of landscape rebound to regions where ice sheet erosion was insufficiently deep to allow for the application of long-lived 360 361 nuclides.

362 Five bedrock samples from above the highest postglacial shoreline are well-clustered and the weighted mean age (and full uncertainty) of  $11.92 \pm 1.53$  ka overlaps with the predicted deglaciation age of 10.8 363 364 ± 0.3 ka (Fig. 4a; Hughes et al., 2016; Stroeven et al., 2016). Removing the youngest age from 365 consideration results in more strongly clustered ages (Fig. 4b) and an older mean weighted age of 11.612.4 ± 1.31 ka, which still overlaps the predicted deglaciation age, within uncertainty. We therefore 366 do not further discriminate between these results. Because derived exposure ages overlap with the 367 predicted deglaciation age, we further infer that the in situ <sup>14</sup>C samples, including those located below 368 the highest postglacial shoreline, within uncertainty, lack significant inheritance from previous 369





and an assumption of no glacial or interglacial erosion (Figure 5). The apparentis lack of inheritance in
samples from above the highest shoreline implies that the last ice sheet advanced over the study area
soon after 35 ka, in accordance with previous inferences for Forsmark (SKB, 2020)Even if the last ice
sheet would have had advanced over the region as late as 28 ka BP, there would only be a very
minornegligible amountinventory of inherited <sup>14</sup>C atoms produced prior to the MIS2 ice advance. An
alternative interpretation is that the last ice sheet advanced more recently but that glacial erosion
during MIS2 was sufficiently deep to remove any nuclide inheritance.

379 Figure 5. Modelled *in situ*<sup>14</sup>C concentration evolution over the last 80 kyr in bedrock surfaces through 380 alternating periods of subaerial exposure and burial by ice sheets during MIS2 and MIS3. 381 histories are modelled from <sup>14</sup>C concentrations in the five samples (BG21-001-BG21-005) from above 382 the highest shoreline. and assume no glacial or interglacial erosion The <sup>14</sup>C development is modelled 383 assuming no glacial or interglacial erosion, continuous exposure to cosmic rays during ice-free periods, and full shielding from cosmic rays (no <sup>14</sup>C production) during periods with ice cover. The points just 384 385 left of the plots display the measured <sup>14</sup>C concentrations for the six sample measurements of the 386 samples (Table 1). (a) Scenario with Schort periods of MIS42 and MIS23 ice cover from 66 to 60 ka BP and from 28 ka BP to the time of deglaciation around 10.7 ka BP. (b) Scenario with Honger periods of 387 MIS42 and MIS23 ice cover from 70 to 57 ka BP and from 35 ka BP to the deglaciation around 10.7 ka 388 BP. Due to the rapid decay of <sup>14</sup>C (with a half-life of 5700 ± 30 5700 years), both scenarios yield similar 389 similar-end-point concentrations of <sup>14</sup>C that overlap,<del>s</del> within uncertainties, <del>with t</del>he measured sample 390 391 concentrations.

Our *in situ* <sup>14</sup>C data from above the highest (postglacial) shoreline demonstrate <del>good itstheir</del> potential
 for this nuclide to help constraining the deglaciation chronology of former ice sheets. This is especially
 true for regions with thin drifttill drapes, abundant bedrock exposures, and sparse moraines outlining

395 successive retreat stages. In Fennoscandia, thin drift-tills conditions occur commonly (cf. Kleman et al., 396 2008) and ice sheet retreat appears to have proceeded uninterrupted inside the Younger Dryas moraine 397 belt (apart from the Central Finland Ice-Marginal Formation; e.g., Rainio et al., 1986; Stroeven et al., 398 2016). Whereas the post-Younger Dryas deglaciation of east-central Sweden is well constrained by clayvarve chronology below the highest postglacial shoreline (Strömberg, 1989) below the highest 399 400 postglacial shoreline, there are vast areas above the highest shoreline that remain poorly constrained by data (Stroeven et al. 2016). In addition to a lack of datable deglacial landforms, this is attributable 401 to glacial erosion of bedrock having frequently been insufficient to remove inventories of long half-life 402 <sup>10</sup>Be and <sup>26</sup>AI (Patton et al., 2022), thereby leaving nuclides inherited from exposure prior to the last 403 glaciation (Heyman et al., 2011; Stroeven et al., 2016). Because of the short <sup>14</sup>C half-life and an 404 405 improved sampling methodology, in situ <sup>14</sup>C may now be a prime candidate nuclide to be included in 406 last deglaciation studies on glaciated cratons, such as the dating of boulders deposited along glacial flowlines; a technique practiced successfully using <sup>10</sup>Be (Margold et al., 2019; Norris et al., 2022). 407

408

#### 409 6. Conclusion

Ten *in situ* <sup>14</sup>C measurements on bedrock are consistent with a RSL curve for Forsmark derived from organic radiocarbon dating of basal sediments in isolation basins and the Fennoscandian Ice Sheet deglaciation chronologies from Stroeven et al. (2016) and Hughes et al. (2016). This study introduces the use of *in situ* <sup>14</sup>C in Fennoscandian Ice Sheet paleoglaciology and outlines a promise of its use as a basis for supporting future shoreline displacement studies and for tracking the deglaciation in areas that lack datable organic material and where <sup>10</sup>Be and <sup>26</sup>Al routinely return complex exposure results.

416

417 Data availability. Data are available in Supplements 1-3. LiDAR data used in the study can
 418 downloaded-are available from https://www.lantmateriet.se

419 Author contributions. BWG and APS initiated the study, with support from KH and JON, and drafted

420 the manuscript. BWG, APS, and AL did the sampling. AL did petrological analyses of the sampled 421 bedrock. NAL completed sample preparation for AMS and provided the results. JH carried out

422 cosmogenic nuclide production rate and exposure age calculations. MWC oversaw the AMS. All

423 authors revised the manuscript.

424 Competing interests. The contact author has declared that none of the authors has any competing425 interests.

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