



using in situ cosmogenic 14C from quartz 2 3 4 Bradley W. Goodfellow1* 5 Arjen P. Stroeven^{2,3} 6 Nathaniel A. Lifton^{4,5} 7 Jakob Heyman⁶ 8 Alexander Lewerentz¹ Kristina Hippe⁷ 9 10 Jens-Ove Näslund⁸ 11 Marc W. Caffee^{4,5} 12 13 ¹Geological Survey of Sweden 14 ²Department of Physical Geography, Stockholm University 15 ³Bolin Centre for Climate Research, Stockholm University 16 ⁴Department of Earth, Atmospheric, and Planetary Sciences, Purdue University 17 ⁵Department of Physics and Astronomy, Purdue University 18 ⁶Department of Earth Sciences, University of Gothenburg 19 ⁷Umweltplanung Dr. Klimsa 20 ⁸Swedish Nuclear Fuel and Waste Management Company (SKB) 21 22 *Corresponding author: bradley.goodfellow@sgu.se 23 24 Abstract 25 In situ 14C in quartz provides a recently developed tool to date exposure of bedrock surfaces up to 26 ~25 000 years. From outcrops located in east-central Sweden, we test the accuracy of in situ ¹⁴C dating 27 against (i) a relative sea level (RSL) curve constructed from radiocarbon dating of organic material in isolation basins, and (ii) the timing of local deglaciation constructed from a clay varve chronology 28 29 complemented with radiocarbon dating. Five samples of granitoid bedrock were taken along an 30 elevation transect extending southwestwards from the Baltic Sea coast near Forsmark. Because these 31 samples derive from bedrock outcrops positioned below the highest postglacial shoreline, they target 32 the timing of progressive landscape emergence above sea level. In contrast, in situ ¹⁴C concentrations 33 in an additional five samples taken from granitoid outcrops above the highest postglacial shoreline, 34 located 100 km west of Forsmark, should reflect local deglaciation ages. The ten in situ 14C

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





measurements provide robust age constraints that, within uncertainties, compare favorably with the 35 36 RSL curve and with the local deglaciation chronology. These data demonstrate the utility of in situ 14C to accurately date ice sheet deglaciation, and durations of postglacial exposure, in regions where 37 cosmogenic ¹⁰Be and ²⁶Al routinely return complex exposure results. 38 39 1. Introduction The pacing of retreat of ice sheets in North America and Eurasia since their maximum expansion 40 41 during the last glaciation remains an active research field (e.g., Hughes et al., 2016; Stroeven et al., 2016; Patton et al., 2017; Dalton et al., 2020, 2023). Understanding the triggers and processes causing 42 the demise of these ephemeral ice sheets yields the best blueprint for understanding the future 43 44 behavior of the Greenland and Antarctic ice sheets in a warming climate. Coupling the behavior of 45 deglaciating ice sheets over the course of the Late Glacial and early Holocene to increasingly precise 46 climate reconstructions and climatic events, requires increased precision in ice sheet reconstructions 47 (e.g., Bradwell et al., 2021). Increased precision can be achieved through a coupling of 48 geomorphological mapping of ice sheet margins (such as moraines, grounding zone wedges, lateral meltwater channels, and ice-dammed lake shorelines and spillways) with numerical field constraints 49 50 from a diverse array of dating techniques (e.g., Stroeven et al., 2016; Bradwell et al., 2021; Regnéll et 51 al., 2023). 52 Ice sheet reconstructions, especially in North America, have attained a high level of detail through 53 radiocarbon dating (Dyke et al., 2002; Dalton et al., 2020). With the advance of offshore imaging of 54 glacial geomorphology (Greenwood et al., 2017, 2021; Bradwell et al., 2021), radiocarbon dating has 55 received a renewed upswing in recent years (e.g., Dalton et al., 2020; Bradwell et al., 2021). However, 56 large tracts of landscape lack radiocarbon age constraints on ice sheet retreat simply due to a lack of 57 datable organic material. Fortunately, optically-stimulated luminescence ages on buried sand layers 58 (e.g., Alexanderson et al., 2022) and cosmogenic nuclide apparent exposure ages on exposed bedrock 59 and erratics have narrowed some of the gaps (e.g., Hughes et al., 2016; Stroeven et al., 2016; Dalton et 60 al., 2023). In studies using cosmogenic nuclides, an 'apparent' exposure age is derived from a simple 61 calculation from the nuclide concentration under consideration (Lal, 1991; Gosse and Phillips, 2001). However, correctly interpreting the exposure age relies on modelling that considers geological factors 62 that can reduce the nuclide concentration relative to the time since initial subaerial exposure (such as 63 64 erosion and burial by glacial ice, water, snow, and/or soil; Gosse and Phillips, 2001; Schildgen et al., 65 2005; Ivy-Ochs and Kober, 2008). Exposure dating is the only technique available in regions where ice sheet erosion has left the surface bare or covered by a thin drape of till. Kleman et al. (2008) show that 66

for Fennoscandia, these conditions are widespread in coastal regions where ice accelerated towards its





69 existing sediment covers. 70 Coastal sectors in formerly glaciated regions provide sites important to the study of paleoglaciology. 71 They offer an abundance of bedrock exposures from which patterns and processes of subglacial 72 erosion can be studied through cosmogenic nuclide exposure dating (e.g., Hall et al., 2020). Also, 73 because of the interplay with postglacial sea level, coastal areas yield data on glacioisostatic rebound 74 that are critical to geodynamic modelling of Earth rheology and thicknesses of former ice sheets (e.g., Lambeck et al. (1998, 2010) and Patton et al. (2017), for Fennoscandian examples). Geodynamic 75 76 models require validation against measurements of vertical crustal motion (Steffen and Wu, 2011), 77 such as those provided by recent global positioning system (GPS) measurements (e.g., Lidberg et al., 78 2010) and postglacial records of crustal rebound afforded by relative sea level (RSL) curves (e.g., Påsse 79 and Andersson, 2005). The construction of RSL curves, detailing the history of land surface emergence from sea level, is traditionally done using either sediments accumulated in isolation basins at different 80 81 elevations above sea level or by dating uplifted gravel beach ridges. Typically, isolation basins, and their 82 sediments, show a progression from marine, to brackish, and finally to freshwater environments as 83 their bedrock sills are uplifted through tidal levels (Long et al., 2011). Histories of land uplift above sea 84 level are documented using micro- and macrofossil analyses of isolation basin sediments and radiocarbon dating on macrofossils (Romundset et al., 2011). Uplifted beach ridges can be radiocarbon 85 86 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 ¹⁰Be 87 exposure dating even though, other than the highest beach ridge, they may be prone to clast 88 89 reworking (Briner et al., 2006; Simkins et al., 2013; Bierman et al., 2018). A distinct advantage of 90 constructing RSL curves using cosmogenic nuclides is that land surface emergence above sea level may 91 be additionally dated from boulders (Briner et al., 2006) or bedrock (Bierman et al., 2018). 92 The potential for cosmogenic surface exposure dating of last ice sheet retreat in recently glaciated low-93 relief cratonic landscapes would seemingly be high because of the frequent outcropping of glacially 94 sculptured quartz-bearing crystalline bedrock. However, the ice sheet may have been either non-95 erosive or erosion was insufficiently deep to remove all the cosmogenic nuclide inventory from 96 previous exposure periods. Apparent ages are therefore often older than indicated by radiocarbon 97 dating (Heyman et al., 2011; Stroeven et al., 2016) because they include a component of nuclide 98 inheritance. Apparent ages younger than indicated by radiocarbon dating can also occur if sampled 99 rock surfaces have been shielded, for example by sediments, following deglaciation. Concentrations of 100 ¹⁰Be and ²⁶Al, in either bedrock or erratic boulders, therefore often reflect complex exposure histories 101 rather than simple deglacial exposure durations (Heyman et al., 2011; Stroeven et al., 2016).

streaming sectors and where wave wash during glacial rebound further thinned or removed pre-





102 In this study we use ¹⁴C produced *in situ* in quartz-bearing bedrock (*in situ* ¹⁴C) because it potentially 103 circumvents an overt reliance on the need for deep erosion (> 3 m) to remove the inherited signal from 104 previous exposure periods (Gosse and Phillips, 2001). The reason for this is that, because of its short 105 half-life of 5700 ± 30 years, nuclide inheritance 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 106 107 is, ca. 5 half-lives (Briner et al., 2014). 108 Some studies assessing changes in glacier and ice sheet extents over Late Glacial to Holocene 109 timescales have used in situ 14C (Miller et al., 2006; Fogwill et al., 2014; Hippe et al., 2014; Schweinsberg et al., 2018; Pendleton et al., 2019; Young et al., 2021; Schimmelpfennig et al., 2022). In 110 111 such studies, in situ 14C has been applied with other nuclides with longer half-lives, in particular 10Be, 112 to unravel complex histories of glacier advance and retreat (e.g., Goehring et al., 2011) and spatial patterns in glacial erosion in mountainous terrain (e.g., Steinemann et al., 2021). However, extensive 113 regions formerly covered by ice sheets are characterized by low relief, low elevation terrain, and the 114 effectiveness of in situ 14C in dating ice sheet retreat in these non-alpine settings and in quantifying 115 116 shoreline displacement from bedrock samples has not been previously assessed. The aim of this study is therefore to validate the use of ¹⁴C formed *in situ* in bedrock as a reliable chronometer by evaluating 117 its performance in duplicating (i) a previously-established Holocene RSL curve based on radiocarbon 118 dating (Hedenström and Risberg, 2003; SKB, 2020) and (ii) the timing of deglaciation above the highest 119 120 (post-glacial) shoreline in nearby east-central Sweden according to reconstructions of deglaciation of the last ice sheet (Hughes et al., 2016; Stroeven et al., 2016). 121

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2. Study Area

Our study is focused on a region that includes low elevation, low relief, Forsmark-Uppland and adjoining higher elevation and relief Dalarna-Gävleborg in east-central Sweden (Fig. 1). This region was selected because Forsmark is the location of a planned geological repository for spent nuclear fuel (e.g., SKB 2022) and therefore also has abundant geologic data relevant to our study. This includes indepth analyses of bedrock and 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).

From spatio-temporal ice sheet reconstructions by Kleman et al. (2008), the study area was glaciated 16-20 times for a total duration of c. 330 ky over the past 1 Ma. The last deglaciation of the study area is well-constrained by two recent reconstructions that differ in their approach (Hughes et al., 2016; Stroeven et al., 2016). The Hughes et al. (2016) reconstruction is explicitly based on chronological constraints, but the Stroeven et al. (2016) reconstruction combines geomorphological constraints for ice sheet margin outlines with chronological constraints. Whereas Hughes et al. (2016) reconstruct ice





136 sheet retreat every 1 ka, and for every ice margin plot its position as "most credible", "min", and 137 "max", Stroeven et al. (2016) present ice margin positions for every 100 years inside the Younger Dryas standstill position (Stroeven et al., 2015). These marginal positions are temporally and spatially defined 138 139 by the "Swedish Time Scale" clay varve record along the Swedish east coast (De Geer, 1935, 1940; 140 Strömberg, 1989, 1994; Brunnberg, 1995; Wohlfarth et al., 1995). From Stroeven et al. (2016), the last 141 deglaciation of the study area occurred 10.8 ± 0.3 ka BP, which overlaps the timing of deglaciation of the study area from Hughes et al. (2016), within uncertainty (Fig. 1). The highest postglacial shoreline 142 143 in east-central Sweden is located at a present elevation of ~200 m a.s.l. in Dalarna-Gävleborg, ~100 km 144 west of Forsmark (SGU, 2015). The exposure duration of bedrock above the highest postglacial 145 shoreline therefore represents the time since local deglaciation. Hence, in situ ¹⁴C ages from bedrock above the highest postglacial shoreline should conform to the reconstructed deglaciation age of 10.8 \pm 146 147 0.3 ka from Stroeven et al. (2016). 148 Below the highest postglacial shoreline, in the Forsmark-Uppland region, the last deglaciation 149 occurred in a marine environment and the landscape has progressively emerged above sea level 150 through postglacial isostatic uplift. A RSL curve constructed from radiocarbon dating of basal organic 151 sediments trapped in isolation basins along elevation transects describes the progressive emergence 152 of the Forsmark-Uppland landscape above sea level (Robertsson and Persson, 1989; Risberg, 1999; Bergström, 2001; Hedenström and Risberg, 2003; Berglund, 2005; SKB, 2020). Ages calculated from in 153 154 situ 14C from bedrock outcrops along an elevation transect would then mirror the Forsmark RSL curve for their corresponding elevations (but be slightly older because of nuclide production through 155 156 shallow water before emergence). 157 A potential complication to the accurate exposure age dating of bedrock surfaces using in situ 14C in 158 east-central Sweden is that the most recent period of ice sheet burial may not have been sufficiently 159 long to decay the in situ 14C inventory inherited from preceding exposure. Here, the extent of the Fennoscandian Ice Sheet during interstadial MIS3 and the timing of ice advance across the Forsmark 160 161 region during late MIS3 are crucially important. Kleman et al. (2020) have identified ice-free conditions 162 around Idre (330 km NW, up-ice, of our study area; Fig. 1) between 55 ka and 35 ka, which implies inundation of our study area by ice after 35 ka. Combined with a well-constrained final deglaciation 163 164 age of 10.8±0.3 ka (Stroeven et al. 2016), it appears that our study area has most recently (during 165 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). 166 167 Consequently, it is possible that in situ ¹⁴C concentrations may reflect subaerial exposure of bedrock in 168 our study area during MIS3 in addition to Holocene exposure, resulting in an offset towards older ages 169 relative to the RSL curve for Forsmark (Hedenström and Risberg, 2003; SKB, 2020) and the deglaciation 170 chronologies of Hughes et al. (2016) and Stroeven et al. (2016).





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3. Methods

3.1. Sampling of bedrock outcrops for in situ ¹⁴C measurement

We used the following sampling strategy to evaluate the accuracy of bedrock exposure ages derived from in situ 14C 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 hydrothermal processes that is likely to occur in major pegmatite intrusions, outcrops located in major deformation zones, and outcrop-scale veins, fractures, and adjacent rock volumes. Consequently, sampling was done on outcrops of metagranitoid from the early-Svecokarelian GDG-GSDG suite that dominates the Bergslagen lithotectonic unit (Stephens and Jansson, 2020). A petrological examination 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 collected using an angle grinder, which permits sampling of hard crystalline bedrock isolated from outcrop edges, fractures, and quartz veins, and consistently limits sample thicknesses to 3 cm. We collected a total of ten samples for in situ 14C analyses. Five of these were collected along a SW-NE 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 14 C can therefore be evaluated against the Forsmark RSL curve. We collected a further five samples from locations above the highest shoreline (Fig. 1a) to determine the age of local deglaciation for comparison with published deglaciation chronologies (Hughes et al., 2016; Stroeven et al., 2016). Sample locations were logged on a 2 m-resolution LiDAR digital elevation model (DEM) displayed in ArcGIS 10 on a tablet computer. A GPS add-in tool in ArcGIS 10 was used to record positional data, within a horizontal precision of 2 m. The elevation of each sample location was extracted from the DEM and has a precision of tens of centimetres. The influence of these minor positional uncertainties on our ¹⁴C calculations is trivial and none of the sample sites is influenced by topographic shielding that could reduce the accumulation of ¹⁴C in bedrock. 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 sampling bedrock that was moss-covered, we cannot be certain that moss mats did not formerly cover the sample sites. Given a compressed thickness of 0.5 cm and an estimated density of 0.7 g/cm³, this

may have contributed to a shielding of the sampled rock surfaces of 0.35 g/cm², which is negligible and

is therefore excluded from our age inferences.





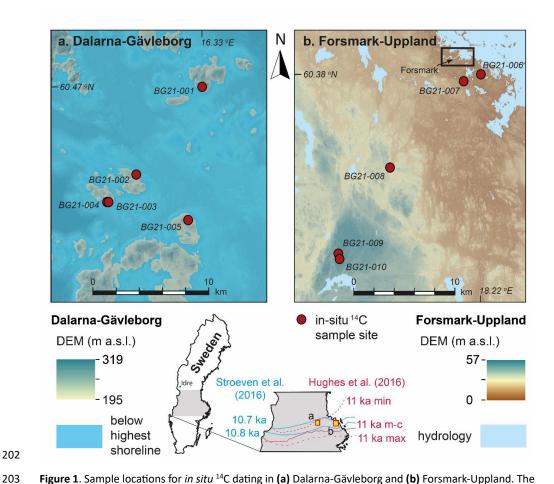


Figure 1. Sample locations for *in situ* ¹⁴C dating in **(a)** Dalarna-Gävleborg and **(b)** Forsmark-Uppland. The five Dalarna-Gävleborg sample sites are located above the highest postglacial shoreline (shown), whereas the five sample sites from Forsmark-Uppland are located below the highest shoreline (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 ka BP (most-credible, minimum, and maximum) retreat isochrones (red) from Hughes et al. (2016). The 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.

3.2. Laboratory preparation for accelerator mass spectrometry (AMS)

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* ¹⁴C were determined from purified quartz separates through automated procedures (Lifton et al., 2023). Approximately 5 g of quartz from each sample was added to a degassed LiBO₂ flux in a re-usable 90% Pt/10% Rh sample boat and heated





to 500 °C for one hour in ca. 6.7 kPa of Research Purity O₂ to remove atmospheric contaminants, which were discarded. The sample was then heated to 1100 °C for three hours to dissolve the quartz and release the *in situ* ¹⁴C, again in an atmosphere of ca. 6.7 kPa of Research Purity O₂ to oxidize any evolved carbon species to CO₂. The CO₂ from the 1100 °C step was then purified, measured quantitatively, and converted to graphite for ¹⁴C AMS measurement at PRIME Lab (Lifton et al., 2023). To test for data reproducibility, sample BG21-002 was randomly selected to undergo laboratory preparation and AMS a second time. Measured concentrations of *in situ* ¹⁴C are calculated from the measured isotope ratios via AMS following Hippe and Lifton (2014).

3.3. Exposure age calculations

The expage calculator version 202312 (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 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 ages are calculated using resulting time-varying ¹⁴C production rates accounting for decay and interpolated to match the measured ¹⁴C concentration. The production rate from muons is calibrated against the Leymon High core ¹⁴C data of Lupker et al. (2015) and the production rate from spallation is calibrated against updated global ¹⁴C production rate calibration data (Schimmelpfennig et al., 2012; Young et al., 2014; Lifton et al., 2015; Borchers et al., 2016; Phillips et al., 2016; Koester and Lifton, 2023). This calibration is done iteratively for spallation and muons to reach convergence, using the expage production rate calibration methods (Fig. 2).

emergence through shallow water. However, burial 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.



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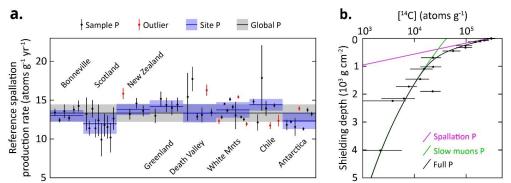


Figure 2. Production rate calibration of ¹⁴C in quartz. (a) Reference spallation ¹⁴C production rate calibration based on data from Schimmelpfennig et al. (2012), Young et al. (2014), Lifton et al. (2015), Borchers et al. (2016), and Phillips et al. (2016), corrected per Hippe and Lifton (2014) and compiled in Koester and Lifton (2023). An uncertainty-weighted production rate is calculated for each of the eight sites. Outliers, which are not included in the uncertainty-weighted production rates, are determined based on the requirement that there should be at least three samples yielding a reduced chi-square statistic (X_B^2) with a p-value of at least 0.05 for the assumption that the individual production rates from a site are derived from one normal distribution. For X_R^2 , but not the uncertainty-weighting, we use the largest of the sample-specific production rate uncertainty based on the ¹⁴C concentration uncertainties and 5% of the sample production rate. This procedure does not punish samples with low measurement uncertainties, which otherwise risk exclusion as outliers. We adopt a global reference spallation 14C production rate of 13.35 \pm 1.13 atoms g^{-1} yr⁻¹, calculated as the arithmetic mean of the eight site production rates with the uncertainty being based on an uncertainty-weighted deviation of all included single sample production rates, excluding outliers. (b) Calibration of ¹⁴C production rate from muons based on the data of Lupker et al. (2015). The calibration is based on the method used in the CRONUScalc calculator (Marrero et al., 2016; Phillips et al., 2016). The figure shows the best fit ¹⁴C concentration profiles produced from spallation, slow muons, and full production. The best fit yields near zero production from fast muons (cf. Lupker et al., 2015). The production rate calibration has been carried out using the expage-202306 calculator in an iterative way to make the global reference spallation 14C production rate converge with the production rate from muons.

4. Results

Inferred ages for the five *in situ* ¹⁴C samples 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* ¹⁴C exposure age curve considering subaqueous cosmogenic nuclide production (Figure 3; Tables 1 and 2). Exposure age uncertainties are large with internal uncertainties (measurement uncertainties;





Balco et al., 2008) of 5-9% and external uncertainties of 12-20% (also including production rate uncertainties, which are high relative to 10 Be (Borchers et al., 2016; Phillips et al., 2016). Apparent exposure ages increase consistently with elevation and match expected ages within uncertainty. The two highest samples have near-identical apparent exposure ages and elevations. However, these samples provide independent ages because they are horizontally separated by 624 m (Figure 1b). There is good agreement between ages inferred from these *in situ* 14 C data and the RSL curve constructed from organic radiocarbon dating of isolation events (Hedenström and Risberg, 2003; SKB, 2020).

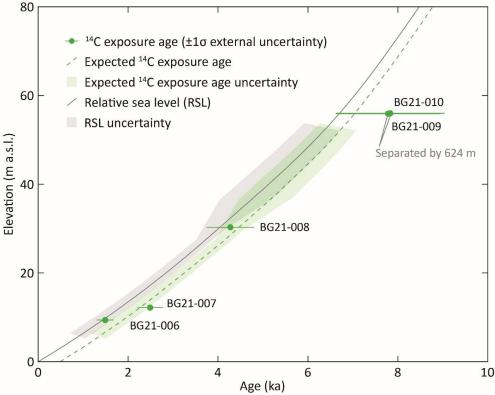


Figure 3. Apparent 14 C exposure ages for five Forsmark samples from below the highest shoreline (Fig. 1b; Table 2) with 16 C exposure ages for five Forsmark samples from below the highest shoreline (Fig. 1b; Table 2) with 16 C exposure ages are calculated assuming the RSL curve is correct, the 14 C 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 G 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* 14 C samples located above the highest shoreline in Dalarna and Gävleborg (Fig. 1a) are shown in Figure 4 and Table 2. The weighted mean age from all five samples is 11.2 ± 1.3 ka. These data display a X_R^2 of 1.78 and a p-value of 0.13 based on 1σ internal uncertainties (Fig. 4a), which does not support a rejection of the hypothesis that the apparent exposure ages represent 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 ± 0.3 ka (Stroeven et al. 2016). Replicate measurements on sample BG21-002 closely agree and an age based on a weighted mean 14 C concentration is shown in Figure 4. Sample BG21-001 provides the youngest apparent age but, because 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 sites. We therefore consider this sample as least likely to provide a reliable age. Removing this sample from consideration indicates that the remaining four sample sites are more clustered, with an older weighted mean age of 11.6 ± 1.1 ka, which displays a X_R^2 of 0.43 and a p-value of 0.73 based on 1σ internal uncertainties (Fig. 4b).

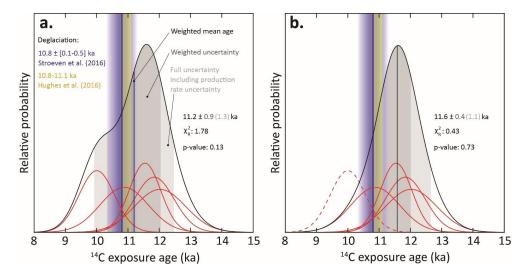


Figure 4. Probability density plots of the exposure ages from samples above the highest shoreline (Fig. 1a; Table 2). The individual samples (red curves) display 1σ internal uncertainty (measurement uncertainty). For the repeat sample BG21-002, the exposure age is calculated with a weighted mean ¹⁴C concentration using a 2% uncertainty. **(a)** The probability density and data for all five samples. For the full set of samples, 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. **(b)** The probability density and data with sample BG21-001 excluded as an outlier. These cosmogenic nuclide ages yield a X_R^2 of 0.43 and a p-value of 0.73 based on internal uncertainties, which again indicate that

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307 they are from the same population. All ages are referenced to the sampling year 2021. The weighted

ages of 11.2 ± 1.3 ka and 11.6 ± 1.1 ka both overlap with the deglaciation age from Stroeven et al. (2016).





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FCEGS #	Ę	Mass Quartz (g)	(µg)	mass C	Mass C	∂ 13 C 3 (%0VPDB)	(70°)	"C/Ctotal (70°)	**C (10° at)	["C] (70° at g")"
146 2	202101960	5.02378	5.0 ± 0.1	393.8 ± 4.8	382.3 ± 4.6	45.9 ± 0.2	0.3399 ± 0.0075	0.3412 ± 0.0079	0.6177 ± 0.0179	1.2296 ± 0.0357
147 2	202101961	5.02383	7.8 ± 0.1	303.3 ± 3.7	294.4 ± 3.6	-44.8 ± 0.2	0.4555 ± 0.0096	0.4623 ± 0.0102	0.6470 ± 0.0181	1.2879 ± 0.0360
148 2	202101962	5.01070	17.6 ± 0.3	303.4 ± 3.7	294.5 ± 3.6	43.9 ± 0.2	0.4633 ± 0.0108	0.4709 ± 0.0113	0.6604 ± 0.0197	1.3180 ± 0.0393
150 2	202201473	5.04116	7.7 ± 0.1	305.3 ± 3.7	296.4 ± 3.6	45.2 ± 0.2	0.4558 ± 0.0135	0.4624 ± 0.0142	0.6519 ± 0.0237	1.2931 ± 0.0470
152 2	202101963	5.05927	11.9 ± 0.2	305.7 ± 3.7	296.8 ± 3.6	-44.6 ± 0.2	0.4618 ± 0.0079	0.4691 ± 0.0083	0.6630 ± 0.0159	1.3105 ± 0.0314
153 2	202101964	5.07578	4.6 ± 0.1	304.5 ± 3.7	295.6 ± 3.6	45.4 ± 0.2	0.4600 ± 0.0127	0.4667 ± 0.0134	0.6566 ± 0.0225	1.2935 ± 0.0444
155 2	202101965	5.06572	5.5 ± 0.1	306.8 ± 3.7	297.8 ± 3.6	45.2 ± 0.2	0.1277 ± 0.0056	0.1172 ± 0.0059	0.1243 ± 0.0101	0.2453 ± 0.0199
157 2	202101966	5.03589	6.9 ± 0.1	309.2 ± 3.8	300.1 ± 3.7	-45.0 ± 0.2	0.1684 ± 0.0051	0.1601 ± 0.0054	0.1922 ± 0.0096	0.3817 ± 0.0191
158 2	202101967	5.07653	4.0 ± 0.1	308.9 ± 3.8	299.9 ± 3.6	-45.4 ± 0.2	0.2357 ± 0.0063	0.2308 ± 0.0067	0.3015 ± 0.0119	0.5938 ± 0.0234
160 2	202101968	5.01906	55.3 ± 0.7	305.6 ± 3.7	296.6 ± 3.6	-38.0 ± 0.2	0.3339 ± 0.0095	0.3368 ± 0.0101	0.4601 ± 0.0170	0.9168 ± 0.0339
161 2	202101969	4.99961	42.2 ± 0.6	306.0 ± 3.7	297.0 ± 3.6	40.1 ± 0.2	0.3320 ± 0.0068	0.3340 ± 0.0072	0.4565 ± 0.0132	0.9130 ± 0.0264
PCEGS # = sample number in the Purdue Carbon Extraction and Graphitization System	nber in the	Purdue C	arbon Ext	raction and	d Graphitiza	ition Systen	٦			
BG21-001 146 202101960 5.02378 5.0±0.1 393.8±4.8 382.3±4.6 45.9±0.2 BG21-002 147 202101961 5.02383 7.8±0.1 393.8±4.8 382.3±4.6 45.9±0.2 BG21-003 148 202101962 5.01070 17.6±0.3 303.4±3.7 294.4±3.6 44.8±0.2 BG21-004 150 202201473 5.04116 7.7±0.1 305.3±3.7 296.4±3.6 45.2±0.2 BG21-005 153 202101963 5.05927 11.9±0.2 305.7±3.7 296.8±3.6 45.2±0.2 BG21-006 155 202101964 5.07578 4.6±0.1 304.5±3.7 295.6±3.6 45.2±0.2 BG21-007 157 202101966 5.03589 6.9±0.1 309.2±3.8 300.1±3.7 45.0±0.2 BG21-008 158 202101967 5.07653 4.9±0.1 309.2±3.8 300.1±3.7 45.0±0.2 BG21-019 160 202101968 5.01906 55.3±0.7 305.6±3.7 296.6±3.6 45.4±0.2 BG21-019	BG21-001 146 202101960 5.02378 5.0±0.1 393.8 ± 4.8 BG21-002 147 202101961 5.02378 7.8 ± 0.1 393.3 ± 3.7 BG21-003 148 202101962 5.01070 17.6 ± 0.3 303.3 ± 3.7 BG21-002R 150 202201473 5.04116 7.7 ± 0.1 305.3 ± 3.7 BG21-004 152 202101963 5.05927 11.9 ± 0.2 305.7 ± 3.7 BG21-005 153 202101963 5.05578 4.6 ± 0.1 304.5 ± 3.7 BG21-006 155 202101965 5.06572 5.5 ± 0.1 306.8 ± 3.7 BG21-007 157 202101966 5.03589 6.9 ± 0.1 306.8 ± 3.7 BG21-008 158 202101966 5.03589 6.9 ± 0.1 308.9 ± 3.8 BG21-009 160 202101967 5.07653 4.0 ± 0.1 308.9 ± 3.7 BG21-009 160 202101969 5.01906 55.3 ± 0.7 305.0 ± 3.7 BG21-009 160 202101969 5.01906 55.3 ± 0	5.02378 5.02383 5.01070 5.04116 5.05927 5.05927 5.06572 5.03589 5.07653 5.01906 4.99961 Purdue C Purdue C	5.0 ± 0.1 7.8 ± 0.1 17.6 ± 0.3 7.7 ± 0.1 11.9 ± 0.2 4.6 ± 0.1 5.5 ± 0.1 6.9 ± 0.1 4.0 ± 0.1 55.3 ± 0.7 42.2 ± 0.6 arbon Ext where VPI	393.8 ± 4.8 303.3 ± 3.7 303.4 ± 3.7 305.3 ± 3.7 305.7 ± 3.7 304.5 ± 3.7 306.8 ± 3.7 309.2 ± 3.8 309.9	382.3 ± 4.6 294.4 ± 3.6 296.4 ± 3.6 296.8 ± 3.6 295.6 ± 3.6 297.8 ± 3.6 299.9 ± 3.6 299.9 ± 3.6 296.6 ± 3.6 296.6 ± 3.6 296.6 ± 3.6 297.0 ± 3.6 297.0 ± 3.6 297.0 ± 3.6	45.9 ± 0.2 44.8 ± 0.2 45.2 ± 0.2 45.4 ± 0.2 45.2 ± 0.2 45.2 ± 0.2 45.2 ± 0.2 45.0 ± 0.2 45.4 ± 0.2 38.0 ± 0.2 40.1 ± 0.2 ition Systen elemnite)		0.3412±0.0079 0.4623±0.0102 0.4709±0.0113 0.4624±0.0142 0.4691±0.0083 0.4667±0.0134 0.1172±0.0059 0.1601±0.0054 0.2308±0.0067 0.3368±0.0101 0.3340±0.0072	0.6177 0.6470 0.6604 0.6519 0.6630 0.6566 0.1243 0.1922 0.3015 0.4601	E 0.0179 E 0.0181 E 0.0197 E 0.0237 E 0.0225 E 0.0225 E 0.0101 E 0.0096 E 0.0170 E 0.0170





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Sample ID	(°)	Long (°)	Elevation (m a.s.l.)	Thickness (cm)	Density (g cm ⁻³)	Shielding factor	Erosion (cm yr ⁻¹)	¹⁴ C ± 1σ (10 ² at g ⁻¹)	¹⁴ C Age ± Unc. ^{Ext.} (± Unc. ^{Int.}) ^a (ka)
BG21-001	60.47432	16.33134	236.5	ω	2.7	_	0	1 230 ± 36	10.0 ± 1.7 (± 0.6)
BG21-002	60.40615	16.22197	212.6	ω	2.7	_	0	1288 ± 36	$11.5 \pm 2.2 (\pm 0.7)$
BG21-002R	60.40615	16.22197	212.6	ω	2.7	_	0	1293 ± 47	$11.6 \pm 2.3 (\pm 0.9)$
BG21-003	60.38459	16.17649	216.3	ω	2.7	_	0	1 318 ± 39	$12.0 \pm 2.4 (\pm 0.8)$
BG21-004	60.38451	16.17440	217.8	ω	2.7	_	0	1 311 ± 31	$11.8 \pm 2.3 (\pm 0.6)$
BG21-005	60.36888	16.30526	248.1	з	2.7	_	0	1294 ± 44	$10.9 \pm 2.1 (\pm 0.8)$
BG21-006	60.38490	18.22308	9.4	ω	2.7	_	0	245 ± 20	$1.5 \pm 0.2 (\pm 0.1)$
BG21-007	60.37892	18.19129	12.2	ω	2.7	_	0	382 ± 19	$2.5 \pm 0.3 (\pm 0.1)$
BG21-008	60.30504	18.04993	30.3	ω	2.7	_	0	594 ± 23	$4.3 \pm 0.5 (\pm 0.2)$
BG21-009	60.22988	17.94989	56.0	ω	2.7	_	0	917 ± 34	$7.8 \pm 1.2 (\pm 0.5)$
BG21-010	60.22431	17.95051	55.9	ω	2.7	_	0	913 ± 26	$7.8 \pm 1.2 (\pm 0.4)$

Table 2. In situ 14C from quartz, Dalarna-Gävleborg and Forsmark-Uppland.

 $^{\text{a}}$ Unc. $^{\text{Ext.}}$ is external uncertainty and Unc. $^{\text{Int.}}$ is internal uncertainty. Both are 1σ



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5. Discussion

The in situ 14C bedrock exposure ages from the Forsmark-Uppland transect (i.e., below the highest 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 landscape emergence, in situ 14C is indicated as having high potential as a chronometer over Late Glacial-Holocene timescales in low relief, low elevation settings. This study adds to precious few demonstrations of the ability of cosmogenic nuclide isotopes to define postglacial landscape emergence above sea level (Briner et al., 2006; Bierman et al., 2018). Briner et al. (2006) present good (visual) congruence with a record of shoreline emergence built from radiocarbon-dated driftwood and fauna by Dyke et al. (1992) using ¹⁰Be measurements on boulders in beaches derived from wavewashed till. Their study also mentions that building a relative sea level curve from pebbles, cobbles and plucked bedrock suffered from inheritance problems, an experience shared by Matmon et al. (2003) while attempting the dating of chert on beach ridges in southern Israel and heeded by Bierman et al. (2018). Bierman et al. (2018) successfully dated landscape emergence on Greenland using ¹⁰Be across a range of settings, including bedrock below the highest shoreline, cobbles from beach ridges at the highest shoreline, and boulders and bedrock above the highest shoreline. They note that 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 Sweden and more widely in Fennoscandia, these requirements are not met either because of cold-based conditions (Patton et al., 2016; Stroeven et al., 2016) or weakly erosive warm-based ice such as at Forsmark (Hall et al., 2019; SKB, 2020), during all or much of glacial time. Cosmogenic nuclide inheritance is therefore a part of the landscape fabric. Bierman et al. (2018) advocate the use of in situ ¹⁴C as a methodology to circumvent inheritance problems. Our study is the first to follow-up on that suggestion, and shows, convincingly, that using in situ 14 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 nuclides. Five bedrock samples from above the highest postglacial shoreline are well-clustered and the weighted mean age (and full uncertainty) of 11.2 ± 1.3 ka overlaps with the predicted deglaciation age of 10.8 ± 0.3 ka (Fig. 4a; Hughes et al., 2016; Stroeven et al., 2016). Removing the youngest age from consideration results in more strongly clustered ages (Fig. 4b) and an older mean weighted age of 11.6 ± 1.1 ka, which still overlaps the predicted deglaciation age, within uncertainty. We therefore do not further discriminate between these results. Because derived exposure ages overlap with the predicted deglaciation age, we further infer that the in situ 14C samples, including those located below the highest postglacial shoreline, within uncertainty, lack inheritance from previous exposure. This implies that the last ice sheet advanced over the study area soon after 35 ka, in accordance with previous inferences



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for Forsmark (SKB, 2020). 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.

Our in situ ¹⁴C data from above the highest (postglacial) shoreline demonstrate good potential for this nuclide to help constrain the deglaciation chronology of former ice sheets. This is especially true for regions with thin drift, abundant bedrock exposures, and lacking moraines outlining successive retreat stages. In Fennoscandia, thin drift conditions occur commonly (cf. Kleman et al., 2008) and ice sheet retreat appears to have proceeded uninterrupted inside the Younger Dryas moraine belt (apart from the Central Finland Ice-Marginal Formation; e.g., Rainio et al., 1986; Stroeven et al., 2016). Whereas the post-Younger Dryas deglaciation of east-central Sweden is well constrained by clay-varve chronology (Strömberg, 1989) below the highest 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 to glacial erosion of bedrock having frequently been insufficient to remove inventories of long half-life ¹⁰Be and ²⁶Al (Patton et al., 2022), thereby leaving nuclides inherited from exposure prior to the last glaciation (Heyman et al., 2011; Stroeven et al., 2016). Because of the short ¹⁴C half-life and an improved sampling methodology, in situ ¹⁴C may now be a prime candidate nuclide to be included in last deglaciation studies on glaciated cratons, such as the dating of boulders deposited along glacial flowlines; a technique practiced successfully using ¹⁰Be (Margold et al., 2019; Norris et al., 2022).

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6. Conclusion

Ten *in situ* ¹⁴C 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* ¹⁴C 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 ¹⁰Be and ²⁶Al routinely return complex exposure results.

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- **Data availability.** Data are available in Supplements 1-3. LiDAR data used in the study can downloaded from https://www.lantmateriet.se
- 431 Author contributions. BWG and APS initiated the study, with support from KH and JON, and drafted

bedrock. NAL completed sample preparation for AMS and provided the results. JH carried out

- the manuscript. BWG, APS, and AL did the sampling. AL did petrological analyses of the sampled
- 434 cosmogenic nuclide production rate and exposure age calculations. MWC oversaw the AMS. All
- 435 authors revised the manuscript.





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