



Evaluating the 11-year solar cycle and short-term ^{10}Be deposition events with novel excess water samples from the EGRIP project

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Abstract. ^{10}Be is produced by the interaction between galactic cosmic rays (GCR) and solar energetic particles (SEP) with the Earth's atmospheric constituents. The flux of GCR is modulated by the varying strength of the magnetic fields of the Earth and the Sun. Measurement of ^{10}Be concentrations from polar ice cores is thus a
20 valuable tool to reconstruct the variations of the geomagnetic field and solar activity levels. The interpretation of ^{10}Be records is, however, complicated by non-production related effects on the ^{10}Be deposition rate caused by climate/weather induced noise. Furthermore, volcanic eruptions have been proposed to lead to short-term ^{10}Be deposition enhancements. In this study, we test the use of excess meltwater from continuous flow analysis (CFA) to measure ^{10}Be , allowing less time-consuming and more cost-effective sample preparation. We compare two
25 records obtained from CFA and discrete samples from the EGRIP S6 firn core, reaching back to 1900 CE. We find that the two records agree well and that the ^{10}Be record from CFA samples agrees as well as the discrete samples with other records from Greenland. Furthermore, by subtracting the theoretically expected GCR-induced signal, we investigate the high-frequency variability of the ^{10}Be records from Greenland and Antarctica after 1951
30 CE, with focus on SEP events and volcanic eruptions. Finally, we use the ^{10}Be records from Greenland and Antarctica to study the 11-year solar cycles, allowing us to assess the suitability of the CFA samples for the reconstruction of solar activity. This result opens new opportunities for the collection of continuous ^{10}Be records with less time-consuming sample preparation while saving an important portion of the ice cores for other measurements.

35 1 Introduction

^{10}Be is produced in the atmosphere by the interaction between high-energy cosmic ray particles (galactic cosmic rays – GCR) and atmospheric atoms. About 65% of atmospheric ^{10}Be nuclides are produced in the stratosphere (Heikkilä et al., 2013; Heikkilä et al., 2009; Masarik & Beer, 1999), where they can be assumed to be well mixed
40 due to an average residence time of 1-2 years (Heikkilä et al., 2008). Upon binding to sulfate aerosols (e.g. Igarashi et al., 1998), ^{10}Be is removed from the atmosphere by wet and dry deposition, and can be measured in environmental archives, such as ice cores from Greenland and Antarctica. Using the ECHAM5 general circulation



model coupled with the aerosol module HAM, Heikkilä et al. (2009) showed that the dominant component of ^{10}Be in polar ice cores is stratospheric ^{10}Be , making up to about two thirds of the signal preserved in ice cores from
45 Greenland and Antarctica. In consequence, they argued that deposition fluxes of ^{10}Be reflect well the changes in the global average ^{10}Be atmospheric production rate.

The flux of GCRs reaching the Earth is modulated by the magnetic fields of Sun and Earth. The production of ^{10}Be thus anticorrelates with the strength of the helio- and geomagnetic fields allowing us to use ^{10}Be from ice cores to reconstruct solar activity (e.g. Beer et al., 1990; McCracken et al., 2004; Muscheler et al., 2007;
50 Steinhilber et al., 2012; Vonmoos et al., 2006), and the variations in the geomagnetic field dipole moment (e.g. Muscheler et al., 2005; Raisbeck et al., 1985; Raisbeck et al., 2006).

In the last decade it has also been shown that solar storms can leave a significant imprint in ^{10}Be from Greenland and Antarctica. These events can, in fact, lead to short-lived peaks in the radionuclide concentrations (Mekhaldi et al., 2015; Miyake et al., 2019; Miyake et al., 2015; O'Hare et al., 2019; Paleari et al., 2022). Mekhaldi et al.
55 (2021) modeled the theoretically expected globally averaged annual production rate of ^{10}Be since the 1950s using neutron monitor data, and the theoretical production induced by ground level enhancements (GLEs), i.e. solar energetic particle (SEPs) events that cause sudden increases in ground-based neutron monitor count-rates. The strongest directly observed GLE - GLE no.5 from 1956 – caused, for instance, an increase of only about 5% in the annual ^{10}Be production rate. Considering that the 11-year solar cycle is estimated to cause a variability of
60 $\pm 15\%$ to $\pm 35\%$ (e.g. Baroni et al., 2011; Paleari et al., 2022; Pedro et al., 2012, Mekhaldi et al., 2021), the signal of modern events cannot be unequivocally distinguished from the variability caused by the 11-year cycle, and climate and local weather influences (Mekhaldi et al., 2021). For the instrumental period no unambiguous SEP signal could be detected in ice cores so far, not even in seasonal ^{10}Be data (e.g. Pedro et al., 2011; Zheng et al., 2020).

65 Moreover, as proposed by Baroni et al. (2011, 2019), peaks in ^{10}Be concentrations in ice cores may sometimes be linked to stratospheric volcanic eruptions. The Agung and Pinatubo eruptions, for instance, are estimated to have caused an increase in ^{10}Be deposition of 66% and 35% relative to the baseline radionuclide concentration in the Vostok ice core from Antarctica (Baroni et al., 2011).

70 Especially for detecting the short-lived spikes caused by SEPs in the past, high-resolution measurements are required. These are often labor-intensive and limited by the availability of sufficient ice-core sample material. In this study, we aim at testing the use of excess meltwater samples usually discarded from continuous flow analysis (CFA) for the measurement of cosmogenic ^{10}Be . This method could provide the potential for high-resolution and cost-efficient ^{10}Be sampling without the need for competing for the valuable ice. We collected the meltwater from
75 the CFA system at the Institute for Climate and Environmental Physics at the University of Bern (Switzerland). The system is used to measure impurities in ice cores (e.g. Erhardt et al., 2022), which requires the use of pristine ice. The system is equipped with a melt head designed to only capture the inner part of the ice core for the measurement of impurities, preventing the ice from being possibly contaminated by the surrounding atmosphere (Erhardt et al., 2019). The outer part can in theory be used for cosmogenic radionuclide measurements as they are
80 less prone to contamination. This technique has already been used to collect a short record of about 40 years from the EGRIP ice core to study an extreme SEP event that hit Earth 9,125 years BP (Paleari et al., 2022). While suitable for the assessment of the existence of the peak in ^{10}Be concentrations, it was not possible to analyze in



depth and ultimately quantify the uncertainties related to this method, as the record was only compared to ice core records from other locations from Greenland and Antarctica. Here, we present a new ^{10}Be record from the S6 firm core from EGRIP (East Greenland Ice core Project, see Fig. 1 for the location), covering the depth from about 4 to 26m, corresponding approximately to the period between 1900 and 2008 CE. To assess the suitability of the CFA samples to measure cosmogenic radionuclides and retrieve a solar signal, ^{10}Be records obtained from CFA and discrete firm samples from the EGRIP S6 core (Zheng et al., submitted) are compared. The use of two ^{10}Be records from the same core allows us to directly assess the quality and possible differences between the sampling methods.

In addition, we investigate the causes of short-term and non-GCR related ^{10}Be variability in the ^{10}Be records from EGRIP S6 and other available records from Greenland and Antarctica. More specifically, we aim to investigate whether we can detect the signal of large SEP events (or GLEs) and volcanic eruptions. Furthermore, we investigate the preservation of the 11-year solar cycle in the ^{10}Be records.

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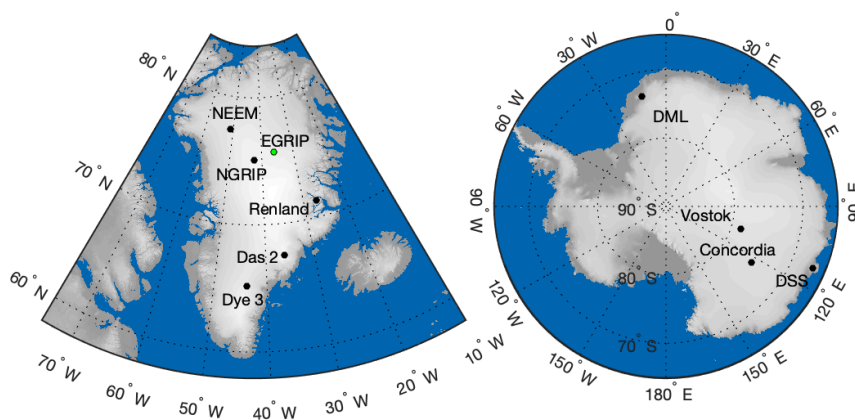


Figure 1. Location of the EGRIP site (marked with a green circle) and other ice core locations from Greenland (left) and Antarctica (right) discussed in this work.

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2 Methods

2.1 Collection and preparation of the CFA samples

The CFA samples were collected during the melting campaign carried out at the Institute for Climate and Environmental Physics at the University of Bern (Switzerland) in October 2019. During the campaign, 36x36 mm vertical sticks of ice of 1m in length were continuously melted on a melt head. The melt head is designed to only inject meltwater from the inner part of the core for measurement (26x26 mm, e.g. Erhardt et al., 2022), while the meltwater from the outer part of the core, otherwise discarded, is pumped towards centrifuge tubes for continuous ^{10}Be sampling. To collect the water, we used 50ml centrifuge tubes where 0.1 mg ^9Be carrier was previously injected.

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The preparation of the samples for Accelerator Mass Spectrometry (AMS) measurements was carried out at the Department of Geology at Lund university (Sweden). Each ~50ml sample was directly precipitated with NH_4OH . After centrifugation, the precipitate was transferred to a quartz crucible. During this step, consecutive samples were paired to obtain samples of ~100g with 0.2mg of ^9Be carrier. The crucibles were then placed into an oven
115 where the temperature is slowly raised to 850°C , allowing the oxidization of $\text{Be}(\text{OH})_2$ to BeO . Finally, 1mg of Niobium was added and the mixture was pressed into target holders for measurements. The measurements were carried out at the Laboratory of Ion Beam Physics at ETH, Zurich (Switzerland). The record was blank corrected ($^{10}\text{Be}/^9\text{Be}$ blank ratios: ~6% of the $^{10}\text{Be}/^9\text{Be}$ samples ratio; average $^{10}\text{Be}/^9\text{Be}$ of the blank samples: 0.007×10^{-12}). The measured $^{10}\text{Be}/^9\text{Be}$ ratios were normalized to the ETH Zurich in house standards S2007N and S2010N57,
120 which were both calibrated relative to the ICN 01-5-1 standard ($^{10}\text{Be}/^9\text{Be} = 2.709 \times 10^{-11}$ nominal) (Christl et al., 2013).

2.2 Depth attribution of the samples

The depth-sample relationship has to be reconstructed since the continuous melting does not allow a precise
125 measurement of the depth range covered by each CFA ^{10}Be sample. During sample collection it is possible to know in which vial there is a transition between two consecutive 1m sections, and the samples in between are assumed to contain equidistant sections of firn in terms of ice-equivalent depth ignoring any short term fluctuations in melt speed. This is probably not accurate as the layer thickness is not constant, which could lead to some uncertainty in the depth attribution of the single centrifuge tubes (likely <6cm, 3-4 months on average).
130 We thus consider this to not affect significantly our results (see Section 3).

2.3 Timescale of the EGRIP S6 core

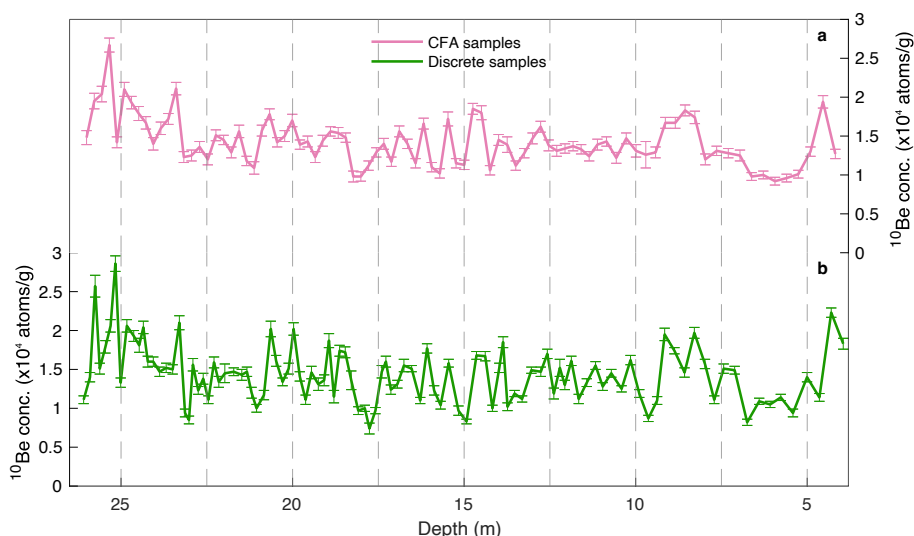
The timescale of the EGRIP S6 core is based on annual layer identification in the impurity record obtained from the CFA measurements collected at the Institute for Climate and Environmental Physics at the University of Bern
135 and an updated age-model proposed by Zheng et al. (submitted). The first chronological marker identified and used to constrain the chronology is a volcanic tie point in 1912 CE. From this tie point, the time scale is adjusted within annual layer counting uncertainty based on work by Zheng et al. (submitted) who identified the timescale offset (i) between the EGRIP discrete firn ^{10}Be data and the theoretical ^{10}Be production rate referred from the sunspot numbers and (ii) between accumulation rates (layer thickness) and precipitation. For consistency, in this
140 study we adopt the updated chronology by Zheng et al. (submitted).

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3 ^{10}Be record from CFA excess water samples



155 Figure 2. ^{10}Be concentration record from CFA excess water samples (panel a) and ^{10}Be concentrations from
discrete firn samples (panel b - Zheng et al., submitted) from the EGRIP S6 core. The records are plotted versus
the depth of the core. The error bars represent the 1σ measurement uncertainty.

The ^{10}Be record from EGRIP S6 CFA excess water samples is shown in pink in Figure 2a. The ^{10}Be concentrations
160 were measured in firn from about 4 to 26 m of depth, encompassing the period 1900–2008, according to the
chronology by Zheng et al. (submitted). The samples correspond to an average length of 25 cm, resulting in an
average time resolution of 1.2 years, with a minimum of 0.7 years and maximum of 2 years depending on annual
layer thickness and firn density. The ^{10}Be concentrations have an average of $1.43 (\pm 0.31) \times 10^4$ atoms/g. Figure
2b shows the ^{10}Be concentrations from discrete firn samples from the same core. The average ^{10}Be concentration
165 of the discrete record is $1.42 (\pm 0.32) \times 10^4$ atoms/g (Zheng et al., submitted). The meltwater samples have, on
average, a measurement uncertainty of 5%, similar to the ^{10}Be measurements carried out on the discrete firn
samples (Zheng et al., submitted).

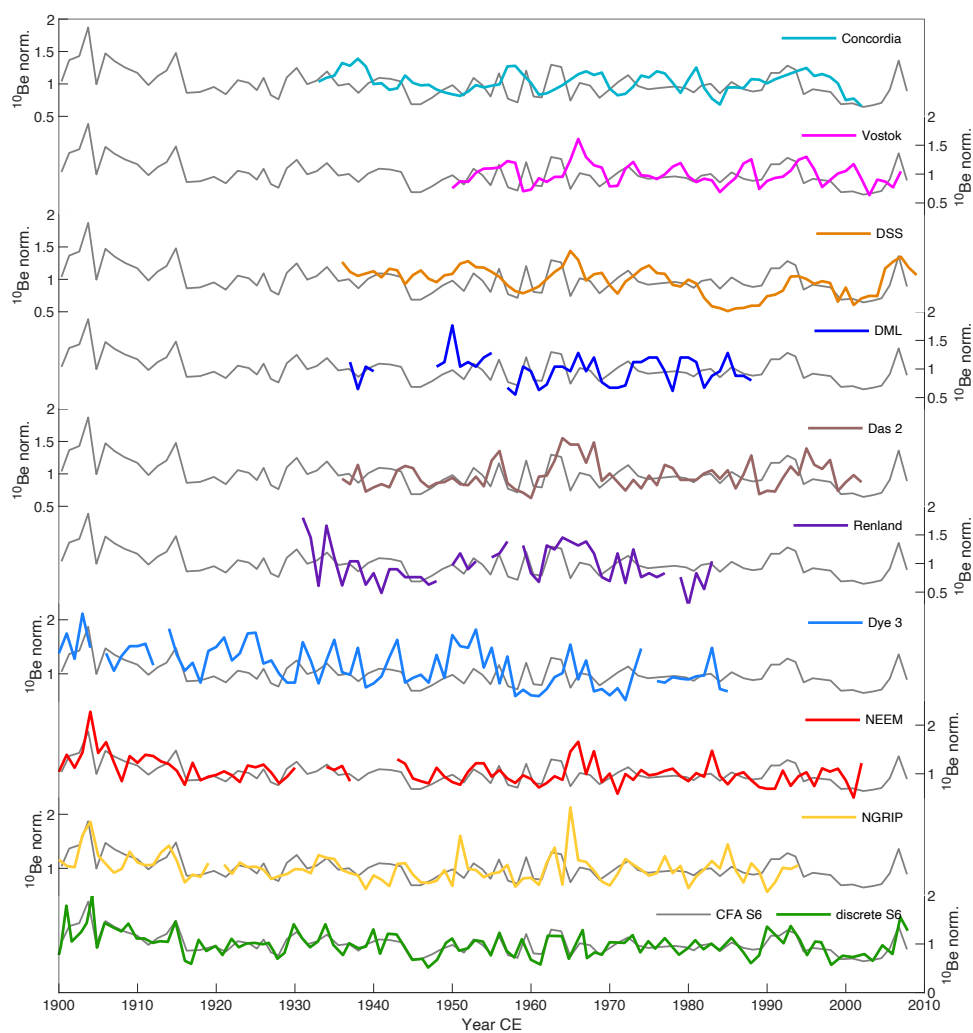
As the sampling rates of the two S6 records are different, the ^{10}Be records need to be virtually resampled to be
directly compared. The ^{10}Be concentrations from CFA samples and from discrete samples were interpolated
170 linearly at steps of 0.1 year and the yearly ^{10}Be concentrations were calculated by averaging over 10 consecutive
datapoints. The two yearly resampled ^{10}Be records agree well ($r=0.77$, $p<0.01$ - computed using a t-test). If we
resample the discrete record to match the resolution of the CFA ^{10}Be record, we obtain a correlation coefficient
of 0.71 ($p<0.01$). It can, however, be pointed out that the first ~ 8 m of the CFA excess water record (between ~ 4
and ~ 12 m of depth) is seemingly more autocorrelated compared to the deeper part of the record and relative to
175 the discrete ^{10}Be record. The sampling method and rate can likely affect the degree to which the two records agree.



¹⁰Be deposition is known to be highly seasonal, with maxima closely related to the enhanced stratosphere-troposphere exchanges (STE) causing increased descent of ¹⁰Be from the stratosphere, where most of the production takes place, to the troposphere (Stohl et al., 2003; Zheng et al., 2020; Spiegl et al., 2022). Moreover, one of the main complications of dealing with CFA systems is the possible smoothing of the signal locked in the ice (Erhardt et al., 2022; Kaufmann et al., 2008; Mekhaldi et al., 2017; Sigg et al., 1994). ¹⁰Be ions may attach to the walls of the plastic tube connecting the melt head to the centrifuge tubes, as ¹⁰Be precipitates in a basic environment. Therefore, it is possible that some atoms may precipitate on the plastic walls and get remobilized at a later stage. Loss of ¹⁰Be to the centrifuge tube walls is considered unlikely since the ⁹Be carrier (slightly acidic) was added to the centrifuge tubes before collecting the samples, and the ¹⁰Be/⁹Be ratio is in theory preserved. It is however most likely that the smoothing in the shallower part of the CFA ¹⁰Be record is caused by analytical issues related to the upward wicking of meltwater in the very porous snow during melting, which affects the shallow most parts of the core.

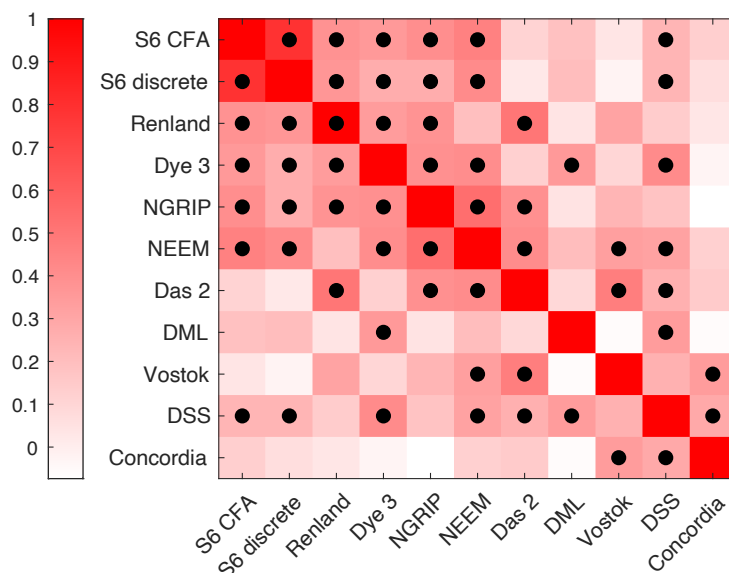
190 **4 Comparison with other records from Greenland and Antarctica**

To investigate whether the CFA excess water samples are suitable to preserve the ¹⁰Be production rate signal as robustly as discrete firn samples, the ¹⁰Be records from EGRIP S6 are compared to other available ¹⁰Be records from Greenland and Antarctica. We compare the ¹⁰Be record from CFA samples to ¹⁰Be records from the NGRIP (Berggren et al., 2009), NEEM (Zheng et al., 2021), Dye 3 (Beer et al., 1990), Renland (Aldahan et al., 1998), Das 2 (Pedro et al., 2012), DML (Aldahan et al., 1998), DSS (Pedro et al., 2012), Vostok (Baroni et al., 2011) and Concordia (Baroni et al., 2011) cores. The NEEM record is available at sub-annual resolution, and thus, we calculated mass-weighted means of each year. Figure 3 shows the comparison between the CFA ¹⁰Be record (grey line) and the other records from Greenland and Antarctica starting in 1900 CE. The degree to which the CFA ¹⁰Be record agrees with the other ¹⁰Be records can be seen in the correlation matrix in Figure 4. For this analysis, we used the CFA and discrete ¹⁰Be records resampled at 1 year resolution, to match the timescale of the other records. To calculate the correlation coefficients, the records were normalized to their mean for the period 1949-1985, representing the time span shared by all records.



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Figure 3. Normalized ^{10}Be records from the EGRIP S6 CFA samples in comparison to the normalized ^{10}Be records from discrete samples (Zheng et al., submitted), NGRIP (Berggren et al., 2009), NEEM (Zheng et al., 2021), Dye 3 (Beer et al., 1990), Renland (Aldahan et al., 1998), Das 2 (Pedro et al., 2012), DML (Aldahan et al., 1998), DSS (Pedro et al., 2021), Vostok (Baroni et al., 2011) and Concordia (Baroni et al., 2011) starting in 1900 CE.



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Figure 4. Correlation matrix (Pearson correlation coefficient) between the EGRIP S6 CFA and discrete samples, Renland, Dye 3, NGRIP, NEEM, Das 2, DML, Vostok, DSS and Concordia since 1900 CE. Dots represent significant correlations ($p < 0.05$).

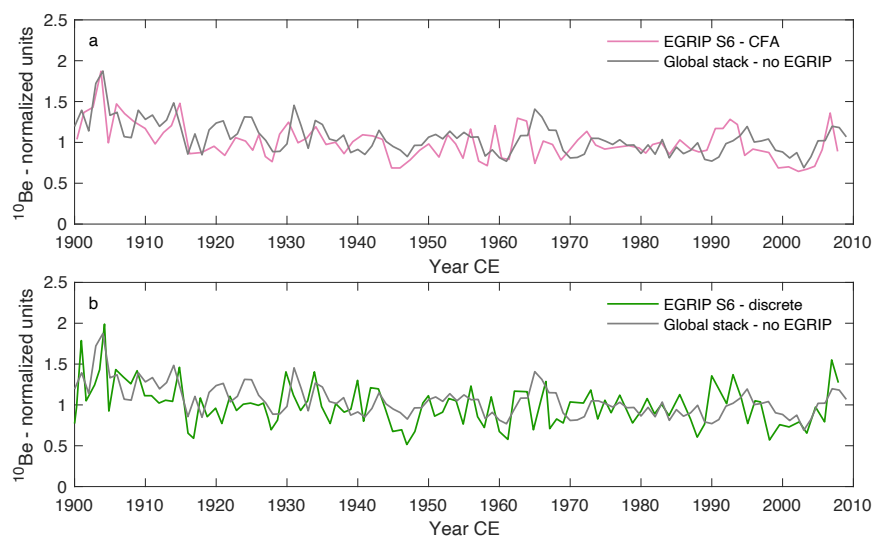
215 In general, the EGRIP CFA ^{10}Be record shows similar but somewhat higher correlation coefficients with the other records from Greenland and Antarctica, compared with the discrete EGRIP ^{10}Be record. While the higher correlation coefficients can likely be attributed to the higher autocorrelation inherent to the smoothing effect of CFA sampling in the shallower part of the record, our results indicate that the signal measured in the CFA samples is reproducing the common radionuclide signal in Greenland and Antarctica as well as the discrete firm samples.

220 The EGRIP S6 records from CFA and discrete firm samples agree well with the ^{10}Be records from Greenland, except for Das 2 (See Figure 4). This could be explained by the very high ^{10}Be concentrations in the Das2 record during the 1960s, corresponding to a pronounced minimum in accumulation rate (Pedro et al., 2012). Moreover, the differences between the EGRIP S6 records and the Das2 record in the early 1990s likely affect the correlation coefficient. Unsurprisingly, we obtain lower correlation coefficients with the ^{10}Be records from Antarctica.

225 Differences between the records could arise from non-production related biases such as measurement and sampling uncertainty, meteorological influences on the transport and deposition of ^{10}Be en-route and on site, as well as local re-distribution of snow by surface winds. For instance, Heikkilä and Smith (2013) found a strong correlations between ^{10}Be deposition and the North Atlantic Oscillation (NAO) in Greenland, with a positive correlation on the east coast and negative on the west coast.

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CFA and discrete ^{10}Be records are then compared with the stack of the normalized ^{10}Be records from Greenland (excluding EGRIP S6 records) and Antarctica (Figure 5). The calculation of a stack with several records from different sites allows us to reduce the noise by isolating their common signal which is expected to more closely reflect the atmospheric production signal.



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Figure 5. Comparison between the global stack calculated using the normalized ^{10}Be records from Greenland and Antarctica (NGRIP, NEEM, Renland, Dye 3, Das 2, DML, DSS, Vostok and Concordia) and the normalized ^{10}Be records from EGRIP S6 CFA (panel a) and discrete (panel b) samples.

240 The CFA ^{10}Be record has a correlation coefficient of 0.59 ($p < 0.01$) with the global ^{10}Be stack, while the ^{10}Be record from discrete firn samples has a correlation coefficient of 0.48 ($p < 0.01$). The records agree particularly well during the period 1900-1960, where we find a correlation coefficient of 0.66 ($p < 0.01$) with the ^{10}Be record from CFA samples and 0.59 ($p < 0.01$) with the discrete ^{10}Be record. The agreement decreases after 1960 ($r = 0.30$, $p < 0.05$ for CFA samples; we do not obtain significant results for the discrete record). The correlation between the
245 EGRIP S6 ^{10}Be records and the stack is improved if the EGRIP S6 records are shifted one year (+1) in the section 1960-2008 ($r = 0.52$, $p < 0.01$ for the CFA ^{10}Be record; $r = 0.32$, $p < 0.05$ for the discrete ^{10}Be record), even though the correlation could be influenced by peaks that may be caused by site-specific influences, such as the peak in the early 1990s, not prominently seen in any of the other ^{10}Be records (see Figure 3).

250 5 Potential SEP and volcanic signals in ^{10}Be

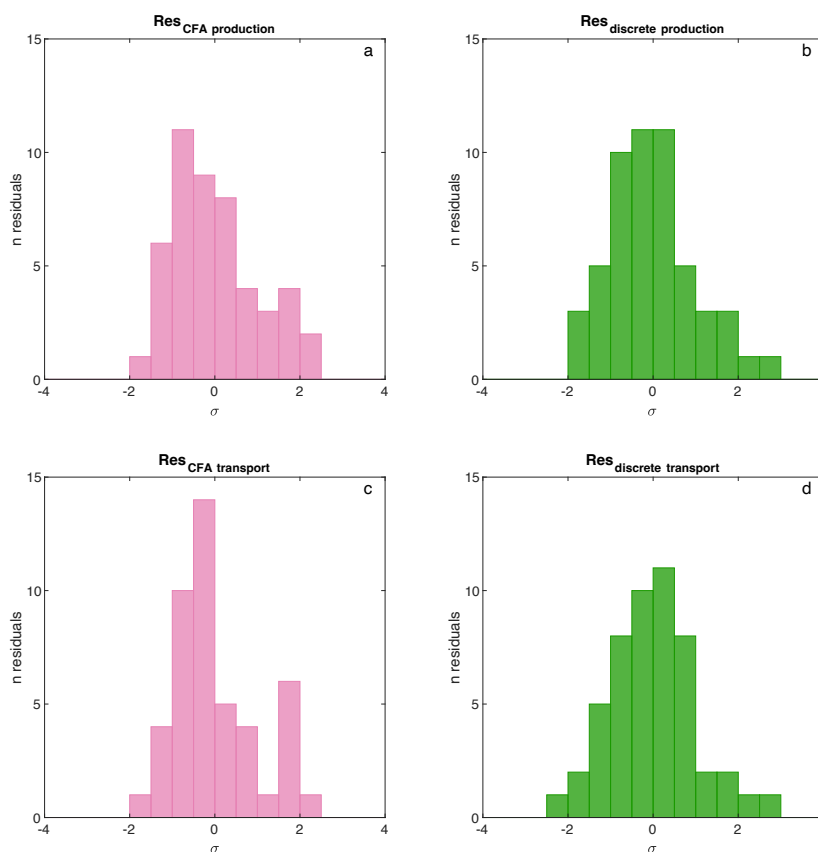
5.1 EGRIP S6 records

255 ^{10}Be records from ice cores are valuable tools to reconstruct solar activity prior to the advent of spaceborne measurements, ground-based instrumental records and sunspot observations. Nevertheless, the interpretation of ^{10}Be records is complicated by the influence of climate, local weather noise (e.g. Pedro et al., 2011) and the stochastic occurrence of volcanic eruptions (Baroni et al., 2011, 2019).

Here, we investigate outliers in the two ^{10}Be records from the EGRIP S6 firn core to assess whether it is possible to infer their cause. To do so, the normalized globally averaged ^{10}Be production rate caused by GCR (Mekhaldi



260 et al., 2021) was virtually resampled at the resolution of the CFA and discrete ^{10}Be records. This resampled
production rate was subtracted from the two normalized ^{10}Be records from EGRIP S6 for the period from 1951 to
2008 assuming that this removes the variability induced by the 11-year solar cycle. We then standardized the
residuals to z-scores, i.e. the number of standard deviations by which the residuals are above or below their mean
value. We repeated the experiment using the modeled normalized ^{10}Be deposition flux over latitudes 60-90N
265 caused by GCR and including the transport modeled using the parametrization from Heikkilä et al. (2009)
(Mekhaldi et al., 2021). From this point on, we will refer to $\text{Res}_{\text{recordname_prod}}$ and $\text{Res}_{\text{recordname_transport}}$. The
distributions of resulting residuals are shown in Figure 6 (CFA samples in pink, and discrete firn samples in green)
and a summary of the results is shown in Table 1.



270 Figure 6. Distribution of the standard scores for the EGRIP S6 ^{10}Be records. Panel a and b show the distribution
of the standardized residuals obtained by subtracting the normalized theoretical ^{10}Be global production rate from
the normalized EGRIP S6 ^{10}Be records since 1951 (CFA ^{10}Be sample residuals in pink, discrete ^{10}Be sample
residuals in green). Panel c and d show the same as panel a and b, but for the ^{10}Be deposition flux including the
transport over latitudes 60-90N (Mekhaldi et al., 2021).



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The distributions of the residuals can be approximated to a normal distribution, with a longer tail towards $+2-3\sigma$. Although we expect some of these results to be exclusively caused by noise inherent to the data, values exceeding 1σ in the years following a stratospheric volcanic eruption and/or a solar storm may be more likely. Several of the major stratospheric volcanic eruptions from the last 70 years have been identified in sulfate records from Greenland in 1957, 1964, 1982-84 and 1991 (Sigl et al., 2015; Zielinski et al., 1997). Some of the eruptions were traced to, for instance, Agung (in 1963), El Chichón (in 1982) and Pinatubo (in 1991). Taking into account timescale uncertainties and the lag due to the transport of ^{10}Be from the stratosphere, we would expect to find values exceeding 1σ in the years of the eruptions and/or 1 to 2 years after the eruptions. Values exceeding 1σ can be identified during years 1956, 1959, 1963, 1965, 1990, 1991, 1992, 1993 and 2006 in the $\text{ReSCFA}_{\text{prod}}$ and in 1959, 1963, 1965, 1990, 1991, 1992, 1993 and 2006 in the $\text{ReSCFA}_{\text{transport}}$. Values exceeding 1σ in 1956, 1959, 1990, 1991 and 1993 can be identified also in $\text{ReS}_{\text{discrete_prod}}$, in addition to years 1967, 1982 and 2007. Values exceeding 1σ can be identified in $\text{ReS}_{\text{discrete_transport}}$ in years 1959, 1967, 1990, 1991, 1993 and 2007. Some of these results are coeval with the timing of the Agung, El Chichón and Pinatubo eruptions, albeit the Pinatubo eruption is generally not particularly prominent in the other ^{10}Be records from Greenland (see Figure 3) and the uncertainty of the S6 chronology may affect our interpretation.

Although it is reasonable to expect values exceeding 1σ to be related to stochastic noise, we want to investigate whether two years characterized by some of the largest GLEs, which occurred in 1956 and 1989 (GLE no.05 and GLEs no.41-45), left any traces in the ^{10}Be records from ice cores. GLE no.05 and GLEs no.41-45 are the events that yielded the most annual production of ^{10}Be over the last 70 years of instrumental data, with an estimated increase of $+5.1\%$ and 4.6% , respectively, in the modeled annual ^{10}Be global production rate (Mekhaldi et al., 2021). The analysis of the standard scores shows that values exceeding 1σ can be identified in $\text{ReSCFA}_{\text{prod}}$ during years 1956 and 1990, and 1990 in the $\text{ReSCFA}_{\text{transport}}$, similarly to $\text{ReS}_{\text{discrete_prod}}$ and $\text{ReS}_{\text{discrete_transport}}$. We find an increase in ^{10}Be concentrations of 19% and 28% during years 1956 and 1990, respectively, in the CFA record, and an increase of 42% and 44% during years 1956 and 1990 in the discrete record. The increases were calculated relatively to the average ^{10}Be concentration over solar cycles 19 and 22 (excluding the targeted years). However, we cannot exclude that the increase in ^{10}Be concentration is caused mostly by the natural variability of ^{10}Be with a more modest contribution of the SEP events and/or volcanic eruptions. Moreover, although the increase in 1990 falls in the $2-3\sigma$ envelope (see S1), the increase in 1956 is in the lower end of the $1-2\sigma$ envelope, showing that expected signals in the 5% range are impossible to unequivocally detect in the radionuclide data.

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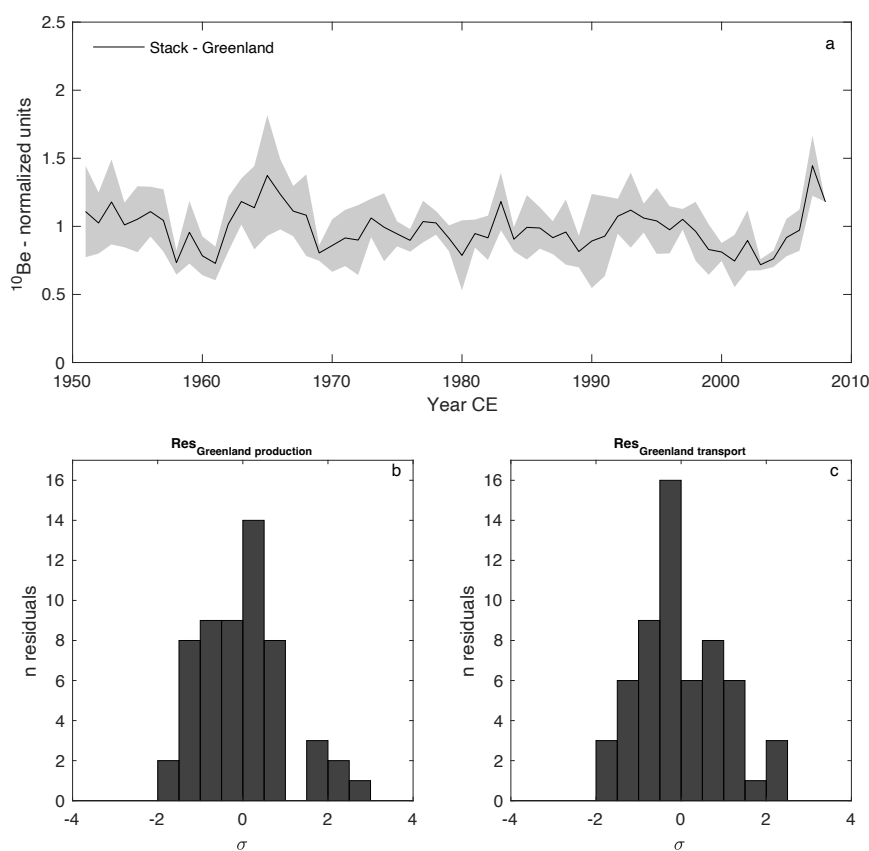
5.2 ^{10}Be stacks from Greenland and Antarctica

In the following section we analyzed the standard scores of the residuals obtained by subtracting the normalized theoretical ^{10}Be production rate (by GCR) from the Greenlandic and Antarctic stacks. As volcanic eruptions may impact ^{10}Be records from Greenland and Antarctica differently, the analysis of the standard scores has been performed separately on the stacks from the two regions.

The stack from Greenland (see Figure 7a) has been calculated using all the records from Greenland from Fig. 3 normalized to their respective mean during the period 1951-1985, representing the time span shared by all the records. Figure 7b shows the distribution of the standard scores calculated on the residuals obtained by subtracting



315 the modeled normalized ^{10}Be production rate (Mekhaldi et al., 2021) from the Greenland ^{10}Be stack (Res_{Greenland_prod}). Figure 7c shows the distribution of the standard scores calculated on the residuals obtained by subtracting the normalized ^{10}Be modeled deposition flux including the transport in the northern hemisphere (Mekhaldi et al., 2021) (Res_{Greenland_transport}).

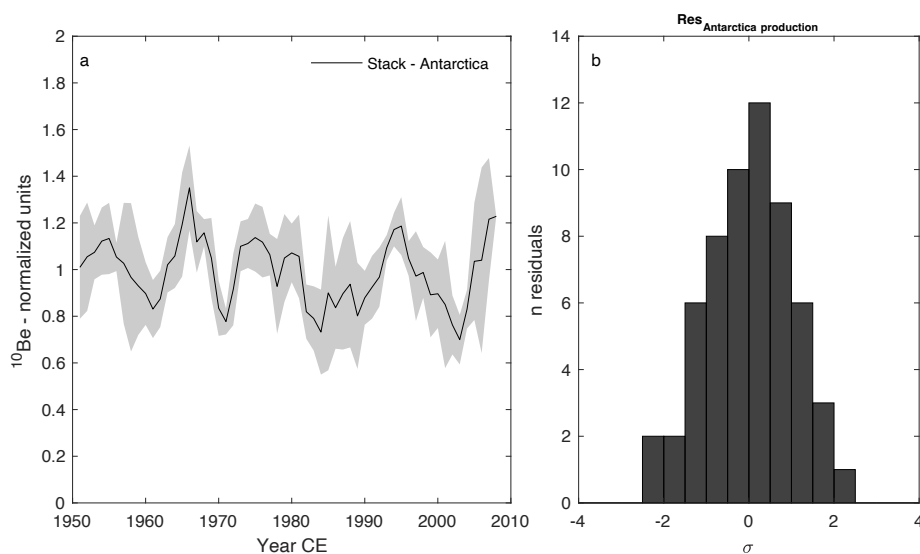


320 Figure 7. Distribution of the standard scores for the difference between the Greenland ^{10}Be stack and the
 theoretical GCR-induced production rate. The stack (panel a) was calculated using the normalized ^{10}Be records
 from Dye 3, NGRIP, NEEM, Das 2, Renland and EGRIP S6 (CFA and discrete) since 1951. Panel b shows the
 distribution of the standardized residuals obtained by subtracting the modeled ^{10}Be global production rate from
 the stack. Panel c shows the same as panel b but using the normalized ^{10}Be deposition flux modeled (taking
 325 transport into account) over latitudes 60-90N (Mekhaldi et al., 2021).

The standard scores are above -2σ , while they present a tail skewed towards $2-3\sigma$, in agreement with the results from the EGRIP S6 records. The years in which the standard scores exceeded 1σ are summarized in Table 1. We find Res_{Greenland_prod} exceeding 1σ during years 1957, 1959, 1963, 1965, 1966, 1968, 1983, 1991, 1992 and 2007.



330 The $\text{Res}_{\text{Greenland_transport}}$ show values exceeding 1σ in 1963, 1965, 1983, 1992, 1993 and 2007. We find an increase
of 7.5% in ^{10}Be concentrations in year 1957 relative to the baseline (average ^{10}Be concentration over solar cycle
19). However, the increase in ^{10}Be concentrations is coeval with a peak detected in the NEEM sulfate record (Sigl
et al., 2013). Therefore, the presence of a ^{10}Be increase in ice cores from Antarctica would lend support to the
possible contribution of GLE no.05 to the ^{10}Be deposited in Greenland and Antarctica in 1957. We carried out the
335 same analysis on the stack calculated from the Antarctic records. We subtracted the normalized global ^{10}Be
production rate from the stack of DSS, DML, Concordia and Vostok normalized ^{10}Be records and calculated the
standard scores. The stack and distribution of the residuals are shown in Figure 8.



340 *Figure 8. The distribution of the standard scores for the Antarctic stack. The stack (panel a) was calculated using
the normalized ^{10}Be records from Vostok, DML, Concordia and DSS from 1951. The standard scores were
calculated on the residuals obtained by subtracting the normalized modeled ^{10}Be global production rate from the
stack. The distribution of the standard scores is shown in panel b.*

We find that years 1957, 1958, 1966, 1967, 1968, 1969, 1980, 1981, 1991 and 1994 exceed 1σ . The increase in
345 1994 can be linked to a possible volcanic influence on the ^{10}Be transport and/or deposition (Baroni et al. 2011).
We do not find any evidence of the El Chichón eruption in the Antarctic stack, although the eruption left an
imprint in some Antarctic sulfate records (Sigl et al., 2015). We find an increase relative to the baseline of 3.9%
in 1957, lower than the increase modeled by Mekhaldi et al. (2021) and the increase recorded in the stack of
records from Greenland.

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355

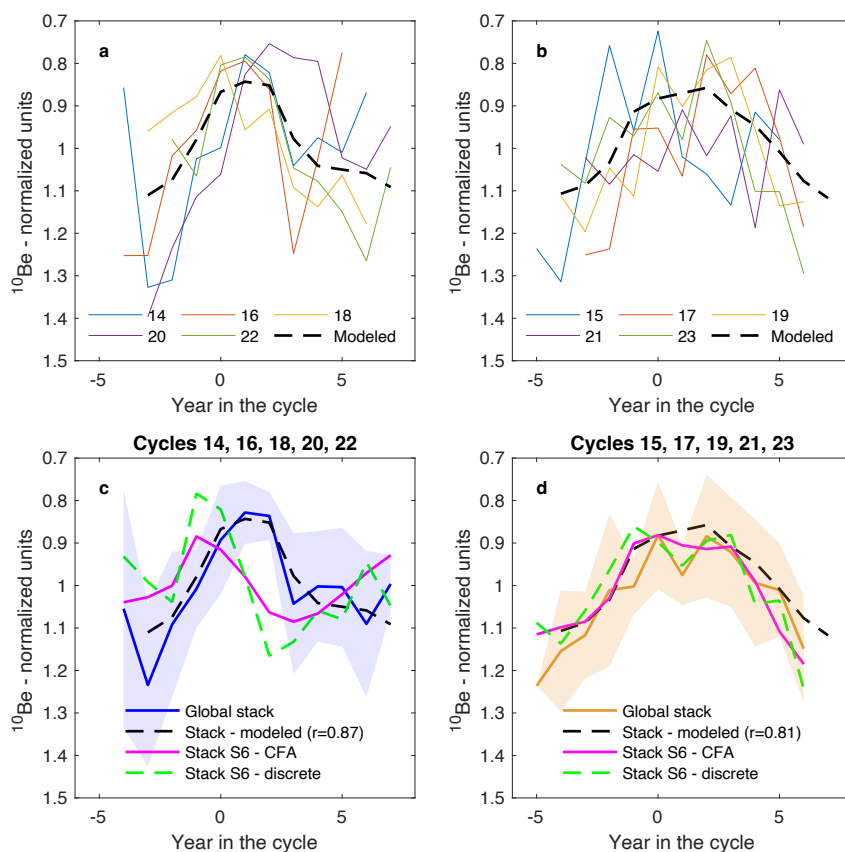
Records	1-2 σ	2-3 σ
¹⁰ Be EGRIP S6 - CFA	1956, 1963, 1965, 1991, 1992, 1993, 2006	1959, 1990
Including transport:	1959, 1963, 1965, 1990, 1991, 1993, 2006	1992
¹⁰ Be EGRIP S6 - discrete	1956, 1959, 1967, 1982, 1991, 1993, 2007	1990
Including transport:	1959, 1967, 1991, 1993	1990, 2007
¹⁰ Be stack – Greenland	1957, 1959, 1963, 1966, 1968, 1991, 1992	1965, 1983, 2007
Including transport:	1963, 1992, 1993	1965, 1983, 2007
¹⁰ Be stack - Antarctica	1957, 1958, 1967, 1968, 1969, 1980, 1981, 1991, 1994	1966

Table 1. Summary of the years during which we find standard scores exceeding 1σ from the GCR-production corrected EGRIP S6 records, Greenland and Antarctic stacks.

360

6 The 11-year solar cycle

To assess the suitability of CFA meltwater samples to reconstruct solar activity, we investigate the presence of the 11 year solar cycle in the data. The wiggles of the 11 year solar cycles are estimated to cause a variability of about $\pm 15\%$ to $\pm 35\%$ in the ¹⁰Be records from Greenland and Antarctica (e.g. Baroni et al., 2011; Heikkilä et al., 2009; Mekhaldi et al., 2021; Paleari et al., 2022). In addition, neutron monitor measurements of the GCR flux to Earth show that the 11-year cycle has a distinct shape (sharper peak versus broader peak) depending on the polarity of the solar magnetic field, which reverses every ~ 11 years (e.g. Lockwood et al., 2001). Bearing this difference in mind, we separate our analysis between even (sharper peak) and odd (broader peak) cycles. The ¹⁰Be records from Greenland (except EGRIP S6) and Antarctica were split into time windows for cycles 14 to 23 and normalized to their respective mean, and therefore used to create a stack for each cycle. Figure 9a and 9b shows the stacks of even (14, 16, 18, 20, 22) and odd (15, 17, 19, 21, 23) solar cycles, respectively, centered around the solar maximum (identified using the sunspot record), indicated as year 0. The stacks of even and odd cycles obtained from the modeled ¹⁰Be production rate from Mekhaldi et al. (2021) are shown as a dashed line for comparison. The data shown in figure 9a and 9b were used to calculate a global stack of even and odd cycles, shown in blue and orange in figure 9c and 9d, respectively. The blue and orange envelopes denote the standard deviation of the records. The global stack is then compared to the modeled stack (black dashed line), CFA stack (magenta) and discrete firn samples stack (green) from EGRIP S6.



380 Figure 9. Panel a and b show the normalized ^{10}Be stack of even (a) and odd (b) cycles including the records from
 the NEEM, NGRIP, Dye 3, Das 2, Renland, DSS, Vostok, DML and Concordia cores, compared to the stacks from
 the modeled global production rate from Mekhaldi et al. (2021) (dashed black line). Panel c shows the global
 stack of even cycles (in blue, the blue envelope represents the standard deviation), the stack of even cycles from
 the discrete samples (dashed green line) and the stack of even cycles from CFA samples (solid magenta line) from
 385 EGRIP S6. The stacks are compared to the even-cycle average of the modeled production rate (dashed black line,
 correlation coefficient with the global stack denoted in the legend). Panel d shows the same for odd cycles. The
 global stack for the odd cycles is shown in orange.

Figure 9a and 9b show that, on average, the 11-year solar cycles are well-preserved in the ^{10}Be records from
 390 Greenland and Antarctica. The stacks from discrete and CFA samples agree very well for both even ($r=0.78$,
 $p<0.01$) and odd cycles ($r=0.92$, $p<0.01$), although the S6 CFA stacks are smoother than the discrete. Moreover,
 the S6 stacks for the odd cycles agree very well with the global and modeled stack. The S6 stacks for the even
 cycles agree well with the global and modeled stack in shape and amplitude, though they are both characterized



by a shift in the timing of the solar maximum. This may be due to the interplay of several factors, such as
395 uncertainties in the timescale, local weather and climate influences and noise inherent to the data. However, CFA
and discrete stacks agree well, showing that the two records show a very similar signal. This result is promising,
as it shows the suitability of CFA samples for solar activity reconstructions.

7 Conclusions

400

In this study, we present the ^{10}Be record obtained from excess water from CFA measurements of the EGRIP S6
firn core. The record, spanning from 1900 to 2008, has been compared to another ^{10}Be record obtained from
discrete firn samples from the same core. Although some smoothing due to analytical issues inherent to the CFA
system may be present in the shallower part of the core, the CFA ^{10}Be record agrees well with the ^{10}Be record
405 from discrete firn samples and with other ^{10}Be records from Greenland.

By subtracting the global ^{10}Be production rate variability caused by the 11-year solar cycle, we investigate the
possible causes of excursions in the EGRIP S6 ^{10}Be records and in the stacks of Greenlandic and Antarctic records.
We find that values exceeding 1σ in the standardized residuals are coeval with the timing of some of the major
volcanic eruptions that are also detected in sulfate records (Zielinski et al., 1997; Sigl et al., 2015), such as the
410 Agung (1963) and Pinatubo (1991) eruptions. Moreover, values exceeding 1σ in the residuals in 1956/1957 and
1990 may be consistent with the occurrence of GLEs no.5 and no.41-45. However, an increase in sulfate
concentrations was detected in records from Greenland according to Sigl et al. (2015), illustrating the challenges
to separate the causes for small radionuclide excursions. We find an increase of 7.5% in the stack from Greenland
and in 3.9% the stack from Antarctica in 1957.

415 Finally, we analyzed how the 11-year solar cycles are preserved in the discussed records. Our results show that
the 11-year solar cycles are well preserved in CFA and discrete ^{10}Be records from EGRIP S6. By stacking even
and odd cycles separately it is possible to see their different shape, as expected according to the flux of the GCR
to Earth. The even and odd cycle stacks from CFA ^{10}Be samples agree well with the stacks from discrete ^{10}Be
samples from EGRIP S6. Our results thus show that CFA excess water samples are suitable for the reconstruction
420 of solar activity. These results open the possibility of collecting continuous and high-resolution ^{10}Be records with
a more time-efficient sampling and preparation, while saving an important portion of valuable ice for other
measurements.

Data availability

425 The ^{10}Be data generated in this study is provided in the supplementary material.

Author contribution

R.M and C.P. designed the project. C.P. sampled and prepared the EGRIP CFA ^{10}Be samples, performed the
analysis, and wrote the manuscript. F.M. contributed to the interpretation of the data. M.C. contributed with the
430 measurement of ^{10}Be samples. T.E. helped with the collection of the EGRIP CFA samples and the interpretation
of the data. M.H. helped with the sampling of the EGRIP S6 core. All authors contributed to the discussion.



Competing interests

435 The authors declare that they have no conflict of interest.

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