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Changes in Holocene climate and the intensity of Southern Hemisphere Westerly Winds based on a high-resolution palynological record from sub-Antarctic South Georgia

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Abstract

Research paper

Sub-Antarctic South Georgia is a key region for studying climate variability in the Southern Hemisphere, because of its position at the core of the Southern Hemisphere Westerly Wind belt and between the Antarctic Circumpolar Current and the Polar Frontal Zone. Here, we present a 5.8-m long high-resolution pollen record from Fan Lake on Annenkov Island dominated by local sub-polar vegetation, with *Acaena* and Poaceae being present throughout the last 7000 years. Palynological and sedimentological analyses revealed a warm late Holocene 'climate optimum' between 3790 and 2750 cal. yr BP, which was followed by a gradual transition to cool and wet conditions. This cooling was interrupted by slightly warmer environmental conditions between 1670 and 710 cal. yr BP that partly overlap with the Northern Hemisphere 'Medieval Climate Anomaly'. Increases in non-native and long-distance pollen grains transported from South America (e.g. *Nothofagus, Podocarpus*) indicate that stronger Southern Hemisphere Westerly Winds over South Georgia possibly occurred during some 'colder' phases of the late Holocene, most notably between c. 2210 and 1670 cal. yr BP and after 710 cal. yr BP.

Keywords

Holocene, palynology, South Georgia, Southern Hemisphere Westerly Winds, sub-Antarctica, vegetation

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Introduction

The mid- to late Holocene climate of the Southern Hemisphere high latitudes is characterised by an early Holocene warm period from 11,000 to 9500 cal. yr BP, a general cooling until 4500 cal. yr BP and a mid–late Holocene warm period from 4500 to 2800 cal. yr BP or later (Bentley et al., 2009). Following this, a cooler period generally persisted until the present rapid regional warming of the last few decades with many regional variations (Hodgson et al., 2009).

The mid–late Holocene warm period in the Antarctic Peninsula region has been attributed to increases in solar insolation (Bentley et al., 2009) and changes in the meridional position of the core of the Southern Hemisphere Westerly Wind (SHW) belt. Understanding the link between climate changes and the position and strength in the SHW in this region is important because changes in the winds can affect upwelling in the Southern Ocean, which in turn affects global CO_2 levels (e.g. Anderson et al., 2009; Björck et al., 2012; Hodgson and Sime, 2010; Le Quéré et al., 2010; Watson and Garabato, 2006). Multiple climate model simulations (Varma et al., 2012) suggest an overall strengthening and poleward shifting trend in the SHW during the course of the mid- to late Holocene corresponding with a cooling trend in Antarctica (Masson et al., 2000). However, validating the model results with reconstructions of the

past position and intensity of the SHW still remains elusive. For example, Varma et al. (2012) note that proxy records from western Patagonia (Moreno et al., 2010) suggest a trend of increasing SHW strength during the past 7000 years, which is not supported by sedimentological and pollen-based reconstructions of precipitation variability in South Patagonia (see Fletcher and Moreno, 2012; Lamy et al., 2010).

These studies highlight a limitation of many high-latitude palaeoclimate records which show considerable variability not only in the timing of inferred warm and cool periods (which is further

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compounded by low resolution age control in many studies; Table 1), but also in differences in the precise climate forcing mechanisms used to explain them. In the core belt of the SHW, the lack of reliable reconstructions is particularly acute because of the limited amount of landmasses from which palaeoclimate records can be obtained. These landmasses consist of the southernmost tip of South America and a small number of islands in the sub-Antarctic region. Even at these locations, local orographic, glaciological and edaphic factors can overprint the regional climate signals.

Here, we present a well-dated, high-resolution pollen record of mid- to late Holocene vegetation and climate change from a lake sediment core from Fan Lake, Annenkov Island, South Georgia (Figure 1). The South Georgia archipelago lies in the core belt of the SHW, and its climate is governed by the latitudinal movement of the Polar Front Zone (PFZ) and the SHW (Bentley et al., 2007), which makes it particularly sensitive to even relatively minor climatic fluctuations. Most existing palaeorecords have been taken from the north and northeastern side of South Georgia (Bentley et al., 2007; Rosqvist and Schuber, 2003), but sites off the southwest coast, such as Annenkov Island, are thought to be particularly responsive because they are not affected by active glaciation, or orographic influences prevalent on the northeastern sites.

The overall aims of our project were (1) to establish when Annenkov Island was last glaciated, (2) to assess whether the timing and extent of any periods of Holocene glacier advance were synchronous or offset with known phases of glacier advance on mainland South Georgia and (3) to test the hypothesis that Holocene palaeoenvironmental change on Annenkov Island is more directly coupled to changes in the SHW circulation compared with sites elsewhere on South Georgia that are influenced by orographic effects. The first two aims are addressed further in forthcoming papers. In this paper, our aims were to identify long-term regional changes in climate and vegetation and to test whether long-distance pollen influx at Fan Lake is related to previously reconstructed changes in SHW circulation.

Site and present day vegetation

South Georgia is located in the South Atlantic Ocean approximately 1300 km east-southeast of the Falklands Islands (Figure 1). The island is ~170 km long with mountains (Allardyce Range) reaching up to 2000 m down the middle of the island (Bentley et al., 2007; Rosqvist and Schuber, 2003). Meteorological records from King Edward Point show well-defined summer and winter seasons (Barrow, 1978) on the north coast of South Georgia with a mean annual temperature of 2°C (Van der Putten et al., 2004) and a mean annual precipitation of around 1400 mm at sea level (Rosqvist and Schuber, 2003). South Georgia is positioned at the maximum northern limit of Antarctic pack ice, and over half the island is permanently covered in glaciers and ice fields (Bentley et al., 2007).

Annenkov Island (54°29'S-55°00'S, 35°30'W-38°30'W; Figure 1) is situated c. 15 km off the mid-south-west coast of South Georgia. Although it rises to 640 m (Olstad Peak; Figure 1c), no permanent snow or ice cover remains (Barrow, 1983). Fan Lake is one of two large permanent lakes on Annenkov Island. The lake is c. 430 m long, 200 m wide and located c. 90 m.a.s.l. on the southeastern corner of Annenkov Island (Pettigrew, 1981). The lake catchment consists of a shallow U-shaped valley surrounded by a series of ridges, which become progressively less vegetated with altitude (Figure 1d). There are several streams entering the lake, possibly changing course across the vegetated fan area through time. There is a small inflow through a peat land covered gravel delta at the western end, and at the eastern end an outflow stream cuts through two prominent sets of bedrock ridges and a landscape of hummocks and hollows filled by peat land. Pettigrew (1981) described glacial till exposed by the outflow stream from Fan Lake, but from our observations, this could also be heavily vegetated weathered bedrock. A steep, partially vegetated scree slope on the northern side of the lake is composed of fine-grained mudstone, which outcrops on a small hill to the north of the fault line. A waterfall-stream also enters the lake on the northern edge.

Cores were extracted from the deepest point of Fan Lake (18 m) and from the centre of a peat land area (henceforth referred to as KH4; 54°29.927'S, 37°02.876'W) *c*. 50 m from the eastern outflow stream. The latter was used to better constrain the local deglaciation history of the site.

The vegetation in the region is dominated by sub-Antarctic herbs and shrubs such as Acaena spp. (Acaena magellanica, A. magellanica × Acaena tenera, A. tenera), Colobanthus spp. (Colobanthus quitensis, Colobanthus subulatus) and Callitriche antarctica (Greene, 1969; Greene and Walton, 1975). A. magellanica herbfields are found in sheltered areas, while Colobanthus spp. is frequent in rock face and fellfield communities on Annenkov Island (Barrow, 1983). Grasslands (Poa flabellata) are abundant throughout South Georgia (Barrow, 1978; Greene, 1969). Ferns such as Polypodium australe, Grammitis kerguelensis, Hymenophyllum falklandicum, Blechnum penna-marina and the clubmoss Lycopodium magellanicum are found in some quantity (Greene, 1969; Greene and Walton, 1975). Montia fontana, a wet flush species, Ranunculus biternatus and C. antarctica, a waterstarwort plant, are found in wetter environments on South Georgia (Barrow, 1983).

One potential problem of reconstructing past vegetation on South Georgia is the selective preservation of pollen produced by some species. For example, while the flowering plants, *Juncus* (*Juncus inconspicuous, Juncus scheuchzerioides*) and *Rostkovia magellanica* are two of the most common native species (Greene, 1969; Van der Putten et al., 2010), the pollen are not preserved in fossil assemblages (Van der Putten et al., 2004).

South Georgia experienced little or no human activity until the establishment of sealing and whaling stations on the north coast of the island during the last c. 150 years (Barrow, 1978). As a result of these human impacts, a number of introduced plant species are present, but in many cases, these are restricted to the areas around the abandoned whaling stations (Osborne et al., 2009). Currently, 62 introduced species have been recorded on South Georgia (Osborne et al., 2009), but they are less pervasive on Annenkov Island because of the infrequency of human visitation. For some species, such as members of the Asteraceae, it remains unclear whether they were introduced or are native to the region (Birnie, 1990; Osborne et al., 2009).

Material and methods

Sedimentology and chronology

Sediment cores were extracted from Fan Lake (Figure 1d) using UWITEC gravity and percussion driven piston corers. Two 18-cm UWITEC gravity cores were extracted to recover the near-surface sediments from the main core site and sectioned at 0.5-cm intervals, with a thick moss layer preventing deeper coring. The UWI-TEC piston corer was deployed to extract deeper sediments in a series of 2-m long offset drives capturing 1.87-1.90m of sediment with 20 cm overlaps to produce a continuous sequence down to 5.8 m, where drilling was stopped by a large rock or bedrock. The cores were photographed (visual and x-radiographic imagery), macroscopically described and analysed for wet density, dry mass (105°C for 12h), organic content (loss-on-ignition (LOI) at 550°C for 2h) and carbonate composition (LOI at 925°C for 2h, multiplied by 1.36 to provide a first approximation of total carbonate) following standard methods (Dean, 1974). Known overlaps, the main changes in lithology defined initially by density, LOI, Barington Instruments MS2G loop sensor (1 cm sensor, 10-s

| Cool/warm event timings (cal. yr BP) | Location | Latitude (S) | Longitude (W) | No. of ¹⁴ C dates | Max. core age (cal. yr BP) | Mean no. of ¹⁴ C ages/1000years | Climate | Evidence | References |
|---|---------------------------------------|-----------------|-------------------|---------------------------------|-------------------------------|---|--|--|----------------------------------|
| After 1000 | Block Lake, South Georgia | 54° 10'17.57" | 36° 43'32.82" | 12 | 7451 | 1.61 | Cooling | Glacial advances | Rosqvist and Schuber (2003) |
| 2200–2000 | Stromness Bay, South Georgia | 54° 11' | 36° 43' | 6 | I 2493 | 0.72 | Colder/wetter | Addition of Bartramia spp. | Van der Putten et al. (2012a) |
| 2200 | South Georgia | 54° 30′–55° 00′ | 35° 30′–38° 30′ | 6 | 9146 | 0.66 | Possible Cooling/Climate deterioration | Vegetation change (Polytrichum–Chorisodontium) | Van der Putten et al. (2009) |
| c. 2120–543 | Beak Island, Ant. Peninsula | 63°36′ | 57°20′ | 20 | 10612 | I.88 | Climate deterioration | Decline in total pigment concentration/lower organic content | Sterken et al. (2012) |
| 2400-1600 | Block Lake, South Georgia | 54° | 36° | 12 | 7451 | 1.61 | Cooling | Glacial advances | Rosqvist and Schuber (2003) |
| c. 2492–1342 | South Shetland Islands, Antarctica | 62°37'43' | 61°08′05″ | 6 | c. 6036 | I.49 | Cold/drier | Extended ice cover | Björck et al. (1993) |
| c. 2546 | Tønsberg Peninsula, South Georgia | 54°09′41″ | 36°40'01 <i>"</i> | 6 | с. 10782 | 0.83 | Wetter conditions | Presence of oligotrophic bog, fresh layer of peat suggests climate deterioration | Van der Putten et al. (2004) |
| c. 4018–1080 | Signy Island, South Orkney Islands | 60°43′ | 45°36′ | 29 | с. 7420 | 3.91 | Climate optimum | Increase biological productivity/high % LOI values | Jones et al. (2000) |
| c. 3509–1662 | South Georgia | 54–55° | 36–38° | 8 | c. 7252 | ⊒ | Cooling | Vegetation disruption/ downslope debris flow | Clapperton et al. (1989) |
| c. 3169–2120 | Beak Island, Ant. Peninsula | 63°36′ | 57°20′ | 20 | 10612 | 1.88 | Mid-late Holocene climate optimum | Increased biological production | Sterken et al. (2012) |
| с. 3332– 2793 | Midge Lake, Livingston Island | 62.6° | 60.5° | Ŋ | c. 4055 | 1.23 | Mild/humid | Increase in <i>Pediastrum</i> , higher organic values | Björck et al. (1991) |
| c. 4375–3121 | South Shetland Islands, Antarctica | 62°37′43′ | 61°08′05″ | 6 | c. 6036 | I.49 | Mild/humid | Increased humidity | Björck et al. (1993) |
| c. 4701 | James Ross Island, Antarctica | 64°10′ | 57°45′ | ß | c. 4463 | 1.12 | More humid/warmer | Increase in Branchinecta gaini | Björck et al. (1996) |
| 4400–3400 | South Georgia | 54° 30′–55° 00′ | 35° 30′–38° 30′ | 6 | c. 9146 | 0.66 | Drier | Decrease in Warnstorfia fontinaliopsis | Van der Putten et al. (2009) |
| 4500- 3500 | Antarctic Peninsula | 63° 20.572′ | 55° 53.195′ | 29 | c. 9620 | 3.01 | Minor glacial retreat and/or decreased sea ice coverage | Minor glacial retreat/decreased sea ice coverage | Michalchuk et al. (2009) |

| Cool/warm event timings (cal. yr BP) | Location | Latitude (S) | Longitude (W) | No. of ¹⁴ C dates | Max. core age (cal. yr BP) | Mean no. of ¹⁴ C ages/1000years | Climate | Evidence | References |
|---|--|-----------------|-----------------|---------------------------------|-------------------------------|---|----------------------------------|--|---------------------------------|
| c. 4896–2546 | Tønsberg Peninsula, South Georgia | 54°09′41″ | 36°40'01″ | 6 | c. 10782 | 0.83 | Drier conditions prevailed | Chorisodontium–Polytrichum bank present | Van der Putten et al. (2004) |
| c. 5353– 4018 | South Georgia | 54–55° | 36–38° | ω | с. 7252 | Ξ | Climatic cooling | Vegetation disruption/ downslope debris flow | Clapperton et al. (1989) |
| 5200- 4400 | Block Lake, South Georgia | 54° 10'17.57" | 36° 43'32.82" | 12 | 7451 | 1.61 | Cooling | Glacial advances | Rosqvist and Schuber (2003) |
| before c. 6370-4692 | Maiviken lakes, South Georgia | 54°15′ | 36°30′ | 4 | с. 6370 | 0.63 | Climate optimum | Peak concentration of pollen and spores | Birnie (1990) |
| 6000- 4500 | Antarctic Peninsula | 63° 20.572′ | 55° 53.195′ | 29 | 9620 | 3.01 | Minor Cooling | Minor glacial readvance/more sea ice | Michalchuk et al. (2009) |
| 6000-5200 | South Georgia | 54° 30′–55° 00′ | 35° 30′–38 30′ | 9 | 9146 | 0.66 | Drier | Decrease in Warnstorfia fontinaliopsis | Van der Putten et al. (2009) |
| 7200- 7000 | Block Lake, South Georgia | 54° | 36° | 12 | 7451 | 1.61 | Cooling | Glacial advances | Rosqvist and Schuber (2003) |
| 7800- 6000 | Antarctic Peninsula | 63° 20.572′ | 55° 53.195′ | 29 | с. 9620 | 3.01 | Mid-Holocene Climatic Optimum | Glacial retreat | Michalchuk et al. (2009) |
| c. 8492–7295 | Stromness Bay, South Georgia | 54° | 36° 41'29.97" | 20 | 18572 | I.08 | Postglacial conditions | Increase in δ^{18} O records | Rosqvist et al. (1999) |
| с. 9380–8520 | Stromness Bay, South Georgia | 54° | 36° 41'29.97" | 20 | 18572 | I.08 | Postglacial conditions | Increase in δ^{18} O records | Rosqvist et al. (1999) |
| c. 10815–9995 | Cumberland-Stromness, South Georgia | 54° 30′–55° 00′ | 35° 30′–38° 30′ | 4 | c. 9998 | 0.4 | Deglaciation | Vegetation colonised ice-free zones/peat growth | Van der Putten et al. (2005) |

Table I. (Continued)

The number of published radiocarbon dates is shown for each record to indicate the density of age control. Non-calibrated radiocarbon dates have been calibrated using the calibration curve SHCal13 in CALIB 7.0.2 (Stuiver et al., 2014).

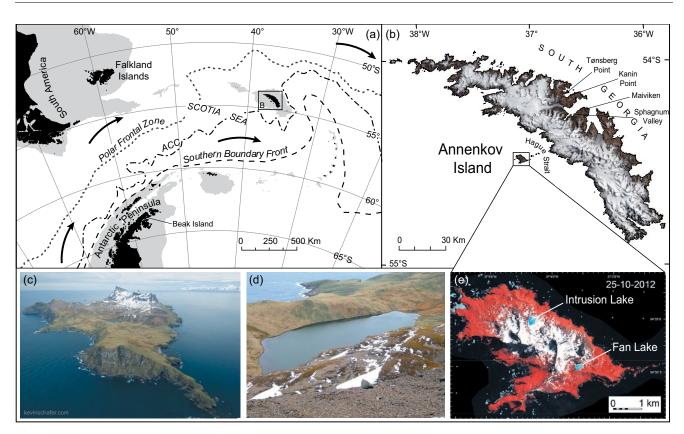


Figure 1. (a) Location map of the South Atlantic, showing the location of the main frontal systems of the Southern Ocean influencing the climate of South Georgia, South America and the Antarctic Peninsula. Landmasses are shaded black; areas less than 500 m deep are shaded grey; ACC: Antarctic Circumpolar Current. (b) LANDSAT image of South Georgia, showing the location of Annenkov Island and place names mentioned in text. (c and d) Fan Lake field site. (e) Satellite image of Annenkov Island. Red colour indicates vegetation cover on 25 October 2012 calculated from very high-resolution Quickbird2 satellite images with an on-the-ground resolution of 2.4 m per pixel. Live green vegetation in each pixel of the satellite image shows up as red in the image because the cell structure of chlorophyll containing cells absorbs red light while reflecting light in the infra-red part of the spectrum (Fretwell et al., 2011).

measurement time) at 2-mm intervals with standard calibration procedures (Gunn and Best, 1998), were used with the position of visible moss layers and key lithological markers, to stack the five core sections into a single stratigraphic sequence and to produce a composite depth sequence.

Lead-210 (²¹⁰Pb) and Ceasium-137 (¹³⁷Cs) dating analysis was undertaken on the upper 11 cm of the surface core using dried and homogenised samples packed into a 40-mm tube to 40 mm depth and left to stand for at least 21 days to allow ²²⁶Ra and ²¹⁴Pb to reach equilibrium. Samples were measured on an Ortec J-shape ultra-low background germanium well detector and remote preamplifier. Data analysis and dating model calculations were undertaken following standard procedures as defined in Appleby (2001). ¹³⁷Cs was at or below detection limits. Unsupported ²¹⁰Pb estimates were derived from both the constant initial concentration (CIC) and constant rate of supply (CRS) method (Table S1, available online) (Appleby and Oldfield, 1978). CRS ages were used in whole core age–depth modelling.

A total of 32 samples for radiocarbon dating were taken from the Fan Lake core. One basal sample from the KH4 peat core was also dated to provide an additional constraint on the timing of catchment deglaciation (Table 2). Calibration of terrestrial radiocarbon ages was carried out in OXCAL v.4.2 (Bronk Ramsey, 2009) using the SHCal13 Southern Hemisphere atmosphere dataset (Hogg et al., 2013). For post-bomb samples, absolute percentage of modern carbon (pMC) data were corrected according to ¹³C/¹²C isotopic ratios from measured pMC, where a 'modern' pMC value is defined as 100% (AD 1950), and the 'present day' pMC value is defined as 107.5% (AD 2010), and calibrated using the SHCal13 SH Zone 1-2 Bomb curve in CALIBomb (Hua et al., 2013; Reimer and Reimer, 2004). As all radiocarbon age errors are less than 50 years, calibrated ages are rounded to the nearest 5 years in Table 2.

A master age–depth model was constructed using a combination of all ²¹⁰Pb CRS ages and 32 radiocarbon ages in a Bayesian age–depth model undertaken in BACON v2.2 (Blaauw and Christen, 2011; Figure 2; Table 2). All ages quoted in text are weighted mean ages produced from the BACON age–depth model rounded to the nearest 10 years (Figure 2; Table 2).

Palynological analysis

Below 92 cm, the core was subsampled at 8-cm increments for pollen analysis. To capture higher resolution changes during the late Holocene, the uppermost ~92cm of the core was subsampled at 3- to 4-cm increments.

Approximately 1–2 g of sediment per sample were run through a 125-µm sieve after treatment with KOH and HCl. Standard laboratory techniques (Fægri and Iversen, 1989) including HF treatment and acetolysis were used to process the pollen samples. A Lycopodium tablet was added to calculate pollen concentrations (Stockmarr, 1971), and samples were mounted with silicone oil. Lycopodium was differentiated between the tablets (Lycopodium clavatum) and the species found on South Georgia (L. magellanicum) because of different reticula (cf. Wilce, 1972). Between 400 and 600 pollen grains were counted per sample, with the exception of the lower portion of the core (400-587 cm), where the pollen influx dropped significantly. Acaena and Poaceae dominate the entire pollen profile with total percentages between 90% and 98%. Barrow (1976, 1978, 1983), Markgraf and D'Antoni (1978), and the pollen reference collection held at Northumbria University was used for pollen and spore identification. Indeterminable

| 1 Subsc-20001 Rest Imp | D D | Laboratory No. | Core | Туре | Sample description | Lith. Unit | Modified depth (cm) | δ ¹³ C V _{PDB} ±0.1 (‰) | U % | Measured pMC | ¹⁴ C age (¹⁴ C yr BP) | ⁴ C yr | OX(ages | OXCAL SHCal ages (cal. yr BP) | al 13 95.4% | OXCAL SHCal13 95.4% 20 calibration ages (cal. yr BP) | ation | Rejection criteria (a–d) and Prob. |
|---|----------|-------------------|--------------------|-------|-------------------------------------|---------------|------------------------|--|-------------|---------------------|---|-------------------|---------------|----------------------------------|-------------|---|-----------------|---------------------------------------|
| UBEC.2011 FMS TM Synthe molationaly 1 4.13 2.95 4.13 5.71 <th></th> <th>min.</th> <th></th> <th></th> <th></th> <th>lean±lσ</th> <th></th> | | | | | | | | | | | | | min. | | | | lean±lσ | |
| UBC-C0913 FMSC EOM Organic mundialicity insert 7 4.35 -2.58 0.0 84.46.0.3 133 1.21 123 | _ | SUERC-20912 | FANSC | Σ | Syntrichia robusta moss | 8 | 0.5 | -29.5 | 4. | 90.74±0.42 | | | | | | | 680±35 | a, b |
| SUBRC-20916 FMNSC TM Spaced most stemal 7 475 -268 207 1286 -296 -916 </td <td>7</td> <td>SUERC-20913</td> <td>FANSC</td> <td>BOM</td> <td>Organic mud/silt/clay</td> <td>7</td> <td>4.25</td> <td>-25.8</td> <td>0.9</td> <td>84.46±0.37</td> <td></td> <td></td> <td>_</td> <td>_</td> <td>_</td> <td>-</td> <td>230±40</td> <td>a, b</td> | 7 | SUERC-20913 | FANSC | BOM | Organic mud/silt/clay | 7 | 4.25 | -25.8 | 0.9 | 84.46±0.37 | | | _ | _ | _ | - | 230±40 | a, b |
| International Interna International International< | m | SUERC-20916 | FANSC ^a | Σ | S. saxicola moss stems/ | 7 | 4.75 | | 20.7 | 128.68 ± 0.59 | | | | | | | -30±I.4 | 0.88 |
| SUBRC-12576 FAN2AI EM Laminared organic muldi 7 5 -2.6.9 1 9.175±0.4 692 4 5 55 655 655 655 610±30 SUBRC-12367 FAN2AI BCM Organic muldishlow 7 10 -2.6.3 -2.8 87.19±0-40 1101 ± 7 915 060 900 903±555 SUBRC-12368 FAN2AI TM Saxiofanoss stemul 7 10 -2.6.3 -2.6.3 455 914±0.39 1101 ± 7 915 0.02 203±15 SUBRC-12368 FAN2AI TM Saxiofanoss stemuls 7 44 -2.5.1 445 93.46.0.3 73 45 73 73 76 76 70 | | | | | leaves | | | | | | | | <u>ו</u> 1 | | | | -13±0.4 | 0.12ª |
| SUBRC-12901 FMNC BOM Organic muddislicity 7 10.25 -289 2.5 8.719±0.40 101 4 3.7 915 1060 960 370±45 BETA-336507 FANJA-1 EM Organic-rich selment 7 31 -26.3 -2.1 445 93.69±0.013 12.3 490 500 <td< td=""><td>4</td><td>SUERC-12578</td><td>FAN2A-I</td><td>BOM</td><td>Laminated organic mud/ silt/clay</td><td>٢</td><td>ß</td><td>-26.9</td><td>_</td><td>91.75±0.4</td><td></td><td></td><td></td><td></td><td></td><td></td><td>610±30</td><td>a, b</td></td<> | 4 | SUERC-12578 | FAN2A-I | BOM | Laminated organic mud/ silt/clay | ٢ | ß | -26.9 | _ | 91.75±0.4 | | | | | | | 610±30 | a, b |
| BETA-315602 ANJA-1 ROM Organic-rich sediment 7 31 -26.3 85.91±0.21 12.00 ± 00 1080 0085 50.015 SURC-12585 FANJA-1 TM Saxicid moss stemal 7 44 -26.1 44.5 93.69.40.41 53 ± 57 740 690 160 65 57.54.10 SURC-12586 FANJA-1 TM Saxicid moss stemals 7 54 -25.1 44.5 93.64.0.39 73 ± 57 740 690 1690 65 973.440 SURC-12586 FANJA-1 TM Saxicid moss stemals 7 53 -25.1 49.5 90.41.40.39 500 150 1700 1704.50 SURC-12586 FANJA-1 TM Saxicid moss stemals 7 53 -25.1 49.3 90.41.40.3 50 170 1704.50 1700 1704.50 1700 1704.51 170 1704.51 170 1704.51 170 170 | S | SUERC-20917 | FANSC | BOM | Organic mud/silt/clay | 7 | 10.25 | -28.9 | 2.5 | 87.19±0.40 | | | | _ | | | 970±45 | a, d |
| SUBRC-12365 FANJA-1 TM Sanciala moss stemaly 7 44 -2.6.1 41.5 9.3.6.9.6.4.0.3 55 490 550 520 520.15 SUBRC-12365 FANJA-1 TM sanciala moss stemaly 7 46 -2.5.8 45.9 9.6.6.0.39 733 47 70 57 740 690 500.15.0 SUBRC-12365 FANJA-1 TM Sanciala moss stemads 7 39 90.6.0.03 73 740 690 500.15.5 SUBRC-12367 FANJA-1 TM Sanciala moss stemads 7 39 90.6.4.0.39 73 740 690 600.430 SUBRC-12367 FANJA-1 TM Sanciala moss stemads 7 39 -2.517 90.3.4.0.24 90 700 700 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 706.4.0 | 9 | BETA-326502 | FAN2A-I | BOM | Organic-rich sediment | 7 | 31 | -26.3 | I | 85.91±0.32 | | | | | | | 385 ± 55 | a, d |
| SURC-1286 FMJA-1 TM Sacold most stems/ leaves 7 46 -258 459 964.03 73 5 740 690 <th<< td=""><td>7</td><td>SUERC-12585</td><td>FAN2A-I</td><td>Σ</td><td>S. saxicola moss stems/ leaves</td><td>7</td><td>44</td><td></td><td>44.5</td><td>93.69±0.4I</td><td></td><td></td><td></td><td></td><td></td><td></td><td>520±15</td><td></td></th<<> | 7 | SUERC-12585 | FAN2A-I | Σ | S. saxicola moss stems/ leaves | 7 | 44 | | 44.5 | 93.69±0.4I | | | | | | | 520±15 | |
| UBRC-12387 FANJA-1 TM Leaves Sourciod moss strands 7 59 -27.9 40.5 87.12 ± 0.3 10.8 3.5 74.5 70.5 775.5 745.5 70.5 775.5 | œ | SUERC-12586 | FAN2A-I | Σ | S. saxicola moss stems/ | 7 | 46 | | 45.9 | 90.6±0.39 | | | | | | | 690±30 | |
| SURC-1258 FAN2A-1 TM Sociolo moss strands 7 59 -2759 405 87/124038 108 4 35 920 1060 955 975440 SURC-12588 FAN2A-1 TM S.sociolo moss strands 7 53 -257 453 90.41±0.39 809 ± 35 650 705 700 703±25 SURC-12589 FAN2A-1 TM S.sociolo moss strands 7 7 33 90.3±0.24 819 ± 35 650 700 703±125 SURC-12590 FAN2A-1 EM S.sociolo moss strands 7 14 -203 -461 7929±035 1864 ± 35 1610 1765 1790±10 SURC-1259 FAN2A-2 EOM Buk organic-rich 7 114 -203 1461 7929±035 1864 ± 35 1615 1765 1765 1765 1765 1765 1765 1765 1765 1765 1765 1765 | | | | | leaves | | | | | | | | | | | | | |
| SUERC-1258 FAN2A-1 TM S. sociola moss strands 7 6.3 -2.6.1 39 9.0.41±0.39 803 ± 35 6.55 745 700 703±35 SUERC-12589 FAN2A-1 TM S. sociola moss strands 7 73 -2.5.7 45.3 90.41±0.3 809 ± 35 6.60 750 700 703±35 BETA-326504 FAN2A-1 BOM Luk organic-rich 7 14 -20.3 46.1 79.24±0.35 186.4 ± 36 6.60 750 700 705±50 SUERC-12579 FAN2A-2 BOM Lairometasus: unknown 7 114 -20.3 46.1 79.24±0.3 186.4 ± 36 1760 1760±50 SUERC-12579 FAN2A-2 BOM Buik organic-rich 7 144 -27.3 244±0.27 248 176 1760±50 1760±50 1760±50 1760±50 1760±50 1760±50 1760±50 176±10±10 176±10±10 176±10±10 | 6 | SUERC-12587 | FAN2A-I | Σ | S. saxicola moss strands | 7 | 59 | | 40.5 | 87.12±0.38 | | | | _ | | | 975 ± 40 | c, d |
| SUERC-1259 FAN2A-1 TM S. saxicla moss strands 7 73 -2.5.7 45.3 90.3 ± 0.24 819 ± 35 6.00 750 700 703 ± 2.6 BETA-326504 FAN2A-1 BOM Buk organic-rich 7 83 -2.5.3 - 7894±0.29 1900 ± 30 1715 1875 1790 1790±40 SUERC-12579 FAN2A-1 BOM Buk organic-rich 7 114 -20.3 46.1 7929±0.33 2118 ± 35 1615 1865 1765 1769 1790±40 SUERC-12579 FAN2A-2 BOM Laminated organic mult) 7 114 -20.3 46.1 7929±0.33 2118 ± 35 1615 1765±105 SUERC-12579 FAN2A-2 BOM Laminated organic mult) 7 144 -27.6 18 4.53 1615 1865 1755 2050 2055±105 BETA-326503 FAN2A-2 BOM Maineted organic-rich <t< td=""><td>0</td><td>SUERC-12588</td><td>FAN2A-I</td><td>Σ</td><td>S. saxicola moss strands</td><td>7</td><td>63</td><td></td><td>39</td><td>90.41±0.39</td><td></td><td></td><td></td><td></td><td></td><td></td><td>700±25</td><td></td></t<> | 0 | SUERC-12588 | FAN2A-I | Σ | S. saxicola moss strands | 7 | 63 | | 39 | 90.41±0.39 | | | | | | | 700±25 | |
| BFTA-325504 FAN2A-1 BOM Bulk organic-rich 7 83 -26.3 - 789±0.02 1 | = | SUERC-12589 | FAN2A-I | Σ | S. saxicola moss strands | 7 | 73 | | 45.3 | 90.3 ± 0.24 | | | | | | | 705±25 | |
| SUERC-1259 FAN2A-2 Bone Taronecusation 7 114 -20.3 46.1 79.29±0.35 186.4 ± 35 1615 186.5 1760 1760±50 SUERC-12579 FAN2A-2 BOM Laminated organic mud/ species small bird 7 114 -27.6 1.8 76.82±0.33 2118 ± 35 1930 2155 2050 2055±55 BETA-326503 FAN2A-2 BOM Laminated organic mud/ silclay 7 114 -27.6 1.8 76.82±0.33 2118 ± 35 1930 2155 2050 2055±55 BETA-326503 FAN2B-1 M Bulk organic-rich 7 149 -25.9 - 73.44±0.27 2480 ± 35 2705 2705 2550±100 SUERC-12593 FAN2B-1 M Polyrichum Juniperinum 7 191 -26.9 47.1 64.76±0.28 3490 ± 35 370 2555 2550±100 SUERC-12595 FAN1C-1 M | 12 | BETA-326504 | FAN2A-I | BOM | Bulk organic-rich | 7 | 83 | -26.3 | I | 78.94±0.29 | | | _ | - | _ | - | 790±40 | |
| SUERC-12579 FAN2A-2 BOM Lanicated organic multion 7 114 -27.6 1.8 7.6.82±0.33 2118 ± 35 1930 1030 <td>2</td> <td></td> <td></td> <td>0000</td> <td></td> <td>٢</td> <td>1</td> <td></td> <td>171</td> <td>70,0407,075</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>740450</td> <td></td> | 2 | | | 0000 | | ٢ | 1 | | 171 | 70,0407,075 | | | | | | | 740450 | |
| SUERC-12579 FAN2A-2 BOM Laminated organic mud/ 7 114 -27.6 1.8 7.682±0.33 2118 ± 35 1930 2155 2050 2055±55 BETA-326503 FAN2A-2 BOM Bulk organic-rich 7 149 -25.9 - 73.44±0.27 2480 ± 30 2355 2705 2515±105 BETA-326503 FAN2A-2 BOM Bulk organic-rich 7 149 -25.9 - 73.44±0.27 2480 ± 30 2355 2490 2515±105 SUERC-12593 FAN1C-1 M Polytrichum juniperinum 7 191 -26.9 45. 47.1 64.76±0.28 3490 ± 35 3710 3720±65 SUERC-12595 FAN1C-1 M Polytrichum moss 6 257.7 -24.6 47.1 64.76±0.28 3490 ± 35 370 370 370±65 SUERC-12595 FAN1C-1 TM S.robustoness stems/ 5 320.7 -31.1 54.5 5367±102 3807±65 3807 ± 35 3 | 2 | 30ENC-1 2370 | 7-070101 | DOILE | species small bird | - | <u>r</u> | | - 0 F | CC.N I (7.6 / | | | | | | _ | | |
| BETA-326503 FAN2A-2 BOM Bulk organic-rich 7 149 -25.9 - 73.44 \pm 0.27 2480 ± 30 2355 2705 2490 2515 ± 105 sediment sediment 7 191 -26.9 45 73.22 \pm 0.32 2360 2715 2355 2555 ± 100 SUERC-12593 FAN2B-1 M Polytrichum juniperinum 7 191 -26.9 45 73.22 \pm 0.32 2504 4 35 2360 2715 2555 ± 100 SUERC-12595 FANIC-1 M Polytrichum juniperinum 7 191 -26.4 47.1 64.76 \pm 0.28 3490 4 35 3505 ± 100 SUERC-12595 FANIC-1 TM P. Polytrichum moss 6 257.7 -24.6 47.1 64.76 \pm 0.28 3490 4 35 365 370 370 \pm 65 350 \pm 65 3505 \pm 100 SUERC-12595 FANIC-1 TM S. robustom stems 5 320.7 21 4 | <u>+</u> | SUERC-12579 | FAN2A-2 | BOM | Laminated organic mud/ silt/clay | ٢ | 114 | -27.6 | 8. I | 76.82±0.33 | | | | | | | 355 ± 55 | |
| SUERC-12593 FAN2B-1 M Polytrichum juniperinum 7 191 -26.9 45 73.22±0.32 2504 ± 35 2360 2715 2555 2550±100 noss moss | 15 | BETA-326503 | FAN2A-2 | BOM | Bulk organic-rich sediment | 7 | 149 | | I | 73.44±0.27 | | | | | | | 515±105 | |
| SUERC-12595 FANIC-I M P, juniperinum moss 6 257.7 -24.6 47.1 64.76±0.28 3490 ± 35 3605 3835 3710 3720±65 stems/leaves SUERC-12596 FANIC-I TM S. robusta moss stems/ 5 320.7 -31.1 54.5 63.67±0.28 36.7 ± 35 3705 4065 3890 385±65 SUERC-12598 FANIC-I TM S. robusta moss stems 4 334.2 -26.2 45.4 64.11±0.28 3571 ± 35 3705 3705 3800 ±60 SUERC-12599 FANIC-I TM S. robusta moss stem 4 335.5 -30.2 56.3 64.39±0.28 3571 ± 35 3705 3705 3800 ±60 SUERC-12599 FANIC-I TM S. robusta moss stem 4 335.5 -30.2 56.3 64.39±0.28 3571 ± 35 360 3705 3765 3765±55 | 16 | SUERC-12593 | FAN2B-I | Σ | Polytrichum juniþerinum moss | 7 | 161 | | 45 | 73.22±0.32 | | | | | | | 550 ± 1 00 | |
| SUERC-12596 FANIC-I TM S. robusta moss stems/ 5 320.7 -31.1 54.5 63.67 ± 0.28 36.7 ± 35 3725 4065 3890 Ieaves leaves 1 S. robusta moss stem 4 334.2 -26.2 45.4 64.11±0.28 3571 ± 35 3690 3795 SUERC-12599 FANIC-I TM S. robusta moss stem 4 335.5 -30.2 56.3 64.39±0.28 3536 ± 35 3640 3875 3765 | 17 | SUERC-12595 | FANIC-I | Σ | P. juniperinum moss stems/leaves | 6 | 257.7 | | 47.I | 64.76±0.28 | | | | | | | 720 ± 65 | q |
| SUERC-12598 FANIC-1 TM 5. robusta moss stem 4 334.2 -26.2 45.4 64.11±0.28 3571 ± 35 3690 3920 3795 SUERC-12599 FANIC-1 TM 5. robusta moss stem 4 335.5 -30.2 56.3 64.39±0.28 3536 ± 35 3640 3875 3765 | 8 | SUERC-12596 | FANIC-I | Σ | S. robusta moss stems/ leaves | ъ | 320.7 | | 54.5 | 63.67 ± 0.28 | | | | | | | 885 ± 65 | |
| SUERC-12599 FANIC-1 TM S. robusta moss stem 4 335.5 -30.2 56.3 64.39±0.28 3536 ± 35 3640 3875 3765 | 61 | SUERC-12598 | FANIC-I | Σ | S. robusta moss stem | 4 | 334.2 | | 45.4 | 64.11±0.28 | | | | | | | 800±60 | |
| | 20 | SUERC-12599 | FANIC-I | Σ | S. robusta moss stem | 4 | 335.5 | | 56.3 | 64.39 ± 0.28 | | | | | | | 765±55 | |

Table 2. Radiocarbon age summary data for Fan Lake sediment core and the basal age from KH4 peat bog core.

| 21 SUERC-12600 FANIC-1 TM S. robusta moss stems/ 4 341.5 22 SUERC-12594 FAN2B-2 M S. robusta moss stems/ 4 340 23 SUERC-12503 FANIC-2 TM S. robusta moss stems/ 4 360 24 SUERC-12603 FANIC-2 TM S. robusta moss stems/ 4 402.7 24 SUERC-12605 FANIC-2 TM S. robusta moss stems/ 4 417.5 25 SUERC-12606 FANIC-2 TM S. robusta moss 2 449.5 26 SUERC-12606 FANIC-2 TM S. robusta moss 2 449.5 27 SUERC-12607 FANIC-2 TM S. robusta moss 2 449.5 27 SUERC-12608 FANIC-2 M Vittia pachyloma moss 2 449.5 28 SUERC-12608 FANIC-2 M Vittia pachyloma moss 2 449.5 29 SUERC-12608 FANIC-2 M S. soxicola moss stems/ 2 466.9 20 SUERC-12608 FANIC-2 </th <th>Unit d</th> <th>depth (cm)</th> <th>0°С УРДВ I U.I (%)</th> <th>ر «</th> <th>pMC</th> <th>BP)</th> <th>ر برک</th> <th></th> <th>ages (cal. yr BP)</th> <th>yr BP)</th> <th>ages (cal. yr BP)</th> <th></th> <th>(a-d) and Prob.</th> | Unit d | depth (cm) | 0°С УРДВ I U.I (%) | ر « | pMC | BP) | ر برک | | ages (cal. yr BP) | yr BP) | ages (cal. yr BP) | | (a-d) and Prob. |
|---|--------|------------------|-----------------------|-------------|------------------------------|------|----------|----------|-------------------|--------------|-------------------|---------------|-----------------|
| SUERC-12600 FANIC-1 TM S. robusta moss stems/ 4 SUERC-12594 FAN2B-2 M S. robusta moss stems/ 4 SUERC-12593 FAN1C-2 TM S. robusta moss stems/ 4 SUERC-12603 FAN1C-2 TM S. robusta moss stems/ 4 SUERC-12604 FAN1C-2 TM S. robusta moss stems/ 4 SUERC-12605 FAN1C-2 TM S. robusta moss 2 SUERC-12606 FAN1C-2 TM S. robusta moss 2 SUERC-12607 FAN1C-2 TM S. robusta moss 2 SUERC-12608 FAN1C-2 TM S. robusta moss 2 SUERC-12608 FAN1C-2 M Vittia pachyloma moss 2 SUERC-12608 FAN1C-2 M Vittia pachyloma moss 2 SUERC-12808 FAN1C-2 M S. robusta moss 2 SUERC-12808 FAN1C-2 M S. robusta moss 2 SUERC-12808 FAN1C S S. coarieola moss stems/ 1 SUERC-1283 FAN1E BS Coarse light | | | | | | | | - | min. | тах. | Median | Mean±lσ | |
| SUERC-12594 FAN2B-2 M S. robusta moss stems/ 4 leaves Beaves leaves 4 SUERC-12603 FANIC-2 TM S. robusta moss stems/ 4 SUERC-12604 FANIC-2 TM S. robusta moss stems/ 4 SUERC-12605 FANIC-2 TM S. robusta moss 2 SUERC-12605 FANIC-2 M S. robusta moss 2 SUERC-12606 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M Vitti pachyloma moss 2 SUERC-12608 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M Vitti pachyloma moss 2 SUERC-12608 FANIC-2 M Vitti pachyloma moss 2 SUERC-12608 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M S. robusta moss 2 <t< td=""><td>4</td><td>341.5</td><td>-29.9</td><td>55.3</td><td>64.77±0.28</td><td>3489</td><td>+1</td><td>35 3</td><td>3605</td><td>3835</td><td>3710</td><td>3715±65</td><td></td></t<> | 4 | 341.5 | -29.9 | 55.3 | 64.77±0.28 | 3489 | +1 | 35 3 | 3605 | 3835 | 3710 | 3715±65 | |
| SUERC-12603 FANIC-2 TM S. robusta moss stems/ 4 Ieaves leaves leaves 4 SUERC-12605 FANIC-2 TM S. robusta moss 2 SUERC-12605 FANIC-2 TM S. robusta moss 2 SUERC-12605 FANIC-2 TM S. robusta moss 2 SUERC-12606 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M Vitti pachyloma moss 2 SUERC-12608 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M Vitti pachyloma moss 2 SUERC-12580 FANIE BS Bulk glacial sediment 1 SUERC-12583 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 BETA-271304 KH4-IC BOM Dark brown organic - | 4 | 360 | -26.4 | 41.6 | 61.03±0.27 | 3966 | +1 | 35 4 | 4235 | 4515 | 4355 | 4355±65 | c, d |
| SUERC-12604 FANIC-2 TM 5. <i>robusta</i> moss 3 SUERC-12605 FANIC-2 TM 5. <i>robusta</i> moss 2 SUERC-12606 FANIC-2 TM 5. <i>robusta</i> moss 2 SUERC-12608 FANIC-2 TM 5. <i>robusta</i> moss 2 SUERC-12608 FANIC-2 M <i>Vittia pachyloma</i> moss 2 SUERC-12609 FANIC-2 M 5. <i>saxicola</i> moss stems/ 2 SUERC-12580 FANIC-2 M 5. <i>saxicola</i> moss stems/ 2 BUIK glacial sediment 1 - <i>sity-clay</i> 1 SUERC-12583 FANIE BS Bulk glacial sediment 1 - <i>sity-clay</i> 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 BETA-271304 KH4-IC BOM Dark brown organic - | 4 | 402.7 | -30.1 | 44.6 | 64.34±0.28 | 3543 | +1 | 35 3 | 3640 | 3885 | 3770 | 3770±55 | |
| SUERC-12604 FANIC-2 IM 5. <i>robusta</i> moss 2 SUERC-12605 FANIC-2 TM 5. <i>robusta</i> moss 2 SUERC-12606 FANIC-2 TM 5. <i>robusta</i> moss 2 SUERC-12608 FANIC-2 M <i>Vittia pachyloma</i> moss 2 SUERC-12609 FANIC-2 M 5. <i>savicola</i> moss stems/ 2 SUERC-12580 FANIC-2 M 5. <i>savicola</i> moss stems/ 2 SUERC-12580 FANIC BS Bulk glacial sediment 1 - sity-clay SUERC-12583 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 BETA-271304 KH4-IC BOM Dark brown organic – | | | | - - - | | | | | LCO | | 0.01 | 12.1201 | - |
| SUERC-125606 FANIC-2 M S. robusta moss 2 SUERC-12607 FANIC-2 M S. robusta moss 2 SUERC-12608 FANIC-2 M Vittia pachyloma moss 2 SUERC-12609 FANIC-2 M S. saxicola moss stems/ 2 leaves SUERC-12580 FANIE BS Bulk glacial sediment 1 - silty-clay SUERC-12583 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 BETA-271304 KH4-IC BOM Dark brown organic – | 4 0 | 7 0 1 1 C. / I + | 196- | 46.6 | 62.37 ± 0.27 64 51 + 0.28 | 3521 | н + | ς κ γ | 2725 2635 | 423U 3855 | 4060 3755 | 3750+60 | D |
| SUERC-12607 FANIC-2 TM S. robusta moss 2 SUERC-12608 FANIC-2 M Vittia pachyloma moss 2 SUERC-12609 FANIC-2 M Vittia pachyloma moss 2 SUERC-12609 FANIC-2 M S. soxicola moss stems/ 2 SUERC-12609 FANIC-2 M S. soxicola moss stems/ 2 SUERC-12580 FANIE BS Bulk glacial sediment 1 SUERC-12583 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 BETA-271304 KH4-IC BOM Dark brown organic - | 2 | 159 | | 41.3 | 64.51±0.28 | 3521 | 1 +1 | | 3635 | 3855 | 3755 | 3750±60 | |
| SUERC-12608 FANIC-2 M Vittia pachyloma moss 2 SUERC-12609 FANIC-2 M Saxicola moss stems/ 2 SUERC-12609 FANIC-2 M Saxicola moss stems/ 2 RUL Bulk glacial sediment 1 - - SUERC-12583 FANIE BS Bulk glacial sediment 1 SUERC-12583 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 BETA-271304 KH4-IC BOM Dark brown organic - | 2 | 162 | | 39.2 | 62.85 ± 0.27 | 3731 | +1 | | 3900 | 4150 | 4030 | 4025 ± 65 | P |
| SUERC-12609 FANIC-2 M S. saxicola moss stems/ 2 leaves leaves leaves leaves 1 SUERC-12580 FANIE BS Bulk glacial sediment 1 SUERC-12583 FANIE BS coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 SUERC-12584 FANIE BS Coarse light grey sandy/ 1 BETA-271304 KH4-IC BOM Dark brown organic - | 2 | 166.9 | I | 57.8 | 64.49±0.28 | 3524 | +1 | 35 3 | 3635 | 3855 | 3755 | 3755±60 | |
| leaves SUERC-12580 FANIE BS Bulk glacial sediment I - silty-clay SUERC-12583 FANIE BS Coarse light grey sandy/ I gravelly clay SUERC-12584 FANIE BS Coarse light grey sandy/ I BETA-271304 KH4-IC BOM Dark brown organic – | 2 | 170.7 | -26.5 | 40.9 | 62.98±0.27 | 3714 | +I | 35 3 | 3885 | 4145 | 4005 | 4005 | |
| SUERC-12580 FANIE BS Bulk glacial sediment I - silty-clay SUERC-12583 FANIE BS Coarse light grey sandy/ I gravelly clay SUERC-12584 FANIE BS Coarse light grey sandy/ I gravelly clay BETA-271304 KH4-IC BOM Dark brown organic – | | | | | | | | | | | | 005 ± 65 | |
| SUERC-12583 FANIE BS Coarse light grey sandy/ 1 gravelly clay SUERC-12584 FANIE BS Coarse light grey sandy/ 1 gravelly clay BETA-271304 KH4-IC BOM Dark brown organic – | _ | 516 | -27.3 | 0.7 | 45.38±0.2 | 6346 | +1 | 35 7 | 7155 | 7320 | 7230 | 7230±50 | P |
| SUERC-12584 FANIE BS Coarse light grey sandy/ 1 gravelly clay BETA-271304 KH4-1C BOM Dark brown organic – | _ | 544 | -27.3 | 0.8 | 51.26±0.22 | 5368 | +1 | 35 5 | 5990 | 6270 | 6105 | 6105±70 | |
| BETA-271304 KH4-1C BOM Dark brown organic | _ | 584 | -27.9 | 0.5 | 42.08±0.19 | 6953 | +1 | 37 7 | 7665 | 7845 | 7740 | 7745 ± 50 | |
| mud (basal) | | I | -22.5 | I | 47.15±0.29 | 6040 | н | 50 6 | 6975 | 6675 | 6835 | 6840±70 | |

was pipeted off from the top of the FANSC core before subsampling in the field. The FAN2A-1 core was packed to minimise disturbance of the flocculent upper sediments during transit. Cores 2A, 2B, IC, IE were used to construct the composite downcore sequence shown in Figure 2.

All radiocarbon age data were included in age-depth modelling to test agreement with the following manually assigned rejection criteria. (1) age reversals where independent ²¹⁰Pb dating evidence suggests radiocarbon ages are 'too old' or reversed, (2) ages from near tops of cores which could have been water washed or disturbed during coring, (3) moss fragments or strands which are more likely to be reworked and (4) improbable age reversals where macrofossil ages occur higher stratigraphically than younger macrofossils and are beyond two-sigma calibrated age range of lowermost macrofossil, or represent bulk inorganic sediment that is more than likely reworked. ^aMost likely age by comparison with Pb-210 data.

Table 2. (Continued)

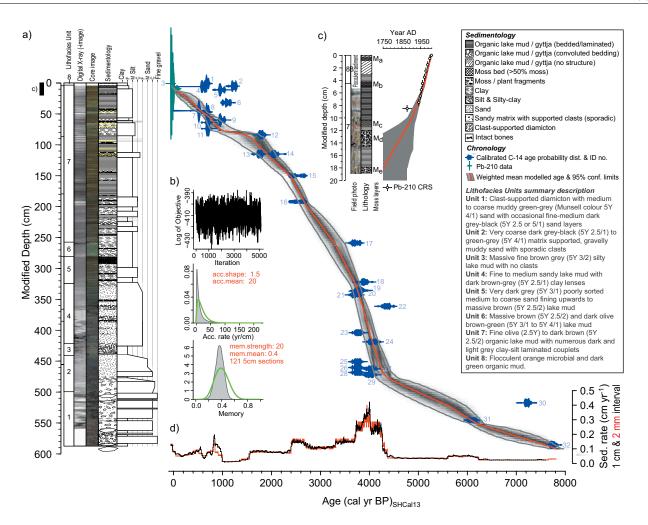


Figure 2. (a) Age–depth model for calculation of down-core radiocarbon ages (see text) compared with visual and x-radiographic core imagery (taken using an ITRAX XRF core scanner; running conditions 50kV, 50mA, 200 ms, 100 µm measurement interval), sedimentology and grain size summary data for the composite Fan Lake core. X-radiographic images are negatives; hence, areas with greater relative density are grey–white and darker areas are more organic or, in Units 1-2, represent areas of the core where clasts were removed before scanning. Age–depth modelling was undertaken in BACON v2.2 (Blaauw and Christen, 2011) using all radiocarbon ages and constant rate of supply (CRS) age model ²¹⁰Pb data (shown as white circles in c; see also Table S1, available online) using the SHCal13 calibration curve (Hogg et al., 2013) with 5-cm segments, 121 segments in total, for 27 million iterations and with every 615th iteration selected. Initial parameters were mean accumulation rate of 20 yr cm⁻¹, accumulation shape = 1.5, mem. strength = 20. When tested, mem. strength values of 10–100 gave essentially the same result, increasing confidence in the model parameters chosen and the reliability of the model. Calibrated age probability distributions with identification numbers are shown in blue. The median best-fit age plot (1-mm interval) in red is superimposed on individual BACON data runs shown as a grey shaded area. Darker grey areas indicate a greater density of individual age–depth model runs. The solid grey line represents the 95% confidence age limits for model runs. Colour versions of the core image can be viewed in the online version of this article; (b) Set up parameters for the BACON age–depth model runs. Colour versions of the core image can be viewed in the online version of this article; (b) Set up parameters for the BACON age–depth model; (c) Pb-210 ages (±1-sigma errors) from the CRS model which were used in the whole core age–depth model (red line and grey shading as described in a; (d) Sedimentation rates fo

and damaged pollen grains were used as an indicator for erosion and fluvial sediment transport, based upon these pollen grains becoming indeterminable because of damage by transportation and sediment removal.

The Tilia program was used to produce percentage and influx pollen diagrams (Grimm, 1990) with pollen assemblage zones delimited by stratigraphically constrained cluster analysis in CONISS (Grimm, 1987). The significance of the zones was assessed using a broken-stick model in the Rioja package (Juggins, 2012). The pollen sum that is used to calculate percentage pollen diagrams is the total of pollen, excluding fungal spores and pollen grains derived from long-distance transport.

Multivariate analysis of the pollen data

Principal components analyses (PCAs) of the Fan Lake pollen data were carried out in PRIMER 6 and PAST (Clark and Gorley, 2006; Hammer et al., 2001). Rare (consistently <1%) pollen taxa were excluded, and remaining pollen percentages were normalised in PRIMER 6 (Clark and Gorley, 2006) prior to

analysis. Confidence intervals (95%) were calculated for the total long-distance taxa (*Ephedra*, Fabaceae undiff., Myrtaceae, *Nassauvia*, *Nothofagus* and Podocarpaceae) using Psimpoll version 4.26 (Bennett, 2008). Through this method, changes in long-distance transport from the Fan Lake record are shown by deviation from the overlapping confidence limits indicated by a mean of 1.496 or greater.

Results

Sedimentology and chronology

The 587-cm Fan Lake core lithology was divided into eight units (Figure 2), from a basal unit (Unit 1) composed of a clast-supported sandy matrix to the uppermost units, which consist of laminated organic lake muds (Unit 7), capped by a living microbial mat and lake mud deposit (Unit 8; Figure 2). Between *c*. 587 and 499.5 cm, Unit 1 consists of a dark grey clast-supported diamicton to a dark greyish brown sandy matrix with clasts. Unit 1 grades into Unit 2 (499.5–442 cm), which is predominantly composed of

dark grey to a very dark greyish brown sandy matrix with clasts. Unit 3 (442–424 cm) is a fine organic lake mud, capped by Unit 4 (424–325 cm), an interbedded, and convoluted unit of sand and fine lake mud. Units 5 and 6 (325–257.5 cm) are geochemically similar, but fine upwards from a dark greyish brown silt-rich silty-clay into a light grey/pale yellow clay-rich silty-clay layer. Unit 7 (257.5–0 cm) comprises finely laminated organic mud, with occasional 2- to 3-cm thick sandy-silt layers. Organic carbon (measured as LOI) increases from 325 cm towards the top of the core, with maximum values present between 250 and 200 cm.

Reworking of older material appears to have affected some of the bulk ages from glacially derived sediments near the base of the core. Within Unit 7, reworking of 'older' moss layers is evident between c. 1500 and 1000 cal. yr BP. In the uppermost 50 cm of the core, the Bayesian age–depth model (Figure 2) favours the 210 Pb data as being a more likely scenario, given the small errors associated with these ages and the age reversals present in radiocarbon data in the top 11 cm of the core. A more detailed assessment of the Fan Lake sedimentology, geochemistry and chronology, and an assessment of the impact of meltwater events on the lake system, will be presented in forthcoming papers.

Palynology

Pollen grains are well preserved through the core with influx continually high between approximately 350 and 150 cm. The major taxa found throughout all the pollen zones are *Acaena* and Poaceae. Pollen of Asteraceae regularly occurs throughout the entire Fan Lake record.

Three major pollen zones (Fan1, Fan2 and Fan3a–e) were defined by the stratigraphically constrained cluster analysis. Fan1, Fan2 and Fan3d–e were statistically significant as assessed using the broken-stick analysis. Other zones defined by the stratigraphically constrained cluster analysis were not significant but retained to aid description and interpretation of the data (Figure 3).

Pollen zone Fan I: 587–340 cm; ~7700–3790 cal. yr BP

Poaceae and *Acaena* are the dominant taxa in this zone with relative percentages of 80% and 10–20%, respectively. A slight increase in *Colobanthus* (2–3%) is found at 384 and 429 cm. *Nothofagus* increases up to 3% at 500 and 515 cm. *Callitriche* and *Polypodium* are found in small quantities (<5%). *Nothofagus* slightly increases between 510 and 480 cm (Figure 3). The total pollen influx reaches 19,348 grains cm⁻² yr⁻¹ at 384 cm (Figure 4).

Pollen zone Fan2: 340–232 cm; ~3790–3050 cal. yr BP

A drop in Poaceae to 65–80% is recorded while *Acaena* increases to 10–35%. Pollen grains derived from long-distance transport (e.g. *Ephedra, Nothofagus*, Podocarpaceae) are found in small quantities (<1%). *Colobanthus* is consistently found at approximately 1% in this zone. *Callitriche* and *Polypodium* are the only other pollen taxa (1–2%) consistently present (Figure 3). Higher pollen influx (5771–13,553 grains cm⁻² yr⁻¹) is observed, along with higher *Acaena* (1493–3867 grains cm⁻² yr⁻¹) and *Callitriche* (11–65 grains cm⁻² yr⁻¹) (Figure 4).

Pollen zone Fan3: 232–0 cm; ~3050 cal. yr BP to present

The pollen zone is characterised by lower *Acaena* (5-30%) and higher Poaceae (70-90%) pollen percentages. *Callitriche*, indeterminable and long-distance transport pollen grains, occur with slightly higher percentages (0-3%) throughout this zone (Figure 3).

Pollen sub-zone Fan3a: 232—200 cm; ~3050—2750 cal. yr BP

Increasing *Acaena* percentages (10–30%) and decreasing Poaceae (70–90%) percentages occur in this zone. *Callitriche, Colobanthus* and indeterminable grains stay between 0% and 1% (Figure 3). Total pollen influx peaks to 13,922 grains cm⁻² yr⁻¹ in this zone with high *Acaena* (up to 4098 grains cm⁻² yr⁻¹) and *Callitriche* (26–56 grains cm⁻² yr⁻¹) (Figure 4).

Pollen sub-zone Fan3b: 200–154 cm; ~2750–2430 cal. yr BP

Poaceae percentages remain from 70% to 80% through this zone, and *Acaena* declines to 17–20%. At the top of this zone, a slight increase in *Nassauvia* (1–2%) is found. *Callitriche* and *Colobanthus* remain around 2% throughout the zone. Asteraceae (1%) and a number of indeterminable undifferentiated grains increase. At 164 cm, a large spike in the unknown sphere or ovoid is observed (Figure 3). Pollen influx begins to drop from 18,262 to 7910 grains $cm^{-2} yr^{-1}$ at the top of this zone. *Callitriche* increases to 85 grains $cm^{-2} yr^{-1}$ (Figure 4).

Pollen sub-zone Fan3c: 154–94 cm; ~2430–1670 cal. yr BP

Poaceae percentages are on average higher compared with zone Fan3d, while *Acaena* percentages decrease. *Colobanthus* and *Callitriche* make up 1% of the total pollen sum, while *Polypo-dium* makes up <1%. *Ranunculus* (1%) first appears in this zone, while an increase in unknown monolete spores (2%) is observed. At 140 cm, a large increase in zygospores occurs, but overall, pollen influx continues to decline. Long-distance transport from South America is represented by pollen of *Ephedra*, Fabaceae, *Nothofagus* and *Podocarpus*. A peak in *Nothofagus* and *Ephedra* influx of 75 grains cm⁻² yr⁻¹ is present at 131.5 cm (Figure 4). A high *Callitriche* influx of 3–75 grains cm⁻² yr⁻¹ is also observed.

Pollen sub-zone Fan3d: 94–33 cm; ~1670–430 cal. yr BP

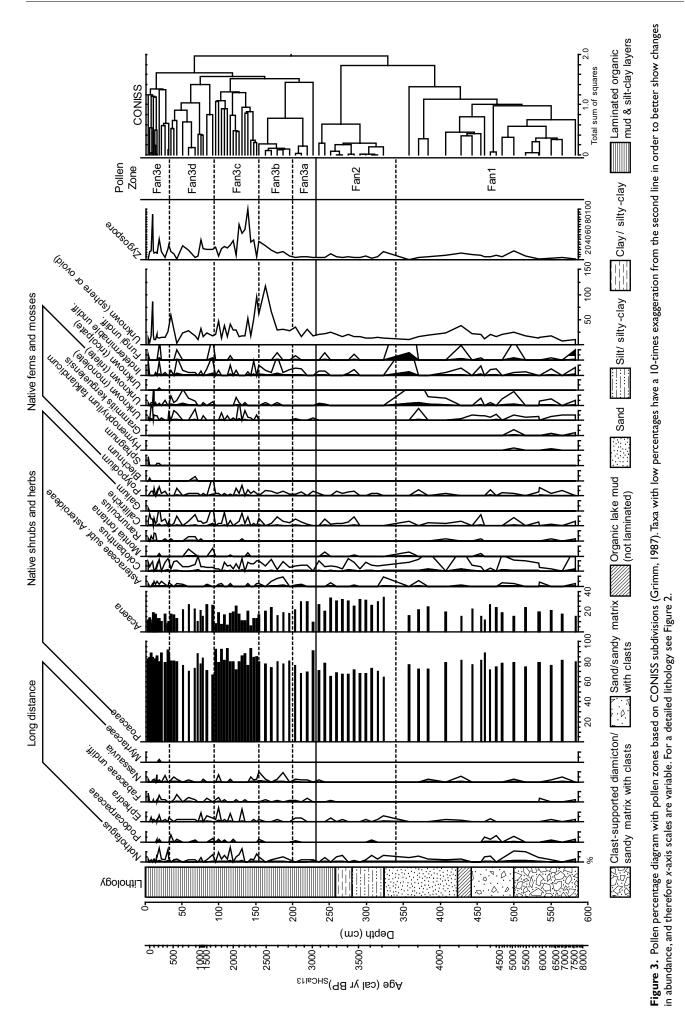
Poaceae decreases to 70–90% of the pollen sum, whereas *Acaena* increases from 10–30%. At 92 cm, a slight increase (3%) is found in *Callitriche, Polypodium*, indeterminable and fungal undifferentiated spores. *Callitriche* is consistently higher in this zone (1–2%) compared with the rest of the core. *Colobanthus* and *Polypodium* taxa are found in small percentages (<5%). *Blechnum* (<1%) is present for the first time in this zone. An increase in pollen influx up to 10,011 grains cm⁻² yr⁻¹ is observed at 60 cm.

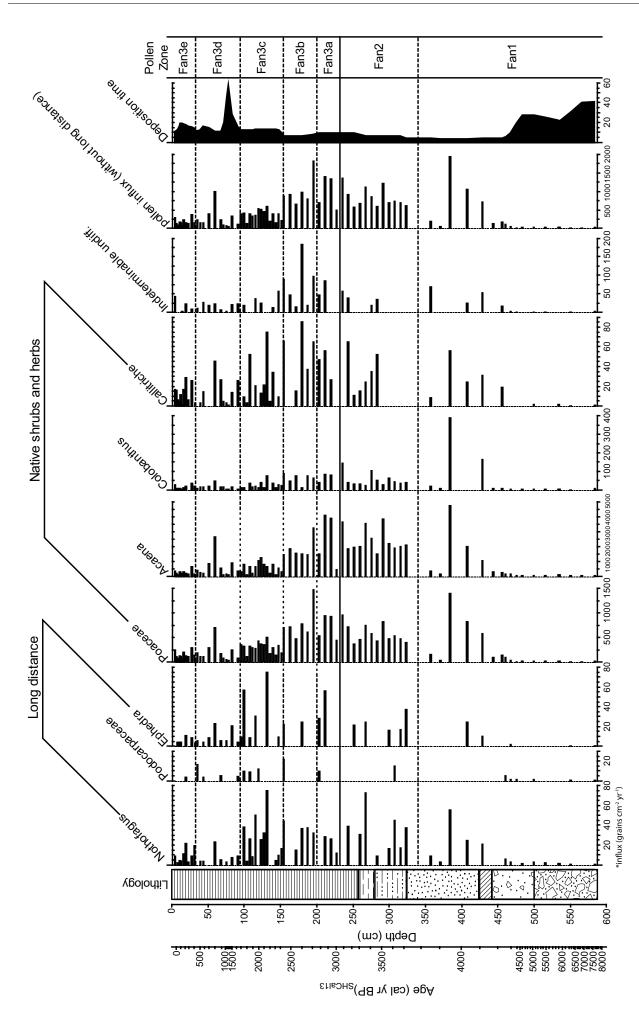
Pollen sub-zone Fan3e: 33–0 cm; ~430 cal. yr BP to recent

Poaceae percentages increase and *Acaena* percentage decreases in this zone. Monolete and trilete spores, indeterminable grains and fungal spores undifferentiated increase to 2%. *Callitriche* and *Colobanthus* remain around 1–2%. *Ranunculus* and *Polypodium* comprise <1% of the pollen sum. *Sphagnum* (<1%) is found in this zone for the first time. *Nothofagus* increases to >1% between 20 cm and 17 cm (Figure 3).

PCA

PCA was used to investigate statistical relationships within the pollen dataset, which could help to identify floral assemblages composed of co-occurring taxa with similar environmental preferences. The first and second PCA axes explain 34% of the total







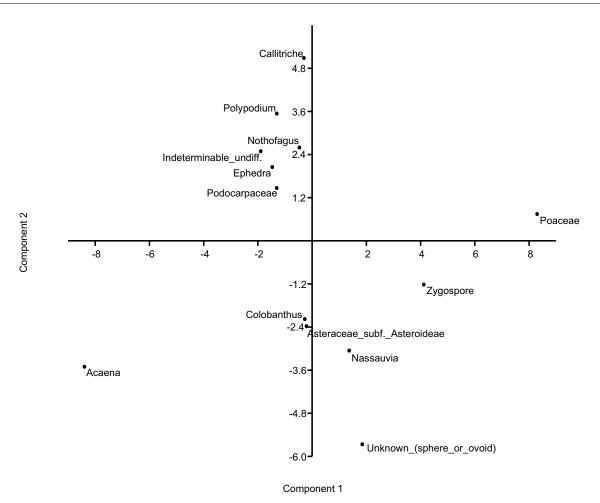


Figure 5. PCA of the Fan Lake pollen data showing the scores for the main pollen types (*Acaena*, Asteraceae, *Callitriche, Colobanthus, Ephedra*, indeterminable, *Nassauvia, Nothofagus*, Podocarpaceae, *Polypodium*, unknown cyst and Zygospore) on the first two principal components. Based on the ecological preferences of these taxa the first axis is interpreted as being related to temperature and the second to precipitation. PCA: principal components analysis.

variation. The first PCA axis explains 20% of the variance and clearly separates the dominant taxa Acaena and Poaceae, providing evidence that they are indicators of different environmental conditions (Figure 5). Although the climate of South Georgia has more than likely remained 'cold' and 'wet' throughout the Holocene, the first PCA axis likely represents a subtle shift in the climate gradient from relatively 'warmer' to 'cooler' conditions, with the second PCA axis indicating a shift along a 'drier' to 'wetter' gradient. The PCA separates the aquatic taxa Callitriche on the second PCA axis from the indeterminable (eroded) pollen grains, possibly indicative of 'wetter' environments, and longdistance taxa (Ephedra, Nothofagus, Podocarpaceae), which are likely related to increasing wind transport. These interpretations are further confirmed by negative sample score values of the first and second PCA axes (Figure 6), indicating slightly warmer and drier environments, and positive sample scores possibly representing cooler and wetter conditions.

Confidence intervals of long-distance taxa

The 95% confidence intervals were plotted for the long-distance taxa (Figure 6) to identify significant changes in longdistance pollen transport (a function of SHW strength) and to separate them from random variations inherent in the percentage counts (Maher, 1972). Confidence limits overlap throughout the majority of the Fan Lake record indicating few significant changes in long-distance pollen transport. However, pollen influx of long-distance taxa such as *Nothofagus* and *Ephedra* is highest in zone Fan3c (Figure 4), and the highest percentage peaks with non-overlapping confidence intervals occur at 99.5 cm (c. 1745 cal. yr BP) and 20 cm (c. 226 cal. yr BP) during periods of potentially colder and wetter climate.

Discussion

Challenges in the interpretation of sub-Antarctic pollen records

Despite the treeless and phanerogam-poor sub-Antarctic vegetation (Van der Putten et al., 2012b) and the low pollen diversity, the Fan Lake record shows interpretable changes in the major pollen taxa (i.e. *Acaena* and Poaceae) and the occurrence of environmental indicator taxa (e.g. *Callitriche*, indeterminable) that can be used, along with sedimentological evidence, to constrain mid- to late Holocene climate changes.

Changes in the relative abundance of pollen from *C. antarctica* can be used as indicators of wetter climatic conditions (Barrow, 1983). Long-distance pollen grain influx from the South American continent also provides evidence for increased pollen transport by the SHW (Schalke and Van Zinderen Bakker, 1971). The reliable identification of warmer and colder climate periods in the Fan Lake record remains challenging, and the alternating dominance of *Acaena* and Poaceae is difficult to interpret. Within the genus, *A. magellanica* is currently the dominant taxon on South Georgia and prefers lowland environments on moist peat or dry scree, while *A. tenera* inhabits high altitude areas (Walton, 1976). The two species could not be differentiated in the pollen records. However, it is most likely that at Fan Lake, the pollen

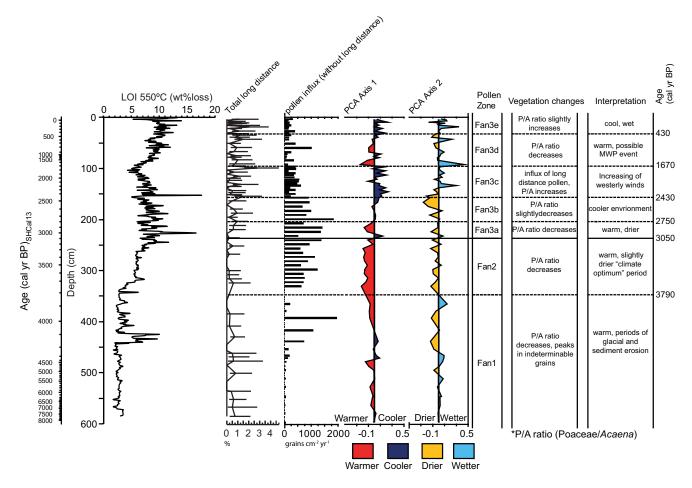


Figure 6. Overview figure showing loss-on-ignition (LOI 550°C) (wt% loss), total long-distance taxa: *Ephedra*, Fabaceae undiff., Myrtaceae, *Nassauvia*, *Nothofagus* and Podocarpaceae plotted with 95% confidence interval, total pollen influx showing the most relevant pollen taxa, sample scores for the first and second principal components of the PCA and general vegetation and climate changes observed in the Fan Lake core throughout the Holocene. PCA: principal components analysis.

was produced by the more abundant species *A. magellanica* that populated the peat around the lake during the Holocene.

The Fan Lake record shows a significant change at c. 2750 cal. vr BP from an Acaena- to a more Poaceae-dominated pollen assemblage. The timing of this change is broadly coincident with late Holocene cooling events identified by many other palaeoenvironmental studies from the sub-Antarctic islands (Table 1; e.g. Bentley et al., 2009; Unkel et al., 2010). Barrow and Lewis Smith (1983) also concluded from a pollen record from Sphagnum Valley, South Georgia, that high Acaena pollen percentages may indicate warmer or drier, and high Poaceae percentages cooler or cool-wetter climates. High Acaena percentages in Holocene pollen records have also been used on the Kerguelen Islands and Marion Island to identify warmer and drier climates, respectively (Scott and Hall, 1983; Young and Schofield, 1973). Despite these previous studies, we still treat the interpretation of Acaena and Poaceae with caution because in the South Indian Ocean sector, Van der Putten et al. (2010) could not demonstrate a relationship between the occurrence of A. magellanica and warmer climates. We therefore consider other proxies such as organic carbon and sedimentological evidence in addition to Acaena and Poaceae to support our palaeoclimate interpretations.

Local lake environments

During the mid-Holocene, Fan Lake was surrounded by sub-Antarctic vegetation mainly consisting of small shrubs and herbs, such as *Acaena*, Asteraceae, *Callitriche* and *Colobanthus*. The presence of Asteraceae is in agreement with findings from Lake Maiviken, South Georgia (Birnie, 1990). Osborne et al. (2009) considered this family as introduced, but we agree with Birnie (1990) in that Asteraceae might be native to South Georgia. The lithology between ~7700 and 3790 cal. yr BP is characterised by poorly sorted sediment such as clast-supported diamicton, a sandy matrix with clasts and sand along with the presence of indeterminable, eroded pollen grains which point to fast flowing waters, erosion and persistent glacial activity in the mid-upper catchment. Increasing temperatures at the beginning of the Holocene probably caused local glacier retreat and increased meltwater input into Fan Lake. Minor and fluctuating increases of Colobanthus between 7590 and 3950 cal. yr BP might indicate short drier periods with reduced precipitation or reduced periods of meltwater input, as this small sub-Antarctic shrub today inhabits drier environments (Convey et al., 1999).

A peak in indeterminable grains, fern and fungal spores at \sim 3860 cal. yr BP suggests a relatively short disturbance and soil erosion, which coincides with an influx of coarse sand and some of the highest sedimentation rates (0.2–0.4 cm/yr) in the core (Figure 2). The subsequent decrease in the percentage of indeterminable grains at \sim 3790 cal. yr BP could indicate a slow progression towards a less glaciated mid–upper catchment and may coincide with the onset of a late Holocene 'climate optimum'. A strong increase in pollen influx and organic carbon suggests higher catchment biomass within lake production between \sim 3230 and 2710 cal. yr BP. A warmer-than-present environment is inferred from an increase in *Acaena* around Fan Lake, probably predominantly residing in the lower altitude moist peat areas.

From c. 2750 cal. yr BP (Pollen sub-zone Fan3b), several changes in the Fan Lake pollen and sediment record indicate the progression towards a cooler and wetter environment in the late Holocene. A sharp drop in total pollen influx suggests decreasing plant biomass. The contemporaneous slight increase in sediment grain size in pollen zone Fan3b along with an increase in indeterminable and eroded pollen grains suggest higher sediment and water inputs possibly from seasonal melting of snowfields in the catchment. Coinciding with this drop in total pollen influx is an increase in Callitriche, suggesting a generally wetter environment from 2750 cal. yr BP onwards. The water-starwort Callitriche, which today colonises moist habitats on South Georgia (Convey et al., 1999), possibly grew in the lower catchment areas of the Holocene Fan Lake where water may have collected in the bogs. The regular flooding might have also caused a general decline in Acaena, which formerly occupied these habitats. Further evidence suggesting a changing lake environment includes the increase in percentages of unknown spheres or ovoids, and the zygospores, which peak at ~2495 and ~ 2260 cal. yr BP, respectively. The origin of both palynomorphs is unknown. However, it is likely that they have been produced within the lake (e.g. by algae), possibly indicating a rapid change in lake ecology.

Two colder/wetter periods can be identified between c. 2750– 1670 and c. 710 cal. yr BP to present if we assume that *Acaena*– Poaceae ratio can be applied to reconstruct temperature and/or precipitation changes (see discussion above). Although a slight trend towards an increase in organic content is observed from c. 710 cal. yr BP to present, a greater influx of *Callitriche* and a low total pollen influx suggest cooler and wetter conditions towards the present. The interpretation of warmer/drier environmental conditions between c. 1670–710 cal. yr BP is supported by higher *Acaena* pollen percentages and a contemporaneous increase in pollen influx and organic content (Figures 3 and 6).

Long-distance signal

Long-distance pollen grains from the South American continent are regularly present in the Fan Lake record. Changes in the abundance of the long-distance grains can aid in determining latitudinal shifts or changes in the intensity of the SHW (Bentley et al., 2009; Markgraf et al., 2003; Moreno et al., 2009). The long-distance signal is dominated by pollen from the southern beech Nothofagus, the conifer Podocarpus and the shrub Ephedra. Nothofagus and Podocarpus are today abundant forest trees in temperate and warm temperate climate zones of South America (Veblen et al., 1995, 1996) but also colonise as shrubs, together with Nassauvia and Ephedra, the sub-Antarctic vegetation of southern South America, for example, southern Patagonia and Tierra del Fuego (Björck et al., 2012; Markgraf et al., 2007; Montade et al., 2012). Because of the wide climate tolerance of these taxa, Holocene climate variability and a subsequent shift of vegetation zones in southern South America are unlikely to have a major impact on the total influx of long-distance pollen grains on South Georgia.

The Fan Lake record shows slight increases in long-distance pollen influx throughout the late Holocene cooling after 2430 cal. yr BP. Small peaks are present between approximately 2210 and 1670 cal. yr BP (Figure 4). The highest abundance of long-distance pollen occurs during the inferred cooler and wetter periods on Annenkov Island. *Ephedra* and *Nothofagus* comprise the majority of the long-distance signal, suggesting that the pollen grains have been transported from the southernmost, sub-Antarctic vegetation of South America (rather than from the *Podocarpus*- and *Nothofagus*-rich cool temperate rainforests).

Climate change

The Fan Lake record provides evidence of warmer conditions in the mid- to late Holocene, which culminates in a period of comparatively higher biomass production between 3790 and 2750 cal. yr BP. With a subsequent gradual change towards slightly cooler and wetter environments after c. 2750 cal. yr BP, the pollen-based climate reconstruction from the Fan Lake record broadly agrees with other South Georgia, Antarctic Peninsula, maritime- and sub-Antarctic records (see Table 1; e.g. Bentley et al., 2009; Unkel et al., 2010).

Previous records from the north coast of South Georgia suggest several cooling and drying events during the mid- to late Holocene, but no strong signal of these changes are present on Annenkov Island. For example, from grey scale density, weight-loss-on-ignition and grain size analyses, Rosqvist and Schuber (2003) reconstructed cooler periods between 7800 and 7400 cal. yr BP, 7200 and 7000 cal. yr BP, 5200 and 4400 cal. yr BP and 2400 and 1600 cal. yr BP. The slight increases in the pollen percentages of the cushion plant *Colobanthus* might indicate short dry events around 7590–3950 cal. yr BP, which roughly coincide with dry periods previously identified by Van der Putten et al. (2004, 2009) through macrofossil analyses at Tønsberg Point (8800–8200 cal. yr BP) and Kanin Point (4400–3400 cal. yr BP) on the north coast of South Georgia.

Rosqvist and Schuber (2003) indicate cooling on South Georgia between 2400 and 1600 cal. yr BP based on glacial advances, although the accuracy of these dates have been questioned (Moreton et al., 2004). The interpretation of a late Holocene climate change at Fan Lake is also supported by lake sediment studies on South Georgia (Lake Maiviken), which suggest the onset of wetter conditions at c. 2750 cal. yr BP (Birnie, 1990). However, recent macrofossil studies from Kanin Point and Tønsberg are more contradictory (Van der Putten et al., 2009). While Tønsberg assemblages suggest a change from wet to extremely wet conditions at 2200 cal. yr BP, Kanin Point seems to have developed from a wet to a dry peat bank environment (Van der Putten et al., 2009). Since the Tønsberg sequence was located in a depression, it was more likely affected by higher precipitation. In contrast, the Kanin Point vegetation responded to a decrease in temperature allowing a Warnstorfia bog to be introduced by Chorisodontium aciphyllum and Polytrichum cf. strictum, indicating wetter environments (Van der Putten et al., 2009). Various lake sediments, pollen and macrofossil proxy data also suggest a cooler period on South Georgia that started at c. 2200 cal. yr BP (Clapperton et al., 1989; Van der Putten et al., 2009, 2012a).

Various sub- and maritime-Antarctic islands and Antarctic Peninsula records closely resemble Fan Lake in showing a late Holocene 'climate optimum', late Holocene cooling and a possible warming overlapping with the 'Medieval Climate Anomaly' ('MCA') (Table 1). At Fan Lake, the warm and stable lake conditions inferred between 3790 and 2750 cal. yr BP overlap with the late Holocene 'climate optimum' from c. 3500 to 1150 cal. yr BP on maritime-Antarctic Signy Island, at 850 km distance (Hodgson and Convey, 2005; Jones et al., 2000). Similar to Fan Lake, multiple lake sediment records from Signy Island suggest a late Holocene 'climate optimum' through LOI and pollen concentration data. This interpretation also coincides with the late Holocene Hypsithermal in the Beak 1 Lake record between 3169 and 2120 cal. yr BP from Beak Island in the Prince Gustav Channel, NE Antarctic Peninsula, as indicated by an increase in sediment accumulation and a peak in organic carbon, along with increased primary production (Sterken et al., 2012). Other examples of a climate optimum starting at around the mid-late Holocene boundary (4200 yr BP), or in the first part of the late Holocene, are described by Bentley et al. (2009) and Hodgson et al. (2009).

From 2710 cal. yr BP onwards, the decline in pollen influx in the Fan Lake record indicates a change towards a generally cooler and wetter climate, which persisted until today. With several interbedded warm and cold periods, the climate on Annenkov Island has been very variable throughout the last 2700 years. The majority of research in the sub-Antarctic puts the start of cooling in the late Holocene at *c*. 2500 cal. yr BP, but uncertainty still remains as to the exact timing (Bentley et al., 2009; see also Table 1).

At the end of the late Holocene, increased *Acaena* pollen percentages, high pollen influx and increased organic flux suggest the existence of a relatively warmer period at Fan Lake which was centred around 1670–710 cal. yr BP. The signature of this extended period of late Holocene warming has been detected at several Antarctic Peninsula (e.g. Bentley et al., 2009) and South American locations (e.g. Kastner et al., 2010; Mauquoy et al., 2004; Neukom et al., 2010) and overlaps with the timing of both the earlier Northern and later Southern Hemisphere expression of a 'MCA' (Neukom et al., 2014). The effects of the 'MCA' have been reported in both hemispheres, but the timing of these events varies widely (Bird et al., 2011; Ledru et al., 2013; Neukom et al., 2010). This led Neukom et al. (2014) to conclude that there is no empirical support for a globally coherent warm phase during the pre-industrial (AD 1000–1850) era.

SHW and climate changes

The influx of non-native long-distance transported pollen grains in fossil records from the sub-Antarctic islands is primarily controlled by wind strength and the latitudinal position of the SHW and is therefore indicative of changes in atmospheric circulation patterns (e.g. Van der Putten et al., 2012b). Climate models and observational data suggest that during cold periods, the SHW shifts towards the equator, whereas during warmer phases (such as today), the movement is polewards (Bentley et al., 2009; Lamy et al., 2001; Toggweiler et al., 2006; Varma et al., 2012). However, how latitudinal shifts of wind belts affected precipitation and wind strength in the sub-Antarctic region (at $c. 50^{\circ}$ S) and whether these wind belts contracted or expanded throughout the Holocene remains debated (Hodgson and Sime, 2010).

Multi-proxy studies on lake sediment cores in SW Patagonia and Chile show increased wind strength along with increases in precipitation during cooler periods in the late Holocene, such as the 'Little Ice Age' ('LIA'), 400–150 cal. yr BP (Bentley et al., 2009; Moy et al., 2008). Increased wind strength and precipitation associated with the SHW could have caused the expansion of *Nothofagus*-dominated woodlands in SW Patagonia and Tierra del Fuego in the late Holocene (Moreno et al., 2009; Moy et al., 2011). However, during the mid-Holocene, Lamy et al. (2001) suggested poleward westerly wind shifts occurred in phase with warmer temperatures, changing to an anti-phased relationship in the late Holocene. As a result of this anti-phasing between the core and the northern margin, the core SHW belt became stronger during the early Holocene and weaker during the late Holocene (Lamy et al., 2010).

In contrast, the pollen record from Fan Lake situated at $c.54^{\circ}$ S provides evidence of a strengthening of the SHW during the late Holocene. Although the overall number of grains per sample and the percentage variations are relatively low, the Fan Lake record suggests that highest transport by westerly winds occurred during late Holocene cold periods at 2210–1670 cal. yr BP and after 710 cal. yr BP.

Conclusion

Most existing palaeoenvironmental records from the Antarctic and sub-Antarctic islands broadly identify a trend from a warm period in the beginning of the late Holocene followed by a prolonged cooling phase, but several uncertainties remain in defining the timing and underlying drivers for these past changes.

The well-dated, 5.8 m-long sediment record from Fan Lake on Annenkov Island, near South Georgia, was used to reconstruct the Holocene vegetation and climate history of the South Atlantic sector of the Southern Ocean at a high temporal resolution. We identified three key phases in the pollen record from Fan Lake: (1) a warm late Holocene 'climate optimum' from *c*. 3790 to 2750 cal. yr BP, with stable, but low, in-lake productivity and high biomass production; (2) a step-change after 2750 cal. yr BP toward cooler and wetter environmental conditions, as indicated by a decline in *Acaena* pollen, a rise of the aquatic plant *Callitriche* and an increase in indeterminable, eroded pollen grains; (3) a return to slightly warmer conditions between 1670 and 710 cal. yr BP, overlapping with some warm events associated with a 'MCA' in the Northern and Southern Hemispheres.

The presence of long-distance transport pollen grains in the Fan Lake record was used to reconstruct changes in atmospheric circulation patterns. Although the overall number of long-distance transport pollen grains in the Fan Lake record was comparatively low, the pollen record suggests a link between cold periods and stronger SHW in the late Holocene at this location.

Further studies are needed to better understand the history of the SHW and their impact on Holocene climate change. We suggest that future research should focus on coring sites on the west coasts of sub-Antarctic islands, close enough to the continental landmasses to show a distinct signal of long-distance pollen grain transport.

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