Volcanic record of continental thinning in Baffin Bay margins: Insights from Svartenhuk Halvø Peninsula basalts, West Greenland

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1 **Volcanic record of continental thinning in Baffin Bay** 2 **margins: insights from Svartenhuk Halvø Peninsula basalts,** 3 **West Greenland**

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28 **Abstract**

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30 We report major, trace elements and Pb, Hf, Nd and Sr isotopes in 61-54 Ma old basalts
31 from Svartenhuk Halvø Peninsula (Greenland). This area corresponds to the
32 northernmost exposure of the West Greenland volcanic province, of which the
33 emplacement marks the continental thinning and breakup process leading to the opening

of Baffin Bay during the Eocene. Its wedge structure displays typical characteristics of inner seaward-dipping-reflectors (SDR) with an exposed volcanic sequence thicker than 7 km. Our results cover the entire volcanic sequence starting with an earliest V1 unit reflecting rather low degree partial melts (transitional basalts), which are geochemically imprinted by continental crust contamination. This unit is followed by much thicker and mainly tholeiitic V2 and V3 basaltic lava piles, and ends with a rather thin V4 unit, consisting of alkali basalts and associated trachytes. Pressures and temperatures of melt extraction were estimated based on major element concentrations and rare earth elements patterns. Our results suggest that melts were extracted from the garnet-spinel transition zone at greater depths for the initial V1 and the final V4 lavas (alkali to transitional basaltic compositions) than for the main V2 and V3 mostly tholeiitic lava piles (at 2 ± 0.5 GPa and $1350 \pm 100^\circ\text{C}$). Isotope signatures suggest that mantle sources of the melts were controlled by the mixing of ambient upper mantle and Icelandic plume-type materials. The proportion of ambient upper mantle involved in the lava source appears to increase together with melting rates and the upward propagation of melting zone during V3 emplacement, suggesting that melting in this area progressed as follows: 1. Initiation of partial melting (low degrees) of deep-seated Icelandic plume-type material (V1 and V2); 2. Upward propagation of the melting area within the shallower mixed upper mantle (V3); 3. Progressive decrease of “ambient upper mantle” involvement, deepening of magma extraction and decrease of partial melting associated with the emplacement of V4 lavas. We show that these changes are consistent with the tectonic stages of continental thinning that preceded the Eocene breakup in Baffin Bay.

1. Introduction

Volcanic passive margins (VPM) characterize continental breakup associated with the emplacement of large igneous provinces (White and McKenzie, 1989; Coffin and Eldholm, 1994). In detail, initial mantle melting is thought to start prior to significant lithosphere extension (Courtillet et al., 1999), leading to the eruption of voluminous flood basalts. Then, focusing of mantle melting combined with lithosphere extension and consecutive syn-magmatic break-up leads to the formation of volcanic passive margins (Geoffroy et al., 2015). The corresponding volcanic sequences, although emplaced subaerially, are usually found as immersed seaward-dipping reflectors (SDR) sequences beneath post-rift sediments, a common occurrence throughout the North Atlantic Igneous Province (NAIP; Saunders et al., 1997). Such sequences, linked to the

opening of the Baffin Bay oceanic basin, are exceptionally well-exposed along the West Greenland coast, where they are of Paleocene-Eocene age and up to 5-7 km thick (Larsen and Pulvertaft, 2000; Larsen and Pedersen, 2009; Larsen et al., 2015; Chauvet et al., accepted). Petrologic and geochemical studies of these volcanic successions as well as those of similar sequences exposed in the East Greenland margin raise several important (and still debated) petrogenetic and tectono-magmatic questions, e.g. (1) do the considerable volumes of erupted picrites indicate abnormally high temperatures of partial melting of plume-related materials (Francis, 1985; Larsen and Pedersen, 2000, 2009; Hole and Millett, 2016); (2) does the heterogeneity of the mantle sources of Paleocene-Eocene mafic magmas foreshadow that of the present-day Icelandic hotspot (Holm et al., 1993; Peate and Stecher, 2003; Barker et al., 2006); and finally (3) does the temporal evolution of melting rates and depths record the successive stages of continental extension and rupture in the East (Larsen and Saunders, 1998; Abdelmalak et al., 2012, 2018) and West Greenland (Tegner et al., 1998; Barker et al., 2006) margins as well as in other parts of the NAIP (Kerr, 1994; Geoffroy, 2005; Meyer et al., 2007; Saunders et al., 2007)?

Svartenhuk Halvø Peninsula represents the northernmost exposure of the West Greenland Volcanic Province. The Svartenhuk wedge typically displays the structural and stratigraphic characteristics of inner-SDR wedges observed on deep seismic reflection profiles from VPM worldwide (e.g. Planke et al. 2000; Stica et al., 2014; Geoffroy et al., 2015). The thickness of the exposed volcanic sequence may reach up to 7 km and at least one third of it is made of picrites (Larsen and Pulvertaft, 2000). The aims of the present study are (1) to analyze magmatic processes such as mantle source heterogeneity, depths and rates of partial melting, and crustal contamination that occurred during the c. 10 m.y. of its emplacement (i.e. from 62.5-61 Ma to 54-53 Ma; Larsen et al., 2015; Chauvet et al., accepted), and (2) to discuss their temporal evolution during the successive stages of the continental stretching and thinning margin system related to the opening of the Baffin Bay oceanic basin.

2. Geological setting

2.1. The Labrador-Baffin oceanic domain and West Greenland Volcanic Province

The Labrador-Baffin axis is part of a wider Phanerozoic divergent system including the NE-Atlantic area between Eurasia and Greenland (Fig. 1). A continental and diachronous rift system developed between Greenland and Canada during the Cretaceous. This a-magmatic extension led to continental breakup with mantle exhumation in the southern Labrador Sea. A now extinct oceanic spreading axis in the Labrador Sea led to the drifting apart of Greenland from North America since the Late Cretaceous (Roest and Srivastava, 1989) or Paleocene (Chalmers and Laursen, 1995) and ceased to operate during the Oligocene (~ magnetochron C13, Roest and Srivastava, 1989). The SW-NE-trending left-lateral Ungava transform-fault system (UFS) connected the Labrador Sea to Baffin Bay farther North (Fig. 1).

In the northern Labrador Sea, the oldest undisputed oceanic magnetic anomaly has a mid-Paleocene age (Chalmers and Laursen, 1995), i.e. C27n (62.2 – 62.5 Ma, according to Ogg, 2012). During the Paleocene, the spreading directions were oriented ~N060E, i.e. sub-orthogonal to earlier rifted structures on the conjugate margins. Both the oceanic fracture zones and magnetic anomalies in the Labrador Sea indicate that a major shift in kinematics occurred during the Early Eocene, between C25 and C24, when spreading directions rotated from N060E to N010E (Roest and Srivastava, 1989; Geoffroy et al., 2001; Fig. 1).

In Baffin Bay, the age of the earliest accretion is still debated since no obvious linear magnetic anomalies can be identified on its floor, although there is a consensus for the cessation of spreading during the Oligocene (Roest and Srivastava, 1989; Oakey and Chalmers, 2012). Two major N-S-trending fracture zones in the central part of the Baffin Bay (Fig. 1) suggest that oceanic spreading may have been coeval with the ~N-S reorientation of the kinematic trend in the Labrador Sea during Eocene (Chalmers and Pulvertaft, 2001; Geoffroy et al., 2001; Suckro et al., 2012; Funck et al., 2012). Other authors (Oakey and Chalmers, 2012; Suckro et al., 2012; Hosseinpour et al., 2013) consider that earlier oceanic-type crust of mid-Paleocene age could exist within Baffin Bay. However, none of these studies took into account the age of continental stretching and thinning recorded along the Baffin Bay continental passive margins.

The conjugate passive margins in Baffin Bay (Fig. 1) are clearly of volcanic type in the South of the basin (Geoffroy et al., 2001; Skaarup et al., 2006; Block et al., 2012). The inner part of the SE-Baffin volcanic passive margin (VPM) is largely exposed from latitudes 69°N to 72°N in the Disko-Svartenhuk area (Fig. 1). An inner-SDR (Planke et al., 2000) developed orthogonally to the regional stretching direction during the

Paleogene (Geoffroy et al., 2001; Abdelmalak et al., 2012) and is fully exposed at Svartenhuk where its total thickness reaches up to 7 km. Recently published Ar-Ar ages (Larsen et al., 2015; Chauvet et al., accepted) that range from 62.5-61 Ma to 54-53 Ma for Svartenhuk basalts provide new constraints for the tectono-magmatic evolution of the Baffin Bay margin system.

2.2. The Svartenhuk Halvø Peninsula

The tectonics of the Svartenhuk SDR has already been investigated (Larsen and Pulvertaft, 2000; Geoffroy et al., 2001; Abdelmalak et al., 2012). Until recently, the age of the regional strain remained a matter of debate due to the lack of age determinations on lavas in this area (see discussion in Larsen and Pulvertaft, 2000). However, new sets of $^{40}\text{Ar}/^{39}\text{Ar}$ ages recently published by Larsen et al. (2015) and Chauvet et al. (accepted) provide new constraints on the timing of melting processes at the scale of the NAIP. In Svartenhuk Peninsula, the thickness of the exposed volcanic sequence may reach up to 7 km (Larsen and Pulvertaft, 2000). Its dense fault network results from a seaward-migrating strain (Geoffroy et al., 2001). The Arfertuarssuk Fault represents the exposed master fault bounding and controlling the development of the Svartenhuk wedge (Fig. 2).

In the present study and in the companion paper by Chauvet et al. (accepted) dealing with the same sample set, the volcanic formations making up the SDR wedge in Svartenhuk Halvø have been subdivided into four successive sequences (labeled V1 to V4 upwards; Fig. 2) (Fig. 2). The boundaries between these sequences are consistent with the geological and petrographic distinctions of Larsen (1983) and Larsen and Grocott (1992).

Coeval volcanics crop out on Disko Island, Nuussuaq Peninsula and Ubekendt Ejland Island, where they form a plurikilometric pile of ultramafic to mafic Paleogene lavas crosscut by sheeted intrusions (e.g. Pedersen et al., 2002). The stratigraphy and age of the W-Greenland basalts were defined in these areas as follows (Storey et al., 1998 and references therein). The Vaigat Fm. consists of a lower sequence dominated by hyaloclastites and pillow lavas overlain by picrites and Mg-rich subaerial basalts. The earliest subaqueous volcanics unconformably overlie the remnants of an Early Cretaceous to Paleogene paleobasin (Fig. 2). The overlying Maligât Fm. is dominated by feldspar-phyric and aphyric basalts associated with a sharp change in lava chemistry (Larsen and Pedersen, 2009). Larsen and Pulvertaft (2000) suggested that the lava

stratigraphy established in the Disko-Nuussuaq area could be applied to the Svartenhuk area. As such, the V1 and V2 formations correlate with the southern Vaigat Formation, while the “Svartenhuk Fm.” (V3 in Fig. 2) would be equivalent to the Maligât Fm. (Fig. 3). In Svartenhuk Halvø, a 1-2 km thick succession of transitional to alkaline basalts (V4 unit) lies above the V3 unit. A stratiform sill-like trachytic body marks the boundary between the V3 and V4 units. Larsen and Grocott (1992) considered these basalts and trachyte as the uppermost part of the Svartenhuk Fm. According to Larsen et al. (2015), this succession belongs to a newly introduced Naqerloq Fm. (Fig. 2), dominated by transitional light rare Earth element (LREE) enriched basalts.

From observations on the Disko and Nuussuaq areas, two main volcanic pulses have been identified and further correlated by radiometric dating and paleomagnetic data (Storey et al., 1998; Riisager and Abrahamsen, 1999; Riisager et al., 2003). The earliest pulse was mid-Paleocene in age and spans chrons 27n and 26r (62.5-59.2 Ma after Ogg, 2012). The second pulse occurred during the early Eocene at C24r (54-57 Ma), i.e., after a hiatus of 2 m.y. or more. An Early Eocene age was obtained from the basaltic succession of the Kanísut Mb. (Storey et al., 1998), that crops out exclusively West of the Itilli fault, and which is seaward tilted at about 15° (Fig. 2, Riisager et al., 2003). These basalts lie unconformably above the Vaigat-Maligat basalts, which are in turn deeply down-faulted westward (Larsen et al., 2015). It is noteworthy that early Eocene ages were also obtained for dykes crosscutting the Paleocene-related series in Disko and Nuussuaq (Storey et al., 1998) and also in the Svartenhuk Peninsula (Geoffroy et al., 2001). Larsen et al. (2015) proposed that the Paleocene magmatic pulse in West Greenland spans not only C27n-C26r but also up to chron C25r (59-57.7 Ma), based on four $^{40}\text{Ar}/^{39}\text{Ar}$ ages ranging from 60.3 ± 1.4 Ma to 58.0 ± 0.6 Ma on Svartenhuk Fm. basalts from the northern part of the Svartenhuk peninsula. This implies that the Svartenhuk Fm. was mostly contemporaneous with the Maligât Fm. but that its upper part was younger. These authors also outlined that the transitional to alkaline basaltic successions exposed along the Arfertuarssuk Fault in Svartenhuk Halvø (V4 unit) and on the western parts of the Ubekendt Ejland, Nuussuaq peninsula and Hareøen are all early Eocene in age (Naqerloq and Erqa Fms). Chauvet et al. (accepted) point out that both in W-Nuussuaq and Svartenhuk the Eocene lavas are part of the syn-extension inner SDR, i.e. that continental crust thinning and stretching still occurred during the Eocene (fig. 2).

2.3. *Svartenhuk volcanic facies*

We investigated and sampled 45 sites mostly located along the coastal cross-section of the Svartenhuk volcanic margin (Fig. 1). Hyaloclastic sub-aquatic breccias and associated pillowed lavas formed in a lacustrine environment (Escher and Pulvertaft, 1995) are characteristic of the lower member of the Vaigat Fm (V1 unit, sampling sites 40 and 38). They are up to 200 m thick and dip 20-30° westwards. These dips are mostly of tectonic origin but may also be due to dipping prograding foresets (Jerram, 2002), which have been identified for instance in the Naajaat syn-volcanic lake, Nuussuaq Basin (Pedersen et al., 1998). Svartenhuk hyaloclastic breccias consist of basaltic glassy clasts and pillow fragments, usually of centimeter- to decimeter-size, set in a matrix of yellowish glass particles altered to clays and palagonite. Locally, preserved pillow piles form cliffs up to 10-15 m high which delineate small high points on the shoreline. Individual basaltic pillows, 10 cm to 1 m large, display typical glassy margins and radial fractures together with flow features consistent with their emplacement over moderate slopes. Attempts to recover geochemically fresh glassy basaltic rims were unsuccessful, but the cores of large pillows proved to be rather well preserved from post-magmatic alteration.

Lava flows from V2, V3 and V4 units were exclusively emplaced in subaerial environments as suggested by lack of pillow lavas and hyaloclastites and common occurrence of pahoehoe surface features. Their most common facies, especially in V3 and V4 units, is that of individual tabular flows, 1 to 10 m thick. They generally show a basal scoriaceous autoclastic breccia, which overlies a reddish thermal metamorphic sole. These breccias are, in turn, overlain by a massive basaltic part often displaying rough columnar-jointing patterns, and less frequently sheet flow patterns. The rather common occurrence, in these massive levels, of subvertical vesicle pipes, sometimes feeding horizontal vesicle sheets (Caroff et al., 2000), suggests that the Svartenhuk Peninsula basaltic magmas were water-rich at the time of their emplacement. Tuff horizons usually less than 50 cm thick are occasionally interbedded between these tabular flows, and probably correspond to subaerial ash and/or lapilli fall deposits. Intercalated reworked (volcanic-sedimentary) deposits are much less common than in Disko and Nuussuaq (Larsen and Pedersen, 2000). They correspond to basaltic pebble conglomerates of probable laharic origin, filling local topographic channels. Their maximal thickness is less than 10 m (sites 44 and 45), and they are sometimes overlain by layered clay horizons less than 1 m thick.

Vaigat Fm picrites (V2 unit) and less commonly Vaigat and Svartenhuk Fms tholeiitic basalts (V2 and V3) display a more specific emplacement facies, i.e. a compound-braided flow facies (Jerram, 2002). The corresponding lava piles, up to 30 m thick (site 39), have a typical grayish color also characteristic of Disko and Nuussuaq picrites, (Larsen and Pedersen (2009), and are usually deeply altered and converted into assemblages of post-magmatic minerals (mostly zeolites and carbonates). They consist of thin (usually less than 50 cm thick) anastomosed individual layers emplaced as sheets and lobes of pahoehoe lava breakouts. These individual lobes are elongated (up to several meters) and usually hollow; all the transitions are observed between meter-size lobes and typical lava tubes. Well preserved small pahoehoe ropy lava “tongues” often occur within the central cavity of these lobes or tubes, and provide good markers of flow emplacement patterns. In V2 and V3 units they point usually upwards due to the tectonic tilting of these sequences. Massive (and geochemically better preserved) porphyritic layers 50 cm to 2-3 m thick are common within the lobe/tube piles. They display intrusive relationships with respect to them (locally chilled margins) and are thought to result from inflation processes within the pahoehoe compound flow sequences (Jerram, 2002; Self et al., 1997). In the studied picritic lava sequences, both pahoehoe lobes/tubes and intrusive (inflation-related) sheets commonly contain large vesicles and vesicle pipes typical of water-rich magma emplacement features (Caroff et al., 2000). These facies are often ascribed to relatively low volume, low viscosity eruptions occurring during early stages of emplacement of continental flood basalts (Jerram, 2002).

Most dykes range in width from 10-20 cm to 2-3 m, and display characteristic jointing patterns perpendicular to their chilled margins (which are glassy to microlitic) and have coarser-grained, often porphyritic, vesicle-rich central parts. These dykes tend to form swarms associated with fracture zones, and composite (multiple) dykes are common. Sills are usually easily distinguished from lava flows by their larger and more regular columnar jointing patterns. The thickest ones (> 10 m) display evidence for gravitational settling of olivine phenocrysts, which accumulate above their lower chilled margin.

3. Sampling and analytical methods

3.1. Sampling strategy

Sampling was conducted in 45 sites mostly located along the coastal cross-section of the volcanic margin (Table 1 and Fig. 2), from the basal lacustrine hyaloclastites and pillow basalts of the Vaigat Fm (labelled V1 in the geochemical discussion below; sites 40 and 38) towards successively: the subaerial picritic and basaltic flows of the main Vaigat Fm (labelled V2; sites 37, 39, 41, 42, 20, 43, 45, 44, 5, 3, 6, 7, 1, 27, 25, 28); then those of the overlying Svartenhuk Fm (labelled V3; sites 4, 2, 36, 35, 8, 31, 32, 11, 10, 34, 12, 9, 33); and finally the trachytic intrusion (site 29) and the subaerial alkali basaltic lava flows from the top of the Svartenhuk Fm, recently attributed to the Naqerloq Fm by Larsen et al. (2015) (site 30, labelled V4 below). Additional samplings were done for V2 (sites 19, 22, 23, 16, 14, 15, 13), V3 (site 26) and V4 (site 21). Major, trace element and isotopic measurements were only carried out on samples from lava flows, dykes and sills which were relatively fresh based on their macroscopic and microscopic features, and from which zeolite- and carbonate-bearing vesicles were removed by handpicking. The corresponding analytical database includes 5 samples from V1, 53 from V2, 28 from V3 and 5 from V4 lava flows, together with 47 intrusive (sills, dikes) samples (Tables 1, 2, 3 and 4). Most of the pillow fragments collected from V1 are altered (Loss on Ignition higher than 1.9 wt%), and a number of compound-braided basaltic flows (pahoehoe lobes and lava tube piles, see below), especially from V2, are too rich in secondary minerals (zeolites and carbonates) to have retained their primary magmatic features. Such samples were primarily collected to investigate these post-magmatic mineral associations found as vesicle fillings and vein/fracture deposits (57 samples from V2 and V3).

3.2. Analytical methods

All samples were subsequently crushed in stainless steel and agate mortars. Major element concentrations were obtained using an ICP-AES Jobin Yvon Ultima 2 at the University of Brest, after a HF-HNO₃ digestion, boric acid neutralization and dilution in nitric acid, as described in Cotten et al. (1995). Trace element concentrations were measured with a Thermo® Element2 HR-ICP-MS in Brest. 15 mg of samples were repeatedly (twice) dissolved at 120°C in 2 mL Teflon Savilex® beakers using in 1mL HF + 0.5 mL HClO₄. After total evaporation, samples were diluted 10000 times using 0.5N HNO₃ and analyzed using both medium and low resolution on the Thermo® Element2 as described by Mougél et al. (2014). All samples were externally calibrated using international standards BCR2, BHVO2 and BIR-1 (Li and Lee, 2006). These

standards demonstrated external reproducibilities better than 5–10% depending on the element and concentration as documented in Supplementary Table 1. Pb, Hf, Sr and Nd isotope compositions were obtained on single fractions (0.5–1 g of samples) leached for one hour at 100°C in 6mL of 6N HCl, and subsequently dissolved in 3mL HF + 1mL HNO₃ following the protocol described by Mougél et al. (2014). Hf, Nd and Pb were purified following the protocol of Blichert-Toft et al. (2005) (fluoride centrifugation, and column chromatography). Sr was separated from the Nd fraction using Eichrom Sr-specific resin following the protocol by Moynier et al. (2010), adapted from Fietzke and Eisenhauer (2006). Pb, Hf and Nd isotopes were analysed using a Thermo® Neptune MC-ICP-MS at Ifremer-Brest. Sr isotopes were obtained using a Thermo Triton TIMS at the University of Brest. 2σ internal errors for Hf and Nd are reported in Table 4 and never exceed the last significant digit (1E-5) in each analysis. External reproducibility of Pb isotopes is estimated based on the repeated measurements since 2008 of a mixed standard solution and never exceeded 60 ppm for ²⁰⁶Pb/²⁰⁴Pb, 150 ppm for ²⁰⁷Pb/²⁰⁴Pb and 300 ppm for ²⁰⁸Pb/²⁰⁴Pb. Measured values were compared with repeatedly analyzed international standards NBS981 (Todt et al., 1996), JMC475 (Blichert-Toft et al., 1997) and La Jolla48 (Tanaka et al., 2000) shown in Supplementary Table 2. Duplicated Sr, Nd, Hf, Pb isotope analyses of one of our samples (S39E1) are shown in Supplementary Table 3.

4. Petrographic and geochemical results

4.1. Petrographic features and lava classification

Lava samples were classified according to their position in the Total Alkali-Silica plot (TAS; Le Bas et al., 1986; Fig 3a). With the exception of a trachytic intrusion from V4 (site 29), all of them plot within or near the fields of microbasalts and basalts (SiO₂ = 43–53 wt%, K₂O+Na₂O=0.1–5 wt%). A large majority of them are silica-saturated or -oversaturated, and plot below the alkaline-subalkaline boundary of Macdonald and Katsura (1964). Therefore, these samples, which include all the massive or pillowed flows of units V1, V2 and V3, the sills and about half the population of dykes can be termed as tholeiitic or transitional picrites and basalts. Alkali basalts plotting above this boundary include the V4 basaltic flows (site 30) and the other half of the analyzed dykes. Some V3 basalts plot near or slightly above Macdonald and Katsura's (1964)

boundary (Fig. 3a) and can be considered as transitional, as well as V1 basalts which plot definitively below this limit but display strong enrichments in highly incompatible elements (see below).

Pillowed tholeiitic basalts from V1 unit display groundmass textures ranging from glassy (devitrified) rims to microlitic cores. They are subaphyric, with 4-8 modal% phenocrysts and microphenocrysts of olivine (mostly altered to smectite and chlorite) and fresh plagioclase ranging in size from 0.1 to 0.5 mm. Their groundmass is variably altered into clays and palagonite-type mineral aggregates. Zeolites and carbonates are found as vesicle- and vein-filling materials.

Picrites and picritic basalts (microbasalts in Fig. 3a) are much more common in subaerial Vaigat Fm flows (V2) than in the overlying main unit of Svartenhuk Fm (V3). They are highly porphyritic, with up to 60 modal% centimeter- to millimeter-size olivine phenocrysts, containing occasional magnesiochromite inclusions, and subordinate (and smaller) plagioclase phenocrysts and microphenocrysts. The percentage of olivine phenocrysts and their size tend to decrease together with decreasing MgO contents of bulk rocks, while the relative abundance of plagioclase increases. Clinopyroxene microphenocrysts are uncommon in picrites, and this mineral follows plagioclase in the crystallization sequence of V2 and V3 lavas. The picritic basalts from these two units are rather similar in thin section. Their groundmass is usually intergranular or microlitic. Neither native iron, nor graphite has been identified in the studied picrites. This suggests a weaker contamination in these samples (originally selected for their primitive features) than that described for some more acidic samples from Disko and Nuussuaq (Larsen and Pedersen, 2009).

Tholeiitic basalts from V2 and V3 are the most common petrographic type encountered in the studied sequences. Their phenocryst contents are highly variable and decrease together with MgO contents of bulk rocks, from 25-20 modal% (porphyritic basalts) to less than 5 modal% (subaphyric basalts). Olivine is followed by calcic plagioclase, and then sometimes by minor amounts of augitic clinopyroxene in their crystallization sequence. Plagioclase phenocrysts dominate over olivine in the most evolved (silica-rich) samples, which are especially common in the Svartenhuk Fm main unit (V3). Clinopyroxene phenocrysts mostly occur in this V3 unit, and are uncommon in V2 basalts. The groundmass of tholeiitic basalts usually display microlitic to intergranular textures. Subdoleritic textures occur in sills and in the cores of thick flows and dykes.

Post-magmatic mineral associations are commonly found in V2 and V3 tholeiitic picrites and basalts. They occur as vesicle- and vein-fillings, and include a variety of zeolites, carbonates (calcite, Mg-calcite, aragonite, dolomite), and clays (smectites, chlorites).

Alkali basalts from Naqerloq Fm. (V4) and from a number of dykes display microlitic to intergranular groundmass textures. They contain about 10 to 15 modal% phenocrysts, which are, by order of decreasing abundance, calcic plagioclase (up to 0.5 cm large), diopsidic clinopyroxene and magnesian olivine. They contain significantly smaller amounts of post-magmatic minerals than V1, V2 and V3 lavas.

The fresh alkali trachyte collected at site 29 (V4 unit) is highly porphyritic, with ~40 modal% centimeter-size sanidine phenocrysts, and lesser amounts of smaller titanomagnetite, olivine and green clinopyroxene phenocrysts. Its groundmass, almost exclusively made of alkali feldspar laths, displays a fluidal texture.

4.2. Major element trends

The Loss on Ignition (LOI) values (Table 2) range from 0.58 wt% to 10 wt%, with an average of 2.8 wt%. These rather high values are accounted for by the common occurrence of post-magmatic water-rich minerals (zeolites and clays) and of carbonates in the studied lavas. The variations of SiO₂ and MgO in the studied basaltic flows (V1 to V4) are plotted against their Ni, CaO and alkali element contents. The higher MgO contents of V2 lavas (Table 2) evidence the dominance of picrites (MgO > 12 wt%) and Mg-rich basalts in the Vaigat subaerial flows. An upward increase of TiO₂ is also observed in the stratigraphic pile: contents higher than 2 wt%, usually considered typical of High-Ti basalts from Large Igneous Provinces (Pik et al., 1998; Shellnutt and Jahn, 2011), are only observed in some V3 evolved tholeiitic and transitional basalts and in V4 alkali basalts (Table 2). Conversely, all V1 and V2 basalts and a majority of V3 tholeiites are typical Low-Ti basalts (TiO₂ < 2 wt%, Table 2). Similar features have been reported by (Larsen and Pedersen, 2009) for the Disko and Nuussuaq areas, where the lower Vaigat Fm is exclusively made of Low-Ti basalts while High-Ti lavas occur within the overlying Maligât Fm (lateral equivalent of Svartenhuk Fm according to Escher and Pulvertaft (1995)).

Major element variations in the studied sequence are dominated (and largely overprinted) by processes linked to accumulation or removal/fractionation of olivine phenocrysts. Indeed, as shown in Figure 3B, addition or removal of magnesian olivine

are consistent with the covariation of CaO and MgO: olivine-rich picritic basalts are CaO-poor while evolved basalts are CaO-rich. These variations are partly accounted for by local gravitational settling effects involving olivine phenocrysts in thick lava flows or sills. However, such shallow processes during and after emplacement are unlikely to occur in thin pahoehoe units and meter-thick tabular flows which make up most of the Vaigat and Svartenhuk Fms sequences. Therefore, fractional crystallization involving olivine separation/accumulation must have occurred at depth, possibly within thick sills or genuine underlying magma chambers. In addition, the least Mg-rich basalts, mostly sampled from dykes, display a co-linear decrease of CaO and MgO (Fig. 3B), that is consistent with concomitant removal/ fractionation of olivine plus augite and/or calcic plagioclase. Such variations might result from differential motions of crystal-rich and crystal-poor liquids during dyke emplacement (“Bagnold effect”, e.g. Komar, 1972), and thus cannot be taken as evidence for deep-seated fractional crystallization involving olivine, augite and plagioclase.

4.3. Trace element variations

Compatible trace elements Ni, Co and Cr, which are hosted by olivine and magnesiochromite, display positive correlations with MgO contents (Fig. 3B). Their contents decrease from picrites to evolved basalts, and their behaviors are mainly controlled by olivine removal/accumulation processes.

Incompatible element concentrations vary in a more complex fashion, as illustrated by Rare Earth Element (REE) plots normalized to CI chondrite (Fig. 4). V1 transitional basalts display the most fractionated REE patterns and the highest contents in Light REE (LREE; e.g. La, Ce), up to 80 times the primitive mantle (PM). V4 alkali basalts also display fractionated REE patterns, although their LREE enrichment remains lower than that of V1 lavas. V2 and V3 REE patterns are fan-shaped, and display variable enrichments in LREE, those of picritic samples being the lowest. The REE patterns of dykes are also fan-shaped, and they encompass the variability found within lava flows (Fig. 4). Alkali basaltic dykes show fractionated patterns and LREE enrichments similar to those observed for V4 alkali basaltic flows. A second family of dykes (and the analyzed sills) has rather flat patterns which resemble those of picrites; and a third dyke family shows moderately LREE enriched patterns similar to those of V2 and V3 evolved tholeiitic basalts. Large Ion Lithophile Elements (LILE: Cs, Rb, Ba, Sr) display somewhat erratic behaviors, with highly variable enrichments or depletions

generating positive or negative spikes in incompatible multi-element patterns normalized to the primitive mantle (Figure 4B). Given the high solubility of these elements into water, such variations can be considered as post-magmatic and attributed to alteration by surface waters and/or percolation of hydrothermal fluids through the studied lava piles. The elements less affected by post-magmatic processes (ie rare earth elements, high field strength elements...) define patterns similar to that of typical OIB (Fig. 5).

Highly incompatible elements (e.g. Th and La) versus MgO (not shown) in V2 and V3 lavas define a rough negative correlation (not shown) consistent with the “dilution” of incompatible elements in bulk rocks as a result of accumulation of incompatible element-poor olivine phenocrysts in picritic and Mg-rich lavas. Compared to V2 and V3, V4 alkali basalts and alkaline dykes display higher Th and La contents (Fig. 4) consistent with their more fractionated REE and multi-element patterns, as well as higher Nb/La ratios (Fig. 5). These features might indicate either lower degrees of partial melting, or a more incompatible element enriched source (with respect to V2 and V3 ones), or a combination of both. Finally, despite their transitional character, some V1 pillowed lavas (together with a few dykes) show the highest La, Th and Pb contents found in the studied lava sequence (Figs. 4 and 5), and consistent with their highly fractionated REE and multi-element patterns. Once again, these features could be attributed to a more enriched source, lower degrees of melting, or both.

4.4. Sr, Nd, Pb and Hf isotopes

Lead isotopes define two distinct trends (Figs. 6 and 7). $^{208}\text{Pb}/^{204}\text{Pb}$ range from 36.7 to 39, $^{207}\text{Pb}/^{204}\text{Pb}$ from 15 to 15.5 and $^{206}\text{Pb}/^{204}\text{Pb}$ from 16.13 to 19.26. A principal component analysis calculated on these Pb isotopes reveals the existence of only two significant principal components (the first one accounting for 94.2% of the total variability, principal component 2 for about 5.3%, and principal component 3 is insignificant). This result indicates that a mixture of three end-members is sufficient to explain the isotope variability in this area (Blichert-Toft et al., 2005). Hafnium isotopes are scattered as well, and feature both radiogenic and unradiogenic signatures ($^{176}\text{Hf}/^{177}\text{Hf}$ from 0.28247 to 0.2832; $\epsilon_{\text{Hf}} = -10.81$ to $+15.95$). They also define a three end-members-pattern corresponding to two distinct mixing trends (Fig. 6). Nd isotopes correlate with Hf data ranging from $^{143}\text{Nd}/^{144}\text{Nd} = 0.51222$ ($\epsilon_{\text{Nd}} = -8.15$) to

$^{143}\text{Nd}/^{144}\text{Nd}=0.51309$ ($\epsilon_{\text{Nd}}=8.9$). Even if most of the samples have “mantle-like” Sr isotopic compositions ($^{87}\text{Sr}/^{86}\text{Sr} < 0.704$), 16 of them display significantly more radiogenic Sr signatures (up to $^{87}\text{Sr}/^{86}\text{Sr} = 0.708$). Ten out of those 16 samples also have unradiogenic Pb ($^{206}\text{Pb}/^{204}\text{Pb} < 17$), and trace element patterns (low Nb, Ta and high Pb contents, Fig. 5) typical of continental crust contamination (Rudnick and Fountain, 1995; Rudnick and Gao, 2003). These samples include all of those collected from the basal unit V1.

The two mixing trends identified in Sr, Pb and Hf isotope spaces, are also visible on diagrams using isotopic and REE ratios (e.g., $^{206}\text{Pb}/^{204}\text{Pb}$, La/Sm, Sm/Yb and La/Yb, Fig. 8). They are also related to the stratigraphic position of the samples, i.e. their height within the lava pile. Indeed, all the samples from basal units V1 and V2 lie within the same trend (Figs. 6 and 7) connecting a component “Sk1” (with low $^{207}\text{Pb}/^{204}\text{Pb}$ and “intermediate” $^{208}\text{Pb}/^{204}\text{Pb}$) to a moderately high $^{208}\text{Pb}/^{204}\text{Pb}$ and $^{206}\text{Pb}/^{204}\text{Pb}$ component “Sk2”. Complementarily, all samples from units V3 and V4, together with those from sills and dykes, plot along a trend connecting “Sk2” to a component “Sk3” (with high $^{208}\text{Pb}/^{204}\text{Pb}$, $^{207}\text{Pb}/^{204}\text{Pb}$ and $^{206}\text{Pb}/^{204}\text{Pb}$).

5. Discussion

5.1. Trace element and isotopic evidence for a minor role of crustal contamination

The highly incompatible elements Nb and Ta display positive or negative peaks in multi-element patterns (Fig. 4B) which are unlikely of post-magmatic origin considering the insoluble character of these High Field Strength Elements (HFSE). Nb/La ratios show a tentative negative correlation with SiO_2 , whereas Pb contents are positively correlated with silica for V1 and V2 lavas (Fig. 5). These features likely reflect crustal contamination effects in these two lowermost units, since silica-rich continental crust is selectively depleted in Nb and Ta and enriched in Pb with respect to elements of similar incompatibility, e.g. La (Tang et al, 2019 and references therein)..

Component Sk1 is clearly distinct from either “typical” mantle-reservoirs (Hart et al., 1992), or any North Atlantic volcanic rocks. Its radiogenic Sr, unradiogenic Hf and Nd, as well as the low Nb, Ta, and high Pb contents (Fig. 5) of lavas plotting close to it (Figs. 6 and 7) suggest that the origin of this component is related to continental crust contamination of mantle-derived melts by materials similar in composition to

Greenland basement late Archean orthogneisses and granodiorites analyzed by Larsen and Pedersen (2009). The fact that these contamination effects are limited to the oldest units V1 and V2 is consistent with the hypothesis of an uprise of the younger melts (constituting units V3 and V4) through a plumbing system coated with earlier basaltic lavas (e.g. Arndt and Christensen, 1992) or exhausted of fertile material by previous melting events (e.g. Meade et al, 2014).

5.2. *Partial melting conditions*

Si and Mg contents can be used as proxies in order to estimate pressure and temperature of melt extraction in the mantle (Lee et al., 2009). Only a limited number of samples are “primitive enough” (i.e. experienced only olivine fractionation), to adequately estimate their pristine major element concentrations by simple olivine addition (Albarède, 1992; Lee et al., 2009). Moreover, some of them, with MgO contents >15 wt%, need to be afterwards corrected for olivine accumulation (back to equilibrium with their olivine phenocrysts Mg#~0.83, Supplementary Table 4) and olivine fractionation (back to equilibrium with olivine of mantle-like composition Mg#~0.89). Thermobarometric estimates thus obtained on ten samples from units V2 and V3 suggest temperatures and pressures of $\sim 1350 \pm 100$ °C and 2.5 ± 0.5 GPa, respectively. These results, although consistent with those recently reported by Hole and Natland (2019), need to be considered with caution, given the number of approximations in the thermobarometric approach, as well as uncertainties concerning the precise geological setting at that time (crustal thickness). The estimated temperature range suggests that the mantle beneath Svartenhuk was not very hot compared with other published evaluations for the NAIP (e.g. 1480-1550 °C, Hole and Millett, 2016; see also Herzberg et al. 2007 and references therein). Our estimated P-T conditions overlap the spinel/garnet transition zone in the mantle (2.2 GPa at 1300 °C, O'Neill, 1981; Robinson and Wood, 1998), and it is therefore probable that Svartenhuk Fm. lavas were extracted from melting columns involving both garnet- and spinel-bearing peridotites, as often proposed for NAIP magmas (Kerr, 1994; Kerr et al, 1999; Saunders et al., 2007; Hole and Millett, 2016).

Even if the “fan-shaped” rare earth elements (REE) patterns observed on figure 4 are partly the result of different rates of partial melting, LREE/HREE fractionation can also be accounted for by the presence of residual garnet in the source (e.g. Frey et al. 1978, Thirlwall et al., 1994). In figure 8 plots, Pb isotopes, La/Sm and Sm/Yb ratios are

540 compared with the average slope of REE spectra quantified by La/Yb ratios. The upper
541 diagram ($^{206}\text{Pb}/^{204}\text{Pb}$ vs La/Yb) reflects the effects of crustal contamination (units V1
542 and V2) on geochemical compositions (see section 5.1). The middle diagram documents
543 the concomitant effects of crustal contamination and degrees of partial melting on
544 LREE/intermediate REE relative abundances. The lower plot provides insights
545 concerning the amount of garnet-bearing peridotites involved in the melting column.
546 Altogether, these three diagrams suggest that:

547 i/ Lavas derived from rather high degrees of partial melting all belong to units V2
548 and V3. Basalts from units V1 and V4 were generated by smaller degrees of
549 mantle melting (as suggested by their alkali to transitional signatures in Figs. 4
550 and 5). Precise amounts of melting are difficult to assert considering the
551 uncertainties concerning the initial composition(s) of mantle source(s).
552 Nevertheless, models of garnet- (shown in red) and spinel-bearing (shown in
553 green) batch lherzolite melting (see Gurenko and Chaussidon, 1995 for details on
554 initial compositions, McKenzy and O'nion, 1991 for partition coefficients; and
555 Thirlwall et al.1993 for mineral proportions in the melting) show that V1 and V4
556 lavas would be derived from melting degrees lower than 5%, and most V2 and
557 V3 lavas from clearly higher ones (Fig. 8).

558 ii/ On average, basalts from units V2 and V3 feature Sm/Yb ratios 50% lower
559 than those of samples from units V1 and V4. This stronger fractionation between
560 intermediate and heavy REE in V1 and V4 suggests a larger involvement of
561 garnet peridotites (and therefore greater melting depths) at the beginning and the
562 end of the magmatic history than during the emplacement of the main V2-V3
563 volcanic sequences. These results are in agreement with previous petrogenetic
564 studies of West Greenland picritic and Mg-rich basalts demonstrating that their
565 main sequences derived predominantly from spinel-bearing peridotites (Gill et al.,
566 1991; Ryabchikov et al., 1998; Larsen and Pedersen, 2000, 2009).

567 Therefore, we propose that small degrees of melting leading to V1 lavas were
568 initiated at depths greater than 75 km below a relatively thick lithosphere. Subsequently,
569 these melts propagated upwards into the spinel peridotite zone in connection with
570 lithospheric thinning. The melting rates increased consistently during this process,
571 forming V2 and V3 lavas. Finally, the melting zone deepened again, and melting rates
572 decreased, thereby generating V4 alkali basalts.

5.3. Mantle source heterogeneity and its temporal evolution

The isotope diversity reported in this study covers a range similar to that previously reported for the neighboring Disko Island (Larsen and Pedersen, 2009) and the only differences concern the extent of crustal contamination (Fig. 5). The least contaminated samples (identified on Figs. 5 and 8) also largely overlap with samples from Iceland and the adjacent ridges, as well as with those from East Greenland (figs. 6 and 7). The isotopic diversity within the Iceland plume has been widely discussed (Chauvel and Hémond, 2000; Thirlwall et al., 2004; Kokfelt et al., 2006; MacLennan, 2008; Kitawaga et al., 2008) and is denoted in figure 7 by both depleted (D, Kitawaga et al., 2008; ID-1 and ID-2, Thirlwall et al., 2004), and enriched (E-1 and E-2, Kitawaga et al., 2008; IE-1 and IE-2, Thirlwall et al., 2004) Icelandic internal components (Fig. 7). In figures 6 and 7, Hf, Sr and Pb compositions of component “Sk2” are very similar to those of OIB and E-MORB related to the Iceland hotspot, and more especially to a mixture of the Icelandic inner components “D” (for depleted) and “E-1” (for Enriched 1) of Kitawaga et al. (2008) or “ID1” – “ID2” (for Iceland Depleted 1 and 2) and “IE1” (for Iceland enriched 1) of Thirlwall et al. (2004).

Alternatively, Sk3 seems to correspond to the Iceland E-2 enriched component of Kitawaga et al. (2008) (or “IE-1” of Thirlwall et al., 2004) that point toward the common component C (Hanan and Graham, 1996) in all isotope spaces. This enriched component has been proposed to be intrinsic to the Icelandic hotspot (Hanan and Schilling, 1997; Kitawaga et al., 2008; Thirlwall et al., 2004; Peate et al., 2010). The presence of distinct Icelandic isotopic components has also been described in Eastern Greenland (Barker et al., 2006). Nevertheless, in that region, the different signatures appear in separate areas (Milne, Rømer and Skræterne), located hundreds of kilometers away from each other. In the present study, we report distinct signatures on samples all coming from a restricted segment (<40 km along the coast) and covering a time span shorter than 8 m.y. Such diversity can therefore hardly be imagined as the result of the tapping of distinct parts of an Iceland type mantle plume. Instead, we suggest that the Sk3 signature in our dataset corresponds to a trend toward the C component of Hanan and Graham (1996). This component has been recognized as being widely disseminated within the upper mantle feeding the Arctic segments of the mid-Atlantic ridges and is therefore believed to correspond to an ubiquitous ingredient of the North Atlantic upper mantle (Blichert-Toft et al., 2005).

Nb, Zr and Y concentrations also suggest consistency between uncontaminated Svartenhuk samples and Iceland plume geochemical signatures. Indeed, Fitton et al. (1997) used the comparison of Nb/Y and Zr/Y in order to decipher whether the magmatic sources of basalts are more likely related to the Iceland plume or to the neighboring MORB mantle. They proposed a ΔNb index $= 1.72 + \log(\text{Nb}/\text{Y}) - 1.92\log(\text{Zr}/\text{Y})$; where ΔNb is negative in MORB samples, but positive in Iceland plume-derived material. Even if this approach has been criticized as possibly biased by differences in pressures of melting (Hanan et al., 2000; Stracke et al., 2003), it is worth noting that most samples from this study feature positive ΔNb values. Indeed, in Figure 9, only some samples from units V2 and V3 display negative ΔNb (Fig. 9), suggesting that the contribution of Icelandic material in lava sources was lower during these intermediate stages than during the emplacement of the earliest and latest V1 and V4 units. Moreover, a plume imprint on geochemical signatures of Svartenhuk lavas is also consistent with the extremely primitive helium isotopic compositions recently published by Péron et al. (2018) for olivines from our V3 samples S33E7 and S35E7 ($R/\text{Ra}=31.0$ and 34.9 , respectively), similar to those previously reported for samples from Disko Island and Baffin Island (Graham et al., 1998; Jackson et al., 2010).

Altogether, these features suggest that the Sk2-Sk3 trend (common to both V3 and V4 units) could be accounted for by a mixture of Icelandic-type material (Sk2) and a mixed ambient upper mantle reservoir (Sk3). The respective proportions of these two components appear to evolve through time, since only samples coming from unit V3 carry isotope signatures suggesting the involvement of significant amounts of C-like material, whereas the uppermost V4 lavas seem to tap preferentially Iceland hotspot material. Furthermore, even if geochemical signatures of V1 and V2 are partly controlled by crustal contamination, they also point solely toward the Iceland-like signature. This sequence of magmatic events is summarized in figures 10 (petrogenetic constraints) and 11 (speculative tectonic interpretation). During the emplacements of basal units V1 and V2 (61 to 59.5 Ma old, Chauvet et al., accepted), initially low, but temporally increasing degrees of melting of enriched Icelandic plume material (displaying Sk2 isotopic signatures) generated basaltic melts at depths starting beneath 75 km, and these magmas were contaminated during their ascent through the continental crust. During V3 emplacement (59.5-57.7 Ma), the melting process affected a wider range of lithologies in the spinel stability field and propagated to neighboring

(shallower) materials from the upper mantle (involving material displaying component Sk3 isotope signatures), with little or undetectable crustal contamination. Finally, during V4 emplacement, i.e. between 57.7 and 53.3 Ma (Chauvet et al., accepted), the melting zone deepened again and was once more mostly restricted to the enriched mantle material (Figs. 10 and 11).

5.4. Baffin Bay breakup and tectonic controls of mantle melting

This sequence of events as reported by combining geochemical and tectonic data can be interpreted as the progressive initiation of decompression melting of an upwelling mantle below a progressively thinning lithosphere in a continental extension setting (Fig. 11). The thinning observed between the emplacement of V1 and V2-V3 lavas likely results from a combination of the tectonic stretching of the upper part of the lithosphere and of the thermal erosion of its base by convecting hot mantle (Fig. 11). Such a combination has been proposed as an explanation of the magmatic history of Mull and Skye islands (Kerr, 1994; Kerr et al., 1999; Saunders et al., 1997). In these islands, as well as in Giant's Causeway basalts (Barrat and Nesbitt, 1996), temporally increasing melting rates and decreasing melting pressures similar to those described in Svartenhuk are observed, despite rather different tectonic contexts characterized by lower extension rates (Meyer et al., 2007). A temporal decrease of lithospheric thickness seems the most likely factor controlling such changes in NAIP basaltic magma compositions (Ellam, 1992; Kerr, 1994).

The decreasing involvement of the ambient mantle (component Sk3) contribution from V3 to V4 (Fig. 10), and the concomitant deepening and decrease of partial melting rates (back to alkali basaltic compositions) in this most proximal part of the volcanic margin, may be (at least partly) attributed to the gradual westward shift of extension with time (Geoffroy et al., 2015; Chauvet et al., accepted). Indeed, the innermost Svartenhuk margin progressively became passive although extension still continued (as attested by syn-tectonic control of V4 emplacement), albeit at lower rates (Chauvet et al., accepted). It is also possible that the accumulation beneath the lithosphere of refractory mantle residues left after melt extraction blocked the ascent of hot and fertile mantle sources and resulted in deeper, lower-degree melting as previously suggested for East Greenland flood basalts (Fram and Leshner, 1993, 1997).

6. Conclusions

Geochemical data obtained on 93 samples of basaltic to trachytic rocks from the Svartenhuk peninsula allow us to document the c. 7 m.y. long volcanic activity accompanying the continental thinning process that preceded the Eocene opening of Baffin Bay, the chronological and tectonic aspects of which are discussed in a companion paper (Chauvet et al., accepted). The sequence of magmatic events started with (i) the deep (>75 km) low degree partial melting of an Iceland plume-like source, followed by (ii) shallower and higher degrees of melting of a mixture of Iceland hotspot and “ambient upper mantle” materials, and finally ended with (iii) lower degrees of melting of a deeper source dominated again by Icelandic-type signatures. The first melting event could have been triggered by the adiabatic ascent of the Iceland plume below a ~75-km-thick lithosphere, inherited from the previous Mesozoic extension. The OIB-type signature of V1 lavas is clear despite the crustal contamination processes that they experienced during their ascent. The following melting events (V2 to V3 lavas) occurred at decreasing pressures and were coeval with syn-magmatic extension in the upper crust (Chauvet et al., accepted) and therefore a subsequent reduction in lithosphere thickness (Figs. 10 and 11). Thus, V2-V3 melts were a direct consequence of the convective upwelling of Icelandic OIB-type mantle in the necked sub-lithospheric area (Boutiller and Keen, 1999), associated with lithospheric thinning resulting from a combination of tectonic stretching and thermal erosion of the Svartenhuk lithosphere. The later decrease in magmatic activity (emplacement of V4 lavas) was concomitant with a progressive focusing of syn-magmatic strain towards the central axis of the proto-Baffin Bay during the Eocene. This shift in extension was likely accompanied by a progressive cooling (and thus, thickening) of the proximal W-Greenland lithosphere over a still convecting asthenosphere. The combined decrease in melting rates and the return to enriched source materials suggest a further deepening of the melting zone, possibly linked to the accumulation beneath the lithosphere of refractory mantle residues.

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Figure captions

Figure 1: Simplified structural map of Greenland and Northern Atlantic Igneous Province, modified after Chauvet et al (accepted). The studied area (red square) is located on the western volcanic margin of Greenland.

Figure 2: The 45 sampling sites (numbers) located on the simplified geological map of Svartenhuk Peninsula (West Greenland), modified after Chauvet et al. (accepted). Locations of cross sections AA', BB' and CC' are shown on the map.

Figure 3: Major element compositions of samples from units V1 (purple triangles), V2 (beige circles), V3 (blue hexagons) and V4 (yellow triangles). A: Total alkali – Silica plot (Le Bas et al., 1986). The thick black line (Macdonald and Katsura, 1964) separates the fields of subalkaline basaltic series (below) from alkali basaltic series (above).

B: CaO vs SiO₂ and Ni vs SiO₂ plots. Compositions range from basaltic to trachytic (the latter in V4 unit only) following olivine and plagioclase + pyroxene fractionation trends. Dashed line identifies primitive Ni concentrations (higher than 400 ppm) and dotted line denotes samples with MgO=12 wt%, consistent with olivine phenocryst compositions (Mg#=0.83, supplementary table 1).

Figure 4: Multielement patterns of Svartenhuk samples. A: Rare earth element (REE) patterns normalized to CI Chondrite (McDonough and Sun, 1995) for samples from units V1 (pink), V2 (beige), V3 (blue) and V4 (yellow). Samples from V1 and V4 have comparable REE patterns with significant light REE enrichments compared with

heavy REE, typical of alkali and transitional basaltic lavas. Samples from units V2 and V3 have more diverse signatures ranging from alkali to tholeiitic (LREE depleted) patterns. B: Multielement plot normalized to Primitive mantle (McDonough and Sun, 1995) compiling all samples (V1=purple lines, V2= beige lines, V3= blue lines, V4= yellow lines) and the median compositions of each group (V1= purple triangles, V2= beige circles, V3= blue hexagons, V4= yellow triangles).

Figure 5: Plot of SiO₂ vs Pb concentrations and Nb/La ratios. Continental crust contamination typically generates SiO₂ and Pb enrichments (Rudnick and Fountain, 1995) associated with Nb depletions (Tang et al, 2019 and references therein) illustrated here by low Nb/La ratios. Crustal contamination trends (black arrows) can be identified within samples from units V1 and V2 only.

Figure 6: Pb, Hf, Sr and Nd isotopic data. Samples from units V1 to V4 (see figure 3 for color codes) are plotted together with basalts from the North Atlantic ridge : blue crosses (Agranier et al., 2005; Blichert-Toft et al., 2005), Iceland : gray squares: (Breddam, 2002; Chauvel and Hémond, 2000; Hanan and Schilling, 1997; Hanan et al., 2000, Kitawaga et al., 2008; Kokfelt et al., 2009; Kuritani et al., 2011; Peate et al., 2010, 2009; Sims et al., 2013; Stracke et al., 2003; Thirlwall et al., 2004; Willbold et al., 2009) and East Greenland: orange crosses (Barker et al., 2006). Samples from this study plot between three end-members materialized by letters Sk1, Sk2 and Sk3. The Sk1-Sk2 trend can be accounted for by crustal contamination (Sk1 features radiogenic Sr, and unradiogenic Pb, Hf and Nd); while Sk2-Sk3 largely overlaps with mantle like signatures (Kitawaga et al. 2008) including North Mid-Atlantic Ridge and Iceland basalts. Its trend in all diagrams appears to point toward the common component C of Hanan and Graham (1996).

Figure 7: Pb isotopic data. Samples from this study are represented together with basalts from Iceland (gray squares, see figure 6 for references) and East Greenland (Barker et al., 2006) (orange crosses). The Iceland plume inner components: D, E-1 and E-2 (Kitawaga et al., 2008; Thirlwall et al., 2004) are also located in this diagram. Apart from continental crust trend (segment Sk1-Sk2), all samples plot within in a triangle delineated by D, E-1 and E-2.

Figure 8: Plots of $^{206}\text{Pb}/^{204}\text{Pb}$, La/Sm, Sm/Yb against La/Yb. Pb isotopes (upper diagram) evidence crustal contamination for some V1 and V2 lavas and mantle-like compositions for most of the sample set. La/Sm as well as Sm/Yb provide insights about melting rates. On the Sm/Yb plot are reported for comparison the results of simple batch melting models considering garnet bearing (red line) and spinel bearing (green line) depleted lherzolite as proposed for Iceland by Gurenko and Chaussidon (1995), partition coefficients by McKenzie and O’Nions (1991) and mineral proportions by Thirlwall et al. (1993). Numbers from 0.02 to 0.1 denote F values. The higher Sm/Yb values observed for most of V1 and V4 lavas suggest a stronger involvement of garnet in their sources.

Figure 9: Plots of $^{206}\text{Pb}/^{204}\text{Pb}$, La/Sm and Sm/Yb against $\Delta\text{Nb} = 1.74 + \log(\text{Nb}/\text{Y}) - 1.92 \log(\text{Zr}/\text{Y})$ (Fitton et al., 1997). ΔNb is considered as a bench marker of Iceland plume affinity. Only samples from units V2 and V3 display negative ΔNb , indicative of the contribution of “normal” upper mantle devoid of plume related enriched component. The thick black arrows illustrate the temporal evolution of Icelandic material contribution from V1 to V4.

Figure 10: Sketch summarizing the four main stages of Svartenhuk volcanism deduced from geochemical and petrologic constraints. Sk1, Sk2 and Sk3 represent the three geochemical reservoirs defined in figures 6 and 7. The tectonic evolution of the lithosphere (crustal thinning) is more precisely depicted in figure 11. During V1 stage, decompression melting affects Iceland plume-like material ascending through the asthenosphere below an already thinned lithosphere. During V2, the ongoing crustal thinning leads to an uprise of the isotherms and therefore to the upward propagation of partial melting and to its increase. During the emplacement of these two first units, geochemical signatures (benchmarked by Pb isotopes on the bottom diagram) can be accounted for by the simple binary mixture of components Sk1 and Sk2. During V3, the melting has propagated within Sk3 reservoir. Finally, during V4, the melting deepens again and is once more mostly restricted to reservoir Sk2. See text for details.

Figure 11: Tentative tectonic sketches depicting Svartenhuk magmatic evolution in relation with the evolution of South Baffin Bay during the Paleogene, at the times of V1, V2-V3 and V4 emplacements. Sk1, Sk2 and Sk3 refer to crustal and mantle

reservoirs discussed in the text and shown in figure 10. Convection geometry is purely conceptual, and based on small scale convection models (Petersen et al., 2018; Simon et al., 2009). In the V1 sketch (top left), the occurrence of Cretaceous sediments, shown in light green suggests that crustal thinning started during the Cretaceous. The melting zones are indicated with coloured dots. Red boxes show the location of our field and petrologic investigations. The shaded areas on the left of some diagrams underline the speculative character of these interpretations when extrapolated to areas located away from our study area.

Table 1: Sample locations. K/Ar and Ar/Ar ages are from Chauvet et al (accepted)

Table 2: Major element data (oxide wt%, total iron as Fe₂O₃, LOI: Loss On Ignition). Analytical methods are presented in the text.

Table 3: Trace element data (ppm). Analytical methods are presented in the text.

Table 4: Pb, Hf, Sr and Nd isotopic data. Analytical methods are presented in the text.

Supplementary Table 1: ICP-MS repeated trace element analyses for standards BHVO2, BCR-2 and BIR1.

Supplementary Table 2: MC-ICP-MS repeated Nd, Hf and Pb isotope analyses of standards JNdi-1, JMC475 and NIST981, respectively.

Supplementary Table 3: Replicated analyses of Pb, Hf, Nd and Sr isotopic compositions of sample S39E1.

Supplementary Table 4: Olivine compositions obtained on Cameca SX100 electron microprobe at Service Commun “Microsonde Ouest” (Plouzané), at 15 kV accelerating voltage with a probe current of 12 nA.

Figure 1

Figure 2

A **B**

Sample names	longitude	latitude	Fm	³⁹ Ar- ³⁹ Ar age K-Ar age (My)
	Lon	lat		
S19E1	-53.974	71.470 V1		
S19E2	-53.974	71.470 V1		
S38E2	-53.936	71.446 V1		
S38E8	-53.936	71.446 V1		
S40E1	-53.899	71.455 V1		59.11 ± 0.83
S40E2	-53.899	71.455 V1		
S40E4	-53.893	71.453 V1		
S40E5	-53.893	71.453 V1		
S40E6	-54.067	71.426 V1		60.38 ± 1.34
S1E1	-54.605	71.370 V2		
S1E2	-54.605	71.370 V2		
S1E3	-54.606	71.370 V2		
S1E4	-54.606	71.370 V2		
S1E5	-54.604	71.368 V2		
S1E6	-54.607	71.369 V2		
S3E1	-54.300	71.386 V2		
S4E1	-54.679	71.363 V2		
S5E11	-54.282	71.395 V2		
S5E5	-54.288	71.394 V2		
S5E6	-54.287	71.394 V2		
S6E10	-54.359	71.374 V2		
S6E11	-54.367	71.378 V2		
S6E6	-54.364	71.378 V2		
S6E9	-54.366	71.378 V2		
S7E10	-54.458	71.371 V2		
S7E11	-54.457	71.371 V2		
S7E4	-54.461	71.372 V2		
S7E5	-54.461	71.372 V2		
S7E6	-54.456	71.370 V2		
S7E7	-54.459	71.371 V2		
S7E8	-54.455	71.370 V2		
S7E9	-54.459	71.371 V2		
S15H4	-54.719	71.438 V2		
S17H1	-54.218	71.835 V2		
S17H2	-54.224	71.834 V2		
S20I4	-54.117	71.425 V2		
S20E10	-54.109	71.422 V2		
S20E15	-54.118	71.425 V2		
S20E17	-54.119	71.425 V2		
S20E7	-54.112	71.425 V2		
S22H3	-54.486	71.417 V2		
S27E10	-54.738	71.356 V2		
S27E12	-54.740	71.359 V2		
S27E13	-54.741	71.359 V2		
S27E15	-54.751	71.356 V2		
S27E17	-54.751	71.356 V2		
S27E18	-54.752	71.359 V2		
S27E20	-54.752	71.356 V2		
S37E1	-55.204	71.389 V2		
S38E10	-53.942	71.446 V2		
S38E6?	-53.936	71.446 V2		
S39E1	-53.974	71.443 V2		
S41E1	-54.066	71.432 V2		
S41E3	-54.066	71.432 V2		
S42E1	-54.074	71.432 V2		
S42E2	-54.074	71.432 V2		
S42E4	-54.082	71.431 V2		
S42E5	-54.082	71.431 V2		61.08 ± 0.56
S43E1	-54.133	71.423 V2		
S43E2	-54.134	71.423 V2		
S43E4	-54.136	71.423 V2		
S43E5	-54.136	71.423 V2		
S43E6	-54.139	71.422 V2		
S44E10	-54.163	71.419 V2		
S44E2	-54.154	71.420 V2		
S44E4	-54.156	71.420 V2		
S44E7	-54.157	71.420 V2		
S44E8	-54.159	71.420 V2		
S44E9	-54.160	71.419 V2		
S45E10	-54.151	71.420 V2		
S45E4	-54.150	71.421 V2		
S45E9	-54.151	71.420 V2		
S2E2	-54.791	71.358 V3		
S2E3	-54.792	71.358 V3		
S2E5	-54.793	71.358 V3		
S2E7	-54.791	71.355 V3		
S2E8	-54.791	71.355 V3		54.98 ± 1.16
S2E9	-54.796	71.358 V3		
S8E10	-55.006	71.438 V3		
S8E13	-55.013	71.437 V3		55.73 ± 2.49
S8E6	-55.004	71.438 V3		
S8E9	-55.003	71.436 V3		
S9E1	-55.201	71.393 V3		
S11E10	-55.102	71.436 V3		
S11E12	-55.104	71.437 V3		
S11E7	-55.110	71.435 V3		
S26H1	-55.086	71.479 V3		54.20 ± 2.88
S26H3	-55.088	71.479 V3		
S33E1	-55.206	71.393 V3		
S33E12	-55.219	71.394 V3		
S33E4	-55.206	71.393 V3		
S33E7	-55.208	71.393 V3		
S33E8	-55.210	71.393 V3		57.78 ± 1.88
S34E2	-55.118	71.401 V3		
S34E5	-55.117	71.403 V3		
S34E6	-55.120	71.399 V3		
S34E7	-55.120	71.399 V3		
S35E1	-54.920	71.408 V3		
S35E11	-54.916	71.410 V3		
S35E12	-54.916	71.410 V3		
S35E3	-54.919	71.408 V3		
S35E5	-54.918	71.409 V3		59.78 ± 0.41 59.15 ± 0.85
S35E7	-54.917	71.409 V3		
S35E8	-54.917	71.410 V3		
S36E1	-54.922	71.404 V3		
S36E4	-54.922	71.403 V3		
S36E5	-54.922	71.403 V3		
S21H1	-55.338	71.541 V4		55.20 ± 0.79 55.86 ± 1.25
S21H2	-55.340	71.541 V4		
S29E2	-55.218	71.495 V4		
S30E1	-55.249	71.502 V4		54.39 ± 0.60 59.07 ± 1.16
S30E2	-55.236	71.503 V4		
S30E3	-55.244	71.503 V4		57.96 ± 1.20
S30E4	-55.241	71.502 V4		58.73 ± 1.53
S11E9	-55.118	71.435 dike		
S13H1	-54.823	71.489 dike		
S13H5	-54.823	71.489 dike		
S14H1a	-54.773	71.438 dike		
S14H2	-54.773	71.438 dike		56.43 ± 1.06
S15H3	-54.719	71.438 dike		
S16H1	-54.635	71.441 dike		
S16H2	-54.635	71.441 dike		
S16H4	-54.635	71.441 dike		
S20E18	-54.108	71.422 dike		55.42 ± 1.91
S22H2	-54.491	71.423 sill		53.85 ± 2.84
S22H4	-54.486	71.417 dike		54.41 ± 0.99
S23H1	-54.456	71.390 dike		
S23H4	-54.456	71.390 dike V3		58.98 ± 0.93
S23H5	-54.456	71.390 dike		
S24H1	-53.783	71.679 Sill Itsako		55.31 ± 1.09
S27E4	-54.752	71.356 dike		55.95 ± 0.88
S27E8	-54.738	71.356 dike		
S41E2	-54.060	71.426 dike		

Sample names	Fm.	wt. %	SiO ₂	TiO ₂	Al ₂ O ₃	FeO*	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	LOI	total
S19E1	V1		44.8	1.38	11.9	11.4	0.2	13.2	10.1	1.57	0.15	0.15	4.27	100.5
S19E2	V1		48.4	1.31	14.1	9.6	0.16	8.0	11.2	2.05	0.31	0.15	4.27	100.5
S38E2	V1		47.2	1.29	14.8	9.5	0.19	7.4	11.5	2.1	0.26	0.15	3.65	99.0
S38E8	V1		51.1	1.29	14.5	9.9	0.16	8.2	10.0	2.13	0.21	0.15	2.03	100.7
S40E1	V1		51.4	1.32	14.6	9.4	0.17	6.2	11.1	2.56	0.59	0.17	1.37	100.0
S40E2	V1		50.4	1.33	14.9	9.4	0.21	6.3	11.4	2.63	0.51	0.17	1.94	100.3
S40E4	V1		51.9	1.37	14.9	9.5	0.16	6.0	10.7	2.73	0.61	0.16	1.07	100.1
S40E5	V1		47.3	1.1	13.7	12.7	0.2	5.4	10.5	2.83	1.06	0.36	0.91	99.5
S40E6	V1		52.5	1.33	15.2	9.5	0.17	6.2	10.6	2.72	0.68	0.16	1.29	101.3
S1E1	V2		43.9	1.45	11.6	10.7	0.16	12.7	9.6	1.93	0.15	0.17	1.1	94.6
S1E2	V2		41.6	0.91	8.1	10.8	0.17	23.4	7.2	0.75	0.05	0.12	5.65	99.9
S1E3	V2		43.1	1.01	9.4	11.1	0.17	21.6	7.0	1.15	0.05	0.12	3.87	99.7
S1E4	V2		45.6	1.44	13.2	10.8	0.18	11.1	11.0	2.09	0.17	0.16	1.12	98.0
S1E5	V2		46.7	1.82	15.6	10.3	0.19	7.3	12.1	2.3	0.37	0.22	2.63	100.7
S1E6	V2		44.4	1.44	12.9	10.8	0.2	10.9	11.0	1.86	0.08	0.16	5.52	100.4
S3E1	V2		47.3	4.11	12.6	15.1	0.25	4.7	9.8	2.91	1.18	0.54	-0.58	99.6
S4E1	V2		45.3	1.66	13.9	11.0	0.18	9.1	11.1	2.49	0.24	0.14	3.63	100.0
S5E11	V2		49.5	1.02	12.7	8.5	0.17	13.0	6.5	1.88	0.99	0.12	3.79	99.1
S5E5	V2		49.0	1.37	13.9	10.1	0.26	8.7	10.1	2.13	0.54	0.15	2.46	99.7
S5E6	V2		49.5	1.56	14.6	9.2	0.17	8.2	10.8	2.52	0.84	0.17	1.96	100.6
S6E10	V2		42.1	3.8	13.0	13.5	0.21	5.9	10.7	2.74	0.72	0.71	2.69	97.6
S6E11	V2		43.1	1.1	9.1	11.3	0.18	21.7	7.7	0.82	0.31	0.13	5.88	102.5
S6E6	V2		43.4	1.04	9.5	11.4	0.17	21.7	7.4	1.24	0.14	0.11	3.02	100.5
S6E9	V2		41.8	1.08	9.0	10.9	0.17	20.9	7.2	0.87	0.26	0.12	5.9	99.3
S7E10	V2		42.7	1.52	11.7	11.3	0.19	13.1	10.6	1.09	0.06	0.16	7.47	101.0
S7E11	V2		41.1	0.99	8.4	10.7	0.17	24.6	6.0	0.6	0.06	0.11	5.96	99.8
S7E4	V2		44.2	1.58	12.4	12.2	0.25	10.8	10.4	2.14	0.12	0.17	4.85	100.4
S7E5	V2		42.1	1.01	7.7	10.4	0.17	26.2	5.8	0.51	0.07	0.12	6.28	101.4
S7E6	V2		44.9	1.51	11.3	11.2	0.19	15.1	9.7	1.4	0.22	0.15	4.36	101.3
S7E7	V2		40.2	1.01	7.4	10.4	0.16	25.1	6.3	0.38	0.07	0.11	7.07	99.3
S7E8	V2		46.4	4.11	12.9	14.8	0.25	4.7	9.9	2.79	1.46	0.59	0.5	100.0
S7E9	V2		45.4	1.26	13.9	10.1	0.15	10.8	11.5	1.58	0.17	0.12	4.46	100.6
S15H4	V2		42.5	0.84	8.5	11.0	0.18	24.7	7.2	1.01	0.13	0.11	3.32	100.7
S17H1	V2		49.1	2.18	15.5	12.4	0.21	4.8	10.9	2.7	0.15	0.23	1.39	100.9
S17H2	V2		48.2	2.16	15.1	12.3	0.2	4.6	10.7	2.59	0.22	0.23	1.83	99.4
S20I4	V2		44.8	1.77	13.1	10.7	0.22	8.7	11.1	2.16	0.24	0.17	3.66	97.7
S20E10	V2?		48.5	2.2	13.6	13.0	0.22	5.8	11.3	2.67	0.29	0.22	0.52	99.8
S20E15	V2		45.4	1.43	10.8	11.3	0.18	16.5	9.2	1.68	0.28	0.16	2.06	100.2
S20E17	V2		43.0	1.02	8.4	12.1	0.18	24.1	6.5	1.06	0.13	0.12	3.87	101.8
S20E7	V2		45.4	1.61	13.9	10.5	0.18	8.8	11.6	2	0.21	0.18	3.64	99.2
S22H3	V2		47.2	2.47	13.1	13.3	0.23	5.9	10.9	2.46	0.35	0.27	0.63	98.2
S27E10	V2		46.7	2.53	14.8	11.3	0.19	6.2	10.7	2.69	0.9	0.35	0.82	98.3
S27E12	V2		43.4	1.1	8.5	11.7	0.18	23.3	7.0	1.12	0.09	0.1	1.5	99.4
S27E13	V2		46.7	1.72	12.9	11.2	0.17	12.6	10.5	2.01	0.12	0.16	1.61	100.9
S27E15	V2		46.7	1.85	13.8	11.0	0.18	7.6	11.6	2.37	0.16	0.2	1.77	98.4
S27E17	V2		47.2	3.35	13.7	14.1	0.23	4.5	9.0	3.44	1.17	0.4	1.07	99.7
S27E18	V2		45.8	1.45	13.3	10.5	0.15	9.8	10.5	2	0.16	0.13	3.13	98.1
S27E20	V2		47.6	2.62	13.5	12.2	0.2	7.4	10.7	2.41	0.6	0.31	0.98	99.9
S37E1	V2		38.4	0.59	5.0	9.3	0.15	32.4	3.1	0.1	0.04	0.07	10.07	100.3
S38E10	V2		44.3	1.37	12.5	11.0	0.19	12.2	10.3	1.68	0.21	0.13	2.67	97.7
S38E6	V2		48.6	1.18	14.5	9.0	0.15	7.8	11.1	1.88	0.19	0.13	5.06	100.6
S39E1	V2		49.5	1.35	14.1	9.1	0.15	6.9	9.4	3.28	0.65	0.16	4	99.6
S41E1	V2		45.5	1.09	12.0	9.9	0.17	13.5	9.0	1.57	0.24	0.11	4.08	98.3
S41E3	V2		49.9	1.32	14.1	9.2	0.16	7.3	10.1	2.1	0.41	0.15	2.64	98.4
S42E1	V2		46.8	1.24	12.8	10.9	0.18	12.3	9.9	1.74	0.22	0.13	3.33	100.7
S42E2	V2		50.4	1.36	14.3	9.5	0.17	7.7	10.6	2.4	0.25	0.15	2.32	100.2
S42E4	V2		47.8	1.21	14.1	9.6	0.19	9.1	10.0	2.8	0.42	0.13	3.1	98.0
S42E5	V2		50.6	1.38	14.4	9.7	0.17	7.8	10.3	2.67	0.42	0.16	1.89	100.5
S43E1	V2		43.2	1.24	11.7	11.3	0.18	13.9	9.5	1.67	0.23	0.13	6.72	101.0
S43E2	V2		42.2	1.2	11.3	11.0	0.16	12.9	8.9	1.43	0.12	0.12	7.91	98.4
S43E4	V2		42.5	1.6	12.5	11.3	0.2	10.7	10.6	1.94	0.08	0.16	7.61	100.6
S43E5	V2		42.8	1.3	12.7	9.9	0.2	11.5	10.5	1.81	0.13	0.13	7.01	99.0
S43E6	V2		44.3	1.53	14.1	11.2	0.2	9.0	11.3	1.98	0.49	0.14	5.46	100.9
S44E10	V2		43.7	1.11	9.1	11.3	0.18	22.3	7.4	1.39	0.11	0.15	3.2	101.1
S44E2	V2		43.6	1.38	9.5	11.3	0.17	17.0	7.6	1.7	0.22	0.15	5.05	99.0
S44E4	V2		43.1	1.94	12.3	11.9	0.2	8.4	10.8	2.45	0.16	0.22	5.63	98.5
S44E7	V2		44.2	1.7	12.8	11.3	0.19	9.4	10.7	2.54	0.07	0.2	6	100.3
S44E8	V2		44.3	1.12	10.1	10.8	0.16	20.3	8.1	1.46	0.09	0.14	2.74	100.6
S44E9	V2		43.1	1.12	9.1	11.3	0.18	21.4	7.3	1.43	0.14	0.15	2.1	98.5
S45E10	V2		44.2	1.74	12.3	12.0	0.18	9.9	10.0	2.09	0.3	0.19	5.97	100.2
S45E4	V2		43.0	0.96	9.7	10.8	0.18	22.2	7.8	1.22	0.08	0.09	2.69	100.0
S45E9	V2		45.1	1.69	12.0	11.4	0.18	11.0	10.1	2.11	0.27	0.18	4.39	99.8
S2E2	V3		43.3	1.54	11.4	10.8	0.17	13.1	9.0	2.13	0.52	0.15	5.82	99.0
S2E3	V3		45.7	2.28	13.5	10.8	0.19	7.0	10.7	3.31	0.25	0.24	2.83	98.0
S2E5	V3		46.2	1.92	13.2	11.1	0.18	9.4	10.7	2.4	0.57	0.2	2.11	99.3
S2E7	V3		47.9	3.97	12.7	14.6	0.23	5.0	9.4	2.63	1.02	0.52	0.34	99.9
S2E8	V3		48.7	3.09	13.7	12.9	0.21	4.9	9.9	3.05	0.91	0.4	0.98	100.2
S2E9	V3		46.4	2.7	13.6	12.3	0.22	6.7	10.3	2.87	0.28	0.25	2.17	99.1
S8E10	V3		44.9	1.36	13.0	11.0	0.18	10.8	10.4	2.1	0.13	0.13	4.3	99.6
S8E13	V3		48.0	1.9	13.5	12.0	0.2	6.7	11.7	2.35	0.24	0.18	0.6	98.7
S8E6	V3		46.2	1.68	14.5	11.4	0.19	7.7	11.8	2.03	0.22	0.18	3.88	101.0
S8E9	V3		43.3	3.72	12.7	14.4	0.21	4.8	8.9	2.39	1.04	0.47	4.71	98.3
S9E1	V3		48.1	2.27	14.0	12.8	0.22	5.6	11.1	2.56	0.33	0.23	0.8	99.4
S11E10	V3		48.4	1.79	13.9	10.0	0.18	6.2	11.6	2.46	0.19	0.17	1.83	97.9
S11E12	V3		47.5	1.83	14.2	10.7	0.17	6.9	12.0	2.26	0.19	0.17	2.4	99.5
S11E7	V3		48.7	1.72	14.0	11.6	0.19	6.6	11.9	2.64	0.16	0.16	2.61	101.7
S26H1	V3		48.8	2.13	14.3	12.5	0.21	6.0	11.5	2.49	0.3	0.22	0.21	99.8
S26H3	V3		47.9	2.58	13.2	12.5	0.22	5.8	10.9	2.41	0.32	0.28	3.09	100.5
S33E1	V3		44.5	1.23	11.7	10.4	0.17	14.3	9.2	2.12	0.79	0.12	5.25	100.8
S33E12	V3		44.3	1.31	13.5	10.7	0.17	10.7	11.0	1.92	0.18	0.12	5.02	100.1
S33E4	V3		41.7	0.85	8.3	11.5	0.18	22.6	6.9	1.49	0.3	0.1	4.69	99.9
S33E7	V3		44.7	1.23	11.2	11.3	0.19	16.4	9.					

Sample name	Fm.	ppm	Li	Be	Sc	Ti	V	Cr	Co	Ni	Cu	Zn	Ga	Ge	Rb	Sr	Y	Zr	Nb	Cs	Ba	La	Ce	Pr	Nd	Sm	Eu	Tb	Dy	Ho	Er	Tm	Yb	Lu	Hf	Ta	Pb	Th	U			
S19E1	V1		4.24	0.42	32.4	7296	263	648	61.3	324	114	77.2	13.4	1.6	1.81	282	19.3	75.5	6.21	0.04	57.4	6.27	157	2.28	11.3	32.7	107	0.62	3.69	3.65	0.72	1.95	0.27	1.84	0.26	2.2	0.41	0.01	1.18	0.6	0.16	
S19E2	V1		6.63	0.64	35.7	7513	288	448	42.3	45.1	41.9	87.3	23.4	1.74	2.12	238	21.6	78.6	90.8	8.68	163	13.6	30	3.89	17.4	41.6	133	0.67	4.13	0.67	0.43	0.77	0.19	2.3	0.3	0.07	0.28	1.28	1.64	0.25		
S19E3	V1		6.49	0.72	34.4	7708	285	401	38.9	33.1	35.7	68.9	20.3	1.59	2.34	231	21.6	78.6	90.8	8.68	163	13.6	30	3.89	17.4	41.6	133	0.67	4.13	0.67	0.43	0.77	0.19	2.3	0.3	0.07	0.28	1.28	1.64	0.25		
S38E8	V1		10.3	0.69	34.2	7682	283	422	39.3	30.9	25.5	76	18.7	1.56	3.98	250	18.9	85	6.61	0.26	146	19.9	21.2	27	12.5	32.8	111	0.59	3.65	0.42	0.65	1.84	0.56	1.56	0.22	2.21	0.32	0.22	2.8	1.45	0.31	
S40E1	V1		5.93	0.8	37	8142	279	292	33.1	16.6	25.1	87.8	20.6	2.03	8.34	410	24.5	114	22.5	0.17	973	517	10.1	15	42.6	6.4	20.9	0.91	6.51	0.96	0.96	2.56	0.36	2.29	0.33	3.26	0.41	0.24	8.5	3.18	0.32	
S40E2	V1		7.4	0.78	37.4	8142	272	297	33.2	18.5	25.5	87.1	20.4	1.87	4.07	395	24.2	112	21.5	0.04	986	476	94.1	10.5	39.4	6.3	19.9	0.86	5.9	0.47	0.49	2.49	0.36	2.18	0.32	3.19	0.42	0.18	7.79	3	0.3	
S40E3	V1		6.01	0.82	36.3	8279	276	284	33.7	17.4	25.4	86.4	20.3	1.94	1.15	200	23.9	114	22.5	0.07	944	94.2	15.7	38.1	6.8	19.9	0.86	5.9	0.47	0.49	2.49	0.36	2.18	0.32	3.19	0.42	0.18	7.79	3	0.3		
S40E5	V1		5.13	1.21	39.4	18260	402	98.8	41.4	77.3	23.4	111	23.4	20.1	19.4	390	31.5	111	25.4	0.23	180	21.3	50.6	6.62	30	6.83	21.3	7.78	5.81	1.09	2.92	0.4	2.5	0.32	0.56	1.38	0.37	2.07	2.14	0.6		
S40E6	V1		9.82	0.79	36.5	7910	276	270	31.5	12.7	22.1	81.3	20.3	1.91	1.53	391	23.7	112	21.1	0.36	1072	482	97.5	107	39.9	5.88	1.86	0.81	5.52	0.46	0.83	2.31	0.32	1.98	0.3	2.97	0.38	1.76	2.8	0.25		
S1E1	V2				31.81			304.93	811.76	62.86	458.99							210.39	19.53	82.26	4.88																		0.35			
S1E2	V2				23.13			1226.42	88.54	1119.58								64.69	13.01	80.23	2.16																			1.58	1.13	
S1E3	V2		2.97	0.38	28.8	5935	221	142.2	6.4	946	84.7	73	13.4	1.11	2.08	105	13.																						0.54	0.17	0.05	
S1E4	V2		35.78		339.56		699.32	57.1	312.05									205.06	21.58	82.65	4.22																			0.41		
S1E5	V2		2.9	0.48	0.62	110202	272	392	46.1	55	42.5	82.1	1.81	1.58	5.44	320	14.5	85.1	14.7	0.07	176	18.4	36.8	45.2	19.6	38.1	15.3	0.61	3.97	3.31	0.63	1.69	0.23	1.68	0.3	2.19	0.23	0.83	2.05	1.14	0.33	
S1E6	V2		36.13		342.79		939.46	46.71	375.77									186.68	21.05	79.51	4.25																			1.58	1.13	
S3E1	V2		4.54	1.36	31.6	22770	413.69	36.46	46.17	40.5	24.1	44.9	22.9	2.39	2.52	144.47	48.62	298.09	45.43	0.28	331.94																			0.92		
S4E1	V2		7.95	0.72	42.5	10140	332	506	55.8	191	167	80.9	19	1.26	6.1	255	24	91.5	47.2	0.03	42.9	5.2	14.2	23.4	12.1	3.61	1.28	0.71	4.3	4.33	0.85	2.31	0.83	0.29	1.28	0.29	0.29	0.69	0.33	0.1		
S5E11	V2		3.72	1.32	26.9	6066	185	1124	51.3	279	43	67.7	17.2	1.04	2.63	157	17.7	120	71.6	0.57	233	15.7	33.3	4.1	16.6	3.57	1.05	0.67	3.62	3.43	0.67	1.83	0.27	1.64	0.24	3.45	0.41			6.62	3.83	0.89
S5E10	V2		6.29	0.81	40.4	8681	301	884	50.4	66.3	55.2	90.2	18.8	1.7	10.5	214	22.3	113	82.7	0.13	141	11.1	24.4	35.1	15.6	38.2	11.5	0.65	4.06	4.1	0.78	2.24	0.31	2.15	0.3	2.84	0.44	1.54	2.62	0.42		
S5E6	V2		9.55	1.09	43	9918	310	1210	28.4	44.9	53.3	76.9	20.4	1.05	1.71	234	23.7	112	97.5	0.16	229	12.1	27.5	38.5	14.7	41.7	13.8	0.74	4.49	4.37	0.85	2.3	0.33	1.98	0.29	3.04	0.48			2.22	1.69	0.41
S6E10	V2		4.28	0.85	30.1	19390	342	51	44.8	47.8	93.7	81.7	18.4	2.24	10.7	339	22.5	95	22.1	0.03	310	17.3	39.4	59.6	25.7	5.97	2.27	0.9	6.16	4.73	0.88	2.25	0.69	1.63	0.25	1.29	0.13	1.05	1.45	0.39		
S6E11	V2		0.402	0.32	24.9	5458	206	1076	17.5	814	83.6	70.7	10.7	1.57	3.99	139	13.3	60.4	7.95	0.1	303	8.62	15.9	2.44	11.6	30.3	1.05	0.5	3.16	3.03	0.58	1.61	0.23	1.54	0.2	2.02	0.62	0.03	0.74	0.75	0.2	
S6E6	V2		2.99	0.24	24	5380	212	1231	78.8	907	88	74.3	12.2	1.57	5	109	142	51.2	29.6	0.05	203	3.23	85.9	14.9	18.2	25.3	0.89	4.9	3.14	0.61	1.7	0.24	1.68	0.23	1.8	0.23	0.42	0.29	0.08			
S6E9	V2		4.5	0.42	30.6	6574	216	1247	84.6	869	105	68.7	13.3	1.1	1.18	147	13.7	67.1	83.2	0.14	479	656	15.3	12.2	12.9	37.7	0.94	4.89	2.84	2.86	0.57	1.55	0.23	1.3	0.19	2.01	0.57			3.73	0.61	0.19
S7E10	V2		2.18	0.36	25.6	6455	203	711	48.2	31.2	45.9	36.7	12.2	1.33	0.51	169	15.5	5.67	0.01	281	5.6	14.2	32.5	15.1	5.09	1.66	0.97	6.06	1.2	3.46	0.69	1.2	0.46	0.22	0.43	0.22	0.38	0.1	0.05	0.11		
S7E11	V2		3.88	0.36	28.4	5723	204	1022	91.3	106.1	67.6	60.4	10.7	1.454	8.57	193	14.7	11.7	86.6	2.38	0.05	4.86	2.74	2.81	0.68	2.74	2.81	0.68	2.74	2.81	0.68	2.74	2.81	0.68	2.74	2.81	0.68	2.74	2.81	0.68	2.74	
S7E4	V2		3.62	0.45	34.8	8725	305	983	73.2	454	97.5	90.8	16.1	1.73	0.83	196	20	77.8	5.98	35.9	60.1	14.8	2.35	12	3.73	11.8	0.86	4	3.84	0.72	2.01	0.28	1.86	0.26	2.33	0.39	0.01	0.53	0.47	0.13		
S7E5	V2		3.48	0.26	30.6	4957	174	988	76.1	116.1	72.9	51.9	7.66	1.39	3.26	72	11.1	51.7	47.7	0.06	186	45.1	108	18.1	9.15	25.8	0.86	4.05	2.78	2.66	0.51	1.45	0.32	1.31	0.18	1.84	0.34	0.17	0.57	0.45	0.11	
S7E6	V2		3.36	0.42	31.5	8064	305	983	73.2	454	97.5	90.8	16.1	1.73	0.83	196	20	77.8	5.98	35.9	60.1	14.8	2.35	12	3.73	11.8	0.86	4	3.84	0.72	2.01	0.28	1.86	0.26	2.33	0.39	0.01	0.53	0.47	0.13		
S7E7	V2		5.1	0.48	25.4	5939	189	2025	88.9	112.3	82.8	61.9	12.9	0.98	4.34	82.8	12.7	58.6	4.7	0.07	14.9	7.5	11.6	11.83	8.7	23.7	0.81	4.64	2.66	2.62	0.51	1.36	0.2	1.17	1.77	0.3			0.61	0.41	0.12	
S7E8	V2		5.13	1.58	29.9	23180	378	51	44.5	44.6	30.4	15.2	2.33	2.38	3.35	342	46.2	29.5	52.8	0.29	377	35.8	76.9	49.9	42.2	94.7	0.304	1.54	6.99	6.87	1.65	4.42	0.62	3.81	0.53	0.67	2.85	3.31	3.65	1.03		
S7E9	V2		5.96	0.73	30.2	12460	480	1095	96.6	529	201	90.3	27.9	1.73	3.68	343	36.8	131	13.5	0.04	152	15.5	34.7	44.8	22.5	5.54	1.97	0.65	6.01	3.56	1.67	1.36	0.38	0.52	3.35	3.99	0.77	0.05	0.26	1.29	0.3	
S15H4	V2		2.22	0.19	23.1	4291	186	1501	83.6	1026	80.7	51	11.45	2.85	1	100	12.1	38.6	14.6	0.01	287	1.81	5.1	5.77	1.97	10.9	0.4	2.38	2.61	0.52	1.49	0.22	1.4	0.19	1.35	0.12	0.03	0.43	0.14	0.04		
S4E1	V2		3.13	0.79	37.5	12700	411	388	41.8	46.1	260	90.9	23.1	1.91	2.41	267	23.9	114	22.5	0.07	944	94.2	15.7	38.1	6.8	19.9	0.86	5.9	0.47	0.49	2.49	0.36	2.18	0.32	3.19	0.42	0.18	7.79	3	0.3		
S17H2	V2		3.74	0.77	36	12700	410	436	40.1	43.7	263	90.6	22.5	2.01	4.2	218	35.9	124	78.9	0.05	53.9	8.77	20.7	15.1	48.1	19.6	0.91	5.98	1.25	3.25	0.46	1.02	0.44	3.38	0.42	0.06	1.33	0.85	0.24			
S20H4	V2		3.22	0.55	36.6	10440	369	564	51.4	306	148	72.1	19.5	1.76	2.12	250	22.7	88.8	5.43		39.4	5.47	23.3	12.54	12.8	3.97																

Sample names	Fm.	²⁰⁸ Pb/ ²⁰⁴ Pb	²⁰⁷ Pb/ ²⁰⁴ Pb	²⁰⁶ Pb/ ²⁰⁴ Pb	¹⁷⁶ Hf/ ¹⁷⁷ Hf	2 σ	ε _{Hf}	¹⁴³ Nd/ ¹⁴⁴ Nd	2 σ	ε _{Nd}	⁸⁷ Sr/ ⁸⁶ Sr
S19E1	V1	37.7907	15.3018	17.7224		0.0000073					
S19E2	V1	37.1751	15.2492	16.9346	0.282636	0.0000048	-4.82				
S38E2	V1	36.9612	15.157	16.6579	0.282607	0.0000042	-5.84				0.70780
S38E8	V1	36.7908	15.1218	16.5386	0.282612	0.0000042	-5.65	0.512302	0.0000063	-6.56	0.70807
S40E1	V1	38.7418	15.4345	18.6374				0.512350	0.0000047	-5.62	0.70636
S40E2	V1	38.7234	15.4236	18.6007				0.511881	0.0000088	-14.76	0.70628
S1E1	V2	37.7479	15.3743	17.9969	0.283198	0.0000041	15.06				0.70338
S1E3	V2	37.8174	15.4329	18.0711	0.283198	0.0000034	15.08				0.70338
S1E4	V2	37.7933	15.3716	18.0389	0.283193	0.0000004	14.89				0.70332
S3E1	V2	38.5017	15.489	18.7265							0.70362
S4E1	V2	37.714	15.4216	18.0089	0.283203	0.0000079	15.23				0.70311
S5E11	V2	37.4929	15.1063	16.915		0.0000028		0.512345	0.0000047	-5.71	
S5E5	V2	37.2793	15.0701	16.9389	0.282652	0.0000036	-4.26				0.70679
S5E6	V2	37.3562	15.0888	17.0592	0.282692	0.0000003	-2.84	0.512506	0.0000044	-2.57	0.70652
S6E6	V2	37.7648	15.4218	18.037	0.283157	0.0000048	13.61				0.70330
S6E9	V2	38.0612	15.4362	18.2839	0.283138	0.0000049	12.94				0.70628
S7E6	V2	37.9231	15.443	18.098	0.283177	0.0000077	14.31				0.70338
S7E7	V2	37.9827	15.4257	18.1173	0.283173	0.0000036	14.19				0.70349
S7E9	V2	38.0256	15.3296	17.9856	0.283193	0.0000005	14.9				0.70363
S15H4	V2	37.5416	15.3724	17.9168	0.283220	0.0000034	15.85				0.70309
S20E15	V2	37.811	15.3603	17.9032	0.283104	0.0000043	11.74				0.70369
S20E7	V2	38.4594	15.3964	18.4259	0.283059	0.0000033	10.16				0.70398
S27E12	V2	37.7582	15.4339	18.099	0.283202	0.0000029	15.21				0.70794
S27E13	V2	37.6621	15.4177	18.0158	0.283211	0.0000044	15.52				0.70315
S27E18	V2	37.8792	15.3751	18.0833	0.283170	0.0000046	14.09				0.70335
S38E10	V2	38.0322	15.3567	18.0635	0.283115	0.0000034	12.12				0.70374
S39E1	V2	37.1137	15.0627	16.4893	0.282501	0.0000046	-9.58	0.512220	0.0000042	-8.15	0.70624
S41E3	V2	36.8255	14.9913	16.1358	0.282531	0.0000048	-8.51				0.70704
S42E2	V2	36.8122	15.0013	16.1661	0.282565	0.0000043	-7.31				0.70665
S42E5	V2	36.8	15.0014	16.1686	0.282574	0.0000027	-7.01	0.512303	0.0000006	-6.54	0.70673
S43E1	V2	37.9194	15.3688	18.0556	0.283116	0.0000061	12.15				0.70355
S43E4	V2	37.8974	15.4283	18.0467	0.283139	0.0000104	12.97	0.512992	0.0000052	6.9	0.70357
S43E6	V2	37.7803	15.3919	17.9208	0.283175	0.0000005	14.25				0.70337
S44E2	V2	37.7681	15.4128	18.0671	0.283165	0.0000053	13.89				0.70330
S44E7	V2	37.9074	15.4131	18.0767	0.283176	0.0000073	14.29	0.513050	0.0000083	8.05	0.70356
S44E8	V2	37.5621	15.3715	17.9248	0.283204	0.0000053	15.26				0.70328
S45E4	V2	37.9674	15.4621	18.1573	0.283230	0.0000005	16.18				0.70326
S45E9	V2	37.8374	15.2977	17.7873	0.283065	0.0000109	10.35	0.512883	0.0000061	4.78	0.70391
S2E2	V3	37.8894	15.4361	18.2036	0.283193	0.0000055	14.88				0.70338
S2E3	V3	38.042	15.4536	18.3113							0.70335
S2E5	V3	37.9368	15.4521	18.202							0.70331
S2E7	V3	38.5979	15.4975	18.94	0.283124	0.0000063	12.46	0.512977	0.0000069	6.62	0.70377
S2E8	V3	38.5267	15.4941	18.808	0.283155	0.0000048	13.56	0.512989	0.0000006	6.84	0.70366
S2E9	V3	37.9398	15.4554	18.2417	0.283172	0.0000045	14.14	0.513054	0.0000044	8.12	0.70822
S8E13	V3	37.9907	15.4015	18.2182	0.283174	0.0000046	14.22	0.513033	0.0000009	7.67	0.70353
S8E6	V3	39.0141	15.5255	19.112	0.283083	0.0000046	11	0.512719	0.0000065	1.54	0.70392
S9E1	V3	38.1598	15.4592	18.4869	0.283191	0.0000051	14.82				
S11E10	V3	38.0766	15.4443	18.4272	0.283203	0.0000004	15.23				0.70348
S26H1	V3	38.2176	15.5662	18.2994	0.283198	0.0000036	15.05				
S26H3	V3	37.9707	15.4411	18.3145	0.283205	0.0000043	15.3	0.513094	0.0000077	8.89	0.70345
S33E12	V3	38.4105	15.4844	18.6862	0.283192	0.0000074	14.86				
S33E7	V3	38.7609	15.4922	19.075	0.283111	0.0000074	11.97				
S35E11	V3				0.283171	0.0000005	14.1	0.513027	0.0000122	7.54	0.70354
S35E12	V3	38.0041	15.453	18.392							0.70336
S35E3	V3	37.9532	15.4402	18.2024	0.283188	0.0000049	14.7				
S35E5	V3	38.0145	15.4499	18.2868	0.283182	0.0000048	14.49	0.513055	0.0000098	8.13	0.70363
S35E8	V3	37.9179	15.4556	18.2058							0.70330
S36E5	V3	37.9807	15.4298	18.3146	0.283186	0.0000046	14.64				
S21H1	V4	38.3919	15.4636	18.6247	0.283150	0.0000035	13.36	0.512990	0.0000084	6.87	0.70367
S21H2	V4	38.0081	15.4547	18.2627	0.283223	0.0000039	15.95				
S29E2	V4	38.3614	15.4604	18.4962	0.283182	0.0000045	14.49				
S30E1	V4	38.4071	15.4795	18.6598	0.283145	0.0000023	13.19	0.512992	0.0000053	6.91	0.70806
S30E4	V4	38.4888	15.5065	18.8252	0.283146	0.0000025	13.23	0.512984	0.0000061	6.76	0.70376
S13H5	dike	37.8156	15.3998	18.1175							0.70332
S14H2	dike	38.5245	15.5106	18.765	0.283142	0.0000047	13.09	0.512985	0.0000009	6.73	0.70364
S15H3	dike	38.4756	15.4871	18.7332	0.283160	0.0000038	13.74				0.70364
S20E18	dike	38.5508	15.4971	18.7477	0.283154	0.0000057	13.52	0.512984	0.0000051	6.71	0.70379
S22H2	sill	37.8774	15.4337	18.2553	0.283213	0.0000047	15.6	0.513095	0.0000171	8.87	
S22H4	dike	38.5058	15.4949	18.7357	0.283153	0.0000048	13.48	0.512984	0.0000051	6.71	0.70611
S23H4	dike V3	38.9386	15.5336	19.2555	0.283128	0.0000052	12.6	0.512822	0.0000066	3.56	0.70380
S23H5	dike	38.812	15.4818	19.1238	0.283153	0.0000045	13.47	0.512909	0.0000007	5.24	0.70370
S24H1	Sill Itsako	37.8957	15.3504	18.0561	0.283164	0.0000037	13.85				
S27E4	dike	38.4799	15.4969	18.8261	0.283130	0.0000043	12.65	0.512994	0.0000071	6.91	0.70375