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Formation age and metamorphic history of the Nuvvuagittuq Greenstone Belt

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The Nuvvuagittuq Greenstone Belt (NGB) in Northern Quebec, Canada, is dominated by mafic and ultramafic rocks metamorphosed to at least upper amphibolite facies. Rare felsic intrusive rocks provide zircon ages of up to ~3.8 Ga (David et al., 2009; Cates and Mojzsis, 2007) establishing the minimum formation age of the NGB as Eoarchean. Primary U-rich minerals that may provide reliable formation ages for the dominant mafic lithology, called the Ujaraaluk unit, have yet to be found. Metamorphic zircons, rutiles and monazites are present in the unit and give variably discordant results with ²⁰⁷Pb/²⁰⁶Pb ages ranging from 2.8 Ga to 2.5 Ga. The younger ages overlap 2686 ± 4 Ma zircon ages for intruding pegmatites (David et al., 2009) and Sm-Nd ages for garnet formation in the Ujaraaluk rocks suggesting this era as the time of peak metamorphism and metasomatism in the NGB, coeval with regional metamorphism of the Superior craton. Sm-Nd data for Ujaraaluk whole rocks scatter about a Sm/Nd vs. ¹⁴³Nd/¹⁴⁴Nd correlation (MSWD = 134) whose slope would correspond to 3.6 ± 0.2 Ga if interpreted as an isochron. This "isochron" is seen to consist of a series of younger \sim 3.2–2.5 Ga slopes for the different geochemical groups within the Ujaraaluk, emanating from a baseline distribution older than 4 Ga. The ¹⁴⁶Sm-¹⁴²Nd chronometer is less affected by metamorphism at 2.7Ga because of 146 Sm extinction prior to ${\sim}4$ Ga. Expansion of the ¹⁴²Nd dataset for the Ujaraaluk and associated ultramafic rocks continues to show a good correlation between Sm/Nd ratio and 142 Nd/ 144 Nd that corresponds to an age of 4388 $^{+15}_{-17}$ Ma. The dataset now includes samples with superchondritic Sm/Nd ratios that extend the correlation to ¹⁴²Nd excesses of up to 8 ppm compared to the terrestrial standard with a total range in ¹⁴²Nd/¹⁴⁴Nd of 26 ppm. The upper Sm/Nd ratio end of the Ujaraaluk correlation is defined by rocks that are interpreted to be cumulates to compositionally related extrusive rocks indicating that this crystal fractionation had to occur while ¹⁴⁶Sm decay was active, i.e. well before 4 Ga. Intruding gabbros give ¹⁴³Nd and ¹⁴²Nd isochron ages of respectively 4115 \pm 100 Ma and 4313 ⁺⁴¹₋₆₉ Ma, also supporting an Hadean age for the gabbros and providing a minimum age for the intruded Ujaraaluk unit. 3.6 Ga tonalites surrounding the NGB, 3.8 Ga trondhjemitic intrusive veins, and a 2.7 Ga pegmatite show a deficit in ¹⁴²Nd compared to the terrestrial standard. These felsic rocks plot to the low Sm/Nd ratio side of the Ujaraaluk isochron and do not show a correlation between their Sm/Nd and ¹⁴²Nd/¹⁴⁴Nd ratios, which can be explained if they are melts of ancient LREE-enriched mafic rocks, such as the Ujaraaluk, with the melting occurring after ¹⁴⁶Sm was extinct. A subset of least disturbed Ujaraaluk samples has coherent isotopic compositions for both short-lived and long-lived Nd isotopic systems giving 143 Nd and 142 Nd isochron ages overlapping within error of 4321 ± 160 Ma (MSWD = 6.3) and 4406 $^{+14}$ –17 Ma (MSWD = 1.0), respectively. This age represents our best age estimate for the Uiaraaluk unit. The NGB thus preserves over 1.6 billion years of early Earth history including an expanse of mafic crust formed in the Hadean.

1. Introduction

The first billion years of Earth history is poorly presented in the rock record with most early terrains being Eoarchean in age, leaving the Hadean Eon almost devoid of available samples. Until recently, the only available samples older than 4.1 billion years old (Ga) were detrital zircons from the Jack Hills conglomerate (\sim 4.4 Ga; Wilde et al., 2001). Although these zircons provide invaluable information about the early Earth, their host rocks

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have been destroyed, leaving no direct samples of Earth's primordial crust. As a result, the nature of Earth's early crust and the processes responsible for its formation are largely conjecture. Other than these detrital zircons, rare occurrences of Eoarchean crust provide the only compositional constraints on the nature of Earth's early crust. These terrains include the Acasta Gneiss, Canada (4.03 Ga; Bowring and Williams, 1999), the Itsag Gneiss Complex, Greenland (3.85-3.60 Ga; Nutman et al., 1996; Nutman and Friend, 2009), the Napier complex, Antarctica (3.95-3.8 Ga; Williams et al., 1986; Simon and Nigel, 2007), the Saglek-Hebron block, Labrador (3.78-3.73 Ga; Collerson, 1983; Schiotte et al., 1989), and the Anshan area, China (3.8 Ga; Song et al., 1996; Liu et al., 2008; Wu et al., 2008, 2009; Nutman et al., 2009). Earth's primitive crust is most likely to have been derived from the partial melting of the mantle to produce an early mafic crust. This is consistent with the Lu-Hf isotopic compositions of the Jack Hills zircons that suggest that their host rock was tonalitetrondhjemite-granodiorite (TTG) derived from the melting of a basaltic Hadean precursor (Blichert-Toft and Albarede, 2008; Kemp et al., 2010).

Investigation of Earth's primitive crust and the timing of its formation have largely focused on felsic rocks because they are the most likely host rocks for the zircons that provide robust geochronological constraints. These felsic rocks, however, cannot have been directly produced by melting of mantle peridotite. Instead, they likely were produced by melting of an older mafic precursor. Using zircon ages from felsic rocks to provide geochronological constraints on suites of mafic rocks requires a very accurate understanding of the geologic relationships between the different lithologies. Stratigraphic relationships between felsic and mafic units are rarely unequivocal in early Archean terrains that generally have been affected by multiple phases of intense deformation and metamorphism. This emphasizes the need to focus on the mafic component in Eoarchean/Hadean terrains in order to constrain the composition and the timing of formation of Earth's earliest crust. Unfortunately, obtaining accurate ages on old terrestrial mafic rocks is difficult due to their lack of zircon, the common recrystallization of their original igneous mineralogy during later metamorphic events, and their generally low concentrations of the incompatible-element-based radiometric systems used for dating. While long-lived isotopic systems such as ¹⁴⁷Sm-¹⁴³Nd, ¹⁷⁶Lu-¹⁷⁶Hf and ¹⁸⁷Re-¹⁸⁷Os are powerful tools to study the evolution of the crust-mantle system through time, Eoarchean/Hadean terrains usually have suffered extensive metamorphism and deformation that can affect these geochronometers through metamorphic/metasomatic redistribution of parent and/or daughter element any time after rock formation. On the other hand, the short-lived ¹⁴⁶Sm-¹⁴²Nd isotopic system (half life = 68 Ma; Kinoshita et al., 2012) is not as susceptible to partial resetting because 146 Sm became extinct prior to ${\sim}4$ Ga. Thus the ¹⁴⁶Sm-¹⁴²Nd system is an ideal tool to probe any potential Hadean terrain. ¹⁴²Nd deficits recently have been identified in the Nuvvuagittug Greenstone Belt (NGB) mafic rocks (O'Neil et al., 2008) suggesting an age of nearly 4.3 Ga, which would make it the only known remnant of Hadean crust preserved on Earth. However, the oldest U-Pb age obtained in zircon from rare intrusive felsic rocks in the NGB is 3817 ± 16 Ma (David et al., 2009). Nevertheless, the NGB offers a unique opportunity to constrain the composition of Earth's early crust and the processes responsible for the formation of crust in the Eoarchean/Hadean.

In this paper, we present an isotopic study combining both Nd long-lived and short-lived systems (147 Sm $^{-143}$ Nd, $T_{1/2}$ = 106 Ga and 146 Sm $^{-142}$ Nd, $T_{1/2}$ = 68 Ma) focusing on the mafic and ultramafic rocks of the NGB to put as many geochronological constraints as possible on the formation and evolution of the main lithology in the belt.

2. Geologic setting

The Nuvvuagittug Greenstone Belt is located in Northern Quebec, Canada within the Hudson Bay terrane of the Northeastern Superior Province (Fig. 1). The Hudson Bay terrane is a TTGgreenstone terrain subdivided into different domains (Tikkerutuk, Bienville, Goudalie, La Grande, southern Douglas Harbour domains) and characterized by Nd model ages generally older than 3.0 Ga (Boily et al., 2009). The old rocks of the terrane are intruded by a series of younger TTG (~2.7 Ga) with low initial ¹⁴³Nd/¹⁴⁴Nd suggesting recycling of older crust (Boily et al., 2009). The Hudson Bay terrane is interpreted to represent a cratonic nucleus of the Northeast Superior Province onto which the more juvenile rocks of the Rivière Arnaud terrane were juxtaposed. The NGB is a $\sim 10 \text{ km}^2$ volcano-sedimentary sequence preserved on the East shore of Hudson Bay approximately 35 km south of the Inuit municipality of Inukjuak. The NGB is dominantly composed of mafic to ultramafic rocks with chemical sedimentary rocks and minor felsic components, all highly deformed and metamorphosed. The map pattern of the NGB (Fig. 1) is dominated by a large-scale North-closing synform plunging moderately to the South. The West limb of this open fold consists of a tight isoclinal South-closing synform with a steeply E-plunging fold axis. The NGB has been metamorphosed to at least upper amphibolite facies. Cates and Mojzsis (2009) proposed that the NGB experienced at least two significant metamorphic episodes around 3.6 Ga and 2.7 Ga based on overgrowth of zircons, with the Neoarchean event reaching 640 °C.

The NGB is dominated by mafic rocks. The most abundant lithology is a heterogeneous, but generally basaltic, group of rocks mainly composed of variable proportions of cummingtonite-biotite-plagioclase-garnet. Due to its important variation in mineralogy, we refer to this lithological unit as the Ujaraaluk unit (formerly called the faux-amphibolite; O'Neil et al., 2007, 2008) that comprises facies that include both cummingtonite-rich rocks and rocks that are dominated by a garnet-biotite paragenesis. Despite this wide range in mineralogical proportions, the different parageneses of the Ujaraaluk unit show a relatively small range of major element composition from basaltic to basaltic andesite (O'Neil et al., 2011a) reinforcing their grouping as a lithological unit. The Ujaraaluk unit can be subdivided into three distinct geochemical groups mainly based on Al/Ti ratios and trace element profiles. These geochemical groups follow a chemical stratigraphy within the NGB with a high-Ti group that has relatively flat normalized incompatible trace element profiles at the base of the tight synform followed by a low-Ti group stratigraphically above. Chemical sediments lie at the transition between the high-Ti and the low-Ti Ujaraaluk groups. The low-Ti Ujaraaluk unit is further subdivided into two subgroups with distinct Al/Ti ratios and trace element profiles referred as the depleted low-Ti group and the enriched low-Ti group based on their relative concentration of incompatible trace elements (O'Neil et al., 2011a). Rocks from the low-Ti Ujaraaluk unit directly above the chemical sediments have lower concentrations of incompatible trace elements and distinct U-shaped rare earth element (REE) profiles, whereas the rocks from the low-Ti Ujaraaluk at the top of the sequence have the most enriched REE profiles (O'Neil et al., 2011a). The Ujaraaluk unit as a whole is interpreted by O'Neil et al. (2011a) to represent hydrothermally altered mafic volcanic crust transitioning from rocks with tholeitic affinitities to rocks with boninitic and calcalkaline affinities deposited in a submarine volcanic environment. This is supported by the presence of deformed, but well-preserved pillows within the Ujaraaluk unit. The SW corner of the NGB contains massive aphanitic basaltic greenstones mainly composed of chlorite-epidote-quartz-plagioclase-actinolite (O'Neil et al., 2007). These greenstones can be divided into the same three compositional groups seen in the Ujaraaluk unit with the same

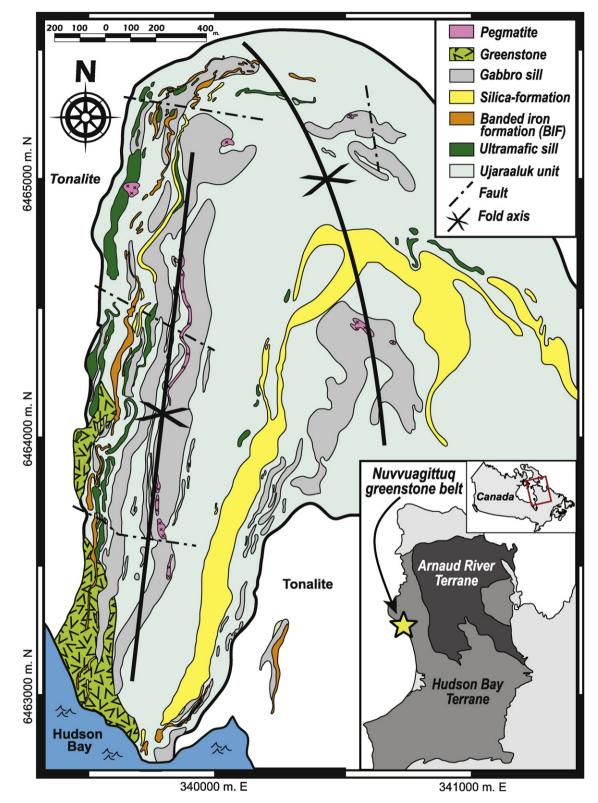


Fig. 1. Geologic map of the NGB. Coordinates are in UTM NAD 27.

geochemical characteristics [Fig. 1 Supplementary Data (SD1)] and following the same stratigraphy within the NGB. They are interpreted to be a lower grade, most likely retrograde, metamorphic equivalent to the Ujaraaluk unit. This interpretation is supported by the presence of these same greenstones to the SE of the NGB where they contain relicts of garnets completely chloriticized that display the same mineral texture observed in the garnet-rich Ujaraaluk rocks.

The Ujaraaluk unit includes ultramafic bodies that can be divided into the same three distinct geochemical groups following the same chemical stratigraphy as the Ujaraaluk mafic rocks (O'Neil et al., 2011a). These ultramafic bodies are interpreted to be ultramafic sills. They mainly consist of serpentine-rich interiors with amphibole-rich margins and locally exhibit amphibole-rich layers within the ultramafic interiors. The composition of the ultramafic rocks follow a well defined olivine-fractionation line and are interpreted to be ultramafic cumulates cogenetic with corresponding mafic Ujaraaluk groups sharing the same geochemical trends and characteristics (O'Neil et al., 2007, 2011a). The Ujaraaluk unit also is intruded by numerous highly deformed gabbroic sills. These mafic intrusions are composed of plagioclase and hornblende. They can reach ~100 m in width and can sometimes be followed continuously over 1 km.

Rare trondhjemitic rocks intruding the southern extremity of the NGB were dated at 3751 ± 10 Ma (Cates and Mojzsis, 2007), 3817 ± 16 Ma (David et al., 2009) and 3774 ± 32 Ma (Darling et al., 2012) indicating that the NGB is at least Eoarchean in age. The NGB is surrounded by a 3661 ± 4 Ga foliated tonalite (David et al., 2009) locally intruding the NGB to the West, itself sheathed by younger (2750 ± 16 Ma) heterogeneous tonalities from the Tikkerutuk domain (Simard et al., 2003). The NGB also is intruded by Neoarchean pegmatitic sheets that provide zircon dates of 2686 ± 4 Ma (David et al., 2009). David et al. (2009) obtained U–Pb ages in zircons from a gabbro associated with the tonalite in the center of the synform and from a plagioclase–quartz–biotite schist. These samples yielded ages of 2693 ± 3 Ma and 3366 ± 2.5 Ma respectively, interpreted to be metamorphic ages.

Zircons from what is interpreted to be a conglomeratic unit in the NGB yield 207 Pb/ 206 Pb ages ranging from 3770 Ma to 3258 Ma (David et al., 2009). These zircons lie along a discordia with an upper intercept of 3787 ± 25 Ma. These conglomeratic units, however, are found only within high-strain zones and always within less than 100 m of the southern NGB-tonalite boundary making their relationship with the adjacent lithologies and exact origin equivocal.

Controversy about the age of the NGB emerged after O'Neil et al. (2008) showed that the Ujaraaluk unit had a deficit in ¹⁴²Nd relative to the terrestrial standard Nd, Unlike most occurrences of variable ¹⁴²Nd in early Earth rocks, the NGB rocks show a correlation between ¹⁴²Nd/¹⁴⁴Nd and ¹⁴⁷Sm/¹⁴⁴Nd that, if interpreted as an isochron yields an age of 4.37 Ga [an age of 4.28 Ga was reported by O'Neil et al. (2008), but all ¹⁴⁶Sm-¹⁴²Nd ages reported in this paper are calculated using the newly determined ¹⁴⁶Sm half life of 68 Ma and recalculated Solar system initial ¹⁴⁶Sm/¹⁴⁴Sm=0.0094 (Kinoshita et al., 2012)]. An Hadean age for the Ujaraaluk unit is supported by a ¹⁴³Nd/¹⁴⁴Nd vs. ¹⁴⁷Sm/¹⁴⁴Nd isochron for intruding gabbros that provides an age of over 4.1 Ga for these intrusive units. While the surrounding and intruding felsic lithologies of the NGB suggest an Eoarchean age, the dominant mafic lithologies yield an Hadean isotopic signature suggesting that the NGB could represent the oldest preserved crust on Earth.

3. Analytical procedures

Weathered surfaces and quartz veins were removed from rock samples prior to crushing. Weathering-free samples were crushed in a steel jaw crusher and then ground to powder in an alumina shatter box. Some samples contain coarse-grained garnet and therefore, to obtain representative whole rock powders, large size samples (15–20 cm) were crushed.

Whole rock samples were digested using a flux-fusion technique. Between 250 and 350 mg of powdered sample was fused in graphite crucibles using ~0.8 g of a mixture of 67% lithium tetraborate-33% lithium metaborate and ~0.02 g of lithium bromide for 20 min at 1050 °C. The molten beads were poured into a Teflon beaker containing 40 ml of 2 M NHO₃ and stirred for 20 min until the glass was completely dissolved. The dissolved samples were filtered to remove flakes of graphite from the crucibles. Aliquots of this solution corresponding to 10% of the dissolved sample were spiked with enriched ¹⁵⁰Nd-¹⁴⁹Sm tracers to measure the Sm and Nd concentrations by isotope dilution. The remaining 90% of the dissolved sample was not spiked and was used for Nd isotope measurements (¹⁴²Nd and ¹⁴³Nd). From both spiked and unspiked aliquots of the dissolution solution, the REE were precipitated by raising the pH of the HNO₃ solution to between 5.5 and 6 by the addition of NH₄OH. An Fe solution prepared from 99.9% anhydrous Aldrich FeCl₂ was previously added to both spiked and unspiked dissolved fractions to facilitate the precipitation. The precipitates were re-dissolved in 2.5 M HCl and loaded onto primary columns $(1 \text{ cm} \times 20 \text{ cm})$ filled with 200-400 mesh AG50W-X8 cation-exchange resin for the REE extraction. Sm and Nd were extracted from this REE fraction using methylactic acid as described in Boyet and Carlson (2005). Both Sm and Nd were extracted from the spiked aliquot with one pass through the methylactic acid columns whereas only the Nd was extracted on the unspiked aliquots but with three passes through the methylactic acid columns in order to remove as much Ce and Sm as possible. Six Ujaraaluk samples also were measured for their unspiked Sm isotopic compositions to check for possible neutron irradiation or meteoritic components in these rocks. All six samples showed Sm isotopic compositions within error of the terrestrial standard (Table SD1). For the samples from barrel B369 and higher (see Table SD2), the Nd fractions for both spiked and unspiked aliquots obtained after methylactic acid columns were then purified by passing the samples through columns filled with 300 mg of Eichrom Ln Resin 50–100 µm. No variation within error of the ¹⁴²Nd/¹⁴⁴Nd ratios has been observed for duplicate samples (Table SD3) that have been purified with Eichrom Ln Resin.

Garnet separates were obtained after sieving of the crushed rock samples to collect a grain fraction between $120 \,\mu\text{m}$ and $75 \,\mu\text{m}$. The grains were washed and garnets were separated from the other grains by magnetic separation. The cleanest garnet grains were then handpicked under a binocular microscope to avoid composite grains. This cleaner garnet fraction was leached following Pollington and Baxter (2010) to remove inclusions and then powdered using an agate mortar/pestle. Garnet samples (~65 mg) were digested in Parr bombs under pressure with a HF–HNO₃ mixture in Teflon containers.

Whole rock samples for Lu–Hf analysis were fused and precipitated using the same procedure as for the Sm–Nd. Samples were spiked with enriched tracers of 176 Lu and 178 Hf. The spike solution was added in the Teflon beaker with 40 ml of 2 M NHO₃ before pouring of the fused bead to assure spike-sample equilibration. Garnet separate samples were spiked for both Sm–Nd and Lu–Hf within the same dissolution. The Hf and Lu were collected and purified following procedures from Carlson et al. (2006).

Sm and Nd isotopic compositions were measured using double Re filaments in the Department of Terrestrial Magnetism (DTM) Thermo-Fisher Triton thermal ionization mass spectrometer. Mass spectrometric procedures followed those described in Carlson et al. (2007). The ¹⁴⁷Sm/¹⁴⁴Nd ratios as well as the Sm and Nd concentrations reported in Table 1 were measured on the spiked aliquots whereas all other reported isotopic data were obtained on the unspiked aliquots. Nd is measured as Nd+ in a two-step dynamic routine that provides static measurements of all Nd isotope ratios and ¹³⁸Ba, ¹⁴⁰Ce, ¹⁴¹Pr, and ¹⁴⁷Sm, and also provides a dynamic measurement for ¹⁴²Nd/¹⁴⁴Nd. Individual runs consist of 540 ratios taken with 8 second integrations per mass step. When the amount of Nd on the filaments allowed, samples were run more than once on the same filament. Samples that were run multiple times from the same filament were combined into one run of 540+ ratios. Data were corrected for instrumental mass fractionation using the exponential law to 146 Nd/ 144 Nd = 0.7219. Total Nd blank for the flux-fusion technique is 2.6 ng. Considering

Table 1

Not isotopic data for the NGB rocks and mineral separates. $\varepsilon Nd_{(i)}$ values were calculated at 4300 Ma for the Ujarraluk and Greenstone, at 4115 Ma for the gabbro, at 3660 Ma for the tonalite, at 3800 Ma for the trodhjemite band and at 2700 Ma for the pegmatite.

Sample	Lithology	[Nd] (ppm)	[Sm] (ppm)	¹⁴⁷ Sm/ ¹⁴⁴ Nd	143 Nd/ 144 Nd $\pm 2\sigma$	$\varepsilon Nd_{(i)}$	$T_{\rm DM}~({\rm Ga})$	142 Nd/ 144 Nd $\pm 2\sigma$	μ^{142} Nd ± 2
High-Ti group	11	10.50	2.01	0 1007	0.510000 + 1	~ •	2.0	1 1 41025 4 + 20	0.0
PC-058	Ujaraaluk	19.56	3.91	0.1207	0.510809 ± 1	6.4	3.8	1.1418354 ± 39	0.0 ± 3.4
PC-129 ^a	Ujaraaluk	8.07	2.10	0.1571	0.511818 ± 2	5.8	3.5	1.1418239 ± 45	-10.1 ± 3.9
PC-132	Ujaraaluk	6.13	1.83	0.1818	0.512386 ± 16	3.2	3.5	1.1418263 ± 66	-8.0 ± 5.3
PC-171	Ujaraaluk	15.34	5.24	0.2073	0.512870 ± 3	-1.6		1.1418386 ± 50	2.8 ± 4.4
PC-180	Greenstone	5.16	1.61	0.1890	0.512572 ± 1	2.8	3.4	1.1418361 ± 58	0.6 ± 5.1
PC-421	Ujaraaluk	4.26	1.41	0.2012	0.512802 ± 2	0.4	4.3	1.1418421 ± 50	5.8 ± 4.4
PC-423	Ujaraaluk	7.16	2.13	0.1810	0.512221 ± 3	0.4	4.3	1.1418317 ± 52	-3.3 ± 4.1
°C-425	Ujaraaluk	4.33	1.11	0.1559	0.512221 ± 3 0.511898 ± 3	8.1	3.2	1.1418289 ± 56	$-5.7 \pm 4.$
PC-429	Ujaraaluk	8.54	2.24	0.1595	0.511765 ± 1	3.5	3.8	1.1418287 ± 37	$-5.9 \pm 3.$
PC-451	Greenstone	11.77	2.75	0.1412	0.511345 ± 1	5.5	3.7	1.1418314 ± 27	-3.6 ± 2.1
PC-220	Ultramafic	4.04	1.46	0.2192	0.513219 ± 5	-1.4	3.0	1.1418422 ± 63	5.9 ± 5.2
PC-018	Ultramafic	1.51	0.61	0.2452	0.513658 ± 8	-7.4	2.7	1.1418445 ± 62	8.0 ± 5.0
PC-033	Ultramafic	3.76	1.28	0.2062	0.512944 ± 1	0.5	4.3	1.1418411 ± 43	5.0 ± 3.0
Depleted low-	Ti group								
PC-131	Ujaraaluk	2.44	0.66	0.1643	0.511875 ± 2	2.9	3.9	1.1418234 ± 58	-10.5 ± 5.0
PC-151	Ujaraaluk	2.55	0.81	0.1919	0.512433 ± 2	-1.6			
PC-184	Greenstone	2.42	0.63	0.1572	0.511848 ± 1	6.4	3.4	1.1418211 ± 54	$-12.6 \pm 4.$
PC-222	Ujaraaluk	2.70	0.81	0.1825	0.512216 ± 2	-0.6	5.4	1.1418264 ± 46	$-7.9 \pm 4.$
PC-227	Ujaraaluk	2.29	0.81	0.2144	0.512753 ± 1	-7.9	4.5	1.1418372 ± 39	$1.5 \pm 3.$
PC-230 ^a	Ujaraaluk	2.44	0.72	0.1793	0.512137 ± 14	-0.3	4.5	1.1418272 ± 9	$-7.2 \pm 0.$
PC-410	Ujaraaluk	2.37	0.65	0.1673	0.511927 ± 3	2.3	4.0	1.1418296 ± 63	-5.1 ± 5.1
PC-412	Ujaraaluk	1.62	0.55	0.2059	0.512601 ± 1	-6.2		1.1418371 ± 47	1.5 ± 4
PC-432	Ujaraaluk	2.86	0.78	0.1662	0.511983 ± 2	4.0	3.7	1.1418286 ± 58	-6.0 ± 5.0
PC-433	Ujaraaluk	2.05	0.59	0.1736	0.512110 ± 3	2.3	3.9	1.1418251 ± 82	-9.1 ± 7
C-214	Ultramafic	0.14	0.06	0.2681	0.512110 ± 3 0.514124 ± 16	-11.1	2.8	1.1418419 ± 71	5.7 ± 6
Enriched low-1	Ti groun								
PC-149 ^a	Ujaraaluk	4.75	1.27	0.1618	0.511691 ± 0.3	0.7	4.3	1.1418230 ± 10	-10.9 ± 0.1
PC-152°	Ujaraaluk	9.41	2.08	0.1343	0.511091 ± 0.5	4.3	3.9	1.1418266 ± 5	-7.7 ± 2
	2						5.5		
PC-162	Ujaraaluk	5.17	1.39	0.1626	0.511602 ± 3	-1.5		1.1418313 ± 38	-3.6 ± 3.0
PC-185	Greenstone	8.30	1.95	0.1421	0.511244 ± 1	3.0	4.0	1.1418182 ± 26	-15.1 ± 2.0
PC-186	Greenstone	6.44	1.50	0.1411	0.511166 ± 1	2.0	4.1	1.1418189 ± 66	-14.5 ± 5.0
C-225	Ujaraaluk	8.99	2.17	0.1462	0.511363 ± 5	3.0	4.0	1.1418187 ± 56	-14.7 ± 4
PC-250 ^c	Ujaraaluk	9.36	2.00	0.1297	0.511050 ± 3	6.2	3.8	1.1418200 ± 43	-13.5 ± 3.0
PC-251 ^a	Ujaraaluk	7.95	1.88	0.1429	0.511177 ± 15	1.2	4.2	1.1418185 ± 21	-14.9 ± 1.0
PC-267 ^a	Ujaraaluk	9.50	2.29	0.1458	0.511225 ± 4	0.5	4.3	1.1418194 ± 90	-14.1 ± 7
PC-275°	Ujaraaluk	10.48	2.34	0.1355	0.510976 ± 5	1.4	4.2	1.1418212 ± 77	$-12.5 \pm 6.$
PC-276 ^a	Ujaraaluk	7.55	2.00	0.1602	0.511557 ± 9	-1.0	4.5	1.1418245 ± 34	$-9.6 \pm 3.$
PC-278	Ujaraaluk	9.43	2.17	0.1399	0.511228 ± 3	3.9	3.9	1.1418180 ± 55	$-15.3 \pm 4.$
PC-282 ^{a,d}	Ujaraaluk	9.58	2.45	0.1543	0.511420 ± 0	-0.4	4.4	1.1418216 ± 49	$-12.1 \pm 4.$
PC-438	Ujaraaluk	7.51	1.65	0.1343	0.511089 ± 3	4.3	3.9	1.1418240 ± 55	$-10.0 \pm 4.$
PC-442	Ujaraaluk	4.39	1.12	0.1544	0.511420 ± 3	-0.5	4.4	1.1418225 ± 68	$-11.3 \pm 6.$
PC-443 ^c	Ujaraaluk	3.15	0.95	0.1825	0.512154 ± 2	-1.8		1.1418271 ± 6	-7.3 ± 2.0
PC-452°	Ujaraaluk	9.36	2.05	0.1377	0.511188 ± 3	4.4	3.9	1.1418227 ± 24	-11.1 ± 3
PC-456							3.9		
	Greenstone	9.54	2.17	0.1378	0.511208 ± 1	4.7		1.1418149 ± 59	-18.0 ± 5.0
PC-532	Ujaraaluk	7.98	2.15	0.1631	0.511935 ± 6	4.8	3.6	1.1418317 ± 34	-3.3 ± 3.1
PC-071	Ultramafic	6.42	1.70	0.1603	0.511894 ± 1	5.5	3.5	1.1418340 ± 42	-1.3 ± 3
PC-075	Ultramafic	2.13	0.61	0.1737	0.512191 ± 1	3.9	3.6	1.1418282 ± 26	-6.4 ± 2.00
PC-137	Ultramafic	4.90	1.49	0.1846	0.512351 ± 3	0.9	4.2	1.1418317 ± 69	-3.2 ± 6.0
PC-142	Ultramafic	7.04	1.94	0.1674	0.512060 ± 3	4.9	3.5	1.1418366 ± 38	1.0 ± 3.0
C-213	Ultramafic	2.04	0.69	0.2034	0.512657 ± 1	-3.6		1.1418406 ± 44	4.6 ± 3
C-213 C-218 ^a	Ultramafic	6.91	2.22	0.1945	0.512569 ± 2	-0.3		1.1418335 ± 13	-1.7 ± 1
PC-307	Ultramafic	2.57	0.81	0.1914	0.512422 ± 1	-1.5		1.1418373 ± 57	1.6 ± 5.0
°C-309	Ultramafic	5.25	1.54	0.1779	0.512258 ± 3	2.8	3.7	1.1418386 ± 52	2.8 ± 4
C-503	Ultramafic	2.26	0.69	0.1856	0.512286 ± 1	-0.9		1.1418319 ± 64	-3.1 ± 5
PC-506	Ultramafic	7.36	2.06	0.1689	0.511995 ± 1	2.7	3.9	1.1418357 ± 31	0.3 ± 2
Central gabbro)								
PC-221 ^a	Gabbro	7.12	2.16	0.1835	0.512395 ± 14	2.1	3.7	1.1418337 ± 42	-1.5 ± 3.0
PC-228 ^a	Gabbro	5.46	1.37	0.1515	0.511532 ± 15	2.3	3.9		
C-248 ^{a,d}	Gabbro	7.20	2.27	0.1903	0.512591 ± 11	2.3	3.5	1.1418318 ± 31	-3.1 ± 2
C-253 ^a	Gabbro	8.34	2.67	0.1932	0.512669 ± 10	2.3	3.4		5.1 ± 2
PC-254 ^a	Gabbro	8.90	2.12	0.1442	0.511331 ± 11	2.3	3.9		
PC-255 ^a	Gabbro	7.47	2.34	0.1893	0.512534 ± 23	1.7	3.7		
PC-259 ^a	Gabbro	6.82	2.29	0.2034	0.512909 ± 10	1.5	3.0		
C-268 ^a	Gabbro	6.58	2.08	0.1908	0.512557 ± 11	1.4	3.9		
C-508	Gabbro	10.42	2.55	0.1481	0.511430 ± 5	2.1	3.9	1.1418260 ± 27	-8.3 ± 2
C-511	Gabbro	8.99	2.26	0.1522	0.511450 ± 5 0.511572 ± 5	2.7	3.9	1.1418265 ± 38	-7.8 ± 3
C-512	Gabbro	10.12	2.50	0.1494	0.511444 ± 6	1.7	4.0	1.1418284 ± 34	-6.2 ± 3
PC-522	Gabbro	2.71	0.80	0.1788	0.512181 ± 6	0.4	4.2	1.1418351 ± 49	-0.3 ± 4
PC-523	Gabbro	7.01	1.65	0.1424	0.511233 ± 7	1.3	4.1	1.1418295 ± 60	-5.2 ± 5
C-J2J									

Table 1 (Continued)

Sample	Lithology	[Nd] (ppm)	[Sm] (ppm)	¹⁴⁷ Sm/ ¹⁴⁴ Nd	143 Nd/ 144 Nd $\pm 2\sigma$	$\varepsilon Nd_{(i)}$	$T_{\rm DM}~({\rm Ga})$	142 Nd/ 144 Nd $\pm 2\sigma$	μ^{142} Nd $\pm 2\sigma$
Felsic rocks									
PC-103 ^a	Tonalite	20.81	3.76	0.1092	0.510245 ± 2	-5.6	4.2	1.1418215 ± 61	-12.2 ± 5.3
PC-104	Tonalite	23.37	3.64	0.0941	0.510049 ± 1	-2.2	3.9	1.1418221 ± 37	-11.7 ± 3.3
PC-105	Tonalite	20.02	3.34	0.1009	0.510199 ± 1	-2.5	3.9	1.1418286 ± 45	-6.0 ± 3.9
PC-106	Tonalite	20.90	3.08	0.0891	0.509932 ± 1	-2.1	3.9	1.1418191 ± 34	-14.3 ± 3.0
PC-169	Tonalite	13.60	2.15	0.0954	0.510119 ± 1	-1.5	3.9	1.1418239 ± 49	-10.1 ± 4.3
PC-286 ^a	Tonalite	20.80	3.74	0.1087	0.510106 ± 2	-8.1	4.4	1.1418169 ± 72	-16.2 ± 6.3
POR23 ^b	Tonalite	21.60	3.66	0.1025	0.510263 ± 8	-2.0	3.9		
POR23-2 ^b	Tonalite	21.10	3.56	0.1017	0.510271 ± 14	-1.5	3.9		
WP59 ^b	Tonalite	25.50	4.09	0.0968	0.510140 ± 8	-1.7	3.9		
WP104 ^b	Tonalite	36.40	4.81	0.0799	0.509737 ± 8	-1.6	3.8		
WP69b ^b	Tonalite	23.30	4.22	0.1096	0.510497 ± 9	-0.8	3.8		
WP67 ^b	Tonalite	14.10	2.20	0.0943	0.510117 ± 9	-1.0	3.8		
PC-101 ^a	Trond. Band	15.14	2.48	0.0989	0.510249 ± 10	1.2	3.8	1.1418249 ± 68	-9.2 ± 5.9
PC-102 ^a	Trond. Band	17.85	2.87	0.0972	0.510173 ± 10	0.6	3.8	1.1418241 ± 64	-9.9 ± 5.6
PC-287	Trond. Band	13.87	2.16	0.0940	0.510090 ± 1	0.5	3.9	1.1418254 ± 34	-8.8 ± 3.0
R18	Pegmatite	7.18	1.50	0.1260	0.510738 ± 1	-12.7	4.2	1.1418230 ± 34	-10.9 ± 3.0
Mineral separa	ate								
PC-230GRT	Garnet	0.40	1.05	1.6053	0.536509 ± 20				
PC-267GRT	Garnet	2.11	1.49	0.4269	0.516282 ± 5				
PC-275GRT	Garnet	2.06	1.55	0.4552	0.516651 ± 7				
PC-276GRT	Garnet	4.30	2.37	0.3333	0.514457 ± 13				
PC-276BT	Biotite	0.52	0.08	0.0938	0.510886 ± 24				
PC-282GRT	Garnet	1.48	0.95	0.3896	0.515415 ± 11				
PC-438GRT	Garnet	1.25	0.67	0.3250	0.514299 ± 14				
PC-536GRT	Garnet	2.33	0.74	0.1924	0.512176 ± 9				
PC-085WR	Gabbro	7.18	2.32	0.1956	0.512722 ± 1				
PC-085PL	Plagioclase	0.53	0.10	0.1146	0.511261 ± 45				
PC-085HBL	Hornblende	9.81	3.22	0.1987	0.512762 ± 24				

^a Data from O'Neil et al. (2008).

^b Data from David et al. (2009).

^c Average μ^{142} Nd value for duplicate samples (see Table SD3).

^d Average μ^{142} Nd value for duplicate samples including data from O'Neil et al. (2008) (see Table SD3).

 2σ errors for ¹⁴³Nd/¹⁴⁴Nd and ¹⁴²Nd/¹⁴⁴Nd are in the last decimal place.

that between 700 and 500 ng of Nd were analyzed for a sample, the Nd blank represents at most 0.5% of the total Nd analyzed. Repeated measurements of the JNdi-1 standard yielded a long-term dynamically measured value of 142 Nd/ 144 Nd = 1.1418354 (*n* = 41, with most JNdi-1 standards analyzed twice on the same filament) with an external reproducibility (2σ , not 2σ -mean) of 4.5 ppm for the period of the study (Table SD4). The chondritic reference values used for ε Nd calculations are; ¹⁴³Nd/¹⁴⁴Nd_{CHUR} = 0.51263, 147 Sm/ 144 Nd_{CHUR} = 0.196 (Bouvier et al., 2008). Values used for Nd model age calculations are present day 143 Nd/ 144 Nd = 0.51301 and ¹⁴⁷Sm/¹⁴⁴Nd = 0.2085 reflecting a non-chondritic bulk Earth composition that today would have the 18 ppm excess ¹⁴²Nd/¹⁴⁴Nd observed in the modern terrestrial mantle compared to average ordinary chondrite (Carlson and Boyet, 2008). Initial ε^{143} Nd and uncertainty were calculated following Fletcher and Rosman (1982). Initial μ^{142} Nd were determined from isochrons and normalized to a depleted mantle (DM) reservoir with 147 Sm/ 144 Nd = 0.2085 and μ^{142} Nd = 0. The decay constant used for ¹⁴⁷Sm is 6.54×10^{-12} yr⁻¹ and 1.02×10^{-8} yr⁻¹ for ¹⁴⁶Sm. For the ¹⁴²Nd analysis, corrections for both Ce and Sm interference were made prior to mass fractionation correction. ¹⁴⁰Ce/¹⁴⁶Nd ratios ranged from 8.4×10^{-7} to 8.3×10^{-4} with 52 out of 62 samples having 140 Ce/ 146 Nd < 10^{-4} . The measured 147 Sm/ 146 Nd ratio during Nd runs averaged 3.1×10^{-6} with only 2 samples exceeding 1×10^{-5} . One sample (PC-220) yielded higher 140 Ce/ 146 Nd and 147 Sm/ 146 Nd ratios of 1.24×10^{-3} and 1.7×10^{-4} respectively.

The ¹⁴³Nd/¹⁴⁴Nd and ¹⁴⁷Sm/¹⁴⁴Nd ratios for PC-085 (whole rock, hornblende and plagioclase separates) as well as the biotite separate for PC-276 were analyzed at the GEOTOP laboratories (Université du Québec à Montréal) following the procedure described in O'Neil et al. (2008)

The μ^{142} Nd [((((¹⁴²Nd/¹⁴⁴Nd)_{sample}/(¹⁴²Nd/¹⁴⁴Nd)_{JNdi-1}) - 1) × 10⁶] for each sample was calculated using the average

(¹⁴²Nd/¹⁴⁴Nd)_{INdi-1} ratio within the same barrel used for sample analysis. In order to compare the ¹⁴²Nd/¹⁴⁴Nd ratios for samples from different barrels, the ¹⁴²Nd/¹⁴⁴Nd ratios for each sample reported in Table 1 were calculated using their μ^{142} Nd re-normalized to the long-term ¹⁴²Nd/¹⁴⁴Nd ratio for JNdi-1 of 142 Nd/ 144 Nd = 1.1418354. The same re-normalization was applied to data from O'Neil et al. (2008) to allow comparisons of both studies. No correlation exists between ¹⁴²Nd/¹⁴⁴Nd and ¹⁴⁰Ce/¹⁴⁶Nd or ¹⁴⁷Sm/¹⁴⁶Nd in samples or standards (Fig. SD3) suggesting that Ce and Sm interferences do not affect the data. Sample PC-220 has higher Sm and Ce interferences (Table SD2) but still does not display any correlation between ¹⁴²Nd/¹⁴⁴Nd and ¹⁴⁰Ce/¹⁴⁶Nd or ¹⁴⁷Sm/¹⁴⁶Nd (Fig. SD3) and has a ¹⁴²Nd isotopic composition consistent with other samples from the same geochemical group. Sample PC-456 has the lowest μ^{142} Nd value (-18; Table 1) but with very small Sm and Ce interferences (Table SD2) and without any correlation between ¹⁴²Nd/¹⁴⁴Nd and ¹⁴⁰Ce/¹⁴⁶Nd or ¹⁴⁷Sm/¹⁴⁶Nd (Fig. SD3) supporting the fact that the measured μ^{142} Nd values are not artifacts due to mass interferences and represent the real ¹⁴²Nd/¹⁴⁴Nd isotopic compositions of the samples. There also is no correlation between the ¹⁴²Nd/¹⁴⁴Nd ratios for the samples and their ¹⁴⁵Nd/¹⁴⁴Nd, ¹⁴⁸Nd/¹⁴⁴Nd, and ¹⁵⁰Nd/¹⁴⁴Nd ratios (Fig. SD4) indicating that the variability in ¹⁴²Nd/¹⁴⁴Nd ratio is not due to an incorrect mass dependency for the fractionation correction. Duplicate ¹⁴²Nd/¹⁴⁴Nd analyses are presented in Table SD3 and Fig. SD2. No variation within error of the ¹⁴²Nd/¹⁴⁴Nd ratios is observed for duplicate samples. Duplicate samples also reproduce data from O'Neil et al. (2008) for the same sample. Table SD5 shows three ¹⁴²Nd duplicate analyses of samples from Isua analyzed previously by Rizo et al. (2011). The ¹⁴²Nd data obtained at DTM that range from μ^{142} Nd of +8 to +12 agree within error with the data reported by Rizo et al. (2011) on the same samples.

Lu-Hf measurements were performed on the DTM Nuplasma multicollector-ICP-MS. Data are fractionation corrected to ¹⁷⁹Hf/¹⁷⁷Hf=0.7325 and normalized to the average value of 176 Hf/ 177 Hf obtained for the JMC-475 Hf standard (n = 10) during the same analytical session as the sample analyses. Values for ¹⁷⁶Hf/¹⁷⁷Hf for the IMC-475 standard ranged from 0.282139 to 0.282172 with an average value of 0.282152 ± 0.000022 (2 σ . not 2σ -mean). All data are reported relative to 176 Hf/ 177 Hf=0.282160 for the IMC-475 standard. Total Hf blank for the flux-fusion technique used for the whole rock measurements is 630 pg. The garnets were digested in Parr bombs where the total Hf blank is 88 pg. Although the Hf blank for the flux-fusion technique is higher, between 250 and 85 ng of Hf were analyzed for the whole rock samples. The Hf blank thus represents 0.7–0.3% of the total Hf analyzed. Table SD5 shows two Lu-Hf duplicate analyses of Isua samples previously reported by Rizo et al. (2011). The samples analyzed at DTM were dissolved using the flux-fusion technique and despite the higher blank, agree with the results reported by Rizo et al. (2011) within error.

U–Pb geochronology of zircons was conducted at the Laboratoire Magmas et Volcans, Clermont-Ferrand (France). Zircons were first identified directly on polished thin sections using a CAMECA SX 100 electron microprobe and then analyzed by laser ablation inductively coupled plasma spectrometry (LA-ICPMS). Analytical techniques for LA-ICPMS analysis are detailed in Table SD6. The method used for the zircon isotope dating with laser ablation ICP-MS followed the method developed for monazite by Tiepolo (2003) and Paquette and Tiepolo (2007). Concordia ages and diagrams were generated using Isoplot/Ex v. 2.49 software package by Ludwig (2001). The concentrations in U–Th–Pb were calibrated relative to the certified contents of the GJ-1 zircon standard (Jackson et al., 2004).

4. Results

4.1. ¹⁴⁷Sm-¹⁴³Nd

4.1.1. Ujaraaluk unit and cogenetic ultramafic rocks

Samples from the three different geochemical groups of the Ujaraaluk unit, including samples from the greenstones, were analyzed for their ¹⁴⁷Sm-¹⁴³Nd isotopic composition along with their respective cogenetic ultramafic sill samples (Table 1). The Ujaraaluk mafic rocks display a wide range of ¹⁴⁷Sm/¹⁴⁴Nd ratios from substantially subchrondritic to slightly superchondritic (0.1257-0.2143) with ¹⁴³Nd/¹⁴⁴Nd isotopic compositions ranging from 0.510976 to 0.513219. The ultramafic sills have higher ¹⁴⁷Sm/¹⁴⁴Nd ratios ranging from 0.1603 to 0.2452 with more radiogenic ¹⁴³Nd/¹⁴⁴Nd compositions from 0.511894 to 0.513658. Sample PC-214 has a significantly more radiogenic isotopic composition compared to the rest of the ultramafic samples with a ¹⁴⁷Sm/¹⁴⁴Nd ratio of 0.2681 and a ¹⁴³Nd/¹⁴⁴Nd ratio of 0.514124. This is a sample of a layer of metamorphic orthopyroxene within an ultramafic sill, which may replace an original magmatic cumulate layer. A ¹⁴⁷Sm-¹⁴³Nd isochron fit to all Ujaraaluk and cogenetic ultramafic samples yields an age of 3598 ± 200 Ma (Fig. 2A) with an initial ε Nd value of -1.0 ± 0.7 . This isochron, however, displays considerable scatter (MSWD = 134) and breaks down into a series of individual correlations with different slopes corresponding to the different geochemical groups within the Ujaraaluk (Fig. 2A). The ¹⁴⁷Sm-¹⁴³Nd correlation lines fit separately for the high-Ti Ujaraaluk, the depleted low-Ti Ujaraaluk and the enriched low-Ti Ujaraaluk, along with their respective cogenetic ultramafic rocks, yield ages of 3257 ± 300 Ma, 2517 ± 240 Ma and 3734 ± 300 Ma respectively (Fig. 2B-D). The isochron for the enriched low-Ti Ujaraaluk samples can be further broken down into 2 distinct slopes with younger ages of 2844 ± 460 Ma and 2685 ± 470 Ma respectively. These ages have been calculated with the exclusion of 2 high-Ti Ujaraaluk (PC-058, PC-451) samples and one depleted low-Ti Ujaraaluk (PC-214) sample. These samples lie well off the best fit lines defined by the remaining samples. We interpret this as reflecting post-magmatic processes that are further discussed in Section 5.2. The Ujaraaluk/ultramafic sills display a wide range of depleted-mantle model ages (T_{DM}) mainly ranging from 3.6 Ga to 4.5 Ga. A few samples have $T_{DM} \ge 4.5$ Ga most likely due to their near depleted-mantle Sm/Nd ratios resulting in an isotopic evolution trend that is sub-parallel to the depleted mantle evolution line. Samples with T_{DM} values between 3.6 Ga and 4.5 Ga have a bimodal distribution with groupings around 3.9 Ga and 4.3 Ga (Fig. 3A).

4.1.2. Central gabbro sills

Gabbro sill samples were collected toward the center of the tight synform. These gabbros display a distinctive fine-scale gneissosity defined by plagioclase-rich and amphibole-rich bands. The gabbros have chondritic to slightly unradiogenic Nd isotopic compositions with ¹⁴⁷Sm/¹⁴⁴Nd ratios ranging from 0.2034 to 0.1424 and ¹⁴³Nd/¹⁴⁴Nd ranging from 0.512909 to 0.511233 (Table 1). They define a relatively good isochron (MSWD=7.2) yielding an age of 4095 ± 130 Ma (*n* = 14) (Fig. 2E). Only one sample (PC-522) lies off the best fit line. An isochron excluding this sample gives an age of 4115 ± 100 Ma (MSWD = 4.8) with an initial ε Nd value of +1.9 \pm 0.3. The gabbro samples in Fig. 2E define 2 clusters in the isochron diagram and hence might represent 2 distinct groups of gabbros. However only fine grained gabbros displaying the same texture and collected along 2 large continuous sill bodies (35-130 m in width) that can be traced continuously in the field over 1.5 km, were selected for this isochron. T_{DM} values for the central gabbro show less variation with the majority of the samples having $T_{\rm DM}$ values between 3.9 Ga and 4.1 Ga (Fig. 3B).

4.1.3. NGB felsic rocks

A new series of felsic rocks from the NGB were analyzed for their ¹⁴⁷Sm-¹⁴³Nd isotopic composition. These include samples from the 3.6 Ga tonalite surrounding the NGB, the thin bands of 3.8 Ga intruding trondhjemite, and the 2.7 Ga pegmatites (Table 1). Tonalite samples and the trondhjemitic band sample show a narrow range of ¹⁴⁷Sm/¹⁴⁴Nd ratios from 0.1009 to 0.0891 and ¹⁴³Nd/¹⁴⁴Nd ranging from 0.510199 to 0.509932. The Neoarchean pegmatite sample has a slightly higher ¹⁴⁷Sm/¹⁴⁴Nd ratio of 0.1260 and a ¹⁴³Nd/¹⁴⁴Nd ratio of 0.510738. These values are within the range of isotopic compositions previously reported by O'Neil et al. (2008) and David et al. (2009) for the Nuvvuagittuq felsic rocks. The range in Sm/Nd ratios for the tonalities analyzed in this study is too narrow to produce a precise isochron, however, the combined tonalite data from this study, data from O'Neil et al. (2008) and data from David et al. (2009), yields an isochron with an age of 3575 ± 450 Ma (Fig. 2F) with an initial ε Nd value of -2.9 ± 0.4 . This isochron is obtained after rejection of 3 samples with higher Sm/Nd ratios that clearly fall off the main trend for the tonalities. Despite the large error due to the small range in Sm/Nd ratios, this age is in agreement with the U–Pb zircon age of 3661 ± 4 Ma for the Nuvvuagittuq tonalities obtained by David et al. (2009). The Nuvvuagittuq tonalities also have T_{DM} values systematically older than their crystallization age ranging between 3.8 Ga and 3.9 Ga, with 2 samples from O'Neil et al. (2008) yielding T_{DM} of 4.2 and 4.4 Ga (Fig. 3C) that suggest derivation from an older crustal protolith.

4.1.4. Mineral separates

The Sm–Nd isotopic compositions for mineral separates are given in Table 1. Fig. 4A shows the isochron ages obtained from the different mineral separates. All garnet/whole rock tie-lines for

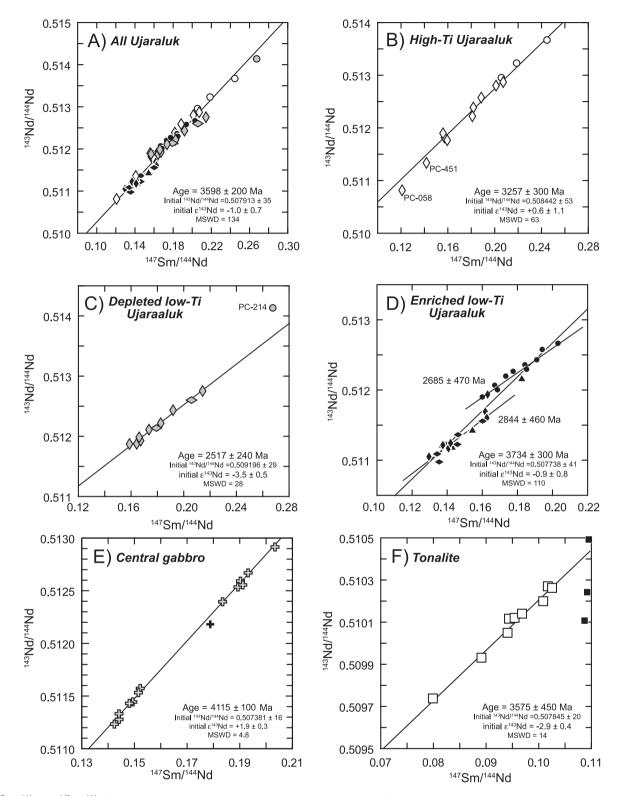


Fig. 2. ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴³Nd/¹⁴⁴Nd isochron diagrams for (A) All NGB Ujaraaluk and cogenetic ultramafic rocks, (B) the high-Ti Ujaraaluk geochemical group and cogenetic ultramafic rocks, (C) the depleted low-Ti Ujaraaluk geochemical group and cogenetic ultramafic rocks, (D) the enriched low-Ti Ujaraaluk geochemical group and cogenetic ultramafic rocks, (E) the central gabbro sills and (F) the tonalites. Symbols: open symbols, high-Ti Ujaraaluk unit and Type-1 ultramafic; gray symbols, depleted low-Ti Ujaraaluk unit and Type-2 ultramafic; black symbols, enriched low-Ti Ujaraaluk unit and Type-3 ultramafic; vertical diamond, garnet-free Ujaraaluk; horizontal diamond, garnet-bearing Ujaraaluk; triangle, cordierite-anthophyllite Ujaraaluk; circle, ultramafic; gray plusses, gabbro; open square, tonalite. Black symbols in Fig. 2E and F are samples not included in the isochron age calculation.

the Ujaraaluk samples (7 garnet/whole samples) yield Neoarchean ages ranging from 2537 Ma to 2735 Ma with negative initial ε Nd values ranging from -9.4 to -12.4, with PC-230 having an initial ε Nd value of -6.5 (Table SD7). An isochron comprising

only garnet from 7 Ujaraaluk samples, including one sample from the lower greenschist metamorphic grade (PC-536 GRT), gives an age of 2619 ± 50 Ma (Fig. 4B) with an initial ε Nd value of -9.3 ± 1.8 . The coarse grained gabbro (PC-085) yields an isochron

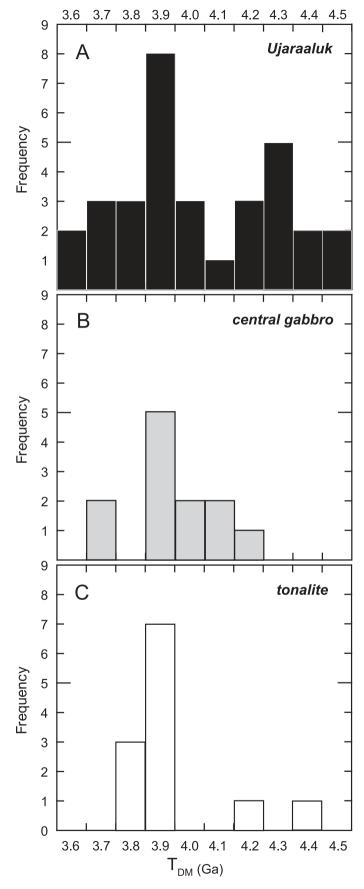


Fig. 3. Histograms of T_{DM} values for (A) the Ujaraaluk unit, (B) the gabbros and (C) the tonalites. Values used for Nd model age calculations are present day 143 Nd/ 144 Nd = 0.51301 and 147 Sm/ 144 Nd = 0.2085.

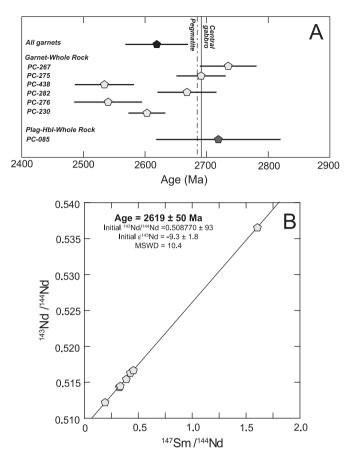


Fig. 4. (A) Sm–Nd isochron ages for garnet separates and mineral/whole rock tielines. Dashed line is the U–Pb age of the intruding pegmatite at 2686 Ma. Solid vertical line is the U–Pb age for zircons from the central gabbros (see Section 4.4). (B) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴³Nd/¹⁴⁴Nd isochron diagram for all garnet separate samples.

age (hornblende–plagioclase–whole rock) of 2719 ± 100 Ma, but with a positive ε Nd value of $\pm 1.8\pm0.5$ because the whole rock gabbro has a near chondritic Sm/Nd ratio. These Neoarchean ages generally overlap within error with the age for the intrusion of the Nuvvuagittuq pegmatite reported by David et al. (2009) of 2686 ± 4 Ma, as does the imprecise whole rock Pb–Pb age of 2.4 ± 0.4 Ga for the Ujaraaluk samples measured in O'Neil et al. (2008). PC-276, in contrast, yields a Proterozoic biotite–whole rock isochron age of 1537 Ma.

4.2. Lu-Hf

Four of the garnet separate samples analyzed for Sm–Nd isotopic composition have also been analyzed for their Lu–Hf isotopic composition, along with their whole rock Lu–Hf isotopic composition (Table SD8). ¹⁷⁶Lu–¹⁷⁶Hf garnet/whole rock tie-lines for the Ujaraaluk samples give a wide range of Proterozoic ages with, PC-230 and PC-275 yielding ages of 1655 Ma and 1599 Ma respectively, whereas PC-282 and PC-438 give younger ages of 1219 Ma and 1157 Ma respectively.

4.3. ¹⁴²Nd

The ¹⁴²Nd isotopic data are presented in Table 1. The ¹⁴²Nd/¹⁴⁴Nd ratios of the Ujaraaluk rocks show a 26 ppm variation ranging from samples having an excess in ¹⁴²Nd (up to +8 ppm) to samples having a deficit in ¹⁴²Nd (up to -18 ppm) relative to the terrestrial standard (Fig. 5). The ¹⁴²Nd/¹⁴⁴Nd ratios for this entire set of samples correlate with their Sm/Nd ratios. This correlation

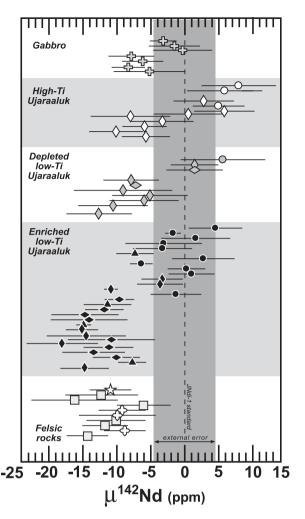


Fig. 5. μ^{142} Nd values for all NGB samples relative to the terrestrial Nd standard. Vertical gray band shows the 4.5 ppm external error. Samples are sorted in order of Sm/Nd ratios for each different geochemical group/lithology. The different geochemical groups/lithologies are separated by alternating white and gray horizontal zones. Symbols as in Fig. 2; open plusses, trondhjemitic bands; open star, pegmatite.

between the ¹⁴²Nd/¹⁴⁴Nd ratios and Sm/Nd ratios also is present when considering the 3 different geochemical groups of Ujaraaluk individually. The largest deficits in ¹⁴²Nd are observed in the enriched low-Ti Ujaraaluk, whereas the sample yielding a μ^{142} Nd of +8 is an ultramafic rock from the high-Ti Ujaraaluk group. The central gabbro samples also exhibit a correlation between their ¹⁴²Nd/¹⁴⁴Nd and their ¹⁴⁷Sm/¹⁴⁴Nd ratios, but with a smaller range in μ^{142} Nd values from 0 to -8 relative to terrestrial standard. All Nuvvuagittuq felsic rocks analyzed, including the tonalities (-6 to -14 ppm), the trondhjemitic band (-9 ppm) and the pegmatite (-11 ppm) yield deficits in ¹⁴²Nd relative to the terrestrial standard (Fig. 5). There is, however, no correlation between the ¹⁴²Nd/¹⁴⁴Nd and the ¹⁴⁷Sm/¹⁴⁴Nd ratios for any felsic lithology. The ¹⁴²Nd isotopic data measured in this study are consistent with the data reported by O'Neil et al. (2008) for every lithology, but expands the range of isotopic compositions exhibiting this correlation. The deficits in ¹⁴²Nd in rocks from the NGB have been confirmed by measurements of the same samples in an independent lab at ETH Zurich (Roth et al., 2011).

4.4. Zircon U–Pb

Zircons from three central gabbro samples have been analyzed by LA-ICP-MS (Table SD9). The analyzed zircons were relatively

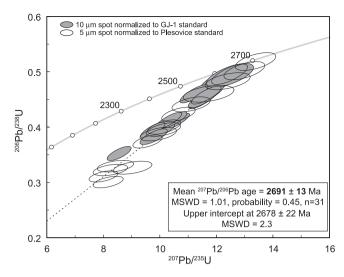


Fig. 6. U–Pb conrcordia diagram for zircons in the central gabbros. Ellipses are 2σ error. Solid gray ellipses are for 10 μ m laser spots normalized to GJ-1 standard and open ellipses are for 5 μ m laser spots normalized to Plesovice standard.

rounded in shape and between 30 μ m and 10 μ m in size with most of them being ~20 μ m. The most concordant zircons (n = 31) yield a ²⁰⁷Pb/²⁰⁶Pb age of 2691 ± 13 Ma (Fig. 6) in agreement with the age of the intrusion of the pegmatite and the metamorphic ages obtained on the mineral separates.

5. Discussion

5.1. Significance of the Ujaraaluk ¹⁴²Nd isotopic composition

 146 Sm decays into 142 Nd with a half-life of $T_{1/2}$ = 68 Ma (Kinoshita et al., 2012). Given this short half life, ¹⁴⁶Sm is effectively extinct by ~4.0 Ga, therefore any deviation of the ¹⁴²Nd isotopic composition from the terrestrial standard has to be produced by fractionation between Sm and Nd that occurred prior to 4.0 Ga. The ¹⁴²Nd data set for the Nuvvuagittuq rocks has been expanded with 30 samples having a resolvable deficit in ¹⁴²Nd compared to the terrestrial standard. Most samples with low ¹⁴²Nd/¹⁴⁴Nd derive from the enriched low-Ti Ujaraaluk group. Within every group of Ujaraaluk, however, the ¹⁴²Nd/¹⁴⁴Nd ratios correlate with Sm/Nd ratios. Any valid model for the origin of the Ujaraaluk unit needs to account for three main features regarding their ¹⁴²Nd isotopic composition. First, it must explain the deficit in ¹⁴²Nd relative to the terrestrial standard. Second, it must account for the 26 ppm range in ¹⁴²Nd observed in the Ujaraaluk and cogenetic ultramafic sills. Third, it must explain the correlation between the Sm/Nd and ¹⁴²Nd/¹⁴⁴Nd ratios for all Ujaraaluk rocks and within each individual geochemical group. Here, we will discuss 3 possible models that could potentially explain the observed ¹⁴²Nd deficits: (1) Derivation of the NGB rocks in the Eoarchean from an enriched mantle formed in the Hadean, (2) Mixing in the Eoarchean with an Hadean enriched mantle reservoir and (3) Formation of the NGB rocks in the Hadean. We then will evaluate which model best accounts for the ¹⁴²Nd isotopic composition of the NGB rocks.

5.1.1. Derivation from an Hadean enriched mantle

Variability in ¹⁴²Nd in relatively young rocks can be inherited from mantle sources whose Sm/Nd ratios were fractionated by differentiation events occurring prior to 4 Ga. A clear example is the presence of a wide range in initial ¹⁴²Nd/¹⁴⁴Nd in <4 Ga old lunar mare basalts (Rankenburg et al., 2006; Boyet and Carlson, 2007). Most terrestrial rocks have a ~20 ppm excess in ¹⁴²Nd compared to ordinary chondrites, which led Boyet and Carlson (2005, 2006)

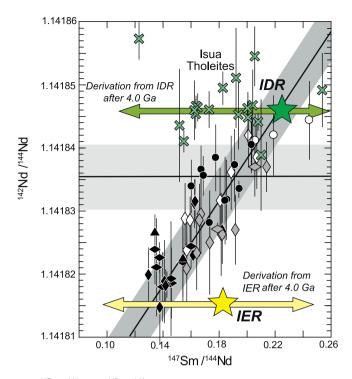


Fig. 7. ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd diagram. Symbols as in Fig. 2; green crosses, Isua samples from Rizo et al. (2011) and Bennett et al. (2007); green star, Isua-like depleted reservoir (IDR) isotopic composition with a ¹⁴⁷Sm/¹⁴⁴Nd ratio = 0.225 and a μ^{142} Nd value of ~+9 relative to terrestrial standard (Rizo et al., 2011; the ¹⁴⁷Sm/¹⁴⁴Nd ratio was re-calculated using the ¹⁴⁶Sm half-life of 68 Ma); yellow star, possible complementary Isua-like enriched reservoir (IER) with a μ^{142} Nd value of ~-15 ppm relative to terrestrial standard to account for the most negative NGB values and a ¹⁴⁷Sm/¹⁴⁴Nd ratio = 0.182, required for an age of differentiation of 4.47 Ga (Rizo et al., 2011). Horizontal black line shows the terrestrial standard value with 4.5 ppm external error gray band. Oblique black line shows the best fit for the Ujaraaluk sample correlation with 4.5 ppm external error being shown by the gray band. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)

to propose that the Bulk Silicate Earth either has a superchondritic Sm/Nd ratio or differentiated within 30 Ma of planet formation to form an early depleted mantle. If the bulk Earth is chondritic in Sm/Nd ratio, then the formation of an early depleted, i.e. high Sm/Nd ratio, reservoir having an excess in ¹⁴²Nd must be coupled with the formation of an early enriched reservoir (EER) that is characterized by a complementary deficit in ¹⁴²Nd (Boyet and Carlson, 2006). Any rock derived from the EER will carry this deficit in ¹⁴²Nd independent of its age. Such rocks will not only have a deficit in ¹⁴²Nd relative to the terrestrial standard, but also relative to ordinary chondrite (i.e. ~–40 ppm relative to the terrestrial standard). The NGB rocks have ¹⁴²Nd deficits up to –18 ppm relative to terrestrial standard and thus no deficit relative to ordinary chondrites. Therefore, the NGB rocks cannot be derived directly from the EER.

Positive anomalies in ¹⁴²Nd are found in amphibolites from Isua in SW Greenland. These ~3.8 Ga rocks have an excess in ¹⁴²Nd (up to +19 ppm) compared to terrestrial standards (Rizo et al., 2011; Boyet and Carlson, 2006; Caro et al., 2003, 2006; Bennett et al., 2007). There is no correlation between the ¹⁴⁷Sm/¹⁴³Nd and the ¹⁴²Nd/¹⁴⁴Nd ratios in the Isua amphibolites (Fig. 7), consequently, the most likely origin of the excess ¹⁴²Nd in these rocks is that their mantle source was formed in the Hadean and was depleted in incompatible elements (i.e. high Sm/Nd ratio) (Caro et al., 2006; Bennett et al., 2007; Rizo et al., 2011). This mantle reservoir was later sampled by an Eoarchean crust forming event that produced melts with variable Sm/Nd ratio as a result of magmatic fractionation, but with ¹⁴²Nd excess inherited from the source. Because ¹⁴⁶Sm was extinct by 3.8 Ga, the rocks formed from these magmas will retain their initially uncorrelated Sm/Nd and ¹⁴²Nd/¹⁴⁴Nd ratios to the present day. An Hadean complementary enriched reservoir to the Isua mantle source would have a deficit in ¹⁴²Nd relative to the terrestrial standard. Rocks derived from such an enriched reservoir during some later melting event would therefore yield negative anomalies in ¹⁴²Nd and thus could have similar isotopic compositions to the NGB rocks. However, partial fusion after 4.0 Ga of an enriched mantle with a deficit in ¹⁴²Nd will produce magmas with deficits in ¹⁴²Nd that need not correlate with Sm/Nd ratio, as shown by the analogous case of the Isua amphibolites (Fig. 7). In other words, all rocks derived from this source would have the same ¹⁴²Nd isotopic composition, regardless of their Sm/Nd ratios. Direct post-4.0 Ga derivation from an enriched mantle reservoir formed contemporaneously with the Isua mantle source cannot account for the 26 ppm variation in ¹⁴²Nd observed in the NGB rocks and more importantly, cannot explain the correlation between the Sm/Nd and the ¹⁴²Nd/¹⁴⁴Nd ratios characteristic of the Ujaraaluk unit.

5.1.2. Mixing of Hadean enriched and depleted mantle reservoirs

There is always the possibility that a correlated array of Nd isotopic composition and Sm/Nd ratio may represent a mixing line between 2 isotopically distinct reservoirs. The minimum age for the Ujaraaluk unit is constrained by the intrusion of the trondhjemitic bands dated at ~3.8 Ga (Cates and Mojzsis, 2007; David et al., 2009). The only way that the Ujaraaluk can have a deficit in ¹⁴²Nd if they indeed formed at 3.8 Ga, is to involve an Hadean enriched component in their petrogenesis. If the Ujaraaluk unit formed at 3.8 Ga from a "normal" depleted mantle (DM) source, mixing with an early enriched reservoir (EER) would induce a ¹⁴²Nd deficit and could yield a correlation between Sm/Nd and ¹⁴²Nd/¹⁴⁴Nd if igneous fractionation did not greatly change the Sm/Nd ratios of the mixed rocks (Fig. 8A). In this case, the correlation between the Sm/Nd and the ¹⁴²Nd/¹⁴⁴Nd ratios would represent a mixing line without any age significance for the Ujaraaluk, at best providing only an approximate age for the time of separation of the EER and the DM. If the Ujaraaluk Sm/Nd vs. ¹⁴²Nd correlation is a mixing line, the ¹⁴⁷Sm/¹⁴⁴Nd ratio of the enriched reservoir must be less than ~0.12, and have a 142 Nd deficit of at least ~-20 ppm relative to terrestrial standard (EER 1 in Fig. 8A). Given that mantle rocks with ¹⁴⁷Sm/¹⁴⁴Nd significantly less than 0.12 are uncommon [even the KREEP component of the Moon, a possible analog to an early terrestrial EER, has a ¹⁴⁷Sm/¹⁴⁴Nd of ~0.168 (Lugmair and Carlson, 1978)], the lowest Sm/Nd ratio end member in the Ujaraaluk would have to be dominated by the enriched end member. One could imagine that the early-formed enriched reservoir might be a strongly LREE enriched Hadean felsic crust, in which case the lowest Sm/Nd ratio end member in the Ujaraaluk would still require more than a 50% by mass contribution from such "crustal contamination" if the mantle-derived end member were from "normal" terrestrial mantle. Given the basaltic major element composition of the low Sm/Nd ratio end of the Ujaraaluk array, the compositional characteristics of the Ujaraaluk rocks do not support the possibility that this enriched mixing end member is Hadean felsic crust. Furthermore, if the ¹⁴²Nd deficits in the Ujaraaluk unit were produced by mixing with an Hadean LREE-enriched reservoir, a correlation between the Nd concentrations and the ¹⁴²Nd isotopic compositions at the time of mixing (presumably 3.8 Ga) would be expected. However, the μ^{142} Nd values of the Nuvvuagittuq rocks show absolutely no correlation with their Nd concentrations for the entire set of data or within the individual Ujaraaluk geochemical groups (Fig. 8B). Moreover, if the ¹⁴²Nd deficits observed in most NGB lithologies, and the Sm/Nd vs. 142Nd/144Nd correlation, were produced by mixing between enriched and depleted mantle sources at 3.8 Ga, a positive correlation between Sm/Nd ratio and initial ¹⁴³Nd/¹⁴⁴Nd_(3.8 Ga) would be expected that would give a

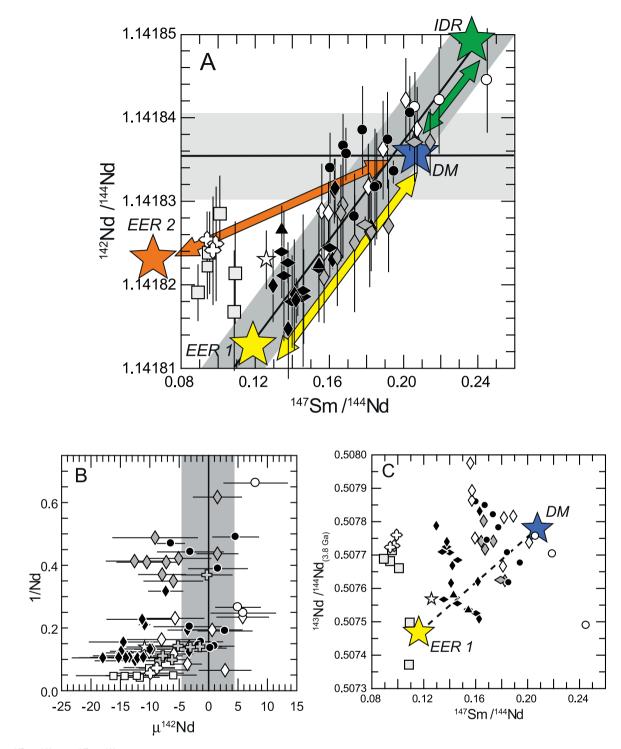


Fig. 8. (A) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd diagram. Symbols as in Fig. 5; blue star, DM; green star, Isua-like depleted reservoir (IDR) isotopic composition required to account for the superchondritic end of the Ujaraaluk correlation; yellow star, EER1 isotopic composition required to account for the subchondritic end of the Ujaraaluk correlation; orange star, EER2 isotopic composition required to account for the NGB felsic rocks. Horizontal black line shows the terrestrial standard value with 4.5 ppm external error. Oblique black line shows the best fit for the Ujaraaluk samples correlation with 4.5 ppm external error being shown by the gray band. (B) Measured μ^{142} Nd vs. 1/Nd concentration. Simple two-component mixing would predict a linear correlation between a low 1/Nd at negative μ^{142} Nd enriched source and a high 1/Nd, positive μ^{142} Nd depleted source. No such correlation is apparent in the data as the different compositional groups primarily all show a similar range in μ^{142} Nd, offset at different Nd concentrations. (C) ¹⁴⁷Sm/¹⁴⁴Nd vs. initial ¹⁴³Nd/¹⁴⁴Nd calculated at 3.8 Ga. Symbols as in Fig. 5. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)

slope with an age somewhere between the time of the mixing event and the formation of the enriched and depleted mantle sources. This would also result in a Sm/Nd vs. 143 Nd/ 144 Nd($_{(today)}$ correlation producing a "fossil" isochron with an age older than 3.8 Ga. A 147 Sm– 143 Nd isochron diagram constructed for the

Ujaraaluk samples using the ¹⁴³Nd/¹⁴⁴Nd ratios calculated at 3.8 Ga (Fig. 8C) displays mostly negative slopes within the individual chemical groups of Ujaraaluk, consistent with the <3.8 Ga ¹⁴⁷Sm-¹⁴³Nd isochron ages calculated for the different chemical groups. One could imagine a positive slope passing through the

midpoints of the negative slopes of the individual chemical units shown in Fig. 8C that could be attributed to formation of the different chemical groups by mixing between Hadean depleted and enriched mantle sources. In this case, however, the negative correlations between the Sm/Nd and 143 Nd/ 144 Nd($_{(3.8 Ga)}$ ratios in the individual chemical groups would have to be attributed to substantial modification of Sm/Nd ratios after 3.8 Ga. However, changes to Sm/Nd ratio caused either by magmatic differentiation at 3.8 Ga, or by metamorphism after 3.8 Ga would not be accompanied by changes in ¹⁴²Nd/¹⁴⁴Nd because ¹⁴⁶Sm was extinct by this time. Consequently, if the lack of a clear positive correlation between Sm/Nd and ¹⁴³Nd/¹⁴⁴Nd_(3.8 Ga) is due to either magmatic or metamorphic modification of Sm/Nd ratio at 3.8 Ga or later, then a similar, or greater, degree of scatter should be observed in the Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd plot, yet this isochron diagram shows a good positive correlation for all Ujaraaluk samples and for each individual chemical group (see Fig. 10).

Although the possibility of mixing between Hadean enriched and depleted sources in the Archean to explain the NGB ¹⁴²Nd isotopic composition cannot completely be ruled out, another complication if all the NGB rocks formed in the Eoarchean is that the Eoarchean felsic rocks, including the only rocks unequivocally dated at 3.8 Ga (i.e. the trondhjemitic bands), do not fall on the same Sm-Nd array defined by the Ujaraaluk samples (Fig. 8A). If the Ujaraaluk Sm/Nd-142Nd array reflects simple two-component mixing between enriched and depleted mantle sources at 3.8 Ga, then the offset of the felsic rocks from this line suggests a second enriched source (EER 2 in Fig. 8A) with much lower Sm/Nd ratio at the same ¹⁴²Nd/¹⁴⁴Nd ratio compared to the enriched source involved in the Ujaraaluk correlation. Moreover, the 2.7 Ga pegmatite would have to acquire its ¹⁴²Nd deficit by mixing with the same enriched source sampled by the Ujaraaluk (EER 1 in Fig. 8A), but over a billion years later, implying not only alternating sampling of different mixed sources (EER 1 and EER 2) in time but also the survival of these enriched reservoirs within the mantle for at least 1.5 billion years. Alternatively, the offset of the felsic rocks to lower Sm/Nd ratio compared to the Ujaraaluk correlation could reflect the magmatic fractionation of Sm/Nd ratio during the production and differentiation of these felsic rocks at various times post 3.8 Ga. If a similar magmatic fractionation of Sm/Nd ratio in the Ujaraaluk occurred, as is suggested by the trace element characteristics of the Ujaraaluk (O'Neil et al., 2011a), then the preservation of a correlation between Sm/Nd and 142Nd/144Nd ratio in a two-component mixture of enriched and depleted mantle sources is surprising if these rocks formed after 146Sm became extinct. At the high Sm/Nd ratio end of the Ujaraaluk correlation, the slight excess in ¹⁴²Nd observed in some NGB ultramafic rocks would require the involvement of an early Isua-like depleted reservoir to produce μ^{142} Nd values as high as +8 (Fig. 8A). Therefore, a mixing model to account for the Sm-Nd systematics of all NGB lithologies would require at least 3 distinct sources including an early Isua-like depleted reservoir and 2 early enriched reservoirs. While such complicated mixing scenarios cannot be ruled out unequivocally, we consider them geologically unlikely.

All three Ujaraaluk geochemical groups follow the exact same Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd correlation (Fig. 10A). O'Neil et al. (2011a) showed that these Ujaraaluk groups have distinct major and trace element compositions suggesting that they were formed in different geological settings and derived from mantle sources with different geochemical compositions. The high-Ti Ujaraaluk unit is consistent with derivation from an undepleted mantle source and fractionation under relatively dry conditions, while both low-Ti Ujaraaluk units appear to be derived from a mantle source that had experienced enrichment of highly incompatible elements followed by magmatic fractionation under elevated water pressures. Prior to the source incompatible element enrichment, the

mantle source of the depleted low-Ti Ujaraaluk unit appears to have been a refractory mantle (e.g. subchondritic Sm/Yb ratios) whereas the enriched low-Ti Ujaraaluk shows flat middle to heavy REE patterns consistent with an initially undepleted mantle source. Given that the three Ujaraaluk groups have distinct major and trace element compositions that imply they were produced from geochemically distinct mantle sources, fractionated under different conditions at low pressure, and possibly were produced in different geological settings, we consider it highly unlikely that the Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd correlation represents a mixing line where all three Ujaraaluk groups could have formed from the same isotopic end-members, producing a single correlation for the three compositionally distinct groups. Moreover, even a complex mixing history between depleted and enriched reservoirs cannot produce the isotopic relationship observed between the Ujaraaluk and their cogenetic ultramafic sills. As shown in O'Neil et al. (2011a), the sills share the same Al-Ti-Zr trends that define the 3 Ujaraaluk compositional groups with compositions controlled primarily by olivine fractionation (Fig. 9A). These compositional features suggest that the ultramafic rocks are cumulate bodies formed by the Ujaraaluk parental magmas. The magmatic accumulation of olivine, and possibly orthopyroxene, will produce cumulates with high Sm/Nd ratios relative to the liquid. If the magmatic accumulation occurred after 4.0 Ga, the ultramafic cumulates would have high Sm/Nd ratios, but retain the ¹⁴²Nd/¹⁴⁴Nd ratios of the Ujaraaluk parental magmas because ¹⁴⁶Sm is no longer extant to produce a change in ¹⁴²Nd abundance. High Sm/Nd ratio cumulates produced after 4.0 Ga would thus be expected to lie to the high Sm/Nd ratio side of the Ujaraaluk Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd array for the same reason that the felsic rocks lie to the low Sm/Nd side of the array if they are partial melts or fractionated liquids from some parental rock/magma lying on the Ujaraaluk array. The same would be true if the ultramafic rocks are interpreted as restites from partial melting that produced the Ujaraaluk parental magmas. If this magma-restite relationship was established at 3.8 Ga, the Ujaraaluk ultramafic rocks would today be offset to high Sm/Nd ratio from the Sm/Nd-¹⁴²Nd/¹⁴⁴Nd correlation shown by the Ujaraaluk mafic rocks. If the ultramafic rocks formed before 4.0 Ga, however, while ¹⁴⁶Sm was still actively decaying, their high Sm/Nd ratios would lead to high ¹⁴²Nd/¹⁴⁴Nd ratios compared to the Ujaraaluk magmas. The Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd correlation between the Ujaraaluk and their cogenetic ultramafic cumulates observed in each of the geochemical groups (Fig. 9B) can only be explained if the mineral fractionation forming the ultramafic cumulates occurred before 4.0 Ga. The isotopic relationship between the Ujaraaluk and their cogenetic ultramafic sills thus supports an Hadean formation age for the mafic/ultramafic rocks of the NGB.

5.1.3. Formation of the NGB rocks in the Hadean

In the simple case involving no metamorphic disturbance and no initial variation in the isotopic composition of the whole rocks, the correlation between Sm/Nd and ¹⁴²Nd/¹⁴⁴Nd ratios in the Ujaraaluk would represent an isochron dating the time of the igneous event that formed these rocks. The Sm-142Nd isochron passes, within error, through the DM isotopic composition consistent with the derivation of the Ujaraaluk from "normal" depleted mantle and does not require the involvement of any "missing" enriched reservoirs or Isua-like depleted reservoirs. The decay of ¹⁴⁶Sm in rocks with Sm/Nd ratios higher than the depleted mantle, primarily the cumulate ultramafic rocks, led to ¹⁴²Nd/¹⁴⁴Nd ratios higher than the modern depleted mantle whereas those with low Sm/Nd ratio ended up with a deficit in ¹⁴²Nd compared to depleted mantle. While O'Neil et al. (2011a) interpreted the relationship of mafic to ultramafic Ujaraaluk rocks as parental magma to cumulate, the ultramafic rocks also could be proposed to represent the residues of the partial melting that produced the

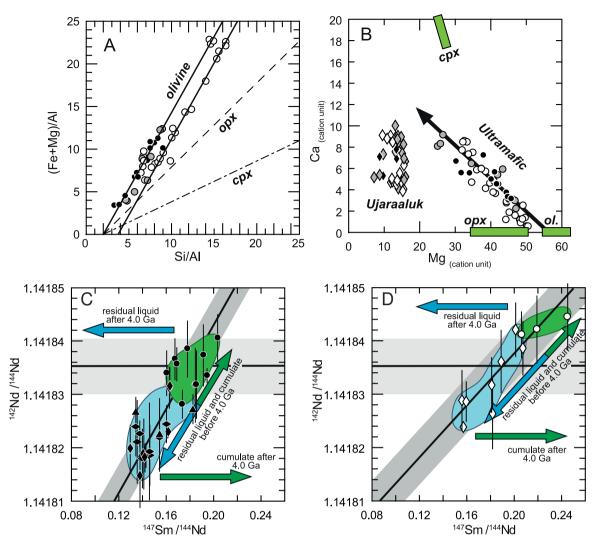


Fig. 9. (A) Pearce-type plot of (Fe + Mg)/Al vs. Si/Al and (B) Mg vs. Ca (cation units) diagram for the ultramafic samples illustrating the primary control of olivine fractionation. (C) ¹⁴⁷ Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd diagrams for the enriched low-Ti Ujaraaluk and (D) high-Ti Ujaraaluk. Symbols as in Fig. 2. Green areas highlight the ultramafic samples. Blue areas highlight the compositionally associated Ujaraaluk unit samples. Gray bands show the 4.5 ppm external error on the terrestrial standard value and the best fit lines. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)

mafic Ujaraaluk magmas. If this hypothetical relationship were established by melting at 3.8 Ga, both magma and restite would today have the same 142 Nd/ 144 Nd because there would be no 146 Sm extant to increase the 142 Nd/ 144 Nd ratios in the high Sm/Nd ratio ultramafic rocks. There is no petrological or geochemical evidence supporting the idea that each of the Ujaraaluk groups is composed of a series of restites and melts. On the contrary, the major and trace element compositional variations of the mafic rocks from the three groups of Ujaraaluk are consistent with fractional crystallization from distinct parental magmas, whereas the composition of the ultramafic samples is consistent with them being cogenetic cumulates to the different mafic parental magmas (O'Neil et al., 2011a). Therefore, the compositional variations of the Ujaraaluk unit and the ultramafic rocks, as well as the Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd ratio correlation are interpreted to be primary features acquired during the emplacement and crystallization of the Ujaraaluk unit.

The ¹⁴⁶Sm–¹⁴²Nd isochron for all Ujaraaluk samples and cogenetic sills corresponds to an age of 4388⁺¹⁵–₁₇ Ma (Fig. 10A) with an initial μ^{142} Nd value of +4.7 ± 2.0 overlapping with the "normal" DM. Fitting all the Ujaraaluk mafic samples only, the isochron gives the same age of 4388⁺¹⁸–₂₄ Ma. Isochrons fit individually to each of the different compositional groups in the Ujaraaluk (Fig. 10B–D)

give ages of 4363 $^{+21}_{-26}$ Ma, 4374 $^{+26}_{-35}$ Ma and 4409 $^{+23}_{-29}$ Ma for the high-Ti, depleted low-Ti, and enriched low-Ti groups respectively. Further support for an Hadean age for the Ujaraaluk unit is provided by the central intruding gabbro whose 146 Sm $^{-142}$ Nd systematics yield an age of 4313 $^{+41}_{-69}$ Ma (Fig. 10E). The Sm/Nd vs. 142 Nd/ 144 Nd correlation for the central gabbros is however not well-defined due to the limited spread in Sm/Nd ratios and the consequent limited variation in 142 Nd/ 144 Nd. Nevertheless, both the 146 Sm $^{-142}$ Nd and 147 Sm $^{-143}$ Nd systematics for the gabbros suggest an Hadean age of emplacement, providing a minimum age for Ujaraaluk unit. The 146 Sm $^{-142}$ Nd isochron for these gabbros also passes through the DM with an initial μ^{142} Nd value of +0.1 \pm 3.1.

5.2. Consequences of metamorphic disturbance

If the NGB rocks were not affected by later metamorphism, then the ¹⁴⁷Sm-¹⁴³Nd and ¹⁴⁶Sm-¹⁴²Nd data should provide concordant ages. The NGB has been extensively affected by metamorphism in the Neoarchean, however, as indicated by the numerous Sm-Nd ages on metamorphic minerals in the Ujaraaluk unit and intruding gabbro ranging between 2537 Ma and 2735 Ma (Fig. 4, Table SD6). These ages overlap the U–Pb zircon age of the intruding pegmatite, the U–Pb ages obtained on metamorphic

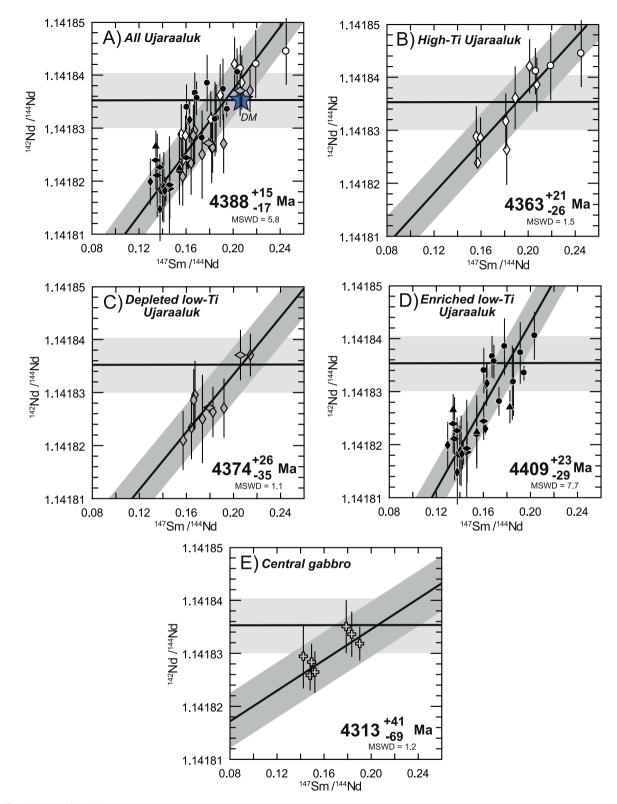


Fig. 10. ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd isochron diagrams for (A) All NGB Ujaraaluk and cogenetic ultramafic rocks, (B) the high-Ti Ujaraaluk geochemical group and cogenetic ultramafic rocks, (C) the depleted low-Ti Ujaraaluk geochemical group and cogenetic ultramafic rocks, (D) the enriched low-Ti Ujaraaluk geochemical group and cogenetic ultramafic rocks, (D) the enriched low-Ti Ujaraaluk geochemical group and cogenetic ultramafic rocks, and (E) the central gabbro sills. Gray bands show the 4.5 ppm external error on the terrestrial standard value and the best fit lines. Symbols as in Fig. 2; blue star, DM. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)

zircon, rutile and monazite in the Ujaraaluk unit (O'Neil et al., 2011b; Darling et al., 2012) and the central gabbros (Fig. 6), as well as the Pb-Pb age defined by whole rock samples of the Ujaraaluk (O'Neil et al., 2008). These results suggest that the peak of the metamorphism occurred at \sim 2.7 Ga, contemporaneous with the

emplacement of the Boizard suite (Simard et al., 2003) of heterogeneous tonalites surrounding the NGB supracrustal rocks and the 3.6 Ga NGB tonalite. Post-magmatic disturbance is also supported by younger Proterozoic Lu–Hf ages obtained in the same garnet dissolutions that provided Neoarchean Sm–Nd ages.

The long-lived ¹⁴⁷Sm-¹⁴³Nd isotopic system can be affected by younger thermal events even at the whole rock scale. The data presented here show clearly that the ¹⁴⁷Sm-¹⁴³Nd system in the NBG samples has been affected by the Neoarchean metamorphism. The ¹⁴⁷Sm-¹⁴³Nd systematics for the different compositional groups of Ujaraaluk give ages between 3734 Ma and 2517 Ma with large errors, significant scatter, and a wide range of initial ε Nd values. The best example of this is the isochron for the enriched low-Ti Ujaraaluk that yields an age of 3734 Ma, but is clearly composed of 2 different groups of samples with 2 different slopes, both of which give Neoarchean ages (Fig. 2D). Small metamorphically induced changes to Sm/Nd ratios at 2.7 Ga will have significant effect on the evolution of 143Nd as 147Sm continues to decay to the present day. The same is not true for the ¹⁴⁶Sm-¹⁴²Nd system, however, because ¹⁴⁶Sm was long extinct by 2.7 Ga. Thus, late metamorphic changes to the Sm-Nd system will affect the ¹⁴⁷Sm-¹⁴³Nd and ¹⁴⁶Sm-¹⁴²Nd results differently. The only way to change the ¹⁴²Nd/¹⁴⁴Nd ratio after ¹⁴⁶Sm has become extinct is to involve mixing with a fluid/melt having a different ¹⁴²Nd isotopic composition. Changes in Sm/Nd ratio after ¹⁴⁶Sm is gone will not affect the ¹⁴²Nd isotopic composition, but will affect the slope of a Sm/Nd vs. 142Nd/144Nd correlation, which we show below will have only minimal affect on the ¹⁴⁶Sm-¹⁴²Nd age obtained.

Fig. 11 shows the effects of a complex metamorphic history on the ¹⁴⁷Sm-¹⁴³Nd system and the robustness of the ¹⁴⁶Sm-¹⁴²Nd system. Here we model two scenarios for a 2.7 Ga disturbance in 4.4 Ga rocks and examine the relative effects on both long-lived and short-lived Sm-Nd isotopic systems. The circles in Fig. 11 represent two undisturbed 4.4 Ga Ujaraaluk rocks, a low Sm/Nd rock (U1, U2, U3) and a high Sm/Nd rock (U4, U5, U6), at three different times, from t = 0, or 4.4 Ga ago (U1 and U4) to today (U3 and U6). In the first scenario (Fig. 11A), at 2.7 Ga (or 1.7 Ga after t = 0), the undisturbed rocks (U2 and U5) are affected by a thermal event and partial melting, producing the ~2.7 Ga pegmatites (yellow stars). The analyzed pegmatite sample has a deficit in ¹⁴²Nd despite the fact that it was produced after 4.0 Ga. Thus the most likely source of the pegmatites is a rock that already contained a ¹⁴²Nd deficit, such as the Uiaraaluk unit. Melting of the Uiaraaluk unit at 2.7 Ga produces restites with higher Sm/Nd ratios (R1 and R3) and mixing between the pegmatitic liquid and the Ujaraaluk also produces rocks with lower Sm/Nd ratios (M1 and M3). In this case, the two hypothetical Ujaraaluk starting samples are subjected to the same melting scenario and produce their own pegmatitic liquids having the same isotopic composition as their respective parent rocks at the time of melting. The restites and mixed rocks evolve to give ¹⁴⁷Sm-¹⁴³Nd isochron ages of 2.7 Ga (M2-U3-R2 and M4-U6-R4). Sm/Nd ratios for the restites were calculated using batch melting equations. The composition of the pegmatite (sample R18) was used as the liquid composition whereas the composition of Ujaraaluk sample PC-129 was used as the original unmelted composition of the low Sm/Nd rock (U2) as it falls close to the ¹⁴²Nd isochron in Fig. 10A and has the same ¹⁴²Nd/¹⁴⁴Nd ratio as the pegmatite. The variation in Sm/Nd ratio obtained for the low Sm/Nd restite (R1) was used to calculate the Sm/Nd ratio of the high Sm/Nd restite (R3). If the restites are produced from 15% melt extraction of both Ujaraaluk rocks, and the mixed rocks are formed from 30% mixing with the pegmatite, this would translate today to a 5% increase in Sm/Nd for the restite (U6-R4) and 6% decrease in Sm/Nd for the mixed rock (U3-M2). Increasing the Sm/Nd ratio of the high Sm/Nd rocks while decreasing the Sm/Nd ratio of the low Sm/Nd rock will have the largest effect on the ¹⁴²Nd isochron age. However, the age of the ¹⁴²Nd isochron only decreases from 4.4 Ga to 4.37 Ga (M2-R4 in Fig. 11C). Different degrees of melting lead to different Sm/Nd ratios but the effect on the ¹⁴²Nd isochron remains small. Using 5% and 25% melting for the low Sm/Nd rock (R5) and the high Sm/Nd rock (R7)

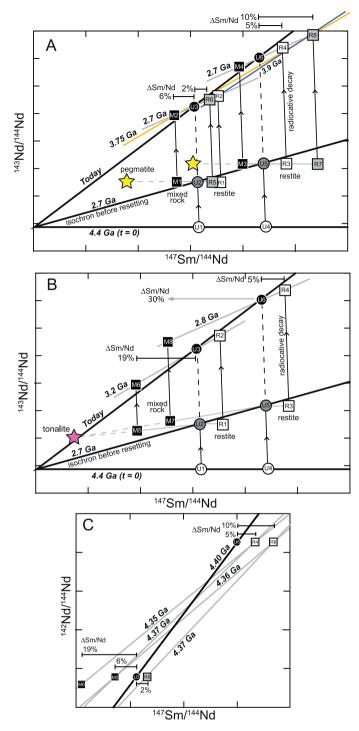


Fig. 11. (A) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴³Nd/¹⁴⁴Nd diagram showing the effect of a 2.7 Ga isotopic disturbance after partial melting of the Ujaraaluk unit and mixing with a pegmatitic liquid. (B) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴³Nd/¹⁴⁴Nd diagram showing the effect of a 2.7 Ga isotopic disturbance after partial melting of the Ujaraaluk unit and mixing with a 3.66 Ga tonalitic liquid. (C) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd diagram showing the effect of the different Sm/Nd variations illustrated in (A) and (B) on the short-lived isotopic system. Symbols: circles, undisturbed Ujaraaluk; R1–R8 squares, restites; M1–M8 squares, mixed rocks, yellow star, pegmatitic liquid; pink star, tonalitic liquid. (Fer interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)

respectively, the ¹⁴³Nd isochron for the restites yields an age of 3.9 Ga (R6–R8), while the ¹⁴²Nd isochron age is lowered to 4.37 Ga. A ¹⁴³Nd isochron passing through the high Sm/Nd restite and the low Sm/Nd mix rock (M2-R8) gives an age of 3.75 Ga, while the ¹⁴²Nd age decreases to 4.36 Ga.

The second scenario (Fig. 11B) involves partial melting of the Ujaraaluk at 2.7 Ga producing the same restites as in Fig. 11A (R1, R3) but the mixed rocks are formed by mixing between the undisturbed rocks and an enriched tonalite (M5, M7). The tonalite isotopic composition used for the model is in the middle of the range of isotopic compositions measured for the Nuvvuagittuq tonalities. This tonalite has a deficit in ¹⁴²Nd and therefore will also affect the ¹⁴²Nd isotopic composition of the mixed rock. A mixture with 30% tonalitic melt produces ¹⁴³Nd isochrons ranging from 2.8 Ga to 3.4 Ga while the age of the ¹⁴²Nd isochron goes from 4.4 Ga to 4.36 Ga. The ¹⁴²Nd isochron is calculated using the high Sm/Nd restite and the low Sm/Nd mix rock (M6-R4) in order to lower the slope of the isochron. Mixing involving more tonalite will have a more important effect on the ages. A mixture with 50% tonalite (Fig. 11B) produces ¹⁴³Nd isochrons ranging from 3.2 Ga (M6-U3-R2) to 2.8 Ga (M8-U6-R4). This degree of mixing with a tonalite would result in a composition that deviates substantially from the mafic compositions of the Ujaraaluk rocks. Even this high extent of tonalite "contamination", however, only lowers the ¹⁴²Nd isochron age to 4.35 Ga (M6-R4 in Fig. 11C). Although there is little evidence that the 3.6 Ga tonalite mixed with the Ujaraaluk at 2.7 Ga, the Ujaraaluk is enclosed by the tonalite at the time of the peak metamorphism and the tonalite represents an obvious potential contaminant. Despite the fact that this tonalitic contaminant would significantly disturb the ¹⁴⁷Sm-¹⁴³Nd systematics of the Ujaraaluk rocks, the ¹⁴²Nd ages are relatively unaffected, reinforcing the robustness of the short-lived isotopic system even if the long-lived system is partially reset.

A few Nuvvuagittug samples show evidence of strong isotopic disturbance consistent with the most extreme cases illustrated in Fig. 11. Two high-Ti Ujaraaluk (PC-058, PC-451) and one depleted low-Ti Ujaraaluk (PC-214) samples lie well off of the Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd correlations defined by the remaining samples. These samples are interpreted as being strongly disturbed. Samples PC-058 and PC-451 exhibit LREE-enriched profiles strikingly different from the rest of the high-Ti Ujaraaluk samples (Fig. 12A). They also have higher Nd concentrations consistent with them being affected by LREE-enriched fluids during post-magmatic alteration or metamorphism. Both PC-058 and PC-451 fall to the low Sm/Nd side of the ¹⁴⁶Sm-¹⁴²Nd isochron (Fig. 12A). Though their Sm/Nd ratios are lower than other high-Ti Ujaraaluk samples, their ¹⁴²Nd isotopic compositions are similar to the least affected high-Ti Ujaraaluk samples with flat REE profiles. Ujaraaluk sample PC-058 exhibits the most extreme offset from the Sm/Nd-142Nd correlation defined by the majority of the Ujaraaluk samples. The composition of this sample can be reproduced by a disturbance similar to the scenario involving 50% mixing with a tonalitic melt (Fig. 11B). Its strong LREE-enriched profile can be reproduced by 50% mixing of an undisturbed high-Ti Ujaraaluk and a Nuvvuagittuq tonalite (Fig. 12B). A 50% mixture of a high Sm/Nd rock and a tonalite produces a 30% change in Sm/Nd (U6-M8), consistent with the difference in Sm/Nd between PC-058 and an undisturbed sample falling on the ¹⁴²Nd isochron (35% variation; Fig. 11B). PC-214 is offset to the high Sm/Nd side of the main Ujaraaluk Sm/Nd-142Nd correlation (Fig. 12A). This sample is an orthopyroxene-rich layer within an ultramafic sill (cogenentic with the depleted low-Ti Ujaraaluk) that exhibits a metamorphic texture with $\sim 120^{\circ}$ grain boundaries and has a strongly depleted REE profile (Fig. 12A). Formation of a metamorphic orthopyroxenite after 4 Ga will increase the Sm/Nd ratio without affecting the ¹⁴²Nd/¹⁴⁴Nd ratio. Fig. 12A shows that this sample has been horizontally shifted to the high Sm/Nd side of the Ujaraaluk array with a ¹⁴²Nd isotopic composition that remained in the same range as the other ultramafic sill values. A $^{\rm 147}{\rm Sm}{\rm -}^{\rm 143}{\rm Nd}$ isochron including ultramafic samples, PC-214 and other pyroxene layers within the same sill yields an age of 2763 Ma (Fig. SD5) supporting the fact that the orthopyroxene layers were formed

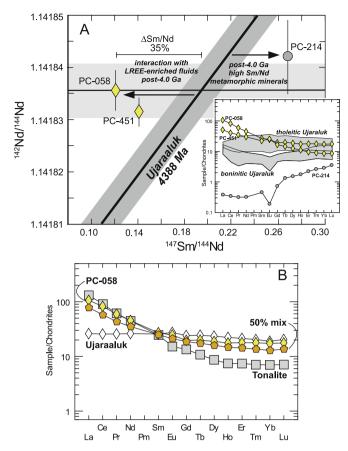


Fig. 12. (A) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd diagram showing the effect of post-4.0 Ga disturbances on some NGB samples. Oblique black line shows the best fit line of Fig. 10A. Gray bands show the 4.5 ppm external error on the terrestrial standard value and the best fit line. Inserted diagram shows the REE profiles of the most disturbed Ujaraaluk samples compared to their respective geochemical groups. (B) REE profiles for a Ujaraaluk sample, a tonalite sample, PC-058 (most disturbed sample) along with the composition of 50% mix between the Ujaraaluk and the tonalite. Chondritic normalization values are from Sun and McDonough (1989).

during the Neoarchean metamorphism. These three strongly disturbed samples (PC-058, PC-451 and PC-214) fall significantly off their respective 147 Sm/ 143 Nd and 146 Sm- 142 Nd isochrons and have been considered as outliers and excluded from both the 143 Nd and 142 Nd age calculations.

In summary, although metamorphism has disturbed the longlived ¹⁴⁷Sm-¹⁴³Nd isotopic system in the Ujaraaluk, the effects on the short-lived ¹⁴⁶Sm-¹⁴²Nd system are not significant. In contrast, the central gabbro sills yield ¹⁴³Nd and ¹⁴²Nd isochrons with Hadean ages between 4.1 and 4.3 Ga, suggesting that the ¹⁴³Nd isotopic system has not been reset to the same extent in the gabbro sills as it has in the Ujaraaluk unit. This may reflect their different lithology. The Ujaraaluk unit is interpreted to be a hydrothermally altered pyroclastic volcanic deposit (O'Neil et al., 2011a), whereas the gabbro sills are massive intrusions that are perhaps more resistant to disturbance by metamorphic fluids/melts. The magnitude of metamorphic disturbance that would turn 4.4 Ga ¹⁴⁷Sm-¹⁴³Nd ages into the scatter of younger, 3.8-2.7 Ga, ages seen in the various compositional groups of the Ujaraaluk would reduce the ¹⁴⁶Sm-¹⁴²Nd ages by only of order 50 Ma or less. Thus, while there is no escaping the conclusion that Archean metamorphism has disturbed the Sm-Nd geochronometer in the Ujaraaluk rocks, the discordance between the ¹⁴⁷Sm-¹⁴³Nd and ¹⁴⁶Sm-¹⁴²Nd results does not necessarily imply that the ¹⁴⁶Sm-¹⁴²Nd ages are significantly inaccurate. In fact, in our opinion, the types of metamorphic disturbance modeled in Fig. 11 provide a simpler and more geologically plausible explanation to the discordant ¹⁴⁷Sm-¹⁴³Nd and ¹⁴⁶Sm-¹⁴²Nd ages than do the complicated source mixing models considered in Section 5.1.2. We suggest that the discordance between ¹⁴³Nd and ¹⁴²Nd ages provides additional support for our conclusion that the Ujaraaluk rocks formed by partial melting of "normal" depleted mantle in the Hadean, rather than by Eoarchean melting of multiple Hadean mantle sources followed by some complicated mixing/fractionation scenario that would obliterate the Hadean ages from the ¹⁴⁷Sm-¹⁴³Nd system, but preserve them in ¹⁴⁶Sm-¹⁴²Nd.

5.3. Origin of the NGB felsic rocks

The Nuvvuagittug felsic rocks include the 3.8 Ga trondhiemitic bands, the 3.6 Ga tonalites and the 2.7 Ga pegmatites, all Archean in age. Despite the fact that they formed after the extinction of ¹⁴⁶Sm. most Nuvvuagittug felsic rocks have a resolvable deficit in ¹⁴²Nd (Fig. 5). Negative ¹⁴²Nd anomalies in Archean rocks indicate that their precursor had a deficit in ¹⁴²Nd at the time of their formation. The obvious candidates for precursors for the Nuvvuagittug felsic rocks are the mafic basement rocks of the Ujaraaluk unit. Adam et al. (2011) conducted melting experiments on samples of the Ujaraaluk unit and obtained partial melts that match the compositions of the Nuvvuagittuq tonalites. In the field, enclaves a few meters wide by tens of meters long of mafic amphibolites are observed within the tonalite. The size and number of these mafic enclaves in the tonalite increase as the contact between the tonalite and the Nuvvuagittuq belt is approached and seams of tonalite occur in the mafic lithologies adjacent to the tonalite contact.

All Nuvvuagittuq felsic rocks plot to the low Sm/Nd ratio side of the ¹⁴²Nd isochron defined by the Ujaraaluk. Partial melting in the Archean of any mafic Ujaraaluk composition will produce a felsic magma with a lower Sm/Nd ratio but with the same ¹⁴²Nd/¹⁴⁴Nd ratio as their parent rock. The Nuvvuagittuq tonalities yield μ^{142} Nd values ranging from -6 to -16 ppm with no correlation between their Sm/Nd and ¹⁴²Nd/¹⁴⁴Nd ratios (Fig. 13) consistent with their derivation from the melting of the mafic Ujaraaluk unit post-4.0 Ga. The Ujaraaluk unit exhibits a wide range of ¹⁴²Nd isotopic

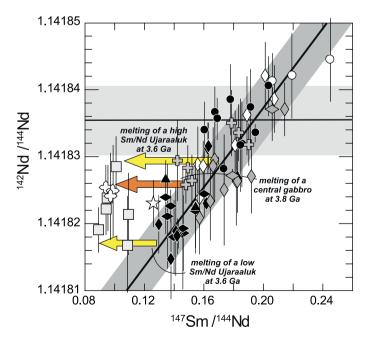


Fig. 13. ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd diagram showing the effect of post-4.0 Ga partial melting of mafic Ujaraaluk to produce the NGB felsic rocks. Gray bands show the 4.5 ppm external error on the terrestrial standard value and the best fit line. Symbols as in Fig. 5.

compositions, thus partial melting of Ujaraaluk rocks would produce tonalites spanning the same range in ¹⁴²Nd/¹⁴⁴Nd, but with lower Sm/Nd ratios which causes the tonalites to be displaced to the left of the Ujaraaluk 146Sm-142Nd isochron. Moreover, a ¹⁴⁷Sm-¹⁴³Nd isochron for the tonalities yields an age of 3575 Ma, consistent with their U–Pb age, but with an initial ε Nd value of -2.9. This negative ε Nd value at 3575 Ma indicates a much older, low Sm/Nd ratio, precursor for the tonalite. In fact, the initial ¹⁴³Nd isotopic composition for the tonalites falls on the isotopic evolution line of a 4.4 Ga mantle-derived mafic precursor (Fig. 14) having an isotopic composition consistent with the Ujaraaluk unit. An Hadean source rock for the tonalites is further supported by their T_{DM} values ranging from 3.8 Ga to 4.4 Ga. The trondjhemitic bands and the pegmatites also have deficits in ¹⁴²Nd with T_{DM} values ranging from 3.8 Ga to 4.2 Ga. These felsic rocks fall on the low Sm/Nd ratio side of the Ujaraaluk ¹⁴⁶Sm-¹⁴²Nd isochron (Fig. 13) suggesting that they also may have been formed by the partial melting of the Ujaraaluk unit or the central gabbros during later events.

5.4. Other Eoarchean components in the NGB

Beside the trondhjemetic bands and the tonalites, zircons from rocks interpreted to be detrital sediments by some authors (David et al., 2009; Cates et al., 2011; Darling et al., 2012) have yielded Eoarchean U-Pb ages. These rocks include quartzites with 3.78 Ga zircons (Cates et al., 2011), putative conglomerates with distinct zircon populations with ages of 3.78 Ga, 3.66 Ga, 3.36 Ga and \sim 2.7 Ga (David et al., 2009) and rocks interpreted to be guartzalbite-clinozoisite metasediments with concordant zircon ages ranging from 3630 to 3680 Ma. The NGB, however, has been subjected to a complex thermal history and extensive metasomatism sometimes accompanied by the formation of younger metamorphic/metasomatic zircon as seen in the central gabbro (Fig. 6). Understanding the exact geological context of a given lithology, its position in the NGB stratigraphy, and the origins of the zircons it contains thus is critical to the correct interpretation of the zircon age constraints. The detrital origin of the quartz-rich rocks in the NGB is difficult to conclusively demonstrate due to their intensive deformation and recrystallization. The conglomeratic units consist of 0.5-2 m thick horizons containing rounded quartz clasts in a biotite-bearing matrix. The quartz grains in the clasts have ¹⁸O-enriched compositions and the trace sulfides in the clasts yield significant mass-independent S isotope fractionation consistent with a sedimentary origin for the quartz and sulfide grains (Kitayama et al., 2011). ¹⁸O-enriched compositions and mass-independent S isotope fractionation only suggest a sedimentary origin for the quartz and sulfide grains contained in the conglomeratic unit, but do not require a detrital origin for of the host rock. On the other hand, the homogeneity in clast composition is consistent with a single local source, most likely the prominent silica-formation described in O'Neil et al. (2007) interpreted to be a chemical sediment. Kitayama et al. (2011) interpret the conglomeratic unit as a structural mélange of restricted local lithologies, in agreement with the fact that they seem to occur principally in high strain zones. This would suggest that the zircons in the conglomeratic unit do not place any age constraints on the age of the emplacement of the mafic units of the NGB. The similarity in isotopic compositions between the conglomeratic units and the quartz-rich rock interpreted by Cates et al. (2011) to be a detrital guartzite, and the fact that they both contain zircons of the same age is consistent with a similar process for their formation and a similar source for the quartz and the sulfide. Thus, we consider that the quartz-rich rocks do not put geochronological constraints on the age of emplacement of the Ujaraaluk unit. Moreover, all 3.8 Ga zircon bearing rocks are found to the south of the NGB within less than 100 m of the contact with the tonalite, often in highly deformed

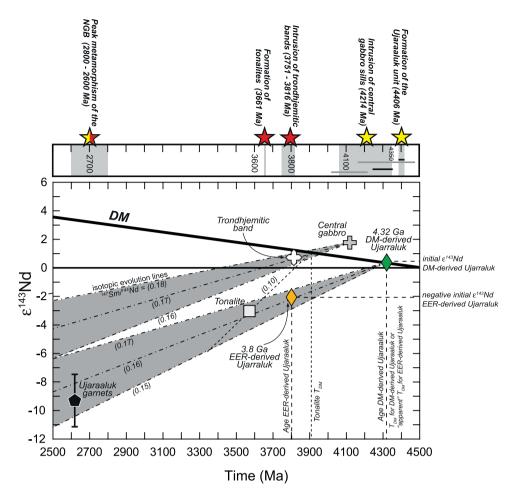


Fig. 14. ε^{143} Nd vs. time isotopic evolution diagram. Symbols: green diamond, 4.3 Ga Ujaraaluk derived from the DM; orange diamond, 3.8 Ga Ujaraaluk derived from the EER; gray plus, central gabbro; open plus, trondhjemitic band; gray square, tonalite; black pentagon, Ujaraaluk garnet. Age and ε^{143} Nd data for the gabbro, the tonalite and the Ujaraaluk garnet are from the isochrons of Figs. 2E, F and 4B respectively. Age and ε^{143} Nd data for the DM-derived Ujaraaluk are from the isochron diagram of Fig. 15A. ε^{143} Nd data for the trondhjemitic band is from the average whole rock isotopic composition of 3 samples (Table 1). The initial ε^{143} Nd value of ~ -2 for the 3.8 Ga EER-derived Ujaraaluk was determined considering a 3.8 Ga EER-derived rock with a 147 Sm/ 144 Nd of 0.16 (average Ujaraaluk value) that would give an "apparent" T_{DM} of ~ 4.3 Ga. Gray areas are 147 Sm/ 144 Nd isotopic evolution lines ranging from 0.18 to 0.16 for the gabbro and from 0.17 to 0.15 for the Ujaraaluk. Symbols on the time scale: red stars are U–Pb ages and yellow stars are Sm/Nd isochron ages. Gray vertical areas represent age intervals for the different NGB lithologies. The age interval for the metamorphism is estimated using mineral separates and U–Pb ages for the different metamorphic minerals and associated rocks. The age interval for the tonalite represents the error on the U–Pb age (David et al., 2009). The age interval for the trondhjemitic band represents the age difference between David et al. (2009) and Cates and Mojzis (2007) data. The age interval for the gabbro anges (see Fig. 15). Gray horizontal lines are 143 Nd isochron ages and errors for the gabbro and Ujaraaluk unit. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of the article.)

and sheared areas. The 3.8 Ga zircons could therefore have been formed during an episode of intense deformation responsible for the formation of shearing and structural mélange zones consistent with the characteristics of the conglomeratic units. Although the \sim 3800 Ma intrusions of trondhjemetic bands place a minimum age on the Ujaraaluk unit, the Ujaraaluk also seems to contain rocks interpreted as metasediments that have 3630–3680 Ma zircon ages (Darling et al., 2012), suggesting deposition on a much older basement. This emphasizes the complex geologic history of the NGB and shows that ages obtained on rare zircon-bearing felsic rocks have to be interpreted with caution in terms of the constraints they provide on the age of the dominant mafic lithologies.

5.5. Proposed time scale for the NGB

In this section, we summarize all geochronological evidence to establish a time scale for the NGB and to constrain its secular evolution (Fig. 14). The minimum age of 3.75–3.82 Ga for the NGB was previously established from U–Pb data on the zircons in the felsic trondhiemitic bands (Cates and Moizsis, 2007: David et al., 2009). However, the NGB also is intruded by gabbro sills that yield ¹⁴³Nd and 142 Nd isochron ages at ~4.2 Ga which pushes back the minimum age of the belt into the Hadean. The Ujaraaluk unit is intruded by the central gabbros and therefore must be older than \sim 4.2 Ga. The only model that adequately explains all aspects of the ¹⁴²Nd isotopic variation within the comagmatic Ujaraaluk and ultramafic sills is the one in which the Ujaraaluk unit formed while ¹⁴⁶Sm was still actively decaying. Therefore, the correlations between the Sm/Nd and ¹⁴²Nd/¹⁴⁴Nd ratios for each of the Ujaraaluk compositional groups may represent isochrons dating the formation of the Ujaraaluk at ~4.4 Ga. The Ujaraaluk unit is surrounded by the 3.66 Ga Nuvvuagittuq tonalities that apparently formed by the partial melting of the Hadean mafic Ujaraaluk unit from which they inherit their ¹⁴²Nd deficit (Fig. 13). The initial ε^{143} Nd value indicated by the ¹⁴⁷Sm-¹⁴³Nd isochron for tonalites falls on the 143 Nd evolution line of the \sim 4.4 Ga Ujaraaluk unit (Fig. 14). The trondhjemitic bands also have a ¹⁴²Nd deficit and may as well have been formed by the partial melting of the Ujaraaluk unit.

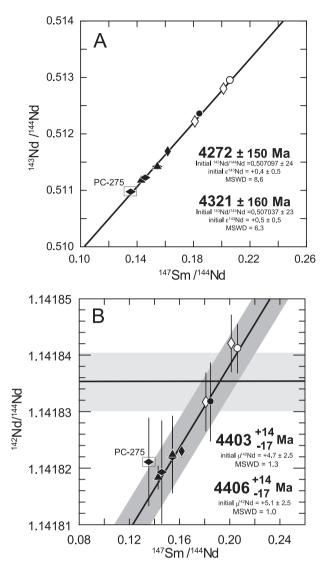


Fig. 15. (A) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴³Nd/¹⁴⁴Nd isochron diagram and (B) ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd isochron diagram for the least disturbed Ujaraaluk samples and cogenetic ultramafic rocks. Gray bands show the 4.5 ppm external error on the terrestrial standard value and the best fit line on the ¹⁴⁷Sm/¹⁴⁴Nd vs. ¹⁴²Nd/¹⁴⁴Nd isochron diagram.

However the average initial ε^{143} Nd value of the 3 trondhjemitic band samples (+0.8) falls closer to the evolution line of the central gabbros suggesting that their source may instead be the 4.2 Ga gabbros with μ^{142} Nd values overlapping with the μ^{142} Nd values for the trondhjemitic bands (Fig. 14). The NGB was then metamorphosed at ~2.7 Ga with the concomitant intrusion of pegmatite and the formation of metamorphic minerals such as garnets, rutiles, zircons and monazites. The isochron for garnet separates gives an initial ε^{143} Nd value falling on the ¹⁴³Nd evolution line of the 4.4 Ga Ujaraaluk unit consistent with their crystallization from a parent rock that was an mafic Hadean rock before metamorphism (Fig. 14).

The 2.7 Ga thermal event also affected the long-lived Sm–Nd isotopic system for some of the Ujaraaluk unit resulting in poorly defined isochrons ranging from 3.7 Ga to 2.5 Ga and a wide range in Sm–Nd model ages (Fig. 3A). A population of Ujaraaluk samples, however, has T_{DM} values near 4.3 Ga (10 samples with 4.2 Ga $\leq T_{\text{DM}} \leq 4.4$ Ga; Fig. 3A). This cluster of samples with Hadean model ages could reflect one of three possibilities: (1) they are derived from a "normal" DM at ~4.3 Ga and have suffered little or no later modification of their Sm–Nd systems, (2) they were

derived from an enriched (negative ε^{143} Nd; EER in Fig. 14) source at 3.8 Ga or later, or (3) their Sm/Nd ratios have been metamorphically disturbed, resulting in erroneously old T_{DM} model ages. In case 1, the ¹⁴⁷Sm-¹⁴³Nd isochron defined by these samples would yield an age of 4.3 Ga with an initial ε^{143} Nd value of \sim +0.5. In both cases 2 and 3, the ¹⁴⁷Sm-¹⁴³Nd isochron for this group of samples would yield a younger (2.5-3.8 Ga) age with a negative initial ε Nd value (Fig. 14). Fig. 15A shows a ¹⁴⁷Sm–¹⁴³Nd isochron for the sub-population of samples with T_{DM} values between 4.2 Ga and 4.4 Ga. This isochron yields an age of 4272 ± 150 Ma with an initial ε^{143} Nd value of +0.4 ± 0.5 consistent with derivation from a "normal" DM at \sim 4.3 Ga and not from an enriched missing source at 3.8 Ga. This suggests that this population of Ujaraaluk samples have the least disturbed ¹⁴⁷Sm-¹⁴³Nd systematics. The ¹⁴⁶Sm-¹⁴²Nd isochron plotting only these least disturbed samples gives an age of 4403^{+14}_{-17} Ma (Fig. 15B), with very little scatter. As for the initial ϵ^{143} Nd value, the initial μ^{142} Nd value of +4.7 ± 2.5 for this isochron is consistent with derivation from the DM. Following the reasoning that post-4.0 Ga disturbance affects the Sm/Nd ratio without affecting the ¹⁴²Nd/¹⁴⁴Nd ratio (i.e. a horizontal shift on the isochron diagram), samples falling further from the ¹⁴²Nd isochron have the most affected ¹⁴⁷Sm/¹⁴⁴Nd and thus the most affected ¹⁴³Nd/¹⁴⁴Nd. In Fig. 15B, one of the least disturbed samples (PC-275) plots further from the isochron than the rest of the data, which we interpret as indicating that it has a slightly disturbed Sm-Nd system. Taking this single sample off the isochrons of Fig. 15, the ¹⁴³Nd age becomes 4321 ± 160 Ma (MSWD = 6.3), whereas the ¹⁴²Nd age becomes only 3 million years older (i.e. 4406^{+14}_{-17} Ma; MSWD = 1.0). While this subset of samples admittedly has been highly screened in an effort to identify the most pristine samples, nevertheless, this set of samples has consistent Sm-Nd systematics for both the long-lived and short-lived systems both of which give overlapping Hadean ages. Thus, our best estimate for the age of the Ujaraaluk unit is 4406^{+14} $_{-17}$ Ma, which makes it the oldest vestige of crust preserved on Earth. The age of the Ujaraaluk unit overlaps those of the oldest Jack Hills zircons (Wilde et al., 2001) and the oldest zircons from the Moon (Nemchin et al., 2009), suggesting the intriguing possibility that the Nuvvuagittug Greenstone Belt may preserve a remnant of the first crust formed on Earth after the giant impact that formed the Moon.

6. Conclusions

The 142Nd data set for the Nuvvuagittuq Greenstone Belt rocks has been increased to more than 70 samples. The range in 142 Nd/ 144 Nd ratios is now 26 ppm (-18 ppm to +8 ppm) and the expanded dataset for the Ujaraaluk rocks and associated ultramafic cumulates continues to show a good correlation between the Sm/Nd and the ¹⁴²Nd/¹⁴⁴Nd ratios for each compositional group. Although a complex mixing model involving an early depleted reservoir and two never-seen-before early enriched reservoirs conceivably could explain this correlation, such a multiple source mixing model cannot account for several characteristics of the Nuvvuagittug rocks. (1) The major and trace element compositions of the Ujaraaluk are not consistent with simple mixing between a single depleted and a single enriched reservoir. (2) All the compositionally distinct Ujaraaluk groups follow the exact same Sm/Nd vs. ¹⁴²Nd correlation. The fact that the composition of these groups indicate their derivation from at least three sources with distinct geochemical features followed by distinct fractional crystallization histories under different conditions makes it unlikely that any simple mixing scenario would simultaneously produce identical Sm/Nd vs. ¹⁴²Nd/¹⁴⁴Nd correlations in all three groups. (3) In a mixing scenario, a correlation between the ¹⁴²Nd and ¹⁴³Nd isotopic compositions with Nd concentration would be expected at the time the mixing occurred, as would "fossil" ¹⁴⁷Sm⁻¹⁴³Nd

correlations giving ages older than the presumed mixing event. None of these features are present in the Ujaraaluk isotopic data. (4) A mixing event or a melt-restite relationship in the Eoarchean cannot explain the Sm/Nd vs. ¹⁴²Nd correlation observed between the mafic Ujaraaluk and their cogenetic ultramafic cumulates. (5) An Eoarchean formation age for the Ujaraaluk cannot be reconciled with the intruding gabbros yielding ¹⁴²Nd and ¹⁴³Nd isochrons with Hadean ages.

All these features, however, can be explained by a simple model where the Sm/Nd vs. ¹⁴²Nd correlation is produced by igneous fractionation of the Sm/Nd ratio at constant Nd isotopic composition at a time when ¹⁴⁶Sm was still actively decaying, leading to ¹⁴²Nd isochron ages of ~4.4 Ga. The least disturbed Ujaraaluk samples yield a 142 Nd isochron age of $4406 + 14_{-17}$ Ma and a 143 Nd isochron age of 4321 ± 160 Ma supporting an Hadean age. Intruding gabbros give overlapping ¹⁴³Nd and ¹⁴²Nd isochron ages with a mean of 4214 ± 140 Ma providing additional support for an Hadean age for the NGB. Therefore, the ¹⁴⁷Sm–¹⁴³Nd and ¹⁴⁶Sm–¹⁴²Nd isotopic compositions in the mafic lithologies of the Nuvvuagittuq Greenstone Belt suggest that it represents the oldest crustal remnant on Earth. Following Hadean formation, the NGB was subjected to a complex structural and metamorphic reworking recording over a billion years of Earth's Archean history. Despite the fact that this metamorphic history has affected the long-lived ¹⁴⁷Sm-¹⁴⁴Nd isotopic system, the short-lived 146Sm-142Nd system has not been significantly affected.

The geological evolution of the NGB can be summarized as follows:

- (1) 4.3–4.4 Ga: Eruption and crystallization of the Ujaraaluk unit and comagmatic ultramafic sills. The Ujaraaluk unit exhibits a transition from basaltic rocks having tholeitic affinities to basaltic rocks that resemble modern boninites to basaltic/andesitic rocks having calk-alkaline affinities.
- (2) 4.2 Ga: Intrusion of gabbro sills in the central part of the NGB.
- (3) 3.8 Ga: Intrusion of trondhjemitic bands produced by the melting of the mafic lithologies. Possible development of shear zones and sutures composed of recrystallized quartz that locally resembles conglomerate texture.
- (4) 3.66 Ga: Formation of the surrounding NGB tonalites by partial melting of the Hadean mafic lithologies.
- (5) ~3.3 Ga: Possible metamorphic event recorded in a small population of zircons (David et al., 2009).
- (6) ~2.7 Ga: Extensive metamorphism concomitant with the intrusion of pegmatites, formation of metamorphic minerals including garnet, rutile, monazite and zircon in the Ujaraaluk unit, orthopyroxene in the ultramafic sills and zircons in the gabbro sills.

Despite the fact that the Nuvvuagittuq Greenstone Belt was formed less than 300 million years after the beginning of Earth's accretion, it shares geochemical affinities with numerous younger Archean greenstone belts. This suggests that the geological processes still responsible for forming crust were established very early in the Earth's history. Nevertheless, evidence for the existence of stable crust shortly after the Moon forming impact is becoming more abundant, with the 4.3–4.4 Ga Nuvvuagittuq Ujaraaluk and the 4.4 Ga Jack Hills zircons being contemporaneous with the formation of the lunar crust at 4.36–4.41 Ga (Borg et al., 2011; Nemchin et al., 2009).

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