SIMS zircon ages and Nd isotope systematics of the 2.2 Ga mafic intrusions in northern and eastern Finland



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Abstract

Using the SIMS, ID-TIMS and Sm-Nd isotopic methods and the electron microprobe, we have studied several differentiated mafic intrusions of the c. 2.2 Ga gabbro-wehrlite association (GWA) from four Paleoproterozoic schist belts and the Archean Kuhmo Greenstone Belt. Back-scattered electron images and electron microprobe analyses revealed that zircon crystals vary from well-preserved to turbid and highly altered with individual grains often displaying irregular, hydrated, CaO-bearing domains. In the most pristine domains, suitable for establishing the crystallization ages, SIMS ²⁰⁷Pb/²⁰⁶Pb ages fall in the range of 2210-2220 Ma, which is consistent with the most concordant ID-TIMS U-Pb ages. One of the studied intrusions that had previously yielded a conventional U-Pb date of less than 2.0 Ga, could be shown by spot analysis to belong to the 2.2 Ga family. In contrast to the well-preserved zircon domains, altered domains exhibit a variable and often strong U-Pb discordance up to 70 % and have distinctly lower ²⁰⁷Pb/²⁰⁶Pb ages. Some zircon grains record isotopic resetting at the time of the Svecofennian orogeny (ca. 1.8-1.9 Ga), while the most discordant ones project in the concordia diagram to late Paleozoic lower intercept ages indicating a relative recent Pb loss. The mineral chemistry of zircon suggests that the leakage of radiogenic Pb can be ascribed to an opensystem behavior related to hydrothermal alteration via action of CaCl₂-bearing fluids.

Common albitization of plagioclase in the GWA intrusions has caused this mineral to behave as an open system with regard to the Sm-Nd isotopic systematics. Despite this uncertainty, our Nd isotopic data indicate that the magma that produced the GWA intrusions in various parts of northern and eastern Finland was isotopically homogeneous and had an initial $\epsilon_{_{\rm Nd}}(2220 \text{ Ma})$ value of c. +0.6 precluding significant upper crustal contamination upon emplacement and subsequent fractional crystallization.

Key words: intrusions, gabbros, granophyre, absolute age, U/Pb, Sm/Nd, zircon, electron probe data, Paleoproterozoic, Northern Finland, Eastern Finland

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1. Introduction

One of the distinct phases of the Paleoproterozoic mafic magmatism in Finland is represented by the c. 2.2 Ga layered sills, which are widely spread in eastern and northern Finland (Vuollo & Huhma, 2005). On the basis of the predominant rock types, Hanski (1986a, b, 1987) assigned these intrusive rocks to the gabbro-wehrlite association (GWA). Later the rocks have also been called karjalites (Vuollo & Piirainen, 1992).

Since the beginning of the 1970s, coarse-grained or pegmatoidal gabbros in the upper parts of the GWA sills have served as favorable targets for dating purposes as they contain abundant zircon (Sakko, 1971; Kouvo, 1977). Occasionally, baddeleyite has also been found in these rocks (e.g. Hyppönen, 1983; Perttunen & Vaasjoki, 2001). Hanski et al. (2001) compiled available ID-TIMS U-Pb analyses of zircon and baddeleyite from the dated GWA sills from eastern and northern Finland. Even though the apparent ID-TIMS ages of the dated intrusions have a spread of more than 200 Ma, they concluded that the time interval of the magmatic event that produced these rocks was probably considerably shorter. The reason for this ostensible discrepancy is the fact that analytical data from individual dating sites often display a considerable discordance on concordia diagrams (see figure 2 in Hanski et al., 2001) and therefore the interpretation of the isotopic results has not always been straightforward. Under the microscope, zircon crystals from the GWA intrusions commonly display a very turbid appearance indicative of a high degree of metamictization, and conventional ID-TIMS studies have revealed that they often have high U contents (e.g. Perttunen & Vaasjoki, 2001; Hanski et al., 2001).

The high sensitivity and high spatial resolution of the ion microprobe provides a powerful tool for *in situ* dating of single zircon grains (e.g. Williams, 1998). By analysis of different domains within zircon crystals, this method allows primary crystallization ages to be determined also for the zircon populations that have experienced various degrees of radiation damage and subsequent partial opening of the U-Pb isotopic system. Six samples from northern and eastern Finland, five gabbros and one granophyre, which differ from each other in terms of the degree of post-magmatic alteration and the geological environment, were selected for in situ zircon dating at the NORDSIM facility in Stockholm. Zircon and/or baddeleyite in all these samples were previously dated using the conventional multi-grain method of isotope dilution thermal ionization mass spectrometry (ID-TIMS). We also report ID-TIMS U-Pb zircon data from two locations. In addition, we used an electron microprobe to see how different domains within zircon grains, as revealed by back-scattered electron (BSE) images, deviate chemically from each other. Only a limited number of Nd isotopic data have so far been published from the GWA intrusions (Huhma et al., 1990; Huhma et al., 1996). In this paper, we present new Nd isotopic data from several intrusions.

2. General characteristics of the GWA sills

Sill-like, gravity-differentiated intrusions of the gabbro-wehrlite association have been found in Karelian schist belts (Northern Karelia, Kainuu, Kuusamo, Peräpohja, Central Lapland) in eastern and northern Finland. Typically the sills occur close to the unconformity between the Archean granite-gneiss basement and overlying Paleoproterozoic sedimentary and volcanic rocks, usually intruding concordantly into "Jatulian" quartzitic metasediments but occasionally shifting their position to the underlying basement (Vuollo & Huhma, 2005).

Individual sills may reach several hundred meters in thickness and they can often be traced for many kilometers along strike. The maximum length of more than 100 km is attained by the Runkaus sill in the Peräpohja Belt. From the bottom upwards, the layered sequence of the sills normally comprises cumulates with the following cumulus mineral assemblages: olivine, olivine–clinopyroxene, clinopyroxene, clinopyroxene–magnetite and plagioclase–clinopyroxene–magnetite. Orthopyroxene may be present in the olivine–clinopyroxene cumulates, but its abundance is always low. The ultramafic cumulates characteristically contain magmatic edenitic hornblende as poikilitic grains. Plagioclase in gabbroic cumulates is commonly altered to secondary albite, but more calcic ones with An up to 47 % have been discovered in some intrusions in Lapland (Hanski, 1987). The parental magma of the GWA intrusions corresponded to hydrous, low-Al tholeiite (Vuollo & Piirainen, 1992), which explains the late appearance of plagioclase as a cumulus phase. The magma was characterized by a low Al_2O_3/TiO_2 ratio (5–6) and differed in this respect from other "Jatulian" mafic magmas. It was also enriched in incompatible elements and had LREEenriched chondrite-normalized REE patterns with (La/Yb)_{CN} of -4–5 (Hanski, 1986a).

3. Description of sampling sites and samples

Previously one GWA intrusion from the Kuusamo Schist Belt has been dated using the SIMS method (Evins & Laajoki, 2001). For our U-Pb and Sm-Nd study, we chose samples from intrusions occurring in four other Paleoproterozoic supracrustal belts in northern and eastern Finland: the Central Lapland Greenstone Belt, Peräpohja Schist Belt, Tahkomäki-Kinahmi schist belt, and North Karelian Schist Belt. In addition, we sampled c. 2.2 Ga intrusions found within the Archean Kuhmo Greenstone Belt. The sampling sites are shown in Fig. 1 and described below in the order of their geographical position from north to south.

3.1. Silmäsvaara, Haaskalehto, Ahvenvaara

There is a string of mafic–ultramafic bodies close to the southern margin of the Central Lapland Greenstone Belt (Fig. 1). Their occurrence can be easily discerned on geophysical maps due to the presence of magnetite-bearing gabbros and variably altered ultramafic cumulates in these bodies. Among them are the Silmäsvaara, Haaskalehto and Ahvenvaara intrusions, which are included in the present study. It is possible that these three intrusions originally belonged to the same sill-like intrusive body, which was injected concordantly into sedimentary rocks of the Sodankylä Group and was later disrupted into separate blocks due to tectonic movements (Hanski & Huhma, 2005).

The *Haaskalehto intrusion* is situated ~20 km west of Sodankylä. It can be regarded as the type occurrence of the GWA in central Finnish Lapland (Lehtonen et al., 1998). On aeromagnetic maps, the Haaskalehto intrusion can be followed along strike for c. 5 km and its thickness is estimated to be c. 0.5 km. The rock sequence is reasonably well exposed and comprises wehrlites, pyroxenites and gabbros. Despite the fact that the sill is in contact with an intrusive granite on its southern side, the original magmatic mineralogy is usually well preserved (e.g. olivine Fo_{72.6-81.6}, plagioclase up to An₄₇; Hanski, 1987).

The Haaskalehto intrusion was chosen for this study because its zircon grains are known to belong to the least altered ones that have been encountered in the GWA intrusions. Tyrväinen (1983) reported a concordant ID-TIMS U-Pb zircon age of 2220 \pm 11 Ma for the gabbro sample A892. The same sample was used for SIMS zircon analyses in this study. In addition, we utilized three gabbroic samples, A1408, 19.1-HSP-78, and 24-HSP-78, together with separated minerals (plagioclase, clinopyroxene, amphibole) for Sm-Nd isotopic analyses.

The eastern end of the *Silmäsvaara intrusion* is located c. 5 km WNW of the western end of the Haaskalehto intrusion. The sill is c. 5 km long and more than 0.5 km thick. The gabbroic rocks are not as fresh as those at Haaskalehto, but the olivine pyroxenites in the lower part of the intrusion contain portions with original magmatic minerals still abundantly present. Sampling for this study was restricted to one olivine-bearing pyroxenite sample, A1430, from which plagioclase and pyroxene were separated for Sm-Nd isotopic analysis.

The *Ahvenvaara intrusion* is located on the southern side of the Pyhätunturi Mountain c. 50 km SSE of Sodankylä. On magnetic maps, it has an appearance of a 2x3 km block that has been rotated by c. 90° relative to the general strike of the surrounding metasedimentary rocks. The exposed rock types of the intrusion are limited to olivine pyroxe-



Fig. 1. Map showing sampling sites in northern and eastern Finland. 2.2 Ga layered sills are shown as black lines. The base map modified after Koistinen et al. (2001). nites and pyroxenites. The Ahvenvaara intrusion provided a fresh olivine pyroxenite sample, A1431, and plagioclase and pyroxene separates for our Sm-Nd isotope study. It is worth mentioning that Räsänen and Huhma (2001) published a precise ID-TIMS U-Pb zircon age of 2222 \pm 6 Ma for the Harjunoja intrusion, which is a GWA-type sill located geographically between the Haaskalehto and Ahvenvaara intrusions.

3.2. Susivaara

From the Peräpohja Schist Belt 14 samples have earlier been used for conventional U-Pb dating of the gabbro-wehrlite association (Perttunen & Vaasjoki, 2001). For this study we chose one of these samples, A865 from Susivaara, which was known to display a significant spread in discordance between separate zircon fractions. The Susivaara sill is located in the western part of the Peräpohja Schist Belt, on the Törmäsvaara map sheet described by Perttunen and Hanski (2003). It forms a c. 100-mthick, weakly differentiated gabbroic body intruded concordantly into quartzites and siltstones of the Palokivalo Formation of the Kivalo Group. The same metasediment unit hosts a larger, more typical layered GWA body, the Kivimaa sill, which has also been dated (see below).

Earlier U-Pb isotope studies on the Susivaara intrusion have revealed that zircon and baddeleyite are discordant whereas sphene is concordant (Perttunen & Vaasjoki, 2001). Two concordant analyses on sphene together with three discordant analyses of baddeleyite (\pm zircon intergrowths) provided an upper intercept age of 2208 \pm 19 Ma. Perttunen and Vaasjoki (2001) reported similar results for the nearby Kivimaa sill: sphene yielded a concordant U-Pb age of 2216 \pm 8 Ma, while zircon and baddeleyite fractions were discordant though close to a chord and indicated an age of c. 2215 Ma.

3.3. Rahasmäki

The Rahasmäki sill is located in the northern part of the Tahkomäki-Kinahmi schist belt, which is a small, 1.0–3.5-km-wide and c. 40-km-long, NS- trending Karelian supracrustal belt surrounded by Archean gneisses (Paavola, 1984). There are only a few outcrops, but aeromagnetic data show that on the present erosion surface, the intrusion forms a folded, U-shaped (synclinal), 100–200-m-thick body emplaced concordantly into quartzites or occurring partly in the Archean basement. The exposed parts consist of coarse-grained metagabbro and minor metapyroxenite.

Paavola (1984) published five U-Pb analyses on zircon from a gabbroic sample, which yielded concordia intercepts at 1967 \pm 24 Ma and 425 \pm 48 Ma. However, the data do not plot exactly on a chord (MSWD = 30), which together with the geological setting of this intrusion and its lithological characteristics resembling those of the GWA intrusions has caused some uncertainty on the real crystallization age of the intrusion. In order to clarify this question, we analyzed zircon grains by SIMS from the same sample, A977, as employed by Paavola (1984).

3.4. Ensilä

The presence of Paleoproterozoic mafic intrusions within the Archean Kuhmo Greenstone Belt was indicated for the first time by Hyppönen (1983) who published age determinations of two mafic dikes from this belt. However, the discordance of the results precluded the calculation of precise crystallization ages. Later Hanski (1982, 1984) performed a comparative study of the intrusive mafic-ultramafic rocks in the Kuhmo Greenstone Belt and the "Jatulian albite diabases" in the Koli area and came to the conclusion that many "Jatulian" mafic intrusive bodies can be found within the Kuhmo Greenstone Belt. For example, a continuous differentiation series from amphibole-bearing olivine cumulates to coarse-grained gabbros, analogous of that displayed by the Koli sill, can be observed near the Ensilä farm, c. 30 km NW of Kuhmo. In addition, the GWA magmatism has generated flow-differentiated mafic-ultramafic dikes within the greenstone belt (Hanski, 1984). In the Ensilä area, the originally horizontal cumulate layering of the gravitydifferentiated bodies dip now by an angle of c. 45°

demonstrating Paleoproterozoic folding of these intrusions and the associated Archean greenstone belt as well. It is peculiar that GWA intrusions have been encountered within the relatively narrow, NS trending greenstone belt, but so far not in the adjacent granitoid basement outside the greenstone belt.

Lithological and geochemical data on the GWA sills in the Ensilä area can be found in Hanski (1984). The sills are spatially associated with the Kellojärvi Ultramafic Complex (Tulenheimo, 1999). This complex was earlier thought to belong to the GWA (Hanski, 1984), but was later shown to be Archean in age and is probably related to the komatiitic magmatism of the greenstone belt (Papunen et al., 1998).

A coarse-grained metagabbro sample, A586, was used for conventional zircon dating. The rock is moderately altered; for example, ilmenomagnetite is replaced partly by biotite and there are small garnet crystals enclosed in plagioclase. The same sample was also used for SIMS dating. Sm-Nd isotopic results on GWA rocks from Kuhmo have previously been reported by Tulenheimo (1999) and are included in our Table 4. They contain eight wholerock analyses of wehrlites, metapyroxenites and metagabbros from the Ensilä area and one wehrlite (TTT-170-96) and its two pyroxene separates from the Arola area, c. 15 km north of Ensilä.

3.5. Koli

One of the best-exposed GWA sills in Finland can be found in the Koli area, Northern Karelia. Detailed descriptions of the stratigraphic sections and geochemical and mineralogical data of the Koli sill have been published from different localities (Piirainen, 1969; Hanski, 1982, 1984; Vuollo & Piirainen, 1992). The sill can be followed for more than 60 km along strike and has a maximum thickness of 340 m. It is partly located in the Archean basement following conformably the contact between the Archean rocks and the overlying Jatulian quartzites and partly emplaced into these quartzites. It cuts the two lowermost sedimentary lithostratigraphic units, the Koli Formation and Jero Formation, but has not been observed to cut the uppermost Jatulian formation, the Puso Formation (Piirainen & Vuollo, 1991). Relatively good exposure has facilitated determination of the detailed stratigraphy of the Koli sill. From the bottom upwards, the following zones have been recognized: wehrlite, clinopyroxenite, magnetite clinopyroxenite, magnetite gabbro, coarse-grained gabbro, granophyre, clinopyroxenite (the upper marginal zone) (Vuollo & Piirainen, 1992).

Sampling for isotopic analysis was performed at a segment of the Koli sill that was emplaced into Archean gneisses. Three samples from three different localities were utilized: samples A587 and A1182 from the main gabbroic cumulate at Savilahti and Kaunislahti, respectively, and sample A1096 from Kaunisniemi, representing granophyre from the most evolved part of the sill. Baddeleyite was analyzed using ID-TIMS from sample A587 and zircon from samples A1096 and A1182. Zircon grains from the latter two samples were also studied using the NORDSIM facility.

Five samples from the Koli sill were analyzed for Sm-Nd isotopes. Whole-rock analyses were performed for two gabbro samples, 48-JIV-85 and A1182, and one granophyre sample labelled A1096a. In addition, Sm-Nd isotopic compositions were determined on plagioclase and pyroxene separates from two clinopyroxenite samples, A1220 and A1221.

4. Analytical methods

4.1. U-Pb isotopic analyses

Procedures for conventional U-Pb analyses followed the method by Krogh (1973) and involved aliquoting the HCl solution and addition of ²⁰⁸Pb/ ²³⁵U isotopic tracer. Measurements were made using a VG Sector 54 mass spectrometer at the Geological Survey of Finland (GSF), Espoo. The performance of the ion counter was checked by repeated measurements of a NBS 983 standard.

In situ U-Th-Pb analyses of zircons were carried out using a Cameca IMS1270 ion microprobe at the Swedish Museum of Natural History, Stockholm (the NORDSIM facility). The analytical pro-

cedure is described in Whitehouse et al. (1997, 1999). The spot diameter for the 4 nA primary O_2^{-1} ion was c. 30 mm, and oxygen flooding was used to improve the ionization of Pb. Calibration of the U/ Pb ratio was based on analyses of the Geostandards zircon 91500, which has an age of 1065 Ma (Wiedenbeck et al., 1995). Correction of the measured isotopic ratios for common Pb was estimated from monitored ²⁰⁴Pb counts and the terrestrial average Pb isotopic composition at 1900 Ma was used for this correction (Stacey & Kramers, 1975). Data reduction was performed using the NORDSIM software written by Martin Whitehouse, and data regressions were carried out using the Isoplot/Ex 2.49 program of Ludwig (2001). Back-scattered electron images were used to select targets for SIMS analyses.

4.2. Sm-Nd isotopic analyses

The Sm-Nd isotopic work was performed at the GSF on a VG Sector 54 mass spectrometer using the freshest samples available. Standard procedures were used for crushing and separation of plagioclase and pyroxene with final purification made by hand-picking when necessary. For Sm-Nd analyses, mineral concentrates were washed ultrasonically in warm 6 N HCl for 30 min and rinsed several times in water. The samples (150-200 mg) were dissolved in HF-HNO₃ using Savillex screw cap teflon beakers or sealed teflon bombs (felsic whole rocks) for 48 h. Mixed ¹⁴⁹Sm-¹⁵⁰Nd spike was added to the sample prior the dissolution. After careful evaporation of fluorides, the residue was dissolved in 6N HCl and a clear solution was achieved. Samarium and Nd were separated in two stages using a conventional cation exchange procedure (7 ml of AG50Wx8 ion exchange resin in a bed of 12 cm length) and a modified version of the Teflon-HDEHP (hydrogen diethylhexyl phosphate) method developed by Richard et al. (1976). The measurements have been made in a dynamic mode using Ta-Re triple filaments. 143Nd/144Nd ratio is normalized to ${}^{146}Nd/{}^{144}Nd = 0.7219$. The average value for the La Jolla standard is ${}^{143}Nd/{}^{144}Nd = 0.511850$ \pm 7 (1 σ , n = 70, triple filament measurements du-

ring 1995–2001). The Sm/Nd ratio of the spike was calibrated against the Caltech mixed Sm/Nd standard (Wasserburg et al., 1981). Based on duplicated analyses, the error in ¹⁴⁷Sm/¹⁴⁴Nd is estimated to be 0.4%. Initial $^{143}\text{Nd}/^{144}\text{Nd}$ ratios and $\epsilon_{_{Nd}}$ values were calculated with the following parameters: λ^{147} Sm = 6.54 x 10⁻¹²a⁻¹, ¹⁴⁷Sm/¹⁴⁴Nd = 0.1966 and ${}^{143}Nd/{}^{144}Nd = 0.51264$ for present CHUR. Depleted mantle model ages (T-DM) were calculated according to DePaolo (1981). Measurement on the rock standard BCR-1 provided the following values: Sm = 6.58 ppm, Nd = 28.8 ppm, ¹⁴⁷Sm/ 144 Nd = 0.1380, 143 Nd/ 144 Nd = 0.51264 ± 0.00002. The blank measured during analyses was: 30-100 pg for Sm and 100-300 pg for Nd. Programs by Ludwig (1991, 2001) have been employed for age calculations.

A few older Sm-Nd analyses measured using an old technique and a non-commercial mass spectrometer (Huhma, 1986) are included in this paper. Compared with more recent analyses, they tend to yield slightly larger errors in ¹⁴³Nd/¹⁴⁴Nd, but based on duplicated newer analyses, are consistent within error.

4.3. Electron microprobe analyses

In order to acquire preliminary data on the relationship between zircon chemistry and the observed U-Pb isotopic and BSE characteristics, different domains of a zircon grain were analyzed for major (Zr, Si) and trace (Hf, Y, Ca, Fe, Mn, U, Th, Pb) elements. Electron microprobe analyses were performed by the wavelength dispersive technique using a Cameca SX100 microprobe at the Geological Survey of Finland in Espoo. The analytical conditions were an accelerating potential of 20 keV, a sample current of 30 nA, and a beam diameter of 1 µm. Synthetic cubic zirconia was employed as the standard for Zr, Y, and Hf, while natural galena was used for Pb, diopside for Si and Ca, rhodonite for Mn, and almandine for Fe. The uranium and thorium standards were pure metals. Back-scattered electron images were obtained with a JEOL JCXA-733 scanning electron microscope (SEM) at the Department of Electron Optics, University of Oulu.

5.1. U-Pb geochronological results

Ion microprobe U-Th-Pb analytical data for zircons are listed in Table 1 and plotted on concordia diagrams in Figs. 3, 6, 8, 10 and 13. In these figures, the error boxes represent 2σ errors of isotope ratios and the uncertainties in the calculated ages are reported at the 95 % confidence level. New conventional ID-TIMS data are presented in Table 2 and, together with the previously published ID-TIMS data from the studied samples, are compared with SIMS results in the above mentioned concordia diagrams.

5.1.1. Haaskalehto (A892)

The Haaskalehto intrusion represents one of the rare locations of the GWA intrusions from which previous conventional U-Pb analyses on zircon plot relatively close to the concordia curve and thus pro-

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vide a reliable age estimate (2220 \pm 11 Ma; Tyrväinen, 1983). This is in harmony with the good preservation of the zircon grains that is evident from the BSE images taken of zircons from sample A892. As shown in Fig. 2, the grains have a weak oscillatory zoning and are remarkably clean without darker, patchy alteration patterns displayed by metamict zircons in the samples from the other study areas (see below).
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Four U-Pb analyses on four zircon grains made by ion microprobe are all concordant and give an age of 2211 ± 6 Ma (Fig. 3) which overlaps the age obtained by the conventional method. The analyses show a low common lead but a fairly large range in Th/U ratio (Table 1).

5.1.2. Susivaara (A865)

In terms of alteration, the studied zircon grains from the Susivaara intrusion represent an extreme case. Most of them are very dark brown and turbid. Backscattered electron images show complex alteration





A892 Haaskalehto



Fig. 3. Concordia plots of U-Pb data of sample A892 from the Haaskalehto intrusion, Central Lapland. SIMS analyses shown as yellow error ellipsoids and ID-TIMS analyses by blue circles without error indication. Note that the scale is different from the other concordia diagrams.

features in most zircon crystals. The altered parts of the grains are characterized by a lowered BSE intensity, i.e. a lower mean atomic number with respect to the apparently unaltered domains (Fig. 4). The BSE contrast between altered and more pristine domains is often obvious (for example, grains n654-02 and n654-03 in Figs. 5 and 4B). Even using the largest magnification of SEM, the darker, grayish domains, such as shown in Fig. 4A, appear homogeneous in BSE images without recognizable intergrowths of zircon and other phases. Some zircon crystals are completely altered to heterogeneous, spongy material (Fig. 4E). The altered domains often contain tiny thorite grains standing up as white specks in BSE images (see below).

Twelve conventional U-Pb analyses on the Susivaara sample A865 published previously by Perttunen and Vaasjoki (2001) are shown in Fig. 6. They are widely scattered. The cleaner zircon fractions and baddeleyite separates plot closer to the concordia curve than the turbid zircon fractions. Two sphene analyses are nearly concordant. Perttunen and Vaasjoki (2001) calculated an age of 2208 ± 19 Ma using baddeleyite and sphene fractions. Including discordant zircon fractions does not change the age but increases the error $(2208 \pm 29 \text{ Ma}, \text{ see Fig. 6})$.

Eleven analyses on eight zircon grains were obtained by ion microprobe. The data are technically good but scattered and discordant (Fig. 6). Interestingly, the well-preserved zircon domains have isotopic compositions relatively close to the concordia curve with ²⁰⁷Pb/²⁰⁶Pb ages of c. 2.2 Ga (n654-02a and n654-03a, Fig. 6), whereas the altered zircon zones exhibit strong U–Pb discordance with lower ²⁰⁷Pb/²⁰⁶Pb ages (ca. 1.7–2.0 Ga) n654-02b and n654-03b, Fig. 6). The U-Th-Pb data show that the altered domains tend to be enriched in Th. For example, in analysis n654-02b the Th/U ratio is 11 (Table 1).

The five most pristine domains provide an upper intercept age of 2198 ± 24 Ma (lower intercept at 154 ± 230 Ma, MSWD = 4.2, Fig. 6), which can be considered the best estimate for magmatic crystallization. The data on this sample do not constrain the age of alteration. However, all analyses on turbid zircon plot on the "younger side" of the chord defined by pristine domains. This suggests a multi-stage lead loss and possibility that major alteration relates to the 1.8–1.9 Ga thermal pulse (Hanski et al., 2001).

Sample/ spot #	[U] ppm	[Pb] ppm	Th/U meas.	f ₂₀₆ %	²⁰⁷ Pb/ ²³⁵ U	±σ %	²⁰⁶ Pb/ ²³⁸ U	±σ %	rho	Disc. % 2ơ limit	²⁰⁷ Pb/ ²⁰⁶ Pb age (Ma)	±σ	²⁰⁷ Pb/ ²³⁵ U age (Ma)	±σ	²⁰⁶ Pb/ ²³⁸ U age (Ma)	±σ
A892, Haasl	calehto,	gabbro)													
n651-01a	574	334	0.88	0.02	8.073	1.9	0.4208	1.9	0.98	**	2216	6	2239	17	2264	36
n651-02a	533	265	0.14	* 0.01	8.080	1.9	0.4234	1.9	0.99	**	2207	5	2240	17	2276	36
n651-03a	724	364	0.23	* 0.01	7.984	1.9	0.4184	1.9	0.99	**	2207	4	2229	17	2253	36
n651-04a	693	451	1.57	0.03	8.026	1.9	0.4198	1.9	0.98	**	2211	6	2234	17	2260	36
A865, Susiva	aara, me	tagabb	ro													
n654-01a	2547	1097	0.07	0.05	7.031	1.1	0.3721	1.1	0.98	-5.9	2190	4	2115	10	2039	18
n654-02a	1858	1282	2.51	0.01	7.112	1.2	0.3760	1.2	0.99	-4.8	2192	3	2126	11	2058	21
n654-02b	1498	654	11.34	1.23	1.725	2.0	0.1231	1.6	0.81	-51.7	1655	21	1018	13	748	11
n654-03a	793	470	2.27	0.18	6.296	1.2	0.3312	1.2	0.96	-16.2	2201	6	2018	11	1844	19
n654-03b	1087	479	5.65	0.88	3.100	1.3	0.1976	1.2	0.88	-37.7	1860	11	1433	10	1163	12
n654-04a	608	280	1.45	0.78	5.162	1.0	0.2999	0.9	0.88	-16.3	2026	8	1846	8	1691	13
n654-05a	1062	289	1.05	2.67	3.191	1.2	0.1911	0.9	0.77	-43.0	1973	13	1455	9	1127	9
n654-06a	1282	665	2.16	0.14	5.566	1.3	0.2978	1.3	0.98	-23.4	2171	4	1911	12	1681	19
n654-07a	1070	650	2.78	0.07	6.052	1.2	0.3233	1.1	0.98	-17.3	2174	4	1983	10	1806	18
n654-07b	455	225	2.62	0.55	4.790	1.2	0.2761	1.1	0.92	-23.2	2040	9	1783	10	1572	16
n654-08a	617	230	4.75	1.64	2.340	1.8	0.1540	1.2	0.71	-46.1	1803	23	1225	13	923	11
A1096, Koli	, granop	ohyre														
n655-01a	125	72	3.58	0.59	4.734	2.0	0.3051	1.6	0.81	-1.5	1841	21	1773	17	1716	25
n655-02a	1841	1307	2.03	0.18	7.310	2.3	0.3928	2.3	0.98	**	2164	9	2150	21	2136	42
n655-03a	735	395	0.99	* 0.01	7.142	1.1	0.3788	1.1	0.96	-3.7	2186	6	2129	10	2071	19
n655-03b	723	360	1.39	0.16	6.198	1.2	0.3376	1.1	0.95	-11.7	2140	7	2004	11	1875	18
n655-04a	1373	504	6.48	0.8	2.341	1.0	0.1628	0.8	0.78	-42.8	1703	12	1225	7	972	7
A1182, Koli	, gabbro															
n656-01a	364	245	2.68	0.97	6.651	1.4	0.3491	1.3	0.92	-11.3	2204	9	2066	12	1930	21
n656-01b	250	79	1.60	7.9	2.895	3.3	0.1972	1.3	0.41	-22.0	1740	54	1381	25	1160	14
n656-02a	1330	156	2.79	2.73	1.166	2.5	0.0683	0.7	0.29	-69.2	2012	41	785	14	426	3
n656-03a	638	396	1.87	0.02	7.029	2.0	0.3674	1.8	0.90	-5.3	2212	15	2115	18	2017	31
n656-03b	834	507	1.89	0.02	6.900	1.7	0.3588	1.6	0.93	-8.9	2221	11	2099	15	1977	27
n656-04a	2139	329	3.89	0.78	1.289	5.0	0.0928	4.9	0.98	-61.2	1637	20	841	29	572	27
A977, Rahas	smäki, n	netagał	obro													
n653-01a	764	438	0.68	0.14	8.118	1.4	0.4298	1.3	0.97	3.1	2190	5	2244	13	2305	26
n653-02a	429	152	0.21	0.58	4.588	1.5	0.3017	1.4	0.93	-2.9	1804	10	1747	13	1700	21
n653-03a	2737	2073	13.38	0.23	2.750	3.9	0.2729	3.8	0.99	46.5	1016	13	1342	29	1555	53
n653-04a	3038	486	1.04	0.62	1.364	1.1	0.1130	0.9	0.78	-48.2	1372	14	873	7	690	6
n653-05a	462	405	4.67	0.51	5.545	1.3	0.3406	1.2	0.89	**	1927	10	1908	11	1890	19
n653-05b	371	359	3.95	0.87	6.430	2.4	0.3899	2.3	0.97	4.5	1950	11	2036	21	2122	42
n653-06a	649	460	4.39	5.3	4.575	3.0	0.2995	1.3	0.43	**	1812	49	1745	26	1689	20
n653-07a	570	396	3.86	0.56	5.040	1.4	0.3102	1.3	0.89	-7.1	1924	12	1826	12	1742	19
n653-08a	567	670	6.63	0.12	7.422	1.2	0.3974	1.2	0.97	**	2170	5	2164	11	2157	21
n653-08b	607	510	7.21	0.51	4.299	1.5	0.2660	1.3	0.85	-19.0	1914	14	1693	12	1521	17
n653-09a	1899	2062	31.51	8.16	2.286	14.7	0.2365	3.4	0.23	**	931	268	1208	109	1368	42
n653-10a	3642	2595	8.85	4.13	3.571	6.6	0.3068	3.0	0.45	0.3	1302	111	1543	54	1725	46
A586, Ensilå	i, metag	abbro														
n657-01a	560	217	0.33	*0.01	5.060	1.4	0.3197	1.3	0.96	-2.2	1876	7	1829	12	1788	21
n657-02a	777	354	11.38	4.22	4.821	1.6	0.2599	1.1	0.69	-29.4	2158	21	1789	14	1489	15
n657-03a	382	264	2.72	0.08	6.930	1.9	0.3615	1.9	0.97	-8.0	2215	8	2103	17	1989	32
n657-04a	1381	825	2.10	0.02	6.605	1.2	0.3468	1.2	0.98	-12.7	2204	4	2060	11	1919	20
n657-05a	900	666	7.87	6.68	4.494	7.4	0.2496	0.8	0.11	-4.5	2106	123	1730	63	1437	11
n657-05b	455	252	3.12	2.48	4.755	3.1	0.2597	2.6	0.85	-26.2	2135	28	1777	26	1488	35

Table 1. Ion microprobe U-Th-Pb analytical data for zircons from 2.2 Ga mafic intrusions.

 f_{206} %: Percentage of ²⁰⁶Pb contributed by common Pb, assuming Pb isotopic composition at 1.9 Ga (Stacey and Kramers, 1975).

 $f_{_{\rm 206}}\%$: *0.01 - insignificant amount of common Pb.

Disc. %: Discordance of data (if > 2σ error of analysis). ** indicates that analysis is concordant within 2σ error.

Sample coordinates: A892: X = 7488.49, Y = 3464.70; A865: X = 7345.14, Y = 2513.90; A586: X = 7132.99, Y = 4456.79; A977: X = 7020.02, Y = 3548.98;

A1096: X = 6991.79, Y = 4498.06; A1182: X = 6992.35, Y = 4497.80.

Coordinates are presented in the Finnish kkj-coordinate system.



A865 Susivaara

Fig. 4. BSE images of zircon grains from the Susivaara sill (sample A865). All annotated ages are ²⁰⁷Pb/²⁰⁶Pb ages, which may differ from upper intercept ages depending on degree of discordance.



A865 Susivaara

Fig. 5. Spots of electron microprobe analyses (see Table 3) with red and green symbols representing "altered" and "fresh" domains, respectively.



Fig. 6. Concordia plots of U-Pb data of sample A865 from the Susivaara intrusion, Peräpohja Schist Belt. SIMS analyses shown as yellow error ellipsoids and ID-TIMS analyses by blue symbols without error indication.

5.1.3. Ensilä (A586)

Most zircon grains in sample A586 from the Ensilä metagabbro are turbid, but BSE images reveal the existence of grains that are less altered or even very clean (Fig. 7). Magmatic zoning is still present in some grains (Fig. 7C). In Fig. 7F is shown the BSE image of an interesting grain having three different types of material: a broad, light-colored, fractured rim, representing the most pristine zircon, encloses a non-fractured and obviously more altered interior, which is composed of a complicated patchwork of light-colored and dark-colored areas. The central part of this grain shows that even though a BSE image reveals two or more contrasting shades in terms of BSE intensity, none of them necessarily represents unaltered zircon.

Five conventional multi-grain U-Pb analyses indicate rather high contents of common lead for

zircon in sample A586. The isotopic data are discordant and heterogeneous providing ²⁰⁷Pb/²⁰⁶Pb ages from 1.95 to 2.12 Ga (Table 2, Fig.8). Also shown in the diagram are U-Pb zircon data reported by Hyppönen (1983) for two gabbroic samples of the same rock association, 12 and 15 km south of our site, respectively (A491 from Hietaperä and A910 from Petäjäniemi). These data are similarly heterogeneous precluding precise age calculation.

The ion microprobe was used for six measurements from five zircon crystals, but due to a high amount of common lead, one analysis (n657-05a) has been omitted from further discussion. Two analyses (n657-03a, n657-04a) from the well-preserved and relatively clear, though unzoned zircon grains yield moderately discordant data with ²⁰⁷Pb/ ²⁰⁶Pb ages of c. 2.2 Ga, whereas two analyses (n657-02a, n657-05b) from more altered zircon grains are significantly discordant (Fig. 8). A discordia forced



A586 Ensilä

Fig. 7. BSE images of zircon grains from the Ensilä gabbro (sample A586). All annotated ages are ²⁰⁷Pb/²⁰⁶Pb ages, which may differ from upper intercept ages depending on degree of discordance. Note the central part of the zircon grain in figure A, containing many thorite inclusions.

	Sample	Mineral	Sample	∍	Pb	²⁰⁶ Pb/ ²⁰⁴ Pb	²⁰⁸ Pb/ ²⁰⁶ Pb			ISOTOPIC R	ATIOS *			Rho**	APPAREN	T AGES / N	a
A586 Ensilis, Kuhmo, gabbro 3.65 4.22 155 321 0.09 0.29816 0.65 6.11988 0.3 0.83 A586B Zr 4.3/+200 3.65 4.22 155 321 0.09 0.29816 0.65 0.11988 0.3 0.83 A586B Zr 4.2.4.3 2.32 515 208 2225 0.16 0.29945 0.65 0.11988 0.3 0.83 A586D Zr 3.8.4.0/HF/ 8.05 1187 6.167 0.55 5.3055 0.65 0.13056 0.3 0.89 A587 Savilabit, Koli, gabbro 8.06 1167 6.26 425 0.54 0.33039 0.65 0.1377 0.3 0.89 A587 Savilabit, Koli, gabbro 7 1209 449 3331 0.17 0.33039 0.65 0.1377 0.3 0.89 A587 Badd 44.3 7 1209 449 33331 0.17 0.32405 0.65 0.1377 0.3 0.89 A5877 Badd 44.3		density/size fraction	weight (mg)	bpm	bpm	measured	radiogenic	²⁰⁶ Pb/ ²³⁸ U	20%	²⁰⁷ Pb/ ²³⁵ U	20%	²⁰⁷ Pb/ ²⁰⁶ Pb	20%		²⁰⁷ Pb/ ²⁰⁶ Pb	²⁰⁶ Pb/ ²³⁸ U	²⁰⁷ Pb/ ²³⁵ U
A586A Zr 44.3/+200 3.65 4.22 155 321 0.09 0.29816 0.65 5.1119 0.65 0.11988 0.3 0.83 A586B Zr 4.2.4.3 2.32 515 208 225 0.16 0.2945 0.65 5.1119 0.65 0.1295 0.33 0.83 A586D Zr 4.4.4.2 8.06 1167 6.26 4.24 0.55 5.3055 0.65 0.13056 0.3 0.89 A586E Zr 3.8.4.0 8.06 1167 6.26 4.24 0.55 5.3055 0.65 0.13056 0.3 0.89 A587C Badd 4.4.3 7 1209 449 3331 0.17 0.32948 0.65 6.1676 0.65 0.1377 0.3 0.89 A587C Badd 4.4.3 7 1209 449 3331 0.17 0.32946 0.65 0.1576 0.65 0.177 0.3 0.89 A587C	A586 Ensi	lä, Kuhmo, gabbro															
A586B $Zr 4.2.4.3$ 2.32 515 208 225 0.16 0.2945 0.65 5.1119 0.65 0.13056 0.33 0.87 A586C Zr 4.0.42 8 712 382 181.1 0.41 0.2952 0.65 5.3055 0.65 0.13056 0.3 0.83 0.87 A586C Zr 4.0.44 8.62 1167 626 425 0.53058 0.65 0.13056 0.3 0.83 0.83 A587 Badd +4.3 7 1209 449 3331 0.17 0.32405 0.65 5.1676 0.65 0.13177 0.3 0.87 A587 Badd +4.3 7 1209 449 3331 0.17 0.32405 0.65 6.1676 0.65 0.1306 0.9 0.97 0.97 A587 Badd +4.3 7 1209 449 4208 0.16 0.65 0.156 0.17 0.97 0.97 A587 Badd +4.3 61,420.0	A586A	Zr +4.3/+200	3.65	422	155	321	0.09	0.29816	0.65	4.928	0.65	0.11988	0.3	0.89	1954	1682	1807
A586C Zr 40-4.2 8 772 382 181.1 0.41 0.2952 0.65 5.3055 0.65 0.13036 0.3 0.89 A586D Zr 38.4.0 8.62 1183 611 234 0.55 0.3089 0.65 5.3746 0.65 0.13177 0.3 0.89 A586D Zr 3.8.4.0 8.06 1167 6.26 4.25 0.33948 0.65 5.3746 0.65 0.13177 0.3 0.89 A587B Badd +4.3 2 1193 431 4208 0.16 0.33357 0.65 5.59504 0.65 0.97 0.97 A587D Badd +4.3 3.2 1193 431 4208 0.16 0.33357 0.65 5.9564 0.65 0.97 0.97 A587C Badd +4.3 3.2 1299 449 3331 0.17 0.32495 0.65 0.65 0 0.15 0.97 A1096 Zr 4.3 <	A586B	Zr 4.2-4.3	2.32	515	208	225	0.16	0.2945	0.65	5.1119	0.65	0.1259	0.33	0.87	2041	1663	1838
A586D Zr 3.8-4.0 8.62 1183 611 234 0.55 0.3089 0.65 5.3746 0.65 0.11276 0.4 0.81 A586E Zr 3.8-4.0/HF/ 8.06 1167 6.26 4.25 0.30898 0.65 5.3746 0.65 0.13177 0.3 0.89 A587A Badd 44.3 7 1209 449 3331 0.17 0.32495 0.65 5.9504 0.65 0.15 0.39 A587B Badd 44.3 3.2 1193 431 4208 0.16 0.32093 0.65 5.9504 0.65 0.979 0.97 A587C Badd 44.2 3.2 1248 467 6439 0.16 0.33357 0.65 5.896 0.65 0.97 0.97 A1096 Zr 43.3 11.2 264 1259 0.74 0.29494 0.65 0.156 0.15 0.97 A10960 Zr 43.100-200 br11 120 0.333357 0.65 5.9294<	A586C	Zr 4.0-4.2	8	772	382	181.1	0.41	0.2952	0.65	5.3055	0.65	0.13036	0.3	0.89	2102	1667	1869
A586E Zr 3.84.0/HF/ 8.06 1167 626 425 0.54 0.33948 0.65 6.1676 0.65 0.13177 0.3 0.89 A587A Badd 44.3 7 1209 449 3331 0.17 0.32093 0.65 6.165 0 0.15 0.97 A587A Badd 44.3 2 1209 449 3331 0.17 0.32393 0.65 6.165 0 0.15 0.97 A587C Badd 44.3 2 1209 449 3331 0.16 0.33357 0.65 6.165 0 0.15 0.97 A587C Badd 44.2 3 12248 467 6439 0.16 0.33357 0.65 6.1458 0.65 0.15 0.97 A1096A Zr 42.43 11.12 261 125 0.33356 0.65 0.156 0.1269 0.125 0.95 A1096A Zr 42.43 0.163 <	A586D	Zr 3.8-4.0	8.62	1183	611	234	0.55	0.30089	0.65	5.3746	0.65	0.1296	0.4	0.81	2092	1695	1880
A587 Savilahti, Koli, gabbro 7 1209 449 3331 0.17 0.32405 0.65 5.9504 0.65 0 0.15 0.97 A587A Badd +4.3 2 1193 431 4208 0.16 0.32305 0.65 5.9504 0.65 0 0.15 0.97 A587C Badd +4.3 2 1193 431 4208 0.16 0.33357 0.65 5.896 0 0.15 0.97 0.97 0.97 0.97 0.97 0.915 0.97 0.915 0.97 0.97 0.95 5.896 0 0.15 0.97 <	A586E	Zr 3.8-4.0/HF/	8.06	1167	626	425	0.54	0.33948	0.65	6,1676	0.65	0.13177	0.3	0.89	2121	1884	1999
A587A Badd +4.3 7 1209 449 3331 0.17 0.32405 0.65 5.9504 0.65 0 0.15 0.97 A587B Badd +4.4.3 2 1193 431 4208 0.16 0.32357 0.65 5.8966 0.65 0 0.15 0.97 A587B Badd +4.4.3 3.2 1193 457 6439 0.16 0.33357 0.65 5.8966 0.65 0 0.15 0.97 A1096 Kamisniemi, Koli, grauophyre 10.4 165 80 1259 0.74 0.29494 0.65 4.8233 0.65 0.11861 0.2 0.95 A1096 Zr 4.4.3 11.1.2 261 125 0.058 0.30458 0.65 4.8233 0.65 0.11861 0.2 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95 0.95	A587 Savi	lahti, Koli, gabbro															
A587B Badd 44.3 2 1193 431 4208 0.16 0.32093 0.65 5.896 0.65 0 0.15 0.97 A587C Badd 44.2/+200 abr 3.2 1248 467 6439 0.16 0.33357 0.65 6.1458 0.65 0 0.15 0.97 A1096 Zr 4.3 10.4 165 80 1259 0.74 0.29494 0.65 4.8233 0.65 0.11861 0.2 0.95 A1096K Zr 4.3.3 11.2 261 125 0.074 0.29494 0.65 4.8233 0.65 0.11861 0.2 0.95 A1096K Zr 4.3.710-200 6.1 190 93 987 0.68 0.30395 1.2 0.05 0.1269 0.2 0.95 A1096D Zr 4.3.710-200 6.1 190 93 987 0.68 0.30458 0.65 0.12699 0.2 0.95 A1082D Zr 4.2.4.2100 10.9 695	A587A	Badd +4.3	~	1209	449	3331	0.17	0.32405	0.65	5.9504	0.65	0	0.15	0.97	2140	1809	1968
A587C Badd $44.2/t$ 200 abr 3.2 1248 467 6439 0.16 0.33357 0.65 6.1458 0.65 0 0.15 0.97 A1096 Kaunisniemi, Koli, granophyre 10.4 165 80 1259 0.74 0.29494 0.65 6.1458 0.65 0 0.15 0.97 A1096 Kaunisniemi, Koli, granophyre 11.2 261 125 1030 0.64 0.30396 1.2 6.13 0.12 0.2 0.99 A1096 M 2 Zr 43.3 10.4 165 80 1259 0.74 0.29494 0.65 0.12069 0.2 0.995 A1096 M 2r $4.32/t-100$ 10.9 695 303 892 0.58 0.30458 1.25 0.12069 0.2 0.95 0.12066 0.2 0.95 0.12066 0.2 0.95 0.12066 0.2 0.95 0.12069 0.2 0.95 0.12069 0.2 0.95 0.1259 0.95 0.12066 0.2 0.95 0.125	A587B	Badd +4.3	2	1193	431	4208	0.16	0.32093	0.65	5.896	0.65	0	0.15	0.97	2141	1794	1960
A1096 X + 4.3 0.65 80 125 0.74 0.29494 0.65 4.8233 0.65 0.11861 0.2 0.95 A1096A Zr +4.3 10.4 10.4 125 0.05 0.054 0.11861 0.2 0.95 A1096A Zr +4.3 11.2 11.2 125 1030 0.64 0.303958 1.2 0.12 0.2 0.95 A1096D Zr +4.3/100-200 abrbh 11.2 190 93 987 0.68 0.303958 0.65 0.12069 0.2 0.95 A1096D Zr 3.4.4.2/100 10.9 695 303 892 0.58 0.28627 0.65 4.7388 0.65 0.12069 0.2 0.95 A1182A Zr 44.2/100 6.9 337 102 308 0.86 0.15999 0.65 0.12396 0.2 0.95 A1182D Zr 44.0/200 6.9 1067 316 277 0.65 0.127407 0.2 0.95	A587C	Badd +4.2/+200 abr	3.2	1248	467	6439	0.16	0.33357	0.65	6.1458	0.65	0	0.15	0.97	2146	1855	1996
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	A1096 Ka	unisniemi, Koli, granophy	re														
A1096B Zr $4.2-4.3$ 11.2 261 125 1030 0.64 0.30396 1.2 5.029 1.2 0.12 0.2 0.99 A1096C Zr $4.3/100-200$ abr2h 6.1 190 93 987 0.68 0.30458 0.65 5.0682 0.65 0.12069 0.2 0.95 A1096D Zr $3.84.2t+100$ 10.9 695 303 892 0.58 0.30458 0.65 0.12069 0.2 0.95 A1182 Kaunishti, Koli, gabbro 10.9 695 303 892 0.58 0.28627 0.65 4.7338 0.65 0.12069 0.2 0.95 A1182A Zr $44.24.2t+200$ 6.3 632 161 275 0.13873 0.65 0.12396 0.2 0.95 A1182D Zr 44.0200 6.3 632 161 275 0.75 0.17373 0.65 0.12407 0.2 <	A1096A	Zr +4.3	10.4	165	80	1259	0.74	0.29494	0.65	4.8233	0.65	0.11861	0.2	0.95	1935	1666	1788
A1096C Zr +4.3/100-200 abr2h 6.1 190 93 987 0.68 0.30458 0.65 5.0682 0.65 0.12069 0.2 0.95 A1096D Zr 38.4.2/+100 10.9 695 303 892 0.58 0.30458 0.65 5.0682 0.65 0.12069 0.2 0.95 A1096D Zr 38.4.2/+100 10.9 695 303 892 0.58 0.28627 0.65 4.7388 0.65 0.12006 0.2 0.95 A1182A Zr 44.2/+200 6.3 633 102 308 0.84 0.15999 0.65 2.7344 0.65 0.12396 0.2 0.95 A1182D Zr 4.4.4.200 6.3 161 275 0.75 0.13873 0.65 2.3744 0.65 0.12407 0.2 0.95 A1182D Zr 4.4.4.200 6.3 161 275 0.755 0.17573 0.65 0.12407 0.2 0.95 A1182C Zr 4.4.4.200 6.	A1096B	Zr 4.2-4.3	11.2	261	125	1030	0.64	0.30396	1.2	5.029	1.2	0.12	0.2	0.99	1956	1710	1824
A1096D Zr 3.84-2/+100 10.9 695 303 892 0.58 0.28627 0.65 4.7388 0.65 0.12006 0.2 0.95 A1182 Zr 44.2/+200 5.5 337 102 308 0.84 0.15999 0.65 2.7344 0.65 0.12396 0.2 0.95 A1182B Zr 44.2/+200 6.3 6.3 161 275 0.75 0.13873 0.65 2.7344 0.65 0.12396 0.2 0.95 A1182D Zr 4.4.4.2/+200 6.3 161 275 0.75 0.13873 0.65 2.3731 0.65 0.12407 0.2 0.95 A1182D Zr 3.4.4.4.200 6.3 167 315 4.83 0.65 3.1122 0.65 0.12407 0.2 0.95	A1096C	Zr +4.3/100-200 abr2h	6.1	190	93	987	0.68	0.30458	0.65	5.0682	0.65	0.12069	0.2	0.95	1966	1713	1830
A1182 Zr $4.2/2$ 337 102 308 0.84 0.15999 0.65 2.7344 0.65 0.12396 0.2 0.95 A1182A Zr $44.2/1+200$ 6.3 632 161 275 0.75 0.13873 0.65 2.7344 0.65 0.12396 0.2 0.95 A1182D Zr $44.4.2/1+200$ 6.3 632 161 275 0.17573 0.65 2.3734 0.65 0.12407 0.2 0.95 A1182C Zr $34.0+200$ 6.3 161 275 0.17753 0.65 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95 0.12715 0.2 0.95	A1096D	Zr 3.8-4.2/+100	10.9	695	303	892	0.58	0.28627	0.65	4.7388	0.65	0.12006	0.2	0.95	1957	1622	1774
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	A1182 Ka	unislahti, Koli, gabbro															
A1182B Zr 4.0-4.2/+200 6.3 6.32 161 2.75 0.75 0.13873 0.65 2.3731 0.65 0.12407 0.2 0.95 A1182C Zr 3.84.0/+200 6.9 1067 315 483 0.66 0.17753 0.65 3.1122 0.65 0.12715 0.2 0.95	A1182A	Zr +4.2/+200	5.5	337	102	308	0.84	0.15999	0.65	2.7344	0.65	0.12396	0.2	0.95	2014	956	1337
A1182C Zr 3.84.0(+200 6.9 1067 315 483 0.66 0.17753 0.65 3.1122 0.65 0.12715 0.2 0.95	A1182B	Zr 4.0-4.2/+200	6.3	632	161	275	0.75	0.13873	0.65	2.3731	0.65	0.12407	0.2	0.95	2015	837	1234
	A1182C	Zr 3.8-4.0/+200	6.9	1067	315	483	0.66	0.17753	0.65	3.1122	0.65	0.12715	0.2	0.95	2059	1053	1435

Table 2. ID-TIMS U-Pb data on zircon and baddeleyite from Kuhmo and Koli.

 \propto Kramers 19/5). and age related common lead (Macey (gu C.0 d'l) Mand Isotopic ratios corrected for fractionation, Error correlation between Pb/U errors

^{**}) Error correlation between Pb/1 Zr = zircon, Badd = baddelevite

= 7132.99, Y = 4456.79; A587: X = 7002.65, Y = 4482.49; A1096; X = 6991.79, Y = 4498.06; A1182: 6992.35, Y = 4497.80. Coordinates are presented in the Finnish kkj-coordinate system. Sample coordinates: A586: X

through these four analyses gives an upper intercept age of 2232 ± 19 Ma. Analysis n657-01a is clearly distinct from the other and provides a nearly concordant ²⁰⁷Pb/²⁰⁶Pb age of c. 1.9 Ga. This analysis also has a lower Th/U ratio (0.3) compared to other data (Table 1, Fig. 8). The grain in question is relatively transparent and, surprisingly, possesses a weak zoning pattern. One possibility is that the grain is an outsider introduced by contamination during sample preparation. On the other hand, it represents the heaviest zircon fraction in the sample $(d > 4.2 \text{ g/cm}^3)$, and the same features are obvious from the conventional data, i.e. analysis A586A from the heaviest zircon fraction has the lowest ²⁰⁷Pb/²⁰⁶Pb age and Th/U ratio (as deduced from the ²⁰⁸Pb/²⁰⁶Pb data in Table 2). Note that dark, low-Z areas within the zircon crystals were not analyzed by SIMS from sample A586 as all the analyzed spots represent relati-

In summary, most of the multi-grain zircon fractions analyzed by the conventional method plot within the area delineated by the spot analyses (Fig. 8). The new data confirm that the Ensilä intrusion belongs to the c. 2.2

vely high-intensity BSE areas (see Fig. 7).

Fig. 8. Concordia plots of U-Pb data of three samples (A491, A586, A910) from Paleoproterozoic mafic intrusive rocks from the Archean Kuhmo Greenstone Belt. SIMS analyses shown as yellow error ellipsoids and ID-TIMS analyses by blue and green circles without error indication.



Ga age group and suggest that the intrusive body has undergone a multi-stage evolution including significant resetting of the zircon U-Pb system at c. 1.8–1.9 Ga. This means that the surrounding Archean greenstone belt has also experienced the same tectono-metamorphic phenomena in the Paleoproterozoic time.

5.1.4. Rahasmäki (A977)

Zircons in the Rahasmäki intrusion are commonly very turbid but, as revealed by BSE images, some of the grains still retain well-preserved zones (Fig. 9). An interesting feature of the conventional data from Rahasmäki, as published by Paavola (1984), is that the heavier zircon grains (density 4.0–4.2 g/cm³) have higher U contents and are more discordant than the lighter ones (3.8–4.0 g/cm³). Normally the situation is reversed (see Table 2).

Twelve U-Pb analyses on ten grains were obtained by ion microprobe. Three analyses with high amounts of common lead (206Pb/204Pb <400) and consequently large errors have been omitted from the concordia diagram and further discussion. The data as a whole are very scattered (Table 1, Fig. 10). Four analyses plot close to the chord defined by the conventional data of Paavola (1984), but two analyses from the most pristine zircon domains are nearly concordant at c. 2.2 Ga (n653-01a, n653-08a). This difference is clearly seen between the two spots in grain n653-08, which exemplifies the complicated alteration pattern seen in BSE images of many crystals (Fig. 9E). One analysis (n653-02a) is slightly on the "younger side" of the chord, but analysis n653-03a plots above the concordia with a ²⁰⁷Pb/ ²⁰⁶Pb age of c. 1.0 Ga. The analysis indicates a very high Th content (c. 4 %), and it is expected that the analytical routine for normal zircon may not be



A977 Rahasmäki

Fig. 9. BSE images of zircon grains from the Rahasmäki sill (sample A977). All annotated ages are ²⁰⁷Pb/²⁰⁶Pb ages, which may differ from upper intercept ages depending on degree of discordance. Note the large, white baddelyite inclusion in the smaller zircon grain in figure A. The largest inclusion in the zircon grain of figure B is thorite.



Fig. 10. Concordia plot of U-Pb data of sample A977 from the Rahasmäki sill, Tahkomäki-Kinahmi schist belt. SIMS analyses shown as yellow error ellipsoids and ID-TIMS analyses by blue circles (without error indication).

valid for such a composition. The white specks in the photomicrograph of grain n653-03a (Fig. 9B) are thorite grains which further highlights the complicated isotopic history of the zircon grain.

In summary, the SIMS results from the Rahasmäki sample suggest magmatic crystallization at c. 2.2 Ga, i.e. c. 200 Ma earlier than indicated by the previous ID-TIMS data (1967 \pm 24 Ma), and a major lead loss from zircon at c. 1.8–1.9 Ga.

5.1.5. Koli (A1096, A1182, A587)

Zircon in all samples is turbid. Especially in the granophyre A1096 it occurs as large, broken, very turbid grains. As shown by the BSE images of Fig. 11, zircon grains from sample A1096 are, together with the A865 zircons from Susivaara, the most altered with a particularly spongy internal texture. Zircon crystals from the gabbro sample A1182 are notably variable in terms of preservation. Some grains are almost devoid of signs of alteration (Fig. 12C), some are traversed by a distinctive alteration band (Fig. 12F), and there also exist grains that are heterogeneously but pervasively altered (Fig. 12B, D). In Fig. 12E, the primary magmatic zoning is still recognizable in a remnant of pristine zircon close to the edge of the grain. The rest of the grain is composed of an irregular network of light-colored and grey secondary zircon enclosing a great number of silicate inclusions.

Four conventional U-Pb zircon analyses from sample A1096 are discordant and plot close to each other yielding ²⁰⁷Pb/²⁰⁶Pb ages of c. 1.96 Ga (Table 2, Fig. 13A). Air abrasion has no effect on the concordance of analysis, which is understandable after inspection of the BSE images (Fig. 11). Three ID-TIMS analyses of baddeleyite from sample A587 are discordant with ²⁰⁷Pb/²⁰⁶Pb ages of c. 2.14 Ga (Fig. 13B). A linear regression through these data gives concordia intercepts at 2170 ± 20 Ma and 253 ± 150 (MSWD = 1.3, n = 3). As shown in Fig. 13B, the three conventional U-Pb analyses made on zircon from the gabbro sample A1182 are extremely discordant and plot close to the chord defined by the baddeleyite fractions.

Eleven spot analyses on zircon were performed by SIMS from samples A1096 and A1182. Two of the analyses (n656-01b and n656-02b) have large proportions of common lead and thus large errors. The data are heterogeneous and show a very large



A1096 Kaunisniemi (Koli)

Fig. 11. BSE images of zircon grains from granophyre of the Koli sill (sample A1096 from Kaunisniemi). All annotated ages are ²⁰⁷Pb/²⁰⁶Pb ages, which may differ from upper intercept ages depending on degree of discordance.

A1182 Kaunislahti (Koli)



Fig. 12. BSE images of zircon grains from metagabbro of the Koli sill (sample A1182). All annotated ages are ²⁰⁷Pb/²⁰⁶Pb ages, which may differ from upper intercept ages depending on degree of discordance.



Fig. 13. Concordia plot of U-Pb data from the Koli sill. SIMS analyses shown as yellow error ellipsoids and ID-TIMS analyses by blue circles (zircon) and crosses (baddeleyite) without error indication. A. Granophyre A1096 from Kaunisniemi. B. Gabbro A1182 from Kaunislahti and gabbro A587 from Savilahti.

variation in the discordance and U content. From the BSE images it is clear that the least discordant analyses (n655-02a, n655-03a, n656-03a and n656-03b; Figs. 11, 12) with ²⁰⁷Pb/²⁰⁶Pb ages approaching 2.2 Ga represent most pristine zircon domains. Regression of four analyses from the granophyre A1096 define intercepts at 2199 \pm 45 and 552 \pm 97 Ma (MSWD = 3.3, Fig. 13A). One SIMS analysis (n655-01a) as well as the conventional data plot distinctly on the "younger" side of the chord, suggesting significant influence of the 1.8–1.9 Ga thermal fluids in zircon. In fact, analysis n655-01a suggests complete resetting of the U-Pb system and/or formation of new zircon at c. 1.9 Ga. This grain also has a much lower U content than the other grains, but surprisingly, in the BSE image (see Fig. 11A), is not very distinct from the other altered zircon crystals from the same sample.

The least discordant data from sample A1182 provide an average 207 Pb/ 206 Pb age of 2211 ± 12 Ma, and the two very discordant points from strongly altered domains together with conventional data manifest near-complete and relatively recent loss of radiogenic lead (Fig. 13B).

The most concordant zircon analyses have a U content of less than 1000 ppm while discordant zircons often display a U content of higher than 1000 ppm (Table 1). However, the relationship between the U content and degree of discordance is not simple as the above generalization does not always hold.

5.2. Electron microprobe analyses of altered zircons

Our electron microprobe data include 33 analyses and are restricted to a single zircon grain from sample A865 representing a gabbro from the Susivaara sill. As shown in Fig. 5, this grain exhibits a typical alteration pattern of the GWA zircons with two distinctly different shades in the BSE image, i.e. darker, patchy zones representing more altered parts and lighter, cleaner zones representing less damaged parts. In the following we refer to these two areas as altered and fresh domains, respectively. The two SIMS analyses, n654-02a and 0654-02b, were made on the same grain and resulted in clearly different ²⁰⁷Pb/²⁰⁶Pb ages of 2192 ± 3 and 1655 ± 21 Ma, respectively (Table 1). In fact, these SIMS spots represent the two extremes in the degree of discordance of the analyses from sample A865 (Fig. 6). As indicated in Fig. 5, we analyzed two profiles by electron microprobe, one crossing a fresh zone and the other an altered zone. These were supplemented by five analyses on the SIMS crater n654-02a and six scattered points elsewhere in the grain. Representative analyses are shown in Table 3. In spite of the limited number of analyses, we believe that the data obtained from this single grain are representative of the main chemical characteristics of altered zircon grains in sample A865 or in the GWA rocks in general.

0.20 3.66 0.19 28.58 0.63 0.25 Ο 23 01.38 0.10 2.32 29.04 0.0457.37 1.35 0.21 0.29 0.03 0.64U 22 0.05 .46 0.13 0.28 0.03 0.27 1.21 0.23 υ 21 0.65 0.25 0.02 2.25 0.07 0.21 υ 20 7.60 1.39 0.03 0.18 0.36 0.03 0.83 0.07 2.35 C 19 4.98 6.04 1.43 0.00 0.041.53 0.15 0.27 18 υ 0.692.58 7.42 0.00 0.03 1.37 0.31 υ 17 6.59 1.59 0.02 0.440.27 0.473.01 0.1701.93 υ 16 0.16 0.13 0.39 0.53 0.25 3.41 9.97 υ 15 $0.42 \\ 0.17$ 2.45 1.540.05 0.31υ 14 0.1056.75 1.65 0.440.39 0.21 2.44 0.13 0.08 70.00 13 Ο 1.52 0.22 1.29 0.00 0.06 0.00 0.30 .61 12 В 0.06 0.10 51.39 0.00 0.02 0.20 В 11 51.93 1.480.20 1.13 0.00 0.07 0.00 0.22 В 0 0.15 51.68 0.00 0.140.00 0.35 6.07 1.01 В 6 0.18 0.09 1.63 0.00 0.00 0.32 52.07 1.03 В ø 31.40 0.06 0.08 52.46 1.640.16 0.00 96.88 0.96 0.01 0.11 В 52.86 0.09 0.00 0.13 0.96 0.00 0.37 7.45 .51 В 9 52.85 0.08 0.00 0.16 7.41 1.640.12 0.80 0.00 В Ś 0.160.89 0.05 0.00 0.31 0.01 4 R 0.330.05 53.63 l.48 0.16 0.00 0.07 0.00 0.91 V $\hat{\mathbf{c}}$ 53.94 0.13 0.00 0.06 0.28 0.01 1.61 0.90 V 2 54.00 1.58 0.10 0.89 0.240.00 0.06 0.00 V Location otal

5 for the location of the profiles and SIMS crater at spot n654-02)

Table 3. Electron microprobe analyses of zircon, grain n654-02, sample A865 (see Fig.

Location: A: SIMS crater; B = profile 1; C = profile 2.

Figure 14 displays the analytical data divided into two groups that represent the "fresh" and "altered" domains. One of the most striking features of the altered domains is their elevated CaO content, which can be used as a simple criterion for distinction between relatively fresh and altered parts of zircon (cf. Geisler & Schleicher, 2000). In the former domains, the CaO content is always very low (≤ 0.02 wt.%), approaching or being below the detection limit of the electron microprobe, whereas in the altered domains, CaO is commonly higher than 1.0 wt.% and reaches a maximum value of 3.6 wt.% (Fig. 14B).

A characteristic feature of the altered domains is the low analytical total varying from 90.0 to 93.5 wt.%. It is likely that the low totals are mostly accounted for by the presence of a water species in the damaged parts of the zircons (cf. Geisler et al., 2003a). Because the Al_2O_3 content was not analyzed by microprobe, we cannot know the exact amount of H_2O in zircons. However, later SEM-EDS analyses of altered zircons in other samples indicate approximate Al_2O_3 contents in the range of 1-2.5 wt.%, which suggest that the water concentrations in altered domains can reach many percentages. The SiO₂ and ZrO₂ contents are lower in the altered domains than in the fresh domains (Fig. 14A) and can be explained by a high degree of hydration. The atomic Si/(Zr+Hf) ratio in both zones overlaps and approaches stoichiometric 1.0, though on average it is slightly higher in the altered domains (0.99–1.07) compared with the fresh domains (0.95–1.03).

The FeO content in fresh domains is commonly less than 0.15 wt.% while in altered domains it is higher and, excluding two analyses (ca. 6 wt.%), falls in the range of 0.15–1.0 wt.% (Fig. 14C). The MnO contents are negligible in fresh zones and also low (<0.1 wt.%) in some analyses from altered zones but rises to values between 0.25 and 0.53 wt.% in some other parts of the altered zones (Fig. 14D). The Y_2O_3 content of the altered domains is commonly less than that in the fresh domains with the averages being 0.37 wt.% and 0.96 wt.%, respecti-



Fig. 14. Electron microprobe data on a zircon grain from the Susivaara sample A865 (for location of data points, see Fig. 5).

centrations of HfO₂ (1.30–1.65 wt.%), TiO₂ (0.03– 0.09 wt.%), and PbO (0.08–0.46 wt.%). The UO₂ content (0.00–0.35 wt.%) is often lower in altered domains but in some spots it is higher than in fresh domains.

SEM-EDS analysis was used to identify the mineralogy of the distinct, bright specks, from a few to 20 mm in size, which commonly occur within altered domains (Figs. 4, 5, 7, 9, 11, 12). This revealed mainly Th and Si (+ U, Zr, Ca) indicating that these mineral grains are composed of thorite (ThSiO₄). A few of the high-BSE intensity crystals turned out to be lead selenide (clausthalite). Baddeleyite was also found (Fig. 9A). A detailed study on the numerous silicate inclusions in zircon grains has not yet been carried out, but the currently available data show the presence of Fe-rich chlorite and sphene.

In summary, based on our limited electron microprobe data, the degree of alteration in zircons seems to correlate positively with measured CaO and inferred H₂O contents and less systematically with concentrations of FeO and MnO. In addition, altered zircon domains often contain tiny thorite grains.

5.3. Sm-Nd results 5.3.1. Intrusions from Central Lapland

The three samples from the Haaskalehto intrusion that were used for our Sm-Nd study are relatively well-preserved gabbros. Some of the Sm-Nd isotopic analyses of Table 4 showing relatively large errors were made in the 1980s using an old technique and a non-commercial mass spectrometer (Huhma, 1986). Ten analyses including two whole rocks (duplicated analyses on both) and clinopyroxene, amphibole and plagioclase separates define an isochron which gives an age of 2119 ± 40 Ma with an initial ε_{Nd} value of +0.8. The relatively large MSWD value of 2.5 suggests some scatter in excess of analytical error. If the two plagioclase analyses are excluded, MSWD decreases to 0.9 and the age from the slope, as defined largely by the pyroxene analyses of sample 19-HSP-78, is 2187 ± 44 Ma $(\varepsilon_{Nd} = +0.7, \text{ Fig. 15})$. This age is more consistent with the U-Pb zircon age of 2220 ± 11 Ma (Tyrväinen, 1983).

Fig. 15. Sm-Nd isotope data for whole-rock samples (gabbro) and mineral separates from the Haaskalehto intrusion. Analytical results on plagioclase are excluded in regression.



Table 4. Sm-No	l isotopic data	on 2.22 Ga	mafic intrusions	(ϵ_{Nd})	calculated	at 2200	Ma).
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Sample	Wr/ mineral	Rock type	Coord X	inates Y	Sm (ppm)	Nd (ppm)	¹⁴⁷ Sm/ ¹⁴⁴ Nd	¹⁴³ Nd/ ¹⁴⁴ Nd	$2 \sigma_{_{\rm m}}$	$\boldsymbol{\epsilon}_{_{Nd}}$
Haadralahta intr	usion Control	Lonland		_	(FF)	·rr/				
10 1 HSD 78	W/r	Cabbro	7/87 53	3466 63	2 70	0.71	0.1678	0.512240	3/1	0.4
19.1-FISF-/0	WI W/# #2	Cabbro	7407.33	2466.62	2.70	9./1	0.16/8	0.512240	- 54 - 11	0.4
19.1-1131-/0 10 LICD 70	WI #2	Cabbro	7407.33	2466.62	2.05	7.00	0.1062	0.512207	20	0.0
19-ПЗР-/8	Cpx #2	Gabbro	/48/.33	3400.03	2.01	/.00	0.2255	0.51508/	20	0.5
19-HSP-/8	Cpx #3	Gabbro	/48/.55	3466.63	2.65	6./0	0.23/6	0.51526/	10	0.5
24-HSP-/8	Wr	Gabbro	/48/.89	3465.80	2.50	7.94	0.1946	0.512646	20	0./
24-HSP-/8	Wr	Gabbro	/48/.89	3465.80	2.51	/./)	0.1955	0.512660	10	0./
24-HSP-/8	Ampn+Cpx	Gabbro	/48/.89	3465.80	4.8/	14.95	0.1968	0.512682	24	0.8
24.1-HSP-/8	Plag	Gabbro	/48/.89	3465.80	0.58	2.48	0.0999	0.5115/9	60	5.0
A1408	Plag	Gabbro	/488.00	3465.00	0.54	2.89	0.1124	0.51150/	10	1.9
A1408	Ampn	Gabbro	/488.00	3465.00	4.92	14.51	0.2049	0.512//8	10	0.5
Silmäsvaara intru	sion, Central I	Lapland								
A1430	Wr	Ol-pyroxenite	7499.00	3440.30	1.88	5.65	0.2007	0.512698	10	0.0
A1430	Cpx ¹⁾	Ol-pyroxenite	7499.00	3440.30	2.01	5.93	0.2049	0.512751	10	-0.2
A1430	Plag ²⁾	Ol-pyroxenite	7499.00	3440.30	0.10	0.72	0.0821*	0.510962	41	-0.1
A1430	Plag #2	Ol-pyroxenite	7499.00	3440.30	0.08	0.62	0.0814*	0.510988	21	0.7
Ahvenvaara intru	sion, Central I	Lapland								
A1431	Wr	Ol-pyroxenite	7429.99	3508.70	1.68	4.76	0.2130	0.512918	12	0.7
A1431	Cpx	Ol-pyroxenite	7429.99	3508.70	1.94	5.36	0.2192	0.513005	10	0.7
A1431	Plag	Ol-pyroxenite	7429.99	3508.70	0.09	0.63	0.0867*	0.511035	25	0.0
A1431	Plag #2	Ol-pyroxenite	7429.99	3508.70	0.08	0.62	0.0827*	0.511014	10	0.8
Koli sill, Norther	n Karelia		,							
A1220	Wr	Pyroxenite	6991.97	4498.35	2.92	10.35	0.1706	0.512326	15	1.3
A1220	Wr #2	Pyroxenite	6991.97	4498.35	2.98	10.55	0.1709	0.512315	10	1.0
A1220	Cpx	Pyroxenite	6991.97	4498.35	2.28	6.51	0.2121	0.512885	32	0.4
A1220	Plag	Pyroxenite	6991.97	4498.35	0.40	2.88	0.0833	0.511224	35	4.7
A1221	Wr	Pyroxenite	6991.96	4498.41	2.67	8.63	0.1869	0.512607	46	2.1
A1221	Wr #2	Pyroxenite	6991.96	4498.41	2.87	9.32	0.1858	0.512548	20	1.3
A1221	Cpx	Pyroxenite	6991.96	4498.41	2.02	5.33	0.2288	0.513134	11	0.5
A1221	Cpx #2	Pyroxenite	6991.96	4498.41	2.00	5.21	0.2323	0.513205	37	0.8
A1221	Plag	Pyroxenite	6991.96	4498.41	1.28	8.69	0.0888	0.511334	24	5.3
48-JIV-85	Wr	Gabbro	6991.81	4498.09	4.97	20.79	0.1447	0.511940	10	1.2
A1096a	Wr	Granophyre	6991.79	4498.06	7.59	36.10	0.1271	0.511663	10	0.8
A1182	Wr	Gabbro	6992.35	4497.80	5.02	19.18	0.1582	0.512122	10	0.9
Arola dike, Kuhn	10 Greenstone	Belt								
TTT-170-96	Wr	Wehrlite	7149.00	4454.00	2.42	8.69	0.1681	0.512271	10	0.9
TTT-170-96	Cpx ³⁾	Wehrlite	7149.00	4454.00	1.91	5.74	0.2016	0.512742	10	0.6
TTT-170-96	$C_{px} # 2^{4)}$	Wehrlite	7149.00	4454.00	2.27	7.13	0.1922	0.512599	10	0.5
Ensilä dikes Kuh	mo Greenston	e Belt	, 1 1,100	1191100	2127	7.120	011/22	0.912999	10	0.5
TTT 82 14 96	W/r	Wahrlita	7132.10	4456 10	1.0	6.61	0.1730	0.512354	10	0.9
TTT 155 06	WI W/r	Wahalita	7132.10	44 30.10	2.22	7.02	0.1739	0.512354	10	0.9
TTT 192 4 06	WI W/n	Dumouronito	7130.70	44554.20	2.23	12 66	0.1/02	0.512129	10	0.7
TTT 01 1A 06	WI W/n	Dimovonito	7131./0	44)4.20	2.49	7 27	0.197	0.512156	10	1.)
TTT 122 2 06	WI W/n	Dimovonito	7132.10	44 30.00	4.70	20.52	0.1920	0.511000	10	-0.5
TTT 26 06	WI W/r	Cabbro	7131.90	4455.10	4./9	20.55	0.1411	0.512211	10	1.0
1 1 1-30-90 TTT 20 00	WI W/-	Gabbro	7122.40	4455.10	5.5 6.17	11.02	0.1/19	0.512511	10	0.6
1 1 1-89-90 TTT 146 14 06	WI W/-	Gabbro	/152.30	4436.20	4.1/	10.21	0.1557	0.512062	10	0.4
1 1 1-140.1A-90	wr	Gabbro	/151./0	4455.80	4.4)	18.38	0.1462	0.311962	10	1.2
Runkausvaara, Pe	eräpohja Schist	Belt (taken from	Huhma et al	., 1990)						
HH-19/82	Wr	Gabbro	7329.80	2544.08	3.90	15.90	0.1505	0.511991	35	0.4
KF597/78	Wr	Gabbro	7329.02	2563.48	4.80	20.50	0.1408	0.511854	46	0.4
KF742/78	Wr	Wehrlite	7329.02	2563.48	2.30	9.06	0.1538	0.512039	40	0.4
KF753/78	Wr	Gabbro	7329.20	2563.30	7.70	36.20	0.1287	0.511629	50	-0.5
A475	Sphene	Gabbro	7334.15	2544.09	52.10	174.40	0.1807	0.512394	50	-0.3
A475	Plag	Gabbro	7334.15	2544.09	1.42	5.74	0.1498	0.511833	45	-2.5

Appreviations: Wr = whole rock, Plag = plagioclase, Cpx = clinopyroxene, Amph = amphibole, Ol = olivine.

Notes on mineral separates: 1) density 3.3-3.4; 2) density 3.63-2.76; 3) magnetic fraction, density 3.25-3.33; 4) not handpicked, abrasion 30 min. The name of the rock type corresponds to the original, unmetamorphosed rock.

Error in ¹⁴⁷Sm/¹⁴⁴Nd is 0.4%, except when marked by *, it is 2%.

 143 Nd/ 144 Nd radio is normalized to 146 Nd/ 144 Nd = 0.7219, error is 2 standard error of the mean in the last sign. digits.

Error in $\boldsymbol{e}_{_{Nd}}$ is normally ±0.4 units.

Coordinates are presented in the Finnish kkj-coordinate system.

Microscopic observations reveal that plagioclase is turbid and yellowish and contains impurities, which were not possible to be removed by handpicking. It is conceivable that metamorphic effects, which are common in Lapland, have also slightly influenced the Sm-Nd system in plagioclase in spite of the relatively well-preserved nature of the rocks at the sampling site.

Mineral separation using well-preserved olivine pyroxenites from the Silmäsvaara and Ahvenvaara intrusions yielded clean fractions of pyroxene and plagioclase. Due to low concentrations of Sm and Nd, the analytical error for plagioclase is relatively large (Table 4). Four analyses from the Silmäsvaara sample A1430 give an age of 2185 ± 35 Ma (ε_{NM} = 0.0, MSWD = 1.07), and the result from Ahvenvaara (A1431) is 2231 ± 27 Ma (ε_{Nd} = +0.8, MSWD = 1.5, Fig. 16). These ages are largely based on plagioclase, since the Sm/Nd ratio in the whole-rock samples is close to that of pyroxene. Compared to Haaskalehto discussed above, plagioclase in the Silmäsvaara and Ahvenvaara samples is very clear and has likely remained closed since its magmatic crystallization. The age of c. 2.2 Ga is compatible with the U-Pb zircon age of 2222 ± 6 Ma from the Harjunoja gabbro (Räsänen & Huhma, 2001), which according to the aeromagnetic map, probably belonged originally to the same intrusive body as the Ahvenvaara intrusion.

5.3.2. Dikes in the Kuhmo Greenstone Belt

A few Sm-Nd analyses have been made on mafic dikes occurring in the Ensilä and Arola areas in the Archean Kuhmo Greenstone Belt. The samples are classified into three groups, wehrlites, metapyroxenites and metagabbros, which all contain largely a metamorphic mineral paragenesis. The results of the Sm-Nd isotope analyses are shown in Table 4. The data are scattered probably due to metamorphic effects (Fig. 17). An attempt was made to use pyroxene to date a wehrlitic sample (TTT-170-96) from Arola, but no pure fraction was possible to obtain due to alteration of pyroxene. Nevertheless, two analyses were made on altered pyroxene concentrates, which together with whole-rock data yield a date of 2120 ± 86 Ma. The average ε_{NA} (2220 Ma) value of +0.7 for the whole data set is similar to that of other GWA intrusions (see below).



Fig. 16. Sm-Nd isotope data for the Silmäsvaara (A1430) and Ahvenvaara (A1431) intrusions. Analytical results on olivine pyroxenites and clinopyroxene and plagioclase separates are regressed separately for both intrusions.



Fig. 17. Sm-Nd isotope data for Paleoproterozoic mafic intrusive rocks from the Archean Kuhmo Greenstone Belt. Samples include two impure clinopyroxene separates and one wehrlitic whole-rock sample from Arola and ultramafic and mafic whole-rock samples from Ensilä. Two regression lines are shown, one based on the Arola analyses and the other on the whole data set.

5.3.3. Koli sill (A1220, A1221)

The Sm-Nd studies from the 300-m-thick Koli lavered sill were focused on the Kaunisniemi section described previously by Vuollo (1988). The mineral separation from two pyroxenites yielded fairly clean fractions of clinopyroxene, but albitic plagioclase is generally turbid and yellowish. The Sm-Nd data are shown in Table 4, which also includes some older analyses having relatively large errors. The twelve analyses available on five whole rocks and mineral fractions do not define an isochron, as shown by the calculated MSWD value of 30. If the two analyses on albitic plagioclase are excluded, an age of 2201 \pm 58 Ma and an initial $\epsilon_{_{Nd}}$ value of + 1.0 can be calculated (Fig. 18). The slightly elevated MSWD value of 3.5 may be due to underestimation of the analytical error (in old analyses), inclusion of samples from several locations in the same regression or most likely due to opening of the Sm-Nd system during metamorphism. The metamorphic effects on albitic plagioclase are obvious, as can be seen from the Sm-Nd age estimate of c. 1.9 Ga calculated for the whole rock-plagioclase pairs of samples A1220 and A1221 (Fig. 18).

The analysis on granophyre provides an initial $\epsilon_{Nd}(2220 \text{ Ma})$ value of +0.8 which is similar to the

cumulates of the main series. This suggests that the granophyre in the upper part of the sill was formed from evolved magma without any significant contamination from the adjacent Archean country rocks.

6. Discussion and conclusions

6.1. Timing of the GWA magmatism

The previous age determinations using the conventional U-Pb method have yielded dates from 1.97 to 2.22 Ga for intrusions that have petrological characteristics and field relationships typical of the gabbro-wehrlite association. This age range probably reflects the degree of preservation of the dated zircons and the GWA magmatism was not such a longlived event (Hanski et al., 2001), but rather could have been even shorter in duration than the resolution of our dating methods. To reach this conclusion has required a large number of conventional zircon analyses, supplemented by occasional baddeleyites, from various parts of eastern and northern Finland. The new data acquired by the SIMS techni-



Fig. 18. Sm-Nd isotope data for samples A1220 and A1221 from the Koli sill. A linear regression of analyses on whole-rock samples and clinopyroxene separates gives an isochron age of 2201 ± 58 Ma, while plagioclase-whole rock pairs yield ages of c. 1900 Ma.

que, which is able to circumvent problems caused by zircon alteration, has confirmed this view. These results together with BSE images and electron microprobe analyses provide us now with a clearer understanding of the large variation of the U-Pb isotopic disturbance in zircon shown by the previous ID-TIMS studies. The most well-preserved zircon grains are found in the GWA intrusions in Central Lapland, and hence the zircon U-Pb ages obtained for the Haaskalehto and Harjunoja intrusions, 2220 \pm 11 Ma and 2222 \pm 6 Ma (Tyrväinen, 1983; Räsänen & Huhma, 2001), can be taken as the most representative determinations of the timing of the GWA magmatism in Finland.

6.2. Alteration of zircon by fluids

Several mechanisms have been proposed to control the degree of U-Pb isotopic discordance in zircon (see the review by Gebauer and Grünenfelder, 1979). One of these is low-temperature hydrothermal alteration of radiation-damaged zircon as originally put forward by Krogh and Davis (1975). Several lines of evidence make the alteration model most appealing in our case, including the high contents of non-formula components (CaO, Al₂O₃, FeO, MnO, H₂O) in zircon grains, their inferred hydrous nature, the general lack of magmatic zoning features (cf. Connelly, 2001), and other physical indications of strong alteration. In Fig. 19, our ion microprobe analyses have been divided into two groups on the basis of the BSE intensity of the analyzed spots. There is an obvious relationship between the average atomic number of zircon (degree of hydration and related alteration) and the degree of discordance as the domains displaying low BSE intensities have experienced the largest amount of Pb loss. Note that a high BSE intensity (light shade) does not necessarily indicate a non-altered nature of zircon. Recrystallized parts of the grains may be even more light-colored than pristine zones. This is well exemplified by Figs. 7F and 12E.

Geisler and Schleicher (2000) have studied the relationship between the chemical composition of some non-metamorphosed zircons and their age discordance. They observed a negative correlation between the CaO content and apparent U-Thtotal Pb ages with the highest, concordant ages obtained for those zircon domains that had CaO contents of less than 0.2 wt.%. In contrast, the dates yielded by the domains having elevated CaO concentrations were drastically lower than the accepted



Fig. 19. Concordia plot of the SIMS U–Pb data divided into two groups on the basis of the BSE intensity of the analyzed spot.

magmatic ages. Geisler and Schleicher (2000) attributed the high contents of Ca and other non-formula elements to ion exchange processes during fluid-induced alteration processes that led to removal of Zr, Si and Pb from the zircon lattice. Later Geisler et al. (2001, 2002, 2003) demonstrated experimentally that in a large temperature range between 175 °C and 650 °C, hydrothermal alteration of partially metamict zircon through an interaction of CaCl₂-bearing solutions can generate zircon domains with elevated CaO contents up to c. 2 wt.%. In these experiments, addition of CaO was accompanied by extensive hydration of the mineral with resultant H₂O contents reaching 8 wt.%. Both experimental data and natural examples show that, due to the incompatibility of Pb2+ in the newly grown zircon phase, radiogenic Pb is easily rejected from the mineral grains during hydrothermal annealing of partially metamict zones, resulting in a strong disturbance of the U-Pb isotope system (Pidgeon et al., 1966; 1973; Krogh and Davis, 1975; Geisler et al., 2003a,b).

Based on our electron microprobe and SIMS data from a single zircon grain from the Susivaara sill, the same kind of relationship between the CaO content and discordance, as observed by Geisler and Schleicher (2000), may be a general feature of altered zircon grains in the GWA intrusions. We suggest that the high CaO contents and high degree of discordance are related to hydrothermal alteration of the zircon grains by $CaCl_2$ -bearing fluids and the low analytical totals of the electron microprobe analyses reflect high H₂O contents of the altered zircon domains.

Apart from the loss of radiogenic Pb and gain of Ca, hydrothermal alteration of metamict zircon can result in significant changes in concentrations of other minor or trace elements such as Sr, Ba, Al, Fe, Mn, Y, REE, and most importantly from the point of view of geochronology, U and Th (Krogh and Davis, 1975; Geisler et al., 2003b). The behavior of REE and Y during the alteration processes may be complicated. For example, Geisler et al. (2003b) observed a positive correlation between the CaO and Y₂O₂ contents in natural altered zircon grains from an Egyptian granite, whereas in our study, the altered domains show an Y₂O₃ content of about a half of that observed in the least altered domains (Fig. 14C). Evidently, a more comprehensive electron microprobe and LA-ICP-MS study of zircon in the GWA rocks is needed.

High U and Th contents induce high a-decay

doses and high degrees of metamictization and, consequently, the altered zircon grains may exhibit a positive correlation between the U and Th contents and the degree of U-Pb isotopic discordance. Figure 20 shows U concentrations as a function of ²⁰⁷Pb/ ²³⁵U as determined by SIMS. Using the ²⁰⁷Pb/²³⁵U ratio as an index of discordance, it can be seen that there is no clear correlation between U content and degree of discordance. The same applies to the Th content and Th/U which varies widely from lower than 1.0 to higher than 10. Identification of independent Th-rich mineral grains (thorite) in the lowintensity BSE zircon domains is noteworthy in this context as their presence indicates secondary mobility of Th and within the spot of the beam, they may strongly affect the Th/U ratio and Th concentration in the analytical results. The lack of correlation between the U content and 207Pb/235U in severely altered zircons was regarded by Cherniak and Watson (2000) as an indication of non-diffusional processes, such as recrystallization and various fluidassisted processes that could significantly alter the Pb isotope ratios and U content in zircon. It is noteworthy that even though major loss of radiogenic Pb from zircons and consequent decrease in ²⁰⁷Pb/ ²³⁵U and ²⁰⁶Pb/²³⁸U seem to have taken place in our

case, hydrothermal alteration may also have lead to gain or loss of U and/or Th.

As plagioclase in the GWA intrusions is almost ubiquitously albitized, the source of CaO of the altered zircons could be envisaged to be the anorthite component of plagioclase liberated during albitization. This is consistent with the most well-preserved and concordant zircon being found in the sample from the Haaskalehto intrusion, one of the GWA intrusions that have a relatively well-preserved primary magmatic mineralogy including labradoritic plagioclase. The CaO released during albitization of plagioclase could also have participated in the change of ilmenomagnetite to sphene, a phenomenon that is often observed in the GWA sills. Alteration of ilmenomagnetite was accompanied by liberation of Fe, which may have been partly consumed in the formation of secondary biotite, but could also have been incorporated into other secondary minerals including altered domains of zircon grains.

The more or less one-stage process described above may seem reasonable, but it is not easily fit with the observed U-Pb isotopic systematics of zircon and sphene. Taken together, the ID-TIMS and SIMS analyses suggest a multi-episodic history for the rocks. First, concordant ID-TIMS sphene ages



Fig. 20. Uranium - ²⁰⁷Pb/²³⁵U relations for zircon grains as determined by ion microprobe. ²⁰⁷Pb/²³⁵U is used as an index of discordance.

of c. 2.2 Ga indicate that some alteration phenomena were very early and took place perhaps already in the late-magmatic stage, though we must add that in the GWA intrusions sphene is not exclusively concordant at 2.2 Ga but partially suffered from later isotopic disturbances together with zircon (see figure 3 in Hanski et al. 2001). Second, some SIMS ages on zircon are nearly concordant at 1.8-1.9 Ga suggesting a resetting event related to the Svecofennian orogeny, which is also supported by our Nd isotopic data on plagioclase from Koli and some other zircon studies from northern Finland (Mertanen et al., 1989). Third, some zircon grains give very discordant SIMS results indicative of a very recent Pb loss as these minerals contain sufficient uranium to have developed higher 206Pb/238U and ²⁰⁷Pb/²³⁵U ratios over time had the major Pb loss occurred much earlier.

6.3. Alteration event

Given a multi-episodic history of zircon in the GWA intrusions, the large scatter in the lower intercept ages of the discordias obtained in this and previous studies becomes understandable. These ages range from c. 150 to 960 Ma and appear to be geologically meaningless, which is in accord with the view that the lower intercept ages in general have no geological significance (e.g. Mezger and Krogstad, 1997). However, lower intercept ages obtained from spot analysis of zircon that contain a high concentration of non-formula components (e.g. CaO) can be sufficiently discordant (close to the lower intercept with concordia) to precisely date a leaching event and relate it to the regional thermotectonic history (Geisler et al., 2001). For example, Geisler et al. (2003b) obtained 100 % discordant SHRIMP ages of c. 20 Ma for hydrothermally altered, Ca-bearing zircons from a c. 620 Ma Egyptian granite mentioned above. They were able to link this alteration event and similar fission track ages of apatite to the mafic magmatism associated with the main rifting phase of the Red Sea. The most discordant analytical points of our SIMS study are not sufficiently close to the concordia to yield a precise time for the suggested relatively recent Pb loss event,

but nevertheless indicate a Paleozoic age of some hundred million years.

It is worth noting that even though the variation in discordance shown by our SIMS data is large, it is not larger than that displayed by the previously published zircon and baddeleyite ID-TIMS analyses from the GWA intrusions (Hanski et al., 2001, figure 2). Figure 21 exhibits all available SIMS and ID-TIMS data on a concordia diagram. The whole data set can be accounted for by various combinations of ancient and relatively recent Pb loss. The huge variation in the 207Pb/235U and 206Pb/238U ratios generates a long array that converges towards a lower intercept age of c. 300 Ma. Some ID-TIMS data points in this array even show a higher degree of discordance than the most discordant in situ SIMS analyses. With the help of the ion microprobe, zircon grains from these most discordant bulk samples could potentially be used to get more precise age constraints on the relatively recent Pb loss deduced from the present zircon data.

Larson and Tullborg (1998) drew attention to the fact that conventional U-Pb zircon data from the Svecofennian domain in Sweden often record late Paleozoic lower intercept ages, which fit with the timing of intense regional Pb mobilization. Similar lower-intercept ages are also common for Paleoproterozoic rocks further east in Finland as exemplified by ID-TIMS measurements from northern Finland in Fig. 22. Larson and Tullborg (1998) attributed the relatively recent lower-intercept ages in the crystalline basement of the shield to leaching of zircons by low-temperature hydrothermal fluids, whose generation was related to burial of the shield beneath upper Paleozoic sedimentary rocks produced by rapid erosion of the Caledonides. Högdahl et al. (2001) observed an almost complete resetting at c. 380 Ma of Paleoproterozoic zircon in a highgrade deformation zone in central Sweden. They suggested that this annealing event affected mostly metamict zircon and was influenced by low-T saline fluids circulating in the basement due to the Caledonian orogeny. Still one reason for heating of the lithosphere of the Fennoscandian Shield in the Paleozoic time may have been the (mantle plumerelated?) magmatic event that generated Devonian



Fig. 21. Concordia plot of the U–Pb SIMS and ID-TIMS data on zircon and baddeleyite of this study and previous studies (after Hanski et al., 2001, figure 2).

Fig. 22. Concordia-based lower-intercept ages for ID-TIMS U-Pb zircon dates of Paleoproterozoic rocks from northern Finland. Also shown as a yellow histogram is the distribution of lower-intercept ages of U-Pb zircon data from the Svecofennian domain in Sweden as published by Larson and Tullborg (1998).

alkaline rocks in the eastern part of the shield (e.g. Downes et al., 2005). It is unclear whether any of the above-mentioned geological processes were responsible for the wide-spread resetting of the U-Pb system in the zircons of the 2.2 Ga GWA intrusions. Anyway, alteration and recrystallization of partially metamict zircon by hydrothermal solutions can take place at fluid temperatures lower than 200 °C and the former existence and timing of such a thermal event may not be easily verified with other isotopic methods than the U-Pb zircon dating.

6.4. Nd isotopic characteristics of the GWA magmatism

Earlier Sm-Nd isotope data on the Runkausvaara sill published by Huhma et al. (1990) yielded an imprecise isochron with an age of 2205 ± 220 Ma and initial ε_{Nd} of 0.1 ± 1.4. Our new, somewhat more precise isochron data give similar results with the initial ε_{Nd} values being consistently between 0.0 and +1.0. Combining all relevant data of a total of 30 analyses gives an age of 2223 ± 28 Ma and an

initial isotopic composition corresponding to ε_{Nd} of +0.6. This age is in good agreement with the zircon data and the initial ε_{Nd} value is thought to be close to that of the parental magma. Figure 23 shows a diagram of calculated initial ε_{Nd} (2220 Ma) values vs. Sm/Nd ratios for mineral and whole-rock samples, constructed using data from the present study and Huhma et al. (1990). It is evident that plagioclase data are very scattered, while pyroxenes and most whole-rock samples give initial ε_{Nd} values close to the above mentioned isochron-based, preferred value of +0.6.

The initial Nd isotopic composition of the GWA magma with a slightly positive initial $\varepsilon_{Nd}(2220 \text{ Ma})$ differs from that of the contemporaneous depleted mantle reservoir ($\varepsilon_{Nd} = -+3$) as defined by the isotopic evolution of DePaolo (1981). This deviation may have resulted from interaction between the magma and crustal rocks. However, our Nd isotopic data as a whole indicate that the magma that produced the GWA intrusions in various parts of northern and eastern Finland was isotopically homogeneous and hence we conclude that

it did not undergo significant upper crustal contamination upon emplacement and subsequent fractional crystallization.

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Fig. 23. ϵ_{Nd} (2220 Ma) vs. ¹⁴⁷Sm/ ¹⁴⁴Nd diagram for whole-rock and mineral analyses. Data from this study and Huhma et al. (1990).

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