The basal dunite of the Precambrian mafic-ultramafic Näränkävaara intrusion: Petrogenetic considerations and implications to exploration

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Abstract
Several mafic-ultramafic layered intrusions were emplaced in the Fennoscandian Shield during wide-spread mantle-sourced magmatism at 2.5–2.4 Ga. The Näränkävaara intrusion (surface area 5 × 30 km²), northeastern Finland, contains a 1.5–2 km thick basal dunite (not dated), and a 1.5 km thick layered series (2436 ± 5 Ma). A newly discovered marginal series between the basal dunite and the layered series indicates that the basal dunite is older, and highlights the need for further study on their relationship. Along its southern basement contact, the basal dunite contains a 200–300 m thick zone of olivine ortho- and mesocumulates, but the bulk of it is composed of olivine adcumulates and lesser olivine-orthopyroxene heteradcumulates. Based on whole-rock geochemistry, the basal dunite is divided into a low-Fe zone (average FeO t 10.2 wt% and Ni 2250 ppm) and a high-Fe zone (average FeO 12.5 wt% and Ni 1700 ppm). Both zones have high MgO (32–47 wt%) and varying Cr (830–5160 ppm) and Al₂O₃/TiO₂ (16–26). Textural and geochemical layering is similar along the 30 km strike of the basal dunite. A LREE-enriched high-MgO basaltic parental magma composition (13–18 wt% MgO) is inferred for the basal dunite from olivine–melt mixing trends in orthocumulates. The dunite exhibits at least two geochemical reversals as well as abundant low-porosity adcumulates, poikilitic chromite, and bimodal olivine, suggesting formation in a high-volume open magmatic system. Significant similarity in major and trace element compositions with the Näränkävaara layered series and the Burakovsky intrusion and Vetreny belt extrusives in Russian Karelia suggests that the basal dunite belongs to the Fennoscandian 2.5–2.4 Ga mafic layered intrusions. An Archean komatiitic origin for the dunite body cannot be completely ruled out, however. Distinct Ni-depletion in olivine is found in the basal dunite from the low-Fe zone to the high-Fe zone (3200 versus 2200 ppm). This depletion does not correlate with Fo contents, which suggests that it is not related to olivine fractionation. The basal dunite may thus have potential for Ni-(Cu-Co-PGE) sulfide mineralization.

Keywords Tonio-Näränkävaara belt · Näränkävaara · Layered intrusion · Dunite · Komatiite · Ni-criticality

Introduction
The Tornio-Näränkävaara intrusive belt (TNB) in northern Finland is critical for orthomagmatic PGE-Ni-Cu-Co and Cr-V-Ti-Fe deposits (Huhtelin 2015; Iljina et al. 2015; Makkonen et al. 2017). The easternmost of these 2.44 Ga layered intrusions (Iljina and Hanski 2005) is the mafic-ultramafic Näränkävaara intrusion (Fig. 1). The geology and mineral potential of the Näränkävaara intrusion is still relatively poorly known (Alapieti 1982; Järvinen et al. 2020). The presence of a large mass of ultramafic cumulates below the layered series of the Näränkävaara intrusion (Fig. 2a) has been known since the 1960’s (Auranne 1969), but the petrogenesis, mineral potential, and relationship of this basal dunite with
the neighboring layered series have not been resolved. More recent mineral exploration projects have reported the basal dunite to be composed of homogeneous, ~2 km thick olivine adcumulate, and to be continuous for the 30 km length of the intrusion (Fig. 2a) (Iljina 2003; Vesanto 2003; Lahinen 2005). The basal dunite exhibits several features commonly thought of as komatiitic, being mostly composed of tightly packed monomineralic olivine adcumulate, sometimes with bimodal olivine and poikilitic chromite (cf. Barnes and Hill 1995). The recent discovery of a marginal series between the Näränkävaara layered series and the basal dunite indicates that the basal dunite must be at least marginally older (Järvinen et al. 2020).

Here, the Näränkävaara basal dunite is hypothesized to represent either a feeder channel cumulate related to the same Fennoscandian 2.5–2.4 Ga magmatism as the 2436 Ma Näränkävaara layered series (Fig. 2a) (Alapieti 1982; Kulikov et al. 2010), or an older komatiitic wall-rock for the layered series extending into the Näränkävaara area from the nearby Archean Takanen or Suomussalmi greenstone belts (Iljina 2003; Papunen et al. 2009; Mikkola et al. 2011). Mantle-derived ultramafic cumulates are known to host orthomagmatic sulfide deposits (e.g. Begg et al. 2010; Barnes et al. 2016a); in the former case it could have potential for magmatic sulfide mineralizations linked to feeder channel cumulates, such as in Jinchuan (Chai and Naldrett 1992), and in the latter case for komatiite-hosted sulfide mineralizations (Konnunaho 2016; Makkonen et al. 2017).

New drilling and mapping was conducted in the Näränkävaara basal dunite during 2017–2018 by the Geological Survey of Finland (GTK) (Halkoaho et al. 2019). Based on these new results and a compilation of pre-existing
data (Alapieti 1982; Iljina 2003; Vesananto 2003; Lahtinen 2005; Akkerman 2008; Telenvuo 2017; Järvinen et al. 2020), we characterize the Näränkävaara basal dunite, model its petrogenesis, and estimate its Ni-(Cu-Co-PGE) potential.

Geological background for the 2.5–2.4 Ga magmatism in the Fennoscandian shield and the Tornio-Näränkävaara belt in Finland

The northern part of the Fennoscandian shield (mainly northern Finland and NW Russia) hosts 20–40 mafic-ultramafic layered intrusions and associated dyke swarms with ages between 2.53–2.39 Ga (Fig. 1) (Alapieti et al. 1990; Amelin et al. 1995; Iljina and Hanski 2005; Bayanova et al. 2009). These intrusions are located mainly along rifted intracontinental paleoplate boundaries in the Archean Karelia and Kola cratons (Bayanova et al. 2009; Kulikov et al. 2010; Tiira et al. 2014).

The five intrusions and intrusion complexes comprising the TNB have an average age of 2.44 Ga (Fig. 1) (Iljina and Hanski 2005). They are typical layered intrusions with a layered and marginal series, and are divided into three compositional groups: (1) mafic-ultramafic, which contain a peridotitic–pyroxenitic lower zone (Näränkävaara and Kemi), (2) mafic, which are dominated by gabbronoritic rocks (Koillismaa), and (3) megacyclic (Penikat and Portimo), which are characterized by macrorhythmic compositional reversals (Alapieti and Lahtinen 2002).

Stratigraphically, the intrusions of the TNB are found at the contacts of late Archean granitoids and overlying Paleoproterozoic supracrustal rocks (Alapieti et al. 1990; Amelin et al. 1995; Iljina and Hanski 2005; Bayanova et al. 2009).
In contrast to other intrusions of the TNB, the Näränkävaara intrusion is completely surrounded by the Archean granite-gneiss basement complex (Fig. 1) (Hölttä et al. 2012). In addition, Näränkävaara is the only intrusion in the TNB associated with voluminous dunitic cumulates (Alapieti 1982; Järvinen et al. 2020).

The parental magmas of the Finnish 2.5–2.4 Ga mafic layered intrusions are thought to be (boninite-like) siliceous high-Mg basalts (Alapieti et al. 1990; Iljina and Hanski 2005). Whole-rock and isotope geochemical systematics (average initial $\varepsilon_{\text{Nd}}$ of $\sim$–2) point to crustal contamination of primitive magmas from a mantle plume source (Puchtel et al. 1997; Hanski et al. 2001; Vuollo and Huhma 2005; Yang et al. 2016; Rämö et al. 2017; Maier et al. 2018). Intrusive and extrusive formations with similar compositions are found in Russian Karelia, for example the komatiitic basalts and dykes of the Vetreny belt, and the mafic-ultramafic Burakovsky layered intrusion (Fig. 1) (Puchtel et al. 1997; Chistyakov and Sharkerov 2008).

The Näränkävaara layered intrusion has been included as a part of the Koillismaa layered intrusion complex (KLIC) (Alapieti 1982). The KLIC comprises (1) the tectonically dismembered western intrusion blocks of the Koillismaa layered intrusion (Karinen 2010), (2) the eastern Näränkävaara intrusion, and (3) a strong positive gravity and magnetic anomaly connecting the first two (dashed line in Fig. 1). Alapieti (1982) interpreted the Koillismaa and Näränkävaara intrusions to be co-genetic, with the Näränkävaara intrusion representing the lowest exposed stratigraphic level of the KLIC. The connecting geophysical anomaly – hitherto not intersected in drilling – was thought to be a magmatic conduit that fed the two intrusions. The recently discovered (likely Archean) Takanen schist belt (Fig. 2a) coincides with the geophysical anomaly (Iljina 2003; Salmirinne and Iljina 2003; Iljina et al. 2006).

**Geology of the Näränkävaara intrusion**

The mafic-ultramafic Näränkävaara intrusion (Fig. 2a) is mostly unexposed but shows as a strong positive magnetic anomaly about 30 km long and 5 km wide, with the basal dunite also being highly and untypically conductive (Niskanen and Jokinen 2018). Gravimetric and magnetometric measurements indicate that the intrusion extends to a depth of 5–10 km (Elo 1992; Salmirinne and Iljina 2003). A large SW–NE trending fault separates the Näränkävaara intrusion into two blocks (NW and SE block).

The intrusion comprises the following three distinct igneous bodies (or series) that are partly or dominantly composed of ultramafic cumulates (Fig. 2a): (1) The southernmost unit consists of a 1–2 km thick basal dunite (not dated thus far) that is composed of dunite and minor peridotite and pyroxenite (Vesanto 2003; Järvinen et al. 2020). This unit is in the focus of the present study. (2) The central, 2436 ± 5 Ma (2σ) old Näränkävaara layered series is composed of a 700 m thick peridotitic–pyroxenitic ultramafic zone, and a 600 m gabbroic–dioritic mafic zone (Alapieti 1982). In the SE block of the intrusion, a poorly developed 10 m thick gabbronoritic marginal series was recently found in the contact between the basal dunite and the layered series, grading into the layered series (Järvinen et al. 2020). (3) A series of smaller elongate intrusions runs parallel to the northern contact of the layered series and composed of poikilitic harzburgite (Alapieti et al. 1979). These northern harzburgites have been drilled (Iljina 2003) but have not been investigated in detail.

In the layered series, the dip of igneous layering in the NW block is 30 ° to the NE, while in the SE block it is 5–15 ° to the SW (Fig. 2a) (Alapieti et al. 1979). In addition to the large fault splitting the intrusion, there are several strike-slip faults along the short axis of the body, and dip-slip faults with strikes along the long axis. Apart from faulting the intrusion is mostly undeformed. The layered series is relatively unaltered, whereas the basal dunite is thoroughly serpentined.

One previous drilling profile located in the NW block of the intrusion intersects the basal dunite, but includes several gaps up to 600 m in thickness. Järvinen et al. (2020) divided the basal dunite along this NW cross section into three olivine cumulate units (see Fig. S1 in electronic supplementary material), from south to north: (1) a harzburgitic olivine mesocumulate transition zone in contact with the basement complex and characterized by distinctly higher contents of incompatible elements compared to the other two units, (2) a homogeneous olivine adcumulate center section characterized by relatively low Cr contents (1000–1500 ppm Cr) compared to the other two units (2000–4000 ppm Cr), and (3) an olivine adc- to mesocumulate unit characterized by coarse poikilitic orthopyroxene, in contact with the layered series. Orientation of igneous layering in the NW block of the basal dunite is poorly constrained, but is assumed to be subvertical at the southern basement contact and to get shallower towards the north, being about 45 ° at the basal dunite–layered series contact. No gabbronoritic marginal series is found at this contact in the NW block, unlike in the SE block.

In addition to the previously mentioned cumulate units, 600 m of homogeneous olivine adcumulate was intersected in one drill hole on the northern side of the layered series in the SE block of the intrusion (indicated as ‘northern dunite’ in Fig. 2a) (Akkerman 2008). This northern dunite has a much lower Al$_2$O$_3$/TiO$_2$ ratio than the main basal dunite body on south side of the layered series (average of 7 versus 21), and has not been found elsewhere.

**Materials and methods**

The materials for this study were obtained during new drilling and mapping activities by the Geological Survey of Finland in
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2017–2018 (Halkoaho et al. 2019). New drilling and sampling is located in the SE block of the Näränkävaara intrusion, and comprises 10 drill holes with 2380 m of drill core and 109 analyzed whole-rock samples (drill holes R1–R10 in Fig. 2b), and 18 surface samples. Pre-existing data has been taken into consideration for purposes of petrographical and geochemical correlations and interpretations, but has generally not been plotted in diagrams for sake of clarity. All sample materials used, and sources used for data compilation, are listed in Table S1.

Full whole-rock analysis results, method descriptions, detection limits, and calibration materials used are listed in Table S2. Explanations of method codes and method descriptions can also be found in the reference Labtium Oy (2015). Major and some minor elements were determined with a Panalytical Axios PW 4400 X-ray fluorescence (XRF) spectrometer from pressed powder pellets (Labtium Oy method code 176X). Platinum, Pd and Au were determined by Pb-fire assay fusion followed by graphite furnace atomic absorption spectrometry (GFAAS; method code 704 U). For the rare-earth elements (REE) and some trace elements, sample solutions were prepared by HF–HClO₄ digestion and lithium metaborate–sodium perborate fusion, followed by inductively coupled plasma mass spectrometry (ICP-MS) with a Thermo Electron iCAP Qc (method 308 M), or optical emission spectrometry (ICP-OES) with a Thermo Electron iCAP 6500 (method 308P), with final results calculated as an average of three replicate analyses. For all analysis methods, duplicate analyses were made for 5% of samples, with a coefficient of variation generally below 5%. All results of whole-rock analyses presented in this study are normalized to volatile-free 100 wt%.

For parental magma modeling, and for calculating whole-rock Mg# [Mg# = Mg/(Mg + Fe²⁺)], whole-rock Fe₂O₃/FeO ratios have been calculated from total FeO (Fe₂O₃) according to the method of Barnes et al. (2007). This method was developed for pure olivine cumulates (komatitites) where Fe₂O₃ contents is assumed to be zero, as Fe³⁺ is incompatible in olivine. A linear increase in whole-rock Fe₂O₃/FeO ratio from 0 to 0.1 is calculated as a function of decreasing modal olivine and increasing liquid component. Olivine mode is proxied using whole-rock MgO contents, so that a Fe₂O₃/FeO ratio of 0.1 results at <25 wt% MgO, and a ratio of 0 at >50 wt% MgO, with a constant linear variation in the ratio in between (Barnes et al. 2007).

Petrography of 208 thin sections were examined with cross-polarised transmitted-light microscopy. Chemical compositions of minerals from the SE block of the Näränkävaara body were analyzed by wavelength dispersive spectroscopy with a Cameca SX100 electron probe micro-analyzer (EPMA) (464 spot analyses from 225 mineral grains from 40 thin sections). Full results along with analyzing conditions, X-ray lines analysed and calibration materials used are listed in Table S2. Standard deviation based on repeated standard measurements is <1%. Chromite analyses have been recalculated assuming stoichiometry to an ideal XY₂O₄ formula according to Barnes (1998) [with X = (Fe²⁺, Ni, Mn, Co, Zn) and Y = (Cr, Fe³⁺, Al)].

Results

Generalities

Eight of the ten new drill holes are located along the southern contact between the basal dunite and the basement complex (R3–R10 in Fig. 2b), and two drill holes are located at the northern contact between the basal dunite and the layered series (R1–2 in Fig. 2b), with surface samples in between (white triangles in Fig. 2b). Based on this new sampling, a new geological map and cross section of the SE block of the Näränkävaara intrusion is presented in Figs. 2b and c. The basal dunite is subdivided into ten units based on differences in petrography and whole-rock geochemistry as described below. These units are numbered 1–10 in Fig. 2b and c, and their characteristics are summarized in Table 1. The cross section in Fig. 2 spans a horizontal length of 2500 m, but includes horizontal gaps of up to 1000 m between sample points due to lack of sampling in the center of the basal dunite. Even with poor sampling density, it is possible to correlate this cross section with the previous cross section from the NW block of the intrusion by similarities in whole-rock compositions and petrography (Fig. S1; Järvinen et al. 2020).

We have aimed to use cumulus terminology descriptively according to Irvine (1982), without genetic connotations to the intrusive or extrusive origin of the rocks. A cumulate consists of varying amounts of accumulated (near) liquidus crystals and trapped intercumulus melt. An adcumulate contains >93 vol% accumulated crystals, mesocumulate 93–75 vol%, and orthocumulate 75–50 vol%. In the basal dunite, adcumulates are typically dunites and meso- and orthocumulates are harzburgites and lherzolites. A heteradegumulate is a cumulate with a poikilitic texture and a very minor component of recognizable trapped intercumulus melt (Wager et al. 1960; Barnes et al. 2016b). The presence of trapped intercumulus melt in olivine and orthopyroxene rich cumulates, like found in Näränkävaara, can usually be recognized from elevated contents of incompatible elements (e.g., Al, Ti, P, Zr) in whole-rock analysis. As such, adcumulate and heteradegumulate rocks tend to exhibit adcumulate chemistries poor in incompatible elements, and orthocumulate rocks tend to exhibit orthocumulate chemistries relatively rich in incompatible elements.

Petrography of the basal dunite

Ortho- and mesocumulates of the SW border zone

A 200–400 m thick border zone composed of olivine ortho- and mesocumulates and minor adcumulates (units #1–2 in
Table 1  Main petrographical and geochemical characteristics of lithological units of the Närinkävaara basal dunite

| Petrography                  | Heights (m)  | Height (m) | Petrography                  | Geochemistry                                                                 | Mg# | Fo in olivine |
|-----------------------------|--------------|------------|-----------------------------|-------------------------------------------------------------------------------|-----|---------------|
| Orthocumulate and mesocumulate Low-Fe (High-Ni) | (0-20)       | 0-250      | Strongly altered contact zone of meta-peridotite, tremolite-rock, soapstone | Olivine orthocumulate grading into mesocumulate with distance from contact; intercumulus consists of pyroxenes, minor plagioclase, biotite, trace chromeite, sulfides; euhedral-subhedral chromeite | 89.1 | 88.0^4 89.8 14.7 16.1 16.4 18.1 |
| Orthocumulate and mesocumulate Low-Fe (Moderate-Ni) | 2            | 0-250      | Orthocumulate chemistry; olivine-melt mixes; moderate Ni (average 2000 ppm) | Orthocumulate chemistry; olivine-melt mixes; moderate Ni (average 2000 ppm) | 87.3 | 87.8 89 13.0 14.5 14.8 16.5 |
| Adcumulate Low-Fe (High-Ni) | 3            | 250-550    | Adcumulate chemistry (incompatible element rich); olivine-melt mixes; high Ni (average 2500 ppm) | Adcumulate chemistry (incompatible element poor); high Ni | 89.5 | 86.9 |
| Adcumulate Low-Fe (Moderate-Ni) | 4            | 550-625    | Orthopyroxene adcumulate (incompatible element rich); low Ni (average 4000 ppm) | Orthopyroxene adcumulate (incompatible element poor); high Ni | 88.5 |
| Adcumulate Low-Fe (?) | 5            | 625-725    | Adcumulate chemistry; low Ni (average 1700 ppm) and Cr (average 1200 ppm) | Adcumulate chemistry; low Ni (average 1700 ppm) and Cr (average 1200 ppm) | 87.2 | 86.3 |
| Adcumulate High-Fe | 6            | 725-750    | Adcumulate chemistry; low MgO and Mg# for olivine adcumulate | Adcumulate chemistry; low MgO and Mg# for olivine adcumulate | 85.7 | 85.1 |
| Adcumulate High-Fe | 7            | 750-1900?  | Adcumulate chemistry, low Ni and Cr | Adcumulate chemistry, low Ni and Cr | 87 | 87.7 87.3 |
| Adcumulate High-Fe | 8            | 1900?-2500 | Adcumulate chemistry; low Ni and Cr | Adcumulate chemistry; low Ni and Cr | 86.6 | 87.6 87.3 |
| Adcumulate High-Fe | 9            | 2275-2252  | Adcumulate chemistry; low Ni and Cr | Adcumulate chemistry; low Ni and Cr | 86.4 | 87.0 86.5 |

1  Average Mg# for unit from whole-rock XRF analyses; Mg# = Mg/(Mg + Fe^{2+}); FeO/Fe_{2}O_{3} ratio calculated according to Barnes et al. (2007), see text.  
2  Olivine composition determined from linear regression of whole-rock Mg/Al vs. Fe/Al (see Makkonen et al. 2017);  
3  Intersection on a MgO-FeO diagram of a whole-rock sample regression line and a model liquid composition in equilibrium with highest Fo olivine found; Parameters Fo_{90.90} and KD 0.30-0.33 were used to calculate compositions of different equilibrium liquids (see text and Fig. 5);  
4  Fo_{88.6} from boulder; EPMA = electron probe micro-analyzer, KD = Olivine-melt Mg-Fe exchange coefficient.
Fig. 2 and Table 1) is found all along the SW contact between the basal dunite and the Archean basement complex. The basement contact is subvertical or dips steeply to the NE (75–90°). The granite-gneiss basement is hornfelsed for a depth of at least 0.5 m, and leached and broken for up to 5 m from the contact. The immediate contact zone is <20 m thick and composed of fine grained metaperidotites and tremolite chlorite schists. The tremolite schists are intensely sheared and have probably acted as a deformational buffer zone for post-emplacement tectonic movements. Degree of alteration rapidly decreases with distance from contact. Meter-scale basement xenoliths and decimeter-scale potassic to dioritic veins derived from partially molten basement are sometimes found up to 250 m away from the contact.

The olivine ortho- and mesocumulates forming the main mass of the border zone (units #1–2) are strongly serpentinized but with undeformed fabrics, showing relict cumulus textures with pseudomorphs after olivine (Fig. 3a and b). Rare samples contain unaltered olivine and pyroxenes. Amount of olivine increases with distance from the basement contact, from about 40–60 vol% to 90–95 vol% over 200–300 m. Olivine grain size is typically between 0.5–3 mm, but intersections with coarser up to 10 mm grains are found in all drill holes. Olivine occasionally defines magmatic lamination (Fig. 3b). An evenly distributed network of intercumulus material is characteristic for these rocks (Fig. 3a and b). Intercumulus consists of orthopyroxene and lesser clinopyroxene, with minor phlogopite and plagioclase. Pyroxenes are poikilitic with grain sizes between 5 and 10 mm (Fig. 3a). Fine-grained euhedral-subhedral chromite typically appears in trace amounts (0.5–2 vol%). Chromite is also found as inclusions in olivine.

Sulfides with primary magmatic textures are very rarely found, whereas trace amounts of fine-grained disaggregated sulfides (metamorphic or remobilized) are common. Trace amounts of very fine-grained Fe-Ni-Cu sulfides were commonly found during preliminary EPMA mineral analyses, either as discrete grains or intergrown with oxides. No Pt or Pd phases were found except for one 1 μm grain.

**Adcumulates and heteradcumulates**

Decrease of intercumulus material to small isolated pockets (typically <3 vol%) marks the beginning of the approximately 2000 m thick main adcumulate mass of the basal dunite (units

![Fig. 3](https://example.com/fig3.png)

**Fig. 3** a-f Plane-polarized light photomicrographs showing thin-sections of main rock-types in the Näränkävaara basal dunite; length of black scale-bar is 5 mm; a Border zone orthocumulate (unit #1); serpentinized cumulus olivine with intercumulus showing dark poikilitic pseudomorphs after pyroxene (sample R6 89.80 m); b Border zone mesocumulate (unit #2) exhibiting magmatic lamination along horizontal plane (sample R7 157.20 m); c Ni-rich olivine adcumulate from low-Fe zone (unit #3) with euhedral and interstitial chromite (sample VJJA-88); d Olivine orthopyroxene heteradcumulate (unit #9) with a dark poikilitic orthopyroxene pseudomorph enclosing tightly packed serpentinized olivine (sample R2 72.30); e Harzburgitic adcumulate interlayer (unit #10) close to contact with layered series (sample R2 30.40 m); f Olivine adcumulate from the NW block of the Näränkävaara intrusion, high-Fe zone, showing bi-modal olivine and interstitial to lobate chromite (correlates to units #6–8 in the SE block; image is a scanned thin section; sample R303 127.43 m)
It is composed of olivine adcumulates and minor orthopyroxenites in the SW (units #3–8), followed by olivine-orthopyroxene heteradcumulate and minor orthopyroxenites in the NE (units #9–10). Olivine adcumulate textures range from subhedral (Fig. 3c) to tightly packed polygonal (lower half of Fig. 3d). Olivine is typically equigranular with grain size between 0.5–3 mm, but rare bimodal olivine is also found in the NW block of the dunite (Fig. 3f). The orthocumulate border zone (units #1–2) most likely grades into the adcumulates, but this boundary is only intersected in one drill hole (R5), where it coincides with a 1 m thick olivine-pегматoidal layer, composed of loosely packed cm-sized subhedral olivine pseudomorphs in a fine grained serpentine groundmass. No other magmatic layering has been identified in the adcumulates, and for the most part the inner structure of the SE block of the basal dunite is poorly constrained.

A ≤ 200 m thick unit of fresh (olivine) orthopyroxene adcumulate (unit #5 in Fig. 2 and Table 1) is found sandwiched between two olivine adcumulate units. This orthopyroxenite unit is notable in two ways. First, it marks the border between the geochemically defined low-Fe and high-Fe zones, as discussed later. Second, a change in chromite textures seems to be associated with the stratigraphic level of this unit. It has been argued that all chromites in ultramafic cumulates are cumulus in origin, i.e., a liquidus phase, regardless of their crystal habit (Barnes 1998; Godel et al. 2013), and thus the terms cumulus and intercumulus are avoided for describing chromite textures. In this study, chromites have been classified as euhedral-subhedral (upper grain in Fig. 4b), interstitial (weakly growing between cumulus grains; lower grain in Fig. 4b, also Fig. 3c), lobate (more extensive fingering between cumulus grains; Fig. 4c), and poikilitic (completely enclosing adjacent grains; Fig. 4d). As labeled in Fig. 2c, below the orthopyroxenite unit (in units #1–4) chromite textures are fine grained euhedral or interstitial, whereas above the orthopyroxenite (in units #6–9) chromite textures are typically lobate or poikilitic. These systematics continue up to the middle of unit #9, where chromite texture changes back to euhedral. In the basal dunite, olivine cumulates with most abundant chromite (up to 1–2 vol%) always show euhedral-subhedral chromite textures. Poikilitic chromite is notably coarser (up to 2 mm) compared to euhedral chromite, but appears in much lower total modal quantities. Units 7–9# contain the coarsest lobate and poikilitic chromites in the basal dunite (Fig. 4c and d).

The heteradcumulate unit (unit #9 in Fig. 2 and Table 1), found along the basal dunite–layered series contact, has a distinct poikilitic texture with sparse pseudomorphic orthopyroxene grains 2–5 cm in diameter enclosing olivine grains (Fig. 3d), but otherwise its petrography is similar to the regular olivine adcumulates. Midway up through the unit, chromite texture changes from coarse poikilitic to more abundant fine-grained euhedral, and continues like this until the layered series contact (Fig. 2c). A relatively heterogeneous 50 m thick unit (unit #10) comprising 0.5–5 m thick interlayers of variable-textured harzburgitic to orthopyroxenitic adcumulates and mesocumulates (Fig. 3e) is found within the heteradcumulate unit, about 100 m before the layered series contact. The change from poikilitic to cumulus orthopyroxene (and also chromite) observed from unit #9 to...

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**Fig. 4** a Whole-rock MgO versus Cr compositions of Närinkääraa basal dunite samples. Olivine adcumulate samples (units #3, 4, 6, 7, 8, and 9) represent either olivine cumulates or olivine chromite cumulates, whereas the SW border zone ortho- and mesocumulate samples (units #1, 2) represent olivine–melt mixtures (±chromite) (Barnes 1998). Labeled fields show (1) modeled basal dunite parental magma composition (“Model PM”), and (2) comparisons to whole-rock compositions from the komatitic basalts of the Vetreny Belt (Puchtel et al. 1997) and the komatitic Kellojärvi ultramafic cumulate complex (Papunen et al. 2009). Colored fields indicate texture of chromites in thin section (see legend): b–c Back-scattered electron images of chromites with thin chromian magnetite rims, number indicates length of picture bottom edge: b lower grain euhedral, upper grain interstitial (sample JSN-87), c lobate (sample JSN-89), d poikilitic (sample VJJA-81)
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unit #10 is in line with the common observation that poikilitic phases precede the appearance of that phase as cumulus (Campbell 1987).

A subvertical tectonized contact between the olivine heteradcumulate (unit #9) and the Näränkävaara layered series is intersected in drill hole R1 at a depth of ~170 m. The contact includes a 10 m thick non-cumulus textured melagabbro-noritic marginal series developed on the side of the layered series (bright red in Fig. 2c) (Järvinen et al. 2020).

Whole-rock chemical compositions

Main results

Results of representative whole-rock analyses of the Näränkävaara basal dunite are presented in Table 2. All samples are MgO-rich (24–47 wt%). Olivine adcumulates contain 45–47.7 wt% MgO, with the exception of one low-Mg# adcumulate unit with only 42 wt% MgO (unit #6 in Table 1 and Fig. 2). In the adcumulates, FeO t ranges between 9.0–13.2 wt%, and increases with decreasing MgO (Mg# 84.4–90.4; Fig. 5). Mesocumulates contain 39–45.5 wt% MgO and orthocumulates 31–38 wt% MgO, but values down to 24 wt% are found in heavily altered (olivine-pyroxenitic?) samples closest to the basement contact. The heteradcumulates in drill holes R1 and R2 (unit #9) typically contain 42–45 wt% MgO, grading down to 32–34 wt% in the orthopyroxene interlayers (unit #10). On Pearce element ratio plots of (Mg + Fe)/Al versus Si/Al, all olivine cumulate and heteradcumulate units plot on straight lines with slopes of 1.7–2.0. This indicates strong olivine control of whole-rock compositions as pure olivine plots on a slope of 2.0.

Based on olivine-compatible element concentrations in whole-rock samples, the basal dunite can be divided into two zones with overlapping MgO contents (Fig. 6a): (1) a more primitive low-Fe zone (units #1–5; downward-triangles in all diagrams), characterized by relatively low FeO t (average 10.2 wt%) and high Ni (average 2250 ppm), with MgO between 32 and 47.3 wt% (Mg# 85.3–90.4); and (2) a more fractionated high-Fe zone (units #6–10; upward-triangles in all diagrams) characterized by relatively high FeO t (average 12.5 wt%) and low Ni (average 1700 ppm), with MgO between 40 and 46.0 (Mg# 85.7–88.6). The low-Fe zone can be further subdivided into a high-Ni subzone with an average of 2500 ppm Ni (units #1 and #3; pink in Fig. 6c and all other diagrams), and a moderate-Ni subzone (units #2 and #4; red in Fig. 6e and all other diagrams) with an average of 2000 ppm Ni. Average Al2O3/TiO2 is slightly higher in the low-Fe zone (22) compared to the high-Fe zone (19), ranging 16–26.

As shown in Figs. 2 and 7, the low-Fe zone (units #1–5) encompasses the ortho- and mesocumulate border zone, the lowermost olivine adcumulates, and the lowermost orthopyroxenite unit. The high-Fe zone (units #6–10) encompasses the rest of the basal dunite up to the layered series contact. The high-Ni subzone of the low-Fe zone has only been intersected in drill holes R6, R7, and R9, and thus does not appear in the geochemical profile in Fig. 7.

The low-Fe and high-Fe zones both have variable Cr concentrations. Chromite texture roughly correlates with whole-rock Cr contents (Fig. 4): poikilitic and lobate chromite is found in high-MgO low-Cr cumulates, and euhedral-subhedral chromite in high-Cr cumulates. Similar MgO-Cr systematics have been described from komatiites (Barnes and Hill 1995; Barnes 1998). In the basal dunite, highest Cr contents (2000–5160 ppm) are found in samples with euhedral and/or weakly interstitial chromite (Fig. 4). Lower Cr contents (<2000 ppm) are found in samples with lobate chromite. Lowest Cr contents (700–850 ppm) are found in samples with poikilitic chromite, which does not appear together with euhedral chromite. Sharp changes in whole-rock Cr contents of about 2000 ppm are associated with a change from anhedral to more abundant euhedral-subhedral chromite (Fig. 7d).

Orthocumulate and heterocumulate compositions

In the adcumulates, olivine-incompatible trace element contents are generally very low or below detection limits. Exception is cobalt (Table 2), which is uniformly higher in the high-Fe zone (average 140 ppm) compared to the low-Fe zone (average 105 ppm).

Variation diagrams in Fig. 6b and d show two linear trends of increasing TiO2 and Al2O3 versus decreasing MgO. The more distinctive one is related to the ortho- and mesocumulates of the border zone (units #1–2; TiO2 < 0.41 wt% and Al2O3 < 9.4 wt%), and the less well-defined to the heteradcumulates and their harzburgitic interlayers (units #9–10; TiO2 < 0.08 wt% and Al2O3 < 1.2 wt%). Regression lines for both trends cross the x-axis at MgO-values between 46 and 48 wt%, which coincides with measured olivine compositions.

For the ortho- and mesocumulates (units #1–2), similar compositional trends with no inflection points are almost invariably found regardless of which elements are plotted against each other (see Fig. 8a and b). This suggests that these rocks are mixtures of cumulus olivine and trapped intercumulus melt. These samples also plot along olivine–melt mixing lines in the MgO versus Ni diagram in Fig. 6c (Barnes and Fiorentini 2012), and, in respect to the moderate-Ni subgroup (unit #2), also in the MgO versus Cr diagram in Fig. 4 (Barnes 1998). This explanation also agrees with petrographic observations, and thus these trends are interpreted as linear olivine–melt mixing lines.

In contrast, in the olivine-orthopyroxene heteradcumulates and their interlayers (units #9–10 in Fig. 6), incompatible elements do not notably increase with decreasing MgO. Instead,
Table 2  Representative results of whole-rock chemical analyses from the Näränkävaara basal dunite, with a modeled parental magma composition for the basal dunite and comparison to the Näränkävaara marginal series gabbronorite

| Zone (subzone) | Low-Fe (high-Ni) | Low-Fe (moderate-Ni) | High-Fe | Marginal Series | Modeled parental magma |
|---------------|------------------|----------------------|--------|----------------|------------------------|
|                | Rocktype³       |                     |        |                |                        |
| Cumulus⁴       | Per             | Serp | Dun | Dun | Dun | Dun | Hzb | Mgbnor | abMCpf |
| Unit #         | 1               | 2    | 3   | 4   | 7   | 8   | 9   | 10   |        |
| Sample⁵        | 1               | 2    | 3   | 4   | 5   | 6   | 7   | 8   | 9      |
| Major oxides (wt%) |        |        |        |        |        |        |        |        |        |
| SiO$_2$        | 45.15           | 46.44 | 42.23 | 41.60 | 43.50 | 41.44 | 41.49 | 46.43 | 54.28 | 50.70 |
| TiO$_2$        | 0.14            | 0.34  | 0.03  | 0.02  | 0.01  | 0.01  | 0.01  | 0.04  | 0.52  | 0.54  |
| Al$_2$O$_3$    | 2.77            | 8.29  | 0.78  | 0.34  | 0.21  | 0.16  | 0.22  | 0.65  | 11.34 | 12.24 |
| FeO$_t$        | 9.85            | 9.65  | 11.49 | 9.99  | 13.46 | 12.29 | 12.46 | 11.34 | 9.42  | 9.23³ |
| MnO            | 0.14            | 0.21  | 0.17  | 0.13  | 0.20  | 0.17  | 0.18  | 0.18  | 0.17  | n.a.  |
| MgO            | 41.61           | 28.83 | 45.16 | 47.69 | 42.58 | 45.49 | 45.13 | 40.49 | 13.79 | 15.59 |
| CaO            | 5.99            | 0.10  | 0.20  | 0.05  | 0.04  | 0.05  | 0.01  | 0.86  | 7.20  | 8.82  |
| Na$_2$O        | 0.08            | 0.07  | <0.03 | <0.03 | <0.03 | <0.03 | <0.03 | <0.03 | 2.33  | 0.10  |
| K$_2$O         | 0.09            | 0.11  | 0.04  | 0.01  | <0.003 | <0.003 | 0.01  | 0.01  | 0.84  | 0.80  |
| P$_2$O$_5$     | 0.02            | 0.07  | 0.01  | 0.01  | <0.006 | <0.006 | <0.006 | <0.006 | 0.10  | n.a.  |
| Trace elements (ppm) |        |        |        |        |        |        |        |        |        |        |
| Cl              | 900             | 200   | 2600  | 1500  | 606   | 900   | 2400  | 1400  | 2300  | n.a.  |
| Cr              | 3620            | 1670  | 1050  | 4210  | 831   | 780.00 | 680   | 3100  | 1820  | 1696  |
| Cu              | 20              | 20    | <20   | <20   | <20   | <20   | <20   | <20   | <20   | 70     |
| Ni              | 2460            | 1220  | 2100  | 2330  | 2023  | 1630.0 | 1610  | 1120  | 530   | 702   |
| S               | 1300            | 400   | 200   | 200   | <100  | <100  | <100  | <100  | 600   | n.a.  |
| Sr              | 20              | 50    | <10   | <10   | <10   | <10   | <10   | <10   | 20    | 210   |
| V               | 30              | 90    | 20    | 20    | <10   | 10.00 | 10    | 30    | 140   | n.a.  |
| Zr              | 10              | 40    | <10   | <10   | <10   | <10   | <10   | <10   | 80    | n.a.  |
| La              | 6.74            | n.a.  | n.a.  | n.a.  | 0.19  | <0.1  | n.a.  | 12.20 | 9.85  |
| Ce              | 13.90           | n.a.  | n.a.  | n.a.  | 0.23  | <0.1  | n.a.  | 24.40 | 20.62 |
| Pr              | 1.65            | n.a.  | n.a.  | n.a.  | <0.1  | <0.04 | n.a.  | 2.83  | 2.57  |
| Nd              | 6.81            | n.a.  | n.a.  | n.a.  | <0.2  | <0.2  | n.a.  | 11.70 | 10.52 |
| Sm              | 1.50            | n.a.  | n.a.  | n.a.  | <0.2  | <0.06 | n.a.  | 2.37  | 2.38  |
| Eu              | 0.32            | n.a.  | n.a.  | n.a.  | <0.1  | <0.03 | n.a.  | 0.67  | 0.48  |
| Gd              | 1.55            | n.a.  | n.a.  | n.a.  | <0.2  | <0.09 | n.a.  | 2.40  | 2.39  |
| Tb              | 0.24            | n.a.  | n.a.  | n.a.  | <0.1  | <0.04 | n.a.  | 0.37  | 0.35  |
| Dy              | 1.43            | n.a.  | n.a.  | n.a.  | <0.1  | <0.06 | n.a.  | 2.29  | 2.31  |
| Ho              | 0.29            | n.a.  | n.a.  | n.a.  | <0.1  | <0.04 | n.a.  | 0.48  | 0.48  |
| Er              | 0.84            | n.a.  | n.a.  | n.a.  | <0.2  | <0.04 | n.a.  | 1.44  | 1.36  |
| Tm              | 0.12            | n.a.  | n.a.  | n.a.  | <0.1  | <0.03 | n.a.  | 0.20  | 0.15  |
| Yb              | 0.77            | n.a.  | n.a.  | n.a.  | <0.2  | 0.04  | n.a.  | 1.36  | 1.20  |
| Lu              | 0.11            | n.a.  | n.a.  | n.a.  | <0.1  | <0.03 | n.a.  | 0.20  | 0.12  |
| Hf              | 1.21            | n.a.  | n.a.  | n.a.  | 2.06  | <0.5  | n.a.  | 1.55  | n.a.  |
| Zr              | 40              | n.a.  | n.a.  | n.a.  | 3.00  | 6     | n.a.  | 60    | 58    |
| Y               | 7.63            | n.a.  | n.a.  | n.a.  | <0.2  | 0.20  | n.a.  | 11.80 | 9.62  |
| Nb              | 7.63            | n.a.  | n.a.  | n.a.  | <0.2  | 0.20  | n.a.  | 11.80 | 9.62  |
| Ta              | <3              | n.a.  | n.a.  | n.a.  | <0.2  | <3    | n.a.  | <3    | n.a.  |
| Rb              | 5.50            | n.a.  | n.a.  | n.a.  | 0.29  | <0.8  | n.a.  | 29.60 | 25.16 |
| Th              | <2              | n.a.  | n.a.  | n.a.  | 1.27  | 2     | n.a.  | 3.40  | n.a.  |
| U               | 0.50            | n.a.  | n.a.  | n.a.  | <0.2  | <0.06 | n.a.  | 0.75  | n.a.  |
| Sc              | 14.2            | n.a.  | n.a.  | n.a.  | 3.30  | 4.1   | n.a.  | 23.8  | 21.9  |
the trend lines coincide with measured olivine and orthopyroxene compositions, and these units can be modeled as linear mixtures of olivine and orthopyroxene. They are thus interpreted as heteradcumulates with no (or only minor) trapped intercumulus melt (e.g. Barnes et al. 2016b).

**Rare earth element compositions**

Chondrite-normalized REE and primitive-mantle-normalized trace element patterns for the basal dunite samples are presented in Fig. 9. All cumulates and heteradcumulates (units #3–10) are very REE-poor, with La between 0.5–1 times chondritic and most HREE below detection limits. The border zone ortho- and mesocumulates (units #1–2) are LREE-enriched with La-values approximately 2–20 times chondritic. Negative Eu and Sr anomalies are likely related to Ca-mobility during serpentinization. Total REE-abundances increase with decreasing MgO, which together with the parallel REE-patterns suggests olivine control (dilution) of REE. The REE patterns are interpreted to mainly reflect the amount and composition of the trapped intercumulus melt.

**Chalcophile elements**

Concentrations of chalcophile elements are generally low in the basal dunite, except for Ni, which must be almost entirely hosted by olivine judging from the overall low concentrations of sulfur (Table 2). In the adcumulates and heteradcumulates (units #3–10), Cu, Pd, Pt, and S are mostly below detection limits. The border zone cumulates (units #1–2) contain on average 480 ppm S (highest 1300 ppm). Most Cu and about half of Pd and Pt analysis results are below detection limits; highest measured Cu is 170 ppm, Pd is 5.7 ppb, and Pt is 10.8 ppb.

**Mineral chemistry**

**Chromite**

Results of representative EPMA mineral analyses are listed in Table 3. All analyzed spinels from the basal dunite (39 samples, 119 grains, 453 analyses) are classified as chromites after the most abundant trivalent ion (Fig. 10), except for two samples that contain three grains classified as spinels. Chromites have compositions with (average in parentheses): Mg# 0.12–0.5 (0.32), Cr# 0.45–0.80 (0.63), Fe^3+/# 0.03–0.51 (0.12), NiO 0.03–0.22 wt% (0.1 wt%), and ZnO 0.08–1.43 wt% (0.4 wt%) [where Cr# = Cr/(Cr + Al) and Fe^3+/# = Fe^3+/(Cr + Al + Fe^2+)]. There is large overlap in compositions, but generally the ortho- and mesocumulates (units #1–2) contain more chromian and ferrian chromites, while the adcumulates (units #3–10) contain more aluminian chromites (Fig. 10). Chromite core compositions are igneous based on constraints of Barnes and Roeder (2001). Chromite compositions overlap with the compositional fields of both layered intrusions and komatiites (Fig. 10) (Barnes and Roeder 2001). No systematic compositional variations between euhedral and lobate–poikilitic chromite grains were found.

Chromites from ortho- and mesocumulates (units #1–2) exhibit a wider range of FeO compositions (5–21.5 wt%) compared to those from other units (average 5.8 wt%) (Fig. 10a and c). In back-scattered electron (BSE) imaging, the chromites do not exhibit magmatic
zoning, but thin chromian magnetite rims with Fe₂O₃ replacing Al₂O₃ are commonly found (Fig. 4b–d) especially in the orthocumulate samples. The orthocumulate chromites also have higher contents of e.g. Mn, V, Zn, and Ni, with abundances of these elements increasing with decreasing whole-rock MgO. Barnes (1998) interpreted similar compositional trends in komatiitic cumulates as post-cumulus equilibration between chromite and trapped intercumulus melt during slow cooling. Compositional variations in chromites along the X-X' cross-section (Fig. 2) are illustrated in Fig. S2.

Olivine

Fresh olivine in the Näränkävaara basal dunite was found in only 13 samples. Relatively abundant fresh olivine is found in some less-altered ortho- and mesocumulate samples (units #1–2), and also in the (olivine) orthopyroxenites (units #5 and 10). In the adcumulates and heteradcumulates olivine has only been found in two samples as small relict cores in otherwise serpentinized pseudomorphs (units #8–9).

The analyzed olivine compositions range between Fo₈₇–₉₈.₆, with two orthocumulate samples extending down to Fo₈₄.₅ (see Tables 1 and 3). Highest measured forsterite content of Fo₉₈.₆ is from an olivine mesocumulate boulder (unit #1); the second highest of Fo₈₈.₅ is found in the lowermost orthopyroxenite (unit #5). Analyzed olivines exhibit three different levels of Ni concentrations (Fig. 11): (1) high-Ni from 0.40–0.37 wt% NiO; (2) moderate-Ni from 0.30–0.28 wt% NiO; and (3) low-Ni from 0.28–0.21 wt% NiO. These levels mirror whole-rock Ni values and the geochemical subdivision presented earlier (Fig. 6e). Compared to a statistical analysis of komatiitic olivine compositions by Barnes and Fiorentini (2012), the Ni-richest olivines from the low-Fe zone (unit #1) can be classified as both “mineralized komatiites” and “barren units in mineralized domains”, while all other olivine analysis results fall in the field of “fresh olivine in unmineralized komatiites”.

Pyroxenes

Orthopyroxene was analyzed from nine samples and clinopyroxene from seven samples (Table 3). Orthopyroxene compositions range between En₈₃.₇–₈₇.₀, Fs₈₇.₇–₁₂.₂, and Wo₂₄–₄₄.₅; with Cr₂O₃ between 0.₅–₀.₇ wt% and NiO between 0.₀₅–₀.₁₁ wt%. Clinopyroxenes range between En₄₆.₇–₅₂.₁, Fs₅₈.₈–₇.₂, and Wo₁₉.₅–₄₄.₉; with Cr₂O₃ between 0.₆₅–₁.₃₀ wt% and NiO between 0.₀₃–₀.₆₆ wt%. Pyroxenes from the low-Fe zone contain uniformly higher NiO compared to those of the high-Fe zone (e.g. average of 0.₄₈ versus ₀.₃₅ wt% NiO in clinopyroxene).

Fig. 5 Whole-rock MgO versus FeO** compositions of Näränkävaara basal dunite samples; FeO** is FeO calculated from total FeO according to Barnes et al. (2007). Pure olivine compositions Fo₉₆–₉₉.₁ are plotted on the right, and model liquids in equilibrium with Fo₉₀ olivines at K_D values of modeled parental magma compositions that results for units #1 and #2 MgO (black bar on x-axis). Field labeled “Model PM” shows the range of modeled parental magma compositions that results for units #1 and #2 using a liquid composition in equilibrium with Fo₉₀ olivine at K_D values 0.₃₀–₀.₃₃; note that this field has been plotted in all other variation diagrams in this paper. Black dashed line connects compositions of olivine and liquid in equilibrium with the Näränkävaara marginal series gabbro-norite composition (Järvinen et al. 2020)
Fig. 6  a–e Whole-rock compositions of Näränkävaara basal dunite samples on variation diagrams; a Dashed line shows the main geochemical division of basal dunite samples into low-Fe and high-Fe zones. Field labeled “Model PM” shows the range of modeled basal dunite parental magma compositions. Note that the trend of the border zone ortho- and mesocumulate samples (units #1–2) coincides with the marginal series gabbronite of the layered series (labeled “MS”), suggesting these rocks are mixtures of cumulus olivine and melt approximately of the marginal series composition. Other labeled fields show measured mineral compositions (Ol, Opx, Cpx) and comparisons to whole-rock compositions from the komatiitic basalts of the Vetreny Belt (Puchtel et al. 1997) and the komatiitic Kellojärvi ultramafic cumulate complex (Papunen et al. 2009).
Two major research questions concern the Näränkävaara basal dunite. First is related to its age and relationship with the neighboring layered series. In the SE-block of the Näränkävaara intrusion, a marginal series gabbronorite separates the dunite from the layered series (Fig. 2). Unless the current structural configuration is fully the result of post-emplacement faulting, this means there has to be an age difference between the two. The basal dunite could either be (1) comagmatic with the Fennoscandian 2.5–2.4 Ga layered series magmatism, with a long enough hiatus in magmatism to allow for the cooling needed for the subsequent formation of the marginal series, or (2) represent an Archean komatiitic wall-rock to the Proterozoic layered series magmatism. Since isotope dating methods have thus far proven unsuccessful, here we aim to constrain the petrogenesis of the basal dunite by studying its parental magma composition and differentiation, and compare our findings to the neighboring layered series and to nearby komatiites. The second major research question is related to the orthomagmatic Ni-potential of the basal dunite as, in any event, it must have formed from a large volume of mantle derived primitive magma(s). Some observations related to the Ni-potential of the basal dunite are discussed at the end of this section.

**Parental magma**

One way to study a possible comagmatic relationship between the Näränkävaara basal dunite and the layered series is to compare their parental magma compositions. The Näränkävaara layered series parental magma has been inferred from a marginal series gabbronorite intersected in drill hole R1 (Fig. 2) (Järvinen et al. 2020). As the composition of the basal dunite parental magma cannot be directly inferred from a chilled margin or spinifex textured rock, instead the parental magma composition has been modeled here using results of whole-rock analyses from orthocumulates. The methods used here have been outlined in detail by e.g. Bickle (1982) and Chai and Naldrett (1992), and are based on the assumption that olivine orthocumulate rocks are a mixture of cumulus olivine and melt in equilibrium with this olivine. First, the Mg# of a melt in equilibrium with the most primitive olivine found in the basal dunite was calculated by using the olivine-melt Mg-Fe exchange coefficient (K_Mg) of Roeder and Emslie (1970). Then, the MgO and FeO contents of the parental magma were calculated from the intersection
Fig. 8  a–e Whole-rock trace element compositions and ratios of basal dunite samples. Field labeled “Model PM” shows the range of modeled basal dunite parental magma compositions; other labeled fields show comparisons to whole-rock compositions from the Näränkäväara layered series (Järvinen et al. 2020), komatiitic basalts of the Vetreny Belt (Puchtel et al. 1997), and the komatiitic Kellojärvi ultramafic cumulate complex (Papunen et al. 2009). Error bars show 1σ relative errors for element ratios, samples with 1σ errors larger than the ratio have been omitted.

Layered series

Star Marginal series (R1 173 m)

Basal dunite - High-Fe zone

- Unit 9 - Olivine-orthopyroxene heteradcumulate
- Unit 8 - Olivine adcumulate
- Unit 6 - Olivine adcumulate

Basal dunite - Low-Fe Zone

- Unit 5 - Orthopyroxenite
- Unit 4 - Olivine adcumulate, moderate-Ni
- Unit 2 - Olivine ortho- and mesocumulate, moderate-Ni
- Unit 1 - Olivine ortho- and mesocumulate, high-Ni
between this equilibrium-melt composition and a regression line fitted through samples composed of olivine-melt mixtures, as illustrated in Fig. 5. As the border zone ortho- and mesocumulates (units #1–2) are interpreted to represent olivine-melt mixtures, they were used for calculating these regression lines. Concentrations of other elements in the parental magma were then estimated in a similar fashion, by fitting regression lines through the ortho- and mesocumulate units and calculating the value of each element at the previously obtained MgO value. Samples taken from immediate contacts with basement or xenoliths were omitted from modeling to reduce scatter in whole-rock sample regression lines, as these near-contact samples show distinctly stronger LREE-enrichment indicative of in-situ contamination (not shown).

The most primitive olivines (Fo88–88.6) are found in the olivine ortho- and mesocumulates of the low-Fe zone (units #1–2). During post-cumulus processes, olivine compositions tend to progressively re-equilibrate towards lower Fo contents as modal intercumulus increases (e.g. unit #2 in Fig. 11) (Barnes 1986). It is thus unlikely that the olivines analyzed from the relatively intercumulus-rich ortho- and mesocumulates represent original compositions. It is also possible that the most primitive olivines have simply been destroyed during serpentinization. Therefore, average olivine compositions have also been inferred from whole-rock compositions. For the adcumulate units, olivine Fo contents can be directly inferred from whole-rock Mg#. The highest whole-rock Mg# of 90.4 is found in the lowermost olivine adcumulates of the low-Fe zone (unit #4 in Fig. 7a and Table 1). A slightly higher Fo content can be inferred from the MgO versus FeO plot in Fig. 5, where these samples plot close to pure Fo89 olivine. In addition, average Fo contents were inferred from molecular ratios of whole-rock Mg/Al and Fe²⁺/Al (see Makkonen et al. 2017), with results listed in Table 1. Using this latter method, results for the high-Ni subzone (unit #1) are between Fo89.90 and for the moderate-Ni subzone (unit #2) between Fo87–89, depending on the drill hole. Results of this method indicate similar olivine Fo contents as regression lines on variation diagrams intersecting pure model olivine compositions (e.g. Figure 5), with the added benefit of the method also being applicable to intercumulus-poor olivine adcumulate units. The most primitive olivine inferred from the basal dunit is thus in the range Fo89–91.

Parental magma compositions have been modeled separately for the high-Ni and moderate-Ni ortho- and mesocumulates (units #1–2) with a liquid composition in equilibrium with Fo89–91, using a K_D of 0.30 and 0.33 (Roeder and Emslie 1970; Toplis 2005). Resulting parental magma MgO varies between 13 and 18 wt% (average 15.6 wt%), depending on the unit used for regression and the parameters used to calculate the equilibrium liquid (Fig. 5 and Table 1). The high-Ni ortho- and mesocumulates (unit #1 in Table 1) always result in a higher parental magma MgO estimate compared to the moderate-Ni ortho- and mesocumulates (unit #2 in Table 1). This difference is related to the slopes of trend lines on the MgO-FeO diagram in Fig. 5, with unit #1 having slightly increasing FeO with decreasing MgO resulting in a higher MgO intercept. As the high-Ni subzone is Ni-richer and also appears to contain slightly more forsteritic olivine (Table 1), it is plausible that it has formed from a slightly more primitive magma compared to the moderate-Ni subzone. An “average” parental magma composition for these two units has been modeled using a Fo89 equilibrium liquid and K_D of both 0.30 and 0.33 (see Fig. 5) – the four resulting parental magma compositions are plotted as the corners of the “Model PM” field in variation diagrams in Figs. 4, 5, 6 and 8; with REE and trace element diagrams in Figs. 8 and 9. An average model parental magma composition is also listed in Table 2.

Overall, the modeled parental magma composition is that of an LREE-enriched high-MgO basalt, highly similar to the marginal series gabbronorite of the Näränkävaara layered...
### Table 3  Representative results of EPMA chemical analyses and calculated cation fractions

| Chromites | Low-Fe | Low-Fe | Low-Fe | High-Fe | High-Fe | High-Fe | High-Fe |
|-----------|--------|--------|--------|---------|---------|---------|---------|
| Unit #    | 1      | 2      | 4      | 6       | 7       | 8       | 9       |
| Rock type | Peridotite | Lherzolite | Dunite | Dunite | Dunite | Dunite | Dunite |
| Cumulus   | oMCb?  | oOCb*a*(pf) | oAC   | oAC   | oAC   | oCA    | oMC(b+) |
| Sample    | 1      | 2      | 3      | 4       | 5       | 6       | 7       |
| Major oxides (wt%) |        |        |        |         |         |         |         |
| SiO₂      | 0.06   | 0.34   | 0.06   | 0.00    | 0.05    | 0.10    | 0.10    |
| TiO₂      | 0.52   | 0.46   | 0.51   | 0.42    | 0.58    | 0.51    | 0.43    |
| V₂O₅      | 0.15   | 0.50   | 0.13   | 0.19    | 0.18    | 0.20    | 0.26    |
| Al₂O₃     | 16.34  | 10.82  | 17.68  | 14.24   | 19.63   | 22.26   | 22.68   |
| Cr₂O₃     | 45.01  | 38.84  | 46.82  | 50.09   | 44.49   | 41.55   | 42.99   |
| Fe₂O₃     | 30.76  | 42.48  | 26.34  | 28.36   | 27.49   | 26.57   | 23.80   |
| MnO       | 0.41   | 0.34   | 0.23   | 0.18    | 0.34    | 0.35    | 0.12    |
| MgO       | 5.26   | 4.04   | 7.50   | 5.36    | 6.53    | 7.51    | 9.23    |
| NiO       | 0.09   | 0.16   | 0.12   | 0.03    | 0.07    | 0.08    | 0.08    |
| ZnO       | 0.73   | 0.33   | 0.21   | 0.52    | 0.33    | 0.70    | 0.22    |
| Total     | 99.33  | 98.30  | 99.59  | 99.42   | 99.69   | 99.84   | 99.92   |
| Calculated cation fractions (apfu)² |        |        |        |         |         |         |         |
| Fe³+/Σtrivalent | 0 10  | 0 31  | 0 10  | 0 05    | 0 05    | 0 06    | 0 04    |
| Cr⁺/Σtrivalent | 0.59  | 0.49  | 0.58  | 0.66    | 0.57    | 0.52    | 0.54    |
| Al⁺/Σtrivalent | 0.32  | 0.20  | 0.32  | 0.28    | 0.38    | 0.42    | 0.42    |
| Mg#       | 0.26   | 0.19   | 0.34   | 0.27    | 0.32    | 0.36    | 0.43    |
| Cr#       | 0.65   | 0.71   | 0.64   | 0.70    | 0.60    | 0.56    | 0.56    |

| Olivines   | Low-Fe | Low-Fe | High-Fe | High-Fe |
|------------|--------|--------|---------|---------|
| Unit #     | 1      | 2      | 8       | 9       |
| Rockname   | Peridotite | Peridotite | Serpentinite | Dunite |
| Cumulus    | oOCba? | o(c)OCb*(a*) | oAC   | oMC(b*c*) |
| Sample     | 1      | 2      | 3       | 4       |
| Major oxides (wt%) |        |        |         |         |
| SiO₂       | 41.21  | 40.43  | 40.48   | 40.63   |
| Al₂O₃      | 0.009  | 0.013  | 0.022   | 0.007   |
| Cr₂O₃      | 0.015  | 0.019  | 0.025   | 0.021   |
| Fe₂O₃      | 11.85  | 11.87  | 11.87   | 11.95   |
| MnO        | 0.204  | 0.186  | 0.187   | 0.185   |
| MgO        | 46.74  | 47.39  | 47.37   | 47.27   |
| CaO        | 0.030  | 0.072  | 0.078   | 0.059   |
| NiO        | 0.405  | 0.309  | 0.278   | 0.245   |
| CoO        | 0.016  | 0.016  | 0.018   | 0.020   |
| Total      | 100.48 | 100.31 | 100.32  | 100.39  |
| Mg#        | 87.55  | 87.68  | 87.68   | 87.58   |

| Pyroxenes  | Low-Fe | Low-Fe | High-Fe | High-Fe |
|------------|--------|--------|---------|---------|
| Unit #     | 2      | 2      | 5       | 8       |
| Rockname   | Peridotite | Peridotite | Bronzitite | Dunite |
| Cumulus    | o(c)OCb*(a*) | o(c)OCb*(a*) | bAC   | oAC    |
| Sample     | 1      | 1      | 2       | 3       |
| Mineral    | Opx    | Cpx    | Opx     | Cpx     |
Table 3 (continued)

| Major oxides (wt%)          |   |   |   |   |
|-----------------------------|---|---|---|---|
| SiO₂                        | 55.97 | 51.59 | 55.80 | 52.53 |
| TiO₂                        | 0.08  | 0.71  | 0.07  | 0.37  |
| Al₂O₃                       | 1.52  | 3.03  | 1.08  | 3.91  |
| FeO             | 7.59  | 4.27  | 7.29  | 3.51  |
| MnO                        | 0.18  | 0.15  | 0.20  | 0.11  |
| MgO                        | 33.05 | 18.30 | 33.74 | 16.85 |
| CaO                        | 1.88  | 20.17 | 1.41  | 20.84 |
| Na₂O                       | 0.00  | 0.44  | 0.00  | 0.72  |
| Cr₂O₃                      | 0.69  | 1.05  | 0.60  | 1.03  |
| NiO                        | 0.09  | 0.04  | 0.07  | 0.03  |
| Total                      | 101.15 | 100.00 | 100.36 | 100.00 |

Calculated cation fractions (apfu)⁶

| Mg#        | 0.89   | 0.88   | 0.89   | 0.90   |
| En         | 85.31  | 51.07  | 86.67  | 48.43  |
| Fs         | 11.20  | 6.87   | 10.72  | 5.83   |
| Wo         | 3.50   | 40.46  | 2.60   | 43.05  |

FeO₅ = total FeO

* Cumulus nomenclature after Irvine (1982), with exceptions that intercumulus phases marked as suffix, minerals <5 vol% mode marked with parentheses, and poikilitic phases marked with asterisk

1 Samples (drill hole & depth): 1) R9 219.3 m; 2) R8 141.0 m; 3) R4 170.7 m; 4) VJJA-2017-57.1 (surface); 5) JSN-89 (surface); 6) VJJA-2018-81.1 (surface); 7) R2 237.9 m

2 Chromite analyses calculated to an ideal ulvospinel formula (4 oxygens) according to Barnes (1998); sum of trivalent cations = (Cr+Ti+V+Al+Fe³⁺); Mg# = Mg/(Mg+Fe²⁺); Cr# = Cr/(Cr+Mg)

3 Samples (drill hole & depth): 1) R6 89.8 m; 2) TTK-4.1 m; 3) VJJA-2017-81.1 m; 4) R2 237.9 m

4 Samples: 1) TTK-4.1 (surface); 2) VJJA-2017-55.2 (surface); 3) VJJA-2017-81.1 (surface)

5 Orthopyroxene = Opx; Clinopyroxene = Cpx

6 6 oxygens per formula unit, En = Mg/(Mg+Fe²⁺+Fe³⁺+Ca+Mn), Fs = (Fe²⁺+Fe³⁺+Mn)/(Mg+Fe²⁺+Fe³⁺+Ca+Mn), Wo = Ca/(Mg+Fe²⁺+Fe³⁺+Ca+Mn)

Fig. 10 a-c Chromite compositions in the Näränkävaara basal dunite, with grey circles showing previous results (Alapieti 1982; Telenvuo 2017). Labeled fields show comparisons to chromites in komatiites in green (Barnes et al. 1996) and to mafic-ultramafic layered intrusions in red (Wilson 1982; Chistyakov and Sharkov 2008; Karinen 2010; Bailly et al. 2011). Results of chromite analyses have been recalculated assuming stoichiometry to an ideal XY₃O₆ formula according to Barnes (1998); Mg# = Mg/(Mg+Fe²⁺); Cr# = Cr/(Cr+Mg); Fe³⁺# = Fe³⁺/(Fe³⁺+Mg+Cr)

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series (Figs. 8 and 9, and Table 2; marginal series plotted as a star in all variation diagrams) (Järvinen et al. 2020). The marginal series gabbronorite has a Mg# of 74 and thus is in equilibrium with Fo 90–90.6 olivine (KD 0.30–0.33), similar to the most primitive olivine inferred from the basal dunite (Fig. 5). These results suggest a comagmatic origin for the basal dunite and the Näränkävaara layered series.

Large uncertainty to the model results presented here is caused by scatter in whole-rock MgO and FeO contents (Fig. 5). Some of this may be related to iron mobility during serpentinization, as secondary magnetite veinlets are relatively common in drill core. Several other significant sources of error are also related to the methods used here, discussed in more detail in e.g. Amdt et al. (2008).

Inner structure and differentiation of the basal dunite

Layered structure

The same succession of cumulates and geochemical variations are found in both the SE and NW blocks of the basal dunite, in two separate cross-sections 25 km apart (Fig. 2a) (this study and Järvinen et al. 2020). In cross-section in both blocks, from SW to NE, ortho- and mesocumulates are followed by adcumulates and finally olivine orthopyroxene heteradcumulates before the layered series contact. Geochemically, both begin with a low-Fe zone followed by a high-Fe zone, with whole-rock and olivine compositions showing decrease in Mg# and Ni contents towards the layered series (Figs. 7a and c; see also Fig. S1). The sequence of high- and low-Cr cumulates (Fig. 7d), reflecting the amount of euhedral (cumulus) chromite, is also the same on both blocks. These observations indicate that the structure of the basal dunite is at least somewhat layered and remains similar over the 30 km strike length of the intrusion.

Ortho- and mesocumulates of the border zone

Olivine orthocumulates are found in both komatiite flows and mafic-ultramafic layered intrusions (Campbell 1987; Hill et al. 1995). Orthocumulate units of similar thickness as in the basal dunite (Fig. 2c) are common along the margins of komatiite flows, with modal olivine increasing inwards with distance from the wall-rock contact (Gole and Barnes 2020). In comparison, the Aganozero-block of the ~2.44 Ga Burakovsky layered intrusion in Russian Karelia (Fig. 1; Amelin et al. 1995) contains a 100–200 m thick olivine orthocumulatic marginal sequence interpreted as a marginal series for the intrusion (Nikolaev and Ariskin 2005), with similar modal and geochemical variation as found in the Näränkävaara ortho- and mesocumulates. As the low-Fe zone represents the most primitive part of the Näränkävaara basal dunite – and as the orthocumulates of the low-Fe zone can be modeled as mixtures between the most primitive olivine found and a trapped melt in equilibrium with this olivine – we suggest that the ortho- and mesocumulate border zone (units #1–2) represents a marginal series of sorts similar to the referred examples. Orthocumulate textures have likely developed by relatively rapid cooling of olivine-bearing magma against a cool wall-rock leading to early crystallization of intercumulus minerals (Campbell 1987).

Adcumulates and heteradcumulates

Over 90% of the ~1.5–2.5 km thick Näränkävaara basal dunite consists of olivine adcumulates (Fig. 2c). Olivine adcumulates are thought to be formed by in-situ crystallization on the crystal-liquid interface, with continuous low-porosity crystal growth facilitated either by diffusion combined with
convective extraction of solute in near-equilibrium low-heat flux conditions (e.g., slow-cooling magma chamber) (Campbell 1987; Walker et al. 1988) or by turbulent flow of low-viscosity magma flushing away the depleted solute from the crystal-melt interface (e.g., komatiite flow) (Hill et al. 1995).

The adcumulate units of the Näränkävaara basal dunite exhibit some features typically described from high-volume turbulent komatiite flows. In the low-Fe zone, adcumulates contain smooth sinusoidal fluctuations in Mg# with height (Fig. 7a, see unit #4 at 350–550 m). Similar fluctuations in komatiitic cumulates have been interpreted as evidence of crystallization in an open system with continuous influx of magmas with variable MgO content (Arndt et al. 2008). In addition, the high-Fe zone adcumulates commonly contain poikilitic chromite (Fig. 4c-d), and rarely also bimodal olivine (Fig. 3f). Both are thought to form in open system turbulent flows in komatiite settings (Barnes 1998; Gole and Barnes 2020). Poikilitic chromite in adcumulates cannot form from trapped intercumulus liquid alone, as the solubility of Cr in basaltic and komatiitic magmas is low (~3000 ppm). Rather, poikilitic chromite is thought to crystallize competitively with olivine in conditions of chromite supersaturation but inhibited nucleation (Godel et al. 2013). Such conditions may exist in freely convecting magma chambers or channels under conditions of low supercooling, where chromite supersaturation is reached only locally and transiently; e.g. in hot turbulent komatiite flows (Barnes 1998). In contrast, euhedral chromite is typically found in more static settings with low cooling rates, e.g. in differentiated flows (Barnes 1998) and in layered intrusions. In Archean komatiites, poikilitic chromite is typically found in adcumulates with Fo92–93 and euhedral chromite below Fo91, with the high Fo contents associated with poikilitic chromite indicating high crystallization temperatures close to chromite liquidus (Barnes 1998). Here, those Fo limits seem to have been shifted down about 3 mol% (Table 1). As far as we know, these are the lowest Fo rocks where poikilitic chromite has been reported. An additional complication is that, unlike in komatites, poikilitic chromite in Näränkävaara appears to be associated with the lowest Fo olivine adcumulates rather than the highest (Table 1), implying that some other factor than temperature has inhibited chromite nucleation. Also, although modeling indicates a relatively high liquidus temperature between 1300 and 1415 °C for the basal dunite parental magma (COMAGMAT 5.5.2 with oxygen fugacity buffered at quartz-fayalite-magnetite; Ariskin and Barmina 2004), that is close to or above chromite liquidus (~1350 °C; Murck and Campbell 1986), it is not clear whether such a dynamic turbulently flowing magma system as prescribed by Barnes (1998) could exist in an intrusive environment. In an intrusive setting, poikilitic chromite has only been described from the Dumont sill in Canada (Duke 1986). Resolving how the poikilitic chromite in Näränkävaara formed is outside the scope of this paper, but its ubiquitous presence in the high-Fe zone adcumulates in association with bimodal olivine suggests the possibility that these adcumulates were at least partly formed in a high flow-through environment.

Whether the adcumulates of the basal dunite were formed in a high-volume system or not, the composition of the in-situ formed olivine should reflect the composition and differentiation of the melt from which it formed. Generally, whole-rock Mg# and Ni (as well as olivine and chromite Ni) decrease from SW to NE towards the layered series contact (Fig. 7), suggesting differentiation by olivine fractionation from Fo90–91 down to Fo87–86. There are, however, at least two distinct breaks or reversals indicated by Mg#. The first break is marked by the lowermost orthopyroxenite unit that separates the low-Fe and high-Fe zones (unit #5 in Fig. 7a), and the second by a relatively low-Mg# adcumulate in the high-Fe zone (unit #7 in Fig. 7a). At least the latter is likely related to an influx of more primitive magma.

Difference in average compositions between the low-Fe and high-Fe zones is quite large in regard to MgO, FeO, and Ni (Fig. 6a and e). The olivine Fo-content at the beginning of orthopyroxene crystallization also differs between the two zones. Orthopyroxene first appears on the liquidus in the low-Fe zone at Fo89 (orthopyroxenite unit #5 in Fig. 7a), and a second time in the high-Fe zone at Fo87 (heteradcumulate unit #9). This indicates slightly different liquid lines of descent, and suggests a more significant change in parental magma composition between the low-Fe and high-Fe zones (fractionation in staging chamber?). The low-Fe zone may represent a single magmatic system, beginning with formation of the border zone orthocumulates (units #1–2), followed by possible open system flow and formation of adcumulates (units #3–4), and ending in closed system pooling, fractionation, and formation of the lowermost orthopyroxenite (unit #5).

Similar olivine orthopyroxene heteradcumulates as in unit #9 (Fig. 2c) are found in a similar position (preceding the layered series contact) in the NW block of the intrusion, 25 km to the NW (Järvinen et al. 2020). In the NW block, the heteradcumulates are in contact with, or grade into, the basal harzburgite of the Näränkävaara layered series. These NW contact zone heteradcumulates and layered series basal harzburgite compositions are plotted as grey crosses in variation diagrams in Figs. 4, 5 and 6. These compositions closely match those of units #9 and #10 in the SE block. At least for the high-Fe zone, this suggests the basal dunite and layered series magmas were following identical liquid lines of descent, having been locked-in to the same olivine-orthopyroxene-(chromite?) peritectic at an olivine composition of approximately Fo87.5.

**Archean or Proterozoic origin?**

The parental magma composition and liquid line of descent of the Näränkävaara basal dunite seem to be strikingly similar to that of the neighboring 2.44 Ga mafic-ultramafic layered series. However, the basal dunite exhibits many features typical of Archean komatiitic cumulates, except with distinctly lower-
Fo olivine as noted above. As the Näränkävaara intrusion is located between the Takenan and Suomussalmi greenstone belts (Fig. 2a), both containing Archean komatiites (Iljin 2003; Lehtonen et al. 2017), the hypothesis that the dunite body could represent an older komatiitic wall-rock for the layered series must also be considered. The nearby Suomussalmi and Takenan komatiites are relatively thin olivine+clinopyroxene cumulates and flows with average $\text{Al}_2\text{O}_3/\text{TiO}_2$ ratios of 17–19. Chondrite-normalized REE patterns are flat or LREE-depleted (Papunen et al. 2009; V. Järvinen unpublished data), except for the Vaara locality which shows roughly similar LREE-enriched patterns as the Näränkävaara basal dunite (Kunnunaho et al. 2013). Vaara is the only nearby komatiite body with similar high-Mg and low-Cr olivine adcumulates as found in Näränkävaara (see Fig. 4), but it is significantly smaller in volume. The uniformity of the inner structure of the Näränkävaara basal dunite along a strike length of 30 km would only be expected in proximal conduit or large scale channel facies komatiite flows (Gole and Barnes 2020). The 2.79 Ga Kellojärvi ultramafic cumulate complex, located about 100 km south of Näränkävaara in the Kuhmo greenstone belt of the Suomussalmi-Kuhmo-Tipasjärvi greenstone complex, is the closest comparison of komatiotic origin to the basal dunite (Papunen et al. 2009; Lehtonen et al. 2016). The Kellojärvi complex is 3–5 km thick and primarily composed of metamorphosed olivine ortho-, meso-, and adcumulates with similar petrography as in Näränkävaara, with the exception of clinopyroxene instead of orthopyroxene. The general lithology of the Kellojärvi complex is also similar, with a < 50 m thick border zone composed of olivine orthocumulate and minor pyroxenites (with basement xenoliths), grading into dunite towards the center of the body (Tulenheimo 1999; Papunen et al. 2009). Highest measured igneous olivine Fo contents (Fo$_{82-89}$) (Tulenheimo 1999) and whole-rock geochemistry are quite similar to Näränkävaara (Fig. 6). Kellojärvi parental magma composition has been inferred from whole-rock compositions to have 23 wt% MgO (Makkonen et al. 2017), and the highest Fo content inferred from whole-rock Mg#'s is about Fo$_{82}$, which are both somewhat higher compared to the estimates from Näränkävaara presented here. These differences, as well as the differences in trace element ratios (Figs. 8 and 9), could be explained by Näränkävaara having formed from a more strongly differentiated and crustally contaminated komatiitic magma.

One observation against a komatiitic origin is that compared to typical komatiitic olivine (Arndt et al. 2008) the olivines from the Näränkävaara basal dunite have distinctly lower Cr$_2$O$_3$ (<0.05 wt% versus 0.15–0.5 wt%) and CaO (<0.15 wt% versus 0.2–0.35 wt%). The low CaO content of olivines in Näränkävaara suggests a high-pressure intrusive environment (Simkin and Smith 1970). The higher degree of serpentinization in the basal dunite compared to the layered series is not thought to be significant evidence for either hypothesis because peridotitic rocks in the layered series are typically strongly serpentinized and fresh orthopyroxenites are in turn found in the basal dunite (unit #5). Overall, the highly similar parental magma compositions, liquid lines of descent, and roughly congruent layered structures between the Näränkävaara basal dunite and layered series argue for a common intrusive origin. There are no other 2.44 Ga layered intrusions in northern Finland with such an extensive basal dunite as the Näränkävaara intrusion, however. Closest correlations are the ~2.44 Ga Burakovsky layered intrusion (Chistyakov and Sharkov 2008) and the komatiitic basalts of the Vetreny belt (Puchtel et al. 1997), both found in Russian Karelia (Fig. 1). The two are considered comagmatic, with parental magmas between 13 and 18 wt% MgO, and similar trace and isotopic compositions (Puchtel et al. 1997; Nikolaev and Ariskin 2005; Kulikov et al. 2010). As shown in Figs. 4, 6, 8, and 9, geochemistry of the Näränkävaara basal dunite is extremely similar to the Vetreny belt extrusives. The Aganzero-block of the Burakovsky intrusion also hosts an up to 6 km thick basal dunite composed of homogeneous olivine adcumulate, with olivine compositions Fo$_{82-89}$ (Nikolaev and Khorov 2003), interpreted to have formed by repeated recharge of high-MgO magma (Chistyakov and Sharkov 2008). In addition, the marginal series orthocumulate in the Aganzero-block is similar to the Näränkävaara border zone, as discussed previously.

Kulikov et al. (2010) proposed that the igneous formations of the Vetreny belt, together with similar formations in Kola and northern Finland, are parts of at least two intermixed large igneous provinces (LIP) spanning 2.5–2.4 Ga, with related intrusive, hypabyssal, and extrusive rocks. The rocks share similar characteristics of a crustally contaminated komatiitic (plume-related) magma, with 9–18 wt% MgO, 0.5–1 wt% TiO$_2$, LREE-enrichment, and an average initial $\varepsilon_{Nd}$ of −2 (intrusions) to −0.4 (extrusives). Isotope data are not available from Näränkävaara, but otherwise the inferred parental magmas for the Näränkävaara layered series (Järvinen et al. 2020), and for the basal dunite presented here, are in line with the results presented for the Vetreny belt (Puchtel 1997; Kulikov et al. 2010).

An open question is whether the komatiitic features indicating low-viscosity turbulent flow can form in an intrusive setting from a high-Mg basaltic magma in equilibrium with −Fo$_{80}$ olivine? We suggest that the Näränkävaara basal dunite represents a (subvolcanic?) sill or chamber with intermittent high volume flow-through of high-MgO basaltic magma possibly evolving towards lower MgO content with time. Formation of a marginal series between the basal dunite and layered series indicates a longer hiatus in magmatism, with again at least three (relatively much smaller) magma pulses forming the Näränkävaara layered series (Järvinen et al. 2020).

Overall, available evidence suggest that the Näränkävaara intrusion formed in an open system dynamic magmatic environment (sill-dyke network?) with several discrete pulses of different magmas with different MgO and Ni contents, but
with a uniform origin. Näränkävaara may have acted as a magmatic feeder channel related to the Koillismaa layered intrusion complex and the “Hidden dyke” (Fig. 1) as suggested by Alapieti (1982), and more generally as a magmatic center related to the Fennoscandian LIPs as suggested by Kulikov et al. (2010). In terms of their composition, volcanic rocks found in the Saari-Kiekki greenstone belt in eastern Finland (Luukkonen 1989) could be viewed as differentiates of the Näränkävaara basal dunite and layered series parental magmas (Järvinen et al. 2020; this study). Although these volcanic rocks have not been precisely dated, they may represent extrusive equivalents of the Finnish 2.44 Ga layered intrusions. Nevertheless, extrusive rocks of this age group have not been definitively identified from Finland, so it is possible that either (1) the magmatic systems never vented, (2) they have been eroded away, or (3) the lavas all flowed into the Karelia-Kola rift now located in Russian Karelia. Although we prefer the origin of the Näränkävaara basal dunite in relation to the layered series in the light of the presented evidence and discussion, we emphasize that its petrogenesis is not settled in the absence of absolute age or other isotopic data, which have proven difficult to acquire from the serpentinitized ultramafic rocks.

Mineral potential for orthomagmatic Ni-(Cu-Co-PGE) deposits

The Näränkävaara basal dunite represents a large volume of ultramafic cumulates derived from a mantle plume source in an area with large trans-crustal structures, and its mineral potential for Ni-(Cu-Co-PGE) is of great interest (Barnes et al. 2016a). Similar Cr-poor olivine cumulates as found in Näränkävaara (Fig. 4) have been found prospective for komatiite-hosted Ni-Cu-PGE deposits in Finland (Konnunaho 2016).

There is a distinct difference in average whole-rock Ni contents between the low-Fe zone (2200 ppm Ni) and the high-Fe zone (1780 ppm Ni) (Fig. 6e). This Ni-depletion is reflected in olivine compositions (Fig. 11; 3200 ppm versus 2150 ppm Ni), chromite (1500 ppm versus 390 ppm Ni), and to a lesser extent also in pyroxenes. Because the Ni-depletion appears in samples with the same Mg# and forsterite contents, it cannot be related to simple fractionation of olivine, even if the trapped liquid shift to lower Fo contents is taken into account (Fig. 10) (Barnes 1986). Rather, this suggests either a more Ni-rich parental magma for the low-Fe zone compared to the high-Fe zone, or that the Ni-depletion is caused by sulfide saturation and segregation at some stratigraphic level.

No significant amounts of magmatic sulfides have been found in the Näränkävaara area. Trace sulfides are commonly found in the NW border zone olivine ortho- and mesocumulates. A basaltic magma becomes sulfur-saturated at approximately 1000 ppm S (Li and Ripley 2005). Assuming total incompatibility of sulfur in olivine, an olivine cumulate with 10 vol% trapped intercumulus melt (and 90 vol% olivine) would, at sulfur saturation, contain approximately 0.10 × 1000 = 100 ppm S. Of the border zone samples (units #1–2), 62 out of 69 are above the detection limit of 100 ppm S, containing between 100 and 1200 ppm S (average of 480 ppm). These samples contain about 75–95 vol% cumulus olivine, suggesting that the trapped intercumulus melt contained in the samples may have been sulfur saturated at the time of emplacement. However, Cu, Pd, and Pt do not correlate with increasing S contents, suggesting that these sulfides equilibrated with a small volume of magma and possibly formed after emplacement.

Pt, and to a lesser extent Pd, behave similarly to other incompatible elements with their concentrations increasing with decreasing cumulus olivine (Fig. 12). Only 8 samples, all from the moderate-Ni orthocumulate (unit #2), are above detection limit for Cu and Pd, with Cu/Pd ratios between 8000 and 40,000 (Fig. 12d). These are above primitive mantle ratio or about 8000 (McDonough and Sun 1995) which suggests sulfur saturation before emplacement. Ratios of Pd/TiO2 and Pt/TiO2 in the basal dunite are generally below primitive mantle ratio (Fig. 12c), also suggesting either previous sulfide saturation or sulfide retention in the source. Few samples from the high-Ni border zone orthocumulates (unit #1) show ratios above primitive mantle, which together with the generally more primitive nature of this unit suggests that it may have formed from a less PGE-depleted magma compared to unit #2. However, as most assays are below or close to detection limits with high relative errors, more precise analyses are required to assess the sulfide saturation history of the basal dunite in more detail.

Conclusions

The Näränkävaara basal dunite is a 1.5–2 km thick series of ultramafic cumulates stratigraphically below the 2.44 Ga Näränkävaara layered series. It contains a 200–300 m thick border zone along the southern basement contact composed of olivine meso- and orthocumulates, but is otherwise composed of olivine adcumulates with less poikilitic olivine-orthopyroxene heteradcumulates and minor orthopyroxenites. Textural and geochemical layering in the basal dunite remains similar for the 30 km strike length of the body. Geochemically, the dunite is composed of a more primitive low-Fe zone along the basement contact (average FeO of 10.2 wt% and Ni of 2250 ppm with Mg# 85.3–90.4), and a more evolved high-Fe zone in contact with the layered series (average FeO of 12.5 wt% and Ni of 1700 ppm with Mg# 85.2–87.7). A distinct Ni-depletion in olivine is found in the low-Fe zone to the high-Fe zone (3200 versus 2200 ppm Ni). This depletion does not correlate with olivine Fo contents, suggesting it is not related to olivine fractionation.
Hence the basal dunite may have potential for Ni-(Cu-Co-PGE) sulfide mineralization.

The basal dunite parental magma composition has been modeled based on olivine–melt mixing lines inferred from the border zone ortho- and mesocumulates. It is an LREE-enriched high-MgO (13–18 wt.%) basalt that exhibits highly similar major and trace element composition with the parental magma of the neighboring layered series.

The presence of abundant low-porosity adcumulates, sometimes with poikilitic chromite and bimodal olivine, together with pyroxenitic interlayers and at least two compositional reversals, suggests formation in a dynamic magmatic setting with high flow-through of picritic magma, interrupted by periods of pooling and differentiation. Differences in compositions of the low-Fe and high-Fe zones imply a change in parental magma composition with time.

The presence of a marginal series between the basal dunite and the layered series indicates that the basal dunite is older than the layered series. The basal dunite is suggested to have formed in an early magmatic feeder channel related to the same Fennoscandian plume-related magmatism that produced the 2436 Ma Näränkävaara layered series and the other layered intrusions of the terrane. However, an alternative origin as an Archean komatiitic wall-rock cannot be ruled out.

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