

Serpentinites of Different Tectonic Origin in an Exhumed Subduction Complex (New Caledonia, SW Pacific)

Natalie H. Raia¹, Donna L. Whitney¹, Christian Teyssier¹, Stéphane Lesimple²

¹Department of Earth & Environmental Sciences, University of Minnesota, Minneapolis, MN 55455, USA.

²Service Géologique de la Nouvelle-Calédonie, Direction de l'Industrie, des Mines et de l'Energie, BP M2 – 98849 Nouméa CEDEX, New Caledonia.

Corresponding author: Natalie Raia (raia0003@umn.edu)

Key Points:

- Two compositionally-distinct types of serpentinites identified in the New Caledonia subduction complex
- Some serpentinites derive from the subducting plate; others from the overlying mantle wedge
- New Caledonia preserves evidence for incorporation and exhumation of mantle hangingwall via the subduction channel

Abstract

Owing to the importance of serpentinites for planetary geochemical and geodynamic processes, there has been much work discerning the origins of their parent rocks, including distinguishing between serpentinites derived from a subducting plate vs. overlying mantle in exhumed subduction complexes. The island of New Caledonia (SW Pacific Ocean) provides a rare window into Cenozoic Pacific subduction processes. The island is unique in exposing both an exceptionally-preserved high-pressure, low-temperature subduction complex and one of the largest supra-subduction zone ophiolites in the world. Previous studies disagree on the origin of serpentinites in the subduction complex. In this study, we analyze twenty-three serpentinites from this subduction complex for whole-rock major and trace element geochemistry and stable isotope (δD , $\delta^{18}O$) compositions. Our data reveal two distinct groups of serpentinites: Group I samples in the northern portion of the complex are pervasively serpentinized, and exhibit enriched heavy rare earth element (REE) compositions and $\delta^{18}O$ between +6.7‰ and 10.2‰. In contrast, Group II serpentinites in the south preserve relict orthopyroxene and olivine, and show depleted trace element compositions and comparatively lower $\delta^{18}O$ values between +5.1‰ and +8.0‰. We interpret Group I serpentinites to derive from downgoing plate mantle, whereas Group II serpentinites derive from overlying mantle wedge, exhibiting remarkable similarity to the REE geochemistry of the structurally-overlying New Caledonia ophiolite. Our results establish the subduction complex in New Caledonia as an unusual natural record of the entrainment and exhumation of mantle from both the overlying mantle wedge and the downgoing plate in an oceanic subduction zone.

Plain Language Summary

The hydration of Earth's mantle produces rocks called serpentinites that are important to chemical cycling within the Earth system. This process, a form of metamorphism, occurs in several types of tectonic settings on Earth. Serpentinites formed in these different settings are imparted with unique geochemical “fingerprints” due to different types of fluids and varied compositions of the original mantle material. In metamorphic complexes that preserve remnants of subduction zones, serpentinites can offer clues to the original tectonic setting and subsequent mechanics of subduction. We analyze stable isotope and whole-rock compositions to determine the origin of the mantle that produced serpentinites found in the metamorphic complex preserved

in New Caledonia (SW Pacific). We discover two distinct groups of serpentinites: one group in the northern portion of the complex likely formed on the ocean floor prior to subduction, and experienced high degrees of fluid alteration as it was metamorphosed. In contrast, a second group of serpentinites in the southern portion of the complex resembles material from the mantle overlying the subducting plate and is less altered. This locality represents an uncommon example, globally, where material from this overlying mantle is entrained and preserved at the surface in an exhumed subduction complex.

1 Introduction

Serpentinites are hydrated fragments of Earth's mantle and commonly occur in exhumed subduction complexes. They are important vehicles in the global cycling of water, carbon, nitrogen, fluid-mobile elements (FME), and halogens, releasing these elements by prograde dehydration at forearc to subarc depths during subduction (e.g., Alt et al., 2013; Barnes et al., 2018; Collins et al., 2015; Deschamps et al., 2011; Halama et al., 2014; Hattori & Guillot, 2003; John et al., 2011; Kerrick & Connolly, 1998; Kodolanyi et al., 2012; Scambelluri et al. 2004, 2019; Tenthorey & Hermann, 2004; van Keken et al., 2011). Serpentinites and their hybridized, metasomatized derivatives formed at depth in subduction zones are critical in producing the distinct chemical composition of arc magmas (e.g., Codillo et al. 2018; Hattori & Guillot, 2003; Marschall & Schumacher, 2012; Nielsen & Marschall, 2017; Shimoda & Kogiso, 2019; Tatsumi, 1986) and contribute significantly to the net redox budget of subducted oceanic plates and the oxidation state of dehydration fluids infiltrating the overlying mantle (Debret & Sverjensky, 2017; Evans, 2012; Evans & Frost, 2021; Evans et al., 2017). The wide stability field of the serpentine mineral antigorite also makes possible the transport of these chemical components deeper into the mantle, past subarc depths (e.g., Hacker, 2008; Hattori & Guillot, 2003; Kendrick et al., 2011, 2017; Scambelluri & Tonarini, 2012; Schmidt & Poli, 1998; Ulmer & Tromsdorff, 1995; Wunder & Schreyer, 1997).

Serpentinites are also critical to the rheology of the subducting zone, and in particular the plate interface (Gerya et al., 2002; Hermann et al., 2000; Rüpke et al., 2004; van Keken et al., 2011). Their high water contents and low density facilitate the exhumation of denser blueschist and eclogite (Guillot et al., 2000; Magott et al., 2020; Schwartz et al., 2001), and serpentinization of the cold “nose” of the mantle wedge carries implications for mantle wedge flow and the

82 minimum depth of decoupling of the subducting plate and overlying mantle (Hilaitet & Reynard,
83 2009; Kerswell et al., 2021; Reynard, 2013; Wada et al., 2008). Furthermore, serpentinites have
84 been correlated with zones of slow slip and intermediate-depth seismicity in subduction zones,
85 possibly associated with dehydration embrittlement resulting from the breakdown of antigorite
86 (Behr & Bürgmann, 2021; Ferrand, 2019; Hacker et al., 2003; Hilaitet et al., 2006, 2007; Hirth &
87 Guillot, 2013; Jung & Green, 2004; Peacock, 2001; Proctor & Hirth, 2015; Toffol et al., 2022).
88 In short, the occurrence, composition, and spatial distribution of serpentinites and associated
89 metasomatic rocks in paleo-subduction complexes reveals information about first-order
90 subduction processes, including the transport and chemical evolution of fluids during subduction
91 and exhumation.

92 In exhumed subduction complexes, mantle material may originate from the downgoing
93 plate or the overriding plate (Figure 1). Prior to subduction, serpentinites form in the (eventual)
94 downgoing plate via hydrothermal circulation at ridge-transform systems where lithospheric
95 mantle underlying the igneous oceanic crust or exposed at/near the seafloor by extension is
96 hydrated by heated seawater (Barnes & O'Neil, 1969; Bonatti, 1976; Cannat, 1993; Cannat et al.,
97 2010; Mével, 2003; Rouméjon et al., 2015). Hydration may also occur as the plate approaches
98 the trench, if water circulates through slab-bend faults (Ranero et al., 2003). In the overriding
99 plate, the mantle wedge experiences hydration from fluid fluxing off the dehydrating downgoing
100 plate (e.g., Bostock et al., 2002; Fyfe & McBirney, 1975; Hyndman & Peacock, 2003).

101 Serpentinites formed in each of these distinct tectonic settings are geochemically distinct
102 owing to differences in (1) mantle protolith composition (i.e., melt depletion and refertilization
103 histories and degrees of fractional crystallization) and (2) the chemistry and nature of fluid
104 alteration, such as fluid-rock ratio and temperature of serpentinization (Figure 1) (e.g.,
105 Deschamps et al., 2010, 2013; Peters et al., 2017). These distinctions are evident in bulk-rock
106 major and trace element geochemistry, stable isotopes (e.g., O, H, Cl, B), and the composition of
107 relict primary minerals (e.g., spinel, pyroxene, and/or olivine) and have been proposed to be
108 diagnostic of the tectonic setting of serpentinization. This has unlocked opportunities to discern
109 the tectonic setting of mantle material exhumed in subduction complexes (Deschamps et al.,
110 2010, 2013; Peters et al., 2017).

111 Studies aiming to determine the tectonic source of formerly subducted serpentinites must
112 grapple with a nuanced, multi-staged history of serpentinization associated with pre-subduction
113 processes and the prograde and retrograde paths of subduction and exhumation (Figure 1).
114 Quantitatively determining the pressure (P), temperature (T), and time (t) conditions experienced
115 by exhumed serpentinites is challenging owing to high-variance phase assemblages, though
116 promising new approaches expand options for unlocking both temperature and time in these
117 rocks (Cooperdock & Stockli, 2016; Schwartz et al., 2020).

118 Complexities notwithstanding, there is a growing body of literature with examples of
119 provenance interpretations of serpentinites in exhumed subduction complexes, such as studies
120 assigning serpentinite associated with blueschist and eclogite to the downgoing plate (e.g.,
121 Cooperdock et al., 2018; Katzir et al., 2000; Li et al., 2004; Scambelluri et al., 1991; Shen et al.,
122 2015). Some studies have further resolved detail within the oceanic realm of the protolith,
123 interpreting exhumed serpentinites to derive from passive margin settings (e.g., Barnes et al.,
124 2014) or from abyssal transform faults (e.g., Cárdenas-Párraga et al., 2017). In many cases,
125 however, non-distinctive geochemical signatures, lack of relict phases, and heterogeneity of the
126 analyzed samples makes this level of resolution untenable. In other regions, the capturing of
127 hangingwall mantle via subduction erosion and other mass transfer mechanisms has been
128 identified (e.g., Bhat et al., 2019; Guice et al., 2021; Hattori et al., 2010; Lazar et al., 2021; Li et
129 al., 2018; Tewksbury-Christle et al., 2021; Wu et al., 2018). In yet other localities, serpentinites
130 from both the downgoing slab and the overlying mantle wedge are interpreted to be present,
131 implying complex slab-mantle interactions at a range of depths (e.g., Barnes et al., 2013; Blanco-
132 Quintero et al., 2011). These results generate discussion of deep tectonic slicing of slabs at depth,
133 degrees of mechanical mixing and styles of deformation (e.g., coherent nappe stacking, block-
134 and-matrix shear zones), and the processes of underplating and interaction with the mantle
135 material of the overriding plate.

136 In this contribution, we discern the tectonic origin and alteration histories of serpentinites
137 entrained in an Eocene high-pressure / low-temperature (HP/LT) subduction complex on the
138 island of New Caledonia (SW Pacific) and explore implications for the subduction and
139 exhumation history of the complex. The island of New Caledonia exposes an extraordinarily
140 complete Eocene subduction-obduction complex, including a high-temperature metamorphic
141 sole representing incipient stages of subduction initiation, an HP/LT subduction complex, and an

obducted supra-subduction zone ophiolite sequence (Maurizot et al., 2020a). These entities are well-studied and world-class sites: the ophiolite comprises one of the world's largest continuous mantle exposures and has provided invaluable insights into upper mantle processes and melt transfer in the lower crust (e.g., Cluzel et al., 2016; Dupuy et al., 1981; Marchesi et al., 2009; Pirard et al., 2013; Secchiari et al., 2016, 2018; Ulrich et al., 2010). The subduction complex exposes slivers of oceanic and thinned continental lithosphere that experienced blueschist and eclogite-facies metamorphism and has served as a location for investigation of subduction zone fluid-rock interaction, volatile recycling, and slab-mantle interactions (e.g., Cluzel, 2021; Spandler et al., 2008; Taetz et al., 2016, 2018). Despite the geodynamic importance of mantle rocks throughout the subduction-obduction complex, studies addressing the distribution, structure, texture, or geochemistry of meta-ultramafic rocks in the HP/LT complex are few (Cluzel, 2021; Fitzherbert et al., 2004; Rawling & Lister, 2002; Spandler et al., 2008). These studies arrive at differing conclusions regarding the source of the ultramafic material entrained in the complex. To evaluate these differences and determine the tectonic provenance and petrogenesis of ultramafic rocks in this subduction complex, our approach integrates field observations, petrographic characterization, whole rock major and trace element geochemistry, and stable isotope (O and H) geochemistry. Our results reveal the existence of two distinct groups of serpentinites, which we interpret to indicate differing tectonic settings for their protoliths.

2 Geologic History of New Caledonia

Located in the SW Pacific Ocean, the main island of New Caledonia exposes a dense array of geologic terranes that record Paleozoic and Mesozoic histories through to present day: a material archive that stands in contrast to many other islands in the Pacific, which are mostly entirely Cenozoic in age and volcanic in origin (Cluzel et al., 2012; Maurizot et al., 2020a; Paris, 1981). The island (~16,000 km²) is a rare emergent portion of the NW-SE trending submarine Norfolk Ridge, a continental sliver located ~200-400 km southwest of active subduction at the New Hebrides trench and comprising part of the largely submerged Zealandia microcontinent (Figure 2a) (Crawford et al., 2003; Dubois et al., 1974; Lafoy et al., 2005; Mortimer et al., 2017).

New Caledonia exposes Late Carboniferous to Early Cretaceous basement rocks deriving from the last accretionary stages of the active southern Gondwana margin (Figure 2b,

‘undifferentiated “basement” terranes’) (Aitchison & Meffre, 1992; Campbell et al., 1985; Maurizot et al., 2020b); unless otherwise noted, we adopt the naming conventions of Maurizot et al. (2020c) for New Caledonian rock units. In the Late Cretaceous, regional tectonic stresses shifted, prompting divergence and rifting associated with the breakup of the Gondwanan supercontinent. In the SW Pacific, the Tasman Sea and additional small ocean basins opened, rifting ribbons of continental crust away from the eastern margin of Gondwana (Figure 2a) (e.g., the submarine Dampier Ridge and Lord Howe Rise, for instance; e.g., Bache et al., 2014; Davies & Smith, 1971; Mortimer et al., 2018).

Plate convergence in the SW Pacific began in the Late Paleocene, with an intra-oceanic NE-dipping subduction zone initiating at ~56 Ma, as recorded by the recrystallization age of high-temperature amphibolite in the basal sole of the ophiolite, and further corroborated by boninite and adakite series dikes that range from 55-50 Ma (Cluzel et al., 2006, 2012). Two distinct packages of differing protolith types and P-T-t paths are recognized to have been subducted and exhumed. The Pouébo Terrane (Figure 2c, d), which subducted first, consists of fragments of oceanic lithosphere, and was followed by the Diahot-Panié Complex (Figure 2c, d), which consists dominantly of metasediments and metavolcanic rocks (Cluzel et al., 1994; Clarke et al., 1997; Maurizot et al., 2020b). The Pouébo Terrane reached peak eclogite facies metamorphic conditions of ~2.2-2.4 GPa and ~550-600°C, equating to ~70-80 km burial depth, by ~44 Myr (Pirard & Spandler, 2017; Spandler et al., 2005; Vitale-Brovarone et al., 2013). The Diahot-Panié Complex ranges in grade from lawsonite-blueschist facies (~0.5 GPa, ~250°C) to eclogite facies (~1.8-2.2 GPa, ~500-550°C) and likely reached peak conditions at ~38 Ma (Cluzel et al., 2010; Pirard and Spandler, 2017; Potel et al., 2006; Vitale-Brovarone et al., 2018). Exhumation is interpreted to have occurred in two stages: the first stage brought the Pouébo Terrane rocks up to ~40-50 km, where they were juxtaposed with the Diahot-Panié Complex. In the second stage, the two terranes exhumed together between 38-36 Myr, with the last gasps of rapid exhumation taking place at ~34 Ma (Baldwin et al., 2007; Vitale-Brovarone et al., 2018). The arrival and partial subduction of the buoyant continental-affinity Diahot-Panié Complex effectively halted subduction and triggered the obduction of a large fragment of overlying fore-arc mantle and oceanic crust from the modern-day Loyalty Basin onto Grande Terre (referred to as the Peridotite Nappe and Poya Terrane, respectively; Brothers, 1974; Coleman, 1967).

2.1 Ultramafic rocks in New Caledonia

Ultramafic rocks occur in several discrete tectonic terranes on the island. We briefly summarize the state of knowledge with respect to the provenance of these mantle rocks, organized into tectonic groups.

2.1.1. The Peridotite Nappe: Massif du Sud and isolated ophiolitic klippen

The southern end of New Caledonia is dominated by the Massif du Sud, one of the largest coherent exposures of mantle in the world (Figure 2b). This body is notably free of significant tectonic overprint and remains attached to non-obducted lithospheric mantle in some areas (Collot et al., 1987; Patriat et al., 2018; Prinzhofer & Nicolas, 1980). Additional remnants of the large overthrust ophiolite dot the west coast of the island as small klippen. Most of the mantle portion of the Nappe consists of harzburgite and dunite, with lherzolite being found in the north (Belep, Poum, and Tiébaghi massifs; Moutte, 1982; Prinzhofer, 1981; Sécher, 1981; Ulrich et al., 2010). The transition to the crustal sequence (pyroxenite, wehrlite, and gabbro) is preserved on the southern portion of the island; the ophiolite lacks an uppermost dike complex and pillow basalts (Maurizot et al., 2020c). The ophiolite is highly geochemically depleted (Marchesi et al., 2009; Prinzhofer & Allègre, 1985), implying high degrees of melt production, and this has made this body, and in particular, the continuous exposures in the Massif du Sud, a globally-important site for studying melt transfer within the mantle and into the lower crust (Marchesi et al., 2009; Pirard et al., 2013; Pirard & Hermann, 2015; Ulrich et al., 2010).

2.1.2. “Basement” ultramafic bodies of Gondwanan affinity

Serpentinities are reported in several Gondwanan basement terranes on the island. Portions of these units experienced local low-grade (greenschist and/or lawsonite blueschist facies) overprinting by Eocene HP/LT metamorphism. In some cases, serpentinites occur amongst swaths of schist and contain tectonically-entrained oceanic crustal components (e.g., metamorphosed pyroxenite, gabbro, basalt, chert). This ultramafic material is proposed to derive from an incoming oceanic plate that reached maximum depths of ~30-35 km beneath the overriding Gondwanan plate (‘Boghen Terrane’; Black, 1993; Cluzel & Meffre, 2002; Guérangé et al., 1977; Maurizot et al., 2020b). In other localities within the basement, serpentinites appear at faulted boundaries between coherent volcanic and abyssal sedimentary units (e.g., chert,

siltstones) (Maurizot et al., 2020b; Meffre et al., 1996). Published geochemical data on these rocks are lacking and a protolith origin for these serpentinites is not known.

2.1.3. Foreland accretionary units

Ultramafic rocks are reported to cross-cut the Montagnes Blanches Nappe, a remnant foreland fold-thrust belt preserved immediately west of the HP/LT complex (Koumac Terrane of Cluzel et al., 1994). Early interpretations link these rocks to the underlying oceanic lithosphere of the folded abyssal Montagnes Blanche Nappe sedimentary units (Brothers, 1974; Maurizot et al., 1989). These rocks have alternatively been interpreted as overthrust remnants of the Peridotite Nappe (Cluzel et al., 1995; Gautier et al., 2016; Maurizot, 2011), though a recent systematic field description of serpentinite occurrences in these units by Cluzel (2021) has pointed out inconsistencies with this hypothesis, instead positing that they are part of the upper plate. We emphasize that these hypotheses are based on field context, and published geochemical studies of the serpentinites are lacking.

2.1.4. Ultramafic and hybrid “blackwall” rocks of the HP/LT complex

Serpentinites occur throughout the HP/LT metamorphic belt in the NE portion of the island (Figure 2b-d). The exposed subduction complex is sprawling, spanning ~ 200 km long and ~20 km wide. The far northern area of the complex is most often referred to as the Pam Peninsula and exposes some of the best-preserved blueschists and eclogites in the complex (Figure 2c). Though observations and descriptions of serpentinite outcrops exist (e.g., Black & Brothers, 1977; Brothers & Blake, 1973; Cluzel, 2021; Lillie, 1975; Maurizot et al., 1989), geochemical and textural studies of these ultramafic rocks are sparse (Fitzherbert et al., 2004; Spandler et al., 2008), and the tectonic origin of this ultramafic material is debated. Spandler et al. (2008) interpret a seafloor origin based on major and trace element geochemistry, stable isotope measurements, and calculated temperatures of serpentinitization for four serpentinite samples in the far NE portion of the complex (Figure 2c). In contrast, Fitzherbert et al. (2004) note the similarity of P-T paths and the existence of serpentinites across mapped terrane boundaries (the Diahot Terrane and Pouébo Terrane of Cluzel et al. (1995) and Fitzherbert et al. (2003)), interpreting this distribution as inconsistent with a downgoing slab origin, and instead indicating interaction with the overlying mantle. This idea can be traced to earlier interpretations that the ultramafic rocks exposed in the metamorphic complex today were part of a larger, now

extensively eroded overthrust “serpentinite sheet” that was incorporated from the hangingwall, based on their consistent existence at the structurally highest levels of the complex (Rawling & Lister, 2002). At a small, weathered massif outcropping in the SE portion of the complex, near the town of Yambé (Figure 2d), Fitzherbert et al. (2004) interpret the ultramafic rocks to derive from hangingwall mantle that was incorporated with the downgoing plate, subducting with it to P-T conditions past the stability of antigorite. Critically, a geochemical link between the serpentinites in the HP/LT complex and the interpretation of an overriding plate or supra-subduction mantle origin is lacking.

3 Sample Localities and Description

Twenty-three serpentinites, one chlorite schist, and one talc schist were collected in the HP/LT complex for the purposes of discerning their tectonic origin and fluid histories (Table S1; Figure 2c, d). For comparison, four serpentinites were collected from outside the HP/LT complex: two from the Boghen Terrane basement unit, one from the serpentinite sole at the base of the Peridotite Nappe, and one from Kalaa-Gomen, a small klippe of the ophiolite on the northwest coast (Figure 2b). Representative thin section photomicrographs are presented in Figures S1-S4.

3.1 Field Context

Serpentinites in the northernmost portion of the HP/LT complex - defined here as an area encompassed by the Pam Peninsula and extending south along the eastern coastal road (RPN7) to the town of Balade - outcrop both as highly sheared and deformed lenses and as weathered boulders on hillslopes (Figure 2c). In highly sheared areas, the serpentinites act as a matrix for rounded pods of metamorphosed mafic, sedimentary, and other ultramafic rocks. With the exception of a small quarry south of Col d’Amos (NC19-14) and a locality in the foothills north of Ouégoa (NC18-26), all samples group spatially within the “Pouébo-Tiarì” unit of Vitale-Brovarone et al. (2018). These samples are entirely encompassed within the “mélange ophiolitique glaucophanite” unit of Maurizot et al. (1989) and fall variably within the “Pouébo metabasite dominates” and “Diahot metabasite dominates” units of Fitzherbert et al. (2004).

Nine samples in the HP/LT complex are located SE of the northern domain samples. A small (~ 0.5 km²) ultramafic body crops out within blueschist and eclogite facies metamafic

rocks near the town of Yambé (Figure 2d). Prior studies reference this ultramafic body as “Yambé” or “Yambé massif” (Fitzherbert et al., 2004; Spandler et al., 2008) and “Pwa Radèn” (e.g., Cluzel, 2021). With permission from Kanak tribal leadership, we adopt the local name given to this ultramafic body, “Poadja.” In some areas of the Poadja Massif, relict magmatic foliation is discernable at outcrop scale. Further south along the coast, two minor serpentinite outcrops (within 1 km of each other) are exposed in shear zones in beach outcrops, juxtaposed with meta-mafic blocks.

3.2 Petrographic Context

Serpentinites in the NE region are characterized by a dominant mineral assemblage of antigorite + talc + magnetite (Table S1; Figures 3a, b and S1) and are > 90% serpentinized, with rare relict chromian spinel (NC19-54), olivine (NC18-26B), and clinopyroxene (NC19-54). Antigorite most commonly occurs as the dominant matrix phase, forming interlocking crystals with variably preserved evidence for mesh texture. Some samples exhibit anhedral fine-grained (<10 μm), recrystallized antigorite intergrown with talc and less commonly, tremolite (e.g., NC18-15C, NC18-15D, NC18-22A). In some cases, well-developed bastite pseudomorph textures are evidenced by clumped aggregates of coarser-grained (~100-200 μm) euhedral antigorite crystals accompanied by euhedral magnetite (~10's of μm) (e.g., NC19-86). Magnetite additionally occurs as a matrix phase as single subhedral grains (~10's of μm up to ~1 mm) or as polymineralic aggregates of smaller euhedral grains (~10's of μm). In deformed samples these aggregates lie within the foliation (e.g., NC19-54, NC18-22A). In rare cases, relict chromite is preserved in the core of grains mantled by ferritchromite and rimmed by magnetite (Figure 3a). Two types of veins assemblages are present as cross-cutting networks: antigorite-magnetite (NC19-86; Figures S1e, j) and antigorite-tremolite (NC19-94; Figure 3b).

Serpentinites in the SE region are characterized by the presence of relict phases (orthopyroxene and olivine) and range between ~50 to ~90% serpentinized (Table S1; Figures 3c-f and S2). Samples from this locality notably lack talc (with the exception of NC19-158), and four samples contain the Fe-Ni alloy awaruite as an accessory phase (Figure 3d). Oxide textures show varying degrees of retrogression from skeletal chromite with magnetite rims (Figure 3c) to chromian magnetite cores complexly replaced by magnetite (Figure 3e). Texturally-late brucite

veins are observed in one sample (NC18-39A). Mesh textures are less developed compared to the complete pseudomorphing observed in the NE (Figure 3f).

4 Methods

4.1 Whole-rock major and trace element geochemistry (XRF, ICP-MS)

Twenty-three serpentinites, one talc schist, and one chlorite schist were analyzed for whole-rock major and trace element compositions. Major element analyses were acquired by lithium tetraborate fusion X-ray fluorescence methods at the GeoAnalytical Laboratory (Washington State University, USA; WSU) and Franklin & Marshall College (Pennsylvania, USA; F&M). Hand-picked fresh chips of each sample were powdered and fused on-site. Additional details on preparation and fusion are reported in supporting information S1. At WSU, major elements were analyzed on a ThermoARL Advant'XP+sequential X-ray fluorescence (XRF) spectrometer using the preparation and analytical procedure described in detail by Johnson et al. (1999) and Kelly (2018). At F&M, major elements were analyzed on a Malvern PANalytical Zetium X-ray fluorescence spectrometer. Trace element concentrations were acquired at WSU and the University of Rhode Island (USA). Digestions and dilution were performed on-site. At WSU, trace elements were analyzed on an Agilent quadrupole ICP-MS using the method of Knaack et al. (1994). Analytical precision is <5% for REE's and <10% for the remaining elements. At University of Rhode Island, trace elements were analyzed on a Thermo X-Series 2 quadrupole ICP-MS following the method of Savov et al. (2005). Interference corrections were applied for TiO on Zn, Ba⁺⁺ on Ga, and CrO on Nb. Analytical precision is <5% for most elements, and precision for Cr and Ni was <2%. Two samples (NC18-15D, NC18-49) were analyzed by all three labs (Table S2, S3).

4.2 Oxygen and hydrogen stable isotope geochemistry

Oxygen and hydrogen isotope compositions of serpentine, talc, and magnetite mineral separates were measured at the University of Texas at Austin using a ThermoElectron MAT 253 mass spectrometer. Serpentine samples were coarsely crushed and handpicked under a binocular microscope to ensure purity of mineral separates. Handpicked serpentine and talc grains were washed with dilute HCl to remove any trace carbonate material. Approximately 2.0 mg of each mineral separate were analyzed using the laser fluorination method of Sharp (1990)

in which samples were heated by a CO₂ laser in the presence of a BrF₅ atmosphere to liberate oxygen. Liberated oxygen was cryogenically purified and analyzed as O₂. Precision and accuracy of oxygen analyses were verified through garnet standard UWG-2 ($\delta^{18}\text{O} = +5.8\text{‰}$) (Valley et al., 1995), in-house olivine standard San Carlos ($\delta^{18}\text{O} = +5.2\text{‰}$), and in-house quartz standard Lausanne-1 ($\delta^{18}\text{O} = +18.1\text{‰}$). All $\delta^{18}\text{O}$ values are reported relative to SMOW, where the $\delta^{18}\text{O}$ value of NBS-28 is +9.7‰. The error on each $\delta^{18}\text{O}$ analysis is $\pm 0.1\text{‰}$, based on the long-term average of standard analyses. Seven samples were run in duplicate to ensure reproducibility and consistency over four analytical runs.

Hydrogen isotope ratios were measured using the method of Sharp et al. (2001). Samples were hand-powdered using an agate mortar and pestle. Approximately 1 mg of each sample were loaded into silver foil capsules, dried under vacuum at 70° C for 24 h, transferred to a Costech zero-blank autosampler, and flushed with He gas. Samples were measured by continuous-flow mass spectrometry using a ThermoElectron TC/EA (high-temperature conversion elemental analyzer) coupled to the ThermoElectron MAT 253 mass spectrometer. Four internationally referenced and certified standard materials (IAEA-CH7, NBS-22, USGS-57, USGS-58) and one in-house working glass standard were analyzed with the samples throughout the run. Raw δD values were corrected for instrumental drift and normalized to SMOW using a calibration curve generated from the measurements of the standard reference materials. Error based on the reproducibility of standards measured in the analytical runs is $\pm 2\text{‰}$. δD values referenced in the text and plotted are the mean of two individual replicate analyses.

4.3 Raman spectroscopy

Raman spectra were acquired on eight serpentinite samples (two samples from Group I, four samples from Group II, one from the Bogen Terrane, and the sample from Kalaa-Gomen). In situ spot analyses on 30 μm polished thin sections were acquired on a Witec Alpha 300R confocal Raman microscope at the Characterization Facility, University of Minnesota. The confocal Raman microscope is equipped with a UHTS300 spectrometer and DV401 CCD detector. Spectra were acquired with a frequency doubled Nd:YAG 532 nm laser, a 1,800 g/mm grating, and a 100x objective. Spot size was $\sim 1\text{ }\mu\text{m}$ in diameter. Each spectra resulted from the average of two 20 s acquisitions to optimize the signal/noise ratio. Two spectral intervals were measured: a low-wavenumber region (100-1200 cm^{-1}) for structural bonding characterization and

a high-wavenumber region (3300- 4050 cm^{-1}) for characterization of hydroxyl groups. A range of relevant textural settings were targeted in each thin section (e.g., serpentine matrix, veins, and pseudomorphs). Data were processed using the WITec Project Five+ software. Raw spectra underwent background subtraction followed by smoothing using a third-order polynomial Savitzsky-Golay filter. Serpentine species were identified by comparison to previously published data (Auzende et al., 2004; Groppo et al., 2006, Petriglieri et al., 2015; Tarling et al. 2018).

5 Results

5.1 Whole rock major elements

Serpentinites from the HP/LT complex display differences in SiO_2 , Al_2O_3 , MgO , and CaO contents and cluster in two distinct geochemical groupings (Table 1; Figure 4). One group of serpentinites ($n = 15$; herein referred to as Group I) is depleted in MgO compared to other serpentinites (herein referred to as Group II, $n = 9$): 34.05 ± 3.34 wt% in Group I versus 40.19 ± 3.20 wt% in Group II. Group I serpentinites contain higher Al_2O_3 contents than Group II (2.34 ± 1.47 versus 0.75 ± 0.34 average wt%, respectively). Group I samples are generally elevated in SiO_2 compared to Group II, though they overlap within uncertainty, with averages of 44.96 ± 5.23 wt% and 41.13 ± 2.13 wt%, respectively. $\text{Fe}_2\text{O}_{3(\text{T})}$ contents do not vary significantly between the two groups, though samples in Group I display a broader range of variation (7.92 ± 1.85 wt % versus 7.38 ± 0.68 wt% for Group II). CaO concentrations in the HP/LT serpentinites overall are <0.5 wt%, with the exception of the three least serpentinized samples (Table 1 and Figure S2); all Group II samples located in the SE portion of the complex (0.53, 0.57, and 1.08 wt%). The ophiolitic reference sample from Kalaa-Gomen (NC19-178) has 0.66 wt% CaO . Samples have a range of loss-on-ignition (LOI) values, from 5.25 to 12.64%, indicating varying degrees of hydration, with an overall average of $10.43\% \pm 2.77\%$. Na_2O , K_2O , and P_2O_5 are close to or below detection limits.

The chlorite schist (NC19-85) is elevated with respect to the talc schist (NC19-169) in Al_2O_3 , $\text{Fe}_2\text{O}_{3(\text{T})}$, and TiO_2 , whereas the talc schist is elevated in SiO_2 and MgO (Table 1). The chlorite schist (NC19-85) has CaO and P_2O_5 contents of 2.47 and 1.88 wt%, respectively, due to the presence of ~5% modal abundance apatite.

5.2 Whole-rock trace elements

Group I and II samples are overall depleted in trace elements with respect to primitive mantle (Figure 5a, b). Both groups show enrichments in Cs and Pb and negative Rb and Nb anomalies. Group I samples show marked Th and U enrichments and a strong negative Sr anomaly compared to Group II. Group I samples show greater overall enrichment in middle to heavy rare-earth elements (M-HREE; Sm, Eu, Gd, Tb, Dy, Y, Ho, Er, Yb, Lu) compared to Group II (Figure 5c-f). Differences in the pattern of light-rare earth elements (LREE; La, Ce, Pr, Nd, Pm) to HREE most clearly distinguish the two groups ($\text{La}_N/\text{Ho}_N = 1.66 \pm 1.33$ for Group I and 6.59 ± 1.92 for Group II; normalized to primitive mantle).

The chlorite schist (NC19-85) and talc schist (NC19-169) samples display distinct enrichments in trace elements. The chlorite schist is enriched in Li, Sc, V, Cu, Zn, Sr, Y, Zr, Nb, Ba, La, Ce, Pr, Nd, Sm, Eu, Tb, Gd, and Dy relative to the talc schist. Conversely, the talc schist is enriched in Cr and Ni relative to the chlorite schist.

5.3 Stable isotope geochemistry

5.3.1 Oxygen isotopes

Serpentine from the high-pressure complex displays a wide range of oxygen isotope values: from +5.1 to +10.2‰ (Table 2 and Figure 6a, b). Group I samples are between +6.7 and +10.2‰, whereas Group II samples are between +5.1 to +8.0‰. The mean serpentine $\delta^{18}\text{O}$ value for Group I is $+8.7 \pm 0.9\text{‰}$ ($n = 15$) and $+5.5 \pm 0.4\text{‰}$ ($n = 9$) for Group II. A Group II sample (NC18-39A) shows a 1‰ increase in $\delta^{18}\text{O}$ between serpentine extracted from the sample interior (+5.4‰) and serpentine from an outer altered rind (+6.4‰). $\delta^{18}\text{O}$ sample replicates reproduced within 0.2‰, with the exception of two samples, NC18-15D and NC18-22A, which show variability of 1.8‰ and 0.9‰, respectively. Comparative samples from the Boghen Terrane, Kalaa-Gomen Massif, and the Peridotite Nappe serpentinite sole have $\delta^{18}\text{O}$ values of +7.5‰ (average of two samples), +5.7‰, and +6.5‰, respectively.

Oxygen isotopes in magnetite were measured in three samples from the NE part of the complex: a serpentinite (NC19-86), a chlorite schist (NC19-85), and a talc schist (NC19-169). The $\delta^{18}\text{O}$ value of magnetite is +1.5‰ in the serpentinite and +2.4‰ in the chlorite schist. In the talc schist, two separate grain size fractions were measured, 125-250 μm and >710 μm . The 125-

250 μm grain size fraction has a $\delta^{18}\text{O}$ value of +4.3‰, whereas the $>710 \mu\text{m}$ has a value of +4.6
Oxygen isotopes in talc from two samples (one from the SE and one from the NE), were
+9.0 and +10.5‰, respectively.

5.3.2. Hydrogen isotopes

δD values of serpentine in the high-pressure complex range from -79 to -29‰ (Figure
6a). Group I sample range between -76 and -29‰ and Group II samples range between -79 and -39‰ . The mean serpentine δD value is $-44 \pm 10\text{‰}$ ($n = 30$) for Group I and $-52 \pm 14\text{‰}$ ($n = 18$)
for Group II. The relatively large standard deviation in both groups derives from three samples
(one in Group I and two in Group II) that have δD values between -79 and -70‰ . Without these
values, the remaining samples from Group I exhibit a more restricted range of δD values from -49
to -32 , with an average of $-41 \pm 5\text{‰}$ ($n = 28$). Group II ranges from -59 to -39‰ , with an
average of $-46 \pm 8\text{‰}$ ($n = 14$). Comparative samples from the Boghen Terrane, Kalaa-Gomen
Massif, and the Peridotite Nappe serpentinite sole have δD values of -81‰ (average of two
samples), -85‰ , and -82‰ , respectively. Across all samples, δD sample replicates reproduced
within 2‰ , on average.

5.4 Raman spectroscopy

Serpentine has three primary structural polymorphs that vary over P - T space: the low- T
form is chrysotile, followed by lizardite and antigorite. Diagnosing the polymorph in exhumed
serpentinites can aid in retrieving information about its prograde and/or retrograde path and
alteration. Serpentine in two samples from Group I and four samples from Group II was found to
be antigorite (only) in contrast to lizardite identified in comparative samples from the Boghen
Terrane basement unit and Kalaa-Gomen massif in the Peridotite Nappe (Figure 7 and S5-6).

5.5 Categorization of New Caledonia HP/LT serpentinites

As presented above, serpentinites in the high-pressure metamorphic complex cluster in
two distinct groups, revealed by petrography, major and trace element, and stable isotope
geochemistry. The significance of these groups and their composition will be discussed in the
next section. The results are summarized as follows:

Group I: >90% serpentinized, common assemblage of antigorite + talc + magnetite, lower, more homogeneous MgO contents (relative to Group II), $\delta^{18}\text{O}$ values between +6.7‰ and 10.2‰, flat M-HREE patterns;

Group II: ~50-90% serpentinized, notable preservation of orthopyroxene and olivine and distinct presence of awaruite, higher MgO contents (relative to Group I), $\delta^{18}\text{O}$ between +5.1‰ and +8.0‰, distinct curved LREE to HREE pattern compared to Group I.

6 Discussion

Prior studies of serpentinites in the HP/LT terrane of New Caledonia have disagreed on the tectonic origin of the mantle protolith (Fitzherbert et al., 2004; Spandler et al., 2008). Our results indicate the existence of two geochemically-distinct groups of serpentinites in the blueschist- to eclogite-facies portions of the complex. This heterogeneity may derive from (1) geochemical differences in the original mantle source material, (2) processes occurring throughout serpentinization, (3) subsequent metamorphism during subduction and exhumation, or (4) a combination of the above.

6.1 Assessing major and trace element mobility

The major and trace element compositions of exhumed serpentinites result from (a) the original composition and melt history of the parent peridotite, (b) fluid-rock reactions occurring at the source location for serpentinization (e.g., seafloor, mantle wedge), and (c) fluid-rock reactions occurring during subduction and exhumation. Extensive study of variably serpentinized abyssal and mantle wedge peridotites and serpentinites in exhumed subduction complexes has demonstrated high-field strength elements (HFSE; Nb, Ta, Zr, Hf) and REE are generally (but not wholly) immobile during serpentinization and aqueous fluid alteration and can be used to discern information about protolith composition and pre-serpentinization magmatic processes (e.g., Deschamps et al., 2013; Kodolányi et al., 2012; Niu, 2004; Parkinson & Pearce, 1998; Savov et al., 2005; Scambelluri et al., 2004). By contrast, major elements and FME (e.g., Li, Pb, U, Cs, Sr, Ba) must be interpreted with additional caution, as serpentinization is not isochemical with respect to some of these elements and/or they are more readily mobilized during fluid interaction over the course of subduction and exhumation (e.g., Cannao et al., 2016; Malvoisin, 2015; Peters et al., 2017). With this in mind, we first assess the degree to which major and trace

elements in the New Caledonia HP/LT serpentinites may have been mobilized, or whether they otherwise retain faithful geochemical signatures of their initial mantle protolith.

Ratios of immobile trace elements (e.g., LREE, HFSE vs HREE; La/Yb, Zr/Yb, Hf/Yb, Nd/Yb) are observed to vary independently of LOI for both groups of HP/LT serpentinites (Figure 8). This result, combined with the observed similarity of intra-group REE patterns for Groups I and II (Figure 5), provides initial confidence in the retention of protolith mantle geochemical signatures for these elements (c.f. Savov et al., 2005). For further assessment, plots of LREE (La, Ce, Pr, Nd) versus HFSE (Nb, Ta, Zr, Hf) elucidate whether a common process or processes were responsible for the enrichment or depletion of these groups of elements (c.f. Niu, 2004). It is worth noting that LREE are more readily mobilized by aqueous fluids than MREE and HREE, so this test is a conservative indicator of trace element and REE immobility.

Group I LREE's are almost entirely decoupled from HFSE (Figure 9). Indications of a once-coupled relationship between the LREE and HFSE are partially retained in a few samples, as indicated by consistent low abundances that correlate somewhat more linearly (but to a degree no longer statistically significant). More often, however, these samples have been overprinted by a process or processes that added LREE (Figure 9). In contrast, statistically significant correlations are observed for Group II serpentinites between the LREE and Zr and Nb ($R_{\text{LREE-Zr}} = 0.942 \pm 0.027$, $R_{\text{LREE-Nb}} = 0.678 \pm 0.024$) (Figure 9). Notably, LREE appear decoupled from Ta and Hf ($R_{\text{LREE-Ta}} = -0.267 \pm 0.037$, $R_{\text{LREE-Hf}} = 0.308 \pm 0.106$). Of the HFSE, Zr and Nb (here coupled with LREE) are lighter in mass than Ta and Hf. Niu (2004) addresses the potential importance of mass-dependent effects on the observed magmatic fractionation of Nb/Ta and Zr/Hf ratios in global abyssal peridotite datasets, proposing that elevated LREE abundances could be produced or enhanced by mass-dependent transfer rates, but notes that observational tests of this hypothesis are necessary. These data possibly corroborate this hypothesis, particularly given that none of the four elements correlate with LOI. However, the possibility that this decoupling reflect later aqueous alteration processes cannot be ruled out, though it is unclear whether subduction zone temperatures would be sufficiently high enough to mobilize these HFSE.

In summary, both groups show variations in trace and REE independent of LOI - a valuable first indicator for retained magmatic trace element signatures. Strong correlation

between LREE and HFSE in Group II samples bolsters this argument. Observed mobility in Group I LREE does not preclude us from utilizing the REE data to ascertain a tectonic setting for the mantle protolith, as there is no demonstrated correlation between the M-HREE with LOI; though care must be taken in interpreting the significance of LREE abundances and their contribution to the overall shape of L-HREE trends.

6.2 Serpentinization processes and fluid interactions

Within the ultramafic rock system, oxygen and hydrogen isotopes have been widely utilized to discern the source of serpentinizing fluids and subsequent post-serpentinization fluid histories (e.g., Alt and Shanks, 2006; Burkhard and O'Neil, 1988; Fröh-Green et al., 1990, 2001; Kyser et al., 1999). Ultimately, stable isotope values are the amalgamated product of several important variables – the stable isotope composition of the interacting fluid, the fluid-rock ratio, the composition of the reactive solid phase(s) in the system, and the fractionation factors between these components and the fluid. Stable isotope data for the HP/LT serpentinites indicate differing serpentinizing fluid compositions and/or T conditions for serpentinization between the two groups of samples. Group II $\delta^{18}\text{O}$ values ($+5.5 \pm 0.4\text{‰}$) cluster tightly around the typical upper mantle value ($+5.4\text{‰}$), indicating little to no additional alteration of the primary mantle signature during serpentinization, subduction, or exhumation (Figure 6). This assessment is supported by recent stable isotope modeling of the ophiolite that indicates limited interaction of slab sediment-derived fluids within New Caledonia's mantle wedge during the initial phases of subduction (Ulrich et al., 2020). Moreover, these authors discern, on the basis of Sr isotopes and low abundances of sediment-derived elements such as As and Sb (Deschamps et al., 2013), that the input of subducted sediments was low at the time of serpentinization of the mantle wedge, and instead the dehydration of altered oceanic crust comprised the dominant fluid source. Though our data do not include measurements of As and Sb, we observe similarly low degrees of elements typically regarded as classic markers of serpentinization in the mantle wedge via shallow dewatering fluids released from slab sediments and lower- T metamorphic dehydration reactions (e.g., Cs, Sr, Rb, Li) (Figure 1). The narrow range of $\delta^{18}\text{O}$ values across all degrees of hydration (as indicated by LOI %) supports a consistent temperature of serpentinization and/or a consistent fluid composition (Figure 6). The close relation between the $\delta^{18}\text{O}$ values of Group II

serpentinites and antigorite within slab-derived tremolite-antigorite veins in the Peridotite Nappe also supports a similar source for serpentinizing fluids.

By contrast, Group I samples show increased and variable $\delta^{18}\text{O}$, which may be interpreted to reflect: (1) interaction with metamorphic fluids with a sedimentary component, (2) differing temperatures of serpentinization in a seafloor environment, or (3) a mix of these two influences. Both end-member explanations are plausible - all but two Group I samples are located within 2 km of interpreted boundaries between the Pouébo Terrane and the sediment-dominated Diahot-Panié Complex. Moreover, significant fluid fluxing is evident for these samples given major element concentrations, which show a clear deviation from the terrestrial array as a function of increasing modal abundances of talc (Figure 6). The development of talc-rich serpentinite and hybrid rock assemblages in the NE portion of this HP/LT complex has been attributed to high fluid flux-driven metasomatism at depth and is the source of a separate study by Spandler et al. (2008).

A suite of discriminant fluid-mobile trace element diagrams complements our interpretation of significantly different fluid and tectonic histories for Groups I and II. FME's in serpentinites are a sensitive recorder of fluid interactions, in part because peridotite protoliths are depleted in these elements, so increases in their concentration are attributable to fluid interactions (e.g., Peters et al., 2017; Scambelluri et al., 2019). Recent re-analysis and refinement of compiled serpentinite data has also demonstrated that these elements are sensitive tracers of the tectonic environments of serpentinization (Peters et al., 2017). For the New Caledonia HP/LT serpentinites, plots of fluid-mobile element enrichments (e.g., Ba, Cs) relative to fluid-immobile elements such as Yb show compelling trends (Figure 10). Ba in Groups I and II (Figure 10a) indicates a mantle wedge origin for Group II samples, which are relatively enriched compared to Group I samples, which fall squarely within the field for mid-ocean ridge serpentinites. This trend is not observed for Cs, which we interpret as becoming enriched in Group I samples during intense fluid flow and deformation, and possible interaction with neighboring sediments from the Diahot-Panié Complex. Cs in Group II falls lower than compiled forearc data, commensurate with our interpretations for limited influence of sedimentary-derived fluids based on stable isotope data. This Cs depletion is apparent in a plot of Cs vs fluid-immobile Yb, though the data in this context better overlap with the range of observed values for drilled or dredged forearc samples, and the Group I samples correlate well with the compiled mid-ocean ridge dataset.

Trends between U and Th enrichments show that Group I samples lack the seafloor U enrichment that defines some (but not all) of the mid-ocean ridge dataset (Figure 10c).

6.3 Tectonic origin of New Caledonia HP/LT-complex serpentinites

Given evidence presented above, we interpret differing tectonic origins for serpentinites in the NE and SE portions of the terrane:

Group I: Found dominantly in the NE part of the complex, these serpentinites are remnants of lithospheric mantle from the downgoing plate (Figure 11a). They experienced progressive serpentinization on or near the ocean floor prior to subduction and then faced high-fluid-rock ratios, metasomatism leading to the development of talc-bearing assemblages, and deformation in the subduction channel. The formation of these rocks was possibly associated with the development of several major shear zones, which facilitated slicing of the complex at depth and have given parts of the Pouébo Terrane the appearance of a *mélange*. This geodynamic interpretation is supported by recent reevaluations of the architecture of the belt (Maurizot et al., 2020c; Vitale-Brovarone et al., 2018) and consistent with our observation of the proximity of serpentinites to major unit boundaries and faults (e.g., NC18-22A, NC18-26B, NC19-63) and their prevalence as matrix hosts for dismembered blocks in km-scale shear zones. It is plausible that the pre-subduction abyssal environment for the future Pouébo Terrane ultramafic rocks was characterized by lower-T conditions, consistent with stable isotope data presented here (Figure 6b). Seismic imaging of the New Caledonia Basin to the west of the island has revealed thinned continental crust interpreted to consist of a significant volume of serpentinized upper mantle and resembling the structure of crust formed during slow, low-T amagmatic seafloor spreading (Klingelhoefer et al., 2007).

Group II: These samples cluster in the southern portion of the complex, comprising the Poadja Massif. Incorporated onto the top of the slab prior to the onshore obduction of the Peridotite Nappe, this slice of mantle wedge was subducted and exhumed intact, with striking preservation of relict minerals and comparatively minor degrees of fluid alteration, as evidenced by $\delta^{18}\text{O}$ isotope values that remain near the pristine upper mantle average and lower degrees of LREE and FME (Cs, U, Pb, and Sr) enrichment relative to other global forearc mantle settings (Figure 11b). We interpret these data to reflect the lack of a strong sedimentary-derived fluid flux during serpentinization of these rocks in their mantle wedge setting, and this is supported by

independent evidence from analyses of slab-derived tremolite vein assemblages at the base of the obducted Peridotite Nappe (Cluzel et al., 2020). This muted enrichment of LREE and FME also indicates that later overprinting by meta-sedimentary derived fluids during transport within the subduction channel was limited. The samples display characteristic REE patterns that remarkably resemble the harzburgitic mantle of New Caledonia's Peridotite Nappe (Figure 11b).

6.4 Relevance for the architecture of the New Caledonia HP/LT complex

In the NE portion of the complex, our combined field observations and geochemical data suggest that Group I (downgoing-slab) serpentinites: (a) overwhelmingly appear in the highest-grade portions of the complex, within the oceanic affinity Pouébo Terrane, and (b) show distinct geochemical markers for high degrees of fluid interaction and deformation during subduction and/or exhumation (i.e., elevated $\delta^{18}\text{O}$ values, decoupled LREE's, enriched FME's such as Cs). These observations are broadly consistent with recent re-interpretations of this subduction complex by Vitale-Brovarone et al. (2018), which posits the existence of two accretionary domains preserved in a nappe-type belt formed by progressive underplating and stacking. The SE portion of the complex is far less studied: the existence of Group II mantle wedge serpentinites carries implications for the structural level of the subduction zone sampled by rocks in the immediate vicinity of the Poadja Massif. The existence of mantle wedge serpentinites adds to independent observations made by Cluzel (2021) of the presence of hangingwall mafic rocks in lower-grade portions of the Diahot-Panié Complex further north. Integrating these observations into a unified subduction complex-scale picture of preserved lithostratigraphy requires further studies. Nevertheless, these observations provide compounding evidence for greater structural coherence within the New Caledonia subduction "mélange" than has previously been posited.

6.5 Implications for slab-mantle wedge interactions

Our finding of mantle wedge material within the exhumed HP/LT complex on New Caledonia provides direct evidence for interaction with the hangingwall mantle in this Eocene subduction zone. Geochemical modeling has demonstrated the importance of the mantle wedge as a reservoir for water and fluid-mobile elements that contribute to arc magma isotope and elemental budgets (e.g., Debret et al., 2019, 2020; Ribeiro & Lee, 2017; Savov et al., 2005, 2007), and there are strong conceptual bases, observational evidence, and modeled predictions

for the incorporation and downward dragging of mantle wedge material in some subduction zones (e.g., Bebout & Barton, 2002; Bebout, 2007; Cloos & Shreve, 1988; Malatesta et al., 2012; Peacock & Hyndman, 1999; Savov et al., 2005). New Caledonia provides an important natural rock record comparison for geodynamic modeling: particularly for outcomes in which exhumation of downgoing plate mantle *and* overlying mantle wedge material is predicted (e.g., Gerya & Stockert, 2006; Gerya et al., 2002; Malatesta et al., 2012). In these cases, the rheology of serpentine and evolving hydration state of the ultramafic material are critical to the locus of the weak, low-viscosity subduction channel and eventual migration of the plate interface upwards into the mantle wedge. The development of talc in the overlying mantle assemblage provides a possible physiochemical mechanism for this migration, and its stability in the ultramafic chemical system may even control the depth of plate decoupling (Peacock & Wang, 2021).

This hypothesis is interesting to consider in light of our observations of relatively undeformed, nearly talc-absent mantle wedge assemblages observed in the SE and deformed, talc-bearing downgoing mantle assemblages in the NE. A local bulk assemblage-induced rheological switch in the mantle wedge and consequent avulsion of the slab-mantle interface could be broadly consistent with the results presented here. In this scenario, deformation rapidly re-localizes in the new region of weakness and a stranded slice of mantle wedge material is incorporated atop the downgoing slab. This mantle may feasibly have already experienced partial high-T serpentinization from fluxed aqueous dehydration fluids (consistent with stable isotope data presented here) and might largely escape prolonged intense deformation, because the subduction interface was localized first below, and then above the mantle wedge slice in rheologically weaker assemblages (consistent with observed petrographic textures and field observations).

Tectonic erosion of the mantle wedge, and the type of HP/LT ultramafic material it produces, contrasts markedly with the processes that retrieve lithospheric mantle under subducted oceanic crust. Slicing and nappe development along lithologic (rheologic) heterogeneities in the slab are commonly invoked as ways by which downgoing mantle may be incorporated into the subduction channel (e.g., Angiboust & Agard, 2010; Guillot et al., 2009; Hermann et al., 2000; Wakabayashi, 1992). During these processes, the mantle is juxtaposed and reacts with sedimentary and mafic components of the slab, as reflected by stable isotope

compositions and Si and trace element enrichments. The data presented here for New Caledonia's NE serpentinites reflect this type of metamorphic history, including their spatial concentration within the oceanic Pouébo Terrane and proximity to inferred boundaries with the Diahot-Panié Complex.

7 Summary

In this study, we resolve two petrographically, geochemically, and spatially distinct groups of serpentinites within New Caledonia's HP/LT subduction complex. Our work reconciles existing disparate interpretations of the tectonic origin of these rocks, with implications for slab-mantle interaction during subduction. In the NE portion of the complex serpentinite geochemistry and field relations indicate a downgoing slab origin. By contrast, the chemistry of serpentinites in the SE points to a mantle wedge origin, indicating incorporation and subduction of upper plate material. The exhumation of serpentinites from these two different geodynamic settings within a single HP/LT complex is not common in the natural rock record, but predicted for certain modeled scenarios as a function of serpentine rheology and hydration state, among other subduction parameters. The mineralogy, chemistry, and spatial distribution of New Caledonia's HP/LT serpentinites hold clues for understanding the architecture of this subduction complex, and more broadly, for the modeled dynamics of deep tectonic slicing of subducted oceanic plates and mass transfer between the upper and lower plates.

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9 Open Research

Data Availability Statement

To ensure the accessibility and discoverability of the samples used in this research, and to align with the National Science Foundation's guidelines of effective data practices, all samples used in this study have been registered with IGSN Global Sample Numbers through the System for Earth Sample Registration (SESAR). SESAR is maintained by the GeoInformatics Research Group of the Lamont-Doherty Earth Observatory. <https://www.geosamples.org/> Accessed 17 February 2022. Whole rock and stable isotope data presented in this paper will be submitted for archival in EarthChem data repository prior to paper acceptance.

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1319 $\text{MgO}-\text{SiO}_2-\text{H}_2\text{O}$ (MSH). *Lithos*, 41(1–3), 213–227. [https://doi.org/10.1016/S0024-](https://doi.org/10.1016/S0024-4937(97)82013-0)
1320 [4937\(97\)82013-0](https://doi.org/10.1016/S0024-4937(97)82013-0)
- 1321 **Figure 1.** Schematic subduction zone cross-section showing relevant locations of
1322 serpentinization and enrichments and loss of fluid-mobile elements during subduction and
1323 exhumation (after Deschamps et al., 2013; Peters et al., 2017). Large semi-transparent arrow
1324 indicates subduction and exhumation path of serpentinites, emphasizing that these rocks can
1325 encounter chemical changes on both the prograde and retrograde paths. Subduction zone after
1326 Guillot et al. (2015).
- 1327 **Figure 2.** (a) Regional map highlighting significant tectonic features of the SW Pacific region
1328 after Maurizot & Collot (2009) and Sutherland et al. (2019). (b) Simplified geologic map of the
1329 main island of New Caledonia showing sample locations and additional localities referenced in
1330 text. Map and units after Maurizot et al. (2020c). (c) Simplified map of Pam Peninsula showing
1331 locations of Group I serpentinites samples (this study) and prior studied samples. Unit names
1332 after Maurizot et al. (2020c) and boundaries modified after Vitale-Brovarone et al. (2018). (d)
1333 Simplified map showing sampled locations at Poadja Massif. Estimated uncertainty for

1334 Fitzherbert et al. (2004) georeferenced raster map locations (c, d) is ~500-700 m and ~200-400
 1335 m, respectively. Solid black lines denote roads, solid blue lines denote streams, and dashed black
 1336 line denotes high-angle normal fault.

1337 **Figure 3.** Backscatter electron images from select high-pressure serpentinites. (a) A relict
 1338 chromite core surrounded by a mantle of ferritchromite and rim of chromian magnetite
 1339 containing inclusions of antigorite (NC19-48). (b) Delicate veins of tremolite and antigorite cross
 1340 cut a pervasively serpentinized matrix (NC19-94). (c) Skeletal chromite replaced by magnetite
 1341 and antigorite (NC18-39B). (d) Inclusions of Fe-Ni alloy awaruite and antigorite in magnetite
 1342 (NC18-39A). (e) Brightness-contrast enhanced image highlighting complex curvilinear
 1343 retrogression of chromian magnetite to magnetite (NC19-154A). (f) Representative texture of
 1344 typical replacement of olivine by antigorite and magnetite via growth from fractures (NC18-
 1345 39A).

1346 **Figure 4.** Whole rock major element ratios of MgO/SiO_2 vs. $\text{Al}_2\text{O}_3/\text{SiO}_2$ for serpentinites and
 1347 hybrid rocks in New Caledonia's HP/LT terrane. Samples are plotted with compiled data for
 1348 harzburgites in the New Caledonia ophiolite (gray circles) (Liu et al., 2018; Mothersole, 2014;
 1349 Mothersole et al., 2017; Ulrich et al., 2010). Also shown is the field for abyssal serpentinites
 1350 from Niu (2004). Depleted mantle value from McDonough and Sun (1995), primitive mantle
 1351 value from Salters and Stracke (2004). "Terrestrial array" line after Hart and Zindler (1986),
 1352 Jagoutz et al. (1979), and Niu (2004).

1353 **Figure 5.** Whole rock trace and REE compositions for New Caledonia HP/LT serpentinites.
 1354 Samples are plotted together (a, b) and separately into respective categories: Group I (c, d) and
 1355 Group II (e, f). Trace element concentrations are normalized to primitive mantle values of Sun
 1356 and McDonough (1989) (a,c,e). REE concentrations are normalized to C1 chondrite
 1357 (McDonough & Sun, 1995) (b,d,f).

1358 **Figure 6.** (a) Hydrogen (δD) and oxygen ($\delta^{18}\text{O}$) stable isotope compositions of New Caledonia
 1359 HP/LT serpentinites compared with existing data for serpentine in the overlying Peridotite Nappe
 1360 (ophiolite groups after Cluzel et al., 2020; Ulrich et al., 2020). Weathered samples and samples
 1361 for which antigorite was not specified in Cluzel et al. (2020) were excluded. Upper mantle values

from Eiler (2001), Kyser and O'Neil (1984), and Matthey et al. (1994). (b) $\delta^{18}\text{O}$ versus loss on ignition (LOI) for HP/LT serpentinites. Group I serpentinites show decreasing $\delta^{18}\text{O}$ with progressive hydration whereas Group II serpentinites exhibit a narrow range of $\delta^{18}\text{O}$ values irrespective of degree of hydration. An arrow denotes the direction of $\delta^{18}\text{O}$ change when serpentine is formed at increasing temperatures in equilibrium with seawater (e.g., Saccocia et al., 2009).

Figure 7. Representative cross-polarized photomicrographs and correlated Raman spectra for (a) two separate matrix sites in Group I sample NC18-09B and (b) one pseudomorph (red) and one vein (blue) in Group II sample NC19-152. Diagnostic Raman peaks for antigorite are highlighted tan vertical bars. All serpentine polymorphs in Group I and II were identified as antigorite (Figures S5, S6).

Figure 8. Variations in ratios of HFSE/HREE (i.e., Yb) with increasing LOI (%) for New Caledonia HP/LT serpentinites. Variations in trace element ratios are independent of LOI value.

Figure 9. Plots of LREE (La, Ce, Pr, and Nd) versus HFSE (e.g., Zr) for all HP/LT serpentinites, with zoomed insets in upper right of graphs to highlight trace element depleted Group II samples. Groups I and II symbols are the same as Figure 8. One sample, NC19-158 (open orange circle), has been excluded from the trendline and computed R values.

Figure 10. Plots for investigation of FME trends among New Caledonia HP/LT serpentinites. Data are plotted against compilations for mid-ocean ridge (MOR) serpentinites (teal circles) and forearc (FA) serpentinites (tan circles) from Peters et al. (2017). (a) Ba/Yb vs. Ba, (b) Cs/Yb vs. Cs, (c) U vs. Th, and (d) Cs vs. Yb.

Figure 11. REE compositions for New Caledonia HP/LT serpentinites. (a) Group I serpentinites plotted versus a global dataset of mid-ocean ridge serpentinites (Peters et al., 2017); (b) Group II serpentinites plotted versus harzburgite compositions from the New Caledonia ophiolite (Marchesi et al., 2009; Secchiari et al., 2020; Ulrich et al., 2010).

Figure 1.

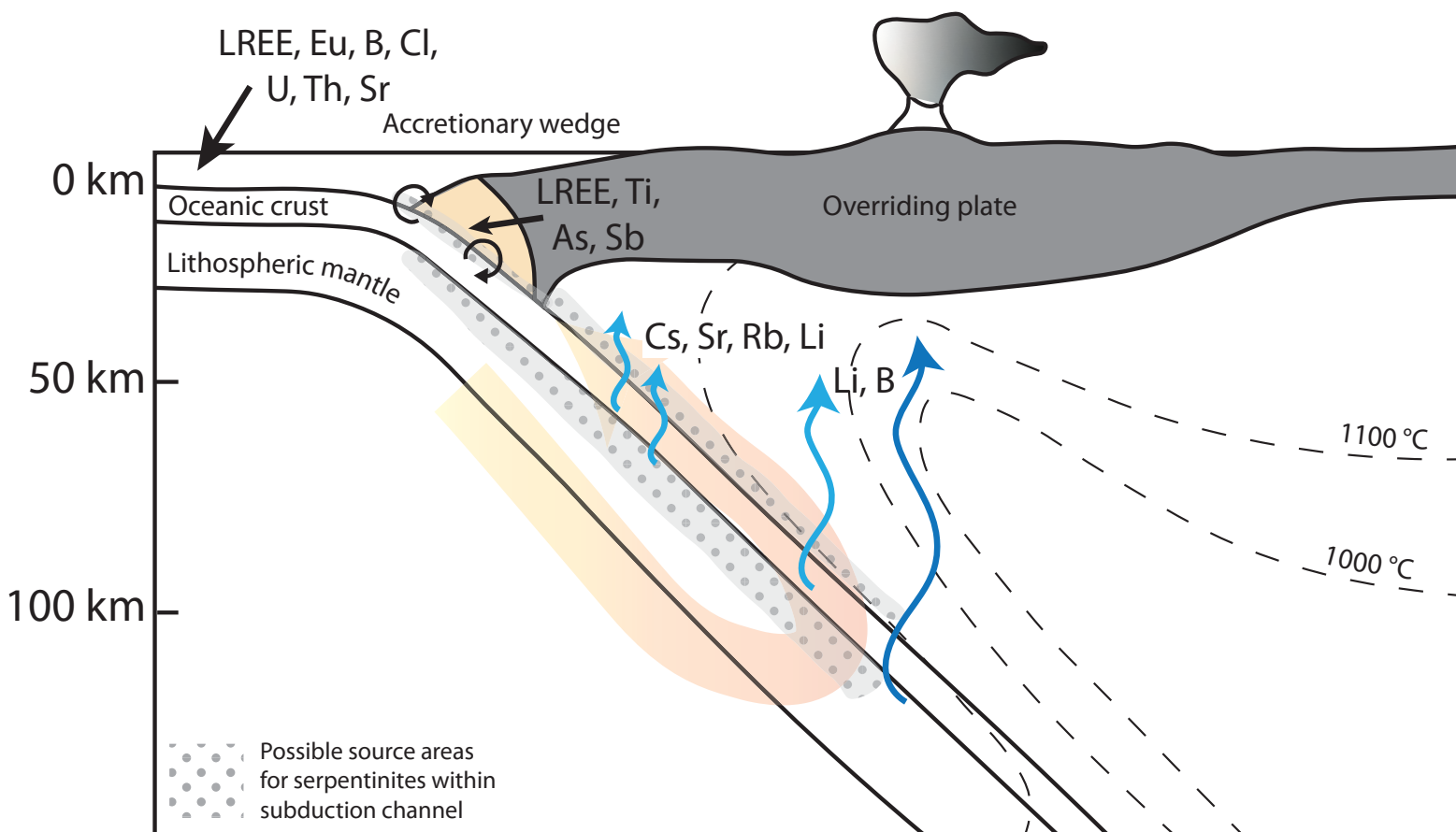


Figure 2.

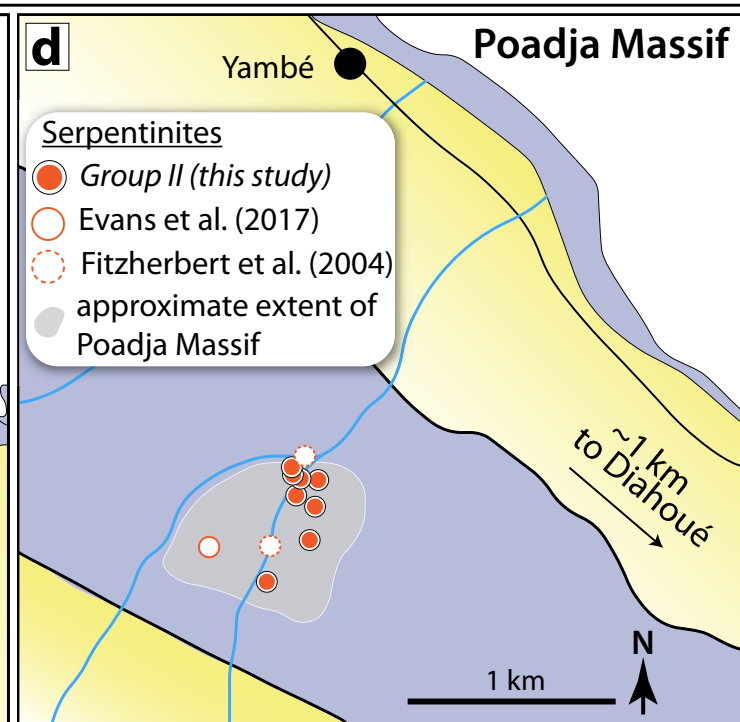
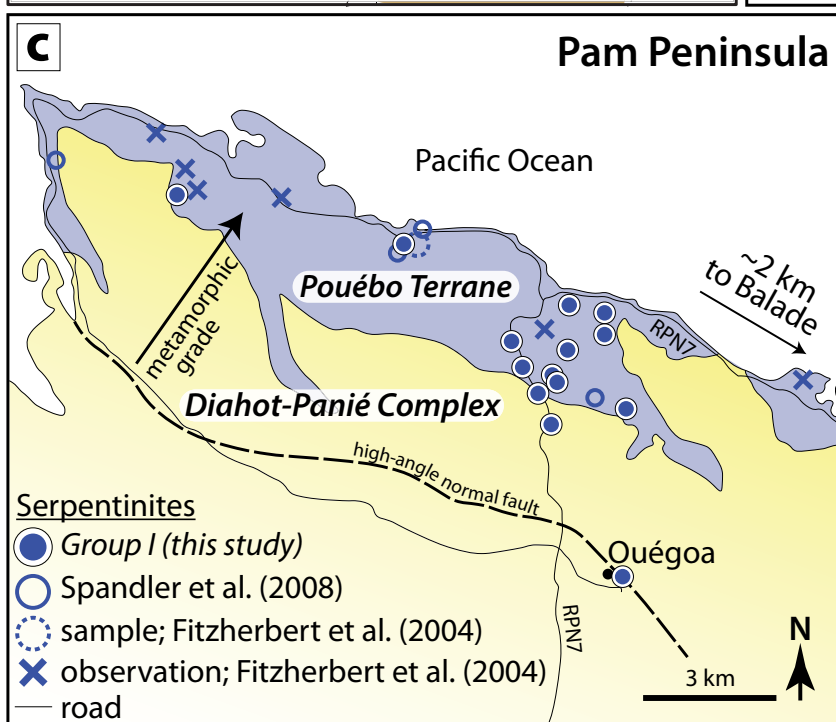
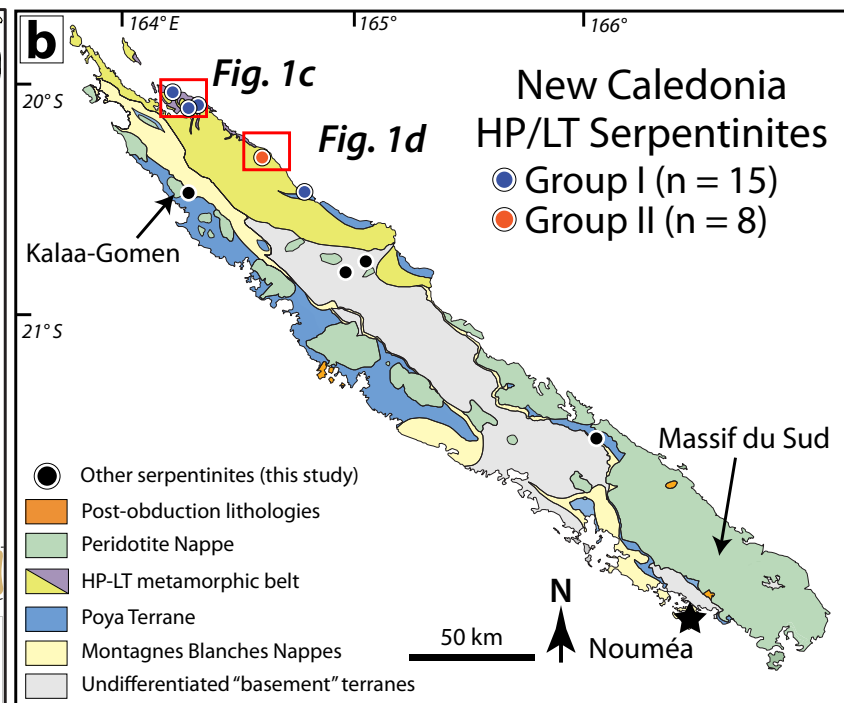
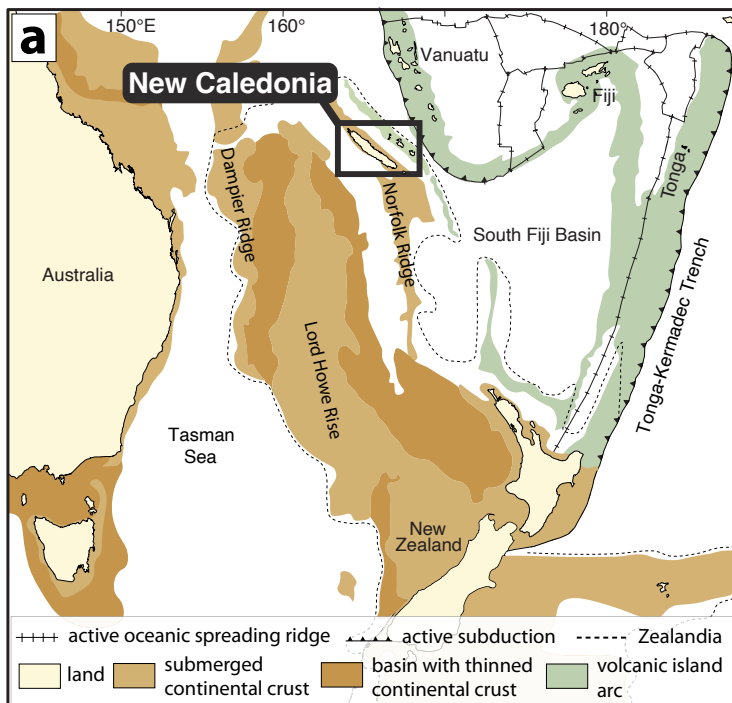


Figure 3.

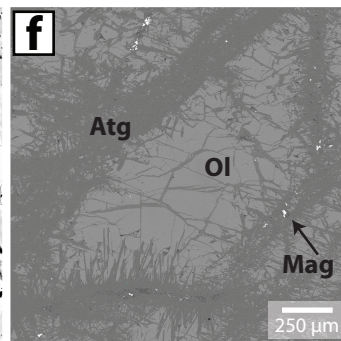
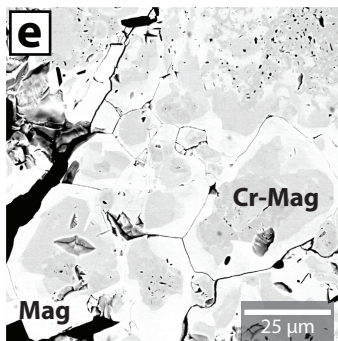
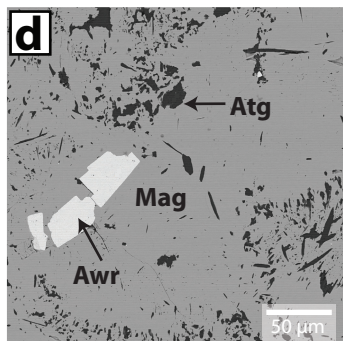
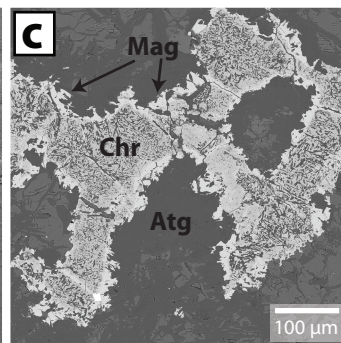
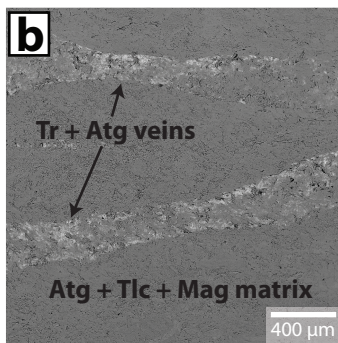
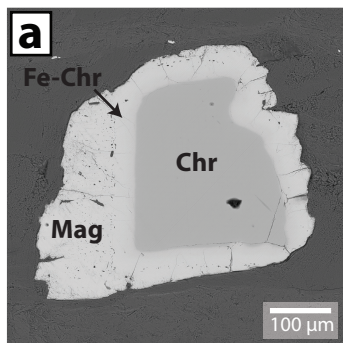


Figure 4.

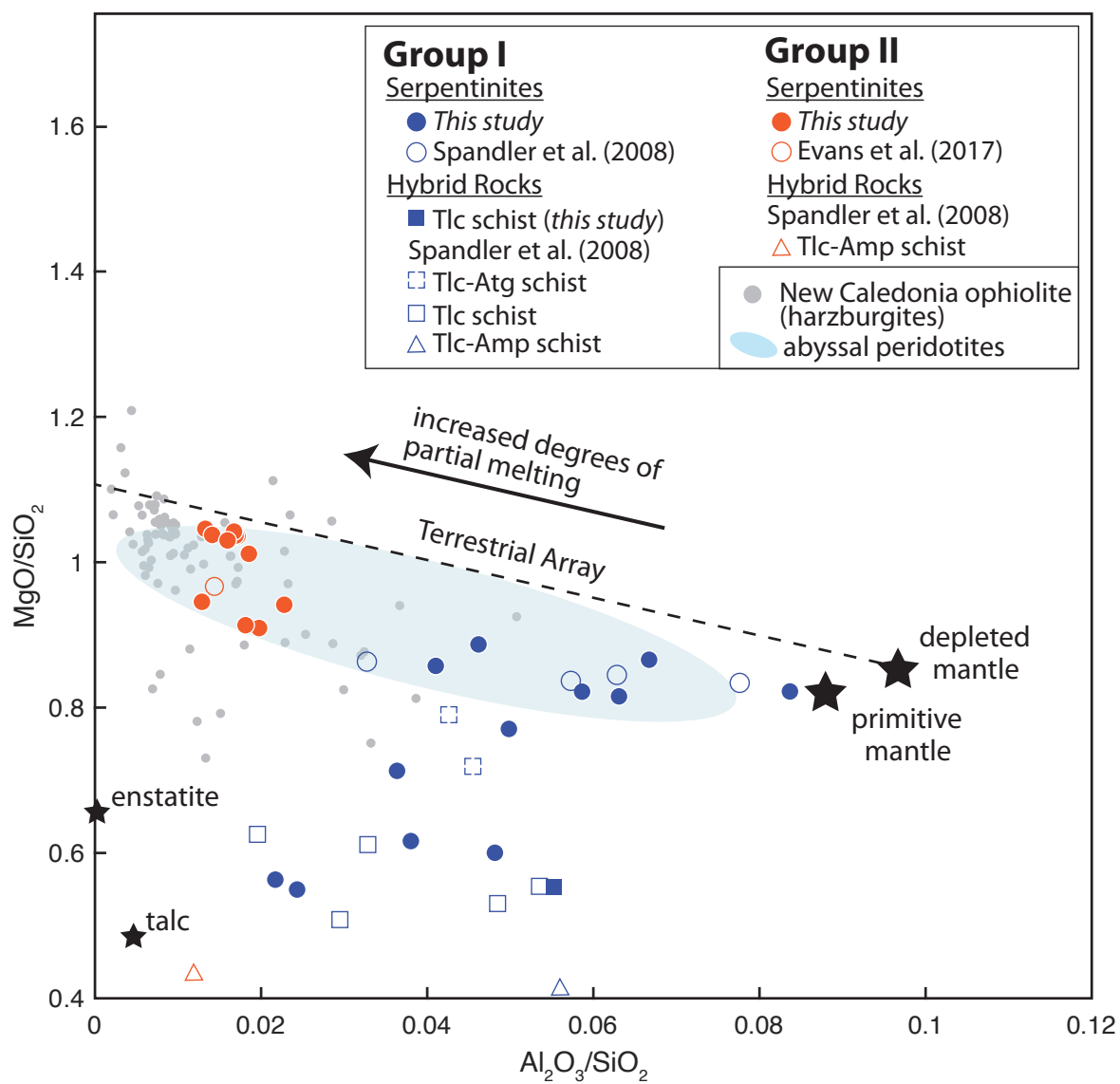


Figure 5.

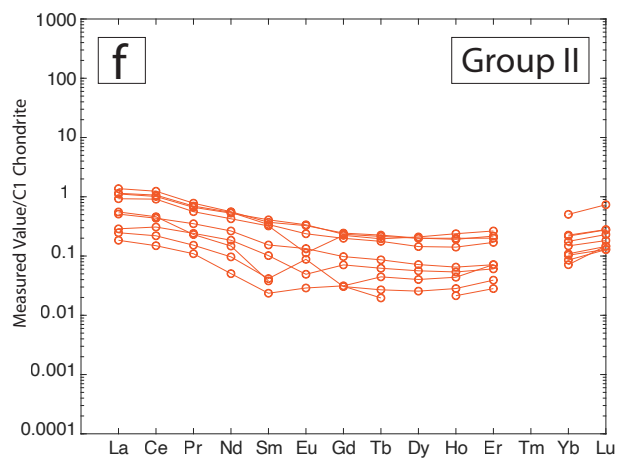
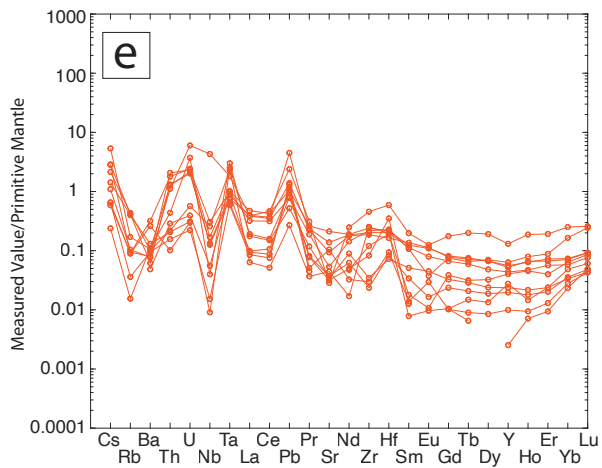
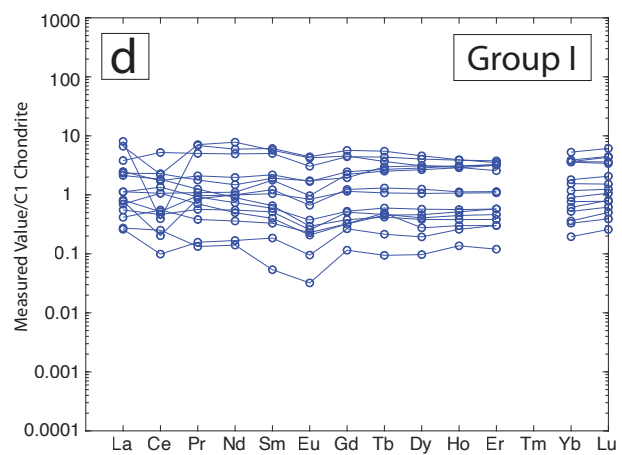
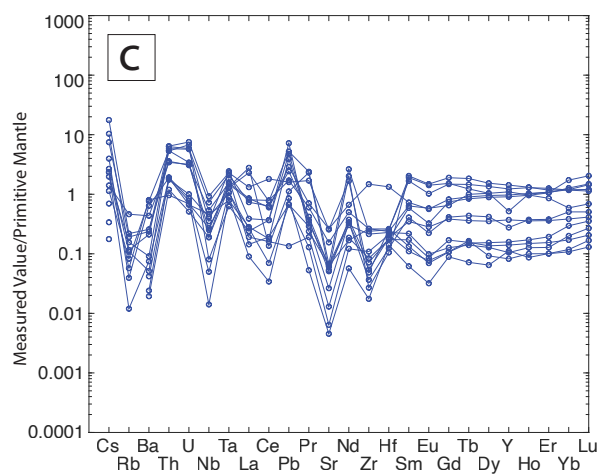
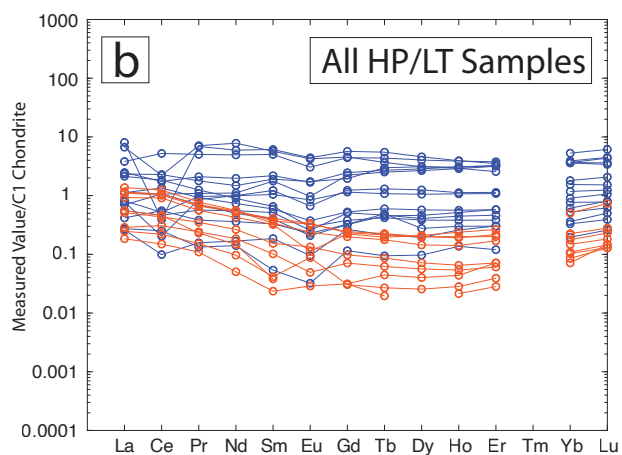
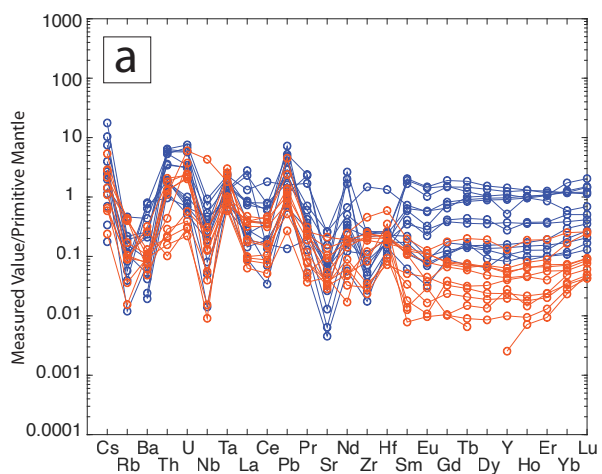


Figure 6.

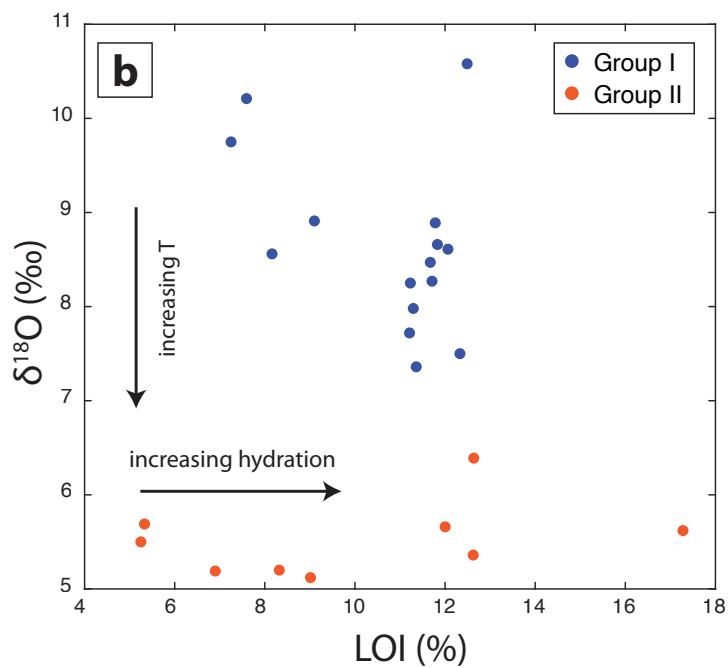
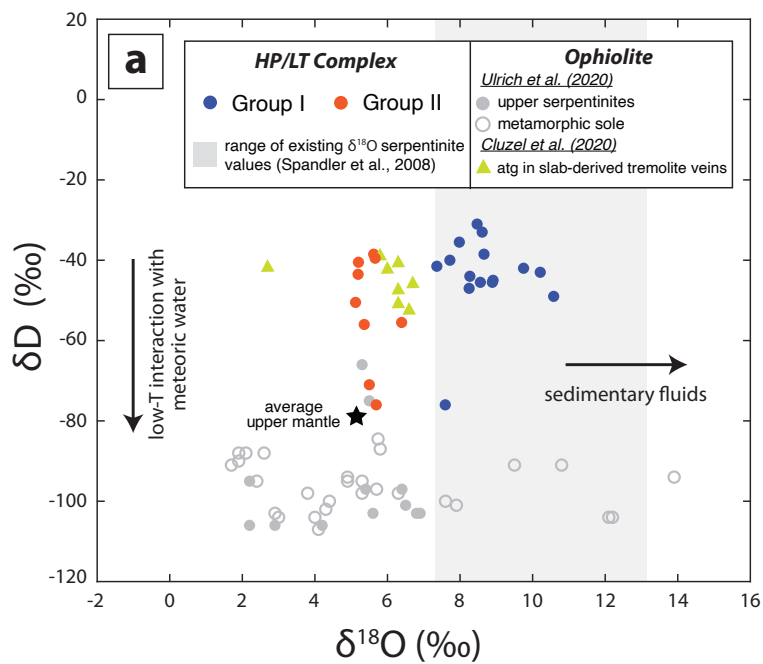


Figure 7.

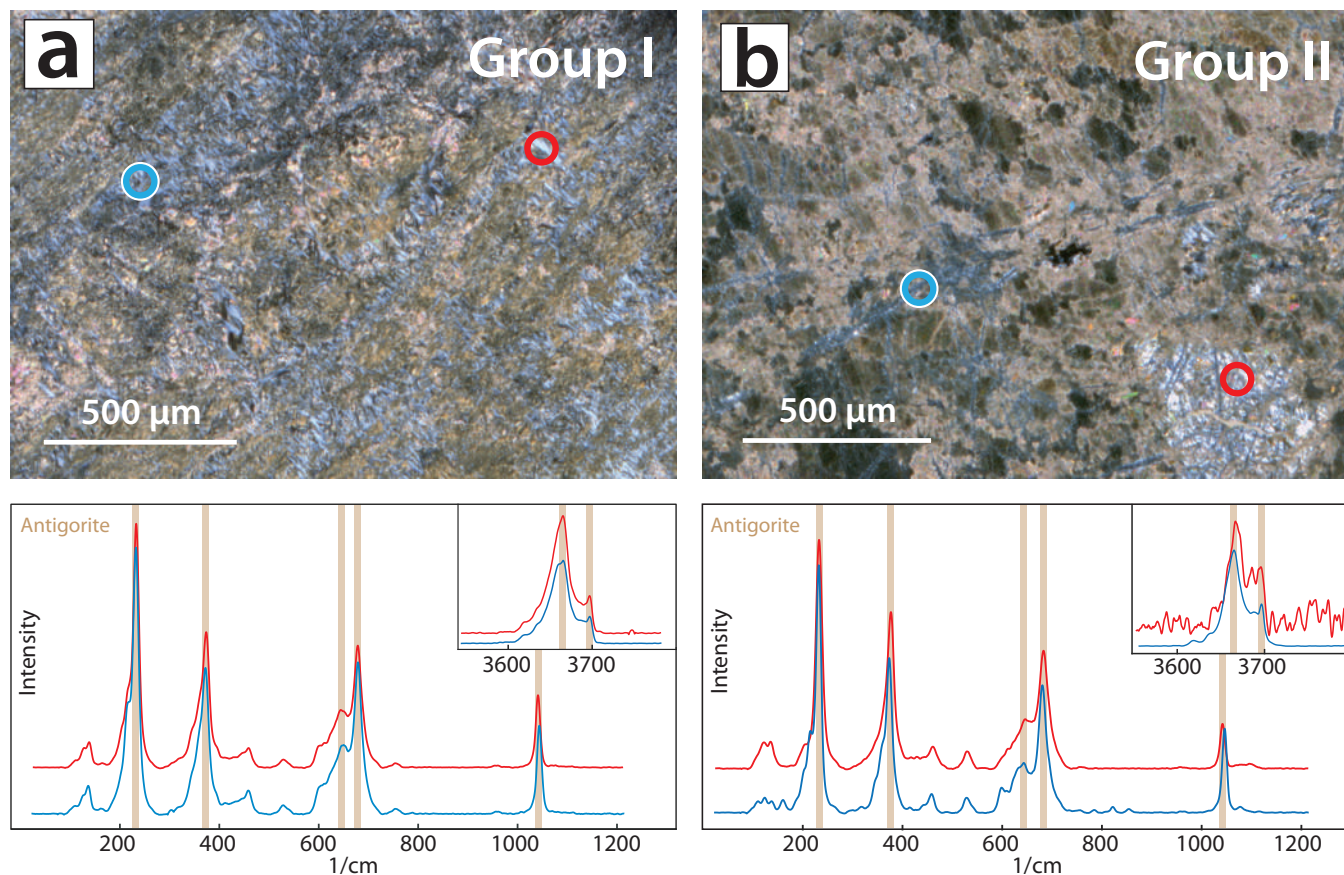


Figure 7. Representative cross-polarized photomicrographs and correlated Raman spectra for (a) two separate matrix sites in Group I sample NC18-09B and (b) one pseudomorph (red) and one vein (blue) in Group II sample NC19-152. Diagnostic Raman peaks for antigorite are highlighted tan vertical bars. All serpentine polymorphs in Group I and II were identified as antigorite (Figure S5).

Figure 8.

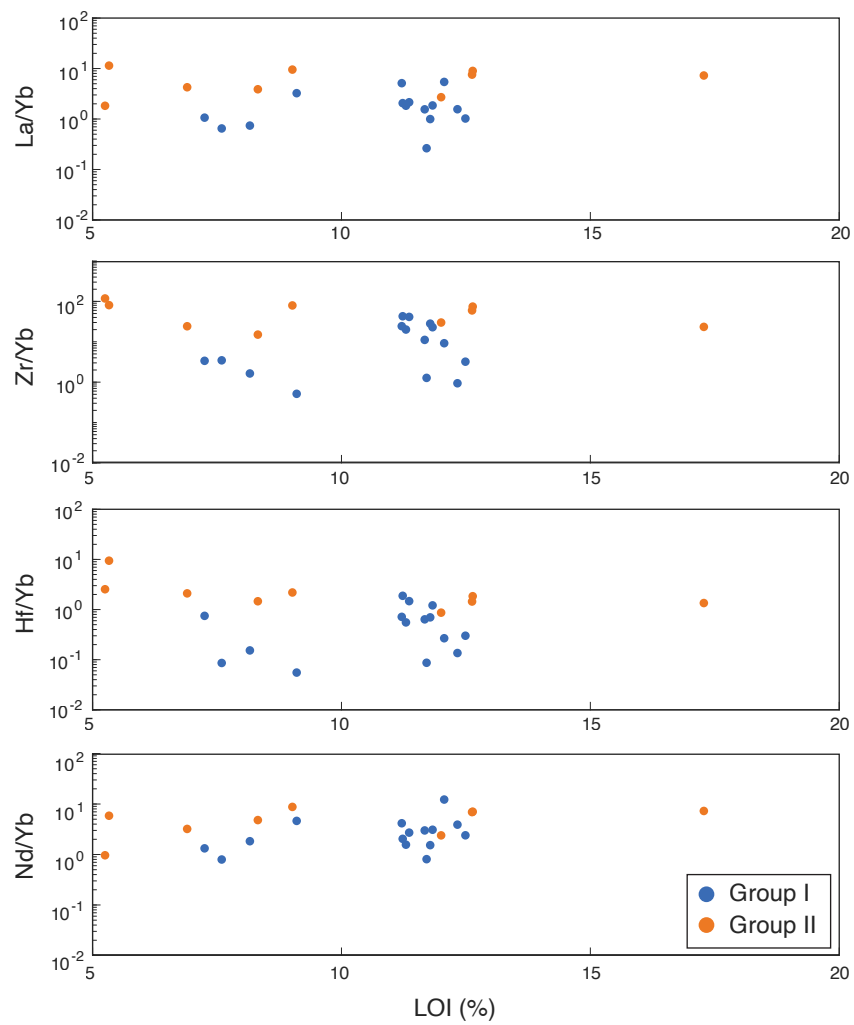


Figure 9.

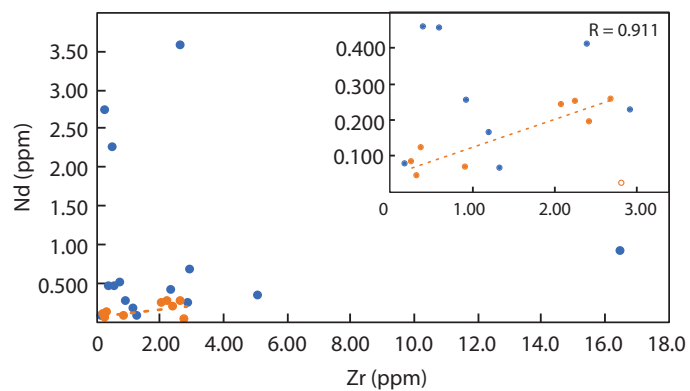
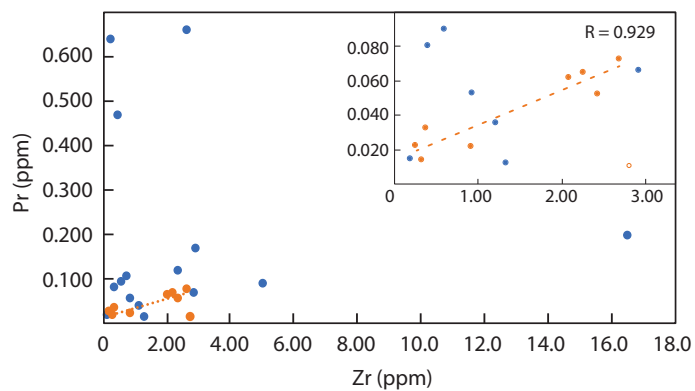
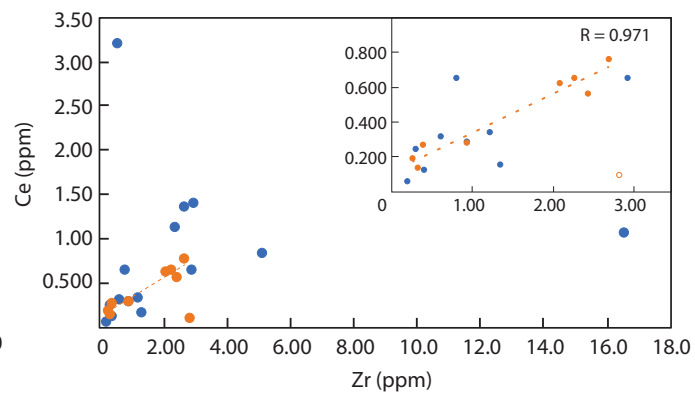
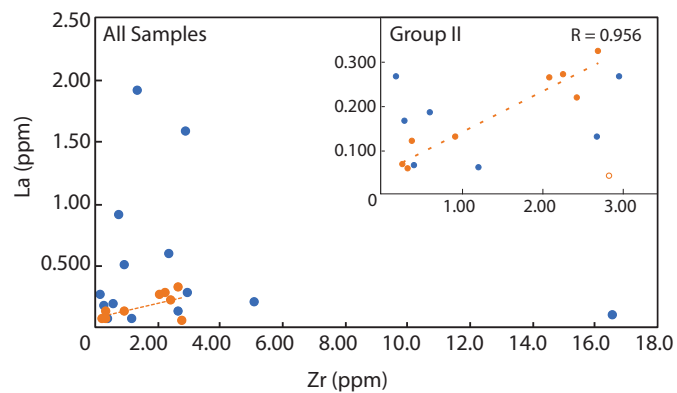


Figure 10.

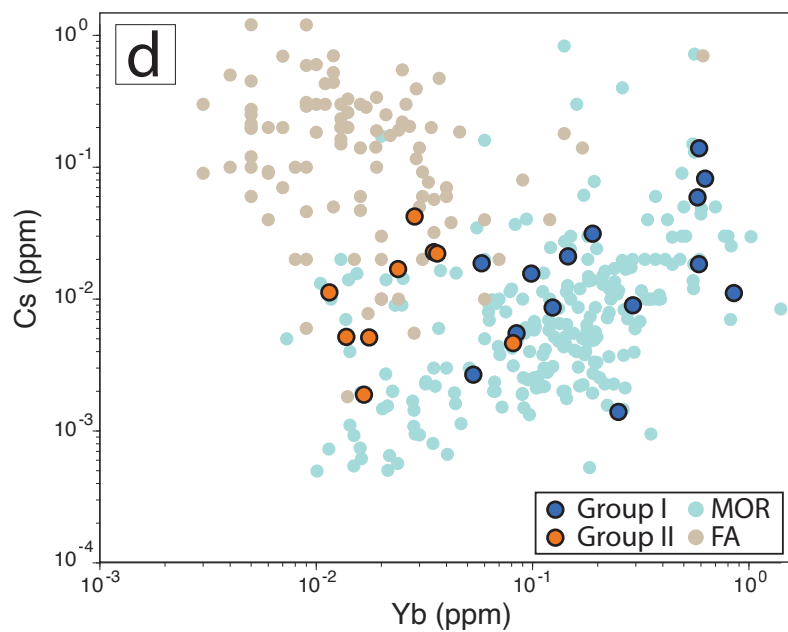
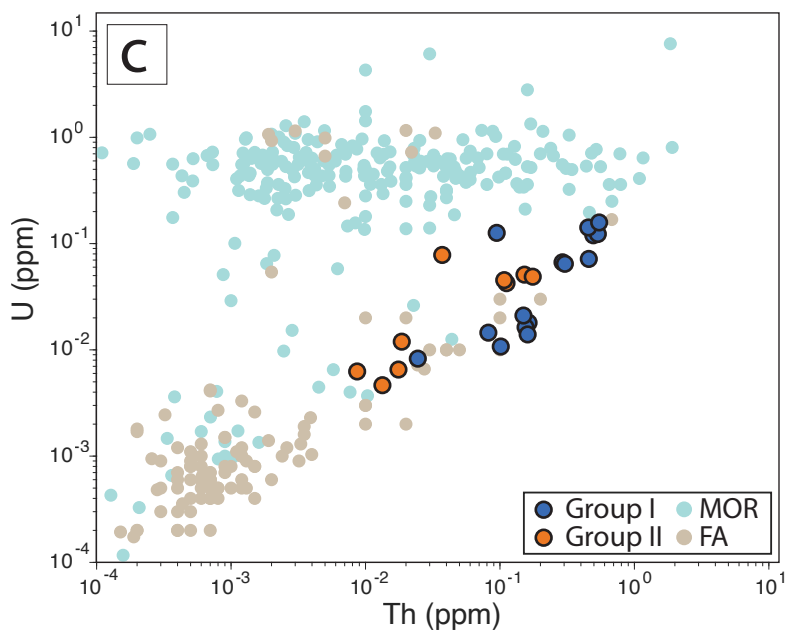
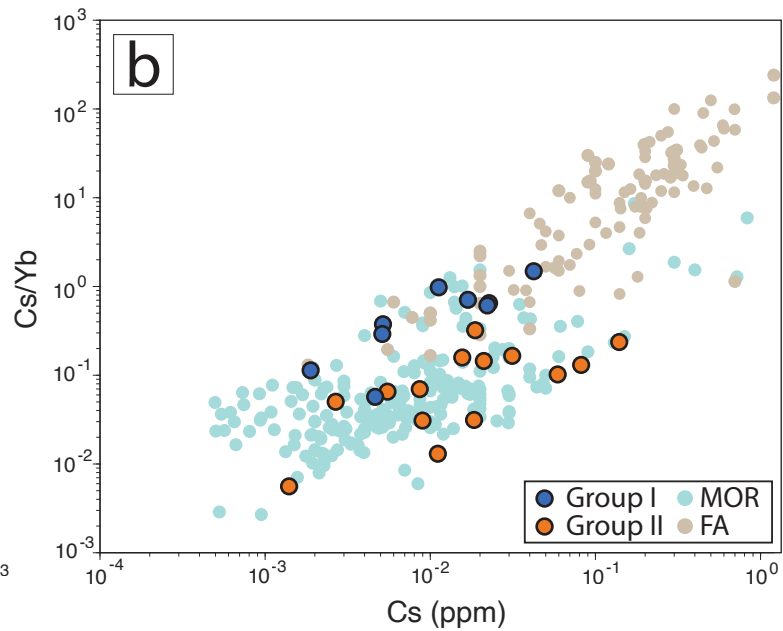
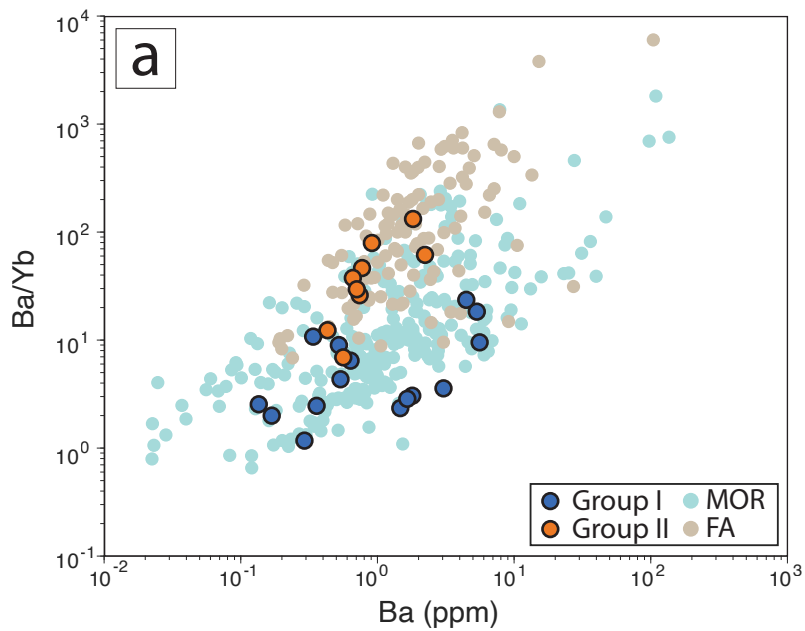


Figure 11.

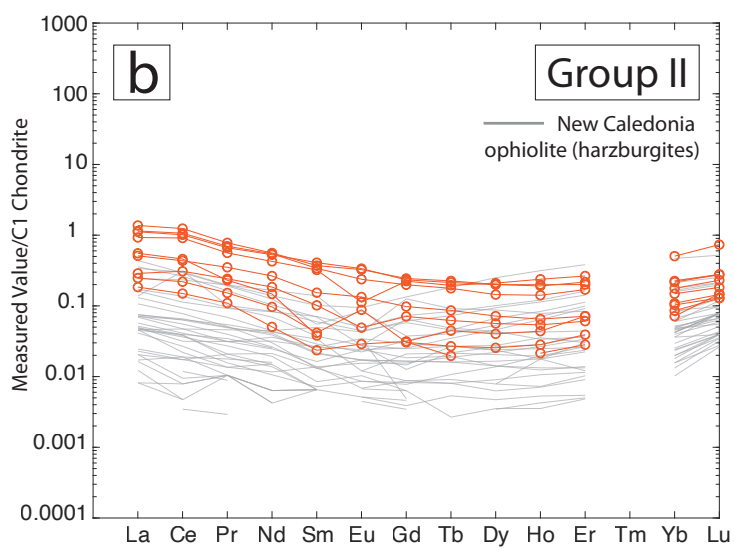
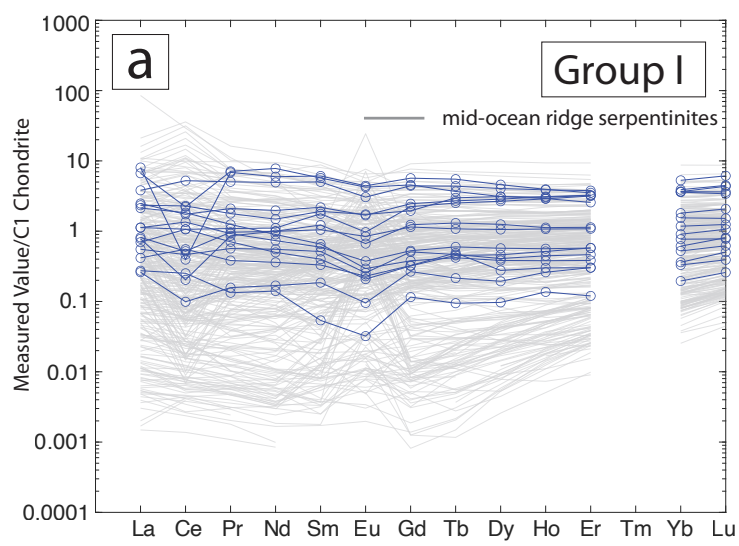


Table 1***Whole rock major and trace element compositions***

Lithology	Serpentinite				
Sample name	NC18-07C	NC18-09B	NC18-15C	NC18-15D	NC18-22A
IGSN	IENHR0002	IENHR0003	IENHR0004	IENHR0005	IENHR0006
Group/Locality	I	I	I	I	I
Major elements (wt%)					
SiO ₂	53.95	43.78	41.90	42.25	41.32
TiO ₂	0.02	0.02	0.02	0.02	0.02
Al ₂ O ₃	1.31	2.18	2.46	1.73	3.46
Fe ₂ O _{3(T)}	6.39	8.56	8.63	7.50	9.63
MnO	0.04	0.11	0.16	0.10	0.14
MgO	29.65	33.74	34.44	36.22	33.97
CaO	0.03	0.02	0.01	<0.01	0.01
Na ₂ O	0.00	0.00	0.00	<0.01	<0.01
K ₂ O	0.01	0.00	0.00	<0.01	<0.01
P ₂ O ₅	0.01	0.00	0.01	<0.01	<0.01
LOI ^a	7.59	11.30	12.06	11.67	11.21
Total	98.98	99.69	99.68	99.50	99.76
Mg#	0.82	0.80	0.80	0.83	0.78
Lab	WSU	WSU	WSU	WSU	WSU
Trace elements (ppm)					
Li	n.d.	n.d.	n.d.	0.0464	n.d.
Be	n.d.	n.d.	n.d.	0.110	n.d.
Sc	7.12	10.4	8.33	7.97	11.5
V	n.d.	n.d.	n.d.	37.3	n.d.
Cr	n.d.	n.d.	n.d.	2252	n.d.
Co	n.d.	n.d.	n.d.	94.8	n.d.
Ni	n.d.	n.d.	n.d.	2384	n.d.
Cu	n.d.	n.d.	n.d.	1.89	n.d.
Zn	n.d.	n.d.	n.d.	29.4	n.d.
Ga	n.d.	n.d.	n.d.	2.73	n.d.
Rb	0.293	0.0691	0.0525	bdl	0.0992
Sr	3.27	1.34	1.43	0.274	1.29
Y	2.36	0.720	4.99	0.637	0.478
Zr	2.95	2.93	2.68	0.938	2.40
Nb	0.180	0.251	0.136	0.0356	0.519
Cs	0.0111	0.0211	0.00898	0.00551	0.0156
Ba	3.04	0.359	5.33	0.168	0.636

La	0.554	0.267	1.58	0.131	0.506
Ce	1.40	0.653	1.36	0.285	1.12
Pr	0.165	0.0655	0.659	0.0526	0.116
Nd	0.677	0.228	3.57	0.253	0.410
Sm	0.283	0.0592	0.845	0.0761	0.0970
Eu	0.0977	0.0117	0.238	0.0144	0.0165
Tb	0.108	0.0163	0.133	0.0168	0.0151
Gd	0.388	0.0638	0.906	0.101	0.0755
Dy	0.765	0.113	0.765	0.100	0.0943
Ho	0.166	0.0281	0.159	0.0244	0.0208
Er	0.515	0.0925	0.411	0.0742	0.0614
Tm	0.102	0.0169	0.0533	0.0122	0.0117
Yb	0.852	0.146	0.291	0.0843	0.0986
Lu	0.151	0.0262	0.0509	0.0152	0.0199
Hf	0.0733	0.0808	0.0779	0.0538	0.0705
Ta	0.0605	0.0998	0.0528	0.0443	0.0864
Pb	0.121	0.512	0.371	0.00955	0.173
Th	0.292	0.495	0.458	0.163	0.534
U	0.0667	0.120	0.0717	0.0180	0.123
Lab	WSU	WSU	WSU	URI	WSU

Note . n.d. = not determined; bdl = below detection limits; F&M = Franklin & Marshall College; URI =
aLOI measured by combustion

NC18-26B IENHR0008 I	NC18-43 IENHR000C I	NC19-14 IENHR000G I	NC19-48 IENHR000I I	NC19-54 IENHR000J I	NC19-63 IENHR000K I	NC19-81 IENHR000L I
38.46	46.20	42.45	54.34	42.52	42.48	51.51
0.22	0.01	0.01	0.01	0.01	0.01	<0.01
6.73	1.68	1.96	1.18	0.84	2.68	2.48
13.39	7.00	6.50	6.30	7.09	8.29	6.83
0.14	0.07	0.10	0.07	0.09	0.12	0.07
29.19	32.94	37.65	30.61	38.67	34.64	30.92
0.44	0.11	0.01	0.02	0.01	0.02	0.14
<0.01	0.18	0.03	0.01	0.03	0.02	0.02
<0.01	0.02	<0.01	<0.01	<0.01	<0.01	<0.01
<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
11.78	12.33	11.83	7.25	11.36	12.49	8.16
100.35	100.54	100.54	99.79	100.62	100.75	100.13
0.69	0.82	0.85	0.83	0.85	0.81	0.82
WSU	F&M	F&M	F&M	F&M	F&M	F&M
n.d.	5.72	0.0154	0.907	1.75	2.08	0.820
n.d.	0.846	0.0985	0.155	0.126	0.236	0.0904
13.8	10.5	6.72	4.03	6.71	11.5	7.75
n.d.	43.0	27.8	25.2	33.7	41.2	45.4
n.d.	2400	2500	2193	2861	2280	2103
n.d.	88.8	91.9	88.0	169	69.6	82.1
n.d.	2209	2299	2368	5877	1942	2288
n.d.	4.09	4.61	2.98	17.1	24.6	2.12
n.d.	39.4	35.9	40.9	106	34.0	34.7
n.d.	4.51	2.00	1.98	1.83	3.18	3.31
0.139	0.122	bdl	0.00760	0.0662	0.0250	bdl
5.54	5.40	0.135	0.0957	0.602	1.39	0.558
4.41	5.59	0.374	0.507	0.595	1.69	1.26
16.6	0.543	1.22	0.196	5.11	0.609	0.409
0.384	0.661	0.134	0.0100	3.07	0.331	0.296
0.139	0.0591	0.00267	0.0187	0.00863	0.0313	0.00140
1.80	1.65	0.135	0.522	0.537	4.47	0.292

0.587	0.903	0.0988	0.0619	0.266	0.194	0.185
1.07	3.21	0.339	0.0607	0.827	0.317	0.124
0.195	0.468	0.0355	0.0146	0.0857	0.0896	0.0798
0.902	2.26	0.165	0.0770	0.336	0.455	0.457
0.323	0.744	0.0489	0.0274	0.0878	0.155	0.181
0.0947	0.172	0.0129	0.00537	0.0211	0.0475	0.0375
0.0975	0.158	0.0177	0.00776	0.0216	0.0390	0.0472
0.489	0.881	0.0646	0.0530	0.105	0.228	0.246
0.702	0.994	0.0680	0.0477	0.141	0.260	0.308
0.169	0.211	0.0164	0.0142	0.0307	0.0586	0.0614
0.520	0.604	0.0482	0.0485	0.0921	0.173	0.182
0.0849	0.0911	0.00814	0.00799	0.0169	0.0268	0.0323
0.589	0.579	0.0532	0.0581	0.124	0.189	0.249
0.107	0.0837	0.00963	0.0124	0.0191	0.0306	0.0374
0.411	0.0786	0.0645	0.0436	0.182	0.0570	0.0382
0.0639	0.0939	0.0745	0.0362	0.0744	0.0322	0.0434
0.204	0.114	0.0482	0.0606	0.169	0.278	0.0475
0.304	0.547	0.101	0.155	0.0946	0.161	0.149
0.0646	0.158	0.0107	0.0163	0.126	0.0140	0.0210
WSU	URI	URI	URI	URI	URI	URI

= University of Rhode Island; WSU = Washington State University.

NC19-86 IENHR000N I	NC19-89 IENHR000O I	NC19-94 IENHR000P I	NC18-39A (core) IENHR0009 II	NC18-39A (rxn) IENHR0009 II	NC18-39B IENHR000A II
41.69	50.75	42.12	38.57	40.64	40.05
0.02	0.01	0.01	0.01	0.01	0.01
0.95	1.93	2.81	0.51	0.52	0.57
7.15	7.10	7.50	7.49	7.29	7.59
0.05	0.09	0.13	0.11	0.12	0.11
39.25	31.28	36.47	40.34	38.42	41.55
0.03	0.02	0.06	<0.01	0.17	0.57
0.03	<0.01	0.03	<0.01	<0.01	<0.01
<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
0.01	<0.01	<0.01	<0.01	<0.01	<0.01
11.23	9.10	11.71	12.62	12.64	9.01
100.41	100.28	100.84	99.66	99.81	99.46
0.85	0.82	0.83	0.84	0.84	0.85
F&M	F&M	F&M	WSU	WSU	WSU
0.0470	0.602	1.60	n.d.	n.d.	n.d.
0.0267	0.124	0.326	n.d.	n.d.	n.d.
8.03	10.7	9.49	6.68	7.12	7.12
30.4	45.2	58.0	n.d.	n.d.	n.d.
2454	2747	2796	n.d.	n.d.	n.d.
52.5	93.9	110	n.d.	n.d.	n.d.
1601	2386	2678	n.d.	n.d.	n.d.
3.68	11.9	0.604	n.d.	n.d.	n.d.
30.0	42.3	40.8	n.d.	n.d.	n.d.
1.33	3.64	4.44	n.d.	n.d.	n.d.
bdl	0.0364	0.0736	0.264	0.0577	0.108
bdl	1.13	1.07	1.99	4.47	2.92
0.183	6.48	4.13	0.238	0.254	0.198
1.35	0.302	0.799	2.09	2.69	2.26
0.0397	0.194	0.0574	0.0947	0.122	0.0900
bdl	0.0184	0.0819	0.0227	0.0222	0.0423
0.339	5.59	1.47	0.431	2.23	0.741

0.0649	1.91	0.165	0.265	0.325	0.271
0.154	0.242	0.652	0.620	0.760	0.654
0.0124	0.637	0.101	0.0616	0.0723	0.0646
0.0643	2.72	0.507	0.242	0.256	0.250
0.00798	0.903	0.262	0.0605	0.0550	0.0503
0.00181	0.249	0.0545	0.0190	0.0183	0.0134
0.00341	0.198	0.0909	0.00773	0.00814	0.00639
0.0230	1.13	0.444	0.0464	0.0485	0.0396
0.0239	1.13	0.655	0.0491	0.0493	0.0355
0.00747	0.215	0.158	0.0109	0.0104	0.00772
0.0192	0.563	0.507	0.0320	0.0347	0.0272
0.00424	0.0848	0.0867	0.00475	0.00558	0.00478
0.0315	0.587	0.627	0.0350	0.0362	0.0285
0.00637	0.0891	0.110	0.00679	0.00687	0.00567
0.0592	0.0325	0.0544	0.0505	0.0669	0.0624
0.0377	0.0546	0.0257	0.0356	0.0416	0.0308
0.0841	0.0793	0.340	0.0981	0.0745	0.320
0.0245	0.0820	0.454	0.112	0.152	0.108
0.00827	0.0145	0.142	0.0424	0.0510	0.0453
URI	URI	URI	WSU	WSU	WSU

NC18-39C IENHR000B II	NC19-42 IENHR000H II	NC19-152 IENHR000Q II	NC19-154A IENHR000R II	NC19-157 IENHR000S II	NC19-158 IENHR000T II	NC18-45 IENHR000D Boghen Terrane
41.61	41.96	40.80	38.86	40.64	41.98	40.26
0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.01
0.75	0.72	0.69	0.72	0.68	0.67	0.41
7.21	7.75	8.28	5.75	7.55	7.91	9.41
0.09	0.11	0.12	0.09	0.11	0.11	0.15
38.01	43.46	42.29	39.31	42.35	43.24	36.43
0.01	0.50	1.08	0.02	0.42	0.53	0.01
<0.01	0.06	0.05	0.04	0.05	0.08	<0.01
<0.01	0.01	<0.01	<0.01	<0.01	0.01	<0.01
<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
12.00	5.33	6.90	17.28	8.32	5.25	13.19
99.70	99.90	100.21	102.07	100.12	99.78	99.86
0.84	0.85	0.84	0.87	0.85	0.85	0.79
WSU	F&M	F&M	F&M	F&M	F&M	WSU
n.d.	0.717	0.481	0.0560	0.180	0.630	n.d.
n.d.	bdl	bdl	0.0160	bdl	bdl	n.d.
7.20	8.75	7.01	6.74	6.01	5.39	9.79
n.d.	21.4	12.1	17.8	19.2	17.6	n.d.
n.d.	2672	1773	2285	2117	1819	n.d.
n.d.	104	97.8	98.4	103	112	n.d.
n.d.	2331	2184	2461	2278	2555	n.d.
n.d.	10.2	2.16	2.36	3.21	2.04	n.d.
n.d.	30.1	33.2	29.3	23.3	29.7	n.d.
n.d.	0.977	1.76	0.933	0.878	0.463	n.d.
0.0598	0.254	0.0568	0.00982	0.0229	0.276	0.0814
1.13	0.670	2.20	0.724	0.788	0.934	2.19
0.289	0.0115	0.0454	0.108	0.0895	0.126	0.197
2.43	0.929	0.334	0.388	0.264	2.82	2.06
0.219	0.00648	bdl	0.0284	0.0109	0.185	0.0892
0.00464	0.0112	0.00517	0.00189	0.00512	0.0169	0.0921
0.562	0.911	1.82	0.772	0.661	0.706	3.88

0.220	0.131	0.0586	0.121	0.0682	0.0437	0.281
0.559	0.282	0.135	0.266	0.189	0.0916	0.556
0.0520	0.0214	0.0141	0.0326	0.0224	0.0102	0.0546
0.194	0.0674	0.0442	0.121	0.0844	0.0231	0.212
0.0477	0.00564	0.00616	0.0227	0.0151	0.00348	0.0367
0.00634	bdl	0.00495	0.00752	0.00277	0.00163	0.0772
0.00705	0.000708	0.000971	0.00312	0.00225	0.00161	0.00446
0.0444	0.00616	0.00608	0.0195	0.0141	0.00625	0.0273
0.0517	bdl	0.00629	0.0176	0.0139	0.00989	0.0308
0.0130	0.00117	0.00154	0.00355	0.00294	0.00239	0.00704
0.0423	0.00451	0.00627	0.0114	0.00971	0.0115	0.0230
0.00821	0.00128	0.00118	0.00187	0.00176	0.00219	0.00417
0.0813	0.0115	0.0138	0.0166	0.0176	0.0239	0.0300
0.0181	0.00363	0.00315	0.00319	0.00357	0.00449	0.00509
0.0705	0.109	0.0290	0.0224	0.0257	0.0607	0.0558
0.0403	0.105	0.0240	0.123	0.0259	0.0952	0.0311
0.0555	0.0684	0.0932	0.0369	0.0576	0.0191	0.860
0.175	0.0176	0.00867	0.0372	0.0134	0.0186	0.123
0.0488	0.00653	0.00626	0.0782	0.00465	0.0119	0.0374
WSU	URI	URI	URI	URI	URI	WSU

NC18-46	NC18-49	Peridotite NC19-178	Chlorite schist NC19-85	Talc schist NC19-169	
IENHR000E	IENHR000F	IENHR000V	IENHR000M	IENHR000U	
Boghen Terrane	Serpentinite Sole	Kalaa-Gomen	I	I	
39.00	42.04	40.51	22.35	54.16	
0.02	<0.01	<0.01	3.54	0.02	
1.05	0.51	0.69	13.43	2.99	
10.22	6.91	7.73	29.21	6.62	
0.15	0.06	0.11	0.24	0.02	
36.38	37.51	40.91	19.23	29.93	
<0.01	0.04	0.66	2.47	0.03	
<0.01	0.03	0.05	0.01	<0.01	
<0.01	<0.01	<0.01	<0.01	<0.01	
0.01	<0.01	<0.01	1.88	0.01	
12.90	13.67	9.77	7.88	6.25	
99.72	100.77	100.43	100.24	100.03	
0.78	0.84	0.84	0.40	0.82	
WSU	WSU	F&M	F&M	F&M	
n.d.	1.91	1.27	7.12	0.162	
n.d.	0.0109	bdl	0.252	0.0814	
11.8	5.60	7.30	35.9	8.51	
n.d.	19.7	26.6	448	49.0	
n.d.	2314	2307	284	2336	
n.d.	95.5	95.7	140	93.1	
n.d.	2563	2069	247	1985	
n.d.	2.55	1.82	59.7	7.40	
n.d.	25.3	82.9	128	43.3	
n.d.	0.645	0.879	8.47	4.28	
0.144	0.00673	0.0809	0.0242	0.167	
1.74	0.677	0.428	36.0	2.27	
0.453	1.08	0.0220	54.8	0.327	
3.12	0.497	0.353	25.0	0.256	
0.155	0.00694	bdl	12.7	0.0267	
0.787	0.00578	0.00336	0.00499	0.0109	
3.50	0.709	0.168	12.9	0.341	

0.296	0.202	0.0183	13.1	0.0523
0.600	0.170	0.0501	35.7	0.192
0.0760	0.0701	0.00444	4.29	0.0302
0.350	0.330	0.0101	20.5	0.156
0.0475	0.100	0.00109	6.42	0.0523
0.0220	0.0264	bdl	1.82	0.00663
0.00897	0.0199	0.000194	1.76	0.0118
0.0493	0.143	0.00120	9.71	0.0693
0.0652	0.110	0.000929	11.4	0.0706
0.0175	0.0273	0.000413	2.30	0.0141
0.0556	0.0752	0.00259	5.50	0.0355
0.00961	0.00983	0.000758	0.662	0.00444
0.0711	0.0544	0.00955	3.44	0.0237
0.0128	0.0109	0.00254	0.453	0.00355
0.0668	0.0437	0.0224	0.738	0.0105
0.0891	0.0525	0.0217	0.641	0.0135
0.362	0.260	0.00374	0.984	0.0827
0.0939	0.0165	0.00392	3.08	0.0223
0.0368	0.00565	0.00129	0.133	0.00457
WSU	URI	URI	URI	URI

Table 2
Oxygen and hydrogen stable isotope data.

Sample	Rock type	Group/Locality	δD_{srp} (‰)	$\delta^{18}O_{\text{srp}}$ (‰)
NC18-07C	serpentinite	I	-45, -41	10.2
NC18-09B	serpentinite	I	-35, -36	8.0
NC18-15C	serpentinite	I	-34, -32	8.6
NC18-15D	serpentinite	I	-33, -29	8.5, 6.7
NC18-22A	serpentinite	I	-40, -40	7.7, 6.8
NC18-26B	serpentinite	I	-44, -46	8.9
NC18-43	serpentinite	I	-75, -77	7.5
NC19-14	serpentinite	I	-38, -39	8.7, 8.8
NC19-48	serpentinite	I	-42, -42	9.8
NC19-54	serpentinite	I	-42, -41	7.4, 7.2
NC19-63	serpentinite	I	-49, -49	10.6
NC19-81	serpentinite	I	-47, -44	8.6
NC19-86	serpentinite	I	-47, -47	8.3
NC19-89	serpentinite	I	-45, -45	8.9, 8.8
NC19-94	serpentinite	I	-44, -44	8.3
NC18-39A (core)	serpentinite	II	-59, -53	5.4
NC18-39A (rxn)	serpentinite	II	-56, -55	6.4
NC18-39B	serpentinite	II	-52, -49	5.1, 5.5
NC18-39C	serpentinite	II	-39, -40	5.7
NC19-42	serpentinite	II	-73, -79	5.7
NC19-152	serpentinite	II	-43, -44	5.2
NC19-154A	serpentinite	II	-40, -37	5.6
NC19-157	serpentinite	II	-42, -39	5.2
NC19-158	serpentinite	II	-70, -72	5.5
NC19-85	chlorite schist	I		
NC19-169	talc schist	I		
NC18-45	serpentinite	Boghen Terrane	-82, -86	8.0
NC18-46	serpentinite	Boghen Terrane	-78, -79	7.0
NC18-49	serpentinite	Serpentinite Sole	-82, -82	6.5
NC19-178	serpentinite	Kalaa-Gomen Massif	-84, -85	5.7

Note. Mineral abbreviations after Whitney and Evans (2010): *Srp* serpentine, *Mag* mag

¹ >710 μm grain size fraction

² 125-250 μm grain size fraction

$\delta^{18}\text{O}_{\text{mag}}$ (‰)

$\delta^{18}\text{O}_{\text{tlc}}$ (‰)

9.0

1.5

2.3, 2.5
4.3¹, 4.6²

10.5

agnetite, *Tlc* talc