

Melt inclusions at MT. Edixon (Antarctica): Chemistry, petrology and implications for the evolution of the Lanterman range

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Fabio Ferri, Bernardo Cesare, Omar Bartoli, Silvio Ferrero, Rosaria Palmeri, et al.. Melt inclusions at MT. Edixon (Antarctica): Chemistry, petrology and implications for the evolution of the Lanterman range. Lithos, 2020, 374-375, pp.105685. 10.1016/j.lithos.2020.105685. hal-02959290

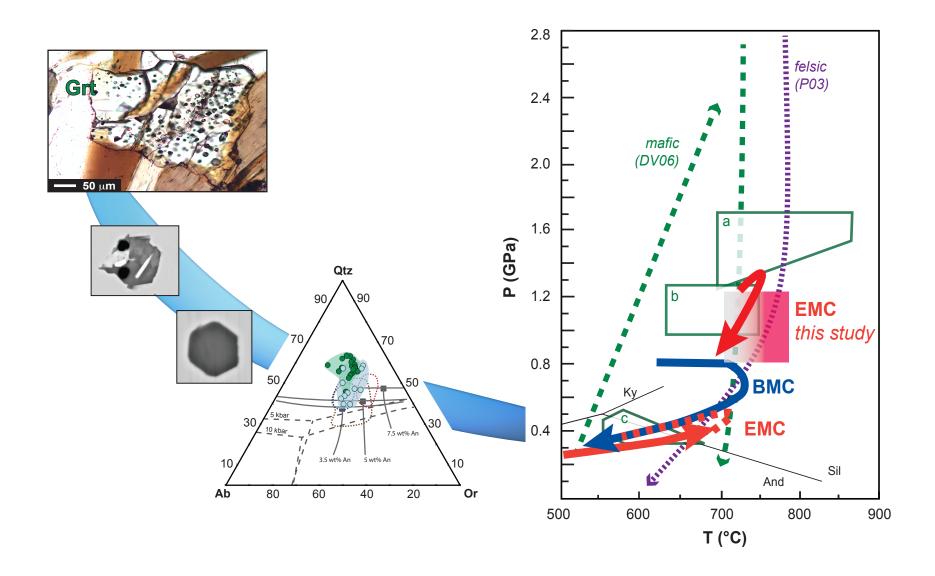
HAL Id: hal-02959290 https://hal.science/hal-02959290

Submitted on 24 Dec 2020 $\,$

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3 4 5 6	2	CHEMISTRY, PETROLOGY AND IMPLICATIONS FOR THE
6 7 8 9	3	EVOLUTION OF THE LANTERMAN RANGE
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- Melt inclusions first discovered in peritectic garnets in paragneiss from the Edixon Metamorphic Complex, Lanterman Range, Antarctica
- Primary glasses from remelting of inclusions are sub-alkaline and peraluminous with 4-5 wt. % H_2O and $CO_2 < 0.1$ wt. %
- Anatexis at the Edixon Metamorphic Complex occurred at different crustal levels



ABSTRACT

The recovery of melt inclusions in peritectic phases of metamorphic rocks brings the opportunity to retrieve the chemical composition of primary melts and infer melt entrapment conditions. Here we report the first recovery of such inclusions in a medium-grade muscovite-biotite-plagioclase-K-feldspar-quartz gneiss from the Edixon Metamorphic Complex (EMC) of the Lanterman Range in Northern Victoria Land, Antarctica.

The most common inclusions are polycrystalline (PI) and fluid (FI) types coexisting in the same clusters and displaying frequent negative crystal shape and textures indicative of a primary origin. The PI contain muscovite + quartz + $H_2O \pm biotite \pm calcite \pm siderite \pm albite$ while the FI are composed by $CO_2 + H_2O + N_2 + CH_4 \pm H_2S$. The PI were remelted in a piston cylinder apparatus at conditions of 740-900°C at 0.8-1.0 GPa with complete remelting taking place at 760-780°C. The remelted glass is sub-alkaline, peraluminous (ASI \approx 1.3-1.5) with CaO \approx 1.2 - 2.1 wt% and K₂O \approx 3.4 - 4.1 wt%, and contains \approx 4.3-5.3 wt% of H₂O and 400-1150 ppm of CO₂.

The stable mineral assemblage is represented by muscovite + plagioclase + quartz + K-feldspar + garnet + melt ± rutile coexisting with excess C-O-H fluid while biotite is retrograde (Ti 1.2 a.p.f.u.). The calculated phase equilibria and compositional isopleths in the MnCNKFMASHT model system delimit the equilibrium conditions to 1.0-1.3 GPa and 680-730°C.

Our study confirms that the EMC at the Lanterman Range is not a homogeneous metamorphic complex and that anatexis could have taken place at different crustal levels during the evolution of the Ross Orogen. Our results indeed suggest that, in the Lanterman Range, the high-pressure eclogitic rocks, the medium pressure amphibolite stage and the low-pressure amphibolite stage shared most of the late exhumation path.

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4 5	50	Key words: melt inclusions, crustal anatexis, nanogranitoids, Lanterman Range,
6 7	51	Edixon Metamorphic Complex
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1. INTRODUCTION

The discovery of primary melt inclusions (MI) hosted in peritectic minerals from high grade metamorphic rocks has been described in recent papers (reviewed by Cesare et al., 2015 and Bartoli et al., 2016) and new examples are continuously adding from typical felsic crustal compositions or new mafic to ultramafic ones (Ferrero et al. 2018). The MI represent a window into the anatectic history of partially melted terranes (Acosta-Vigil et al., 2010) and may provide key microstructural and compositional information on crustal anatexis, like mechanisms and nature of the melting process, conditions of entrapment and melting, timeframes of melt production and segregation (Cesare et al., 2015). In the absence of any petrographical evidence of anatexis as pseudomorphs after melt films (Holness et al., 2011), the MI become a unique tool to infer past melting events throughout the metamorphic history of a rock. When MI coexist with fluid inclusions (FI), they are able to shed light on the fluid/melt/rock interactions during anatexis, and recent works demonstrated that melting processes are more commonly fluid-present than traditionally expected (Carvalho et al., 2018). The combination of MI and FI studies can also help understanding the exchange, transport and recycling of volatile species (especially CO₂) in crustal evolution or during subduction.

In Antarctica, the presence of felsic-nanogranite inclusions produced after cooling of MI have been documented in garnets from the Lutzow-Holm Complex (East Antarctica, Hiroi et al., 2019). Here we report the first recovery of preserved MI in metapelitic garnets from the Edixon Metamorphic Complex from the Lanterman range in Antarctica. These inclusions turn out to be an essential tool to identify anatexis in rocks lacking further evidence of partial melting and, together with the petrological study of their host rocks, provide new clues to the geodynamic evolution of this crystalline basement.

2. SAMPLE DESCRIPTION

2.1 Geological setting

The Ross orogenic belt is one of the Earth's great Phanerozoic mountain belts resulting from convergence along the paleo-pacific margin of Antarctica in Gondwana land (Cawood, 2005; Talarico et al., 2004). The Northern Victoria Land (Fig. 1A, B) represents one of the major segments of the Ross orogenic belt (e.g., Federico et al., 2006, see summary by Stump, 1995) and comprises the Lanterman-Mariner suture (Ricci and Tessensohn, 2003, and references therein), a tectonic structure where peculiar HP-UHP metamorphic rocks have been reported (Ricci et al., 1996; Palmeri et al., 2011).

From east to west, the Northern Victoria Land can be divided into three contrasting tectonometamorphic terranes with NNW-SSE structural trends separated by major faults (Bradshaw and Laird, 1983): 1) the Robertson Bay Terrane, 2) the Bowers Terrane and 3) the Wilson Terrane (Fig. 1B). The Wilson Terrane represents the palaeo-Pacific margin of Gondwana in the region and consists of intermediate- to high-P metamorphic rocks with local eclogites close to the Lanterman-Mariner suture, (Grew et al., 1984; Di Vincenzo et al., 1997; Ricci et al., 1997), and of low-P metamorphic rocks (Palmeri et al., 1991) with local remnants of a polymetamorphic granulite complex (Talarico et al., 1995). The Wilson Terrane was extensively intruded by the Granite Harbour Intrusive Complex, an orogenic association of Cambro-Ordovician granitoids showing arc affinity (Bomparola et al., 2006; Kleinschmidt and Tessensohn, 1987; Borg et al., 1990; Ghezzo et al., 1989).

The Lanterman-Mariner suture separates the Wilson Terrane and the Bowers Terrane and is best exposed in two areas, namely the Lanterman Range to the north and the Mountaineer Range to the south. From west to east, three different lithotectonic units have been identified in the Lanterman Range area of the Lanterman-Mariner suture (Talarico et al., 1998): i) the Edixon Metamorphic Complex, ii) the Bernstein Metamorphic Complex and iii)

the Gateway Hills Metamorphic Complex (Fig. 1C). The Edixon Metamorphic Complex mainly consists of micaschists and gneisses containing bands and lenses of calc-silicate rocks, with minor amphibolites in the southernmost outcrops of the range. This complex underwent medium- to high-grade metamorphism and partial melting under low-P conditions and shows close similarities to the internal sequences of the Wilson Terrane, with the presence of migmatites and abundant granitoids of the Granite Harbour Intrusive Complex. The Bernstein Metamorphic Complex consists of micaschists and gneisses affected by medium- to highgrade metamorphism under intermediate-P conditions. In contrast to the Edixon Metamorphic Complex, calc-silicate rocks, migmatites and granitoids are rare (Ghiribelli, 2000; Talarico et al., 1998). The Gateway Hills Metamorphic Complex is characterized by abundant mafic and ultramafic rocks, including well-preserved eclogite-facies rocks, with minor gneisses and quartzites (Di Vincenzo et al., 1997; Ghiribelli et al., 2002; Palmeri et al., 2003, 2007, 2009, 2011). There are no granitoids in this complex (Talarico et al., 1998). Recently Di Vincenzo et al. (2016) established that two eclogite types coexist along the same ridge at the Lanterman Range: i) medium-grained, "colder" and undeformed eclogites, that recorded both prograde and peak conditions at nearly 530 Ma (U-Pb zircon age) during ongoing regional calc-alkaline magmatism, and ii) finer-grained, "hotter" and slightly deformed eclogites, that recorded only metamorphic peak conditions at nearly 500 Ma and hints of the subsequent retrogression (\approx 480 Ma), coincident with the outboard accretion of an island arc (Bowers Terrane) onto the Gondwanan margin.

The evolution of the Lanterman Range (Capponi et al., 1999, 2002; Crispini et al., 2007; Di Vincenzo et al., 2014, 2016; Federico et al., 2010) can be summarized in a westover-east thrusting at ca. 500 Ma, followed by sinistral strike-slip shearing at 480–460 Ma ago, both under amphibolite to greenschist-facies metamorphism. Large-amplitude folding of

the late Ross or Lachlan orogens in the late Ordovician and Cenozoic brittle tectonics terminated the evolution of the suture.

2.2 Petrography and bulk composition

The studied rock belongs to the Edixon Metamorphic Complex of the Lanterman Range, was sampled at SE of Mt. Edixon (71° 50'23.597"S, 163°39'41.35"E) and stored in the PNRA (Programma Nazionale Ricerche in Antartide) rock repository at the Museo Nazionale dell'Antartide (Siena, Italy) as specimen 24-12-94GO5 (from hereafter GO5). The rock is a fine-grained psammitic gneiss composed of biotite, muscovite, garnet, quartz, plagioclase, K-feldspar, minor ilmenite, apatite and tourmaline, and free of Al₂SiO₅ polymorphs (Fig. 2). Muscovite and biotite define the main foliation and show mm-sized crystals with no evidence of alteration. Quartz is abundant and occurs in coarse crystals or tiny aggregates of subgrains with segmented borders showing evidence of bulging. Garnet is scarce, with average diameter $< 400 \,\mu\text{m}$ and exceptionally up to $\approx 1 \,\text{mm}$. The sample displays evidence for extensive down-temperature replacement of garnet by biotite (Figure 2B, C). Overall, the mineralogical composition and texture of the studied rock would suggest a medium grade metapsammite. Typical microstructural evidence of melt presence such as grains with euhedral crystal faces or pseudomorphs after melt films (e.g., Holness et al., 2011) are not present, and only the occurrence of melt inclusionsin garnet (Fig. 2B, C) attests for the former anatectic history of the rock.

3. METHODS

3.1 Analytical techniques

After MI recognition by optical microscopy, the mineral phases and the inclusions were first characterized in detail on polished thin sections by back-scattered electron images (BSEs) obtained with FESEM Sigma Zeiss at the ICMATE - CNR of Padova and with SEM CamScan MX3000 at the Department of Geoscience of the University of Padova.

The mineral phases and the glass of the inclusions were investigated by Micro-Raman spectroscopy using the HORIBA Jobin Yvon LabRAM HR 800 located at the Institute of Earth and Environmental Science, University of Potsdam. The machine is equipped with a Peltier-cooled multichannel CCD detector and coupled with an Olympus BX41 petrographic microscope. An air-cooled NdYAG laser was used for excitation ($\lambda = 532$ nm, laser power on the sample of 2–3 mW) with a grating of 300 lines/mm, slit width and confocal hole set to 100 and 200 µm, respectively. The Raman spectra of the solid and fluid phases were acquired with a 100× objective in the wavenumber range 100–4000 cm⁻¹, integrating 3 repetitions of 60 s with spectral resolution of 10 cm⁻¹.

The composition of mineral phases and glasses was determined using a Jeol JXA 8200 Superprobe at the Dipartimento di Scienze della Terra, Università di Milano (Italy) using natural and synthetic silicates and oxides as standards. Analytical parameters for minerals were 15 kV accelerating voltage, 5 nA current, counting time of 30 s on peak and 10 s on background. For glass analyses, the beam diameter was reduced to 1 μ m and the counting time to 10 s on peak and 2 s on background. Sodium, K, Al, and Si were the first analyzed elements to limit element loss. Concentrations were corrected against leucogranitic glass standards with comparable H₂O contents (Morgan and London, 2005). Details concerning the application of correction factors and the composition of the standard glasses are given by Ferrero et al. (2012) and Bartoli et al. (2013a, 2013b).

169 The analysis of H and C in glasses was performed using the Cameca Nano Secondary 170 Ion Mass Spectrometry 50 (NanoSIMS) installed at Muséum National d'Histoire Naturelle in 171 Paris (France). Polished experimental capsules with MI exposed on the garnet surface and 172 standard glasses were mounted in Indium (Aubaud et al., 2007). MI were identified by 173 collecting secondary ion images of Si, K and Fe. For every analysis, we first performed a pre-174 sputtering step on a $3\times3 \text{ }\mu\text{m}^2$ surface area for 2 min with a 400 pA primary Cs⁺ beam to

remove gold coating, surface contamination and reach a steady-state sputtering regime. Then a primary beam of 37 pA was used for data acquisition. Data were acquired by rastering a $3 \times$ $3 \,\mu\text{m}^2$ surface area and collecting only ions from the inner $1 \times 1 \,\mu\text{m}^2$ (beam blanking mode) to reduce surface contamination. Each analysis stacked about 200 cycles, each cycle being 1.024 s long. The ⁶OH⁻ (used as a proxy for H₂O), ²⁸Si⁻, ³⁹K¹⁶O⁻ and ⁵⁶Fe¹⁶O⁻ were recorded simultaneously in multicollection mode. We checked that ${}^{16}\text{OH}^{-/28}\text{Si}^{-}$ ratio was stable during MI analyses (see Supplementary material). Secondary ions were collected by electron multipliers with a dead time of 44 ns. Mass resolution was set to 10000. One inclusion was large enough for replicated analyses. For NanoSIMS calibration we used a 5.5 wt% H₂O-bearing leucogranitic glass from Acosta-Vigil et al. (2003), a 4.3 wt% H₂O- bearing leucogranitic glass from Behrens and Jantos (2001) and an anhydrous leucogranitic glass from Morgan and London (2005). The standards used for the determination of CO₂ contents were four trachyandesitic standards (STR 9, 10, 11 and 13) from the Stromboli volcano experimentally doped in carbon and water by Bureau et al. (2003). H and C concentrations were recalculated as H₂O and CO₂, respectively. Data corrections, using the aforementioned calibration, and error calculations were performed using the R program following the procedure described in Bartoli et al. (2014) and Thomen et al. (2014). During the session, the vacuum in the analysis chamber remained between 2.5 and 5×10^{-10} Torr.

3.2 Mineral chemistry

The mean compositions of minerals are reported in Table 1 (Supplementary Material). Garnet is almandine-rich (\approx 57 mol. % Alm) with slight growth zoning. From core to rim, the pyrope content decreases from 12 mol. % to 8-10 mol. % while grossular and spessartine components increase from 2 mol. % to 4 mol. %, and from 25 mol. % to 28-30 mol. % respectively. The almandine content is constant throughout the crystal. 2 200

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 The white mica is potassic ($X_{Ms} = 0.92-0.95$) with Si ranging from 3.00 to 3.04 a.p.f.u. (11 oxygens), Al ranging from 2.76 to 2.72 a.p.f.u. and X_{Mg} of 0.41-0.45. The TiO₂ content (0.03 a.p.f.u.) is typical of amphibolite-facies subsolidus muscovite (e.g., Cesare 1999, compare with Redler et al., 2011). The composition of the white mica is similar to that reported by Di Vincenzo et al. (2001) in sample BT5 and BT2 collected from the Edixon and Bernstein Metamorphic Complexes respectively and classified by the Authors as type-2 muscovite with selected area diffraction (SAED) pattern of 2M1 polytypes.

Biotite shows no systematic variations with microstructural occurrence, and has a fairly constant composition, with Ti of 0.11-0.13 a.p.f.u. and X_{Mg} of 0.44-0.45. Also for biotite the titanium content is typical of amphibolite-facies subsolidus conditions (Tajčmanová et al., 2009. Plagioclase composition has 15-23 mol. % anorthite, 76-84 mol.% albite and ca. 1 mol.% K-feldspar. Alkali feldspar is almost pure K-feldspar with albite <1-3 mol.%.

3.3 Remelting of inclusions

The following nomenclature will be used: 'remelted' inclusions are those in which the crystalline assemblage was completely melted to a single glassy phase, but melt + vapor homogenization was not achieved; 'rehomogenized' are those inclusions that appear as a homogeneous glass without bubbles (although former accidentally-trapped minerals may be present). Experimental remelting of the melt inclusions was performed on hand-picked separates of garnets obtained from crushed rock pieces sieved at < 500 μ m size. The garnet fragments were loaded into Au capsules filled with dry powder of silica or chemically pure CaCO₃. The capsules were loaded in a salt + MgO + graphite assembly and experiments were performed in single stage piston cylinder apparatus at the Department of Geosciences of the University of Milan at condition from 740°C to 900°C at 0.8 and 1.0 GPa, with duration of 21-68 hours (Table 2). The temperature was controlled by a K-type thermocouple with uncertainty of ± 5°C. To avoid decrepitation of inclusions, quenching was performed at

experimental pressure and load was released after achieving ambient temperature. After the experiments, the capsules containing garnet + SiO_2 were mounted in epoxy and polished to expose rehomogenized inclusions, whereas the capsules containing garnet + $CaCO_3$ were first opened and immersed in diluted HCl to dissolve calcite and recover the garnet grains. The garnet grains were mounted in epoxy and polished (more experimental details in Bartoli et al., 2013a and Ferrero et al., 2015).

4. RESULTS

4.1 Microstructures of melt and fluid inclusions

The inclusions preserved in the starting material and after experiments were characterized on the basis of optical microscopy BSE imaging, EDS elemental mapping and micro-Raman spectroscopy, and distinguished in fluid (FI), polycrystalline (PI) also called nanogranitoids (Cesare et al., 2015) and partially crystallized (= former melt, MI).

Inclusions have irregular distribution in some garnet hosts, but they tend to cluster at the rims of large crystals or in the core of tiny crystals (Fig. 2B, C). These microstructures are indicative of a primary entrapment (according to the classification of Roedder, 1984), and suggest that inclusions were mostly trapped during the final stage of garnet growth. The FI and PI coexist in the clusters suggesting that they belong to the same fluid inclusion assemblage and that entrapment took place during the same stage of garnet growth in a condition of fluid-melt immiscibility. Most FI and PI inclusions are < 10 μ m in diameter, with isometric (negative crystal) shapes, equally abundant and have comparable size range. In PI, typical phases are albite, K-feldspar, quartz, muscovite, biotite, calcite, siderite, rutile and ilmenite. The FI contain a C-O-H fluid composed of H₂O and CO₂ (Fig. 3A). The presence of H₂O is marked by the broad band between 3100 and 3600 cm⁻¹, the CO₂ by the presence of 247 the Fermi diad (peaks at 1285 cm⁻¹ and 1387 cm⁻¹), with peak distance (Δ) corresponding to 248 CO₂ density of < 0.1 g cm⁻¹ (see Wang et al., 2011).

After the experiments, the inclusions were carefully inspected to infer the temperature of entrapment from approach to remelting equilibrium (Fig. 4). At 740°C, the MI have well defined negative crystal shapes and contain glass (yellow arrows in Fig. 4) together with abundant unmelted crystals of quartz, biotite, muscovite, albite and ilmenite, indicating that remelting was incomplete (i.e., they are partially-remelted inclusions). At 760°C, albite and muscovite are completely resorbed, biotite is less frequent, and quartz is still common with typical round shape, suggesting partial resorption. Few inclusions are completely remelted to a homogeneous glass with presence of a shrinkage bubble. At 780°C experiments show the best approach to equilibrium with > 40-50% remelted inclusions, displaying regular walls, negative crystal shape and absence of offshoots. The presence of resorbed quartz is rare. At 810°C, completely or partly remelted inclusions display offshots (white arrows in Fig. 4) and the presence of secondary minerals (i.e., formed during experiments or upon quenching) as aggregates of tiny acicular Bt. Remelted inclusions represent > 70% of the total. At 850°C and 900°C almost all inclusions are remelted or rehomogenized but show evidence of overheating (see Cesare et al., 2015; Bartoli et al., 2013c; Acosta-Vigil et al., 2016) as indicated by i) presence of lobate-cuspate walls at the contact with the host garnet, ii) frequent offshoots, iii) growth of secondary garnet at the inclusion walls, iv) segmented straight walls after assimilation of the Grt (at 900°C). Ilmenite and rutile as trapped phases become less common at T> 810°C and almost absent at 900°C. In the Raman spectrum shown in Figure 3C, the remelted glass has a broad band at ≈ 3400 cm⁻¹ related to the presence of H₂O, while the shrinkage bubble within the glass contains a fluid mixture of $CO_2 + N_2 + CH_4$ (Fig. 3B).

4.3 Major elements composition of remelted glasses

In order to determine the composition of the melt produced during anatexis of the metapsammite, and to infer the temperature of entrapment of inclusions, we conducted EMP glass analyses on completely or partially remelted glasses to explore their potential interaction with the host garnet as function of experimental temperature (Table 3). We excluded analyses with clear signal of contamination either by minerals preserved in the inclusions or by host garnet e.g. $SiO_2 < 60$ wt% and > 80 wt%, or $Al_2O_3 > 20$ wt% or $(FeOt + MgO + TiO_2) > 7$ wt%. The difference to 100% of the totals may reflect not only the fluid content but also interaction of the microprobe beam with the shrinkage bubble if present below the polished surface of the MI (see discussion below on H₂O-CO₂ content), so results of the analyses were recalculated on anhydrous basis (100 wt%). Figure 5 compares the chemical composition of the glasses recovered in all the experiments. The average MnO vs. FeO₁ content is respectively 0.5 wt% and 1.6-1.7 wt% up to 780°C, and systematically increases from 810°C to 900°C. The FeO_t + MgO + TiO₂ is 1-3 wt% up to 780°C with SiO₂ increasing from 66 wt% at 740°C to 72-76 wt% at 760°C and 75-79 wt% at 780°C. At T > 810°C, the FeO_t + MgO + TiO_2 is inversely proportional to SiO_2 (lower SiO_2 , higher $FeO_1 + MgO + TiO_2$). The CaO vs. SiO₂ has similar behavior for most temperatures with the SiO₂ increasing from 760°C to 780°C and decreasing from 780°C to 810°C, 850°C and 900°C (Fig. 5C). No temperature dependence is visible on the Na₂O vs. SiO₂ plot, except at 900°C where the SiO₂ is systematically lower for comparable Na₂O contents. The Al₂O₃ is inversely proportional to SiO₂ at 760-810°C but values get dispersed at T> 850°C resulting in an ASI number > 1, with the exception of few inclusions remelted at 850-900°C. The systematic variations reported in Fig. 5A-C and E may result from contamination of melt by mixtures of Grt-Qz and Feld-Qz (see directions of exchange vectors reported in Fig. 5B and E) suggesting that host interaction

is not negligible at T> 810°C. As a consequence, we consider the glass compositions obtained
 at 760-780 °C as most representative of the anatectic melt trapped within garnets.

In Fig. 6 the composition of glasses remelted at 760-780°C is plotted on the pseudoternary normative diagrams Qz-Ab-Or and An-Ab-Or. Compositions at 760°C include remelted glasses with and without daughter quartz. The MI cluster above the cotectic lines at 0.5-1.0 GPa, which is consistent with the displacement of cotectic lines towards Qz apex as effect of increasing Al and Ca content in the granite model system (see Bartoli et al., 2016). The composition of MI at 760°C differ by lower SiO₂ which is related to the widespread presence of relict unmelted quartz in the inclusions. However, few glasses at 760°C overlap with glasses at 780°C suggesting that equilibrium melt composition are presumably achieved within the interval 760-780°C and represent the closest approximation to the original anatectic melt. Displacement of the MI composition from the minimum melt in the Qz-Ab-Or model system has been explained by Wilke et al. (2017) as due to the presence of extra components in natural systems, mostly Ca, Fe and Ti. Wilke et al. (2017) report a displacement of the eutectic point to Or-enriched composition at similar Qz amount when melts have $\approx 3.5-7\%$ normative An. Such An content corresponds to that of remelted glasses clustering at 2-7% with few analyses up to 11-16%. The MI show variation in the Qz/feldspar ratio as indicated by their distribution along the directories Ab/Or = 0.38 and Ab/Or = 0.5collinear with the Qz apex. Such trends differ from those of Bartoli et al. (2016) who reported compositional alignments of remelted granitic glasses parallel to the Qz-feldspar join and interpreted them as due to slow diffusion of Si and Al compared to alkalis in granitic melts (Acosta-Vigil et al., 2006, 2017). In the An-Ab-Or projection, most glass compositions plot in the granite field close to the boundary with Qz-monzonite, with few Qz-monzonite compositions. Figure 6 also reports the composition of MI recovered in garnet of metasediments from Ojén (Bartoli et al., 2013b, 2013c) and in plagioclase of Grt-Bt-Sil

enclaves from El Hoyazo (Acosta-Vigil et al., 2010; Baroli et al., 2016) equilibrated at similar metamorphic conditions (700°C and 0.5 GPa for Ojén, 700-750°C and 0.5-0.7 GPa for El Hoyazo, see below). In terms of Ab/Or ratio, the composition of glasses remelted at 760°C and 780°C are similar but those at 780°C have higher normative Qz and An. The glasses plot in the calcic field of the modified alkali-lime index (MALI = $Na_2O + K_2O - CaO$) vs. SiO₂ (wt%) after Frost et al. (2001) and in the rhyolite field of the TAS classification (Fig. 7A, B).

4.4 Volatile content of remelted glasses

The H₂O and CO₂ concentration in glasses was determined by nanoSIMS only on MI remelted at 760°C and 780°C (Table 4 – Supplementary Material) and performed on selected homogeneous portion of the glass exposed on the surface of polished garnets.

The H₂O content of individual analyses is reported in Fig. 8 and is between 2.2-8.0 wt% at 760°C and 3.0-6.5 wt% at 780°, while the corresponding CO_2 is 250-850 ppm at 760°C and 500-2200 ppm at 780°C. In Fig. 8B and C the MI analyses are grouped according to host Grt and display similar variability except for garnets ANT9-1 and ANT8-1 displaying higher scatter for H₂O and CO₂ respectively. The averages at 760°C and 780°C (white circles in Fig. 8B, C) are similar for H₂O with 4.3 ± 1.7 wt% at 760°C, and 5.3 ± 1.0 wt% at 780°C, and slightly differ for CO₂ with 407 \pm 184 ppm at 760°C, and 1147 \pm 519 ppm at 780°C. In Figure 3C the measured H₂O in remelted glasses is compared with maximum and minimum water contents expected in model primary granitic melts (Holtz et al., 2001) and will be discussed below. The Fig. 3D compares the H₂O contents with the values predicted for H₂Oundersaturated haplogranitic melts (Holtz et al., 2001) while the solidus is from thermodynamic modelling (see below).

4.5 Thermodynamic modelling

Constraints on the P-T conditions of melting and inclusions entrapment were obtained by phase equilibria modelling (Fig. 9). Calculations were done in the model chemical system

MnNCaKFMASHT using the Perple X software (Connolly, 2009) with the thermodynamic database of Holland and Powell (2003 revision). We used the solution models of White et al. (2007) for melt, Holland and Powell (1998) for garnet, Tajčmanová et al. (2009) for biotite, Coggon and Holland (2002) for white mica, Newton et al. (1980) for plagioclase, and Thompson and Hovis (1979) for K-feldspar. The bulk rock composition of the rock was obtained from XRF analysis and is reported as inset in Fig. 9. The bulk composition has no -residual character and is considered representative of the original protolith with no necessity of melt reintegration (see Bartoli, 2017) because i) melt inclusions are the only petrographic evidence of anatexis (absence of melt pseudomorphs), ii) the composition of white mica, and that of biotite as well, is comparable to that of primary muscovite from amphibolite-facies subsolidus metapelites in the area (Di Vincenzo et al., 2001). The amount of H₂O used in the calculation was assumed to be the L.O.I. (loss of ignition) from the XRF analysis (=1.5 wt%). The presence of $CO_2 + H_2O \pm carbonates$ (calcite and siderite) in PI and FI point to a C-O-H fluid coexisting with the rock and melt during anatexis (Connolly and Cesare, 1993). Slight variations in H₂O activity (tested from 0.8 to 1.0 but not reported here) resulted in limited displacement of the solidus at P > 0.5 GPa, where the reaction slope is high and positive (see later and Fig. 9).

The petrographic study shows that the stable mineral assemblage is represented by Ms $+ Pl + Kf + Grt + Qz + Liq \pm Rt$ coexisting with excess C-O-H fluid as indicated by the presence of FI and PI within the same cluster. In the pseudosection of Fig. 9, such assemblage is stable at P > 0.5 GPa within a triangular field limited by the fluid-present solidus curve and by the Ms-out/Als-in curve to temperatures of 690°C < T < 820°C. The Fig. 10 (Supplementary Material) reports the calculated compositional isopleths for garnet, plagioclase and biotite. In larger garnets from cores to rims, the X_{Pyp} slightly increases and X_{Sps} slightly decreases suggesting a \approx 50°C temperature increase during the garnet growth,

while smaller garnets are homogeneous with composition similar to that of large garnet rims. The X_{Alm} in garnet has a slightly reversed tendency (higher temperature in the core and lower at the rim). The combination of garnet isopleths with the X_{An} in plagioclase, delimit a P-T space corresponding to approximately 680°C to 730°C at 1.0-1.3 GPa (Fig. 9). The low Ti content in biotite (0.12 a.p.f.u.) is not compatible with such temperatures and supports the petrographic observation of post peak recrystallization of biotite at $T < 600^{\circ}C$.

The calculated isomodes for the phases biotite, garnet and melt reported in Fig. 9B and muscovite (nor reported for clarity) indicate that biotite and muscovite are partially consumed while limited amount of garnet is formed during progressive metamorphism at P-T evolving from condition 1 to conditions 2 in Fig. 9. Modelled muscovite decreases from 12 to 8 wt%, biotite from 22 to 20 wt% and garnet increases from 1.5% to 2-4 wt%. The melt produced is limited to 2-4 wt%. The calculated isomodes are in agreement with the modal phase proportions constrained by mass balance calculation and corresponding to 7.9 wt% of muscovite, 28.1 wt% of biotite, 3.4 wt% of garnet and \approx 2 wt% of melt.

5. DISCUSSION

5.1 Conditions of MI entrapment

The present study is an example of successful remelting of inclusions that adds to previous case studies of metasedimentary rocks undergoing anatexis (e.g., Cesare et al, 2015; Bartoli et al., 2016; Tacchetto et al., 2019; Carvalho et al., 2018). Previous works report MI entrapped under fluid present (Bartoli et al., 2013b) or fluid-absent melting of felsic (e.g., Cesare et al., 2015) or ultramafic rocks (Ferrero et al., 2018), and in a range of temperatures from upper-amphibolite to UHT conditions. The present case study yields new data on the onset of low-T anatexis of metasedimentary crust under possibly fluid-present conditions, which are likely to be more common than previously thought (see Carvalho et al., 2018 and discussion therein).

The P-T-fluid conditions of melt entrapment within the host mineral can be reconstructed combining the microstructures produced during remelting experiments at increasing temperature with the proportion of remelted inclusions, the compositional variation of major elements in glass and the thermodynamic modelling. In remelting experiments, microstructural changes include i) type of minerals coexisting with glass (daughter minerals and/or trapped phases), ii) presence of offshoots (suggesting decrepitation), iii) presence of irregular boundaries between MI and Grt (indicative of host interaction), iv) crystallization of new phases (indicative overheating), v) appearance of zoning/growth structures (indicative of intense host interaction). Since glasses remelted at 780° are mostly free of daughter minerals, have no visible cracks/offshoots, and display limited or no chemical interaction with the host garnet (e.g., low $MnO + FeO_t$ in Fig. 5), they can be confidently interpreted as representative of entrapment conditions. However, at 760°C, where most inclusions have quartz which has not been re-homogenized and hence lower SiO₂, few remelted inclusions resemble in texture and composition those typically recovered at 780°C. One may question about the effect of kinetics on remelting conditions and the counter play of temperature and run duration on the achievement of equilibrium for remelting. Preliminary experiments at 780°C performed at different durations (hours to days, unpublished data) seem to confirm that run duration has an effect on remelting and that the chemical signature of the glasses recovered at lower experimental duration partly overlaps (though not completely) with the composition of homogeneous glasses recovered at 760°C. Even though the kinetics dilemma is out of the scope of this work, it is reasonable to consider the interval 760-780°C at 0.8-1.0 GPa as representative of the entrapment conditions. Temperatures above 810°C are not representative due to evident and intense glass-host interaction. Thermodynamic calculations indicate that the wet solidus of the rock locates at $\approx 690-710^{\circ}$ C between 0.5 GPa and 1.5 GPa, and that the stability field of the assemblage Ms + Pl + Kf + Grt + Qz + Liq \pm Rt is between 700-780°C.

The temperature intervals indicated by compositional isopleths of Grt and Pl and the mineral isomodes (Fig. 9) are more consistent with \approx 700-730°C and \approx 730-760°C respectively. The discrepancy between calculated and experimental conditions is maximum 60-80°C and can be ascribed to diverse sources of uncertainty in phase equilibria calculations, such as the quality of thermodynamic data and, in turn, the quality of mineral and melt solution models (White et al., 2011). For example, the current melt model requires improvements to reproduce properly natural processes (see discussion in Bartoli et al., 2013b), as already pointed out in previous works (e.g., Bartoli et al., 2013a and 2016; Grant, 2009 and White et al., 2011).

5.2 Fluid regime and volatile content of the primary melt

There are few lines of evidence suggesting that partial melting of the investigated rocks occurred in the presence of a fluid phase, including i) the coexistence of primary FI and PI within the same cluster, and ii) the major element composition of primary melt.

The abundant FI coexisting with PI in the same cluster indicate the presence of a C-O-H fluid dominated by H₂O and CO₂. Graphite was not observed within the FI but carbonates (calcite and siderite) are common within garnets. Recently Carvalho et al. (2018) discussed the recovery of C-O-H bearing FI in metasedimentary rocks from Ivrea at amphibolite to granulite conditions and argued that "a (graphitic) rock undergoing melting will always show the presence of a free C-O-H fluid, in variable amounts and with variable a_{H2O} depending on pressure, temperature and the bulk composition of the system". According to Carvalho et al. (2018), the fluid regime during anatexis might be characterized by CO_2 -dominated fluid in processes dominated by dehydration of OH-bearing phases. In the Edixon Complex, where the fluid was likely H₂O dominated, conditions of entrapment are compatible with the beginning of partial melting and potentially with both muscovite and biotite fluid-present conditions. The calculated isomodes suggest that muscovite and biotite are consumed at the solidus to produce limited amounts of melt and peritectic garnet, always in the absence of

aluminum silicate. Since the muscovite fluid-present melting (without biotite) would produce
peritectic aluminum silicate instead of garnet (Weinberg and Hasalova, 2015), the rock
assemblage is more consistent with a muscovite + biotite fluid-present melting reaction
(Weinberg and Hasalova, 2015). Increasing a_{H2O} should reduce garnet stability (Conrad et al.,
1988; Gardien et al., 2000), but the experiments of Alonso-Perez et al. (2009) on Si-rich melts
indicated the opposite, i.e. that stability of peritectic garnet increases with increasing H₂O in
melt.

The H₂O content of glasses at 760°C is more scattered (≈2-8 wt%) than at 780°C (≈4-7 wt%) but in both cases it is close to the values predicted for H₂O saturation at 0.8 GPa for haplogranitic melts (\approx 7.6 wt%; Holtz et al., 2001, Fig. 8D). The lower H₂O contents of some glasses may be explained as effect of heterogeneous distribution of the fluid (see discussion in Bartoli et al., 2014) and/or by the presence of a diluting carbonic species in the COH fluid, namely CO₂. The H₂O content of glasses determined as difference to 100 from EMPA analyses may yield similar contents, though Acosta-Vigil et al. (2016) report deviations up to 30%. In Table 3, the average difference to 100 total decreases with increasing experimental temperatures from ≈ 14 wt% at 740°C to $\approx 12-11$ wt% at 760-780°C and ≈ 10 wt% at 810-900°C. All values overestimate the measured H₂O in glasses. Possible explanations may account for the presence of light and incompatible elements as B, which is abundant in first anatectic products or micropores and would be consistent with the presence of tourmaline in the rock. Concerning other volatiles, the Cl and F content of the glasses was below the detection limits (Table 3). This inconsistency requires further investigation.

The maximum CO_2 content of the remelted glasses from NanoSIMS (with all C calculated as CO_2) is 200-500 ppm at 760°C and 500-2000 ppm at 780°C (Fig. 8A). Such difference may be explained considering that i) the CO_2 solubility in felsic melts slightly increases with increasing T at least at 0.5 GPa (Tamic et al., 2001), ii) the CO_2 solubility and

speciation are affected by the glass composition, with total CO_2 increasing with increasing SiO₂ and decreasing with increasing Na₂O, and molecular CO₂ prevailing on the $(CO_3)^{2^2}$ species in SiO₂ rich- and Na₂O poor- compositions (Brooker et al., 1999). At 780°C, glasses have higher SiO₂-CaO and lower Na₂O with respect to glasses produced at 760°C, so the CO₂ speciation is expected to be dominated by molecular CO₂. Even though most of the analyzed MI appeared homogeneous and bubble-free on polished surface, the presence of shrinkage bubbles deep within the inclusion cannot be excluded. Since the fluid phase preserved in the shrinkage bubble is mostly CO₂-rich (Fig. 3B) in agreement with Wallace et al. (2015), this may account for an additional CO₂ reservoir. If (slightly) higher resorption of the shrinkage bubbles takes place at increasing temperature, this may explain the higher CO₂ content of glasses remelted at 780°C vs. 760°C.

Granitic melt produced by water-present melting of metasedimentary rocks are expected to have higher ASI values than those produced by muscovite or biotite dehydration melting (Acosta-Vigil et al., 2003), to be enriched in Na₂O and CaO (Weinberg and Hasalova, 2015), and to contain higher proportion of MgO and FeO at increasing water content or water activity (e.g. Conrad et al., 1988; Holtz and Johannes, 1991) at constant P–T (Patiño Douce and Beard, 1996). This is partly consistent with the glass composition at 780°C, with ASI value of 1.57, MALI (= Na₂O + K₂O - CaO) value of \approx 4.0 ± 1.0 wt%, CaO content of \approx 1.27 ± 0.7 wt.% and FeO_{tot} + MgO > 2.1 wt%, supporting the hypothesis of water-saturated partial melting.

5.3 Anatexis and evolution of the Edixon Metamorphic Complex

The tectonometamorphic history of the Edixon Metamorphic Complex and its role in the evolution of the Wilson Terrane has been primarily constrained by investigations along the western Lanterman Range but also southward within a wider belt including Salamander Range and Niagara Icefalls-Mountaineer Range regions which are exposed, as the Lanterman

Range, close to the Wilson-Bowers tectonic boundary (e.g., Ghiribelli, 2000; Talarico et al., 1998; 2004). In general, the EMC contains no record of an early HP metamorphic stage, but preserves a broad metamorphic zonation from low/medium grade to high grade up to anatexis. However, previous studies showed the presence of rocks with contrasting P-T paths in different localities of the EMC (Castelli et al., 2003; Ghiribelli, 2000; Talarico et al., 1998, 2004). Ghiribelli (2000) recognized, at the western Lanterman Range and at the Salamander Range, that the rocks of the EMC show close similarities in lithological assemblages and metamorphic field gradients with other metasedimentary sequences characterized by an andalusite-sillimanite prograde counterclockwise path with isobaric cooling, and abundant Grt-Crd migmatites outcropping at the Deep Freeze Range (Palmeri et al. 1991, 1994; Palmeri, 1997). The latter low-pressure belt (> 200 km wide) located in an internal position of the Wilson Terrane is characterized by the presence of the Granite Harbor Intrusive complex (Borg et al., 1990; Ghezzo et al., 1989). The extensive and prolonged magmatic underplating of the Granite Harbor Intrusives bodies in the Wilson Terrane (from 520-530 to 500 Ma, Giacomini et al., 2007; Rocchi et al., 2004) possibly triggered, in different times, the anatectic event and, at the same time, promoted the juxtaposition of migmatites with contrasting P-T paths likely formed during the same Ross orogenic cycle (Palmeri, 1997).

In other localities, as at the Mountaineer Range region, a different P-T path has been proposed (Castelli et al., 2003; Talarico et al., 1998), involving an early decompression path following a moderate thickening, magmatic intrusions between 500 to 480 Ma (the Granite Harbour intrusives), and a final cooling-unloading path has been inferred (Castelli et al., 2003).

516 In this paper, the MI preserved in peritectic garnet and phase equilibria modelling
517 demonstrate that the portion of the EMC close to Mt. Edixon experienced anatexis above
518 700°C and up to 760-780°C at ≈1.0 GPa, corresponding to the P-T conditions of entrapment.

In the NW part of the Lanterman Range (Carnes Crag) Grt-Crd migmatites yield T-P conditions at $\approx 700^{\circ}$ C and ≈ 0.5 GPa (Ghiribelli, 2000). Different conditions, however, were found in the southern portion of the EMC (Mountaineer Range), where the highest metamorphic grade shows temperature at $\approx 700-750^{\circ}$ C and pressure from ≈ 0.5 up to 0.6 GPa (Castelli et al., 2003). The spread in low to medium pressure rock types and contrasting P-T paths at the boundary with the Wilson-Bowers terranes is not uncommon (Ricci et al., 1997; Ricci and Tessensohn, 2003), and supports the interpretation that anatexis could have taken place at different crustal levels during the evolution of the tectono-metamorphic stages of the Ross Orogen.

The latter is in keeping with the multiple and diachronous generations of white micas evidenced by Di Vincenzo et al., (2001) in the different metamorphic complexes of the Lanterman Range. The different white micas have been interpreted as testifying different P-T deformation stages developed during the evolution of the three metamorphic complexes (Di Vincenzo et al., 2001).

The white micas in the investigated sample GO5 have no evidence of retrogression, they are texturally and compositionally similar to type-2 white mica of Di Vincenzo et al., (2001), in fact they grow along the main amphibolite-facies foliation, and show a light enrichment in celadonitic component (Si = 3.1 a.p.f.u. and Ti = 0.022 a.p.f.u.). Moreover, data collected in this paper indicate that white micas are compatible with fluid-present melting as reconstructed by combination of thermodynamic modelling and MI remelting experiments. Since the composition of white micas in the Wilson Terrane evolves with time and the type-2 mica dates to ca. 480-490 Ma, we infer that the latter age could be referred to melting in the Lanterman Range.

Even though it is not possible to reconstruct a proper P-T-relative time path based on the single sample GO5, the equilibrium conditions reconstructed from the remelting

experiments and thermodynamic modelling allow extending the evolution of the EMC up to ca. ≈ 1.0 GPa at 760-780°C. The latter data suggest two possibility: i) the EMC at the Lanterman Range is not an homogeneous metamorphic complex characterized by a low-pressure metamorphic imprint as postulated previously (Ghiribelli, 2000) but, similar to the HP-UHP belt (GHMC), it represents a mélange of different crustal slices; ii) the outcrop, from which the study sample comes from could be the extension towards the southern Lanterman Range of the intermediate-pressure belt of the BMC. Whatever the solution of the latter problem is, combining (Fig.10) geochronological data collected at the Lanterman Range (Di Vincenzo et al., 2016 and references therein), previous P-T paths (Di Vincenzo et al., 2001; Ghiribelli 2000; Talarico et al., 2004) and the results for sample GO5, the docking between the BMC and/or the different tectonic slices of the EMC might have been before 480 Ma. The latter age is widespread in the area and it was interpreted as the Early Ordovician final "closure" of micas at the Lanterman Range (Di Vincenzo et al., 2001) and mainly referred to the exhumation process of the different complexes, whereas in the inner Wilson Terrane that age correspond to the youngest undeformed calc-alkaline granitoids. Thus, in the Lanterman Range, the high-pressure eclogitic rocks (GHMC), the

medium pressure amphibolite stage (BMC/EMC pro parte) and the low-pressure amphibolite
 stage (EMC pro parte) shared most of the late exhumation path.

AKNOWLEDGEMENTS

This work was funded by the CaRiPaRo project Makearth (Cassa di Risparmio di Padova e Rovigo) to FF, the grant SIR RBSI14Y7PF (Italian Ministry of Education, University, Research) to OB, the PNRA (PdR 13/B2.07) to RP and by the German Federal Ministry for Education and Research and the Deutsche Forschungsgemeinschaft (Project FE 1527/2-1 and FE 1527/2-2) to SF.

Rock sample used in this study was collected by Gianni Capponi e Giacomo Oggiano and is available at the PNRA (Programma Nazionale Ricerche in Antartide) rock repository located at the Museo Nazionale dell'Antartide (Siena, Italy).

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FIGURE CAPTIONS

Figure 1 – Location of the Ross orogenic belt in the Transantarctic Mountains
(modified from Godard and Palmeri, 2013). A) Location map; B) geological sketch map of
Northern Victoria Land (RBT, Robertson Bay Terrane; BT, Bowers Terrane; WT, Wilson
Terrane; LR, Lanterman Range; SR, Salamander Range; MR, Mountaineer Range); C)
geological map of the Lanterman Range (after Ghiribelli, 2000) and location of the sampling area.

Figure 2 – Microphotographs of sample GO5 under optical microscope, plane
polarized light (PPL). (A) Typical microstructure with muscovite and biotite growing along
the main foliation, abundant plagioclase and quartz, garnet of ca. 400 µm. Cluster of
inclusions (FI and PI) (B) along the rims of a large crystal of garnet, or (C) in the core of a
tiny crystal of garnet partly replaced by down-grade biotite.

Figure 3 –Representative Raman spectra of phases and volatile species in: A) fluid inclusion; B) shrinkage bubble in remelted glass; C) remelted glass.

Figure 4 – Microstructures of MI under scanning electron microscope (BSE images)
after remelting experiments in piston cylinder apparatus at increasing temperature: 740°C partially homogenized MI with glass (yellow arrows), quartz, biotite, muscovite, albite and
ilmenite; 760°C – glassy MI with rounded quartz (= partial resorption); 780°C - glassy MI
with regular walls, negative crystal shape and crystals of Ilm; 810°C - homogenized MI with

961 local offshots (white arrows) and secondary acicular Bt; 850°C - homogenized MI with
962 lobate-cuspate walls, offshots (white arrows), and secondary Grt suggesting overheating and
963 host interaction; 900°C - glassy MI with segmented straight walls after assimilation of host
964 Grt (inset).

Figure 5 – Harker diagrams showing the major element concentrations (wt%) of melt inclusions in crustal enclaves after remelting experiments at 740°C to 900°C (data in Table 3).
(A) MnO vs. FeOtot. (B) FeOt + MgO + TiO₂ vs. SiO₂ with mixing vectors between the mineral pairs Grt-Qz and Feld-Qz (see text for details). (C) CaO vs. SiO₂. (D) Na₂ O vs. SiO₂.
(E) Al₂ O₃ vs. SiO₂ with mixing vectors between the mineral pairs Grt-Qz and Feld-Qz. (F) ASI vs. SiO₂.

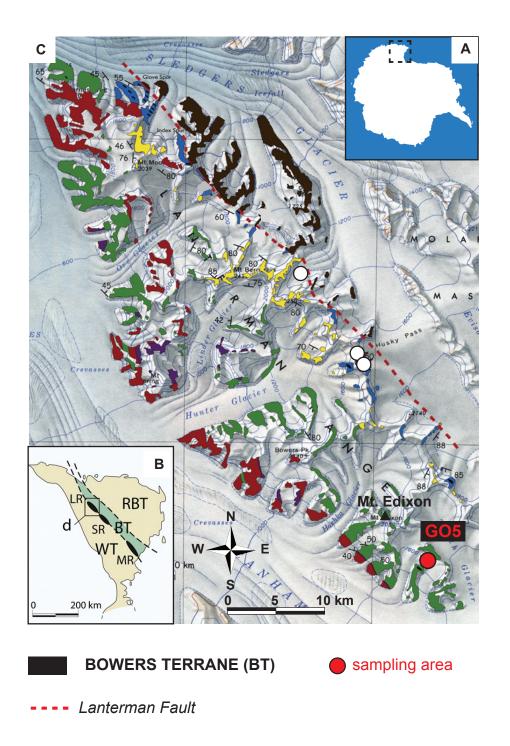
Figure 6 – CIPW Qz-Ab-Or and An-Ab.Or diagrams showing the normative compositions of all analyzed melt inclusions. Blue and green circles are for glassy inclusions remelted at 760°C and 780°C respectively. Grey dotted lines show the eutectic points and cotectic lines for the subaluminous haplogranite system at $a_{H2O} = 1$ and different pressures (Holtz and Johannes 1992; Luth et al. 1964; Huang and Wyllie 1975). Grey circles and lines represent the effect of different amounts of normative melt An at constant 100 MPa pressure (data from James and Hamilton, 1969). References to experimental glasses from Ojén (Bartoli et al., 2013c) and El Hoyazo (Acosta-Vigil et al., 2007) where "met." refers to metatexites, and "diat." to diatexite.

Figure 7 – Composition of analyzed melt inclusions in (A) modified alkali-lime index (MALI = Na₂O + K₂O – CaO) vs. SiO₂ (wt%) after Frost et al. (2001); (B) total alkali vs. SiO₂ diagram. Symbols as in Fig. 6.

Figure 8 – H₂O and CO₂ concentration of re-homogenized melt inclusions in experiments performed at 760°C (blu circles) and 780°C (green circles) at 0.8-1.0 GPa measured by NanoSIMS. White circles in (B) and (C) are averages of analyses at 760°C and 780°C. Horizontal bars are one standard deviation on average values. (D) Comparison of H₂O with expected values from Holtz et al. (2001), cross symbols are averages at $760^{\circ}C$ (green) and 780°C (blue). Figure 9 – (A) P–T section for GO5 calculated in MnNCaKFMASHT system (bulk composition from XRF analysis). (B) Isomodes of biotite, garnet and melt. The green shaded area in B defines the calculated modal composition consistent with mass balance calculations. Fig. 10 (Supplementary Material) – Compositional isopleths for X_{Pyr} , X_{Alm} and X_{Spn} in garnet, X_{An} in plagioclase, X_{Mg} and Ti (a.p.f.u.) in biotite calculated in MnNCaKFMASHT system (bulk composition of GO5 from XRF analysis). Fig. 11. P-T trajectory path for prograde to retrograde path of eclogite and felsic rocks from Lanterman Range (modified from Di Vincenzo et al., 2016). Blue box represents the estimated P-T conditions for partial melting in the Edixon Complex. **TABLE CAPTIONS** Table 1 (Supplementary Material) – Mean compositions (wt%) of minerals from GO5 (standard deviations within brackets).

Table 2 – Experimental conditions of piston-cylinder remelting experiments.

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	anhydrous basis. Numbers in parentheses refer to 1σ standard deviation.
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9 ₁₀ 1015	Table 4 (Supplementary Material)- H ₂ O and CO ₂ concentrations measured in remelted
	Table 4 (Supplementary Materiar)- 1120 and CO ₂ concentrations measured in remented
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WILSON TERRANE (WT)



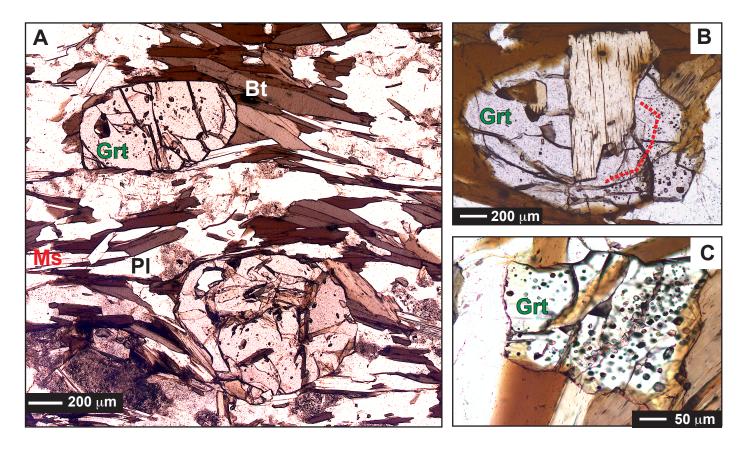


Figure 2

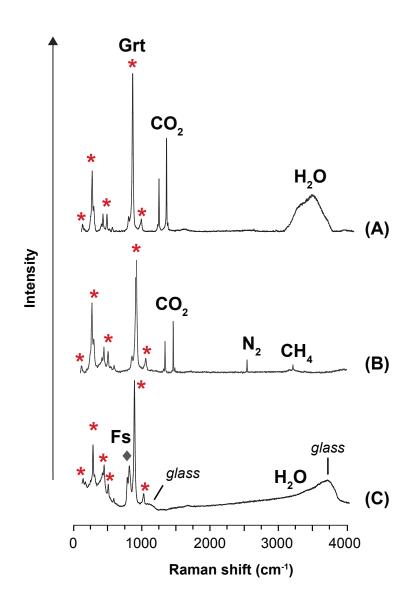


Figure 3

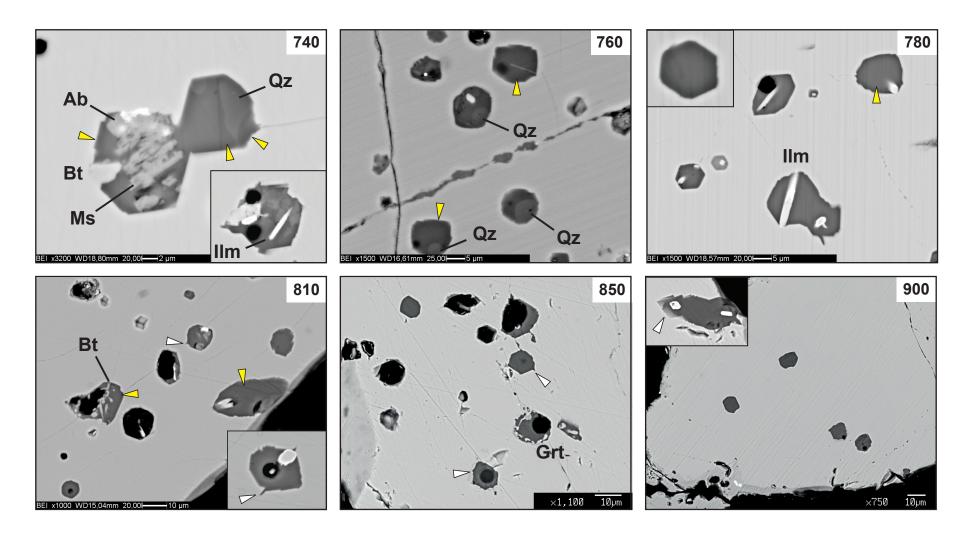


Figure 4

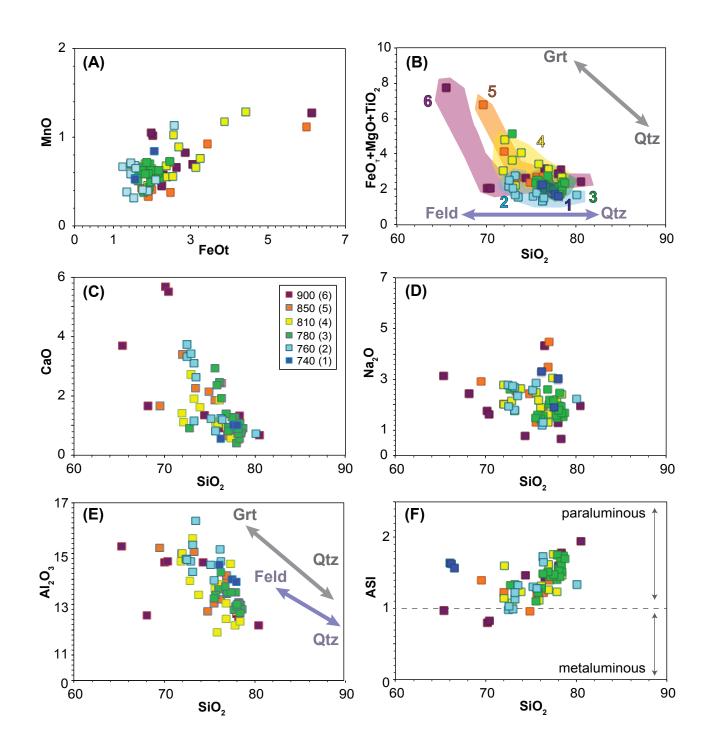


Figure 5

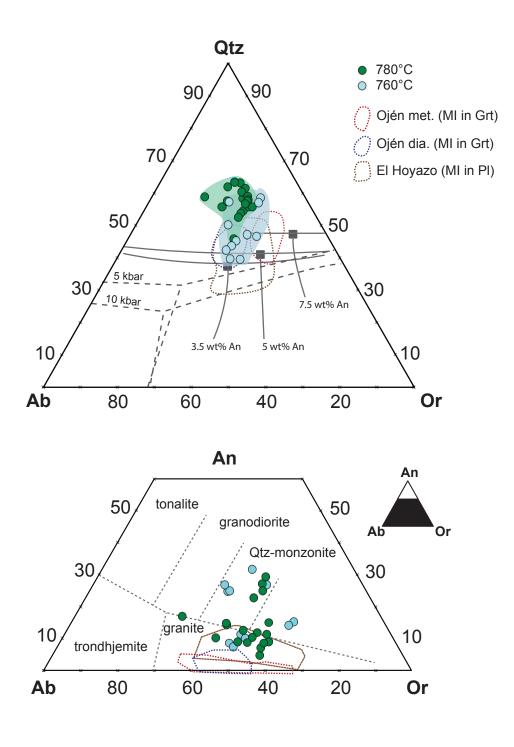


Figure 6

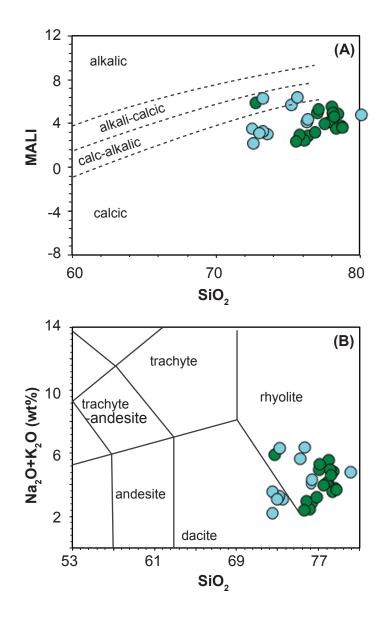


Figure 7

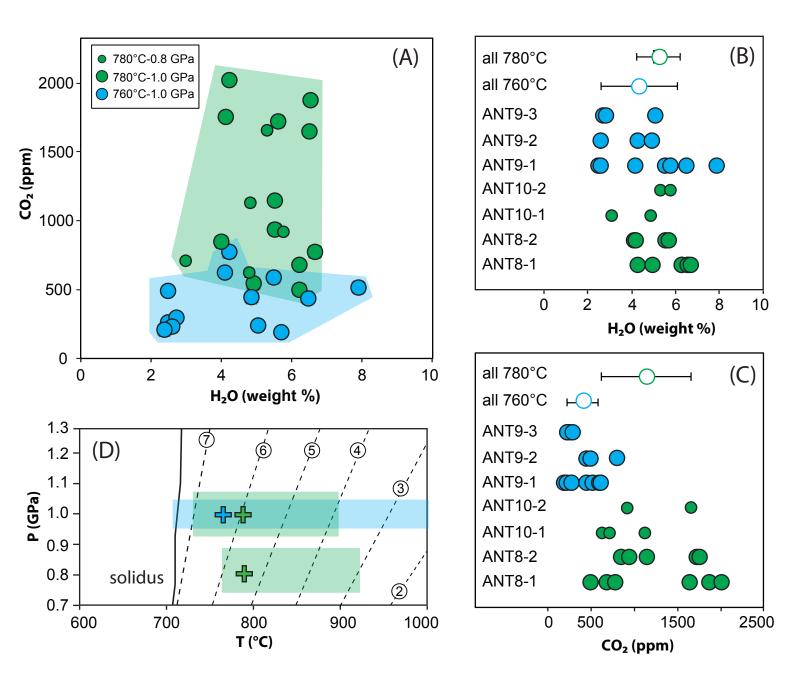


Figure 8

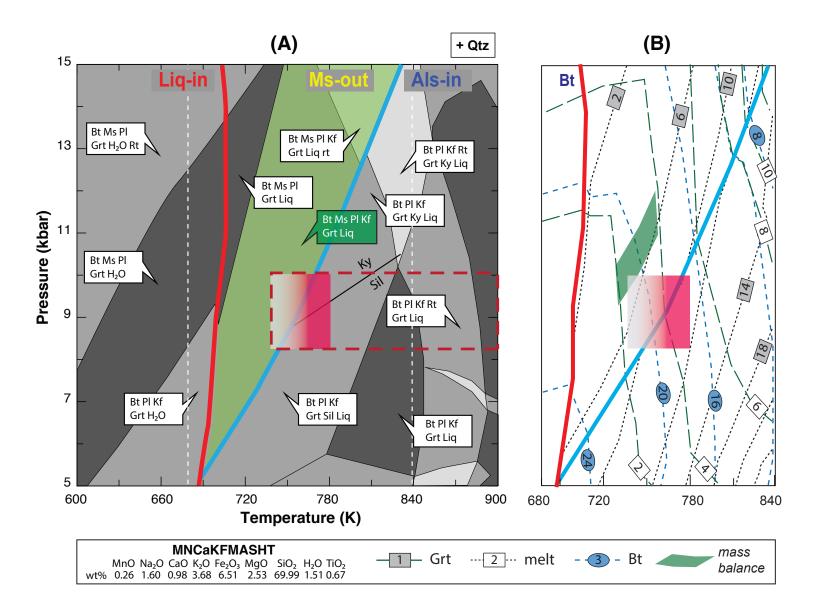


Figure 9

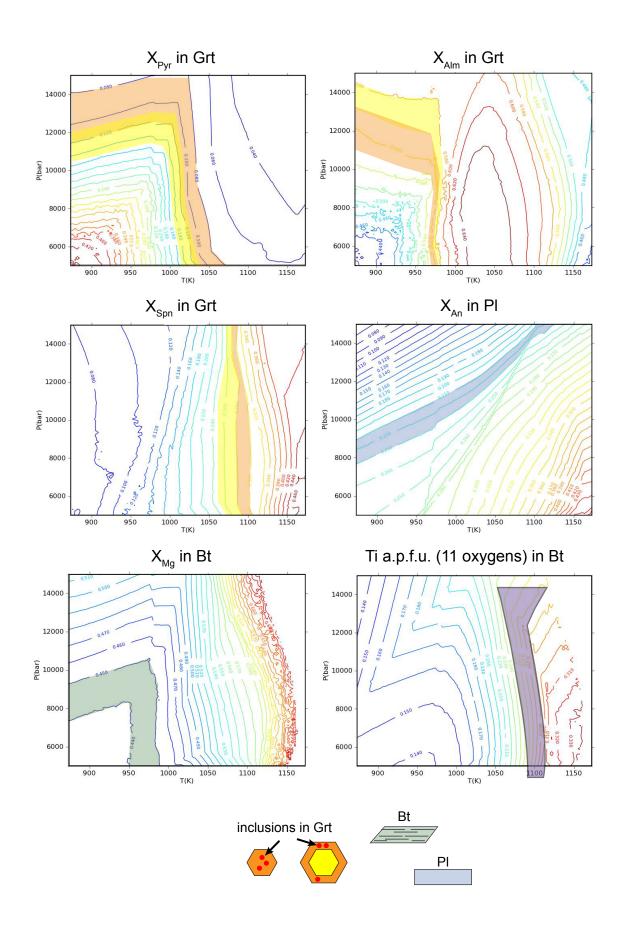


Figure 10 (Supplementary Material)

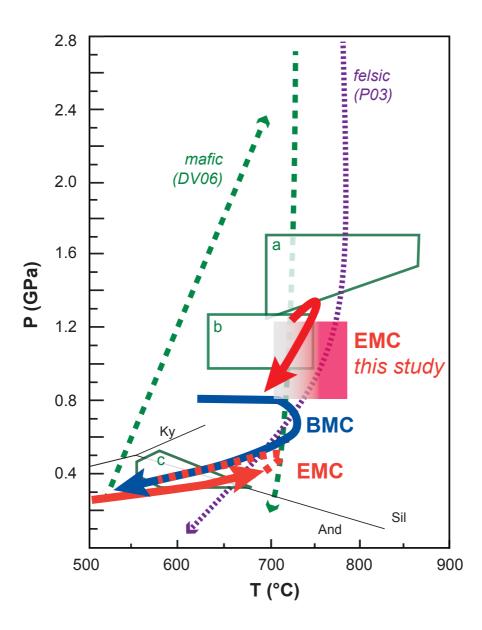


Figure 11

(n. analyses))		Garnet	(6)				Musco	ovite (8)			Bioti	te (9)		P_{i}	lagio	clase (8)		Kfeldspar (7)
		core	CERD		rim	CED				CTD	<u> </u>			CED	·			CIED	. <u> </u>		CED
5:0		mean SiO ₄	STD	iO ₅	mean	STD	SiC SiO ₈	mean	i SiO ₉	STD	SiC SiO ₁₁	mean	8:0	STD	SiC SiO ₁₄	mean	SiO ₁₅	STD	SiC SiO ₁₇	mean SiO ₁₈	STD
SiO ₂	SiO ₃				SiO ₆								SiO ₁₂								
TiO ₂	TiO ₃	TiO_4		iO ₅	TiO ₆		TiC TiO ₈		TiO ₉		TiC TiO ₁₁		TiO ₁₂		TiC TiO ₁₄		TiO ₁₅		TiC TiO ₁₇	TiO ₁₈	
Al_2O_3	Al_2O_4	Al_2O_5		l_2O_6	Al_2O_7		$Al_2 Al_2O_9$		Al_2O_{10}		$Al_{2'}Al_{2}O_{12}$		Al_2O_{13}	3	$Al_2 Al_2O_{15}$		Al_2O_{16}		$Al_2 Al_2O_{18}$		19
FeO	FeO	FeO		eO	FeO		FeC FeO		FeO		FeC FeO		FeO		FeC FeO		FeO		FeC FeO	FeO	
MnO	MnO	MnO		InO	MnO		Mn MnO		MnO		Mn MnO		MnO		Mn MnO		MnO		Mn MnO	MnO	
MgO	MgO	MgO		1gO	MgO		Mg MgO		MgO		Mg MgO		MgO		Mg MgO		MgO		Mg MgO	MgO	
CaO	CaO	CaO		aO	CaO		Ca(CaO		CaO		Ca(CaO		CaO		Ca(CaO		CaO		Ca(CaO	CaO	
Na ₂ O	Na ₂ O	Na ₂ O		a_2O	Na ₂ O		Na_2Na_2O		Na ₂ O		Na_2Na_2O		Na ₂ O		$Na_2 Na_2 O$		Na ₂ O		Na ₂ Na ₂ O	Na ₂ O	
K ₂ O	K_2O	K ₂ O	K	$_{2}O$	K ₂ O		K ₂ (K ₂ O		K_2O		K ₂ (K ₂ O		K_2O		K ₂ (K ₂ O		K_2O		K ₂ (K ₂ O	K ₂ O	
Total		0.00	(0.64)		0.00	(0.63)		0.00)	(0.40)		0.00		(0.25)		0.00)	(0.46)		0.00	(0.46)
Si		2.98	(0.01)		2.99	(0.01)		3.04	Ļ	(0.05)		2.74		(0.03)		2.80)	(0.03)		3.01	(0.02)
Ti		0.00	(0.00)		0.00	(0.00)		0.03	;	(0.01)		0.12		(0.01)		0.00)	(0.00)		0.00	(0.00)
Al		2.00	(0.01)		1.99	(0.02)		2.71		(0.07)		1.63		(0.04)		1.21		(0.03)		1.00	(0.02)
Fe ³⁺		0.06	(0.02)		0.05	(0.02)		0.00)	(0.00)		0.00		(0.01)							
Fe ²⁺		1.69	(0.09)		1.67	(0.03)		0.12	2	(0.01)		1.30		(0.02)		0.00)	(0.00)		0.00	(0.01)
Mn		0.76	(0.04)		0.83	(0.04)		0.00)	(0.00)		0.04		(0.00)		0.00)	(0.00)		0.00	(0.00)
Mg		0.35	(0.09)		0.29	(0.04)		0.10)	(0.02)		1.04		(0.01)		0.00)	(0.00)		0.00	(0.00)
Ca		0.17	(0.01)		0.16	(0.03)		0.00)	(0.00)		0.00		(0.00)		0.21		(0.02)		0.00	(0.00)
Na		0.00	(0.00)		0.00	(0.00)		0.06	5	(0.01)		0.01		(0.00)		0.77		(0.03)		0.01	(0.01)
Κ		0.00	(0.00)		0.00	(0.00)		0.87	'	(0.01)		0.92		(0.02)		0.01		(0.00)		0.97	(0.01)
xMg(Fetot)		0.16	(0.04)		0.14	(0.02)		0.45		(0.01)		0.44		(0.01)							
Al(IV)								0.96		(0.05)		1.26		(0.02)							
Al(VI)								1.75	i	(0.03)		0.37		(0.03)							
Grossular		0.11	(0.03)		0.09	(0.01)															
Almandine		0.54	(0.03)		0.55	(0.01)															
Pyrope		0.05	(0.00)		0.05	(0.01)															
Spessartite		0.25	(0.01)		0.27	(0.02)															
Anorthite																0.21		(0.02)		0.00	(0.00)
Albite																0.78		(0.03)		0.01	(0.01)
Orthoclase																0.01		(0.00)		0.99	(0.01)

Garnet compositions normalized to 12 oxygens, muscovite and biotite compositions normalized to 11 oxygens, plagioclase and feldspar compositions normalized to 8 oxygens, STD standard deviation

#	matrix	T(°C)	P(GPa)	time (hrs)
ANT7	CaCO ₃	740	0.8	46
ANT9	CaCO ₃ and SiO ₂	760	1.0	48
ANT8	SiO ₂	780	0.8	68
ANT10	SiO ₂	780	1.0	67
ANT5	CaCO ₃	810	0.8	23
ANT3	CaCO ₃	850	0.8	22
ANT4	CaCO ₃	900	0.8	21

Temp. (No. Analyses)	740°C	(3)	760°C	(10)	780°C	(18)	810°C	(14)	850°C	(13)	900°C	(9)
	mean	^c STD	mean	STD	mean	STD	mean	STD	mean	STD	mean	STD
SiO ₂	77.28	(0.97)	74.13	(1.55)	77.48	(1.00)	75.52	(2.30)	71.58	(7.96)	73.54	(5.23)
TiO ₂	0.02	(0.02)	0.10	(0.11)	0.10	(0.10)	0.08	(0.12)	0.22	(0.21)	0.57	(1.30)
Al_2O_3	14.13	(0.37)	14.83	(0.63)	13.22	(0.40)	13.49	(1.14)	15.59	(4.38)	13.63	(1.20)
FeO	1.72	(0.28)	1.69	(0.40)	1.92	(0.23)	2.59	(0.83)	3.81	(2.12)	3.25	(1.65)
MnO	0.63	(0.18)	0.57	(0.24)	0.60	(0.09)	0.76	(0.25)	0.75	(0.34)	0.85	(0.28)
MgO	0.12	(0.06)	0.11	(0.05)	0.19	(0.14)	0.20	(0.10)	0.64	(0.66)	0.21	(0.12)
CaO	0.86	(0.26)	2.18	(1.16)	1.24	(0.71)	1.29	(0.59)	1.50	(0.91)	2.44	(1.99)
Na ₂ O	2.75	(0.74)	2.22	(0.63)	1.83	(0.35)	2.18	(0.47)	1.90	(1.25)	2.00	(1.17)
K ₂ O	2.48	(0.70)	4.15	(0.40)	3.42	(0.54)	3.89	(0.87)	4.00	(2.38)	3.51	(1.57)
total	100.00		100.00		100.00		100.00		100.00		100.00	
^a total _{EMPA}	85.70	(0.81)	87.86	(2.20)	88.97	(1.10)	90.06	(1.60)	86.53	(1.33)	89.29	(2.14)
^b H ₂ O(diff)	14.30	(0.99)	12.14	(1.87)	11.03	(1.18)	9.94	(1.71)	13.47	(5.81)	10.71	(2.25)
ASI	1.61	(0.04)	1.27	(0.26)	1.50	(0.17)	1.35	(0.21)	1.59	(0.51)	1.33	(0.42)
AI	0.07	(0.00)	0.07	(0.02)	0.06	(0.01)	0.06	(0.01)	0.08	(0.04)	0.06	(0.01)
MALI	4.38	(0.59)	4.19	(1.46)	4.01	(0.98)	4.79	(0.86)	4.40	(2.56)	3.07	(1.55)

^a totals from EMPA, ^bH₂O by difference to 100%, ^c standard deviation

sample	H ₂ O (wt%)	STD	CO ₂ (ppm)	STD							
remelting $T = 780^{\circ}$ C; mean SiO ₂ = 68.27 wt. %											
ANT8_1_1	6.66	(0.09)	774.39	(19.19)							
ANT8_1_2	4.90	(0.06)	541.17	(19.04)							
ANT8_1_3	4.22	(0.05)	2019.91	(39.55)							
ANT8_1_4	6.52	(0.09)	1878.96	(36.74)							
ANT8_1_5	6.48	(0.09)	1648.30	(31.55)							
ANT8_1_6	6.21	(0.08)	680.84	(16.56)							
ANT8_1_7	6.21	(0.08)	496.32	(17.25)							
mean	5.89	(0.94)	1148.56	(670.27)							
ANT8_2_1	5.52	(0.07)	939.69	(19.50)							
ANT8_2_2	5.52	(0.07)	1148.56	(22.02)							
ANT8_2_3	4.01	(0.05)	849.99	(18.09)							
ANT8_2_4	4.11	(0.05)	1750.82	(34.39)							
ANT8_2_3	5.63	(0.07)	1725.19	(37.19)							
mean	4.96	(0.82)	1282.85	(429.48)							
ANT10_1_1	4.83	(0.06)	1128.05	(22.70)							
ANT10_1_2	2.98	(0.05)	710.32	(17.85)							
ANT10_1_3	4.80	(0.05)	624.46	(17.12)							
mean	4.20	(1.06)	820.94	(269.41)							
ANT10_2_1	5.73	(0.07)	916.62	(20.10)							
ANT10_2_2	5.28	(0.06)	1661.12	(34.52)							
mean	5.51	(0.32)	1288.87	(526.44)							
remelting $T = 76$	$0^{\circ}C$; mean SiO ₂ =	= 65.02 wt.	%								
ANT9_1_1	2.47	(0.05)	265.23	(18.28)							
ANT9_1_2	2.40	(0.05)	209.09	(18.51)							
ANT9_1_3	6.44	(0.09)	443.40	(17.03)							
ANT9_1_4	7.88	(0.13)	512.97	(17.69)							
ANT9_1_5	5.49	(0.07)	588.63	(18.11)							
ANT9_1_6	4.11	(0.05)	613.04	(18.78)							
ANT9_1_7	5.72	(0.08)	188.34	(19.18)							
mean	4.93	(2.04)	402.96	(180.26)							
ANT9_2_1	4.21	(0.05)	770.47	(19.33)							
ANT9_2_2	2.50	(0.05)	489.78	(16.15)							
ANT9_2_3	4.87	(0.06)	443.40	(17.42)							
mean	3.86	(1.22)	567.89	(176.97)							
ANT9_3_1	2.71	(0.05)	296.96	(17.88)							
ANT9_3_2	2.61	(0.05)	228.61	(18.55)							
ANT9_3_3	5.03	(0.06)	248.14	(19.26)							
mean	3.45	(1.37)	257.90	(35.20)							
STD = standard devi		× /	-	× /							

STD = standard deviation