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Interplay between Fluid Extraction Mechanisms and Antigorite Dehydration Reactions (Val Malenco, Italian Alps)

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ABSTRACT

We investigate the feedback between antigorite dehydration reactions and fluid extraction mechanisms through the analysis of field-scale structures and microtextures formed by antigorite dehydration reactions producing olivine and talc in the Bergell intrusion contact aureole at Alpe Zocca (Malenco Unit, Northern Italy). The reactions, which resulted in the replacement of serpentinites by metaperidotites, occurred under quasi-static conditions. The main reaction front, which defines the equilibrium isograd, is an ∼150 m wide zone composed of variably reacted rocks with an irregular distribution at scales ranging from a few centimeters to a few meters. Veins composed of the pro-grade mineral assemblage occur downstream (<100 m) of this front. They are often surrounded by centimeter- to decimeter-scale dehydration reaction zones that propagate into the serpentinite wall-rock. Olivine in the metaperidotites and partially reacted serpentinites has a crystallographic preferred orientation (CPO) correlated with the antigorite CPO, with [010]_{Ol} axes parallel to [001]_{Atg}. This CPO is accompanied by a shape-preferred orientation (SPO) that marks the foliation in both rock types. Olivine crystals in Ol–Tlc dehydration veins also have SPO and CPO that define a jackstraw texture within the plane of the vein. They are elongated parallel to [001] within the vein plane and have their [010]_{Ol} axes normal to the vein. We interpret the olivine–talc assemblage in the veins as resulting from dehydration reactions at lower temperature than the equilibrium conditions owing to fluid extraction from the wall-rock into the veins. The jackstraw texture indicates fast kinetics, with the crystal orientation controlled by anisotropic growth under a fluid pressure gradient. We interpret the foliated metaperidotites as formed at near equilibrium conditions, with pervasive fluid extraction from the metaperidotite by viscous metamorphic compaction. Olivine CPO in these rocks may result from topotaxy, oriented growth in the presence of an anisotropic (foliation-controlled) fluid flow, and/or solid reorientation of the anisometric olivine crystals accommodated by the deformation of the weaker talc (+ chlorite) matrix during compaction.

Key words: serpentinite; dehydration reactions; fluid migration; microstructures; crystal preferred orientations; effective pressure; fluid pressure; compaction

INTRODUCTION

Dehydration reactions produce a significant net decrease of the solid volume as well as a large volume of fluids, resulting in a fluid-filled porosity. The low permeability of crystalline rocks surrounding dehydration fronts at middle crust and greater depths (e.g. Manning & Ingebritsen, 1999) should prevent efficient large-scale pervasive fluid migration through an interconnected pore network (e.g. Leclère et al., 2018). In the absence of fluid extraction, increase in fluid pressure decreases the effective pressure (difference between the confining pressure and the fluid pressure) and, by consequence, slows down the reaction rates (e.g. Llana-Fúnez et al., 2007, 2012). Successful models of pervasive fluid flow during replacement reactions in the crust and mantle require metastability of the solid matrix in contact with
the percolation aqueous fluid (i.e. coupled dissolution–precipitation driven by chemical fluid–rock disequilibrium interaction; Putnis, 2002; Putnis & Putnis, 2007; Putnis & John, 2010; Etschmann et al., 2014; Plümpner et al., 2017b). In such situations, it has been shown that chemical potential gradients along the fluid-filled nanoporosity (diffusio-osmosis) and those related to electrical charges (electro-osmosis) may be as important as fluid pressure gradients in controlling fluid flux (Kar et al., 2016; Plümpner et al., 2017a). However, aqueous fluids released by a homogeneous lithological column being dehydrated are not expected to be significantly far from equilibrium when brought in contact with the surrounding unreacted protolith. Furthermore, dehydration reactions have the potential to produce a significant volume of fluid-filled porosity over discrete temperature ranges (i.e. discontinuous dehydration reactions) and, hence, fluid-filled pore spaces well beyond nanoporosity (Bedford et al., 2017). Dissolution–precipitation alone is therefore not an efficient mechanism for fluid extraction during dehydration reactions.

Alternative models for fluid migration during dehydration reactions rely on the deformation of the solid matrix or on a hierarchical channelling of an initially heterogeneous distribution of fluid-pockets (Miller et al., 2003; Plümpner et al., 2017b). Under the high-pressure and high-temperature conditions prevailing in the middle and lower crust and mantle, viscous compaction has been proposed as an efficient draining mechanism (Connolly, 1997, 2010). When the time scale for viscous compaction is shorter than the time scale of fluid production, fluid migration takes place through porosity waves that propagate independently of the reaction front (Connolly, 1997, 2010; Connolly & Podladchikov, 1998, 2013, 2015). Alternatively, if the time scale for compaction is larger (cooler and lower pressure conditions), the rocks remain rigid on the time scale of the reaction and fluid overpressure may result in fracturing. The widespread occurrence of synmetamorphic veins in the upper crust (e.g. Etheridge et al., 1983) is usually taken as evidence for the latter process (Nakashima, 1995; Flekkøy et al., 2002; Okamoto et al., 2017).

Numerical models dealing with fluid migration by porosity waves and/or hydrofracturing are increasingly more sophisticated (e.g. Flekkøy et al., 2002; Malvoisin et al., 2015), but experimental and natural observations constraining the contribution of these two processes in different geological settings remain elusive. Yet they should result in contrasting magnitudes of the difference between the transient fluid pressure and the confining pressure, which in turn should affect the reaction rates (Llana-Fúnez et al., 2007, 2012). Viscous compaction should be associated with low effective pressure (fluid pressure approaches lithostatic pressure), whereas hydrofracturing is the result of a transient overpressure (fluid pressure higher than lithostatic) that rapidly evolves towards high effective pressure owing to an increase in permeability coeval with crack propagation and connection to external reservoirs. Because effective pressure influences reaction kinetics and, by consequence, grain growth processes (Lasaga, 1989; Miller et al., 2003; Wang & Wong, 2003; Hildyard et al., 2011; Llana-Fúnez et al., 2012; Leclère et al., 2018), the microstructural investigation of dehydrated rocks (Padrón-Navarta et al., 2010b, 2011, 2015; Clément et al., 2018; Dilissen et al., 2018) is an appealing tool to deduce the fluid migration mechanisms active in nature.

The purpose of this work is to investigate the mechanisms of fluid migration during antigorite dehydration by studying macro- and microstructures over a 1 km² exposure of partially dehydrated antigorite-schist in the Malenco unit (Northern Italy). Serpentinites from this locality were dehydrated under quasi-static conditions in response to the intrusion of the Bergell tonalite (Evans & Trommsdorff, 1970, 1974a; Trommsdorff & Evans, 1972, 1974, 1977, 1980; Montrasio & Trommsdorff, 1993; Trommsdorff & Nievergel, 1983; Mellini et al., 1987; Worden et al., 2001; Trommsdorff et al., 2000, 2005). They offer therefore a unique opportunity to follow an undisturbed sequence of antigorite (Pfeifer et al., 1993) dehydration reactions at the kilometer scale and track the mechanisms of fluid extraction during these reactions.

**GEOLOGICAL SETTING**

The Malenco unit is located at the transition between the lower Austroalpine Margna and the upper Penninic Suretta nappes (Fig. 1a; Trommsdorff & Evans, 1972; Peretti et al., 1992; Müntener & Hermann, 1996; Hermann et al., 1997; Trommsdorff et al., 2005). It comprises a fossil lower crust–upper mantle section mainly composed of gabbros, peridotites, serpentinites, ophiocarbonates, and lower crust high-grade metapelites (Müntener & Hermann, 1996; Hermann et al., 1997). The ultramafic rocks from the Malenco unit are interpreted as part of the Adriatic subcontinental lithosphere exhumed by rifting in the Jurassic up to exposure at the seafloor of the Piemont Ligurian Ocean (see the dashed grey pressure–temperature path in Fig. 1b from Hermann, 1997; Müntener, 1997; Müntener et al., 1999).

During exhumation and exposure at the ocean floor, the ultramafic rocks were serpentinitized and ophicarbonates were formed in them, both within fractures and as platform sediments (Trommsdorff & Evans, 1977). The assemblage of ultramafic rocks, mafic rocks, and ophicarbonates was subducted to moderate depths in the late Cretaceous to early Tertiary, as recorded by regional Alpine metamorphism at greenschist- to epidote-amphibolite-facies conditions (Fig. 1b, peak temperature and pressure estimated at 450°C and 5.5 kbar; see Hermann, 1997; Hermann et al., 1997; Müntener, 1997; Trommsdorff et al., 2000). This subduction-related metamorphism led to transformation of the chrysotile-bearing serpentinitized peridotites to antigorite-schists with a mineral assemblage of Atq + OI + Di + Chl + Mag (abbreviations as given by Whitney & Evans, 2010) with Ti-clinohumite, brucite, and sulfides as accessory phases (Trommsdorff & Evans, 1974; Mellini et al.,
Fig. 1. (a) Geological setting of the Malenco Unit, Italian Alps (after Trommsdorff & Evans, 1974). Labelled lines P, B and OL are isograds for non-ultramafic lithologies (P, pumpellyite-out, coincident with the chrysotile/lizardite to antigorite reaction; B, biotite-in; OL, oligoclase-in) and numbers 1–4 are isograds for ultramafic lithologies (see text). Isograds 1a and 1b are from Peretti et al., (1992), where indentation indicates the side at which olivine is stable. Ultramafic rocks include peridotites and serpentinites. The studied area (Alpe Zocca) is indicated by a small black rectangle and is shown enlarged in (c). (b) A representative pseudosection in the CaO–FeO–MgO–Al2O3–SiO2–H2O system (fluid-saturated), showing isograds 1a and 1b related to the Cretaceous convergence (Alpine metamorphism) and isograds related to the contact metamorphism of the Bergell intrusion (Trommsdorff et al., 2000, and references therein). The reaction lizardite to antigorite and brucite is from Evans (2004). Also shown, as a dashed grey P–T path, is the pre-Alpine retrograde metamorphism. It should be noted that isograd 4 is not well defined, as anthophyllite occurs only sporadically in veins. CdA corresponds to the peak metamorphic conditions of Cerro del Almirez (Spain) discussed in the text. (c) Map of the studied area (from Trommsdorff et al., 2005, with slight modifications) indicating the position of the contact metamorphism isograds 2 and 3 (the latter corresponds to an ~150 m wide zone composed of variably reacted rocks with an irregular distribution at scales ranging from a few centimeters to a few meters), the studied dehydration veins, and sampling locations. The black line marks the profile along which the crystal preferred orientation (CPO) data presented in Figs 10, 11 and 13a have been projected.
Mineral assemblages containing anthophyllite and enstatite associated with reactions (4) and (5), respectively, occur sporadically within 100 m from the intrusion (Trommsdorff & Evans, 1972), thus making it difficult to trace the isograds (Trommsdorff & Connolly, 1996). They were not investigated in the present study.

\[ 2\, \text{Mg}_3\text{Si}_4\text{O}_10(\text{OH})_2 + \text{Mg}_2\text{Si}_4\text{O}_8 = \text{Mg}_7\text{Si}_8\text{O}_{22}(\text{OH})_2 + 3\, \text{H}_2\text{O}. \] (4)

\[ \text{Mg}_3\text{Si}_8\text{O}_{22}(\text{OH})_2 + \text{Mg}_2\text{Si}_4\text{O}_8 + 2\, \text{Mg}_2\text{Si}_2\text{O}_6 + \text{H}_2\text{O}. \] (5)

Figure 1b shows a temperature-pressure phase diagram (pseudosection) in the CaO–FeO–MgO–Al\text{2}O\text{3}–SiO\text{2}–H\text{2}O (CFMASH) system for a Ca-poor antigorite serpentine whole-rock composition (sample MA16-51A). A serpentinite hosting an Ol–Tlc vein, only the serpentinite wall-rock composition is used in the calculation (Table 1).

The phase diagram was computed using Perple_X (6.8.6, Connolly, 2009), the thermodynamic database of Holland & Powell (2011), the compensated-Redlich-Kwong (CORK) equation of state for the fluid and the solid solution models used by Padrón-Navarta et al. (2013; see their Table 1). The ferric iron is at present difficult to model because of the lack of a solid solution model accounting for Fe\text{3+} in antigorite, therefore the effective bulk rock was recast by subtracting 3 wt % of magnetite (e.g., López Sánchez-Vizcaino et al., 2005). Chlorite is an additional product phase of some of the reactions listed above owing to the solubility of aluminium in the reactant antigorite through a Tschermak’s exchange (e.g., Padrón-Navarta et al., 2013). Figure 1b also shows the high-pressure antigorite-out reaction, so far only reported in Cerro del Almirez, Spain (Trommsdorff et al., 1998; Padrón-Navarta et al., 2011):

\[ \text{Mg}_4\text{Si}_8\text{O}_{22}(\text{OH})_2 = 14\, \text{Mg}_2\text{Si}_4\text{O}_8 + 20\, \text{Mg}_2\text{Si}_2\text{O}_6 + 31\, \text{H}_2\text{O}. \] (6)

**METHODS**

Detailed sampling and structural mapping of a 1 km × 1 km area was undertaken in the northeastern part of the contact aureole of the Bergell intrusion, along the western side of the Val Ventina (Alpe Zocca, Northern Italy, Fig. 1a and c) to investigate the evolution of antigorite schists to olivine–talc (Ol–Tlc) metaperidotites and, in Ca-rich domains, of diopside-bearing antigorite schists to tremolite-bearing antigorite schists through reactions (2) and (3). Foliation planes (and when possible lineations) of serpentinites and metaperidotite were measured to map possible structural
Table 1: Representative whole-rock compositions and modal mineralogy of different types of ultramafic rocks in Alpe Zocca (Malenco unit)

<table>
<thead>
<tr>
<th>Type</th>
<th>Sample</th>
<th>S MA16-17</th>
<th>S+ MA16-51A</th>
<th>Tr-S MA16-36</th>
<th>S-r MA16-42</th>
<th>Vein MA16-51C</th>
<th>M MA17-34</th>
<th>M MA16-48</th>
<th>Js MA16-12A</th>
</tr>
</thead>
<tbody>
<tr>
<td>Major elements determined by XRF (wt %)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>SiO₂</td>
<td>42.02</td>
<td>41.66</td>
<td>45.97</td>
<td>45.94</td>
<td>44.23</td>
<td>46.24</td>
<td>42.36</td>
<td>43.59</td>
<td></td>
</tr>
<tr>
<td>TiO₂</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td></td>
</tr>
<tr>
<td>Al₂O₃</td>
<td>0.99</td>
<td>1.82</td>
<td>1.02</td>
<td>2.20</td>
<td>1.35</td>
<td>1.15</td>
<td>1.39</td>
<td>0.79</td>
<td></td>
</tr>
<tr>
<td>Cr₂O₃</td>
<td>0.36</td>
<td>0.41</td>
<td>0.43</td>
<td>0.39</td>
<td>0.35</td>
<td>0.47</td>
<td>0.53</td>
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</tr>
<tr>
<td>Fe₂O₃</td>
<td>2.53</td>
<td>3.59</td>
<td>4.16</td>
<td>0.87</td>
<td>2.61</td>
<td>3.08</td>
<td>4.97</td>
<td>2.16</td>
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<tr>
<td>FeO</td>
<td>4.46</td>
<td>2.38</td>
<td>2.54</td>
<td>4.20</td>
<td>4.48</td>
<td>3.90</td>
<td>5.25</td>
<td>7.12</td>
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<tr>
<td>MnO</td>
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<td>0.20</td>
<td>0.27</td>
<td>0.21</td>
<td>0.20</td>
<td>0.29</td>
<td>0.31</td>
<td>0.26</td>
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<tr>
<td>MgO</td>
<td>37.35</td>
<td>38.32</td>
<td>34.76</td>
<td>37.54</td>
<td>39.15</td>
<td>39.79</td>
<td>41.57</td>
<td>41.31</td>
<td></td>
</tr>
<tr>
<td>CaO</td>
<td>0.01</td>
<td>0.08</td>
<td>5.12</td>
<td>0.50</td>
<td>2.16</td>
<td>0.24</td>
<td>0.08</td>
<td>0.61</td>
<td></td>
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<tr>
<td>Na₂O</td>
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<td>0.00</td>
<td>0.02</td>
<td>0.02</td>
<td>&lt;0.01</td>
<td>&lt;0.01</td>
<td>&lt;0.01</td>
<td>&lt;0.01</td>
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<tr>
<td>LOI</td>
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<td>11.10</td>
<td>5.30</td>
<td>8.04</td>
<td>4.70</td>
<td>4.30</td>
<td>2.70</td>
<td>2.60</td>
<td></td>
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<tr>
<td>Mode (mass-balance wt %)</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Antigorite</td>
<td>98</td>
<td>94</td>
<td>19</td>
<td>51</td>
<td>—</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Olivine</td>
<td>—</td>
<td>2</td>
<td>29</td>
<td>20</td>
<td>39</td>
<td>47</td>
<td>68</td>
<td>70</td>
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<tr>
<td>Tremolite</td>
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<td>38</td>
<td>4</td>
<td>10</td>
<td>6</td>
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<tr>
<td>Chlorite</td>
<td>—</td>
<td>—</td>
<td>10</td>
<td>8</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>7</td>
<td></td>
</tr>
<tr>
<td>Talc</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>16</td>
<td>47</td>
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<td>20</td>
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<td>3</td>
<td>4</td>
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<td>3</td>
<td>2</td>
<td>4</td>
<td>1</td>
<td></td>
</tr>
</tbody>
</table>

*Ferric iron was calculated from the difference of the total iron measured by XRF and the measured ferrous iron.

**Types:** S, Atg-serpentinite; Tr-S, tremolite-bearing serpentinite; S-r, partially reacted serpentinite; M, pervasive metaperidotite; Js, jackstraw-textured Ol–Tlc vein. LOI, loss on ignition.

Petrographic analyses were carried out by optical microscopy to investigate microstructural relationships between antigorite and its dehydration products and the presence or not of viscoplastic deformation of the dehydration products. For geographically oriented samples, thin sections were prepared from sections oriented parallel to the vertical east–west plane looking from the south (i.e., the south direction points upward from the thin section surface and the long axis of the thin section is parallel to the east–west vertical plane). This orientation was chosen because the foliation predominantly dips towards the east in the study area. Non-oriented samples were prepared normal to the foliation plane, when visible. A few additional thin sections were prepared parallel to the foliation plane. All thin sections were polished with diamond paste, followed by colloidal silica for 4 h, and by a last stage of chemical polishing with colloidal silica in a vibratory polisher (Veritas Mineral Laboratory, Vancouver, Canada), using lithium borate fusion coupled to inductively coupled plasma emission spectrometry (ICP-ES). Additional analyses were conducted on selected samples to quantify FeO content (and Fe₂O₃ by difference) by the titration technique.

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Orientation data were expressed as Bunge Euler angles and processed with the MATLAB toolbox MTEX (Hielscher & Schaeben, 2008; Bachmann et al., 2010; Mainprice et al., 2014). Grain detection was performed with a mis-orientation threshold of 15°. Grains composed by <3 pixels were removed from the dataset. Orientation distribution functions for all phases were calculated using a ‘de la Vallée Poussin’ kernel with a constant half-width of 10°. Pole figures of antigorite and olivine were plotted in the lower hemisphere using the grain average orientation and a constant colour bar (0–4 multiples of a uniform distribution, m.u.d.) for easy comparison between samples. The intensity of the crystal preferred orientation (CPO) is characterized by the J-index (Bunge, 1982). The symmetry of the olivine CPO is characterized in terms of symmetry of the orientation distribution of the [010] and [001] axes by the BC-index. This index is calculated similarly to the BA-index (Mainprice et al., 2014) according to the following expression:

\[
\text{BC index} = \frac{1}{2} \left[ 2 - \left( \frac{P_{010}}{P_{010} + G_{010}} \right) - \left( \frac{G_{001}}{G_{001} + P_{001}} \right) \right] (7)
\]

where \( P \) and \( G \) are the point and girdle indices of Vollmer (1990), respectively. If the [010] axes of olivine define a perfect point concentration and the [001] axes form a girdle, the BC-index value is zero. If the [001] axes of olivine define a point concentration and the [010] axes form a girdle, the BC-index value is unity. CPOs with orthorhombic symmetry (point concentrations of both [010] and [001] axes) are characterized by BC-index values of 0.5.

FIELD OBSERVATIONS

Serpentinites and pervasive metaperidotites

Within the study area, from east to west, serpentinites evolve to fully reacted metaperidotites (Fig. 1c). The two isograds corresponding to reactions (2) and (3) crop out at c. 1230 m and at c. 770 m from the intrusion, respectively. The observed distance of the two isograds from the contact with the Bergell intrusion is consistent with the predictions by the numerical model of Trommsdorff & Connolly (1996): 1110–1310 m and 500–710 m, respectively.

The serpentinite is a greenish schistose fine-grained rock composed of antigorite with variable amounts of olivine, diopside, tremolite, magnetite, and chlorite (Trommsdorff & Evans 1972). The intensity of the schistosity varies spatially. Massive serpentinites with poorly defined foliation is not uncommon, but in most of the studied area a foliation could be measured in the field (Fig. 2). This foliation is marked by the shape-preferred orientation (SPO) of antigorite (Fig. 3a). Locally, magnetite forms elongated aggregates marking a lineation in the foliation plane. Metamorphic olivine appears as red–orange (on altered surfaces) or dark millimeter- to centimeter-scale crystals with rounded or prismatic shapes. In addition to olivine, serpentinites may contain metamorphic diopside, which typically occurs as white millimeter-scale prismatic crystals, or tremolite, which occurs as white acicular millimeter-scale crystals (Fig. 3a) replacing metamorphic diopside close to the contact with the pluton. The presence of diopside and tremolite indicates a locally more fertile (lherzolitic, Ca-rich) protolith. Spatial distribution of Ca-free and Ca-rich serpentinites does not show any regular arrangement. This observation is confirmed by larger scale mapping of the Malenco serpentinites (Lafay et al., 2019). Serpentinites with relict mantle clinopyroxene, which occurs as dark green millimeter-scale crystals (referred to as ‘dusty’ diallage or clinopyroxene by Trommsdorff & Evans (1972)), are common on the eastern side of the valley (Lago Pirola) and restricted to the eastern side of isograd 2 in our study area.

The strike of the serpentinite foliation gradually rotates from a NNW orientation steeply dipping (~80°) towards the ENE close to the isograd of reaction (2) to a more NNE strike orientation when approaching the isograd of reaction (3) (Fig. 2). Locally, foliations with opposite dip directions are observed; they are associated with mesoscale open folds or crenulations. The dip of the foliation plane in the serpentinites is opposite to the one determined by Hermann & Muntener (1996) on the eastern side of the Val Ventina valley (inset in Fig. 2), constraining the axis of a kilometer-scale fold (NNE sub-horizontal, subparallel to the trend of the Val Ventina; Trommsdorff et al., 2005). Lineations marked by the elongation of magnetite aggregates in serpentinites show variable dip (Fig. 2), in contrast to the lineations reported by Hermann & Muntener (1996) on the eastern side of the valley, which are dominantly NW sub-horizontal.

The serpentinites are progressively transformed to olivine–talc (± tremolite ± chlorite) metaperidotites through reaction (3) (Fig. 1b and c). Ol–Tlc (±Tr ±Chl) rocks are less schistose and more massive than antigorite schists, but a weak foliation can still be recognized in the field. The transition between serpentinites and metaperidotites is smooth. It occurs over a c. 150 m wide domain (indicated by the colour gradient in Fig. 1c), composed of partially dehydrated rocks characterized by layers, patches, or irregularly shaped septa of Ol–Tlc (±Tr ±Chl) rocks intercalated with serpentinites (Fig. 3b).

Metaperidotites are defined by the presence of more than 90 vol. % of product phases from dehydration reaction (3): Ol–Tlc (±Tr ±Chl). They appear as dark grey roughly foliated rocks in fresh surfaces, but have reddish to whitish tones on altered surfaces (Fig. 3c and d). Fine-grained metaperidotites (with submillimetric olivine) predominate. These rocks often show a rough compositional layering marked by alternating Ol-rich and Tlc-rich domains (Fig. 3c). They may display a foliation owing to the shape-preferred orientation (SPO) of olivine and a lineation marked by the alignment of magnetite aggregates (Fig. 3c). Locally, metaperidotites are
coarse-grained containing centimeter-scale olivine crystals associated with white talc flakes and dark grey chlorite crystals (Fig. 3d). Coarse-grained metaperidotites are usually foliated. No clear spatial arrangement between fine-grained and coarse-grained metaperidotite as a function of the distance from the Bergell intrusion was observed within the study area. This observation contrasts with the increase in grain size with increasing distance from the Bergell intrusion proposed by Lafay et al. (2019) based on larger scale mapping of the Malenco serpentinites. The strike of the metaperidotite foliation is remarkably similar to that of the serpentinites close to the Ol–Tlc isograd (Fig. 2). The lineation tends, however, to have steeper plunges in the metaperidotites than in serpentinites (Fig. 2).

Metamorphic (dehydration) veins
A striking feature of the study area (Alpe Zocca) is the common occurrence of olivine-bearing veins with variable sizes and shapes (Trommsdorff & Evans, 1974; Trommsdorff & Connolly, 1996). Two types of olivine-bearing veins have been identified: Ol–(±Tr) veins and Ol–Tlc veins, which are concentrated just before (downstream of) the isograds of reactions (2) and (3), respectively (Figs 1c and 2). The veins are dominantly oriented N60–70° with a subvertical dip; they most often crosscut the serpentinite foliation (Fig. 2).

The most distant veins from the intrusion are Ol–(±Tr) veins (Figs 1c and 4a). These veins are centimeters to decimeters wide and may usually be followed for a few meters. They contain olivine and occasionally...
Ol + Tr, most often as centimetric crystals, although millimeter-size crystals also occur. At the outcrop scale, multiple subparallel or interconnected veins with a spacing of tens of centimeters are observed with sharp limits at the outcrop scale (Fig. 4a). Locally conjugate pairs of veins were observed. Ol–Tlc veins are notably observed in the serpentinites just downstream of the Tlc-in isograd and in the transition domain, which is characterized by partial (patchy) transformation of the serpentinites into Ol–Tlc metaperidotites [reaction (3); Fig. 1c]. In the field, Ol–Tlc veins appear dominantly reddish, with white patches between olivine grains corresponding to talc (Fig. 4b–f). Ol–Tlc veins are tens of centimeters wide and several meters long. Some veins display an outward growth of tabular olivine crystals oriented perpendicular to the vein trend (Fig. 4b). Ol–Tlc veins have a wide range of morphologies from channel-like branching shapes to more planar structures (Fig. 4b and c). Locally, they occur as multiple subparallel veins, forming locally en echelon overlaps, or have criss-crossing structures with conjugate or more complex configurations. The thinnest Ol–Tlc veins are formed by discontinuously aligned centimeter-long olivine crystals; this arrangement results in a boudinaged-like aspect of the vein, but olivine crystals in the vein have prismatic shapes (Fig. 4c).

Ol–Tlc veins display a wide variation in morphology and size of olivine crystals. However, most Ol–Tlc veins are composed of coarse (>1 cm) prismatic to blade-like olivine crystals, with the long axes arranged in criss-cross, radial, or parallel patterns within the vein plane (Fig. 4e). This texture has been previously described as ‘jackstraw olivine’ texture (Evans & Trommsdorff, 1974a; Snoke & Calk, 1978; Bakke & Korneliussen, 1986). It is most characteristic when viewed in the plane of the vein, where the long axes of the olivine crystals are concentrated (Fig. 4e). The coarsest olivine grains observed in the study area are up to 20 cm long, but olivine crystals up to 1 m long, associated with talc and magnesite, have been reported in the Malenco ultramafic unit by Evans & Trommsdorff (1974a). Veins with significantly smaller olivine sizes (0.5–1 cm), but also

Fig. 3. Representative mesoscopic textures. (a) Tr-bearing serpentinite with a poorly defined foliation (sample MA16-34). (b) Partially reacted serpentinite from the ∼150 m wide reaction front between the serpentinites and pervasive metaperidotites with Ol–Tlc patches of variable dimensions, from several tens of centimeters wide (below the hammer) to a few centimeters wide in the right side of image. (c) Fine-grained Ol–Tlc metaperidotite with a weak lineation marked by the elongation of magnetite aggregates (sample MA17-39). (d) Coarse-grained Ol–Tlc–Chl metaperidotite with a coarse foliation and compositional layering marked by alignment of talc-rich domains and a shape-preferred orientation of olivine (MA16-48).

Ol + Tr, thus most often as centimetric crystals, although millimeter-size crystals also occur. At the outcrop scale, multiple subparallel or interconnected veins with a spacing of tens of centimeters are observed with sharp limits at the outcrop scale (Fig. 4a). Locally conjugate pairs of veins were observed.
displaying a jackstraw texture, have also been observed. In the metaperidotites, some structures are interpreted as Ol–Tlc veins based on their coarser olivine grain size and the change in texture (mainly the olivine SPO, which is jackstraw-like) relative to the host metaperidotite (Fig. 4f).

In addition, Ol–Tlc veins are commonly texturally composite. Composite Ol–Tlc veins are characterized by a central domain in which olivine has a jackstraw texture surrounded by a reaction zone in which olivine crystals are elongated at high angle to the vein, but parallel to the serpentinite foliation in the wall-rock. These reaction zones may be one olivine crystal wide (≤1 cm, Fig. 4b) up to 10–20 cm wide (Figs 4d and 5). The wider ones may display finger-like shape limits (Fig. 4d) or be planar at the hand sample scale (Fig. 5). Reaction fronts are texturally similar to pervasive metaperidotites. More rarely, Ol–Tlc veins display a jackstraw-textured domain in sharp contact to the serpentinite, with no associated reaction front. The contact with the serpentinite is irregularly shaped, but no more than a few millimeters wide.

PETROGRAPHY
Serpentinites and pervasively reacted metaperidotites
Ca-free serpentinites are mainly composed of fine-grained antigorite crystals (Fig. 6a and b). They also
contain isolated olivine porphyroblasts with irregular, but sometimes elongated shapes (up to 2 cm long). The SPO of antigorite and occasionally the elongation of olivine porphyroblasts marks the foliation. Serpentinites also contain magnetite crystals, which form elongated aggregates marking a lineation.

Ca-rich serpentinites are also foliated (Figs 3a and 6c). At the microscopic scale the foliation is marked by the orientation of antigorite and of acicular crystals of tremolite, which are ~200 μm long. Olivine in Ca-rich serpentinites occurs as aggregates of rounded fine-grained crystals (<10 μm in diameter) or as larger porphyroblasts (>200 μm long) with irregular shapes (Fig. 6d). Locally, crystals of olivine in textural equilibrium with tremolite are observed (Fig. 6d), indicating that they result from reaction (2). Magnetite forms aggregates, which have highly variable shapes and are sometimes aligned, marking a lineation (Fig. 6c).

Partially reacted serpentinites are composed of variable amounts of antigorite and Ol + Tlc ± Tr in diffuse patches. They often contain coarse olivine crystals (up to 1 cm long and 300 μm wide) either isolated or in aggregates. Both the individual olivine crystals and the aggregates are elongated in the foliation plane defined by the antigorite shape-preferred orientation (Fig. 6e and f). Talc occurs either associated with olivine (Fig. 6f) or as diffuse layers parallel to the foliation in the

Fig. 5. (a) Macroscopic image and photomicrographs of a composite Ol–Tlc vein with a jackstraw-textured central domain, a wide reaction zone (selvage), and a centimeter-wide fingered contact with the serpentinite wall-rock (MA17-47). The reaction zone displays elongated olivine marking a foliation parallel to the foliation of the serpentinite wall-rock. (b) Pole figures showing the crystal preferred orientation of antigorite in the serpentinite wall-rock and of olivine in the reaction zone (contours) and in the jackstraw-textured central domain of the vein (red points). Inverse pole figure showing the rotation axes accommodating intragranular misorientations in olivine within the reaction front.
antigorite matrix. Olivine crystals do not display fully developed crystallographic faces, but have highly irregular grain boundaries (Fig. 6f). They contain common antigorite or tremolite inclusions and are sometimes poikilitic.

Metaperidotites have highly variable microstructures, characterized by changes in shape and size of olivine and in the modal proportion of talc. Most often, metaperidotites are composed of a fine-grained assemblage of small prismatic olivine crystals (~20 µm, aspect ratio <2 on average) and talc flakes, whose alignment marks a poorly developed foliation (Fig. 7a). Magnetite aggregates up to 5 mm long are aligned in the foliation, marking a lineation (Fig. 7a). These aggregates are composed of rounded magnetite crystals intercalated with domains enriched in olivine and/or talc, which are oriented normal to the aggregate elongation, similar to extensional cracks (Fig. 7b). Antigorite-rich patches, usually a few millimeters wide, are observed in many metaperidotites at the thin-section scale.

Variations in olivine grain size and shape (probably associated with the orientation of the crystal) occur in a diffuse manner at the millimeter to centimeter scale, as illustrated in Fig. 7c, where a fine-grained metaperidotite with olivine <20 µm in diameter grades into a medium-grained peridotite with olivines >100 µm long.
Fig. 7. Optical photomicrographs under cross-polarized light of representative textures of pervasive metaperidotites from Alpe Zocca (a, c, e, g) and corresponding enlarged areas (b, d, f, h). (a) Fine-grained Ol–Tlc metaperidotite with aggregates of magnetite crystals elongated in the foliation plane (S) (MA17-34). (b) Detail of magnetite aggregates showing extensional crack-like features filled by talc. Olivine grain size is on average ~20 µm. (c) Medium-grained Ol–Tlc metaperidotite with a foliation affected by open folds and a bimodal olivine grain-size distribution (either higher or lower than 200 µm in length) (MA17-31). (d) Medium-grained elongated olivine with an SPO marking the foliation plane. Talc crystals are occasionally bent between olivine grains. (e) Medium-grained pervasive metaperidotite with elongated olivine crystals defining a foliation plane parallel to the compositional layering defined by alternating olivine-rich and talc-rich bands. (f) Detail illustrating the coexistence of undulose extinction and irregular overgrowths in olivine grains (MA17-36). (g) Coarse-grained Ol–Tlc metaperidotite (sample MA16-48, shown in Fig. 3d). Olivine crystals show distinct undulose extinction. (h) Detail showing the serrated, dendritic-like overgrowths at the edge of olivine crystals as well as chlorite and talc flakes bent around olivine crystals.
that are often slightly elongated, marking a poorly defined foliation. Metaperidotites with medium-grained olivine (millimeter-size crystals) frequently have a clear foliation marked by the orientation of prismatic olivine crystals and talc flakes (Fig. 7e). Coarse-grained metaperidotites are composed of centimeter-long olivine crystals with prismatic or, more rarely, skeletal shapes flattened in the foliation plane (Fig. 7g). Independently of grain size, olivine crystals usually show undulose extinction and talc or chlorite crystals are often bent around olivine grains (Fig. 7d and h). Olivine crystals have fingering contacts with talc, owing to millimeter-wide dendritic-like overgrowths (Fig. 7f and h), but more regular contacts with other olivine crystals (Fig. 7g).

Metamorphic veins
Ol–(±Tr) veins have usually sharp contacts with the surrounding serpentinite at the macroscopic scale, but the contacts are always diffuse at the thin-section scale. Ol veins are mainly composed of centimeter-scale olivine crystals (Fig. 8a), but occasionally they may be constituted of fine-grained rounded to polygonal olivine crystals. In thin (centimeter-wide or smaller) coarse-grained veins, olivine crystals in the centre of the vein are elongated parallel to its trend, whereas olivine crystals growing outward from the vein are dominantly elongated at high angles to the vein trend and parallel to the foliation of the serpentinite (Fig. 8b).

Jackstraw-textured Ol–Tlc veins are characterized by prismatic, blade-like or, more rarely, skeletal olivine crystals arranged in a criss-cross or radial pattern (Fig. 8c–f). Antigorite is never present in jackstraw-textured Ol–Tlc veins. Olivine crystals usually show fingering contacts with talc (dendritic-like overgrowths, Fig. 8d and f). Olivine–olivine contacts tend to be polygonal (Fig. 8c). Olivine crystals tend to be coarser (pluri-centimetric) in veins containing interstitial carbonates (Figs 4d and 8c), as noted by Evans & Trommsdorff (1974a). Olivine in carbonate-free veins is most often millimetric to centimetric (Fig. 8d–f), but locally very large prismatic crystals may be observed (Fig. 8f). Talc is typically fine-grained and randomly oriented (Fig. 8d–f). Olivine crystals are usually undeformed (no undulose extinction, nor sub-grains). In some veins, olivine grains display locally undulose extinction, but preserve the fingering grain boundaries with talc (Fig. 8e and f). In these veins, talc crystals are bent around olivine (Fig. 8e and f). Ol–Tlc veins enclosed in the coarse-grained metaperidotites of the pervasively reacted domain share many similarities with those in the serpentinites and partially reacted serpentinites, but their limits are difficult to define (Fig. 8f).

BULK-ROCK COMPOSITIONS
Bulk-rock compositions of representative samples (Table 1) are displayed in a projected MgO–SiO₂–H₂O (MSH) ternary diagram using CSpace (Torres-Roldán et al., 2000). Mineral and bulk-rock analyses were projected from the average composition of tremolite, chlorite, and magnesite and the exchange vector Fe₁₋₂Mg₁, Mn₁₋₂Mg₁ and Ni₁₋₂Mg₁ (Fig. 9). In addition to measured bulk-compositions, the projections also include calculated bulk compositions based on modal compositions estimated from the EBSD maps and on mineral compositions measured by EPMA (average composition of antigorite, olivine, talc, tremolite, chlorite, and diopside for representative samples is shown in Supplementary Data Tables 1S–6S; supplementary data are available for downloading at http://www.petrology.oxfordjournals.org). Measured and calculated compositions for serpentinites plot close to the antigorite pole (Fig. 9); the dispersion results from variable olivine contents. Pervasive metaperidotites plot on the talc–olivine tie-line (Fig. 9). The observed range in talc and olivine proportions (i.e., relative proportion of the projected components MgO vs SiO₂) is within the projected MgO/ SiO₂ ratio in the serpentines (antigorite with a variable proportion of olivine; blue field in Fig. 9). Jackstraw-textured Ol–Tlc veins globally plot in the same field as the pervasive metaperidotites. Some reactional walls of large veins such as the one illustrated in Fig. 5 show evidence for SiO₂ depletion, which moves the bulk-rock composition along the antigorite–olivine tie-line (Fig. 9). Such zones can be interpreted as selvages or alteration halos characteristic of extreme fluid channelization (e.g. Ague, 2011). Calculated compositions of partially reacted serpentinites plot inside the Atg–Tlc–Ol field (Fig. 9). Some of these lithologies might correspond to a true divariant (or higher) mineral assemblage in the full chemical system, but in most cases the three-phase assemblage is simply related to an incomplete dehydration of antigorite (i.e. heterogeneous distribution of metastable antigorite patches or septa; see Frost, 1975).

CRYSTAL PREFERRED ORIENTATIONS
Serpentinites and pervasive metaperidotites
An outstanding feature of the study area is the consistency of the orientation of the foliation in the serpentinites and in the pervasive metaperidotites. We investigate the potential crystallographic inheritance from antigorite to olivine by comparing the CPOs of both minerals at the thin-section scale in samples recording different degrees of transformation through reactions (2) and (3). In the serpentinites and partially reacted serpentinites, antigorite is dispersed in the matrix (Fig. 6a–f). In the metaperidotites, it occurs as discontinuous patches or septa in the dominantly Ol–Tlc ± Tr rock-mass. Figure 10 illustrates the evolution of the relationship between antigorite and olivine CPOs along a profile approximately normal to the contact with the Bergell intrusion, shown as a black line in Fig. 1c. All pole figures were projected in the geographical reference frame (inset in Fig. 10). When measured in the field, the orientation of the foliation is indicated
by a continuous line in the [001]_Atg and [010]_Ol pole figures. The plane normal to the [001]_Atg and [010]_Ol maxima is indicated as a dashed line for comparison.

Antigorite and olivine in all studied samples have a clear CPO. The strength of the CPO of the two minerals is usually correlated. Weak antigorite CPOs are associated with weak olivine CPOs and vice versa (Fig. 10). Antigorite in the serpentinites and in small unreacted patches in the pervasive metaperidotites shows similar CPOs (Fig. 10). [001]_Atg is concentrated normal to the foliation, but with some dispersion in the plane normal to the [010]_Atg maximum. [010]_Atg is dispersed in a girdle subparallel to the foliation plane with a maximum in this plane. [100]_Atg usually shows more dispersed orientations, but also tends to form a girdle subparallel to the foliation plane. When a lineation could be inferred from the orientation of magnetite aggregates, it did not show a clear correlation with either [100]_Atg or [010]_Atg.

In most talc-bearing metaperidotites, olivine displays a strong concentration of [010]_Ol axes at low angle to the [001]_Atg maximum and at high angle to the foliation, when the latter could be measured in the field (Fig. 10).
[001]$_{\text{Ol}}$ and [001]$_{\text{Ol}}$ axes usually form girdles normal to the [010]$_{\text{Ol}}$ maximum, with poorly defined mutually orthogonal maxima within the girdle (e.g. samples MA17-31 and MA16-45 in Fig. 10). This CPO pattern corresponds to relatively low BC-indices ($<$0.35, Fig. 11a). Other samples (e.g. MA16-43 in Fig. 10) display more orthogonal olivine CPO patterns, characterized by three orthogonal maxima, resulting in higher BC-indices ($>$0.5, Fig. 11a). In these samples, [001]$_{\text{Ol}}$ is usually better oriented than [100]$_{\text{Ol}}$. The remainder of the samples have olivine CPO patterns intermediate between these two end-members. This CPO is associated with an SPO: olivine crystals are usually prismatic, elongated parallel to [001] and flattened parallel to [010]. However, the difference in elongation between [010] and [100] is probably small, as crystals oriented with [001] normal to the thin section are roughly isometric. When present in pervasive metaperidotites (e.g. MA17-31), tremolite has [001] axes forming a girdle in the foliation plane and [100] axes aligned normal to the foliation plane, parallel to the [010]$_{\text{Ol}}$ maximum.

In talc-free serpentinities, the relation between olivine and antigorite CPO is variable. Serpentinite MA16-08, sampled close to the isograd from reaction (3), shows, like the metaperidotites, concentration of [010]$_{\text{Ol}}$ axes at low angle to the [001]$_{\text{Asg}}$ maximum (Fig. 10). However, serpentinites MA16-36 and MA16-34, sampled farther from the Bergell intrusion contact, in which olivine probably results from reactions (2) and/or (1b), respectively, have [001]$_{\text{Ol}}$ axes, rather than [010]$_{\text{Ol}}$, oriented subparallel to the [001]$_{\text{Asg}}$ axes.

There is neither correlation nor anti-correlation between the strength of olivine CPO and the distance to the Bergell intrusion (Fig. 10). This observation is corroborated by the analysis of a larger number of metaperidotites (30 samples) with variable grain sizes, which shows that neither the intensity of the olivine CPO (J-index) nor its symmetry (BC-index) varies in a systematic way with increasing distance to the intrusion (Fig. 11). Axial-[010] olivine CPO (BC-index <0.35) and orthorhombic olivine CPO (BC-index between 0.35 and 0.65) predominate (Fig. 11a). Axial-[001] textures (BC-index >0.65) are rare. There is also no clear relation between the olivine CPO intensity or symmetry and grain size, although fine-grained metaperidotites tend to have dominantly weak fabrics (J-index < 2, Fig. 11b).

**Metamorphic (dehydration) veins**

Figure 12 illustrates the CPO of olivine and, when present, tremolite, in dehydration veins as well as the antigorite CPO in the serpentinite from the wall-rock at the contact with the vein. There is no correlation between the CPO of the antigorite outside the vein and the olivine or tremolite CPO in the vein, except in those rare cases where the vein cuts the serpentinite foliation at low angle (e.g. MA17-16).

In talc-free veins, olivine grains are either weakly oriented (MA17-16) or, because of the coarse grain sizes, the number of grains analysed is too low (MA17-47 in Fig. 5 and MA17-10 and MA17-17 in Fig. 12) to define a representative CPO (the multiple maxima correspond to large individual crystals, with some growth-related internal mis-orientation). However, the [010] axes of olivine tend to align at high angle to the plane of the vein (indicated by a continuous great circle in the [010]$_{\text{Ol}}$ pole figures in Fig. 12), whereas [100] and [001] axes form wide girdles at low angle to this plane. This relation between olivine CPO and vein orientation is better...
developed in olivine in jackstraw-textured Ol–Tlc veins (Fig. 12). It is consistent with the observation that jackstraw olivines are elongated in the plane of the vein, but have their long axis, which is usually parallel to [001], oriented in a criss-cross pattern within the plane of the vein (Fig. 4d and e).

The change in olivine CPO and SPO patterns between the veins and the pervasive metaperidotite is reproduced at smaller scale in the composite Ol–Tlc veins. Figure 5 illustrates the variation in microstructures and olivine and antigorite CPO and SPO in a well-developed composite vein characterized by a central

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Fig. 10. Pole figures of antigorite and olivine from representative samples along the profile shown in Fig. 1c. Shaded pole figure is for tremolite. Orientations (average orientation of the grain) were plotted in lower hemisphere projections in the geographical reference frame contoured at one multiple of uniform distribution intervals. The continuous line indicates foliation planes measured in the field, whereas dashed lines mark the planes normal to the [001]Atg and [100]Ol or [010]Ol maxima. S, Sr and M refer to serpentinite, partially reacted serpentinite, and metaperidotite, respectively. Phase proportions of each sample are indicated by the grayscale bars at the right of the stereoplots. Frames highlight the correlated axes of antigorite and olivine. The number of grains included is indicated as thousands (k). It should be noted that for olivine, the profusion of inclusions makes difficult to correctly identify the grains, therefore the number of grains is probably overestimated.
The foliation plane in the reaction zone rotates by \( \angle_{C24} \) extinction. The rotation axes accommodate this trend of the vein. Similarly to some pervasive metaperidotites, fingered contact with the serpentinite (Fig. 5a). Millimeter scale, the reaction front of the vein has a dislocation. Samples were classified as fine-grained (average olivine grain size <200 \( \mu \)m), medium-grained (between 200 \( \mu \)m and 1 cm), and coarse-grained (>1 cm).

Domain with irregular width (up to 2 cm) composed of coarse-grained Ol–Tlc–Tr with a jackstraw texture surrounded by a reaction zone c. 10 cm wide. The reaction zone is formed by a foliated metaperidotite composed of tabular olivine crystals intercalated with talc. The foliation in the metaperidotite, defined by the SPO of olivine and talc, is parallel to the foliation plane defined by the SPO of antigorite in the serpentinite wall-rock and both are highly oblique to the vein trend (Fig. 5a). At the millimeter scale, the reaction front of the vein has a diffuse, fingered contact with the serpentinite (Fig. 5a). The foliation plane in the reaction zone rotates by \( \sim 20^\circ \) from the contact to the serpentinite to the centre of the vein. EBSD analyses in the serpentinite and in the reaction zone show strong CPO of both antigorite and olivine, with a similar orientation of the [001]_{Agt} and of the [010]_{Ol} axes normal to the foliation in the serpentinite and metaperidotite, respectively (Fig. 5b). The low number of olivine grains that could be analyzed in the central part of vein, which has a jackstraw texture, does not allow definition of a statistically meaningful CPO, but [010]_{Ol} axes are dominantly at high angle to the general trend of the vein. Similarly to some pervasive metaperidotites, olivines in the reaction zone show undulose extinction. The rotation axes accommodating this intracrystalline mis-orientation are \(<u0w>\) crystal axes, with some predominance of [100] (Fig. 5b).

**Olivine intracrystalline misorientation**

Variations in orientation of the crystalline lattice within a crystal (\( \leq 15^\circ \)) result essentially from stored dislocations. Clear intragranular mis-orientations around coherent rotation axes are indicative of deformation by dislocation creep. Thus, we analysed the variation of the mean (at the sample scale) intragranular mis-orientation of olivine in pervasive metaperidotites and in veins as a function of the grain size and of distance to the contact with the Bergell intrusion, as well as the rotation axes accommodating these mis-orientations to characterize the potential viscoplastic deformation of olivine (Fig. 13). Mean intragranular mis-orientations \( \geq 4^\circ \) are observed only within 900 m from the contact with the Bergell intrusion, in the domain affected by reaction (3). However, the lack of high mean intragranular mis-orientations farther from the intrusion may be a bias from the less dense sampling of olivine-rich rocks in this domain. Otherwise the mean intragranular mis-orientation in olivine seems to be uncorrelated with distance from the intrusion. Coarse- and medium-grained metaperidotites (Fig. 13c) have olivine with highly variable average intragranular mis-orientations (from 1 to 6°). Fine-grained metaperidotites have a lower range of average mis-orientations (from 1 to 3°), but these lower values may result from the lower number of orientation data per grain. Olivine crystals in veins have average intragranular mis-orientations in olivine overlapping with those of medium- to coarse-grained pervasive metaperidotites, but average intragranular mis-orientations in olivine in veins are on average lower (Fig. 13a). The spatial distribution of intragranular mis-orientations in olivine in the veins is also usually more heterogeneous (coexistence of deformed and undeformed grains; Fig. 13d) than in the pervasive metaperidotites (Fig. 13c). The mis-orientation data are consistent with petrographic analyses: olivine in metaperidotites and veins displays a large range of microstructures, from barely free of intragranular deformation (Fig. 8c and d) to well-developed undulose extinction associated with bent talc crystals (Figs 7e–h and 8e, f). The rotation axes accommodating intragranular mis-orientations in olivine in pervasive metaperidotites and in veins usually differ (Fig. 13b). They are dominantly \(<u0w>\) with a maximum at [100] in pervasive metaperidotites, similar to those in the reaction zones of composite veins (Fig. 5), whereas in jackstraw-textured domains of Ol–Tlc veins, rotation axes accommodating intragranular mis-orientation in olivine are mainly \(<0vw>\) with predominance of [010] axes. Some pervasive metaperidotites have rotation axes accommodating intragranular mis-orientations in olivine intermediate between these two dominant cases, with two maxima, one parallel to [100] and the other parallel to [010].

![Fig. 11.](image-url) (a) Symmetry (BC-index) and (b) strength (J-index) of the olivine CPO as a function of distance from the Bergell intrusion. Samples were classified as fine-grained (average olivine grain size <200 \( \mu \)m), medium-grained (between 200 \( \mu \)m and 1 cm), and coarse-grained (>1 cm).
DISCUSSION

A major observation in the study area is the occurrence of the prograde assemblages formed by reaction (2) and notably by reaction (3) in two contrasting modes: (1) pervasive metaperidotites, most often weakly foliated, separated from the unreacted serpentinites by a c. 150 m wide zone characterized by a patchy mixture of variably reacted rocks; (2) downstream of this ‘isograd’, metaperidotites within dehydration (replacement) veins, which often have centimeter- to decimeter-scale reaction fronts or selvages propagating towards the unreacted serpentinite wall-rocks. Microstructures and CPO of the reaction products in the two modes of occurrence differ. We interpret the variation in mode of occurrence and in microstructure of the products of the antigorite dehydration reactions as resulting from different fluid extraction processes and associated reaction conditions. Mode (1) records the progressive and pervasive transformation of the antigorite schists into metaperidotite at near equilibrium conditions, whereas mode (2) records local displacement of the reactions towards lower temperature conditions owing to the 

Fig. 12. Olivine pole figures from representative veins and antigorite pole figures from wall-rock serpentinite. Shaded pole figure is for tremolite. Orientations (average orientation of the grain) were plotted in lower hemisphere projections in the geographical reference frame contoured at one multiple of uniform distribution intervals. For the Oi–Tr vein, olivine orientations were not contoured because too few grains could be measured owing to the coarse grain size. The continuous black line indicates the orientation of the plane of the vein measured in the field and the green line marks the foliation plane in the serpentinite wall-rock.
formation of extensional veins, which acted as high-permeability channels allowing for effective draining of the system.

**Pervasive metaperidotites: fluid drainage by viscous compaction**

Pervasive peridotites formed by progressive and pervasive transformation of the antigorite schists into metaperidotite at near equilibrium conditions, as indicated by the development of a 150 m wide partially reacted domain (Fig. 1c) composed of variably reacted rocks with an irregular, patchy distribution at scales ranging from a few centimeters to a few meters (Figs 3b and 6e, f), and by the systematic preservation of small patches of antigorite in the metaperidotites (Fig. 7c). Growth of elongated olivine over schistose serpentinite and persistence of unreacted serpentine septa over 20 m were also documented in low-pressure contact metamorphism of ultramafic rocks from Paddy-Go-Easy pass in the Central Cascades (Frost, 1975). The persistence of unreacted serpentinite is never observed in the dehydration veins.

In a recent experimental dehydration study (gypsum to bassanite reaction), Leclère et al. (2018) showed that high effective pressure resulted in sharp dehydration fronts separating fully reacted from non-reacted rocks, whereas wide reaction fronts preserving significant amount of unreacted gypsum are produced when the effective pressure is low. Low effective pressure during viscous metamorphic compaction would result in reaction kinetics close to equilibrium, because the reaction affinity remains low when fluid pressure approaches lithostatic pressure (Dahlen, 1992; Padron-Navarta et al., 2011; Dilissen et al., 2018). Grain growth is then controlled by interface-coupled dissolution–precipitation processes (Putnis, 2002; Putnis & Putnis, 2007). This may explain the parallelism between the foliation in the serpentinites and metaperidotites (Fig. 2), which results in a consistent fabric at the kilometer scale, across the reaction front. It may also account for the CPO relations between olivine and antigorite in the metaperidotites (Fig. 10), which are consistent with those observed during topotaxial replacement of olivine by antigorite (Boudier et al., 2010).

Under such conditions, viscous compaction has been proposed as an efficient draining mechanism (Connolly, 1997, 2010). The pervasive metaperidotites show evidence for solid-state deformation following their crystallization. Olivine displays undulose extinction (Figs 7 and 13c). Talc and chlorite crystals are bent

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**Fig. 13.** (a) Mean intragranular olivine mis-orientation as a function of the distance from the Bergell intrusion. Veins are included in the plot; they are mostly coarse-grained. No correlation is observed with distance from the intrusion. (b) Inverse pole figures of intragranular mis-orientation axes for olivines (2–10') from a representative pervasive metaperidotite (MA16-45) and from a jackstraw-textured Ol-Tlc vein (MA16-51C). (c, d) Intragranular mis-orientation maps for olivine (mis-orientation relative to the mean orientation M2M) for the two samples.
(Fig. 7d and h). However, olivine crystals preserve prismatic shapes and even the delicate fingering contacts with talc (Fig. 7). The latter observation suggests that the solid-state deformation is weak and cannot be responsible for the foliation displayed by these rocks (Figs 3c, d and 7). Yet local variations in foliation orientation observed at the outcrop and sample scale may have resulted from macroscopic-scale compaction. Similar deformation microstructures were documented and attributed to syn-metamorphic viscous compaction in metaperidotites composed of olivine + enstatite + chlorite (± tremolite) with granofels texture in Cerro del Almirez (Padrón-Navarta et al., 2011, 2015; Dilissen et al., 2018). These metaperidotites, formed by antigorite dehydration at higher pressures and temperatures (16–19 kbar and 680 °C, Fig. 1b), display additional evidence for viscous compaction such as shear-induced inversion of orthoenstatite to low clinoenstatite, which allow the stresses involved in this process to be constrained to a few tens of MPa (Padrón-Navarta, 2015; Clément et al., 2018). These stress levels are consistent with predictions of numerical models of viscous metamorphic compaction during dehydration (Connolly, 1997).

Weak deformation of olivine by dislocation creep under such low stresses, and at the low-temperature conditions inferred for the study area (500–600 °C), might be possible, as the metamorphic olivine crystalized under fluid-saturated conditions and had initially high densities of point and planar defects (clinohumite lamellae and/or talc inclusions), but low dislocation densities. Moreover, at the beginning of compaction, stresses may be locally high, as the contact area between olivine grains is small. Stresses decrease with increasing compaction as the contact area increases. Intragranular mis-orientation in olivine in the metaperidotites is associated with rotations around <u0w> axes, mainly [100], and less often around [010], which is nevertheless the dominant rotation axis accommodating intragranular mis-orientations in vein olivine (Fig. 13b). At low temperature and high stresses and in wet conditions, the dominant slip systems in olivine are [001](110) and [001](100); accumulation of dislocations in these systems should produce rotations around [100] and [010] (e.g. Raleigh, 1968; Mackwell et al., 1985; Demouchy et al., 2013, 2014; Idrissi et al., 2016).

### Dehydration Veins: Reaction Triggered by Deformation-Induced Chemical Disequilibrium

Replacement veins resulting from dehydration reactions in open systems have long been known to form at lower temperatures than those expected for the same reactions to occur in a closed system, where the H2O fluid pressure equals the confining pressure (e.g. Fyfe et al., 1958; Coombs et al., 1959; Bruton & Helgeson, 1983; Coombs 1993). Dehydration can indeed proceed at isothermal and isobaric conditions (constant lithostatic conditions) if fluid pressure is allowed to decrease (e.g. Fyfe et al., 1958). The chief variable in such open systems is the chemical potential of the H2O component, which needs to be lower than what corresponds to fluid saturation for the given confining pressure (Bruton & Helgeson, 1983). The seminal work of Greenwood (1961) on the dehydration of analcime using Ar as inert gas to reduce the chemical potential of H2O is an early and elegant experimental confirmation of this model. Joint-controlled replacement reactions resulting in dehydration of the host-rocks at the centimeter to decimeter scale are indeed not unusual in the brittle crust (e.g. Coombs et al., 1959; Coombs, 1993). They have also been described in eclogite-facies conditions as ‘prograde-metamorphic dehydration fluid conduits’ or dehydration veins owing to an influx of an H2O-unsaturated fluid (e.g. Beinlich et al., 2010; John et al., 2012; Taetz et al., 2016, 2018).

The most likely origin for the dehydration veins observed in the present study is a sudden change in the H2O chemical potential of the system as the result of the formation of extensional fractures, which produced a local pressure decrease and hence a chemical potential gradient. The reduction of the H2O chemical potential in the crack may have been triggered by either (1) infilling of the fracture by external fluids rich in other components such as CO2 (e.g. Evans & Trommsdorff, 1974b) or (2) release of H2O-aqueous fluid by dehydration of antigorite, aiming to homogenize the chemical potential gradient created across the crack. The first mechanism may have played a role locally, as some veins have magnesite. Isobaric (3.5 kbar), fluid-saturated T-X(CO2) infilling of the fracture by external fluids rich in other components such as CO2 (e.g. Demouchy et al., 2016) have also been described in eclogite-facies conditions as ‘prograde-metamorphic dehydration fluid conduits’ or dehydration veins owing to an influx of an H2O-unsaturated fluid (e.g. Beinlich et al., 2010; John et al., 2012; Taetz et al., 2016, 2018).

Before the formation of the fractures the serpentinites may have been water-saturated, owing to the chemical and structural adjustment of antigorite under increasing temperature conditions. Wunder et al. (2001) experimentally showed that the antigorite polysome changes from 18 to 14 when temperature is increased from 450 to 650 °C at 30 kbar. This change in polysomatism results in the release of up to 0.19 wt % of H2O (cf. Shen et al., 2020). Such a decrease of the antigorite
polysome number with increasing metamorphic grade (from \(m = 17–18\) at 420 ± 30°C to \(m = 14–18\) at 520 ± 20°C, at 3.5 kbar) has been documented in Alpe Zocca (Mellini et al., 1987). Yet water saturation is not a requirement for triggering the dehydration reactions below the equilibrium conditions.

Olivine-bearing veins in serpentinites are classically described as dehydration embrittlement veins (Nishiyama, 1989; Strating & Vissers 1991; Healy et al., 2009; Dunkel et al., 2017). However, the systematic absence of talc or tremolite in the serpentinites hosting dehydration veins in the present study indicates that the fractures predate reactions (2) and (3). In consequence, they cannot have been produced by dehydration embrittlement phenomena associated with fluid overpressure owing to these reactions. They also cannot have formed by the channel-forming reactive porosity model proposed by Plümper et al. (2017b) to account for the olivine-bearing veins in Erro–Tobbio serpentinites (Ligurian Alps, Italy), as the characteristic hierarchical drainage structure, with a clear fluid-flow polarity direction (tree root-like arrangement), reported by Plümper et al. (2017b) is not observed in our study area, where unconnected parallel or conjugate-like arrangements of olivine-bearing veins are conspicuous (see Fig. 4a–c). A possible explanation for the dissimilarity between the two cases is the meter-scale chemical homogeneity of the reacting Val Malenco foliated antigorite-serpentinites, which contrasts with the small-scale compositional heterogeneity of the undeformed massive serpentized peridotites in Erro–Tobbio (Scambelluri et al., 2001; Strating & Vissers, 1991; Hermann et al., 2000; Plümper et al., 2017b). Indeed, the numerical model developed by Plümper et al. (2017b) to support the channel-forming reactive porosity mechanism is strongly dependent on the chemical heterogeneity of their input mid-ocean ridge serpentinite peridotite at all scales. We propose therefore that the fractures formed in response to an external process and that it is their effect on the pressure field that triggered the reaction. The remarkable common orientation of most veins at high angle to the isograds (~N70° nearly vertical, Fig. 2) suggests that they might have originated as extensional fractures associated with tangential extensional stresses produced in the host-rock by the Bergell intrusion (e.g. Koide & Bhattacharji, 1975). It is worth noting that previous interpretations of the olivine-bearing veins in the serpentinized peridotites of Erro–Tobbio also proposed a major role for external tectonic stresses in their formation (see fig. 3 of Strating & Vissers, 1991).

The observed variability in the veins’ morphology probably records different activity time lapses. A fluid pressure lower than lithostatic has to be maintained for the reaction to progress, but this situation is mechanically unstable. The porosity production and total volume change resulting from dehydration on the surface of the extensional crack walls might produce a transient fluid-filled porosity and transient permeability, but reaction progress will eventually be halted by mineral precipitation and unavoidable collapse of pores maintaining sub-lithostatic pressures. Very thin veins (Fig. 4a–c) probably record a pulse-like activity. In contrast, development of reaction fronts up to tens of centimeters wide requires that the vein worked as a high-permeability channel for a finite time, draining the fluids as the reaction progressed into the serpentinite host-rock (Figs 4d and 5). The process allowing for the maintenance of these high-permeability channels under lithostatic pressures of c. 3.5 kbar is not fully understood. However, all these veins have a well-developed core with a coarse-grained jackstraw olivine texture. The criss-cross arrangement of the stiff (at these low temperatures) olivine crystals may have played an essential role in avoiding full compaction of the vein for a given period of time.

Bucher (1998) presented a model based on a crack–reaction–seal mechanism to account for olivine and tremolite veins replacing massive dolomite marble close to, or as enclaves in, the Bergell intrusion. These veins were generated during contact metamorphism in response to an important influx of metasomatizing SiO2-rich fluids, probably derived from the cooling intrusion. They are typically surrounded by symmetrical reaction zones that resemble those of the composite Ol–Tlc veins in this study (Figs 4a, d and 5). These reaction zones show either straight (tremolite veins) or highly irregular (olivine veins) boundaries. This change in morphology was interpreted as the result of predominance of surface-controlled reaction kinetics over diffusive processes for straight reaction fronts, and the reverse for irregular fronts (Bucher, 1998). The crossover in time between the two mechanisms is temperature-dependent. This is consistent with the observation that Ol–(±Tr) veins, situated farther from Bergell intrusion contact display straight and sharp limits (Fig. 4a), whereas Ol–Tlc veins are often composite (Figs 4d and 5). Time scales for the propagation of the reaction fronts inferred by Bucher (1998), based on the observed thickness of reaction zones, are of several hundreds of years for the sharp tremolite veins formed at 450°C [similar to the Ol–(±Tr) veins studied here] and up to a thousand years for the diffuse olivine veins formed at 550°C (similar to the Ol–Tlc veins in our study area).

The morphology and size of olivine crystals in jackstraw-textured Ol–Tlc dehydration veins is rather outstanding given the low temperatures (≤500°C) at which they were crystallized. This olivine texture has been repeatedly observed in metamorphic peridotites produced by dehydration reactions resulting in olivine associated with talc or enstatite (Matthes, 1971; Evans & Trommsdorff, 1974a; Collerson et al., 1976; Hietanen, 1977; Snoke & Calk, 1978; Nilsson, 1985; Bakke & Korneliussen, 1986; Trommsdorff et al., 1998; Akinin, 1994; Padrón-Navarta et al., 2010a,b, 2011; Dilissen et al., 2018). This olivine morphology might reflect a particular growth kinetics linked to high effective fluid
pressure in the dehydration veins. In unconstrained magmatic systems (free growth), feathery to hopper olivine crystal shapes are indeed known to result from fast growth in response to undercooling (Donaldson, 1976; Faure et al., 2003, 2006). Extrapolation of these results to fluid-mediated solid-state reactions is not obvious, but a possible explanation for development of similar textures in dehydrating ultramafic systems would be fast growth owing to an increase of the affinity of the reaction in response to effective fluid drainage in the veins. The coarse olivine grain size in veins (Figs 4e and 8c) also suggests high time-integrated fluid–rock ratios. The marked change in grain size from the jackstraw-textured core of the composite veins to selvages (Fig. 5) may therefore record fluid focusing in the vein.

There is a marked resemblance between the olivine microstructure, SPO and CPO in the reaction zones associated with veins (Fig. 5) and in the pervasively dehydrated metaperidotite. Olivine crystals are systematically elongated parallel to (001) and flattened parallel to [010], marking the foliation, and intragranular misorientation is mainly accommodated by rotation around <u0w> axes with dominance of [100] (compare Figs 5 and 13b). This suggests that the processes controlling olivine grain growth are similar and that viscous compaction was active in both domains.

The propagation of reaction zones generated by dehydration veins might lead to pervasive dehydration of serpentinites. Reminiscent of dehydrating veins included in the pervasive metaperidotite are relatively common. They have clear evidence of macroscopic deformation (but limited strain) suggesting that the formation of the veins predated the pervasive transformation of antigorite to the talc + olivine assemblage. However, it is unlikely that the entire pervasive metaperidotite domain formed by progression of the reaction zones associated with veins. The thickness of the reaction fronts in the composite veins, which does not exceed a few tens of centimeters, is consistent with slow propagation (velocities normal to the fracture orientation of the order of 5 cm ka⁻¹; Bucher 1998) and short vein lifetimes. The thermal model of Trommsdorff & Connolly (1996) suggests that rocks at 890 m from the intrusion contact [a position intermediate between the isograds of reactions (2) and (3)] attained a maximum temperature of 520 °C and began cooling < 100 kyr after the intrusion. Closer to the intrusion, maximum temperatures were higher and cooling slower. However, even under such conditions, the vein spacing should be of the order of 10 m or less to account for pervasive replacement of the serpentinites. Although it is difficult to map the veins in the pervasive metaperidotites, their spacing is significantly larger than this value.

Observations indicating auto-retrogression (dehydration) of the highest-grade assemblages in the pervasive domain close to the intrusion, in particular anthophyllite retrograded to talc (Trommsdorff & Evans, 1972), imply that the time scale for fluid extraction in the pervasive domain was of the order of, or lower than, 100 kyr. These time scales are consistent with those predicted for viscous metamorphic compaction in numerical models (e.g. Connolly, 1997). If fluid production rates can be accommodated by viscous metamorphic compaction, the existence of highly permeable domains such as fractures does not perturb the generation of porosity waves that remain the main fluid extraction mechanism (see fig. 5 of Connolly, 2010).

Olivine SPO and CPO: indicators of fluid flow?

The jackstraw-textured veins, the associated reaction zones, and the pervasive metaperidotite show a strong correlation between olivine SPO and CPO. Olivine crystals are systematically elongated parallel to [001]. However, their orientation varies. In the veins, [001]Ol axes are contained in the vein plane and [010]Ol axes are normal to it (Fig. 12), whereas in the veins’ reaction zones and in the pervasive metaperidotite [001]Ol axes are contained in the foliation plane and [010]Ol axes are normal to it (Figs 5 and 10). Because the orientation of the veins is usually at a high angle to the foliation (Fig. 2), the CPO and SPO in the veins are also often roughly normal to those in the reaction zones and pervasive metaperidotites. Within the jackstraw-textured domains, the olivine CPO is independent of the antigorite fabric in the reacting wall-rock (Fig. 12), whereas in the veins’ reaction zones and in the pervasive metaperidotite there is a strong correlation between the olivine and antigorite CPO.

The olivine CPO in the pervasive metaperidotite has a strong correlation with the antigorite CPO. This relation may be produced by epitaxial growth of olivine on antigorite with the relation [001]Ol // [010]Atg // [001]Atg, which is one of the two orientation relations (No. 2) originally documented by transmission electron microscopy during olivine hydration to form antigorite (Boudier et al., 2010). The other orientation relation (No. 1), [001]Ol // [001]Atg // [010]Atg, described by Boudier et al. (2010), is documented in some Ca-rich serpentinites, but not in the metaperidotites (Fig. 10). Orientation relations between coexisting antigorite and olivine have been recognized in olivine- and antigorite-bearing serpentinites from the eastern part of Val Malenco, outside the contact metamorphism aureole [where olivine is formed by reactions (1a) and (1b) using EBSD data (Jung, 2009, 2011; Liu et al., 2018; Morales et al., 2018). Liu et al. (2018) observed evidence for the relationship [010]Ol // [001]Atg (No. 2), which is the dominant one in our study area. Previous data for olivine from Jung (2009, 2011) for the same lithology outside the contact aureole show either [001]Ol or [010]Ol at high angles to the foliation. However, the bulk fabric of the olivine in his study is rather oblique to the antigorite foliation, making it difficult to establish a clear crystallographic relationship. Morales et al. (2018) described the two antigorite-olivine topotaxy relations originally defined by Boudier et al. (2010) and two new
Implications for dehydration reactions and fluid transport in subduction zones

The jackstraw-textured dehydration veins described here were not produced by hydrofracturing (embrittlement owing to local increase in pore-fluid pressure). They were also not formed by coalescence of a fluid-filled microporosity produced by spatially heterogeneous triggering of the dehydration reactions. They were produced by external tectonic processes, which transiently and locally allowed for the dehydration reactions to occur at lower temperatures than the ‘equilibrium’ ones. These dehydration veins are well developed in Malenco probably as a result of the particular conditions at which the antigorite dehydration occurred: shallow depths and, hence, low lithostatic pressures (3.5 kbar), and anomalous thermal and stress fields related to the Bargell intrusion. Such veins are not, for instance, observed in Cerro del Almirez, where antigorite dehydration occurred at subduction-like conditions (Padrón-Navarta et al., 2011, Fig. 1b). This suggests that the veins in Val Malenco probably do not represent major features allowing for transport of antigorite dehydration fluids in subduction zones. However, these structures highlight the strong interplay between dehydration reaction kinetics (and by consequence fluid production rates) and deformation-induced variations in fluid pressure, which in a subduction environment may occur in response to hydrofracturing or to propagation of porosity waves. Such a coupling between enhanced fluid extraction rates by hydrofracturing and antigorite dehydration kinetics was, for instance, proposed based on the variations in texture of the metaperidotites in Cerro del Almirez (Padrón-Navarta et al., 2010b, 2011).

Prograde olivine in Malenco has weak to moderate but clear SPO and CPO, indicating that even if dehydration occurs under static conditions, it might contribute to seismic anisotropy in the mantle wedge. However, the present results imply that the interpretation of this seismic anisotropy is not straightforward. First, the data in the veins show that the olivine CPO is not controlled by the pre-existing antigorite CPO, but that it may result from anisotropic growth in the presence of fluid pressure gradients and, hence, record fluid flow patterns. An important role of fluid flow in the formation of fabrics in prograde metaperidotites has also been inferred in Cerro del Almirez, where an increase in elongation and change in orientation of the magnetite aggregates from the serpentinites to the metaperidotites, and a correlation between the magnetite lineation and the orientation of [001]_{Ol} in the metaperidotites, were documented (Dilissen et al., 2018). These observations open a new path for mapping fluid flow in subduction zones.

Moreover, even if topotaxial growth plays a role in the formation of olivine CPO, the relation between the antigorite CPO and the deformation reference frame is not always clear. Fast S-wave polarization normal to the flow direction is observed only if [100]_{Ol} is
concentrated normal to the flow direction. This orientation may be produced by the topotactic relation $[001]_{\text{Ol}}$ $(100)_{\text{Ol}} || (010)_{\text{Atg}} (001)_{\text{Atg}}$, which is not observed in the Malenco pervasive metaperidotites. It might also be produced by the topotactic relation $[001]_{\text{Ol}}$ $(010)_{\text{Ol}} || (010)_{\text{Atg}} (001)_{\text{Atg}}$. If the $[010]_{\text{Atg}}$ was aligned in the flow direction before the pervasive dehydration. Based on the assumption that the $[010]_{\text{Atg}}$ axes correspond to the macroscopic lineation in the serpentinites (and therefore shear direction), some researchers have classified the resulting prograde olivine fabric as B-type (Jung, 2011; Nagaya et al., 2014; Liu et al., 2018; but see Morales et al., 2018). However, most often both $[100]_{\text{Atg}}$ and $[010]_{\text{Atg}}$ are dispersed in a girdle in the foliation plane of the serpentinites, with no strong point maximum of $[010]_{\text{Atg}}$ (Fig. 10; Padroñ-Navarta et al., 2012; Soda & Wenk, 2014; Kern et al., 2015; Dilissen et al., 2018). Furthermore, when SPO data for magnetite are available (Dilissen et al., 2018) there is no clear correlation between the concentrations of $[100]_{\text{Atg}}$ or $[010]_{\text{Atg}}$ and the lineation defined by magnetite aggregates.

CONCLUSIONS

Macro- to microstructures observed in the Malenco unit in proximity to the Bergell contact aureole record two different modes of fluid extraction during dehydration reactions. Dehydration veins allowed transient, rapidly focused, fluid flow as the result of a crack–reaction–seal mechanism, allowing for dehydration to occur locally at temperatures lower than the equilibrium conditions. This resulted in fast anisotropic olivine grain growth under high time-integrated fluid–rock ratios, leading to the development of jackstraw textures, characterized by a criss-cross arrangement of prismatic olivine crystals elongated parallel to the $[001]$ direction. The olivine $[001]$ axis and $(010)_{\text{Ol}}$ plane tend to be systematically oriented within the vein plane. In addition, some veins developed reaction zone propagating outwards from the initial crack. In these reaction zones, olivine crystals are oriented with $[001]_{\text{Ol}}$ dominantly normal to the vein and $(010)_{\text{Ol}}$ normal to the foliation in the serpentinite wall-rock. Both observations suggest oriented growth of olivine controlled by fluid pressure gradients, with orientation of the fast growth direction $[001]_{\text{Ol}}$ along the maximum fluid pressure gradient.

However, it is unlikely that fluid extraction by the veins played a significant role in the pervasive dehydration of serpentinites observed within c. 770 m of the contact with the Bergell intrusion, for the following reasons: (1) this transformation is characterized by a wide reaction front of c. 150 m, composed of partially reacted rocks, where the intensity of the reaction bears no clear spatial relation to dehydration veins; (2) full transformation of the serpentinites in metaperidotites by progression of reaction zones around veins would imply a decimeter-scale spacing of veins, which is not observed in the field. We interpret therefore the foliated metaperidotites as formed at near equilibrium conditions, with extraction of the fluids produced by viscous metamorphic compaction. Evidence for limited solid-state deformation of the metaperidotites after their crystalization includes undulose extinction in olivine crystals that preserve nevertheless their prismatic shapes and delicate fingering contacts with talc, and bent talc and chlorite crystals. The SPO and CPO of olivine in the pervasive metaperidotites and in the reaction zones around the veins could be inherited from the CPO of antigorite in the serpentinite, via epitaxial or topotaxial growth, or be controlled by fluid flow with a higher permeability parallel to the foliation. The present results suggest that metamorphic olivine SPO and CPO may represent potential markers of fluid flow direction during dehydration reactions.

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SUPPLEMENTARY DATA

Supplementary data are available at Journal of Petrology online.

REFERENCES


