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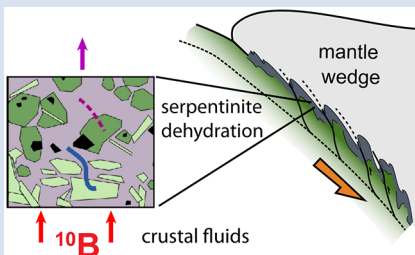
Metamorphic olivine records external fluid infiltration during serpentinite dehydration

E. Clarke¹, J.C.M. De Hoog^{1*}, L.A. Kirstein¹, J. Harvey², B. Debret³



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Abstract



We used boron (B) isotope systematics of co-existing olivine and serpentinite to study deep fluid flow in subduction zones. Metamorphic olivine produced by serpentinite dehydration at sub-arc conditions from high pressure ophiolites in the Western Alps contains significant concentrations of B (2–30 µg/g) with a high $\delta^{11}\text{B}$ values (+9 to +28 ‰), whilst co-existing serpentinite has 2–50 µg/g B with $\delta^{11}\text{B} = +6$ to +24 ‰. Boron isotope fractionation between olivine and its precursor serpentinite ($\Delta^{11}\text{B}_{\text{ol-srp}} = \delta^{11}\text{B}_{\text{ol}} - \delta^{11}\text{B}_{\text{srp}}$) is highly variable, which indicates significant isotopic disequilibrium between these minerals. Importantly, samples with B-enriched olivine have low $\Delta^{11}\text{B}_{\text{ol-srp}}$ (down to –9 ‰), evidence that olivine grew in the presence of a mixture of serpentinite-derived fluids and external fluids with $\delta^{11}\text{B}$ of ca. +6 to +15 ‰. The

composition of these external fluids is consistent with those from subducting sediments and altered oceanic crust at 50–80 km depth, and at least 15–45 % fluid addition. Our work shows that large scale slab fluid infiltration and fluid-mobile element transport accompanies serpentinite dehydration in subduction zones.

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Introduction

The recycling of lithosphere-hosted volatile and fluid-mobile elements (e.g., H₂O, CO₂, B, halogens) back into the convecting mantle *via* subduction has been crucial for Earth's evolution. Down-dragged mantle wedge serpentinites and hydrated oceanic mantle are water-rich (up to 13 wt. % H₂O), and increasingly recognised as important sources of water and fluid-mobile elements (Deschamps *et al.*, 2013; Scambelluri *et al.*, 2019). However, many details of serpentinite dehydration in subduction zones are still poorly understood, including the relative roles of mantle wedge *vs.* ocean floor serpentinites (Martin *et al.*, 2020) and B isotope fractionation during serpentinite breakdown (De Hoog *et al.*, 2014). Previous work on B behaviour during de-serpentinisation in subduction settings focussed on whole rock geochemistry (Harvey *et al.*, 2014; Cannò *et al.*, 2015), which obscures the detailed record of fluid-rock interaction preserved in individual mineral phases.

Metamorphic olivine is formed during breakdown of brucite or antigorite (high pressure serpentine) at ~400 °C and ~650 °C, respectively (Scambelluri *et al.*, 2004) and is B-rich (up to 100 µg/g; Scambelluri *et al.*, 2004; De Hoog *et al.*, 2014) compared to primary mantle olivine (<0.11 µg/g; Ottolini *et al.*, 2004). The incorporation of B in metamorphic olivine uniquely records fluid processes during serpentinite dehydration and allows insight into the composition, evolution and origin of these fluids.

Here we present *in situ* $\delta^{11}\text{B}$ data of co-existing serpentinite and metamorphic olivine from high pressure serpentinites,

which show significant disequilibrium between the two minerals. We interpret this as evidence for the infiltration of externally derived fluids during serpentinite dehydration, attesting to complex fluid pathways near the slab-wedge interface.

Samples and their Geological Setting

To capture the dehydration process, we selected eight serpentinite samples with phase assemblages between the brucite-out and antigorite-out phase transitions with 20–40 % modal olivine from three ophiolites in the Western Alps (Fig. 1). The Monviso Lago Superiore Unit (LSU) is a shear zone which underwent lawsonite-eclogite facies metamorphism and likely represents a serpentinite channel on the interface between the subducting plate and overlying mantle (Guillot *et al.*, 2004). The Zermatt-Saas ophiolite represents relict oceanic lithosphere of the Mesozoic Tethys ocean subducted to ~60–70 km: samples were obtained from near Zermatt and from Val d'Aosta. The Erro-Tobbio metaperidotites in the Voltri massif were subducted to 2.0–2.5 GPa and 550–600 °C and represent a subduction channel domain. See [Supplementary Information](#) for further sample details.

Boron Isotope Systematics of Olivine and Serpentine

Boron isotope ratios and concentrations were measured by SIMS at the Edinburgh Ion Microprobe Facility. Matrix-matched

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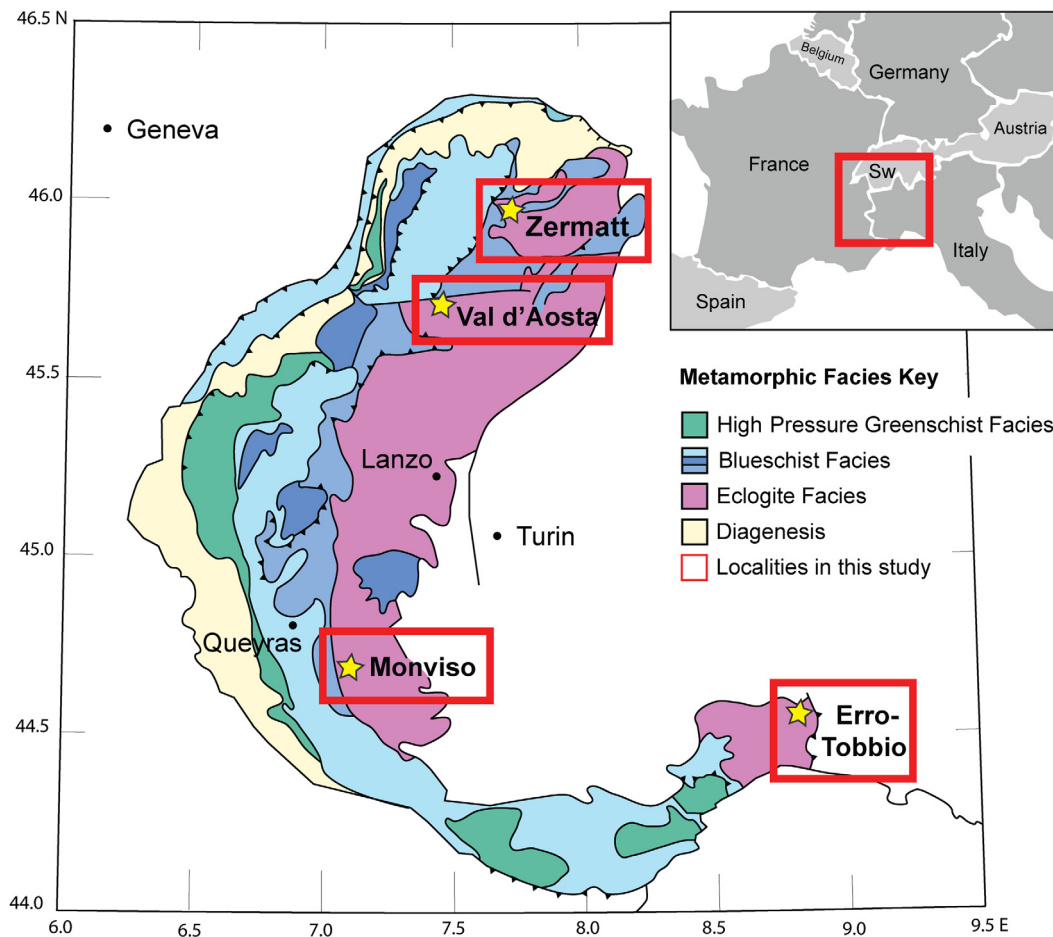


Figure 1 Metamorphic facies map of the Western Alps (after Bousquet *et al.*, 2012) showing study areas and sample locations.

standards including serpentine and olivine with known B isotope compositions were used for calibration. Typical uncertainty for B isotope analyses is ± 1.6 ‰ (1 σ). Further analytical details and the complete dataset can be found in the [Supplementary Information](#).

Olivine and serpentine show high $\delta^{11}\text{B}$ (+6 to +28 ‰) in all samples (Fig. 2). Minor phases present in some samples include pyroxene, chlorite and clinohumite. Clinohumite has similar [B] as olivine (4.8 to 6.5 $\mu\text{g/g}$), but very low modal abundance. Pyroxene and chlorite have low [B] (<2 $\mu\text{g/g}$) compared to olivine and serpentine and can be ignored for the overall B budget.

On average, Monviso metamorphic olivine shows lower $\delta^{11}\text{B}$ and higher [B] ($\delta^{11}\text{B} = +8$ to +14 ‰, [B] ~ 16 $\mu\text{g/g}$) than co-existing serpentine ($\delta^{11}\text{B} = +18$ to +21 ‰, [B] ~ 6 $\mu\text{g/g}$). Erro-Tobbio olivine has slightly higher $\delta^{11}\text{B}$ and [B] ($\delta^{11}\text{B} +18$ to +20 ‰, [B] ~ 11 $\mu\text{g/g}$) than co-existing serpentine ($\delta^{11}\text{B} +16$ to +17 ‰, [B] ~ 10 $\mu\text{g/g}$). Zermatt olivine shows a wide range of $\delta^{11}\text{B}$ (+16 to +28 ‰) and is higher on average than serpentine ($\delta^{11}\text{B} = +6$ to +21 ‰), but lower [B] (~ 4 $\mu\text{g/g}$) in olivine compared to serpentine (~ 8 $\mu\text{g/g}$). The Val d'Aosta sample has higher [B] in olivine *vs.* serpentine (22 *vs.* 10 $\mu\text{g/g}$) but somewhat lower $\delta^{11}\text{B}$ (+17 to +19 ‰ in olivine, +16 to +21 ‰ in serpentine).

The most striking aspect of our dataset is the large variation in B isotope fractionation between olivine and serpentine ($\Delta^{11}\text{B}_{\text{ol-srp}}$; $\delta^{11}\text{B}_{\text{ol}} - \delta^{11}\text{B}_{\text{srp}}$ of co-existing olivine and serpentine), which shows that olivine is often not in isotopic equilibrium with

serpentine, despite being its daughter product. Our dataset also shows a clear covariation between $\Delta^{11}\text{B}_{\text{ol-srp}}$ and B-enrichment in olivine ($[\text{B}]_{\text{ol/srp}}$: $[\text{B}]_{\text{ol}}/[\text{B}]_{\text{srp}}$): Zermatt metamorphic olivine is depleted in B compared to co-existing serpentine but has heavier $\delta^{11}\text{B}$ (Fig. 3). Monviso and Val d'Aosta olivine, however, is enriched in B compared to the co-existing serpentine but has lower $\delta^{11}\text{B}$. Erro-Tobbio olivine lies between these two extremes with roughly similar [B] and $\delta^{11}\text{B}$ in olivine and serpentine. As the measured olivine is metamorphic and grew upon serpentine dehydration, protolith composition and heterogeneity have no bearing on $[\text{B}]_{\text{ol/srp}}$ and $\Delta^{11}\text{B}_{\text{ol-srp}}$, therefore processes operating during olivine growth are needed to explain these observations.

Fluid Infiltration during Serpentine Dehydration

No partitioning data for B between serpentine and olivine ($D_{\text{B}}^{\text{ol/srp}}$) is available, but a value of 3–5 for $D_{\text{B}}^{\text{fluid/residue}}$ was recorded during serpentinite dehydration experiments with olivine as the main reservoir for B in the residue (Tenthorey and Hermann, 2004). Given this, mass balance dictates that $D_{\text{B}}^{\text{ol/srp}} \sim 0.6$ –0.9 (see [Supplementary Information](#)), and higher values indicate excess B in olivine. Samples from Zermatt show $[\text{B}]_{\text{ol/srp}}$ 0.4–0.9, close to the expected equilibrium value. This is accompanied by $\Delta^{11}\text{B}_{\text{ol-srp}}$ of +7.4 to +14.6 ‰ (Fig. 3). In most silicates, including serpentine, B^{3+} substitutes for Si^{4+} or Al^{3+} in tetrahedral (IV) coordination (Hervig *et al.*, 2002; Pabst *et al.*, 2011). In mantle

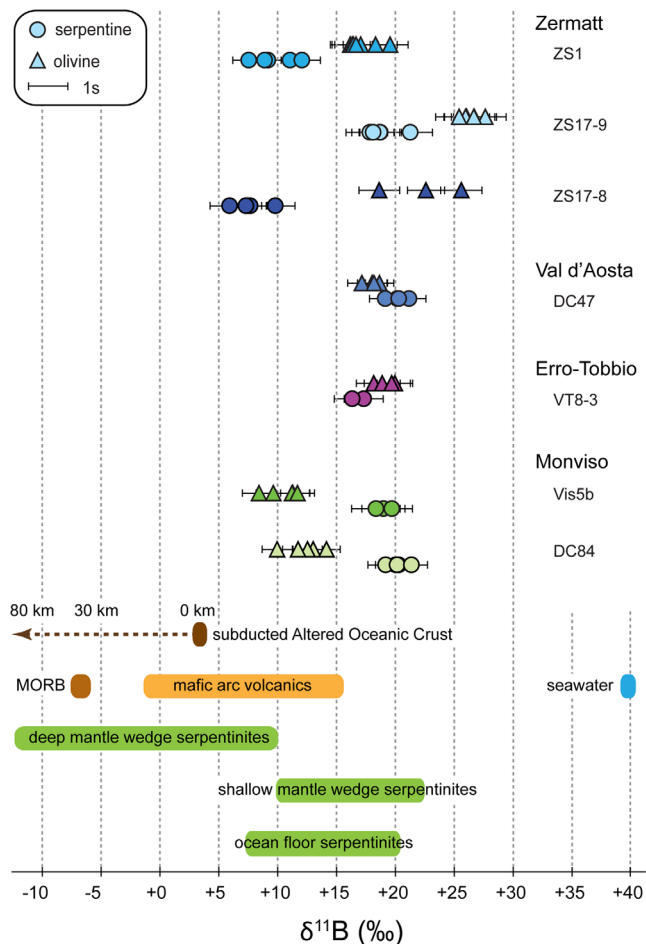


Figure 2 $\delta^{11}\text{B}$ data for olivine and serpentine in our samples. Error bars represent 1 σ uncertainties and include calibration uncertainties. Also indicated are B isotope compositions of various reservoirs relevant to subduction zones (data sources: Tonarini *et al.*, 2011; De Hoog and Savov, 2018 and references therein; Martin *et al.*, 2020).

silicates such as clinopyroxene and olivine, B also substitutes for Si^{4+} but occurs in trigonal (III) coordination as a BO_3 group (Hålenius *et al.*, 2010; Ingrin *et al.*, 2014). As ^{11}B preferentially resides in trigonal and ^{10}B in tetrahedral coordination (Kowalski *et al.*, 2013), olivine with trigonal B will show a heavier B isotope signature than co-existing serpentine with tetrahedral B. For the Zermatt samples, $\Delta^{11}\text{B}_{\text{ol-srp}}$ is close to the experimentally determined equilibrium B isotope fractionation between III and IV phases, which decreases from $\Delta^{11}\text{B}_{\text{ol-srp}} = +12.5$ to $+8.5$ ‰ with T increasing from 450 to 650 °C (Kowalski *et al.*, 2013).

Therefore, we conclude that Zermatt metamorphic olivine was in equilibrium with internal fluids released during dehydration of serpentine, with no evidence for contamination by external fluids. This conclusion is supported by REE and Sr isotope data (Gilio *et al.*, 2019) and consistent with the geological setting of the Zermatt samples, which come from a thick (~2 km) coherent section of subducted ocean lithosphere. The range in $\Delta^{11}\text{B}_{\text{ol-srp}}$ of $+7.4$ to $+14.6$ ‰ is probably related to metamorphic olivine forming at different temperatures during antigorite-out and brucite-out reactions, respectively.

The majority of metamorphic olivines from localities other than Zermatt have $[\text{B}]_{\text{ol/srp}} > 1$ and $\Delta^{11}\text{B}_{\text{ol-srp}} < +5$ ‰, *i.e.* more B and ^{10}B in olivine than is expected for growth during equilibrium dehydration of serpentine (Fig. 3). The most likely source of

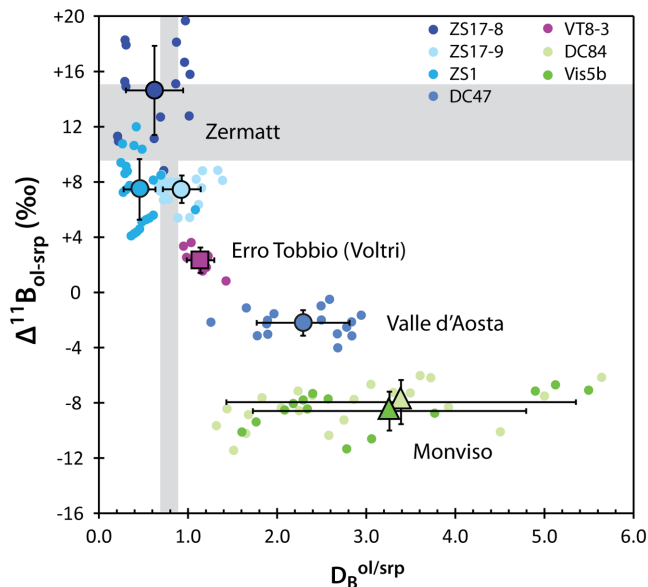


Figure 3 Diagram comparing boron isotope systematics of co-existing metamorphic olivine and serpentine. $[\text{B}]_{\text{ol/srp}} = [\text{B}]_{\text{olivine}} / [\text{B}]_{\text{serpentine}}$, and $\Delta^{11}\text{B}_{\text{ol-srp}} = \delta^{11}\text{B}$ of olivine – $\delta^{11}\text{B}$ of serpentine. Large symbols with 1 σ error bars indicate averages for each sample, whereas small symbols represent all possible olivine-serpentine pairs for each sample (see Supplementary Information for details). A broadly negative correlation between $[\text{B}]_{\text{ol/srp}}$ and $\Delta^{11}\text{B}_{\text{ol-srp}}$ is observed. Grey bars indicate equilibrium fractionation of boron and its isotopes at relevant *P-T* conditions.

excess B is external fluid, as no other B-rich phases that dehydrate at 450–650 °C were present. Scarce clinohumite is B-rich but stable at the prevailing *P-T* conditions (Shen *et al.*, 2015) and its breakdown products (ilmenite, rutile) were not observed. Thus, we envisage a model where fluid from serpentine dehydration mixed with infiltrating external fluid, and metamorphic olivine grew in equilibrium with this mixed fluid. Serpentine did not re-equilibrate with this fluid due to sluggish kinetics at these low temperatures (450–650 °C) (*e.g.*, Lafay *et al.*, 2019). Mass balance calculations indicate that these external fluids had $\delta^{11}\text{B}$ of $+6$ ‰ (Monviso) to $+15$ ‰ (Erro-Tobbio) depending on the B concentration of the external fluids (Fig. S-6) and assuming mildly alkaline pH conditions for both internal and external fluids (Galvez *et al.*, 2016; Debret and Sverjensky, 2017). This matches the composition of fluids derived from a mixture of altered oceanic crust and sediments from 50–80 km depth (Tonarini *et al.*, 2011; Yamada *et al.*, 2019), consistent with the depth of the serpentinites in the former subduction zone. The external fluids comprise at least *ca.* 15–45 % of the total fluid budget of the dehydrating serpentinites, which are minimum estimates, as any partial re-equilibration of serpentine would lead to underestimating the amount of external fluid.

Metasomatism near the Subduction Interface

External fluids may permeate dehydrating serpentinites without the need for large scale faulting, as seen in Erro-Tobbio, which only shows metre scale vein networks and deformation. During serpentine dehydration there is a rapid increase in porosity and rock permeability allowing fluid to drain from the rock without the need for brittle failures and opening of fault pathways (Tenthorey and Cox, 2003; Plümper *et al.*, 2017). The creation of such porosity also provides pathways for significant volumes of B-rich external fluids to infiltrate.



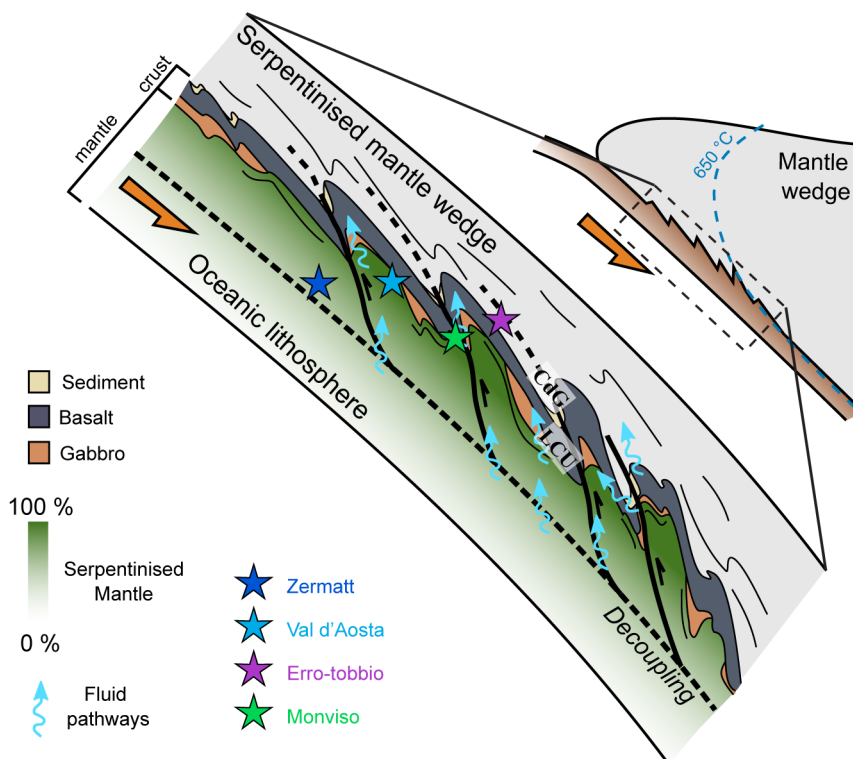


Figure 4 Schematic diagram depicting proposed fluid pathways and setting within the plate interface for serpentinite localities in this study after Gilio *et al.* (2019), where LCU and CdG indicate positions of Lago di Cignana and Cima di Gagnone HP ophiolites from that publication.

The trend towards higher $[B]_{ol/srp}$ with lower $\Delta^{11}B_{ol-srp}$ indicates an increasing influence of B-rich external fluids from Zermatt to Erro-Tobbio to Valle d'Aosta to Monviso. This trend correlates with the position of the serpentinite bodies relative to the slab-mantle interface during peak metamorphic conditions (Fig. 4). The Monviso ophiolite and Erro-Tobbio massif are thought to represent serpentinite domains between the subducting slab and the mantle wedge (Guillot *et al.*, 2004; Scambelluri and Tonarini, 2012); a prime location to be infiltrated by fluids derived from the subducted slab. Shear zone associated faulting provides many additional pathways for fluid flow throughout the Monviso ophiolite, leaving the serpentinites exposed to extensive interaction with external fluids (Angiboust *et al.*, 2014; Gilio *et al.*, 2020). The Val d'Aosta sample is considered part of the Zermatt-Saas ophiolite but at Val d'Aosta the serpentinite is thinner (~0.5 km) and surrounded by crustal lithologies (metasediments and metabasalts). This sample shows elevated $[B]_{ol/srp}$ and lower $\Delta^{11}B_{ol-srp}$ compared to the other Zermatt samples, likely related to its proximity to other lithologies and increased exposure to dehydration fluids.

The micro-scale process where metamorphic olivine grows from a mixture of internal and external fluids would not be detected using whole rock $\delta^{11}B$ data, as serpentinites in this study have positive whole rock $\delta^{11}B$ (e.g., Scambelluri and Tonarini, 2012) whilst being variably affected by external fluid infiltration. In contrast, subduction mélange serpentinites and metaperidotites with negative $\delta^{11}B$ suffered more extensive contamination and extreme $\delta^{11}B$ overprinting consistent with their close association with large proportions of crustal rocks and sediments (Cannaò *et al.*, 2015; Martin *et al.*, 2020).

Conclusions

Metamorphic olivine is a sensitive recorder of the fluid composition from which it grew, and recorded extensive external fluid infiltration during subduction dehydration of Alpine serpentinites,

which are sinks and carriers of crust-derived fluid mobile elements. External fluid infiltration was most prominent in serpentinites close to the subduction interface and amounted to up to 45 % of the total dehydration fluid budget. The study of B isotope systematics of texturally co-existing olivine and serpentine provides a powerful tool to study fluid flow deep in subduction zones, and reveals processes not visible in whole rock $\delta^{11}B$ compositions.

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Additional Information

Supplementary Information accompanies this letter at <https://www.geochemicalperspectivesletters.org/article2039>.



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