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Can Pseudotachylytes Form via Fracture-Induced Decompression Melting Under Hydrous Conditions?

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Frictional rock sliding and resultant shear heating along a fault plane are proposed as the necessary conditions to generate earthquake-related pseudotachylytes. However, frictional melting alone is energy expensive, requiring large temperature increases of several hundreds to thousands of degrees. Using the example of the pseudotachylyte structures of the Balmuccia Peridotite Massif (Ivrea-Verbano Zone, Alps, Italy), a minimum temperature increase of up to ~540°C for frictional melting has been proposed under isobaric and anhydrous conditions. Such conditions are however inconsistent with the moderate temperature increases, of up to $\sim 400^{\circ}$ C, and diminishing pressures, less than 0.7 GPa, required to form the observed A-type pseudotachylyte structures, with ultramafic composition corresponding to the melting of specific mineral phases as a function of Carboniferous Period pressure and temperature conditions. Here we show that pseudotachylytes could be produced by fracture-induced decompression melting under hydrous conditions, that favor the formation of immiscible liquids derived from melting Al-Cr spinel and orthopyroxene with suspended clinopyroxene minerals in this immiscible melt. We propose thermodynamic calculations that constrain phase stability in the pressure-temperature space of the Balmuccia lherzolite under both anhydrous and hydrous conditions (0.5 to 1 wt.% H₂O). We illustrate that the Al-Cr spinel+orthopyroxene composition of the pseudotachylyte is consistent with lower pressure conditions than those of the initial peridotite prior to fracturing. These thermodynamic calculations help determine the pressure-temperature path of pseudotachylyte formation, not only favored by frictional heating but also by pressure drop (0.3–0.9 GPa) following dilation related to rock fracturing. Our results call for a reassessment of the origin of many pseudotachylytes formed in the lower crust and upper mantle. We show that fracture-induced decompression melting under hydrous conditions can be a viable mechanism that assists frictional melting by reducing the temperature rise from ambient temperature to melting temperature by 18% to 74%. A similar process may be significant in producing other pseudotachylytes during tectonic movement of lithospheric blocks in the deep crust and upper mantle.

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1 Introduction

Pseudotachylytes are fossil remnants of ancient earthquakes generated along fault planes in lithospheric rocks during tectonic processes (Sibson, 1975; Austrheim and Andersen, 2004; Andersen and Austrheim, 2006; Lin, 2007; Di Toro et al., 2009). Pseudotachylytes help decipher the mechanisms of coseismic fault lubrication and slip-weakening (Spray, 1995; Magloughlin and Spray, 1992; Kanamori et al., 1998; Bouchon and Ihmlé, 1999; Hirose and Shimamoto, 2005; Di Toro et al., 2006a, 2011; Yao et al., 2023) associated with the propagation of seismic ruptures in the lithosphere. There is a general

consensus that pseudotachylytes are formed when frictional heat, abrasion, and comminution (Tsutsumi and Shimamoto, 1997a; Menegon et al., 2017, 2021; Spang et al., 2024; Aldrighetti et al., 2025) induce melting of the rock during rapid/fast seismic fault slip; slower slip does not produce sufficient heat concentrations to generate a melt (McKenzie and Brune, 1972; Lin and Shimamoto, 1998; Nielsen et al., 2008; Del Gaudio et al., 2009; Moris-Muttoni et al., 2022). Therefore, pseudotachylytes are solidified earthquake-derived frictional melts (Michalchuk et al., 2023). The melt also rapidly solidifies/quenches, typically to glassy material, i.e., pseudotachylyte (Wenk, 1978; Sibson, 1980; Spray, 1992; Trouw et al., 2010; Ferrand et al., 2018; Dai

et al., 2022), which is thus considered to be a robust benchmark of past seismic slip events (paleoseismicity; Sibson 1975). Relatively unaltered pseudotachylyte veins found in exhumed lithospheric massifs testify to the occurrence of seismic ruptures even in deep portions of the lithosphere where rocks tend to have a dominantly viscous response to tectonic stresses (Obata and Karato, 1995; Dunkel et al., 2021).

Yet questions remain about how pseudotachylytes are generated in the deep lithosphere, where high differential stresses (>0.4 GPa; Obata and Karato 1995; >1 to 6 GPa; Scambelluri et al. 2017; Campbell et al. 2020; Pennacchioni et al. 2024; Menegon et al. 2021; Toffol et al. 2024) are required for faulting. Laboratory experiments simulating dynamic fault strength and frictional melting processes under ambient conditions (Spray, 1987, 1992, 1995; Shimamoto and Tsutsumi, 1994; Tsutsumi and Shimamoto, 1997a,b; Hirose and Shimamoto, 2005; Hirose, 2005; Di Toro et al., 2006a,b; Del Gaudio et al., 2009) have highlighted that coseismic slip velocities of ~ 1 m/s and large displacements of tens of meters produce microstructures similar to those found in natural pseudotachylytes hosted in mantle rocks (Lin and Shimamoto, 1998). In these tests, rock weakening is first associated with breaking of grain asperities and pulverization forming gouge material. Then, rapid (flash) temperature increase leads the rock to partially melt above the solidus temperature or completely melt above the liquidus temperature depending on the maximum temperature achieved: 1200°C to over 1650°C in peridotites (Andersen and Austrheim, 2006) and up to 1392 to 1850°C inferred from the melting point of end-member mineral phases (olivine, pyroxenes, and spinel) in the peridotite (Del Gaudio et al., 2009). In this case, thermal runaway of several hundreds to thousands of degrees in quasi-steady-state, infinite-like ductile localization that weakens rocks by producing a molten shear band that lubricates the fault and sustains rupture propagation (Ogawa, 1987; Karato et al., 2001; Braeck et al., 2009; Spang et al., 2024) may be a driver of earthquakes at great depth (Green and Houston, 1995; Zhan, 2020). The frictional melt rapidly flows and continuously coats the slip surfaces while the fault slips at velocities of 2 to 10 m/s (Lin and Shimamoto, 1998; Ferrand et al., 2018). Following the host rock stress relaxation producing heat (Spang et al., 2024), frictional or flash melting thus results in lubrication of the slip surface and weakening. But both suction into open fractures, and solidification of melt when shear heating reduces with heat loss due to cooling on lubricated surfaces (Bjornerud and Magloughlin, 2004), can lead to rock strength recovery and sliding arrest (Ferrand et al., 2018). Recent experiments using cubic-anvil apparatus have simulated triaxial deformation of synthetic serpentinized peridotite at 1.1 GPa and have shown that rapid (0.1 to 10 seconds) and large temperature rises (>1250°C) can occur even with extremely small seismic slip on the order of 10 microns on tiny microfaults to generate pseudotachylytes (Ferrand et al., 2017).

To date, experiments have supported the assumption that pseudotachylytes can form from the frictional melting of all the mineral phases along the fault plane in deep-seated (>40 km) deformational instabilities in relatively low-temperature environments (e.g., 700 to 900°C in mantle peridotites; Kelemen and Hirth 2007; Ueda et al. 2008, 2020; Souquière and Fabbri 2010). However, this mechanism is inconsistent with the actual geochemical composition of natural pseudotachylytes produced by melting selected mineral phases (starting from the minerals of low melting point; Shand 1916; Spray 1992) as a function of pressure (P), temperature (T), and under hydrous conditions. Specifically, the large temperature increases at isobaric conditions assumed for frictional melting do not explain the resulting pseudotachylyte composition produced at moderate temperatures ($\sim 1300-1450$ °C) and lower pressures (<0.7 GPa). We have inspected representative glassy pseudotachylytes in peridotites, by coupling a combination of micro- to nanostructural and geochemical analyses with thermodynamic modelling, to test the feasibility of rapid decompression melting under hydrous conditions. Decompression melting refers to a process in which a rock body, held at approximately the same temperature, experiences reduction in confining pressure (adiabat) until it achieves its solidus temperature above which the rock partially melts (McKenzie and Bickle, In this study, we consider fracture-induced 1988). decompression melting as a rapid geological process stimulated by a sudden pressure drop due to dilation, resulting from accommodating shear of rough surfaces or gouge without microscale fragmentation in Mode II/III fractures formed when Coulomb's criterion is met, and accommodating slip parallel to the crack surface (Jaeger and Cook, 1979; Samuelson et al., 2009; Brantut, 2019). This fracturing mode might result in internal P reduction that leads the rock system (and the minerals therein) to surpass their solidus. In these types of fractures leading to the formation of pseudotachylytes, decompression would aid efficient generation of melts so less heat has to be produced by frictional work. However, whether partial melting due to decompression occurs or not depends on the petrology of the rock system and, thus, on two factors: where the rock system is originally positioned in P-T space, and the phase diagram topology, including the position of the rock solidus temperature modulated by the available water concentrated in hydrous minerals, nominally anhydrous minerals, and/or fluids along grain boundary regions within the rock (Stünitz et al., 2024). Here we show that pseudotachylytes in peridotitic rocks in the lower crustal/upper mantle domain can be produced by partial melting instead of total melting of the original rock system along the fault plane. We propose that decompression-driven partial melting can be a crucial complementary mechanism assisting frictional melting during the generation of pseudotachylytes.

2 Geological Setting and History

2.1 The Balmuccia Peridotite Massif in the Ivrea-Verbano Zone

A well-studied geological site where well preserved pseudotachylytes from the deep crust are exposed is the Balmuccia Peridotite Massif in the Ivrea-Verbano Zone (Southern Domain of the Western Alps, Italy) (Obata and Karato, 1995; Ueda et al., 2008, 2020; Souquière and Fabbri, 2010; Souquière et al., 2011; Menegoni et al., 2024) (Figure 1). The Balmuccia Massif is one of the largest peridotite slivers in the Ivrea-Verbano Zone and is one of the best exposed peridotites in the world. It is an elongated NNE-SSW trending lens of 4.5 km in length, 0.5 km in width, and 1.1 km in height (Shervais, 1979) that stretches from the surface down to 3-km depth with a downward broadening shape, that steeply inclines at $\sim 60^{\circ}$ towards the west with a gently inclined boundary of 30° to 40° towards the east (Ryberg et al., 2023). The Balmuccia peridotite is considered detached from the original Adriatic or Adria mantle (Quick et al., 1995) because it is bounded by extensional high-temperature shear zones (Brodie and Rutter, 1987; Brodie et al., 1992; Rutter et al., 1993) while being encapsulated in the deep lower crust made of Paleozoic granulite-facies metasedimentary units (Kinzigite and Stronalite Formations; Schmid and Wood 1976; Schnetger 1994; Redler et al. 2012) and the Lower Permian Mafic Complex (Basischer Hauptzug or Mafic Formation) of the Ivrea-Verbano Zone (Zingg, 1983; Shervais and Mukasa, 1991; Quick et al., 1995, 2002; Handy et al., 2005; Rutter et al., 2007; Souquière and Fabbri, 2010) (Figure 1). Specifically, the Balmuccia Peridotite Massif is bounded to the west by a shear zone affecting the units of the Mafic Complex and Kinzigite Formation, and to the east by a <100-m wide zone of the "contact series" consisting of pyroxenites and meta-gabbronorites (Shervais and Mukasa, 1991). The Balmuccia peridotite is the least depleted (or most fertile) and least metasomatized ultramafic body (Voshage et al., 1990; Sossi et al., 2018) along with the Baldissero and Premosello bodies in the exposed continental crustal section of the Ivrea-Verbano Zone (Capedri et al., 1977; Ernst, 1978, 1981; Rivalenti and Mazzucchelli, 2000). The Balmuccia Massif comprises spinel lherzolite, which is the main lithology that dominates 85% of the ultramafic body (Rivalenti et al., 1975; Shervais, 1979), and minor spinel harzburgite and dunite, accompanied by pyroxenitic bodies (emplaced at 433±75 Ma; Mazzucchelli et al. 2009) including websterites that are Cr-diopside-bearing and Al-augite-rich (Lensch, 1971; Shervais, 1979; Sinigoi et al., 1983) (Figure 2).

2.2 Structural Components of the Lherzolite

The foliations of the lherzolite strikes N-NE in the southern sector of the massif, approximately 005° , but strikes N-NW, approximately 350° , in the northern sector (Figure 1; *Boudier et al.* 1984). Dips range from 75-85°W in the southern sector to 45-75°W in the

northern sector. Some authors suggest that the original orientation of the foliation was close to horizontal, based on the hypothesis that the entire crustal section of the Ivrea-Verbano Zone was uplifted and tilted through rotation along a N-S trending horizontal axis (*Rivalenti et al.*, 1975; *Zingg and Hunziker*, 1990; *Handy et al.*, 1999). A lineation defined through the orientation of accessory spinel grains in the plane of the foliation mainly plunges 45° to 65° towards 005° in the southern sector of the massif and mainly plunges 10° to 15° towards 350° in the northern sector (Figure 1; *Shervais* 1979; *Boudier et al.* 1984).

The lherzolite is commonly cut by planes of displacement, or macroscale "faults" mainly with <10 cm offset, which are associated with pseudotachylyte structures (Figure 2). The macroscale faults observed at the outcrop scale in the sampling location in this study (Figure 1) are perpendicular to the measured foliation strike and lineation trend (Boudier et al., 1984) as reported by Ferrand et al. (2018). Based on field observations, these macroscale faults appear to be oriented NE-SW with less common W-E and NW-SE strikes with dips of $>60^{\circ}$ (Figure 2) where field measurements were possible along planar face of the outcrop (see photo inset in Figure 1). NE-SW and NW-SE macroscale faults are observed to locally offset the clinopyroxene dikes by less than 4.5 cm. The offsets indicate both apparent sinistral and dextral shear sense (Figure 2).

The lherzolite displays a protoequigranular tabular texture associated to the strong foliation in the northeastern portion of the Balmuccia Massif while the southwestern sector of the Massif is characterized by a porphyroclastic texture of olivine and orthopyroxene (Garuti, 1977; Garuti and Friolo, 1979; Shervais, 1979; Boudier et al., 1984). In both rock textures, olivine always has a D-type fabric (Garuti, 1977). The lherzolite (60 to 70% olivine, 20 to 25% orthopyroxene, 12 to 13 vol.% clinopyroxene, <1 to 3 vol.% spinel, and <1 vol.% amphibole) as the main lithology of the Balmuccia Massif is composed of forsteritic olivine, diopsidic clinopyroxene (ranging from Wo₅₀En₅₀Fs₀ to Wo₅₀En₄₅Fs₅) in contact with enstatitic orthopyroxene $(Wo_1En_{88}Fs_{11})$, Fe^{2+} -bearing spinel and Al-spinel with up to 9 wt.% Cr₂O₃ (Al-Cr spinel) located at triple points as well as inclusions in olivine (Garuti and Friolo, 1979; Shervais, 1979; Sinigoi et al., 1983; Skrotzki et al., 1990, 1991).

2.3 Equilibration History of the Peridotite Massif

The original peridotite massif is thought to have equilibrated at T=1100– 1200° C and P=1.3–2.0 GPa (Shervais, 1979) in the mantle during Variscan subduction processes (320–350 Ma) involving the subduction of the Rheic oceanic crust, the deposition of metasediments in an accretionary prism, and their burial, imbrication, and prograde metamorphism (Wyatt et al., 2022) to amphibolite and granulite facies (Schmid and Wood, 1976; Redler et al., 2012). During the

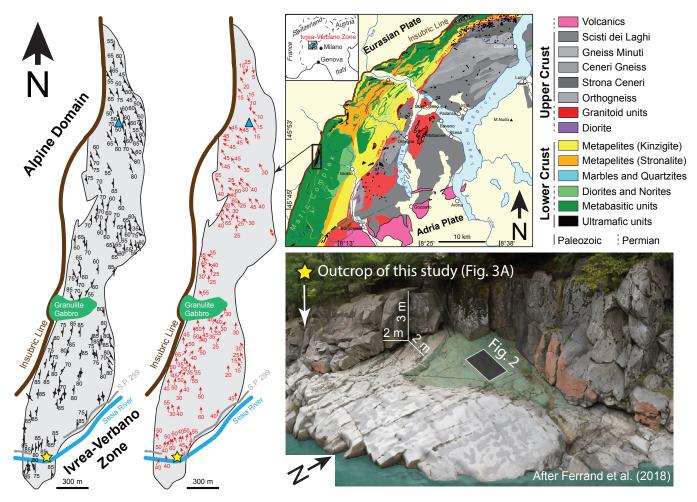


Figure 1 - Geologic context of the Ivrea-Verbano Zone and Balmuccia Peridotite Massif. Geological map of the Ivrea-Verbano Zone (Alps, Italy) (modified after Brack et al. 2010, based on the works of: Rutter et al. 1999; James 2001; Schaltegger and Brack 2007) with the Insubric Line (brown line) dividing the Adria plate (southeast) from the Eurasian plate (northwest) and major pre-Alpine fault zones (red lines). The inset image shows the location of the Ivrea-Verbano Zone in Italy (modified after Rutter et al., 2003). The outline of the Balmuccia Peridotite Massif is shown, with measurements of foliation strike and dip (black symbols) and lineation trend and plunge (red arrows), measured principally though the orientation of spinel grains (Shervais, 1979; Boudier et al., 1984). Note that many measurements of foliation also include deformed Cr-websterite dikes in the same orientation (Boudier et al., 1984). We do not report the internal lithological subdivision (lherzolite, dunite, and pyroxenites) of the Balmuccia Peridotite Massif as shown in the work of Shervais (1979), structural data related to fault and dip, fold axis, destral and sinistral minor fold axis as shown by Shervais (1979) and Boudier et al. (1984), and major discontinuities within the ultramafic body as revealed by the drone-based photogrammetry of Menegoni et al. (2024). The Insubric Line (thick brown line) includes the Tonnello Fault in proximity of the southern portion of the Balmuccia Peridotite Massif. The yellow star indicates the sampling location of the lherzolite outcrop characterized by the presence of pseudotachylyte structure. The location of the outcrop of this study (Figure 3A; N 45°49'13.0", E 8°09'15.6", elevation of 553 m a.s.l.) is located in the upper portion of the rock outcrop investigated by Ferrand et al. (2018). The view of the outcrop from the right bank of the Sesia River is from Ferrand et al. (2018) devoid of the original sketches which were removed using the 3.5 model of the Generative Pre-trained Transformer (OpenAI Inc., USA). The green-shaded area of the outcrop is from Ferrand et al. (2018) indicating the eroded injection plane displaying the remnants on the bottom surface of the tensile fracture. In this outcrop portion, manual field drawing was applied using a tracing paper in which structural and lithological features were recorded at an accuracy down to the subcentimeter scale (see inset grey rectangle of 2.73 x 2.03 m, which is detailed in Figure 2). The outcrop face targeted in this study (indicated by the yellow star) is perpendicular to the polished outcrop surface and has a W-E trend. The light blue triangle indicates the Cima di Lavaggio, the summit point of the Balmuccia Peridotite Massif. S.P. = Strada Provinciale.

late stage of the precursor Variscan orogeny, relatively homogenous mantle deformation occurred at the scales of a few kilometers to a few tens of kilometers (*Drury et al.*, 1991). During the Carboniferous Period (300–320 Ma), the Balmuccia peridotite was exhumed from the mantle and emplaced in granulite-facies metasedimentary units in the lower crust, within the Ivrea-Verbano Zone (*Lensch*, 1971; *Rivalenti et al.*, 1975; *Ernst*, 1978; *Shervais*, 1979; *Shervais and Mukasa*, 1991), probably

at T = 720-900°C and P = 0.9-1.1 GPa (*Handy et al.*, 1999).

More specifically, the exhumation of the mantle peridotite likely occurred $\sim 316-275$ million years ago (*Peressini et al.*, 2007; *Klötzli et al.*, 2014) during the initial period of magmatic underplating (*Voshage et al.*, 1990) producing magmas derived from the hydrous mantle source (*Berno et al.*,

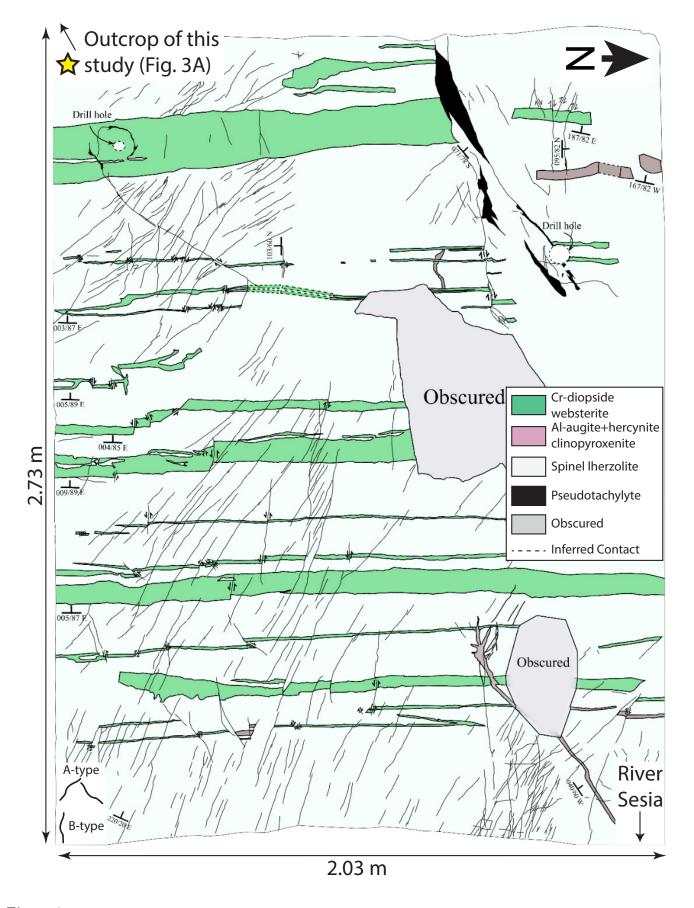


Figure 2 - Digital illustration of a field tracing of a representative portion of an outcrop of the Balmuccia Peridotite Massif located in the north bank of the Sesia river. The reported geological features are accurate to the subcentimeter scale.

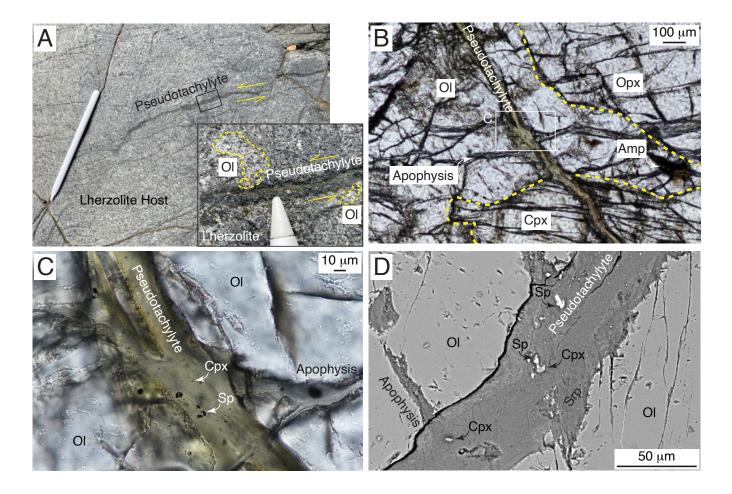


Figure 3 - A-type Pseudotachylyte in Balmuccia lherzolite. A) Field view of the pseudotachylyte in lherzolite host with central serpentinized portion (pencil length = 176 mm) showing 1-cm displacement of olivine minerals (highlighted by yellow dashed contours). B-C) Pseudotachylyte viewed using the unpolarized light with optical microscope. D) SEM-based BSE image of pseudotachylyte with dispersed clinopyroxene and rims of spinel and possible serpentine. Abbreviations: Amp = amphibole, Cpx = clinopyroxene, Ol = olivine, Opx = orthopyroxene, Sp = spinel, Srp = serpentine.

2020). During exhumation, deformation of the mantle rock was accommodated in anastomosing networks of centimeter- to hundred-meter-wide shear zones that may also serve as sites for late-stage fluid percolation (Drury et al., 1991). The peridotite and adjacent metasedimentary granulite complex were then exhumed during the Permian transfersion (270–290 Ma; Handy et al. 1999; Petri et al. 2019) and Triassic-Jurassic rifting of the Piemont-Liguria Ocean (180-230 Ma; Handy and Stünitz 2002). During this period, the peridotite experienced a high-temperature (~1000°C) and low stress (~ 20 MPa) intrusion into the lower crust as manifested in the coarse-grained dynamically recrystallized olivine (Skrotzki et al., 1990) and in the deformation microstructures in the shear zones bordering the pseudotachylytes (Jin et al., 1998). Then, during the Tertiary Alpine collision, the peridotite experienced brittle deformation along the Periadriatic Lineament or Insubric Line (the tectonic boundary separating the African promontory Adria from the Eurasian plate with a dextral sense of shear suggested by the tectonic line gently dipping to the north; Zingg and Hunziker 1990; Figure 1) and crustal overprinting during tilting of the crustal section of the Ivrea-Verbano Zone ($\mathit{Handy\ et\ al.}$, 1999; Rutter et al., 2007).

The second deformation event evidently occurred at relatively low temperature ($\sim 650^{\circ}$ C) and high regional or overburden stress (~300 MPa) (Skrotzki et al., This deformation event is recorded in the 1990). fine-grained dynamically recrystallized rims around the matrix grains, in the subgrain size and dislocation density in olivine within the porphyroclastic domain, and in clinoenstatite lamellae in orthopyroxenes (Skrotzki et al., 1990, 1991). The Cenozoic deformation preserved traces of the pre-Alpine fabrics and mineral assemblages in the Balmuccia lherzolite (Schmid et al., 1989). During these tectonic processes ranging from the Variscan orogen collapse to the Alpine orogenesis, the peridotite sliver of the Balmuccia Massif experienced macroscale fault displacements with common production of pseudotachylytes (Techmer et al., 1992; Obata Based on the combination and Karato, 1995). of geochronology and field relations, two groups of pseudotachylyte structures are found in the Balmuccia Massif: an A-type group (350–270 Ma; including Groups 1 and 2 pseudotachylytes based on the classification of Ueda et al. 2020) largely attributed to the Carboniferous stages of the Variscan orogeny or during the subsequent Permian trans-tension at 550-900°C and 0.6-1.2 GPa, and a B-type group (<70 Ma; Group 3 of Ueda

et al. 2020) related to the Alpine orogeny at <600°C (Wiederkehr et al., 2008, 2009) and <0.3 GPa (Souquière and Fabbri, 2010; Souquière et al., 2011; Ferré et al., In this study, we focus on the A-type pseudotachylyte structures, which seem to represent the features that best record deep earthquake processes in the upper mantle/lower crustal domain. pseudotachylytes characteristically crop out as <1-mm to 2-cm thick structures having sharp boundaries with the host lherzolite (Figures 2 and 3A). Most of the A-type pseudotachylyte structures strike between 140° and 230° and dip more than 65° towards the east (Figure 2).

Methods 3

Fieldwork 3.1

Fieldwork was carried out to identify, characterize, and sample A-type pseudotachylytes at the outcrop scale in the Balmuccia Peridotite Massif. We collected field observations at the outcrops of lherzolites along the north bank of the Sesia river at N 45°49'13.0" and E 8°09'15.6" (as measured using a Garmin eTrek 10), about 60 m south of the Strada Provinciale 299, located halfway between the localities of Balmuccia (SSW) and Isola di Vocca (NNE). Rock sampling was conducted by selecting and chipping (with hammer and chisel) five pseudotachylytes (and the host rock around) displaying a thickness of <5 mm, devoid of alteration/devitrification, with clear evidence of mineral phase displacement (possibly as a result of one single event of displacement with no continuous reactivation of the same fracture zone), and devoid of apophyses at the scale of the rock outcrop. Such a selection helped mitigate a number of complexities for the forthcoming geochemical and microto nanostructural analyses and ensured the identification of the mineral phases and the petrological assessment of frictional melting versus decompression melting. In particular, the limited thickness of the pseudotachylyte on the millimeter scale represents an ideal case where near-equilibrium conditions can be achieved during rock friction, melting, and pseudotachylyte formation, with efficient heat transfer in a limited mass of material, mass conservation (without major apophyses), and identifiable unmelted and melted components in the pseudotachylyte and the surrounding fractured rock. Hence, near-equilibrium conditions are petrologically quantifiable using thermodynamic constraints. The selection of pseudotachylyte devoid of alteration/devitrification was based on the material color and smooth appearance. In particular, dark/black color of the whole pseudotachylyte or the outer rims of it are suggestive of ultramafic material possibly glassy without greenish or reddish coloration, which tends to be associated to geochemical modifications induced by serpentinization and/or oxidation occurring after the generation of the pseudotachylyte (Trouw et al., 2010).

Analytical Investigations 3.2

Rock textural analysis was conducted using different techniques. The selection of the pseudotachylyte to inspect analytically was based on the glassy appearance of the material via optical microscopy. observations of three (out of five) pseudotachylytes in thin section were acquired using a Nikon Eclipse LV100N POL petrographic microscope at the Department of Geology of the University of Georgia (USA). Microstructural analysis of the same rocks was conducted using backscattered electron (BSE) imaging at 15 kV accelerating voltage and 1 nA beam current on a Thermo Fisher Scientific Teneo field emission scanning electron microscope (SEM). We then focused on one single sample of pseudotachylyte showing the largest thickness of the inner unaltered glassy portion for geochemical characterization via secondary electron (SE) imaging using a Hitachi SU-9000 low voltage scanning transmission electron microscope (STEM) (San Gabriel et al., 2024) at the Georgia Electron Microscopy Center of the University of Georgia. Major element analysis of the experimental glasses was performed on the same pseudotachylyte with a JEOL JXA-8350F HyperProbe (EPMA) at the Institute of Earth Sciences of the University of Lausanne (Switzerland), using a 5 nA beam current and 1 μ m beam diameter for nominally anhydrous minerals (clinopyroxene, orthopyroxene, olivine, and oxides), and 10 μ m beam diameter for amphibole and glassy pseudotachylyte. The counting times on the peak and background positions were 30 and 15 s for Al, Ti, Ca, Mg, and Mn and 10 and 5 s for Si, Fe, Na, K, P, and Cl, respectively. Natural and synthetic standards were used and data were ZAF-corrected (Armstrong, 1988), including secondary glass standards of hydrous (5.7 wt.% H₂O) basalt (Müntener et al., 2001). X-ray distribution maps of Si, Al, and Mg were acquired using the Oxford large area windowless energy dispersive spectroscopy (EDS) of the STEM at 30 kV and 10 μ A. Retrieval of standard chemical formulas for minerals from chemical analyses is based on spreadsheet recalculations here reported: https://serc.carleton.edu/ research_education/equilibria/mineralformulaerecalcu lation.html.

Calculations of Melt Polymerization

Using the pseudotachylyte glass composition measured via EPMA, we estimate the first-order approximation to the degree of polymerization of the original melts forming the pseudotachylytes using the numbers of non-bridging oxygens per tetrahedra described by the NBO/T parameter (Giordano et al., 2008). The NBO/T parameter is calculated according to the formulation proposed by Mysen (1988) by assuming that SiO_2 , TiO₂, Al₂O₃, and Fe₂O₃ are network formers and total iron (FeO_T) is split equally between FeO and Fe₂O₃ (Giordano and Dingwell, 2003; Giordano et al., 2006). Such a calculation is applied under anhydrous conditions. Under hydrous conditions, we are not able to assess the exact amount of H₂O dissolved in the glass based on the volatile by-difference method from EPMA analysis of pseudotachylyte composition (Fleet

and Pan, 1995) due to the unavoidable inclusion of the serpentinized olivine rims or devitrified borders of the pseudotachylyte during EPMA measurements which yield for water concentrations higher than those expected for the unserpentinized portion of the pseudotachylyte.

3.4 Mass Balance Calculations

The evaluation of the melted components (produced via melting of original mineral phases) contributing to the bulk composition of pseudotachylyte is conducted using a combination of chemical analyses of each mineral and glass phase (using EPMA analysis of mineral phases and STEM-based EDS analysis of glass phase), the volumetric fraction of each phase in the pseudotachylyte (through inspection using optical microscopy, SEM-based BSE imaging, and STEM-based SE imaging), and the set of mass balance calculations based on the algorithm of Albarede and Provost (1977) and the linear algebra formulation described by *Draper* and Johnston (1992). The system of equations in the mass balance is represented by the matrix expression B, in which A is the matrix of phase compositions, x is the vector of the weight proportions of each phase, and B is the vector representing the bulk composition of the pseudotachylyte on the anhydrous basis. Mass balance calculations (including propagation of errors from microscale EPMA analysis and nanoscale EDS-based data) were set up to four coexisting phases (olivine, orthopyroxene, clinopyroxene, and Al-Cr spinel) to evaluate the minerals and their proportions contributing to the bulk composition of pseudotachylyte. This mass balance calculation is also accompanied by the isocon analysis (Grant, 1986, 2005) based on the Grensens' (Gresens, 1967) equation quantifying the composition-volume relations between measured pseudotachylyte (on the anhydrous basis) and pseudotachylyte obtained by melting olivine, orthopyroxene, clinopyroxene, and spinel in different modal proportions. This approach further helps evaluate the conservation of the chemical components during the melting process forming the pseudotachylyte and the efficiency of partial melting of minerals contributing to the composition of the pseudotachylyte.

3.5 Calculations of Diffusion of Heat, Water, and Chemical Elements

In the discussion section (section 5.2), we estimate the timescales of diffusion of heat, major elements, and H₂O to evaluate whether pseudotachylytes are formed under equilibrium or disequilibrium conditions. the latter scenario, within the duration of formation of pseudotachylyte, we expect to observe evidences of incomplete melting, crystallization, inefficient heat transfer, and/or mass loss that underline potential disequilibrium conditions. Both heat and chemical diffusion are described by the first law of Fick (Fick, 1855) that relates the diffusive flux to concentration (of heat or a chemical component) in a spatial derivative under the assumption of steady state. Unidimensional (1D) heat or thermal diffusion timescale with a characteristic length scale (which is equivalent to the thickness of the pseudotachylyte considered in this study) and thermal diffusivity are estimated using the Fourier number (baron de Fourier, 1822). Diffusion of chemical components in melt during the melting process is described by the second law of Fick (Fick, 1855) that predicts how diffusion causes the concentration to change with time. Water diffusion is described by the simple approximation of the H₂O diffusion length scale $l = \sqrt{t \times D_{H_2O}}$ (Watson, 1982; Baker, 1991) with t as timescale and D_{H_2O} as water diffusivity determined using the pressure-dependent formulation of Ni and Zhang (2008). The formation of equilibrium conditions of glassy pseudotachylyte occurs when the following conditions are met: i) the thermal diffusion timescale is shorter than the frictional melting duration (0.1 to 10 seconds; Ferrand et al. 2018), which allows the rapid and complete melting of the specific mineral phases during thermal runaway (Spang et al., 2024) at certain pressure and temperature conditions; and ii) the timescale of chemical diffusion of both major elements and water in the melt is longer than the duration of melting and cooling processes leading to pseudotachylyte formation, implying that crystallization and modification of the associated glass composition do not occur in the pseudotachylyte.

3.6 Thermodynamic Modelling

To the goal of offering a thermodynamically rigorous petrological assessment of pseudotachylyte formation in the discussion (sections 5.3-5.4), we used the software package Perple_X (Connolly, 1990, 2005) (version 6.8.1) to compute stable mineralogy and phase diagrams (i.e., pseudosections; Connolly and Petrini 2002) as a function of P-T for the Balmuccia anhydrous and hydrous (0.5–1.0 wt.%) lherzolite composition (Hartmann and Hans Wedepohl, 1993) by Gibbs free energy minimization (Connolly, 2009). We used the thermodynamic database labelled as HP02, which includes the dataset of end-member properties of Holland and Powell (1996) and solid-solution models for olivine—O(HP), garnet—Gt(HP) (Holland and Powell, 1996), pyroxenes—Cpx(HP) and Opx(HP) (Holland and Powell, 1998), plagioclase—Pl(I1,HP) (Holland and Powell, 2003), spinel—Sp(HP) (Jamieson and Roeder, 1984), and amphibole—Amph(DHP) (Dale et al., 2000). We explore the different stability fields of melt-free and melt-bearing mineral assemblages divided by the respective anhydrous and hydrous solidus temperatures in a wide range of pressure (0.0001 to 4 GPa) and temperature (700 to 1600°C). The P-T space in these phase diagrams allows to summarize the P-T evolution of the Balmuccia lherzolite through time (from 433 to 180 Ma) in a large depth range from 124 km to surface using the hydrostatic/lithostatic pressure equation P = $P_0 + \rho g h$, where P_0 is the atmospheric pressure, ρ is the spinel lherzolite density of 3300 kg/m^3 , g is the average gravitational acceleration of 9.81 m/s², and his depth. The spinel lherzolite density is derived from previous Perple X calculations of Pistone et al. (2020) reporting estimates ranging from 3305 to 3325 kg/m³ in the pressure range of 0.9-1.6 GPa at <700°C.

In the chosen P-T space, we explore and compare the conditions of frictional melting and decompression melting forming A-type pseudotachylyte between 700°C, below which the lherzolite is a "frozen" lithology (Ewing et al., 2015; Smye et al., 2019) using an average continental geotherm from Christensen and Mooney (1995), and 1600°C chosen just above the reference peritectic reaction temperature of 1557°C at 0.0001 GPa of forsterite-enstatite in the forsterite-SiO₂ binary phase diagram (Bowen and Andersen, 1914; Creig, 1927). Within the proposed P-T space, we expect that the formation of A-type pseudotachylyte occurs between 0.9 and 1.1 GPa, which is the originally proposed pressure range of formation of A-type pseudotachylytes (Handy et al., 1999; Ueda et al., 2008), and lower pressure conditions at which decompression melting should assist frictional melting process.

4 Results

The microstructural analysis shows that a microscale pseudotachylyte cross-cuts the primary minerals (olivine and pyroxenes) of the lherzolite (Figure 3B-C), displacing them sinistrally with an offset of ~ 1 cm based on the displacement of light-colored olivine (see yellow dashed lines surrounding the olivines in Figure 3A). The central portion of the pseudotachylyte appears to be mainly glassy owing to its transparent to weakly absorbent aspect to transmitted light using the petrographic microscope (Figure 3C). Potentially serpentinized olivine rims or devitrified pseudotachylyte boundaries are found at the periphery of the pseudotachylyte in contact with the host rock and contain accessory spinel along the olivine-pseudotachylyte contacts (Figure 3D). Serpentine is related to late-stage fluid percolation accompanying deformation at lower temperatures (200–400°C) during Alpine orogenesis (Handy and Stünitz, 2002; Ueda et al., 2020); thus, serpentine is not associated to the original A-type pseudotachylyte composition. Microscopic apophyses (50 to 200 μm long) are found along the primary pseudotachylyte feature (Figure 3B-D) and are suggestive of limited mass transport (<5% of the pseudotachylyte vein area observed in thin section; Figure 3B) from the main pseudotachylyte A few suspended grains of clinopyroxene are found in the central portion of the preserved glassy pseudotachylyte (Figure 3D) in agreement with previous observations of the same minerals in amorphous pockets within ultramylonite-like veins found in the Balmuccia peridotite (Ferrand et al., 2018). Accessory amphibole located at the boundaries of primary minerals cross-cut by pseudotachylytes (Figure 3B) are associated to percolation of hydrous (H₂O-bearing) mafic melts in the peridotite from sublithospheric mantle metasomatized during subduction processes in the Variscan prior to the Carboniferous emplacement of the peridotite in the lower crust (Mazzucchelli et al., 2009), thus also prior to the formation of A-type pseudotachylytes. Minerals in the lherzolite (forsteritic olivine, diopsidic clinopyroxene [Wo₄₉En₄₇Fs₄], enstatitic orthopyroxene [Wo₁En₉₀Fs₉], Al-Cr-spinel, and pargasitic amphibole) were identified

through quantitative analyses of major elements using EPMA (see supplementary data at *Pistone*, 2025). On the anhydrous basis, the chemical composition of the pseudotachylyte is mainly made by SiO₂ (48 wt.%) and MgO (44 wt.%) followed by FeO_T (6.5 wt.%), and a limited concentration of Al₂O₃ (1 wt.%), while the same composition is CaO-poor (0.1 wt.%) along with MnO (0.1 wt.%), TiO_2 and Na_2O (<< 0.1 wt.%), and has no K₂O (see supplementary data at *Pistone*, 2025). Nanoscale analysis of the glassy portion of the pseudotachylyte (devoid of the serpentinized olivine rims or devitrified pseudotachylyte borders reported in Figure 3D) reveals droplets of amorphous materials separated by curvilinear contacts (Figure 4) and the presence of distinctive nanometer clinopyroxene minerals displaying sharp corners or edges suspended in the glass matrix (Figures 3C–D and 4). Through the geochemical inspection, we identify that the matrix appears to be made of two glasses derived from melting Al-Cr spinel and orthopyroxene, respectively, as observed in the chemical profiles and maps of Si, Al, and Mg (Figure 4). These chemical analyses reveal that the glass composition is best described by a coordination of 12 oxygens per formula unit, which is inconsistent with the stochiometric oxygen coordination expected for the cation occupancy in minerals such as spinel (4 oxygens per formula unit) and orthopyroxene (6 The excess number of oxygens per formula unit). oxygens is suggestive of presence of unmeasured volatile component. The presence of volatile components (e.g., H_2O) is also suggested by low EPMA totals (~85 wt.%) for pseudotachylyte compositions (see supplementary data at Pistone, 2025), although such as EPMA totals are acquired at microscale with defocused beam (10 μ m) used for amorphous phases and thus are partly affected by the presence of serpentinized olivine rims or devitrified pseudotachylyte material included in Under anhydrous conditions, the two the analysis. glasses are characterized by NBO/T values of: 2.36 for depolymerized orthopyroxene glass (due to large concentration of network modifiers such as MgO) and 0.17 for polymerized Al-Cr spinel glass (due to the presence of network formers such as Al_2O_3). hydrous conditions, the NBO/T parameter would increase with increasing NBO in presence of H₂O added into the melt (Mysen, 2014). The isocon analysis showcases that the slope of the isocon of all major elements with concentrations >1 wt.\% is 1 ± 0.1 , except for Al₂O₃ and FeO_T having isocon slopes of 0.4 ± 0.1 and 0.8 ± 0.1 , respectively. Both isocon slopes for Al_2O_3 and FeO_T are sensitive to the low spinel modal proportion (<1 vol.%), in the studied lherzolite.

5 Discussion

5.1 Nanoscale Analysis Reveals the Fusible Mineral Components Forming Pseudotachylytes

At nanoscale, we identify two melted components (Al-Cr spinel and orthopyroxene) forming the glass matrix of the pseudotachylyte and one unmelted component

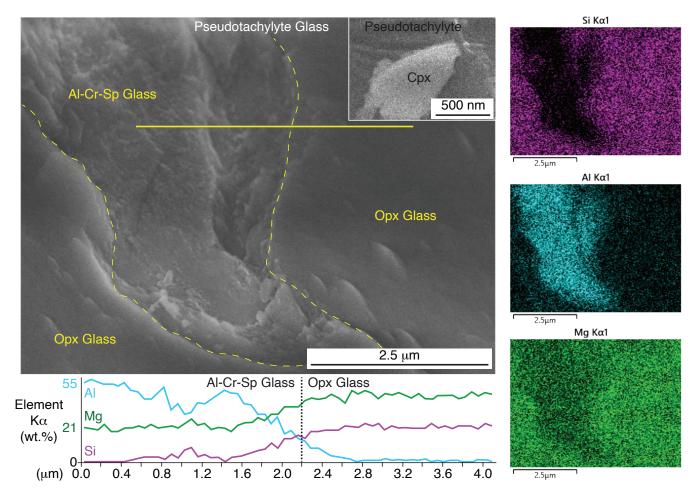


Figure 4 – Nanoscale view of pseudotachylyte. STEM-based SE image of pseudotachylyte glass and EDS analyses by applying a transect (shown as a yellow solid line) of individual analysis spots and X-ray distribution maps of Si, Al, and Mg K α from the Al-Cr spinel glassy droplet to the orthopyroxene glass droplet. The two droplets are divided by yellow dashed lines. The stippled vertical line in the chemical plot marks the point where the yellow horizontal line intersects the dashed curve in the above image. Same abbreviations as in Figure 3.

represented by clinopyroxene. Nanometer-sized clinopyroxene minerals suspended in the glass matrix may have formed by mechanical abrasion along the fault plane, yielding their angular shapes. However, they are unlikely to have experienced protracted shearing within a granular gouge as this would have resulted in rounded shapes (*Ujiie et al.*, 2007). Also, crystallization of clinopyroxene in the pseudotachylyte melt upon cooling appears to be unlikely under either anhydrous conditions, which hamper fast crystallization rates, or hydrous conditions, which tend to suppress nucleation of new minerals (Mollo and Hammer, The nanoscale analysis better corroborates 2017). which fusible minerals contribute to the formation of pseudotachylytes and helps identify whether melting is congruent (a phase melts to a liquid with the same composition as the original mineral) or incongruent (a phase melts to a liquid with a composition different from the original mineral and produces a new mineral of different composition to the original mineral). Our nanoscale analysis of the glass matrix does not reveal the presence of either a melted olivine component or olivine nanocrystals. Olivine could be an expected mineral phase involved in the peritectic reaction (incongruent melting) of orthopyroxene with residual

melt with increasing temperature (Bowen, 1928). If we had applied only microscale analysis, the average bulk composition of the pseudotachylyte analyzed in the Balmuccia lherzolite (see supplementary material at Pistone, 2025) should be matched by melting 60 vol.% olivine, 27 vol.% orthopyroxene, 12 vol.% clinopyroxene, and 1 vol.% Al-Cr spinel using the set of mass balance calculations and isocon analysis described in section 3. However, the pseudotachylyte found in the Balmuccia lherzolite displays serpentinized olivine rims or devitrified pseudotachylyte glass, especially along the margins of the pseudotachylyte vein (Figure 3D), which have been unavoidably included in the EPMA analyses, as testified by the low EPMA totals (see supplementary material at *Pistone*, 2025). Therefore, in contrast to previous microscale analyses (Obata and Karato, 1995) and our microscale analyses of pseudotachylytes, nanoscale analysis can sufficiently distinguish the compositional heterogeneities within the glass matrix and the suspended mineral phases therein (Figure 4). In this study, nanoscale analysis helps to discriminate the components that melted alone/once (Al-Cr spinel and orthopyroxene) from unmelted components (clinopyroxene suspended in the glass; Figure 4; and olivine surrounding the pseudotachylyte; Figure 3) and assess the "unserpentinized" composition of the pseudotachylyte. We do not find a melted olivine component anywhere in the nanoscale inspection. The absence of CaO in the pseudotachylyte composition along with the presence of clinopyroxene grains suggest the exclusion of clinopyroxene as a melted component contributing to the pseudotachylyte geochemistry. We thus propose that congruent melting of orthopyroxene and Al-Cr-spinel occurred during pseudotachylyte formation.

5.2 Are Pseudotachylytes the Result of Equilibrium or Disequilibrium Melting?

To date, pseudotachylytes have been interpreted as rocks formed under disequilibrium conditions caused by rapid cooling of melts formed rapidly during rupture and seismic slip of rocks (Sibson, 1975; Allen, 1979; Maddock, 1986, 1992; Magloughlin, 1989; Bossière, 1991; Spray, 1992, 1993; Camacho et al., 1995; O'Hara and Sharp, 2001; Andersen and Austrheim, 2006). However, this conventional interpretation is counterintuitive as disequilibrium cannot be explained only in terms of most effective transfer of kinematic/elastic energy to thermal energy linked to frictional heat (Spang et al., 2024), which then leads to frictional fusion and generation of pseudotachylyte. This efficient conversion of kinematic energy into heat is satisfied when slip velocity approaches seismic rate causing the dynamic coefficient of friction to fall to zero (Di Toro et al., 2004) and when heat diffusion proceeds within a timescale that is shorter or identical to the pseudotachylyte formation via melting the original rock undergoing fracturing and high-temperature friction. Thus, determining whether a phase (glass or mineral) is formed under equilibrium or disequilibrium conditions depends on the rate at which equilibrium is attained along the P-T-time path (Powell, 1978). In the case of formation of glassy pseudotachylyte in the Balmuccia lherzolite, frictional or flash melting is a rapid process (on the order of 0.1 to 10 seconds; Ferrand et al. 2018) that allows original mineral phases to melt. The formation of glassy pseudotachylytes under equilibrium conditions must include: a) the efficient diffusion of heat through the fracturing region where the pseudotachylyte forms, b) the efficient transfer of geochemical components from minerals to melt via frictional melting, c) the efficient diffusion of H₂O that helps suppress the solidus and thus, the minimum temperature required to melt the minerals forming the pseudotachylyte, d) the concomitant inefficient diffusion of chemical components that allows to identify the mineral phases fused into the pseudotachylyte melt, e) the very rapid cooling to quenching that freezes the glassy pseudotachylyte composition. Here is the detailed description of each aspect related to the generation glassy pseudotachylytes under equilibrium conditions:

a) The short timescale of frictional melting (0.1 to 10 seconds; Ferrand et al. 2018) forming pseudotachylytes impedes their crystallization. The efficiency of frictional melting of selective minerals without generating new

crystallites is driven by efficient thermal diffusion and large degree of undercooling that suppresses the nucleation and growth of new minerals. Thermal diffusion during pseudotachylyte formation occurs in 0.02 to 2 seconds along the pseudotachylyte-filled space of 10 μ m to 1 mm (Figure 3) as thermal diffusivity in a melt is $<5 \times 10^{-7}$ m²/s at >900°C (*Romine et al.*, 2012).

- b) Chemical multicomponent diffusion from mineral to melt during melting becomes increasingly fast with the diffusion profile in the crystal shortening with the progress of the mineral dissolution into melt (Zhang, 2008, 2010).
- c) Water diffuses faster than any other major element by about three orders of magnitude (Ni and Zhang, 2008). The in-melt diffusion of H_2O derived from hydrous minerals (amphibole; Figure 3B) and fluids commonly located along mineral grain boundaries (Stünitz et al., 2024) helps depress the system solidus and liquidus and impede crystallization by largely increasing the degree of undercooling (Hort, 1998) as H₂O diffusion takes about 100 seconds with H₂O diffusivity of $\sim 10^{-10}$ m^2/s at limited concentrations (0.5–1 wt.%) at 1000 to 1400°C (Van Der Laan and Wyllie, 1993; Ni and Zhang,
- d) Once the mineral phases melt at lower solidus temperature under hydrous conditions favored by efficient H₂O diffusion in the pseudotachylytic melt, the diffusion of the chemical components derived from each mineral results slow and inefficient because the diffusion of major elements such as Si, Al, and Mg having diffusivities of 3×10^{-12} m²/s to 9×10^{-12} m²/s in a melt (Van Der Laan and Wyllie, 1993) is slower than both H₂O and heat diffusion by three to five orders of magnitude (Hofmann, 1980; Watson, 1981; Sparks and Marshall, 1986) and, thus, slower than the rate of formation of a pseudotachylyte through melting and cooling stages.
- e) Then, rapid cooling (accompanied by a large degree of undercooling that hampers crystallization) to ambient temperature on the scale of seconds to minutes following frictional melting (O'Hara and Sharp, 2001; Andersen and Austrheim, 2006; Ferrand et al., 2018) "freezes" the newly formed melt in the form of glass.

Therefore, the combination of efficient melting of fusible minerals and impeded crystallization during the short timescale makes the chemical composition of pseudotachylyte glass an informative snapshot of the P-T conditions of formation of pseudotachylyte under near-equilibrium conditions achieved by efficient thermal diffusion and mineral dissolution within the timescale of rock heating, melting, and solidification into glassy pseudotachylytes. This "frozen melt" records the last snapshot of P-T conditions following frictional melting envisioned as a dominant T increase under isobaric conditions as it has been proposed to date (McKenzie and Brune, 1972; Sibson, 1975, 1980; Spray, 1992, 1995; Obata and Karato, 1995; Kanamori et al., 1998; Bouchon and Ihmlé, 1999; Hirose and Shimamoto, 2005; Di Toro et al., 2006a, 2011; Kelemen and Hirth, 2007; Lin, 2007;

Ueda et al., 2008, 2020; Del Gaudio et al., 2009; Spang et al., 2024).

The timescale of solidification of 5-mm thick pseudotachylytes here investigated is on the order of 2 seconds, sufficient to freeze the newly formed pseudotachylyte without undergoing crystallization upon Based on the equation 3 reported in cooling. Andersen and Austrheim (2006) derived from Turcotte and Schubert (2002) detailing the solution of the Stefan problem (Stefan, 1891) applied to cooling silicate liquids, we estimate the timescale of solidification of pseudotachylyte from the Balmuccia lherzolite partial melting using these parameters: the thermal diffusivity of 5×10^{-7} m²/s (Romine et al., 2012), the latent heat of melting of 5.8×10^5 J/kg, the heat capacity of 1793 J/kg/°C (Lesher and Spera, 2015), and a temperature rise of 400°C (hydrous conditions) to 540°C (anhydrous conditions) as estimated using Perple X (see sections 5.3 and 5.4) to determine the dimensionless variable of 0.81 quantifying the latent heat liberated at the solidification boundary conducted vertically upward (Turcotte and Schubert, 2002), away from the interface between partially molten pseudotachylyte and unmelted fractured rock.

We identify the two chemical components derived from the melting of mineral phases of Al-Cr spinel and orthopyroxene as two distinctive immiscible glassy droplets in the pseudotachylyte matrix (Figure 4). The slow diffusion of major elements from Al-Cr spinel and orthopyroxene allows the geochemical distinction of the two minerals as melt/glassy components through the EDS analysis (Figure 4). Furthermore, the short timescale of melting and cooling processes impedes the efficient blending or complete miscibility (Lesher and Spera, 2015) of Al-Cr spinel and orthopyroxene components. The slow chemical diffusion with respect to the short timescale of formation of pseudotachylyte hampers the generation of mineral nuclei displaying dendritic habit due to large undercooling caused by rapid cooling (i.e., quenching), heat loss or inefficient chemical diffusion through the different melt components (Fenn, 1977; Van Der Laan and Wyllie, 1993; Webber et al., 1997; Nabelek et al., 2010; Pistone et al., 2016). In addition, the large discrepancy in polymerization (i.e., viscosity) of the two melts with NBO/T of 2.36 for depolymerized orthopyroxene melt and 0.17 for polymerized Al-Cr spinel, respectively, favor their immiscibility (Jarvis et al., 2021) and distinction through geochemical analysis (Figure 4).

5.3 Petrological Assessment of Pseudotachylyte Formation Under Anhydrous Conditions

The glassy pseudotachylyte from this study is devoid of pulverized grains derived from mechanical friction, new mineral phases upon cooling, and/or cryptocrystalline phases suspended in the glass matrix at very fine spatial scale (on the order of $<2 \mu m$; Figure 4). We only identify two distinctive glasses with suspended nanometer-sized clinopyroxene crystals (Figure 4). These observations

highlight that the formation of glassy pseudotachylytes can be considered under near-equilibrium conditions based on the quantitative comparison between the timescales of heat diffusion, chemical diffusion, and solidification and the timescales of rapid melting and cooling required for pseudotachylyte formation (see section 5.2). Under such conditions we can attempt to apply thermodynamic modelling that helps reconstruct the pressure (P) and temperature (T) pathway of pseudotachylyte formation based on the collected observations at the micron (Figure 3B–D) and submicron scales including our new compositional constraints of the pristine or "unserpenitinized" pseudotachylyte glassy components (Figure 4). Specifically, we can apply thermodynamic calculations to reconstruct stable mineralogy and phase diagrams for the lherzolite composition as a function of P-T conditions experienced by the Balmuccia Massif in the pre-Alpine orogenesis period from 433 to 180 Ma (Figures 5–6). We remind the reader that, during this period, A-type pseudotachylytes were generated (Souquière and Fabbri, 2010; Souquière et al., 2011; Ferré et al., 2017). These are most likely related to the peridotite exhumation during the Carboniferous Period (*Ueda et al.*, 2008).

Below 700°C, the exhumed Balmuccia peridotite can be considered petrologically "frozen" (Ewing et al., 2015; Wyatt et al., 2022) with an average geotherm of $\sim 18^{\circ}$ C/km for a ~ 40 -km thick continental crust made of felsic granulites (using the temperature coefficient of 5×10^{-4} km/s/°C derived from average heat flow and seismic compressional wave profiles; Blackwell 1971; Lachenbruch and Sass 1977; Christensen 1979; Christensen and Mooney 1995; Berno et al. 2020), closely related to the Paleozoic lower crust made of granulite-facies metapelites in the Ivrea-Verbano Zone Herein, we evaluate the conditions of (Figure 1). pseudotachylyte formation in P-T space using the composition of the pseudotachylyte glass (made of Al-Cr spinel and orthopyroxene) with suspended/unmelted clinopyroxene (Figures 3-4) and olivine (Figure 3). All the phases identified within and around the pseudotachylyte glass help reveal which phases effectively melted and those that did not melt during the frictional melting process. Using the anhydrous (devoid of H_2O) and hydrous (0.5–1.0 wt.% H_2O) scenarios (Figures 4–5), we aim to test two end-member processes: pure frictional melting under isobaric T increase as envisioned in previous studies (McKenzie and Brune, 1972; Sibson, 1975, 1980; Spray, 1992, 1995; Obata and Karato, 1995; Kanamori et al., 1998; Bouchon and Ihmlé, 1999; Bjornerud and Magloughlin, 2004; Hirose and Shimamoto, 2005; Di Toro et al., 2006a, 2011; Kelemen and Hirth, 2007; Lin, 2007; Ueda et al., 2008, 2020; Del Gaudio et al., 2009; Souquière and Fabbri, 2010; Souquière et al., 2011; Kirkpatrick and Rowe, 2013; Ferrand et al., 2018; Fondriest et al., 2020; Dunkel et al., 2021; Dai et al., 2022; Moris-Muttoni et al., 2022), and fracture-induced decompression melting, newly proposed in this study.

If we consider the Balmuccia lherzolite as an anhydrous system (Figure 5), the formation of

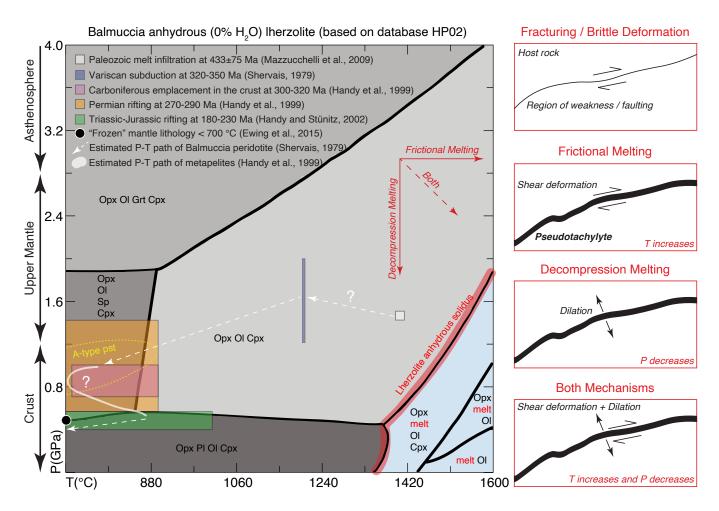


Figure 5 – P–T plot of Balmuccia anhydrous lherzolite. Perple_X-based P–T plot of Balmuccia lherzolite throughout different processes from Paleozoic to Jurassic (highlighted by dashed white arrows). Stability fields in light blue contain melt phase with the position of the anhydrous rock solidus temperature highlighted in red. The area delimited by dashed yellow line indicates the P–T conditions of A-type pseudotachylyte stability originally proposed by previous studies ($Handy\ et\ al.$, 1999; $Ueda\ et\ al.$, 2008). Sketches and red arrows indicate the conditions of both frictional and decompression melting in P–T space. For better display, the boundaries between different mineral assemblages are improved by manually interpolating the "step edges" of the original lines generated by the Perple_X software. Same abbreviations as in Figure 3, plus: Grt = garnet, melt = ultramafic melt, Pl = plagioclase, pst = pseudotachylyte.

pseudotachylytes during the Carboniferous Period (T = $720-900^{\circ}$ C and P = 0.9-1.1 GPa; Handy et al. 1999; Ueda et al. 2008) requires intense friction and heat production to partially melt the minerals in the lherzolite along the fault plane. In this case, for pure frictional melting, a T increase of ~ 540 °C is needed from the starting 900°C to surpass the anhydrous rock solidus temperature (Figure 5). Elastic fracture opening or shear-induced dilatancy will induce a P-drop (Samuelson et al., 2009; Nüchter, 2017; Brantut, 2019) during rock friction-induced heating (with temperature increase from the pink area of Carboniferous emplacement toward the anhydrous lherzolite solidus reported in Figure 5). This could be total (i.e., to 0 Pa), but is likely to be less because melt can migrate into the fractures and partly mitigate the pressure drop. A pressure drop does not appear to influence heat transfer (Everts and Meyer, 2018) required to induce partial melting and form pseudotachylytes. This migration of melts into open fractures becomes especially efficient and rapid when the produced melts are highly depolymerized (such as orthopyroxene liquid with NBO/T = 2.36) and,

thus, characterized by very low viscosity (Mysen, 1988) and in presence of high compaction length of the pore space within the fractured rock (Liang et al., 2010). For a P drop of 0.3–0.7 GPa, frictional melting only requires a T increase of $\sim 440^{\circ}$ C from 900°C to surpass the anhydrous rock solidus temperature (Figure 5). However, under anhydrous conditions, the melt-bearing mineral assemblage does not match the composition of the natural pseudotachylyte found in the Balmuccia lherzolite (Figure 4). Furthermore, anhydrous conditions that make the rock solidus lying at significantly high temperatures (>1340°C at 0.0001-0.6 GPa; Figure 5) should favor rock comminution forming fine-grained polyphase aggregates in the form of "pulverized" granular pseudotachylytes and ultramylonites (Menegon et al., 2017, 2021), whose rheology differs greatly from the generation of glassy pseudotachylytes (Figures 3-4) via partial melting considered in this study.

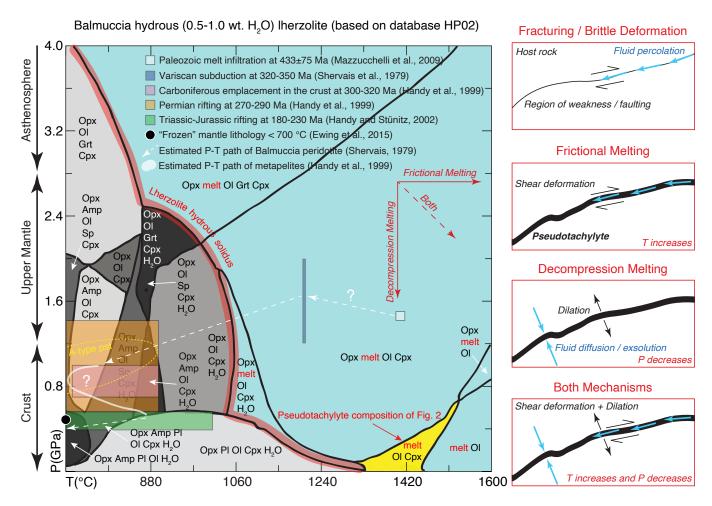


Figure 6 – P–T plot of Balmuccia hydrous (0.5-1.0 wt.% H₂O) lherzolite. Perple_X-based P–T plot of Balmuccia lherzolite throughout different processes from Paleozoic to Jurassic (highlighted by dashed white arrows). Stability fields in light blue contain melt phase with the position of the hydrous rock solidus temperature highlighted in red. The yellow field highlights that the melt composition corresponds to the pseudotachylyte composition (Al-Cr spinel- and orthopyroxene-like chemistry) with suspended clinopyroxene minerals of Figure 4 and surrounding olivine minerals of Figure 3. The area delimited by dashed yellow line indicates the P–T conditions of A-type pseudotachylyte stability originally proposed by previous studies (Handy et al., 1999; Ueda et al., 2008). Sketches and red arrows indicate the conditions of both frictional and decompression melting in P–T space in the presence of fluids (in light blue). For better display, the boundaries between different mineral assemblages are improved by manually interpolating the "step edges" of the original lines generated by the Perple_X software. Same abbreviations as in Figures 3 and 5, plus: H_2O = water as exsolved fluid.

5.4 Petrological assessment of pseudotachylyte formation under hydrous conditions and the role of decompression melting

We expect that the Balmuccia lherzolite must contain limited $\rm H_2O$ concentrations (0.5–1 wt.%) distributed in hydrous minerals (amphibole; Figure 3B), nominally anhydrous minerals, and/or shear zones of the mantle rock in which fluid percolation is assisted by deformation. The presence of water during the formation of pseudotachylytes is confirmed by previous studies reporting that relatively unaltered pseudotachylytes are hydrous glasses (up to 1.5 wt.% $\rm H_2O$) containing vesicles, suggestive of fluid exsolution during pseudotachylyte melting at temperatures and pressures at which excess water is not soluble in the melt (Obata and Karato, 1995; O'Hara and Sharp, 2001; Ferrand et al., 2018). Presence of water in the bulk lherzolite rock expands the P-T conditions under which the rock system can

be partially molten as water helps suppresses the rock solidus to lower T conditions in a wide range of pressure (Figure 6). In this scenario, the pseudotachylyte can form by a much more limited T increase (only 160° C above 900°C) via pure frictional melting. However, to obtain a pseudotachylyte matching the chemical composition derived from melting both Al-Cr-spinel and orthopyroxene (at T = 1300-1530°C and P < 0.5GPa; Figure 6), decompression must aid frictional melting. Dilational jogs along deep crustal faults could be sites of near-total pressure release and, thus, decompression melting (Bjornerud and Magloughlin, 2004) or even boiling (forming bubbles/vesicles as documented by Obata and Karato 1995) at sufficiently low pressure (Williams, 2019). Previous work proposed that decompression melting is localized and only applies to significant depth (relevant to eclogite facies) with significant pressure drops (>1 GPa) (Bjornerud and Magloughlin, 2004). However, here we show that fracture-induced decompression melting can occur at P

<1 GPa in the granulite facies in the lower crustal section of the Ivrea-Verbano Zone (Figure 1). Indeed, a decompression of 0.3-0.9 GPa and T increase of 400°C are both required to generate Al-Cr-spinel and orthopyroxene liquids composing the ultramafic composition of the pseudotachylyte (Figure 6). This observation also supports that selective melting of minerals is more petrologically rigorous (Dobson et al., 2021) than complete melting of the total rock as previous studies suggest based on microscale chemical analysis of pseudotachylytes (Obata and Karato, 1995; Ferrand et al., 2018) or based on both experimental and computational simulations that illustrate how rocks experience very high temperatures due to frictional heating (Del Gaudio et al., 2009; Lin and Shimamoto, 1998; Ferrand et al., 2018; Woo et al., 2023; Spang et al., 2024). Compared to the anhydrous scenario in which T increase via frictional melting is up to $\sim 540^{\circ}$ C (Figure 5), water derived from hydrous minerals or external fluids facilitates both shear localization and pseudotachylyte generation (Lin and Shimamoto, 1998; O'Hara and Sharp, 2001; Andersen and Austrheim, 2006; Ferrand et al., 2018). Indeed, the resulting T increase during frictional melting is reduced by 100 to 400°C corresponding to a temperature decrease of 18% to 74% (Figure 6), depending on the amount of frictional heat and pseudotachylyte melt generated in the faulted rock.

5.5 Does Overpressure Favor the Pseudotachylyte Formation?

Recently, other researchers have proposed that pseudotachylyte apophyses should be formed from overpressurized frictional melt with a pressure of 0.5 GPa greater than the lithostatic pressure (Zhong et al., 2021). Melt overpressure can be caused both by volume expansion due to melting and by heating of the faulted rocks (Sibson, 1975; Swanson, 1989, 1992; Zhong et al., 2021). However, this process of melt overpressurization, which is opposite to decompression, does not reproduce the geochemical composition of the pseudotachylyte inspected in this study (Figure 4). Thus, melt overpressure appears to be unfeasible within our thermodynamic constraints (Figures 55-6). Decompression melting might still be a primary process that occurs soon after fracture opening that results in pressure drop rather than pressure increase. Furthermore, such a decompression driven by fracture dilation may also be associated to dynamic stress drops related to lubrication effects of the pseudotachylyte melt as observed experimentally (Di Toro et al., 2009, and references therein). Decompression melting may also prevent other effects of dilatancy, such as pressure gradients that could cause melt to flow away from where it is generated (O'Hara and Sharp, 2001; Di Toro et al., 2005; Andersen and Austrheim, 2006; Ujiie et al., 2007), especially if the liquids produced are highly polymerized and viscous (thereby sluggish to flow) as they derive from mineral phases enriched in network formers such as Al_2O_3 from Al-Cr spinel.

Conclusions

In summary, decompression associated with dilation during fracture opening in presence of water derived from hydrous minerals and fluids from grain boundaries is a crucial process facilitating the generation of pseudotachylytes in the lithosphere upon fault rupture, slip, and resultant shear heating. Water as free fluid or initially bonded in minerals suppresses the solidus of the system to lower temperature when released. Decompression melting allows pseudotachylyte to be composed of amorphous mineral phases that are consistent with relatively lower P and higher Tconditions than those of the pre-failure host rock. We have demonstrated that seismic dilation can contribute to the generation of pseudotachylytes. With the analyses presented in our study, pseudotachlylytes can be the consequence of a seismic rupture facilitated by water and decompression, so that less shear heating than has been proposed to date is required.

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Author contributions

M.P. conceived of the idea and wrote the manuscript. M.P., V.G.T., and M.W.O. conducted fieldwork and M.P., M.R., and E.F. conducted rock sampling. the analytical investigations. All authors discussed the results and edited the manuscript.

Data availability

Data were acquired at the Georgia Electron Microscopy Center of the University of Georgia and at the electron probe micro-analyzer laboratory of the University of Data and supplementary material are published in a public repository: https://doi.org/10 .5281/zenodo.16884827.

Competing interests

The authors declare no competing interests.

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