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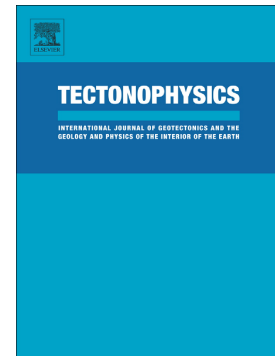
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*Large-scale strain localization induced by phase nucleation in mid-crustal
granitoids of the south Armorican massif*

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Abstract

The reduction of grain size is believed to play a critical role in strain localization to form shear zones. Although many mechanisms have been proposed, the source of grain size reduction remains debated. The South Armorican Shear Zone (SASZ) is a crustal-scale strike-slip fault that deforms granitoids at mid-crustal conditions. The SASZ records the transition from protolith to ultramylonite, representative of increasing ductile shear strain. To investigate the evolution of strain localization, the different states of deformation were studied using a combination of detailed microstructural, chemical and electron backscatter diffraction analyzes. Increasing strain from protolith to ultramylonite resulted in (1) grain size reduction, (2) the development of interconnected monophasic layers of mica and incipient mixed-phase zones composed of phengite-quartz \pm K-feldspar in protomylonite and low-strain mylonite, and (3) the formation of fine-grained mixed-phase zones composed of K-feldspar-

quartz ± phengite in high-strain mylonite and ultramylonite. We propose that the causes for interconnection of mica are the formation of cracks in the protolith combined with fluid-assisted nucleation. The latter process also plays a major role in phase mixing, as attested by the precipitation of K-feldspar at triple junctions in quartz-rich layers and in fine-grained tails of inherited K-feldspar porphyroclasts in ultramylonite. The transition from quartz-rich layers and mixed-phase zones is accompanied by a strong dispersion of the quartz lattice preferred orientation. These microstructural and textural evidences suggest that phase nucleation is the major process behind phase mixing, possibly accompanied by the action of grain-boundary sliding to open cavities during deformation. Instead of a single process, we therefore highlight a succession of weakening processes in the evolution of the SASZ, starting with the crystallization of mica as a first weakening material, and then evolving with the formation of very fine-grained mixed-phase zones made principally of feldspar, quartz and phengite.

Keywords

Strain localization; phase mixing; dislocation creep; diffusion creep; phase nucleation

1. Introduction

Strain localization in ductile shear zones is commonly characterized by the development of protomylonite, mylonite and fine-grained ultramylonite with increasing strain (e.g. Kilian et al., 2011; Linckens et al., 2011). As accommodating a large part of strain in large-scale structures, they play a crucial role for the dynamics of mountain belts, subduction zones or rifting. However, their occurrences have been considered as resulting from one or several processes of rheological weakening, the origins of which remain today very elusive (e.g. Stünitz and Fitz Gerald, 1993; Fliervoet et al., 1997; Fossen and Cavalcante, 2017). Investigating the processes responsible for large-scale strain localization is essential to understanding the behaviour of the lithosphere-scale structures over geological timescale.

Several processes have been proposed and first introduced in Poirier (1980) and White et al. (1980) to account for a local weakening during deformation. They include the occurrence of (1) metamorphic reactions (2) geometric softening (3) chemical softening (4) a change in deformation mechanism owing to grain size reduction (5) shear heating and/or (6) pore fluid pressure effects. In polyphase rocks, the presence of different phases with various strengths may also be an important cause of ductile strain localization, as they produce a framework of interconnected weak layers (e.g. Jordan, 1988; Handy, 1990; Shea and Kronenberg, 1993; Dell'Angelo and Tullis, 1996; Lonka et al., 1998; Stewart et al., 2000; Holyoke and Tullis, 2006). Of further interest is the mechanism related to grain size reduction as it systematically occurs in ductile shear zones (e.g. White et al., 1980; Rutter and Brodie, 1988; Herwegh et al., 2011; Kilian et al., 2011; Montési and Hirth, 2003; Platt, 2015). The role of grain size reduction in strain localization thus deserves a special attention. Moreover, grain size reduction may give rise to substantial weakening when the mean grain size is small enough for grain size-sensitive (GSS) creep to control the bulk rock rheology (e.g. Boullier and Gueguen, 1975; Etheridge and Wilkie, 1979; Rutter and Brodie, 1988; Stünitz and Fitz Gerald, 1993; Fliervoet et al., 1997; De Bresser et al., 1998; Kruse and Stünitz, 1999; Kenkmann and Dresen, 2002; Précigout et al.,

2007; Raimbourg et al., 2008; Linckens et al., 2011; Kilian et al. 2011; Viegas et al., 2016). Nevertheless, the sources of grain size reduction still remain poorly constrained and require further detailed studies of shear zones, particularly for polyphase rocks through large-scale structures.

In this contribution, we investigate the microstructural features of a granitoid mylonitized within a major crustal shear zone of South Brittany, Western France. This shear zone corresponds to the historical place of first recognition of S-C and S-C' fabrics (Berthé et al., 1979). Here, we focus on the strain gradient outlined by a gradual grain size reduction from undeformed protolith to ultramylonite. Using light microscope, cathodoluminescence (CL), backscattered electron (BSE) imaging along with electron backscatter diffraction (EBSD) analyzes, we describe the progressive evolution of the microstructure and associated changes in active deformation processes.

2. Geological setting

The French Armorican massif corresponds to one of the largest pieces of the Paleozoic Variscan orogenic belt, which extends throughout western and central Europe (e.g. Matte, 1991, 2001). The Variscan belt resulted from the collision between two major continental domains, the Gondwana continent to the south and the Laurussia continent to the north.

The structural pattern of the Armorican massif is dominated by late-Carboniferous W-E to NW-SE striking, dextral strike-slip shear zones, namely the North Armorican shear zone (NASZ) and South Armorican shear zone (SASZ) (e.g. Watts and Williams, 1979; Jégouzo, 1980; Matte, 1986; Tartèse et al., 2011). In the Armorican massif, the formation of crustal-scale shear zones is associated with widespread extension and magmatism during the collapse of the Variscan orogenic belt (e.g. Berthé et al., 1979; Jégouzo, 1980; Vignerresse and Brun, 1983; Ballèvre et al., 2009). Macrostructural studies demonstrated that deformation localized over time towards the contact between syntectonic plutons and host rocks (e.g. Berthé et al., 1979). From a paleogeographic point of view, the Armorican massif is divided into three main distinct parts, namely the Léon, North-Central and South-Armorican

domains, all bound by major tectonic contacts including suture zones such as the Nort-sur-Erdre deformed zone (Fig. 1a, e.g. Cartier and Faure, 2004; Faure et al., 2005, Ballèvre et al., 2009). These domains drastically differ from each other by their metamorphic grade and structural style (e.g. Gumiaux et al., 2004; Faure et al., 2008; Ballèvre et al., 2009; Turrillot et al., 2011; Augier et al., 2015).

The SASZ extends over hundreds of kilometers and represents a major structural zone that separates the North-Central and South Armorican domains (Fig. 1a). The SASZ is divided into two main branches roughly oriented WNW-ESE for the northern branch and NW-SE for the southern branch (Fig. 1a; e.g. Cogne, 1960; Jégouzo, 1980). The southern branch separates the South Armorican domain in two parts: 1) the Ligerian Domain to the North characterized by Early Paleozoic ophiolites, high-pressure mafic and felsic rocks (Ballèvre et al., 1994; Bosse et al., 2000), and 2) the coastal South Armorican domain, which exhibits metamorphic and granitic rocks (Brown and Dallmeyer, 1996; Turrillot et al., 2009, 2011; Augier et al., 2015).

The present study focuses on the southernmost parts of the Ligerian Domain along the Southern branch of the SASZ, where Neoproterozoic to Lower Paleozoic metasediments are intruded by Carboniferous granitoids traditionally described as synkinematic intrusion (Fig. 1b; e.g. Berthé et al., 1979; Tartèse and Boulvais, 2012). These granitoids are affected by a variable strain intensities, giving rise to local mylonites and ultramylonites (e.g. Berthé et al., 1979). This is notably the case for the Questembert granite, where strain increases southward and reaches a maximum in the core of the Southern branch of the SASZ (Fig. 1b; e.g. Augier et al., 2011; Tartèse et al., 2012).

The studied samples were collected along a section through a satellite body of the Questembert granite massif (Fig. 1b-c). This section represents a strain gradient from undeformed or slightly deformed host rocks of the SASZ (domain 1) towards the most deformed, ultramylonitic sample in the core of the SASZ (domain 5). Accordingly, the grain size decreases and the foliation progressively transposes into the shear plane from domain 1 to domain 5 (Fig. 1c). This section has been also completed by another sample collected in the Lescastel quarry to allow comparison with very recent results of Bukovská et al. (2016) (Fig.1b).

3. Material and Methods

Thin sections were cut parallel to the stretching lineation and perpendicular to the foliation (XZ plane). Microstructural and chemical evolution of the different finite strain states were studied using optical microscopy, cathodoluminescence (CL) imaging, electron backscatter diffraction (EBSD), and electron probe micro-analysis (EPMA).

3.1. Analytical Methods

To perform a CL analysis, the sections were placed in a vacuum chamber at Argon pressure of $\sim 60 \times 10^{-3}$ mbar. They were irradiated by a cold cathode source electron gun provided by OPEA (France). Standard tension-current beam conditions were maintained at ~ 13 - 15 kV and 90 - 110 μ A, respectively. The angle between the gun axis and thin section plane was set to 18° . The chamber was placed on a simplified optical mount designed to minimize light absorption on the beam pathway. The optical stage was equipped with a Retiga 2000R 1394a cooled camera (captor size 1 inch, resolution 1600×1200 pixel², square pixel size 7.4 μ m). Acquisition time was fixed to ca. 10 s for quartz and micas and ca. 2 s for feldspars.

EBSD was carried out with an EDAX PEGASUS EDS/EBSD system and OIM DC 6.4 software (manufacturer EDAX, Mahwah – USA) at the BRGM (Orléans, France). The polishing procedure of thin sections includes chemico-mechanical polishing using first diamond paste of 0.25 μ m followed by colloidal silica. The operating conditions involved an accelerating voltage of 20 kV, a working distance of 15 or 18 mm, and a probe current of ~ 6 nA. The EBSD data were processed to produce orientation maps and equal-area lower-hemisphere pole figures plotted as ‘one-point-per-grain’ in order to give the same weight to all grains independently of their size. The grains are defined so that their boundaries involve a minimum misorientation angle of 10° . The iso-contours on pole

figures were plotted using the OIMTM – EDAX software. Lattice preferred orientation (LPO) strength was quantified using the texture (J) and misorientation (M) indices (Bunge, 1982; Skemer et al., 2005). Pole figures and J-index were collected considering a Gaussian half-width angle of 10°.

Mineral compositions were collected using a CAMECA SX Five EPMA at the ISTO-BRGM (Orléans, France) on carbon-coated thin sections (20 nm thickness). The analyzes were carried out with an accelerating voltage of 15 kV, a beam current of ~6 nA and a beam size of ca. 1 µm.

3.2. Quantitative Microstructural Study

Scanning electron microscope/backscattered electron (SEM/BSE) and light microscope images have been used to determine grain size distributions of quartz. Within each strain domain, grains in quartz-rich areas have been digitized manually. To ensure statistical representation of data, about 22 000 quartz grains were processed. The grains were then analyzed using the software ImageSXM® (<http://www.ImageSXM.org.uk>) to quantify the grain size, grain contact frequency (GCF) and shape preferred orientation (SPO).

Two types of source images were used to quantify the grain size: transmitted light microscope images when grains were easily distinguishable and SEM/BSE images when the grain size was too small, including zones of mixed phases in mylonites and ultramylonites. The number weighted grain size was defined as the diameter of equivalent circular diameter ($d_{\text{equ}} = 2x \sqrt{\frac{\text{area}}{\pi}}$). The grain contact frequency (GCF) between quartz-quartz (Q-Q), feldspar-feldspar (F-F) and quartz-feldspar (Q-F) was evaluated using the method described by Heilbronner and Barrett (2014). In this method, GCF between grains is expressed as deviation from a random phase distribution. Results of GCF allowed us to define spatial distributions of grains as clustered or ordered, corresponding to an increasing degree of phase mixing. For a two-phase mixing where grains are randomly distributed, Heilbronner and Barrett (2014) define pQQ, pFF, and pQF as the predicted amount of quartz/quartz, feldspar/feldspar and quartz/feldspar boundaries, respectively. For a two-phase mixing pQ and pF are defined as phase

proportion where $pQ + pF = 1$. The random phase distribution that separates clustered and ordered domains is described by the following equations as a function of phase proportions:

$$p_{QQ} = pQ \cdot pQ = pQ^2$$

$$p_{FF} = pF \cdot pF = (1 - pQ)^2$$

$$p_{QF} = pQ \cdot pF + pF \cdot pQ = 2 pQ (1 - pQ)$$

The fraction of Q-F contacts is higher than a random distribution for the “ordered” case, and the relationship is opposite for the “clustered” case. Accordingly, the fractions of Q-Q and F-F contacts are lower and higher than a random distribution for ordered and clustered cases, respectively (Heilbronner and Barrett, 2014). In other words, ordered/clustered distribution implies a higher/smaller mixing degree of phases and a larger/smaller fraction of interphase boundaries than random distribution. In addition, the SPO analysis was performed using the PAROR method (Panozzo, 1983), and the preferred orientation of each grain shape distribution was plotted in a rose diagram. The aspect ratio b/a (where a and b are the long and short axis of best-fit ellipses approximating the grain shape) was also determined to document the bulk grain shape anisotropy.

3.3. Bulk-rock compositions

Bulk-rock chemical analyzes of the samples investigated and used in pseudosection calculations are given in Table 1. Analyzes were carried out in the *Centre de Recherches Pétrographiques et Géochimiques* (CRPG-Nancy) by the *Service d'Analyse des Roches et des Minéraux* (SARM). Phase relationships were determined using the *Perple_X* thermodynamic software (Connolly, 2005), version 6.7.9 with the thermodynamic database of Holland and Powell (2004). These relationships were modeled in the NCKFMASH system (Na₂O-CaO-K₂O-FeO-MgO-Al₂O₃-SiO₂-H₂O) with water in excess and pure albite phase excluded (only solid solution of plagioclase).

4. Results

4.1. Microstructural evolution - qualitative approach

4.1.1. Strain facies across the strain gradient

The Questembert granite is traditionally described as a large-scale synkinematic intrusion (e.g. Berthé et al., 1979; Tartèse and Boulvais, 2010). However, microscopic-scale observations showed no evidence for the presence of melts during deformation even in low-strain domains (Bouchez et al., 1992). It is mainly composed of quartz (qtz), K-feldspar (kf), plagioclase (pl) and white mica (wm) with minor amounts of biotite (bt), apatite (ap), chlorite and kaolinite. Solid-state deformation is characterized by a strain gradient from a quasi-undeformed protolith to a km-scale ultramylonite where grain size is strongly reduced (Fig. 1c). This aspect is considered in terms of strain gradients based on microstructural observations, including grain size and evolution of porphyroclast/matrix proportions as major parameters. We thus use changes in grain size and proportion of porphyroclast/matrix with respect to the protolith as proxy to define several domains of strain intensity, such as described in the following sections. In order to avoid ambiguity when considering quartz aggregates and their possible polymineralic character, the term “quartz-rich aggregates/layers” will be used in this text when referring to the domains of quartz with a volumetrically minor amount of second phases. In addition, the term “nucleation” will also be used to denote the processes involving changes in chemical composition, unlike dynamic recrystallization.

4.1.1.1. Low-strain protolith

The protolith contains abundant porphyroclasts of K-feldspar, plagioclase, mica and quartz, without discernible shape-preferred orientation. Quartz grain boundaries have variable geometries,

from slightly curved to serrated (Fig. 2a). These quartz grains show weak deformation features such as sweeping or patchy undulatory extinction (Fig. 2b). In this low-strain protolith, both biotite and white micas are present in a comparable amount. These micas are either undeformed or slightly kinked and bent. Quartz-rich aggregates are associated with cracks along grain boundaries filled with biotite and white mica that form essentially at the edges of porphyroclasts (Fig. 2c). Small grains of white mica are also distributed in plagioclase as rod-like cross-sections (Fig. 2d). Plagioclase grains form large porphyroclasts locally twinned, rarely bent, and affected by fractures (Fig. 2b). K-feldspar porphyroclasts are also frequently fractured. Some of the fractures that crosscut feldspar porphyroclasts are filled by quartz, plagioclase, white mica and biotite (Fig. 2b). K-feldspar porphyroclasts are strongly perthitic, with plagioclase exsolution lamellae (Fig. 2e). Commonly, plagioclase nucleates in many different places within and at the rim of K-feldspar grains (Fig. 2d-e). Both types of feldspar are almost undeformed, such as shown by smooth undulatory extinction and limited chemically-driven nucleation at grain boundaries.

4.1.1.2. Medium-strain protomylonite

The protomylonite samples are microstructurally and geographically close (Fig. 1b) to the one collected by Bukovská et al. (2016). Three main different types of mica microstructures are identified in the protomylonite: (1) small aligned white mica grains form disconnected trails along the grain boundaries in quartz-rich layers (Fig. 3a-b), (2) some thin and short monophasic layers of mica form an anastomosing network connected by trails of very fine-grained recrystallized grains inherited from mica porphyroclasts (Fig. 3c-d), and (3) a few fine-grained polyphase layers mainly composed of mica and quartz (Fig. 3e). These mica-quartz-rich polyphase layers are subparallel to the foliation, cross-cut quartz-rich layers and wrap around feldspar porphyroclasts (Fig. 3e). The polyphase layers increase the proportion of matrix compared with the protolith. With respect to the overall size of recrystallized quartz grains, the quartz grain size is considerably smaller in these high-strain narrow zones, where elongated white micas are present in large amounts (Fig. 3e). Very few chlorite grains in the layers are also observed.

4.1.1.3. High-strain mylonite

The mylonite is composed of a network of polyphase layers wrapping up isolated relic domains in which inherited large porphyroclasts are still visible (i.e., microlithons; Fig. 4a). Some feldspar porphyroclasts are fragments resulting from fracturing of initially larger grains. While the feldspar porphyroclasts are smaller in size and in volume fraction than those in the protomylonite, biotite and monophase layers of mica are no longer present. With respect to the protomylonite, the amount of mica-quartz-rich polyphase layers increases (Fig. 4a-b-c). Within these polyphase layers, white mica grains are strongly aligned and become interconnected (Fig. 4b-c). A large amount of apatite is also documented in these deformed zones (Fig. 4b). The pervasively recrystallized quartz grains exhibit a shape-preferred orientation nearly parallel to the mylonitic foliation (Fig. 4c). The clasts of plagioclase and K-feldspar locally occur in these zones in small quantities (Fig. 4b-c).

The mylonite is also distinguished from the protomylonite by the incipient development of other fine-grained polyphase layers enriched in quartz and K-feldspar (Fig. 4d-e). While micas are scarce in these layers, isolated grains of K-feldspar exist at triple junctions of quartz grains (Fig. 4f). Within these polyphase layers, plagioclase grains occur as clasts. Fracturing is dominant in plagioclase although a marginal nucleation of plagioclase promotes the formation of fine grains at the boundaries of the old grains. In some places, the remaining K-feldspar porphyroclasts are completely rimmed and replaced by plagioclase (Fig. 4g).

These types of fine-grained polyphase layers isolate domains where the grain size is much larger and where less deformed porphyroclasts can still be recognized. Outside of the fine-grained polyphase layers, the quartz-rich layers are highly elongated subparallel to the foliation and their grains reveal patchy or sweeping undulose extinction patterns and the presence of subgrains (Fig. 4h). Locally, new small grains of quartz recrystallize within older, larger grains of quartz (Fig. 4i).

4.1.1.4. Very high-strain ultramylonite

The ultramylonite is largely composed of a succession of foliation-forming parallel layers of fine grains that embed few small clasts, forming a clast-in-matrix microstructure. These latter are plagioclase grains (Fig. 5a-b), and more rarely K-feldspar and white mica grains (Fig. 5a-c). Tails around K-feldspar clasts are composed of fine-grained quartz and K-feldspar (Fig. 5c). The fine-grained parallel layers are either composed of quartz-rich layers, or of polyphase layers of quartz, K-feldspar and white mica (Fig. 5d). The vast majority of the ultramylonite consist of these polyphase layers, while quartz-rich layers occur sparsely in the form of elongated domains parallel to the foliation.

4.1.2. Microstructural evolution across the strain gradient

The prominent microstructural features of the investigated strain gradient are the grain size reduction, the formation of monophase layers of mica and polyphase layers. With increasing strain, coarse-grained mineral phases like feldspar, quartz and mica are replaced by fine-grained layers (Fig. 4-5-6). The grain size reduction of coarse-grained feldspar occurs predominantly by fracturing and nucleation of new grains with a different mineralogical composition than the protolith (Fig. 6). New grains are mostly distributed along the margins of old magmatic feldspar porphyroclasts, or within the tails of porphyroclasts (Fig. 6). The pink K-feldspar that composes the fine-grained mixed zones in the most strongly deformed domains is the same as the one that composes the rims of the coarse-grained light blue K-feldspar present in the protolith and the protomylonite (Fig. 6). Similarly, the plagioclase grains present in the ultramylonite are also observed as reaction products of the coarse-grained plagioclase of the protolith. These CL properties of K-feldspar and plagioclase support the connection of the microstructures throughout the strain gradient: high-strain domains (ultramylonite and mylonite)

are therefore interpreted as the evolution, with increasing strain, of low strain domains (protolith and protomylonites).

4.1.3. Layers of mixed phases

The transition from protolith to ultramylonite is characterized by the development of fine-grained polyphase layers, i.e. fine-grained mixed-phase zones (Fig. 7), which include three different types depending on their phase content (Fig. 7a-c-e). The mixed-phase zones also contain plagioclase, but only locally and in very small amounts (hence not shown in images of Fig. 7). The first type (type I) involves the presence of white mica and quartz in equal proportion with minor K-feldspar (Fig. 7a-b). While scarce in the protomylonite, this mixed-phase assemblage is abundant in the anastomosing network of the mylonite polyphase layers. The two other mixed-phase zones (types II and III) are composed of quartz + feldspar and contain smaller amounts of white mica. They are very similar and simply differ by the nature of the major phase, either represented by K-feldspar (Fig. 7c-d) or quartz (Fig. 7e-f). Types II and III mixed-phase zones are observed in high-strained zones of the mylonites (domain 4), and most of the polyphase layers in ultramylonite. The micas form a partially interconnected network in type I mixtures (Fig. 7b), but they form isolated grains in types II and III (Fig. 7c-f). All fine-grained mixed-phase zones may also contain few very small clasts of mica, plagioclase, or K-feldspar.

In all of these zones, the grains of white mica show a SPO with a mean long axis subparallel to the foliation and an aspect ratio of ~ 0.32 (Fig. 7b-d-f). The quartz grains are preferentially elongated parallel to the foliation in type I mixture (aspect ratio of ~ 0.46 ; Fig. 7b), but they have a much more equant shape in type II and III mixtures (aspect ratio of ~ 0.72 ; Fig. 7d-f). The small grains of K-feldspar only present in type II and III mixtures also have more equant shape with aspect ratios of ~ 0.72 and ~ 0.60 , respectively (Fig. 7d-6f).

4.2. Microstructural evolution - quantitative approach

The micro-textural study detailed below is based on digitized maps of each strain domain (Fig. 8), particularly in quartz-rich areas. Figure 8 shows the main microstructural evolution with increasing strain, including the development of quartz-rich layers and monophasic layers of mica in low-strain domains (Fig. 8b) and mixed-phase zones in higher strain domains (Fig. 8c-d-e).

4.2.1. Grain size

Overall, there is a significant decrease in quartz grain size (equivalent diameter) with increasing strain (Fig. 9a). While the median of quartz grain size is 170 μm in domain 1 (protolith) and 120 μm in domain 2 (protomylonite), it decreases in domains 3 and 4 (mylonites) with respective median sizes of 114 μm and 38-29 μm . A gradual narrowing of the grain size distribution is observed with a wide range of size in the weakly deformed domains (1 and 2) and a rather narrow distribution for the mylonitic domains (3 and 4). In mylonites, these grain sizes correspond to the quartz-rich layers in low-strain domains where no phase mixing occurs. In contrast, where mixed-phase zones are abundant in high-strain domains, the median equivalent diameter of quartz grains is strongly reduced to 5.4 in the mylonite and 2.3 μm in ultramylonite (Fig. 9b). In these latter, a bimodal distribution of quartz grain size is observed (Fig. 9b). This is caused by the variable grain size distribution between polyphase mixed zones and preserved monophasic aggregates in these high-strain domains. It is also worth to note that the recrystallized grain size of quartz in mixed-phase zones is similar and stable in mylonite and ultramylonite. The proportion of mixed-phase zones is more important in ultramylonite than in mylonite.

4.2.3. Grain Contact Frequencies (GCF)

The GCF between Q-Q, F-F and Q-F are shown in figure 9 as a function of the amount of boundary surface by these phases (Heilbronner and Barrett, 2014). Quartz grains are distributed in a clustered fashion for all domains (Fig. 10a). The GCF of quartz grains in the strongly deformed samples (strain domains 3 to 5) are closer to a random distribution than the ones in the weakly deformed samples (strain domains 1-2). With increasing strain, this implies an increase of the degree

of mixing of quartz grains with other phases. Accordingly, the feldspar grains are organized in the ordered field for all domains (Fig. 10b), but once again, there is an evolution with strain. In the weakly deformed sample (strain domain 1), the GCF between feldspar grains are closer to the random distribution curve. Thus, with increasing strain (strain domains 2 to 5), the GCF between feldspar grains move away from the random distribution curve, suggesting an increase in the mixing degree of feldspar grains with the other phases. Regarding the GCF between quartz and feldspar grains, the weakly deformed samples (strain domains 1-2) are localized in the clustered domain, while the more strongly deformed samples (strain domains 3 to 5) are located in the ordered domain, notifying again an increase in the mixing degree of quartz and feldspar grains with increasing strain.

4.2.4. Quartz Lattice Preferred Orientation (LPO)

Using equal-area lower-hemisphere pole figures, we show in Fig. 11 the quartz LPO of mylonitic (Fig. 11a-b) and ultramylonitic layers (Fig. 11c-d). In the mylonite, the quartz-rich layer exhibits a moderate LPO (Fig. 11a). The pole figure of quartz c-axes [0001] shows an incomplete crossed girdle (Fig. 11a). In the ultramylonite, a moderate quartz c-axes [0001] single girdle is developed at a large angle to the foliation and follows the sense of shear (Fig. 11c). The a-axes [11-20] of these quartz-rich layers are scattered and show locally a minor maximum close to the lineation. All of these distributions are strongly dispersed first in fine-grained quartz located at the rim of the layers (Fig. 11a-c), and in fine-grained polyphase mixed zones of both the mylonite and ultramylonite (Fig. 11b-d). The decrease of fabric strength is outlined by the evolution of the J-index and M-index from quartz-rich layers ($J = 3.59$ and 2.51 ; $M = 0.13$ and 0.11) to polyphase mixed zones ($J = 1.63$ and 1.56 ; $M = 0.05$ and 0.02) of the mylonite and ultramylonite, respectively (Fig. 11).

5. Mineral chemistry

The bulk-rock and mineral chemical compositions are summarized in Table 1 and 2. and in the isocon diagram of Fig. 12 (Grant, 1986). The average concentration of an element in the protomylonite, mylonite and ultramylonite (C^a) has been compared to the average concentration of this element in the protolith (C^0). All major element concentrations plot close to the line of constant concentration ($C^a = C^0$), or close to the lines of 10% gain ($C^a = 1.1C^0$) and 10% loss ($C^a = 0.9C^0$). Thus, the comparison between the strain domains studied indicates the lack of major chemical variations during progressive mylonitization. Nevertheless, a slight enrichment in K_2O - P_2O_5 - MgO and depletion in Na_2O in the mylonite/ultramylonite compared to the protolith is reported (Fig. 12).

The chemical composition of white mica shows that the recrystallized grains are richer in celadonite component than the large magmatic white micas (Fig. 13a). Thus, the large magmatic white mica grains correspond to muscovite, whereas the fine recrystallized grains of white mica (especially abundant in mixed-phase zones) are phengites. The feldspar grains have a narrower compositional range (Fig. 13b-c), but within this narrow range we can observe a tendency of the recrystallized grains to be slightly richer in An-component for plagioclase (Fig. 13b). The K-feldspar in mixed-phase zones is richer in Or-component (Fig. 13c). The plagioclase grains also appear to have zoning patterns with a blue core and a brown rim in terms of cathodoluminescence colors (Fig. 5c), but no chemical variation is identified with the EPMA (Table 2).

To better characterize the P-T conditions based on mineral compositions, we further show two pseudosections in Fig. 14. Over a large range of pressures, temperatures and parageneses, K-feldspar, quartz, white mica, and plagioclase are all stable, i.e., the assemblage observed in high-strain domains. Nevertheless, in all of the predicted parageneses, additional accessory minerals are present in particular to incorporate Fe and Mg. Above 320-350°C, biotite is stable, whereas below this temperature range, chlorite is the stable phase. The predicted amount of these accessory minerals is very low (below 1% for chlorite and 2% for biotite; Fig. 14a), reflecting the very low concentration in Fe and Mg in the bulk-rock. The parageneses observed do not include any of the accessory minerals predicted by the pseudosection (clinozoisite, chlorite and biotite). On the other hand, we observed the destabilization of biotite in the evolution from the protolith to mylonite/ultramylonite, suggesting that

deformation occurred at temperature lower than the biotite-out reaction. The plagioclase in fine-grained polyphase layers shows compositional range between 3 and 8% of anorthite component. The molar isopleths in white mica (phengite) show that Si content of phengite is dependent on both, pressure and temperature and evolves from 3.06 per formula unit (p.f.u) in the high temperature/low-pressure side to 3.26 (p.f.u) in the low temperature/high-pressure side (Fig 13b). In addition, the X_{Mg} ratio ranges between 0.36 and 0.43 (Fig. 14b). Phengite compositions do not show considerable variation in Si content between the strain domains. Nevertheless, there is an important range of composition (Si p.f.u) from 3.16 to 3.23 and X_{Mg} molar ratio from 0.37 to 0.41 (Fig. 14b).

6. Discussion

6.1. P-T conditions of deformation

Across the South Armorican Shear Zone, previous petrological studies have shown that deformation occurred under greenschist to lower amphibolite facies metamorphic conditions, and strain localization took place below the low temperature biotite isograd (300-350°C; e.g. Gapais and White, 1982; Bukovská et al., 2016). Our analyzes show a constant composition of the phengite in mixed zones across the strain gradient (Fig. 13a), which suggest constant P-T conditions during deformation, as the rock composition is also relatively constant.

Given the rock chemical composition, the P-T conditions of deformation are also constrained by the Fe/Mg-bearing phases (Fig. 14). Nevertheless, their predicted abundance is very small (clinozoisite/chlorite: below 1%, biotite: below 2%) (Fig. 14a). Based on their observation of chlorite in layers, Bukovská et al. (2016) defined the stability field for deformation as associated with the parageneses of chlorite, plagioclase, white mica, clinozoisite, microcline and quartz. The absence of clinozoisite in their microstructural study is explained by the substitution of the latter by another Ca-bearing phase such as apatite (Bukovská et al., 2016). Considering our microstructural observations, this is consistent with the large amount of apatite documented in the deformed zones (e.g. Fig. 4b). In the actual parageneses of our deformed samples, biotite is absent, while large biotite grains were

present in the magmatic assemblage. On the other hand, chlorite is also in most cases absent in our high-strain samples and all Fe and Mg are completely taken up by the phengite, as can be approximately inferred from its modal abundance and composition (Table 1-2). When chlorite is present, its textural relationship with the mylonitic assemblage is rather unclear, casting some doubt on the stability of chlorite during deformation. There is therefore much uncertainty on the P-T conditions of deformation, even if the destabilization of biotite from the magmatic paragenesis and the intersection of compositional isopleths suggest that deformation occurred at temperatures below the biotite-out reaction curve, i.e., below ~ 320-350°C, similar to the conclusions of Gapais and White (1982) and Bukovská et al. (2016) (Fig. 14).

6.2. Mechanisms of grain size reduction from protolith to ultramylonite

Dynamic recrystallization is a major process for grain size reduction during plastic flow (e.g. Etheridge and Wilkie, 1979; Behrmann, 1985; Fliervoet and White, 1995; Kruse and Stünitz, 1999; De Bresser et al., 2001; Kenkmann and Dresen, 2002; Stipp et al., 2002), provided that the conditions of deformation in terms of grain size, temperature, pressure and differential stress are so that the rock deforms in the dislocation creep regime (Hirth and Tullis, 1992; Passchier and Trouw, 2005). The pervasive occurrence of dynamic recrystallization is demonstrated in quartz microstructures (e.g. Fig. 4h), and results in strong grain size reduction (Fig. 9). The recrystallization microstructures are characterized by plastically elongated quartz-rich layers subparallel to the foliation (Fig. 4h). Their grains reveal patchy or sweeping undulose extinction patterns and the presence of numerous subgrains. More rarely, small bulges and new grains of quartz are developed along grain boundaries of old quartz grains (Fig. 4i). These observations involve subgrain rotation recrystallization (SGR) as the dominant dynamic recrystallization process of quartz, with a minor and local contribution to bulging recrystallization (BLG; e.g. Drury and Urai, 1990).

Grain size reduction can also result from (1) the nucleation of new grains of different composition (e.g. Kruse and Stünitz, 1999; Kenkmann and Dresen, 2002; Kilian et al., 2011; Herwegh

et al., 2011; Platt, 2015; Précigout and Stünitz, 2016), (2) melt-rock reactions (Dijkstra et al., 2002), and/or (3) cataclasis/fracturing (e.g. Hippler and Knipe, 1990; Stünitz et al., 2003; Kruhl et al., 2007; Viegas et al., 2016). Owing to the lack of melt-rock reactions in shear zones formed under greenschist facies conditions, melting cannot be considered for grain size reduction. In these conditions, feldspar is generally expected to be mechanically strong, and grain size reduction is usually associated with nucleation, fracturing and reaction (e.g. Simpson and Wintsch, 1989; Hippertt, 1998; Tsurumi et al., 2003; Ree et al., 2005; Menegon et al., 2006).

The effect of nucleation of new grains to reduce the grain size is twofold. First, in itself, the nucleation of small grains is achieved at the expense of larger grains, which are affected by dissolution (Kilian et al., 2011), e.g. K-feldspar formed at triple junction in quartz aggregates (Fig. 4f). Second, nucleation of phase A within a cluster of grains of phase B result in pinning phase B grain boundaries (Fig. 3a-b; e.g. Kruse and Stünitz, 1999; Herwegh and Jenni, 2001; Herwegh and Berger, 2004; Song and Ree, 2007; Brodhag and Herwegh, 2010; Linckens et al., 2015; Hunter et al., 2016; Cross and Skemer, 2017; Gilgannon et al., 2017). In the early stages of deformation, quartz-rich layers reveal the presence of tiny white mica grains located along quartz grain boundaries (Fig. 3a-b). The small micas formed at grain boundaries inhibit the motion of quartz grain boundaries (Fig. 3a-b; e.g. Herwegh and Jenni, 2001; Linckens et al., 2015; Gilgannon et al., 2017). Such pinning acts against grain growth and maintains a small grain size (e.g. de Ronde et al., 2005; Linckens et al., 2015). This combined effect of nucleation to decrease the grain size is apparent in mylonites and ultramylonites, where the grain size of quartz is significantly larger in domains essentially composed of quartz compared to domains composed of quartz and feldspar (Fig. 9).

Feldspar grain size reduction is mainly induced by a combination of fracturing and nucleation, such as replacement reaction of K-feldspar by plagioclase (Fig. 4g). This combination leads to the breakdown of the large porphyroclasts into smaller fragments (Fig. 6). Fracturing of feldspar and nucleation microstructures were described in natural shear zones deformed under greenschist facies conditions (e.g. Fitz Gerald and Stünitz, 1993; Ree et al., 2005) similar to the ones in this study (Fig. 2b). In addition to its role of reducing the grain size, fracturing also enables fluid access and transport

of elements required for nucleation to reduce the grain size (Fitz Gerald and Stünitz, 1993). Consequently, the dominant grain size controlling processes are dynamic recrystallization (quartz), nucleation of new grains with a different composition (feldspar and mica) and fracturing (feldspar).

6.3. Formation of a weak mica network

Mica is typically viewed as the weakest phase in granitoid rocks (e.g. Tullis and Wenk, 1994). The micas in the protolith are present as undeformed or only slightly bent magmatic phases. Locally, cracks filled with mica crosscut quartz-rich layers and typically occur at the tip of large magmatic micas (Fig. 2c). With increasing strain, the protomylonite exhibits newly formed micas that tend to organize into proto-layers, defined as an alignment of several micas parallel to each other and cutting across quartz-rich layers or precipitate along quartz grain boundaries (Fig. 3a-b). When located at quartz grain boundaries, these micas locally induce pinning of grain boundaries (e.g. Fig. 3a). With further evolution, an anastomosing network of monophase layers composed of mica starts to form (Fig. 3c). The onset of interconnection occurred where trails of very fine-grained phengites connect the large inherited grains of micas (Fig. 3c-d).

The formation of an interconnected weak layer microstructure requires the coalescence of the weak phases. The experiments of Dell'Angelo and Tullis (1996) investigated weak phase interconnection of quartz, while that of Holyoke and Tullis (2006) was more focused on mica interconnection. The latter proposes that stress concentration and brittle deformation in strong phases such as feldspar may induce weaker grains to become interconnected. Although the brittle deformation is mostly localized in feldspar porphyroclasts with a large number of fractures (Fig. 2b), cracks filled with mica are also observed in quartz-rich layers (e.g. Fig. 2c). These cracks occurred essentially at the tips of large mica grains where a phase strength contrast between quartz and mica is present.

According to Holyoke and Tullis (2006), the stress concentration adjacent to weak phases may induce local brittle deformation, and could explain the formation of mica cracks crosscutting the quartz-rich layers. These cracks which are best preserved in low-strain domains, with limited ductile overprint, may enhance fluid access and then nucleation of grains that act as shear band precursors (e.g. Bukovská et al., 2016; Wehrens et al., 2016). Brittle deformation as evidenced by fractures is thus considered here as an initial important mechanism for localization of deformation, although its importance later decreases with a switch in dominant plasticity (e.g. Fousseis and Handy, 2008). The nucleation of micas in cracks and along quartz grain boundaries combined with the initial plasticity of old grains could lead to the interconnection of these weak phases. The low-temperature conditions of the SASZ deformation are in agreement with such an interplay between brittle and plastic deformation active at greenschist conditions (e.g. Passchier, 1984; Fousseis and Handy, 2008; Wehrens et al., 2016).

6.4. Phase mixing with increasing strain

6.4.1. Evidence for nucleation of new grains

The transition from weakly deformed samples to highly deformed ones is accompanied by grain size reduction and a gradual dispersion of quartz grains from aggregates to mixed-phase zones (Fig. 9). In the protolith, the protomylonite, and in some low-strain domains in mylonite, the quartz is arranged in quartz-rich layers (Fig. 3c-4h), while in high-strain domains in mylonite and ultramylonite the quartz is intimately mixed with other phases (Fig. 7c-e). Indeed, there is an increase in mixing concomitant with an increase of strain (Fig. 10). There is no or limited evolution between the mylonite and ultramylonite, neither between the protolith and protomylonite. The sharp increase towards mixing corresponds micro-structurally to the increase in polyphase mixed zones within mylonites (e.g. Fig. 4a-c-d). It is more than likely that between quartz-rich layers with small amounts of white mica and the first documented polyphase layers, i.e. fine-grained mixed zone rich in white mica and quartz, there must be transitions (Fig. 3b-e). The appearance of monophasic layers of white mica may point towards such transitional microstructures (Fig. 3c).

Dynamic recrystallization during plastic deformation, as a result of either SGR or grain boundary bulging, may reduce the grain size, but it is not expected to induce alone phase separation (Kilian et al., 2011). An additional process is required such as e.g. nucleation of new grains in dilatant sites to induce a spatial rearrangement of mineral phases, giving rise to phase mixing. Evidence of nucleation are first observed in the weakly deformed domains where fine-grained white micas occur along quartz grain boundaries (Fig. 3a-b), and in the form of small oriented particles in feldspar (Fig. 2d). Micas also filled fractures in feldspar clasts and quartz aggregates (Fig. 2b). Evidence of phase nucleation also includes the occurrence of isolated K-feldspar grains located at triple junctions in quartz-rich layers and in newly formed fine-grained mixed-phase zones (Fig. 4e-f-g; e.g. Kilian et al. 2011), and the local formation of tails of fine-grained quartz and K-feldspar, which nucleate around K-feldspar clasts in the ultramylonite (Fig. 5c). Thus, quartz aggregates are progressively disintegrated by dispersed feldspar with increasing strain. In addition, the formation of mixed-phase zones within clasts tails may be due to the nucleation of new phases (Fig. 5c; e.g. Kruse and Stünitz, 1999). These microstructures are in agreement with the increasing amount of grain contact between quartz and feldspar grains with increasing strain (Fig. 10). Phase nucleation is also consistent with the contrasted quartz LPO observed between quartz-rich layers and mixed-phase zones (Fig. 11). In these mixed phase zones, the quartz LPO is dispersed with respect to adjacent quartz-rich layers, which could be the result of enhanced nucleation in mixed-phase zones (Fig. 11). A similar process has been already documented in several experimental and natural studies to cause grain size reduction, loss of LPO and phases mixing (e.g. Kilian et al., 2011; Précigout and Stünitz, 2016; Cross and Skemer, 2017).

6.4.2. Driving force for nucleation

The formation of fine-grained polyphase assemblage by nucleation of new grains may be enhanced by chemical reactions (e.g. Stünitz and Tullis, 2001) and/or grain-boundary sliding (GBS; e.g. Linckens et al., 2014; Linckens et al., 2015). The nucleation of new grains with a different composition such as e.g. small and scattered white mica in dilatant sites may attest for intergranular fluid circulation (Fig. 3a-b; Fig. 4f; e.g. Oliot et al., 2014). The fluid circulation allows the transport of

dissolved elements required for nucleation. In the case studied here, chemical variations – even small – are systematically observed between the porphyroclasts and newly formed grains (Fig. 13). Thus, a purely mechanical phase mixing is considered as very unlikely because of these chemical variations, implying instead mass transfer via fluids. Element transport is manifested by the slight enrichment of K_2O - P_2O_5 - MgO and depletion of Na_2O with increasing strain (Fig. 12). All domains of the strain gradient from protolith to ultramylonite contain white mica. White mica porphyroclasts show quite homogeneous muscovite-rich composition, whereas fine-grained white mica is enriched in the Mg-celadonite-component (phengite; Fig. 13). This gradient of reaction appears coeval with the investigated strain gradient and may explain the slight enrichment in MgO in the bulk composition. The enrichment of K_2O and depletion of Na_2O may be a result of the nucleation of new Or-rich, K-feldspar and An-rich plagioclase with increasing strain. However, the variations in composition are too small to be interpreted with any level of confidence. Therefore, we suggest that the scale of mass transport is limited in order to maintain a quite homogeneous bulk composition, but it allows the local transport of elements required for chemically-driven nucleation.

To achieve the progressive breakdown of quartz aggregates with increasing strain by nucleation of new phases, GBS is an additional process that may be invoked. GBS is one important deformation mechanism that commonly combines with other mechanisms to deform any fine-grained mixed-phase zones (e.g. Raj and Ashby, 1971; Ashby and Verrall, 1973; Passchier and Trouw, 2005; Langdon, 2006; Warren and Hirth, 2006; Svahnberg and Piazzolo, 2010; Miranda et al., 2016). When the displacement along grains is not fully accommodated, this may induce the opening of cavities into which the material can diffuse (e.g. Fliervoet et al., 1997; Fuisseis et al., 2009; Kilian et al. 2011; Platt, 2015; Précigout and Stünitz, 2016; Précigout et al., 2017; Gilgannon et al., 2017). Nucleation of new grains in cavities inhibits grain growth and may promote the activity of grain size sensitive creep (e.g. Herwegh et al., 2011; Menegon et al., 2015; Gilgannon et al., 2017). In our case, several observations are consistent with a component of GBS during deformation and following nucleation in cavities, such as (1) the dispersion and weakening of pre-existing moderate LPO (Fig. 11; Jiang et al., 2000; Bestmann and Prior, 2003), (2) the relatively equant grain shape of quartz and K-feldspar (Fig. 7d-f;

Passchier and Trouw, 2005), (3) the increase in mixing degree of quartz and feldspar grains with increasing strain due to e.g. phase nucleation (Fig. 10c; Kruse and Stünitz, 1999), and (4) the homogeneous mixing in the alternating fine-grained polymineralic layers of ultramylonite (Fig. 5a; Fliervoet et al., 1997) From the above, it is concluded that the main driving forces for nucleation are due to the combination of chemical reactions and grain-boundary sliding under the presence of fluids.

6.5. Succession of deformation mechanisms and weakening processes

6.5.1. Evolution from protolith to protomylonite

Increasing strain from protolith to protomylonite resulted in the development of an interconnected network of monophase layers of mica (Fig. 3c). The initial deformation of old magmatic mica grains by kinking, bending and recrystallization along the edges suggests a process of crystal plastic deformation (Holyoke and Tullis, 2006). This process combined with the nucleation of micas in cracks and along quartz grain boundaries may have promoted the interconnection of these weak phases in the protomylonite. The formation of a network of mica-bearing monophase layers have also been described in several granitic rocks deformed under greenschist to amphibolite facies conditions (Berthé et al., 1979; Gapais, 1989; Vernon et al., 2004; Johnson and Brown, 2004; Bukovská et al., 2016; Hunter et al., 2016, Wehrens et al., 2016, 2017) and is a major process for the weakening of the rock and the localization of deformation (e.g. Holyoke and Tullis, 2006; Menegon et al., 2008; Montési, 2013; Gardner et al., 2017). In addition, the reaction progress forming weak material as phengite in high-strain polyphase layers and ultramylonite is also expected to promote the localization of shear deformation, particularly when local grain size reduction is involved (Fig. 4a; e.g. Keller et al., 2004; Fossen and Cavalcante, 2017).

A recent study characterized the initiation and evolution of shear bands in the Questembert granite (Bukovská et al., 2016). According to their study, the formation of a network of mica shear bands is the main microstructural change responsible for weakening. Our observations show that the

formation of a network of mica is only an early stage in the evolution, already achieved in low-strain mylonites (Fig. 8b-c). Following this stage, mylonites and ultramylonites are affected by a large decrease in grain size and increase in mixing zones (Fig. 4-5). These late stages of microstructural evolution and their rheological consequences are addressed hereafter.

6.5.2. Evolution from protomylonite to ultramylonite

6.5.2.1. General features

Increasing strain led to the formation of high-strain mylonite and ultramylonite characterized by the disintegration of most of large feldspar porphyroclasts and coarse-grained quartz-rich layers (Fig. 8e). This evolution results in the development of well-mixed fine-grained zones with a few preserved quartz-rich layers, forming alternating layers in the ultramylonite (Fig. 5-7). This is consistent with studies showing that in quartzo-feldspathic rocks deformed under greenschist facies conditions, mylonites and ultramylonites are composed largely of fine-grained mixed-phase zones (e.g. Stünitz and Fitz Gerald, 1993; Fliervoet et al., 1997). The fine-grained mixed-phase zones are commonly observed in the form of homogeneous mixing of all phases (e.g. Fitz Gerald and Stünitz, 1993; Fliervoet et al., 1997; Tsurumi et al., 2003), or in the form of alternating layers of fine-grained mixed-phase zones (e.g. Ree et al., 2005; Ishii et al., 2007). The geometrical organization of these zones in our study is close to the latter observations. Nevertheless, their compositions differ from published studies (Behrmann and Mainprice, 1987; Fitz Gerald and Stünitz, 1993; Fliervoet et al., 1997; Tsurumi et al., 2003; Ree et al., 2005; Ishii et al., 2007), where mixed zones are principally composed of plagioclase + K-feldspar with less proportion of quartz \pm mica \pm other minerals.

6.5.2.2. Temporal sequence of protolith-protomylonite-mylonite-ultramylonite

According to Augier et al. (2011), the most deformed ultramylonitic domain in the core of the shear zone presents locally less deformed rock portions that correspond to lower-strained domains. Identifying these relics of less deformed rocks suggest that the ultramylonite domains are the evolution, with increasing strain, of low-strain domains. An additional evidence in favour of this evolution with strain is revealed by CL-colors and textures. The pink feldspar grains and the dark blue/purple plagioclase porphyroclasts present in the ultramylonite are already observed as reaction rim products, in the protolith and protomylonite (Fig. 6). Furthermore, the homogeneous bulk rock composition (Table. 1; Fig.12) and the constant composition of the new nucleated phengites throughout the strain gradient (Fig. 13a) suggest that the protolith is identical in all strain facies. Consequently, high-strain domains (ultramylonite and mylonite) are interpreted as the evolution, with increasing strain, of low strain domains (protolith and protomylonites).

6.5.2.3. Preserved quartz-rich layers and fine-grained mixed zones in high-strain mylonite and ultramylonite

The well-developed SPO in the preserved quartz-rich layers in mylonite and ultramylonite, combined with the typical SGR recrystallization microstructures and the dominance of prism- $\langle a \rangle$ and rhomb- $\langle a \rangle$ active slip systems suggest dislocation creep as the dominant deformation mechanism (Fig. 11a-c; Passchier and Trouw, 2005). As indicated by the nearly random LPO of fine quartz grains at the rim of the layers, the quartz-rich layers are locally dismembered and incorporated in the polyphase mixed zones (Fig. 11a-c). Since we consider the ultramylonite and its fine-grained polyphase layers as the most evolved domains of the studied strain facies, the remaining quartz-rich layers in this domain must be considered as relics of an original mylonitic fabric before total disaggregation (e.g. Gilgannon et al., 2017). Although the quartz LPO in quartz-rich layers near the mixed zones is initially moderate, the evolution of the LPO fabric between these two domains suggests that during the disaggregation the quartz LPO dispersed (Fig. 11).

The polyphase mixed zones are characterized by a very small grain size (Fig. 9b), a weaker LPO relative to quartz-rich layers (Fig. 11), a higher degree of phase mixing (Fig. 7-9), and an equant shape of quartz grains (Fig. 7). Based on these observations, we argue that deformation in the fine-grained mixed-phase zones is dominated by grain size-sensitive (GSS) mechanisms, including a contribution of GBS (e.g. Behrmann and Mainprice, 1987; Rutter and Brodie, 1988; Olgaard, 1990; Linckens et al., 2011). GBS contributes to the dispersion of phases, reflected in their anticlustered distribution (Fig. 10c), the homogeneous mixing in the alternating layers (Fig. 5a), and to the weakening of the LPO in mixed-phase zones (Fig. 11; e.g. Behrmann and Mainprice, 1987; Stünitz and Fitz Gerald, 1993; Kilian et al., 2011). Some plagioclase clasts are also preserved and scattered throughout the microstructure in the ultramylonite (Fig. 5a-b). These plagioclases are characterized by a grain shape mostly rounded with convex grain boundaries, which is one evidence for a mechanically strong phase (e.g. Naziri et al., 1975; Stünitz and Fitz Gerald, 1993). Such microstructures are observed in the study of Lonka et al. (1998) and may reflect the progressive rotation of plagioclase clasts as rigid body in a ductile fine-grained zone during diffusion-accommodated GBS. The increase in GSS contribution to overall strain is in turn a weakening process (e.g. Boullier and Gueguen, 1975, Kerrich et al., 1980; De Bresser et al., 2001; Stünitz and Tullis, 2001; Menegon et al., 2015; Précigout and Stünitz, 2016) resulting in further strain localization in the deforming domain.

7. Conclusions

An evolution in microstructures and deformation mechanisms occurred across the different finite strain domains developed across the South Armorican Shear Zone in the Questembert granite. The microstructural, chemical and EBSD analyzes allow us to draw the following conclusions:

1. The transition from protolith to ultramylonite is associated with grain size reduction, which occurred by dynamic recrystallization, nucleation of grains with a different composition and fracturing.

2. The transition from protolith to ultramylonite is also associated with a large increase in phase mixing. Phase mixing is mainly controlled by nucleation, whose driving force involves a combination of chemical reactions and grain-boundary sliding/dilatancy.
3. A reduced grain size may have triggered a transition in deformation mechanism with increasing strain. The quartz-rich layers, which display moderate LPO patterns, are deformed predominantly by dislocation creep, whereas the loss of LPO in polyphase fine-grained mixed zones, where strain has been strongly localized, suggests that diffusion creep and GBS became the dominant deformation mechanism. GBS is also highlighted by the occurrence of isolated K-feldspar grains located at triple junctions, and contributing to the dispersion of phases.
4. The early brittle deformation in low-strain domains, such as cracks in quartz aggregates, enhances fluid access and chemical reactions/nucleation to reduce grain size. A consequence of the formation of cracks and fluid-assisted nucleation is the interconnection of weak mica monophase layers in protomylonite.
5. With increasing strain, the mica-bearing layers evolve first into an interconnected network of fine-grained polyphase layers enriched in quartz and phengite in protomylonite and mylonite, and then by layers of fine-grained polyphase mixing composed of K-feldspar, quartz and phengite homogeneously mixed in mylonite and ultramylonite.
6. The strain weakening of the rock is a two-step process and the nature of the weak material, mica then fine-grained polyphase mixed zones that can deform by diffusion-accommodated grain boundary sliding, changes along with strain.
7. Although the pressure-temperature path of the shear zone is poorly constrained, the constant composition of the mica throughout the strain gradient suggests that the succession in weakening processes is not related to changing P-T conditions but to the accumulation of strain itself.

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Table captions

Table. 1 Bulk rock chemical compositions of the four studied grades of deformation.

Table. 2 Representative chemical composition of K-feldspar, plagioclase and white mica between porphyroclast and grain in mixed zones.

Figure captions

Fig. 1. (a) Schematic map of the main domains forming the Armorican Massif, modified from Gumiaux et al., 2004. The studied area is delineated by the red rectangle. (b) Simplified geological map of the studied area modified from Augier et al., 2011. The location of the samples used is highlighted with colored dots. (c) Pictures of the samples used in this study, showing the different stages of deformation, from weakly deformed sample (domain 1) to the most deformed one (domain 5).

Fig. 2. (a) Quartz grains with polygonal to lobate grain boundaries. (b) Plagioclase porphyroclast with mechanical twinning and fractures filled by plagioclase (white arrows), and quartz aggregates crosscut by cracks filled with biotite (black arrows). (c) Crack filled with biotite crosscutting quartz aggregates (black arrows). (d) Rod-like cross-sections of white mica in the plagioclase. Also note that the

plagioclase grain replaces that of the K-feldspar. (e) Large porphyroclast of K-feldspar showing the presence of oriented plagioclase exsolution. K-feldspar is progressively replaced by plagioclase grains (white arrows). All cross-polarized light micrographs. qtz: quartz, pl: plagioclase, kf: K-feldspar, wm: white mica, bt: biotite.

Fig. 3. (a-to-e) Cross-polarized light pictures showing the microstructures observed in protomylonite. (a) Small rod-like cross-sections of recrystallized white mica located at joints of two quartz grains. Note also the restricted growth of quartz by this white mica. Black arrow indicates the direction of growth. (b) Arrangement of small and oriented crystals of white mica (black arrows) in the quartz-rich layer, which define the foliation. (c) Thin monophasic layers of mica crosscut quartz-rich layers and start to connect. (d) Connection of white micas through trails of very fine-grained white micas (black arrows). (e) Narrow zone of recrystallized grains forming polyphase layer composed principally of quartz and white mica. The latter are elongated preferentially parallel to the recrystallized zone margins. qtz: quartz, pl: plagioclase, kf: K-feldspar, wm: white mica, bt: biotite.

Fig. 4. (a-to-h) Coupled cross-polarized light micrographs, cathodoluminescence (CL) and SEM-BSE images representative of microstructures observed in mylonites. (a) Fine-grained polyphase layers (yellow dotted outline) rich in quartz and white mica, with locally few K-feldspar and plagioclase. (b) Close-up of the mixed-phase zone showing the partial interconnection of white micas intimately mixed with very fine-grained quartz grains. (c) Details of (b) showing the very fine-grained mixing composed at grain scale of quartz and white mica. (d) Polyphase layers rich in K-feldspar and quartz, with small quantities of white micas and plagioclase. (e) Grain size reduction of K-feldspar by nucleation in quartz essentially along grain boundaries and at triple junctions (black arrows). (f) K-feldspar located in triple junction of quartz grains. (g) Irregular rim of plagioclase around a K-feldspar porphyroclast. (h) Quartz-rich layers strongly oriented, which show the development of subgrains (white arrows). (i) Small new quartz grains appearing along grain boundaries and within old grains (black arrows). qtz: quartz, pl: plagioclase, kf: K-feldspar, wm: white mica, ap: apatite.

Fig. 5. (a-to-e) Coupled cathodoluminescence (CL) and SEM-BSE images representative of microstructures documented in ultramylonite. **(a-b)** Ultramylonite textures made of intercalated bands of fine-grained polymineralic layers. Clasts are white mica (black), albite (purple) and apatite (yellow), while the pink, dark brown and black colors are due to K-feldspar, quartz and white mica, respectively. **(c)** The tails of a K-feldspar clast are filled by K-feldspar and quartz. **(d)** Layers of variable composition with quartz-rich layers and fine-grained polyphase layers rich in quartz, K-feldspar and white mica. Bigger plagioclase grains remain strong in form of clasts. **(e)** Close-up of the polyphase layers showing the thin white mica grains elongated subparallel to the lineation. qtz: quartz, pl: plagioclase, kf: K-feldspar, wm: white mica, ap: apatite.

Fig. 6. Cathodoluminescence (CL) images showing the evolution of feldspar microstructures with increasing strain. On CL images, quartz is dark blue-brown, initial K-feldspar is bright blue, new K-feldspar is pink, white mica is black, initial plagioclase is brown and new plagioclase is dark blue/purple. qtz: quartz, pl: plagioclase, kf: K-feldspar, wm: white mica.

Fig. 7. (a-c-e) SEM-BSE images representative of the different type of mixed-phase zones and their respective manually digitized maps **(b-d-f)**. Particle (PAROR) orientation of each phase is presented in a polar plot. **(a)** Mixed-phase zone rich in white mica and quartz, locally associated with K-feldspar and plagioclase. **(c)** Mixed-phase zone rich in K-feldspar, quartz, white mica, and minor plagioclase. **(e)** Mixed-phase zone similar to **(c)** with a higher proportion of white mica and where the quartz is the dominant phase. qtz: quartz, kf: K-feldspar, wm: white mica, ap: apatite.

Fig. 8. (a-to-e) Manually digitized maps of individual grains and phase mixing of each domain of studies are used as a basis for the quantitative microtextural analyzes. Mixed-phase zones develop in the mylonite and ultramylonite stage (domains 3 to 5), as an anastomosing network of fine-grained polyphase layers (in the mylonites) or as foliation-forming layers of fine-grained mixing (in the ultramylonite). Note that no distinction is made between K-feldspar and plagioclase for the first four domains; the yellow color refers to all types of feldspar. qtz: quartz, pl: plagioclase, kf: K-feldspar, wm: white mica.

Fig. 9. Quartz grain size evolution. **(a)** The results are shown in box-and-whisker diagram. Individual boxes were limited by upper and lower quartiles, and within it the median was defined. This grain size evolution is obtained from cross-polarized light images, as the grain size is sufficiently large. **(b)** Evolution of quartz grain size between quartz-rich layers and mixed-phase zones for mylonite and ultramylonite. While mylonitic quartz grain size was determined using SEM-BSE images, we used the EBSD to document the grain size in the ultramylonitic domain.

Fig. 10. (a-to-c) Graphs showing the probability of contact between two phases, Q=quartz and F=feldspar, for varying surface contact percentages of Q and F. The results can be compared to the curves showing the theoretical values for random distribution p_{QQ} **(a)**, p_{FF} **(b)** and p_{QF} **(c)**. **(d)** Examples of maps used for the GCF analysis. The maps of all boundaries, phase Q, phase F and contact surfaces between Q and F are successively represented.

Fig. 11. (a-to-d) Comparison of quartz c-axis [0001] and a-axis [11-20] preferred orientation, for both quartz-rich layers and mixed-phase zones, in mylonite **(a-b)** and ultramylonite **(c-d)**. There is a systematic decrease in fabric strength (J-index and M-index) between the monomineralic quartz-rich layers and the fine-grained mixed-phase zones in both mylonite and ultramylonite. Related microstructures to these pole figures are also presented. The quartz-rich layers in both mylonite and ultramylonite show a preferred orientation of grains **(a-c; black arrows)**. All images: BSE pictures. All pole figures in the upper hemisphere and equal area. qtz: quartz, pl: plagioclase, kf: K-feldspar, phg: phengite.

Fig. 12. Isocon diagram comparing the chemical composition of the protolith (C^0) with those of protomylonite, mylonite and ultramylonite (C^a). The blue line ($C^a=C^0$) represents the constant concentration, while the black lines represent 10% gains and 10% losses of components in C^a compared to C^0 .

Fig. 13. (a-b-c) Chemical composition of white mica, plagioclase and K-feldspar. **(a)** Celadonite-Muscovite-Pyrophyllite ternary diagram shows composition of white mica grains from different stages

of deformation. (b-c) Albite-Orthoclase-Anorthite combined ternary diagram for both plagioclase and K-feldspar composition is also presented for different stages of deformation.

Fig. 14. (a) P-T pseudosection calculated in the CNKFMASH system with the following solution models: Ti-Fe-Mg-Mn-biotite with compound formation (Powell and Holland, 1999), white mica (Coggon and Holland, 2002; Auzanneau et al., 2010), feldspar (Holland and Powell, 2003), and chlorite (Holland et al., 1998). Isopleth plots of the amount of chlorite, clinozoisite and biotite are also superimposed. (b) Representation of Si and X_{Mg} content in white mica isopleth plots and the compositional isopleths of anorthite components in plagioclase. The blue area represents the intersection of the compositional isopleths of Si and X_{Mg} in white mica and anorthite components in plagioclase.

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Table. 1 Bulk rock chemical compositions of the four studied grades of deformation

Sample type	Protolith	Protomylonite	Mylonite	Ultramylonite
Wt% oxide				
SiO ₂	75.57	73.88	72.95	74.37
TiO ₂	0.11	0.08	0.18	0.13
Al ₂ O ₃	14.29	14.94	15.39	14.96
Fe ₂ O ₃	0.81	0.89	0.79	0.78
MnO	0.02	0.03	0.01	0.01
MgO	0.18	0.19	0.20	0.21
CaO	0.53	0.61	0.47	0.53
Na ₂ O	3.37	3.65	2.96	2.69
K ₂ O	4.43	4.40	5.08	5.30
P ₂ O ₅	0.39	0.37	0.48	0.50
Total	99.70	99.04	98.51	99.48
Molar content				
SiO ₂	1.2577	1.2296	1.2141	1.2378
TiO ₂	0.0014	0.0011	0.0023	0.0016
Al ₂ O ₃	0.1402	0.1465	0.1509	0.1467
Fe ₂ O ₃	0.0051	0.0056	0.0049	0.0049
MnO	0.0002	0.0004	0.0001	0.0001
MgO	0.0045	0.0047	0.0050	0.0052
CaO	0.0095	0.0109	0.0084	0.0095
Na ₂ O	0.0544	0.0589	0.0478	0.0434
K ₂ O	0.0470	0.0467	0.0539	0.0563
P ₂ O ₅	0.0014	0.0013	0.0017	0.0018
Total	1.5212	1.5056	1.4891	1.5072

Table. 2 Representative chemical composition of K-feldspar, plagioclase and white mica between porphyroclast and grain in mixed zones

Type	K-feldspar			Plagioclase			White mica	
	Porphyroclast	New grain	Grain in mixed zone	Clas t core	Clast rim	Grain in mixed zone	Porphyroclast	Grain in mixed zone
Wt% oxide								
SiO ₂	66.24	64.86	64.66	67.62	68.76	66.27	45.72	46.63
TiO ₂	0.06	0.00	0.03	0.00	0.00	0.00	0.52	0.35
Al ₂ O ₃	18.12	18.13	17.79	19.74	19.87	19.77	33.42	30.27
FeO	0.00	0.00	0.12	0.08	0.00	0.00	31.37	3.17
MnO	0.00	0.01	0.00	0.03	0.03	0.00	0.00	0.08
MgO	0.02	0.00	0.00	0.00	0.03	0.00	0.81	1.55
CaO	0.00	0.00	0.00	0.96	0.69	1.49	0.00	0.00
Na ₂ O	0.55	0.54	0.43	10.97	11.30	10.66	0.75	0.16
K ₂ O	15.91	16.16	16.22	0.16	0.08	0.18	10.44	11.05
Total	100.90	99.70	99.25	99.56	100.75	98.37	93.03	93.26
Formula								
Si	3.04	3.01	3.01	2.98	2.99	2.96	3.12	3.21
Ti	0.00	0.00	0.00	0.00	0.00	0.00	0.03	0.02
Cr	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Al	0.98	0.99	0.98	1.03	1.08	1.04	2.69	2.46
Fe	0.00	0.00	0.01	0.00	0.00	0.00	0.08	0.25
Mn	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Mg	0.00	0.00	0.00	0.00	0.00	0.00	0.08	0.16
Ca	0.00	0.00	0.00	0.05	0.03	0.07	0.00	0.00
Na	0.05	0.05	0.04	0.94	0.95	0.92	0.10	0.02
K	0.93	0.96	0.96	0.01	0.00	0.01	0.91	0.97
Total	5.00	5.00	5.00	5.00	5.00	5.00	7.00	7.00
XMg molar							0.51	0.39

Note. XMg molar = Mg/(Mg+Fe)

Highlights

- Grain size reduction and phase mixing increases with increasing strain.
- Phase nucleation is the major process behind phase mixing.
- Phase nucleation is driven by chemical reactions and grain-boundary sliding.
- Succession in weakening processes related to the accumulation of strain itself.

ACCEPTED MANUSCRIPT

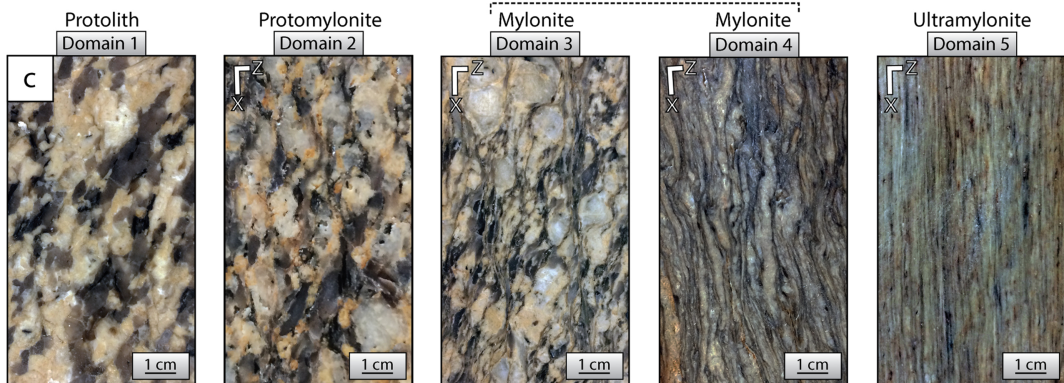
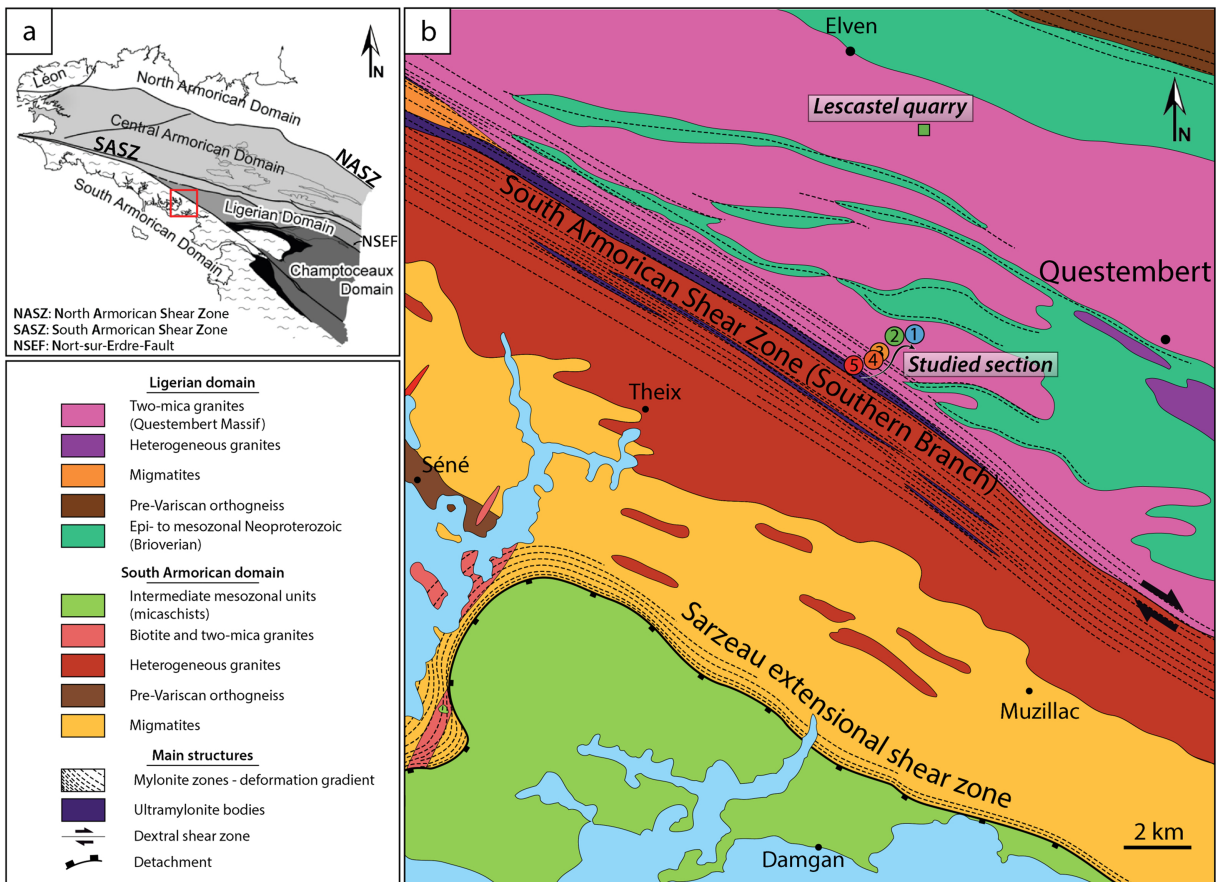


Figure 1

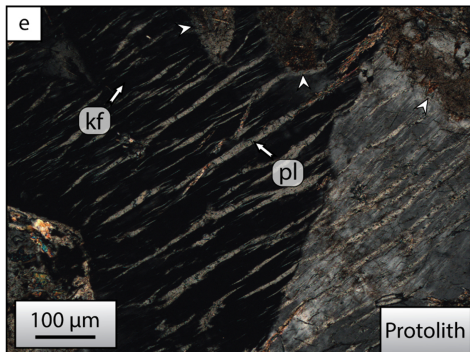
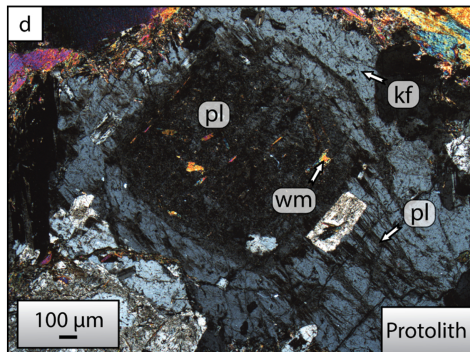
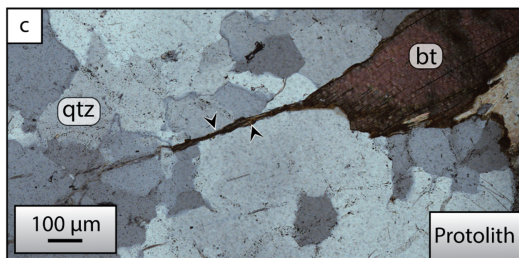
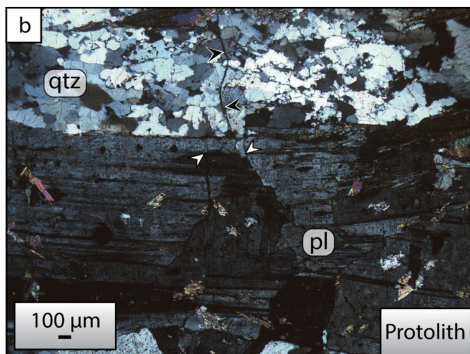
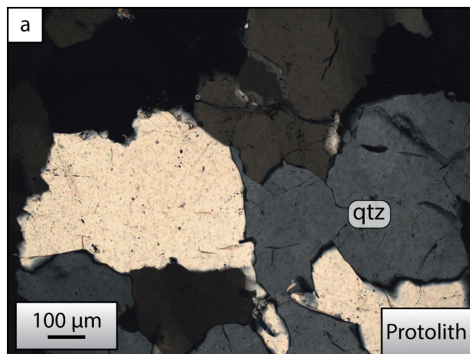


Figure 2

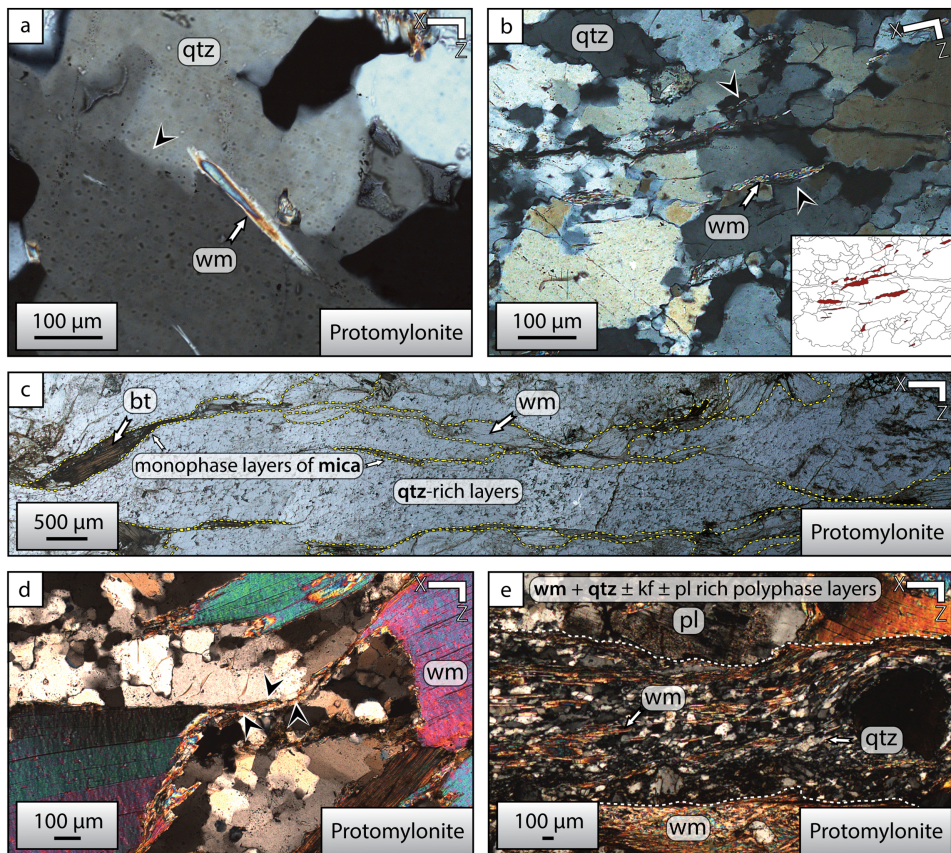


Figure 3

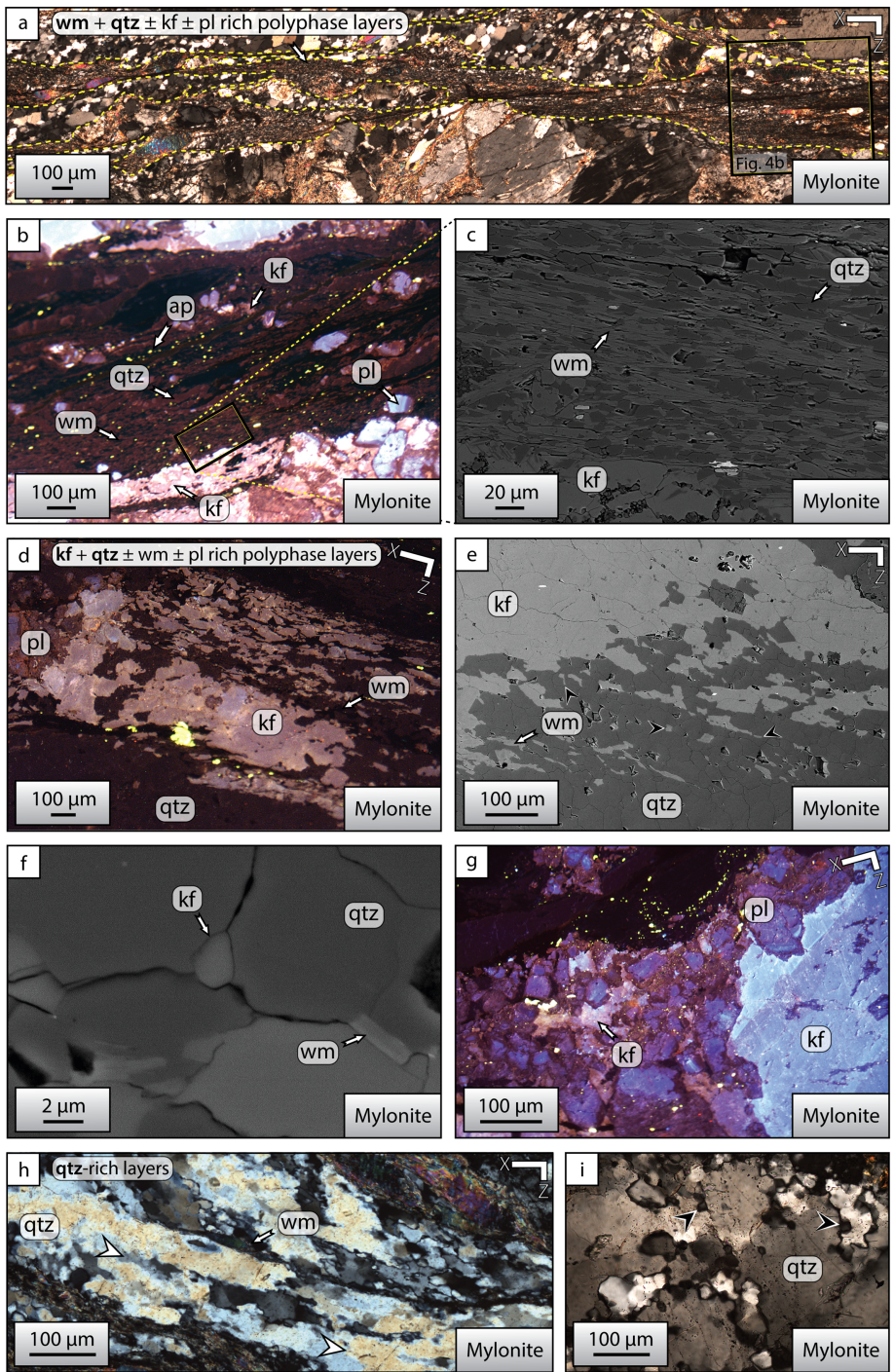


Figure 4

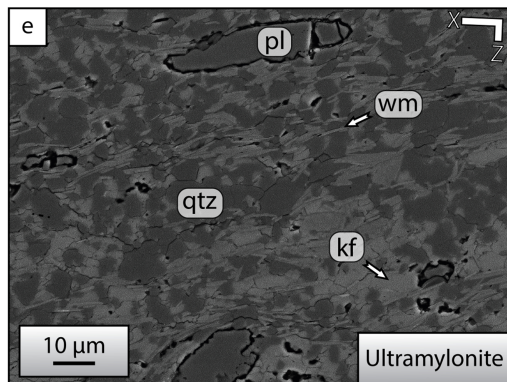
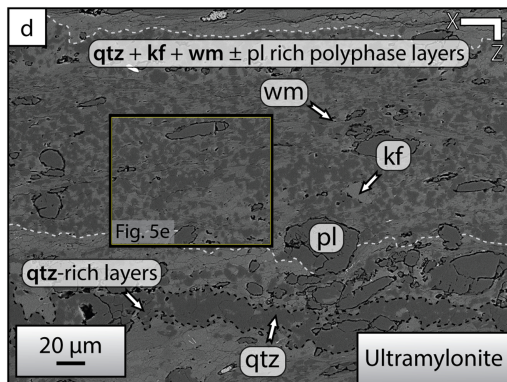
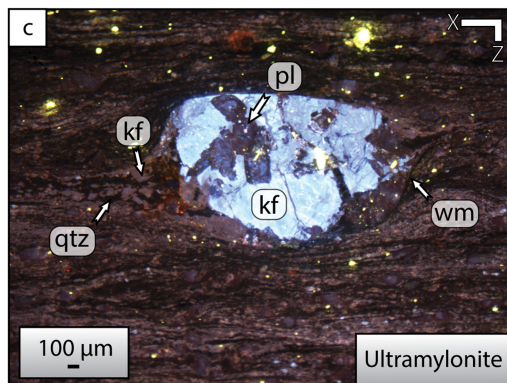
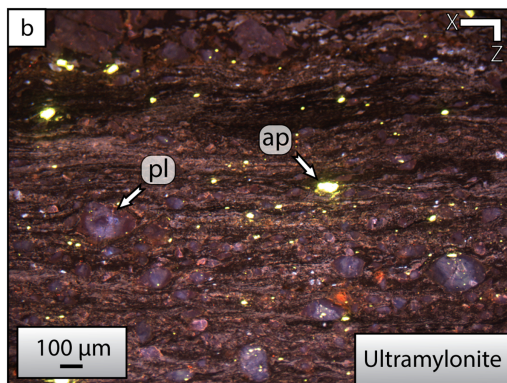
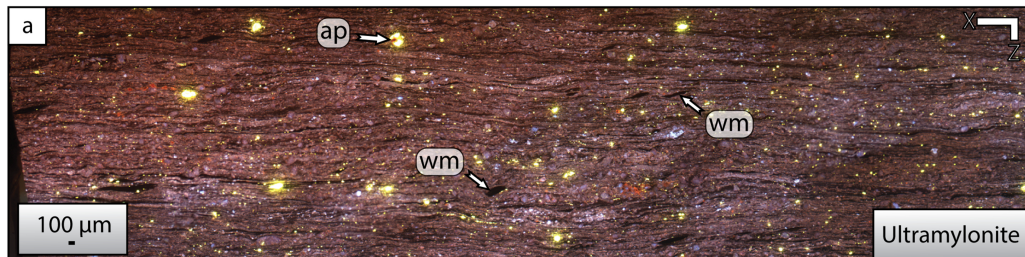


Figure 5

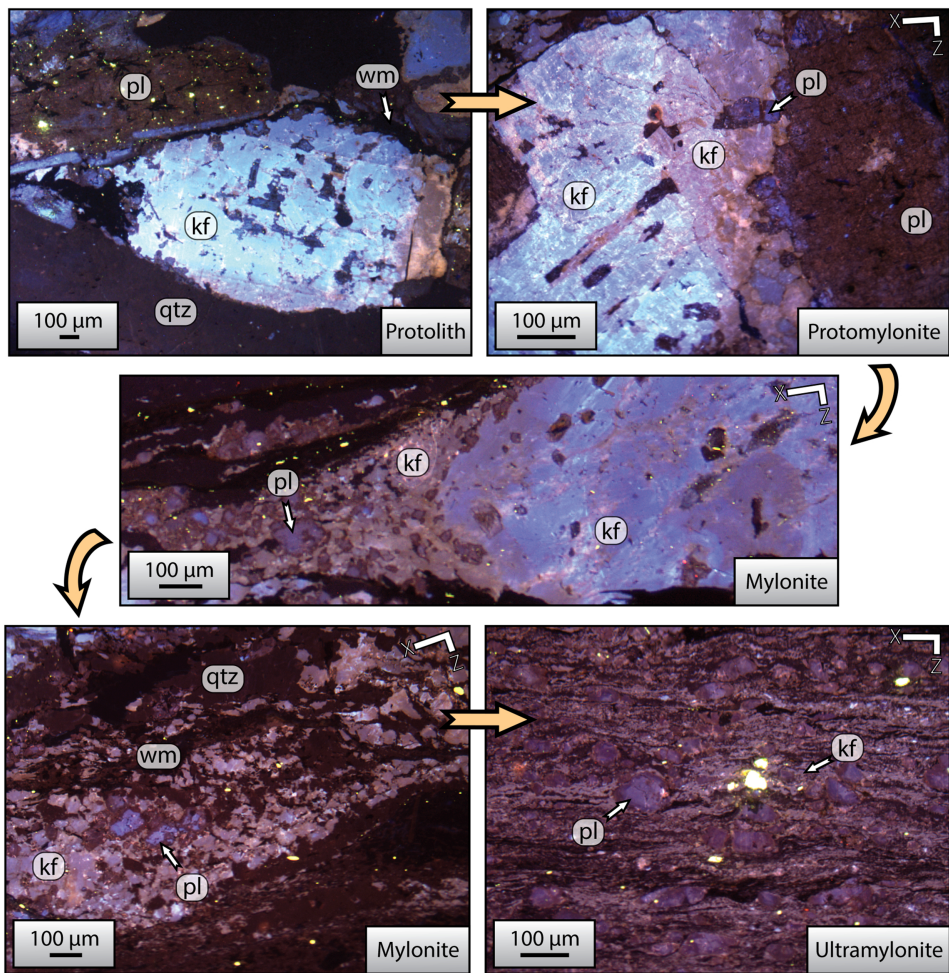


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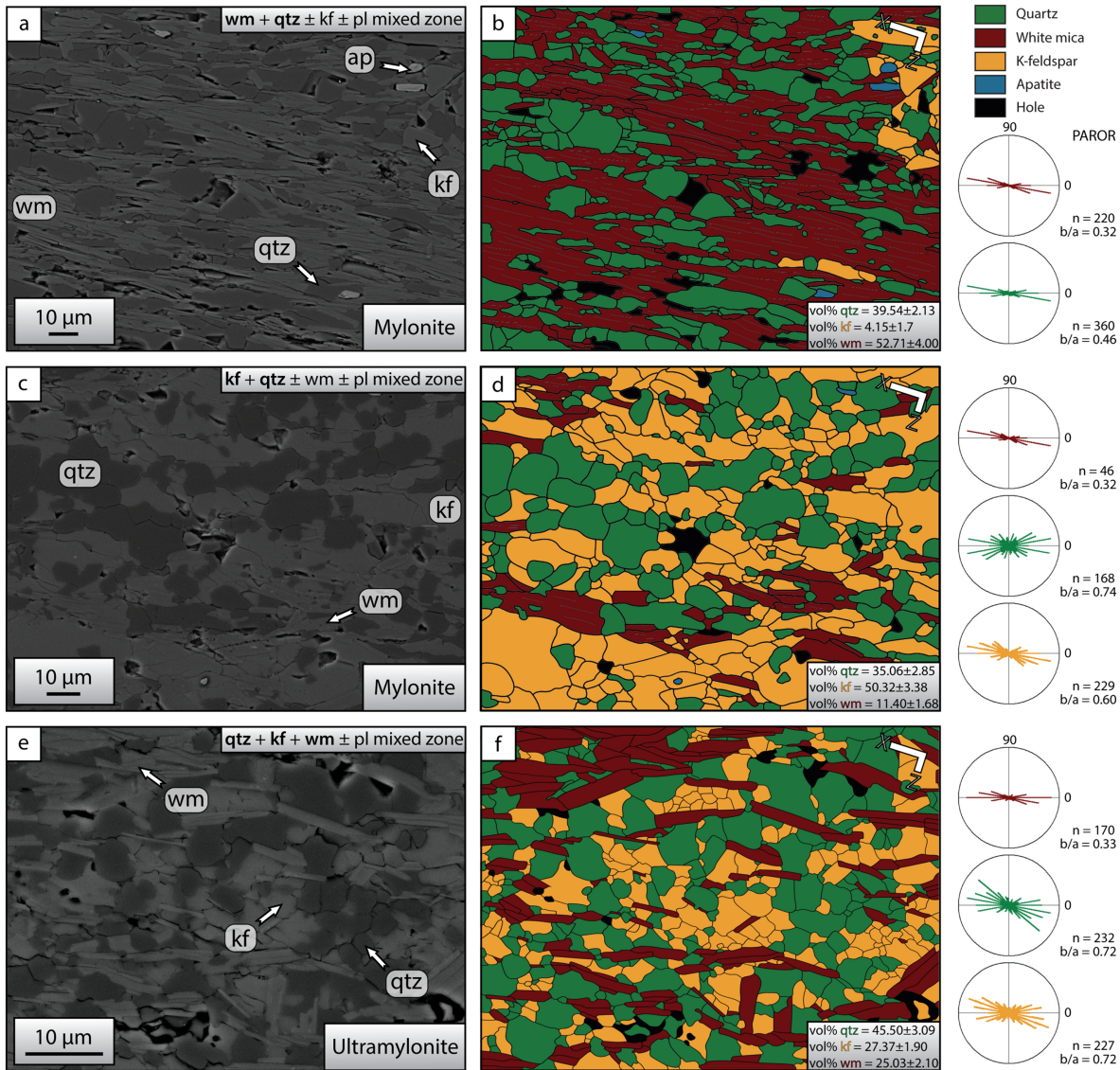


Figure 7

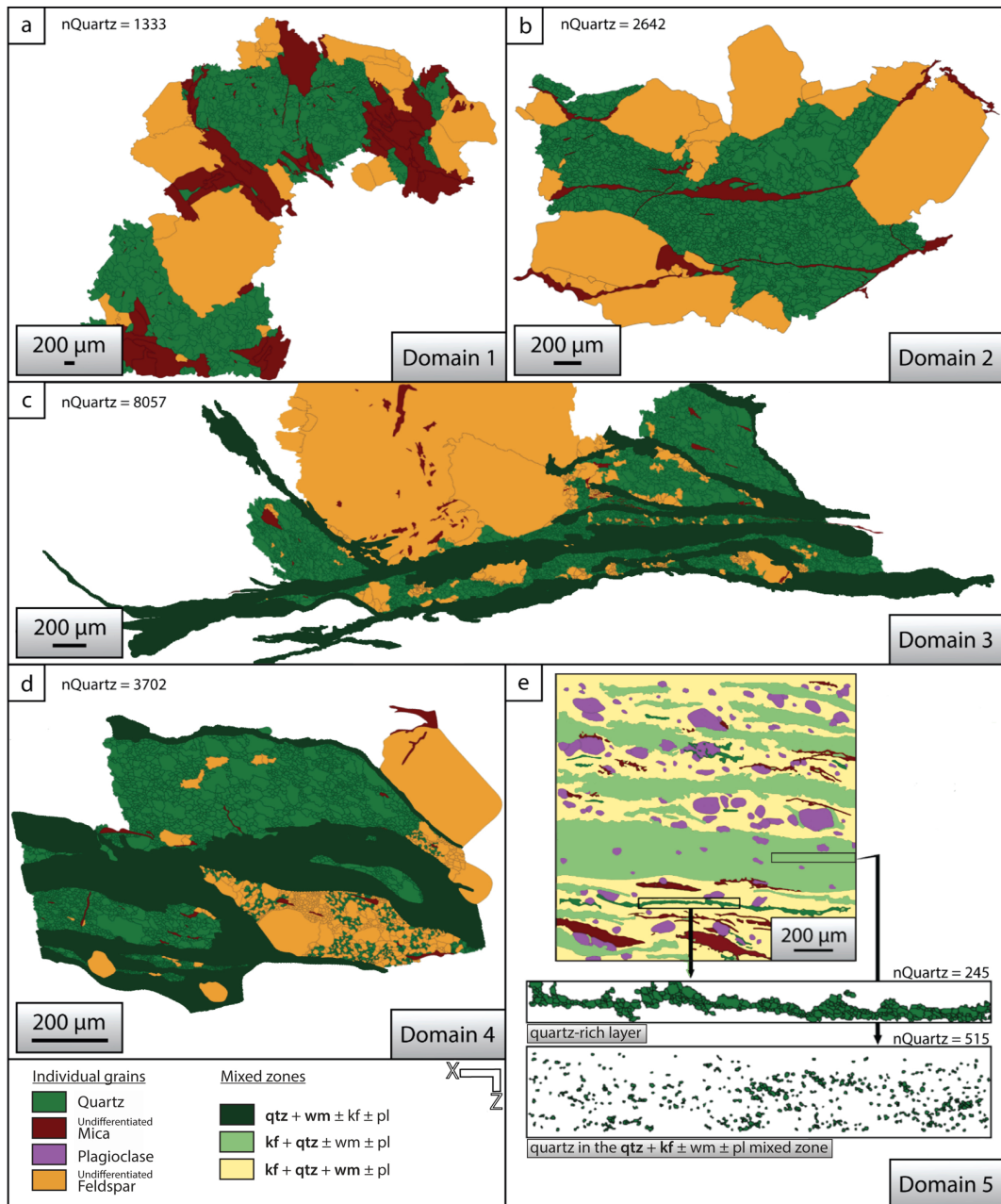


Figure 8

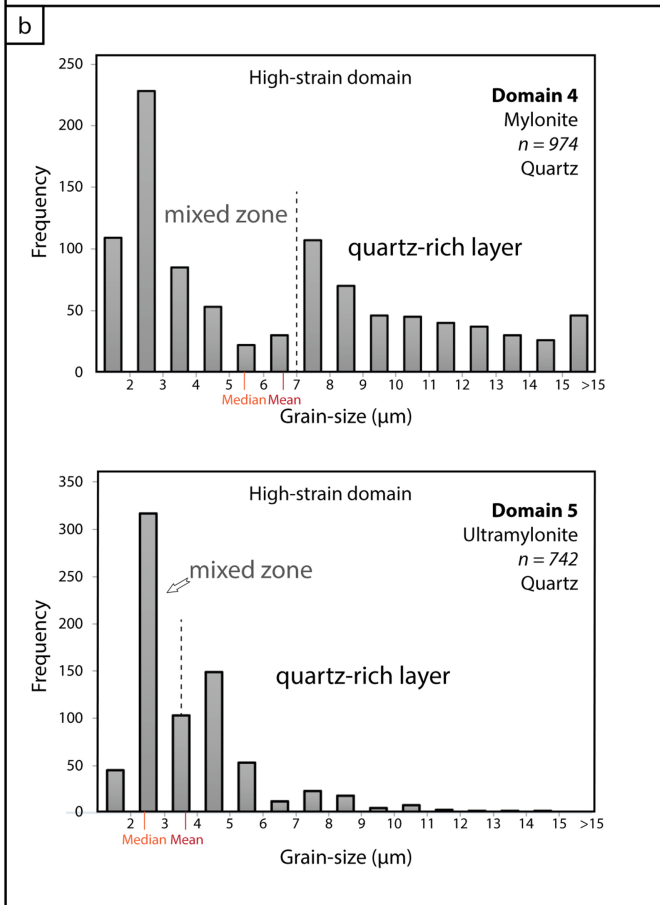
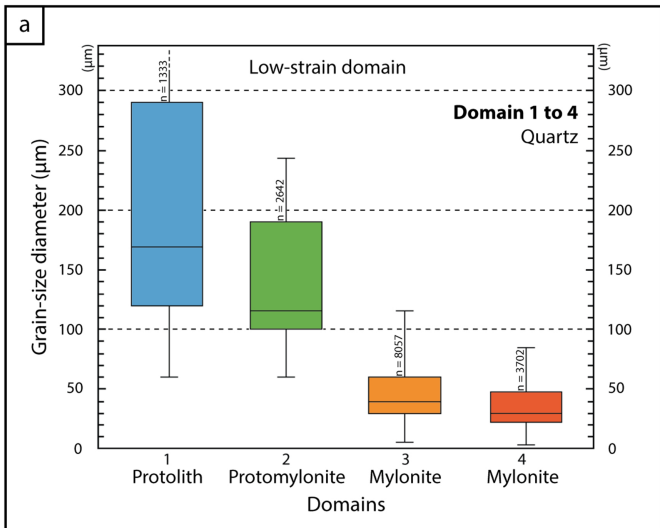


Figure 9

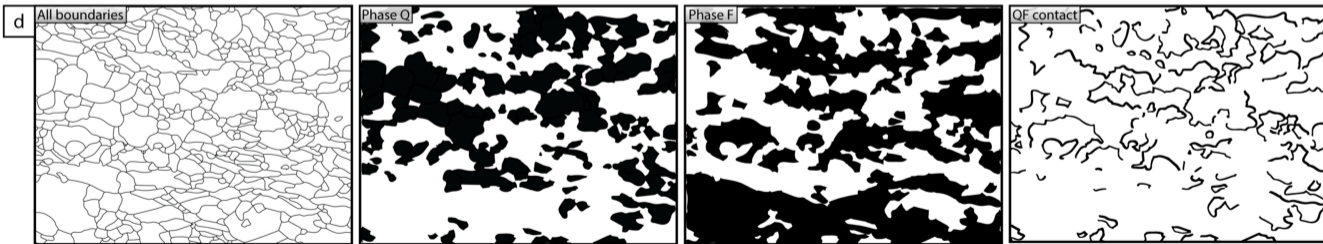
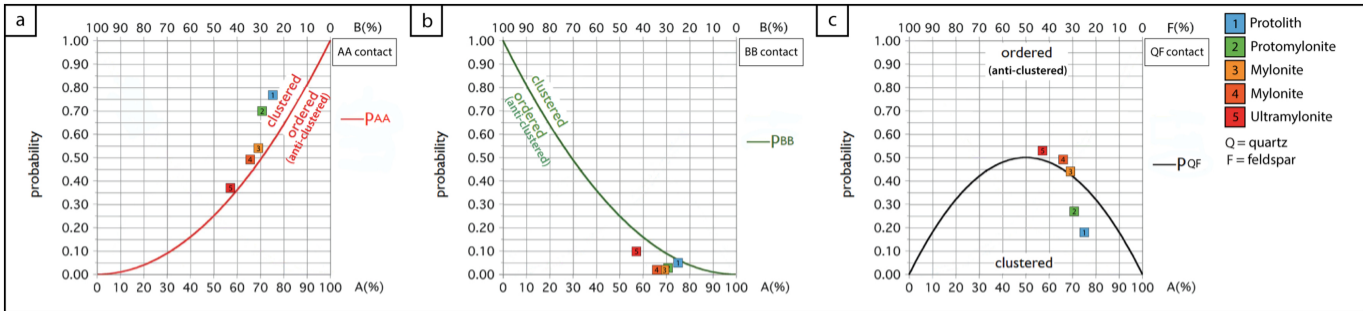


Figure 10

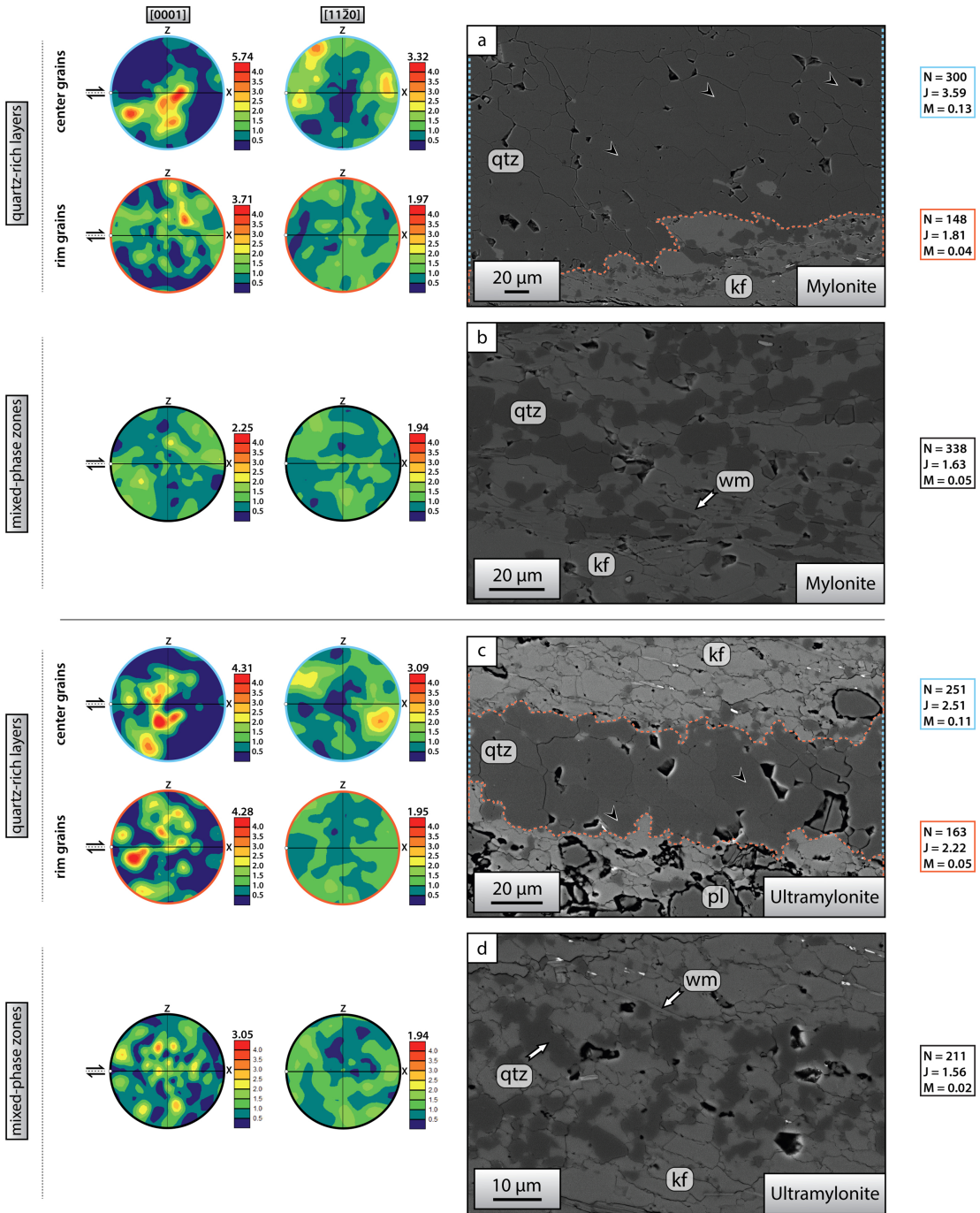


Figure 11

Ca - protomylonite - mylonite - ultramylonite

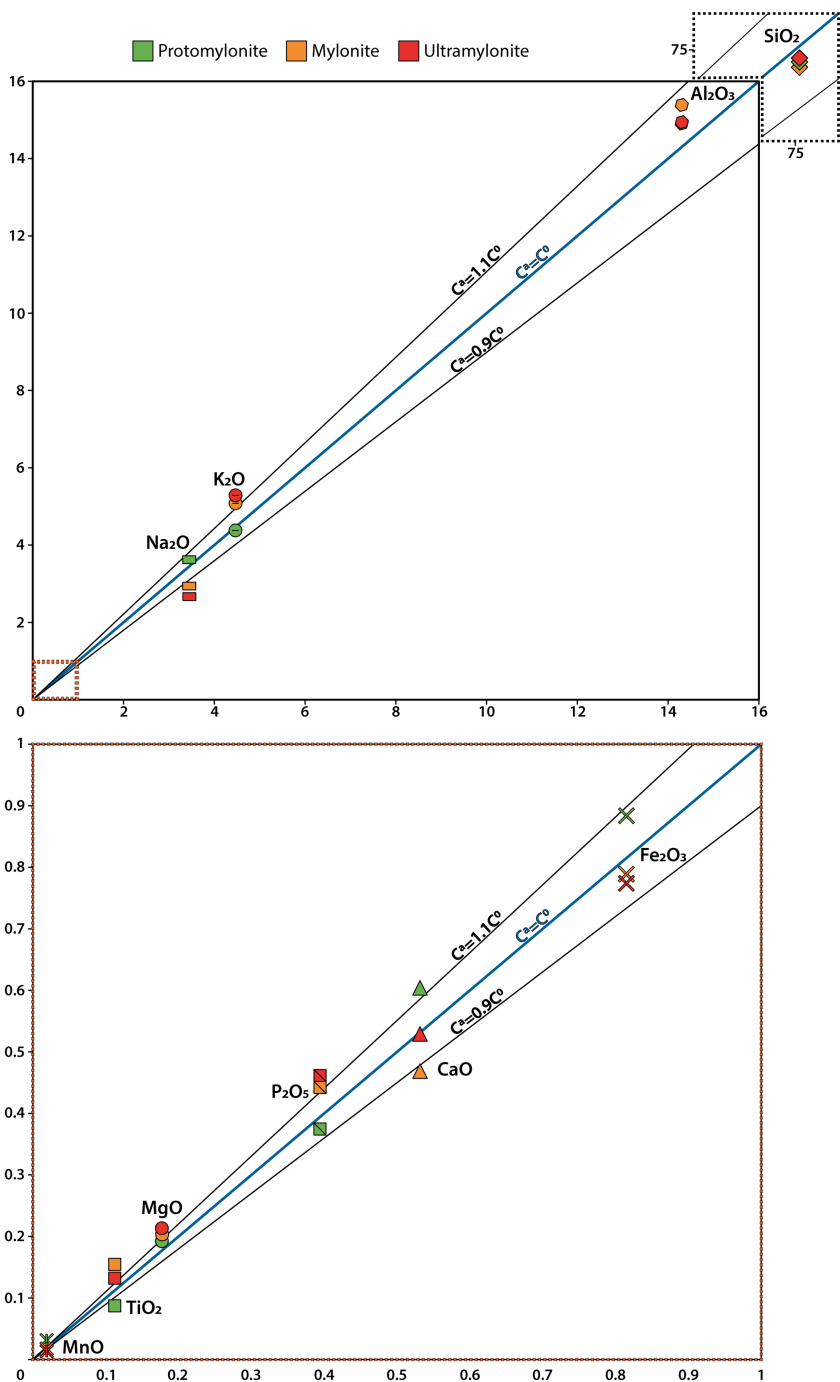


Figure 12

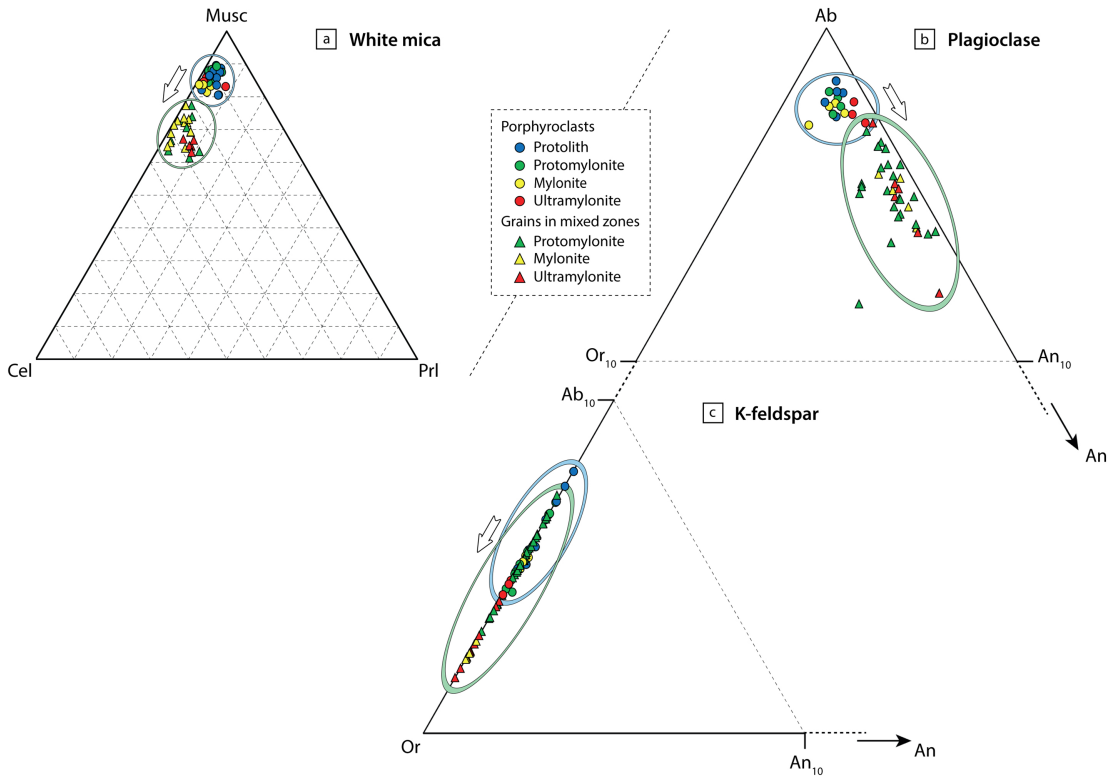


Figure 13

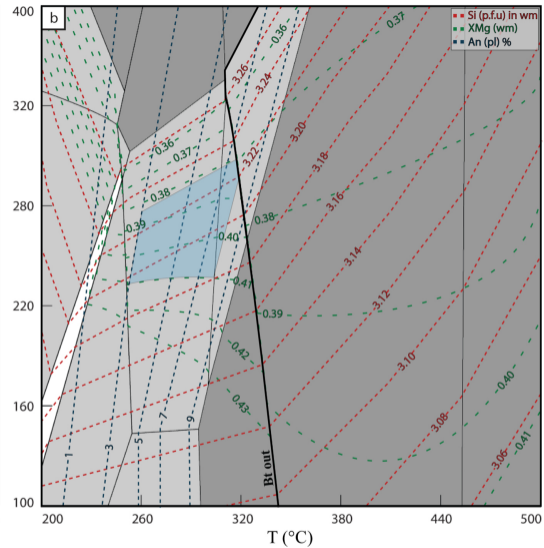
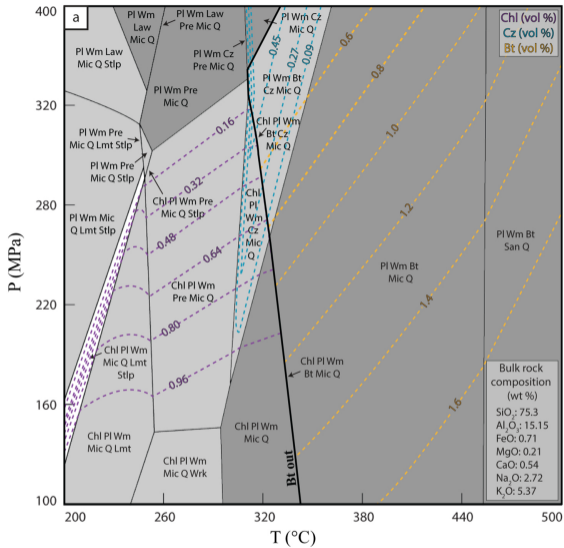


Figure 14