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The effect of strain on the crystallinity of carbonaceous matter: application of Raman spectroscopy to deformation experiments

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#### **0-Abstract**

Raman spectroscopy, applied to the carbonaceous matter (CM) present in fault zones, is a potentially powerful tool to aid in the reconstruction of slip conditions, provided the respective effect of heating and strain can be disentangled. For this purpose, we have carried out static heating and deformation experiments on low-grade sediments containing disseminated CM particles and tracked the evolution of CM crystalline ordering (crystallinity) using Raman spectroscopy. Heating at high pressure (200 MPa), at varying temperatures (500 to 800°C) and durations (3 to 1080h) resulted in an inc east d crystallinity, which was observed by an increase in the intensity ratio of the Discrete Band over the Graphite Band (Intensity Ratio R1 parameter) of the Raman spectra The experimental relationship between R1, T and duration was compiled into a law of contution, which was then used to design the deformation experiments. Experiments were performed in a Paterson rig at 150 MPa or in a Griggs-type rig at 1 GPa. We observed two types of deformation microstructures: (1) purely brittle ones, such as breccia along on pranes and (2) ductile ones, such as foliated layers, resulting from grain reorientation and comminution along with dissolution precipitation. In all deformation microstructures, an increase in R1 is visible and interpreted to be the result of higher crystallinity of c rbo aceous particles. Such increase is negligible in breccia, but is significant, up to 40%, in ductile deformation zones. The slow strain rates prevailing in ductile deformation prevented any local temperature increase, therefore the increase in crystallinity of the CM can be solely attributed to deformation. Thus, in natural fault zone, the increased crystallinity of CM reported in the literature and interpreted to be the result of frictional heating during earthquakes might rather be the result of strain localization without transient temperature increase.

#### 1-Introduction

Determining the degree of crystallinity of the carbonaceous material is possible through Raman Spectroscopy of Carbonaceous Matter (RSCM) analyses. Upon heating, the degree of crystallinity increases by thermal maturation through the carbonization (low-grade to greenschist-facies metamorphism) and graphitization stages (amphibolite- to granulite-facies metamorphism) (Beyssac et al., 2002; Lahfid et al., 2010; Oberlin, 1984). Qualitative correlations first linked the degree of metamorphism to CM lattice organization (Pasteris and Wopenka, 1991; Wada et al., 1994). More recently, quantitative calibration of these correlations resulted into the development of geothermometers based on the RSCM parameters, which are applitable to a temperature range of 150 to 650°C (Aoya et al., 2010; Beyssac et al., 2002; Kouketsu et al., 2014; Lahfid et al., 2010; Rahl et al., 2005). These geothermometers are widely used in ancient subduction and collisions zones studies (Beaudoin et al., 2015; Clerc et al., 2015; Lanari et al., 20.2).

Amongst regional-scale temperature trends, anon. clous RSCM spectra, i.e. these strongly deviating from the trend, are sometimes observed (Kec' r et al., 2020; Barzoi, 2015; Barzoi and Guy, 2002; Moris-Muttoni et al., 2022). For example, the ratio of the two main peaks of the Raman signal is increased by up to 40% in fault zoren from the Shimanto Belt (Moris-Muttoni et al., 2022) and by ~20% in a high strain zone in the everturned limb of a fold in the Alps (Kedar et al., 2020). A possible interpretation of these anomalies involves the effect of strain, which was mainly studied on well-crystallized CM in the advanced graphitization processes of natural (Kedar et al., 2020; Nakamura et al., 2015; Suchy et al., 1997) or experimental (Bonijoly et al., 1982; Bustin et al., 1995; Ross et al., 1992; Ross and Bustin, 1990) samples. An additional explanation for these anomalies relies on transient and local heating, caused by frictional slip during earthquakes (Furuichi et al., 2015; Ito et al., 2017; Ujiie et al., 2021). The case of anomalous RSCM signals in fault zones is particularly problematic, as they potentially combine the contribution of both strain and transient heating to CM crystallinity (Figure 1).

To explore the respective effects of heating and strain on CM crystallinity in low-grade samples, i.e. conditions of metamorphism at or below 300-350°C, we have carried out static heating experiments, in

order (1) to determine the relevant Raman spectra features that can be used to track crystallinity changes with heating, (2) to define the kinetic law that connects such features to temperature and time and (3) to define the appropriate conditions for deformation experiments that constitute the other part of the present work. These deformation experiments were exploratory, so rather than a systematic parametric study over a narrow domain of conditions, we considered a large range of experimental conditions and geometries. In particular, we used two different apparatuses, the Paterson rig and the Griggs-type rigs, enabling a large range of conditions of confining pressure to be explored. These experimental conditions led to a variety of microstructures, which vere then systematically analyzed by RSCM in order to connect deformation processes with CM evolution.

#### 2-Starting material and Methods

#### 2.1 Starting material

Samples used for static heating in autoclaves and deformation experiments have been selected from the Shimanto Belt, on the island of Kyushu and Jopan, where the first-order thermal structure across the belt (over ~100km width) has already been studied by RSCM (Raimbourg et al., 2014; Raimbourg et al., 2017; Raimbourg et al., 2019). Two hortogeneous shales, belonging to turbidites formations, have been sampled and used as starting numerical. The choice of shales was motivated by the fact that this material contains relatively abund at particles of carbonaceous matter (Raimbourg et al., 2017) and Raman spectra anomalies detected in fault zones were often found in shale-rich host rock (e.g., Ujiie and Kimura, 2014, Meneghi ii et al., 2010).

The maturity of the carbonaceous particles in the starting materials has been investigated using RSCM, vitrinite reflectance (%VR) and RockEval pyrolysis (all methods, except for RSCM, which is described below, are presented in the Supplementary Materials S2). Chemical and mineralogical compositions of the whole rock have been obtained by X-ray fluorescence (XRF) and diffraction (XRD) analyses, respectively. Microscopy techniques have not been used due to the small size of particles in both samples.

The first starting material, 18NOB01A, is from very low-grade metamorphic turbidites in the Miyazaki Group in the Shimanto Belt (Supplementary Materials S2). The Miyazaki Group is a Pliocene forearc basin composed of a thick sequence of turbidites (Murata, 1997; Shuto, 1961). The selected material has been sampled from a layer of shale in the turbidite unit and is composed of a fine assemblage of Quartz – Albite – K-feldspar – Calcite – Mullite – Illite – Pyrite. Semi-quantitative estimation based on the Rietveld method shows that the majority of minerals present are Quartz and Albite (ca. 60%), whereas calcite represents about 5% of the rock and the rest is composed of clays (ca. 35%). Average vitrinite reflectance values obtained for 18 NOB01A is  $0.54 \pm 0.04\%$  and corresponds to a temperature of  $84 \pm 7$  °C based on the calibratio 1 of Barker (1988) (Table 1) and a thermal maturity between the end of the diagenesis and the beginning of the catagenesis. RSCM parameters have been measured but the evolution of the  $\mathbb{R}$ ,  $\mathbb{R}$  man parameters at these low degrees of maturity is still controversial.

The second starting material, HN78, comes from 'urbuites belonging to the Hyuga Coherent unit in the Shimanto Belt, which was buried and 'commed within the accretionary wedge. The selected material was sampled from a layer of shale within turbidites. It is very similar in terms of lithology to 18NOB01A but is considered to be varigher metamorphic grade. This sample is composed approximately of 45% Quartz – 1.5% Albite – 33% Illite – 5% Chlorite based on the XRD results and the semi-quantitative Rietveld method. The chemical composition is given in Table 1. In this sample, the CM has reached a Chernal maturity at the transition between the metagenesis and the metamorphism. The calculated temperature obtained using the RSCM calibration of Lahfid et al. (2010) is approximately 201 ± 11°C. No vitrinite reflectance could be measured.

From RockEval pyrolysis, the carbonaceous particles in both 18NOB01A and HN78 belong to Type III, i.e., originate from small debris of terrestrial plants.

#### 2.2 Analytical methods

#### **2.2.1 RSCM**

Raman Spectroscopy of Carbonaceous Matter (RSCM) has been used to constrain the crystallinity of the carbonaceous material (CM) present as micron-size particles disseminated in the rock samples. Analyses were carried out using the ReniShaw InVIA microspectrometer, at ISTO-BRGM, Orléans, using an argon laser beam of 514 nm focused on the sample by a DM2500 Leica microscope equipped with x100 objective. These apparatuses are coupled with Renishaw software Wire 4.0. Acquisition time was 30 to 200 seconds (spectrometer opening time was ~5 s with 4.3 spectra accumulation). Laser power was maintained at less than 0.5 mW on the surface of the courte to avoid heating of CM grain during analyses. To avoid defect bands generated during thin section polishing all analyses were performed on subsurface grains by focusing the laser beam, through transparent or translucent minerals (Ammar and Rouzaud, 2012; Katagiri et al., 1988; Fasteris, 1989). For the static heating experiments, around 20 particles have been measured consecutive is spatially located and associated with a deformation microstructure, so that various across the sample or across the different microstructures can be determined.

Raman spectra processing was as ne using PeakFit v4.12. To carry out the linear baseline correction, the baseline fit was obtain d by selecting the data points in the ranges 900-1000 and 1900-2000cm<sup>-1</sup>. The Raman spectra of the CM considered here is composed of two maxima, one around 1600cm<sup>-1</sup> (the graphite-like "G band"), corresponding to highly-ordered carbon and one around 1350cm<sup>-1</sup> (the defect "D band"), corresponding to disordered carbon (Beny-Bassez and Rouzaud, 1985; Lahfid et al., 2010; Sadezky et al., 2005; Tuinstra and Koenig, 1970; Wang et al., 1990).

To interpret Raman spectra, several parameters were assessed. First, on baseline-subtracted spectra, the position and intensities of the two maxima (the D and G bands) were determined. Second, we applied peak deconvolution procedure to the baseline-subtracted spectra, following Lahfid et al. (2010). This approach is well adapted to low metamorphic grade material and consists of a 5 band

(D1, D2, D3, D4 and G) deconvolution using Lorentzian functions. D4+D1+D3 peaks form the D band, while the G band is composed of D2 + G peaks.

During the transformation from low-grade carbonaceous matter to perfect graphite, Raman spectra evolution can be decomposed in two successive steps. In the low temperature realm (i.e. 200-330°C), the D band increases in area and intensity with increasing metamorphic grade (Lahfid et al., 2010), while at higher temperature, the D band decreases in area and intensity with increasing grade (Beyssac et al., 2002). In the present study, parameters used to track the CM evolution during static heating and deformation experiments are the D- and G-band positions, full width at half maximum of the D1 and G bands (FWHM), R1 intensity ratio (Beyssac et al., 2002) and datan e between the D and G bands (i.e. RBS in Henry et al. (2019)). Of particular interest in the present work is the evolution of R1 (Figure 4), as it shows anomalies in experimentally and no uran'y faulted rocks (Muirhead et al., 2021), which still remains to be understood. In addition, peak area ratios used for thermometric calibrations (Beyssac et al., 2002; Lahfid et al., 2010), recalting from peak deconvolution, have also been analyzed.

All raw and processed Raman spectra ar . vanable at https://doi.org/10.5281/zenodo.8321278.

#### 2.2.2 Electron microscopic imaging

At ISTO, Orléans, Scanning Elect on Microscopy (SEM) analyses were performed by using a SEM Merlin Compact Zeiss, equipped with an EDS Bruker Quantax detectors, at 15kV. EDS maps were acquired at 15kV with a dwell time of 850µs and summed over 9 scans.

#### 2.3 Experiments

#### 2.3.1 Hydrostatic heating experiments

Hydrostatic thermal maturation has been performed in autoclaves in order to understand the CM evolution under constant temperature and pressure during the carbonization and early graphitization stages. Powdered samples of the two shales, 18NOB01A and HN78, sealed in gold capsules, were matured at a pressure 150MPa and a temperature of 500 to 800°C for a duration between 3 hours to 45 days. The heating rate was set at 4.5°C/min. Additionally, distilled water was added to some

experiments. The details of the sample preparation and experimental techniques are presented in Supplementary Materials S1. After completion of the experiments, the CM crystallinity was estimated by RSCM analyses. All the results obtained are presented in Table 2.

#### 2.3.2 Deformation experiments

2.3.2.1 Objectives and overview: This study includes 8 Paterson Rig apparatus and 4 solid-medium Griggs apparatus experiments (Figure 1, Figure 2), performed at low (150MPa) and high (1GPa) confining pressures, respectively. The aim of these experiments is to reproduce experimentally the structures of deformations shown in Figure 1, i.e. faults with features of brittle deformation (such as fault core gouge) coexisting with more distributed, ductile, in formation structures (such as the progressively elongated clasts), as well as to analyze the corresponding Raman signatures. For this purpose, we explored experimental conditions wherein (1) both brittle and ductile deformation microstructures could be observed and (2) contrasting Raman features are observed, enabling the effect of deformation on CM crystallinity to be carter random.

Laboratory rates of deformation are by definition orders of magnitude larger than natural ones; therefore, to compensate for this, large type ally >700°C) experimental temperatures are necessary in order to avoid purely brittle deformation and to reach the onset of ductile deformation. However, during deformation and static hearing experiments (see the section 4.1) at high temperatures (700 or 800°C), the CM Raman spectra tends asymptotically towards an endmember shape, which corresponds to the end of the carbonization processes, such that the effect of deformation vanishes.

As a result of this delicate compromise to obtain simultaneously a large variety in microstructures and contrasting Raman signature across the experimental sample, a large range of conditions were tested on two distinct deformation apparatuses. In many instances, deformation behavior was mostly brittle and deformation was extremely localized on slip planes.

With Paterson Rig apparatus, 5 experiments were carried out in simple shear geometry and 3 in co-axial mode on cylinders. With the Griggs apparatus, 2 experiments have been carried out at constant deformation rates in the "Tullis-modified" Griggs apparatus and 2 in the new Griggs apparatus at

constant external load. Deformation experiments were performed on intact or powdered shale 18NOB01A. Experimental conditions are detailed in Table 3.

2.3.2.2 Experimental procedure in the Paterson rig: The deformation experiments were carried out on the Paterson rig available at Orléans (see details in Paterson (1990) and Figure 2). The confining pressure was applied by Argon up to a pressure of 100MPa before heating. The temperature was increased to the target temperature at a rate of 4.5°C/min, after which the confining pressure is adjusted at 150 MPa. When the temperature and pressure were stable, the load-cell was run in order to begin the deformation. Deformation experiments were carried out at \$\frac{1}{2}\$0 MPa, from 600 to 700°C, for 3 to 42 hours and at constant displacement rates of 0.1 to 0.0. \( \text{ur} \) /s. One 72h experiment was conducted at a constant load, corresponding to \$\times 10^{-7} \text{s}^{-1}\$ strain retain At the end of the experiment, the oven was turned off and the temperature dropped below 100 C in few minutes. At this temperature, the autoclave was then depressurized slowly to prevent accompression cracks. After the experiments, the experimental assembly was removed from the autoclave and the sample was cut and prepared for analysis.

#### 2.3.2.3 Experimental procedure in the Grigos-type rig:

The Griggs-type deformation experiments were carried out on two different apparatuses, both present in Orléans. The Griggs-type rig. are either a Tullis-modified design (Marti et al., 2018; Pec et al., 2012) or a more recent design ("new Griggs" (Précigout et al., 2018)). The Tullis-modified rig or the new Griggs were used to perform constant strain-rate (at ~2 10<sup>-5</sup>s<sup>-1</sup>) and constant external load experiments, respectively. In the latter type of experiments, the applied nominal load is fixed, but the actual load on the sample increases with deformation as the surface of overlap of the two pistons decreases. The confining pressure was applied by either a KI or NaCl confining medium at conditions of deformation at 400°C or 600°C, respectively. The heating rates were 16°C/min for the Tullis-modified design (experiments T609BM and T610BM), and 30°C/min for the "new Griggs" (experiments OR87BM and OR92BM).

In the constant load experiments, two sets of conditions were tested. In OR87BM, the sample was left at 600°C and 1 GPa for 17 hours before deformation proceeded. Several steps of increasing load were then applied, and the strain rate for the highest external load (500 MPa) was ~10<sup>-5</sup>s<sup>-1</sup>. In OR92BM, the sample was left at 600°C and 1 GPa for 1/2 hour before setting the external load (at 500 MPa), resulting in a much higher deformation rate on the order of 5.10<sup>-4</sup>s<sup>-1</sup>.

#### 3. Results

#### 3.1 Static heating experiments

Of all the parameters studied (Figures 3 and 4, Supplementary M, terials S3), the ones that show the most significant and straightforward relationship to heating, for the different temperatures applied, are the ratios of areas of fitted peaks RA1 (Lahfid et al., 2013)-(Figure 3a), as well as the intensity ratio R1 (Figure 3b). In particular, the latter parameter follows a monotonic (i.e., f(x) varies only in one direction over the interval of interest  $x \in [x1, x2]$ ) increase in the three highest temperatures (out of the four tested ones).

#### 3.2 Deformation experiments

#### 3.2.1 Mechanical behaviour

Paterson experiments (Figure 5 and Table 3): In the two experiments containing porous alumina and having a connection of the powdered sample to a reservoir of water with controlled pressure (PP629 and PP630), the sample deformation only consisted of compaction, until deformation shifted to the alumina pistons. These two experiments are therefore not considered in the following. To avoid dehydrating the sample upon heating and transform of the material into hard and undeformable ceramics, all the other Paterson experiments used non-porous pistons.

In all experiments, including shear and coaxial geometries (with the exception of the constant load experiment (PP649)), stick-slip events, characterized by a sharp stress drop and rapid forward motion of the load cell, occurred (Figure 5). The differential stress before the initial stick-slip event in the coaxial experiments PP638 and PP641 is close to the Goetze criterion ( $\sigma_1 - \sigma_3 = P_{conf} = 150 \, MPa$ ),

considered as the onset of brittle deformation (Kohlstedt et al., 1995). In shear experiments, the differential stress before the initial stick-slip event is much more variable, from 60 to 130 MPa. The differential stress after stick slip is also variable, from ~30 to 70MPa.

Griggs experiments (Figure 6 and Table 3): The two, constant strain-rate experiments (T609BM and T610BM) show contrasting mechanical behavior, as a result of applied temperatures of 400 and 600°C, respectively. At 400°C (T610BM), the differential stresses increase to values of ~930 MPa, close to the Goetze criterion (1GPa), before a significant and progressive weakening down to 630 MPa. At 600°C (T609BM), the differential stresses increased in notonically throughout the experiments, while remaining at values of below 400 MPa. The locating curve in the experiment at 600°C has a much lower slope than in the experiment at 400°C.

In the constant load experiments, large differences arose is a result of the duration of the heating stage at 600°C and 1 GPa without any differential stress, which precedes the deformation stage. For a similar nominal load of 500 MPa, the experiment statically heated for 17h (OR87BM) was by a factor of ~50 more viscous than the experiment statically heated for ½ hour at the same conditions before deformation (OR92BM).

#### 3.2.2 Microstructures

Low pressure experiments (Lawrson rig):

In all shear geometry experiments, slip along the boundary between the piston and the sample was observed. In sample PP633, breccia zones, characterized by a large and heterogeneous grain size reduction, and a large porosity compared to undeformed domains, are distributed through the sample (Supplementary Material S4). In sample PP636, along the piston-sample boundary, a µm-thick layer is present, where grain size reduction proceeded down to the submicron scale and porosity has disappeared. Further away from the boundary, a ~5-10 µm thick layer is constituted of clasts of the original material, embedded in clays developing a foliation (defined by phyllosilicate preferred orientation), which asymptotically merges into the boundary layer. Further away, deformation is no longer visible.

In coaxial experiments (PP638 and PP641), conjugate fractures are observed. They consist, in both experiments, in straight slip surfaces cutting across the whole samples. In PP638, the slip plane is surrounded by breccia layers with a thickness of  $\sim 100 \, \mu m$ , asymmetrically developed on both sides, or localized in step-overs of the slip plane (Supplementary Material S4).

In PP641, which was deformed at a strain rate 10 times lower than PP638, a high strain, foliated layer developed along the slip plane. The foliation is defined as the preferential orientation of elongated clasts of quartz/feldspar, and elongated domains of phyllosilicates (Figure 7c and e). This foliation is at a low angle to the slip plane and is asymptotically curved towards. In the foliated layer, the grain size is significantly reduced and the porosity is decreased, with page 2t to undeformed material. In certain cases, breccia zones are also present locally and are page 11.1 to the high strain foliated layer along the slip planes (Figure 7c).

High pressure experiments (Griggs-type rig):

All the high-pressure experiments were carried out in shear geometry. Unlike the low-pressure experiments described above, the deformation is not localized along the sample-piston boundary, but along a fracture zone at a low angle of such boundary, with a synthetic sense of motion (Supplementary Material S4). Present along the fracture plane is a breccia layer, a few microns- to a at most a few tens of microns-thick and characterized by a large grain size reduction and a large porosity compared to the host rock.

Additionally, in three of the four samples (TM610BM, OR87BM, OR92BM), ductile deformation is also present within a foliated layer that can reach several hundreds of micrometers of thickness, located between the fracture/slip plane and the undeformed host rock. The foliation is defined by the preferential elongation of phyllosilicates, quartz/felspar and pyrite aggregates (Figure 8). Towards the slip plane, within the ductile deformation zone, a high strain layer is present locally, as is apparent by the deflection of the foliation towards the slip plane and a decrease in grain size (Figure 8).

#### **3.2.3 Raman**

One potential bias of deformation experiments is the actual temperature field. Heat conduction across the experimental samples potentially leads to temperature gradients across the sample between the lower and the upper alumina piston (Pec et al., 2016), and thus these gradients might influence the RSCM parameters. To assess this possibility, we determined whether any R1 variations were present and if they were systematic with respect to the distance to upper and lower alumina piston.

At low pressures (Paterson rig), sample PP649, which was deformed to bulk strain larger than 10% and without localization of the deformation, shows R1 values of 0.622+/-0.025 (standard deviation), i.e. minimal variations and no systematic trend with respect to the position within the sample. As a result, for Paterson deformation experiments, comparisons of R1 values across the whole sample can be performed.

Conversely, in high pressure experiments (TM609BM), measurements in similar, undeformed material near the upper and lower alumina pistons showed systematic R1 differences (0.804+/-0.082 for top vs. 1.013+/-0.075 for bottom), consistent with a compensature gradient across the sample (because of the heat conduction described above), and as a consequence R1 values could not be connected solely to strain. As a result, for Griggs-type experiments we focused only on carbonaceous particles across relatively small distances (of the order of a few hundreds of µm).

#### 3.2.3.1 Breccia zones

Few measurements we. a cyrried out in breccia zones, because of the difficulty finding appropriate particles that could be analyzed according to the protocol. In the lower pressure sample PP633, a slight increase in R1 (~+5%) is observed compared to undeformed host rock in the two piston-sample boundaries, wherein fracturing is pervasive (Figure 9). In a higher-pressure sample, a fractured zone contains CM particles with a R1 ratio that is slightly higher, by 3% on average, than the undeformed host rock nearby (Figure 10). Given the many parameters, external to CM crystallinity, which may influence the Raman spectra, in particular the increase in sample fluorescence in fine-grained zones, it is highly questionable whether strain itself has an influence on CM crystalline organization.

#### 3.2.3.2 Ductile strain zones

A significant increase in R1 is observed from undeformed zones to foliated, high strain domains next to the slip planes, irrespective of the low or high-pressure conditions (Figures 10 and 11, Supplementary Materials S5). In these three examples, the increase ranges from +4 to +41% (the values are defined as averages over microstructural domains). In samples where gradients of strain are visible as variations in the grain size and the preferred orientation of the microstructural elements, the increase in R1 is correlated with the increase in strain (Figures 10 to 12). The largest increase, +41%, is observed in the low-porosity high-strain zone of sample OR92BM (Figure 10). From all the possible parameters computed from RSCM spectra, R1 is the parameter that thows the most systematic and the largest variation as a function of strain (Figure S10).

#### 3.2.4 Chemical changes in the high-strain zones

EDS compositional mapping was performed in zones with deformation gradients in low pressure sample PP641. Clasts in the undeformed domains are composed of quartz, albite and K-feldspar, whereas the matrix is composed of illite and chi'one' (Figure 13). In the high-strain layer adjacent to the slip plane, the clasts are elongated and derine a foliation bending towards the slip plane. These elongated clasts, originating from felds par grains, contain both K and Na, i.e. have an intermediate composition between albite and K-fer spar. The distribution of these two elements is relatively heterogeneous. The BSE imaging of these chemically mixed zone revealed the comminution of the original feldspar, along with a very large microporosity.

#### 4. Discussion

# 4.1 Kinetics of maturation in static heating experiments

#### 4.1.1 Relevant Raman parameters to track crystallinity increase during heating

In both nature and experiments the increase in temperature affecting CM particles leads to an evolution in chemical and crystalline structure and is reflected in changes in the Raman spectrum (Beyssac et al., 2002; Lahfid et al., 2010; Li et al., 2006; Pasteris and Wopenka, 1991; Wopenka and Pasteris, 1993). Accordingly, a wealth of Raman parameters have been proposed as proxies of temperature, such as peak area ratios (Beyssac et al., 2002; Lahfid et al., 2010), peak width (Kouketsu et al., 2014), amplitude ratios of the peaks resulting from the raw spectrum deconvolution (Ito et al., 2017) or amplitude ratio of the maxima of the raw spectrum (Kanek and Hirono, 2018, 2019; Kaneki et al., 2016).

In the static heating experiments shown here (Figure 3, figure S3 and Table 2), several parameters, such as D1 FWHM, RA1 and R2 area ratios, and R1 intensity ratio, evolve as a function of heating intensity and duration. Amongst these different parameters, the R1 intensity ratio (Figure 4) is the best proxy for CM crystallinity. Justifications for this conditions are, first, the R1 is the parameter that shows monotonic variations, except for the experiment with HN78 (Figure S3), enabling the derivation of quantitative laws of evolution. Second, the correlation of R1 with temperature is demonstrated from earlier studies, both in a tural metamorphic samples (Kouketsu et al., 2014; Lahfid et al., 2010) and in experimentally statically heated ones (Kaneki et al., 2016; Li et al., 2006; Muirhead, 2012). According to Last al. (2006), the increase in R1 reflects the relative increase, upon heating, of the concentrations of aromatic rings having six or more fused benzene rings. Finally, in experiments of deformation R1 intensity ratio is the most widely reported variable, enabling the effect of deformation on CM crystallinity to be studied (see Table 5 for the compilation of available studies). Please note that we have used the intensities defined as maxima on the full spectrum (Kaneki and Hirono, 2018, 2019; Kaneki et al., 2016) (Figure 4), and not as amplitude of the peaks obtained through peak fitting procedure (e.g., Ito et al. (2017)).

#### 4.1.2 Kinetic model of Raman spectrum evolution upon heating

As discussed above, the R1 ratio evolution is monotonic in the range of experimental heating conditions considered. To derive the kinetic laws that link the R1 parameter to duration of heating and

temperature, we followed the method presented in Nakamura et al. (2017, 2020) for higher degrees of crystallinity (i.e., graphitization stage). Previous studies for simple evolution proposed the use of a power rate model or the Johnson-Mehl-Avrami (JMA) model (Huang, 1996; Muirhead, 2012); however, carbonization, similar to graphitization, is not a simple chemical reaction, but the result of simultaneous reactions (formation of basal structural units, devolatilization and crystal growth). Therefore, in order to model the complex chemical evolution of the CM, we preferred to use the method proposed by Nakamura et al. (2017), using a sigmoidal function with the following equation:

$$R1(T,t) = R1_{min} + \frac{(R1_{max} - R1_{min})}{1 + \left[\frac{t_{half}}{t}\right]^h} \tag{1}$$

where from the immature, original state  $(R1_{min})$ , a sharp volution in R1 is observed (at high temperature) over few hours, after which R1 remains close  $\kappa$  R1<sub>max</sub>. In this equation,  $t_{half}$  and h are the inflection point and the order of reaction, called "Iill coefficient" of the sigmoid function, respectively. The parameter  $t_{half}$  could be described following an Arrhenius relation as:

$$t_{half} - A exp\left(\frac{-m}{T}\right)$$
 (2)

Combining the equations (1) and (2) we obtained:

$$R1(\vec{r}, t) = R1_{min} + \frac{(R1_{max} - R1_{min})}{1 + \left[\frac{A \exp\left(\frac{-m}{T}\right)}{t}\right]^{h}}$$
(3)

Where  $R1_{min}$  and  $R1_{max}$  are the minimum and maximum R1 values, respectively; A is the intercept and m the slope of the Arrhenius plot; T and t are the temperature and the time of heating, respectively.

The full procedure is explained in Supplementary Materials S6. Results of the fitting procedure applied to static heating experiments are shown in Table 4 and Figure 14 for the two samples analyzed. The parameters for sample HN78 are close to the ones obtained in Moris-Muttoni et al. (2022), but are more reliable as they are calculated from a larger experimental dataset. Form Figure 14, one can also observe that the initial heating stage (at rates of 4.5°C/min) introduces negligible

errors, because before reaching high temperatures, such as 700°C or above, the evolution of R1 is very slow compared to the duration of heating.

Finally, these kinetic laws can be used in the deformation experiments described hereafter to predict the contribution of sole heating to the evolution in the crystalline organization of the CM.

#### 4.2 Deformation processes

We analyze here first what types of deformation processes were operating in the experiments we performed, and which natural conditions of deformation these experiments are relevant for.

Two apparatuses were used, with contrasting conditions of confined pressure. At low pressure, the principal slip zone is localized along the sample-piston bouldary, whereas at high pressure, strain is accommodated by a synthetic fracture zone cutting through the sample. In spite of these different geometries, the microstructures observed are relatively similar, as we systematically observed a breccia zone along the main slip interface. The breccia layers, defining the slip plane, are characterized by a large grain-size decrease with respect to starting material, clasts of variable size but with angular shapes and small aspect ratios, and a large porosity (Figures 7 and 8, Supplementary Material S4).

We also observed, along certain slip surfaces, high-strain foliated layers. These layers are characterized by small grain stre, elongated and preferentially orientated grains or grain aggregates, and a reduced microscopic porosity (Figure 8). At smaller scale, in the elongated feldspar aggregates present in the high-strain layer, pervasive grain comminution, accompanied with a widespread submicron porosity, is visible (Figure 13).

Pec et al. (2016) analyzed the semi-brittle flow of granitoids in high pressure experiments, at lower temperature but higher strain rates, than the experiments described here. In their experiments, where the starting grains are two orders of magnitude larger than in ours, localized zones of deformation form, first Riedel shear zones then macroscopic shear-plane parallel ones. The sequence of processes starts with ductile deformation of the grains accompanied by pervasive grain fracturing, followed by the formation of an amorphous layer in the highest-strain zone. The pervasive grain fracturing in the

plagioclase in Pec et al. (2016) is accompanied by chemical exchanges and enrichment in K, which the authors interpret as reflecting the operation of dissolution-precipitation. The chemical patterns in plagioclase from the high strain zone reported here suggest that the same processes are active.

Deformation processes have therefore resulted in the formation of two types of microstructures, characterized by contrasting porosity. In some cases these microstructures are next to each other, irrespective of the pressure (Figure 7, low pressure, and Figure 8 bottom, high pressure), raising questions as their relative contribution to total strain, or their simultaneity. In the example of PP641, the relative contribution of the breccia and the high strain foliated layer is unraveled by the thin layer of pyrite connecting the two large fragments of the original grain or bo h side of the slip zone (Figure 7b). The thin layer of pyrite, parallelized to the slip plane, is within the high strain foliated zone, suggesting that most of the total strain is accommodated there and not on the breccia layer next to it. Furthermore, in PP641 deformation experiments, as well as in all low-pressure, constant strain-rate experiments, the macroscopic stress-strain curve shows the alternation of two types of behavior, either stick-slip events or steady-state creep (Figure 7). Stick-slip events and creep accommodated a small and a large fraction of total strain, respectively. This comparison between the macroscopic behavior and the strain accommodated suggests that receia formed during stick-slip events, whereas high strain foliated zones formed during cre o. This interpretation is consistent with the microphysical models and the frictional viscous rhed ogy (Den Hartog and Spiers, 2014; Niemeijer, 2018; Niemeijer et al., 2002), wherein the marourn tures are interpreted to be the result of the competition between chemically-induced, dissuition-precipitation mediated, compaction and strain-induced porosity creation. In the highest strain-rate zone, on the slip plane itself, high porosity breccia forms because dissolution-precipitation is slower than strain-induced grain fracturing and dilatancy. In the layers next to the breccias, strain-rate is lower, dissolution-precipitation reactions in combination to grain-size reduction lead to rearranging the grains and closing up of the pores.

The case of high-pressure experiments is more problematic, as breccia and high strain foliated layers are both present (Figure 8), whereas no stick-slip events occurred. In the high-pressure experiments by Pec et al. (2016) reported above, or in Marti et al. (2017), porous breccia such as shown in Figure 8-

bottom are absent, while they observed zones of very high, ductile strain similar to the high strain foliated layers described in the present study. A peculiarity of the experiments carried out here, compared to Pec et al. (2016) or Marti et al. (2017), is the abundance of hydrous clays, such as smectite-illite, which may release water during the deformation at high temperature. Water release is a factor favorable to the formation of breccia, as attested by hydraulic fracturing structures observed in nature (e.g., Jebrak (1997)). Nonetheless, further experimental work is necessary to decipher the factors that control the types of microstructures that form during high-pressure deformation experiments.

In summary, the concomitant operation of brittle-fracturing and do solution-precipitation operating in our experiments (Figure 13), similar to the observations in Peter 2. (2016), support the interpretation of semi-brittle deformation regime. This regime of defor nation prevails in the seismogenic range of depths, i.e., the upper part of the crust where temperature conditions are lower than ~300-350°C (Sibson, 1977; Hyndman et al., 1997; Scholz, 1979). The high-strain foliated microstructures formed in the experiments of the present study which the ductile structures and microstructures of deformation present in the corresponding ration of natural conditions (e.g., Raimbourg et al., 2019).

# 4.3 Effect of deformation on Raman signature and crystallinity of carbonaceous matter

#### 4.3.1 Strain or shear he. in 5.

Deformation, irrespective of its rate, releases some energy, hence raises locally the temperature. The amplitude of the local temperature rise needs to be evaluated, in order to discriminate the influence of deformation from the temperature onto the Raman signature of carbonaceous matter. In the deformation experiments we performed, order-of-magnitude calculations (Supplementary Material S7) show that the effect of temperature on the Raman signal is limited to low-pressure experiments, where stick-slip events occurred. Furthermore, this effect is restricted to a layer at most  $\sim 1 \mu m$ -thick along the slip surface.

As a consequence, except for these very restricted zones, all the Raman signal variations observed in the present study can be safely connected to strain and not to temperature effect. In particular, the large increase in R1 observed in large-strain zones, within a layer of ~50 µm thick (Figure 11), cannot be attributed to transient heat pulses during stick-slip events and is therefore the result of strain. Semi-brittle, ductile deformation, similar to what can be observed in the "seismogenic" range of conditions in nature, has an undisputable and large effect on the increase in crystallinity of CM particles.

#### 4.3.2 Interpretation of R1 increase in terms of crystallinity of carbonaceous particles

The static heating experiments reported in Section 4.1, similarly to other experimental studies (Deldicque et al., 2016; Hirono et al., 2015; Li et al., 2006), are consistent with an increase in R1 with increasing temperature. The deformation experiments reported in section 4.2 show that strain has a similar effect to temperature and tends to increase R1, in comparison to undeformed material.

All the Raman spectra analyzed in the present work is examposed of five characteristic bands, typical of the low T conditions (up to ~330°C) that correspond to the carbonization stage (Lahfid et al., 2010). In this range of T conditions, increase in metamorphic grade results in an increase in the relative height of the D band (Rahl et al., 2003) which reflects an increased structural ordering of the carbonaceous matter. The increase in R1 observed in deformation zones (Figures 10 and 11) therefore reflects an increased crystallinity. Thus, strain has a catalytic effect on the process of carbon ordering.

For T above 330°C (the gr. phi'ızation stage), on the contrary, increased metamorphism results in the decrease in the height and amplitude of the defect band (Beyssac et al., 2002; Kouketsu et al., 2014). This two-stage evolution, which led to define several Raman-based geothermometers depending on the T range considered (Beyssac et al., 2002; Lahfid et al., 2010), restricts the application of the present study to the relevant T range, namely the low range, where carbonization operates.

#### 4.3.3 Comparison with other experimental studies

Many high-velocity experiments have tried to reproduce the evolution of CM crystallinity in fault zones as a result of seismic slip (Table 5). Invariably, these experiments have shown that at seismic slip velocities (~m/s), an increase in CM crystallinity in zones of highest slip is observed for low-grade

starting CM (Aubry et al., 2018; Bustin et al., 1995; Furuichi et al., 2015; Kaneki and Hirono, 2018, 2019; Oohashi et al., 2011; Savage et al., 2018). This conclusion does not hold for higher grade CM material evolving within the graphitization stage, as contrasting evolution, i.e., increase (Kuo et al., 2017) or decrease in crystallinity (Kaneki and Hirono, 2019; Kirilova et al., 2017), is observed. We therefore restrict the discussion here to the processes affecting low-grade CM material evolving within the carbonization stage, similar to the present study.

In all the high-velocity experiments cited above, a large temperature increase is predicted, and the evolution of CM towards higher crystallinity that is observed is interpreted as the result of this flash heating within the principal slip zone. Nonetheless, this effect of temperature does not exclude the possible complementary effect of strain.

Several friction studies have carried out tests at different sing velocity, encompassing seismic (~m/s) to subseismic (~100µm to 50mm/s) slip rates. These friction experiments are consistent with a large influence of the slip velocity, and the CM participes plution at low slip rates is non visible (Oohashi et al., 2011). Note, that these friction experiments are carried out at ambient temperature and low (~1-10MPa) normal stress.

Slow strain rate experiments have and been carried out at high pressures and temperatures in a Griggstype apparatuses, similar to the experiments of the present study (Bustin et al., 1995; Wilks et al.,
1993). It has been shown in these experiments that strain promotes the reorientation, alignment and
coalescence of the basal structural units made of polyaromatic layers, the reduction of the porosity,
and overall an evolution of carbonaceous matter towards graphite structure. This evolution, supported
by TEM and XRD anlayses, is recorded in the Raman signal by an increase in R1 ratio similar to the
one observed in the present study. Bustin et al. (1995) note the discrepancy between the different
evolution (TEM and XRD vs. Raman), where the latter show an apparent increase in the defect band,
whereas the two other techniques show an increased organization of the crystalline structure. We
remark here that this increase in R1 with increasing maturity, although its atomic-scale interpretation
is unclear, is exactly the same as the one observed in natural rocks in the carbonization stage

(Kouketsu et al., 2014; Rahl et al., 2005), and was also observed experimentally as a result of heating (Li et al., 2006).

Slow strain-rate experiments therefore confirm the effect of strain on enhancing the crystallinity of low-grade CM, similar to the present study. Furthermore, the comparison of different experimental conditions highlights the major role of confining pressure and temperature: at room conditions, experiments are unable to reproduce the effect of strain, which is observable only under the conditions of high pressure and temperature that are relevant to natural shear and fault zones. Finally, a direct effect of pressure on enhancing CM maturation is also apparent from the comparison of Paterson and Griggs-type experiments (150MPa vs. 1 GPa of confining pressure) carried out at 600°C. In undeformed zones, at 1 GPa for 20hrs, (OR92BM), R1 reaches 0.69, while at 150 MPa for 42hrs (PP641) and 75hrs (PP649), R1 reaches only 0.62.

#### 4.3.4 Comparison with natural studies

In naturally deformed samples, numerous str dies have focused on the characteristic of CM particles in zones that concentrates deformation, in order to decipher the deformation processes. For low-grade CM material in the range of temperature corresponding to carbonization, most studies have recorded an increase in crystallinity in zones of strain, based on the RSCM signal (Ito et al., 2017; Kaneki et al., 2016), biomarkers (Savage and rollissar, 2019; Savage et al., 2014) or vitrinite reflectance (Suchy et al., 1997) (Table 5). The est dies were, for the largest fraction of them, focused on fault zone material, hence they are dyzed rocks that localized both strain and (potentially) heat production/temperature rise. While temperature was considered as the main factor causing the increase in the crystallinity of CM particles within fault zones (Ito et al., 2017; Kaneki et al., 2016; Savage and Polissar, 2019; Savage et al., 2014), the results of the present experimental study show that the Raman anomalies in fault zone rocks are ambiguous and difficult to interpret, as both temperature AND strain have an influence on CM crystallinity and the associated Raman signature. This complexity is illustrated in the datasets of natural fault zones ranging from low-grade up to greenschist-facies conditions studied by Muirhead et al. (2021), where the authors reported an inconsistent evolution with either a decrease or an increase in R1 between fault rock and undeformed host rock. Muirhead et

al. (2021) suggests that the effect of strain is a function of the pre-deformation crystallinity of the CM particles.

Natural slow strain-rate structures of deformation have focused much less attention and corresponding studies are much fewer. Nonetheless, the three studies reported in Table 5 have all concluded to an increased crystallinity of carbonaceous matter zones of localized ductile strain (Kedar et al. 2020, Petroccia et al., 2022; Suchy et al., 1997), over a large range of temperature conditions. The interpretation of this increased crystallinity is nevertheless controversial, as Petroccia et al. (2022) and Suchy et al. (1997) invoke shear heating rather than strain, to account for the anomaly in Raman signature. In Kedar et al. (2020), a zone of concentrated strain with the overturned limb of a fold was characterized by a decrease in R1, but also interpreted as a super-related increase in maturity. The starting material in Kedar et al. (2020) has spectra whose shape is transitional between the carbonization and graphitization stages, hence the interpretation of R1 variations is not straightforward.

As a summary, if natural, slow strain-rate struct, res of deformation show a systematic increase in CM crystallinity, the anomaly in Raman signal in high strain-rate structures such as fault zones is still relatively unclear. Beside strain, strain-rate itself may have a variable effect on CM crystallinity and Raman signal. This is illustrated in higher grade material during the graphitization stage, where two contrasting trends are obserted and are a function of the type of deformation processes. In ductile shear zones, where strain-rate is low, strain is accompanied by increased CM crystallinity (Petroccia et al., 2022), whereas in fault zones, with cataclasites, ultraclasites and pseudotachylites, cutting across high-grade rocks, a decrease in CM crystallinity was reported (Nakamura et al., 2015). In the former case, the strain energy of viscous deformation is interpreted as promoting graphitization and in the latter case, high-velocity deformation promoted amorphization and decrease in crystallinity. How this relates to the carbonization stage is not known, but it is possible that slow, aseismic deformation and very rapid, seismic deformation have different influence on CM evolution.

#### 4.4 RSCM geothermometry applied to naturally deformed rocks

Beyond its application to geothermometry (Bellanger et al., 2015; Berger et al., 2020; Delchini et al., 2016; Girault et al., 2020; Raimbourg et al., 2014), RSCM has also been applied to deformation zones in order to decipher the past temperature field, particularly the occurrence of flash heating during earthquakes (Kouketsu et al., 2017; Savage and Polissar, 2019). The present study demonstrated the effect of strain, independently of temperature, to influence the RSCM signal in low grade CM particles typically present in fault cores cutting across sedimentary terranes. Even more, high-resolution R1 profiles across several natural fault zones showed that R1 is principally controlled by strain, and not temperature (Moris-Muttoni et al., 2022). For these reasons, usin r RSCM to reconstruct the past temperatures experienced within natural fault zones is questionable

Similarly, the effect of deformation to enhance crystallinity is combistacle to the use of RSCM as a geothermometer in naturally strained rocks. The influence of deformation is significant, as illustrated in Figures 10 and 11. Although many field examples provide ample exposures of undeformed rocks, the issue is still central for rocks that are pervasively deformed, such as tectonic mélanges. This is for instance the case in the Japanese Shimanto Prat, where several units of tectonic mélanges show an anomaly in the crystallinity of carbonaccous matter, as inferred from vitrinite reflectance data (Sakaguchi, 1999). A geological scenario with local thermal anomalies might well account for the higher CM crystallinity in tectonic meaninge units, but the high localization of strain in these units, and its effect of on CM crystallinity, is in alternative explanation.

#### **5 Conclusions**

The static heating experiments carried out in this work have shown that the evolution of CM is best tracked by the Intensity Ratio Raman parameter R1. The increase in R1 upon heating, from a low-grade starting material, follows the same trend as the one observed in natural metamorphic series. It is therefore interpreted as reflecting an increase in the structural ordering, within the range of evolution corresponding to carbonization.

The ductile deformation zones obtained experimentally at low and high pressure are also characterized by a large increase in R1, reflecting a higher crystallinity than undeformed material. These results

point to the influence of strain, at slow strain-rate and without local temperature increase, to reorganize CM crystalline structure.

The influence of strain on CM crystallinity potentially undermines all the conclusions drawn so far in many studies from the literature on naturally faulted rocks, where the anomaly in Raman signature of CM (compared to undeformed host rock) has been interpreted as reflecting coseismic shear heating during past earthquakes. As both heating and strain drives carbonaceous matter towards higher structural ordering, then the evolution presented in Figure 1 may equally be the product of intense shear heating and large amounts of finite strain.

Finally, our experimental results highlight the importance of performing experiments at high P and T to mimick natural processes at depth. In this way the effect of strain only, which becomes significant at confining pressures of 150 MPa, is the highest in experiments at 1GPa and is not visible in friction experiments carried out at a few MPa of normal stresses, can be determined.

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# **Table captions**

Table 1: Characterization of chemistry and degree of crystallinity of the carbonaceous material of the starting materials (18NOB01A – HN78).

- Table 2: Summary of the RSCM parameters measured for all static heating experiments for the non-metamorphic (18NOB01A) and very-low grade (HN78) starting materials. SD stands for standard deviation. In the range 320-330°C, both geothermometers were extrapolated.
- Table 3: Details of the deformation experiments, resulting microstructures and Raman characteristics.
- Table 4: Parameters obtained from the fitting procedure of R1 evolution for static heating experiments on 18NOB01A and HN78. Time t is in minutes and temperature T is in °C.
- Table 5: Summary of CM properties in natural and experimental deformation zones. Note the difference in the treatment of Raman spectra, in particular the ratio of Disordered (D band) to Graphite band (G band): R1 is calculated from the ratio of the intensities of the raw spectrum (see Figure 4), while ID1/ID2 stands for the ratio of the intensities of the function. D1 and D2 used to deconvolute the raw spectrum. Peak area ratio R2 is similarly defined in Beyssaz et al. (2002) as the area ratio of the D band over G band, based on the 4-bands deconvolution, adapted to the graphite-like, highly-ordered carbonaceous matter. For the poorly-ordered carbonaceous matter within the carbonization stage, the increase in crystallographic ordering is reflected carbonaceous matter within the graphitization stage, the increase in crystallog aphic ordering is reflected in a decrease in both intensity or area ratio of the D band over the G band

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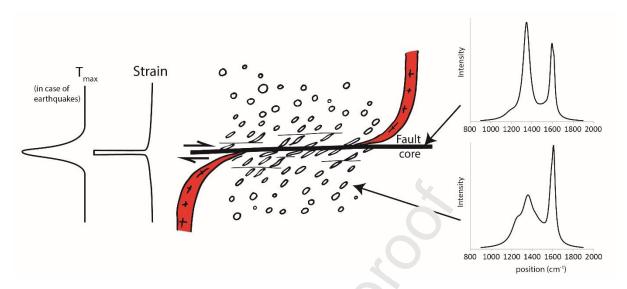


Figure 1: Raman spectra of carbonaceous particles show anomalies, corresponding to higher crystallinity, in the core of fault zones (for example Mor's-1, futtoni et al., 2022). We are considering here conditions of deformation and metamorphism ser aining to the brittle field, up to brittle-ductile transition, hence regional temperatures in the range 50-350°C approximately. As illustrated on the left plots, fault cores are zones of both high struin and possibly high Tmax, i.e. maximum experienced temperature (if the fault slipped during ear you kes). While the role of temperature on Raman is, at least qualitatively, well established, the role of train is not known, and constitutes the object of the present study.

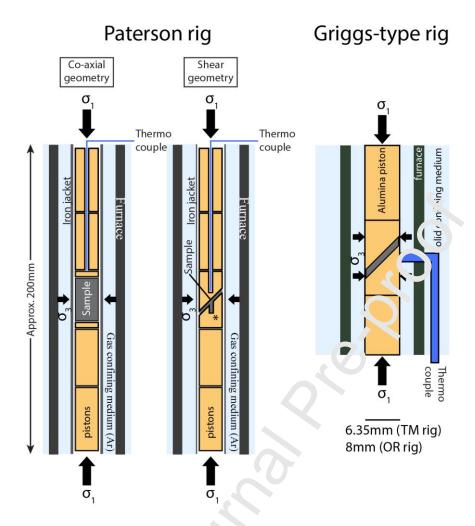


Figure 2: Deformation experiment setup in the Paterson and Griggs-type rigs, with gas and solid confining media, respertively. In shear geometry with the Paterson rig, a cylinder of sample material (shown as \*) is added to the 15°-cut sample to serve as an undeformed reference.

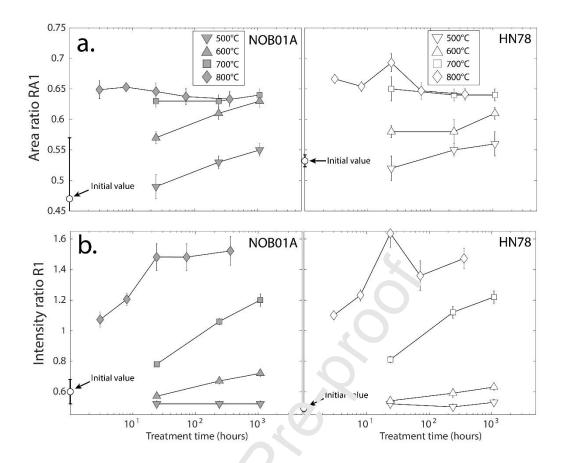


Figure 3: Evolution of the RSCM parameters carough the carbonization stage for 18NOB01A and HN78. **a.** RA1 area ratio (Lahfid et al., 2013). **b.** Intensity Ratio R1. The error bar is the standard deviation associated with the 20 CM partial s/s/spectra analyzed on each experimental sample.

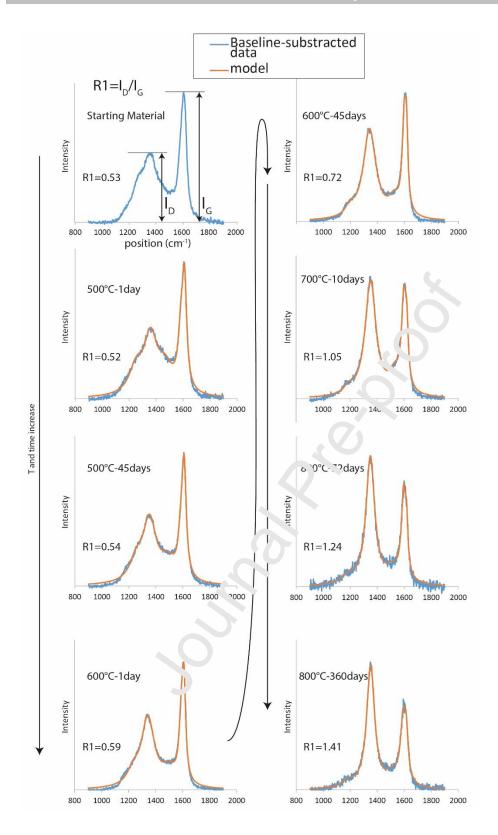


Figure 4: Representative spectra (baseline-substracted and modelled as a result of peak-fitting procedure) showing the evolution with increased duration and temperature of static heating, for sample 18NOB01A. Intensities of the peaks vary between ~1000 and ~5500 counts between the different samples.

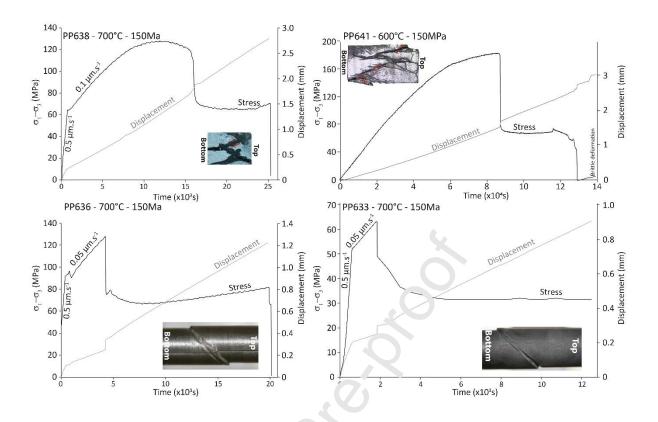


Figure 5: Paterson rig experiments. Differe, tal stress  $\sigma_1$  -  $\sigma_3$  and axial displacement curves as a function of time for coaxial (PP638 and PP641) and shear (PP633 and PP636) deformation experiments. Deformed samples photographs of axial sections for PP638 and PP641 and to the external jacket for PP633 and PP335 In all photographs  $\sigma_1$  is horizontal and  $\sigma_3$  vertical.

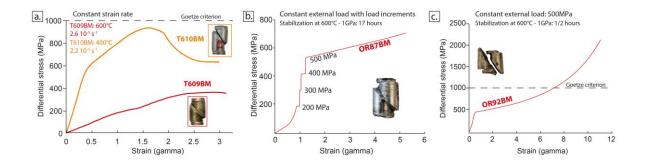


Figure 6: Griggs-type rig experiments. (a) Constant strain rate experiments, at 400 and 600°C, (b) and (c), constant load experiments at 600°C. In (b), the sample was left for 17 hours at 600°C-1GPa before external loading with incremental steps from 100 to 500MPa. In (c), the sample was left for 1/2 hour at 600°C-1GPa before external loading at 500MPa. All experiments at 1GPa confining pressure. In constant load experiments, the actual differential stress on the samp. The recalculated a posteriori as the stressed surface varies as a function of strain.

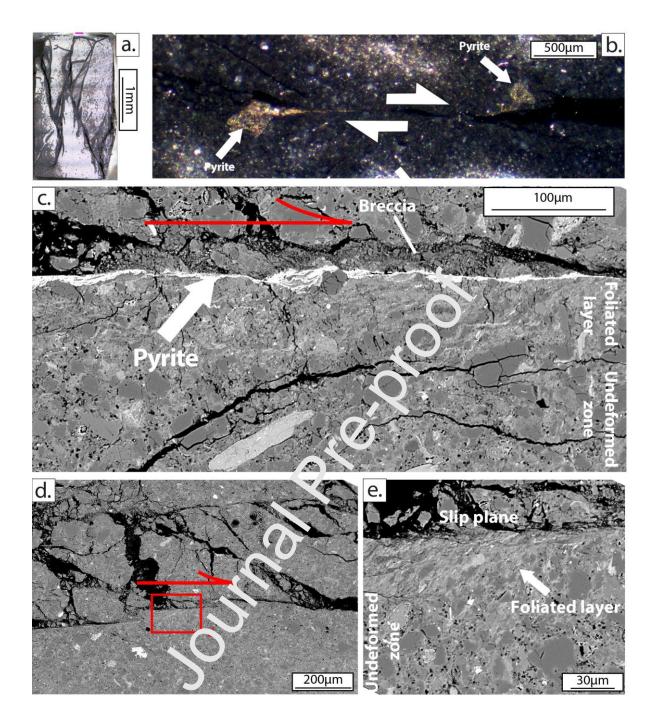


Figure 7: Deformation microstructures observed for the Paterson Rig experimental sample PP641 deformed in coaxial geometry. **a:** cross-section of the whole sample, showing the conjugate fracture planes. **b.** Offset of a pyrite grain by a slip plane. **c-d-e:** The slip zone is bordered by a foliated layer, less than 50m-thick. In c, the slip zone is underlined by a bright layer of pyrite originating from the large pyrite grain and one side of the slip plane is composed of a breccia, the other one of a foliated zone. Note in c and e the decrease in porosity in the foliated zone compared to undeformed host-rock or to breccia. e is a close-up view of d. a: photograph, b: optical microscopy in reflected light, c to e: SEM-BSE imaging.

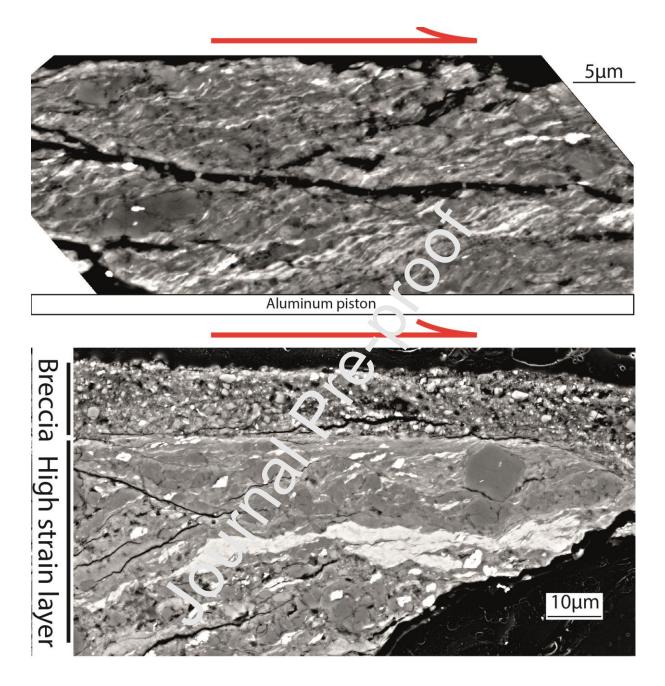


Figure 8: High-strain layers in the vicinity of the main slip planes (shown by red arrows), in OR92BM (top) and TM610BM (bottom). These layers contain plyllosilicates defining a foliation at low-angle to the main slip plane. The grain size is of the order of the micron or lower, and the porosity is strongly reduced. Note in the bottom picture the juxtaposition of the high strain layer with a thin layer of breccia showing submicron-sized clasts and a much larger porosity. All images from SEM-BSE imaging.

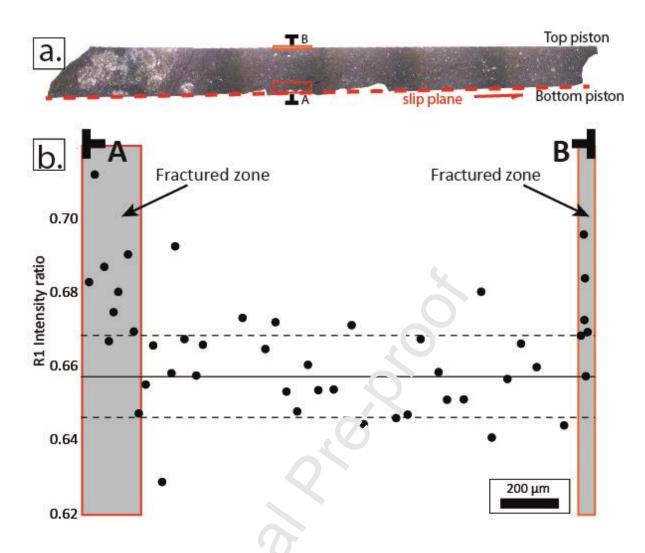


Figure 9: RSCM results obtained for the sample PP633. **a.** Optical microscopy image in reflected light of sample PP633 showing the in calcon of the RSCM profile and the two deformed zones. Slip was localized on the lower piston-sal calculated be boundary (dashed red line). **b.** Intensity ratio profile across the sample.

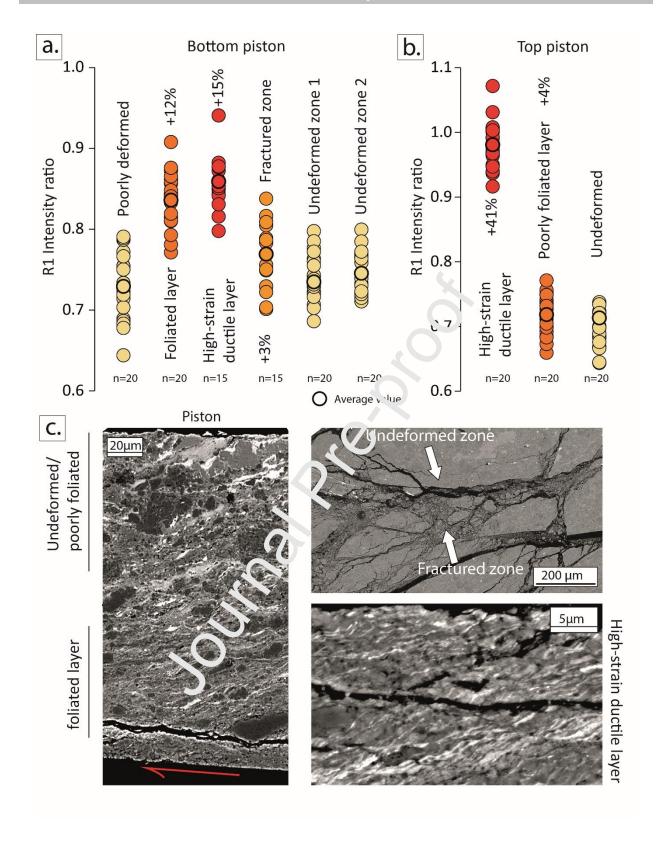


Figure 10: RSCM results obtained for the sample OR92BM. **a. and b.** Diagram representing the R1 evolution in the different microstructural zones defined. A systematic increase in R1 is observed as a function of strain (the domains analyzed are in the vicinity of bottom piston (a) or top piston (b)) **c.** BSE imaging of the different microstructures.

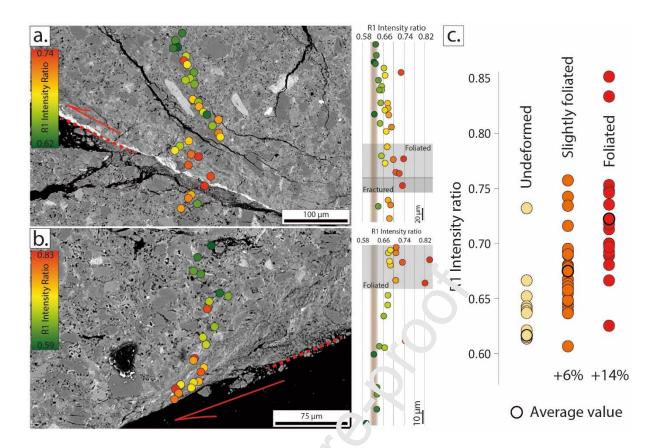


Figure 11: RSCM results obtained for the sample PP641. **a. and b.** high-resolution R1 profiles obtained along the strained zones developed during the experiment. The slip surfaces are shown by a dashed red line, and the sense of shear is increated by the arrows. **c.** R1 diagram by zones showing the evolution of the R1 in the undeformed, slightly foliated and foliated zones.

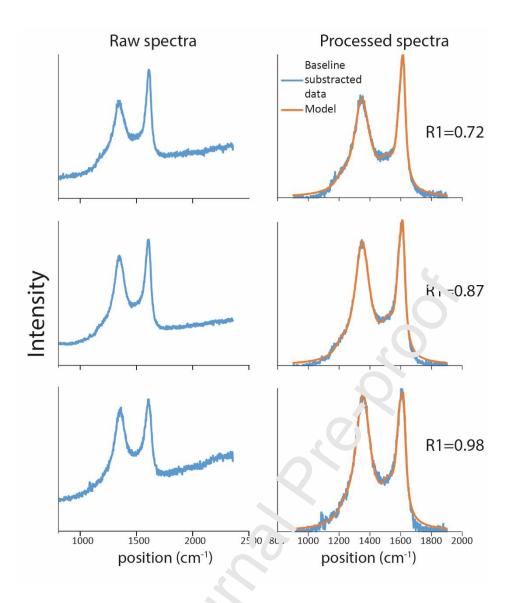


Figure 12: Representative RSCM spectra obtained for the sample OR92BM. Left: raw spectra; Right: Baseline-substracted spectra, and their modelling as a result of peak-fitting procedure. R1 calculated on the modelled spectra shows large variations as a function of strain (Figure 10). The CM particles analyzed are located through a section across an undeformed (top), foliated (middle) and intensely foliated (bottom) zones.

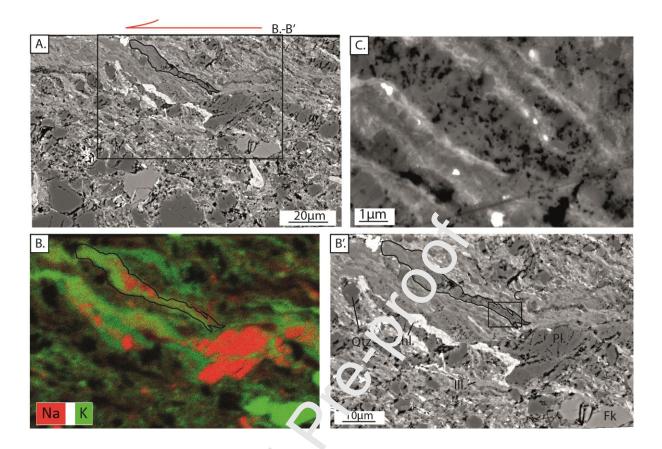


Figure 13: A. Large-scale view showing the elongation of the different mineral phases in a high strain layer adjacent to the slip plane. Note a'so the increase in nanoporosity in the quartz and feldspar aggregates in the high-strain layer B.-D'. While away from the high strain layer, K-feldspar and Plagioclase have composition of pure endmember minerals, in the high-strain-zone, the comminuted feldspar have a composition in volving both Na and K. C. Close-up view of the comminuted feldspar, where the very large poros ty due to fragmentation is visible. A, B' and C: BSE images, B EDS map for Na and K. Sample PP6-1. In B', Qtz, Pl, Fk, Ill and Chl stand for Quartz, Plagioclase, K-feldspar, Illite and Chlorite, respectively.

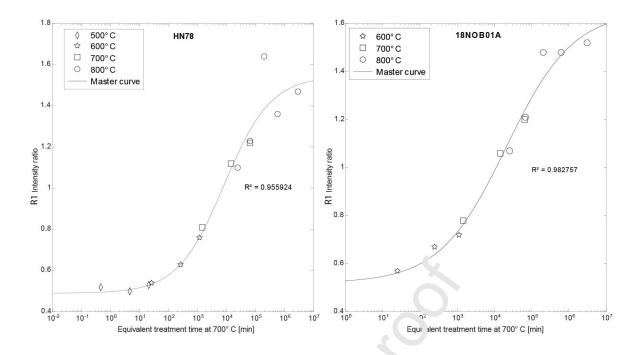


Figure 14: Master curves resulting for the superposition movel for 18NOB01A (a) and HN78 (b), following (Nakamura et al., 2017, 2020)

Table 1

	Raman s	spectrosc	ору								Vitrinite Reflecta		Rock Eval pyrol yse
Sample	D1 band (cm <sup>-1</sup> )	D1 band FW HM	G band (cm <sup>-</sup>	G ban d FW HM	R2 ratio	RA1 ratio	R1 Intens ity Ratio	Width ratio	D/G dist. (cm <sup>-1</sup> )	Estimat ed temper ature (°C) [a]	Rm%	Estimat ed temper ature (°C) [b]	Tmax
18NO B01A HN78 [a] Lahfid	1358 $\pm 7$ 1350 • $\pm 2$ d et al., 20	105 ± 13 116 ± 6 10; [b] Ba	1610 ± 7 1603 ± 3 arker,	69 ± 18 52 ± 5	$0.48 \pm 0.09$ $0.53 \pm 0.01$	0.47 ± 0.10 0.53 ±0.01	0.60 ± 0.08 0.49 ± 0.02	1.66 ± 0.62 2.25 ±0.14	252 ± 7 254 ± 2	- 201 ± 11	0.54 ± 0.04	84 ± 7°C	437°C -
1988		, , ,											

**************************************	Loss on ignitio	5:02	T:O2	Al2	FοO	MaO	MnO	CaO	Na.`	W20	P205	503
wt%	n	SiO2	TiO2	03	FeO	MgO	MnO	CaO	<u> </u>	K2O	P2O5	SO3
18NO B01A	4.36	64.08	0.72	16.7 3	5.82	1.9	0.04	0.29		3.26	0.23	0.17
HN78	8.38	58.77	0.66	14.8 7	4.81	2.06	0.07	4.67	1.68	2.83	0.15	0.61

Table 2

Sa mp le	Te mp era tur e (°C )	T i m e ( h o u rs )		S D	D 1 b a n d	נ	G b 1 c m - 1	S D	b a n d F W H M	S D	R 2 r a t i o	S D	R A 1 r a t i o	S D	R 1 In te ns it y R at io	S D	W i d t h r a ti o	S D	D / G d i s t	S D	G ss / D 2 di st. (c m -1)	S D	Es ti m at ed te m p. [a J[ b]	S D
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	0	4	3	3	1	4	6	0		6					0.				2	3	2	2		
			6		1		0		6		4	0	4	0	5	0	8	2	4		5.			
			0	8	4	8	8	7	1	2	9	2	9	2	2	2	7	2	8	6	7	6	-	-
	50	2	1	1	1	6	1	0		1	0	0	0	0	0.	0	2	0	2	1	2	0		
	0	4	3		1		6		5						5				5		3.			
	[1]	[	5	4	0	1	0	4	4	4	5	0	5	0	0	0	0	1	2	9	5	9	-	-

	1	5				7				0	2	0	2		2	3	2						
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50 0	1 0 8	1 3 4	2	1	5	1 6 0	0	4 9	3	0 . 5	0 0	0 . 5	0 . 0	0. 5	0 0	2 4	0 . 0	2	2	1 9.	3	22	1 3
60 0	0 2 4	9 1 3 4	0	8 1 2	1	8 1 6 0	5 0 .	4	6	8 0 6	1 0 . 0	5 0 5	1 0 . 0	2 0. 5	2 0 . 0	2 2 5	9 0 . 1	0 2 6	5 0 .	7 1 9.	1	24	6 9
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[1] wet: addition of 2 wt% H2O; [2] wet: addition of 10 wt% H2O [a] Lahfid et al., 201; [b] Beyssac et al., 2002

Table 3

Ex pe ri m en ts	Name	D ef or m at io n m od e	Pi st o n s	D ur ati	Te m pe ra	Co nfi ng ry su re	Typotmaterial	Final height (mm) ? r axia experiments, final hickness (mm	Initia I assembly diameter (mm)	Di sp la ce m en t ra te	To tali sp la ce m en t	S ti c k - sl i p	S s · d is p l.	Stressdrop	Peakdiffeential stess (beforestick-slip if occured)	Co ndi tio ns of po re flu id	De for ma tio n mi cro str uct ur es	RamanResults
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								) fo r sh ea r ex pe ri m en ts				4					
Pa te rs on Ri g	P P 6 2 9	Sh ea r	Porousalumina	5 ho ur s	70 0 C	√5 ¬ M Pa	c. o w d e r	0. 8	1 5	0. 1 μ m. s <sup>-1</sup>	1. 4 m m	N o		3 2 0 M P a	Po re fluid connected with water, ke pt at a low pressure	Compaction of the same of the	N o ne

																th e alu mi na pis to ns, wh ich loc ali ze d th e def or ma tio n)	
P P 6 3 0	Sh ea r	P or o u s al u m in a	6 ho ur s	70 0° C	15 0 M Pa	Polder	1. 6	1 5	0. 1 μ m. s <sup>-1</sup>	3. 5 m m	N o	-	-	3 8 0 M P a	Po re flui d co nn ect ed wit h wa ter , ke pt at a lo w	Co mp act ion onl y (de hy dr ati on of cla y ma de th e sa	N o ne

		m in a					-0						a		ess ur e	pis to n	t ro ck , 0. 71 9 ± 0. 06 4 al o ng th e sli p su rf ac e (+ 4 %) 0.
P P 6 3 3	Sh ea r	Non-porous alumina	4 ho ur s	70 0° C	15 0 M Pa	Poyder	1. 8	1 5	0. 05 μ m. s <sup>-1</sup>	1. 0 m m	Y e s	4 5 μ m	3 0 M P a	6 4 M P a	No co ntr ol of po re flui d pr ess ur e	Br ec cia zo ne s (~2 00 µ m-thi ck, wit h he ter og en eo us gra	0. 65 6 +/ - 0. 01 2 in th e h os t ro ck , 0. 67 3 +/ - 0. 01 5 (math of the color

								in comminution, large posity various distributed with the sample + Vitre ous displane Cold not	ax va lu e) , i.e . + .6 % in crea se in br ec ci at ed la ye r al o ng th e sli p pl an e
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																be ob ser ve d with SE M) alo ng pis to n	0.
P P 6 3 6	Sh ea	Non-porousalumina, roughened contact	6 ho ur s	70 0° C	15 0 M Pa	Powder	2. 1	1 5	0. 05 μ m. s <sup>-1</sup>	1. 6 m m	Y e s	9 O μ m	5 O M P a	1 2 8 M P a	No control of pore fluid pressure	High strain lay er (~1	0. 78 4 +/ - 0. 05 in th e 1 μ m - th ic k hi gh st ra in la ye r, 0. 72 6+ /- 0. 3 in ~2 0

pl a n e e ( )
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																s wh er e th e po ros ity is de cre as ed	0.
P P 6 4 1	Co - ax ial	Non-poous aum in a	42 ho ur s	6, 0° C	15 0 M Pa	l n t a c t	31	1 5	0. 01 5 μ m. s <sup>-1</sup>	3. 0 m m	Y e s	1 1 0 µ m	1 3 5 M P a	1 8 1 M P a	No control of pore fluid pressure	Conjugat e sh ear zo ne s (~ 0 m thi ck) + sli p pla ne	62 4 +/ - 0. 01 7 in th e u n de fo r m ed zo ne , 0. 79 +/ - 0. 05 1 in th e sh ea r

															zo ne : +1 4 % in cr ea se in th e sh ea r zo ne
P P 6 4 9	Co - ax ial	Non-porous alumina	72 ho ur s	6 5 0	15 /) M Pa	l n t a c t	29	1 5	Cont n n n n n n n n n n n n n n n n n n	2. 3 m m	N o	-	No control of pore fluid pressure	Ho me ne ou s sh ort eni	IR 0. 62 2+ /- 0. 02 5 in th e w h ol e sa m pl e

Ex pe ri m en ts	N a m e	D ef or m at io n m od e	A p p ar at u s	Ty pe of di sp la ce m en t	Ti m e	Te m pe ra tu re	T y p e o f m a t e ri a l	Pr es su re	P e a k st r e s s	Ap pli ed no mi na l lo ad	Di sp la ce m en t	S tr ai n r a t e	G a m m a	P t- j a c k e t r u p t u r e	L o ca liz e d d ef .	De for ma tio n mi cro str uct ur es	Ra ma n Re sul ts
So lid med iu m Grig gs ty pe pa pa ratu s	T 6 0 9 B M	Sh ea r	T ul lis - m o di fi e d	Co ns ta nt sti n ra te	34 ho 'ır s	60 0° C	l n t a c t	10 00 M Pa	3 5 0 M P a		1. 5 m m	2 · 5 7 E 1 0 5 s 1	3 . 1	N o	Yes	Mi cro br ec cia with inc rea se d porosity, 20 to 50 μ m thick, along	IR 0.8 31+ /- 0.6 7; No diff ere nce bet we en und efor me d and bre ccia ted zon e; larg e gra die nts in IR bet we

							Q								th e sli p pla ne. No visi ble foli ate d sh ear zo ne	en upp er and low er pist ons: 0.8 04+ /- 0.0 82 for top part vs. 1.0 13+ /- 0.0 75 for bot tom part
T 6 1 0 B M	Sh ea r	T ul lis - m o di fi e d	Co ns ta nt str ai n ra te	72 ho ur s	40 0° C	l n t a c t	10 00 M Pa	9 5 0 M P a	ı	2. 3 m m	2 . 2 4 E 1 0 5 S 1	3 . 0	Y e s	Yes	Mi cro - br ec cia , wit h inc rea se d po ros ity, alo ng th	IR 0.5 86+ /- 0.0 37 No diff ere nce bet we en und efor me d, bre ccia ted and she ar zon e

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														vel op ed in br oa d zo ne s ar ou nd th e sli p pla ne.	
O R 8 7 B M	Sh ea r	N e w g e n er at io n	Co ns ta nt lo ad	5 ho ur s	60 0° C	l n t a c t	10 00 M Pa	20 0, 30 0, 40 0 an d 50 0 M Pa	3. 1 m m	9 · 9 E 1 · 0 · s · 1 (f o r 5 0 0 M P a )	5 . 3	N o	Y es	S-C du ctil e str uct ur es + a ma in sli p pla ne is pr es en t; mo	IR 0.7 19+ /- 0.0 35 in the und efor me d zon e, IR 0.8 58 +/- 0.0 58 in the hig h stra in zon

										stl y brecialo ng si p pla e + fe w μ m thick strongly he are done to ng he e si p pla ne	e: +19 % incr eas e in the she ar zon es	
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	O R 9 2 B N	Sh ea r	N e w g e n er at io n	Co ns ta nt lo ad	20 ho ur s	60 0° C	l n t a c t	10 / Ni Pa		50 0 M pa	8. 1 m m	5 · 0 E 1 0 4 s 1	1 6	Yes	Y es	A mains in plane, companie d by she are zone s (~100 μ m thick) + strongly she are zone γ 0μ m thick, ver y	IR 0.6 92+ /- 0.0 28 in the und efor me d zon e, IR 0.9 79 +/- 0.0 36 in the high stra in zon e: +41 % incr eas e	
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Table 4

	HN78	18NOB01A
$R1_{min}$	0.49	0.52
R1 <sub>max</sub>	1.5450	1.6523
h	0.5407	0.4847
m	-1.6733 x 10^4	-2.1392 x 10^4
In A	-15.791	-21.339

Table 5

Refe renc e	Type of occu rren ce	Defo rme d mat erial	Orga nic matt er anal yzed	Type of siructure of deform ation	Initia I matu rity of orga nic matt er	Range of organ ic matte r evolu tion (in terms of Rama n signal	Metho d of observ ation	Evolu tion of organ ic matte r prope rties in strain zones (quali tative )	Evolutio n of organic matter properti es in strain zones (quantita tive)	Delta crystal lograp hic orderi ng in strain zone	Effec t of local heat pulse
Sav age et al. 201 4	Natu ral	Shal es	Orga nic com pou nds in the	Kodiak pseudo tachyly te, Alaska, US	220- 270° C (Rajic et al., 2023	Carbo nizati on	Gas chrom atogra phy- mass spectro scopy on extract	-	Increase in the proporti on of diamond oids/n- alkanes (=increas ed	Positiv e	Infer red by the auth ors

							ed organic molecu les		maturati on)		
Sav age et al. 201 9	Natu ral	Sand ston es	All orga nic com pou nds in the rock	Punchb owl strike- slip fault, US	Less than 120° C of maxi mum temp eratu re of buria I (Polis sar et al. 2011 )	Carbo nizati on	Gas chrom atogra phy- mass spectro scopy on extract ed o gailic mulacu les	300	Methylp henanthr ene isomers ratio MPI-4 increase in principal slip zone	Positiv e	Infer red by the auth ors
Mui rhe ad et al. 202	Natu ral	shal es and phyll ites	CM parti cles	Fault zones in variour sedime ntary terral 9	Varia ble, from lov. gr, de to g. ee nschi st-facie s conditions	Carbo nizati on	RSCM		Inconsist ent evolutio n: either increase or decrease in R1	Non syste matic	possi ble
Ked ar et al. 202 0	Natu ral	marl s	CM parti cles	Large- scale folding	sub- gree nschi st facie s	Carbo nizati on to graph itizati on	RSCM		Decrease in R1 in the overturn ed limb of a fold	Positiv e	Not infer red by the auth ors; the sole effec t of defor mati on is invok

											ed
Ito et al. 201 7	Natu ral	shal es	CM parti cles	Pseudo tachyly te in the Shiman to Belt, Japan (Mugi mélang	T~16 0°C (Rai mbo urg et al. 2019	Carbo nizati on	RSCM	<u></u>	Increase in ID1/ID2	Positiv e	Infer red by the auth ors
Kan eki et al., 201 6	Natu ral	Turb idite s	CM parti cles extr acte d from the rock	Pseudo tachyly te in the Shiman to Belt, Japan (Nonok awa fromation)	~210 - 230° C (Rai m, n ai, e a'. 2019	Carbo nızati on	RSCM	intens ity of IR signal of C-H and arom atic C-C signal in defor med zone comp ared to host rock	Slight Increase in ID1/ID2, although the spectra are noisy	Positiv e	Infer red by the auth ors
Ujiie et al. 202	Natu ral	Pela gic mud ston es	CM parti cles	pseudo tachyly te and ultraca taclasit e	~220 ℃	Carbo nizati on	RSCM		Increase in ID1/ID2	Positiv e	Infer red by the auth ors
Suc hy et al. 199	Natu ral	lime ston es, marl s and grey wack	Vitri nite	Greens chist- facies shear zones	250- 350° C (RSC M Girau It et al.	Graph itizati on	Optical micros copy + vitrinit e reflect ance	Graph itizati on of vitrini te partic les in plasti	-	Positiv e	Infer red by the auth ors

		es			2020			c strain zones			
Petr occi a et al. 202 2	Natu ral	Quar tzite s and phyll ites	CM parti cles	Shear zones in the Varisca n Belt in Sardini a, Italy	420- 520° C	Graph itizati on	RSCM		Decrease in R2 from undefor med to the core of the shear zone	Positiv e	Infer red by the auth ors
Nak amu ra et al. 201 5	Natu ral	Bioti te- mus covit e schis ts	CM parti cles	Catacla sites, ultraca taclasit es and pseudo tachyly tes in the Hidaka Metam orphic Belt, Japan	340- 600° C	Graph itizati on	RSCR1 + LaRTER1	300	Increase in R2 in zones of brittle deforma tion	Negati ve	Not infer red by the auth ors; the sole effec t of defor mati on is invok ed

Ref ere nce	Type of occu rrenc e	Defor med mate rial	Org anic mat ter anal yzed	Type of struct ure of defor matio n	Initi al mat urity of orga nic mat ter	Rang e of organ ic matt er evolu tion (in term s of Rama n signal )	Metho d of observ ation	Evolu tion of orga nic matt er prop ertie s in strai n zone s (qual itativ e)	Evolutio n of organic matter properti es in strain zones (quantit ative)	Delta matu ratio n in strai n zone	Efectofoce – heat pulse	Exper imen tal condi tions
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Sav age et al. 201 8	High veloc ity fricti on expe rime nts (~m/s)	Woo dford Shale , from centr al Okla hom a	Org anic com pou nds in the rock	Princi pal slip zone of ~200µ m of thickn ess	Rela tivel y imm atur e	Carb oniza tion	Gas chrom atogra phy- mass spectr oscopy on extract ed organi c molec ules	?	methylp henanth rene isomers ratio MPI-4 increase (=increa se in maturity ) in principal slip zone, as a function of work density and power density	Posit ive	ye s	5 to 10M Pa norm al stress , water - satur ated condi tions at room temp eratu re
Kiril ova et al. 201 7	Medi um veloc ity fricti on expe rime nts (~1 to 100 µm/s )	Com pacte d and heat ed grap hite powd er	Gra phit e parti cles	Princi pal slip zone, comp acted near the slip surfac e, with higher porosi ty away from it	Crephit E Dow der	Grap hitiza tion	RSCM	?	Increase in R2 in zones of brittle deforma tion	Nega tive	?	5 to 25M Pa norm al stress , room temp eratu re and room humi dity
Kan eki and Hir ono , 201 9	High veloc ity fricti on expe rime nts (~m/s)	CM powd er	CM parti cles	Homo geneo us defor matio n, witho ut description of the micros	Vari able , fro m low grad e CM to grap hite	Varia ble	RSCM + HRTE M + IR	?	Increase in R1 for low to interme diate maturity CM, decrase in R1 for graphite	Posit ive in low to inter medi ate matu rity CM, nega	ye s	norm al stress es of 0.5, 1.0, 1.5 and 2.0 MPa, slip rate

				tructu res of defor matio n			30			tive in grap hite		of 1 m/s and slip dista nce of 15 m unde r water - satur ated condi tions at room temp eratu re
Ito et al. 201 7	High veloc ity fricti on expe rime nts (~m/s)	Natu ral shale s	CM parti cles	Grain comm inutio n and meltin g within a pseu otal. Vlyte	Low - grad f (T'1) 67°C	Carb Sniza tion	RSCM	?	Increase in ID1/ID2	Posit ive	ye s	2.7 to 13.3 MPa norm al stress , room temp eratu re and humi dity
Fur uic hi et al. 201 5	High (~m/s) and low (0.15 mm/s) veloc ity fricti on experime nts	Hemi pelag ic muds tone from the Nank ai Trou gh artifi cially mixe d with low	CM parti cles	kelati vely thick foliate d zone in wet tests, princi pal slip zone in dry tests at low and high velocit	Very low grad e - mod ern sedi men ts at max ~1k mbs f	Carb oniza tion	RSCM	?	Increase in ID1/ID2 in high velocity friction experim ents, no change in low velocity friction experim ents	Posit ive	ye s	Dry and wet tests were cond ucted at a slip rate of 1.3 m/s and a norm al stress (on)

		grad e brow n coal from the Kum ano Basin		Y								of 2.0 MPa with displa ceme nt of ~11 m. In additi on, one dry test was perfo rmed at a low slip rate of 0.15 mm/s and low $\sigma_n$ of 1.0M Pa with displa ceme nt of ~0.3 m
Aub ry et al. 201 8	Tri- axial defo rmat ion test, with stick- slip even ts on a fract ure plan e	West erly grani te with a saw- cut fract ure plane	Carb one coat ing on the fract ure plan e	Fault plane with molte n surfac e	è	Carb oniza tion	RSCM	ŗ	Increase in R1 in strained samples	Posit ive	ye s	Roo m temp eratu re, confi ning press ure of 45, 90 and 180M Pa, displa ceme nt rate

												of 3μm/ s
Bus tin et al. 199 5	Grigg s- type HP HT shea r and coaxi al expe rime nts	Anthr acite coal from Penn sylva nia and a high volati le bitu mino us coal from Nova Scoti a.	ide m	?	Low grad e	Carb oniza tion then graph itizati on	TEM+X RD+Ra ma 1	XRD and TEM show an evolution stranned samp les towards a grap hite-like structure	Increase in R1 in strained samples	Posit ive	n o	Confining press ures of 0.6, 0.8 or 1.0 GPa, temp eratures between 400° C and 900° C, strain rates of 10° 5/s or 10° 6/s
Oo has hi et al. 201 1	Rota ry shea r expe rime nts, 50 mm/ s to 1.3 m/s	Powd ers of com merci al amor phou s carbo n	ide m	Princi pal slip zone of ~50µ m thickn ess	Amo rpho us	-	XRD+T EM	XRD and TEM show an evolu tion in strai ned samp les towa rds a grap hite-like struc ture, only in	?	Posit ive	ye s	Norm al stress es of 0.5 to 2.8 MPa, equiv alent slip rates (Ve) of 50 mm/s to 1.3 m/s, and displa ceme nts of 0.09

								high- veloc ity and unde r N2 atmo sphe re expe rime nts				to 118 m. Atmo spher e of N2 or of air
Kuo et al., 201 7	Rota ry shea r expe rime nts, 0.00 03 m/s (subs eismi c) and 3 m/s (seis mic)	Fault brecc ia from the Long mens han fault zone	CM parti cles	Princi pal slip zon of '200 t t'.ickn ess	T~3 50°C (fro m t'in ak se nof D2, see Kou kets ou et al. 201 4)	Grap hitiza tion	Raman +TEM	TEM show s the cryst alliza tion of grap hite parti cles only in expe rime nts at fast veloc ity and with out adde d wate r	Decreas e in R1, for slow and fast slip, irrespec tive of water saturati on level	Posit	ye s	Norm al stress es of 8.5 MPa, slip rates of 0,3 mm/s, and displa ceme nts of 3 m, at room temp eratu re and humi dity and, by the additi on of 0.5 g of distill ed water , at water - damp ened

												condi tions
Kan eki and Hir ono , 201 8	Rota ry shea r expe rime nts, slip rate of 0.00 1 m/s + subs eque nt heati ng unde r N2 or vacu um	Mixt ure of quart z and lignit e powd er	CM parti cles	?	Imm atur e type -I.' koro	Ca b niza tion	RSCM + IR	Decr ease in aliph atic and arom atic C-C bond s as a resul t of shear only	No deciphe rable Raman spectru m as a result of shear only	Posit	?	Roo m humi dity, equiv alent slip rate of 0.001 m.s <sup>-1</sup> (Hiro se & Shim amot o, 2005) , 10m of total displa ceme nt at norm al stress es of 1 and 3 MPa unde r air condition.

#### **Author Contribution**

Benjamin Moris-Muttoni: Conceptualization, Investigation, Writing - original draft, Acquisition,

Analysis and interpretation of Raman spectra

Hugues Raimbourg: Conceptualization, Writing - original draft, Supervision;

Rémi Champallier: Experiments

Romain Augier: Interpretation of Raman spectra

Abdeltif Lahfid: Interpretation of Raman spectra

Emmanuel Le Trong: Numerical modelling

Ida Di Carlo: SEM imaging

#### **Declaration of interests**

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

Hugues Raimbourg reports financial support was provided by French National Research Agency.

#### Highlights

- Using Raman we studied carbonaceous matter (CM) evolution with strain and temperature
- Kinetics of evolution were derived from high-pressure experiments of static heating
- High-pressure deformation experiments show the large effect of strain on Raman of CM
- Raman anomalies in natural fault zones can result from strain and not temperature