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## Quartz rheology constrained from constant-load experiments: Consequences for the strength of the continental crust

Subhajit Ghosh<sup>a,\*</sup>, Holger Stünitz<sup>a,b</sup>, Hugues Raimbourg<sup>a</sup>, Jacques Précigout<sup>a</sup>

<sup>a</sup> Institut des Sciences de La Terre d'Orléans (ISTO) - Université d'Orléans, CNRS, BRM, UR7327, 45071, Orléans, France  
<sup>b</sup> Department of Geology, University of Tromsø, Bransveien 201, 9001, Tromsø, Norway

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## ABSTRACT

The mechanical properties of quartz are fundamental to control the plastic behaviour of the continental crust. Our understanding of quartz rheology is still limited in the following respects: i) the large variability of flow law parameters in the earlier literature (stress exponent  $n = 4$  to  $\leq 2$  and activation energy  $Q = 120$  to  $242$  kJ/mol), and ii) the difficulty to identify the rate-limiting deformation mechanism, if several mechanisms are operating simultaneously. These two issues are connected and cannot be resolved separately. The present study has carried out constant-load experiments to constrain the flow law parameters of quartz. A new generation hydraulically-driven Griggs-type apparatus has been employed, resulting in reproducible mechanical data, even at very low strain rates ( $10^{-8}$  to  $10^{-9}$  s $^{-1}$ ; so far, closest to the natural ones). Furthermore, the  $Q$ -value in constant load experiments can be estimated without prior knowledge of the  $n$  value. Our new  $n$  ( $= 2$ ) and  $Q$  values ( $= 110$  kJ/mol) are fairly low. We calculated an  $A$ -value of  $1.56 \times 10^{-9}$  /MPa/sec. Microstructural analysis suggests that the bulk sample strain in our experiments is achieved by crystal plasticity, i.e., dislocation glide with minor recovery by sub-grain rotation, accompanied by grain boundary migration. Micro-cracking helps to nucleate new grains. It is inferred that strain incompatibilities induced by dislocation glide are accommodated by grain boundary processes, including dissolution-precipitation creep and grain boundary sliding. These grain boundary processes are responsible for the  $n$ -value that is lower than expected for dislocation creep ( $n \geq 3$ ). The new flow law can consistently estimate strain rates (especially at low stresses) in excellent agreement with documented natural case studies and predicts a rapid drop in strength of quartz-bearing rocks in the continental crust below a depth of  $\sim 10$  km or at a temperature of  $\sim 300^\circ\text{C}$  and higher.

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## 1. Introduction

Quartz is one of the most important rock-forming minerals of the lithosphere. Due to its abundance, quartz rheology (i.e., polycrystalline quartz aggregate) in the presence of water forms the basis for constructing crustal strength profiles that are crucial for constraining plate tectonic processes and modelling the long-term flow behaviour of the continental crust (e.g., Kohlstedt et al., 1995). Therefore, several experimental studies in the last four decades (e.g., K&T84  $\sim$  Kronenberg and Tullis, 1984; Koch et al., 1989; P&L90  $\sim$  Paterson and Luan, 1990; L&P92  $\sim$  Luan and Paterson, 1992; G&T95  $\sim$  Gleason and Tullis, 1995; Hirth et al., 2001; S&T03  $\sim$  Stipp and Tullis, 2003; Stipp et al., 2006; R&B04  $\sim$  Rutter and Brodie, 2004; Kidder et al., 2016; Fukuda et al., 2018) focused on

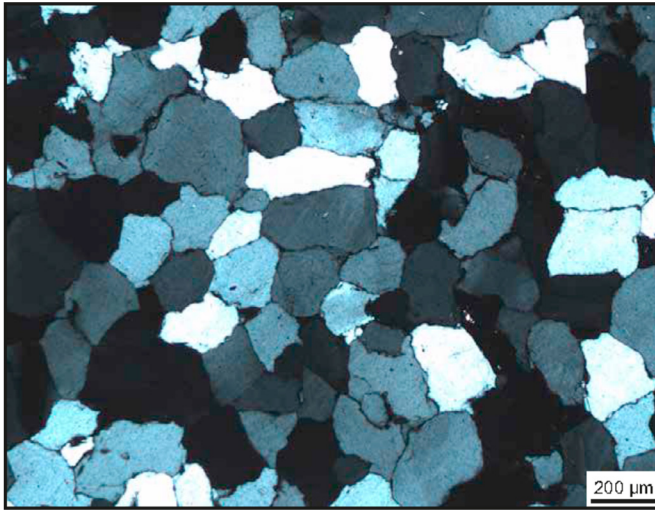
establishing a creep flow law of polycrystalline quartz aggregate (e.g., natural and synthetic quartzites, powders of natural single crystals and sand) based on the following equation:

$$\dot{\epsilon} = A \left( \frac{\sigma^n}{d^p} \right) \exp \left( - \frac{Q}{RT} \right), \quad (1)$$

where,  $\dot{\epsilon}$  is the strain rate,  $A$  is dimensionless constant,  $Q$  is the activation energy,  $R$  is the gas constant,  $T$  is absolute temperature,  $\sigma$  is the differential stress,  $d$  is the grain size,  $n$  is stress exponent and  $p$  is the grain size exponent. Theoretical models of creep have shown different values of  $n$  and  $p$  for different creep mechanisms. For example,  $n = 1$  and  $p = 2$  or  $3$  indicate lattice diffusion (Nabarro-Herring) creep (Nabarro, 1948; Herring, 1950) or grain boundary diffusion (Coble) creep (Coble, 1956), respectively. Conversely,  $n \geq 3$  and  $p = 0$  generally correspond to the GSI dislocation creep process. In addition, Kohlstedt et al. (1995) introduced a water fugacity term ( $f_{\text{H}_2\text{O}}$ ) in the equation (1) to account for the effect of  $\text{H}_2\text{O}$  on the creep rates at a certain pressure-temperature

\* Corresponding author.

E-mail address: subhajit.ghosh@univ-orleans.fr (S. Ghosh).



**Fig. 1.** Cross-polarized light-microphotograph of the representative undeformed Tana quartzite showing mosaic-like microstructure without any evidence of internal deformation like undulatory extinction.

(P-T) condition. Accordingly, the equation (1) for the dislocation creep can be written as follows, where  $r$  is water fugacity exponent:

$$\dot{\epsilon} = A\sigma^n f_{H_2O}^r \exp\left(-\frac{Q}{RT}\right). \quad (2)$$

Earlier experimentally determined creep law parameters of polycrystalline quartz by different groups show many discrepancies (Table 1) that result in significant variations of predicted crustal strengths (Kohlstedt et al., 1995; Stipp et al., 2002b). The  $n$ -value vary from 4 to  $<2$ , while the  $Q$  value vary from 120 to 242 kJ/mol, in addition to large differences in the  $A$ -value. Due to the power-law nature of equation (2), minor changes in flow law parameters can lead to significant differences in the predicted crustal strength when extrapolated to geological conditions (Hirth et al., 2001; Lusk et al., 2021; Tokle et al., 2019). In several cases the frequently used experimentally-derived flow laws can overestimate the strength of a particular region. For example, Boutonnet et al. (2013) showed that the  $\dot{\epsilon}$  calculated by any of the earlier flow laws from stresses derived from S&T03 piezometer (without the Holyoke and Kronenberg, 2010 correction) are slower than the well-constrained natural rates in the Ailao Shan–Red River (ASRR) strike-slip shear zone.

The starting material for experiments varied considerably among the earlier studies (e.g., novaculite, different varieties of quartzites, single-crystal or quartz powder to synthetic silica gel/acid), and the initial grain sizes showed more than one order of magnitude difference (Table 1). The finest grain size samples are expected to show a contribution of grain size sensitive (GSS) diffusion creep as is found in polycrystalline mineral aggregates of olivine, or clinopyroxene (Yabe et al., 2020; Ghosh et al., 2021). Importantly, different interpretations for the grain-scale deformation mechanisms, from climb-controlled dislocation creep to a combination of dislocation and diffusion creep for quartz were postulated among different studies (G&T95; Richter et al., 2018; Fukuda et al., 2018; Nègre et al., 2021). In other words, it still remains difficult to constrain the intra- and inter-grain processes responsible for plastic deformation and related flow law parameters.

In order to constrain a robust flow law for wet quartzite, we have generated novel high-resolution data through constant-load experiments at high pressure and high temperature. We used a new generation Griggs-type apparatus with natural coarse-grained ( $\sim 200 \mu\text{m}$ ) polycrystalline Tana quartzite (Fig. 1) to perform mechanical testing by load-stepping. This method has two main ad-

vantages compared to strain rate ( $\dot{\epsilon}$ ) stepping tests: (i) The equilibrium between load,  $\dot{\epsilon}$ , and microstructure is achieved faster (Kohlstedt, personal communication), and (ii) The  $Q$ -value can be determined without previous knowledge of the  $n$ -value. The second point is particularly important for the extrapolation of laboratory data to lower temperature conditions in naturally deformed rocks. The wide range of  $Q$ -values in the existing literature have triggered the work by Lu and Jiang (2019), which attributed such variations to an effect of confining pressure. Another advantage of load-stepping is that the bulk strain of the sample is smaller than in  $\dot{\epsilon}$ -stepping tests, so that geometrical problems due to change in the sample shape are minimized. We compare our results with  $\dot{\epsilon}$ -stepping experiments using the same Tana quartzite to understand the effect of much larger final strain, hence, significant recrystallization (Pongrac et al., 2022). A comparison between our newly established flow law with previous results provides the opportunity to further explore the effects of using different apparatuses, choice of the sample, and the deformation mechanism on the mechanical properties of wet quartzite. Finally, we will discuss the implication of our flow law on crustal strength.

## 2. Methodology

### 2.1. Deformation tests

Load-stepping experiments during coaxial shortening were performed on natural quartzite samples in a new generation Griggs-type solid-medium (NaCl) apparatus (Fig. 2a) at ISTO (Orléans, France, Précigout et al., 2018). Cylindrical samples of 8 mm diameter were core-drilled from the starting material and dried at  $110^\circ\text{C}$  for one day.  $\dot{\epsilon}$  and  $\sigma$  for each step were obtained from the steady-state portion of the mechanical curves (see supplementary text S1).

The hit point (Fig. 2b) in the solid-medium apparatus is a major source of error in stress measurements (Holyoke and Kronenberg, 2010). When stopping the displacement of the deformation actuator, the load can decrease below the hit point (supplementary text S1). In order to account properly for such a load decrease (affecting the stress at slow displacement rates), the displacement was stopped overnight after the hit point and before the first load increment. This decrease is labelled as ‘relaxation’ in Figs. S1 and S2. The stress correction by Holyoke and Kronenberg (2010) was not applied to the data, because the use of solid medium salt involves a subtraction of 48 MPa, resulting in zero or negative  $\Delta\sigma$  values for some of our low stress experiments. We also estimated the magnitude of the axial stresses related to different displacement rates of the  $\sigma_1$  piston (during our load-stepping) due to viscous drag and/or friction with the confining medium (see supplementary text for details). These values are two orders of magnitude lower than the error estimated for the hit point construction, and therefore, inconsequential for our mechanical analysis.

### 2.2. Starting material and sample characterization

As the starting material, we used a pure ( $>99\%$  silica), natural quartzite without evidence of prior micro-scale deformation (Fig. 1) from the ELKEM quarry in Austertana (Northern Norway). The mean grain size is  $\sim 200 \mu\text{m}$  and the  $\text{H}_2\text{O}$  has been measured as of the order of  $\sim 1700 \text{ H}/10^6 \text{ Si}$  for grain interiors and  $\sim 2100 \text{ H}/10^6 \text{ Si}$  for grain boundaries in a previous study (Nègre et al., 2021). Further details on chemical and microstructural characterizations of the starting material can be found in Nègre et al. (2021) and Pongrac et al. (2022). Deformed samples were cut through the centre, parallel to the compression direction, and then characterized using Polarized light microscopy (LM), SEM-cathodoluminescence (CL) and Electron backscatter diffraction (EBSD) (see supplementary text S1).

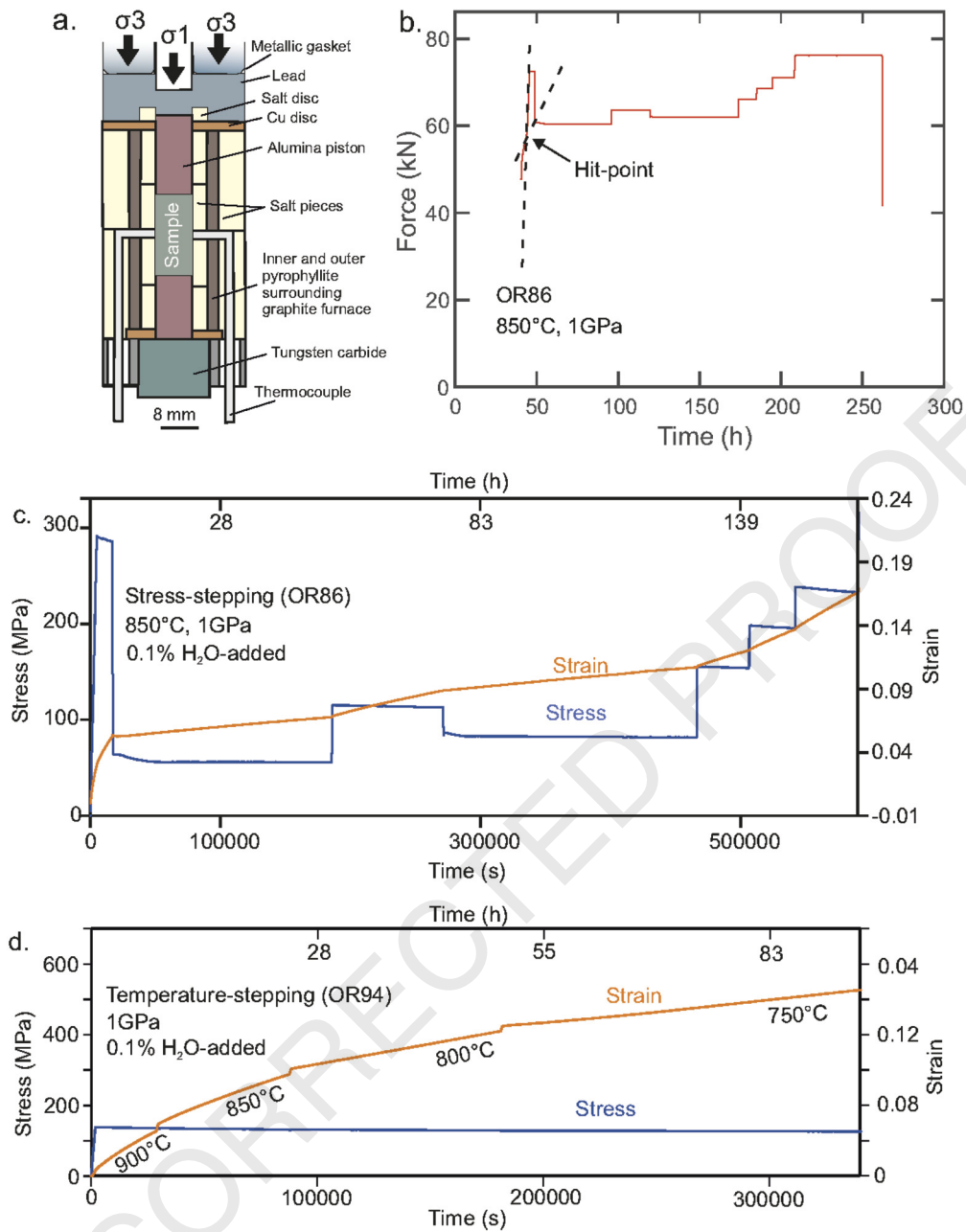
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**Table 1**  
Previously determined flow law parameters and the experimental conditions (also see Table S1).

Source	Experiment apparatus	Sample characterization	Deformation condition ( $T$ °C and $P_c$ GPa)	Initial grain size ( $\mu\text{m}$ )	$A$ ( $\text{MPa}^{-n} \text{s}^{-1}$ )	Activation energy $Q$ ( $\text{kJ mol}^{-1}$ )	Stress exponent $n$	Grain size exponent $p$	Water fugacity exponent $m$
Kronenberg and Tullis (1984)	Solid salt medium	Heavitree quartzite: As-is (4000 – 5000 H/10 <sup>6</sup> Si $\approx$ 0.05 wt.%).	650 to 1200; 0.90 to 1.17	211	-	170 – 220	2.9 – 3.2	-	-
	Solid salt medium	Heavitree quartzite: 0.4 wt.% water added.	600 to 900; 1.53 to 1.6	211	-	120 $\pm$ 10	2.7 $\pm$ 0.3	0.18	-
	Solid salt medium	Heavitree quartzite: vacuum dried at 1000-800 °C for 12 hrs.	919 to 1119; 0.95-1.45	211	-	300	4	-	-
Jaoul et al. (1984)	Solid salt medium	Arkansans novaculite: As-is.	1019 to 1269; 1.45	1.2	-	210 $\pm$ 10	-	-	-
	Solid salt medium	Arkansans novaculite: 0.4 wt.% water added.	600 to 950; 0.89 to 1.60	3.6 – 4.9	-	120 – 150	2.5 $\pm$ 0.07	0.18	-
	Solid salt medium	Heavitree quartzite: As-is	900 to 650, 1.5	211	$3.11 \times 10^{-4}$	171 $\pm$ 8 (CaCO <sub>3</sub> )	2.3 $\pm$ 0.1	-	-
	Solid salt medium	Heavitree quartzite: 0.39 wt.% water added.	800, 1.5	211	$2.91 \times 10^{-3}$	150.6	1.8 $\pm$ 0.2	-	-
Koch et al. (1989)	Solid salt medium	Heavitree quartzite: vacuum dried at 1010 °C for 5.5 hrs.	890, 1.5	211	-	-	3.3 $\pm$ 0.4 (CaCO <sub>3</sub> )	-	-
	Solid dehydrated talc	Simpson quartzite: As-is (642 $\pm$ 47 H /10 <sup>9</sup> Si $\approx$ 0.009 wt.%).	750 to 900; 1 to 1.25	210	$1.16 \times 10^{-7}$	134 $\pm$ 32	2.72 $\pm$ 0.19	-	-
	Solid hydrated talc	Simpson quartzite: H <sub>2</sub> O from wet talc.	750 to 900; 1 to 1.25	210	$5.05 \times 10^{-6}$	145 $\pm$ 17	2.61 $\pm$ 0.15	-	-
Paterson and Luan (1990)	-	-	-	-	$6.5 \times 10^{-8}$	135	3	-	-
Luan and Paterson (1992)	Gas medium	Silica acid (relatively pure): less than 0.4 wt.% water.	827 to 1027; 0.3	20 (~10 – 30)	$4.0 \times 10^{-10}$ (Modified to $10^{-7.20 \pm 0.52}$ )	152 $\pm$ 71 (Modified to 121 $\pm$ 13)	4 $\pm$ 0.8 (Modified to 3)	-	-
	Gas medium	Silica gel (impure): at least 0.1 – 0.2 wt.% water.	977 to 1027; 0.3	90 (30 – 95)	-	150	2.3 $\pm$ 0.3 (2.01 – 2.6)	-	-
Gleason and Tullis (1995)	Molten salt medium	Black Hills quartzite: As-is (~0.15 wt.%).	900 to 1100; ~1.5 to 1.7	100	$1.1 \times 10^{-4}$	223 $\pm$ 56	4 $\pm$ 0.9 (Modified to 3.6)	-	-
Hirth et al. (2001)	-	Previous Experiment + natural observation	-	-	$6.3 \times 10^{-12}$	135 $\pm$ 14	4	-	1
Post et al. (1996)	Solid salt medium	Black Hills quartzite: 0.2 – 0.1 wt.% added.	900; 0.7 to 1.7	100	-	-	-	-	>2 (using $n = 3.9$ )
Rutter and Brodie (2004)	Gas medium	Crushed Brazilian single crystal quartz: low water content (~10 to 20 H/10 <sup>6</sup> Si).	1000 to 1200; 0.3	12 – 20	$1.2 \times 10^{-5}$	242 $\pm$ 24	2.97 $\pm$ 0.29	-	-
Fukuda et al. (2018)	Solid salt medium	Natural quartz powder: 3,500 ppm H/Si $\approx$ 0.25 wt.% of adsorbed water.	600 to 750; 1.50	9.5 – 12	-	129 $\pm$ 33	2.9 to 5.2	-	1.0 $\pm$ 0.2
	Solid salt medium	-	800 to 950; 1.50	9.5 – 12	$1.0 \times 10^{-2.97}$	183 $\pm$ 25	1.7 $\pm$ 0.2	0.51 $\pm$ 0.13	1.0 $\pm$ 0.2
Richter et al. (2018)	Non-coaxial Solid salt medium	Crushed Alpine cleft quartz: 0.2 wt.% H <sub>2</sub> O added.	800 to 1000; 1.5	< 100	$3.1 \times 10^{-4}$	~170	1.8 – 2	~1.08	-
Nègre et al. (2021)	Solid salt medium	Tana quartzite: 0.1 wt.% added.	900; 0.6 to 2	~200	-	-	~2	-	~1

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**Fig. 2.** Mechanical data and the configuration of the experiment. a) Schematic diagram of the experimental set-up in the new generation Griggs-type deformation apparatus. b) A typical load-stepping mechanical force versus time curve to constrain the hit point. c) A typical stress-strain-time data obtained during stress-stepping experiment at 850 °C, 1 GPa (OR86). d) A typical stress-strain-time data obtained during temperature-stepping (900 to 750 °C) at constant load (OR94).

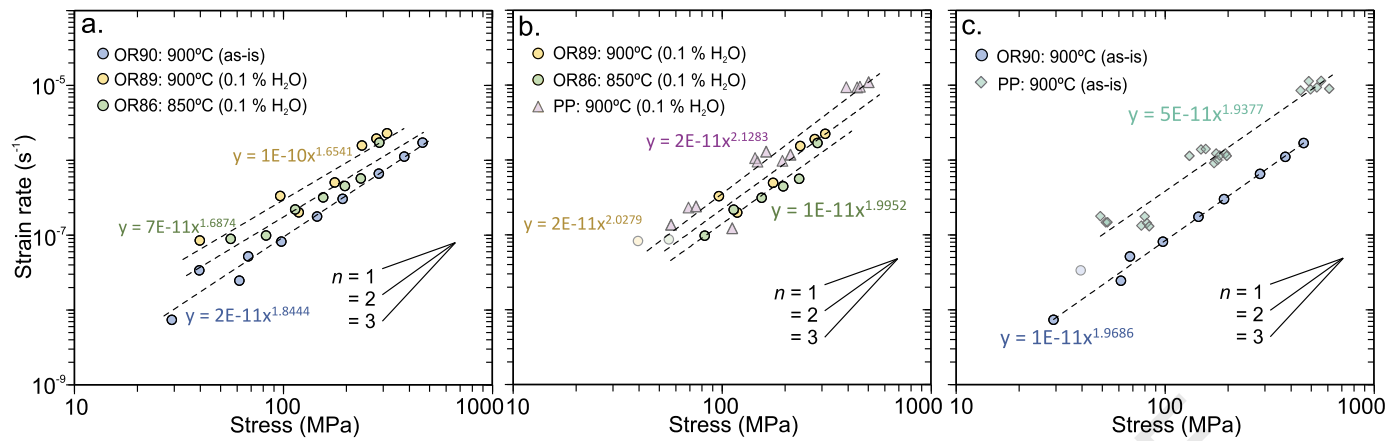
### 3. Experimental results

#### 3.1. Mechanical data

In Fig. 2b and 2c, we provide a typical stepped-load test (with 0.1 wt.% of H<sub>2</sub>O added) at a constant temperature of 900 °C and a confining pressure of 1 GPa. As expected, strain increases faster per time with increasing load on the sample. We see a rapid change in strain measurements during the rapid change in the applied  $\sigma$  (i.e., load/area) from one stress-step to another. The slopes of the strain-time curves need some period of time to attain a steady state condition (Fig. S1). We only use the last part of the strain-time curve, where steady-state has been achieved. In our load-stepping experiments, we have used the conventional hit point as well as the lowest point after relaxation and have carried out the mechan-

ical data analysis with both values as hit point (Figs. S1-S2, 2d). There is no systematic effect on the  $n$  or  $Q$ -values.

The curves in figure S1 show similar characteristics for both as-is and H<sub>2</sub>O-added experiments, although the as-is sample is ~2–5 times stronger than the H<sub>2</sub>O-added sample at 900 °C (Fig. 3a). Depending on the number of steps and  $\dot{\epsilon}$  at each step, the bulk strain in our experiments ranges up to a maximum value of ~16%. Experimental conditions and results from each stress-step from corresponding experiments are sequentially shown in Tables 2a, 2b. During the as-is experiment (OR90), we repeated the best stress-steps acquired during increasing (39.59 and 67.89 MPa) and then during decreasing load (61.50 and 29.22 MPa) to identify the effect of time and microstructural evolution during the experimental run time. The 67.89 and 61.50 MPa steps show similar creep rates, which are within the error generally given for the Griggs-type instruments (Holyoke and Kronenberg, 2010). However, for



**Fig. 3.** Determination of stress exponent ( $n$ ) in a log-log space. Co-axial strain-rate-stepping experiments using the conventional Griggs apparatus are plotted as PP (Pongrac et al., 2022). a) Differential stresses and strain rates for all our experiments ( $H_2O$ -added and as-is ones) are plotted. The slope of the global fit indicates the  $n$  value. Only the  $H_2O$ -added experimental data are shown. Global-fit is calculated without considering the transparent data points in OR86 (green colour) and OR89 (yellow colour). c) Only the as-is experimental data are shown. Global-fit is calculated without considering the transparent data point in OR90 (blue colour). (For interpretation of the references in this figure(s), the reader is referred to the web version of this article.)

**Table 2a**

Experimental results of stepped load tests.

Experiment no.	$T$ (°C)	$P_c$ (GPa)	$H_2O$ (%)	$H$ (mm)	$\sigma$ (MPa)	$\dot{\epsilon}$ ( $s^{-1}$ )	Final $\epsilon$ (%)
OR86	850	1	0.1	14.86	287.28	1.73E-06	~16.60
					55.90	8.94E-08	
					113.76	2.20E-07	
					82.63	9.89E-08	
					154.84	3.19E-07	
OR89	900	1	0.1	14.64	96.56	3.34E-07	~15
					39.83	8.47E-08	
					237.28	1.57E-06	
					278.59	1.95E-06	
					312.75	2.29E-06	
OR90	900	1	as-is	14.91	39.59	3.39E-08	~6.6
					67.89	5.20E-08	
					97.71	8.21E-08	
					144.84	1.77E-07	
					61.50	2.46E-08	
					29.22	7.40E-09	
					191.98	3.05E-07	
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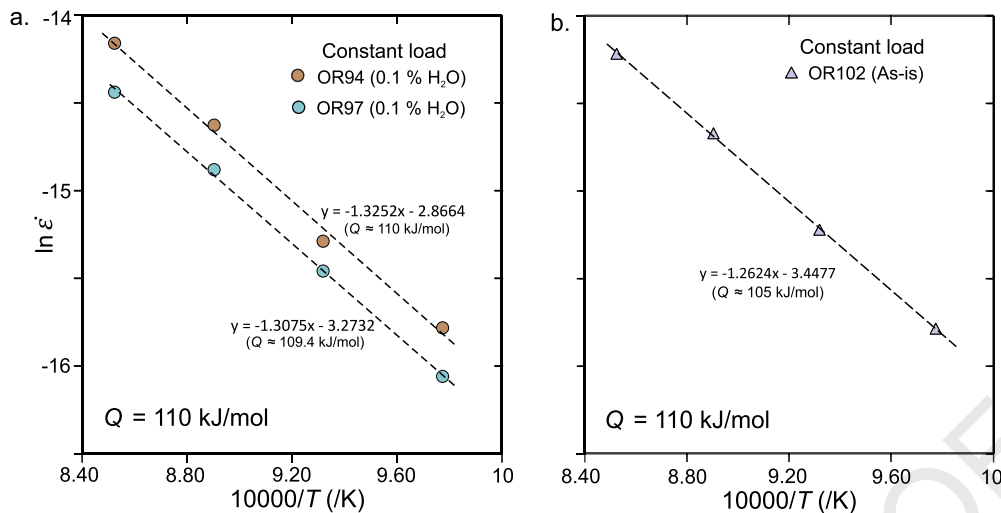
**Table 2b**

Experimental results of stepped heating in constant-load tests.

Experiment no.	$T$ (°C)	$P_c$ (GPa)	$H_2O$ (%)	$H$ (mm)	$\sigma$ (MPa)	$\dot{\epsilon}$ ( $s^{-1}$ )	Final $\epsilon$ (%)
OR94	900	1	0.1	15.02	136.32	7.08E-07	~10.50
	850				132.77	4.44E-07	
	800				130.34	2.29E-07	
	750				126.67	1.40E-07	
OR97	900	1	0.1	15.04	153.51	5.36E-07	~9.80
	850				150.21	3.44E-07	
	800				147.86	1.93E-07	
	750				143.80	1.06E-07	
OR102	900	1	as-is	15.04	146.48	6.69E-07	~10.25
	850				142.71	4.25E-07	
	800				139.21	2.45E-07	
	750				135.76	1.39E-07	

the lowest stress-steps (29.22 and 39.59) the difference in  $\dot{\epsilon}$  is relatively substantial (4.6 times) compared to the difference of  $\sigma$  (1.35 times).

In addition to load-stepping experiments, we have performed three  $T$ -stepping experiments at constant load: two with 0.1 wt.%  $H_2O$  added and one as-is (Table 2b). The mechanical data for the



**Fig. 4.** Arrhenius plots of  $\ln(\dot{\epsilon})$  against inverse of temperature ( $10000/T$ ) for both the  $\text{H}_2\text{O}$ -added (a) and as-is (b) Tana quartzite. The determination of the  $Q$ -value without previous knowledge of the  $n$ -value is a great advantage of the stress-stepping-tests. Our  $Q$ -value of 110 kJ/mol is the lowest  $Q$ -value determined so far from solid-medium piston-cylinder (Table 1). However, it is in the same range of values as the studies by Kronenberg and Tullis (1984) and Paterson and Luan (1990).

two  $\text{H}_2\text{O}$ -added experiments are identical, even the slope of the individual stress-steps have similar patterns (Figs. 2d, S2). At the highest  $T$ -step, the overall slope of the strain curve is slightly convex, while it becomes slightly concave at the lowest  $T$ . The  $\sigma$  values also decreased slightly ( $\sim 10$  MPa) with time as a consequence of the increase in the surface area of the deformed sample (as the load is constant). We used the hit point after relaxing overnight to calculate the  $Q$ -value. Noticeably, all the data points for the OR94 with lower stress-values are  $\sim 0.76$  times faster than the OR97 with slightly higher stress-values, under the exact same condition of deformation (Tables 2a, 2b). Such minor differences in  $\sigma$  and  $\dot{\epsilon}$  values are likely related to the inherent mechanical error, for example, in consistently choosing the hit point. In addition, the oil pressure in the hydraulic-rams is temperature-sensitive, and some variation may be related to small room temperature variations during deformation.

### 3.2. Determination of $n$ and $Q$

$\sigma$  and  $\dot{\epsilon}$  for all the experiments were plotted in a log-log diagram to calculate the  $n$ -value (Fig. 3). The slope of the global-fit indicates a  $n$ -value of  $\sim 1.7$  for  $\text{H}_2\text{O}$ -added and  $\sim 1.85$  for the as-is sample (Fig. 3a). We noticed that the lowest stress points (55.90, 39.83 and 39.59 MPa) measured at the beginning of the load-stepping experiments exert a strong control on the overall global fit. If we do not consider these lowest stress data points, then all the global-fits approach  $n \approx 2$  (Figs. 3b, c). A recent communication (Pongrac et al., 2022) has reported  $n \approx 2$  from co-axial  $\dot{\epsilon}$ -stepping experiments using the same Tana quartzite. These experiments (plotted as PP in Fig. 3b) were performed in a conventional Griggs apparatus located at the University of Tromsø. The samples deformed in Orléans are slightly stronger than samples deformed in Tromsø under the same P-T conditions (Fig. 3b, c).

We plot all of our temperature-stepping data in the Arrhenius  $\ln(\dot{\epsilon}) - T^{-1}$  space (Fig. 4). To calculate the  $Q$ , we used the equation (2) in the following form.

$$\ln(\dot{\epsilon}) = -\left(\frac{10^4}{T} \times \frac{Q}{10^4 R}\right) + \ln(Af_{\text{H}_2\text{O}}^n \sigma^n) \quad (3)$$

In this way, the  $Q$ -value can be calculated independently of the  $n$ -value. The similar slope of the global-fit for both  $\text{H}_2\text{O}$ -added experiments can be well fitted with a single Arrhenius equation with similar values of activation energy (i.e.,  $Q = 110$  kJ/mol) (Fig. 4a).

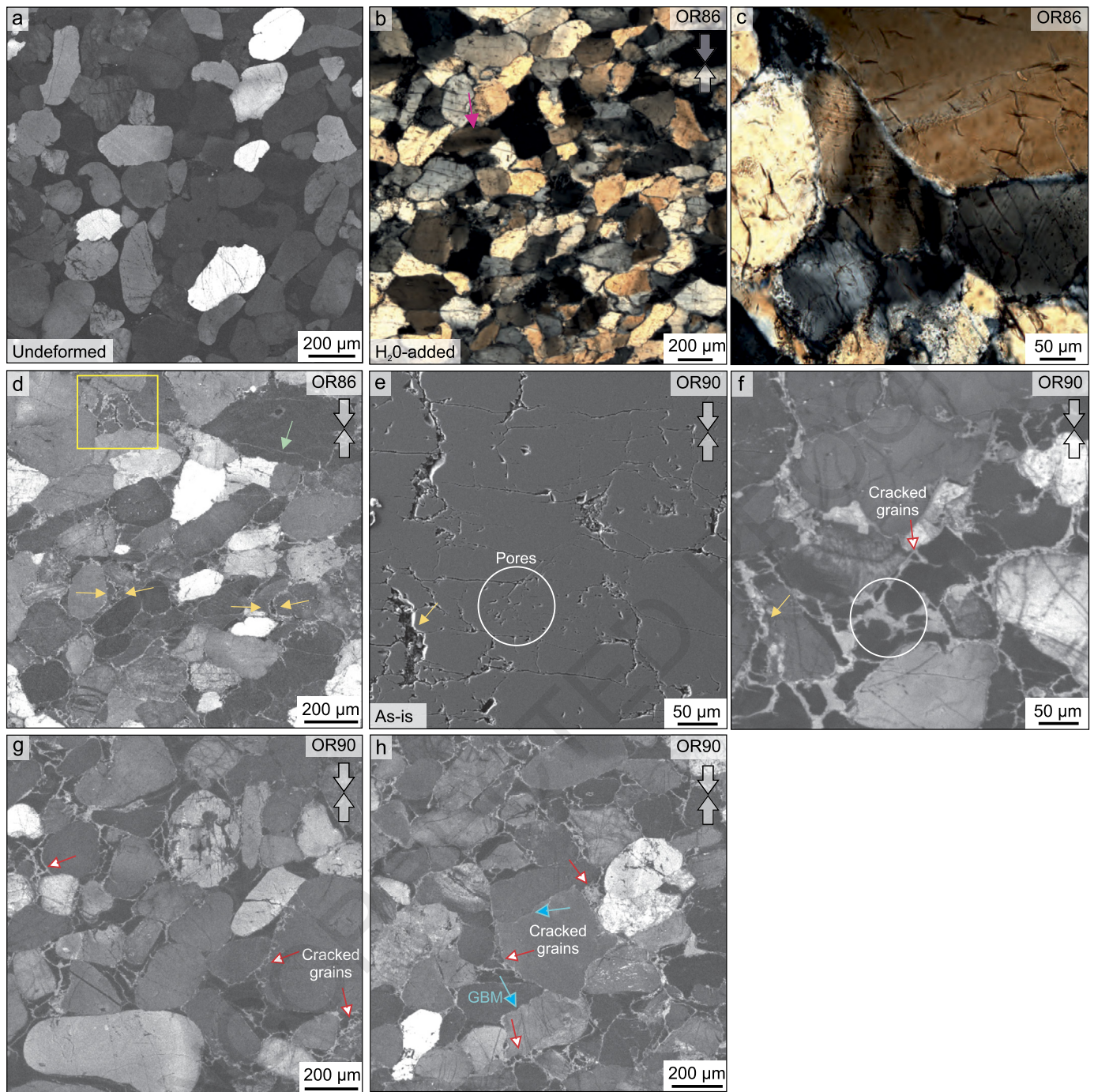
In order to make sure whether the choice of hit point can affect the  $Q$ -value or not, we have calculated  $Q$  using the conventional hit point, too (Fig. S2). There is no difference in the result. But the  $Q$  is slightly ( $\sim 5$  kJ/mol) lower for the as-is sample (Fig. 4b). Considering the inherent errors associated with Griggs-type apparatuses, such minor differences are not significant. Therefore, we consider the  $Q$ -value as the same for the  $\text{H}_2\text{O}$ -added and as-is samples.

### 3.3. Microstructural observations and interpretations

CL-imaging of the undeformed Tana quartzite shows that the individual quartz grains are composed of cores of rounded detrital quartz grains with various colours and intensity, surrounded by a dark authigenic  $\text{SiO}_2$  cement (Fig. 5a). The cement is in crystallographic continuity with the sand grains on which it grows epitaxially (Fig. 1). Due to the filling of the pore space by authigenic  $\text{SiO}_2$  cement, no visible porosity was detected by the LM or SEM. Detailed microstructural characterizations and  $\text{H}_2\text{O}$  content of the deformed (Tana) samples have been presented in two earlier studies (Nègre et al., 2021; Pongrac et al., 2022). Therefore, in the present contribution, we will only summarize the microstructural observations and put the observed grain-scale deformation mechanism into context with our new flow law data.

Under the LM, both, the as-is and  $\text{H}_2\text{O}$ -added samples show evidence of crystal-plastic deformation features, such as undulatory extinction, deformation lamellae, and sub-grains. We can observe some intragranular micro-cracking (Fig. 5c), unrelated to the unloading cracks. Furthermore, a new material with a bright luminescence appears surrounding original quartz grains and in the cracked regions (Figs. 5d-h), indicating the operation of dissolution-precipitation creep (DPC) processes in the presence of fluid. Fluid migration along the grain boundaries and cracks makes them visible in CL-images. Cracked regions typically form along the boundaries of quartz grains (Figs. 5g, h, 6a). Larger, CL-dark fragments can be seen within lighter luminescent material, preserving their angular shape (Fig. 5f, 6a). We notice that these zones are sometimes associated with porosity development (Fig. 5e). In addition, we can identify mode-I cracks (marked by yellow arrow) parallel to the compression direction (Figs. 5e). Such 'axial splitting' cracks typically form at low confining pressures, most likely early in the experimental procedure, before attaining the deformation conditions (Pongrac et al., 2022). These mode-I cracks are



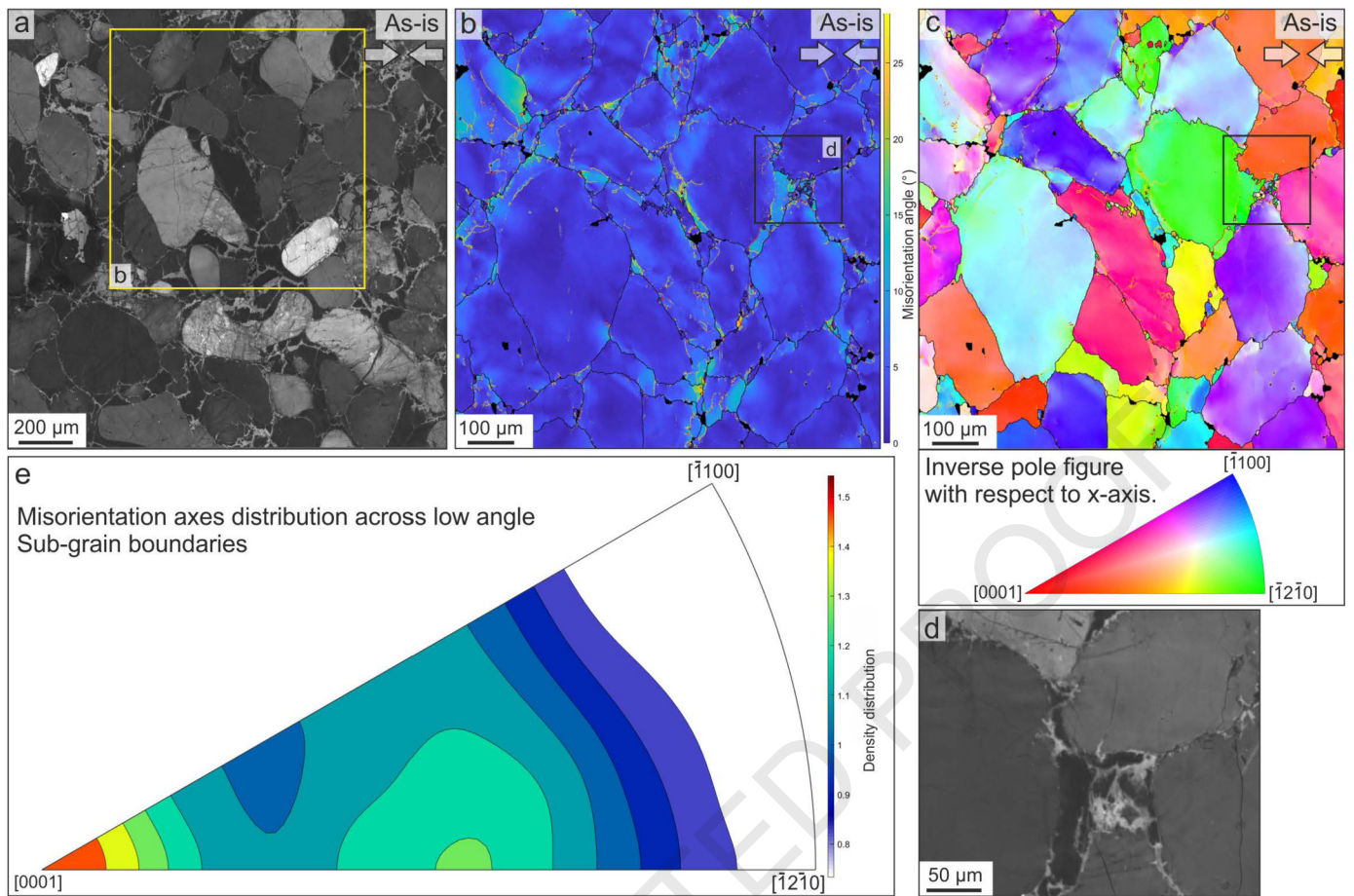


**Fig. 5.** Microstructural observations from deformed Tana quartzite under a light microscope (LM) and scanning electron microscope (SEM). a) SEM-CL-imaging of the undeformed Tana quartzite for comparison with the deformed samples. Individual quartz grains are composed of cores of rounded detrital quartz grains, surrounded by a dark luminescent crystalline authigenic silica cement. b, c) Under the LM, deformed H<sub>2</sub>O-added samples show strong evidence of crystal-plastic deformation features, including undulatory to patchy extinction, deformation lamellae, grain boundary undulations, and subgrain formation. d-h) Compression direction is vertical. In the deformed sample, thin bright luminescence materials are formed along the cracks and boundaries of the quartz grain. Yellow arrows indicate mode-I cracks, while the green arrow indicate unloading cracks. These cracks are healed by silica during the deformation. Such places are associated with porosity generation, as shown in the secondary electron image, marked by a white circle (e). Red arrows show cracked grains, which are formed along the grain boundaries and within the dark cement material. Blue arrows indicate features similar to grain boundary migration (GBM). A careful inspection reveals that these wavy grain boundaries are marked by thin white luminescent material, which indicates the mobility of grain boundaries.

volumetrically small (<1%) and discrete. Typically, all cracks lack any offset. As a result, these cracks hardly accommodate any strain and should not have a substantial effect on the bulk strength of the samples. Lobate grain boundaries, indicative of grain boundary migration (GBM) is also associated with the appearance of the bright luminescence (Fig. 5h).

EBSD mapping reveals some degree of crystal plasticity in the original quartz grains, giving rise to the development of low-angle boundaries (Figs. 6b, c). Such low-angle inner boundaries (marked in yellow) correspond to grains with intense patchy undulatory extinction under the LM. The distorted crystal structures can develop into discrete sub-grain boundaries (red lines) (Fig. 6b). Most of these low-angle boundaries are typically found near the grain





**Fig. 6.** Corresponding CL and EBSD images from the deformed as-is sample (OR90). The compression direction is horizontal. We plot the following three types of boundaries in our EBSD maps: i) high-angle grain boundaries (black lines) defined by misorientation angle  $>10^\circ$ , ii) low-angle boundaries defined by misorientations between 2 to  $10^\circ$  and including sub-grain boundaries (red lines) and inner boundaries (yellow lines). While sub-grain boundaries form closed polygons, the yellow inner boundaries do not. a) The yellow box on the CL image indicates the location of the EBSD mapping. b) Misorientation analysis reveals a high degree of internal (“mis2mean” function in MTEX) crystal plastic deformation in the original quartz grain. c) Individual grains are drawn based on their crystallographic orientation with respect to the compression direction ( $\sim$  East). Crystallographic orientations of grains relative to each other are given by the inverse pole figure scheme. d) The location of the CL image is marked by a black rectangle in the previous figures. The fine-grained materials at the junction of the four original grains might have been identified as the product of dynamic recrystallization. However, the corresponding CL image reveals that this fine-grained material has developed due to cracking in the cement. e) Misorientation axes distribution of low-angle boundaries indicate dominant prism  $\langle a \rangle$  slip with minor rhomb  $\langle a \rangle$  contribution.

boundary regions (Fig. 6b). A few of these low-angle boundaries correspond to micro-cracks in the CL image (Fig. 6a). A comparison between sub-grain boundaries along with the IPF colour map (Fig. 6c) indicates the formation of incipient recrystallized grains along grain boundaries by progressive sub-grain rotation. We report a mean recrystallized grain-size of  $6.23 \mu\text{m}$ , following our EBSD data treatment (see supplementary for procedure). Misorientation axes distribution of low-angle boundaries (sub-grains) suggests that prism  $\langle a \rangle$  slip is statistically dominant with subordinate rhomb  $\langle a \rangle$  slip (Fig. 6e). In addition to sub-grain rotation, a portion of such new grains were originated by micro-cracking (Fig. 5c). The new grains developed by cracking are identified by their bright luminescence and the associated porosity development.

## 4. Discussion

### 4.1. Constitutive equation

We found that the values of  $n = 1.7$  or  $2$  and  $Q = 110 \text{ kJ/mol}$  represent our wet and as-is quartzite mechanical data well (Figs. 2, 3). As mentioned above, the low-stress data (e.g.,  $39.59 \text{ MPa}$  for the as-is experiment in Table 2a) measured at the beginning of the stress-stepping experiments exerts a strong control on the slope of the global-fit that indicates  $n < 2$ . Such low flow stress regimes

with  $\dot{\epsilon}$  on the order of  $10^{-8}$ – $10^{-9} \text{ s}^{-1}$  have not been achieved earlier with coarse-grained quartzite (Table 2a). A few studies like the ones of S&T03, R&B04, and Richter et al. (2018) reported low flow stresses of 30 to 66 MPa, but the associated  $\dot{\epsilon}$  are on the order of  $10^{-7}$  to  $10^{-6} \text{ s}^{-1}$ . For the as-is experiment, we repeated the step at low stress (i.e., 29 MPa), after conducting several steps at higher stresses. Interestingly, this data point conforms with the  $n \approx 2$  global-fit. A  $\pm 30 \text{ MPa}$  error is generally associated with traditional Griggs-type apparatus that may influence the precision of the low-stress data. Jaoul et al. (1984), also discarded the first lowest-stress increment data from their load-stepping experiments because of lower accuracy.

Using the equation (2), the pre-exponential term ( $A$ ) is calculated for all the  $T$ -steps from  $\text{H}_2\text{O}$ -added samples, reported in Table 2b, considering  $n = 2$  and  $f_{\text{H}_2\text{O}}^r$ . The  $f_{\text{H}_2\text{O}}$  is calculated for a known P-T (Tony Wither’s fugacity calculator), assuming an  $a_{\text{H}_2\text{O}}$  of 1 and the water fugacity exponent ( $r$ ) of 1 (Kohlstedt et al., 1995; Fukuda et al., 2018). Thereafter, we take the average of those  $A$ -values, which is  $1.56 \times 10^{-9} / \text{MPa} \cdot \text{sec}$ . The  $A$ -value for the as-is sample will be 2-4 times lower. The modified flow law for the 0.1 wt.%  $\text{H}_2\text{O}$ -added quartzite is as follows:

$$\dot{\epsilon} = 1.56 \times 10^{-9} / \text{MPa} \cdot \text{sec} \cdot \sigma^2 \cdot f_{\text{H}_2\text{O}} \cdot \exp\left(-\frac{110 \text{ kJ/mol}}{RT}\right). \quad (4)$$

Nègre et al. (2021) showed that the water fugacity exponent,  $r$  can be 0.92 for  $n = 2$ . In that case, the  $A$ -value in our flow law will be  $2.83 \times 10^{-9}$  /MPa/sec.

#### 4.2. Comparison with previous studies: non-dimensional plots

After establishing the constitutive equation for the Tana quartzite, we can use it as a reference to compare the original mechanical data from experiments reported in the literature. For this purpose, we normalized all the original mechanical data ( $\sigma$  and  $\dot{\epsilon}$ ) to non-dimensional stress ( $\sigma_0$ ) and strain rate ( $\dot{\epsilon}_0$ ) ratios:

$$\sigma_0 = \frac{\sigma}{\sigma^{*ref}} \quad \text{and} \quad \dot{\epsilon}_0 = \frac{\dot{\epsilon}}{\dot{\epsilon}^{*ref}} \quad (5)$$

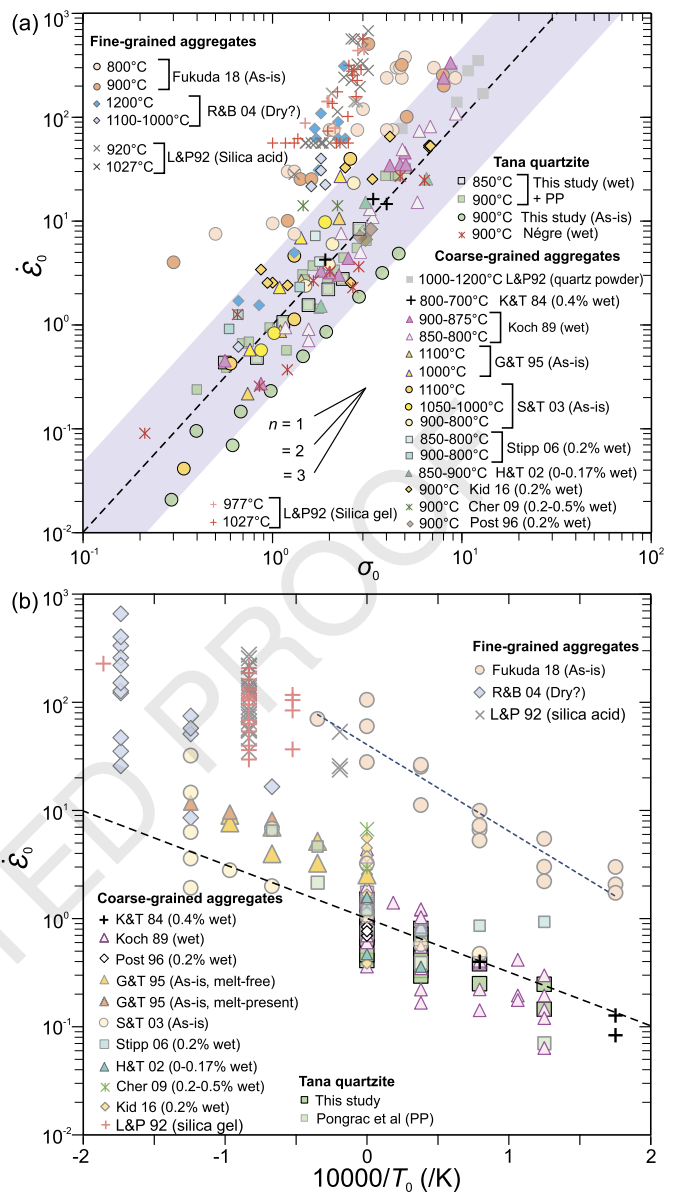
Here, the reference strain rate ( $\dot{\epsilon}^{*ref}$ ) is calculated from our new wet-quartzite flow law (Eq. (4)) by substituting the  $T$  with temperature-values,  $f_{H_2O}$  with those values at which each original mechanical data was acquired and the stress-term ( $\sigma$ ) with the  $\sigma^{*ref}$ . We used a fixed value of 100 MPa as  $\sigma^{*ref}$  in this calculation.

In order to compare the effect of temperature on the creep rates with those given in the existing literature, we created an Arrhenius space of non-dimensional strain rate ( $\dot{\epsilon}_0$ ) and the inverse of temperature ( $T_0^{-1}$ ):

$$\dot{\epsilon}_0 = \frac{\dot{\epsilon}}{\dot{\epsilon}^{*ref}} \quad \text{and} \quad \frac{10000}{T_0} = \frac{10000}{T} - \frac{10000}{T^{*ref}} \quad (6)$$

Here, the reference strain rate ( $\dot{\epsilon}^{*ref}$ ) is calculated from our flow law (Eq. (4)) by substituting the stress-term ( $\sigma$ ) with original reported  $\sigma$ -values and the temperature ( $T$ ) is substituted by  $T^{*ref}$ . We used a fixed value of 1173 K as  $T^{*ref}$ . The  $f_{H_2O}$  is calculated for the  $T^{*ref}$  and the pressure at which each original mechanical data was acquired. It should be noticed that  $T_0^{-1}$  is not a dimensionless ratio in this plot. Previously, the non-dimensional plots were used in comparing mechanical data of olivine and diopside (Yabe et al., 2020; Ghosh et al., 2021). This method differs from using reported flow law parameters to compare results among different studies, where mechanical data were obtained from different stresses, strain rates, temperatures and pressures. To make a rational comparison with our experimental condition, we only included the results from studies that were conducted in a coaxial geometry at  $\geq 750$  °C, and that satisfy the Goetze's criterion ( $\Delta\sigma < P_{conf}$ ).

Our wet-quartzite creep data are transformed to  $\sigma_0$ ,  $\dot{\epsilon}_0$  and  $T_0^{-1}$  plots along the reference line (corresponding to the slope of the stress exponent and the activation energy, respectively) in Fig. 7, which confirms the capability of equation (5) and (6) to express our original mechanical data in a predicted manner. In this way, we are able to compare the mechanical data from the earlier studies to our creep data. All the data show a power-law relationship in a non-dimensional  $\sigma_0 - \dot{\epsilon}_0$  space (Fig. 7a). Clearly, creep results obtained from natural coarse-grained ( $\sim 100$ - $200$   $\mu\text{m}$ ) quartzites broadly overlap with or follow our data. Even the absolute strengths are essentially comparable (within a factor of  $\sim 5$  times the  $\dot{\epsilon}$ , indicated by the shaded region in Fig. 7a), considering a  $\pm 30$  MPa accuracy of the conventional solid-medium apparatus (Holyoke and Kronenberg, 2010). Most of the fine-grained ( $\sim 10$ - $20$   $\mu\text{m}$ ) samples from Fukuda et al. (2018), R&B04, and synthetic samples from L&P92 are weaker compared to the coarse-grained natural samples (except coarse silica gel samples of L&P92). Similar observations are also made for the  $\dot{\epsilon}_0 - T_0^{-1}$  plot in the Arrhenius space (Fig. 7b), where data obtained using natural quartzite overlap with our results. The overlap indicates a comparable effect of temperature on the  $\dot{\epsilon}$  and shows that our low  $Q$ -value (110 kJ/mol; Table 1) is consistent with results of K&T84, Koch et al. (1989), and P&L90. It is surprising to find a lower temperature dependency



**Fig. 7.** Comparisons of our newly developed dislocation creep law for wet quartzite with the results of previous studies in non-dimensional and semi-non-dimensional spaces. To make a rational comparison with our experimental condition, we only included the results from studies that were conducted in a coaxial geometry at  $\geq 750$  °C, and satisfying the Goetze criterion. a) Plot of nondimensionalized strain rate ( $\dot{\epsilon}_0$ ) as a function of nondimensionalized stress ( $\sigma_0$ ), calculated from the original mechanical data ( $\sigma$  and  $\dot{\epsilon}$ ) as reported in the previous literature. b) Nondimensionalized strain rate ( $\dot{\epsilon}_0$ ) as a function of inverse temperature ( $T_0^{-1}$ ). Acronyms used in these plots are as follows. K&T84: Kronenberg and Tullis (1984); Koch 89: Koch et al. (1989); L&P92: Luan and Paterson (1992); Post 96: Post et al. (1996); G&T 95: Gleason and Tullis (1995); H&T 02: Heilbronner and Tullis (2002); S&T03: Stipp and Tullis (?); R&B04: Rutter and Brodie (2004); Stipp 06: Stipp et al. (2006); Cher 09: Chernak et al. (2009); Kid 16: Kidder et al. (2016); Fukuda 18: Fukuda et al. (2018); Nègre: Nègre et al. (2021).

(similar to our data) for the melt-free samples of G&T95, although they reported a higher  $Q$ -value of 223 kJ/mol. Their melt-present samples indicate an even lower slope for a reported  $Q$ -value of 137 kJ/mol (Fig. 7b).

On average the strength of the as-is Tana sample is higher than the 0.1 wt.%  $H_2O$ -added ones (Fig. 7a). This situation is also observed in earlier studies (K&T84; Jaoul et al., 1984). Interestingly, the as-is (G&T95; S&T03) and 0.2 wt.%  $H_2O$ -added (Stipp et al., 2006) data using the Black hills quartzite do not show such discernible effect of  $H_2O$  on the  $\dot{\epsilon}$  with increasing temper-



ature (Fig. 7a). The common thread between these latter studies is that they were performed in a molten-salt-assembly and contrasts with Black hills quartzite data from deformation experiments in a solid-medium apparatus (Kidder et al., 2016). At a lower P-T (900 °C, 1.3 GPa) they plot closely to the above-mentioned studies, possibly due to the retention of added H<sub>2</sub>O (0.2% wt.%). Diffusion of H<sub>2</sub>O from quartz is very slow at an experimental timescale (Kronenberg et al., 1986; Gerretsen et al., 1989), but H<sub>2</sub>O can still be transferred between the grains and grain boundary regions by microcracking. Additionally, the molten salt shows a complete solid solution with H<sub>2</sub>O above 150 MPa pressure (Bodnar et al., 1985), so that the melt in the confining medium is not buffered for H<sub>2</sub>O. Given the ease of hydrogen-diffusion through the Pt jacket, it is possible that H<sub>2</sub>O is effectively lost from the samples in molten salt assemblies at high temperature. Indeed, even a minor amount of melt (present in G&T95) along the grain boundaries might preferentially attract water, and dry the primary grains (Jaoul et al., 1984). The H<sub>2</sub>O content measured by Stipp et al. (2006) from deformed as-is and vacuum-dried specimens show an overall decreasing trend with increasing temperature that may support the conclusion of a drying effect of the molten-salt-assembly. This conclusion may appear inconsistent if several data points from molten-salt-medium experiments are compared with our Tana quartzite data, yielding slightly weaker or overlapping results. However, considering the differences between the apparatus, experimental conditions (e.g., confining pressure), initial sample composition (e.g., the water and impurity content), and normalization method incorporating fugacity estimations, it is apparent that the bulk strengths of those samples are not much different from our results. It emerges from the systematic strength difference that we observe between the 'H<sub>2</sub>O-added' and 'as-is' Tana samples that it is more reasonable to compare the results of as-is and H<sub>2</sub>O-added samples only from molten-salt experiments conducted at the same condition (i.e., 1.5 GPa) in the same apparatus.

It is noteworthy that the earlier  $\dot{\epsilon}$ -stepping experiments (e.g., Nègre et al., 2021; Pongrac et al., 2022) required higher strain to achieve steady-state conditions. Yet, we can express all the previously obtained experimental results from different groups using coarse-grained natural quartzites based on similar  $\dot{\epsilon}$  dependencies on  $\sigma$  and  $T$  as our mechanical data. More significantly, it seems that a larger amount of strain, which should result in more recrystallized material (including different regimes in Hirth and Tullis, 1992) in the earlier studies does not have a great effect on the overall strength of the material, although this needs to be tested in future studies. These observations are indicative of the operation of the same underlying rate-limiting creep mechanism irrespective of the degree of recrystallization (at least in the order at which these studies were conducted). Furthermore, the same  $n$ -values at considerably high total strain in Pongrac et al. (2022) indicate that our  $n$ -values at lower strain are not the result of transient creep.

Contrarily, a number of data from fine-grained samples (especially L&P92; R&B04) are significantly weaker than coarse-grained samples in our non-dimensional plot, indicating some additional effect due to fine grain size. The  $T$ -stepping data of Fukuda et al. (2018) show a steeper slope (Fig. 7b), indicative of a higher  $Q$ -value, as originally reported. Fukuda et al. (2018) and Richter et al. (2018) recognized a mixed mode of deformation mechanism for their fine-grained samples that includes the contribution of diffusion and/or grain boundary sliding (GBS) and dislocation creep. Following their comprehensive study, it is also reasonable to assume the same for the fine-grained powder of R&B04 and silica acid originated samples of L&P92. Although the silica-gel-originated samples show a larger grain size (90  $\mu\text{m}$ ), a contribution from grain boundary processes was also interpreted due to the presence of impurities at the grain boundaries (L&P92).

#### 4.2.1. Interpreting higher $n$ -values from earlier literature

Recently, Tokle et al. (2019) and Lusk et al. (2021) compiled and reviewed earlier experimental studies in order to obtain usable flow parameters. They proposed two distinct correlations with the experimental conditions at which those studies were performed. According to Tokle et al. (2019), the higher  $n$ -values correspond to the lower differential stresses and the lower  $n$ -value to higher stresses, and both can be linked to activated slip systems. However, in this way they are essentially treating all the earlier studies, utilizing different techniques, apparatus, and materials for experiments in the same way. On the other hand, Lusk et al. (2021) implied that low confining pressures (<0.56 GPa) resulted in higher  $n$ -values and higher pressures (0.7–1.6 GPa) in lower ones. As noted by the authors, the low confining pressure conditions are restricted to Paterson-type gas medium apparatus (Table 1), in contrast to Griggs-type apparatus for high confining pressure experiments. The G&T95 study at  $\sim 1.5$  GPa confining pressure and  $n = 4$ , using a molten-salt-assembly in the Griggs-type apparatus is an exception in this trend. Moreover, at high confining pressure (1 GPa), Fukuda et al. (2018) showed that the  $n$ -value determined at low temperature (600–750 °C) and higher stress range is substantially higher (2.9 to 5.2) for fine-grained quartz, indicating power law breakdown (Richter et al., 2018). A careful inspection reveals that the lower  $n$ -value studies can be connected with high-confining pressure ones, but exclusively using solid-salt-assemblies (Table 1). More complication can arise from the fact that any theoretical and statistical re-analysis (Fukuda and Shimizu, 2017; Tokle et al., 2019; Lusk et al., 2021) of earlier experimental studies results in a substantially different flow law parameters from what is originally reported (see supplementary text S2, S3). Therefore, a re-evaluation of the wet quartzite GSI flow law for coarse-grained polycrystalline aggregates, considering the above-mentioned contradictions, is needed.

We tried to verify the  $n = 4$  from G&T95 by plotting their reported creep data in the  $\log \sigma - \log \dot{\epsilon}$  space, but the  $n$ -value came down to  $\sim 3.6$  (Fig. S3a). In the original study, the authors did not include the highest stress points, assuming a different deformation mechanism was active at the higher stress condition. Nonetheless, their result has not been reproduced by any subsequent as-is or H<sub>2</sub>O-added studies. Interestingly, the high  $n$ -value is similar to the vacuum-dried natural quartzites from solid-medium apparatus (Table 1; K&T84; Jaoul et al., 1984). As mentioned in the above section, in our non-dimensional space (negating the effect of temperature) (Fig. 7a), the molten-salt-medium experiments with a given material (i.e., Black hills quartzite), show no effect of added H<sub>2</sub>O. We suspect that the molten-salt-assembly might effectively dry the sample during the experiment, causing a similar strength between the as-is and H<sub>2</sub>O-added samples. As interpreted in Jaoul et al. (1984), the high  $n$ -values ( $>3$ ) in these dried samples were caused by the transition to brittle behaviour leading to power law breakdown (cf. Richter et al., 2018). A similar explanation for the increase in the  $n$ -value from 1.7 to 5 is given for the fine-grained samples from Fukuda et al. (2018) deformed under high stress and low-temperature conditions (cf. Richter et al., 2018).

The values of  $n \approx 3$  from fine-grained wet-samples deformed under high- $T$  condition in a Paterson apparatus are difficult to interpret as compared to Fukuda et al. (2018) with  $n = 1.7$  (Table 1). The driest sample (i.e., crushed fine-grained Brazilian single-crystal quartz with added H<sub>2</sub>O) that has been deformed by R&B04 to construct a 'wet' quartzite flow law has reported  $n = 2.97$ . Using the same type of gas apparatus, the  $\dot{\epsilon}$ -stepping data from the synthetic silica-acid-derived polycrystalline quartz of L&P92 produces a value of  $n = 4$  (modified to 3 by Fukuda and Shimizu, 2017) without considering the Goetze's criterion. According to this criterion, the  $\sigma_1$  higher than the confining pressure results in partly brittle deformation, even in high  $T$ . Noticeably, the L&P92 flow law cannot

1 predict its original creep data (Fig. S8). A steeper slope is observed  
2 for the Paterson experiments in our non-dimensional comparison  
3 (Fig. 7a). The combination of low confining pressure and the dry-  
4 ing effect of the argon gas as the confining medium (similar to  
5 the molten-salt-assembly) in the Paterson apparatus may lead to  
6 partial brittle behaviour (Kohlstedt et al., 1995) that could explain  
7 the high  $n$ -value for these fine-grained samples (cf. Richter et al.,  
8 2018). A few data from Paterson experiments (R&B 2004) at the  
9 lowest stress condition plots very close to our results. This obser-  
10 vation supports our interpretation that with increasing stress the  
11 deformation will transition to a brittle behaviour under low con-  
12 fining pressure.

13 In summary, we interpret that a  $n \geq 3$  for coarse-grained (from  
14 molten-salt-medium and 'vacuum-dried' samples of solid-medium)  
15 and fine-grained quartzite (from gas-medium and solid-medium  
16 apparatus), resulted from the onset of brittle mechanisms. The  
17  $n \leq 2$  for coarse-grained (this study) and fine-grained (9.5-12  $\mu\text{m}$ ;  
18 Fukuda et al., 2018) quartzites resulted from power-law deforma-  
19 tion due to retention of water under the high confining pressure  
20 (satisfying Goetze's criterion) in solid-medium Griggs apparatus.

#### 21 4.3. Deformation mechanism of wet quartzite

22 Commonly climb-controlled dislocation creep is inferred for the  
23 plastic deformation of quartz, requiring a theoretically expected  
24  $n$ -value of 3-5 (Karato, 2008; Paterson, 2012). A  $n$ -value of  $\leq 2$  (Ta-  
25 ble 1) requires that other processes than climb are rate-limiting for  
26 the deformation. Crystal-plastic deformation features like undula-  
27 tory patchy extinction, deformation lamellae, kink bands, and mi-  
28 nor recovery by sub-grain formation/rotation and recrystallization,  
29 limited grain boundary migration accompanied by dissolution-  
30 precipitation and micro-cracking are observed in the microstruc-  
31 tures of our deformed samples (Figs. 5, 6). In order to explain  
32 the low  $n$ -values, a contribution of diffusion creep in addition to  
33 dislocation creep has been inferred (Fukuda et al., 2018; Richter  
34 et al., 2018; Nègre et al., 2021, Pongrac et al., 2022). However,  
35 diffusion creep mechanisms are sensitive to grain size (Herring,  
36 1950; Nabarro, 1948). Thus, a grain-size-dependence is expected  
37 if diffusion creep is operating together with dislocation creep, and  
38 the contribution of diffusion creep should increase with decreasing  
39 grain size. Moreover, our study reports similar  $n$ -values as Fukuda  
40 et al. (2018), who have used fine grained quartz powder. Addi-  
41 tionally, the amount of recrystallization in our samples is less than  
42  $\sim 5\%$ , so that the grain size has remained coarse, indicating that  
43 the contribution of diffusion creep remains difficult to explain.

44 According to the von Mises criterion, at least five independent  
45 sets of slip systems are needed to achieve homogeneous deforma-  
46 tion dominantly by glide mechanism in a polycrystalline material  
47 (Karato, 2008). However, the contribution from other modes of  
48 deformation mechanisms like diffusion creep and/or grain bound-  
49 ary sliding (GBS) can relax this condition (Hutchinson, 1976), and  
50 the deformation can be achieved by activating fewer slip systems.  
51 Such a combination of processes was inferred for wet ice (Goldsby,  
52 2006; Kuiper et al., 2020), where  $n$ -values of 1.8 and 2.4 are ex-  
53 plained by the combination of basal slip (dislocation glide) and  
54 GBS (Goldsby and Kohlstedt, 2001). In that case, the strain in-  
55 compatibilities and hardening from dislocation glide are accommo-  
56 dated by grain boundary processes without activating other harder  
57 slip systems. For GBS-accommodated basal glide only a weak or  
58 no grain size dependence is observed in ice-experiments (Goldsby,  
59 2006). Recently, Tokle et al. (2019) also suspected a GBS com-  
60 ponent in the overall deformation of quartz aggregates. However,  
61 their interpretation of dislocation accommodated grain boundary  
62 sliding (disGBS) would imply a grain size dependence, whereas  
63 the  $n$ -values of  $\sim 2$  are observed in both, coarse (this study) and  
64 fine-grained material (Fukuda et al., 2018; Richter et al., 2018).

65 In addition, it is evident from CL-images that DPC was active at  
66 our experimental conditions (cf. Nègre et al., 2021; Pongrac et  
67 al., 2022). The presence of  $\text{H}_2\text{O}$  and dissolution-precipitation may  
68 weaken the grain boundary region and promote strain accommod-  
69 ation by sliding along boundaries. Recently, a comparable mech-  
70 anism was inferred in olivine (Samae et al., 2021), where grain  
71 boundary amorphization promotes GBS, leading to the viscosity  
72 drop in mantle rocks. In analogy to ice (and perhaps olivine), we  
73 propose that strain is achieved in coarse-grained wet quartzite by  
74 dislocation creep (mainly glide), accommodated by grain boundary  
75 processes including DPC and GBS. It has already been shown from  
76 similar experiments to ours that the bulk sample shortening is ac-  
77 commodated primarily (within a few percent errors) by the plastic  
78 strain of individual quartz grains (Nègre et al., 2021). Consequently,  
79 DPC and GBS both appear to act as serial processes accommodating  
80 strain incompatibilities during dislocation glide where an insuffi-  
81 cient number of slip systems operates.

82 Summarizing, the  $n = 2$  for quartzite is attributed to a com-  
83 bination of dislocation glide, GBS and DPC as serial processes. The  
84 increased solubility of  $\text{SiO}_2$  in  $\text{H}_2\text{O}$  at elevated P-T (Manning, 1994,  
85 2018) makes the DPC mechanism very efficient at our experimen-  
86 tal conditions and may in part explain the strength difference be-  
87 tween our as-is and  $\text{H}_2\text{O}$ -added samples. The inverse relation of  
88 wet quartzite strength to increasing pressure may also be a con-  
89 sequence of increased DPC activity (Nègre et al., 2021). So far, we  
90 do not observe a grain-size-sensitivity for the operation of grain  
91 boundary processes (GBS+DPC).

#### 92 4.4. Geological applications

93 In order to validate the laboratory flow laws, it is necessary to  
94 compare available natural data with the predicted rheology from  
95 experiments. This is commonly done by calculating strain rates ( $\dot{\epsilon}$ )  
96 for given stress and temperature values. Stresses usually are ob-  
97 tained using recrystallized quartz grain size ( $d$ ) piezometry, and  
98 deformation temperatures, usually from petrology of synkinematic  
99 mineral assemblages. As mentioned in the introduction, sometimes  
100 the experimentally derived flow laws can underestimate the ge-  
101 ologically reasonable  $\dot{\epsilon}$ , and therefore tend to overestimate the  
102 strength of the crust (e.g., Behr and Platt, 2011; Boutonnet et al.,  
103 2013). Here, we test the validity of our wet quartzite flow law  
104 (equation (4)) against three well-studied geological regions, where  
105 the geological  $\dot{\epsilon}$  can be estimated (Table 3): i) Eastern Tonale line,  
106 a mylonite zone of the Italian Alps (Stipp et al., 2002a, 2002b);  
107 ii) Whipple fault of the Whipple Mountains metamorphic com-  
108 plex in eastern California (Behr and Platt, 2011); and iii) the Ailao  
109 Shan-Red River (ASRR) strike-slip shear zone in southwest China  
110 (Boutonnet et al., 2013). In all of our calculations, we used the  
111 S&T03 piezometer to calculate the  $\sigma$  from the  $d$  value (without  
112 any 3D correction).

113 The Tonale Line is a dominantly strike-slip fault that displaces  
114 rocks along the Adamello pluton, which acts as a local heat source  
115 producing a contact aureole across the fault zone. Considering a  
116 displacement of 20 to  $\sim 180$  (100-150) km over 3 to 6 Ma (see  
117 Table S2), a  $\dot{\epsilon}$  range of  $1.60 \times 10^{-12}$  to  $2.11 \times 10^{-13} \text{ s}^{-1}$  can be  
118 estimated for a  $\sim 1$  km wide fault zone. Using the  $d$  and  $T$  provided  
119 in Stipp et al. (2002b), our flow-law predicts an  $\dot{\epsilon}$  range of  $1.16$   
120  $\times 10^{-12}$  to  $1.40 \times 10^{-13} \text{ s}^{-1}$ . Similarly, for the extensional Whip-  
121 ple fault from  $d$  and  $T$  values provided by Behr and Platt (2011)  
122 we can predict a  $\dot{\epsilon}$  range of  $8.10 \times 10^{-12}$  to  $1.48 \times 10^{-13} \text{ s}^{-1}$ ,  
123 where a natural  $\dot{\epsilon}$  of  $1.60 \times 10^{-11} \text{ s}^{-1}$  was inferred by Stockli et  
124 al. (2006). A well-constrained natural  $\dot{\epsilon}$  of  $3.5 \pm 0.5 \times 10^{-14} \text{ s}^{-1}$   
125 was measured from the three sets of synkinematic dikes (Sassier  
126 et al., 2009) in the ASRR fault zone. Using the  $d$  and  $T$  data from  
127 Boutonnet et al. (2013), we predict  $\dot{\epsilon} = 2.10 \times 10^{-13}$  to  $2.33$   
128  $\times 10^{-13} \text{ s}^{-1}$ . Considering the uncertainties associated with  $d$ ,  $P$ ,



**Table 3**  
Strain rates calculated from natural shear zones (also see Table S2).

Source	Grain size ( $\mu\text{m}$ )	$\sigma$ (MPa)	$T$ ( $^{\circ}\text{C}$ )	$P_c$ (MPa)	$\dot{\epsilon}$ (This study) ( $\text{s}^{-1}$ )	$\dot{\epsilon}$ (Hirth) ( $\text{s}^{-1}$ )	$\dot{\epsilon}$ (R&B) ( $\text{s}^{-1}$ )	$\dot{\epsilon}$ (L&P) ( $\text{s}^{-1}$ )
Stipp et al. (2002b)								
BLG								
27-3	5.5	173.31	304.8	300	1.40E-13	9.36E-14	1.84E-19	6.59E-15
27-1	7.8	131.34	318.4	300	1.56E-13	6.74E-14	2.94E-19	4.50E-15
26-2	8.7	120.43	345.6	300	4.56E-13	2.07E-13	2.57E-18	1.24E-14
25-3	14.3	81.18	369.4	300	5.52E-13	1.36E-13	5.49E-18	7.64E-15
25-1	15.5	76.15	383	300	8.39E-13	2.01E-13	1.31E-17	1.07E-14
24-4	24.3	53.30	400	300	7.61E-13	1.00E-13	1.55E-17	5.17E-15
SGR								
13-1	58	26.72	451	300	1.06E-12	4.81E-14	5.82E-17	2.21E-15
23-1	61	25.67	461.2	300	1.34E-12	5.94E-14	9.58E-17	2.68E-15
14-5	74	22.02	495.2	300	2.59E-12	1.01E-13	4.16E-16	4.36E-15
15-2	84	19.91	515.6	300	3.63E-12	1.28E-13	9.01E-16	5.40E-15
GBM								
21-2	219	9.31	560	300	2.31E-12	2.19E-14	8.03E-16	8.86E-16
16-3	354	6.36	563	300	1.16E-12	5.19E-15	2.98E-16	2.09E-16
Boutounet et al. (2013)								
SGR								
YY33	62	25.34	425 $\pm$ 40	130 $\pm$ 80	2.10E-13	7.32E-15	4.83E-18	6.99E-16
YY35	58	26.72	425 $\pm$ 40	130 $\pm$ 80	2.33E-13	9.05E-15	5.65E-18	8.64E-16
Behr and Platt (2011)								
PW77	135	13.66	544	560	8.10E-12	1.54E-13	2.81E-15	2.68E-15
PW71	73	22.26	500	480	5.58E-12	2.28E-13	1.03E-15	5.28E-15
PW3	53	28.70	500	480	9.27E-12	6.31E-13	2.19E-15	1.46E-14
PW25	80	20.70	500	480	4.82E-12	1.71E-13	8.28E-16	3.95E-15
PW101	75	21.79	478	430	2.49E-12	8.71E-14	2.46E-16	2.42E-15
PW104Q	68	23.55	475	430	2.71E-12	1.09E-13	2.66E-16	3.00E-15
PW44	43	33.88	405	340	4.28E-13	2.35E-14	6.68E-18	1.03E-15
PW87	52	29.14	416	360	5.05E-13	2.21E-14	9.89E-18	8.68E-16
PW75b	33	41.80	406	340	6.80E-13	5.73E-14	1.35E-17	2.49E-15
PW80b	18	67.63	406	340	1.78E-12	3.92E-13	5.62E-17	1.70E-14
PW34b	24	53.82	387	320	5.18E-13	6.37E-14	6.69E-18	3.15E-15
PW24b	36	39.01	376	310	1.74E-13	1.04E-14	1.09E-18	5.44E-16
PW79b	16	74.26	350	280	1.91E-13	3.42E-14	8.10E-19	2.20E-15
PW24a	11	99.98	357	290	4.87E-13	1.66E-13	3.65E-18	1.00E-14
PW34a	7	143.12	308	240	8.70E-14	4.07E-14	1.10E-19	3.65E-15
PW75a	8	128.73	308	240	7.04E-14	2.67E-14	8.05E-20	2.39E-15
PW79a	6	161.74	331	260	3.65E-13	2.66E-13	1.47E-18	1.97E-14
PW80a	5	186.93	308	240	1.48E-13	1.19E-13	2.44E-19	1.06E-14

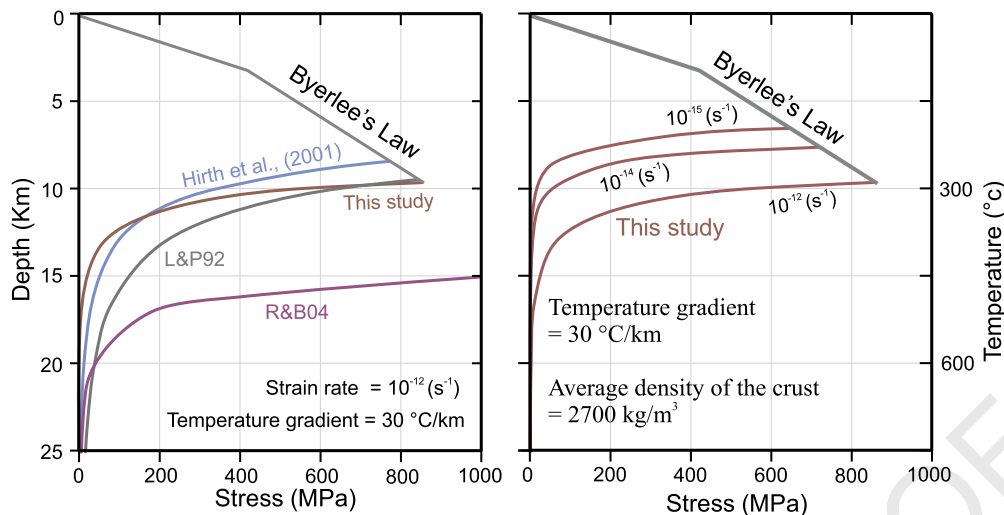
and  $T$  calculations, our new flow law predicts  $\dot{\epsilon}$  values that are in better agreement with constraints from natural observations than the previous flow laws (Table 3). In the case of the Tonale line strike slip zone, where natural constraints imply a constant  $P$  and a constant displacement rate across the zone, it is evident that our flow law provides the most consistent results irrespective of a change in  $T$  and  $\sigma$ . At high- $T$  and low- $\sigma$  conditions, the Hirth et al. (2001) and especially the other flow laws significantly overestimate the strength of the Tonale shear zone. On the contrary, our new flow law with  $n = 2$  predicts reasonable strain rates even at low stress conditions (Table 3). These observations provide an argument against a  $n$ -value transition with increasing stress (Tokle et al., 2019).

The stress versus depth profile (assuming the average density of the crust as  $2700 \text{ kg/m}^3$ ) predicted by our flow law shows the onset of plasticity ( $\sim$  brittle to viscous transition) is expected at 9–10 km depth corresponding to  $\sim 300^{\circ}\text{C}$ , considering a  $\dot{\epsilon} = 10^{-12} \text{ s}^{-1}$  and a geotherm of  $30^{\circ}\text{C/km}$  (Fig. 8). This is similar to what is usually inferred from seismological observations as the down-dip limit of the seismogenic crust (e.g., Scholz, 1998). The flow law of this study shows a distinct shape of the stress versus depth profile in comparison to the previous studies (Fig. 8). The shape of the curves derived from previous flow laws indicate a more gradual change in strength of the crust with depth. In contrast, the flow law of this study predicts a rapid drop in strength below a depth of  $\sim 10 \text{ km}$  ( $\sim 300^{\circ}\text{C}$ ). At shallower conditions, the Hirth et al. (2001) flow law predicts a lower strength in comparison to our flow law.

However, after the onset of plasticity (considering our flow law), the stress values decrease to lower strengths than the other flow laws, indicating a maximum strength of the continental crust that is restricted between 9 to 10 km and a low strength of mid- to lower continental rocks modally dominated by quartz.

## 5. Conclusions

Constant load experiments have yielded new flow law parameters for coarse-grained ( $\sim 200 \mu\text{m}$ ) wet quartzite, giving rise to a stress exponent of  $n \approx 2$  and an activation energy of  $Q = 110 \text{ kJ/mol}$ . Microstructural analysis shows that the sample strain in the wet-quartzite is achieved by grain-scale crystal-plastic processes like dislocation glide with limited recovery, accompanied by grain boundary migration and micro-cracking. Dislocation glide is the main strain producing process. Grain boundary processes, including dissolution-precipitation and grain boundary sliding are inferred to accommodate strain incompatibilities arising from the operation of insufficient slip systems. In the presence of  $\text{H}_2\text{O}$ , the grain boundary processes (GBS + DPC) become crucial for the overall plastic deformation. We interpret that a  $n = 4$  value (G&T95; L&P92) in the earlier literature resulted in part from the onset of brittle mechanisms, caused potentially either by drying effects of the sample (molten-salt-assembly) or lower confining pressure in a gas apparatus. In non-dimensional space, our new flow law can express the previous experimental data (performed using a Griggs apparatus) from natural coarse-grained samples ( $\sim 100-$



**Fig. 8.** A comparison between stress versus depth profile (assuming the average density of the crust as  $2700 \text{ kg/m}^3$  and a geotherm of  $30 \text{ °C/km}$ ). Our flow law predicts lower strength than popular flow laws at  $\dot{\epsilon} = 10^{-12} \text{ s}^{-1}$ . A strength variation at geologically viable strain rates is shown using our flow law. Yield stress envelopes are drawn based on the Byerlee's law (Byerlee, 1978).

200  $\mu\text{m}$ ) within a factor of  $\sim 5$  times the strain rate. Finally, our flow law can estimate strain rates in excellent agreement with documented natural case studies and inferred natural values. In contrast to the previous studies, our flow law suggests that the strength of the continental crust is consistently lower at temperatures above  $\sim 300 \text{ °C}$  when controlled by quartz rheology.

#### CRediT authorship contribution statement

**Subhajt Ghosh:** Conceptualization, Methodology, Experiments, Data curation, Visualization, Writing, Original draft preparation. **Holger Stünitz:** Conceptualization, Methodology, Writing. **Hugues Raimbourg:** Conceptualization, Methodology, Writing, Resource. **Jacques Précigout:** Experiments, Data curation, Writing.

#### Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

#### Data availability

I have shared the link to the data.

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The  $f_{\text{H}_2\text{O}}$  can be calculated from Tony Wither's fugacity calculator (<https://www.esci.umn.edu/people/researchers/withe012/fugacity.htm>), assuming the water pressure approximates the confining pressure during the deformation. Raw mechanical and EBSD data are available in Zenodo (<https://doi.org/10.5281/zenodo.5901073>). We gratefully acknowledge the support from both LabEx VOLTAIRE (LABX-100-01) and EquipEx PLANEX (ANR-11-EQPX-0036) projects. SG acknowledged Leif Tokle for discussions. We thank G. Hirth and A. Kronenberg for their very constructive and insightful reviews that have substantially improved this manuscript. We are also thankful to J. P. Avouc for handling this manuscript.

#### Appendix A. Supplementary material

Supplementary material related to this article can be found online at <https://doi.org/10.1016/j.epsl.2022.117814>.

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## Highlights

- Flow law parameters ( $n = 2$ ,  $Q = 110$  kJ/mol) for quartz aggregate in the presence of water are determined.
- The coarse-grained polycrystalline quartz is deformed dominantly by intra-grain plastic deformation.
- Strain incompatibilities due to the small number of active slip-systems are accommodated by grain boundary processes.
- Grain boundary processes includes dissolution-precipitation and grain boundary sliding.
- Our new creep law consistently predicts lower strength of the continental crust in the viscous regime.

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