

1 **Effect of water and rock composition on re-strengthening of cohesive faults during the**  
2 **deceleration phase of seismic slip pulses**

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10

11 **Abstract**

12 The elastic strain energy release rate and seismic waves emitted during earthquakes are  
13 controlled by the on-fault temporal evolution of the shear stress during rupture propagation.  
14 High velocity friction experiments highlighted that shear stress on the fault surface evolves  
15 rapidly during seismic slip pulses. This temporal evolution of shear stress is controlled by both  
16 fault weakening at seismic slip initiation and re-strengthening rate towards the end of slip.  
17 While numerous studies focused on fault weakening, less attention was given to co-seismic re-  
18 strengthening processes. Here we performed 53 friction experiments (normal stress  $\leq 30$  MPa,  
19 slip-rate  $\leq 6.5$  m s<sup>-1</sup>) imposing constant slip acceleration and deceleration (7.8 m s<sup>-2</sup>), on  
20 cohesive Carrara marble (99% calcite) and micro-gabbro (silicate-built rock) under dry, vacuum  
21 and water pressurized conditions. Microstructural observations showed that micro-gabbro  
22 accommodated seismic slip by bulk melting of the sliding surfaces, whereas Carrara marble by  
23 coupled decarbonation and grain-size dependent crystal plastic processes. Under room humidity  
24 conditions and low imposed power density (i.e., product of normal stress per slip rate), re-  
25 strengthening rate during the deceleration stage was up to  $\sim 17$  times faster in marble than in

26 micrograbbro. In the latter, the re-strengthening rate increased slightly with the power density.  
27 The presence of water enhanced further this trend. On the contrary, in marbles the re-  
28 strengthening rate decreased drastically with power density and in the presence of water. Our  
29 experimental observations highlighted the first order importance of the mineralogy and  
30 rheology of the slip zone materials and, to a second order, of the presence of water in controlling  
31 co-seismic re-strengthening of faults during seismic slip deceleration.

32

33 Keywords: Earthquake, Friction, Re-strengthening, Ground motions, Water effect.

34

35 1) Introduction:

36 Earthquakes are the result of sudden slip along faults, which releases the elastic strain  
37 and gravitational energy stored in the wall rocks [Reid, 1910; Scholz, 2002]. Damage associated  
38 with earthquakes is due to the radiation of elastic waves during seismic rupture propagation  
39 [Stein and Wyssession, 2009, Marty et al., 2019] and to dynamic stress concentration at the  
40 rupture front [e.g., Andrews, 2005]. The intensity and the frequency of seismic waves (ground  
41 motions) are controlled by the rupture length and by the abrupt variations in rupture and slip  
42 velocities along fault during earthquakes [Brune, 1970; Madariaga, 1976]. Seismic rupture (~  
43 km/s) and slip velocities (~ m/s) depend in particular on the temporal evolution of the shear  
44 stress along the fault during sliding (i.e., fault strength weakening and re-strengthening) [Ida,  
45 1972; Ruina, 1983]. Friction experiments, performed with rotary shear machines designed to  
46 impose on rock specimens slip and slip rates typical of natural earthquakes, proved that  
47 experimental faults weakened due to several rock-type and environmental-dependent thermally  
48 activated weakening mechanisms, including melt lubrication, grain-size dependent processes,  
49 thermal and thermomechanical pressurization of pore fluids, etc. [Tsutsumi and Shimamoto,  
50 1997; Di Toro et al., 2004; Han et al., 2010; Rice, 2006; Ferri et al., 2010; Di Toro 2011; Green

51 et al., 2015; De Paola et al., 2015; Spagnuolo et al., 2015; Violay et al., 2015]. Field studies of  
52 exhumed natural seismogenic faults support the evidence of rapid weakening and fault  
53 lubrication during earthquakes [Sibson, 1975; Di Toro et al., 2006]. The rheology of the fault  
54 is related to the physical state (e.g., liquid vs. solid) of the slipping zone [Rice, 2006], the slip  
55 rate and slip acceleration [Di Toro et al., 2004; Niemeijer et al., 2011; Chang et al., 2012], the  
56 rock composition [Di Toro et al., 2011; Goldsby and Tullis, 2011; Green et al., 2015], the  
57 presence of water [Violay et al., 2013; 2014], the ambient and local temperature [Lockner et  
58 al., 1986, Verberne et al., 2015], and in presence of non-cohesive rocks, on the gouge grain  
59 size [De Paola et al., 2015]. Indeed, in cohesive rocks, slip is highly localized from the very  
60 initiation of sliding and the rise of the temperature in the slipping zone increases with slip rate,  
61 effective normal stress and square root of the duration of sliding [e.g., Rice, 2006, Aubry et al.,  
62 2018]. In cohesive rocks, the abrupt increase in temperature favors “flash” heating and  
63 weakening mechanisms [Goldsby and Tullis, 2011] which may eventually evolve, for some  
64 silicate-built cohesive rocks, in bulk melting [Hirose et al., 2005; Niemeijer et al., 2011; Violay  
65 et al., 2013; Aubry et al., 2018]. On the contrary, in the presence of gouges, some slip is required  
66 before strain localization is achieved [Beeler et al., 1996; Marone 1990; Smith et al., 2015;  
67 Rempe et al., 2017; Pozzi et al., 2018]. As a consequence, heat production is more distributed  
68 in the slipping zone and other processes including thermal (if fluids are already present in the  
69 pores of the slipping zone) and thermomechanical (if fluids like CO<sub>2</sub> and H<sub>2</sub>O are released by  
70 the breakdown of the minerals of the slipping zone) pressurization might be favored before bulk  
71 melting of the slipping zone occurs [e.g., Rice, 2006; Segall and Rice, 2006; Brantut et al.,  
72 2008; Ferri et al., 2010; Acosta et al., 2018].

73 While the physical parameters controlling fault weakening have been extensively  
74 studied, little attention has been dedicated on the processes associated to frictional re-  
75 strengthening observed during slip deceleration in seismic velocity pulses [Del Gaudio et al.,

76 2009; Chang et al., 2012; Liao et al., 2015; Proctor et al., 2014; Sone and Shimamoto, 2009].  
77 However, fault strength evolution, i.e both strength weakening and re-strengthening, coupled  
78 with electrodynamic rupture propagation contributes actively to the slip rate evolution during  
79 seismic faulting and to the release of seismic waves [Sone and Shimamoto, 2009]. In addition,  
80 co-seismic re-strengthening processes controls the energy budget of earthquakes, as well as the  
81 transition from crack-like to pulse-like rupture, which implies a strong fault re-strengthening  
82 after the passage of the rupture front [Beeler and Tullis, 1998]. Few mechanisms of fault re-  
83 strengthening have been proposed so far: (1) temporal variation of stress conditions on the  
84 rupture interface during the passage of the rupture front (e.g., punctual pore fluid pressurization)  
85 [Lykotrafitis et al., 2006], (2) spatial variation of the initial stress field (barrier model) [Peyrat  
86 et al., 2001, Latour et al., 2011], (3) high dependence of friction to sliding velocity, i.e., "self-  
87 healing" behavior during sliding velocity deceleration [Beeler and Tullis, 1998; Sone and  
88 Shimamoto, 2009; Proctor et al., 2014; Perrin et al., 1995; Zheng and Rice, 1998]. Here, we  
89 present friction experiments that, by reproducing seismic slip conditions, aim at understanding  
90 the effect of rock composition and water content on fault re-strengthening during slip  
91 deceleration. In particular, we focused on cohesive rocks (calcitic marble and microgabbro)  
92 where strain is localized from slip initiation to the final re-strengthening stage and the  
93 temperature evolution in the slipping zone is simpler than in case for non-cohesive rocks  
94 (gouges). Moreover, the analysis of the data reported here exploit the achievement of the so-  
95 called "steady-state" shear stress conditions in the experiments. This achievement requires slips  
96 of several centimeters to tens of centimeters depending on the applied effective normal stress  
97 [Di Toro et al., 2011]. Consequently, the results presented here should be valid for moderate to  
98 large in magnitude earthquakes, though steady-state conditions might not ever be achieved in  
99 nature [Liao et al., 2015]. The experimental evidences suggest that rock composition and

100 environmental conditions play a pivotal role in fault re-strengthening during seismic faulting,  
101 with dramatic consequences in the radiation pattern of the emitted seismic waves.

102

## 103 2) Methods

104 We performed 53 high speed friction experiments on hollow cylinders (30/50 mm  
105 internal/external diameter) of carbonate-built rock (Carrara marble, 99% calcite) and silicate-  
106 bearing rock (micro-gabbro) (for sample preparation, see Nielsen et al., 2013). Carbonate-built  
107 rocks and gabbros often host earthquakes sequences in nature, as attested by geological  
108 evidence [Sibson, 1975]. The experiments were performed with SHIVA, a rotary shear machine  
109 installed at the HP-HT INGV laboratories in Rome (see sup.mat.Table 1). SHIVA was equipped  
110 with two brushless engines (max power 300 kW) and an air actuator (2000 kg amplified to 5000  
111 kg thanks to a lever) in a rotary shear configuration to slid the two contacting hollow rock  
112 cylinders under the desired conditions (Di Toro et al., 2010). Experiments were performed  
113 either in the presence of pressurized liquid water, room humidity or under vacuum ( $10^{-4}$  mbar).  
114 In the experiments with liquid water, SHIVA was equipped with a pressurizing system which  
115 consisted of (1) a fluid pressure vessel (i.e., the samples were fully immersed in water), (2) a  
116 membrane pump with a 30 cm<sup>3</sup> fluid capacity, (3) a pressure multiplier that imposes up to 15  
117 MPa of fluid pressure ( $P_f$ ), (4) a pressure regulator and, (5) valves and pipes [Violay et al.,  
118 2013]. Normal stress ( $\sigma_n$ ) and pore pressure ( $P_f$ ) (drained conditions) were kept constant during  
119 experiments to target values ranging between 10 and 40 MPa and 0 (nominally dry) and 15  
120 MPa, respectively. To mimic the sliding velocity at a given point of the fault during propagation  
121 and arrest of seismic slip, we imposed a trapezoidal slip velocity function by imposing constant  
122 acceleration and deceleration ( $7.8 \text{ m s}^{-2}$ ) and target slip velocities ranging from  $1 \text{ m s}^{-1}$  to  $6.5 \text{ m}$   
123  $\text{s}^{-1}$ . Total slip ranged from 0.83 m to 18.37 m.

124 Mechanical data (axial load, torque, axial displacement, and angular rotation) were  
125 acquired at a frequency up to 25 kHz (for description of the installed instrumentation, their  
126 calibration and acquisition rates, see Niemeijer et al., 2011). Normal stress, shear stress ( $\tau$ ) and  
127 slip velocity ( $V$ ) were computed from experimental measurements following Tsutsumi and  
128 Shimamoto [1997] and Di Toro et al. [2010].

129

### 130 3) Results

131 In all the experiments, once the slip velocity function was applied, the apparatus and the  
132 sample initially deformed elastically until the static friction coefficient  $\mu_p$  ( $\mu = \tau/\sigma_n^{eff}$  with  
133  $\sigma_n^{eff} = \sigma_n - P_f$ ) was overcome and slip initiated (Fig. 1). Then, the friction coefficient decreased  
134 exponentially with slip over a weakening distance  $D_w$  down to a so-called steady-state friction  
135 coefficient  $\mu_{ss}$ . Towards the end of the experiment, during slip velocity deceleration, the friction  
136 coefficient recovered with time and slip to a final value  $\mu_r$  at the offset of slip (Figs.1a-b). The  
137 effects of rock composition and environmental conditions on fault weakening and  $\mu_p$ ,  $\mu_{ss}$  and  
138  $D_w$  were discussed in Violay et al. [2013 and 2014]. As well as for initial fault weakening [Di  
139 Toro et al., 2011], the recovery of the friction coefficient during the deceleration stage of the  
140 velocity pulse is a function of the sliding velocity (Figs.1c-d) [Goldsby and Tullis, 2011;  
141 Passelègue et al., 2014; Proctor et al., 2014]. The value of  $\mu_r$  ranged from 0.08 to 0.63 for  
142 Carrara marble and 0.11 to 0.37 for micro-gabbro, respectively, and independently of (1) the  
143 environmental conditions, (2) the final slip and (3) of the initial power ( $\sigma_n^{eff} V$ ) imposed on the  
144 fault (Figs. 2 and 3). To investigate the physics of the re-strengthening processes, we computed,  
145 for each experiment, the friction re-strengthening rate  $\dot{\mu} = (\mu_r - \mu_{ss})/(t_r - t_{ss})$ , where  $(t_r - t_{ss})$  is the  
146 deceleration duration from end of the steady-state to the end of slip (Figs. 1a-b). At constant  
147 acceleration and deceleration rate, in the experiments conducted at low initial power density

148 ( $\sigma_n^{eff} V < 20 \text{ MW m}^{-2}$ ) and under both room humidity and vacuum conditions, the re-  
149 strengthening rate was  $\sim 17$  times faster in Carrara marble (from  $2.00$  to  $3.68 \text{ s}^{-1}$ ) than in micro-  
150 gabbro (from  $0.02$  to  $0.16 \text{ s}^{-1}$ ) (Fig 4). In both rock types, re-strengthening rate was initial power  
151 density-dependent and work density independent (i.e., slip independent) (Fig.4). In Carrara  
152 marble, independently of the environmental conditions, the friction re-strengthening rate  
153 decreased with increasing power density (Figs. 3-4). This trend was amplified in pressurized  
154 water conditions, and at the largest power densities tested, re-strengthening rate became almost  
155 negligible (Fig. 4a). The opposite behavior was observed in micro-gabbro. The re-strengthening  
156 rate slightly increased with the power density, and this effect was amplified in presence of  
157 pressurized water (Fig. 4b). Moreover, in Carrara marble the re-strengthening rate was about  
158 two times (at low power density) slower in the presence of pressurized liquid water than under  
159 room and vacuum conditions (Fig 4). Instead, in micro-gabbro, the frictional re-strengthening  
160 rate was about three times faster in the presence of pressurized liquid water than under room  
161 humidity and vacuum conditions (Fig 4). Therefore, pressurized liquid water had an opposite  
162 effect on the frictional re-strengthening rate of Carrara marble with respect to the one of micro-  
163 gabbro.

164       Regarding the microstructures, mineralogy and geochemistry of the slipping zones and  
165 slip surfaces recovered after the experiments, we refer to previous studies performed (1) on  
166 several samples from the experiments presented here [Violay et al., 2013; 2014] and (2) on  
167 slipping zone produced in experiments conducted on similar rocks (e.g., gabbro, basalts and  
168 Carrara marbles) under very similar deformation conditions [Han et al., 2010; Di Toro et al.,  
169 2011]. In the case of micro-gabbro, the slipping zone consisted of a continuous ca.  $200 \mu\text{m}$  thick  
170 layer of a quenched melt (see composition in Table 2), independently of the presence or absence  
171 of liquid water (Fig. 5a) [Violay et al., 2014; 2015; Nielsen et al., 2008; 2010; Niemeijer et al.,  
172 2011; Giacomel et al., 2018]. In the case of the Carrara marble, the slipping zone after

173 experiments conducted under vacuum and room humidity consisted of a 50 to 100  $\mu\text{m}$  thick  
174 layer of nano- to micro-grained, poorly cohesive material made of calcite and, to a minor extent,  
175 lime (Violay et al., 2013; Spagnuolo et al., 2015) (Fig. 5b). Unfortunately, due to the poorly  
176 cohesive nature of the nano- to micro-grained deformed layer, most of the slipping zone was  
177 flushed away during sample recovery in the experiments performed in the presence of liquid  
178 water. However, the few micro-fault patches recovered from the slip surface showed the  
179 presence of nano- to micro-grained slipping zones [Violay et al., 2013; 2015] and evidence of  
180 calcite decarbonation attested by the presence of vacuoles within the grains. Violay et al.,  
181 [2013] also showed evidence of  $\text{CO}_2$  increase in the fluids recovered after high-speed friction  
182 experiments performed on carbonate bearing rocks. We infer that also in the case of the  
183 experiments performed with Carrara marble, the slipping had the same microstructure  
184 independently of the presence or absence of liquid water.

185

#### 186 4) Discussion

##### 187 4.1) *Effect of water on friction re-strengthening rate*

188 The different effect of the presence of pressurized liquid water on the frictional re-  
189 strengthening rate of micro-gabbro and Carrara marble suggested that different micro-physical  
190 processes operated during the deceleration phase of the slip pulse, as clearly supported by  
191 microstructural evidence (Fig. 5). It is well-known that fault surfaces of micro-gabbro melt with  
192 seismic slip under room humidity and vacuum conditions [Tsutsumi and Shimamoto, 1997;  
193 Hirose and Shimamoto, 2005; Nielsen et al., 2008; Niemeijer et al., 2011], but also in the  
194 presence of liquid water [Violay 2014a; 2014b]. On the contrary, slipping zones of Carrara  
195 marble did not record microstructural evidence of frictional melting. Instead, the slipping zones  
196 were made of sub-micrometer to nanometer in size grains [Spagnuolo et al., 2015]. These  
197 microstructures can be associated to grain-size dependent crystal plastic processes [Green et al.,

198 2015; De Paola et al., 2015; Spagnuolo et al., 2015]. Similarly to what occurred during the  
199 initial weakening stage, was the rheology of the materials building the slipping zone that  
200 controlled the final frictional re-strengthening. In our experiments, the rheology of the slipping  
201 zone depended on the physical state of the sheared materials (melt vs. nano-grains), on the slip  
202 and strain rate, on the normal stress, as well as on the environmental conditions and, for Carrara  
203 marble, on grain size.

204 In the case of micro-gabbro, the viscosity and thickness of the melt layer and its  
205 extrusion rate from the slipping zone controlled the viscous strength of the experimental fault  
206 [Nielsen et al., 2008]. Violay et al., [2014a] showed that the initial fault weakening by frictional  
207 melts was delayed by the presence of liquid water that cools the asperity contacts. Here, we  
208 hypothesize that frictional re-strengthening might be enhanced by an increase of the viscous  
209 strength of the melt layer due to water-cooling. To test this hypothesis, we implemented a Finite  
210 Element Methods numerical model which included the rock specimens and the steel-made  
211 pressure vessel (Fig. 6 a, b, c). The slipping zone was modeled as a 200  $\mu\text{m}$ -thick layer (red in  
212 color in Fig. 6, a), consistently with the typical thickness of the solidified melt layers recovered  
213 at the end of the experiments [Nielsen et al., 2008], made by melt plus water or melt plus air  
214 with a constant volume ratio ( $\phi$ ) ranging from 0.5 to 0.9 in agreement with microstructural  
215 observations [Brown and Fialko, 2002, Violay et al., 2014a; 2014b] (Fig. 5). The melt  
216 temperature during slip deceleration was computed using the mechanical data of the samples  
217 sheared under vacuum conditions as a reference. In fact, under vacuum conditions, most of the  
218 frictional work was dissipated into heat (the contribution to wear and rock fragmentation was  
219 negligible, see Niemeijer et al., 2011) and very limited heat was lost by radiation. Consequently,  
220 the heat flux  $Q$  was a function of shear stress and slip rate that evolved with time  $t$  (Fig. 1) and  
221 varied along the sample radius  $r$ :

$$222 \quad Q(r, t) = 0.5 \cdot \tau(t) \cdot V(r, t) \quad (\text{Eq.1}).$$

223 For experiments performed under room humidity conditions and in the presence of water, we  
 224 imposed that the melt layer was cooled by air or water (2D heat diffusion model), respectively.  
 225 Then, as representative of the entire slipping zone, we used the estimated temperature achieved  
 226 by the friction melt at 2/3 of the sample radius. The properties of the slipping zone (indicated  
 227 by the subscript *eff* for effective) were considered as a linear combination of the thermal  
 228 properties of the fluid (air or water) and those of the melt (Table 1). Therefore, the “effective”  
 229 thermal diffusivity in the slip zone was:

$$230 \quad \alpha_{eff} = K_{eff} / (\rho C)_{eff} \quad (Eq.2)$$

231 where

$$232 \quad K_{eff} = (1 - \phi) K_R + \phi K_f \quad (Eq.3)$$

233 and

$$234 \quad (\rho C)_{eff} = (1 - \phi) \rho_R C_R + \phi \rho_f C_f \quad (Eq.4)$$

235 with  $K$ , the thermal conductivity ( $\text{W m}^{-1} \text{K}^{-1}$ ),  $C$  the specific heat capacity ( $\text{J kg}^{-1} \text{K}^{-1}$ ),  $\rho$  the  
 236 density ( $\text{kg m}^{-3}$ ) and  $\phi$  the liquid fraction in the melt. Index  $R$  and  $f$  are related to the rock and  
 237 fluid properties, respectively. Then, the heat diffusion is:

$$238 \quad \rho \cdot C_{eff} \cdot \frac{\partial T}{\partial t} = \nabla \cdot (k_{eff,i} \cdot T) \quad (Eq.5)$$

239 where  $i$  represents the two different materials (water/air and rock) in the model of Fig. 6. Values  
 240 of thermal conductivity, specific heat capacity and density are reported in Table 1. The melt  
 241 viscosity ( $\eta$ ) was computed with the viscosity simulator for silicate melts of Giordano et al.,  
 242 [2008] (we did not take into account the crystal fraction in the melt). The model predicts the  
 243 non-Arrhenius temperature dependence of viscosity for naturally-occurring silicate melts at  
 244 atmospheric pressure:

$$245 \quad \log(\eta) = A + \frac{B}{T(K) - C} \quad (Eq.6)$$

246 where  $A$  is a constant independent of composition and  $B$  and  $C$  are adjustable parameters  
247 depending on melt composition. The glass chemical composition was published by Violay et  
248 al., [2014, sup mat item 4] and reported in Table 2. Numerical modelling results highlighted  
249 that the melt viscosity, because of cooling of the melt, increases with decreasing sliding velocity  
250 (Fig. 6), in agreement with the logarithmic increase of the “friction coefficient” during the  
251 deceleration stage (Figs. 1 and 3). In addition, the increase of melt viscosity during slip  
252 deceleration was significantly faster in experiments performed in the presence of pressurized  
253 water than under room humidity conditions, suggesting that water-cooling was an efficient  
254 mechanism of re-strengthening. Moreover, the efficiency of water-cooling was proportional to  
255 the fluid-to-melt ratio in the slipping zone, and therefore of the temperature of the melt before  
256 deceleration. Indeed, higher was the melt temperature at the steady-state, which was slightly  
257 proportional to the imposed power density, faster was the increase of melt viscosity during slip  
258 deceleration. Instead, given the same conditions, the observation that the re-strengthening rate  
259 was independent of slip (Figs. 4d) was related to the large shortening rate of the rock specimens  
260 during frictional melting (mm per meters of slip, Violay et al., 2014). At steady-state conditions,  
261 the isotherms were almost fixed in space while the rock specimen passes through them and got  
262 melted and extruded (see discussion in Nielsen et al., 2008). As a consequence, once steady-  
263 state had been achieved (always the case for the experiments presented here), during  
264 deceleration and melt cooling, the wall rocks were at similar temperatures independently of the  
265 cumulated slip and the re-strengthening is independent of slip.

266 In the case of Carrara marble, the slower re-strengthening rate in the presence of water  
267 than under room humidity conditions (Fig. 4a) cannot be explained by water cooling of the  
268 slipping zone or by thermal fluid depressurization (i.e., fluid pressure decrease after a short  
269 period of expansion caused by frictional heating). Indeed, neither melt was produced nor  
270 pressurization was observed during these experiments where fluid pressure was kept constant

271 under drained conditions [Violay et al., 2015]. Moreover, both melt lubrication and thermal  
272 pressurization would lead to a faster re-strengthening in presence of water than under room  
273 humidity conditions. However, this is at odds with the experimental evidence: re-strengthening  
274 rate is faster under vacuum conditions (Fig4). Note also that calcite decarbonation as potential  
275 re-strengthening mechanism can be ruled out in our experiments. This mechanism would lead to  
276 faster re-strengthening at high power density (i.e., higher temperature), which it is just the  
277 opposite of our experimental evidence.

278 We suggest that the slower recovery of the frictional strength in the presence of liquid water  
279 during slip deceleration can be due to the strain rate sensitive, grain size dependent crystal  
280 plasticity of calcite in the presence of water (e.g. grain boundary sliding aided by diffusion  
281 creep) [Rutter, 1974, Schmidt et al., 1980, 1987, Walker et al., 1990, Renner et al., 2002].  
282 Decarbonation reaction can probably help this mechanism because it triggers the formation of  
283 nanoparticles allowing the activation of grain size dependent processes. Despite relevant  
284 differences between our experimental protocol and those reported in the above-mentioned  
285 studies (high pressure - high temperature triaxial tests performed at very low strain rates,  $10^{-6}$ -  
286  $10^{-4} \text{ s}^{-1}$ , compared to those achieved in our experiments,  $10^2$ - $10^4 \text{ s}^{-1}$ ), a comparison with those  
287 studies may help to understand better the mechanisms accommodating deformation in our  
288 experiments. At high temperature ( $> 400^\circ\text{C}$ ), calcite strength is temperature and strain rate  
289 dependent [Rutter, 1971, 1974], and governed by crystal-plastic and grain size-dependent  
290 deformation mechanisms (intracrystalline plasticity and grain boundary sliding diffusion-  
291 assisted plasticity). Similar deformation mechanisms were inferred to be operating at sub-  
292 seismic and seismic slip rates in calcite [Verberne et al., 2015; De Paola et al., 2015; Spagnuolo  
293 et al 2015]. Rutter [1972] showed that calcite strength is inversely proportional to water content  
294 due to the reduction of the surface energy at calcite grain boundaries [Boozer et al., 1962] and  
295 its strain rate sensitive behavior decreased with increasing water content. Rutter [1974] had also

296 shown that water effect is relatively small in Carrara marble. If crystal-plastic and grain-size  
 297 dependent processes were triggered during high velocity friction experiments, they would  
 298 therefore result in a negative dependence of strain-rate on residual flow stress, and may explain  
 299 the inhibition of the frictional re-strengthening in Carrara marbles especially at large power  
 300 densities. In fact, for larger power densities, higher temperatures should be achieved in both the  
 301 slipping zone and wall rocks. This should be the case especially once the so-called steady-  
 302 steady conditions were achieved (note that, because of the power-law decay in Carrara marble,  
 303 “steady-state” conditions remain an approximation, see Nielsen et al., 2016). In fact, because  
 304 of the absence or negligible shortening of the marble specimens, contrary to what happens in  
 305 micro-gabbro, heat diffused in the wall rocks. Hotter wall rocks and the different deformation  
 306 mechanism involved would result in slower cooling of the slipping zone, especially if compared  
 307 to the frictional melting case (compare Fig. 4a with 4b). In the presence of water, the efficiency  
 308 of grain size dependent and diffusion-controlled processes should be enhanced due to lower  
 309 activation energy in presence of water, explaining the lower re-strengthening rate. However,  
 310 the slipping zone was flushed away at the end of slip preventing sample recovery for micro-  
 311 analytical investigations [Violay et al., 2013, 2014]. To test this hypothesis, we performed a  
 312 one dimension-time dependent diffusion temperature model fully coupled with grain-size  
 313 diffusion creep flow law (Fig.7). Calcite thermal properties are reported on Table 1. The  
 314 slipping zone was modeled by a 100  $\mu\text{m}$ - layer composed of 100 nm grain size calcite, in  
 315 agreement with microstructural observations [Violay et al., 2013, 2015] (Fig.5). The heat flux  
 316 and diffusion were computed following Eq.1 and 5. The predicted flow stress for diffusion  
 317 creep plasticity can be modelled by the constitutive flow law (Eq. 8, 9).

318

$$\frac{\partial T}{\partial t} = \frac{1}{2} \tau(t) V(t) + \frac{k}{c_p \rho} \frac{\partial^2 T}{\partial x^2} \quad (\text{eq.7})$$

319

$$\tau(t) = \left( \frac{\dot{\gamma}}{\left( A \cdot D^{-b} e^{-\frac{H}{RT(t)}} \right)} \right)^{\frac{1}{n}} \quad (\text{eq.8})$$

320 
$$\dot{\gamma} = \frac{1}{l} \frac{\partial V}{\partial t} \quad (\text{eq.9})$$

321 where  $\dot{\gamma}$  is the shear strain rate,  $A$  the pre-exponential factor,  $H$  the apparent activation energy  
322 for creep,  $R$  the gas constant,  $T$  the temperature,  $\tau$  the shear stress at time step ( $t$ ),  $n$  the stress  
323 exponent,  $D$  the grain size with  $b$  the grain size exponent,  $l$  is the layer thickness, and  $V(t)$  is  
324 the slip velocity function.

325 Following [De Paola et al., 2015], under room-humidity conditions,  $b=2$ ,  $n=1$ ,  $H=217$  kJ mol<sup>-1</sup>,  
326  $A=9.55 \cdot 10^5$  (s<sup>-1</sup> · Bar<sup>-n</sup>),  $R=8.314$  J.K.mol<sup>-1</sup>. In presence of water, we supposed a reduction  
327 of the activation energy ( $H$ ) of about 20% , with  $H=176$  KJ kJ mol<sup>-1</sup> (see discussion in Rutter  
328 1972). Flow stress estimates based on Eqs.7,8,9 (Fig.7) showed that grain boundary sliding  
329 aided by diffusion mechanism can explained both the strength weakening at the beginning of  
330 the experiments, the steady state friction and the final re-strengthening. Moreover, a decrease  
331 of the activation energy of only 20% under pore fluid conditions [Rutter, 1972] may justify the  
332 faster re-strengthening observed under dry rather than under pore fluid conditions (Fig. 1c).

333

#### 334 *4.2) Implications for natural earthquakes*

335 Our experiments demonstrated that the slip deceleration can result in rapid fault re-  
336 strengthening, up to 10 to 90% of the initial peak stress, depending on the rheology of the  
337 material building the slip zone which depends on lithology and environmental conditions (Fig.  
338 1). Because the friction coefficient is strongly velocity-dependent [Sone and Shimamoto, 2009],  
339 the magnitude of the re-strengthening depends mainly on the imposed deceleration rate [De  
340 Gaudio et al., 2009]. The extrapolation of these results to natural earthquakes can be gained by  
341 (1) comparing the friction re-strengthening rate between experiments performed with different  
342 rock types and under environmental conditions, and (2) using the dependency of the re-  
343 strengthening rate with the power density.

344 Here below, we estimate the slip-velocity, acceleration and power density of natural  
345 earthquakes and compare them to the imposed parameters during our experiments. Slip-  
346 velocity functions during earthquake rupture propagation are obtained by inverting ground  
347 motion waveforms. Slip acceleration and deceleration during earthquakes are  $\sim 1-10 \text{ m s}^{-2}$  (and  
348 larger, Tinti et al., 2005) and slip rates, on average,  $\sim 1 \text{ m s}^{-1}$  [Heaton, 1990]. Therefore,  
349 acceleration, deceleration as well as slip-velocity imposed in our experiments are somehow  
350 comparable to those of natural earthquakes. The main differences between tests and  
351 earthquakes are in the shape of the velocity function (trapezoidal in our case) and on the  
352 normal stress imposed. Indeed, crustal earthquakes nucleate between 5 and 15 km depth,  
353 possibly at effective normal stresses of  $\sim 100-200 \text{ MPa}$  [Zoback and Harjes, 1997], and at  
354 power densities perhaps up to ten times higher than those imposed in the experiments  
355 discussed here. Extrapolation of our results to realistic stresses conditions suggest that in  
356 cohesive carbonate-bearing rocks, frictional re-strengthening during slip-velocity deceleration  
357 is probably almost negligible whereas in cohesive silicate bearing-rocks, re-strengthening  
358 processes are highly deceleration-dependent, especially in the presence of water.

359 Indeed, our results demonstrate that, due to the large expected power densities during  
360 natural earthquakes, small variations in the slip velocity could induce strong variations in  
361 friction promoting further changes in the slip velocity history due to co-seismic re-  
362 strengthening, especially in the case of calcite-built rocks. Moreover, the frequency content of  
363 the radiated energy is affected by the abruptness of velocity changes (emission of higher  
364 frequency waves is expected under abrupt decelerations) and thus, by the re-strengthening rate.  
365 Based on our results, cohesive silicate-bearing faults that undergo to frictional melting, also in  
366 the presence of liquid water [see natural case discussed by Brantut and Mitchell, 2018] will  
367 have the largest re-strengthening rates at the highest power densities (5-15 km depth) and should  
368 promote more intense high frequency radiation during slip deceleration. However, according to

369 Fig. 4, the re-strengthening rates in micro-gabbro are much smaller than those achieved in  
370 calcitic-built cohesive rocks. In the latter rocks, if grain-size dependent processes are activated,  
371 intense high frequency radiation should occur especially at low power densities (perhaps  
372 corresponding to shallow depths, < 2 km) and dry conditions, when re-strengthening rates are  
373 the highest (Fig. 4).

374

## 375 5) Conclusions

376 We performed a series of experiments simulating seismic slip under different  
377 environmental conditions (vacuum, room humidity and pressurized water) on two common  
378 cohesive crustal rocks (calcitic Carrara marble and micro-gabbro). In general, independently of  
379 the environmental conditions, the fault re-strengthening rate at the end of slip is up to one order  
380 of magnitude larger in Carrara marble than in micro-gabbro, especially at low power densities  
381 (Fig. 4). This large difference in re-strengthening rates is due to the different on-fault  
382 deformation processes operating during seismic slip: crystal plastic and grain-size dependent  
383 for Carrara marble, melt lubrication for micro-gabbro. The two deformation mechanisms have  
384 different constitutive equations and dependence with temperature. With increasing power  
385 density, which may correspond to increasing crustal depths, the fault re-strengthening rate  
386 becomes almost negligible for Carrara marble whereas it slightly increases for micro-gabbro.

387 We also found some intriguing second order differences in the magnitude of the re-  
388 strengthening rate for the two rock types. These differences are due to the environmental  
389 conditions which impact on the efficiency of the particular co-seismic deformation mechanism  
390 of the rock. Under vacuum and room-humidity conditions, fault re-strengthening rate at the end  
391 of simulated seismic slip in Carrara marble is significantly faster than in the presence of  
392 pressurized water, especially at low power densities (Fig. 4). Instead, in micro-gabbro, fault re-

393 strengthening rate is faster in the presence of pressurized water than under room humidity and  
394 vacuum conditions (Fig. 4). We interpreted these well-reproducible second order variations in  
395 the re-strengthening rate as the consequence of the change in rate-dependent plasticity in the  
396 presence of water for Carrara marble and water-cooling of the frictional interface for micro-  
397 gabbro. Our results suggest that both rock composition and presence of water affect the elastic  
398 strain energy release rate and the seismic waves radiation pattern during rupture propagation.

399

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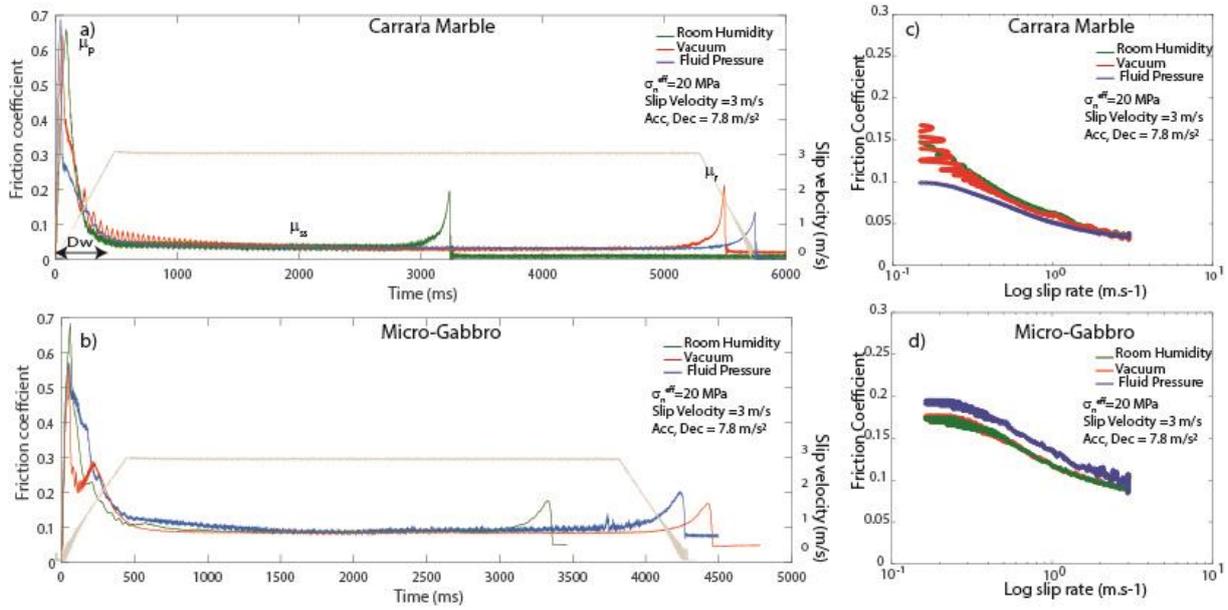
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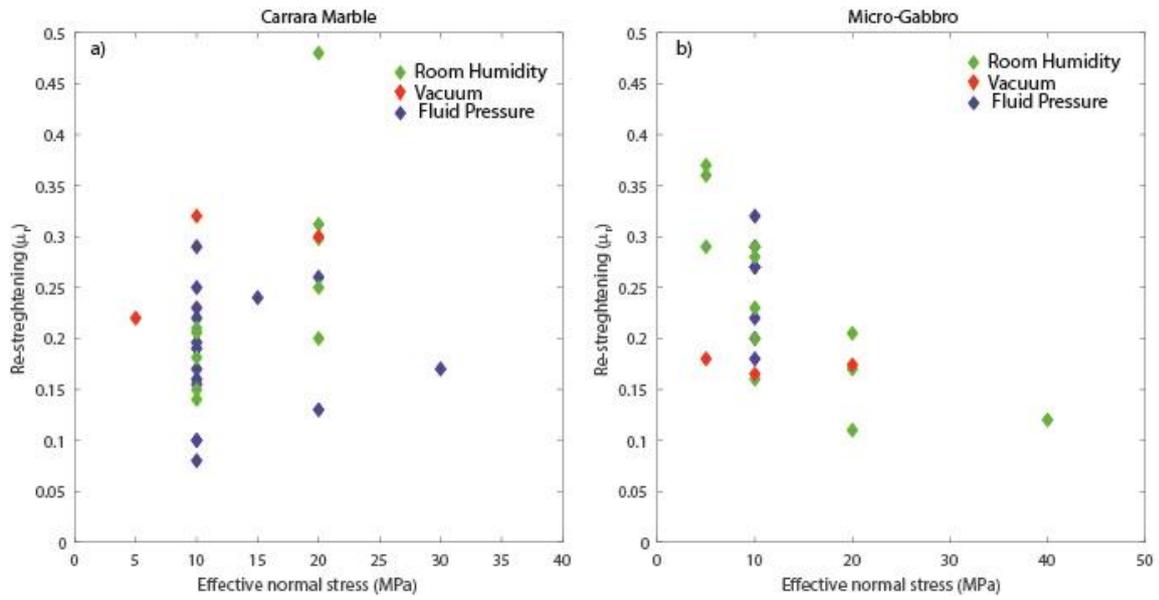
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576 **Figures:**



578

579 **Figure 1.** Evolution of the friction coefficient with time and slip rate for Carrara marble and  
 580 micro-gabbro slid at seismic slip rates. **(a) and (b)** Evolution of the friction coefficient with  
 581 time in Carrara marble and micro-gabbro. **(c) and (d)** Evolution of the friction coefficient  
 582 measured during deceleration versus log of the sliding velocity. The experiments were  
 583 conducted at target velocity  $V_t = 3 \text{ m s}^{-1}$ , acceleration and deceleration  $= 7.8 \text{ m s}^{-2}$ , and  $\sigma_n^{eff} =$   
 584 20 MPa (effective normal stress  $= \sigma_n - P_f$ ). Blue curves (S615 and S567): fluid pressure  
 585 experiments  $\sigma_n = 25 \text{ MPa}$ ,  $P_f = 5 \text{ MPa}$ ; green curves (S307 and S555): room humidity  
 586 experiments; red curves (S614 and S585): vacuum experiments were run at  $P_{vacuum} = 10^{-4} \text{ mbar}$ .  
 587 Two examples of sliding velocity function are drawn in grey.



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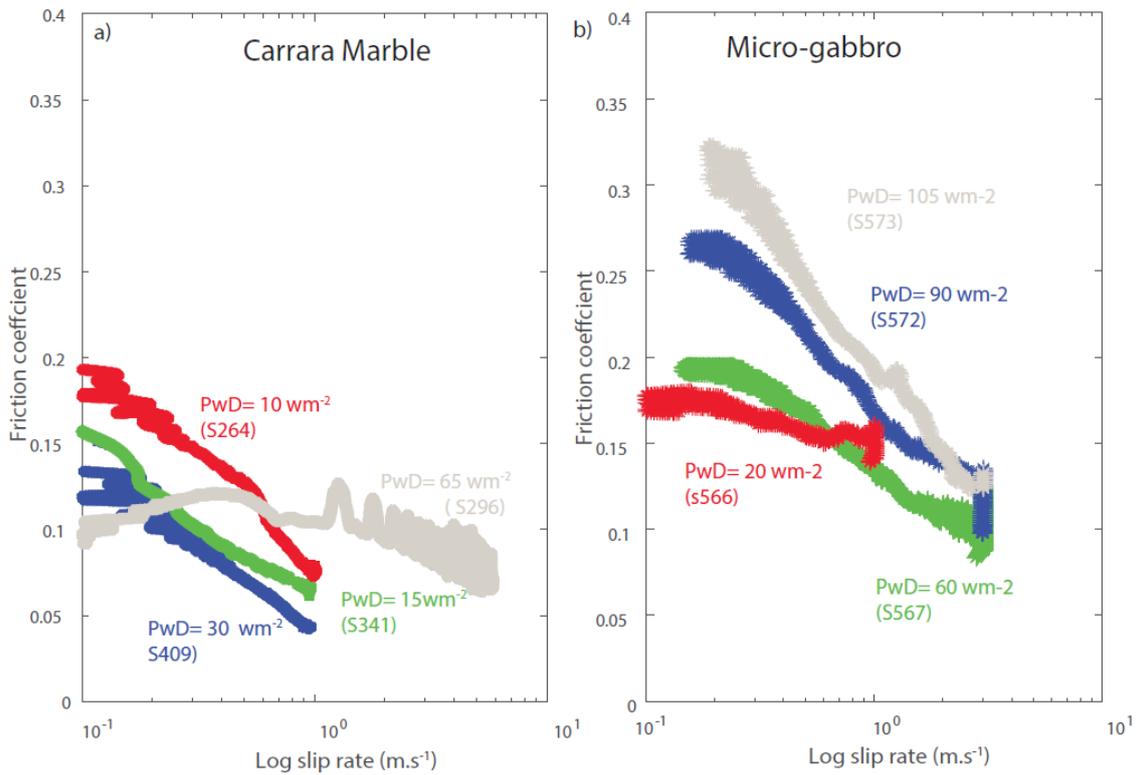
589 **Figure 2.** Influence of  $\sigma_n^{eff}$  on the re-strengthening for (a) Carrara Marble and (b) micro-gabbro.

590 Blue dots: fluid pressure experiments; green dots: room humidity experiments; red dots:

591 vacuum experiments  $P_{vacuum} = 10^{-4}$  mbar.

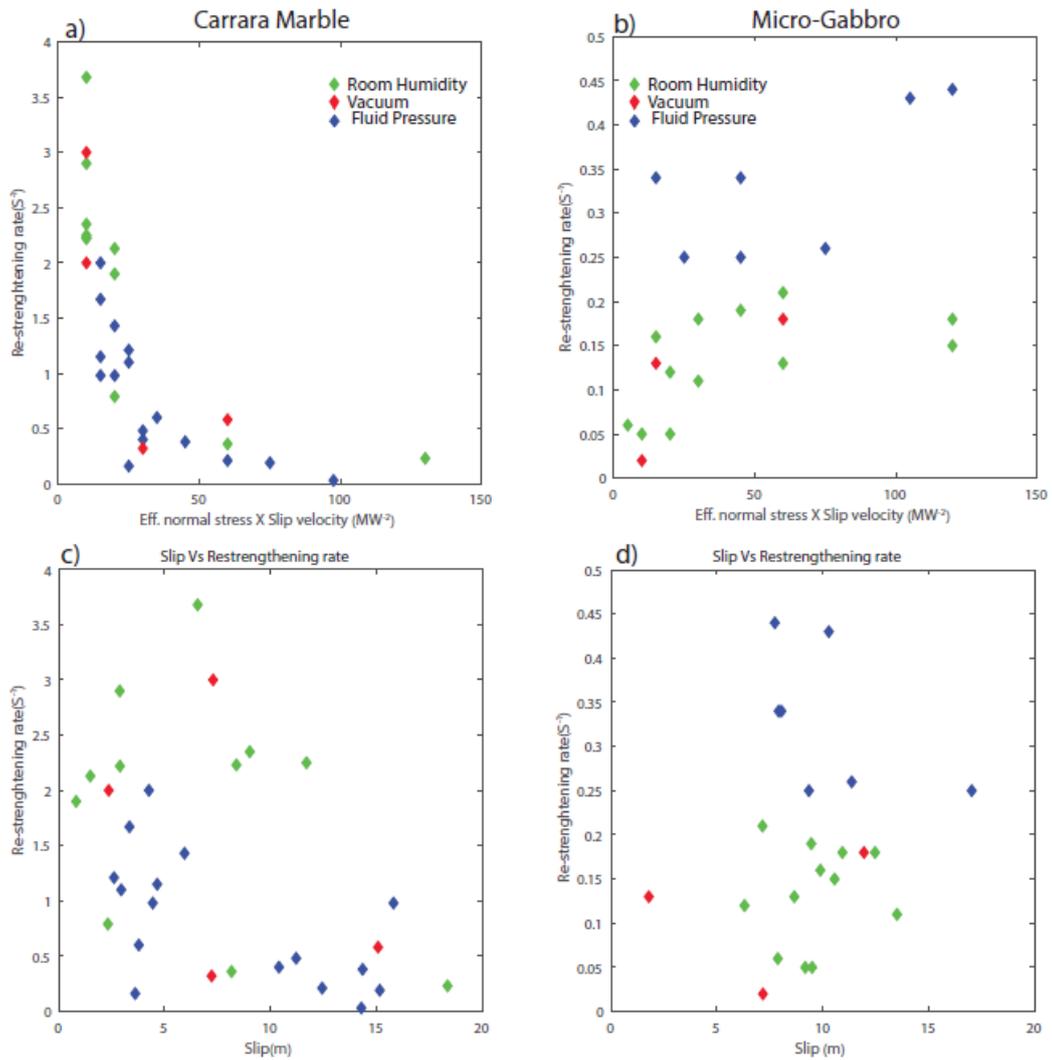
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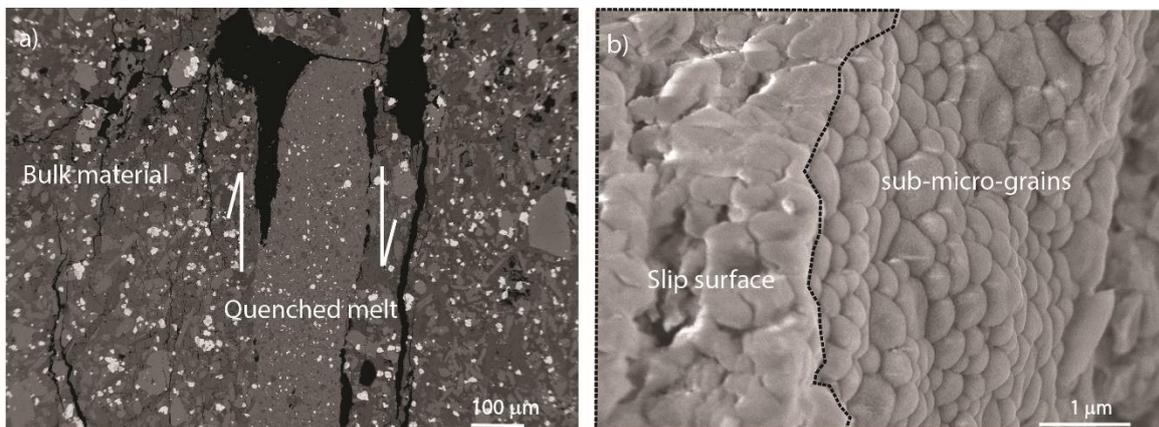
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595 **Figure 3.** Evolution of the friction coefficient during slip deceleration versus log of the sliding  
 596 velocity for **(a)** Carrara marble (S409, S341, S296, S264) and **(b)** micro-gabbro (S566, S567,  
 597 S572 and S573). Experiments were conducted at different power density ( $\sigma_n^{eff} \cdot Vt$ ). Target  
 598 velocity ( $Vt$ ) ranging from 1 and 6.5 m s<sup>-1</sup>, acceleration and deceleration = 7.8 m s<sup>-2</sup>, and  $\sigma_n^{eff}$   
 599 from 10 to 30 MPa (effective normal stress=  $\sigma_n - P_f$ ). The experiments were all performed under  
 600 fluid pressure conditions ( $P_f = 5$  MPa).



601

602 **Figure 4.** Influence of power density **(a,b)** ( $\sigma_n^{eff} \cdot Vt$ ) and slip **(c, d)** on the friction re-  
 603 strenghtening rate. Blue dots: fluid pressure experiments; green dots: room humidity  
 604 experiments; red dots: vacuum experiments  $P_{vacuum} = 10^{-4}$  mbar.

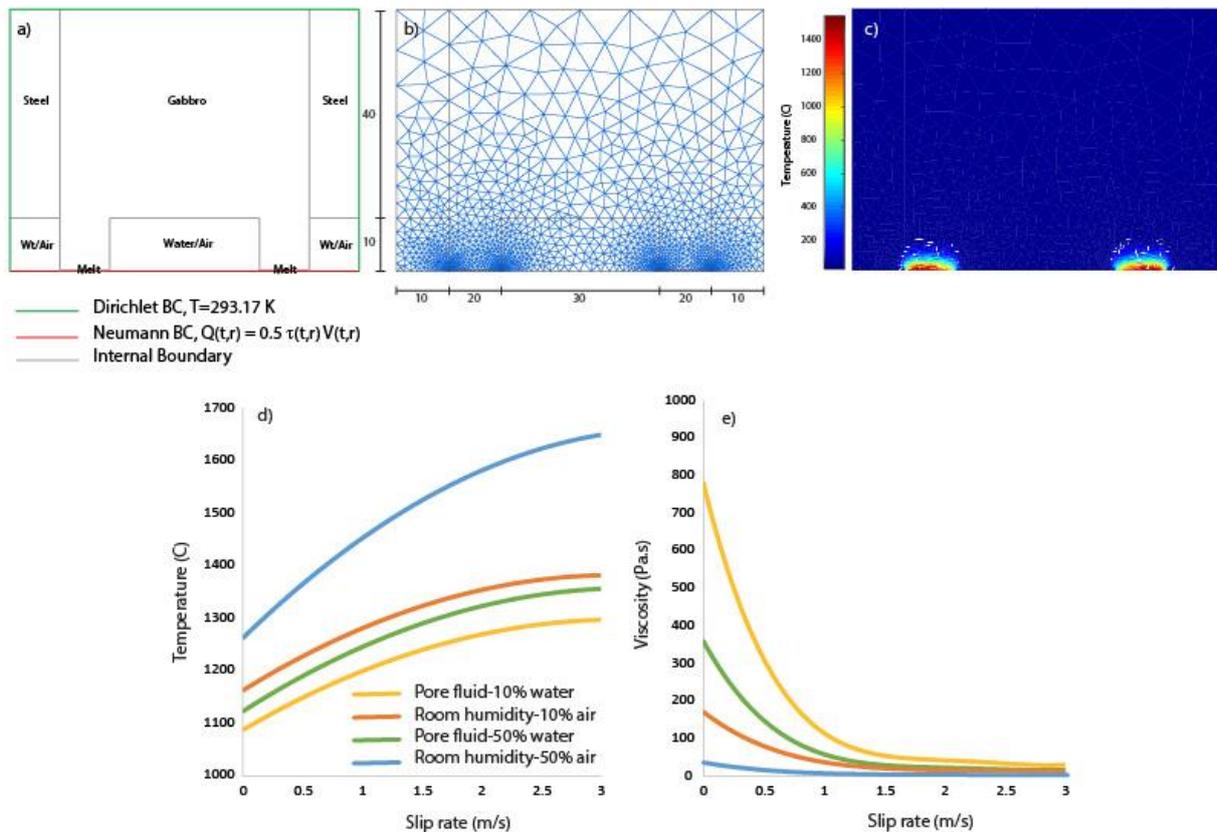


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606 **Figure 5.** Microstructural observations of experimental slip surfaces and slipping zones after  
 607 shearing. **(a)** Slipping zone after an experiment performed on micro-gabbro (s585,  $V = 3 \text{ m s}^{-1}$ ,  
 608  $\sigma_n = 20 \text{ MPa}$ , Vacuum conditions see also Violay et al., 2014). The slipping zone is made by  
 609 quenched melt (i.e., glass matrix, see composition in Table 2) which wraps grains of plagioclase  
 610 and pyroxene that survived from frictional melting (Scanning Electron Microscope, back scatter  
 611 electron image). **(b)** Slip surface after an experiment performed on calcitic Carrara marble  
 612 (s614,  $V = 3 \text{ m s}^{-1}$ ,  $\sigma_n = 20 \text{ MPa}$ , Vacuum conditions, see Violay et al., 2014). The slip surface  
 613 is made by micro- to nano-grains of calcite and lime. Small decarbonation vacuoles decorate  
 614 the recrystallized calcite grains (Scanning Electron Microscope, secondary electron image).

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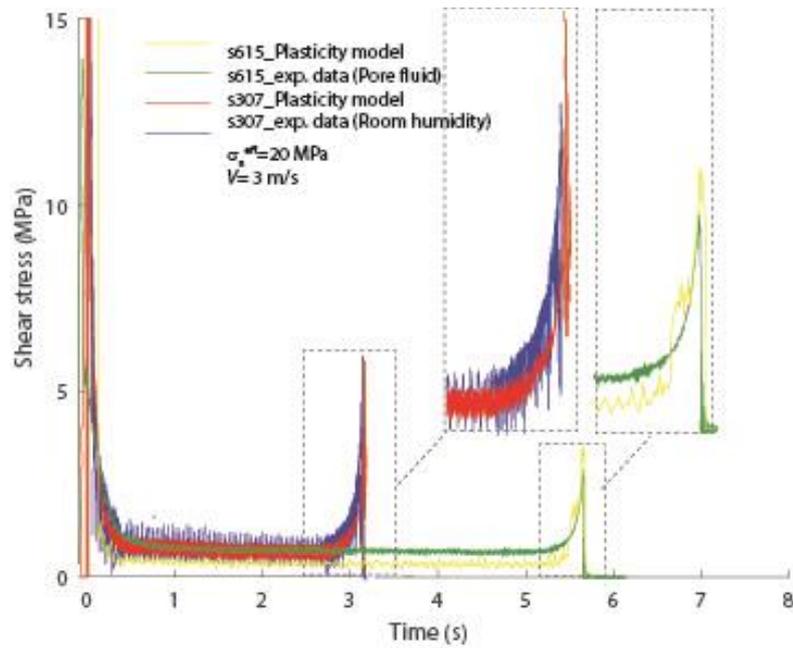
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618 **Figure 6.** FEM 2D time dependent heat diffusion model. **(a,b)** Model geometry with description  
 619 of the Boundary Conditions and mesh geometry. **(c)** The snapshot of the temperature  
 620 distribution at the end of the experiment s585 conducted on gabbro. **(d)** Temperature evolution

621 of the slip zone during slip deceleration ( $t_r-t_{ss}$ ). (e) Melt viscosity evolution during slip  
 622 deceleration.

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626 **Figure 7.** Results of the 1D time dependent heat diffusion model fully coupled with diffusion  
 627 creep flow law for both room humidity (blue and red curves) and pore fluid (green and yellow  
 628 curves) conditions.

629

|                                      | Micro-Gabbro | Carrara Marble | Water | Air  | Melt  | Steel |
|--------------------------------------|--------------|----------------|-------|------|-------|-------|
| K [ $10^{-6} \text{ m}^2/\text{s}$ ] | 0.8          | 1.48           | 0.15  | 1.5  | 0.344 | 4.2   |
| $\rho$ [ $\text{kg}/\text{m}^3$ ]    | 2990         | 2700           | 1000  | 1.2  | 2591  | 2000  |
| C [ $\text{J}/(\text{kgK})$ ]        | 949          | 700            | 4200  | 1000 | 1484  | 460   |

630 **Table 1.** Thermal properties of the fluid (air and water) and micro-gabbro used in the FEM

631 numerical model, as well as the thermal properties of Carrara marble used for the coupled

632 diffusion model and plasticity flow law.  $K$  = thermal conductivity,  $\rho$  = density,  $C$  = specific  
 633 heat.  $\Phi$  the liquid fraction.

|                                |              |       |      |
|--------------------------------|--------------|-------|------|
| Sample                         | Micro-Gabbro | s585  |      |
| Phase                          | crystalline  | glass | s.d. |
| #                              |              | 7     |      |
| Al <sub>2</sub> O <sub>3</sub> | 14.27        | 15.68 | 0.74 |
| Na <sub>2</sub> O              | 4.49         | 4.26  | 0.91 |
| CaO                            | 9.22         | 10.21 | 1.43 |
| SiO <sub>2</sub>               | 45.48        | 45.22 | 0.90 |
| K <sub>2</sub> O               | 2.71         | 2.71  | 0.57 |
| MgO                            | 7.15         | 6.88  | 0.57 |
| P <sub>2</sub> O <sub>5</sub>  | 1.88         | 1.14  | 0.14 |
| FeO                            | 10.28        | 8.31  | 0.34 |
| TiO <sub>2</sub>               | 3.02         | 2.88  | 0.38 |
| MnO                            | 0.19         | 0.17  | 0.05 |
| Total                          | 98.69        | 97.46 |      |

634 **Table 2.** Chemical composition of the micro-gabbro (XRD) and of the glass for experiment  
 635 S585 performed under vacuum conditions (Electron Microprobe Analysis). The analysis does  
 636 not close to about 100% because only Fe<sup>2+</sup> was determined.