

## Brittle-to-viscous behaviour of quartz gouge in shear experiments

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In order to study the microstructure development across the brittle-viscous transition and to derive the corresponding flow laws, we performed shear experiments on quartz gouge in a Griggs-type deformation apparatus. The starting material is a crushed quartz single crystal (sieved grain size  $<100 \mu\text{m}$ ) with 0.2 wt% water added. The experiments were conducted at temperatures between 500 °C and 1000 °C at confining pressures of 0.5 GPa, 1.0 GPa or 1.5 GPa. Four strain-rate-stepping experiments were conducted at strain rates between  $\sim 2.5 \times 10^{-6} \text{ s}^{-1}$  and  $\sim 2.5 \times 10^{-4} \text{ s}^{-1}$ . Other experiments were conducted at constant strain rates of  $\sim 2.5 \times 10^{-6} \text{ s}^{-1}$ ,  $\sim 2.5 \times 10^{-5} \text{ s}^{-1}$ ,  $\sim 2.5 \times 10^{-4} \text{ s}^{-1}$  and  $\sim 2.5 \times 10^{-3} \text{ s}^{-1}$ .

At high confining pressure, the strength of the samples decreases with increasing temperature for all strain rates. The largest decrease occurred between 650 °C and 700 °C at shear strain rates of  $\sim 2.5 \times 10^{-5} \text{ s}^{-1}$ . At the same time, the pressure dependence of strength is positive for  $T \leq 650 \text{ °C}$  while an inverse pressure dependence is observed at  $T > 650 \text{ °C}$ . For  $T < 700 \text{ °C}$ , the friction coefficient decreases slightly with increasing temperature, from 700-1000 °C it shows a strong temperature dependence.

Between 650 °C and 700 °C at shear strain rates of  $\sim 2.5 \times 10^{-5} \text{ s}^{-1}$  a change in the deformation process occurs from one dominated by cataclastic flow to one dominated by crystal plasticity. The microstructure reveals a less abrupt transition in terms of operating processes, because brittle and viscous processes are equally active around 650 °C. With increasing temperature the volume fraction of recrystallised grains increases, and at 900 °C - 1000 °C recrystallisation is nearly complete at strains of  $\gamma \sim 3$ . The crystallographic preferred orientation of the c-axis evolves from a random distribution at low temperatures towards two peripheral maxima at intermediate temperatures. At high temperatures the c-axis show a single Y-maximum.

At high temperature, the stress exponent is  $n = 2.1 \pm 0.2$ . The activation energy  $Q$  is  $193 \pm 12 \text{ kJ/mol}$  at strain rates of  $10^{-5} \text{ s}^{-1}$ , at faster strain rates the activation energy drops down to  $Q = 119 \pm 12 \text{ kJ/mol}$ . This small stress exponent at high temperatures indicates a combination of deformation processes (diffusion in very fine grained material and dislocation creep in coarser grained material). At lower temperatures the n-value is significantly higher ( $n \sim 8$ ) indicating the beginning of the power-law-breakdown.

Interestingly, the mechanical data indicate a sharp transition between brittle- and viscous-dominated regimes around 650 °C to 700 °C, while the microstructure reveals a smoother transition over a wider temperature range. The relatively low stress exponent of  $n \sim 2$  clearly suggests the activation of diffusion creep deformation processes.