

Effects of dissolution–precipitation creep on the frictional properties of opal gouge at low-temperature hydrothermal conditions

*Nakanishi Tomoya², Kyuichi Kanagawa¹, Michiyo Sawai¹

1. Graduate School of Science, Chiba University, 2. Faculty of Science, Chiba University

In order to examine the effects of dissolution–precipitation creep on the frictional properties at low-temperature hydrothermal conditions, we conducted triaxial friction experiments on opal gouge at a confining pressure of 150 MPa, a pore water pressure of 50 MPa, and temperatures (T) ranging from room T to 200°C, and at displacement rates (V) changed stepwise among 0.1155, 1.155 and 11.55 $\mu\text{m/s}$. We then fitted the friction data for each step change in V by the rate- and state-dependent friction constitutive law, and obtained the optimized ($a - b$) value, i.e., an indicator of frictional stability, at each V .

The results show that steady-state friction coefficient μ_{ss} increases with increasing T , from 0.64 at room T to 0.67 at 200°C, which is consistent with slip hardening behavior observed at higher T s. Microstructural observations reveal that significant grain interlocking and porosity reduction occur in the gouge layer sheared at higher T s. Thus increasing gouge lithification with increasing T , which is promoted by thermally activated dissolution–precipitation creep, is likely responsible for increasing μ_{ss} with increasing T .

Our results also show that ($a - b$) value tends to decrease with increasing T or decreasing V at $T \geq 50^\circ\text{C}$. Decreasing ($a - b$) value with decreasing V at a given T is likely due to increasing gouge lithification and hence μ_{ss} with decreasing V , which is promoted by dissolution–precipitation creep favored at lower V s. At a given V , a value does not change much while b value increases with increasing T , which results in decreasing ($a - b$) value with increasing T . Increasing b value with increasing T implies that more strength recovery occurs when V is stepped down, which is also ascribed to increasing activity of dissolution–precipitation creep. Because ($a - b$) value does not change with V at room T , dissolution–precipitation creep was not active at room T .

At a given V , the transition from $a - b > 0$ to $a - b < 0$ occurs with increasing T , but the transition T is also dependent on V , because ($a - b$) value is dependent on both T and V as described above; $T < 50^\circ\text{C}$ at $V = 0.1155 \mu\text{m/s}$, $50^\circ\text{C} < T < 100^\circ\text{C}$ at $V = 1.155 \mu\text{m/s}$, and $T > 100^\circ\text{C}$ at $V = 11.55 \mu\text{m/s}$. Our results suggest that increasing activity of dissolution–precipitation creep with increasing T or decreasing V promotes decreasing ($a - b$) value and hence the transition from stable aseismic faulting with $a - b > 0$ to unstable, possible seismic faulting with $a - b < 0$.

Keywords: frictional properties, opal gouge, low-temperature hydrothermal conditions, dissolution–precipitation creep