Redox heterogeneities in the Archean mantle inferred from mineral inclusions in Siberian E-type diamonds: implications for volatile speciation and diamond formation

GIULIA MARRAS¹, DENIS MIKHAILENKO², CATHERINE MCCAMMON³, SONJA AULBACH⁴, ALLA LOGVINOVA⁵, SERENA DOMINIJANNI⁶ AND VINCENZO STAGNO⁷

¹Department of Earth Sciences, Sapienza University of Rome ²Sobolev Institute of Geology and Mineralogy of the Siberian Branch of the RAS, Novosibirsk, Russia

³Bayerisches Geoinstitut, Universität Bayreuth, Germany

⁴Goethe-Universität Frankfurt

⁵Institute Geology and Mineralogy SB RAS

⁶Institut de Minéralogie de Physique des Matériaux et de

Cosmochimie, Sorbonne Université, Paris, France

⁷Department of Earth Sciences, Sapienza University of Rome, Rome, Italy

Presenting Author: giulia.marras@uniroma1.it

The cycle of volatiles among Earth's reservoirs is related to the oxygen fugacity (fo_2) recorded by mantle rocks, which controls the mobility and composition of volatile-rich fluid/melt. Eclogites formed by the subduction of ancient oceanic crust (protolith) offer insight into the fate of early Earth's volatiles in depth. According to oxy-thermobarometric estimates, most eclogites fall into the diamond stability field equilibrated with CH₄-H₂O dominated C-O-H fluids (Stagno and Aulbach 2021). However, the primary redox state of the subduction-modified eclogite source could be masked by interaction with (metasomatic) circulating fluids through time, which likely acted themselves as growth media for diamonds. Mineral inclusions trapped in eclogitic diamonds can provide pristine information on the redox conditions of the eclogite source.

We investigated nineteen eclogitic diamonds from Udachnaya (Russia) with both exposed (after polishing) and trapped paired garnet and clinopyroxene inclusions (three touching), and crystals extracted from the host rock of one diamond. We combined mineral chemistry (major elements and vanadium) with oxy-thermobarometric and H₂O estimates, to reconstruct the nature of the inferred Archean protolith and redox evolution along subduction, with implications for fluid circulation and diamond formation. We employed EPMA for major elements and V concentration analysis, while Fe³⁺/ Σ Fe was measured by synchrotron Mössbauer source spectroscopy. The OH⁻ bands were detected in clinopyroxene by micro-FTIR supported by X-ray tomography to determine the water content.

Pressure and temperature of equilibrium are 4.9-7.6 GPa and 1080-1350 °C. The Fe³⁺/ Σ Fe is up to 0.07 for garnet and 0.26 for clinopyroxene, while V concentration in the reconstructed bulk rock varies from ~150 to 450 ppm. The calculated log*fo*₂ ranges from -4 to -0.4 log units (FMQ). Notably, we observed similar

 $log fo_2$ and V bulk rock for inclusions in diamond and corresponding minerals in the host eclogite, indicating that (metasomatic) diamond formation did not affect $log fo_2$, as proposed by Aulbach and Stachel (2022). Our findings show that inclusions recorded heterogeneous redox conditions, indicating that diamonds likely grew from diverse fluids/melts from CH₄-H₂O to CO₂-H₂O dominated.

Stagno and Aulbach (2021). In Magma Redox Geochemistry (eds R. Moretti and D.R. Neuville)

Aulbach and Stachel (2022). Nature Communications, 13(1), 1924