

Pfaffenbergite, $\text{KNa}_3(\text{Al}_4\text{Si}_{12})\text{O}_{32}$, a polymorph of sodic feldspar isostructural with kokchetavite and wodegongjieite, found in crystallized melt inclusions in metamorphic garnet

SILVIO FERRERO^{1,*}, SOFIA LORENZON², ROBERTO BORRIELLO^{2,3}, ALESSIA BORGHINI⁴, RICHARD WIRTH⁵, ANJA SCHREIBER⁵, RICO FUCHS⁶, PATRICK J. O'BRIEN⁶, EDWARD S. GREW⁷, AND ENRICO MUGNAIOLI²

¹Department of Chemical and Geological Sciences, University of Cagliari, I-09042 Monserrato, Italy

²Department of Earth Sciences, University of Pisa, I-56126 Pisa, Italy

³Department of Environmental Sciences, Informatics and Statistics, Ca' Foscari University of Venice, I-30172 Mestre (VE), Italy

⁴Faculty of Geology, Geophysics and Environmental Protection, AGH University of Kraków, 30-059 Kraków, Poland

⁵Helmholtz-Zentrum Potsdam, Deutsches GeoForschungsZentrum (GFZ), D-14473 Potsdam, Germany

⁶Institute of Geosciences, University of Potsdam, D-14476 Potsdam, Germany

⁷School of Earth and Climate Sciences, University of Maine, Orono, Maine 04469, U.S.A.

ABSTRACT

Pfaffenbergite, $\text{KNa}_3(\text{Al}_4\text{Si}_{12})\text{O}_{32}$, is a new mineral found in nanogranitoids included in garnets from HP and UHP eclogites in the Saxo-Thuringian Zone of the Bohemian Massif (Saxony, Germany). The occurrence of a new mineral phase was initially inferred from its unique micro-Raman spectrum, characterized by a very strong vibrational mode at 412 cm^{-1} and minor peaks at 105, 832, 130, and 470 cm^{-1} . Subsequently, three-dimensional electron diffraction revealed that pfaffenbergite is a hexagonal mineral, crystallizing in space group $P6/mcc$, isostructural with kokchetavite (KAlSi_3O_8) and wodegongjieite [$\text{KCa}_3(\text{Al}_7\text{Si}_9)\text{O}_{32}$]. These two sheet silicates have feldspar stoichiometry and chemical composition, and pfaffenbergite corresponds chemically to an unmixed binary K-Na feldspar. Microstructural and experimental constraints suggest that it formed as a result of melt crystallization during cooling, after entrapment within metamorphic garnets. The precise pressure-temperature conditions of formation of pfaffenbergite are currently unknown. However, they must have been below the P - T conditions of entrapment of the inclusions, i.e., 1000–1050 °C and 2.2–4.5 GPa, as determined in the present work, based on the values estimated for each individual case study.

We interpret pfaffenbergite as a metastable phase crystallizing rapidly in a silicate melt enclosed in a small pore under non-equilibrium conditions, also based upon previous studies reporting the occurrence of other metastable phases in the same or neighboring nanogranitoids (kokchetavite, kumdykolite, dmisteinbergite, etc.). The increasing number of findings of metastable phases in the last years suggests that these minerals are more common than expected. We propose that it is even possible they may represent rock-forming minerals in natural rocks that experienced rapid cooling/rapid crystallization, for instance, lavas and ignimbrites, along with experimental products involving silicate melts.

Keywords: Pfaffenbergite, nanogranitoids, polymorph, feldspar, wodegongjieite, Bohemian Massif, Waldheim