NEW MINERAL NAMES

Indialite

AKIHO MIYASHIRO AND TOSHIMICHI IIYAMA, A preliminary note on a new mineral, indialite, polymorphic with cordierite. *Proc. Japan Acad.*, **30**, 746–751 (1954).

X-ray spectrograms of natural cordierite differ, especially in the range of 2θ between 29° and 30° for Cu K α radiation from those of synthetic α -Mg₂Al₄Si₅O₁₈ and β -Mg₂Al₄Si₅O₁₈. Alpha-Mg₂Al₄Si₅O₁₈ is hexagonal with a 9.782 Å, c 9.365 Å. The so-called "synthetic iron cordierite" (Schairer and Yagi, Am. J. Sci., Bowen Vol., 471-512 (1952)) is the corresponding hexagonal α -Fe₂Al₄Si₅O₁₈, with a 9.860 Å, c 9.285 Å. Beta-Mg₂Al₄Si₅O₁₈, synthesized by F. R. Boyd of the Geophysical Laboratory, is also hexagonal with a 9.792 Å, c 9.349 Å. Both of these give patterns similar to, but distinctly differing from those of the two naturally occurring polymorphs of cordierite which are orthorhombic (or possibly monoclinic), pseudohexagonal.

It has now been found that the "cordierite" described by Venkatesh, Am. Mineral., 37, 831–948 (1952), 39, 636–646 (1954) is actually the alpha-form, which is now named indialite. X-ray powder data are given; they are indexed and lead to a hexagonal cell with a 9.812 Å, c 9.351 Å, indicating a ratio of Mg: Fe⁺² of 7:3. The mineral occurs in sedimentary rocks fused by the burning of a coal seam in the Bokaro coalfield, India. The burning was due to some natural cause.

The name is for India.

MICHAEL FLEISCHER

Pavonite

E. W. NUFFIELD, Am. Mineral., 39, 409-415 (1954).

Unnamed Magnesium Borate

R. M. THOMPSON AND J. A. GOWER, Am. Mineral., 39, 522-524 (1954).

Callaghanite

C. W. BECK AND J. H. BURNS, Am. Mineral., 39, 630-635 (1954).

Ortho-antigorite

G. W. BRINDLEY AND O. VON KNORRING, Am. Mineral., 39, 794-804 (1954).

M. F.

Berillite

M. V. KUZMENKO, Berillite-a new mineral. Doklady Acad. Nauk. S.S.S.R., 99, 451-454 (1954) (in Russian).

Berillite occurs as fine spherulites 2–3 mm. in diameter or as crusts up to 2 mm. thick in cavities. Color white, luster silky. Soft. Berillite is optically biaxial, negative, with $n_{\rm s}$, $\alpha = 1.541 \pm 0.003$, $\beta = 1.553 \pm 0.002$, $\gamma = 1.560 \pm 0.002$, 2V small, less than 45°. Extinction parallel, elongation positive. G.=2.196.

Analysis on 250 mg. by M. E. Kazakova gave SiO₂ 34.10, TiO₂ tr., Al₂O₃ 1.63, Fe₂O₃ 0.12, MgO tr., BeO 40.00, CaO 0.50, Na₂O 2.42, H₂O⁺ 18.95, H₂O⁻ 3.25; sum 100.97%. (Sum given as 100.47 in original. M.F.). This corresponds closely to Be₃SiO₄(OH)₂ \cdot H₂O (neglecting the Al₂O₃, CaO, Na₂O, and H₂O⁻. M.F.). Spectrographic study showed moderate lines of Ba and Mn and weak lines of P and Cu. A dehydration curve is given; about 12% H₂O is lost up to 350°.

X-ray powder data are given. The strongest lines have spacings (Å? Kx?) and intensities: 4.10 10, 2.34 10, 3.64 9, 1.351 8, 3.39 7, (3.19, 2.90) 7, 2.12 7, 2.03 7, 1.937 7, 2.703 6.

Berillite occurs in the center of a zoned pegmatite that cuts aegirine lujavrite. The border zone consists mainly (70-80%) of aegirine and arfvedsonite; these decrease in amount towards the center and microcline and natrolite increase. The center zone consists mainly of natrolite with relicts of very coarse sodalite (hackmannite) and with fine-grained albite. Berillite occurs rarely in cavities and coating albite and epididymite. The locality is not stated (could be Kola Peninsula? M.F.)

DISCUSSION: The name as given is a direct transliteration, but perhaps the spelling beryllite would be preferable.

M. F.

Shcherbakovite

E. M. Es'KOVA AND M. E. KAZAKOVA, Shcherbakovite—a new mineral. *Doklady Akad*. *Nauk. S.S.S.R.*, 99, 837–841 (1954) (in Russian).

Analysis (by M.E.K.) gave SiO₂ 40.61, $Al_2O_3 0.76$. (Nb, Ta)₂O₅ 10.44, rare earths none, TiO₂ 17.91, ZrO₂ 1.23, Fe₂O₃ 1.080, FeO 0.47, MnO 0.04, MgO 0.26, CaO 0.82, Na₂O 5.82, BaO 6.22, K₂O 12.29, Cl 0.30, H₂O⁺ 0.54, H₂O⁻ 0.10, F none; sum 99.61%. Check determinations by O. P. Ostrogorskaya gave Na₂O 5.69, K₂O 12.64%. This corresponds to (K, Na, Ba)₃ (Ti, Nb)₂(Si₂O₇)₂, with K+Ba:Na nearly 2:1. Spectroscopic study by L. N. Indichenko showed also moderate V and Sr, weak Sn, and traces of Pb and Cu. The mineral is insoluble in HNO₃ and HCl, partly soluble in H₂SO₄ on heating.

Shcherbakovite occurs in monoclinic crystals, elongated prismatic, length up to 1.5-2 cm., width 0.3–0.8 cm., and thickness 0.05–0.2 cm. Two-circle goniometric measurements are given for forms a(100), b(010), c(001), m(110), $S(\overline{1}01)$, and n (hk0, uncertain, nearest to (310)). Only a few terminated crystals were observed. The measurements give a:b:c = 0.70:1:?, beta 126° 33″. Two cleavages were observed under the microscope. X-ray powder data are given; the strongest lines in Å were: 2.90 10, 1.081 10, 2.64 7, 1.688 7 1.576 6.

The mineral is dark brown with luster vitreous, greasy on fracture surface. Brittle. H.= $6\frac{1}{2}$, G.=(pycnometer) 2.968. It is biaxial, negative, with ns, α =1.707, β =1.745, γ =1.776, 2V 82°. Elongation positive. Distinctly pleochroic, Z brownish-yellow, Y yellow, X pale yellow. Opaque in hand specimen to translucent on thin edges.

The mineral occurs in a pegmatite in alkalic rocks (rischorrite), associated with and including natrolite-pectolite aggregates. Other minerals present include albite, K feldspar, astrophyllite, apatite, galena, sphalerite, and molybdenite. The locality is not given specifically, but one table is headed "Shcherbakovite from Khibina Tundra." The name is for D. I. Shcherbakov, Russian mineralogist and geochemist.

M. F.