

SHORT COMMUNICATIONS

MINERALOGICAL MAGAZINE, DECEMBER 1977, VOL. 41, PP. 531-4

Amphibole in the Mayo Belwa meteorite: first occurrence in an enstatite achondrite

EXTRATERRESTRIAL examples of the amphibole richterite $[\text{Na}_2\text{CaMg}_5\text{Si}_8\text{O}_{22}(\text{OH},\text{F})_2]$ have previously been reported from two chemical group IA iron meteorites, Wichita County (Olsen, 1967) and Cañon Diablo (Olsen *et al.*, 1973), and from the enstatite chondrite, Abee (Douglas and Plant, 1968). This paper reports an occurrence of richterite in an enstatite achondrite.

On 3 August 1974 a meteorite fell at Mayo Belwa ($8^\circ 58' \text{N.}$, $12^\circ 5' \text{E.}$) in the Adamawa district of Nigeria, and was quickly recovered. An examination of the mineralogy and chemistry revealed it to be an enstatite achondrite which had been heavily shocked (Graham *et al.*, 1977). The 4.85 kg stone is an imperfect ovoid (24 cm \times 13 cm) with a thin transparent fusion crust. On the surface, and throughout the body of the stone, are numerous irregular vugs up to 1 cm in diameter in which richterite occurs as bundles of radiating fibrous needles up to 3 mm long. The vugs are lined with a white, sugary crystalline crust of mainly enstatite and diopside with some cristobalite.

A few amphibole needles were isolated and a powder diffraction pattern obtained using a 114.6 mm diameter Debye-Scherrer camera. The results are shown in Table I compared with the data obtained by Olsen (1967) for Wichita County richterite. Using a refinement programme based on the method of Hess (1951), and developed by Dr. R. J. Davis (Min. Dept. BM(NH)), the following cell parameters were obtained: a $9.845 \pm 0.01 \text{ \AA}$, b $17.999 \pm 0.02 \text{ \AA}$, c $5.258 \pm 0.01 \text{ \AA}$, β $104.41 \pm 0.005^\circ$, V 902.45 \AA^3 , and $a \cdot \sin \beta$ $9.535 \pm 0.01 \text{ \AA}$. These dimensions are very similar to those obtained by Huebner and Papike (1970) by refinement of Olsen's (1967) data, and suggest that the amphibole is fully fluorinated.

Four amphibole needles, some 1 mm in length, including a needle identified by X-ray diffraction as richterite (X-ray no. 18758), were analysed by electron-microprobe. The methods and results are given in Table II, together with analyses of the three previously known meteoritic amphiboles (Olsen *et al.*, 1973). The Mayo Belwa amphibole shows both intra- and inter-grain homogeneity, and compositions (Table II) are closer to that of ideal richterite than any analyses of natural occurrences of this amphibole quoted in the literature (Deer, Howie, and Zussman, 1963). Grains from the white crystalline crust lining the vugs were also analysed and found to be two distinct pyroxenes, both near end-members, with sharp boundaries one against the other. Enstatite (96_{en} ; 3.6_{di}) cores are surrounded by diopside (89.9_{di} ; 8.2_{en} ; 2.0_{jd}) rims in apparent structural continuity. Their compositions correspond with those expected for diopside and enstatite coexisting in equilibrium according to experimental data determined for the system $\text{MgSiO}_3\text{--CaMgSi}_2\text{O}_6$ (Boyd and Schairer, 1964). This sequence of crystallization is compatible with that found by Grigor'eva *et al.* (1971) in melting experiments in the system $\text{Ca}_2\text{Mg}_5\text{Si}_8\text{O}_{22}\text{F}_2\text{--Na}_2\text{Mg}_6\text{Si}_8\text{O}_{22}\text{F}_2$. The observed order of formation and compositions of the pyroxenes lining the vugs thus suggest that they crystallized from a liquid melt rather than from the vapour phase.

TABLE II. *Electron-microprobe analyses of meteoritic amphiboles*

	1	2	3	4	5	6
SiO ₂	58.23	58.09	57.2	57.9	57.8	Number of ions on the basis of 24 (O,F)
TiO ₂	0.21	0.18	2.2	0.01	1.5	
Al ₂ O ₃	0.15	0.17	0.38	0.15	0.4	
Cr ₂ O ₃	0.07	0.06	3.5	0.00	0.7	Si 8.029
FeO	0.01	0.01	0.44	0.17	0.5	Al — } 8.03
MnO	0.02	0.00	0.05	0.00	0.00	Al 0.027
MgO	24.20	24.34	20.9	26.1	22.4	Ti 0.019 } 5.05
CaO	6.69	6.73	3.7	5.5	5.8	Mg 5.009
Na ₂ O	7.22	7.21	8.3	7.3	7.5	Ca 0.995 } 1.00
K ₂ O	0.48	0.47	1.2	0.36	0.6	Na 1.928
Cl	0.02	0.03	n.f.	n.f.	n.f.	K 0.082 } 2.01
F	3.91	3.91	3.6	4.6	2.3	F 1.707 } 1.71*
Sum	101.21	101.20	101.5	102.1	99.5	
Less O ≡ F	1.61	1.61	1.5	1.9	0.9	
Total	99.60	99.59	100.0	100.2	98.6	

1. Mayo Belwa richterite; grain from X-ray mount no. 18758.
 2. Mayo Belwa richterite; average of three other grains.
 3. Cañon Diablo richterite
 4. Abee richterite
 5. Wichita County richterite
 6. Molecular formula of Mayo Belwa amphibole 2.
- } Analyses from Olsen *et al.* (1973).

Electron-microprobe analyses made in the Department of Mineralogy, BM(NH); using analysed minerals and pure metals as standards, and the results were corrected according to the method outlined by Sweatman and Long (1969) and Mason *et al.* (1969). F by electron-microprobe in the Department of Geology and Mineralogy, University of Oxford, using a fluorite standard.

Total Fe expressed as FeO.

n.f. not found.

* It is assumed that the amphibole is completely fluorinated; this low figure is due to analytical constraints on accuracy for this element.

enstatite achondrites, which suggests that little or no partial melt has been lost due to the shock episode.

Acknowledgements. The authors thank the Director of the Geological Survey of Nigeria (Kaduna South) for making the material available; the Department of Geology and Mineralogy, University of Oxford, for the use of their Microscan 9 for fluorine determinations; Drs. J. E. Chisholm, A. L. Graham, and R. Hutchison for advice and critically reading the manuscript.

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[Manuscript received 17 November 1976]

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MINERALOGICAL MAGAZINE, DECEMBER 1977, VOL. 41, PP. 534-7

Formation of analcime in the Dippin sill, Isle of Arran

THE Dippin sill is a Tertiary differentiated basic intrusion, 43 m thick, occurring in the SE. part of the Isle of Arran, Scotland. The sill is composed of different varieties of analcime olivine dolerite, particularly crinanite in which the analcime occurs most commonly as clear, wedge-shapes in the interstices between plagioclase laths. The interstitial analcime is characterized by low SiO_2 and Na_2O and high CaO (e.g. $\text{Ne}_{66.4}\text{Ks}_{0.2}\text{Qz}_{24.3}\text{An}_{9.1}$, wt%) and is occasionally associated with small (< 0.1 mm) nepheline grains, which are comparatively SiO_2 -rich (e.g. $\text{Ne}_{73.4}\text{Ks}_{8.1}\text{Qz}_{16.9}\text{An}_{1.6}$). Analcime also occurs along innumerable small cracks in plagioclase and as a major constituent of large patches, up to 1 cm across, in which it may be associated with brown fibrous natrolite and radiating aggregates of other zeolites. Rocks from the chilled margins of the sill are characteristically rich in lath-shaped analcime pseudomorphs (after plagioclase) and the analcime-bearing patches; the abundances of both decrease inwards from the contacts.

There can be little doubt that the analcime present as pseudomorphs and that in the patches was formed by the action of hydrothermal solutions at subsolidus temperatures. However, the origin of the wedge-shaped, interstitial analcime is more problematical. Many petrologists have interpreted the textural relations as being indicative of a late-stage 'primary' mineral, i.e. crystallized from the melt (e.g. Scott, 1916; Tyrrell, 1928; Harker, 1954, fig. 45b; Wilkinson, 1958). However, the analcime stability curve at low pressures in the system Ne-Ab- H_2O (Greenwood, 1961; Kim and Burley, 1971) lies well below the solidus for an alkaline olivine basalt and below the nepheline-alkali-feldspar minimum melting curve (fig. 1). Indeed, Roux and Hamilton (1976) showed that in the Ne-Ab- H_2O and Ne-Or-Ks- H_2O systems the assemblage analcime plus melt occurs only at pressures between 5 and 13 kb and between 640° and 600°C . Geological evidence suggests that the Dippin sill is unlikely to have been emplaced at a depth of more than 3 km below the contemporary surface, in which case it is likely to have crystallized at a confining pressure of less than 1 kb. Thus the interstitial analcime must be secondary unless its comparatively high Ca content has substantially increased its thermal stability (Roux and Hamilton, 1976). We have tested this possibility by carrying out experiments on a fresh crinanite sample (AC 649) in which the bulk of the analcime is of the interstitial variety.

Rock powder and glass made from this by melting at 1200°C for 15 hours at a log P_{O_2} of -6.4 and a pressure of 1 bar were used as starting materials in the experiments. Samples of each were sealed with excess water in separate gold capsules and were crystallized at pressures from 0.5 to 2.0 kb for 4-6 days; no attempt was made to buffer the P_{O_2} . The presence of analcime in the