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 $[Na_2CaMg_5Si_8O_{22}(OH,F)_2]$ 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 ($8^{\circ} 58'$ N., $12^{\circ} 5'$ 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 ($24 \text{ cm} \times 13 \text{ cm}$) 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: $a 9.845 \pm 0.01$ Å, $b 17.999 \pm 0.02$ Å, $c 5.258 \pm 0.01$ Å, $\beta 104.41 \pm 0.005^{\circ}$, V 902.45 Å³, and $a.sin\beta 9.535 \pm 0.01$ Å. 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 I 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 (96·4en; 3·6di) cores are surrounded by diopside (89·9di; 8·2en; 2·0id) 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 $MgSiO_3$ —Ca $MgSi_2O_6$ (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 $Ca_2Mg_5Si_8O_{22}F_2 - Na_2Mg_6Si_8O_{22}F_2$. 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.

	Wichita County Olsen, 1967)		o Belwa (th	is study)		Wichita County (Olsen, 1967)		Mayo Belwa (this study)			hkl
I/Io d _{obs} (Å) (visual)		I/Io d _{obs} (Å) (visual)		d_{calc} (Å)	hkl	$\frac{1/Io}{\text{(visual)}} d_{\text{obs}} (\text{\AA})$		I/Io d _{obs} (Å) (visual)		$d_{\rm calc}$ (Å)	
	0.6	20	9.03	9.000	020	70	2.160	60	2.164	{ 2·164	132
70	8.605	6.	0	9		,			•	2:161	261
70	8.393	60	8.47	8.426	110		0	10	2.134	2.134	<u>3</u> 32
30	4.864	40	4.85	4 ^{.8} 57	IIĪ	40	2.048	40	2.021	2.042	202
50	4.493	50	4.20	4.200	040	10	2.020	40	2.018	2.018	351
70	4.160				_			40	2.004	∫ 2.001	40 <u>2</u>
30	4.004	30	4.00	{ 4·015	20Ĩ			т-		1.999	370
30	• •	-	•	3.987	III					1.9591	42 <u>2</u>
50	3.850	40	3.867	3.861	13Ī	10	1.952	30	1.9580	{ 1.9573	190
				{ 3 [.] 379	131					1.9504	152
100	3.376	80	3.389	3.372	041					1.9320	421
				3.368	150			30	1.9305	{ 1.9305	26ž
80	3.267	70	3.276	3.272	240					{ 1·9286	352
100	3.125	90	3.132	3.130	310	20	1.893	40	1.8982	1.8964	510
		20	3.026	3.017	311	IO	1.857	40	1.8617	∫ 1·8614	242
80	2.944	60	2.948	2.944	22 I	10	1.05/	40	1.901	<u>)</u> 1·8591	19Ï
40	2.800	30	2.808	2.809	330			20	1.8328	1.8332	44Ž
100	2.709	100	2.706	2.702	151	10	1.794	20	1.7965	1.7966	191
70	2.281	50	2.587	2.585	061					{ I·7422	203
90	2.522	70	2.527	2.522	202			20	1.7423	{ I·7414	512
	2.382		2.280	{ 2·384	400					1.7399	113
20	2.302	30	2.389	2.383	350					(1.6789	28ž
				{ 2·332	35Ī		- (-(- (1.6783	133
50	2.320	50	2.328	2.319	42 Ī	30	1.676	40	1.6774	1.6741	313
50	2.286	40	2.290	2.291	171					1.6741	39Ī
40	2.258	40	2.266	2.266	312	6-	- (6.	- (1 6514	461
20	2.203	20	2.205	2.200	242	60	1.620	60	1.6530	1.6487	511
	5		5			Plus 27 addi- tional spacings to 0.7882 Å			Plus 60 a spacings 0.7810 Å		5

TABLE I. X-ray diffraction patterns of Mayo Belwa and Wichita County richterite

The three previously recorded occurrences of meteoritic richterite (Olsen *et al.*, 1973) are in association with metal: in the Type I enstatite chondrite, Abee, the amphibole occurs within 'podlike segregations of metal', and Wichita County and Cañon Diablo are both coarse octahedrites. The enstatite achondrites, however, contain little metal (< 0.2 wt%) suggesting that the Mayo Belwa richterite crystallized in a near metal-free environment. Its chemistry reflects this: Fe, Cr, and Mn are present in trace amounts, if at all, whereas the earlier described occurrences contain significant amounts of these elements. Furthermore, its composition is much closer to that of 'end-member' richterite than any of the other three examples.

This chemical 'purity' of the amphibole can be explained by the operation of a virtual distillation process within the meteorite. Fluor-richterite had presumably been formed as a residual phase during the meteorite's pre-shock history. Enstatite, plagioclase, and this existing fluor-compound were mobilized during shock melting, producing fluorine-filled vugs lined with a liquid melt. After crystallization of enstatite followed by diopside, reaction of the sodium-enriched melt with the fluorine vapour produced the radiating sodic amphibole needles, leaving a small amount of residual cristobalite. The bulk analysis of Mayo Belwa is typical of the

	I	2	3	4	5	6	
SiO ₂	58.23	58.09	57.2	57.9	57.8	Number of ions on th	
TiO ₂	0.51	0.18	2.2	0.01	1.2	basis (of 24 (O,F)
Al_2O_3	0.12	0.12	0.38	0.12	0.4		
Cr_2O_3	0.07	0.06	3.5	0.00	0.7	Si	8.029
FeO	0.01	0.01	0.44	0.12	0.5	Al	<u>-</u> 8.03
MnO	0.05	0.00	0.02	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	5.2	5.8	Mg	5.009
Na ₂ O	7.22	7.21	8.3	7.3	7.5	Ca	0.995 1.00
K₂Ō	0.48	0.47	I·2	0.36	0.6	Na	1.928
CĨ	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	I·707 I·7I*
Sum	101.51	101.50	101.2	102·1	99 [.] 5		
Less $O \equiv F$	1.61	1.61	1.2	1.9	0.9		
Total	99.60	99.59	100.0	I00·2	98·6		

TABLE II. Electron-microprobe analyses of meteoritic amphiboles

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 Analyses from Olsen et al. (1973).

5. Wichita County richterite

6. Molecular formula of Mayo Belwa amphibole 2.

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.

Department of Mineralogy (British Museum (Natural History)), Cromwell Road London SW7 5BD A. W. R. BEVAN J. C. BEVAN J. G. FRANCIS

REFERENCES

Boyd (F. R.) and Schairer (J. F.), 1964. Journ. Petrology, 5, 275-309.

Deer (W. A.), Howie (R. A.), and Zussman (J.), 1963. Rock-Forming Minerals, 2. Longmans, London.

Douglas (J. A. V.) and Plant (A. G.), 1968 (abstr.). Meteoritics, 4, 166.

Grigor'eva (L. F.), Chigareva (O. G.), Mikirticheva (G. A.), and Krupenikova (Z. F.), 1971. Phys. and Chem. of Asbestos Minerals. 2nd Internat. Conf. Louvain Univ. Paper 1:6.

Graham (A. L.), Easton (A. J.), and Hutchison (R.), 1977. Mineral Mag. 41, 487.

Hess (J. B.), 1951. Acta Cryst. 4, 209-15.

Huebner (J. S.) and Papike (J. J.), 1970. Amer. Min. 55, 1973-92.

SHORT COMMUNICATIONS

Mason (P. K.), Frost (M. J.), and Reed (S. J. B.), 1969. Nat. Phys. Lab. I.M.S. Report 1. Olsen (E.), 1967. Science, **156**, 61-2. — Huebner (J. S.), Douglas (J. A. V.), and Plant (A. G.), 1973. Amer. Min. **58**, 869-72. Sweatman (T. R.) and Long (J. V. P.), 1969. Journ. Petrology, **10**, 332-76.

[Manuscript received 17 November 1976]

C Copyright the Mineralogical Society.

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, wedgeshapes in the interstices between plagioclase laths. The interstitial analcime is characterized by low SiO₂ and Na₂O and high CaO (e.g. Ne_{66·4}Ks_{0·2}Qz_{24·3}An_{9·1}, wt%) and is occasionally associated with small (< 0·I mm) nepheline grains, which are comparatively SiO₂-rich (e.g. Ne_{73·4}Ks_{8·1}Qz_{16·9}An_{1·6}). Analcime also occurs along innumerable small cracks in plagioclase and as a major constituent of large patches, up to I 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-H₂O and Ne-Or-Ks-H₂O 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 I 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 -6.4and 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

534