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Study of minerals from the pegmatites of the Nellore mica-belt, Andhra Pradesh, India Part III—Biotite

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SUMMARY. The biotites in the pegmatites are Fe²⁺ type. X-ray analyses indicated that they are 1M or 3T polymorphs. In the colour of the biotite iron plays an important role. From the Fe/(Fe+Mg) ratios, the analysed biotites are assumed to have crystallized between 760 and 500 °C.

In the pegmatites of the Nellore mica-belt, biotite occurs in border, wall, and outerintermediate zones; it is black in thick sheets, brown in thin flakes, and yellowishbrown in films, and ranges in size from a few millimetres to tens of centimetres across. The specific gravity is from 3.1 to 3.3. It is pseudo-uniaxial, β from 1.620 to 1.635.

Inclusions of ruby muscovite occur along the basal cleavage planes in biotite books (fig. 1).

Chemical analyses of two biotites, with structural formulae, are given in table I. When the percentages of Mg, $R^{3+}(Al+Fe^{3+}+Ti)$, and $(Fe^{2+}+Mn^{2+})$ of the octahedral layer in the biotites (B1, B2) are plotted in the triangular diagram of Foster (fig. 11, p. 25, 1960), they are in Fe²⁺ biotite field.

The unit cell parameters of the two chemically analysed biotites calculated from powder data are given in table I; the reflections can be indexed

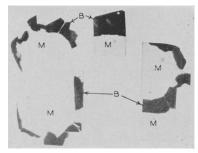


FIG. 1. Biotite-muscovite intergrowths. B, biotite; M, muscovite. $\times \frac{1}{2}$.

equally well on a 1M or 3T cell (Yoder and Eugster, 1954).

According to Hall (1941) the colour of biotite depends on the content of FeO (total), MgO, and TiO₂. When the analysed biotites are plotted on the triangular diagram of Hall (fig. 1, p. 30) a brown colour is indicated, which is the observed colour in flakes. Hayama (1959) considered TiO₂ and the ratio $Fe_2O_3/(Fe_2O_3+FeO)$ the main factors; in his diagram the biotite (B1) falls in the region of yellowish- to greenish-brown, whereas (B2) is in the dark brown to brown region. These plottings indicate that iron plays an important role in the colour of the biotite is mainly caused by electron transfer between Fe^{3+} and Fe^{2+} . But the effect of other elements must also be considered.

Based on the theoretical work of Fersman (1960), the biotite in the area of study, Fe^{2+} -rich type, is assumed to have formed between 700 and 600 °C. According to

Wones (1963), the $Fe^{2+}/(Fe^{2+}+Fe^{3+})$ ratio may serve as an indicator of variations in the oxygen fugacity of the environment in which a biotite crystallizes; this ratio for the two biotites (table II) is near to Ni–NiO (0·10) and Fe_2SiO_4 – Fe_3O_4 – SiO_2 (0·05) buffer conditions (Wones and Eugster, 1965). The Fe/(Fe+Mg) ratios of the analysed biotites

	BI B2		Number of ions on the basis of 12 (O, OH, F) B1 B2		Number of ions on the basis of 11 oxygens B1 B2		
	I·44 20·91 8·12 — 0·82	36.12 19.36 1.68 2.27 20.80 7.85 0.17 0.75 0.43 9.32 0.93 	Si Al ^{ivi} Fe ³⁺ Ti Fe ²⁺ Mg Mn Ca Na K OH F O	$\begin{array}{c} 2 \cdot 509 \\ 1 \cdot 491 \\ 0 \cdot 609 \\ 0 \cdot 200 \\ 0 \cdot 086 \\ 1 \cdot 378 \\ 0 \cdot 965 \\ \hline \\ 0 \cdot 057 \\ 0 \cdot 057 \\ 0 \cdot 836 \\ 0 \cdot 124 \\ 1 \cdot 2000 \\ 1 \cdot 2000 \\ 0 \cdot 960 \\ 0 \cdot 960 \\ 0 \cdot 124 \\ 1 \cdot 2000 \\ 1 \cdot$	$\begin{array}{c} 2\cdot865\\ 1\cdot135\\ 0\cdot673\\ 0\cdot103\\ 0\cdot136\\ 1\cdot382\\ 0\cdot927\\ 0\cdot011\\ 0\cdot062\\ 0\cdot063\\ 0\cdot945\\ 0\cdot945\\ 0\cdot494\\ -\\ 11\cdot506\\ \end{array}\right\}1\cdot068$	$\begin{array}{c} 2 \cdot 396 \\ 1 \cdot 604 \\ 0 \cdot 410 \\ 0 \cdot 192 \\ 0 \cdot 082 \\ 1 \cdot 232 \\ 0 \cdot 915 \\ \end{array} \right) 2 \cdot 921 \\ 2 \cdot 921 \\ 2 \cdot 921 \\ 0 \cdot 915 \\ 0 \cdot 915 \\ 0 \cdot 915 \\ 0 \cdot 925 \\ 0 \cdot $	$\begin{array}{c} 2 \cdot 682 \\ 1 \cdot 318 \\ 0 \cdot 372 \\ 0 \cdot 096 \\ 0 \cdot 128 \\ 1 \cdot 293 \\ 0 \cdot 867 \\ 0 \cdot 011 \\ \end{array} $ $\begin{array}{c} 2 \cdot 767 \\ 2 \cdot 767 \\ 0 \cdot 011 \\ 0 \cdot 058 \\ 0 \cdot 059 \\ 0 \cdot 885 \end{array} $ $\begin{array}{c} 1 \cdot 002 \\ 0 \cdot 885 \\ 1 \cdot 002 \\ 0 \cdot 885 \end{array}$
a b	5·312 9·225	5·300 9·210)	· · · · · ·	+0.610 +0.228
ς β	10°230 100° 3'	10·160 99° 3′					+0·991 +1·060

TABLE I. Analyses of biotites from Madiga Inam mine, Kalichedu, Nellore

TABLE II. Temperature of crystallization of biotites

	Analyse	d biotites	Synthetic biotites*		
	BI	B2	Ni–NiO buffer	$Fe_2SiO_4-SiO_2Fe_3O_4$ buffer	
$\overline{Fe^{3+}/(Fe^{2+}+Fe^{3+})}$	0.1283	0.06717	0.10	0.02	
Fe/(Fe+Mg)	0.7968	0.7945	0.765	0.765	
d_{060}	1.2372	1.5350	1.2211-1.2218	1.5509-1.5528	
Temperature	—	<u> </u>	700–600 °C	760–500 °C	
Pressure		_	2070 bars	2070 bars	

* Wones, 1963.

plotted in the figure of Wones and Eugster (fig. 4, p. 1244, 1965) indicate a crystallization temperature in the range of 760–500 $^{\circ}$ C at 2070 bars pressure, depending upon the oxygen fugacity. It may be concluded that the biotite under study crystallized between 760 and 500 $^{\circ}$ C.

Heinrich (1946), Nockolds (1947), Gower (1957), Engel and Engel (1960), and Foster (1960) studied the variation in composition of biotites in relation to their parent rock type. In the diagram of Engel and Engel (fig. 7, p. 30, 1960), in the diagram

of Gower (fig. 5, p. 150, 1957), and in that of Foster (fig. 12, p. 33, 1960), both analysed biotites fall in the granite field.

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REFERENCES

ENGEL (A. E. J.) and ENGEL (C. G.), 1960. Bull. Geol. Soc. Amer. 71, 1-58.

[FERSMAN (A. É.)] Ферсман (А. Е.), 1960. Избранные труды [selected works], 6. Изд. Акад. наук СССР, Moscow.

FOSTER (M. D.), 1960. U.S. Geol. Surv. Prof. Paper, 354 B.

FYFE (W. S.), 1964. Geochemistry of Solids: An Introduction. New York (McGraw-Hill).

Gower (J. A.), 1957. Amer. Journ. Sci. 255, 142-50.

HALL (A. J.), 1941. Amer. Min. 26, 29-34.

HAYAMA (Y.), 1959. Jour. Geol. Soc. Japan, 65, 21-30.

HEINRICH (E. Wm.), 1946. Amer. Journ. Sci. 244, 836-48.

NOCKOLDS (S. R.), 1947. Ibid. 245, 401-20.

WONES (D. R.), 1963. Amer. Min. 48, 1300-21.

— and EUGSTER (H. P.), 1965. Ibid. 50, 1228-72.

YODER (H. S.) and EUGSTER (H. P.), 1954. Geochimica Acta, 6, 157-85.

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