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## References

- AKIMOTO, S. AND T. KATSURA (1959) Magnetochemical study of the generalized titanomagnetite in volcanic rocks. *Geomag. Geoelectr.* 10, 69–90.
- AOKI, K. (1959) Petrology of alkali rocks of the Iki Islands and Higashi-matsuura District, Japan. Sci. Rep. Tohoku Univ. Ser. III, 6, 261–310.
- —— (1963a) The kaersutites and oxykaersutites from alkalic rocks of Japan and surrounding areas. Jour. Petrology 4, 198–210.
- (1963b) On the anorthoclase feldspars of the alkalic rocks from the Iki and adjacent Islands. Jour. Japan. Assoc. Mineral. Petrol. Econ. Geol. 50, 77-80 (in Japanese, Engl. abs.).

(1964) Clinopyroxenes from alkaline rocks of Japan. Am. Mineral. 49, 1199-1223.

- BASTA, E. Z. (1960) Natural and synthetic titanomagnetites (the system Fe<sub>3</sub>O<sub>4</sub>-Fe<sub>2</sub>TiO<sub>4</sub> -FeTiO<sub>3</sub>). Neues Jb. Mineral., Abh., Bd. 94, 1017-1048.
- DEER, W. A., R. A. HOWIE AND J. ZUSSMAN (1962) Rockforming Minerals. Vol. 5, Nonsilicates. Longmans Co. Ltd., London.
- KATSURA, T. AND I. KUSHIRO (1961) Titanomaghemite in igneous rocks. Am. Mineral 46, 134-145.
- KAWAI, N., S. KUME AND S. SASAJIMA (1954) Magnetism of rocks and solid phase transformation in ferromagnetic minerals. Proc. Japan. Acad. 30, 588,
- MATSUI, K. (1958) Explanatory text of the geological map of Japan, Katsumoto, Gononra and Ashibe. *Geol. Surv. Japan* (in Japanese, Engl. abst.).
- VINCENT, E. A., J. B. WRIGHT, R. CHEVALLIER AND S. MATHIEU (1957) Heating experiments on some natural titaniferous magnetites. *Mineral. Mag.* 31, 624-655.
- WRIGHT, J. B. (1959) Some further heating experiments on natural titaniferous magnetites. Mineral. Mag. 32, 32–37.

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## CELL DIMENSIONS AND SPACE GROUP OF TAMARUGITE

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Tamarugite, NaAl(SO<sub>4</sub>)<sub>2</sub>·6H<sub>2</sub>O, is a secondary mineral formed from the oxidation of sulfides, usually under arid conditions. Optical examination of crystals found with sideronatrite in a sample from Mina de la Compania, Sierra Gorda, Chile (U. S. National Museum, Smithsonian Cat. No. R6287) revealed  $\alpha$ =1.485,  $\beta$ =1.487 and  $\gamma$ =1.498, all ±0.002. These thus agree with the values,  $\alpha$ =1.484,  $\beta$ =1.486,  $\gamma$ =1.497, all ±0.001—reported by Gordon (1940). Similarly, the morphological and physical properties observed in the tamarugite of this present study confirm those reported by Gordon (1940). Published *x*-ray data on tamarugite were not found by the writers. Thus it was necessary to determine

d(obs)	$I/I^1$	d(calc)	hkl	d(obs)	$I/I^1$	d(calc)	hkl
12.538	16	12.612	020	2.544	8	2.545	091
7.034	6	7.033	110	2.521	4	2.522	0.10.0
6.339	2	6.333	120	2.509	2	2.513	$15\bar{2}$
5.530	8	5.523	130	2.462	9	2.462	062
4.923	6	4.923	031	2.350	2	2.350	232
4.557	4	4.563	12T	2.333	4	2.330	$0 \cdot 10 \cdot 1$
4.424	3	4.413	111	2.329	6	2.328	311
4.370	4	4.374	041	2.300	6	2.299	321
4.223	100	4.223	121	2.281	3	2.281	242
4.207	80	4.204	060	2.269	4	2.269	281
4.157	12	4.155	150	2.228	4	2.226	290
3.964	32	3.955	131	2.189	7	2.188	1.11.0
3.861	2	3.866	141			2.190	311
3.647	59	3.646	160	2.116	3	2,115	262
3.355	3	3.351	151	2,102	6	2.102	0.12.0
3.266	3	3.267	201	2.060	2	(2.060	182
3.170	10	3.167	240			2.059	092
3.162	12	3.163	221	2.045	3	2.044	361
3.153	21	3.153	080	2.025	2	(2.024	003
3.036	5	3.036	002			2.024	272
2.949	2	2.952	022	1.00.000	1	1.968	033
		(2.896	180	1,969		1.967	322
2.899	23	2.901	241			1.904	$14\overline{3}$
		2.902	171	1.904	3	{1.903	$1 \cdot 10 \cdot \overline{2}$
2.876	5	2.879	112			1.902	113
2.823	2	2.824	122			1.876	1.13.0
2.766	5	2.761	260	1.876	2	1.877	371
2.739	3	2.740	132			1.877	2.11.1
2.733	4	2.736	042	1.831	2	1.831	400
2.711	7	2.705	112			1.830	0.11.2
2.604	5	2.602	052	1.702	3	1.702	$1 \cdot 12 \cdot \overline{2}$
		∫2.580	261			1.703	083
2.582	5	2.579	181	1.679	2	1.679	460

TABLE 1. INDEXED POWDER DATA FOR TAMARUGITE

the cell dimensions and space group of tamarugite as a first step in a contemplated crystal-structure analysis.

Weissenberg photographs of the h0l, h1l, h2l, and hk0 reflections for copper K $\alpha$  radiation were made for two single crystals of tamarugite, both from specimen R6287. The indexing of these photographs showed that the only systematic extinctions were of the type  $k \neq 2n$  for (0k0). Accordingly the space group is  $P2_1$  or, if the crystal is centrosymmetrical,  $P2_1/m$ . The latter agrees with the 2/m crystal class reported by Palache *et al.* (1951) on the basis of morphologic data. Accurate lattice parameters were obtained by use of a least squares FORTRAN IV program written by Fang and Wolf following the scheme proposed by Main and Woolfson (1963) in which  $\alpha_1$ - $\alpha_2$  separations in the high angle regions of zero level Weissenberg photographs are utilized. The cell parameters were then slightly refined so as to minimize the difference between d(calc) and d(obs) for the powder diffraction data. The cell parameters thus obtained are:  $a=7.35_3$ ,  $b=25.22_5$ ,  $c=6.09_7$  and  $\beta=95^{\circ}12'$ . Ungemach (1935) reports  $\beta$  to be 94°50' on the basis of morphologic data.

The amount of tamarugite in specimen R6287 was too small for a powder pattern to be obtained other than the powder photograph earlier made by Robinson (1962). However, the U. S. National Museum kindly supplied two specimens of tamarugite—Nos. C4703 from Cerro Pintados, Chile and R9117 from Alcaparrosa, Chile—for which powder patterns were obtained in a Norelco wide-angle diffractometer using CuK $\alpha$  radiation. The powder patterns for these two specimens appeared indistinguishable. The pattern obtained from a  $1/8^{\circ}$  (2 $\theta$ ) per minute scan of the Cerro Pintados material, internally calibrated with spinel, was satisfactorily indexed on the basis of the cell parameters reported above (Table 1). The observed *d* spacings for the Cerro Pintados pattern closely matched those measured by Robinson (1962) from a powder photograph of tamarugite collected from the Sierra Gorda specimen.

The axial ratios calculated from the lattice parameters here reported that is, 0.2915:1:0.2417—compare with those of Ungemach (1935) namely, 0.2919:1:0.2415.

The calculated density of tamarugite, assuming Z=4, is 2.066 gm/cm<sup>3</sup>. The measured density, by the flotation method, is  $2.06\pm0.01$  gm/cm<sup>3</sup>. Gordon (1940) and Ungemach (1935) report the density of tamarugite to be 2.07 gm/cm<sup>3</sup>.

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## References

- GORDON, S. G. (1940) The identity of lapparentite with tamarugite. Ac. Nat. Sc. Philadelphia. Not. Nat., 57, 1–9.
- MAIN, P. AND M. M. WOOLFSON (1963) Accurate lattice parameters from Weissenberg photographs. Acta. Cryst. 16, 731-733.
- PALACHE, C., H. BERMAN AND C. Frondel (1951) The System of Mineralogy. Vol. 2, 7th ed. John Wiley and Sons, New York.
- ROBINSON, P. D. (1962) An x-ray study of sideronatrite. M.S. thesis, Southern Illinois Univ.
- UNGEMACH, M. H. (1935) Sur certains minéraux sulfatés du Chili. Bull. Soc. Franc. Mineral. 58, 97–221.