

SCAWTITE FROM CRESTMORE, CALIFORNIA

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ABSTRACT

An occurrence of scawtite has been discovered in the contact zone at Crestmore, California. This is the first on record for California, and the fourth known anywhere, to date. The mineral is in bundles of thin tabular crystals, strongly resembling in habit and mode of occurrence the original find at Scawt Hill. Crystals are in general poor, but it has been possible to make approximate measurements on selected individuals, and to confirm these with x-ray study of single crystals. The forms present are as follows: {100}, {110}, {120}, {130}, {010}, and {101}. The *a*-pinacoid is always dominant, with the prisms narrow and poorly developed. The mineral is monoclinic, probable space group *I* 2/*m*, with the following unit cell dimensions:

$$\begin{aligned} a_0 &= 10.22 \text{ \AA}, & b_0 &= 15.42 \text{ \AA}, & c_0 &= 6.70 \text{ \AA} \\ a:b:c & & 0.6628:1:0.4345 & & \beta &= 100^\circ 29' \end{aligned}$$

Powder photographs, which can be completely indexed, show the following spacings and intensities for the stronger lines: 3.04 Å—10, 3.01 Å—5, 2.49 Å—5, 2.236 Å—5, 1.898 Å—6.

The crystals have been oriented with Tilley's {001} as {100}, giving simpler indices for the forms, and a less oblique unit cell.

OCCURRENCE

Specimens collected on the 910 foot level of the Commercial quarry, Crestmore, California, showed a colorless mineral in thin veins in a matrix of massive diopside-wollastonite-spurrite rock. Where the veins are incompletely filled, crystals of this mineral are abundantly developed. Examination showed these crystals to be nearly square tablets with one edge always bevelled, and almost invariably occurring in sub-parallel aggregates, exactly like bundles of shingles. Chemical and optical tests showed the mineral to be scawtite, which has been found in similar crystalline groups only at Scawt Hill, County Antrim, Ireland, Tilley (1930), and Ballycraigy, Larne, N. Ireland, McConnell (1954). One other occurrence is known, but only as microscopic lath-like grains in a thin section of contact zone material from south of Neihart, Montana, Taylor (1935).

The Crestmore occurrence matches closely that described by Tilley, the mineral being one of the latest in the hydrothermal series, and having precisely the same habit. In general, the scawtite is the latest mineral to form in the veins in which it appears, overlying and enveloping calcite crystal. However, in at least one specimen, small bead-like calcite crystals were noted perched on the surface and edges of the scawtite. In another, delicate tufts of white needles and blades of an unidentified mineral occur closely associated with the scawtite, and seem to be essentially

contemporaneous with it. Many scawtite veins show a zone of radiating clusters of a bladed unidentified mineral usually along one side of the vein, upon which the scawtite has been deposited. The blades of the unknown mineral are extremely thin obliquely rhombic plates, with extinction parallel to the longer edge of the rhomb.

Associated with the massive scawtite in one or two of the veins, are fine-grained sugary aggregates of bultfonteinite, which also appears in similar massive veins of afwillite, which was recently identified from Crestmore by Switzer and Bailey (1953). A description of this mineral, with *x*-ray study, is in preparation, and will be presented shortly.

PROPERTIES AND MORPHOLOGY

Sections parallel to the broad plates show parallel extinction, with the acute bisectrix emerging just outside the field of view, and giving a positive figure. $2V$ is large (Tilley notes $74^\circ \pm$.) A fair cleavage is present parallel to $\{010\}$, and sections with this orientation show an extinction angle of approximately 28° . Tilley records a cleavage parallel to $\{100\}$, but this may well be a parting, due to twinning on this face. Many rod-like inclusions with much lower index were observed in some sections. Indices of refraction of the Crestmore material check satisfactorily with recorded values.

The Crestmore scawtite is readily soluble in HCl, with moderate effervescence, leaving a residual skeleton of silica. It gives good microchemical tests for SiO_2 and CaO. In the closed tube the mineral decrepitates, and gives off water. Before the blowpipe it is infusible, but whitens and glows when heated intensely. No strong flame coloration was observable.

Using the large face as the front pinacoid, the following forms were observed: $\{100\}$ always present and dominant, but usually somewhat uneven, and showing thin slabs of subsidiary crystals, $\{010\}$ usually present, very shiny; $\{101\}$ always present, normally shiny, and almost always giving multiple reflections, due to the aggregation of many flat crystals; $\{130\}$ narrow, but often present as one or two faces; $\{120\}$ and $\{110\}$ narrow, and rarely present. Two faces of $\{011\}$ are present on one crystal. It was possible to select crystals which were simple enough to measure on the reflecting goniometer, although without any great degree of precision. Only doubtful numerical indices could be assigned to the prisms from these measurements, but these could be confirmed from the *x*-ray measurements. Using these *x*-ray data, the angle table for observed forms has been calculated, and is presented below in Table 1.

Actual measurements for rho of $\{101\}$ range from $39^\circ 15'$ to $43^\circ 40'$,

with best probable values of about 42° . For $\{130\}$ several readings gave $26^\circ 33'$ or $27^\circ 18'$; for $\{120\}$ $40^\circ 12'$; and for $\{110\}$ $56^\circ 31'$.

TABLE 1. SCAWTITE ANGLE TABLE

Monoclinic, Space Group $I 2/m$ $a_0:b_0:c_0=10.22:15.42:6.70$, $\beta=100^\circ 29'$ $a:b:c=0.6628:1:0.4345$ $p_0:q_0:r_0=0.6556:0.4273:1$ $r_2:p_2:q_2=2.3406:1.5344:1$ $p_0'=0.6667$, $q_0'=0.4345$, $e_0'=0.1850$

	ϕ	ρ	ϕ_2	$\rho_2=B$	A	C
<i>b</i> 010	$90^\circ 00'$	$0^\circ 00'$	—	$0^\circ 00'$	$90^\circ 00'$	$90^\circ 00'$
<i>a</i> 100	90 00	90 00	0 00	90 00	79 31	0 00
<i>k</i> 130	27 05	90 00	0 00	27 05	85 14	62 55
<i>l</i> 120	$37^\circ 29\frac{1}{2}$	90 00	0 00	$37^\circ 29\frac{1}{2}$	$83^\circ 37\frac{1}{2}$	$52^\circ 30\frac{1}{2}$
<i>m</i> 110	$56^\circ 54\frac{1}{2}$	90 00	0 00	$56^\circ 54\frac{1}{2}$	81 12	$33^\circ 05\frac{1}{2}$
011	23 04	25 17	79 20	66 52	80 22	23 08
<i>d</i> 101	90 00	40 25	49 35	90 00	29 54	49 35

One poor prism face with $\phi=18^\circ 03'$ was observed. This might doubtfully be $\{290\}$ (calculated $\phi=18^\circ 50'$).

Transformation formula, Murdoch/Tilley = $00\bar{1}/010/100$.

X-RAY STUDY

By careful selection it was possible to isolate two crystals which were satisfactory for single crystal x-ray measurement, although even these were multiple, but with orientations so slightly varied as not to make trouble. One of these was rotated about c , and the other about b , so that all three directions could be adequately measured. Rotation, equator and first-layer line Weissenberg photographs were taken with filtered iron radiation, and the three cell-dimensions were calculated from these. Translations on the b and c axes were measured on the rotation photographs, and the layer-line pictures measured, using both top and bottom halves of the films. From these measurements, using as many as four or five orders of diffraction, and correcting for lack of uniformity in crystal dimensions, the values of a_0 , b_0 and c_0 were calculated. The values thus arrived at are as follows:

$$a_0 10.22 \pm 01 \text{ \AA}, \quad b_0 15.42 \pm 01 \text{ \AA}, \quad c_0 6.70 \pm 01 \text{ \AA}.$$

μ , measured directly on the Weissenberg film, and checked on the Schneider construction, is $79^\circ 31'$, so that beta is $100^\circ 29'$. Systematic

TABLE 2. X-RAY POWDER DATA FOR SCAWTITE

Cu K α radiation, Angstrom units

d/n		hkl	I	d/n		hkl	I
Obs.	Calc.			Obs.	Calc.		
8.40	8.418	110	1	1.791 Fe	1.801	451	1
7.75	7.71	020	$\frac{1}{2}$		1.789	053	
6.05	6.06	011	3	1.778 Fe	1.782	521	$\frac{1}{2}$
	6.037	101		1.756	1.761	532	$\frac{1}{2}$
5.03	5.11	200	2		1.743	253	
	5.103	101		1.682	1.683	550	$\frac{1}{2}$
4.76	4.778	210	$\frac{1}{2}$	1.640	1.637	620	1
	4.753	121			1.638	014	
4.56	4.576	130	4	1.610	1.618	224	2
4.22	4.256	121	3		1.611	024, 253	
	4.232	211			1.613	631	
	4.211	220			1.609	204	
4.05	4.052	031	$\frac{1}{2}$	1.589 Fe	1.590	314	$\frac{1}{2}$
3.83	3.81	040	2		1.584	512	
3.57	3.594	240	3	1.577	1.576	602	$\frac{1}{2}$
	3.584	211		1.566 Fe	1.572	114	$\frac{1}{2}$
3.357 Fe	3.35	002	$\frac{1}{2}$	1.560	1.553	291	$\frac{1}{2}$
3.24	3.24	112	3	1.543	1.547	381	2
	3.274	310			1.542	0.10.0	
3.04	3.03	022	10	1.530	1.537	282	$\frac{1}{2}$
	3.057	240			1.526	334	
	3.076	141		1.508	1.515	413, 044	1
					1.510	134	
3.017 Fe	3.018	202	5		1.509	404	
2.97 Fe	2.982	321	1	1.486 Fe	1.488	651	$\frac{1}{2}$
	2.963	212		1.479	1.477	1.10.1	1
2.91	2.92	112	$\frac{1}{2}$	1.457	1.459	701	1
	2.90	241			1.460	613	
2.837	2.81	222	$\frac{1}{2}$	1.443	1.442	224	1
2.79	2.806	330	4	1.375	1.376	314	$\frac{1}{2}$
	2.793	051		1.319			1
2.56 Fe	2.57	060	$\frac{1}{2}$	1.270			1
	2.561	312		1.244			$\frac{1}{2}$
	2.555	400		1.237			$\frac{1}{2}$
2.543	2.55	202	2	1.228			$\frac{1}{2}$
	2.526	251		1.212			$\frac{1}{2}$
2.495	2.505	042	5	1.1515			$\frac{1}{2}$
	2.472	411		1.132			$\frac{1}{2}$
2.41	2.422	222	$\frac{1}{2}$	1.111			$\frac{1}{2}$
2.35	2.365	251, I61	$\frac{1}{2}$	1.102			$\frac{1}{2}$
2.307 Fe	2.318	332	$\frac{1}{2}$	1.094			$\frac{1}{2}$
2.285 Fe	2.288	260	$\frac{1}{2}$	1.0887			$\frac{1}{2}$
2.24	2.251	431	5	1.0687			$\frac{1}{2}$
	2.259	341		1.0578			$\frac{1}{2}$
2.224 Fe	2.232	103	1				$\frac{1}{2}$
2.199 Fe	2.200	402	$\frac{1}{2}$	1.0375			$\frac{1}{2}$
	2.197	152		1.008			$\frac{1}{2}$
2.175	2.174	013	$\frac{1}{2}$.9836			$\frac{1}{2}$
2.164	2.151	170	$\frac{1}{2}$.9277			$\frac{1}{2}$
2.13	2.141	213	2	.8882			$\frac{1}{2}$
	2.1395	312		.7890			$\frac{1}{2}$
2.107	2.107	440	3	.7780			$\frac{1}{2}$
	2.036	431					$\frac{1}{2}$
2.007 Fe	2.012	303					$\frac{1}{2}$
1.938	1.944	451	$\frac{1}{2}$				$\frac{1}{2}$
1.898	1.927	080	6				$\frac{1}{2}$
	1.891	271					$\frac{1}{2}$
1.871	1.876	213	1				$\frac{1}{2}$
	1.872	530					$\frac{1}{2}$
1.817	1.811	262	$\frac{1}{2}$				$\frac{1}{2}$

extinctions in the chosen orientation, leaving only $h+k+l$ even, indicate a body-centered lattice. Other possible settings, with beta about 107° , or Tilley's orientation, with beta $115^\circ 24'$ would give a base centered lattice, but the first was chosen as following a common custom of employing the unit cell with least possible obliquity. In addition this orientation gives simpler indices.

Powder photographs were taken with copper and with iron radiations and can be satisfactorily indexed. In the following, Table 2, giving the spacings and intensities, the symbol Fe indicates the presence of multiple lines shown on the copper film as single, or broadened lines, which have been resolved by the iron radiation. For example, a very strong, broad line on the copper film at about 3.04 \AA is resolved in the iron into a definite triplet, 10, 5 and 1 in intensity.

Independently of the author, Mr. J. D. C. McConnell (this journal, pp. 510-514), working in the laboratory of the Department of Mineralogy at Cambridge University, has completed a study of scawtite from the Scawt Hill and Ballycraigy occurrences. Our results are in close agreement, and are interesting as showing practically identical conditions of formation, habit, etc., in two widely separated localities.

His observation of water in the mineral, overlooked in the original description, has been confirmed by the author for the Crestmore occurrence.

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