

Calciohilairite: $\text{CaZrSi}_3\text{O}_9 \cdot 3\text{H}_2\text{O}$, the calcium analogue of hilairite from the Golden Horn batholith, northern Cascades, Washington

RUSSELL C. BOGGS

Department of Geology, Eastern Washington University, Cheney, Washington 99004, U.S.A.

ABSTRACT

Calciohilairite, $\text{CaZrSi}_3\text{O}_9 \cdot 3\text{H}_2\text{O}$, occurs associated with microcline, quartz, albite, fluorite, chlorite, bastnaesite, zircon, and malachite in miarolitic cavities in peralkaline granite of the Eocene Golden Horn batholith, northern Cascades, Washington. The crystals are white to pale blue, trigonal, up to 2 mm long, and show the forms a' $\{11\bar{2}0\}$, a $\{2\bar{1}\bar{1}0\}$, r $\{10\bar{1}2\}$, and $-r$ $\{01\bar{1}2\}$. The probable space group is $R32$; with $a = 20.870(4)$ and $c = 16.002(4)$ Å. The Mohs' hardness is 4, and there is a conchoidal fracture and no cleavage. The density is $2.68(2)$ g/cm³ (meas.), 2.74 g/cm³ (calc.). The optical properties are uniaxial (–), $\epsilon = 1.619(1)$, $\omega = 1.622(1)$. The X-ray powder-diffraction pattern, which is very similar to that of hilairite ($\text{Na}_2\text{ZrSi}_3\text{O}_9 \cdot 3\text{H}_2\text{O}$), shows the following strongest eight lines [d (Å), I, hkl]: 5.99, 12, 202; 5.23, 100, 220; 3.14, 14, 422; 3.02, 34, 205, 600; 2.606, 8, 440, 612, 244; 1.817, 31, 381, 725, 734; 1.671, 17, 930, 482; 1.668, 15, 482, 384. An average of six electron-microprobe analyses yields the empirical formula



The name is for the Ca content and the relationship to hilairite.

INTRODUCTION

The Eocene Golden Horn batholith drops out over 310 km² in the northern Cascade Mountains in Washington. Four granite types occur in the batholith, a peralkaline arfvedsonite granite, an alaskitic border granite, a hypersolvus biotite granite, and a subsolvus biotite granite (Stull, 1969; Boggs, 1984a, 1984b). Several suites of uncommon minerals occur in miarolitic cavities and small pegmatites in the different granite types and have been described by Dunn et al. (1977) and Boggs (1980, 1984a, 1984b).

Calciohilairite is the third new species to be described from the Golden Horn batholith. The previously described species are zektzerite, $\text{NaLiZrSi}_6\text{O}_{15}$ (Dunn et al., 1977) and okanoganite, $(\text{Na}, \text{Ca})_3(\text{Y}, \text{Nd}, \text{Ce})_{12}\text{Si}_6\text{B}_2\text{O}_{27}\text{F}_{14}$ (Boggs, 1980). All three species occur in miarolitic cavities in the batholith, zektzerite and okanoganite in the arfvedsonite granite and calciohilairite in the border granite (Boggs, 1984a, 1984b). The mineral has been accepted and the name approved by the IMA Commission on New Minerals and Mineral Names. The name is for the Ca content and the relationship to hilairite ($\text{Na}_2\text{ZrSi}_3\text{O}_9 \cdot 3\text{H}_2\text{O}$). Co-type material has been deposited in the National Museum of Natural History, Smithsonian Institution, Washington, D.C. (N.M.N.H. no. 161901 and 162182) and in my own collection (no. T990 to T996).

OCCURRENCE

Calciohilairite has been found in two occurrences in boulders of border granite on a talus slope on the north side of Liberty Bell Mountain, Okanogan County, Wash-

ington (lat 48°31'4"N, long 120°39'7"W). Both occurrences have been collected by Robert M. Boggs, who provided the material for this study. The first occurrence, discovered in September 1982, consisted of white altered trigonal prismatic crystals up to 2 mm associated with microcline, quartz, albite, fluorite, and chlorite in a zone of small miarolitic cavities. The second, discovered in October 1984, consisted of smaller (up to 1 mm) white to pale blue unaltered crystals in a larger isolated miarolitic cavity associated with microcline, quartz, albite, bastnaesite, zircon, chlorite, and malachite. Less than 500 mg of material has been collected from the two occurrences. Other minerals found in miarolitic cavities in the border granite in the same area but not associated with calciohilairite include allanite, kinosite, gadolinite, and beta-fergusonite (Boggs, 1984a, 1984b).

PHYSICAL PROPERTIES

The material from the first find consisted of white altered crystals with a hardness of about 1. Owing to the altered nature of these crystals, it was not possible to measure other physical properties. The crystals from the second occurrence are unaltered and thus permit their physical and optical properties to be measured. These unaltered crystals have a Mohs' hardness of 4. The crystals do not show any cleavage. The density is $2.68(2)$ g/cm³ (measured by flotation in heavy liquids) or 2.74 g/cm³ (calc.). The crystals are uniaxial (–), with $\epsilon = 1.619(1)$ and $\omega = 1.622(1)$. The crystals have a vitreous luster and show no fluorescence under either long-wave or short-wave ultraviolet light.

TABLE 1. X-ray powder-diffraction data for calciohilairite and hilairite

Calciohilairite (altered)				Calciohilairite (unaltered)				Hilairite*		
<i>l</i>	<i>d</i> _{obs}	<i>d</i> _{calc}	<i>hkl</i>	<i>l</i>	<i>d</i> _{obs}	<i>d</i> _{calc}	<i>hkl</i>	<i>d</i> _{obs}	<i>l</i>	<i>hkl</i>
12	5.989	5.990	202	12	5.993	6.003	202	6.00	60	012
100	5.228	5.217	220	100	5.231	5.225	220	5.28	100	110,003
				4	3.917	3.916	104			
								3.64	5	104
				6	3.197	3.186	511			
14	3.139	3.141	422	20	3.137	3.146	422	3.17	50	122
						3.134	134			
34	3.017	3.017	205	30	3.013	3.016	600	3.05	40	300
		3.012	600			3.013	152			
4	2.890	2.898	125	7	2.900	2.898	520	2.994	30	024
		2.894	520					2.639	30	200,033
8	2.606	2.609	440	9	2.607	2.610	612	2.604	5	214
		2.606	612			2.603	244			
		2.598	244							
3	2.386	2.385	434	4	2.393	2.395	262	2.362	5	116
						2.390	434			
				4	2.172	2.173	217	2.197	10	042
						2.173	354			
								2.135	15	134
3	2.128	2.128	722	3	2.050	2.048	182			
		2.124	624							
5	2.009	2.009	075	4	2.005	2.007	327	2.027	30	232
		2.008	900			2.007	544			
		2.007	642							
5	1.980	1.988	018	5	1.976	1.976	265	1.996	30	410,306
		1.973	265			1.975	820	1.866	5	226
		1.972	820					1.852	10	324
31	1.817	1.823	381	4	1.820	1.826	381	1.820	5	208
		1.817	725			1.821	725			
		1.813	734			1.816	437			
		1.740	645							
5	1.738	1.739	660	4	1.744	1.743	645	1.759	40	330
		1.738	292			1.742	660			
								1.719	5	128
17	1.671	1.671	930					1.688	20	422
		1.670	482							
15	1.668	1.670	482					1.661	20	054
		1.668	384							
				3	1.640	1.637	0.11.1			
						1.597	933			
3	1.589	1.591	10.2.2	4	1.594	1.595	835	1.607	10	152
		1.589	574			1.593	850			
		1.588	277			1.593	10.2.2			
								1.584	10	244
3	1.558	1.560	11.1.1					1.562	<5	318,1.0.10
		1.558	2.1.10					1.524	5	600
		1.556	295					1.516	10	514
								1.476	10	342
3	1.461	1.461	862							
		1.460	494							
3	1.443	1.443	10.1.6	4	1.446	1.449	592	1.464	30	520
						1.445	10.1.6			
						1.442	648			
								1.441	10	238,2.1.10
								1.373	10	612
								1.344	5	508,1.3.10
								1.319	15	440,606
								1.288	10	072,532

* Chao et al. (1974).

Applying the Gladstone-Dale relationship with the constants of Mandarino (1981) and the observed average composition gives a K_C of 0.2274. The measured indices of refraction and density give a K_P of 0.2317. This gives a value of the compatibility expression $1 - (K_P/K_C)$ of -0.0189 , which indicates superior compatibility among density, indices of refraction, and chemical composition.

X-RAY AND MORPHOLOGICAL CRYSTALLOGRAPHY

The crystals are very similar in habit to those of hilairite and show only four forms. These are two trigonal prisms a' $\{11\bar{2}0\}$ and $'a\}$ $\{2\bar{1}\bar{1}0\}$ and a right and left rhombohedron r $\{10\bar{1}2\}$ and $-r$ $\{01\bar{1}2\}$. No twinning, as seen in hilairite, was observed. Figure 1 is a drawing of a typ-

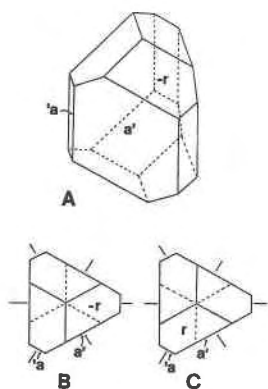


Fig. 1. Crystal drawings of calciophilairite. Forms shown are a' {01 $\bar{1}$ 2}, $'a'$ {2 $\bar{1}$ 10}, r {10 $\bar{1}$ 2}, and $-r$ {01 $\bar{1}$ 2}. (A) Left-handed crystal. (B and C) Projections down the c axis of a left- and right-handed crystal, respectively.

ical crystal along with two projections down c showing the development of right- and left-handed crystals. This interpretation of the morphological crystallography differs from that described for hilairite by Chao et al. (1974), who described the morphology in terms of a hexagonal prism {11 $\bar{2}$ 0} and a rhombohedron {01 $\bar{1}$ 2}. All hilairite crystals seen by me show a habit very much like that shown in Figure 1. This suggests that out of the possible space groups reported by Chao et al. (1974), the space group of both calciophilairite and hilairite is most likely $R\bar{3}2$. This space group was confirmed for hilairite by the crystal-structure determination of Ilyushin et al. (1981).

The X-ray powder-diffraction pattern of calciophilairite, obtained on a Phillips diffractometer using $\text{CuK}\alpha$ radiation ($\lambda = 1.54178 \text{ \AA}$) and quartz as an internal standard, is very similar to that of hilairite. Table 1 shows the data for calciophilairite (both the altered and unaltered material) compared with the data from Chao et al. (1974) for hilairite. The powder pattern obtained on the altered crystals shows an unidentified line at 7.17 \AA . This line is not present in the pattern of an unaltered crystal obtained using a Gandolfi camera and is attributed to an unidentified phase in the altered material. Initially, a least-squares refinement of the powder data for the altered material (based on the relationship to hilairite and Weissenberg

photographs) gave cell dimensions of $a = 10.415(4) \text{ \AA}$ and $c = 16.01(3) \text{ \AA}$. Later single-crystal diffractometer work, using the unaltered material (Subrata Ghose, pers. comm., July 1987), indicated that the above cell was a subcell and that the actual cell has dimensions of $a = 20.887(4) \text{ \AA}$ and $c = 15.914(3) \text{ \AA}$. A refinement of the powder data for the altered material based on this new cell gave cell dimensions of $a = 20.870(4) \text{ \AA}$ and $c = 16.002(4) \text{ \AA}$. A refinement of powder data from the unaltered material (obtained both with a Gandolfi camera and a Picker diffractometer) gave cell dimensions of $a = 20.90(1) \text{ \AA}$ and $c = 16.05(2) \text{ \AA}$. The poorer quality of this refinement is due to the lack of enough unaltered material to obtain a better pattern.

COMPOSITION

Crystals of calciophilairite were embedded in epoxy, ground to expose their centers, and coated with 200 \AA of C. Analyses were done by a combination of wavelength-dispersive (for Na) and energy-dispersive techniques on an ARL EMX electron microprobe equipped with a Tracor Northern TN-2000 energy-dispersive analyzer. Energy-dispersive spectra were corrected for peak overlap using the program SUPERML, and the combined energy- and wavelength-dispersive data were corrected for generation, absorption, and fluorescence effects using a modified version of the program ZAF (both programs supplied by Tracor Northern). The results of six analyses are shown in Table 2. Water was not analyzed for owing to the altered nature of most of the crystals and the small amount of material available. An average of these analyses gives the empirical formula $(\text{Ca}_{0.89}\text{Na}_{0.03}\text{Cu}_{0.03})(\text{Zr}_{1.14}\text{Fe}_{0.01})(\text{Si}_{2.82}\text{Al}_{0.09})\text{O}_9 \cdot 3.30\text{H}_2\text{O}$, or ideally $\text{CaZrSi}_3\text{O}_9 \cdot 3\text{H}_2\text{O}$ with minor substitution of Na^+ and Cu^{2+} for Ca^{2+} . Water was determined by difference. The Si:Zr ratio is consistently slightly lower than the ideal 3.00 (in the range of 2.80–2.90). This lower ratio could be accounted for in two ways. The first would be a systematic error in the determination of the Si:Zr ratio, and the second that some of the Zr is substituting for Ca. The first of these seems to be ruled out by the fact that other Zr-bearing silicates analyzed along with calciophilairite (zektzerite, sogdianite, calcium catapleiite, and elpidite) do not show such systematic errors (Boggs, 1984a).

TABLE 2. Composition of calciophilairite (wt%)

CaZrSi ₃ O ₉ ·3H ₂ O	1	2	3	4	5	6†	
Na ₂ O	—	0.20	0.22	0.24	0.13	0.20	0.19
CaO	13.56	11.25	10.74	10.70	11.62	11.41	14.10
CuO	—	0.19	0.77	1.12	0.42	0.74	n.d.
FeO*	—	0.03	0.70	0.03	0.12	0.09	0.00
SiO ₂	43.58	38.81	39.03	39.74	41.16	41.37	37.35
ZrO ₂	29.79	31.64	32.37	33.39	33.58	32.02	33.97
TiO ₂	—	0.09	0.09	0.04	0.04	0.02	0.00
Al ₂ O ₃	—	2.61	2.59	1.06	0.05	0.28	0.00
Sum	86.93	84.82	86.51	86.32	87.12	86.13	85.61
H ₂ O	13.07	15.18**	13.49**	13.68**	12.88**	13.87**	14.39**
Sum	100.00	100.00	100.00	100.00	100.00	100.00	100.00

* Total Fe as FeO.

** H₂O by difference.

† Altered material.

The second possibility is supported by the fact that the sum of Ca, Na, and Cu is less than one atom calculated with respect to nine oxygens, and the additional Zr brings this total close to one and improves the site occupancies in the Ca and Zr sites. This assumption gives the empirical formula $(\text{Ca}_{0.89}\text{Zr}_{0.15}\text{Na}_{0.03}\text{Cu}_{0.03})(\text{Zr}_{0.99}\text{Fe}_{0.01})(\text{Si}_{2.82}\text{Al}_{0.09})\text{O}_9 \cdot 3.30\text{H}_2\text{O}$. The substitution of Zr^{4+} for Ca^{2+} may seem somewhat unlikely, but their ionic radii do not differ greatly (0.79 and 0.99 Å, respectively), and similar substitutions are known to take place in, for example, the pyrochlore group, where Na^+ , Ca^{2+} , Pb^{2+} , U^{4+} , etc., substitute for one another. This substitution would lead to a generalized formula of $(\text{Ca},\text{Na},\text{Zr},\text{Cu})_{1-2}\text{ZrSi}_3\text{O}_9 \cdot 3\text{H}_2\text{O}$ for calciohilairite.

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