

EUDIDYMITÉ FROM SEAL LAKE, LABRADOR, NEWFOUNDLAND

E. H. NICKEL

*Mineral Sciences Division, Mines Branch,
Department of Mines and Technical Surveys, Ottawa, Canada.*

ABSTRACT

Eudidymite is an accessory mineral in the soda-rich paragneiss at Seal Lake, Labrador. Its chemical composition is as follows: Na₂O 12.20%, K₂O 0.39%, BeO 10.15%, SiO₂ 73.56%, and H₂O 3.62%; total 99.92%. The unit cell is slightly smaller than that of the Langesundfjord eudidymite; other physical and optical properties also show minor differences. These differences are believed to be related to variation in the Be:Si ratio in eudidymite.

INTRODUCTION

Eudidymite, NaBeSi₃O₇OH, is a rare mineral that has been reported in alkaline syenites from only a few parts of the world, *viz.* Langesundfjord in Norway (Nordenskiöld, 1887), Narsarsuk in Greenland (Flink, 1901), the Kola Peninsula in the U.S.S.R. (Shilin & Semenov, 1957), and the Tallask Mountains in the Kirghiz Republic of the U.S.S.R. (Kozlova, 1962). The discovery of eudidymite at Seal Lake, Labrador was mentioned briefly in a recent paper by Nickel & Charette (1962).

The geology of the Seal Lake area has been described by Evans & Dujardin (1961) and Heinrich & Deane (1962). According to them, the host rock is a soda-rich paragneiss associated with volcanics and alkaline syenite. The principal minerals in the paragneiss are albite and arfvedsonite. Accessory minerals found to date include aegerine-augite, quartz, barylite, pyrochlore, neptunite, schizolite, sphalerite, galena, apatite, eudidymite, and a new mineral that appears to be the niobium analogue of astrophyllite (a paper on this last mineral is in progress). A number of other accessory minerals have not yet been identified, and work on these is continuing.

The Seal Lake eudidymite was discovered in diamond-drill core by Mr. R. A. Dujardin, geologist with Rio Tinto Canadian Exploration Limited, and was identified as such by the author. The mineral was subsequently found in other samples from the deposit.

GENERAL DESCRIPTION

Eudidymite in macroscopically visible amounts has been observed in only a few samples, where it forms relatively coarse-grained segregations up to about an inch in diameter. In these segregations it has a rather foliated appearance due to the micaceous cleavage characteristic of the mineral, and to the sub-parallel arrangement of the individual grains. The largest cleavage surfaces observed were about 5 mm in diameter.

The eudidymite in other samples from the Seal Lake deposit appears to occur chiefly as disseminated grains. In these samples it could not be recognized either in hand samples or in thin-sections because of its similarity to albite, one of the dominant minerals in the rock. However, it was identified in immersion mounts made from heavy-liquid separates of the samples.

PHYSICAL AND OPTICAL PROPERTIES

The Seal Lake eudidymite is white in the hand sample, and colourless in thin section in transmitted light. The refractive indices, measured on oriented cleavage fragments using calibrated refractive index liquids and sodium light are as follows: $\alpha = 1.544$, $\beta = 1.545$, $\gamma = 1.549$. $2V$, measured on the universal stage, is $+23^\circ$. The extinction angle $Z \wedge c$ is 54° .

The mineral has perfect, almost micaceous, cleavage parallel to (001), and exhibits twinning parallel to this plane. The specific gravity, as determined by the heavy-liquid suspension method, is 2.578. This is in fairly good agreement with the calculated specific gravity of 2.595.

The refractive indices and birefringence of eudidymite are similar to those of albite. Consequently, it is very difficult to distinguish the two minerals, both in hand sample and in thin-sections. They can, however, be discriminated in immersion mounts with suitable refractive index liquids, particularly if phase contrast is used.

CHEMICAL COMPOSITION

1.6 grams of relatively pure eudidymite were obtained from one of the coarse-grained segregations by crushing a portion of the sample and subjecting it to repeated heavy-liquid separations. The resulting eudidymite concentrate was chemically analysed. The results of the analysis (Table 1) show that, except for the presence of K_2O , the composition of this eudidymite is very close to the theoretical eudidymite composition, also given in Table 1. The Langesundfjord eudidymite deviates from the theoretical

TABLE 1. CHEMICAL COMPOSITION OF EUUDYDMITE

	Seal Lake, Labrador* (This paper)	Langesundfjord, Norway** (Nordenskiöld, 1887)	Theoretical composition (NaBeSi ₃ O ₇ OH)
Na ₂ O	12.20%	12.45%	12.64%
K ₂ O	0.39	—	—
BeO	10.15	10.89	10.20
SiO ₂	73.56	72.65	73.49
H ₂ O	3.62	3.81	3.67
Totals	99.92	99.80	100.00

*ANALYSTS: D. J. Charette and Miss E. M. Penner, Analytical Chemistry Subdivision.

**AVERAGE of two analyses.

composition and from the Seal Lake analysis by its appreciably higher BeO and lower SiO₂ contents.

X-RAY DIFFRACTION ANALYSIS

The x-ray powder diffraction data for the Seal Lake and Langesundfjord euudymites are shown in Table 2. The powder data for the latter are in general similar to those published elsewhere for Langesundfjord euudymite (Christophe-Michel-Lévy, 1961; Kozlova, 1962); the differences that do exist are relatively minor and are probably due to differences in measuring technique. The data given here were obtained from careful measurement of 114.6 mm Debye-Scherrer film, corrected for shrinkage.

A line-by-line comparison of the powder diffraction data in Table 2 shows that, with few exceptions, the *d*-values for the Seal Lake mineral

TABLE 2. EUUDYDMITE X-RAY POWDER DIFFRACTION DATA

Line No.	Langesundfjord, Norway		Seal Lake, Newfoundland		<i>hkl</i>	<i>d</i> (calc.)
	<i>I</i> (est.)	<i>d</i> (Å)	<i>I</i> (est.)	<i>d</i> (Å)		
1	2	6.76	2	6.77	002	6.79
2	6	6.35	7	6.35	110	6.32
3	2	6.10	3	6.09	200	6.10
4		—	1	5.98	111	6.01
5	1	5.48	2	5.47	111	5.47
6		—	1	5.18	202	5.19
7	2	4.95	2	4.94	112	4.93
8	2	4.35	3	4.37	112	4.36
9	3	3.928	3	3.914	113	3.912
10	5	3.687	3	3.684	020	3.686
11		—	2	3.640	311	3.637
12		—	1	3.565	{310 021}	{3.562 3.557}
13	4	3.487	4	3.483	113	3.480
14	8	3.398	9	3.394	004	3.393
15	1	3.324	1	3.318	204	3.320

TABLE 2 (cont'd)

Line No.	Langesundfjord, Norway		Seal Lake, Newfoundland		$\frac{hkl}{hkl}$	d (calc.)
	I (est.)	d (Å)	I (est.)	d (Å)		
16	2	3.234	2	3.243	022	3.239
17	10	3.163	10	3.160	221	3.160
					220	3.155
18	8	3.074	8	3.066	114	3.155
					402	3.070
					400	3.052
19	6	2.999	6	3.001	222	3.006
					221	2.993
20	1	2.928	1	2.915	312	2.914
21	6	2.848	6	2.849	023	2.856
					114	2.846
22	2	2.754	2	2.747	223	2.752
					222	2.735
23	1	2.715	1	2.706	204	2.705
24	3	2.603	3	2.598	404	2.597
25	2	2.560	2	2.562	402	2.564
26	3	2.501	3	2.495	024	2.496
27	2	2.456	2	2.452	224	2.467
28	2	2.416	1	2.418	223	2.450
					130	2.409
29	2	2.396	2	2.393	131	2.391
30	1	2.365	2	2.361	115	2.389
					422	2.359
31	—	—	1	2.307	206	2.308
32	2	2.263	2	2.262	132	2.306
					006	2.262
33	—	—	1	2.235	132	2.237
34	1	2.221	1	2.216	116	2.218
35	2	2.183	2	2.186	025	2.186
					224	2.181
36	—	—	1	2.156	133	2.169
					514	2.154
					424	2.123
37	2	2.125	2	2.123	316	2.120
					331	2.118
38	—	—	1	2.084	133	2.086
					332	2.083
					116	2.051
39	2	2.056	2	2.053	404	2.040
					331	2.041
40	5	2.014	5	2.011	134	2.009

The data were obtained from fibre-mounted samples irradiated with iron-filtered cobalt radiation ($\lambda = 1.78890$ Å); camera diameter 114.6 mm. Intensities were estimated visually.

are smaller than those of the corresponding lines in the Langesundfjord mineral. This indicates that the Seal Lake eudidymite has a smaller unit cell. Its unit-cell parameters, calculated from the powder data by a least-squares procedure, are $a = 12.568$ Å, $b = 7.371$ Å, $c = 13.976$ Å, and $\beta = 103^\circ 47'$. A comparison of these values with those of the Langesundfjord eudidymite given by Ito (1947), and shown in Table 3, shows that the a and c values of the Seal Lake eudidymite are appreciably smaller.

DISCUSSION

The physical and optical properties of the eudidymite from Seal Lake exhibit small but distinct differences from those of the Langesundfjord mineral (Table 3).

TABLE 3. COMPARISON OF THE PHYSICAL AND OPTICAL PROPERTIES OF EUDIDYMITE FROM SEAL LAKE AND LANGESUNDFJORD

	Seal Lake, Labrador (This paper)	Langesundfjord, Norway*
Refractive indices		
α	1.544	1.545
β	1.545	1.546
γ	1.549	1.551
Birefringence ($\gamma - \alpha$)	0.005	0.006
$2V (+)$	23°	30°
Extinction angle ($Z \wedge c$)	54°	58.5
Specific Gravity	2.578	2.553
Unit-cell parameters		
a (Å)	12.568	12.62
b (Å)	7.371	7.37
c (Å)	13.976	13.99
β	103°47'	103°43'

*Optical data and specific gravity from Brögger (1890); crystallographic data from Ito (1947).

The chief compositional differences between the two minerals are in their BeO and SiO₂ contents. In the Seal Lake eudidymite these constituents are quite close to the theoretical values, but in the Langesundfjord mineral, the BeO content is appreciably greater, and the SiO₂ content appreciably less than the theoretical values. This suggests that some of the silicon has been replaced by beryllium in the Langesundfjord eudidymite.

Because of the similarity of their ionic radii, substitution of silicon by beryllium is quite common in some silicates, especially in micas and vesuvianite (Beus, 1956). Because beryllium has a lower valency than silicon, the resulting charge imbalance must be corrected. The most common mechanism in silicates appears to be replacement of divalent oxygen by monovalent fluorine or hydroxyl. Since no fluorine has been reported in the eudidymite analyses, it may be assumed that the charge imbalance has been corrected by the substitution of hydroxyl for some of the oxygen. Recalculation of the three eudidymite analyses available shows that both samples of the Langesundfjord eudidymite contain excess Be and OH and are deficient in Si and O, as expected (Table 4).

This substitution would explain the observed differences in physical properties between the Seal Lake and Langesundfjord minerals. Sub-

TABLE 4. ATOMIC PROPORTIONS IN EU DIDYMIT

	Seal Lake, Labrador	Langesundfjord, Norway		Theoretical
		1	2	
Na	0.97	1.00	0.97	1.00
K	0.02	—	—	—
Be	1.00	1.09	1.04	1.00
Si	3.01	2.93	2.98	3.00
OH	0.99	1.04	1.03	1.00
O	7.01	6.94	6.98	7.00
	13.00	13.00	13.00	13.00

stitution of hydroxyl for oxygen would be expected to create an expansion of the unit cell and an increase in refractive indices, both of which are characteristic of the Langesundfjord eudidymite. The specific gravity would be increased slightly by substitution of oxygen by hydroxyl, but this would be more than balanced by the larger decrease due to substitution of silicon by beryllium and by the effect of the larger unit cell. This is also in accordance with the observed fact that the specific gravity of the Langesundfjord mineral is substantially lower than that of the Seal Lake eudidymite.

In summary, it may be concluded that partial substitution of silicon and oxygen by beryllium and hydroxyl does occur in eudidymite, and that this substitution accounts for observed variations in physical and optical properties.

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