	3-dimen-	2-	-dimension	1-dimensional			
	sional hkl	hOl	hk0	0kl	0k0	001	h00
Structural Morphological	a*, b*, c* a*, b*, c*	2a*, c* 2a*, c*	a*, b* a*, b*	b*, c* b*, c*	2b* 4b*	c* c*	2a* 4a*
				or b*, 2c*			

Table 2. Reciprocal-Lattice Periods for Structure and Bond Assemblage (Morphology)

latter, it is (a, b). As to the yz projection, the period of the bond assemblage could be either (b, c) or (b, c/2). The 1-dimensional conclusions are tentative: the periods of the linear projections on y, z, x, which are b/2, c, and a/2, for the structure, probably are b/4, and c, a/4, for the bond assemblage. (Note that, in the yz projection, the projection lines are parallel to the x axis; in the z projection, the projection planes are parallel to xy.) The morphological data of hodgkinsonite are not sufficient to warrant more definite conclusions.

Note added in proof: The structure of hodgkinsonite has now been published by Rentzeperis (Zeit. Krist. 119, 117–138, 1963).

REFERENCES

- DONNAY, J. D. H. AND G. DONNAY (1961) "Assemblage liaisons" et structure cristalline, Compt. Rend. Acad. Sci. Paris, 252, 908-909; see also, 252, 1982-1983, 1961; 253, 291-292, 1961.
- AND D. HARKER (1937) Généralisation de la loi de Bravais, Compt. Rend. Acad. Sci. Paris, 204, 274-276.
- Palache, C. (1935) The minerals of Franklin and Sterling Hill, Sussex County, New Jersey, U. S. Geol. Survey Prof. Paper 180.
- Rentzeperis, P. J. (1958) The unit cell and space group of hodgkinsonite, Acta Cryst. 11, 448.
- ROBERTS, W. M. B. AND F. M. QUODLING (1962) X-ray, optical, and morphological observations on hodgkinsonite from Franklin Furnace. *Mineral. Mag.* 33, 343-346.

THE AMERICAN MINERALOGIST, VOL. 49, MARCH-APRIL, 1964

A CHROME-NICKEL PENNINE FROM SERPENTINITE, JAMBUR CHROMITE MINES, MYSORE STATE, INDIA

S. VARADARAJAN, Banaras Hindu University, India.

The bluish green variety of chlorite occurring in the form of small veins in the serpentinite mass, adjacent to chromite veins, in the Jambur

¹ Present address: Dept. Geol., Central College, Bangalore, India.

Chromite mines (N. 12°58′45″-12°59′25″: E. 76°28′15″-76°28′37″), Hassan district, Mysore State, India, is found to be a variety of penninite containing both chromium and nickel. The lithology of the Jambur chromite block is characterized by a lensoid serpentinite mass (960 m. ×360 m.) intruding into hornblende amphibolite trending in a NNW-SSE direction. Tabular chromite veins up to 5 m thick (mostly mined out), striking N. 80° E. and dipping N. 10° W. at high angles, occur in a serpentine mass. The veins of the mineral, generally about 1 cm thick, occur around the chromite veins particularly in the footwall of the Sundara Murthy pit. Grains up to 5 mm in diameter lie parallel to the walls of the vein.

PROPERTIES

The mineral is micaceous in habit with perfect basal cleavage. In places it is pseudohexagonal in nature. In thin flakes the mineral is pale bluish green but deeper colored in aggregates. The specific gravity is 2.65.

Optical characteristics of the mineral were determined on the universal stage.

The pleochroism is distinct with X=Y= pale bluish green, Z= pale pinkish orange. Extinction is straight. Uniaxial to biaxial; in biaxial grains 2V is as large as 12°. The optic sign is positive; optic elongation negative. Z is almost normal to (001); optic plane is (010). Dispersion moderately strong, with r < v. α and β determined by immersion method in sodium light and γ index determined by the method of Fergusson and Peacock (1943) are:

$$\alpha = 1.579, \ \beta = 1.580, \ \gamma = 1.584; \ \pm 0.002.$$

 $\gamma - \alpha = 0.005 \pm 0.002.$

X-RAY STUDY

Brindley's investigation of the mineral (pers. comm.) has yielded the following results (Table I).

The results are similar to the data recorded by Lapham (1958, p. 937) for chromian chlorites.

The basal spacing of the mineral is 14.19 ± 0.02 Å. On the basis of the equation $d(001)=14.55\times2.9X$ (Brindley, 1960, p. 270) deducing X=0.89, the estimated tetrahedral ions are (Si_{3.11}Al_{0.89}). This falls in the pennine field of Tschermak's classification and also that of Hey (1954, p. 280). The *b* parameter of the mineral is 9.44 Å.

Brindley (pers. comm.) points out that this value of b (beyond 9.40 Å) is high for chlorite and (if correct) may indicate a considerable percent-

Basal Planes	Intensities	d(Å). (obs.)	d values (obs.)		
001	40	14.0	14.0		
002	85	7.08	14.16		
003	95	4.75	14.25		
004	100	3.562	14.25		
005	34	2.853	14.27		
006	2	2.830	14.28		
007	10	2.040	$ 14.28\rangle 14.29\pm0.0\rangle$		
008	1-2				
009	2	1.591	14.32		
00, 10	5	1.431	14.31		

age of iron or similar ions. Brown and Bailey (1963, p. 52) have recorded a b value of 9.427 Å for a chromian pennine from Erzincan mines, Turkey, having $Cr_2O_3=9.30$, FeO=1.40 and $Fe_2O_3=0.05$ (weight percentages). The mineral investigated here contains FeO=2.40%, $Fe_2O_3=0.50\%$ (Table II).

Table II. Chemical Analysis and the Number of Cations in the Unit Cell of the Cr-Ni Pennine

Oxides	Wt. %	Mol. Prop.	Oxygen atoms	Hydrous basis 18 Oxygen atoms Tschermak (1891)	Anhydrous basis 14 Oxygen atoms Hey (1954)
SiO ₂	32.99	.5498	1.0996	3.14	3.14(0.00
Al ₂ O ₃	14.12	. 1382	.4146	1.58	$\frac{3.14}{1.58}$ 0.86
Cr ₂ O ₃	1.56	.0160	.0480	0.18	0.18 (0.72)
Fe ₂ O₃	0.50	.0030	.0090	0.03	0.03
FeO	2.40	.0330	. 0330	0.19	$0.19 \ 0.21$
MnO	0.18	.0028	.0028	0.02	$0.02 \int_{0.21}^{0.21}$
MgO	32.98	.8247	.8247	4.72	4.72
CaO	0.50	.0090	.0090	0.05	0.05
Na ₂ O	0.07	.0011	.0011	0.01	0.01
K_2O	0.05	.004	.0004	0.005	0.005
H_2O^+	12.66	.7033	. 7033	8.02	
H_2O^-	0.86				
TiO ₂	0.17	.0024	.0048	0.014	0.014
NiO	0.38	.0048	.0048	0.03	0.03
P_2O_5	0.07	.0005	.0025	0.006	0,006
Total	99.49			18.015	9.995

Analyst: S. Varadarajan.

CHEMICAL COMPOSITION

A pure sample of the mineral, devoid of any inclusions, was analyzed according to the methods of Shapiro and Brannock (1956) using G-1 and W-1 as standards (U.S. Bureau of Standards). The chemical composition and the number of cations calculated for the unit cell (hydrous basis 18 oxygen atoms; anhydrous basis 14 oxygen atoms) are given in Table II.

The structural formula for the mineral is:

$$Na_{.01}Ca_{.05}(Mg_{4.72}Al_{.72}Cr_{.18}Fe_{.21}{}^{2}Fe_{.03}{}^{3}Ni_{.03})(Si_{3.14}Al_{.86})O_{10}(OH)_{8.02}$$

or

It is to be noted that the mineral contains 1.56% Cr₂O₃, and 0.38% NiO. The number of tetrahedral ions calculated from the chemical analysis is Si_{3.14} and Al_{0.86}. These correspond very well with those estimated by Brindley on x-ray data. Further, the proportion of Si:Al = 1.99:1 is within the range of pennine (Niggli, p. 96).

DISCUSSION AND CLASSIFICATION

The mineral investigated is a normal chlorite (Nelson and Roy, 1958, p. 707) with a basal spacing of 14.29 ± 0.02 Å. It is a pennine with Si = 3.14 and Si:Al=1.99:1; with 1.56% Cr₂O₃ it is a chromian pennine (Hey, 1954, p. 280; Lapham, 1958, p. 953).

But the refractive indices of the mineral are low for the percentage of Cr₂O₃ that it contains (Lapham, 1958, p. 940) and also for the atomic percentage (calculated for 10 cations) of 7.6 for

$$\frac{\text{Fe} + \text{Mn} + \text{Cr}}{\text{Fe} + \text{Mn} + \text{Cr} + \text{Mg}}$$

(Albee, 1962 p. 865). The presence of nickel introduces discrepancies and tends to lower the refractive index as is clear from the varieties of chromium chlorites, with and without nickel, investigated by Lapham (1958, p. 940), given in Table III.

TABLE III

Variety	Wt. % Cr ₂ O ₃	Wt. % NiO	Refractive Indices			
			α	β	γ	$\gamma - \alpha$
No. 2 Lapham	1.14		1.577	1.578	1.581	.004
No. 23 Lapham	2.90	0.47	1.568	1.569	1.573	.005
No. 25 Lapham	3.08	0.10	1.573	1.574	1.578	.005
Mineral investigated	1.56	0.38	1.579	1.580	1.584	.005

It is clear from Table III that the refractive indices of the mineral are higher than would be expected for the percentage of Cr₂O₃ and NiO that it contains. But the chromian chlorite series, with varying proportions of NiO, has not yet been investigated sufficiently to justify any generalizations in this respect.

ACKNOWLEDGMENTS

The author is indebted to Dr. I. C. Pande for his guidance during the work and for critically reading through the manuscript. The x-ray study of the mineral was carried out by Dr. G. W. Brindley, Pennsylvania State University, and his comments on the text are gratefully acknowledged. The present work was carried out during the period of the fellowship awarded by the National Institute of Sciences of India.

REFERENCES

ALBEE, A. L. (1962) Relationship between chemical composition and physical properties of chlorites. Am. Mineral. 47, 851-870.

Brindley, G. W. (1960) X-ray identification and crystal structures of Clay Minerals, 2nd ed.: Mineral. Soc., London.

Brown, B. E. And S. W. Balley (1963) Chlorite polytypism: II-crystal structure of one-layer chromian chlorite. *Am. Mineral.* 48, 42-61.

Fergusson, D. R. and M. A. Peacock (1943) Measurement of the principal indices of refraction in micaceous minerals by immersion on tilting stage. *Am. Mineral.* 28, 561-570.

HEY, M. H. (1954) A new review of chlorites. Mineral. Mag. 30, 277-292.

LAPHAM, D. M. (1958) Structural and chemical variation in chromian chlorites. Am. Mineral. 43, 921-956.

Nelson, G. W. and R. Roy (1958) Synthetic chlorites and their structural and chemical constitution. Am. Mineral. 43, 707-725.

NIGGLI, P. (1954) Rocks and Minerals; W. H. Freeman Co., San Francisco.

Shapiro, L. and W. W. Brannock (1956) Rapid analysis of silicate rocks. A contribution to geochemistry, U. S. Geol. Survey Bull. 1036-C.

THE AMERICAN MINERALOGIST, VOL. 49, MARCH-APRIL, 1964

ON THE OCCURRENCE OF ASBESTOS IN CHAMOLI, U.P., INDIA

DHARAM PRAKASH AND IQBAL K. DALELA, Directorate of Geology & Mining, U.P., Lucknow, India.

Introduction

Asbestos (Bowles, 1959) mineralization was noted in Kandhara (30° 26′ 10″:79° 5′ 0″) and Jalai villages (30° 27′ 20″:79° 5′ 0″) on the left bank of Mandakini river in the Garhwal Himalaya (Narain, 1954). During 1960–61 Dalela was associated with the prospecting party (Prakash,