

DISTRIBUTION OF SOME MINOR ELEMENTS IN THE NIPISSING DIABASE

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ABSTRACT

Nine cross-sections of Nipissing diabase sheets in the Cobalt-Gowganda region were sampled and analyzed for nickel, cobalt, copper, chromium, silver, and in most cases, for barium, strontium, and boron. Nickel and chromium decrease, and cobalt increases with fractionation as expressed by the mafic index. Copper, silver, and arsenic generally increase with height above the bottom contacts of the intrusions; highest values are attained in varied-texture diabase and in the vicinity of the top contact. Although the total nickel content of the diabases is lowered with fractionation, leach tests indicate that the amount of nickel present in "sulphide" form does not necessarily decrease. Of the diabases examined in the present study, the New Lake basin at Cobalt is by far the most enriched in silver.

INTRODUCTION AND PREVIOUS STUDIES

Cross-sections of the Nipissing diabase in the Cobalt-Gowganda region were sampled and analyzed for cobalt, nickel, copper, silver, and in most cases, a few additional elements. The sections traversed include three at Cobalt, three in Henwood township, and one each in South Lorrain, Elk Lake, and Gowganda.

The relevance of trace element data in understanding the origin of ore deposits is very commonly difficult to assess. However, it seemed worthwhile obtaining the factual data on the Nipissing diabase in order to supplement the limited studies that have been done in the past and to evaluate the published geochemical data which have been used to support various theories of genesis of the Cobalt ores. The earlier studies had focused primarily on the abundance of silver in diabase and granophyre.

The difficulties in the early attempts at determining silver in trace amounts in the diabase from the Cobalt area are summarized by Moore (1934). Subsequently, on the basis of 23 additional determinations, Moore (1956) concluded that granophyric diabase contains more silver than ordinary diabase. Spectrographic analyses of diabases from the Cobalt area are given in the theses by Hriskevich (1952) and Coldwell (1967). Although the theses do not include analyses for silver, Fairbairn *et al.* (1953) analyzed Nipissing diabases from several localities for several elements, including silver. Their range of 0.9 to 7.6 ppm Ag is markedly

higher than the maximum of 0.24 ppm reported by Boyle (1968a) for 36 samples from the Stone-Eplett No. 3 drill hole in Henwood township. These differences emphasize the difficulties in obtaining accurate data for silver. The present study initially had the advantage that relatively large numbers of samples were to be processed in the same laboratories in a short time interval, and the problem of comparing inter-laboratory data and coping with changes in analytical procedures were therefore to be minimal. As will become evident later, this goal was only partly achieved. However, the second advantage was that most of the samples analyzed in the present study are from cross-section traverses of the diabase sheets, so that the trends of trace element variation in the differentiated intrusions are more meaningful than absolute abundances of the minor elements.

SAMPLE PREPARATION AND ANALYSIS

All the diabase hand specimens were cut with a diamond saw to representative blocks approximately $1 \times 2 \times 4$ cm. Larger specimens, particularly of coarse-grained or varied-texture diabase, were sawn into thin slabs

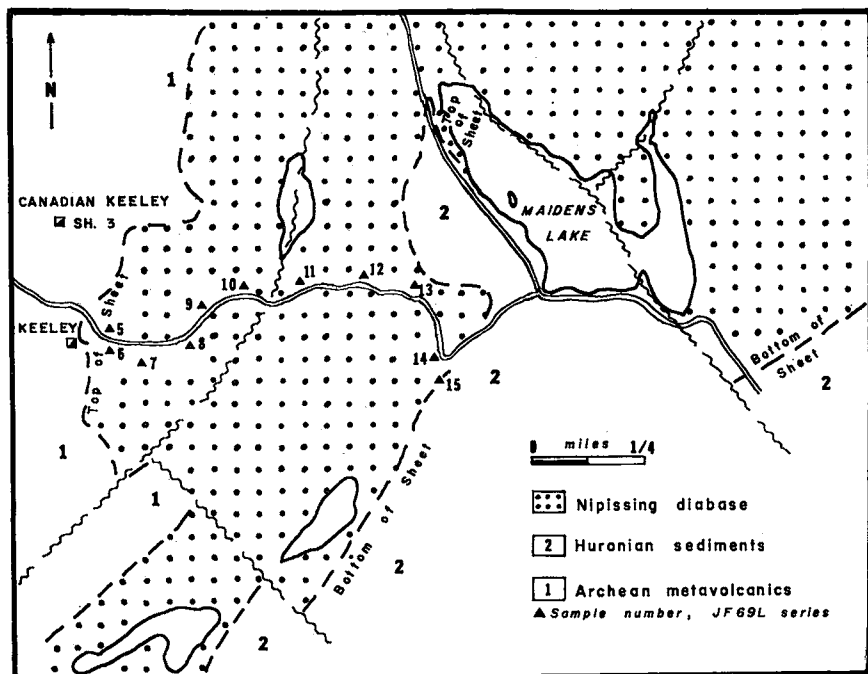


FIG. 201. Location of diabase samples in South Lorrain township.

so that representative blocks of material could be selected for the analysis. Classification of the material into diabase types was largely based on hand specimen identification, but about a third of the samples were also examined in thin sections. Because the various diabase varieties are gradational into each other, there was in some cases difficulty in precisely labeling individual hand specimens. The problem is, however, not a serious one if the gradational character of the diabase varieties is borne in mind.

Strontium analyses of the rocks were obtained by x-ray fluorescence techniques directly on the -150 mesh powders. Unless otherwise specified, all other elements were determined by optical spectrographic analysis by K.A. Church and staff of the Geological Survey. The elements Ba, Cr, Ni, Cu, and Co are considered by the Geological Survey's spectrographic laboratory to be accurate to within ± 15 per cent, and B and Ag to within ± 30 per cent. The results of the present study and the spectroscopic data compiled for rock standards by Flanagan (1969), indicate that these figures may be more realistically regarded in most cases only as an indication of the analytical precision.

TABLE 81. TRACE ELEMENT CONTENT (PPM) OF THE SOUTH LORRAIN DIABASE SHEET.

IF69L-	Rock type	Ni	Co	Cu	Sr	Ba	Cr	B	Ag
1	upper absolute chill	94	31	97	100	300	190	6	0.29
2	fine-grained quartz	85	32	120	70	170	150	11	0.12
3	diabase, progressively	66	31	94	100	150	170	16	0.18
4	coarser	68	33	93	100	130	170	10	0.17
5	varied-texture diabase	48	50	220	70	140	20	13	0.28
6-1		20	35	210	80	54	NF	6	0.22
6-2		25	21	48	90	280	NF	17	0.15
7		20	25	8	70	110	NF	13	0.52
8		NF	34	34	100	240	NF	16	0.38
9		40	34	100	100	160	20	9	0.14
10	transitional to	NF	37	50	110	180	NF	9	0.13
11	hypersthene diabase	NF	37	46	80	180	NF	<5	0.14
12		22	47	9	120	68	NF	12	0.16
13	hypersthene	50	28	84	110	130	20	8	0.17
14	diabase	200	32	55	110	97	170	8	0.58
15	basal chill	100	27	77	90	110	170	15	0.15

Analyses for this and succeeding tables are by K. A. Church and staff, spectrographic laboratories, GSC.

NF: not found; the detection limit for Ni, Co, and Cr in all tables is 10 ppm.

DIABASE ANALYSES

South Lorrain

The distribution of the Nipissing diabase in South Lorrain has been outlined in recent mapping by McIlwaine (1968). For the present study the diabase was sampled along a cross-section extending from the Keeley property to the southeast end of Maidens Lake (Fig. 201). The top of the sheet is known from mine workings to dip westward at about 45° . The diabase contact with overlying Archean volcanics is exposed near the shaft shown on the Keeley property. The bottom contact overlying Huronian sediments is exposed at sample position 15 (Fig. 201). The cross-section of the diabase is therefore equivalent to the elevated rim of an undulating sheet. However, the assumption made in the earlier days of the South Lorrain camp that the diabase formed a simple dome-shaped structure no longer appears to be valid. A more complex structure is indicated by the appearance of the top of a diabase sheet overlain by Huronian sediments at the northwestern end of Maidens Lake.

The trace element abundances for 16 samples of the diabase are given in Table 81. Both the top and bottom chilled contacts are extensively altered, with the feldspars being saussuritized and the pyroxenes replaced

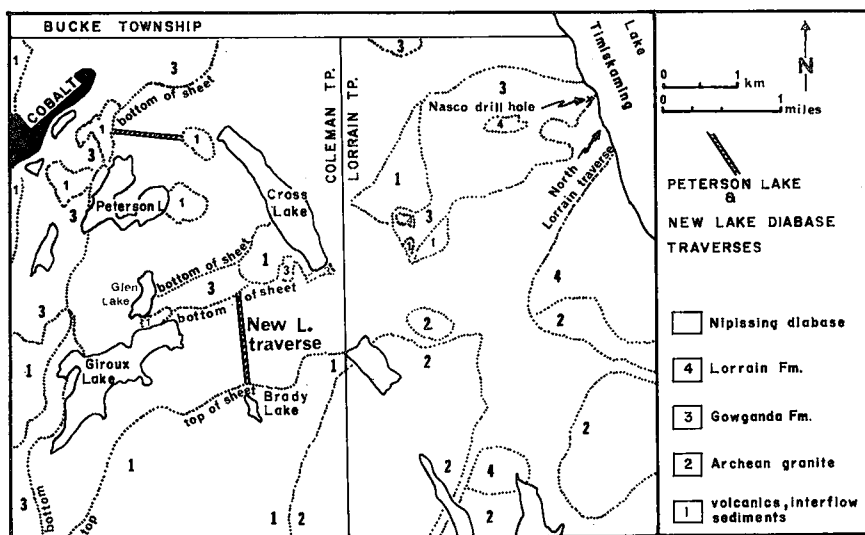


FIG. 202. General locations of the North Lorrain, New Lake, and Peterson Lake diabase samples. The position of the Nasco drill hole studied by Colwell (1967) is also shown.

by amphibole and chlorite. The two samples from position 6 are pinkish granophyric diabase from the varied-texture zone.

Cobalt area

North Lorrain basin — The southeastern portion of the North Lorrain diabase basin was traversed along the shore of Lake Timiskaming (Fig. 202), where the sheet intrudes Huronian sediments. Insufficient information is available to permit drawing a cross-section of the diabase in this area, but the sheet as a whole is known to be roughly basin-shaped. The principal ambiguities are the unknown thickness and dip of the diabase along the southern portion of the basin. If the basin shape prevails, then the southernmost rim of the sheet is the bottom contact. Sampling of the intrusion was begun at the bottom of the sheet (Table 82). Although the

TABLE 82. TRACE ELEMENT CONTENT (PPM) OF THE NORTH LORRAIN DIABASE (SOUTHERN PART) ALONG LAKE TIMISKAMING.

JF69T-	Rock type	Ni	Co	Cu	Sr	Ba	Cr	B	Ag
23	Top of sheet transitional to quartz diabase	68	33	110		140	160	9	0.25
22	varied-texture (22-18)	56	34	74		120	96	10	0.11
21		60	30	130	70	140	170	17	0.21
20		90	36	110	90	140	59	9	0.14
19		<20	37	150	70	230	NF	11	0.19
18		<20	41	86	60	170	NF		
17	quartz diabase	NF	17	21	110	42	NF	11	0.40
16	granophyre	NF	31	11	60	140	NF	<5	0.19
15	varied-texture,	<20	42	120	70	150	NF	5	0.12
14	transitional to quartz	23	36	140	100	160	NF	9	0.19
13	diabase with visible	29	31	75	90	180	NF	7	0.34
12	micropegmatite (15-4)	29	34	180	100	160	NF	15	0.15
10		26	31	79	110	200	<20	8	0.36
9		28	34	84		240	<20	13	0.16
7		54	30	130	120	160	<20	10	0.16
6		39	26	76	120	300	<20	13	0.13
4	7 m from contact	84	28	87		35	120	13	0.15
2	5 m from contact	83		110		150	210	6	0.24
1	chill (<2 m from contact)	99	52	170		110	240	13	0.06

NF : not found ; blanks : not determined

absolute chill was not found, the diabase contact with Lorrain arkose was traceable to within less than 2 meters. The diabase in this vicinity contains irregular, steeply-dipping aplite stringers up to 15 cm wide. Northward toward the top of the sheet, there is a fairly rapid progression toward medium-grained diabase containing visible micropegmatite. The succession is then to well-defined varied-texture diabase and granophyre without any intervening section of hypersthene diabase. Without delving into alternative explanations, the present writer's interpretation is that the bottom contact may be relatively flat along most of the basin, but at the southeastern rim the sheet turns upward along a steep to vertical contact. The zone of hypersthene diabase accumulation would therefore intersect the contact at a greater depth and at a relatively obtuse angle rather than the acute one which should prevail if the basal contact had a continuous shallow dip.

The results of the trace element analyses of the traversed section are given in Table 82. With the exception of 1, 2, 17, and 23, all samples are varied-texture or contain visible micropegmatite. Sample 2 is a normal quartz diabase, and 23 is transitional from varied-texture to quartz diabase near the roof contact. The appearance of quartz diabase slightly past the mid-point of the traverse (no. 17) is worthy of mention because the rock is recognizably distinct in the field and was also identified by Colwell (1967) in the core of a deep drill hole on the Nasco property (Fig. 202). The unit as logged by Colwell is approximately 200 feet (~ 60 m) thick, with the upper transition to varied-texture diabase occurring approximately 180 feet (55 m) vertically below the top contact. Extrapolation from the drill hole to the approximate position of the quartz diabase in the traverse area gives the unit an apparent dip of roughly 15 degrees north. Although it has not been studied in detail, the material appears to be a conformable unit within the sheet. Almost all the rocks, including the central quartz diabase, have undergone intense deuteritic alteration.

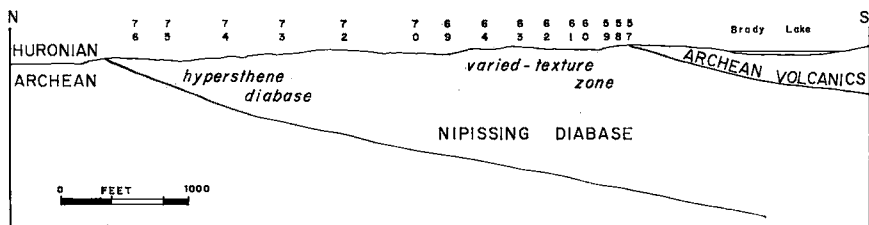


FIG. 203. Cross-section and sample positions of the New Lake diabase traverse shown in Figure 202. Horizontal and vertical scales are identical.

New Lake basin — The north end of the New Lake diabase basin was sampled along a cross-section extending from Brady Lake to the Kerr Lake arch (Fig. 202). The configuration of the diabase sheet and the approximate position of the analyzed samples are shown in Figure 203. Results for the trace element analyses are given in Table 83. Numerous silver properties occur within the basin, the production being surpassed only by deposits associated with the Peterson Lake basin to the northeast.

Peterson Lake basin — The Peterson Lake diabase was traversed northeastward from the upper contact at the Colonial mine to the lower contact near the north end of Peterson Lake (Fig. 202). The upper and lower absolute chills were not obtained, but both contacts are traceable to within a meter or two.

Hriskevich (1952) has studied the petrological variation of the diabase in the vicinity of the Colonial mine shaft, and Colwell (1967) has spectrographically analyzed samples representing a section through the diabase at the Rix Athabasca property, the latter being about 1000 feet (305 m) east of the Colonial shaft. A cross-section through these points

TABLE 83. TRACE ELEMENT CONTENT (PPM) OF THE NIPISSING DIABASE SHEET,
NEW LAKE BASIN, COBALT.

JF69C-	Rock type	Ni	Co	Cu	Sr	Ba	Cr	B	Ag
57	roof quartz diabase	90	46	160	100	140	160	12	0.36
58	varied-texture	64	32	100	100	210	98	15	0.49
59	diabase	78	44	160	110	160	150	11	0.93
60		35	21	80	110	180	<20	10	1.2
61		36	27	79	110	180	<20	12	0.44
62		73	45	150	100	130	<20	7	3.1
63		23	35	99	140	190	NF	13	0.80
64		26	29	61	150	200	NF	7	0.83
65		NF	38	130	130	98	NF	9	0.43
69		28	37	120	120	190	NF	12	0.40
70		94	51	94	60	64	40	9	0.81
71	hypersthene	60	29	110	110	130	23	14	0.61
72	diabase	64	34	140	230	130	<20	11	0.21
73		82	38	160	110	74	39	14	0.24
74		92	NF	150	110	100	150	6	0.39
75		75	25	100	100	98	99	5	0.51
76	(60 m from contact)	76	34	87	110	120	210	13	0.42

NF: not found

Henwood township

The nature of the diabase in Henwood township has already been dealt with a length (p. 50). As was previously mentioned, two separate intrusions are considered to be present, an upper basin-shaped sheet, and a lower sheet which is believed to be relatively flat-lying. They are respectively referred to as the Henwood diabase basin and the Henwood Lower sheet. The rim (Flank) of the diabase basin is exposed on surface, and the basin part was intersected in the Eplett vertical drill hole. An earlier inclined hole collared nearby also passed through the diabase basin. This inclined hole is called the Stone-Eplett No. 3. The relationships of the drill holes and the diabase sheets are schematically shown in Figure 205; detailed descriptions are given by Thomson (1966, 1968).

Core from the Stone-Eplett No. 3 hole was not seen by the present writer, but extensive geochemical data obtained from the diabase were kindly made available through the courtesy of R. W. Boyle. These data (Table 85) are useful for examining the trace element variations within

TABLE 84. TRACE ELEMENT CONTENT (PPM) OF THE PETERSON LAKE DIABASE, COBALT.

JF69C-	Rock type	Ni	Co	Cu	Sr	Ba	Cr	B	Ag	As*
30	roof quartz	91	45	160		30	99	6	0.28	70
31	diabase	75	37	100	110	230	<20	49	0.25	<4
32		100	38	59		230	NF	48	0.14	<4
33	varied-texture	75	36	92	130	190	NF	12	0.16	<4
36	diabase	66	36	59		130	170	9	0.58	
37		81	30	74		190	130	5	0.19	<4
38		110	35	110	120	100	300	7	0.14	<4
39		85	42	180	100	120	24	9	0.12	<4
41		39	31	87	90	150	<20	6	0.14	<4
42	hypersthene	93	31	88	140	69	320	NF	0.08	<4
43	diabase	210	31	200	100	57	500	<5	0.11	
44		71	32	140		74	25	NF	0.12	<4
45		76	29	120	80	80	41	NF	0.12	<4
46		130	35	120	80	70	310	NF	0.10	<4
47		120	29	98	90	73	330	NF	0.14	<4
48	(olivine-bearing)	130	42	57	110	86	400	NF	0.13	<4
49		130	36	67	80	240	360	10	0.24	<4
50		110	28	68	100	100	360	5	0.17	<4
51		110	32	80	100	140	310	5	0.14	<4
53	basal quartz	110	34	84	140	210	280	11	0.15	
54	diabase	130	42	47	120	140	200	9	0.21	<4
55		130	42	110		250	230	12	0.12	<4

* Colourimetric analysis by Geochemistry Section, GSC. NF: not found; blanks: not determined.

TABLE 85. TRACE ELEMENT ANALYSES OF NIPISSING DIABASE IN THE STONE-EPLETT NO. 3 DRILL HOLE, HENWOOD DIABASE BASIN¹

Depth ²	Rock type	Ni	Co	Cu	Ag	As	Sb	S*
4	Quartz diabase, progressively coarser	120	25	560	0.24		1.5	0.10
18		110	35	88	0.20		1.0	0.12
31		110	30	100	0.20		1.5	0.05
57		95	30	46	0.08		1.0	0.08
75		110	35	60	0.17		1.0	0.04
101	varied-texture diabase	60	35	220	0.21		1.0	0.16
112		160	35	44	0.19		1.0	0.15
127		150	30	100	0.24		1.0	0.04
136		25	30	96	0.07			0.06
156		95	45	40	0.06	2		0.08
160	hypersthene diabase,	100	40	60	0.07	4		0.14
170		80	30	88	<0.05			0.08
170		55	30	130	0.08			0.05
180		55	35	110	0.16			0.04
193		160	40	130	0.06			0.05
195		170	40	92	0.15			0.07
228		150	30	88	0.15			0.02
251		120	20	84	0.13			0.05
273		150	30	80	0.14			0.03
315		150	30	80	0.12			0.03
352	quartz diabase, progressively finer	150	35	68	<0.05			0.02
369		110	30	64	0.09			0.03
374		160	40	72	<0.05			0.03
379		110	40	44	0.09			0.03
421		160	35	68	<0.05			0.04
463		170	35	72	<0.05		3.2	0.06
511		160	35	64	<0.05			0.04
556		130	40	60	<0.05			0.03
584		160	40	60	<0.05			0.05
593		160	40	64	0.06			0.04
625	quartz diabase, progressively finer	150	30	64	<0.05			0.04
682		160	35	72	<0.05			0.05
699		150	35	68	0.07			0.06
716		150	40	110	0.24			0.05
727		160	40	76	<0.05			0.08
728	bottom contact	170	40	88	0.19			0.04
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¹ Samples obtained through the courtesy of Robert Thomson, Ontario Dept. Mines. Data courtesy R. W. Boyle.

² Vertical depth in feet from the top contact.

* All data in ppm except for sulphur, which is in per cent. Arsenic values are < 2 ppm, and antimony < 1 ppm except where specific values are given. Silver determined by emission spectrography and sulphur by a combustion method; all other elements determined by colourimetric methods by Geochemistry Section, GSC.

this particular core, but only the trends of variation are regarded as being comparable with newer analyses obtained in the present study. Except for the sulphur determinations, analytical procedures have been revised since the obtaining of the Stone-Eplett data.

Trace element results for the diabases traversed on surface and intersected in the Eplett vertical drill hole are given in Tables 86, 87 and 88.

TABLE 86. TRACE ELEMENT CONTENT (PPM) OF THE HENWOOD LOWER SHEET.

Footage	Rock type	Mafic Index	Ni	Co	Cu	Sr	Ba	Cr	Ag
6186	quartz diabase, progressively coarser	59.5	120	66	310	170	110	93	0.07
6202			57	36	110	160	140	33	0.14
6208			63		150	140	140	27	0.88
6215			79	51	150	180	100	67	0.09
6222			59	39	110		130	20	0.09
6240	varied-texture diabase	65.0	61	38	120	140	100	23	0.14
6259			70	54	230	140	120	20	0.15
6273			85	41	260	130	120	38	0.19
6287			55	37	150	120	140	20	0.18
6293			51	37	140	130	160	20	0.09
6305		66.8	34	44	130	130	140	NF	0.29
6320			80	44	140	130	170	97	0.12
6325			27	35	150	100	120	NF	0.14
6359			100		230	110	79	120	0.35
6384			84	35	140	150	190	98	0.14
6429	hypersthene diabase	57.7	48	41	440	180	110	20	0.33
6438			88	48	32	105	140	110	0.12
6469			90	40	140	130	150	72	0.12
6498			110	46	200	160	130	110	0.16
6530			110	37	150	99	130	140	0.15
6580		51.9	110	37	180	93	150	210	0.09
6613			150	40	220	92	160	310	0.08
6670			96	31	100	110	140	370	0.07
6780			86	38	100	84	140	190	0.06
6816			96	39	160	100	150	170	<0.05
6856	quartz diabase, progressively finer	54.4	120	45	130	100	130	190	0.16
6899			180	49	140	97	150	120	0.06
6900	albitized chill	57.2	180	62	150		180	180	<0.05

Mafic index : $\frac{\text{FeO} + \text{Fe}_2\text{O}_3}{\text{FeO} + \text{Fe}_2\text{O}_3 + \text{MgO}} \times 100$ (see p. 67 for reference to rock analyses).

NF : not found ; blanks : not determined

TABLE 87. TRACE ELEMENT CONTENT (PPM) OF THE HENWOOD DIABASE BASIN.
ANALYSES ARE FOR THE DEPRESSED (BASIN) PART INTERSECTED
IN THE VERTICAL EPLETT DRILL HOLE.

Footage	Rock type	Mafic Index	Ni	Co	Cu	Sr	Ba	Cr	Ag	As
1672	quartz diabase, progressively coarser	87.9	NF	47	200	50	63	<20	0.30	<3
1673			20	110	58	40	72	NF	0.55	<3
1675			110	210	28	<30	13	160	0.13	350
1680	varied-texture diabase	58.6	79	47	180	70	120	150	0.18	<3
1684		59.1	72	39	140	80	120	100	0.24	<3
1685			86	45	65	80	110	160	0.08	<3
1701		57.6	83	38	140	80	100	57	0.13	<3
1718		54.1	78	32	150	80	130	210	0.15	<3
1800		52.7	84	30	93	100	110	340	0.10	<3
1821		51.5	100	40	140	70	100	420	0.16	<3
1826		58.9	66	31	140	70	100	<20	0.23	<3
1831		67.3	63	49	110	85	140	33	0.23	
1836			64	55	200	90	140	NF	0.19	<3
1847	hypersthene diabase		82	37	110	100	100	110	0.15	<3
1850		52.4	87	30	99	100	110	330	0.11	<3
1882		50.3	110	41	130	75	70	370	0.28	<3
1907			47	47	210	90	26	83	0.13	<3
1937			87	28	97	100	110	330	0.07	<3
1945		50.0	98	33	110	90	71	270	0.10	<3
1980			99	25	75	80	130	470	0.09	<3
2002			120	30	110	110	62	330	0.06	<3
2024		45.7	110	29	110	90	55	340	0.14	<3
2066		43.2	120	32	150	100	57	560	0.15	<3
2115	quartz diabase, progressively finer		110	30	65	80	60	540	0.09	<3
2125		45.5	110	31	85	90	70	450	0.12	<3
2201		44.9	120	32	91	100	62	550	0.12	<3
2301		44.4	130	39	79	90	59	550	0.11	<3
2351		44.3	130	39	65	100	68	650	0.18	<3
2372			98	29	75	90	97	580	0.07	<3
2398			110	33	75	100	47	520	0.07	<3
2419		49.7	130	34	77	90	86	500	0.08	<3
2447		49.7	120	36	110	120	76	390	0.14	<3
2475		53.6	110	40	110	90	88	190	0.10	<3
2482	albitized chill		88	28	85	190	34	180	<0.05	<3
2491			110	40	110	100	88	190		5
2493		63.5	100	30	470	<30	220	78	0.14	8

Arsenic determined by colourimetric methods by Geochemistry Section, GSC.
NF: not found; blanks: not determined

In the Flank section the hypersthene zone is not readily identifiable in hand specimens. Most material is either quartz diabase or varied-texture diabase, but on the basis of thin section studies the two samples near the bottom of the sheet are transitional toward hypersthene diabase.

TABLE 88. TRACE ELEMENT CONTENT (PPM) OF THE HENWOOD DIABASE BASIN. ANALYSES ARE FOR THE ELEVATED RIM (FLANK) OF THE SHEET EXPOSED ON SURFACE.

JF-	Rock type	Mafic Index	Ni	Co	Cu	Sr	Ba	Cr	B	Ag	S%
HF-13	fine-grained quartz	58.4	77	34	100	100	130	250	7	0.10	0.12
HF-12	diabase, top of sheet	59.2	77	35	120	105	86	190	<5	0.12	0.05
HF-11	varied texture	61.8	83	41	130	110	91	180	<5	0.08	0.07
HF-10	diabase with	65.4	52	40	99	100	170	<20	9	0.10	0.09
HF-9	abundant amphibole	78.1	<20	41	130	100	180	NF	7	0.33	0.13
HF-7	(11-4)	67.4	33	49	130	105	160	NF	7	0.09	0.06
HF-6		67.4	56	55	180	110	130	NF	8	0.10	0.07
HF-4		67.2	37	37	130	105	190	NF	10	0.07	0.06
HF-3	hypersthene-	59.6	64	28	110	120	100	100	13	0.09	0.06
HF-1	bearing diabase	53.2	150	37	120	110	140	270	14	0.11	0.04

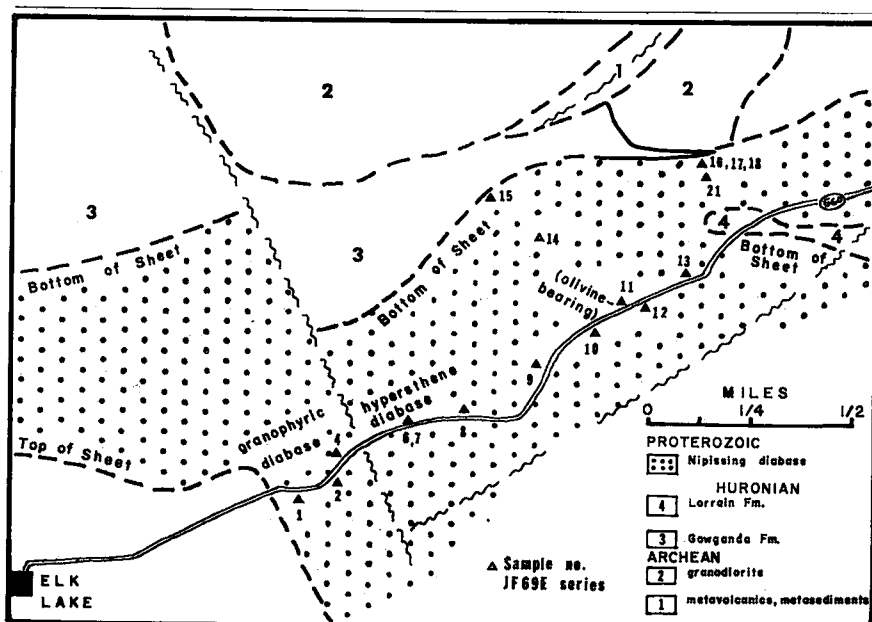


FIG. 206. Location of the Elk Lake diabase samples (geology by MacKean 1964).

Elk Lake

The area of Nipissing diabase sampled at Elk Lake is shown in Figure 206. Along the east-west section there is a relatively abrupt change from granophyric and varied-texture diabase to hypersthene diabase which is in part olivine-bearing. On Ontario Department of Mines preliminary map P. 239, MacKean (1964) has shown a northwest fault which offsets the diabase for about 800 metres. The movement has apparently brought hypersthene diabase from the lower part of a sheet into juxtaposition with granophyric diabase from near the upper part of a sheet.

The results of the trace element analyses of the diabase are given in Table 89. The very fine-grained nature of sample 13 indicates that it is

TABLE 89. TRACE ELEMENT CONTENT (PPM) OF THE DIABASE AT ELK LAKE.

JF69E-	Rock type	Ni	Co	Cu	Sr	Ba	Cr	B	Ag
1	granophyric diabase (1-4)	NF	18	13	70	50	NF	6	0.22
2		NF	16	57	40	160	NF	8	0.12
3		NF	22	64	80	250	210	15	0.12
4		NF	33	130	80	160	170	29	1.5
5	hypersthene diabase (5-12)	50	34	86	120	190	130	19	0.10
6		140	31	170	120	110	200	23	0.13
7		92	27	170	130	130	42	16	0.12
8		62	28	100	120	150	130	19	0.13
9		79	30	90	130	78	230	7	0.06
10		200	40	62	110	60	190	7	0.14
11		84	27	55	110	140	320	18	0.09
12		250	42	60	100	60	120	9	0.11
13	fine-grained quartz diabase	81	33	110	120	160	130	7	0.11
14	hypersthene diabase	200	43	84	90	110	170	12	0.14
15	fine-grained quartz diabase	110	35	94	110	150	180	14	0.16
16	chilled quartz diabase	43	35	190		140	97	6	0.16
17	chilled quartz diabase		37						
18	1 m from contact	44	37	87	130	190	97	12	0.16
19	20 m from contact ; qtz. diabase	45	40	280	130	340	74	19	1.6
20	50 m from contact ; qtz. diabase	43	37	94	130	290	96	25	0.40
21	hypersthene diabase	110	37	84	110	210	190	26	0.24

NF : not found ; blanks : not determined

within a few metres of the bottom contact and suggests that the sheet is relatively flat-lying in this vicinity. Also of interest is the apparent iron-richness of the olivine; an x-ray determination of the mineral's composition in sample 10 gave Fo 60.

Miller Lake basin

The Miller Lake diabase at Gowganda was traversed in the area shown in Figure 207. The basal diabase was not found in outcrop, the first sample being of hypersthene diabase. Zoning in pyroxenes, typical in most of the Nipissing diabase, is exceptionally pronounced in the samples from this vicinity.

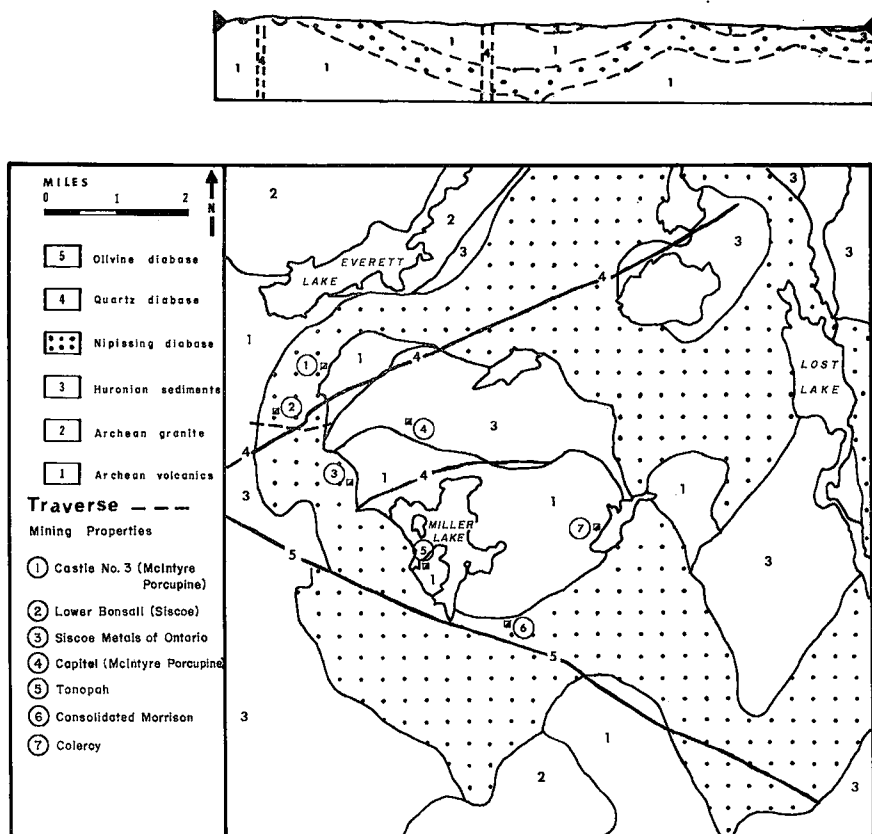


FIG. 207. Location of the Miller Lake sampling traverse. Geology and cross-section reproduced from Moore (1956).

The hypersthene diabase passes upwards into the varied-texture zone, but in many samples the variable grain size is not present and the rock could be simply classified as a quartz diabase. Outcrops from which samples 10 to 13 inclusive were taken are of varied-texture diabase, but the hypersthene-bearing variety reappears towards the top of the sheet. The results of the trace element analyses are given in Table 90.

Chilled Nipissing diabase

Most Nipissing diabase chills are extensively altered. In order to supplement the data obtained from traverses across the diabase basins, several additional chill zones were also analyzed. Samples as large as a foot in diameter were taken from the contacts and slabbed with a diamond saw in order that the least fractured and altered material could be selected. The results of the analyses are given in Table 91. The average Ni, Co, and Cu contents in chilled Nipissing diabases in the Cobalt area were

TABLE 90. TRACE ELEMENT CONTENT (PPM) OF NIPISSING DIABASE,
WEST SIDE OF MILLER LAKE BASIN, GOWGANDA.

JF69G-	Rock type	Ni	Co	Cu	Sr	Ba	Cr	B	Ag
18	roof quartz	64	42	130	100	290	110	23	1.4
17	diabase (18-17)	25	34	130	100	340	<20	16	0.49
16	hypersthene	130	52	130	100	200	300	8	0.14
15	diabase (16-14)	100	33	84	30	140	440	5	0.26
14		130	42	41	90	84	420	6	0.11
13	quartz and varied-	23	35	140	100	190	NF	5	0.40
12	texture diabase	100	31	71	110	130	430	5	0.15
11	(13-6)	130	53	190	90	170	150	8	0.14
10		150	NF	140	140	120	430	5	0.12
9		27	47	180	140	110	<20	7	0.20
8		21	39	140	100	150	<20	6	0.19
7		28	45	140	90	110	<20	9	0.11
6		27	39	140	70	190	<20	8	0.39
5	hypersthene diabase (5-1)	28	160	540	90	31	490	5	1.3
4		130	34	63	100	150	350	5	0.10
3		150	43	77	90	190	250	14	0.09
2		130	55	220	110	180	230	34	0.25
1		93	32	90	110	220	210	17	0.15

determined by Hriskevich (1952) to be 139 ppm Ni, 46 ppm Co, and 76 ppm Cu. Colwell (1967) estimated the average to be 81 ppm Ni, 44 ppm Co, and 96 ppm Cu. No average values have been assigned in the present study.

Granophyric diabases

Moore (1956) concluded that granophyric diabase contains more silver than ordinary diabase, and this was checked in the present study by analyzing 16 additional samples of coarse-grained and granophyric diabases (Table 92). Most of the samples are from around Elk Lake, this area being notable in that granophyres are common, but economic veins are rare.

In addition to the data given in Table 92, mention should also be made of the results reported by George (1968) for the exploration program carried out on the Argentium Silver property in Whitson township, approximately 18 miles (29 km) south of Elk Lake. In this program a series of north-south lines 400 feet (~120 m) apart were sampled at 25-foot intervals, and 281 samples of diabase were analyzed by atomic absorption spectroscopy. From the description given by George (1968), there is no question that well-developed granophyric diabase is present (though the proportion is not known). The arithmetic mean content of silver in the diabase was 0.7 ppm, the range being from 0.0 to 7 ppm. No vein was visible in the 7 ppm sample area. For cobalt the maximum was 400 ppm and the mean 34 ppm; for nickel 160 ppm and 43 ppm respectively.

ELEMENT VARIATIONS

Chromium and nickel

Chromium shows some well-defined trends which are readily apparent from inspection of the trace element data: (1) samples from the elevated portions of sheets, such as the North Lorrain, South Lorrain, and New Lake diabases, are characterized by being relatively low in chromium; (2) the depressed or basin parts of sheets typically have a thick zone of hypersthene diabase which is rich in chromium; (3) in all the intrusions the over-all trend is for chromium to decrease toward the interior of a sheet where the most differentiated diabase varieties occur. Thus, the lower parts of sheets which contain abundant orthopyroxene ($\pm$ olivine) generally contain 200 to 400 ppm Cr whereas lower values are present in the quartz and varied-texture diabases. This type of distribution is entirely consistent with the geochemically known relationship that chromium is

TABLE 91. TRACE ELEMENT CONTENTS (PPM) OF CHILLED NIPISSING DIABASE.

Area	Locality	No.	Position	Ni	Co	Cu	Ba	Cr	B	Ag	As
Cobalt	East side of Nicol Lake basin ;	68-143	8 cm from top contact	130	48	150	76	250		0.12	<4
	contact with Lorrain granite	68-149	15 cm from bottom contact	91	34	110	110	190	10	0.09	<4
		68-144	45 cm from top contact	82	31	100	100	190	7	0.15	<4
		68-145	~ 2 m from top contact	69	30	100	130	200	5	0.41	<4
		68-150	middle of sheet	56	32	100	170	62	9	0.15	<4
Cobalt		68-151	middle of sheet	40	36	120	170	20	10	0.23	<4
	Silver Summit property ;	69-46B	absolute chill	68	32	110	130	160	9	0.17	<4
	bottom contact	69-46M	12 cm above 46B	72	29	110	120	170	5	0.11	<4
	with Cobalt sediments	69-46T	25 cm above 46B	81	30	100	91	190	5	0.18	<4
	Nipissing RL 407, 300 m west of 407 shaft ; bottom contact with Cobalt sediments	69-385T	20 cm above contact (partly altered and albitized) ~ 15 m above contact	100	37	38	33	230	5	0.27	<4
Cobalt		69-386		160	70	97	130	220	14	0.13	<4
	Bucke tp. ; bottom contact with Cobalt sediments	68-1	albitized basal chill	90	36	12	140	230	NF	0.11	<4
		68-2	20 cm above 68-1	99	35	39	29	260		0.11	<4
		68-6	30 cm above 68-1	84	34	88	140	190	<5	0.31	<4
		68-5	50 cm above 68-1	82	36	79	150	170	10	0.30	<4
Gowganda		68-7	2 m above 68-1	76	30	110	140	180	6	0.66	<4
	Lower Bonsall mine, Miller Lake basin ; bottom contact with Archean	68-332	albitized absolute chill	84	40	87	330	140	6	0.48	15
		68-322B	6 cm above contact	90	34	110	460	140	12	0.50	10
		68-333	1 m above contact						8	0.12	8
		68-335	5 m above contact	110	28	60	450	150	21	0.47	<4
		69G-20	absolute chill	80	62	120	77	130	10	0.75	
	Bottom contact in drill core, ~ 800 ft. (245 m)	69G-21	15 cm above contact	150	120	250	75	160	13	0.41	
	south of Lower Bonsall shaft	69G-22	35 cm above contact	70	31	74	140	230	14	2.1	
		69G-23	Archean "intermediate" flow, 8 cm below contact of 69G-20	96	26	170	280	360	11	0.13	
		69G-24	15 cm below contact	110	28	97	290	360	11	0.07	
	Top contact with Archean, Siscoe mine, near No. 6 shaft, 350 level	69G-25	30 cm below contact	96	27	110	190	320	11	0.13	
		69G-19		90	43	110	210	110	12	0.58	19

TABLE 92. TRACE ELEMENT CONTENTS (PPM) OF COARSE-GRAINED DIABASE AND GRANOPHYRE FROM THE COBALT-GOWGANDA REGION.

Area	Locality	No.	Type	Ni	Co	Cu	Ba	Sr	Cr	B	Ag	As
Cobalt	Lot 15, conc. III Coleman township	68-33	granophyre	49	19	57	610		39	NF	0.22	<4
	West side of Anima Nipissing Lake	68-462	granophyric	15	45	54	140	90	NF	6	0.12	<3
	Lot 4, conc. IV James township	68-207	varied texture, granophyric	30	48	70	110		NF	10	0.47	8
	Shane Lake, lot 11, conc. IV,	68-289	granophyric	19	51	140	190	70	<20	<5	0.32	3
	James township	68-282	leucocratic	14	21	1000	110		NF	NF	0.31	<3
Elk Lake		68-284	granophyre	18	44	17	17	30	<20	<5	0.06	<3
		68-285	granophyre	12	25	90	90	30	NF	<5	0.90	38
	Silverclain Lake	68-343	granophyric	60	76	130	580		<20	6	0.57	25
	Mickle township	68-344	coarse,	43	54	48	200	90	<20	8	0.14	<3
	claim MR 16319		not granophyric									
Gowganda	Claim MR 12890	68-346	varied texture	37	54	100	220	70	<20	10	0.56	15
	Claim MR 16294	68-347	granophyric	21	28	87	160	<30	NF	5	4.0	8
	Claim MR 16294	68-350	granophyric	19	10	240	50	30	<20	<5	>5	8
	Drill core, Cotley	68-351	granophyric	18	34	18	410	80	<20	13	0.22	3
	Mines property	68-352	granophyric	18	11	55	98	30	<20	<5	>5	88
Gowganda	Morrison property	68-322	granophyric	56	54	200	870		<20	12	0.70	<3
	Coleroy property	68-315	granophyre	15	37	24	71		NF	<5	0.76	20

NF : not found ; blanks : not determined ; As : determined colourimetrically, with limits stated as < 3 and < 4 ppm as indicated.

incorporated into early-formed spinels and pyroxenes. With progressive fractionation, therefore, chromium is depleted as is shown by a strong correlation with the mafic index. Among the various phases of the Nipissing diabase, granophyres typically have the highest mafic index and are characteristically low in chromium.

The relationships described above are exemplified by the diabase in Henwood township. For those rocks which have been chemically analyzed there is a good correlation between the mafic index and chromium content. In the Lower sheet (Fig. 208) the initial rapid drop in Cr content levels off at about 100 ppm when the mafic index is between 57 and 62, and then falls to < 20 ppm Cr before an index of 65 is reached. This levelling off appears to be absent in the Henwood diabase basin; in the depressed part of the sheet, chromium reaches a maximum of over 600 ppm in the lower part of the hypersthene diabase zone and decreases toward the more salic differentiates and the quartz diabase margins. Although there is a scatter in the points (Fig. 208), the break evident in the Lower sheet seems to be absent in the Upper sheet. That the scatter in the individual points probably reflects a movement, during crystallization, of mafic material toward the depressed part of the sheet is suggested by the over-all lower mafic index and higher chromium content of the basinal rocks as compared with those of the elevated Flank.

Average nickel abundances in quartz-normative tholeiites are generally in the 70-76 ppm range (Prinz 1967). The element is typically enriched in early olivines and pyroxenes, and markedly decreases with fractionation.

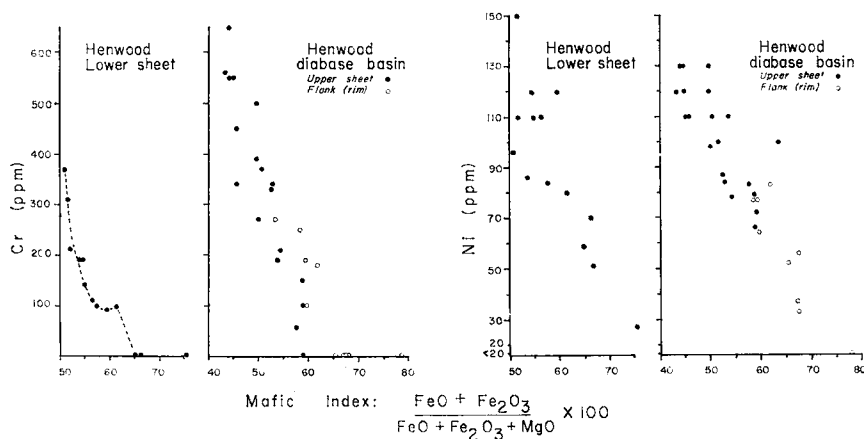


FIG. 208. Variation of Cr and Ni in the Henwood diabbases.

Nickel abundances in the Henwood intrusions are related to the mafic index as shown in Figure 208. In both the diabase basin and the Lower sheet, the highest nickel contents are present in rocks having the lowest (least fractionated) mafic index. These rocks are typically the hypersthene diabases. In the Flank section of the diabase basin most of the rocks are relatively high in mafic index and correspondingly low in nickel.

In addition to the data accumulated by the present writer, two cross-sections of the diabase at Cobalt were spectrographically analyzed by Colwell (1967). These sections are from the Nasco property in the North Lorrain basin (Fig. 202) and the Rix-Athabasca property (about 305 m east of the Colonial as shown in Fig. 204). Colwell's data have been used to construct Figure 209, which shows Ni versus mafic index for the two sections. The mafic index in this case is the ratio of total iron as FeO divided by the sum of this FeO plus MgO (with the product multiplied by 100).

In the present study the mafic index is not available for most of the diabase sheets which have been analyzed for their trace element content. It was, however, previously shown in Figure 208 that a relationship between chromium abundance and mafic index is present in the diabases, and on this basis chromium and nickel should also be inter-related. This is shown

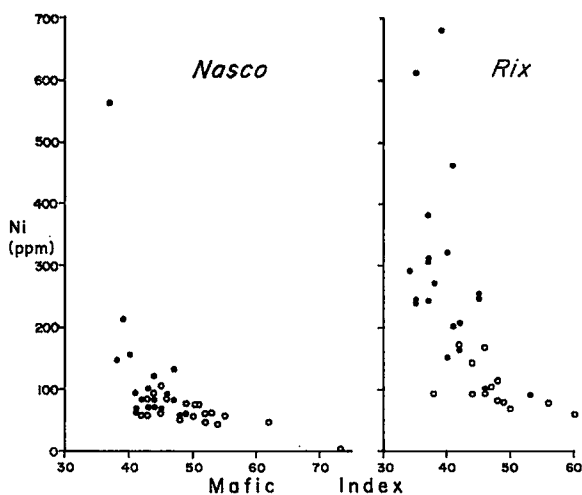


FIG. 209. Variation of nickel with mafic index in the Nasco drill core and Rix-Athabasca cross-section, Cobalt. Open circles represent samples above the hypersthene diabase zone; filled circles are samples extending from the top of this zone to the bottom contact.

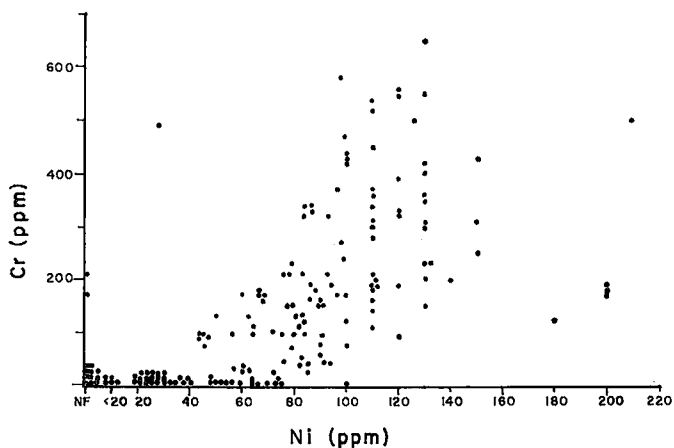


FIG. 210. Relationship of nickel and chromium in diabase in the Cobalt-Gowganda region.

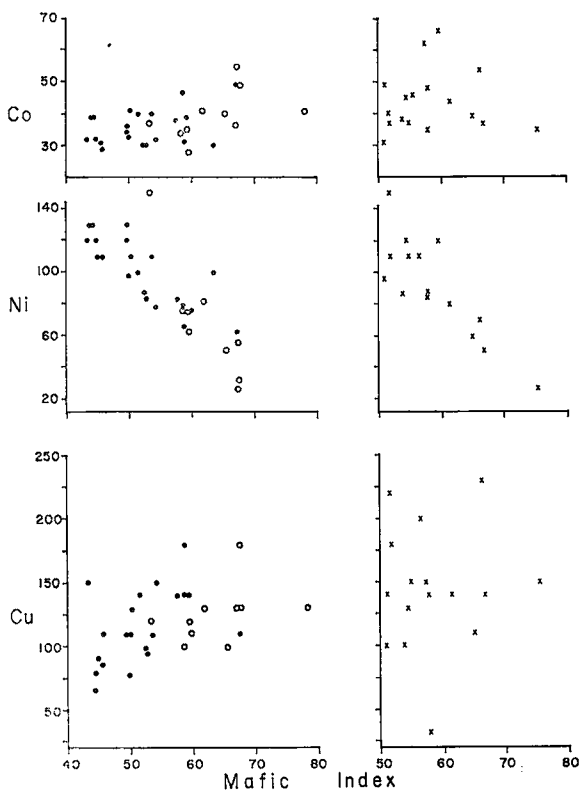


FIG. 211. Cobalt, nickel, and copper (ppm) versus mafic index of the Henwood diabase basin (left) and Lower sheet (right).

in Figure 210. The over-all impression is that nickel is closely related to the early rock-forming minerals, especially the oxides and pyroxenes.

Because alteration is almost, if not always, present in the chilled diabases of the region, no specific trace element values have been assigned to the chills. Nickel abundances in most cases, however, fall within a range of 75 to 100 ppm. The apparent normality of nickel in some thoroughly albitized chills (Table 91) is surprising.

Cobalt

The average cobalt content of the Nipissing diabase is less than 40 ppm. Most of the values higher than this were obtained from varied-texture diabases and quartz diabases. The exception to the generalization is the Miller Lake diabase at Gowganda, where half the analyzed samples (Table 90) exceed 40 ppm Co.

Cobalt decreases with fractionation in most mafic intrusions (Wager & Mitchell 1951; Nockolds & Allen 1956; Tiller 1959; McDougall & Lovering 1963; Rösler & Voland 1965; Greenland & Lovering 1966). The variation in cobalt with mafic index in the Henwood intrusions is shown in Figure 211. In the Lower sheet the two highest values (66 and 62 ppm) occur at the top and bottom contacts. In the Henwood diabase basin, both

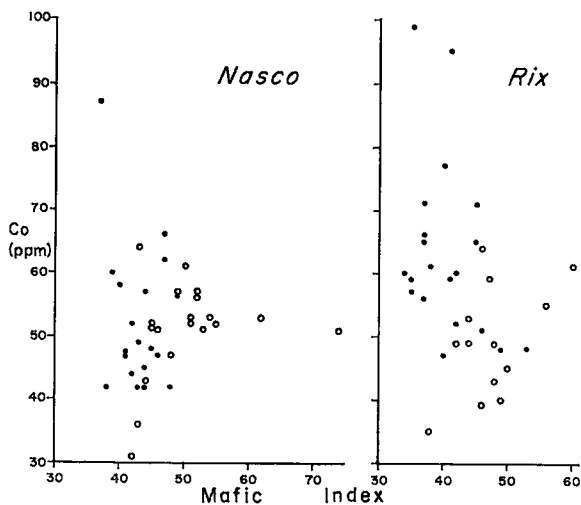


FIG. 212. Cobalt versus mafic index in single cross-sections of the Nasco and Rix-Athabasca diabases, Cobalt. Open circles represent samples above the hypersthene diabase zone; filled circles are samples from the top of this zone to the bottom contact.

the Upper sheet and Flank sections show a slight increase in cobalt with increasing mafic index. The most salic parts of the intrusion are thus enriched in cobalt, this being opposite to what is normally found.

Colwell's (1967) data for the Nasco and Rix sections have been plotted against the mafic index as was done for nickel. Considerable scatter is present in the Nasco data (Fig. 212), but the Rix section more clearly

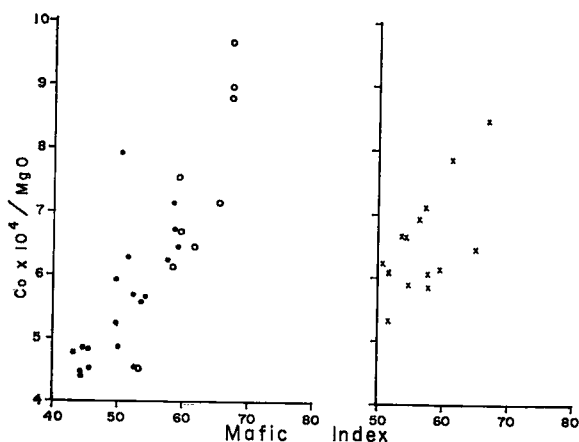


Fig. 213. Variation of Co/MgO with mafic index in the Henwood diabase basin (left) and Lower sheet (right). Filled circles are samples from the Upper sheet intersected in the Eplett vertical drill hole; open circles are Flank samples.

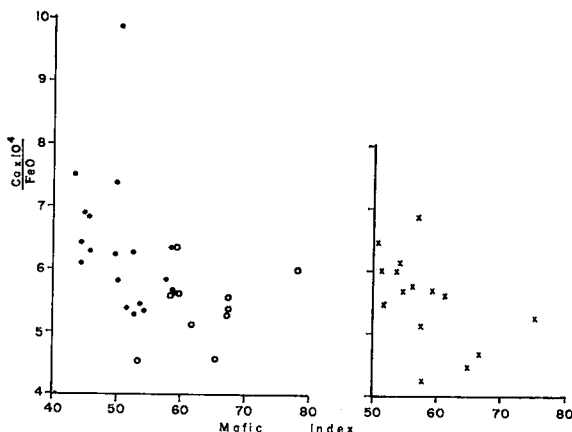


Fig. 214. Variation of Co/FeO with mafic index in the Henwood diabase basin (left) and Lower sheet (right).

shows that the main trend is for cobalt to decline with progressive fractionation. Because this is not the trend found in the Henwood diabases, data from the rocks occurring in the upper parts of the Rix and Nasco sections were distinguished from those in the lower parts; in Fig. 212 the rocks above the hypersthene diabase zone are shown as open circles. The resultant differences indicate that cobalt does not behave uniformly throughout the intrusions.

The association of cobalt with magnesium and iron in the Henwood diabases is shown in Figures 213 and 214. In both the Lower sheet and diabase basin, a more coherent trend is present with magnesium than iron. The slope of the Co/MgO trend steepens to the extent that it is almost vertical at high mafic index values characteristic of varied-texture and granophyric diabase. Throughout the Cobalt-Gowganda region individual specimens of these rock types have been found to be either relatively high or low in cobalt, presumably reflecting the presence or absence of erratically-distributed sulphides which are very commonly megascopically visible. The possibly significant aspect is that high cobalt values occur more frequently in the varied-texture zone than in hypersthene diabase.

Copper

McDougall & Lovering (1963) and Greenland & Lovering (1966) have shown that copper in Tasmanian dolerites markedly increases to a peak in silicic dolerites and then decreases in the granophyric phases. Insufficient data are available to explain the nature of the copper behaviour of the Nipissing diabases, but as a general rule the element has a somewhat irregular distribution because of its common occurrence as a sulphide phase. Some Nipissing granophyres contain megascopic disseminated grains of chalcopyrite, but in others the copper content is very low.

Copper in the Henwood township Lower sheet is considerably erratic with respect to the mafic index (Fig. 211), and position within the sheet (Fig. 216). In the Henwood diabase basin, however, the over-all trend is for copper to increase as the mafic index increases. The scatter along this trend is considerable (Fig. 211). On the other hand, plots of copper versus height in the diabase basin show that some relatively orderly features are present. In the depressed part of the basin intersected by the Stone-Eplett No. 3 and Eplett drill holes, copper increases toward the top of the sheet (Figs. 215, 217). The sequence in the sheet is for copper to decrease upwards as the basal quartz diabase passes into the hypersthene diabase; not until the upper half of the sheet is reached is there any marked change, but as the varied-texture zone is approached and entered, the

frequency of relatively high copper values increases and the over-all distribution becomes considerably more erratic. This pattern of behaviour also occurs at the upper contact, and is presumably an indication of copper sulphide crystallization. Although relatively few points are available for the Flank section, a slight enrichment in copper appears to be present in this part of the intrusion.

Silver

Reference has already been made to the earlier work by Moore (1934, 1956), Fairbairn *et al.* (1953), and Boyle (1968) on silver determinations in the Nipissing diabase. Boyle (1968) has summarized and added to the scanty data available for silver abundances in igneous rocks. In addition, Nesterenko *et al.* (1969) have reported that traps of the Siberian platform

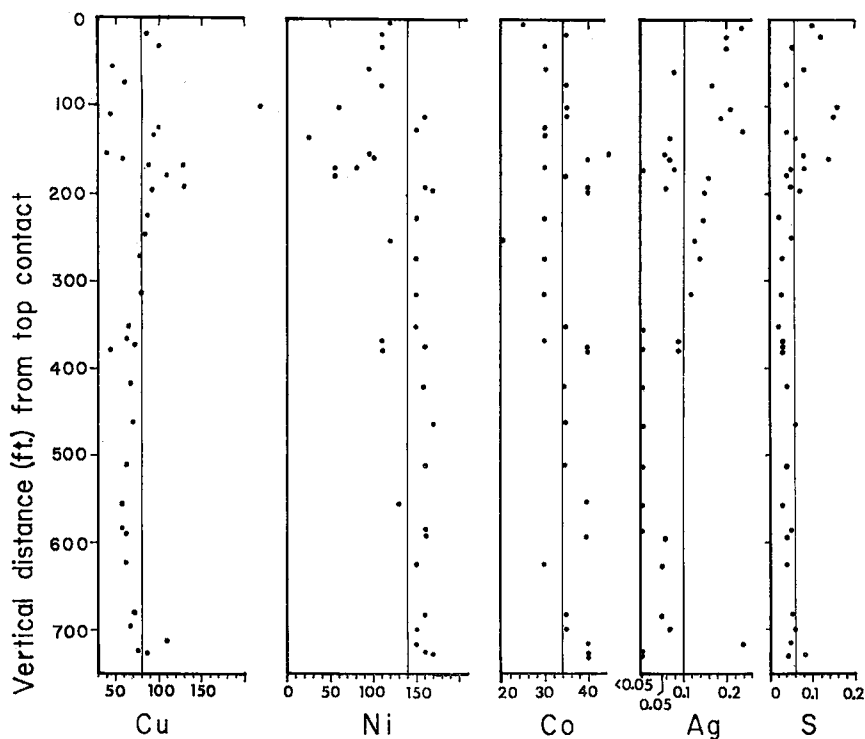


FIG. 215. Variation in Cu, Ni, Co, Ag (ppm) and S (%) in the Stone-Eplett No. 3 drill core, Henwood township. Vertical reference lines are arbitrarily positioned.

contain an average of 0.037 ppm silver, with 0.028 ppm in basalts and 0.042 ppm in gabbro-dolerites.

As has also been mentioned, absolute data for the Stone-Eplett No. 3 drill hole are not directly comparable with results obtained in this study because of interim revisions in the analytical procedures. The trace element trends should nevertheless remain valid. Silver variation in the Stone-Eplett No. 3 drill core is shown in Figure 215. The profile is similar to that of copper, with both elements being least abundant in the hypersthene diabase zone and increasing toward the top of the sheet.

In the Henwood Lower sheet (Fig. 216) the increase in silver toward the top is again evident. Maximum abundances occur in the varied-texture zone, and a single anomalously high value is present near the top contact.

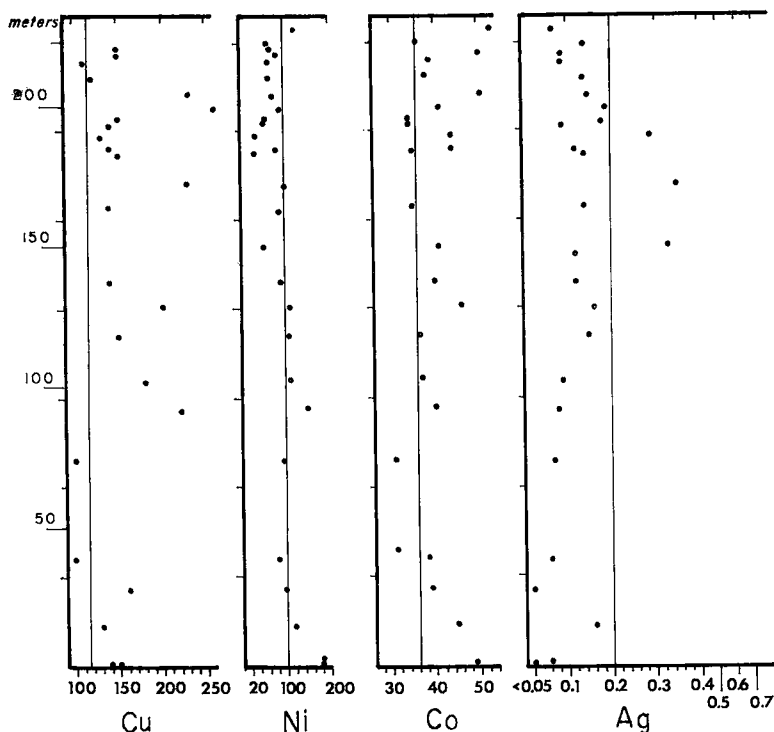


FIG. 216. Variation in Cu, Ni, Co, and Ag (ppm) with height above the bottom contact of the Henwood Lower sheet (subsidiary scale in hundreds of feet). Vertical reference lines may be used for comparison with data for the overlying diabase basin (Figure 217).

The distribution of silver in the Henwood Upper sheet is relatively orderly (Fig. 217), with maximum abundances occurring near the top of the sheet. The trends obtained from the Eplett and Stone-Eplett No. 3 drill holes are approximately congruous. Silver abundance in the Flank section is evidently greatest in the most fractionated rock, with the over-all abundance being similar to that of the Upper sheet. Within the limitations of the sampling, silver in the basin as a whole appears to have reached its peak abundances not only in the varied-texture zone, but also at the top contact of the depressed part of the sheet.

Among the silver profiles of other intrusions in the region, that of the New Lake basin at Cobalt is outstanding. The general trend is again for silver to be least abundant in the hypersthene diabase zone (Table 83),

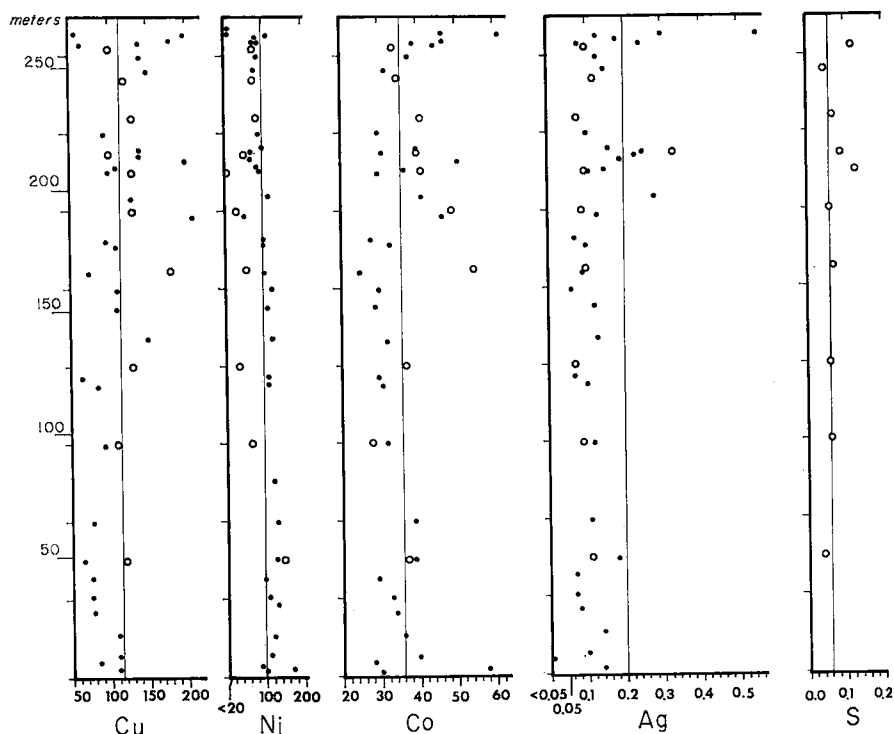


FIG. 217. Variation in Cu, Ni, Co, Ag (ppm) and S (%) with height above the bottom contact of the Henwood diabase basin (subsidiary scale in hundreds of feet). Vertical reference lines are at the same positions as those for the Lower sheet (Figure 216). Open circles represent samples from the Flank section and filled circles represent samples from the Upper sheet as interested in the Eplett drill hole.

but in all parts of the intrusion silver abundances are at a relatively high level, particularly in the varied-texture zone. In the adjacent Peterson Lake basin, however, the silver values are normal to low. The meaning of these variations is not readily evident, but it is clear that the first necessity is to obtain additional profiles along new traverse routes in order to determine whether the trends found here are consistent, characteristic features.

Antimony and arsenic

The average content of antimony in igneous rocks is 0.2 ppm, and 0.1-0.2 ppm in basic rocks (Wedepohl 1969). Data for antimony in the Nipissing diabase are available only for samples of the Stone-Eplett No. 3 drill core. The results (Table 85) indicate that the average antimony content of the diabase is less than 1 ppm, and most of the element is concentrated near the top of the sheet.

The average content of arsenic in igneous rocks is 1.5 ppm As (Wedepohl 1969). Arsenic in the Skaergaard intrusion (Esson *et al.* 1965) averages less than 1 ppm As in all rock types. The arsenic contents of the Nipissing diabbases are also low. Of twenty specimens from the Peterson Lake basin, Cobalt, the only one above the 4 ppm lower limit of detection is JF 69C-30 (Table 84). This specimen, which is close to the upper contact, was found to contain 70 ppm As. In the Stone-Eplett No. 3 and Eplett drill holes, only six analyses are above the detection limit; two occur in the varied-texture zone (Tables 85 and 87), and three near the diabase contacts. Detectable arsenic is common in granophyric diabase (Table 92).

Sulphur

Sulphur analyses are available only for the Henwood diabase basin. The element remains fairly constant in the lower half of the Stone-Eplett No. 3 drill core (Fig. 215), and reaches its maximum abundance in the varied-texture zone near the top of the sheet. The few analyses available for the Flank indicate that sulphur has a similar distribution but is slightly enriched in comparison to the depressed part of the sheet. Sulphur thus appears to have migrated to the most fractionated and the most elevated parts of the intrusion.

Strontium and rubidium

The strontium values in the diabase sheets throughout the region are relatively uniform in that nearly all determinations fall within the range

of 50 to 150 ppm Sr. In a drill hole which passed through the sheet in the Miller Lake basin, Gowganda, Fairbairn *et al.* (1969) found Sr to be fairly constant at about 140 ppm. Samples collected from the same sheet in the present study give lower Sr values (Tables 90), but the data are comparable relative to the much higher median of 400 ppm Sr for quartz-normative tholeiites surveyed by Prinz (1967).

Strontium in diabase is generally considered to substitute in calcium-rich minerals, especially the calcic plagioclase feldspars. Accordingly, the Nipissing granophyres, though they contain large amounts of plagioclase, are not enriched in strontium because they contain albite as the principal feldspar. There is, on the other hand, no obvious coherence between Sr and Ca in the Henwood diabbases (Fig. 218), a situation which has also been noted by Gunn (1966) in Antarctic tholeiites. In the Henwood Lower sheet strontium does increase with fractionation measured in terms of the mafic index (Fig. 219), but there are also strong indications that strontium abundances differ from sheet to sheet. This is especially evident in Figure 220, which shows that the Henwood Lower sheet contains more strontium than the overlying diabase basin regardless of the prevailing rock type. This difference is considered to be additional evidence that the diabase basin and Lower sheet are separate intrusions.

Rubidium analyses have not been tabulated, but the data for the Henwood diabbases are shown in Figure 221. The element increases in abundance toward the tops of the sheets. As is the case with strontium, the values obtained in this study are lower than those reported by Fairbairn *et al.* (1969) for the Miller Lake sheet; in all cases, however, the

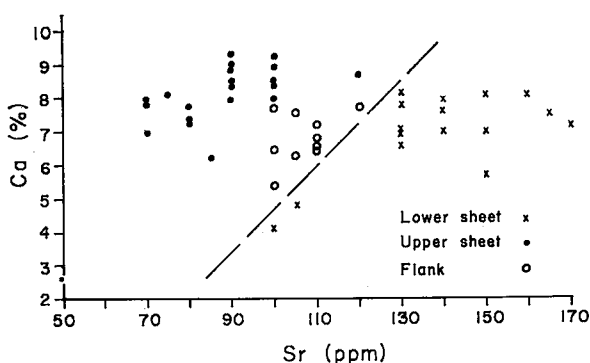


FIG. 218. Calcium versus strontium in the Henwood diabbases. The dashed line separates data for the Lower sheet from data in the overlying diabase basin (Upper sheet plus Flank).

trend is for Rb to increase in abundance with height. In the Henwood sheets, the general trend of rubidium and potassium coherence is evident from Figure 221. Although rubidium enters the K-position in micas in preference to feldspars, the previously demonstrated sympathetic increase in K_2O with increasing Ab contents of the plagioclases may account for a part of the potash and concomitant rubidium increases with height.

Barium

Although barium values in the Nipissing diabase throughout the Cobalt-Gowganda region commonly seem to be erratic, the New Lake and Peterson Lake sheets show that in some cases there is a fairly well-defined increase in barium with fractionation. This relationship is particularly evident in the Henwood diabase basin (Fig. 219). The Upper sheet has the lowest Ba/ K_2O ratio and shows the closest coherence with potassium (filled circles, Fig. 222). For rocks containing less than 1 per cent K_2O , barium in the Lower sheet is consistently more abundant than in the Upper sheet. The Flank section contains more barium than the depressed part of the basin, with this effect being in part attributable to the fact that more plagioclase of intermediate composition occurs in the Flank. Substitution of Ba for K in such feldspars may be important as they are relatively enriched in K_2O .

Boron

Analyses for boron in the present study may seem to be an incongruous choice in view of the limited number of elements investigated. However,

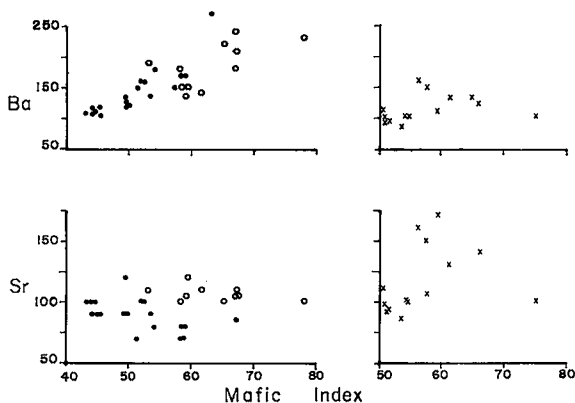


FIG. 219. Variation of strontium and barium with differentiation in the Henwood Lower sheet (right) and diabase basin (left, with solid circles representing the Upper sheet and open circles the Flank).

the boron silicate axinite has been noted in several ore-bearing veins, and the mineral also occurs in post-ore faults and in pre-ore veins in the Archean. The behaviour of boron with diabase fractionation was therefore considered to be germane.

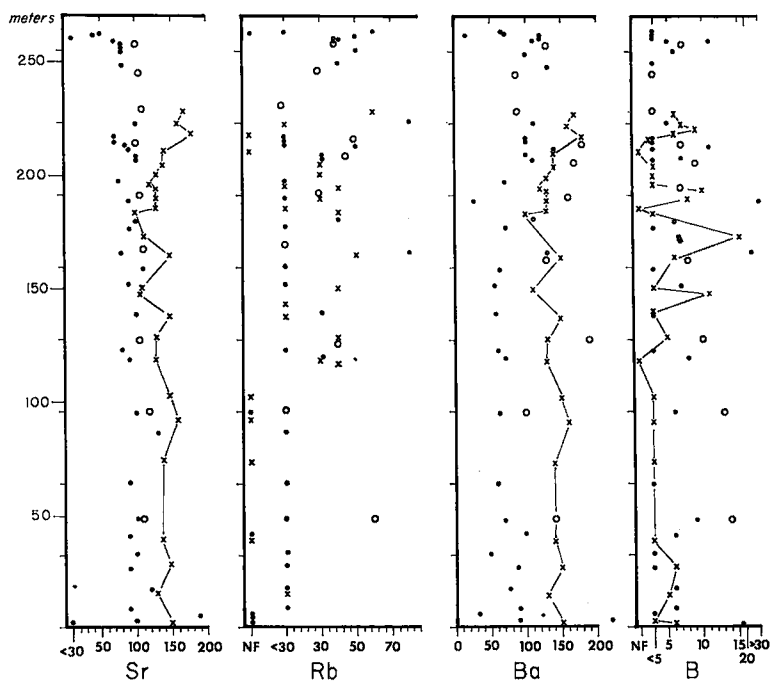


FIG. 220. Sr, Rb, Ba, and B (ppm) versus height above the bottom contacts of the Henwood diabases (subsidiary scale in hundreds of feet). (Lower sheet x; Upper sheet •; Flank O).

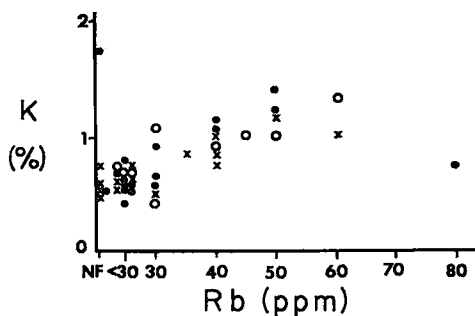


FIG. 221. Rubidium versus potassium in the Henwood diabases. Symbols are as given for Figure 220.

In the Palisades sill, Walker (1969) noted that boron was concentrated in residual melts, with values above 20 ppm occurring only in rocks of later fractionation stages. In the diabases of the Cobalt-Gowganda region a single general pattern seems to be absent. In some intrusions, such as the Peterson Lake diabase, boron does show an increase toward the top of the sheet. A slight upward increase is also present in the Henwood sheet (Fig. 220), but is not readily apparent in the Upper sheet of the overlying basin. The Flank, however, is enriched in boron. This enrichment lessens upward, and in this respect the trend is similar to that of Al_2O_3 in the Flank. The over-all behaviour of boron in the Henwood intrusions suggests that the element is not incorporated into the early-formed minerals. At a middle stage of fractionation, however, the element is rapidly extracted from the magma, probably as (BO_4) replacing (AlO_4) in the intermediate plagioclases.

SIGNIFICANCE OF THE MINOR ELEMENT DISTRIBUTION

The principal purpose of the study of the minor elements in the Nipissing diabase was to determine whether the diabase could be reasonably eliminated as a potential source of the metals present in the ore veins.

No element has remained static during the fractionation of the diabase. Because more chemical and minor element information is available for the Henwood diabases, discussion of the variations has centered on these intrusions. The gross relationships for many elements nevertheless appear to be generally applicable to the other diabase basins in the region. The information from this reconnaissance study indicates that the most pro-

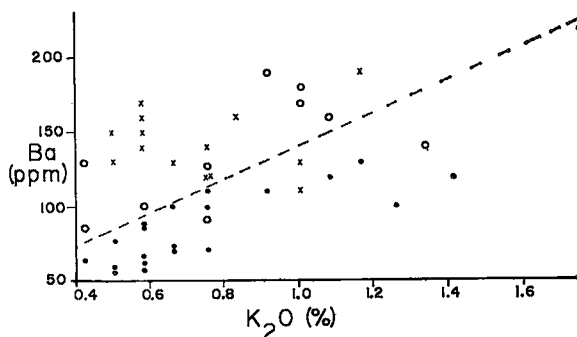


FIG. 222. Barium versus potassium in the Henwood diabases. Symbols are as given for Figure 220. All points for the Upper sheet (filled circles) fall below the dashed line.

fitable results in future work might be obtained by focusing additional attention on the intrusions at Cobalt, namely, the New Lake and Peterson Lake basins.

The most abundant constituents of the ore minerals are cobalt, nickel, iron, copper, silver, arsenic, and sulphur. Antimony is also commonly present, but it is not a principal component and is unimportant in comparison to arsenic. Bismuth is present in some veins, but its distribution has not been studied, partly because the element is inconsequential relative to the above components. Although iron was not discussed in this paper, its variation in the Henwood intrusions is shown in Figures 26 and 27 (p. 68 & 69). For the important ore elements, the basic pattern which has emerged from the present study can be summarized as follows :

1. Of the ore elements Co, Ni, Fe, Cu, Ag, As, S, and Sb, all except Ni either increase with diabase fractionation or have migrated to the upper parts of the intrusions.

2. The most common places for enrichment anomalies in the diabbases are near the contacts, and in the varied-texture zone.

Although it is recognized that the data for antimony in particular are sparse, the above-cited relationships indicate that all of the ore elements were mobile during diabase fractionation. The distribution of the elements has, however, been governed by two somewhat different, though related, processes. The first of these may be broadly referred to as the initial magmatic stage in which the main rock-forming minerals crystallize, and the second is the deuteric to hydrothermal stage in which many of the original minerals are partly altered or changed to new minerals which are commonly hydrous and characteristically represent lower temperature species. It has been shown that nickel methodically decreases with fractionation, and this behaviour militates against its being concentrated in late-stage veins. There is, however, the problem of accounting for the nickel content of granophyric diabase and the presence of erratic values in the varied-texture zones of the intrusions. This type of variability initially suggests that some nickel may be present in "sulphide" form. To test this possibility, the ascorbic acid leach method (Smirnova & Al'mukhamedov 1967 ; Smirnova *et al.* 1968) was used on the diabase samples from the Peterson Lake basin, Cobalt. The leach method used in this study is described in complete detail in Cameron *et al.* (1970). The treatment utilizes an ascorbic acid-hydrogen peroxide mixture which preferentially attacks sulphides and native metals rather than silicates and oxides.

The results of the leach treatment of the Peterson Lake samples are given in Table 93. Because it was necessary to determine nickel in the resulting solutions by atomic absorption spectroscopy, the nickel contents

TABLE 93. ASCORBIC ACID-HYDROGEN PEROXIDE LEACH TESTS OF DIABASE FROM THE PETERSON LAKE BASIN, COBALT.

Sample	HF-HClO ₄ Attack	Ascorbic Acid-H ₂ O ₂ Leachable Fraction	Sample	HF-HClO ₄ Attack	Ascorbic Acid-H ₂ O ₂ Leachable Fraction
JF69C-	Ni (ppm)	Ni (ppm) %	JF69C-	Ni (ppm)	Ni (ppm) %
30	113	66 58	46	181	80 44
31	113	82 73	47	181	52 29
32	130	48 37	48	202	64 32
33	116	48 41	49	146	64 44
37	126	62 49	50	142	58 41
38	133	56 42	51	172	59 34
39	129	60 47	52	192	50 26
41	81	45 56	53	181	42 23
42	160	51 32	54	184	48 26
44	124	50 40	55	175	47 30
45	155	56 36			

Analyses by Geochemistry Section, GSC.

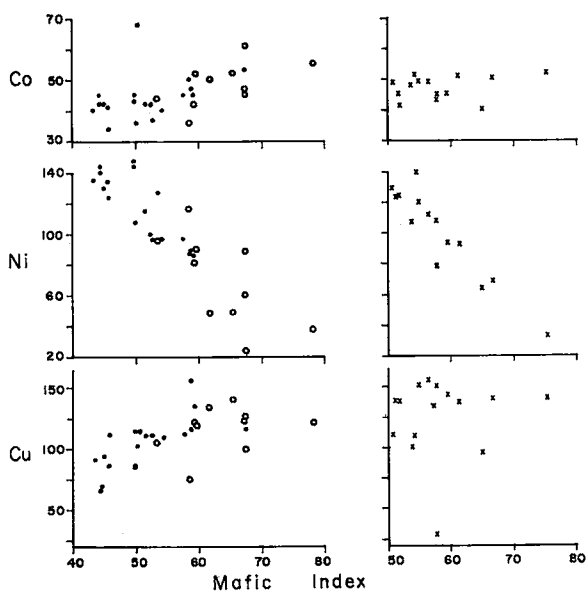


FIG. 223. Variations in mafic index versus Co, Ni, and Cu (ppm) in the Henwood diabases, with trace elements determined by atomic absorption analyses.

(Upper sheet • ; Flank O ; Lower sheet x).

of the untreated samples were also determined by an $\text{HF-HNO}_3\text{-HClO}_4$ attack followed by atomic absorption spectroscopy. Without exception, the nickel results obtained by the A.A. method were higher than those obtained by optical spectroscopy. The new results ranged from +11 to +52 per cent higher, the average being +22. Because of this, the Henwood diabbases were re-analyzed by the A.A. method, but no significant changes in the trends occurred. The scatter of points was in some cases reduced (Fig. 223), but the distribution patterns in the Lower sheet and overlying basin remain essentially unchanged.

The results of the leach tests on the Peterson Lake samples are shown in Figure 224. Total nickel in the rock decreases with fractionation, but the percentage of the total which is present in "sulphide" form actually increases toward the top of the intrusion. The net effect is that the amount of nickel in "sulphide" form is fairly constantly around 50 to 60 ppm throughout the sheet. Thus sulphides account for most of the nickel in granophyric and other salic varieties of the diabase. The distribution of this element therefore reflects both the initial extraction of the element from the magma into the early-formed silicates and oxides, and also the

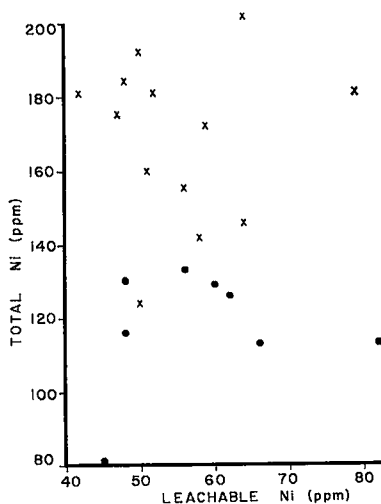


FIG. 224. Total nickel versus leachable nickel in the Peterson Lake diabase, Cobalt. Samples above the hypersthene diabase zone are shown by a solid circle; those in and below the hypersthene zone by an x.

relatively increasing importance of the sulphide form with progressive fractionation in the broadest sense.

In the Henwood diabases the indications are that the distribution of the ore elements is strongly affected by the configuration of the intrusion. The variation of Fe, Cu, Ni, Co, Ag and S in the diabase basin is most interesting because the information available shows that these elements have behaved differently; thus Ag may have possibly "peaked" near the top of the depressed part of the sheet (Fig. 217). Although the situation with respect to "sulphide" nickel is uncertain, the other elements seem to have susceptible to migration toward the elevated rim of the intrusion. This upward trend is readily apparent for Co, and is obvious for Fe.

All writers who have considered the relative proportions of the various elements in the ore veins have had difficulty in agreeing upon an "average" abundance of the major components. Bastin (1939), however, made much of the fact that at Cobalt the proportion of cobalt to nickel was greater than one, and hence the ratio was atypical of ores in basic rocks but more akin to ores associated with salic intrusions. This was one of several reasons why Bastin argued that the ores came not from the Nipissing diabase, but from a more salic magmatic parent.

The cobalt-nickel ratios of the diabase intrusions in Henwood township are shown in Figure 225. Fleischer (1968) has shown that Ni/Co normally decreases with fractionation, and this is also the trend in the Henwood intrusions; cobalt here eventually exceeds nickel in the most salic differentiates and the Co/Ni ratios ascribed to the ores are attained.

Metal abundance in the rocks intruded by the Nipissing diabase have not been investigated in the present study, but analyses of these have

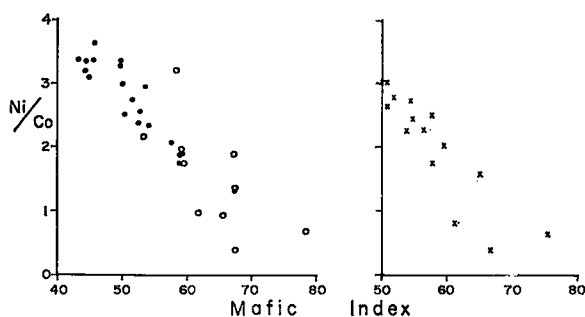


FIG. 225. Variation in Ni/Co ratios with the mafic index of the Henwood diabase basin (left) and lower sheet (right).

been published by Boyle (1968a, 1968b), Boyle *et al.* (1969), and Boyle & Johnston (1970). The present writer is grateful to Dr. Boyle for making available the samples used in compiling the data shown in Table 94. In connection with the previous discussion of the Stone-Eplett No. 3 drill core, it has already been mentioned that interim revisions in the analytical procedures for silver have occurred; the composite samples listed in Table 94 were therefore re-analyzed for silver in order to obtain a direct comparison with the diabases of the present study. The results obtained from two splits from each composite sample are given in Table 94. The data show that silver abundances in Archean volcanic and pyritiferous sedimentary rocks are not much different from those in the Nipissing diabase.

It was stated at the beginning of this section that the principal purpose of this study was to determine whether the Nipissing diabase could be eliminated as a potential source of the ore elements. It has been shown that the metals present in the ore veins are also present in the diabase. More important, it has been shown that the relevant elements have been mobile and have been concentrated in certain parts of the intrusions. It is concluded that the Nipissing diabase is a potential source of the elements found in the veins.

TABLE 94. ANALYSES OF ROCKS FROM COBALT AND SOUTH LORRAIN (AFTER BOYLE 1968a).

Description of samples	Ni	Co	Cu	Zn	Pb	Ag	As	Sb	S	Ag*
Composite sample of Keewatin greenstone, South Lorrain ¹	130	67	330	95	<5	0.30	15	2.5	1700	0.09 0.11
Composite sample of Keewatin greenstone, Cobalt area	105	69	120	215	<5	0.19	2	0.5	775	0.09 0.07
Composite sample of Cobalt conglomerate and greywacke, Cobalt area	60	25	20	20	<5	<0.05	<2	<1	180	0.05 <0.05
Composite sample of Keewatin black pyritiferous slate, schist, and greywacke, Cobalt area ²	160	74	420	2400	75	0.83	140	12.5	~6%	0.34 0.30

All values in ppm except where noted.

¹ Contains visible sulphides, including some chalcopryrite.

² Hudson Bay mine area, Cobalt.

* Re-analyzed for comparison with the diabases of the present study.