The pyroxenes of the alkaline igneous complexes of eastern Uganda

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Summary. Chemical analyses, some optical data, and specific gravities are given for 57 pyroxenes from the alkaline rocks of Uganda. They are members of the diopside-hedenbergite-acmite series, the more diopsidic varieties occurring in rocks of the ijolite series and the more acmitic in syenites and carbonatites. TiO₂ is highest in those pyroxenes from rocks that do not contain melanite. The refractive indices and specific gravities are controlled by the Fe^{2+}/Mg ratio in the diopsidic pyroxenes and by the Fe^{3+} content in the acmitic members. Paragenesis is discussed briefly.

THIS study of the pyroxenes forms part of an investigation of the alkaline complexes of eastern Uganda. Although considerable variations in optical properties obviously reflect variations in chemical compositions, not only are they difficult to determine accurately owing to the intense colour, but the existing graphs relating optical properties and chemical composition are based on largely inadequate data.

It has been usual to refer to the sodic pyroxenes as aegirine-augite and to regard them as members of the series diopside-aegirine (acmite). In the pyroxenes investigated, however, the third end member, hedenbergite, is of equal importance to the other two.

Mode of occurrence of the pyroxenes. The alkaline complexes of eastern Uganda (fig. 1) consist of two groups: an older group, of presumed Cretaceous age, in which only intrusive rocks are present—these are the complexes of Sukulu, Tororo, Bukusu, Sukululu, and Budeda; and a younger group, of Tertiary age, represented by a series of dissected volcanoes, Kadam, Elgon, Napak, Moroto, and Toror (King and Sutherland, 1960). At Napak, erosion has revealed a central plug of plutonic rocks, while at Toror, the volcanic rocks have been entirely removed, and only the intrusive equivalents are found.

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The chief rock types represented in the complexes are:

Volcanic: melanephelinite-nephelinite (nephelinite series); phonolitetrachyte.



FIG. 1. Sketch map showing the positions of the alkaline complexes in Eastern Uganda. Solid black, intrusive complexes; lined areas, volcanic rocks.

Plutonic: pyroxenite-melteigite-ijolite-urtite (ijolite series)-carbonatite; nepheline-syenite-syenite; and fenite (granite)-syenitenepheline-syenite.

The principal, and often the only mafic mineral in all of these rocks is a sodic pyroxene, although its abundance varies widely from pyroxenite in which it is almost the sole mineral to the urtites in which it is only sparingly present.

In the volcanic rocks, there is a marked contrast between the diopsidic pyroxenes of the nephelinites and the very aegirine-rich pyroxenes of the phonolites and trachytes. Because of this obvious distinction, and the comparative difficulty of separating the rather small amount of fine-grained pyroxene, a detailed study of the volcanic rocks has been deferred, and in the present account, attention is devoted to the pyroxenes of the plutonic series in which a continuous range of compositions is represented. The pyroxenes of the ijolite series are particularly important, since in these rocks a complicated sequence of events can be recognized (King and Sutherland, 1960; King, 1965). This sequence is: pyroxenite-melteigite-ijolite-pegmatite-urtite-wollastoniteurtite.

In addition, the pyroxenes of the nepheline-syenites, carbonatites, and fenites have been studied, but less comprehensively. The majority of the pyroxenes analysed are from the ijolite complex of Lokupoi (Napak), the ijolites and syenites of Budeda, and the carbonatites of Tororo. Four pyroxenes were also analysed from Semarule in Bechuanaland where alkali syenites and fenites occur (King, 1955). It has also been of interest to include reference to two new analyses made of pyroxenes from the Loch Borolan area of northwest Scotland and four from the alkaline rocks of Sørøy, northern Norway. A total of 63 new analyses of pyroxenes from alkali suites have been carried out in the present investigation. Previously published analyses of comparable pyroxenes are few in number, of which a high proportion are aegirinerich (Sabine, 1950; Pulfrey, 1950; Deer, Howie, and Zussman, 1963).

Chemical composition of the pyroxenes. Zoning is often common in the pyroxenes, and it was necessary therefore to select samples in which zoning was slight or absent. This restricted to some extent the range of samples available. The methods of analysis used were variations of the gravimetric, colorimetric, and flame photometric methods in common use. The results of the analyses, all by R. C. Tyler, are given in table I.

In figs. 2a and b, plots of $(Mg + Fe^{2+} + Mn)/Ca$ and $(Na + K)/(Fe^{3+} + Ti + Al^{iv})$ are shown from the calculations of the analyses to the formula $X_m Y_{2-m}(Si,Al)_2O_6$.

In fig. 3, the pyroxenes have been plotted in terms of the three principal end-members, diopside, hedenbergite, and acmite. The comparatively small amounts of the other constituents justifies the recalculation of the proportions of diopside, hedenbergite, and acmite to 100 %, whereby the compositions of the pyroxenes are shown as points. On this diagram, all the pyroxenes from the African alkaline rocks lie within a curved band extending from near diopside to near acmite. In the most acmitic pyroxenes, the hedenbergite content commonly exceeds that of diopside.

Among minor elements, the content of titanium in the pyroxenes shows

TABLE I. Analyses of pyroxenes from Budeda (B), Bukusu (Bu), Napak (N), Tororo (To) (all in Uganda), Semarule, Bechuanaland (K), Loch Borolan, Scotland (Bo), and Sørøy, Norway (SB).

	0 SIO ₈		0 A1203	7 FeO	2 MnO	0 MgO	5 CaO	6 Na.0	0 K.0	0 P _z O _s	- H ₂ 0+	3 Total	80 Si	20 Aliv	74 Alvi	26 Ti	58 Fe ³⁺	28 Fe ²⁺	12 Mn	79 Mg	68 Ca	77 Na.	70 K	15 X	77 Y	00 Z	21 a	۲0 v	۰° A: [001]	3 S.G.		ith melanite. te.		
N30	50.8	800	0.0	0.0L	8-0-	9 4 9 <u>6</u>	1.91	9.00	1-4	0.1	J	0.66	-+ 1.9	0.0	0.0	0.0	0.5	0.0	0.0	0.5	9·0	0. 19	0-0	1.0	6-0	2.0	1-7	1.7	20	3.4		l ijolite wi th melanit	0.043.	
N23	50-30	0-39	0.91	9-54	0.52	5.84	18.49	4.49	0.27	0.06]	100.49	1.941	0-058	I	0.011	0.269	0.309	0.016	0.336	0-767	0.334	0.012	1.113	0.941	2.000	1.719	1.759	21°	3.34	syenite.	rse grained I ijolite wit	4. Fe ^{3+(jv)}	L.
Bu414	51.22	TG-0	4-30	7.85	0.28	11.21	23.08	1.54	0.51	none		100-50	1.943**	I	i	[0.080	0.249	600-0	0.634	0.939	0.114	0.023	1.076	0.972	2.000	1.690	1.731	1	3.40	, Cancrinite 4. Melteigite	Medium-coa Fine-grained	lso Tiv 0-01	so Tiv 0.00
B282	49-42	cU-L	15-33	9-76	0.64	3.14	14.54	6.15	0.43	trace	[100.46	1-954	I	1	I	0.340	0.316	0.021	0.181	0.601	0.460	0.023	1.084	0.948	2.000	1.735	1.770		3-42	B282, Bu41	N23, N30,	• ** (A	++ A]
B101	46-08	TC:2	13-36	10-96	0.53	5.48	17-27	3.63	0.21	0.24	I	100.27	1.836	1	ł	I	0.312	0.366	0.017	0.326	0.721	0.278	0.010	1.009	1.021	2.000	1.720	1.758	25°	3.40		/ke.		
B96	50-39	01.1	13.85	10-25	0.32	3.16	12.30	6.25	0.19	0.06	0.04	100-36	1-945	0.055	0.052	0.035	0.404	0.332	0.012	0.131	0.508	0.468	600.0	0.985	1.016	2.000	1.740]	28°	3.48	h nevheline.	ite syenite d	iv) 0-057.	iv) 0.090.
B68	50-83	0.07 1-36	8.86 86.8	60.6	0.48	5.85	18.50	3.52	1.42	trace	[100.58	1.957	0.043	0.017	0.019	0.254	0.293	0.016	0.336	0.763	0.263	0.069	1.095	0.935	2.000	1.717	1.773	22°	3.38	e. tic fenite wit	eline cancrin red ijolite.	⁻ 0.030, Fe ³⁺ (7 0.074, Fe ³⁺
B38	49-59	0.91 2.43	7.88	12.05	0.48	5.23	17.61	3.87	0.45	0.24	1	100.34	1.934	0.066	0.047	0.014	0.230	0.394	0.016	0.305	0.720	0.291	0.023	1.034	1.006	2.000	1.719	1.760	22°	3.36	B38, Ijolite B68, Sveni	B96, Neph B101, Alte	§ Also Tiv	I Also Tiv
B33	49.10	0.50	8.60	10.90	0.42	7.37	20.06	2.60	0.48	0.02]	100.75	1.904§	600.0	1	1	0.194	0.354	0.014	0.427	0.834	0.196	0.023	1.053	0.989	2.000	1.705	1.746	20 1 °	3.40				
B29	49-12 0-40	0-18	9.48	10.50	0.42	6.76	19.12	2.62	66-0	0.14]	99-73	1.932_{1}^{+}	0.009	I	1	0.232	0.345	0.014	0.397	662.0	0.199	0-047	1.045	0.988	2.000	11711	1.753	21_{2}°	3-39				034.
B28	49-30 0-79	0.27	9.08	10.45	0.40	6.88	19.12	2.40	1.04	0.16]	99·88	1.929	0.014	1]	0.234	0.341	0.014	0.402	0.795	0.183	0.052	1.030	0.991	2.000	1.711	1.753	20^{16}	3.39		teigite.		3, Fe ^{3+(iv)} 0-
B7	50-29 1.93	none	12.84	10.94	0.34	4.68	15.26	4.36	0.84	trace)	100.78	1.955*]]]	0.373	0.355	0.012	0.271	0.635	0.327	0.042	1.004	1.012	1.990	1.725	1.780	L] 22 1	3.45	ljolite. Melteigite.	Melteigite. Biotite Mel	so Ti ^{iv} 0-03	so Tiiv 0-02;
	Si0 _s	Al.O.	Fe.0.	FeO	0uM	MgO	CaO	Na_2O	$\mathbf{K_{20}}$	P ₂ 0°	н 1 1 1 1 1 1	Total	Si	Aliv	Alvi	E	Fe ³⁺	Fe ²⁺	Mn	Mg	°.	Na	R	×	، بد	2	в	۔ ح	A: [00.	S.G.	B7,] B28,	B29, B33,	* A]	t Al

	N32	N35C	N35F	N48	N52	N62	N93	N95	N102	N103C	N103F	N107A	
03	50-00	51.10	50.73	49.81	50.40	50.29	50.75	50-30	51 - 36	49-07	49.83	50.10	SiO.
10,	1.10	1.36	2.35	0.75	0.60	0.86	1.01	1.44	1.08	2.49	1.99	0.59	Tio.
ul203	1.46	4.09	3.93	2.21	0.82	4.03	1.09	1.53	4-14	1.51	2.39	none	$AI_{s}O_{s}$
re203	7-70	3.10	4.49	6.57	5.87	2.90	4.37	24.17	21-14	29.69	26.70	15.02	$Fe_{a}O_{a}$
le0	13.90	3.71	3.97	26.7	9.17	6.07	4.32	5-75	5.00	0.34	1-44	11.35	FeO
4n0	0.71	0.11	0.19	0.22	0.27	0.13	0.18	0.44	0.16	0.29	0.45	0.46	MnO
lg0	2.23	12.24	11.32	8.75	9.47	11.91	12.56	0.78	0.66	1.73	1.06	3.32	MgO
a0	19.24	21.90	21.40	21.39	20.56	21.67	23·88	5.93	6.75	2.61	2-47	13.12	CaO
Va ₂ O	3.63	1.10	1.55	2.22	2.34	1.84	1.23	7-59	69-6	11.81	10.62	5.62	Na_2O
č_0	0.43	0.28	0.18	0.34	0.44	0.35	0.20	1.86	0.49	0.94	1.49	0.85	$\mathbf{K}_{s}\mathbf{O}$
2.05	none	0.01	0.03	0.15	0.32	none	none	0.30	0.03	none	0.13	0.04	$P_{2}O_{5}$
$I_2O +$		I		0.02	1	1	1	none	0.02	[[!	$H_{s}O+$
rotal 1	00·40	00-66	100.14	100-40	100.26	100.05	99-29	100.09	100-52	100.48	98-57	100.47	Total
. 19	1.963	1.908	1.879	1.902	1.929*	1.880	1-910†	1.954	1.955	1.8861	1.944	1.967	Si
١liv	0.037	0.098	0.121	0.098	0.037	0.120	0.050	0.046	0.045	0.069	0.056	none	Aliv
LI vi	0.029	0.081	0.053	0.003		0.059	1	0.054	0.142		0.052	1	Alvi
E	0.033	0.038	0.065	0.021		0.025	1	0.042	0.030	0.026	0.059	0.016	Ţ
re ^{s+}	0.227	0.085	0.125	0.188	0.154	0.081	0-111	0.704	0.604	0.859	0.783	0.427	Fe ³+
fe ²⁺	0.455	0.116	0.122	0.255	0.294	0.188	0.136	0.187	0.160	0.011	0.047	0.372	Fe^{2+}
(In	0.024	0.004	0.007	0.007	600.0	0.004	200-0	0.014	0.005	600.0	0.014	0.014	\mathbf{Mn}
Иg	0.130	0.680	0.626	0.500	0.540	0.663	0.706	0.047	0.037	660.0	0.061	0.193	Mg
ja Ja	0.809	0.875	0.850	0.867	0.851	0.867	0.964	0.247	0.277	0.109	960-0	0.552	Ca
Na	0.278	0.081	0.111	0.165	0.175	0.130	0.091	0.569	0.713	0.881	0.802	0.429	N_{a}
м	0.024	0.013	600.0	0.018	0.023	0.017	0.009	0.093	0.023	0.046	0.075	0.042	K
	1.110	0.969	0.970	1.050	1.049	1-014	1.064	606.0	1.013	1.036	0.973	1.023	X
Ł	0.898	1.004	0.998	0.974	166.0	1.020	0.960	1.048	0.978	1.004	1.016	1.006	А
	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2
×	[1.677	1.684	1.699	1.688	1.696	1.686	1.759	1.763	1.765	1.768	1.736	8
2	1	1.704	1.720	1.740	1.737	1.731	1.714	I	ļ	1		1.760	2
A:[001]	j	$16\frac{1}{2}^{\circ}$	16°	20°	19°	14ځ°	15°	33°	33°		l	26°	A: [001]
S.G.	3.46	3.31	3.33	3-35	3-35	3.35	3.37	3.50	3.49	3.50	3.48	3.47	S.G.
N32, U	Jrtite with	wollastonite			N52, Medit	im-grained ij	olite.		N102	, Cancrinite s	syenite.		
N35C,	Medium-gr	ained pyrox	tenite.		N62, Melte	igite.			N103	C, Coarse-gra	vined alkali s	yenite.	
N30F, N48, N	1jouve. Iedium-gra	ined ijolite.			N95, Coars N95, Canci	e-grained 1joi cinite syenite.	nte.		N103	.F., Fine-gram A, Nepheline	ied alkalı sye syenite.	nute.	
		* Alee T	This 0.010 Th	01010 (1/18-	-14 +	- mity 0 000	- FO O V	+	A last maint of o		100 0 mitu		
		I OSTV	11 0.019' F	e**(IV) U-U16.	. T AB	6 TJ 0-029,	Fe*+(IV) U-UL.	г. 1 ғ	ALSO 11-104	§ A	10-0 ATL 081		

	51-80 SiO ₂	COT 20.7	1:91 A1203	1-92 Fe2U3	4.28 FeO	0.18 MnO	13-84 MgO	23-46 CaO	0.96 Na ₂ O	$0.07 K_2 O$	$0.06 P_2O_5$	none H ₂ O+	100-16 Total	1-916 Si	0-084 Aliv	- Alvi	0-047 Ti	0.053 Fe ³⁺	0.133 Fe ²⁺	0.007 Mn	0.763 Mg	0.929 Ca	0-067 Na	0.004 K	1.000 X	1.003 1	2:000 Z	1.660 x	1.710 2	10° A: [001	3-32 S.G.			t Also Ti ^{iv} 0-007.
	50-61 0 20	00-N	2·14	4-12	5.46	0.07	12.94	22.06	1.58	0.28	0.01	ł	99-95	1.898^{+}	0.095	ł	0.013	0.117	0.171	0.002	0.724	0.886	0.113	0.013	1.012	1.027	2.000	1.681	1.720	17°	3.36	ined ijolite. ined ijolite.	-an	v) 0-015.
	50·20	18-0	none	5.66	7.66	0.26	9.85	22.09	1.84	1.44	0.02		66-66	1-957*	none	I	I	0.146	0.246	600.0	0.561	0.869	0.138	0.069	1.076	0.962	2.000	1.687	1.730	$16\frac{1}{2}^{\circ}$	3.41	Aedium-grai Medium-grai	cursiuse rjo Purjaite.	0.028, Fe ³⁺ (i
	51.00	97-0	2-39	4.95	6.49	0.19	9.94	21.87	2.12	0.21	0.05	[96-66	1.927	0.073	0-031	0.020	0.141	0.204	0.007	0.560	0.885	0.154	600-0	1.048	0.963	2.000	1.693	1.733	18_{2}°	3.38	N155, N N159, N	N170, 7	Also 'Ti'v 0
	49.60	16-0	2.36	2.80	8.71	0.26	10.29	21.88	2.11	0.49	none	I	99.35	1.901	0.099	0-007	0.025	0.083	0.278	0.007	0.587	0.899	0.156	0.023	1.078	286.0	2.000	1.687	1.730	19_{2}°	3· 1 0)6 . *
	50.20	0-97	68.0	18.60	8.58	0.42	2.88	10.42	6.61	0.00	0.08	none	100.55	1.952**	0.042		0.022	0.542	0.278	0.014	0.166	0.428	0.500	0.051	6.079	1.022	2.000	1.738		Į	3.45	d ijolite.		lso Ti ^{iv} 0-00
	50.50	0.95	2.80	15.48	7.02	0.15	4.22	12.36	6.71	0.33	0.05	0.03	100-60	1.928	0.072	0-052	0.028	0.445	0.225	0.005	0.241	0.507	0.495	0.018	1.020	0.996	2.000	1.733	1.780	1	!	ained ijolite arse-graine	yemte. ned ijolite.	A **
	51.00	0.85	0.73	4.50	6.71	0.20	11.72	22.26	1.66	0-80	none	I	99-83	1-931	0.032			0.115	0.212	0.006	0.662	0.903	0.123	0.009	1.035	0.995	2.000	1.698	1.723	174°	3.41	t, Coarse-gra	Nepheline s Coarse-graii	+(iv) 0.012.
	51.39	0.80	1.86	4.34	6.98	0.18	10-95	22.45	1 +	80-0	0-03	1	100.50	1.928	0.079	0.000	0.023	0-121	0.219	0.007	0.613	0.902	0.104	0.005	1.001	0.992	2.000	1.687	1.737	16°	3.33	N120R N120V	N121, N152,	^v 0·025, Fe ³
OTTN	07.16	0.43	2.12	7.05	9.43	0.28	5.87	18.59	8-8- 18-8-	1.03	none	l	18-69	1-980	0.090	0-020	0-012	0-203	0.302	0-00	0.337	0.764	0.245	0.051	1.060	0.940	2.000	1.713	1.760	20°	3-47		unite.	¶ Also Ti ⁱ
TTT	50-59	().63	2.04	8.02	9.42	0.36	6-45	17-85	3.40	04.0	1 - 12		86-66	1.040	0.051	610.0	310-0 810-0	0.231	0-303	0.012	0.370	0.736	0.254	0.060	1.050	0.976	2.000	1-715	1.766	164°	3.44	te ijolite.	e with mela	0.110.
1114	47.99	1.68	none	9.39	5.37	0-19	12.00	91.19	59.1	0.40	0-91	none	100.13	1.8491	1 210 1	anni	[]	0.162	0.173	0-007	0.687	0.872	0.120	0.023	1.015	1.029	2.000	1.680	1.709	16°	3.35	vined melani h melanite.	rained ijolit ned ijolite. jolite.	18. Fe ^{3+(iv)}
ONTH	51.40	0.30	2.10	3.63	8.92	0.94	0.03	90.40	01.1	07.1	61.1	ļ	98-73	1,072	200 U	020.0	000.0	600-0	0.286	0.007	0-517	0.842	0-111	0.055	1.008	0.995	2.000	1.696	1.734	174°	3.36	Coarse-gra Ijolite wit	Medium-g Fine-grair Variable i	so Ti ^{iv} 0·04
	SiO_2	Tio.	Al _* O _*	Fe.O.	TeO.	MnO	MeO			1020		H,0+	Total		A Tiv	AL:	Ĩ	т. Празн	Fro2+	Mu -	Ma	۳ د ا	Na.	K	X	A		5	5	A · [001]	S.G.	N108, N114,	71117 N118 N119 0	II AI

TABLE I (cont.)

VISO 11. 0.000

|| Also Ti^{iv} 0.048, Fe³⁺(iv) 0.110.

	^{,3+(} iv) 0-007.	^{iv} 0.059, Fe	¶ Also Ti	0-052.)-022.	0, Fe ^{s+} (iv) 3, Fe ^{s+} (iv) (6 Tl ^{iv} 0-020 0 Tl ^{iv} 0-028	II Als † Als	iv 0-023. iv 0-001.	§ Also Ti* Also Ti	054.	, Fe ³⁺ (iv) 0	to Thiv 0-023 to Thiv 0-001.	‡ Als ** Als	
	autre.	ute caroun	утохепе ара	T021, F		te.	9, Pyroxem 4, Urtite.	N52 N58				ained ijolite. gmatite.	, Coarse-gr , Ijolite-pe	N514 N516
	atite.	ring carbon	arbonate-sye yroxene-bea	To14, C			8, Ijolite. 0, Ijolite.	N52 N52	-St	e and wolls	ith melanit	ned ijolite w	, Fine-grai lite.	N511 ton
		;	enite.	$To1, Sy_{0}$			7A, Ijolite.	N51			e.	rrained ijolit	. Medium-	N178
s.c.	3.40	3.49	3.49	3-47	3.40	3.38	3.32	3.38	3.49	3.42	3·38	3.43	3:45	S.G.
A: [001	25°	31°	29°	26°	1	16°	$13\frac{1}{2}^{\circ}$	16°	20°	l	19°	20°	1 19%	A · [001
~	1				1	1.729	117-1	1.729	1.763	1.769	1.733	1.760	1.743	5 5
α	1.743	1.758	1.756	1.740	l	1.689	1.692	1.689	1.704	1.718	1.699	1.705	1-710	38
7	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	1 10
2	0.991	$066 \cdot 0$	1.060	0.954	0.893	1.017	0.960	0.983	0.978	0.942	0.995	1.003	0.976	414
X	666·0	0.984	1.926	1.069	1.084	1.024	1.114	1.061	1.024	1.073	1.042	1-024	1-035	4 14
K	0.033	0.019	0.042	0.061	0.014	0.018	0.013	0.031	0.084	0.037	0-050	0.046	0.024	e M
N_{a}	0.534	0.665	0.602	0.453	0.260	0.131	0-141	0.150	0.167	0.242	0-164	0.185	0.000	n N N
Ca	0.432	0.300	0.282	0.555	0.810	0.875	096-0	0.880	0.773	0.202	0.928	0.708	0.000	8 2 8
Mg	0.163	0.137	0.093	0.208	0.228	0.609	0.732	0.655	0.449	*10.0	0.695	0-011	900-0 0.998	N N
л Мл	017.0	200-0	102-0	0-000	0-327	0.230	0.168	0.191	0.383	0.369	0.269	0.306	0.370	Fe^{2+}
ге ²⁺	0.000	207-0	0-723	0.501	0.246	0-063	0.053	0.119	0.140	0.233	0.137	0.184	0.165	Fe^{3+}
TI T	0	0.032	0.036		0.037	0.032		0.011		0.016	0.018	0.038	0.024	E
Alvi	I		I	!	0.036	0.076	1			0.012	0.037	0.010	0.022	Alvi
Aliv	1	0.035	0-065	0.056	0.038	0.141	1	0-050	600.0	0.039	0.073	0.115	0-020	Aliv
Si	1-950†	1.964*	1.934**	1.894	1.962	1.859	1.928[1.9278	1.9141	1.961	1-927	1.885	1.980	7
Total	99-23	99-47	100.18	100.61	100.15	100.74	99-95	99·88	100.44	100.61	100.46	100-55	99.25	Total
+ 0° =]		l	Ì		1	1	1	1	1]		$H_{2}O +$
г ₂ 0° 1	trace	10.0	trace	0.53	none	0.04	0.04	0.12	none	0.14	0.11	0.17	0.44	$P_{s}O_{s}$
D°.4	0.64	0.40	0.86	1.25	0.25	1.05	0.28	0.69	1.70	67.0	1.02	0.91	0.43	K.0
Na ₂ O	00-2	8.89	8.01	5.94	3.50	1.80	1.91	2.04	2.21	3.21	2.25	2.53	2.79	Na.0
c_{a0}	10.26	7.23	6.11	13.91	19.59	21.74	23.72	21.83	19.24	19.26	20.52	19-48	19.60	CaO
MgO	2.79	2-37	1.62	3.56	3.97	10.90	13.00	11.57	7.72	5.17	9-29	7-93	6.63	MeO
MnO	0.37	0.22	0.22	0.31	0.60	0.19	0.20	0.19	0.39	0.45	0.29	0.36	0.27	MnO
FeO.	6.41	3-33	6.04	6.79	10.14	7.35	5.30	6.07	11.84	11.34	8.49	09-6	11-25	FeO 3
Fe ₂ O ₃	21.20	24.32	24.76	17.03	8.40	2.16	3.75	4.17	6-75	90·6	4.73	6.39	5.52	Fe.O.
$\overline{AI_{s}0_{s}}$	none	27.0	1.46	1.23	1.62	4.94	none	1.16	0.20	1.13	2.37	2.77	0-96	ALO.
TiO.	0.95	1.13	1.31	1.89	1.29	1.07	0.72	1.23	62-0	0.57	10.00	±9.00	06.00	
SiO_{4}	49.61	50.80	49-79	48.18	50.79	49.50	51.03	50-81	40.60	50.40	10.03	00.01	10101	0.0
	To21	To17	To14	Tol	N584	N529	N520	N518	N517A	N516	N514	N511	N178	

	si0 ₃	TiO ₂	$\underline{A}1_2 \underline{O}_3$	Fe ₂ O ₃	FeO	MnO	MgO	CaO	$Na_{s}O$	K.O	$P_{s}O_{s}$	$H_{2}O_{+}$	Total	is	Aliv	Alvi	Ŀ	Fe ³⁺	\mathbf{Fe}^{2+}	Mn	Mg	Ca	Na	K	X	Y	Z	8	~	Á: [001]	S.G.			
SB945	50-93	0.45	1.64	13.54	6.49	0.94	5.65	14.37	6.08	0.21	0.16	[100-46	1.951	0.049	0-025	0.014	0.392	0.207	0.030	0.322	0.583	0.451	600.0	1.043	0.976	2.000	1.735	1.767	24°	3.45			
SB190	50-99	0.52	1-97	10-69	69.9	0.59	6.69	15.94	5.19	0.40	0.08	1	99-75	1.954	0-046	0.042	0-016	0.309	0.214	0.018	0.382	0.647	0.387	0.018	1.052	0.981	2.000	1.720	1.748	22°	3.45	venite.		
SB78	49-20	0.49	2.38	8-24	11.99	2.07	5.16	16.78	3.36	0.87	0.16	I	100.70	1.921	0-079	0.029	0.014	0.244	0.392	0.068	0.300	0.694	0.253	0.042	0.939	1.047	2.000	1.728	1.770	24 } °	3.47	onatite. snite. Vepheline sy		
SB	51.20	0.27	1.51	6.10	10.8	1.03	8.66	19.54	2.69	0.78	0.17	!	96-66	1.962	0.038	0.031	200.0	0.175	0.256	0.035	0.495	0.794	0.198	0.037	1.029	666.0	2.000	1.716	1.750	ļ	3.43	SB, Carb SB78, Fe SB190, 1 SB190, 1	r (01700	
B_{0330}	50-90	0-88	1.00	4.66	8-05	0-27	10.51	21.66	1.23	0.45	0.14	1	99-72	1.938**	0.046		600.0	0.133	0.256	600.0	0.597	0.877	0.092	0.023	0.992	1.004	2.000	1.688	1.730	164°	3.37			
B_0270	52.00	0.47	1.85	97-6	4.22	0-23	9.74	16.02	3·82	1.37	0.22	ł	02-66	1.963	0.037	0.045	0.014	0.277	0.134	0.007	0.549	0.642	0.281	0.068	166.0	1.026	2.000	1.711	1.741	18°	3.40	enite.		
K440	48.93	0.57	0.03	87.8	96-ZT	19-0	12.9	19.71	2.90	0.23	0.15	none	101-59	1.9461	1	1	1	0.221	0.424	0.021	0.351	0.773	0.221	600·0	1.003	1.017	2.000	1.710	1.760	23°	3.38	te. llieren in sy borolanite.		
K353	50-31	0.37	none	0.	0.47	12.0	11.22	22-47	1.86	0.42	trace	[100-06	1-919	1	I	I	0.123	0-206	0.007	0.637	0.920	0.138	0.018	1.075	0.973	2.000	1.682	1.729	16°	3.33	Joarse syeni yroxene sch Xenolith in Cromaltite		
K320	52.15	0.76	11.0	16.07	01-Z	47.0	0.84	3.09	10.75	0.22	0.27	1	99-55	2.013	0.005	I	0.023	0.839	0.070	200.0	0.049	0.129	0.803	600.0	0.940	0.993	2.013	1.765	l	33] °	3.46	K353, (K440 P Bo270, Bo230	60000	
K62	49.10	0.37	0-20	80.7	TR-TT	10.0	21.7	20.25	2.41	0.48	0.22	попе	100-46	1.926	600.0	1	[0.178	0.391	0.016	0.417	0.834	0.183	0.023	1.040	1.002	2.000	1.707	1.753	1830	3.36			
T0585	50-05	0-79	2:01	70.#T	90.0	00.0	2.94	13.80	5.25	0.18	0.06		99-59	1.943	0.057	0.060	0.023	0.420	0.294	0.021	0.170	0.574	0.397	200-0	0-980	0.988	2-000	1.720	1.770]	3.48	carbonatite		ļ
T_{0572}	50-86	0.71	0.39 16.70	71.01	#T. /	64-D	4.92	11-96	6.32	0.98	0.17		100.50	1.962	0.038	0.004	0.021	0.487	0.229	0.014	0.248	0.487	0.473	0.046	1-006	1.003	2-000	1.736	[24°	3.46	uite. bonatite. xenolith in	enite.	
T_{043}	51.43	0-37	12-0	6.45	66.0	40 D	60.7 1	10-43	6.40	0.65	trace	1	100-39	1.988	600-0		600·0	0.604	0.211	0.030	0.150	0.432	0.478	0-033	0.943	1-004	z-000	1.741	I j	27°	3.40	Syenitic fer Coarse car Pyroxene venite pegr	Banded sye	
	SiO ₃	out	H ₂ O ₃	FeO.	MnO	N ₂ O		Cau	Na ₂ O	K20	$\mathbf{P_{20_{5}}}$	$H_{2}0+$	Total	Si	Aliv	Alvi	Ti	Fe ³⁺	Fe²+	Mn	Mg	ca Ca	Pa	×:	¥	× 1	N	8	~	A: [001]	S.G.	To43, To572, To585, K62, S	K320,	

TABLE I (cont.)

§ Also Ti^{iv} 0.012, Fe³⁺(iv) 0.053. ‡ Also Ti^{iv} 0.003.

|| Also Ti'v 0.011, Fe³⁺(iv) 0.070.

** Also Tiiv 0.016.

T Also Ti'v 0.016, Fe³⁺(iv) 0.038.



FIG. 2. *a* (top). Plot of $(Mg + Fe^{2+} + Mn)/Ca$ for all the newly analysed pyroxenes. *b* (bottom). Plot of $(Na + K)/(Fe^{3+} + Ti + Al^{iv})$ for all the newly analysed pyroxenes. \blacktriangle Budeda; \odot Bukusu; \times Napak; \spadesuit Tororo; \bigcirc Semarule; \triangle Sørøy.

dependence on the presence or absence in the rock of other titanium bearing minerals, notably melanite garnet. Titanium is greatest in the pyroxenes from rocks where such minerals are absent (see fig. 4). Manganese increases with increase in the hedenbergite molecule.



FIG. 3. Pyroxene analyses plotted with diopside + hedenbergite + acmite recalculated to 100 %. Symbols as for fig. 2.

In general, however, titanium, manganese, and also aluminium, vary more in pyroxenes from different regions and localities than among pyroxenes from the same locality; this is shown in table II.

Physical and optical properties of the pyroxenes. The diopsidic pyroxenes are almost colourless; a green colour becomes apparent with an increase in the hedenbergite molecule but becomes particularly noticeable as the content of acmite rises. The variation is readily apparent in zoned crystals. There is some evidence, however, that intensity of colour does not depend solely on the content of the major constituents. It is possible that minor elements have a significant effect not only on the intensity of the colour but also on the shade; for example the pyroxenes from the African complexes are yellowish green in colour, for which the relatively high titanium may be responsible. Other comparable pyroxenes, e.g. Loch Borolan and Sørøy, are a much brighter green.



FIG. 4. Content of TiO_2 in the analysed pyroxenes from Napak. Open circles = rocks with melanite, black circles = rocks without melanite.

More difficult to explain is the occurrence of a yellowish-brown variety among the most sodic pyroxenes. Intensity of pleochroism increases with acmite content, but it remains weak in the rather strongly coloured pyroxenes of intermediate composition.

Specific gravities of the pyroxenes increase with the Fe^{2+}/Mg ratio and with the replacement of CaMg by NaFe³⁺. The same chemical factors are responsible for increases in refractive indices and A : [001] angles (see King, 1962).

Physical and optical properties may be used to determine the approximate chemical compositions of pyroxenes in the diopside-hedenbergiteTABLE II. Analyses recalculated as molecules % of diopside, hedenbergite, acmite, and CaSiO₃ or MgSiO₃. P₂O₅ calculated out first as apatite taking CaO required from the total. The table also includes actual weights % of MnO, TiO₃, and Al₂O₃ to show variation in these constituents between complexes.

	(MnO	0.20	0.19	0.26	0.07	0.18	0.27	0.36	0.29	0.45	0.39	0.19	0.20	0.19	0.60	0.31	0.22	0.22	0.37	7 6·0	0.43	0.66	0.51	0.24	0.21	0.61	0.23	0.27	1.03	2.07	0.59	0.94	
	eight %	Al ₂ O ₃	2.36	2.39	none	2·14	1.91	0.96	2.77	2.37	1.13	0.20	1.16	none	4.94	1.62	1.23	1.46	0.77	none	0.21	0.89	2.51	0.20	0.11	none	0.03	1.85	1.00	1.51	2.38	1.97	1.64	
	3	Tio.	16.0	0.75	0.97	0.68	1.68	0.80	1.33	0.58	0.57	0.79	1.23	0.72	1.07	1.29	1.88	1.31	1.13	0.95	0.37	0.71	6.79	0.37	0.76	0.37	0.57	0.47	0.88	0.27	0.49	0.52	0.45	
	(laSiO ₃	1.91	10-60	7-78	2.74	2.43	5.33	2.68	2.60	10.94	8.53*	2.78	4.33	2.13	24-69	13-08	1.97*	4-96	4.49	4.61	0.23	8·81	2.67	1.24	8.67	1.10*	0.44*	1.83	1.76	none	none	2.83	
%		cmite (17-36	15.57	15-47	13.05	7.51	22·21	22·31	20.76	26.16	21.53	16.30	12.78	17-44	14-42	46-74	67-92	69-50	56-75	53-37	$51 \cdot 12$	41.41	19.54	86·80	14.46	21.53	33-32	11.03	22-96	30.21	38·30	43.99	
Molecular	enhero-	ite A	26-20	20-12	23-85	16·21	L3-68	35-54	80-79	26-40	35-17	39-21	18.87	5.88	22-56	35-84	10.12	06-12	11-58	22-22	25-23	23-96	32·15	38-20	8.19	18-99	H-53	13-59	26-65	27-58	45·81	25-32	22-42	
	ion- Hed	ide	4-41 2	3.65 2	2.86 2	1-96 J	6.32	6-86	4·16 2	D-20 2	2.66	0-68	I -99	E 00-2	2 98·1	4-20 S	9-08 2	8.13	3-94 1	6-55 2	6-74 2	4-65 2	7-62	9-54	3·72	7.83	5.82 4	3.05 1	0.43 2	2-67	3-94 4	6-34 2	5 69-0	
	Sneetmen D	number	N.152 5	N.155 5:	N.159 55	N.163 67	N.170 7(N.178 3(N.511 44	N.514 5(N.516 2'	N.517A 30	N.518 6	N.520 67	N.529 57	N.584 24	To.1 19	To.14 8	To.17 13	To.21 16	To.43 10	To.572 2.	To.585 17	K.62 3:	K.320	K.353 57	K.440 31	Bo.270 51	Bo.330 6	SB 4	SB.78 2	SB.190 3	SB.245 30	
		0 U	0.34	0.40	0-42	0.42	0-48	0.48	0.32	0-53	0-64	0-28	0.52	0.32	112-0	0-11	0.19	0.22	0-27	0.13	0·18	0-44	0.16	0-29	0.45	0.46	0.24	0.19	0.36	0·28	0.18	0.20	0.15	0.49
	eight %	Al ₂ O ₃ 1	none	0.27	0.18	0.20	2.43	1.36	2-39	none	none	none	1.28	2.06	1.46	4-09	3-93	2.21	0.82	4-03	1.09	1.53	4·14	1.51	2.39	none	2-10	none	2.04	2.12	1.86	0.73	2.80	0.80
i	M	Tio ₂	1.23	0.78	0.40	1.00	0.51	0.67	1.16	2.51	1.05	0.51	0.39	0.87	1.10	1.36	2.35	0.75	0.60	0.86	1.01	1-44	1.08	2.49	1.99	0.59	0.30	1.68	0.63	0.43	0.80	0.85	0.95	0.07
	(CaSiO ₃	0.46	5.79	2.47	4·81	1.97	10.75	1.40*	2.70	7.98	3.82	14.38	5.72	20.74	7.16	9.40	10.60	0.22	11.15	10.21	none	7.62	1.11*	2.45*	2.69*	2.97	9.03	4.83	9.36	5.80	1.71	3.78	0.7.4
ur %		Acmite	36-56	22.26	23.30	20.56	31.12	30-35	47.19	29.99	42.55	12.74	31.25	33-85	15.31	0.70	12.38	17.30	18.18	18.52	11.26	72-00	72.06	88.49	84.05	44-49	16.43	13.02	29.87	27.86	10.70	14.98	47-75	54.50
Ë,	13	e e				~	4	5	33	Η·	·14	-98	£1.€	2-93)·12	2-46	3·30	24·66	28·34	18.10	13-13	22-04	16.40	1.99	26-2	36.82	28-98	6.17	28-63	69-08	22-31	20-54	23·63	00.07
Molec	adenhero	ite	36-32	33.63	$36 \cdot 30$	34.35	38 [.]]	80 80	34	36	32	23	õ	õ	ŭ	H	-											1	G	60	6.4			
Moleci	Dion. Hadenherd	side ite	26.61 36.32	38-29 33-63	37-87 36-30	40.23 34.33	28·72 38·1	30-77 28-0	17·04 34·	31·11 36	17-28 32	59-43 23	25-20 2(27-44 35	13-82 5(70-64 1:	64-91 1	47-86	53.22	52.20	$65 \cdot 44$	5.13	3-93	8-41	5-57	15.96	51-58	61-70 1	35-36 2	31-99 8	61-15 2	62.70	24.81	80.7L

* MgSiO₃, not CaSiO₃, as in all other instances.

acmite series. Since, however, the pattern of variation in all properties is similar, and depends on the same chemical factors, the composition obtained will be only approximate (see figs. 5 and 6).



Fig. 5. Diagram showing the variation in specific gravity and the α refractive index of the newly analysed pyroxenes. The positions on diopside-acmite and diopside-hedenbergite are taken from data in Deer, Howie, and Zussman, vol. 2 (1963).

Relation between composition of pyroxene and composition of rock. Petrographic examination shows that the pyroxene of the melanephelinites is a nearly colourless or pale green diopsidic variety. The pyroxene is greener in the more leucocratic rocks, and it is common to find the phenocrysts mantled by narrow zones very rich in acmite. In the phonolites and trachytes, however, a deep green acmitic pyroxene is usual, and the mineral is further characterized by a strongly prismatic or even acicular habit. Occasionally, paler diopsidic cores are present. It is to be inferred that temperature of crystallization (as well as concentration of sodium and iron) is a major factor in determining the composition of the pyroxene, the more acmitic varieties crystallizing at lower temperatures. The sharp compositional difference between diopsidic phenocrysts and their acmitic mantles and ground-mass crystals suggests that these correspond to crystallization under pre-extrusive



FIG. 6. Diagram showing the variation in the angle A: [001] for the newly analysed pyroxenes. Determinations by B. Collins, B.Sc., Bedford College.

and post-extrusive conditions respectively. Experimental work by Bailey (1963) has shown that the upper temperature-stability limit of acmite increases rapidly as pressures fall to normal. It may be noted that the mineralogy of the nephelinites is very largely represented by the phases in the system acmite-nepheline-diopside investigated by Yagi (1963) and the system albite-nepheline-diopside-acmite-water (Nolan, 1966). The appearance of olivine and melilite as primary crystalline phases over certain compositional ranges at one atmosphere pressure is significant since these are also common minerals in the volcanic rocks.

The pyroxenes of the ijolite series have been studied more fully, and

here a continuous range from near diopside to about $\text{Di}_{15}\text{He}_{55}\text{Ac}_{30}$ is shown. Fig. 7 shows that a rather close correlation exists between the composition of the pyroxene and the sequence pyroxenite-melteigiteijolite-urtite. Field and petrographic studies of the ijolite series shows



FIG. 7. Diagram showing the variation of the compositions of the newly analysed pyroxenes according to rock type. F = fenite; P = pyroxenite; T = turjaite. Symbols as for fig. 2.

a highly complex sequence of events, which is only very simply represented by the series from pyroxenite to urtite. Critical chemical characteristics of the sequence are expressed by plotting the ratio of total iron oxide to magnesia against that of lime to magnesia as shown in fig. 8 (see also King, 1965). Generally, the hedenbergite-acmite content of the pyroxenes increases in successively later members of the sequence.

Where zoning in the pyroxenes of the ijolite series is shown, it is of two kinds. One is a simple increase in acmite content towards the margins of the crystals corresponding to successive stages of growth.



FIG. 8. Plot of log $(FeO+Fe_2O_3)/MgO$ against log CaO/MgO showing that the hedenbergite + acmite content of the pyroxenes increases in successively later members of the Tjolite Series. Symbols in the acmite-diopside-hedenbergite

The second is more patchily developed, and although the acmite content is usually higher towards intergranular boundaries, it is evidently a zoning of metasomatic type, for it is often prominently developed in grains that are adjacent to later veins formed by members of the ijolite sequence (see Napak 120 Rock and Napak 120 Vein).

The compositional range of pyroxenes is extended continuously almost to pure acmite in the nepheline and cancrinite syenites and carbonatites.

The pyroxene series corresponds primarily to a gradient of temperature of formation, the continuity of the series reflecting much greater uniformity of pressure under plutonic conditions than the sudden variations that can be inferred for the volcanic rocks.

The compositions of the pyroxenes in the fenites associated with the alkaline complexes show very wide variations from near diopside to near acmite. Only limited chemical data are yet available, but field and petrographic observations suggest that, although temperature of formation is an important factor, the concentration of sodium and iron may here be of equal importance.

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