MINERAL CHEMISTRY OF CLINOPYROXENE AND FELDSPARS IN THE NEOPROTEROZOIC ALKALINE DYKES OF THE BANGALORE DISTRICT, KARNATAKA, INDIA

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The unmetamorphosed Neoproterozoic dykes of the continental intraplate alkaline magmatism studied in Southern Karnataka consist of analcime-tinguaites, tinguaites, bostonites and solvsbergites. The analcime-tinguaites, tinguaites and bostonites display pyroxene trends typical of alkaline rocks; the cores of the phenocrysts containing more diopside whereas the rims contain more hedenbergite and aegirine and the groundmass clinopyroxenes have the highest aegirine contents.

The pyroxene trends of the rocks are similar, indicating that the melts they formed from had similar fugacities of oxygen and/or similar peralkalinities. The absence of reverse zoning indicates that the melts were not intermixed.

The K-feldspar is either K-rich and Na-poor sanidine or orthoclase and is zoned with respect to sodium and sometimes also barium. The plagioclase is almost pure albite.

Key words: dykes, tinguaite, bostonite, solvsbergite, clinopyroxene, K-feldspar, electron probe data, Proterozoic, Neoproterozoic, Karnataka, India.

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INTRODUCTION

Although alkaline rocks account for less than 1% of all igneous rocks, they have been described in all continental areas, where they may be classified into two tectonic settings: 1) continental rift valley magmatism, which has produced most of alkaline rocks and 2) continental intraplate magmatism, which is the second important producer of alkaline rocks but its activity and siting are not subject to obvious tectonic control (Fitton and Upton 1987).

The stabilization of the Karnataka craton took



Fig. 1. Locations of the Karnataka alkaline dykes studied, their names and rock contents.

place between the intrusions of the late Archaean Closepet granite and the unmetamorphosed Archaean/Palaeoproterozoic dolerites, the ages of which are 2500 Ma (Allen et al. 1986) and $2420 \pm$ 246 Ma (Ikramuddin and Stueber 1976), respectively. The completely unmetamorphosed alkaline rocks, which will be described in this paper, intruded some 1500 Ma after this stabilization. as revealed by their Rb-Sr ages of 832 ±40 Ma (Ikramuddin and Stueber 1976). Since these unmetamorphosed alkaline rocks are not associated with continental rifting, they fall into the category of continental intraplate magmatism. Although there are more than 200 dykes in the area, they have been relatively little studied (see references in Ikramuddin 1974) which may be partly due to their modest sizes: they are usually less than 3 m wide and 15 m to 8 km long. Ikramuddin (op. cit.) studied the whole rock chemistry and Rb-Sr-isotopes of these rocks in the Bidadi-Harohalli area and verified their extreme richness in potash (up to 14.59 % K₂O), their high initial 87Sr/86Sr ratio, high K, Rb and Ba and low Mg, Ni and Cr. He concluded that they were not derived from carbonatitic or nephelinitic magma but may have been formed by partial melting of a phlogopite-rich assemblage or by a reaction between the magma and mantle rocks.

This paper, which is based on the laboratory work of Makkonen (1991), concentrates on the chemistry of the main minerals of the alkaline dykes, that occur some 50 km southwest of the Bidadi-Harohalli area. A summary of the geochemical and petrological study of these intrusions is given by Devaraju et al. (1989).

Sampling

For this work 12 unweathered samples were studied. They were collected from the dykes, the distributions of which are shown in Fig. 1. The total amount of dykes is 10, including Chellipura and Arekattedoddi, which are composite structures (Table 1).

Analytical methods

The microprobe analyses were performed at the

Department of Electron Optics, University of Oulu, using a JEOL JCXA 733 microanalyser. The Kfeldspars in samples DJK-39A and DJK-25 were analysed at the University of Bonn, Germany.

PETROGRAPHY

The rocks studied are analcime-tinguaites, tinguaites, bostonites and solvsbergites (Fig. 1., Table 1.). The names recommended by IUGS (Le Maitre 1989) for these rocks would be phonolite, alkali feldspar syenite and trachyte, respectively, but the traditional names are used in order to convey information on the special textures and mineralogies of the dykes. Chemically these rocks are phonolites and trachytes (Devaraju et al. 1989), while they range in texture from porphyritic to equigranular, being most commonly porphyritic. The main minerals are clinopyroxene, K-feldspar, plagioclase and analcime (Table 1). The phenocrysts of the porphyritic rocks consist of clinopyroxene and Kfeldspar, while the groundmass minerals are clinopyroxene, K-feldspar, plagioclase and analcime. The last of which, the only feldspathoid found in the samples, is described in another paper (Devaraju et al., in prep.).

The groundmasses of the porphyritic rocks are fine-grained and hypidiomorphic-granular and occasionally display a flow structure. Their phenocrysts range between 0.3 and 8 mm in size. Equigranular specimens are medium-grained (1-2 mm).

The analcime-tinguaites and tinguaites display a tinguaitic texture, while a bostonitic texture is typical of the bostonites. The bostonites and solvsbergites are markedly altered. The accessory minerals include titanite, calcite, haematite, ilmenite, pyrrhotite, biotite, epidote, magnetite, rutile, apatite and zircon.

PYROXENES

Occurrence

Subhedral, zoned clinopyroxene phenocrysts with pale brown and occasionally also greenish cores

ROCK	DYKES	MINERALS	
		Phenocrysts	Groundmass
Analcime- tinguaite	Chellipura Madarhalli	K-feldspar Clinopyroxene	K-feldspar Clinopyroxene Plagioclase Analcite
			Accessories: sphene haematite ilmenite pyrrhotite calcite zircon
Tinguaite	Kodihalli Ankachari Doddi Konasandra Ajahunse Arekattedoddi Hosadoddi	K-feldspar Clinopyroxene	K-feldspar Clinopyroxene Plagioclase Accessories: sphene biotite epidote
			calcite zircon magnetite rutile apatite
Bostonite	Chellipura Arekattedoddi	K-feldspar Clinopyroxene	K-feldspar Plagioclase Clinopyroxene
			Accessories: ilmenite haematite epidote carbonate rutile sphene apatite
Solvsbergite	Devasandra Dodsadenhalli	K-feldspar Clinopyroxene (altered)	K-feldspar Clinopyroxene Accessories: Plagioclase magnetite rutile epidote

Table 1. The rocks of the Karnataka alkaline dykes studied and their mineral contents.

surrounded by pleochroic green rims (Fig. 2) are common in the analcime-tinguaites, tinguaites and bostonites, whereas the pyroxene phenocrysts in the solvsbergites are totally altered. Sometimes there is also a distinct mantle zone between the core and the rim. The zoning of the pyroxene is normal or oscillatory, and none of the samples studied displays reverse zoning. The boundaries between the zones are sometimes sharp, but gradational zoning is more common. The cores of the clinopyroxene phenocrysts are sometimes euhedral. The optical orientation (c Λ X) varies from -33 degrees in the extreme diopsidic cores to +4 degrees in the rims that are richest in aegirine.

The groundmass clinopyroxenes are green, anhedral or subhedral needles and laths or, in the tinguaites, occur as a fine-grained mass. The grains having the highest aegirine content display an optical orientation (c Λ X) of +5 degrees.

Mineral chemistry

Analcime-tinguaites. In terms of diopside, hedenbergite and aegirine end members, the compositional variation in the cores of the pyroxene phenocrysts of the analcime-tinguaites is $Di_{82.7}Hd_{15.0}Ae_{2.3} - Di_{39.5}Hd_{43.6}Ae_{16.9}$ (Table 2). That of the extreme aegirine-bearing rims is from $Di_{0.8}Hd_{26.0}Ae_{73.2}$ (Chellipura dyke) to $Di_{0.2}Hd_{10.6}Ae_{89.2}$ (Madarhalli dyke) and that of the groundmass clinopyroxenes from $Di_{7.0}Hd_{30.2}Ae_{62.8}$ to $Di_{1.2}Hd_{25.0}Ae_{73.8}$.

Tinguaites. The cores of the pyroxenes in the tinguaites (Table 2) of Arekattedoddi and Hosadoddi are $Di_{68.7}Hd_{26.9}Ae_{4.4}$ and $Di_{87.3}Hd_{11.6}Ae_{1.1}$,



Fig. 2. Photomicrograph of zoned clinopyroxene phenocrysts. Single polar. Field of view is about 1 mm x 1.4mm.

Table 2. Results of the analyses.

	1	2	3	4	5	6	7	8	9
wt%									
SiO ₂	49.52	50.21	52.06	50.26	50.12	53.17	50.89	49.10	48.95
TiO	0.79	2.12	0.37	0.32	0.36	0.25	0.40	0.63	0.76
Al Ó,	2.18	1.85	0.73	1.07	0.92	1.04	1.68	2.07	2.00
Fe _O	5.73	20.97	0.84	6.50	10.92	0.00	4.16	7.34	11.01
FeO	12.55	6.47	8.47	11.22	11.13	5.08	10.41	10.85	9.91
MnO	0.95	0.94	0.24	0.81	0.95	0.16	0.51	0.97	1.01
MgO	6.29	0.19	12.48	6.92	4.04	15.21	9.00	6.47	6.39
CaO	19.04	7.85	22.26	18.23	15.65	22.68	20.49	19.11	19.13
Na O	2 32	9.45	0.62	2 77	4 46	0.50	1.72	2.41	2.64
K O	0.02	0.03	0.00	0.00	0.01	0.00	0.00	0.02	0.00
$\mathbf{X}_{2}\mathbf{O}$	0.02	0.12	0.01	0.00	0.12	0.03	0.06	0.08	0.06
$v_{2}O_{3}$	0.00	0.12	0.01	0.09	0.12	0.05	0.00	0.00	0.00
Total	99.45	100.20	98.08	98.19	98.68	98.12	99.32	99.05	101.8
G 1.C	1.1	1		1.4					
Sructural form	nula base	ed on 6 o	oxygens	and 4 c	ations				
Sructural form	nula base 1.930	ed on 6 o	oxygens 1.986	and 4 c	ations 1.977	1.990	1.953	1.919	1.906
Sructural form Si Al	nula base 1.930 0.100	ed on 6 o 1.946 0.084	0.033	and 4 c 1.972 0.049	ations 1.977 0.043	1.990 0.045	1.953 0.076	1.919 0.095	1.906 0.091
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Sructural form Si Al Ti Fe ³⁺	nula base 1.930 0.100 0.023 0.168	ed on 6 d 1.946 0.084 0.062 0.612	1.986 0.033 0.011 0.024	and 4 c 1.972 0.049 0.009 0.192	1.977 0.043 0.011 0.324	1.990 0.045 0.007 0.000	1.953 0.076 0.012 0.120	1.919 0.095 0.019 0.216	1.906 0.091 0.023 0.252
Sructural form Si Al Ti Fe ³⁺ Fe ²⁺	nula base 1.930 0.100 0.023 0.168 0.409	ed on 6 o 1.946 0.084 0.062 0.612 0.210	1.986 0.033 0.011 0.024 0.270	and 4 c 1.972 0.049 0.009 0.192 0.368	1.977 0.043 0.011 0.324 0.367	1.990 0.045 0.007 0.000 0.159	1.953 0.076 0.012 0.120 0.334	1.919 0.095 0.019 0.216 0.355	1.906 0.091 0.023 0.252 0.323
Sructural form Si Al Ti Fe ³⁺ Fe ²⁺ Mn	nula base 1.930 0.100 0.023 0.168 0.409 0.031	ed on 6 o 1.946 0.084 0.062 0.612 0.210 0.031	1.986 0.033 0.011 0.024 0.270 0.008	and 4 c 1.972 0.049 0.009 0.192 0.368 0.026	1.977 0.043 0.011 0.324 0.367 0.032	1.990 0.045 0.007 0.000 0.159 0.005	1.953 0.076 0.012 0.120 0.334 0.017	1.919 0.095 0.019 0.216 0.355 0.033	1.906 0.091 0.023 0.252 0.323 0.033
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9. Chellipura, bostonite, groundmass, (DJK-13)



Fig. 3. Clinopyroxenes of a. analcime-tinguaites, b) tinguaites and c) bostonites plotted (crosses) on Di-Hd-Ae -diagrams and compared with pyroxene trends from other alkaline rocks. Undersaturated: 1) Auvergne (Varet 1969), 2) Lovozero (Bussen and Sakharov 1972), 3) Itapirapua (Gomes et al. 1970), 4) Uganda (Tyler and King 1967), 5) Morotu (Yagi 1966), 6) South Qoroq centre (Stephenson 1972) 10) Ilimaussaq (Larsen 1976). Oversaturated: 7) Alkalibasalts of Japan (Aoki 1964), 8) Pantellerites (Nicholls and Carmichael 1969), 9) Nandewar (Abbott 1969), 11) Skaergaard (not alkaline) (Brown and Vincent 1963).

respectively, while the cores at Kodihalli are hedenbergite. The zoning at Ankachari Doddi, Konasandra and Ajahunse is weaker, and the cores consist of aegirine-augite of compositions varying . between $Di_{34.1}Hd_{41.9}Ae_{24.0}$ and $Di_{6.6}Hd_{31.2}Ae_{62.2}$. The rims of the phenocrysts are aegirine-augite, as at Kodihalli ($Di_{5.7}Hd_{29.3}Ae_{65.0}$), or aegirine. The clinopyroxene grains in the groundmasses of the tinguaites are aegirine-augite as at Ajahunse $(Di_{16.9}Hd_{45.5}Ae_{37.6})$ or aegirine as in the samples from Hosadoddi $(Di_{0.4}Hd_{13.7}Ae_{85.9})$ and Ankachari Doddi $(Di_{1.0}Hd_{8.8}Ae_{90.2})$.

Bostonites. The pyroxene phenocrysts in the

bostonitic unit of the Chellipura dyke have three main zones, a core, a mantle surrounding the core and an outermost rim, the compositions of which are $\text{Di}_{81.0}\text{Hd}_{15.6}\text{Ae}_{3.4}$, $\text{Di}_{51.7}\text{Hd}_{35.3}\text{Ae}_{13.0}$ and $\text{Di}_{39.8}\text{Hd}_{40.9}\text{Ae}_{19.3}$, respectively (Table 2). The pyroxene composition in the bostonitic unit of the Arekattedoddi dyke varies from $\text{Di}_{29.9}\text{Hd}_{39.7}\text{Ae}_{30.4}$ in the core to $\text{Di}_{1.8}\text{Hd}_{28.7}\text{Ae}_{69.5}$ in the rim. The groundmass pyroxene microliths of the bostonites in these dykes are aegirine-augite and aegirine.

Solvsbergites. The pyroxene in the solvsbergite samples is intensely altered, and only few analyses could be performed, while that in the Devasandra dyke groundmass has a composition of $Di_{3.0}Hd_{10.5}Ae_{86.5}$.

Summary. The compositions of the pyroxenes are plotted on a Di-Hd-Ae diagram in Figure 3. The

analcime-tinguaites, tinguaites and bostonites display pyroxene trends typical of alkaline rocks (Stephenson 1972, Larsen 1976, Mitchell and Platt 1978, Platt and Woolley 1986, Woolley and Platt 1986). The cores of the phenocrysts contain more diopside, whereas the rims and the groundmass clinopyroxenes contain more hedenbergite and aegirine. The trends are affected by the oxygen fugacities and/or peralkalinities of the magmas which the pyroxenes crystallized from (Larsen 1976, Mitchell and Platt 1978, Platt and Woolley 1986, Woolley and Platt 1986). Similar pyroxene trends in the analcime-tinguaites, tinguaites and bostonites of the Bangalore district indicate that their melts had similar oxygen fugacities and/or similar peralkalinities. The absence of reverse zoning indicates that the magmas were not intermixed.

The pyroxenes are also heterogeneous with respect to titanium content, the cores and mantles of



Fig. 4. Photomicrograph of zoned K-feldspar phenocryst. Crossed polars. Field of view is about 4 mm x 1.5 mm

the phenocrysts usually containing less titanium than the rims and groundmass grains. A low Ti content may be attributed to a high formation temperature (Thompson 1974) and indicates that the cores and mantles crystallized at a higher temperature than the rims and groundmass grains, which crystallized quickly after intrusion of the magma.

The zoning displayed by the pyroxene is related to its non-equilibrium crystallization and metastable incorporation of Ti and Al in its structure. The compositions of the aegirines studied support the suggestion of Ranlov and Dymek (1991) that there is a solid solution between aegirine (NaFeSi₂O₆) and 1 to 26 mol% NaFe_{0.0-0.5}Ti_{0.0-0.5}Si₂O₆.

K-FELDSPARS

Occurrence

In terms of the 2V-angle, the K-feldspar of the dykes is sanidine in the tinguaites and solvsbergites and also in the analcime-tinguaites excluding the Madarhalli dyke, in which the K-feldspar is orthoclase, as also in the bostonites.

The K-feldspar is usually saussuritized, and Carlsbad twinning is common. The phenocrysts are subhedral or euhedral and often mantled by albite. Zoning is common in the phenocrysts of every rock type, but is best developed in those of the tinguaites (Fig.4). The groundmass K-feldspar is subhedral to anhedral.

Mineral chemistry

The orthoclase components (Table 3 and Fig. 5) show the majority of the sanidines to be K-bearing sanidines, as they contain more than 80% orthoclase (Smith and Brown 1988). Their anorthite content is mostly less than 0.2%.

In terms of element chemistry, the K-feldspars are zoned in sodium and sometimes also in barium. Barium is enriched in the cores of the K-feldspar phenocrysts, but the zoning of sodium varies. In some of the samples the groundmass grains and the Table 3. Results of the analyses.

Representative analyses of K-feldspars in the Karnataka alkaline dykes.

	1	2	3	4	5	6
wt%						
SiO ₂	65.01	64.45	64.32	64.77	61.40	64.11
TiO,	0.00	0.01	0.00	0.00	0.02	0.00
Al,Ó,	18.03	17.66	17.83	17.89	18.02	17.54
FeO	0.37	0.03	0.01	0.07	0.87	0.71
MgO	0.00	0.00	0.00	0.00	0.00	0.00
CaO	0.00	0.30	0.01	0.00	0.00	0.00
Na ₂ O	1.25	0.37	0.35	0.23	0.53	0.44
K,Õ	14.00	15.70	15.64	16.91	15.16	16.29
BaO	n.d.	n.d.	n.d.	0.36	3.73	0.72
Total	98.71	98.59	98.22	100.24	99.75	99.83

Structural formula based on 32 oxygens

Si	12.067	12.066	12.068	12.033	11.863	12.021
Гі	0.000	0.001	0.000	0.000	0.003	0.000
A1	3.944	3.896	3.943	3.917	4.103	3.876
Fe	0.058	0.004	0.002	0.011	0.141	0.111
Mg	0.000	0.000	0.000	0.000	0.000	0.000
Ca	0.000	0.060	0.003	0.000	0.000	0.000
Na	0.450	0.133	0.126	0.083	0.199	0.160
K	3.314	3.751	3.744	4.008	3.736	3.897
Ba	n.d.	n.d.	n.d.	0.026	0.280	0.053

Molecular per cent of end-member molecules

Or	88.1	95.1	96.7	98.0	95.0	96.1
Ab	11.9	3.4	3.2	2.0	5.0	3.9
An	0.0	1.5	0.1	0.0	0.0	0.0

1. Kodihalli, tinguaite, core of phenocryst, (DJK-18)

2. Arekattedoddi, tinguaite, core of phenocryst,(DJK-39)

3. Chellipura, bostonite, phenocryst, (DJK-13)

4. Arekattedoddi, bostonite, (DJK-39A)

5. Devasandra, solvsbergite, core of phenocryst, (DJK-25)

6. Devasandra, solvsbergite, rim of phenocryst, (DJK-25)

outer zones of the phenocrysts contain more sodium than the cores, but in other samples the relation is reversed.

PLAGIOCLASE

The plagioclase which forms anhedral grains in the groundmass or mantles K-feldspar is almost pure albite $(Ab_{98,9,99,7})$ (Fig. 5). Its iron content is low.

CONCLUSIONS AND DISCUSSION

The similar pyroxene trends found in the analcimetinguaites, tinguaites and bostonites of the Bangalore district prove that the melts had similar fugacities of oxygen and/or similar peralkalinities. The absence of reverse zoning indicates that no intermixing of the magmas took place.

The clinopyroxene and K-feldspar phenocrysts crystallized at a higher temperature, before the magmas intruded into their present locations, whereas the groundmass minerals, i.e. clinopyroxene, K-feldspar, albite and analcime, crystallized rapidly after intrusion of the magmas.

The consistent less aegirine contents of cores as compared to rims of pyroxene phenocrysts and groundmass pyroxenes suggest that the magma from which they were formed was initially less alkaline.

The K-feldspars of the alkaline dykes are zoned in sodium and sometimes also in barium. Barium is enriched in the cores of the K-feldspar phenocrysts, but the zoning of sodium varies.



Fig. 5. Feldspars of the Karnataka alkaline dykes plotted on a Ab-An-Or -diagram.

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