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THE EVOLUTION OF YEMEN BASALT IN RELATION TO THE RIFT DEVELOPMENT AND THE CHEMISTRY OF ITS MINERAL CONSTITUENTS

M.A.MATTASH*, GY.BUDA**

ABSTRACT

Tertiary volcanics cover a wide area in Yemen, approximately one tenth of the national territory. They occupy the majority of the western part of the country. Relative crustal thinning (affected by intensive tectonic activity) resulted in continental rifting (Late Oligocene - Early Miocene), through which magmas ascended and formed the Yemen Highlands and Plateaux. The Yemen Trap Series has mainly been controlled by the NW-SE faulting trend, which is parallel with the Red Sea trend, or less commonly (NE-SW) with the Gulf of Aden trend.

The Late Miocene-Quaternary volcanics are located in the coastal plain of the Gulf of Aden and in the central parts of Yemen. Yemen volcanics are composed of basic, to intermediate and acidic rocks.

Basic rocks are represented by basalts, which form the lower part of the volcanic complexes, whereas the acidic types [rhyolites, volcanic glasses and pyroclastics] occupy the upper part of the Yemen Trap Series volcanics.

This paper deals with the chemistry of the basaltic rocks and with the composition of their main constituents (olivine, pyroxene and plagioclase). The basaltic rocks have alkali-subalkali characters. The clinopyroxene phenocrysts show concentric and hour-glass zoning that can be regarded as a good indication of the magma evolution. The composition of the plagioclase ranges between labradorite-bytownite (for the phenocrysts, An_{87-63}) and andesine-labradorite (for the plagioclase in the groundmass, An_{61-40}). The olivine phenocrysts have higher forsterite content (Fo₉₀₋₇₅) in their cores; but these values decline at the rims as well as in the groundmass (Fo₇₅₋₄₀).

• + •• = DEPARTMENT OF MINERALOGY L. EOTVOS UNIVERSITY 1088 MUZEUM KRT. 4/A BUDAPEST-HUNGARY

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INTRODUCTION

The origin and evolution of Yemen volcanic rocks is regarded as the result of the rifting (synrift, Late Oligocene-Early Miocene) and postrift (Miocene up to Recent) phases of the formation of the Red Sea and the Gulf of Aden (in which oceanic accretion started).

In the Late Eccene, a widespread emergence of the greater part of the Arabian platform occurred. It was probably at that time that Gulf of Aden rifting was initiated, as the Indian Ocean spreading center propagated eastward (Haitham, et al., 1990).

This study was carried out to investigate the mineralogy, petrology, and structural geology of Yemen volcanics. Our study covers both the Yemen Trap Series (YTS, Late Oligocene-Early Miocene), and the Yemen Volcanic Series (YVS, Late Miocene-Recent) and their evolution in relation to the rift development (Fig. 1). In this paper we deal only with the basic rocks and their analyses related to the chemical composition of the basalts and their main constituents (olivine, pyroxene and plagioclase). We have also correlated the generation and development of these volcanic complexes by using the data obtained from seismic and drilled wells in the coastal plains of the Red Sea and the Gulf of Aden. By using microprobe, chemical and optical analyses *etc..*, we attempt to evaluate the chemical characters and the magma evolution of the investigated volcanics.

STRATIGRAPHY, TECTONIC SETTING AND GENESIS

The volcanic rocks of Yemen can be divided into two series *i.e.* Yemen Trap Series (YTS) and Yemen Volcanic Series (YVS).

The YTS had been developed through the period from Late Oligocene to Miocene (synrift-Shihr group).

It is composed of thick volcanic rocks (lava flows, dikes, sills etc.) mainly of basaltic composition, which form the lower part of the series, with intercalations of volcanoclastic deposits.

Genetically, the YTS had been associated with Tertiary continental and oceanic rifting and with the geodynamic of the Afro-Arabian lithosphere plates.

Numerous Tertiary intrusive bodies, mainly of alkali granites and syenites, have also been reported in many localities (particularly in the western part of the study area *i*/ with the Red Sea main trend) and they are closely associated with the volcanic complexes.

In Early Miocene, differentiated alkaline volcanic complexes poured out on to the continent 1 (W-SW) part of the country. The thickness of these volcanic complexes ranges from some metres in the eastern part of YTS to 2000 m in the western part or more in the central part of the area.

Huge amounts of basaltic flows built the lower part of the volcanic complexes as a consequence of continental crustal extension.

In the Red Sea coastal plain, faults caused strong subsidence in Oligocene time, which generated W-dipping blocks. Tilted blocks in the Gulf of Aden dip S. They have been affected by intensive tectonism (fault system) of ESE-WNW and NNE-SSW orientation and are filled with volcanic rocks mainly of synrift traps (Fig. 2).

The YVS is found on the coastal plain of the Gulf of Aden from the west (southeastern part of the Red Sea) Perim Island, Jabal-Kharaz passing onto Imran - Little Aden - Aden to the east (Shuqrah and Bir Ali etc..). It was generated and developed through the postrift stage (upper part of Shihr Group) and it is considered to be of Late Miocene to Recent time. It is also found in other localities throughout the country (Sanaa-Amran, Dhamar-Rada and Sirwah-Marib). Volcanic cones, domes, sheets and lava flows are the characteristic features of this series. It is composed mainly of basaltic lava flows and pyroclastics, except the Al-Lisi volcano (5 km E of Dhamar city), which is completely consisting of acidic lava flows (rhyolite, rhyolitic glass and ash). There is a sharp unconformity between the two series.

In the coastal plain of the Gulf of Aden (e.g. Imran well) thick continental basaltic traps occur, outpoured during aerial deposition, dated 18 \pm 6 Ma (Elf, 1990) with the same characteristic features of those known onshore. Such trap series is generated by synrift extensional fractures, which poured out huge amounts of lavas during a short time respectively. Partly we agree with the idea related to the mode of this magma



FIGURE 1 - SIMPLIFIED GEOLOGICAL MAP OF YEMEN



Fig. 2 Stratigraphic column illustrates Yemen Trap and Yemen Volcanic Series (Modified from Elf 1990).



Fig. 3 Geological sections correlate the distribution of Yemen volcanics in the offshore of the Gulf of Aden (a) and the Red Sea (b). The distribution of the volcanic rocks is also shown from the seismic data of drilled well in the Gulf of Aden offshore (c).

eruption but on the other hand there are some field evidences which also prove the central volcanic activity. The dyke swarms within the YTS have the same mineral composition with the adjacent outcrops, and the large extents of the pyroclastics (*e.g.* ignimbrites) without lateral variation in their thicknesses suggest fissural eruption. On the other hand the difference in the mineral constituents between the basaltic lava flows and the adjacent basaltic dikes and the presence of radial dike swarms support central volcanic activity (*e.g.* Final Report, 1988). Therefore, both fissural and central eruption can be assumed.

There is a tectonic contact between a continental crust block and the synrift deposits. During the Tertiary extensional phase, the continental crust had been stretched and broken. The resulting fault worked as a great crustal fault during the synrift stage.

Based on the data obtained from two wells offshore of the Gulf of Aden (Imran well) and Red Sea (Al-Janbiyah well) the thickness of the volcanic units may attain several kms, probably >3 kms (Fig. 3).

The upper part of the trap series is represented by the stratigraphic level of the break-up of the continent [(12-10 Ma)(Fig. 4) (After E!f Aquatine, 1990)]. Traps (Late Oligocene) are deposited on a first synrift sedimentary units.

An increase in heat flow happened which can be related to the beginning of the Tertiary rifting stage (40-12 Ma) and started decreasing from 12 Ma up to recent time (Fig. 4 After Elf Aq., 1990).

Based on isotope age data reported by many authors (e.g., Civette et al., 1978; Capaldi et al., 1983, 1987; Final Report, 1988; Chiesa et al., 1989) and also based on our radiometric age data [(16.6 \pm 1.8 Ma for the upper part of Jabal-Rabbiy (Ibb); 27.5 \pm 2 Ma for the basalt of Al-Dalil village; 31.6 Ma for the basalt of Al-siian village SW from Ibb city); 1992 this paper] it can be inferred that the volcanic activity in the YTS started about 30 Ma ago and continued for 11 Ma, Juring which differentiated rock types were erupted in different phases.

YTS can partly be correlated with some Oligocene-Miocene volcanic sequences of Alaji, Termaber, Galile, and Trap Series, in Ethiopia (Zanattin et al. 1980a; Zanattin 1992); and also with the continental alkali basalt flows of Harrat As Sirat and Harrat Hadan in Saudi Arabia (Fleck et al. 1973; Arno et al. 1980a). Such a correlation is including both age and chemical affinities.

After a long interval (Late Miocene-Quaternary) the extension continued and a widespread volcanic eruption was created along the coastal plain of the Gulf of Aden and in some other localities throughout the country (e.g. Sanaa-Amran, Sirwah-Marib, Dhamar-Rada etc..). This series was concentrated between 10 Ma and 5 Ma, e.g. Jabal-Kharaz gave 8.8 Ma (Final Report, 1988). Such eruptions were central-vent composite stratovolcanoes, e.g., Jabal Kharaz, and Jabal Shamsan at Aden city. The respective wide hiatus (9-10 Ma) allows us to make such a nomenclature division between the above mentioned series.

Further volcanic complexes are known as recent volcanic activities such as those found in Bir Ali, and AL-Lisi volcano in the vicinity area of Dhamar Province. Furthermore historical volcanoes have also been reported.

NATURE OF OCCURRENCE AND PETROGRAPHY OF THE BASALTS

Among the volcanic rocks of Yemen, basalts are predominant. They occur as 2-5 m lava flows. Columnar joints are also found, and in some localities exceed 50 m (e.g. Al-Dalil village). They also occur as dikes ranging from 0.1 m to 1-2 m in thickness and may attain several hundred metres in length. Thick basaltic lava flows, can be found in several places, e.g., in Dhala, Museimir, Am-Shatt etc., but the maximum thickness (200 m) was observed in the central part of Sanaa.

Basalts are characterized mostly by porphyritic textures (holocrystalline, ophitic and pilotaxitic) and massive structures, but amygdaloidal and vesicular (porous) structures are also reported. Amygdules are filled by zeolites, calcedony, carbonate, chlorite. In some cases the porous structures are devoid of any secondary minerals; particularly in the volcanics of the younger stages (e.g. Shuqrah, Marib *etc.*) they can attain up to 40% of the rock volume.

PETROCHEMISTRY OF THE YTS BASALT

Chemical classification (Final Report, 1988; Chiesa et al. 1989; Mattash et al., 1990) of the Yemen volcanic rocks is illustrated on the total alkali versus silica (TAS) diagram (after Le Bas et al., 1986)(Fig. 5a). It reveals that samples from Yemen Trap Series and Yemen Volcanic Series, lie along a trend of alkalic differentiation. The samples are plotted in the basalt field. According to the field boundary of Irvine and



Fig.5 The petrochemistry of YTS basalts is plotted in (a) the total alkali-versus silica diagram (after Le Bas et al. 1986), (b) AFM diagram after Macdonald and Katsura, 1964. ----- Hawaiian tholeiitic trend, ----- Hawaiian alkalic trend, and (c) Ti-Zr-Y diagram.

Barager (1971), basic members of Yemen volcanic rocks are characterized by alkali basalt (most of the samples of the northern regions), and by subalkaline basalt (most of the samples of the southern regions); only few samples occur along this field boundary. The more differentiated samples (undersaturated) are plotted in the tephrite - basanite field (Ba), in which they characterize the volcanics of the north regions and restricted to them only. This is also proved by AFM diagram (Fig. 5b), in which rocks from northern regions of YTS point out some differences in the alkali content in comparison with rocks from the southern part. On the other hand few samples are plotted in the hawaiite field (S₁); but the more differentiated rocks form a trend line from mugearite (S₂ field), to benmorite (S₃ field). The samples which lie on the basalt - hawaiite and tephrite - hawaiite fields are considered here as hawaiite.

Basalts from both mentioned localities are plotted in the Ti-Zr-Y diagram, the plotted values fall within plate basalt (WPB) field except few samples of transitional character (Fig. 5c).

MINERAL COMPOSITION

Olivine, pyroxene and plagioclase are the most abundant constituents and frequently form phenocrysts exceeding in some samples 30-50 % of the bulk.

Ilmenite, magnetite, apatite and less frequently biotite are the common accessories. The secondary minerals are represented by chlorite, epidote, serpentine, iddingsite, carbonate, and rarely amphibole (hornblende).

The glass phase is very rare and was only found in a very few samples, particularly in those lavas related to the Q-Recent volcanics.

In this study we are only focusing on the main constituents.

OLIVINE

Based on microprobe analyses (*Table 2*) olivine phenocrysts have higher forsterite content (Fo₉₀₋₇₅ in their cores); whereas these values decrease at the rims and within the groundmass (Fo₇₅₋₄₀). According to the optical study olivines range from 0.1 mm to 5 mm, forming euhedral to subhedral grains. In some samples olivines are partly altered to serpentine (Fig. 6a) and "iddingsite" (Fig. 6b). Olivines have *mg*-numbers ranging from 0.39 to 0.91.

The Mg content of Marib basalts has the highest value (MgO = 49.13%) among the analysed basaltic rocks of Yemen. Their mg-numbers (0.91) are also high, therefore, they may be considered to be of deeper origin perhaps from the upper mantle. They are similar in their composition to those olivine xenocrysts described by Kurat (1971), Dietrich and Poultidis (1985).

Taking into consideration the lowest contents obtained from the Aden area (mg-numbers = 0.39), which indicates a shallow magma origin in relation to Marib (mg-number = 0.91), the latter can be regarded as an indirect evidence of deeper magma origin, in which the continental crust becoming thinner E-ward (Balhaf, Al-Gawf-Marib etc.).

The mg-number values decrease at the rims and in the groundmass.

The NiO content slightly increases with increasing MgO (Fig. 7a), and the CaO, MnO and FeO are decreasing. The MnO content ranges between 0.19-2% (southern part of YTS) and it reaches the maximum in the Fe-enriched olivines and Mg-depleted particularly at the rimes of some olivine crystals (e.g. Dhala and Am-Shatt areas in the southern part of Yemen Trap Series), but the MnO (0.12-0.63 wt%) decreases (Fig. 7b) through the connecting line between Ibb and Sanaa.

The CaO content in relation to FeO has linear distribution, but this distribution becomes concentrated with decreasing FeO content (Fig. 7c, southern part of Yemen Trap Series)

PYROXENE

Their grain size ranges from 1 mm to 7 mm. Some of the crystals have simple twinning. Zoning commonly occurs, but concentric zoning and hour-glass structure are not uncommon (Fig. 8 a,b,c,d).

Based on microprobe analyses (Table 1), pyroxenes show a compositional variation from diopsidic to salitic and to augitic (Fig. 9a).

In the light of the diagram drawn by Leterrier et al. (1982), the majority of our pyroxenes compositions plot within the alkali basaltic clinopyroxene (Fig. 9b). According to their Cr, Mg, Fe and Al contents they can also be subdivided as follows: Chromian Ferroan DIOPSIDE, Aluminian Chromian Ferroan DIOPSIDE,

;	NT-2	MT-5 ²) NT-6	4 HT-14	5 HT-15	6 NT-18	7 MT-19	6 HT-27
p O	13.80	11.75	13.44	12.88	13.06	13.79	14.92	14.95
203	3.73	5.35	2.80	4,57	8.26	5.38	5.13	2.88
0	24.17	24.15	20.30	24.67	22.26	47.89	24.07	49.76 20.32
02	1.87	2.96	1.65	2.30	1.67	1.57	1.77	1.25
90 10	0.00	0.09	0.21	0.00	0.10	0.11	0.00	0.34
203	7.77	0.00	0.00	0.00	0.05	8.50	6.72	10.62
20	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
ija.	100.08	99.83	100.22	100.33	99.64	99.74	100.63	100.12
T site	1.805	1.746	1.842	1.767	1.734	1.746	1.744	81 1 24,252
VIV	0.144	0.236	0.124	0.201	0.266	0.214	0.222	1.855
3+	0.031	0.016	0.034	0.032	0.000		0.034	0.018
TAL:	2.000	2.000	2.000	2.000	4.000	2.000	2.000	2.000
Mi site				-	0.097	0.023		
3+	0.083	0.086	0.065	0.099	0.074	0.023	0.143	
14+	0.052	0.044	0.047	0.067	0.047	0.044	0.143	0.076
-3+	0.007	-		-	0.001	0.004	0.015	
2+	0.766	0.661	0.754	0.716	0.725	0.767	0.793	0.831
82+	0.092	0.169	0.134	0.118	0.056	0.064		0.058
DTAL:	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000
M2 site	-	_		-	_			
g2+ m2+	0.036	0.021	0.174	0.014	0.109	0.103	0.024	0.179
n2+	-	0.003	0.007		0.003	0.003	-	0.011
TAL:	0.964	0.976	0.819	0.986	0.885	0.893	0.947	0.811
AT#:	4.038	4.034	4.033	4.044	4.025	4.011		
NUM:	5.999	6.000	6.001	6.000	6.000	5.994	4.060	4.032
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5102 Ca0 T102 Ma0 Fe0 Cr203	нт- нт- нт- нт- нт- нт- нт- нт- нт- нт-	-6 - aluminian -14 - aluminian -15 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - 3100 	Magnaeium-rich) Ferroan DiOPSII subsilicic Perr Parroan DiOPSII chromian subsil (Magnasium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.90 0.39 9.03 0.00	AUGITE DE Coan DIOPSIDE DE (icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 46.91 23.21 1.60 0.13 6.66 0.27	13.02 1.97 13.02 1.97 13.124 20.92 0.79 1.02 11.12 0.15	19 - alumitian 27 - alumitian 14 2367 15.14 3.64 50.69 22.53 1.04 0.16 6.38 0.28	15 2576 1.5 2576 1.3.48 3.55 49.56 21.47 1.73 0.28 9.27 0.03	Licic Ferroan AUGITE 16 2627 13.20 3.73 49.30 21.84 1.96 0.25 9.29 0.02
Si02 CaO Ti02 NaO FeO Cr203 Na20 Sum T site	9 THLAH 23.54 5.65 46.79 23.64 20.64 7.44 0.17 0.60 99.76	-6 - aluminian -14 - aluminian -15 - aluminian -19 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - 3luminian -27 - 3luminian	Magnasium-rich) Ferroan DiOPSI subsilicic Perr Parroan DiOPSI (Magnasium-rich 11 2557 13.76 3.19 50.62 21.18 0.59 50.62 21.18 0.59 0.00 0.00 0.46	AUGITE DE Coan DIOPSIDE DE Licic Ferroan DI 12 2246b 13.77 4.51 46.91 1.66 0.13 6.46 6.27 0.51 \$9.57	NT- HT- DPSIDE 13 2409 13.02 1.97 51.24 20.92 1.02 1.12 0.15 0.56 100.81	19 - aluminian 27 - aluminian 14 2367 15.14 3.44 50.89 22.53 1.04 0.15 4.25 0.50 0.50 100.36	15 2576 13.48 3.55 49.56 21.47 1.73 0.28 9.26 9.26 0.67 0.67	LGIC Ferroan AUGITE 16 2627 13.20 3.73 49.30 21.84 1.96 0.23 5.29 0.02 0.69 100.28
5102 Ch0 T102 Nm0 Fe0 Cr203 Nm20 Sum T site 514+	нт- нт- нт- нт- нт- нт- нт- нт- нт- нт-	-6 - aluminian -14 - aluminian -15 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - aluminian -27 - 31 -31.41 	Magnaeium-rich) Ferroan DIOPSII subsilicic Perr Parroan DIOPSII chromian subsil (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.90 0.39 9.03 0.00 0.46 99.63	AUGITE DE Totan DIOPSIDE DE (icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 48.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.016	NT- HT- DPSIDE 13 2409 13.02 1.97 51.24 20.92 0.79 1.02 11.12 0.15 0.58 100.81 1.909	19 - eluminian 27 - eluminian 14 2367 15.14 30.89 22.53 1.04 6.38 0.50 100.36 1.866	15 2576 13.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.47 3.73 0.28 9.27 0.03 0.67 100.04 3.845	Licic Farroan AUGITE 1.20 15.20 13.20 2.73 49.30 21.84 1.96 0.25 9.29 0.02 0.69 100.28 1.833
5102 Cm0 T102 Mm0 Fe0 Cr203 Mm20 Sum T site Si4+ AllY	9 THLAH 23.54 5.65 46.79 23.64 20.64 7.44 0.17 0.60 99.76	-6 - aluminian -14 - aluminian -15 - aluminian -19 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - 3luminian -27 - 3luminian	Magnasium-rich) Ferroan DiOPSI subsilicic Perr Parroan DiOPSI (Magnasium-rich 11 2557 13.76 3.19 50.62 21.18 0.59 50.62 21.18 0.59 0.00 0.00 0.46	AUGITE DE Coan DIOPSIDE DE Licic Ferroan DI 12 2246b 13.77 4.51 46.91 1.60 0.13 6.64 6.27 0.51 99.57 1.816 0.184	NT- HT- DPSIDE 13 2409 13.02 1.97 51.24 20.92 1.02 1.12 0.15 0.56 100.81	19 - aluminian 27 - aluminian 14 2367 15.14 3.44 50.89 22.53 1.04 0.15 4.25 0.50 0.50 100.36	15 2576 13.48 3.55 49.56 21.47 1.73 0.28 9.26 9.26 0.67 0.67	Licic Ferroan AUGITE 16 2627 13.20 3.73 49.30 21.84 1.96 0.25 9.29 0.02 0.69 100.28 1.833 0.143
5102 Ch0 T102 Nm0 Fe0 Cr203 Nm20 Sum T site 514+	нт- нт- нт- нт- нт- нт- нт- нт- нт- нт-	-6 - aluminian -14 - aluminian -15 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - aluminian -27 - 31 -31.41 	Magnaeium-rich) Ferroan DIOPSII subsilicic Perr Parroan DIOPSII chromian subsil (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.90 0.39 9.03 0.00 0.46 99.63	AUGITE DE Totan DIOPSIDE DE (icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 48.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.016	NT- HT- DPSIDE 13 2409 13.02 1.97 31.24 20.92 1.97 31.24 20.92 1.97 3.02 1.97 3.02 1.97 3.02 1.57 0.58 1.000.81	19 - eluminian 27 - eluminian 14 2367 15.14 30.89 22.53 1.04 6.38 0.50 100.36 1.866	15 2576 13.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.48 3.47 3.73 0.28 9.27 0.03 0.67 100.04 3.845	Licic Farroan AUGITE 1.20 15.20 13.20 2.73 49.30 21.84 1.96 0.25 9.29 0.02 0.69 100.28 1.833
SiO2 Ch0 TiO2 NmC Fe0 Cr203 Nm2C Sum T site Si4+ All Fm3+ TOTAL: NL sit	нт- нт- нт- нт- нт- нт- нт- нт- нт- нт-	aluminian -14 - aluminian -15 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - aluminian -27 - 3100 -13.41 	Magnaeium-rich) Ferroan DIOPSII subsilicic Perr Parroan DIOPSII (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.90 0.39 9.03 0.00 0.46 99.63 1.694 0.106 -	AUGITE DE Totan DIOPSIDE DE (icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 48.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.000	NT- HT- DPSIDE 13 2409 13.02 1.97 51.24 20.92 0.79 1.02 11.12 0.15 0.58 100.81 1.909 0.084 0.005	19 - aluminian 27 - aluminian 14 2367 15.14 1.44 50.89 22.53 1.04 0.16 6.38 0.50 100.36 1.666 0.134 - 2.000	15 2576 13.48 3.55 49.56 21.47 1.73 0.28 9.27 100.04 1.645 0.155 - 2.000	16 16 16 16 16 16 16 2627 13.20 13.20 149.30 25.84 1.96 0.25 9.29 0.02 0.69 100.28 1.813 0.143 0.004
SiO2 CmO TiO2 NmO PeO Cr203 Nm2C Sum T site Si4+ Aliv Fu3+ TO7AL: NI sit	нт- нт- нт- нт- нт- нт- нт- нт- нт- нт-	-6 - aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - aluminian -27 - 3100 -10 -2247b 	Magnaeium-rich) Ferroan DioPSII subsilicic Perr Perroan DioPSII chromian subsil (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.98 0.39 9.03 0.06 0.46 99.63 1.694 0.106 2.000 0.034	AUGITE DE Totam DIOPSIDE DE (icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 48.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.000 2.000 0.014	HT- hT- hT- hT- hT- hT- hT- hT- h	19 - eluminian 27 - eluminian 14 2367 15.14 30.89 22.53 1.04 0.50 100.36 1.866 0.134 2.000 0.015	15 2576 13.48 15 2576 13.48 1.55 2576 13.48 2.55 49.56 21.47 1.73 0.28 9.27 0.03 0.67 100.04 1.55 -2.000 0.001	Licic Farroan AUGITE 1.20 1.20 1.20 2.73 49.30 2.84 1.96 0.25 9.29 9.29 0.69 100.28 1.833 0.143 0.004 2.000
SiO2 ChO TiO2 NhO FeO Cr203 Na20 Sum T site Si4+ AliV Fu3+ TOTAL: N1 sit AlVI Fe3+	NT- NT- NT- NT- NT- NT- NT- NT- NT- NT-	-6 - aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -27 - aluminian -27 - aluminian -27 - 3luminian -27 - 3	Magnaeium-rich) Ferroan DIOPSII subsilicic Perr Parroan DIOPSII (Magnesium-ricc) 11 2557 13.70 3.19 50.62 21.18 0.90 0.39 9.03 0.00 0.46 99.03 1.094 0.106 2.000 0.034 0.050	AUGITE DE Coan DIOPSIDE DE Licic Ferroan DI 12 2246b 13.77 4.51 46.91 2.221 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.000 2.000	NT- HT- DPSIDE 13 2409 13.02 1.97 51.24 0.79 1.02 1.97 0.15 0.54 100.01 1.909 0.081	19 - aluminian 27 - aluminian 14 2367 15.14 1.44 50.89 22.53 1.04 0.14 6.38 0.28 0.50 100.36 1.666 0.134 2.000 0.015 0.090	15 2576 13.48 3.55 49.56 21.47 1.73 0.28 9.27 100.04 1.645 0.155 - 2.000 0.001 0.105	Licic T4rroan AUGITE 1.2627 1.20 1.20 2.1.84 1.96 0.25 9.29 0.02 0.69 100.28 1.83 1.020 1.63 1.004 2.000
SiO2 Cao TiO2 NaC FeD Cr2003 Na20 Sum T site Si4+ AlTV Fe3+ TOTAL: NI sit AlVI Pa3+ Ti4+	HT- HT- HT- HT- HT- HT- HT- HT- HT- HT-	aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -27 - al	Magnaeium-rich) Ferroan DioPSII subsilicic Perr Perroan DioPSII chromian subsil (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.98 0.39 9.03 0.06 0.46 99.63 1.694 0.106 2.000 0.034	AUGITE DE Totan DIOPSIDE DE (icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 48.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.000 2.000 0.014 0.105	NT- HT- DPSIDE 13 2409 13.02 1.97 51.24 0.79 1.02 1.97 1.02 1.12 0.58 100.01 1.999 0.084 0.022 0.094	19 - eluminian 27 - eluminian 14 2367 15.14 3.04 50.89 22.53 1.04 0.50 100.36 1.866 0.134 2.000 0.015 0.990 0.015 0.990	15 2576 13.48 3.55 49.56 13.48 3.55 49.56 21.47 1.73 0.28 9.27 0.03 0.67 100.04 1.645 0.155 - 2.000 0.001 0.105 0.048	Licic Farroan AUGITE 1.20 1.20 1.20 1.20 1.20 1.20 1.20 2.25 9.29 9.29 9.29 0.02 0.69 100.28 1.813 0.143 0.143 0.004 2.000
5102 Ca0 Ti02 Ma0 Cr203 Ka20 Sum T Site Site T Site Site T Site Site T Site Site T Site Site T Site Site T Site Site T Site Site T Site Site Cr203 Sum	NT- NT- NT- NT- NT- NT- NT- NT- NT- NT-	-6 - aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -27 - a	Magnaeium-rich) Ferroan DIOPSI subsilicic Perr Parroan DIOPSI (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.90 0.46 99.03 0.00 0.46 99.03 1.094 0.106 - 2.000 0.034 0.050 0.034	AUGITE DE Coan DIOPSIDE DE (icic Ferroan DI 12 2246b 13.77 4.51 46.91 2.221 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.184 0.184 0.184 0.184 0.109 0.045 0.004	HT- hT- hT- hT- hT- hT- hT- hT- h	19 - aluminian 27 - aluminian 14 2367 15.14 1.44 50.89 22.53 1.04 0.14 6.38 0.28 0.50 100.36 1.666 0.134 2.000 0.015 0.090	15 2576 13.48 3.55 49.56 21.47 1.73 0.28 9.27 0.63 0.67 100.04 1.645 0.155 - 2.000 0.001 0.105 0.048	16 16 16 16 2627 1.20 1.70 1.37 4.30 2.1.84 1.96 0.25 9.29 0.02 0.69 100.28 1.83 1.00 1.43 0.064 2.000 0.104 0.055 0.001
5102 Da0 Tri02 Mm0 Cr203 Sum T Site Site Site T Site Site T Site Site T Site Site T Site Site Site Site Site Site Site Site	HT- HT- HT- HT- HT- HT- HT- HT- HT- HT-	aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -27 - al	Magnaeium-rich) Perroan DioPSII subsilicic Perr Perroan DioPSII chromian subsil 11 2557 13.76 3.19 50.82 21.18 0.98 0.39 9.03 0.06 0.46 99.83 1.894 0.106 2.000 0.034 0.027 0.766	AUGITE DE Totan DIOPSIDE DE (icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 48.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.000 2.000 0.014 0.105	HT- HT- DPSIDE 13 2409 13.02 1.97 51.24 0.15 0.52 0.059 1.02 1.12 0.15 0.54 100.01 1.909 0.005 2.000 0.004 0.022 0.004 0.723 0.167	19 - alumiiilian 27 - alumiiilian 14 2367 15.14 3.44 50.89 22.53 1.04 0.14 6.38 0.28 0.50 100.36 1.866 0.134 - 2.000 0.015 0.029 0.0090 0.029 0.0090 0.029 0.0090	15 2576 13.48 3.55 49.56 13.48 3.55 49.56 21.47 1.73 0.28 9.27 0.03 0.67 100.04 1.645 0.155 - 2.000 0.001 0.105 0.048	Licic Farroan AUGITE 1.20 1.20 1.20 1.20 1.20 1.20 1.20 2.25 9.29 9.29 9.29 0.02 0.69 100.28 1.813 0.143 0.143 0.004 2.000
5102 Da0 Tri02 Mm0 Cr203 Sum T Site Site Site T Site Site T Site Site T Site Site T Site Site Site Site Site Site Site Site	HT- HT- HT- HT- HT- HT- HT- HT- HT- HT-	-6 - aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -27 - a	Magnaeium-rich) Ferroan DIOPSI subsilicic Perr Parroan DIOPSI (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.90 0.46 99.03 0.00 0.46 99.03 1.094 0.106 - 2.000 0.034 0.050 0.034	AUGITE DE Totan DIOPSIDE DE [(icic Ferroan DI) AUGITE 12 2246b 13.77 4.51 48.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.000 2.000 0.014 0.109 0.045 0.045 0.045	HT- hT- hT- hT- hT- hT- hT- hT- h	19 - eluminian 27 - eluminian 14 2367 15.14 50.89 22.53 1.04 0.50 100.36 1.866 0.134 2.000 0.015 0.090 0.029 0.006	15 2576 13.48 15 2576 13.48 1.55 2576 13.48 1.55 2576 13.48 1.55 2576 13.48 1.73 0.28 9.27 0.03 0.67 100.04 1.55 -2.000 0.001 0.105 0.048 0.001 0.748	16 2627 1.20 2627 1.20 2.73 49.30 2.84 1.96 0.25 9.29 100.28 1.833 0.104 0.055 0.000 0.732
5102 Ca0 TiO2 Man0 Cr2003 Man0 Cr2003 Mal0 Sum T site Siat Nal7 Fult AlVI Fult AlVI Sid Fult AlVI Sid Fult AlVI Fult Fult Fult Fult Fult Fult Fult Fult	нт- нт- нт- нт- нт- нт- нт- нт- нт- нт-	-6 - aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -19 - aluminian -19 - aluminian -10 -2247b 13.41 5.77 47.04 2.19 0.65 0.54 0.54 0.552 -2.600 0.041 0.252 -2.600 0.041 0.252 -2.600 0.041 0.252 -2.600	Magnaeium-rich) Ferroan DOPSI subsilicic Perr Parroan DOPSI (Magnaeium-ricc) 11 2557 13.76 3.19 50.69 20.69 0.59 0.39 0.39 0.39 0.39 0.39 0.39 0.39 0.3	AUGITE DE Toon DIOPSIDE DE 122246b 13.77 4.51 4.51 4.51 4.51 4.51 6.64 0.27 0.51 99.57 1.816 0.184 0.000 0.014 0.045 0.062 1.000	HT- HT- DPSIDE 13 2409 13.02 1.97 35.24 20.22 0.79 1.02 1.02 1.02 1.02 1.02 1.02 1.02 1.02	19 - 100011100 27 - 110011100 14 2367 15.14 3.44 50.94 22.53 1.04 5.24 0.25 1.04 5.25 1.04 5.25 1.04 5.25 0.26 1.05 100.36 1.866 0.134 -2.000 0.035 0.039 0.031 1.000	15 2576 13.48 3.55 49.56 21.47 1.73 0.26 9.26 100.04 1.845 0.155 - 2.000 0.051 0.051 0.055 0.051 0.055 0.051 0.055	Licic Farroan AUGITE 1.20 3.73 49.30 2.184 1.96 0.25 9.29 0.02 0.69 100.28 1.813 0.143 0.143 0.004 2.000 - 0.104 0.732 0.106 1.000
5102 CR0 T102 MBD Tr02 MBD Tr02 SNM T site S14+ AllY FR3+ T07AL: NI sit S14+ T07AL: NI sit FR3+ T14+ CC3+ T07AL: NI sit T07AL: NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI sit R04+ NI Sit R04+ NI Sit R04+ NI SIT SIT SIT SIT SIT SIT SIT SIT SIT SI	NT- NT- NT- NT- NT- NT- NT- NT- NT- NT-	-6 - aluminian -14 - aluminian -15 - aluminian -15 - aluminian -19 - aluminian -27 - a	Magnaeium-rich) Ferroan DOPSI subsilicic Perr Parroan DOPSI (Magnesium-ricc) 11 2557 13.76 3.19 50.62 21.18 0.90 0.46 99.83 1.894 0.106 	AUGITE DE (con DIOPSIDE DE (icic Ferroan DI 12 2246b 13.77 4.51 46.91 23.21 1.60 0.13 6.66 0.27 0.51 99.57 1.816 0.184 0.000 2.000 0.014 0.109 0.045 0.006 0.036	NT- HT- DPSIDE 13 2409 13.02 1.97 51.24 0.79 1.02 11.12 0.58 100.81 1.909 0.086 0.005 2.000 0.054 0.022 0.024 0.024 0.022 0.167 1.000 0.090	19 - alumiiilian 27 - alumiiilian 14 2367 15.14 3.44 50.89 22.53 1.04 0.14 6.38 0.28 0.50 100.36 1.866 0.134 2.000 100.36 1.866 0.134 2.000 0.029 0.00	15 2576 13.48 3.55 49.56 21.47 1.73 0.28 9.27 0.03 0.67 100.04 1.645 0.155 - 2.060 0.001 0.748 0.097 1.000 0.087	16 2627 1.20 1.02 1.02 1.01 1.01 1.00 1.03 1.000 1.000 1.000 1.000 1.000 1.000 1.000 1.000 1.000 1.000 1.000 1.000 1.000
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Table 1. Representative microprobe analyses (average of 150 point analyses) of clinopyroxene from Yemen volcanics.

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Fig.6 Scanning electron images of olivine phenocrysis. The alteration to serpentine can be seen through the cracks as well as at the rims. Fo_{81} the average content in the core, Fo_{61} the average content at the rim (a), and (b) shows the alteration of olivine to iddingsite.Magnification: 54X, +N.



Fig. 7 Variation of NiO (a) and MnO (b) versus mg-number and (c) shows the variation of CaO versus FeO, in the olivines of Yemen alkali basalts.

<image>

Fig.8 Scanning electron images of zoned (a,b) and hour-glass structure (c,d) of clinopyroxene phenocrysts. The numbers shown on the crystals indicate the points analysed by microprobe.



Fig.9 Composition of clinopyroxenes of Yemen alkali basalts (a) and (b) shows that the analysed phenocrysts are plotted within the alkali basaltic clinopyroxene (see text).

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Fig. 10 Scanning electron images of titanaugite phenocryst. The average content of the TiO₂ in the core is 1.07%, and the average at the rim is 2.23%.



Fig. 11 (a) The variation of Cr versus mg-nuber, (b) shows the Al/Ti content and (c) illustrates the Ti versus mg-number.

Aluminian Chromian Subsilicic Ferroan DIOPSIDE, Aluminian Ferroan DIOPSIDE, Aluminian subsilicic Ferroan DIOPSIDE, Aluminian Titanian Subsilicic Ferroan DIOPSIDE, [Magnesium-rich] Augite and Aluminian [Magnesium-rich]'Augite. It is worth to mention that the augite is the most common and it is mostly euhedral. Titanaugite was also observed mainly at the rims of the pyroxene grains (Fig. 10). In some samples, pyroxene has been partly altered to chlorite, carbonate, epidote, and rarely uralitized (hornblende).

The Cr content of the clinopyroxene increases with increasing mg-numbers and attains the maximum value through the interval (0.7-0.9 mg-numbers). The Cr_2O_3 content in the core of some pyroxene crystals (Aluminian Chromian Ferroan DIOPSIDE) ranges up to 1.09 wt%, but the Cr_2O_3 content decreases at the rims and within the groundmass (Fig. 11a).

However, Dobosi et al. (1991) in their study found that the Cr content increases in those sectors having Tienrichment. The results found in our study are not in conformity with their findings, since our analyses showed that the Cr content decreases with increasing Ti. Some of our pyroxenes (Ti-poor sectors) are also having more Si, Mg and less Fe and Al.

 Al_2O_3 and TiO_2 have wide variation ranging between 0.40-5.34% for TiO_2 and 1.39-8.88% for Al_2O_3 . The relationship between Al and Ti distribution is given in (Fig. 11b). It shows that the Al content increases with increasing Ti.

The Ti/Al ratio can be considered as an approximation for a qualitative indication of the pressure prevailing during crystallization (Clark *et al.*, 1962; Kushiro, 1969). According to most of our analyses of the pyroxenes, the Ti/Al ratio varies between 0.1-0.2, which indicates high pressure origin. This ratio is higher at the rims of the pyroxene crystals rather than in their cores, a particular characteristic of the titanaugite varieties, indicating lower pressure than that of the above mentioned pyroxene cores. On the other hand, the high Al^{Vi} content and high Al^{Vi}/Al^{IV} ratio also indicates high pressure origin (Aoki and Kushiro, 1968; Wass, 1979). This ratio in our samples is between 0.30 and 0.40 in the cores und <0.26 at the rims and within the groundmass. The latter value indicates lower pressure respectively. The majority of the Ti contents are increased with decreasing mg-numbers, with exception of few samples (Fig. 11c).

PLAGIOCLASE

Their grain sizes range from 1 mm to 8 mm. Albite, carlsbad and pericliva twinns are characteristic features. Zoning is also common (Fig. 12 c,d).

Based on microprobe analyses (*Table 3*), the plagioclase composition ranges between labradorite-bytownite. The highest An contents in our studied samples reached An_{87-63} , in which the lower value characterizes the An content at the rims (Fig. 12 a,b).

Plagioclase within the groundmass has an average composition of laboradorite-andesine (An_{61-40}) , but in some samples the An content declines (andesine composition). The plagioclase has partly been altered into carbonate and epidote.

The groundmass also consists of augite (being strongly altered into chlorite) and euhedral crystals of olivine \pm opaque minerals (ilmenite \pm magnetite).

CONCLUSIONS

Yemen volcanics are parts of an extensive elongate outcrops associated with the Red Sea, Gulf of Aden and Afar (Ethiopia) rifts. They have been generated in the Late Oligocene-Early Miocene (synrift phase) and in the Late Miocene to Recent (postrift phase), and are mainly composed of basaltic rocks. The YTS emplaced in Oligocene and reached the maximum activity in Early Miocene times, in which both fissural and central eruptions took place. These volcanic complexes had been affected by faulting mainly of NW trend (Red Sea trend) and E-NE trend (Gulf of Aden trend). According to the available radiometric age data the Tertiary intrusive bodies (alkali granite, syenite *etc.*.) are closely associated with the Tertiary volcanics (YTS).

Yemen alkali olivine basalts have generated by fractional crystallization (differentiation). Duda and Schmincke, 1985; Bedard *et al.*, 1988; Dobosi, 1989 showed in their study that zoned pyroxenes are the best records of magma evolution. In our samples, zoning and hour-glass structure are common and characteristic features of the analysed pyroxenes, this gives good information about the evolution of the basaltic magma. The chemical composition of the pyroxenes proves that the magma had been crystallized and developed in deep (elevated pressure) and shallow magma chambers. The ratio of Al to Ti in the clinopyroxenes, the wide



Fig.12 (a) An-Ab-Or ternary diagram illustrates the composition of the feldspars from Yemen alkali basalts, (b) scanning electron images of feldspar phenocryst having an average content of An_{78} in the core and An_{63} at the rim. Zoning is also shown by the scanning electron images (c) and by theoptical microscope (d). Magnification : 34X + N.

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range of Mg content of the olivine phenocrysts, and the gap of the An content among the plagioclase phenocrysts indicate deep and shallow magma origin respectively.

According to the recent data obtained from seismic and drilling through the coastal plain (offshore) of the Gulf of Aden (Elf Aqu., 1990), and of the eastern side of the Red Sea (Total, 1989), it can be concluded that the thickness of the volcanics (mainly of basaltic composition) may attain more than 3000 m below the sea level.

YTS can partly be correlated in both age and chemical characters with some younger volcanic sequences of Late Oligocene-Miocene in Ethiopia and Saudi Arabia. On the other hand the Late Miocene - Quaternary volcanics of Yemen (YVS) are corresponding to those volcanics reported from Afar volcanic field (Ethiopia) and to those volcanic fields reported from the southern and western parts of Saudi Arabia. They are in accordance in age and in chemical trends.

According to the optical study of many basaltic samples, the presence of olivine in a stable form, and the availabality of titanaugite indicate an alkaline character for these rocks. In general the Yemen volcanics mostly show an alkaline trend, but tholeiitic affinity is partly represented by some basaltic rocks of the YTS e.g. the south-westernmost area (Am-Shatt). It is the nearest part (of the YTS) to the Gulf of Aden, which is probably affected and generated at that time in which the western part of the Gulf of Aden (rift phase) had been generated.

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