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Petrology and geochemistry of bimodal volcanic rocks of Southern Lake Hayk area, northwestern Ethiopian plateau: implication for their petrogenesis

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ABSTRACT: This study presents and integrates field, petrological, and whole-rock geochemical (major and trace elements) data of the volcanic rocks from the Lake Hayk area to understand their petrogenesis. The study area's major lithological components include lower and upper basalt, rhyolitic lava, rhyolitic tuff, rhyolitic ignimbrite, and unwelded tuff. Petrographic analysis suggests that felsic rocks are dominated by quartz and well-developed sanidine (K-feldspar) phenocrysts with glassy groundmass, whereas mafic volcanic products are characterized by aphyric to porphyritic textures with the olivine and plagioclase dominant phenocryst. The area constitutes bimodal composition of flood basaltic to rhyolitic rock with scarce intermediate composition. Basalts have low Rb/Nb = 0.5–0.58, La/Nb = 0.88–1.06 and high TiO₂ = 2.08–3.04, basaltic andesite have higher Rb/Nb = 2.7, La/Nb = 1.81 and low TiO₂ = 1.96 and rhyolite Rb/Nb = 0.97–1.69, La/Nb = 0.51–1.08 and lower TiO₂ = 0.41–0.71. The positive Ba and negative K anomalies testify amphibole mantle source. The basalts are characterized by low CaO/Al₂O₃ ratios (0.71–0.97) and relatively less fractionated and flat HREE patterns with (TbN/YbN = 1.75–2.33) chondritic values. This suggests a mantle source mostly containing spinel rather than garnet. Rhyolites are characterized by a steep negative correlation in bivariate plots of MgO, Fe₂O₃, TiO, and CaO against SiO₂ and positive anomaly of Ta with slight Nb trough. This suggests that Fractional Crystallization is the major process for the genesis of rhyolitic rocks, rather than crustal contribution, partial melting, and assimilation, producing rhyolitic rocks.

Key words/phrases: crustal contamination, fractional crystallization, major and trace elements, mantle source, petrogenetic process

INTRODUCTION

Continental flood volcanism (CFV) is associated with rupturing of the Earth's lithosphere. As continental breakup progresses into developed rifting, the geochemical characteristics of the lavas typically change to those of oceanic lavas (ocean island basalts, OIBs; and mid-oceanic ridge basalts, MORBs), implying that upper mantle elements are more involved (Pik et al., 1999). Although continental flood basalt (CFB) provinces are typically associated with deep-origin hot mantle plumes, higher mantle and crustal sources play a significant role in their formation (Richards et al., 1989). Nevertheless, the dating of relationships between rifting leading to ocean-floor formation and continental flood basalt (CFB) formation are complex and can vary in space and time (Courtillet et al., 1999; Hawkesworth et al., 1999). Several petrogenetic models have been presented to explain the formation of silicic volcanic rocks in mafic igneous provinces. Rhyolites and associated silicic volcanic rocks (60 - 81 wt% SiO₂) are generated by a combination of processes. (i) Assimilation and fractional crystallization (AFC) of

mafic magma with significant crustal contribution or melting of continental crust (Dereje Ayalew et al., 2002; Dereje Ayalew and Gezahegne Yirgu, 2003, Haldera et al., 2020), and (ii) Parental mafic magma fractional crystallization with minimal crustal contamination (Dereje Ayalew et al., 2002; Haldera et al., 2020). According to Haldera et al. (2020), rhyolites derived by significant fractional crystallization have a modest to non-existent Nb-Ta anomaly, the occurrence of clinopyroxene phenocrysts, and steeply negative slopes in bivariate major oxides plots against SiO₂. Further, Dereje Ayalew et al. (2002) have documented that the plateau rhyolites are geographically correlated with the basalts, low- TiO₂ and high- TiO₂ rhyolites occurring in association with low- TiO₂ and high- TiO₂ flood basalts. Although the area is largely basaltic, the Oligocene volcanic products of the northern Ethiopian plateau show a significantly bimodal distribution of basalts and rhyolites (Dereje Ayalew and Gezahegne Yirgu, 2003). As noted for other continental flood basalt provinces, the coincidence of basalt and rhyolite provinciality suggests that the rhyolites were derived either by fractional crystallization of the associated basalt magma types or by partial

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According to conventional classification, the Ethiopia's Tertiary volcanic sequence is divided into four groups based on their ages and stratigraphic position. From bottom/old to top/young, these are known as the Ashange, Aiba, Alaje, and Termaber Formations (Mohr, 1983). Based on their geochemical makeup the flood basalts are further divided into three groups: low Ti basalts (LT), high Ti1 basalts (HT1), and high Ti2 basalts (HT2) (Pik et al, 1998 & 1999). According to Dereje Ayalew (2019) the ages of marginal rhyolites may be classified into four categories: Oligocene trap rhyolites (28.5–30.8 Ma, on average 30 Ma), early Miocene rhyolites (17.4–23.5 Ma, on average 20 Ma), Rhyolites from the late Miocene to Pliocene epochs (between 8 and 4 Ma), as well as Quaternary rhyolites.

Sampling and analytical technique

15 representative rock samples were prepared for petrographic characterization and 10 samples were prepared for major and trace element analysis. To remove the weathered part from the surface of the rock samples, the selected pieces were broken by hammer into the desired sizes. Following drying, the samples were crushed to 70% of less than 2 mm, riffled to separate the bulk samples, and the splits were ground to 93.3 % of less than 75 microns. The powdered samples were submitted to Australian Laboratory Science (ALS), to determine the concentration of major and trace elements. Inductively Coupled Plasma Atomic Emission Spectrometry (ICP-AES) was used to examine major elements, whereas Inductively Coupled Plasma Mass Spectrometry was used to examine trace elements (ICP-MS). Besides, multi-element four acid digestions 81(ME-4ACD81) were used for base metal determinations such as Cr, Co, Ni, and V. In addition, loss on ignition (LOI) at 1000 °C is determined by an instrument WST-SEQ. The precision of all measured samples is within the confidence level, with major elements like MgO and Na₂O having precisions of 0.1 to 0.2 % and 0.01 %, respectively, especially in comparison to SiO₂, Al₂O₃, Fe₂O₃, CaO, and K₂O.

Other than transitional metals, the precision of the most of trace elements, including REE, is better than 0.6 %. The precision is better than 2% for transitional metals including Co, Cu, Zn, Sc, and Ni.

RESULTS

Geology and petrography

The study area which constitutes Oligocene volcanic rock (flood basalts 31 Ma to shield volcano 25Ma) (Hoffmann et al., 1997; Kieffer et al., 2004; Dereje Ayalew et al., 2019). The

mappable rock units that cover the study area are lower basalt, upper basalt, rhyolite lava, and rhyolitic tuff (Fig.3A). The study area constitutes different volcanic rock units with different texture, mineral assemblages which shows their origin, evolution, and nature of magma generation. Volcanic rock in the study area are flood basalt (70-190m) In the form of sheet, low viscosity magma covering the bottom part or overline by rhyolitic lava(10 – 230 m) and shield volcanoes composed of successive layers of lava flows and pyroclasts and its thickness is (360 m). Flood volcano in the study area includes porphyritic basalt and aphyric basalt and welded rhyolite overlies flood basalt and at the top there is shield volcano constitutes intercalation of strongly welded rhyolite, less welded rhyolite, ignimbrite, amygdaloidal basalt, shield basalt, and glassy rhyolite (fig.4). The flood volcano separated with shield volcano by reddish colored paleosol (near to 30 cm thickness). Farther volcanic rocks of the study area can be classified into mafic and felsic based on their color and visible mineralogical composition. The mappable rock units that are found in the study area are lower basalt, upper basalt, rhyolitic lava, rhyolitic tuff, and friable tuff. Furthermore, the study area is associated with geological structures such as joints and faults. Petrographically analyzed samples (fig.5) were selected based on their stratigraphic position and lithologic variation. The porphyritic basalts observed in the study area constitute phenocrysts of clinopyroxene, plagioclase, and olivine with a groundmass of microcrystals of plagioclase, pyroxene, and opaque mineral (Fe-Ti oxide) with porphyritic and seriate textures majority of the samples. Rhyolite rock units, in addition to basalt rock samples, are characterized by a vitrophyric to porphyritic texture with phenocrysts of quartz and sanidine, and groundmass of volcanic glass, as well as rock fragments (fig.5). Petrographically, Olivine-Clinopyroxene Phyric Basalt rock unit is characterized by its highly porphyritic texture with phenocrysts of dominantly by olivine and secondly by clinopyroxene and embedded in groundmass very fine plagioclase and opaque mineral (Fe-Ti oxide). With a modal percentage of 70.51 – 76.8% phenocrysts of 36.88 - 41.5% olivine and 33.63 - 35.3% clinopyroxene and 15.3 – 23.2 % of groundmass of 8.6 - 9.5% microcrystalline plagioclase and 6.7 - 14.7% opaque mineral. The mineral grain shows a seriated texture because of the presence of a wide range of grain size variations from very fine plagioclase and clinopyroxene grain to coarse-grained olivine and clinopyroxene. Furthermore, olivine and clinopyroxene phenocrysts have a subhedral to euhedral shape, with irregular cracks and modest iddingsite alteration along the fissures in the olivine grain. Aphyric basaltic rocks found in the

study area which is aphyric in hand sample and petrographically microcrystalline in thin section when analyzed under petrographic microscope. These rocks have a dark gray color in hand samples, and crystals that are undetectable to the naked eye have a glassy appearance. In thin section, they contain few phenocrysts with some indistinguishable mineralogy in groundmass. The rock consists of <7% phenocrysts of euhedral to subhedral plagioclase and clinopyroxene. This unit is dominantly composed of microcrystal of plagioclase and with groundmass of pyroxenes and opaque mineral. Petrographically, Plagioclase Phyric Basalt rock unit is characterized by its highly porphyritic texture with phenocrysts of plagioclase and embedded in the groundmass of microcrystals of plagioclase, clinopyroxene, and opaque mineral (Fe-Ti oxide). The modal composition of this rock unit is dominated by phenocrysts of plagioclases the mineralogical modal percentage is ~75.5% and ~24.3% groundmass, which are composed of 71.5 - 81.3% plagioclase phenocryst with a groundmass of 5 - 18.7% clinopyroxene and 9.7 - 20% opaque mineral. Plagioclase, clinopyroxene, and opaque mineral microcrystalline textures account for the majority of the groundmass. In addition, plagioclase phenocrysts show subhedral to euhedral shape and occur as elongated penetrative

twinning nature. Petrographically analyzed rhyolite rock samples such as T1S6, T4S5, T2S5, and T3S4. These rhyolitic rock samples show vitrophyric, Porphyritic, glassy, and banding texture flow. Phenocrysts are alkali feldspar (sanidine) and quartz minerals embedded in the groundmass of mainly volcanic glass and very few opaque minerals. Sample T3S4 shows that flow banding microstructure and felsic texture with phenocrysts (10%) and groundmass (90%). Sample T1S6, T2S5, and T3S4 show similar petrographic characteristics. The rhyolite sample that petrographically analyzed shows porphyritic texture with 29.5-65.2% phenocrysts and, 34.87-0.44% groundmass /volcanic glass. Compositionally, 18.5 - 27.5% sanidine, 3 - 35.7% quartz, 1% plagioclase, 4.2-8.9% opaque, and 30.8-62.3% a groundmass of microlithic sanidine, quartz, volcanic glass, and opaque. Once more, sample a similar petrographic characteristic. Rhyolitic Ignimbrites: One sample T3S1 from the pyroclastic flow deposit is classified as welded tuff/ignimbrite. It has up to 21.25% nepheline, 3.75% clinopyroxene, perlitic crack 16%, 10.24% rock fragments, and 54.4% volcanic glass. Crystals are euhedral sanidine and anhedral clinopyroxene and quartz and also contain skeleton-like glass shards.

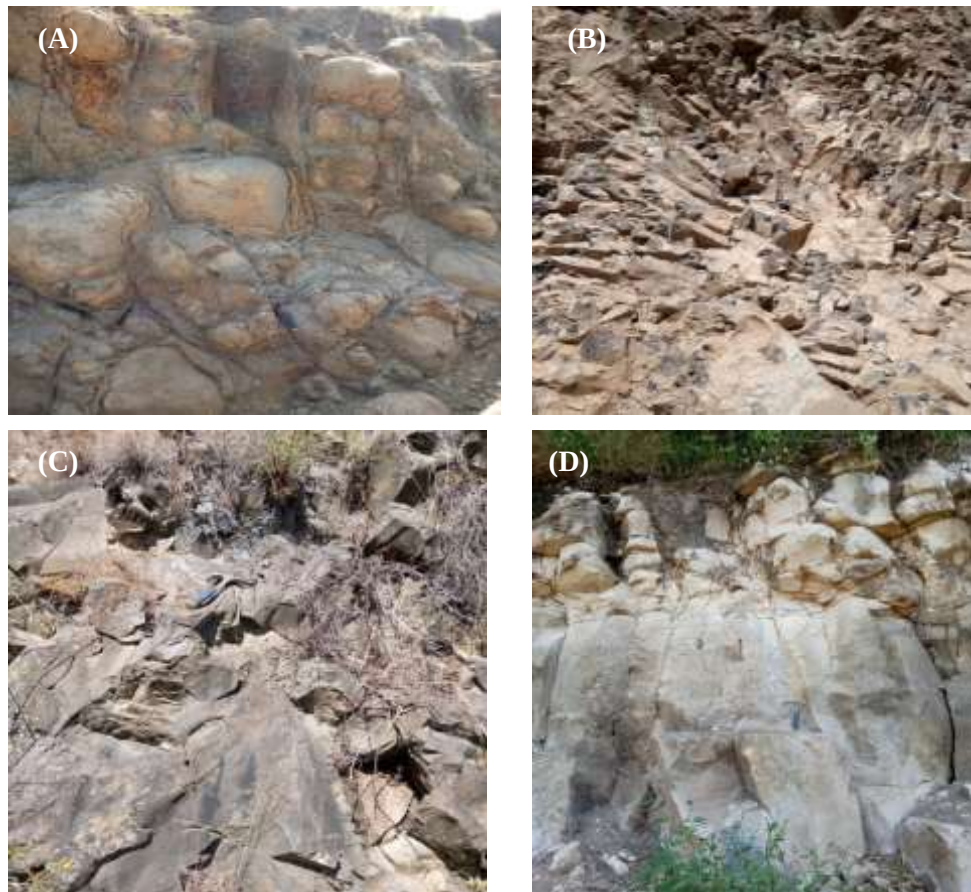


Figure 2. Field photographs of Lake Hayk area (A) Lobe structure solid cores. (B) Complex columnar jointing observed at the top of rhyolite sequence. (C) Aphyric basalt (D) Rhyolite rock unit with vesicles at top part. Note hammer for scale.

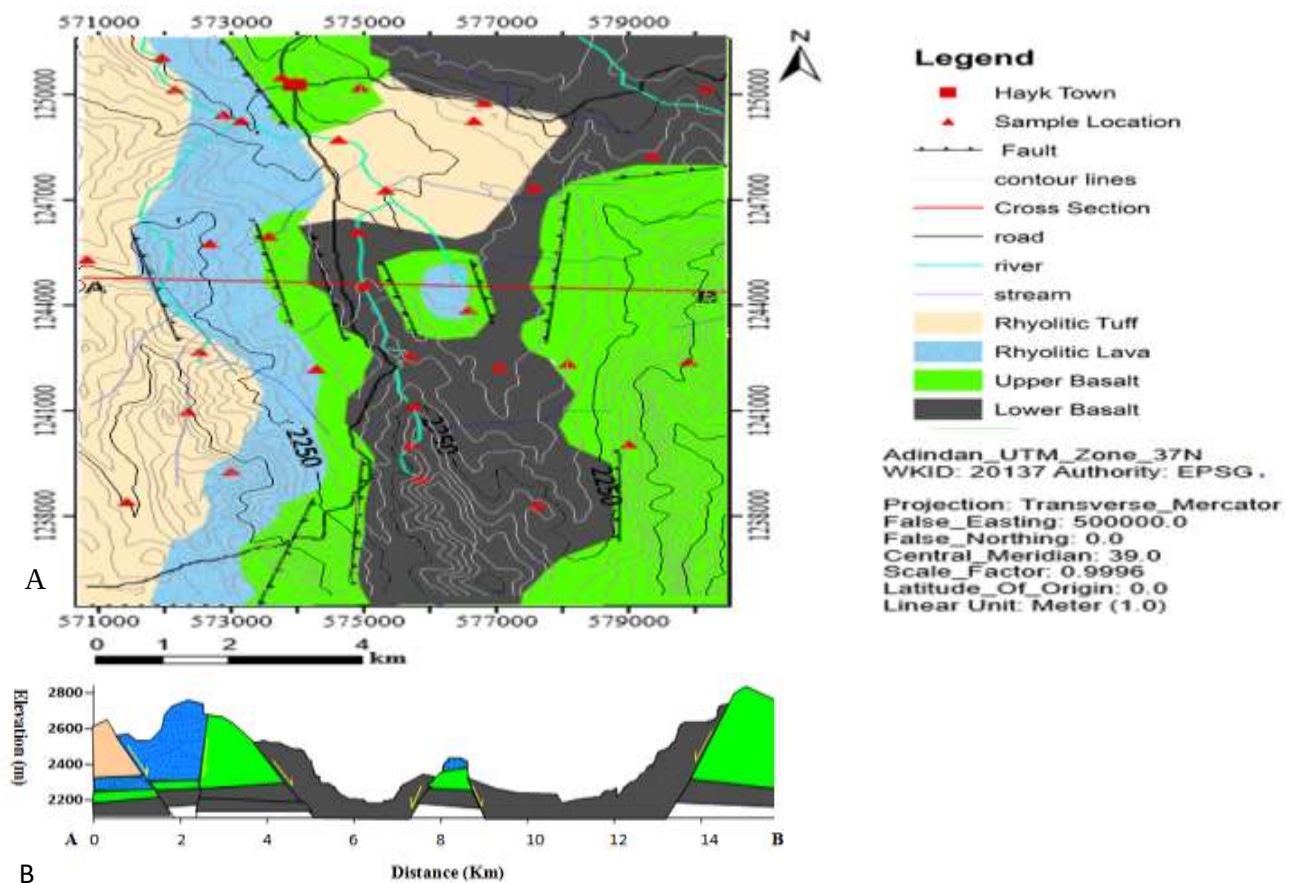


Figure.3. (A), Geological map of the study area and (B), schematic geological cross section (A to B) of Lake Hayk area.

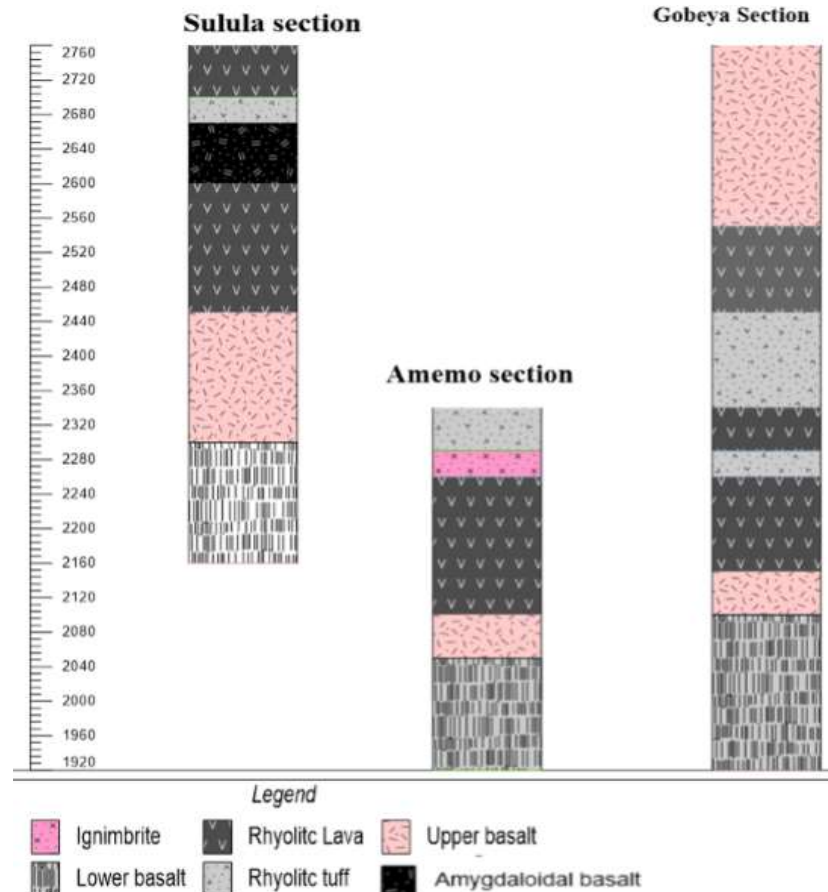


Figure 4. Volcanic stratigraphic section of the study are.

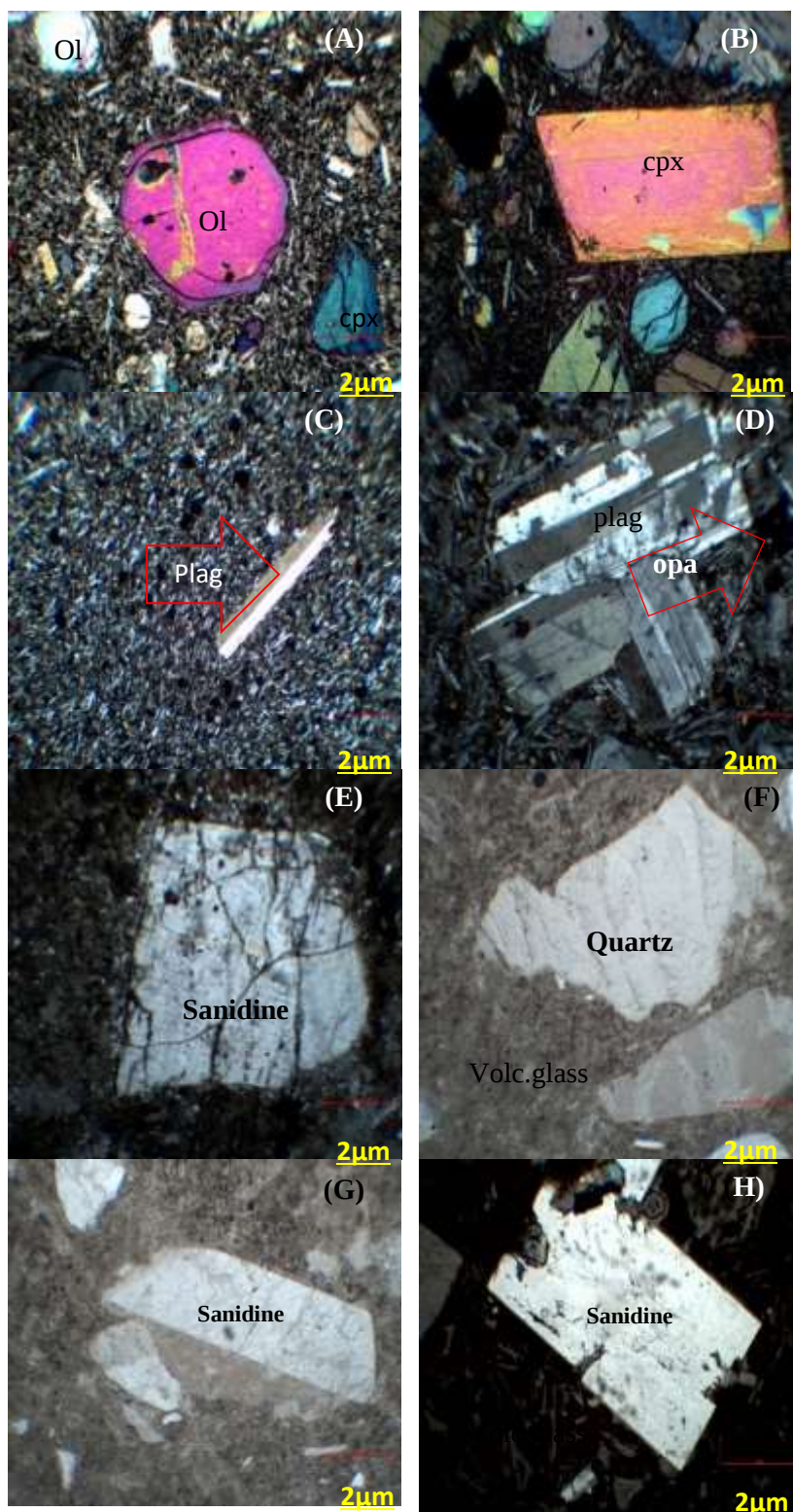


Figure.5. Photomicrographs of representative textures and modal mineralogy of basalts and rhyolite (A), Subhedral to euhedral-shaped phenocrysts of megacryst olivine (ol) and embedded in a groundmass are dominantly fine-grained plagioclase laths and fine-grained clinopyroxene (CPX) with striate texture. (B), Euhedral shaped phenocrysts of clinopyroxene with fine to coarse-grained laths and olivine embedded by fine-grained plagioclase groundmass. It shows a seriate texture. (C), Intersertal texture of aphyric basalt with elongated plagioclase embedded in microcrystals of plagioclase and opaque mineral. A sample is from the upper flood basalt (2207 masl). (D), phenocryst of plagioclase (plag) embedded in opaque mineral (Fe-Ti oxides), clinopyroxene, and fine plagioclase groundmass, it has glomerophyric texture, subhedral shape with oscillatory zoning. (E), a euhedral phenocryst of alkali feldspar (sanidine) embedded in volcanic glass (Volc.glass) and microlithic sanidine and quartz. (F), subhedral quartz megacryst phenocryst with a groundmass of volcanic glass and opaque minerals (G), euhedral sanidine and anhedral quartz phenocrysts embedded in volcanic glass and microlithic sanidine and quartz (H), Rectangularly developed euhedral shape sanidine, subhedral clinopyroxene and rock fragment with groundmass of volcanic glass.

Table 1. Major element wt. % and trace element ppm data of Lake Hayk volcanic rocks.

Sample	T1S1	T2S3	T7S5	T7S1	T3S2	T3S6	T1S3	T3S4	T5S2	T2S1EA
SiO ₂	45.5	48.4	48.5	55.9	63.7	66.7	69.7	71.3	72.3	77
TiO ₂	3.04	2.08	2.37	1.96	1.04	1.08	0.41	0.54	0.71	0.47
Al ₂ O ₃	11.9	15.25	14.25	15.15	15.55	13	14.65	11.95	13.65	10.5
Fe ₂ O ₃	13.1	11.9	12.25	11.15	5.29	5.36	2.99	3.49	2.72	2.17
FeOt	11.79	10.71	11.03	10	4.76	4.83	2.691	3.2	2.25	1.95
MnO	0.21	0.18	0.18	0.17	0.18	0.24	0.06	0.08	0.07	0.07
MgO	9.16	6.16	7.56	3.94	1.35	0.92	0.16	0.25	0.25	0.08
CaO	11.85	11.75	9.93	7.61	3.1	1	0.22	0.28	1.04	0.1
Na ₂ O	2.31	2.66	2.62	3.34	5.22	4.69	3.76	4.4	4.57	2.66
K ₂ O	0.88	0.45	0.81	1.88	2.58	4.48	4.09	5.02	4.21	5.53
P ₂ O ₅	0.53	0.28	0.32	0.29	0.26	0.26	0.03	0.09	0.1	0.01
LOI	1.49	1.07	1.5	0.44	1.57	1.55	3.07	1.04	0.68	1.01
Total	100.1	100.27	100.4	101.9	99.98	99.39	99.19	98.48	100.39	99.61
#Mg	58.31	50.87	55.24	41.41	33.8	25.56	9.6	12.53	6.87	6.08
Sc	33	32	29	25	8	8	4	5	6	6
V	337	301	287	268	23	41	10	18	11	14
Cr	490	110	410	30	<10	10	<10	<10	<10	<10
Co	53	44	46	34	3	4	1	1	2	1
Ni	135	51	97	15	<1	13	4	2	3	1
Cu	159	96	40	31	1	6	1	1	1	<1
Zn	113	103	108	109	117	161	122	163	108	176
Ga	20.5	20.6	19.7	20.2	24.5	31	29.9	34.4	25.1	28.8
Rb	20.6	8.5	10.9	40.2	67.8	102	111.5	123	124.5	136
Sr	546	471	405	342	455	222	21.5	35.1	151	8.6
Y	25.6	23.7	25.3	29.9	39.2	56.8	95.7	90.6	57.2	18.8
Zr	247	146	161	215	452	849	893	1010	623	1190
Yb	1.89	2.26	2.42	2.91	3.76	5.01	10.2	8.1	5.99	3.33
Nb	35.4	17	19.7	14.9	51.6	100.5	114.5	120	73.7	117
Cs	0.33	0.11	0.07	0.22	1.26	0.66	0.81	0.76	1.01	0.61
Ba	256	176.5	260	501	738	774	430	343	698	105
La	31.1	18.1	19.5	27	52.7	88.2	106	104	79.6	59.8
Ce	66.4	39.3	42.1	54.9	110.5	190.5	103.5	219	149.5	92
Pr	9	5.5	5.61	6.98	13.8	24.3	28.4	28.7	19.25	10.1
Nd	37.8	24.4	24.6	30.5	57.4	99.5	110	116.5	74.3	39.9
Sm	8.1	5.8	5.63	6.56	11.15	20.5	21.8	24.9	14.25	7.57
Eu	2.33	1.88	1.9	1.88	3.09	5.28	3.12	4.82	2.66	1.44
Gd	6.82	5.81	5.81	6.48	10	16.45	18.7	21.1	12	5.46
Tb	0.94	0.88	0.88	0.97	1.42	2.31	2.99	3.27	1.9	0.76
Dy	5.45	5.13	4.98	5.52	7.51	12.1	17.8	17.9	10.6	3.89
Ho	0.91	0.92	0.93	1.09	1.39	2.19	3.36	3.28	2.02	0.77
Er	2.36	2.6	2.6	3.11	3.97	5.85	10.15	9.1	5.93	2.28
Tm	0.31	0.35	0.35	0.43	0.58	0.81	1.52	1.29	0.85	0.42
Yb	1.89	2.26	2.42	2.91	3.76	5.01	10.2	8.1	5.99	3.33
Lu	0.3	0.29	0.34	0.42	0.56	0.69	1.45	1.16	0.86	0.57
Hf	5.8	4	4	5.7	11.2	20.7	22.1	24.7	15.7	28.1
Ta	2.1	0.8	0.9	0.9	3.3	6.1	6.8	7.4	4.4	8.1
Pb	<2	<2	2	7	10	5	15	14	13	15
Th	2.9	1.5	2.1	2.93	7.2	11.65	17	16.05	14.3	16.9
U	0.86	0.43	0.43	0.48	2.39	4.04	3.4	5.23	3.38	1.18
Eu/Eu*	0.98	1.00	1.01	0.89	0.90	0.88	0.48	0.65	0.63	0.62

Major elements

Diagrams with their descriptions and interpretation of major elements are based on the recalculated volatile free basis results. The analytical results for both major and trace elements are presented in Table.1. Diagrams with their descriptions and interpretation of major elements are based on the recalculated volatile free basis. Based on SiO₂ versus Na₂O + K₂O (wt %) (TAS) Lake Hayk volcanic rocks are classed as basalt, basaltic andesite, trachydacite, and (following Le Bas et al, 1986). Most of the samples

fall on the boundary line of transitional to sub alkaline/tholeiitic field indicating that mostly transitional to sub-alkaline/tholeiitic affinity. Major elements against silica SiO₂ are illustrated below in Harker variation diagrams (Fig.8). The examined samples loss of ignition (LOI) ranges from 0.44 to 1.57 %, showing that the samples are fresh; whereas one sample (T1S3) has a LOI of 3.07 %, (table.1) Therefore, only one sample could be used to determine how post-emplacement alteration affected elemental mobility.

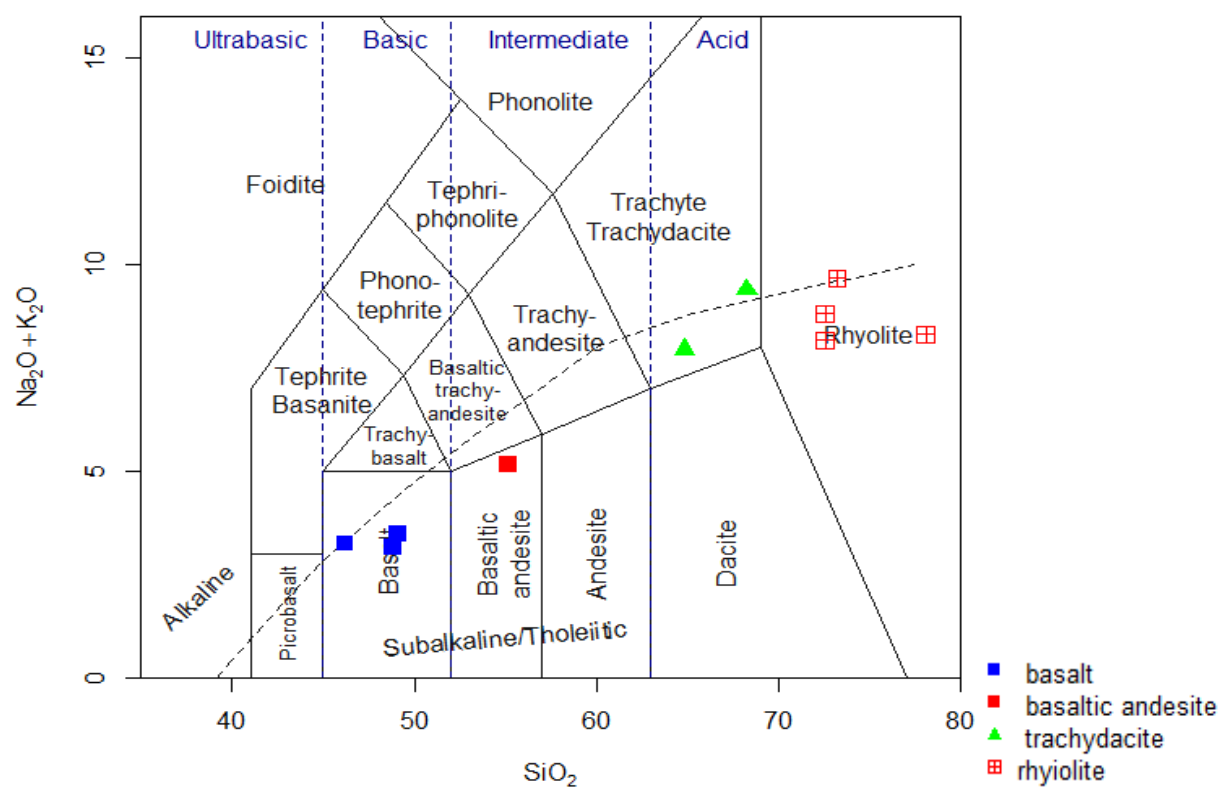


Figure 6. TAS classification diagram of Lake Hayk volcanics.

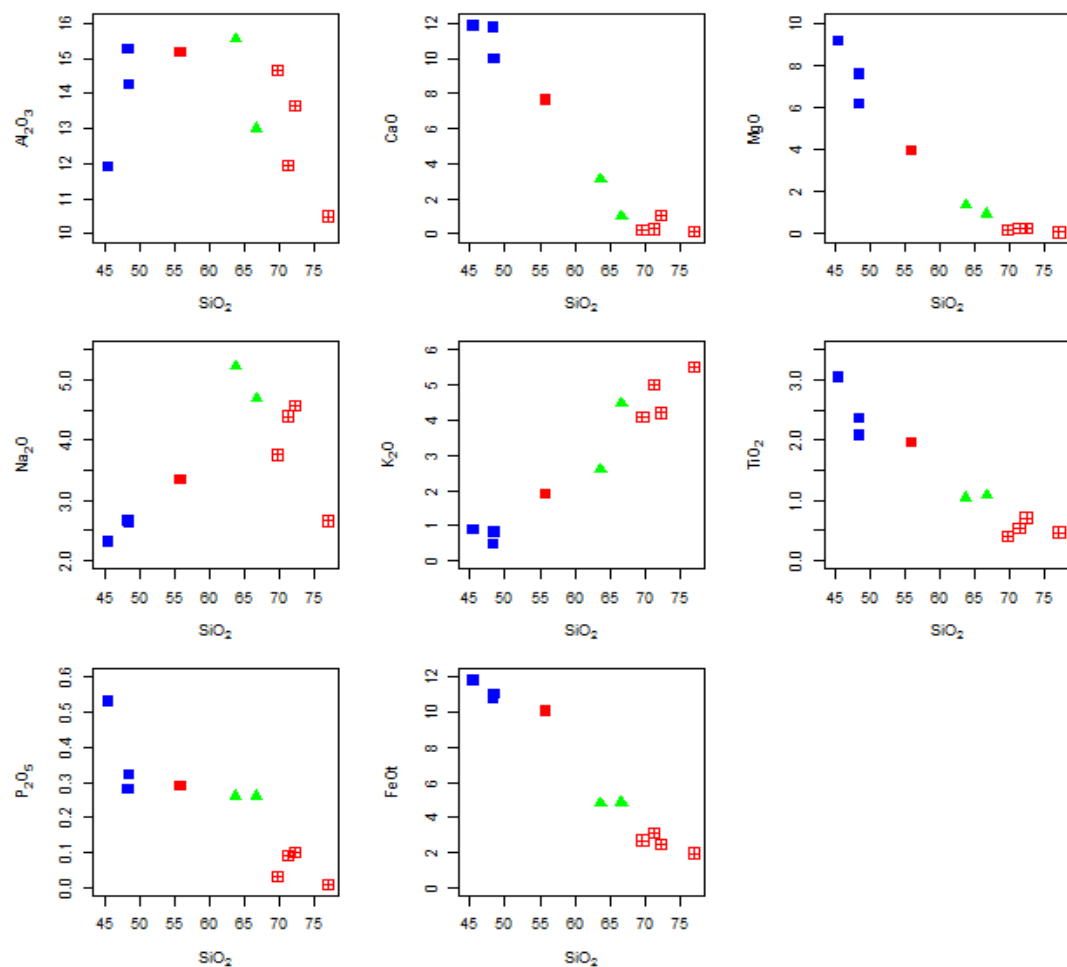


Figure 7. Harker variation diagrams of major element against SiO_2 . Symbols are the same as in Fig.6.

Trace elements

The most common incompatible trace elements are large ion Lithophile (LIL)/ (Cs, K (not shown), Rb, Eu, Ba, and Sr) and high field strength (HFS) elements (e.g. Zr, Hf, Ti, Nb, Th, U, Ce, Ta, and Pb). In Fig. 9 diagrams of incompatible trace element (ppm) variation as a function of SiO₂ demonstrate a clear trend of magma evolution. K and Rb have significant positive trend as a function of SiO₂, but Sr has a strong negative correlation trend. The variation diagrams of HFSE (U, Hf, Y (which is not shown in the diagrams below) Th, La, Zr, Nb,) against SiO₂ plotted below (Fig.9) show positive correlations variably with well-defined evolution trend. Plots of Th, U, La, Zr, Hf, and Nb as a function of SiO₂ show a

negative correlation up to ~50 wt% of SiO₂ then gradually increase and positive correlation with SiO₂ (50-70wt%), but narrow variation for those incompatible elements as silica increase, and La and U negatively correlated with SiO₂ after ~70. wt%; whereas, Pb against SiO₂ show positive correlation and Y gradually increase up to ~65wt% and in between ~65wt% - ~70 wt% sharply increase and positively correlated; whereas after > 70wt% inflected and negatively correlated. All compatible elements (such as Ni, V, Cr and Co) in the variation diagrams show negative correlation plotted as a function of SiO₂ wt %. A compatible element shows almost similar characteristics to corresponding major elements having negative correlation e.g. Fe₂O₃, MgO, CaO, and TiO₂.

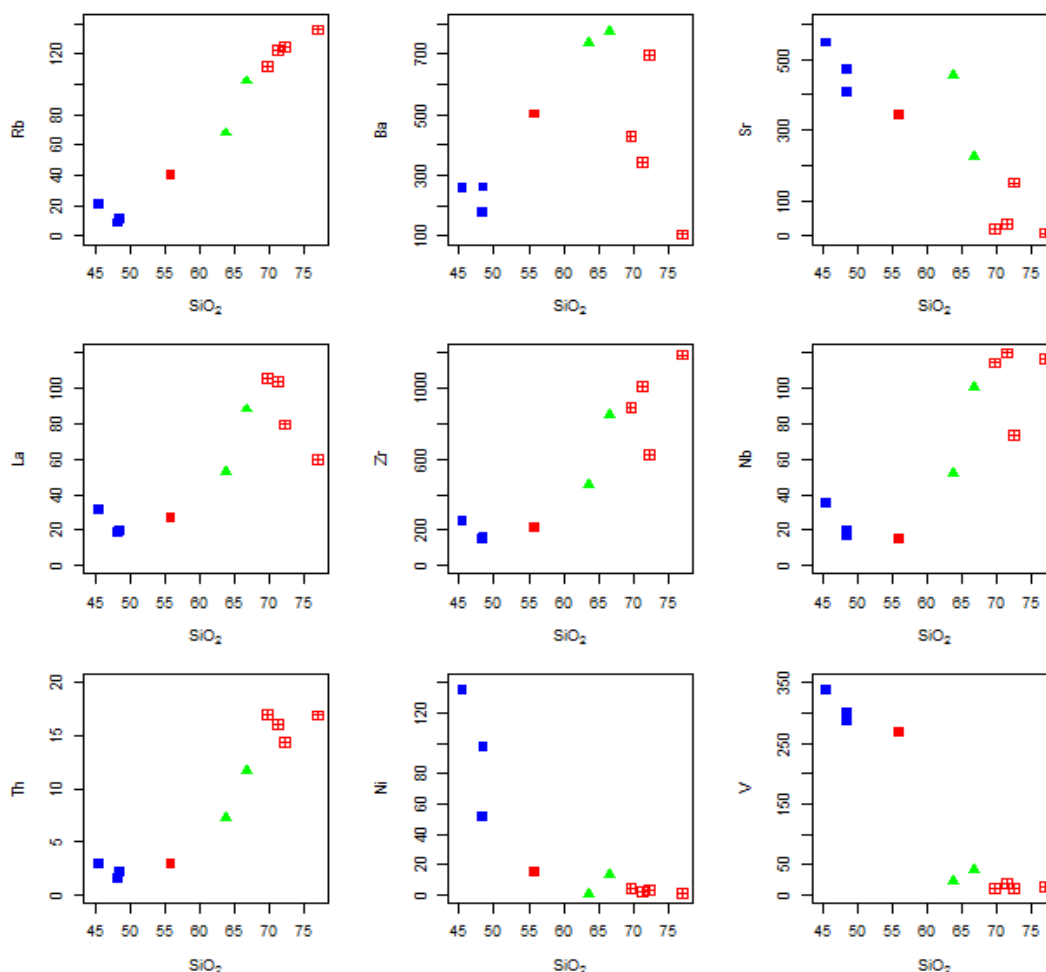


Figure 8. Harker variation diagrams of trace elements against SiO₂. Symbols are the same as in Fig.6.

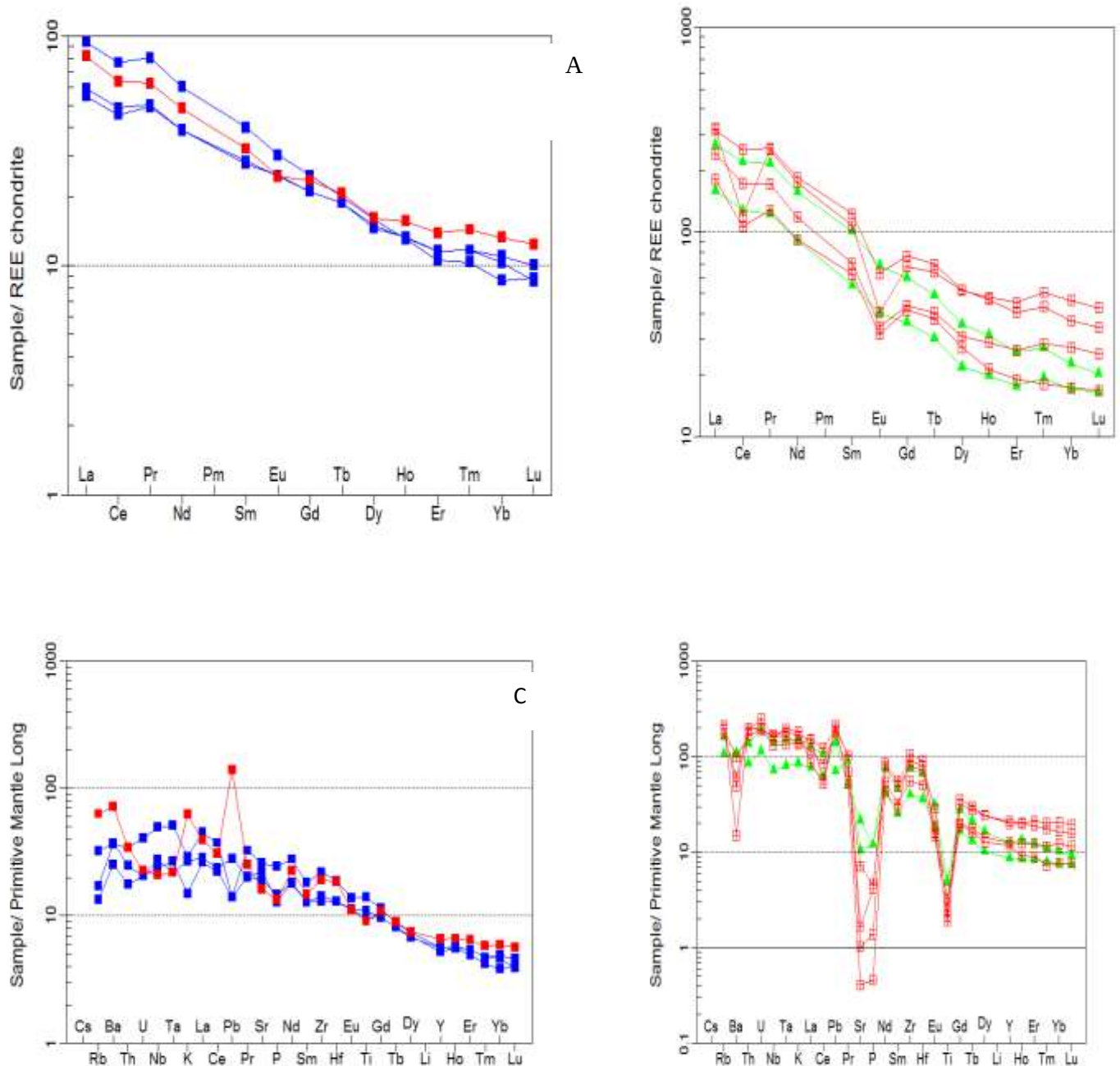


Figure.9. A,B are Chondrite normalized REE pattern of basaltic to basaltic andesite and trachydacite to rhyolite respectively of Lake Hayk analyzed samples. The normalization values are from Nakamura,(1974). C, D are Primitive mantle normalized multi-element variation diagram (Sun and McDonough, 1989) of basaltic to basaltic andesite and trachydacite to rhyolite respectively of Lake Hayk analyzed samples. Symbols are the same as in (Fig.6).

The Spider diagram of mafic to felsic rocks is plotted in fig.9 A and B. All the basaltic and rhyolitic samples are parallel and trachydacite to rhyolitic samples show sub-parallel patterns. The REE chondrite normalized diagram for samples shows a likely pattern of OIB basalt which is enriched in LREE and depleted in HREE. The

normalized value of Eu/Eu^* is 0.98 -1.01 for mafic/ basalt, 0.88-0.89 for basaltic andesite to trachydacite, and 0.48-0.65 for acidic/rhyolitic samples. Basaltic rock show slight negative anomaly of Ce and also, There is one rhyolite sample (T1S3) showing highly depleted Ce than the other and also which has higher LOI (3.07).

DISCUSSION

Geochemically associated Oligocene basaltic erupted at 31 Ma and rhyolitic volcanic rocks ranges in age from 28.5–30.8 Ma around Lake Hayk are belonging to a bimodal suite. Geochemical data, such as major and trace elements, are used to study the petrogenetic evolution of Lake Hayk flood basalt (31 Ma) and differentiation of rhyolite volcanic rocks. In this section discusses the petrogenetic progression of basalt to rhyolite volcanic rocks in detail. The examined basalt samples from Lake Hayk are characterized by wide ranges in compatible trace element contents. Ni (51-135 ppm), Cr (110-490 ppm), MgO (6.16 - 9.16) wt%. However, primary magma in equilibrium with a typical upper mantle mineral assemblage is characterized by their higher compatible trace element contents such as Ni (>400–500 ppm), Cr (>1000 ppm), and MgO content (10–15 wt. %) (Frey et al., 1978; Hess, 1992). Hence, the values of compatible element and MgO in the studied basalts are lower than primary magma suggesting that they do not represent primary magmas and have undergone fractionation of olivine and clinopyroxene. In addition to this the concentrations of basalt, basaltic andesite, trachydacite, and rhyolitic volcanic rocks contains Sr (405-546 ppm), (345ppm), (222 - 455ppm), and (8.6-151 ppm) and for V (287-337 ppm), (268 ppm) (23-41) and (10-18 ppm) respectively for each rock unit. These suggest there is less fractionation of plagioclase and Fe–Ti oxides in basalt than basaltic andesite to rhyolite (Dereje Ayalew and Gibson, 2009).

Fractional crystallization

The mineralogical modal composition of analyzed samples shows a wide range of textural and mineralogical compositions. Porphyritic basalt, aphyric basalt, rhyolite, and ignimbrite are the volcanic rocks that are petrographically examined which are stratigraphically from bottom to top. Different mineral phases of phenocrysts, microphenocrysts, and groundmass levels with different modal percentages are presented in the rock units. The phenocrysts of porphyritic basalt are olivine and clinopyroxene with a groundmass of microcrystals of plagioclase and opaque mineral (Fe–Ti oxide) which is the bottom part of the study area overlain by aphyric basalt which is composed of plagioclase, some pyroxene with a groundmass of opaque mineral and this aphyric basalt which is overlain by strongly welded rhyolite rocks unit which constitutes of alkali feldspar (sanidine), quartz with a groundmass of volcanic glass and very less opaque mineral it intercalated with less welded rhyolite, ignimbrite (composed of nepheline, rock fragments, and volcanic glass) and

with aphyric basalt in some part of the study area (mostly western part). The textural arrangements, the zoning in plagioclase, alteration effects in olivine, clinopyroxene and plagioclase, the euhedral-Subhedral-anhedral shape of phenocryst phase with variations in size, are very common in petrographic analyzed flood basalt rocks and euhedral sanidine, nepheline, and subhedral to anhedral quartz in silicic rocks. Furthermore, the analyzed volcanic rock show differences in crystal shape which implying the order of magma crystallization as euhedral-Subhedral which is the first crystallized minerals, and anhedral crystals crystallized at the last. Plagioclase is normally zoned with a compositional change from the high-temperature mineral phase at the core (Ca-plagioclase) and to low-temperature composition mineral at its rim (Na-plagioclase) and it shows complex zoning, polysynthetic twin, the inclusion of clinopyroxene, overgrowth in olivine, and sieved texture in plagioclase which is the effect of alteration. This sequence can be explained as the enclosed mineral grain olivine is assumed to be the first to crystallize followed by clinopyroxene, plagioclase, K-feldspar (sanidine), quartz, and, nepheline which support /complementary with fractional crystallization trend in geochemical variation plots as discussed below. Generally the trend which show decrease in mafic mineral and increasing of felsic mineral from flood basalt to shield volcano show the decrease of magma influx (Krans et al., 2018). Basaltic rock fractional crystallization can be accessed through the plot of SiO₂ versus major oxides (Winter, 2001; Cox, 1979). With increasing SiO₂ content MgO, Fe₂O₃, CaO, TiO₂ and P₂O₅ display negative trend implying fractionation of Mg- Fe- Ca- Ti and P-bearing minerals such as olivine and/or clinopyroxene, plagioclase, Fe–Ti oxides, and apatite respectively. Al₂O₃ positively correlated from basalt to trachydacite, which suggest the accumulation of plagioclase, and in rhyolite, suite shows a negative correlation with SiO₂ which implicates the fractionation of Al-bearing mineral of plagioclase. K₂O is a positive trend with increasing SiO₂, this testifies the accumulation of alkali feldspar (sanidine is common phenocryst in rhyolite). Na₂O show a positive correlation ~70% of SiO₂ and negative correlation after trachydacite which suggest the fractionation of Na-feldspar fractionation. The model of fractional crystallization for some compatible (Ni and Cr) and incompatible elements (Fig.11) of calculated trends show sharp decrease in Ni and Cr is slight as compared with Ni; whereas incompatible elements and slightly increasing up to %F = 0.4 and sharply increase up to %F = 0.

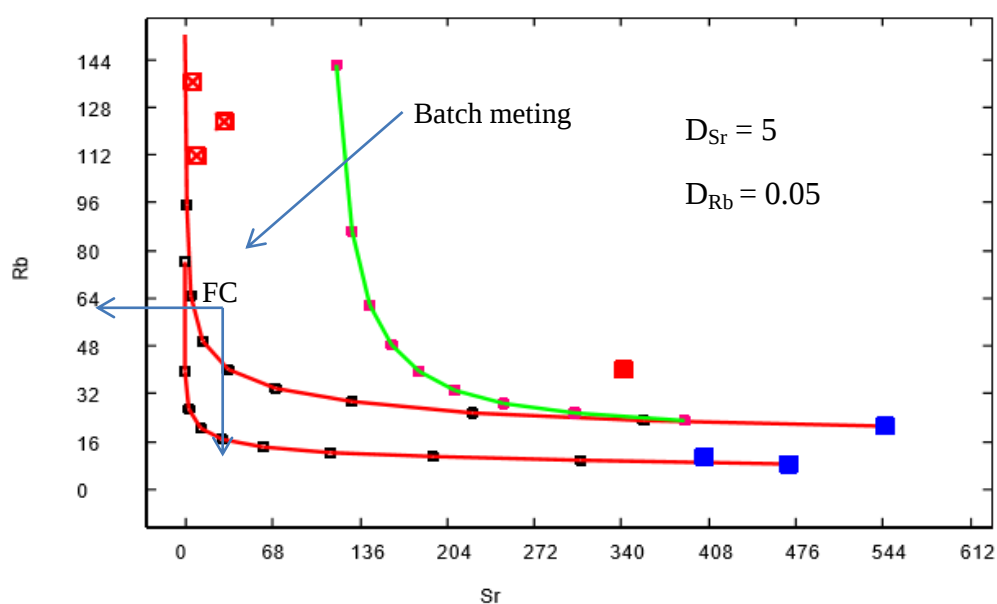


Figure 10. Variation diagram of fractional crystallization (FC) and batch melting model of compatible (Sr) vs. incompatible (Rb) trace element for lake Hayk volcanic rocks. Basalt T1S1 which is less evolved rock ($SiO_2 = 45.5$) has been assumed as the starting composition for both fractional crystallization and batch melting. The partition coefficient for the model is $5 = Sr$ and $0.05 = Rb$ (after Ewart and Griffin, 1994). Symbols the same as in (Fig.6.).

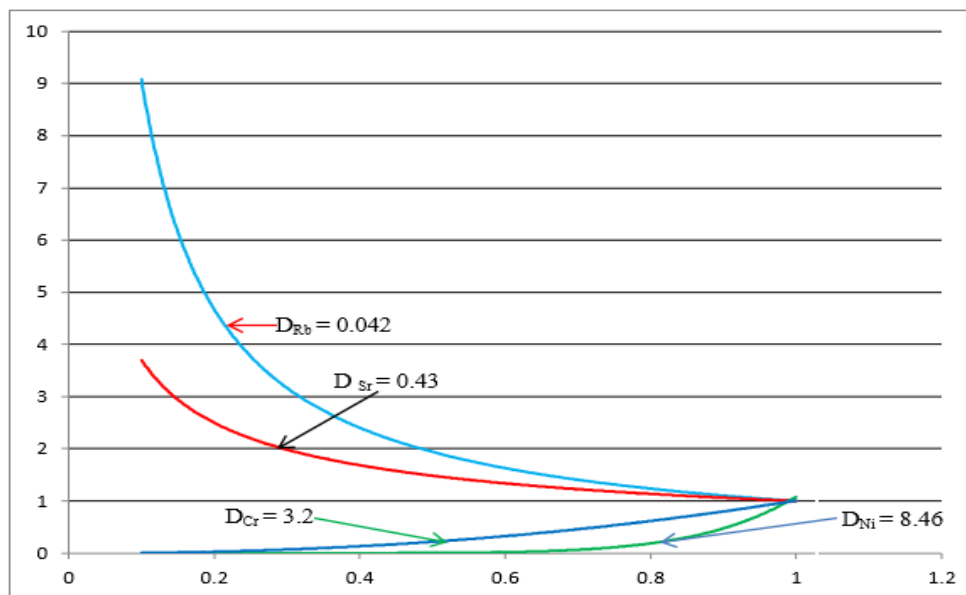


Figure 11. Fractional crystallization model for the most primitive basaltic (T1S1) rock of Lake Hayk. The weight fraction of residual liquid $F\%$.

Table 2. Major element calculated mass balance.

	From	To	Chemical compositions				%Solid	R
	T2S3	T7S1	Olivine	clinopyroxene	Plagioclase	Opa	Fract.	
SiO ₂	48.4	55.9	40.66	47.15	50.71	0		0.003
TiO ₂	2.08	1.96	0	3.38	0	24.91		0.397
Al ₂ O ₃	15.25	15.15	0	4.78	30.67	1.46		0.037
FeO _{tot}	10.71	10	23.44	10.99	0.54	71.73	60.51	-0.156
MnO	0.18	0.17	0.35	0.19	0	0.64		-0.027
MgO	6.16	3.94	34.88	11.86	0.18	1.24		0.069
CaO	11.75	7.61	0.76	21.09	14.72	0		-0.082
Na ₂ O	2.66	3.34	0	0.54	3.05	0		-0.412
K ₂ O	0.45	1.88	0	0	0.14	0.01		0.34
P ₂ O ₅	0.28	0.29	---	---	---	---		-0.169
Fract. Solid wt. %			-42.5	-32.25	-21.74	-3.48	ΣR² =	0.51
	From	To	Chemical compositions				%Solid	R
	T7S1	T3S6	Olivine	Clinopyroxene	Alkali	Opaque	Fract.	
SiO ₂	55.9	66.7	39.87	49.06	51.92	0.51		-0.093
TiO ₂	1.96	1.08	0.03	0	0.77	50.02		-0.311
Al ₂ O ₃	15.15	13	0	32.14	1.85	0		0.547
FeO _{tot}	10	4.83	14.06	0.27	32.19	46.37	43.41	0.025
MnO	0.17	0.24	0.22	0	0	1.44		-0.014
MgO	3.94	0.92	45.38	0.2	0	0.46		0.271
CaO	7.61	1	0.25	15.38	0	0.71		-0.583
Na ₂ O	3.34	4.69	0.04	2.57	12.86	0		0.58
K ₂ O	1.88	4.48	0.01	0.17	0.19	0		0.27
P ₂ O ₅	0.29	0.26	---	---	---	---		-0.169
Fract. Solid wt. %			-14.61	-57.99	-19.67	-7.72	ΣR² =	1.3
	From	To	Chemical compositions				%Solid	R
	T3S6	T3S4	Olivine	Clinopyroxene	Sanidine	Opaque	Fract.	
SiO ₂	66.7	71.3	29.84	48.46	68.31	0		0.26
TiO ₂	1.08	0.54	0	0.46	0	22.57		-0.15
Al ₂ O ₃	13	11.95	0	0.2	18.74	0.28		-0.24
FeO _{tot}	4.83	3.1	64.69	28.6	0.38	74.56		0.312
MnO	0.24	0.08	4.73	1.7	0	2.44	29.47	-0.115
MgO	0.92	0.25	0.43	1.77	0	0.12		0.017
CaO	1	0.28	0.31	18.4	0.07	0		-0.203
Na ₂ O	4.69	4.4	0	1.42	9.17	0		0.651
K ₂ O	4.48	5.02	0	0	3.33	0.01		-0.325
P ₂ O ₅	0.26	0.09	---	---	---	---		-0.207
Fract. Solid wt. %			-5.89	-9.6	-77.74	-6.77	ΣR² =	0.872

In order to test fractional crystallization hypothesis major elements mass balance calculated (Table 2). Major element modeled based on the calculation of mass balance by following steps from the most mafic to felsic rocks (Stormer and Nicholls, 1978). The mineral chemistry of basaltic rock is taken from northwestern Ethiopia flood basalt (Pik et al., 1998) and basaltic andesite to rhyolite rock is from Boseti volcanic complex of main Ethiopian rift (Ronga et al., 2009), this choice is due to minerals of rift margin mineral chemistry is not available. The calculated mass balance indicates that the evolution modeled from basalt (T2S3) to basaltic andesite (T7S1) by 60.51% solid fractional crystallization with sum of the calculated residual (R₂) of 0.51. The evolutions from basaltic andesite (T2S3) to trachydacite (T3S6) by 43.41% of solid fractional crystallization with sum of the calculated residual (R₂) of 1.3 and trachydacite (T3S6) to rhyolite (T5S2) by 29.47% of solid fractional crystallization with sum of the calculated residual (R₂) of 0.8721. The variation diagram of major element against silica of the

calculated composition and the analyzed/parent composition fit linear to curvilinear line which suggest the fractional crystallization is the major process.

Crustal contamination

The studied samples of lake Hayk basalts, have a positive anomaly of Nb and Ta (with exception of a slight trough in a sample (T7S5 slight trough at Ta)) and trachydacite to rhyolite show slight trough at Nd and positive spike at Ta on the primitive mantle normalized multi-element variation diagrams (McDonough and Sun, 1989) as illustrated in the previous section (fig.10B), and this testifies there is minimal crustal contamination in basalt (with exception of T7S5) than trachydacite to rhyolite (Rollinson., 1993). Basaltic andesite (T7S1) show a negative anomaly of Nb and Ta with high Rb and Th which suggest that the involvement of upper crust or contaminated by upper crust (Dereje Ayalew et al., 2019). Basalt have low Rb/Nb = 0.5 - 0.58, La/Nb = 0.88 - 1.06 and higher TiO₂ = 2.08 - 3.04, basaltic andesite

have higher $Rb/Nb = 2.7$, $La/Nb = 1.81$ and low $TiO_2 = 1.96$ and rhyolite $Rb/Nb = 0.97-1.69$, $La/Nb = 0.51 - 1.08$ and lower $TiO_2 = 0.41 - 0.71$ this suggest that there is more crustal involvement in basaltic andesite as compared to basalt and rhyolite; whereas lower in basalt as relative to rhyolite (Rollinson, 1993; Dereje Ayalew and Gezahegne Yirgu, 2003). Further, MgO against incompatible trace elements such as Th, La, and U

have a negative correlation as shown below (Fig.12), this testifies that there is insignificant continental crust involvement (Zhang et al., 2009). Furthermore, Ce/Pb has a positive correlation with MgO (fig.12B), this suggest that crustal contamination is minimal towards basaltic rock which have maximum concentration MgO and Ce/Pb, whereas felsic rocks are contaminated (Dereje Ayalew et al., 2016; Haldera et al., 2020).

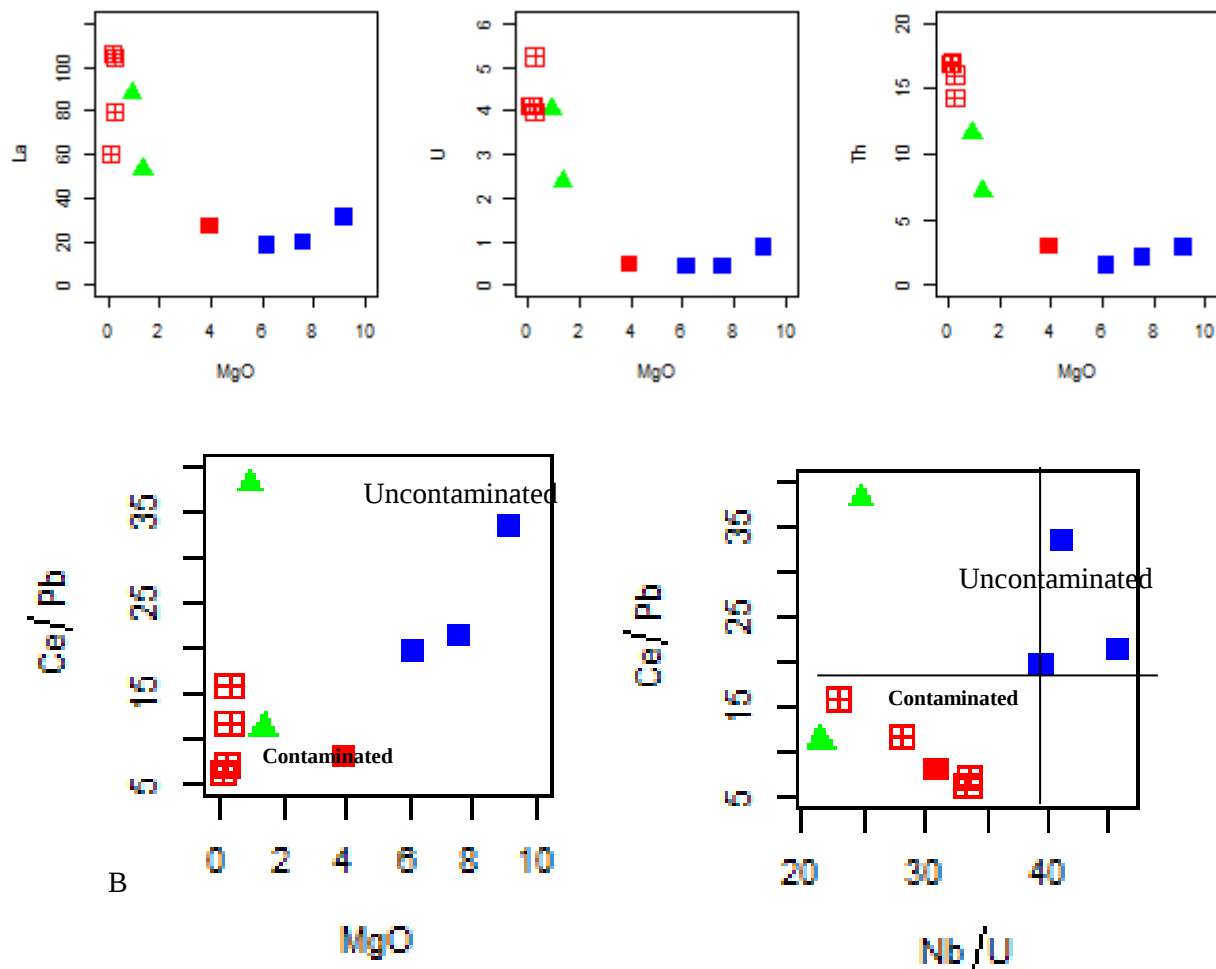


Figure 12. MgO against for selected incompatible trace elements (La, U, Th) diagrams of the Lake Hayk volcanic rocks. Symbols same as in fig.6.

Magma generation and source rock characterization

The basalt rock unit from lake Hayk is characterized by low CaO/Al_2O_3 ratios (0.71–0.95) and in primitive mantle normalized multi-element variation diagram (Sun and McDonough, 1989) HREE relatively no fractionation/flat patterns with ($TbN/YbN = 1.75-2.33$) (fig.9c and). This value most likely suggests a mantle source containing spinel rather than garnet and the slight fractionation of HREE show the slight effect of garnet (Dereje Ayalew et al., 2016). Further, the basalts in the studied area have positive anomaly

of Ba and a negative anomaly of K (positive anomaly at (T7S5)) in comparison to other primitive mantle normalized multi-element variation diagrams (Fig.9 C). Such positive and negative anomaly of Ba and K respectively, testify related to amphibole and phlogopite (for K positive sample (T7S5) in the mantle source (e.g., Jung et al., 2005, 2012; Furman and Graham, 1999; Dereje Ayalew et al., 2006; Rooney et al., 2014). Additionally, basalt samples have $Na_2O > K_2O$, such observation suggests that the source is formed through melting of the amphibole-bearing mantle (Rosenthal et al., 2009). Based on the above

observation the lake Hayk basalts were derived from the melting of amphibole-bearing spinel peridotite sources rather than Phlogopite bearing source (Dereje Ayalew et al., 2016). Amphibole may also occur in the lithospheric mantle, because it is not stable in the sub-lithospheric mantle source of asthenosphere or mantle plume (Class and Goldstein, 1997).

Petrogenesis of rhyolites

To explain the petrogenesis of rhyolitic lava and associated rhyolitic tuff, rhyolitic ignimbrite, glassy rhyolite, and pumice (60-81 wt% SiO₂), several petrogenetic models describing the origin of silicic volcanic rocks in mafic LIPs have been proposed. These include: (i) fractional crystallization of parental mafic magmas (e.g., Ayalew et al., 2002; Mahoney et al., 2008; Melluso et al., 2008; Natali et al., 2011; Cucciniello et al., 2019) (ii) re-melting of underplated basaltic intrusions (Cleverly et al., 1984; Harris and Erlank, 1992; Miller and Harris, 2007), (iii) assimilation of crustal material by parental mafic magmas and simultaneous fractional crystallization (AFC) (Chazot and Bertrand, 1993; Baker et al., 2000; Melluso et al., 2001; Dereje Ayalew and Gezahegne Yirgu, 2003; Peccerillo et al., 2003; Mahoney et al., 2008; Sheth and Melluso, 2008; Cucciniello et al., 2011), and (iv) melting of continental crust (Tian et al., 2010; Shellnutt et al., 2012; Huang et al., 2015). Significant fractional crystallization produces rhyolites with steep negative slopes in bivariate major oxides plots and a modest to no Nb-Ta anomaly (Haldera et al., 2020), whereas strong crustal input produces rhyolites with significant

Nb-Ta negative anomalies. Hence, the rhyolites of Lake Hayk have steep negative correlation in bivariate major oxides such as MgO, Fe₂O₃, TiO₂, and CaO against SiO₂, as well as a positive anomaly of Ta with a slight Nb trough, implying that rhyolites formed by fractional crystallization of mafic magma with insignificant crustal contribution. Further, rhyolites formed by crustal melting have an Rb/Nb ratio >10; whereas, the analyzed rhyolitic samples Rb/Nb < 1.69 this indicates that partial melting of the continental crust did not contribute to the formation of rhyolites ((fig.14b). Plot of TiO₂ vs. SiO₂ (Fig.7) for volcanic rock of lake Hayk shows negative correlations and a positive correlation of HFSE such as Zr vs. Rb, Zr vs. Hf, Zr vs. Nb, and Zr vs. Ta forming a single trend that show the genetic relationship or co-magmatic nature of the basaltic and rhyolitic rocks around Lake Hayk area (Haldera et al., 2020). This suggests that rhyolites are related to the basalts by Fractional crystallization as illustrated in (fig.10). The Zr/Nb ratio of basalt to rhyolite vary between 7.01-8.76 and basaltic andesite (T7S1) and one rhyolite (T1S2 EA) sample Zr/Nb = 14.43 and 10.17 respectively. The little variation of Zr/Nb ratios in basalt to rhyolites indicates that fractional crystallization has been the major process in their history of petrogenesis (Barberi et al., 1975). The only exceptions are basaltic andesite and one rhyolite sample, which have a different array (high Zr/Nb of T2S1 EA = 10.17 and T7S1 - 14.43), suggesting derivation from a depleted plume head source (Ferguson et al., 2010).

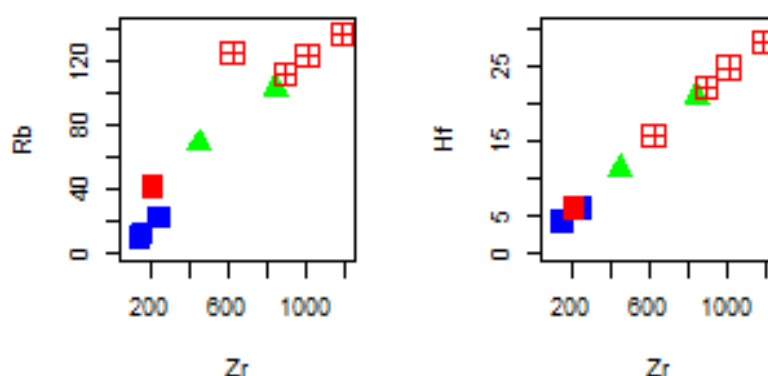


Figure 13. Incompatible against incompatible trace elements variation diagrams (ppm) of Lake Hayk, analyzed samples. Symbols the same as in (Fig.6).

Rhyolites that have (Gd/Lu)N ratio (>2.2), suggest that the parental melts may have been generated at higher depths in the garnet stability field (Haldera et al., 2020). The ratio of (Gd/Lu)N of rhyolite in the study area varies between 1.59 - 2.48. Hence the rhyolite sample (T3S4 GdN/LuN = 2.24 and T1S1 EA GdN/LuN = 2.48) from Lake Hayk which has (Gd/Lu)N ratio (>2.2) implicate that parental melts may have been generated at higher depths in the garnet stability field and the other two sample (T5S2 Gd/Lu)N = 1.72 and T1S3 GdN/LuN = 1.59) which have (Gd/Lu)N ratio (<2.2) are from spinel stability field. The Rb against Sr bivariate plot of rhyolite (Fig. 14 A), shows a large variation in Sr (8.6 - 151ppm) with limited variation in Rb (111.5 - 136ppm) abundances. Accordingly, such highly incompatible against

highly compatible trace element trend suggests that the rhyolite of the Lake Hayk area related to erupted basaltic magmas by the fractional crystallization process than partial melting (Dereje Ayalew et al., 2019; Haldera et al., 2020).

Eventually, the stratigraphy and petrography which supports the above interpretation because stratigraphically basaltic rock units are unconformably overlain by rhyolitic rock unit of different age separated by paleosol, and petrographically from bottom to top are as follow; olivine, clinopyroxene, plagioclase, (very low), k-feldspar (sanidine) and quartz which show they originated within the same magma chamber and that the rhyolitic volcanic rocks were derived by fractional crystallization of the mafic/basaltic magma.

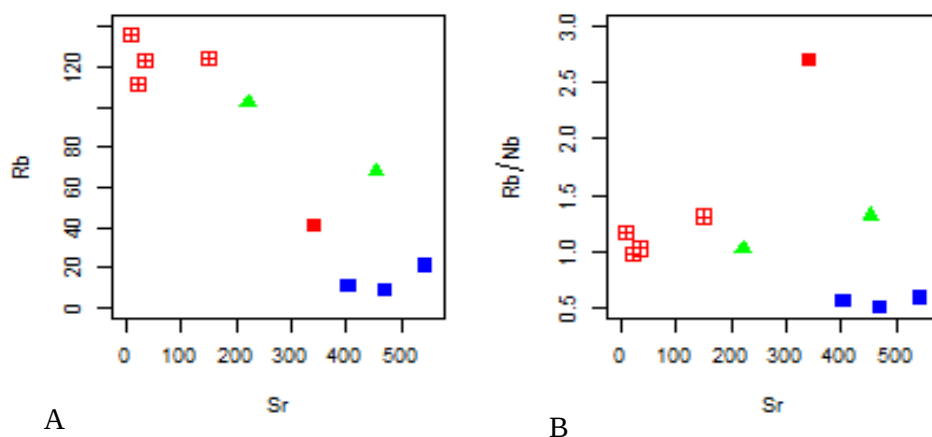


Figure 14. Variation diagrams of (A), Sr vs. Rb and (B), Sr vs. Rb/Nb of Lake Hayk area volcanic rocks. Symbol the same as (fig. 6).

Comparison of lake hayk primary data with previous works

The Lake Hayk area has a bimodal composition with scarce intermediate rock unit. As a comparison, four samples (two basalt, one rhyolite, and one trachyte) from the Gugufu shield volcanics (Kiffer et al., 2004) and eleven samples from Wegel Tena, Kobo, and Hara (Ayalew et al., 2019). The outcomes of the samples are generally consistent with the primary data studied in the current study approach. Ten samples from Lake Hayk and fifteen samples from previous samples were combined for comparison in various diagrams. Both major and trace Variation Elements diagrams of the investigated primary basalts of Lake Hayk plotted against

previous basalts and primary felsic sample plotted against previous felsic sample demonstrate that the signatures are most likely the same. When compared to previously reported Gugufu shield and marginal Hara, Kobo and Wegel Tena volcanic suites of geochemical data obtained by previous investigations, the current study does not demonstrate a significant difference in composition. With the exception one basalt from Gugufu, almost all samples from Lake Hayk fall near the transition between alkaline and sub alkaline fields. As shown in (fig.15) below, the other basalt, trachyte, and rhyolite samples are alkaline, as well as one trachydacite, basaltic trachyte, and rhyolite sample from Lake Hayk, Hara, and Kobo.

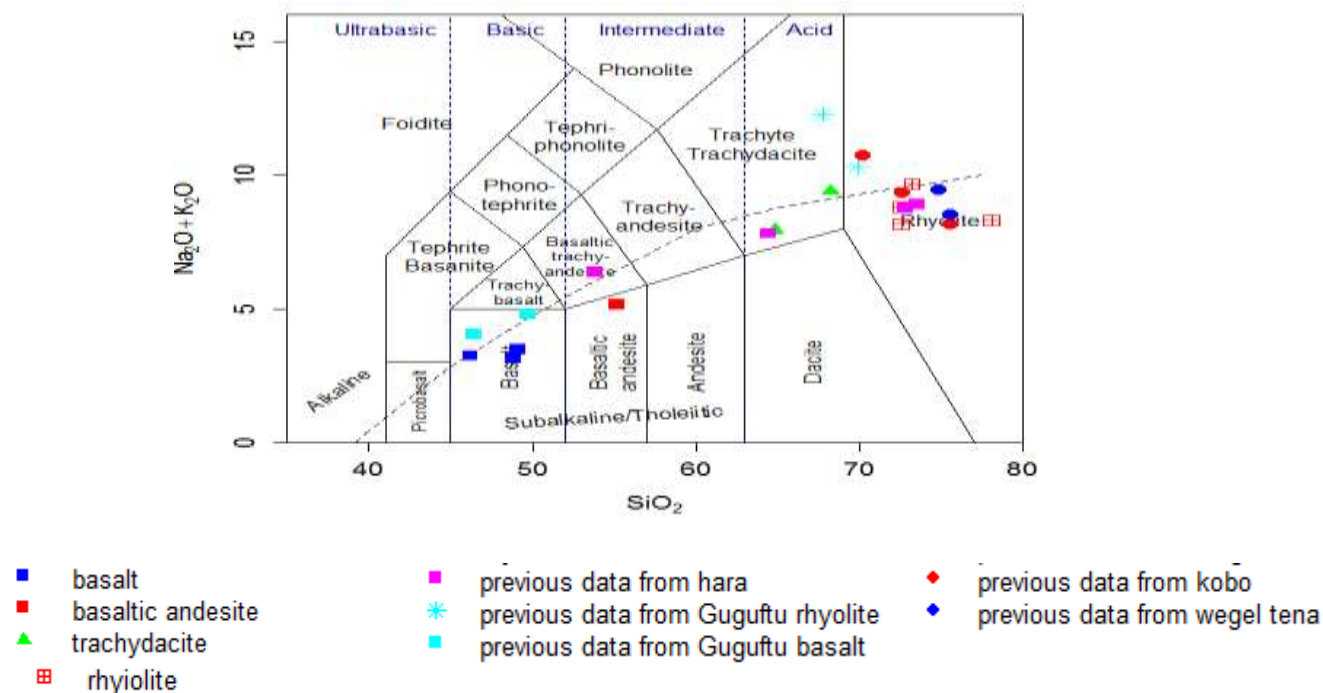


Figure 15. TAS classification diagrams of Lake Hayk flood basalts and rhyolite for composition of Lake Hayk basalt to rhyolite from the previous works.

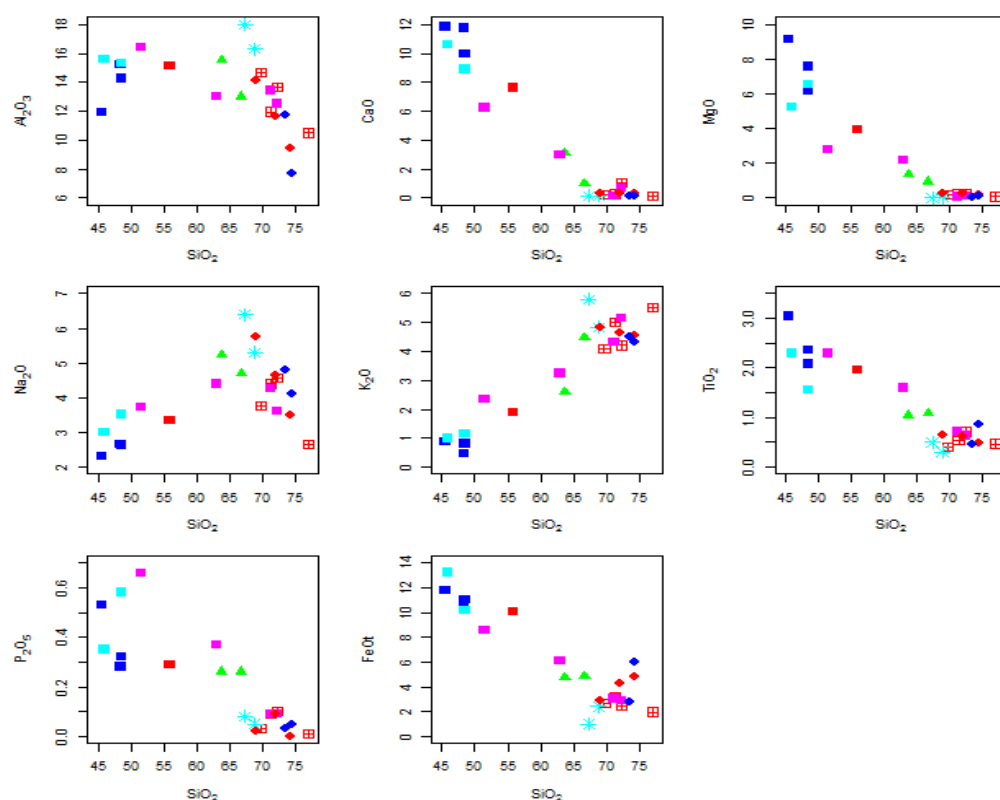


Figure 16. Variation diagrams of major element contents as a function of SiO_2 for comparison of Lake Hayk volcanic rock with previous work. Symbol the same as (fig. 15).

The variation diagram of major element against SiO_2 (fig.16) for both primary and previous data show the same trend (primary basalt with previous basalt, primary basaltic andesite with previous basaltic andesite and primary felsic with previous felsic). Both primary and previous samples show steep negative trend of these compatible trace elements and the accommodating major elements (Fe, Mg, Ca); show pronounced evolution trend starting from basalt-intermediate rhyolite suites. The available primary data show Lake Hayk rhyolites show similar signature of major and trace element abundance with Hara, kobo, and Wegel

Tena silicics analyzed in previous works. Basaltic systems show Sr depletion in response to the fractionation of plagioclase and or crustal assimilation (Haliday et al, 1991); and Rb increases corresponding to the more or less vertical trend (see Fig. 6.10). Incompatible trace elements like Sr drops to very low concentrations both in Hara, Gugufu and Lake Hayk felsic rocks; whereas strongly incompatible trace elements like Rb, Nb, and La showing very high concentration. Previous work results have presented that, the felsic rocks of Hara formation display similar patterns with Lake Hayk felsic rock.

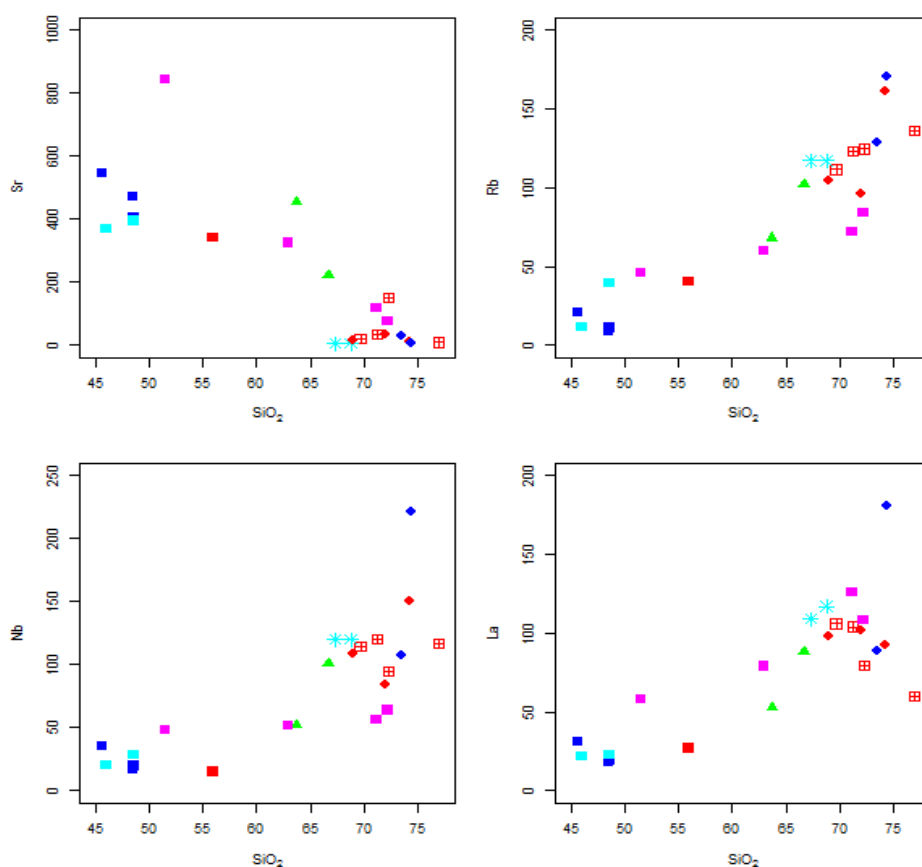


Figure 17. Trace element variation diagrams show the variation of incompatible trace elements against SiO_2 for composition of Lake Hayk basalt to rhyolite to the previous work. Symbols are the same as in Fig.15.

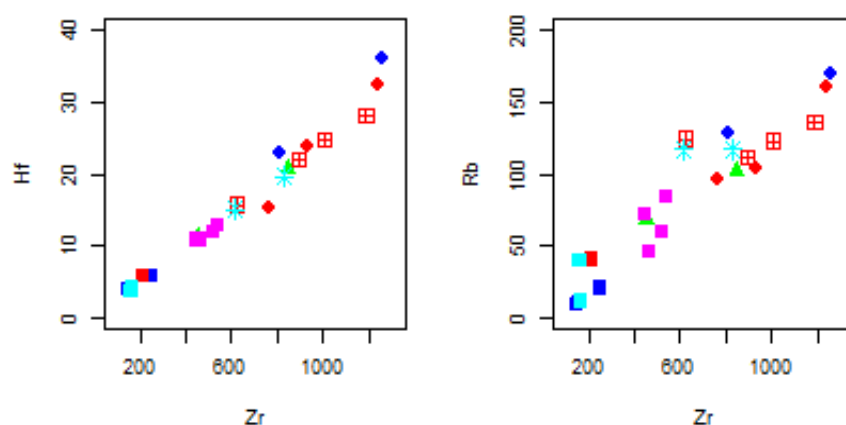


Figure 18. plots of Rb against Zr variation diagram which shows comparison of Lake Hayk volcanic rock with previous work. Symbol the same as fig .15.

Volcanic rock of lake Hayk, Wegel Tena, Gugufu and Hara plots of TiO_2 vs. SiO_2 (Fig.16) shows negative correlations and a positive correlation of HFSE such as Zr vs. Rb, Zr vs. Hf, Zr

vs. Nb, and Zr vs. Ta forming a single trend that show the genetic relationship or co-magmatic nature of the basaltic and rhyolitic rocks those area.

Table 3. 1: Trace element and TiO_2 of the Lake Hayk area volcanic rocks compared with other marginal rhyolites and Gugufu shield volcano.

	Rock type based on TSA classification	Rb/Nb	La/Nb	TiO_2	Source of data
Hara	Basaltic trachyte	0.96	1.21	2.29	Ayalew et al.,(2019)
	Trachydacite	1.16	1.52	1.60	
Gugufu	Rhyolite	1.13-1.19	1.49-2.25	0.57 - 0.71	Kieffer et al.,(2004)
	Basalt	0.57-1.41	0.81-1.07	1.56 - 2.29	
	Trachydacite	0.98	0.91	0.51	
	Rhyolite	0.98	0.98	0.53	
Kobo	Rhyolite	0.95 - 1.14	0.91-1.21	0.63 -0.65	Ayalew et al.,(2019)
Wegel Tena	Rhyolite	0.77-1.21	0.81- 0.83	0.46 - 0.86	Ayalew and Yirgu,(2003)
Lake Hayk	Basalt	0.5-0.58	0.88-1.06	2.08-3.04	present study of this work
	Basaltic andesite	2.70	1.18	1.96	
	Trachydacite	1.01-1.32	0.88-1.02	1.04-1.08	
	Rhyolite	0.97-1.69	0.51-1.08	0.41-0.71	

Furthermore, Lake Hayk rhyolites unit have the same Rb/Nb (0.51-1.08) and La/Nb (0.41-0.71) ratios to that of Kobo, Wegel Tena, and Gugufu rhyolite rock units; whereas relatively lower than Hara rhyolite rock unit, which implicate that in Kobo, Gugufu, and Lake Hayk rhyolites unit there is only limited crustal material was involved in their genesis or minimal amount of crustal contamination than Hara rhyolite. Generally this work supports the previous work.

CONCLUSIONS

The following conclusions have been drawn based on the, field observations, petrographic and geochemical data obtained in this study:

➤ Major and trace element geochemistry data testify that volcanic rocks of Lake Hayk are a bimodal suite, with scarce intermediate composition. The Oligocene basalts and rhyolite rocks are transitional to tholeiitic/sub alkaline

affinity and rhyolite rocks are peraluminous (exceptionally one rhyolite rock sample (T3S4) is peralkaline).

➤ There is minimal to no crustal contamination in basalt (with exception of T7S5) relatively to trachydacite to rhyolite.

➤ Basaltic andesite shows evidence of involvement of upper crust or contaminated by upper crust.

➤ The source for volcanic rock from lake Hayk area is the amphibole-bearing mantle. Parental melts for two rhyolites may have been generated at higher depths in the garnet stability field and for two other samples (T5S2 and T1S3 from spinel stability field).

➤ The parallel pattern of basalt and rhyolite in chondrite normalized REE pattern, Nb, Hf, Ta, and Rb against Zr, and flood basalt and rhyolitic flow range in age from 28.5 to 31 Ma do to these evidences basaltic to rhyolitic rocks generated from the same magma source/ co-magmatic nature.

➤ The rhyolitic rock units of Lake Hayk area are formed by fractional crystallization of mafic parental magma rather than assimilation of continental crust.

➤ Comparison of geochemical content Lake Hayk bimodal volcanic rock with the plateau formations shows that basalts have a similar geochemical pattern with HT1 type continental flood basalt whereas the trace element ratios of the Lake Hayk area rhyolites agree with Kobo, Gugufu, and Lake Hayk rhyolites unit.

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