

Research paper

Cretaceous alkaline volcanism in south Marzanabad, northern central Alborz, Iran: Geochemistry and petrogenesis



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ABSTRACT

The alkali-basalt and basaltic trachy-andesites volcanic rocks of south Marzanabad were erupted during Cretaceous in central Alborz, which is regarded as the northern part of the Alpine-Himalayan orogenic belt. Based on petrography and geochemistry, en route fractional crystallization of ascending magma was an important process in the evolution of the volcanic rocks. Geochemical characteristics imply that the south Marzanabad alkaline basaltic magma was originated from the asthenospheric mantle source, whereas the high ratios of $(La/Yb)_N$ and $(Dy/Yb)_N$ are related to the low degree of partial melting from the garnet bearing mantle source. Enrichment pattern of Nb and depletion of Rb, K and Y, are similar to the OIB pattern and intraplate alkaline magmatic rocks. The K/Nb and Zr/Nb ratios of volcanic rocks range from 62 to 588 and from 4.27 to 9 respectively, that are some higher in more evolved samples which may reflect minor crustal contamination. The isotopic ratios of Sr and Nd respectively vary from 0.70370 to 0.704387 and from 0.51266 to 0.51281 that suggest the depleted mantle as a magma source. The development of south Marzanabad volcanic rocks could be related to the presence of extensional phase, upwelling and decompressional melting of asthenospheric mantle in the rift basin which made the alkaline magmatism in Cretaceous, in northern central Alborz of Iran.

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1. Introduction

Plutonic or volcanic alkaline rocks are typically associated with continental rifting or intraplate continental and oceanic settings that typically related to the partial melting of asthenospheric mantle (McKenzie and Bickle, 1988; White and McKenzie, 1989; Wilson and Downes, 1991; Ernst et al., 2005; Ganguly, 2005; Lustrino and Carminati, 2007).

Fractional crystallization, magma mixing and crustal contamination are the main processes in the evolution of magmas (Best, 1970; Wilson, 1989; Hall, 1996; Rollinson, 1998; Mc Birney, 2006). The effect of these processes in magmatism in orogenic belts, could be cleared by geochemistry and petrology studies.

Marzanabad area in northern part of central Alborz is a part of Alpine-Himalayan orogenic that formed in Cretaceous with well developed sub-marine alkaline volcanic sequences. The volcanism associated with Barremian–Aptian and Cenomanian limestone layers clarified Cretaceous age for volcanic rocks (Cartier, 1971; Sussli, 1976; Vahdati Daneshmand and Nadim, 2001), while there are no corresponding radiometric ages available.

In Marzanabad area, the outcrops of volcanic rocks, allow us to investigate the Cretaceous volcanism of Alpine Himalayan orogenic belt in Iran. Little is known on Paleozoic and Mesozoic igneous activity in central Alborz, whereas only few aspects are known for the Cenozoic activity. However, neither whole-rock geochemistry nor microprobe or isotopic data have been published for the volcanic rocks of Marzanabad area. In this study, major and trace element, microprobe and Sm-Nd isotopic data are used to constrain the role of fractional crystallization and mantle source in the petrogenesis of alkaline magma from Marzanabad area, central Alborz.

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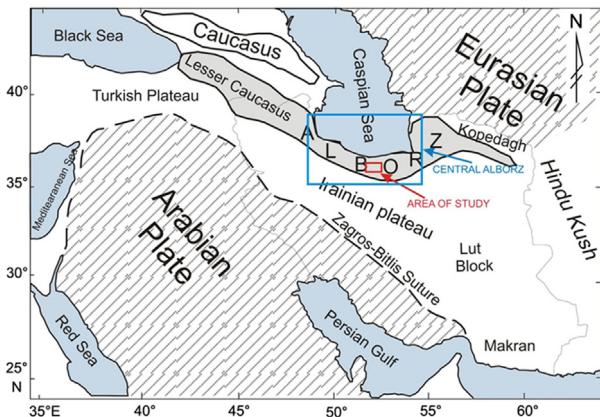


Figure 1. General tectonic map of Iran with Arabian and Eurasian plate and the location of studied area in central Alborz zone.

Results of this study may help to have a better understanding of Cretaceous alkaline magmatism that produced the alkaline volcanic rocks in Alpine Himalayan orogenic belt in central Alborz.

2. Geodynamics background

Central Alborz was a part of the Gondwana plate in early Paleozoic. It separated from Gondwana during Ordovician to Silurian and then collided with Eurasian plate in Triassic, causing the Paleotethys Ocean closure to the north, and the formation of the Neotethys Ocean to the south (Stocklin, 1974; Berberian and King, 1981; Stampfli et al., 1991). After the Triassic collisional event, along both sides of the Paleotethys Ocean, intracontinental compressions were initiated and accompanied by deposition of coal bearing Shemshak Jurassic formation (Berberian, 1983). Compressional tensions coupled with some regional extensional pressures, which were caused by the convergence of Arabian and Eurasian plates (Zanchi et al., 2006).

In central Alborz, Mesozoic era began with the deposition of detrital carbonate sediments which is continued with deposition of Shemshak Formation in late Triassic. In the studied area, extensional phases were started in upper Triassic associated with alkaline igneous activity. This alkaline magmatic phase can be assumed as an evidence for intracontinental tectonic setting related to a rift system in central Alborz during late Triassic (Furon, 1941; Steiger, 1966; Taraz, 1974; Annells et al., 1975; Nabavi and Seyed emami, 1977; Kristan-Tollmann et al., 1979; Berberian and King, 1981; Berberian, 1983; Völlmer, 1987; Faurevelet and Eftekhar Nezhad, 1992; Sabzehei, 1993; Brunet et al., 2003; Seyed emami, 2003; Shahidi, 2005, 2008; Nazari and Shahidi, 2011).

In central Alborz, extensional movements started at the same time with Rhaetic rift volcanism and deposition of coal-bearing Shemshak Formation, in Mesozoic era (Berberian, 1983; Nazari et al., 2004). The extensional phases developed some regional rifts which caused volcanism and plutonism activities in central Alborz (Berberian, 1983).

Soffel and Förster (1984) believed that the extensional movements lead to separation of the central Iranian plate from Eurasia plate, during Jurassic. Lithospheric ruptures, tensions and extensions with ascending of asthenospheric plumes and their partial melting caused the development of the rift system in Alborz zone from middle Jurassic to lower Cretaceous. Extension and the rift development could not persist more than few million years since the tectonical movements of Maastrichtian (Laramide) terminated the extension period.

3. Regional geology

Alborz Mountains are a geological structural zone in north of Iran considered as part of the northern margin of the Alpine-Himalayan orogenic belt (Fig. 1). Alborz block is connected to Caucasus Mountains in the northwest and is bounded by Hindu Kush Mountains in the east (Zanchi et al., 2006), that was part of the Gondwana plate in early Paleozoic. During Ordovician to Silurian, Alborz separated from Gondwana and finally collided with the

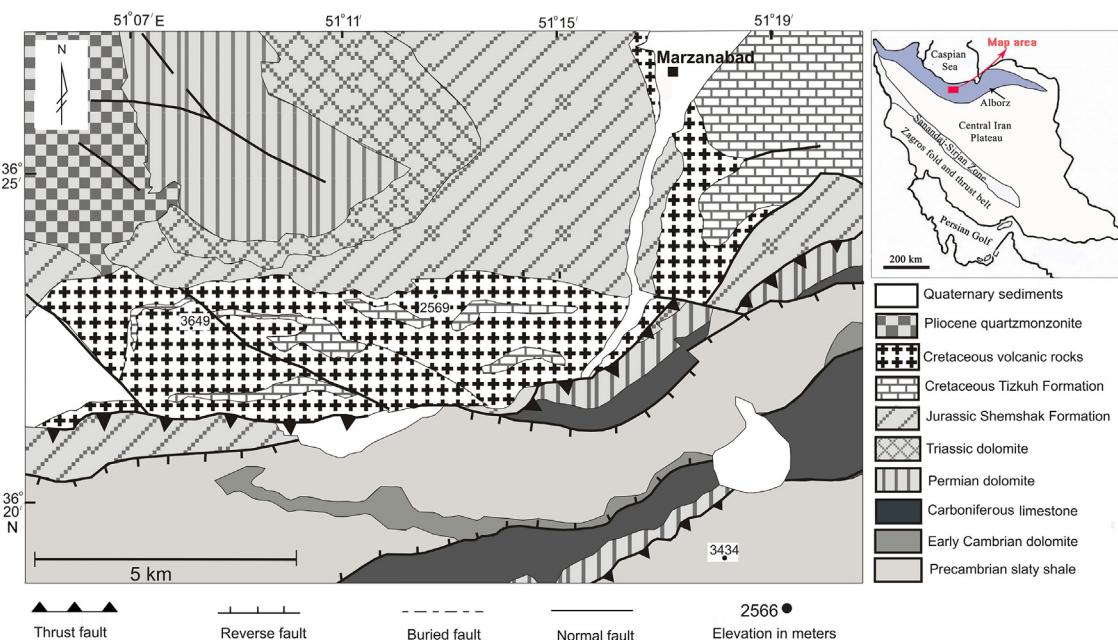


Figure 2. Geological map of the study area with the position of analyzed samples. Modified from the 1:100,000 Marzanabad geological map of Iran.

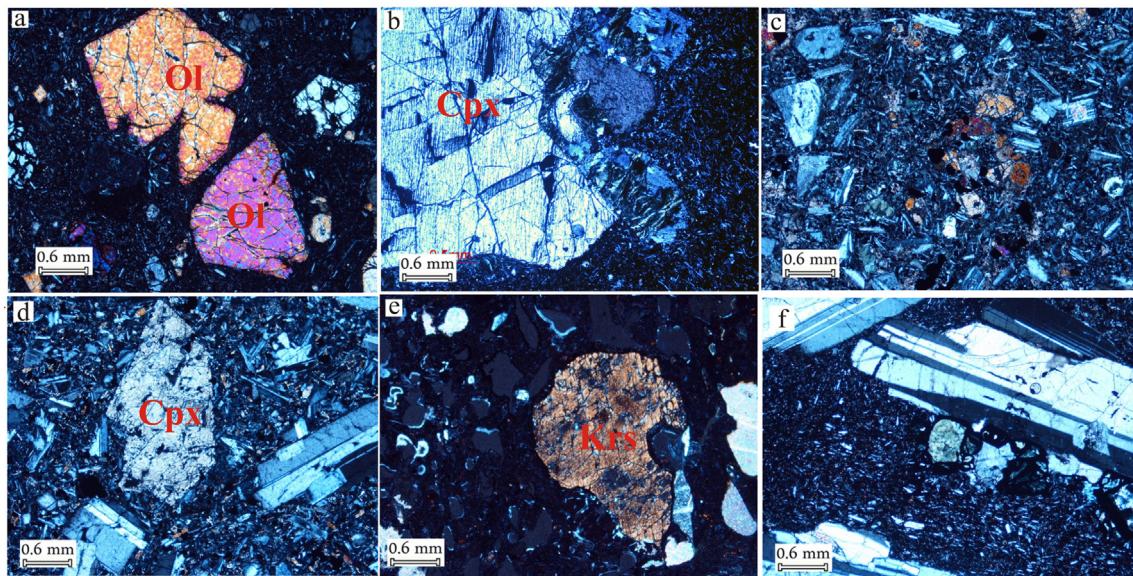


Figure 3. (a) Phenocrysts of olivine (Ol) in alkali-olivine basalts; (b) megacryst of pyroxene (Cpx) in alkali-olivine basalts; (c) glomeroporphyritic texture in Cpx of alkali-basalts; (d) phenocrysts of Cpx and plagioclase in basaltic trachy-andesites; (e) kaersutite (Krs) amphibole in basaltic trachy-andesites; (f) megacryst of plagioclase in basaltic trachy-andesite rock.

Eurasia plate in Triassic (Alavi, 1991; Stampfli, 1996, 2000). So, the Alborz block can be considered as an active mountain belt which is the consequence of Gondwana and Eurasia plate collision in Triassic (Stocklin, 1974; Berberian and King, 1981; Sengör et al., 1988; Sengör, 1990; Stampfli et al., 1991; Sengör and Natalin, 1996; Guest et al., 2006). In Paleozoic and early Mesozoic, regional extensional events are reported by different studies in all basement of Iran including Alborz block, central Iranian plateau, Kopeh Dagh zone, Zagros zone, Lut block and Sanandaj-Sirjan zone (Berberian, 1979; Berberian and King, 1981; Berberian et al., 1982; Zanchi et al., 2006).

During Mesozoic in central Alborz, extensional phases are started along with the occurrence of upper Triassic rift volcanism.

(Rhaetic) and deposition of shaly, coal-bearing Shemshak Formation by the Triassic collision between Arabian and Eurasian plates (Berberian et al., 1982). During upper Jurassic and lower Cretaceous, in the study area, shallow marine limestones deposited and accompanied with eruptions of alkaline basalts (Barrier and Vrielinck, 2008; Nazari and Shahidi, 2011).

The study area is located in the northern part of Alborz structural zone that is called central Alborz (Fig. 1). Central Alborz is composed of Paleozoic to Quaternary sedimentary rocks, wide range of Paleozoic to Quaternary volcanic rocks and some outcrops of metamorphic rocks in the northern border line. Intrusions are also common in this area, with the probably age of Paleocene and Oligocene (Fig. 2).

Table 1

Representative microprobe analysis of olivines in south Marzanabad volcanic rocks (oxides are in wt.% and elements are in apfu). Calculation is based on four oxygen.

Sample	CA53	CA53	CA53	CA53	CA53	CA53	CA53	CA53	CA53	CA53	CA53	CA53	CA53
Note	Core	Core	Core	Rim	Rim	Rim	Rim	Core	Rim	Core	Core	Core	Core
Na ₂ O	0	0	0.01	0	0.02	0	0.02	0	0	0	0.01	0.01	0.02
MgO	37.17	38.27	36.68	34.61	33.68	42.8	44.13	44.25	43.48	44.38	46.56	47.41	
Al ₂ O ₃	0.04	0.1	0.02	0.01	0	0.06	0.04	0.06	0.03	0.07	0	0.03	
SiO ₂	37.13	37.16	37.11	36.68	36.36	38.62	38.45	38.72	38.81	39.34	40.01	40.27	
CaO	0.31	0.34	0.3	0.31	0.45	0.31	0.24	0.23	0.21	0.12	0.27	0.25	
TiO ₂	0.05	0.07	0.04	0.03	0.06	0	0.02	0.01	0	0	0.02	0	
Cr ₂ O ₃	0.02	0.03	0.02	0	0	0.02	0.01	0	0.02	0.02	0.1	0.04	
MnO	0.43	0.37	0.48	0.58	0.56	0.29	0.2	0.23	0.29	0.21	0.15	0.21	
FeO	25.82	24.02	25.83	28.69	28.98	18.09	17.47	16.56	17.02	16.85	13.27	12.44	
Total	100.97	100.5	100.76	101.15	100.17	100.27	100.65	100.06	99.86	100.99	100.38	100.67	
Si	0.98	0.97	0.98	0.98	0.98	0.98	0.97	0.98	0.99	0.99	0.99	0.99	
Ti	0	0	0	0	0	0	0	0	0	0	0	0	
Al	0	0	0	0	0	0	0	0	0	0	0	0	
Cr	0	0	0	0	0	0	0	0	0	0	0	0	
Fe ²⁺	0.57	0.53	0.57	0.64	0.65	0.39	0.37	0.35	0.36	0.35	0.28	0.26	
Mn	0.01	0.01	0.01	0.01	0.01	0.01	0	0	0.01	0	0	0	
Mg	1.46	1.5	1.44	1.38	1.35	1.63	1.67	1.67	1.65	1.66	1.72	1.74	
Ca	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0	0.01	0.01	
Na	0	0	0	0	0	0	0	0	0	0	0	0	
Ni	0	0	0	0	0	0	0	0	0	0	0	0	
Total	3.02	3.02	3.02	3.02	3.02	3.01	3.03	3.02	3.01	3.01	3.01	3.01	
Fo	71.62	73.66	71.3	67.82	67.01	80.58	81.65	82.45	81.75	82.26	86.08	86.98	
Fa	27.91	25.94	28.17	31.54	32.35	19.11	18.13	17.31	17.95	17.52	13.76	12.8	
Mg [#]	71.96	73.96	71.68	68.26	67.45	80.83	81.83	82.65	82	82.44	86.22	87.17	

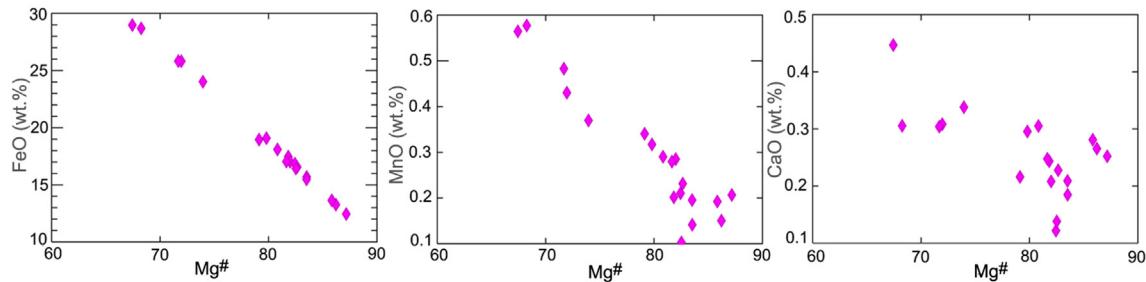


Figure 4. FeO, MnO and CaO versus Mg[#] in the olivines of south Marzanabad volcanic rocks.

A sequence of Precambrian to Cretaceous sedimentary rocks with some unconformities and hiatuses are observed in the study area. In 1971 Cartier suggested a collection of Cretaceous carbonate sediments and volcanic rocks as Chalus Formation in north Marzanabad that locates on Jurassic Shemshak Formation with angular unconformity. [Cartier \(1971\)](#) regarded 5 members for this formation. The first member consists of lower Cretaceous basic volcanic rocks which covered by the Tizkouh Formation (Aptian–Cenomanian orbitolina-enriched limestone). However, volcanic rocks which covered the Tizkouh Formation belong to upper Cretaceous and include other members of Chalus Formation ([Cartier, 1971](#)). In some places, limestone layers with large amount of Orbitolina and other Cenomanian–Coniacian fossils, are observed among volcanic rocks ([Vahdati Daneshmand and Nadim, 2001](#)).

4. Analytical methods

Samples with the most possible fresh chips were selected and powdered in an agate mill. X ray fluorescence (XRF) for major and

some trace elements (Ni, Co, Cr, V, Sr, Ba and Rb) was analyzed on powder pellets, using a wavelength-dispersive automated Philips PW 1400 spectrometer at the Department of Earth Sciences (Ferrara University), based on the method by [Franzini et al. \(1975\)](#) and [Leoni and Saitta \(1975\)](#). Trace and rare-earth elements were analyzed by inductively coupled plasma mass spectrometry (ICP-MS) at the same place, using a VG Plasma Quad2 plus. Mineral compositions were measured at the Department of Mineralogy (Padua University) with a Cameca–Camebax electron microprobe (fitted with three spectrometers) at accelerating voltage of 15 kV, and specimen current of 15 nA, using natural silicates and oxides as standards. The beam was enlarged to a diameter of 5 μm and the counting times were set to 10 s for both peak and background and the data correction was performed using PAP methods ([Pouchou and Pichoir, 1984](#)).

Isotopic analyses on whole rocks were carried out at the Department of Mathematics and Geosciences (Trieste University); whole-rock was leached with hot 6 N HCl and then digested with HF-HNO₃. Strontium and REE were separated by using standard

Table 2

Representative microprobe analysis of pyroxenes in south Marzanabad volcanic rocks (oxides are in wt.% and elements are in apfu). Calculation is based on six oxygen.

Sample	RD29	RD2	RD29	CA53	CA53	CA53	RD107	RD107	RD107	RD107	RD107	
Note	Rim	Middle	Core	Core	Middle	Rim	Core	Middle	Rim	Core	Middle	Rim
Na ₂ O	0.43	0.4	0.43	0.59	0.59	0.52	0.29	0.37	0.37	0.35	0.45	0.6
MgO	15.15	14.96	15.13	13.07	13.19	12.93	14.57	13.65	12.25	13.85	13.15	11.66
Al ₂ O ₃	6.42	6.54	6.31	8.56	8.28	6.08	3.86	3.33	6.92	3.25	2.7	2.25
SiO ₂	49.19	48.85	48.61	44.94	45.83	46.35	48.45	49.45	45.54	49.75	49.77	47.77
K ₂ O	0.01	0.01	0	0	0.01	0	0	0	0	0	0	0
CaO	21.23	21.05	21.18	21.67	22.08	22.32	21.33	21.44	21.82	22	21.94	22.09
TiO ₂	0.84	0.82	0.79	2.57	2.44	2.63	1.53	1.63	2.84	1.82	1.95	1.72
Cr ₂ O ₃	0.53	0.66	0.62	0.81	0.79	0.35	0.03	0.05	0	0	0.04	0.03
MnO	0.13	0.14	0.17	0.11	0.1	0.13	0.2	0.21	0.2	0.2	0.21	0.23
FeO	5.96	5.93	5.9	6.25	6	6.66	8.64	9.24	8.95	8.29	9.39	12.6
Total	99.9	99.36	99.16	98.56	99.29	97.97	98.9	99.37	98.89	99.51	99.6	98.97
Si	1.8	1.8	1.8	1.68	1.7	1.75	1.82	1.86	1.72	1.86	1.87	1.82
Ti	0.02	0.02	0.02	0.07	0.07	0.07	0.04	0.05	0.08	0.05	0.06	0.05
Al	0.28	0.28	0.27	0.38	0.36	0.27	0.17	0.15	0.31	0.14	0.12	0.1
Cr	0.02	0.02	0.02	0.02	0.02	0.01	0	0	0	0	0	0
Fe ³⁺	0.09	0.08	0.1	0.13	0.11	0.1	0.13	0.08	0.12	0.06	0.06	0.19
Fe ²⁺	0.1	0.11	0.08	0.07	0.07	0.11	0.14	0.21	0.17	0.2	0.23	0.21
Mn	0	0	0.01	0	0	0	0.01	0.01	0.01	0.01	0.01	0.01
Mg	0.83	0.82	0.83	0.73	0.73	0.73	0.81	0.76	0.69	0.77	0.74	0.66
Ca	0.83	0.83	0.84	0.87	0.88	0.91	0.86	0.86	0.88	0.88	0.88	0.9
Na	0.03	0.03	0.03	0.04	0.04	0.04	0.02	0.03	0.03	0.03	0.03	0.04
Total	4	4	4	4	4	4	4	4	4	4	4	4
Al ^{IV}	0.2	0.2	0.2	0.32	0.3	0.25	0.17	0.14	0.28	0.14	0.12	0.1
Al ^{VI}	0.08	0.09	0.07	0.06	0.07	0.03	0	0	0.03	0	0	0
Mg [#]	89.42	88.38	91.37	91.62	90.86	86.48	83.05	78.02	80.52	79.37	74.31	65.06
Ti/Al	0.08	0.08	0.08	0.19	0.19	0.28	0.25	0.31	0.26	0.36	0.46	0.49
Wo	0.47	0.48	0.48	0.52	0.52	0.52	0.47	0.51	0.48	0.48	0.51	0.5
Fs	0.06	0.05	0.05	0.04	0.07	0.06	0.12	0.1	0.11	0.13	0.12	0.14
En	0.47	0.48	0.47	0.43	0.42	0.42	0.42	0.4	0.42	0.4	0.37	0.37

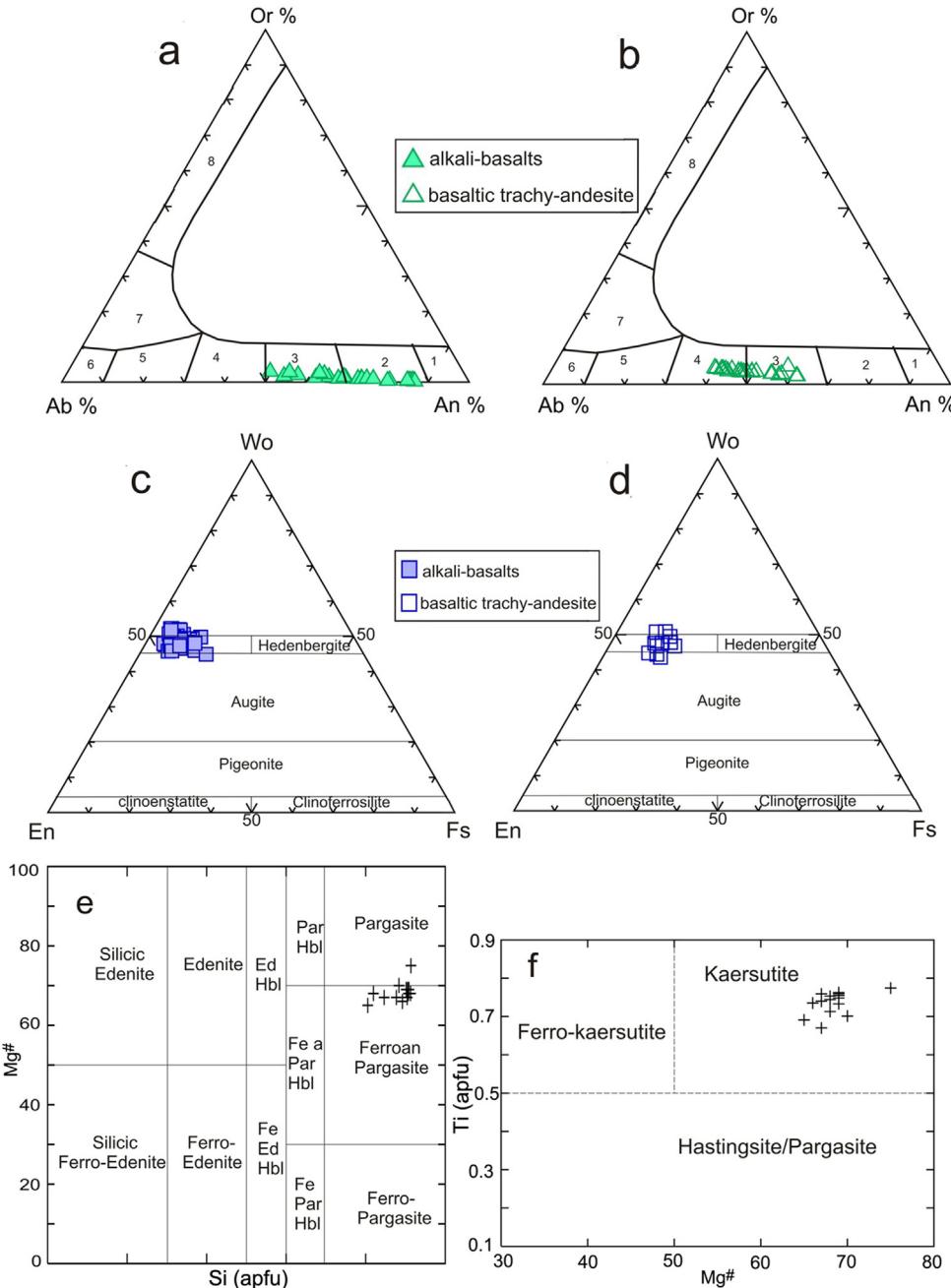


Figure 5. (a, b) Pyroxene composition of the south Marzanabad volcanic rocks in the ternary classification diagram from Morimoto et al. (1988). 1—Anorthite, 2—Bytonite, 3—Labradorite, 4—Andesine, 5—Oligoclase, 6—Albite, 7—Anorthoclase, 8—Sanidine. (c, d) Plagioclases composition of south Marzanabad volcanic rocks in the ternary diagram. (e, f) Composition of amphiboles in south Marzanabad basaltic trachy-andesite rocks which represents pargasite, ferroan pargasite and kaersutite composition (Leake et al., 1997).

cation exchange chromatographic columns with an AG 50W-X8 resin using 2.5 N HCl for Sr and 6 N HCl for the REE. Nd was separated from the other REE by using HDEHP-coated Teflon columns and 0.12 N HCl. The Sr and Nd isotope ratios were corrected by normalizing to $^{86}\text{Sr}/^{88}\text{Sr} = 0.1194$ and $^{146}\text{Nd}/^{144}\text{Nd} = 0.7219$ and using standards SRM 987 and JNd-1 for Sr and Nd, respectively.

5. Petrography

Based on petrographical studies, south Marzanabad volcanic rocks are classified into two groups: alkali-basalts and basaltic trachy-andesites.

5.1. Alkali-basalts

Alkali-basalts are dark gray to dark green in hand specimen and show massive structures. The rocks are mainly alkali-basalts and alkali-olivine basalts with mostly porphyritic texture. Phenocrysts of alkali-olivine basalts are olivine, pyroxene and minor plagioclase, while the matrix consists of pyroxene, plagioclase and accessory apatite and opaque oxide minerals (Fig. 3a). Olivine phenocrysts mostly altered to iddingsite. Porphyritic alkali-basalts contain pyroxene and plagioclase as phenocrysts with glomeroporphyric texture in some pyroxenes (Fig. 3b,c). Matrix of Alkali-basalts contains plagioclase and pyroxene with apatite, alkali feldspar and opaque accessory minerals.

Table 3

Representative microprobe analysis of plagioclases in south Marzanabad volcanic rocks (oxides are in wt.% and elements are in apfu). Calculation is based on 8 oxygen.

Sample	RD109	RD109	RD109	RD109	RD150	RD150	RD107	RD107	CA24	CA24	CA24
Note	Core	Rim	Core	Core	Rim	Core	Middle	Middle	Core	Core	Rim
Na ₂ O	1.48	3.8	1.64	3.5	5.33	3.59	6.18	5.66	4.14	5.72	6.08
MgO	0.05	0.1	0.08	0.1	0.06	0.07	0.05	1.03	0.11	0.03	0.01
Al ₂ O ₃	33.67	29.86	32.97	30.57	28.01	30.39	26.55	25.29	29.52	27.57	26.99
SiO ₂	45.6	51.14	46.26	50.19	54.84	49.99	56.68	54.61	52.14	55.51	56.32
K ₂ O	0.03	0.24	0.07	0.16	0.5	0.17	0.72	0.67	0.39	0.61	0.73
CaO	17.51	13.24	16.82	14.1	10.12	13.84	8.61	8.25	12.48	9.54	8.64
TiO ₂	0.02	0.15	0.03	0.06	0.15	0.1	0.07	0.05	0.14	0.07	0.08
Cr ₂ O ₃	0	0	0.02	0.01	0.02	0.01	0.03	0.02	0.05	0.02	0
MnO	0	0.01	0	0	0	0.04	0	0.05	0	0	0.01
FeO	0.44	0.82	0.45	0.71	0.53	0.54	0.3	1.66	0.57	0.25	0.19
Total	98.78	99.2	98.32	99.35	99.4	98.62	99.12	97.23	99.41	99.26	98.98
Si	2.13	2.35	2.16	2.31	2.49	2.31	2.57	2.54	2.38	2.52	2.56
Ti	0	0.01	0	0	0.01	0	0	0	0	0	0
Al	1.85	1.62	1.82	1.66	1.5	1.66	1.42	1.39	1.59	1.47	1.44
Cr	0	0	0	0	0	0	0	0	0	0	0
Fe	0	0	0	0	0	0	0	0	0	0	0
Mn	0	0	0	0	0	0	0	0	0	0	0
Mg	0	0.01	0.01	0.01	0	0	0	0.07	0.01	0	0
Ca	0.88	0.65	0.84	0.69	0.49	0.69	0.42	0.41	0.61	0.46	0.42
Na	0.13	0.34	0.15	0.31	0.47	0.32	0.54	0.51	0.37	0.5	0.54
K	0	0.01	0	0.01	0.03	0.01	0.04	0.04	0.02	0.04	0.05
Total	5.01	5.01	5	5.02	5.01	5.02	5.01	5.04	5.01	5.01	5.01
Ab (%)	13.25	33.7	14.97	30.71	47.35	31.62	54.15	53.12	36.68	50.2	53.65
An (%)	86.58	64.91	84.62	68.35	49.71	67.42	41.7	42.77	61.04	46.28	42.11
Or (%)	0.16	1.38	0.41	0.95	2.94	0.96	4.15	4.11	2.28	3.52	4.24

5.2. Basaltic trachy-andesite

Basaltic trachy-andesites are porphyritic, aphyric and less often amygdaloidal that contain pyroxene, plagioclase and minor amphibole (Fig. 3d,e). In some samples, the grain size of plagioclase reaches up to 3 cm (Fig. 3f). Matrix contains plagioclase and pyroxene, the accessory minerals are opaque, alkali feldspar and apatite. Biotite is observed in some samples which can be the alteration product of mafic minerals.

6. Mineral chemistry

6.1. Olivine

Representative compositions of olivines in south Marzanabad volcanic rocks, are displayed in Table 1. The olivine cores of alkali-olivine basalts have higher MgO and lower FeO, CaO and MnO rather than rims (Fig. 4). Their cores and rims reveal Fo₇₁₋₈₆ and Fo₆₇₋₈₁ composition respectively.

Table 4

Representative microprobe analysis of amphiboles in south Marzanabad volcanic rocks (oxides are in wt.% and elements are in apfu). Calculation is based on 22 oxygen.

Sample	CA24	CA2	CA24	CA24							
Na ₂ O	2.54	2.55	2.53	2.52	2.54	2.58	2.54	2.16	2.1	2.56	1.39
MgO	11.68	11.61	11.64	11.62	11.76	11.3	11.85	11.76	11.21	11.35	11.73
Al ₂ O ₃	13.6	13.74	13.88	13.86	13.32	13.78	13.29	13.51	14.07	14.02	13.78
SiO ₂	39.4	39.06	38.83	38.98	41.06	39.15	41.18	38.95	39.79	38.95	38.64
K ₂ O	1.13	1.1	1.09	1.08	1.09	1.1	1.01	1.01	1.04	1.06	1.05
CaO	10.55	10.45	10.48	10.53	10.22	10.5	10.82	10.73	10.15	10.08	10.59
TiO ₂	6.68	6.62	6.74	6.73	6.54	6.64	6.32	6.27	6.03	6.8	6.96
Cr ₂ O ₃	0.02	0.03	0	0	0.02	0.03	0.02	0	0	0	0.03
MnO	0.18	0.16	0.13	0.13	0.17	0.17	0.17	0.19	0.16	0.15	0.19
FeO	12.18	12.81	12.4	12.69	12.37	12.98	11.96	12.14	12.16	12.57	12.12
Total	97.95	98.13	97.73	98.13	99.09	98.22	99.16	96.72	96.7	97.55	96.45
Si	5.81	5.75	5.73	5.73	5.95	5.77	5.99	5.79	5.88	5.74	5.73
Al ^{IV}	2.19	2.25	2.27	2.27	2.05	2.23	2.01	2.21	2.12	2.26	2.27
Al ^{VI}	0.17	0.13	0.14	0.13	0.23	0.16	0.27	0.16	0.34	0.18	0.13
Ti	0.74	0.73	0.75	0.74	0.71	0.74	0.69	0.7	0.67	0.75	0.77
Cr	0	0	0	0	0	0	0	0	0	0	0
Fe ³⁺	0.27	0.43	0.39	0.41	0.3	0.34	0.09	0.42	0.43	0.45	0.67
Fe ²⁺	1.23	1.14	1.14	1.15	1.2	1.26	1.37	1.09	1.08	1.1	0.83
Mn	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.01
Mg	2.57	2.55	2.56	2.55	2.54	2.48	2.57	2.61	2.47	2.5	2.59
Ca	1.67	1.65	1.66	1.66	1.59	1.66	1.69	1.71	1.61	1.59	1.66
Na	0.73	0.73	0.72	0.72	0.71	0.74	0.71	0.62	0.6	0.73	0.4
K	0.21	0.21	0.21	0.2	0.2	0.21	0.19	0.19	0.2	0.2	0.2
Total	15.39	15.37	15.38	15.38	15.3	15.39	15.4	15.33	15.21	15.32	15.08
Mg [#]	0.68	0.69	0.69	0.69	0.68	0.66	0.65	0.7	0.7	0.69	0.69

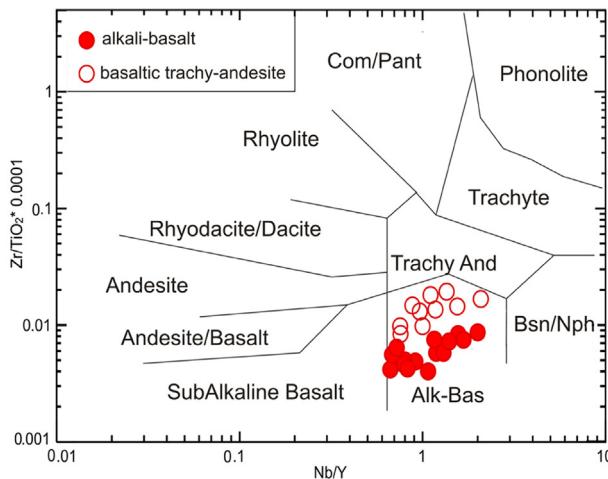


Figure 6. Classification of south Marzanabad volcanic rocks according to [Winchester and Floyd \(1977\)](#).

Table 5

Representative analysis of south Marzanabad volcanic rocks (oxides are in wt.% and elements are in ppm).

Sample	CA21	CA18	RD14	RD114	CA23	RD109	RD150	RD107	RD57	RD101	RD111	RD113	RD104
SiO ₂	48.08	42.91	47.75	42.45	46.53	45.92	45.27	53.37	51.18	44.54	46.42	46.42	46.9
TiO ₂	1.74	2.98	2	3.25	2.14	2.25	2.05	1.56	1.98	3.19	2.39	2.39	2.4
Al ₂ O ₃	18.86	14.1	14.83	15.22	17.29	17.37	15.83	18.16	17.51	15.11	17.92	17.92	17.86
Fe ₂ O ₃	12.1	17.4	12.12	17.16	12.64	12.89	12.89	9.24	9.6	16.35	13.39	13.39	13.48
MnO	0.13	0.3	0.15	0.18	0.2	0.22	0.19	0.16	0.23	0.16	0.2	0.2	0.22
MgO	5.11	7.26	10.31	7.33	7.92	7.07	11.68	2.92	4.45	7.18	4.6	4.6	5.1
CaO	10.61	10.67	9.32	9.82	10.42	10.42	8.92	8.84	8.56	9.79	10.98	10.98	8.99
Na ₂ O	2.33	1.57	2.08	2.14	2.29	3.3	1.75	4.85	5.4	2.19	3.13	3.13	3.94
K ₂ O	0.74	1.18	0.97	0.83	0.29	0.25	1.03	1.26	0.52	0.82	0.59	0.59	0.73
P ₂ O ₅	0.31	0.64	0.47	0.63	0.29	0.31	0.39	0.64	0.57	0.67	0.37	0.37	0.38
Sum	100	99	100	99	100	100	100	101	100	100	100	100	100
LO.I	2.3	1.9	2.3	3.5	2.3	2.1	3	0.7	1.2	2.1	1.4	1.8	2
Ba	236	440	292	428	188	201	280	455	349	374	322	288	230
Co	46	60	51	58	53	60	57	47	33	55	47	41	44
Cr	41	200	222	161	136	153	218	6	3	152	44	48	47
Cu	62	52	22	54	56	25	99	8	29	56	26	19	25
Ga	18	21	16	23	15	17	18	18	17	21	21	20	19
Hf	3	7	5	6	3	5	4	4	5	8	5	5	5
Nb	13	21	36	38	18	20	29	50	41	43	22	22	19
Nd	19	38	27	46	19	17	20	34	37	48	28	30	25
Ni	10	170	99	103	37	35	94	0	0	100	9	10	10
Rb	4	10	4	0	0	0	3	11	2	0	3	0	6
Sc	34	25	24	21	30	30	24	8	13	23	35	36	38
Sr	350	656	652	765	562	677	531	663	737	788	479	486	672
Th	4	6	4	6	5	5	4	4	6	7	5	5	6
V	355	316	213	261	306	341	238	109	150	258	401	393	377
Y	21	23	23	32	23	25	25	24	35	34	35	34	31
Zn	99	99	70	108	75	77	85	99	105	94	92	97	102
Zr	81	143	165	186	99	110	153	402	358	368	153	149	130
La					25.5								
Ce					29								
Pr					3.76								
Sm					3.92								
Eu					1.39								
Gd					3.68								
Tb					0.64								
Dy					3.1								
Ho					0.61								
Er					1.55								
Tm					0.23								
Yb					1.29								
Lu					0.18								
Hf					1.88								
Ta					1.01								
U					0.47								
Sample	RD142	RD29	CA53	RD7	CA20	RD125	RD26	RD140	RD127	RD56	RD57	CA52	RD100
SiO ₂	43.82	46.93	43.86	44.28	52.18	54.08	48.36	49.89	44.97	49.49	51.18	51.8	44.01
TiO ₂	2.88	2.26	3.26	2.11	1.6	1.49	2.2	1.79	2.97	2.05	1.98	2.66	3.19

6.2. Clinopyroxene

Representative compositions of pyroxenes are shown in [Table 2](#). Pyroxene is the main mineral in both alkali-basalts and basaltic trachy-andesites. Their cores are enriched of Mg, Si and Cr, and depleted of Fe, Ti, Al, Ca and Mn.

According to pyroxene classification diagram ([Morimoto et al., 1988](#)), all analyzed samples are located in diopside field ([Fig. 5a,b](#)). Composition of pyroxenes in alkali-basalts and basaltic trachy-andesites ranges from Wo_{47.9}Fs_{4.5}En_{47.6} to Wo_{49.8}Fs_{12.6}En_{37.6} and from Wo_{44.8}Fs_{10.7}En_{44.5} to Wo_{49.5}Fs_{13.6}En_{36.9} respectively. Both TiO₂ and Al₂O₃ show the greatest variation among oxides. TiO₂ ranges from 0.7 to 2.94 wt.% in alkali-basalts and from 1.53 to 3.44 wt.% in basaltic trachy-andesites. Al₂O₃ varies from 2.89 to 9.47 wt.% in alkali-basalts and from 2.25 to 6.91 wt.% in basaltic trachy-andesites. The composition of pyroxenes in groundmass is almost similar to the composition of pyroxene phenocrysts rims, but shows rather higher values of SiO₂ and FeO.

(continued on next page)

Table 5 (continued)

Sample	RD142	RD29	CA53	RD7	CA20	RD125	RD26	RD140	RD127	RD56	RD57	CA52	RD100
Al ₂ O ₃	16	16.94	13.37	15.72	18.18	18.09	18.05	17.68	14.52	18.1	17.51	16.88	15.07
Fe ₂ O ₃	15.28	12.93	14.27	12.83	10.15	9.05	12.21	11.29	15.9	11.21	9.6	10.52	16.83
MnO	0.18	0.2	0.17	0.17	0.18	0.16	0.17	0.17	0.18	0.28	0.23	0.14	0.16
MgO	9.62	5.82	10.77	11.83	3.27	3.24	3.48	5.18	8.08	3.64	4.45	4.59	6.96
CaO	7.95	11.42	10.01	9.68	8.41	6.34	11.05	8.91	8.63	8.83	8.56	7.76	10.02
Na ₂ O	1.77	2.91	3.18	1.73	3.97	5.76	3.26	3.33	1.85	5.47	5.4	3.28	2.21
K ₂ O	1.74	0.27	0.46	1.23	1.35	1.24	0.92	1.01	2.11	0.47	0.52	1.61	0.89
P ₂ O ₅	0.75	0.3	0.66	0.41	0.71	0.55	0.29	0.73	0.78	0.57	0.57	0.77	0.67
Sum	100	100	100	100	100	100	100	100	100	100.1	100	100	100
L.O.I	2.4	1.5	2.4	2.4	0.6	1.6	1.3	1.6	2.1	2.3	1.2	3.3	1.8
Ba	376	208	782	324	518	461	375	514	559	296	350	821	352
Co	49	54	67	59	21	28	41	33	52	47	33	50	51
Cr	12	142	453	248	0	12	46	1	149	25	3	41	153
Cu	59	34	77	92	3	12	42	15	51	20	29	42	55
Ga	19	16	17	18	18	18	19	19	23	20	17	21	22
Hf	6	4	6	3	4	6	4	4	8	5	5	7	7
Nb	56	19	61	35	28	29	19	24	24	31	41	40	42
Nd	40	19	33	19	39	33	19	40	54	28	37	64	54
Ni	37	36	197	117	0	2	10	0	86	6	0	22	103
Rb	8	1	2	5	16	13	12	3	18	0	2	33	0
Sc	17	37	31	27	10	8	31	13	18	18	13	14	25
Sr	886	392	573	325	608	1023	303	718	1235	1006	738	823	792
Th	6	5	8	4	3	5	5	3	7	4	6	9	6
V	193	327	307	242	88	150	357	137	239	261	150	199	253
Y	28	24	32	25	27	30	31	27	29	24	35	34	34
Zn	81	71	82	80	81	82	85	99	103	92	105	74	96
Zr	248	104	290	151	255	194	132	156	124	155	358	359	366
La	13.7	40.1	23.8	13.7	47.6	21.2	34.4						40.5
Ce	78	29	44	78	93	41	69						82
Pr	8.97	3.82	4.8	10.37	11.18	5.54	8.96						10.15
Sm	6.58	3.89	4	9.68	8.3	5.38	7.6						8.1
Eu	2.04	1.38	1.33	3.1	2.59	1.67	2.68						2.54
Gd	5.81	3.64	3.87	8.16	7.18	5.07	6.92						7.19
Tb	0.86	0.63	0.64	1.24	1.08	0.89	1.12						1.13
Dy	3.74	3.02	3.02	5.01	4.67	4.39	5.17						5.09
Ho	0.68	0.59	0.58	0.83	0.84	0.9	1.01						0.94
Er	1.69	1.51	1.45	1.84	2.05	2.41	2.6						2.36
Tm	0.24	0.22	0.21	0.23	0.28	0.38	0.39						0.33
Yb	1.3	1.22	1.15	1.17	1.55	2.22	2.22						1.84
Lu	0.18	0.17	0.16	0.15	0.21	0.32	0.32						0.26
Hf	4.54	1.69	2.16	2.31	5.14	2.48	2.65						3.89
Ta	3.27	0.97	1.56	2.48	2.85	1.04	1.47						2.2
U	1.51	0.48	0.68	1.27	1.35	1.05	0.92						0.9

6.3. Plagioclase

Representative compositions of plagioclases are listed in Table 3. In Ab-An-Or ternary diagram, they present wide range of composition from andesine to bytownite (Fig. 5c,d). Na₂O and K₂O increase from core toward rim, whereas CaO decreases. The composition of plagioclases in alkali-basalts varies from Ab_{13.2}-An_{86.59}Or_{0.16} at core to Ab_{47.35}An_{49.71}Or_{2.93} at rim and in basaltic trachy-andesites ranges from Ab_{36.67}An_{61.04}Or_{2.27} at core to Ab_{55.56}An_{39.86}Or_{4.57} at rim.

6.4. Amphibole

Representative compositions of amphiboles are presented in Table 4. Amphiboles of basaltic trachy-andesite samples, plot in the fields of pargasite and ferroan pargasite. They represent the kaersutite composition due to high Ti content [Ti (apfu) > 0.5] (Leake et al., 1997) (Fig. 5e,f). No systematic variation or zoning pattern observed in Kaersutite phenocrysts.

7. Whole rock chemistry

7.1. Major element geochemistry

South Marzanabad volcanic rocks are compositionally alkali-basalts and basaltic trachy-andesite according to Winchester and Floyd (1977) classification diagram (Fig. 6).

SiO₂, MgO and Mg[#] [$Mg^{\#} = 100 \times Mg/(Mg + Fe^T)$] ranges from 43.45 to 52.36 wt.%, 2.92 to 11.81 wt.% and 36 to 64.63, respectively (Table 5). According to Frey et al. (1978) primitive magmas have Mg[#] > 68 and Ni > 320 ppm. The most nearer sample to primitive composition has Mg[#] = 64.63 and Ni = 117 ppm. The low Ni content in this sample could signify the fractionation of olivine and Cpx (Wörner, 1999).

Basaltic trachy-andesites have higher SiO₂, Al₂O₃ and Na₂O + K₂O with lower MgO content compared to alkali-basalts (Fig. 7a–c). In CaO versus MgO and SiO₂ versus Al₂O₃ diagrams, both decreasing and increasing trends are observed (Fig. 7d,h). MgO versus CaO/Na₂O diagram, shows positive trend in south Marzanabad volcanic rocks (Fig. 7f).

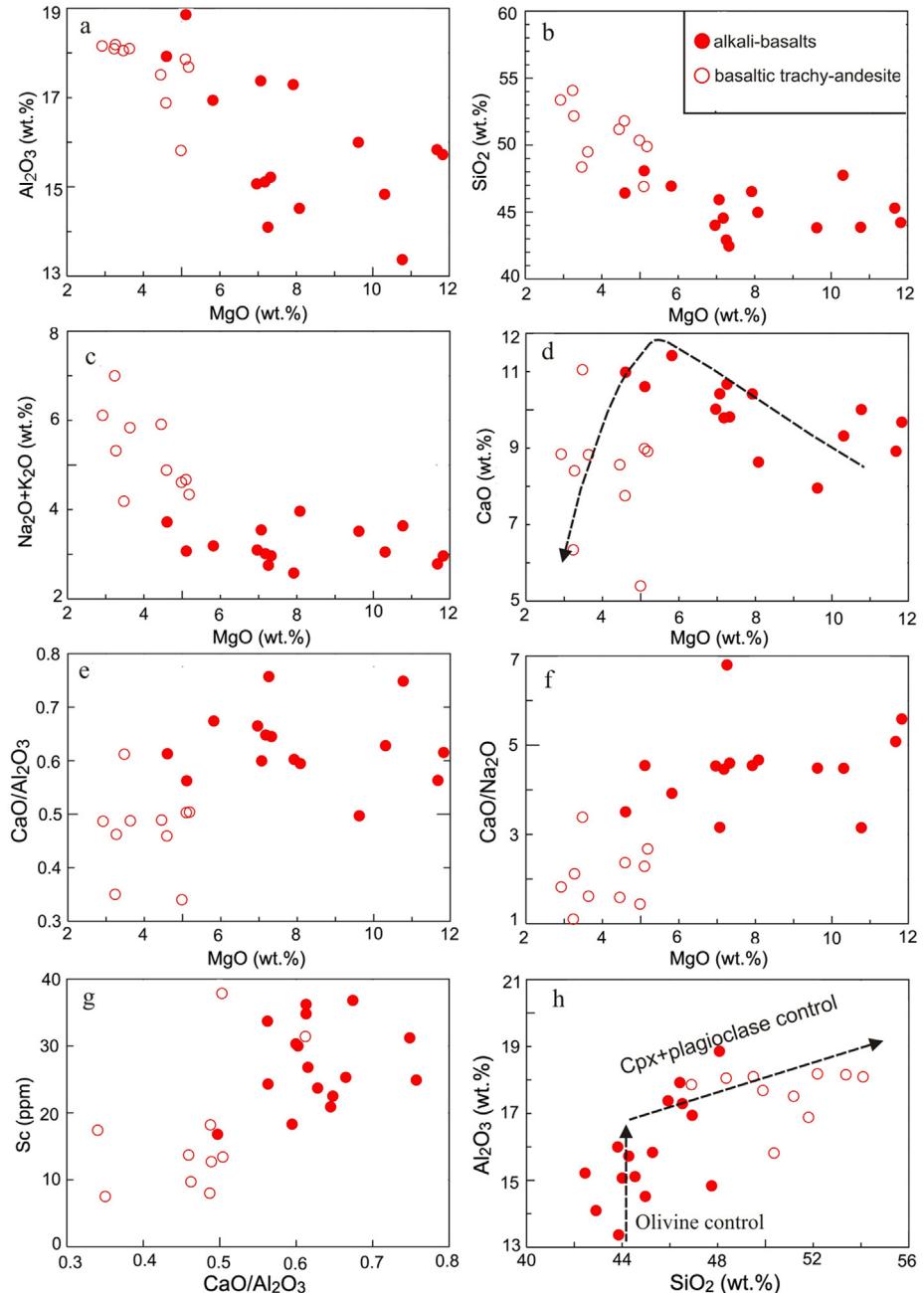


Figure 7. (a) MgO versus Al_2O_3 , (b) MgO versus SiO_2 , (c) MgO versus $\text{Na}_2\text{O} + \text{K}_2\text{O}$, (d) MgO versus CaO, (e) MgO versus $\text{CaO}/\text{Al}_2\text{O}_3$, (f) MgO versus $\text{CaO}/\text{Na}_2\text{O}$, (g) $\text{CaO}/\text{Al}_2\text{O}_3$ versus Sc, (h) SiO_2 versus Al_2O_3 in south Marzanabad volcanic rocks.

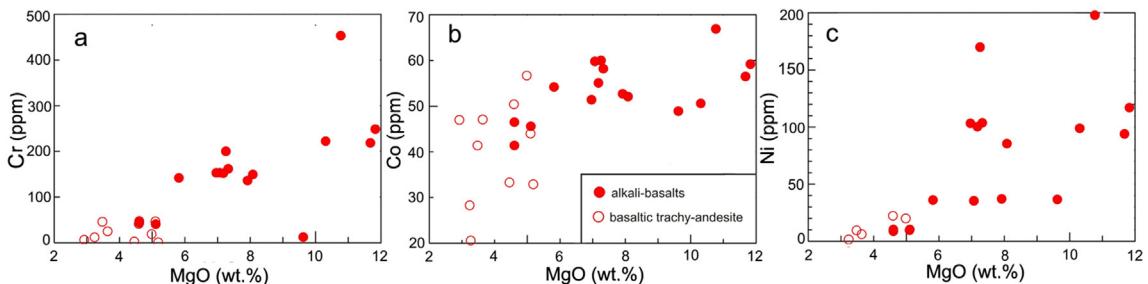


Figure 8. (a) MgO versus Cr, (b) MgO versus Co, (c) MgO versus Ni in south Marzanabad volcanic rocks.

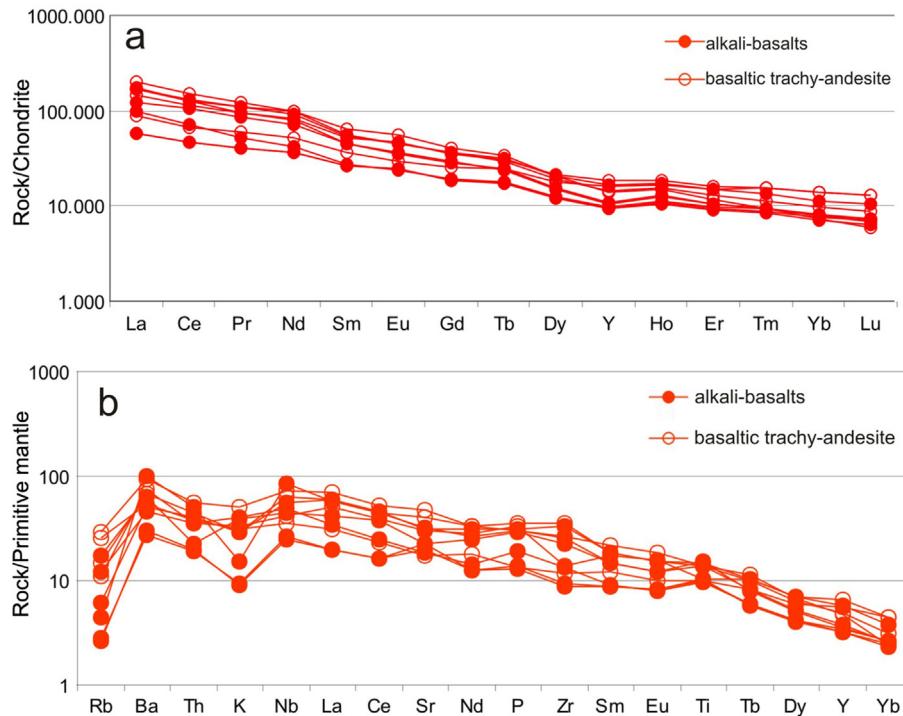


Figure 9. (a) Chondrite normalized diagram for south Marzanabad volcanic rocks, normalized values are from McDonough and Sun (1995). (b) Primitive mantle normalized diagram for south Marzanabad volcanic rocks, normalized values are from McDonough and Sun (1995).

7.2. Trace element and isotopic geochemistry

Ni, Co and Cr represent positive correlation with increasing MgO in both groups of alkali-basalts and basaltic trachy-andesites (Fig. 8). They respectively vary from 453 to 12 ppm, from 198 to 9 ppm and from 67 to 46 ppm in alkali-basalts and from 47 to 0 ppm, from 22 to 0 ppm and from 47 to 21 ppm in basaltic trachy-andesites.

South Marzanabad volcanic rocks are enriched in LREE respect to HREE and show the similar trend with OIB basalts and continental alkaline volcanic rocks (Beccaluva et al., 2002; Bianchini et al., 2002; Beccaluva et al., 2009) (Fig. 9a). They show enrichment of Nb and depletion of Rb, K and Y (Fig. 9b).

The $(\text{La}/\text{Yb})_{\text{N}}$ ratio in both groups of rocks almost is similar and ranges from 7.00 to 20.93 with an average of 13.53. The $(\text{Dy}/\text{Yb})_{\text{N}}$ ratio almost is similar in all samples and varies from 1.30 to 2.80 with an average of 1.81. Small positive Eu anomalies ($\text{Eu}/\text{Eu}^* = 1.00\text{--}1.12$) are displayed in all samples, except one basaltic trachy-andesite sample, that shows small Eu negative anomaly (0.97).

Isotopic analyses of south Marzanabad volcanic rocks are represented in Table 6. Studied samples locate in the depleted part relative to bulk earth composition regard to their low $^{87}\text{Sr}/^{86}\text{Sr}$ (0.70370–0.704387) and high $^{143}\text{Nd}/^{144}\text{Nd}$ (0.51266–0.51281)

(Fig. 10). They represent the isotopic composition of OIB field. More evolved sample from basaltic trachy-andesite group with low Mg[#] has more radiogenic $^{87}\text{Sr}/^{86}\text{Sr}$ and less radiogenic $^{143}\text{Nd}/^{144}\text{Nd}$.

8. Discussion

8.1. Fractional crystallization

Based on petrographical studies (Section 4), south Marzanabad volcanic rocks reveal the evolutionary trend.

For both alkali-basalts and basaltic trachy-andesites, negative correlations between SiO₂, Al₂O₃ and Na₂O + K₂O with increasing MgO, can indicate the derivation of volcanic rocks from the same parental magma, during fractional crystallization (Fig. 7a–c). In CaO versus MgO diagram, negative trend for MgO values more than 7 wt.%, implies the fractionation of olivine and remaining CaO in magma, whereas the fractionation of Cpx and plagioclase assumes CaO and causes positive trend for MgO values less than 7 wt.% (Rollinson, 1998) (Fig. 7d).

The correlation trend between MgO and CaO/Na₂O can reveal Cpx fractionation (Herzberg and Zhang, 1996; Machado et al., 2005) (Fig. 7f). SiO₂ versus Al₂O₃ diagram (Fig. 7h) displays olivine

Table 6

Rb, Sr, Nd and Sm concentrations and present-day Nd and Sr isotopic ratios for south Marzanabad volcanic rocks.

Sample	$^{87}\text{Sr}/^{86}\text{Sr}$	2σ	Rb (ppm)	Sr (ppm)	Rb/Sr	$^{87}\text{Rb}/^{86}\text{Sr}$	$^{143}\text{Nd}/^{144}\text{Nd}$	2σ	Sm (ppm)	Nd (ppm)	Sm/Nd	$^{147}\text{Sm}/^{144}\text{Nd}$
RD29	0.703703	0.0033	1.14	392.12	0.0029	0.0084	0.51275	0.0044856	3.89	16.9	0.23	0.14
CA53	0.704075	0.0037	1.91	573.12	0.0033	0.0096	0.512811	0.00507	6.6	32.9	0.2	0.12
RD7	0.703786	0.004	5.22	325.34	0.016	0.0464	0.51275	0.0050707	4.00	19.3	0.21	0.13
RD26	0.704387	0.0037	11.16	303.39	0.0368	0.1064	0.512666	0.0050715	5.38	24	0.22	0.14

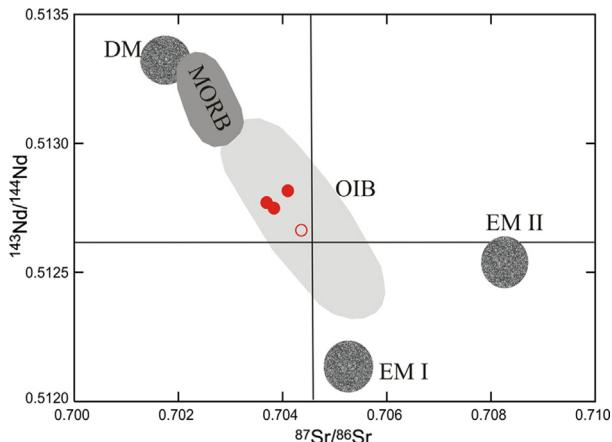


Figure 10. Sr-Nd isotopic diagram for south Marzanabad volcanic rocks. The two types enriched mantle (EMI and EMII), depleted MORB mantle (DM) and OIB field are also shown in this diagram.

fractionation in alkali-basalts and Cpx or plagioclase fractionation in the basaltic trachy-andesites (Wilson, 1989).

Basaltic trachy-andesites show the lowest Ni, Cr and V, which indicate olivine and Cpx were important fractionating mineral phases in the petrogenesis of south Marzanabad volcanic magmas. Basaltic trachy-andesites reveal higher Ba/Nb, K/Nb, Zr/Nb and Rb/Nb than alkali-basalts that display the effect of fractionation process in their evolution. The average values of Ba/Nb, K/Nb, Zr/Nb and Rb/Nb ratios in alkali-basalts and basaltic trachy-andesites are 12.79, 273, 6, 0.14 and 15.31, 316, 8, 0.36 respectively.

Decreasing Al_2O_3 with increasing MgO and small Eu anomalies indicate that plagioclase was not a major fractionating mineral phase during the evolution of the magma, pointing that the fractionation of studied rocks could occur in depth >15 km which is correlated with pressures >5 kbar in lower crust (Jung et al., 2006).

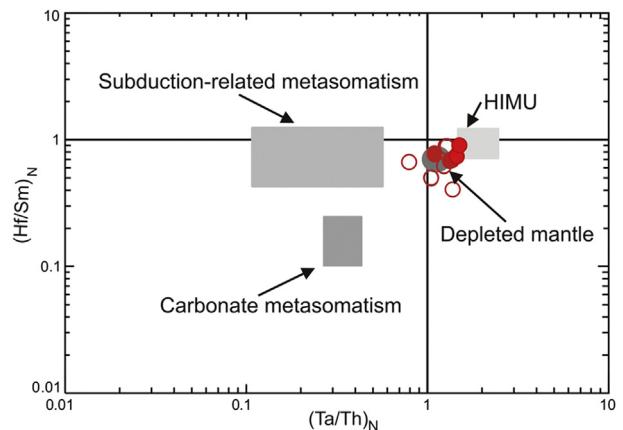


Figure 12. South Marzanabad volcanic rocks on the diagram of $(\text{Ta/Th})_N$ and $(\text{Hf/Sr})_N$ from Wang et al. (2006). All samples locate in the field of depleted mantle.

The MELTS software is used for reconstructing the fractional crystallization process in south Marzanabad volcanic rocks. MELTS is based on the work of Ghiorso and Sack (1995) and Asimow and Ghiorso (1998). In present model, the sample RD7 was assumed as a source magma that is the most primitive and undifferentiated alkali-basalt with the highest Ni and Cr. Based on this model the major element composition of alkali-basalts could be explained by almost 30% of fractional crystallization, whereas basaltic trachy-andesites could be explained by up to almost 50% of fractional crystallization (Fig. 11).

8.2. Crustal contamination and source magma

The K/Nb ratio in the studied samples varies from 62 to 588 and is averagely higher in evolved samples. This ratio ranges from 62 to 588 in alkali-basalts with an average of 234 and from 100 to 540 in basaltic trachy-andesites with an average of 316. Based on Taylor

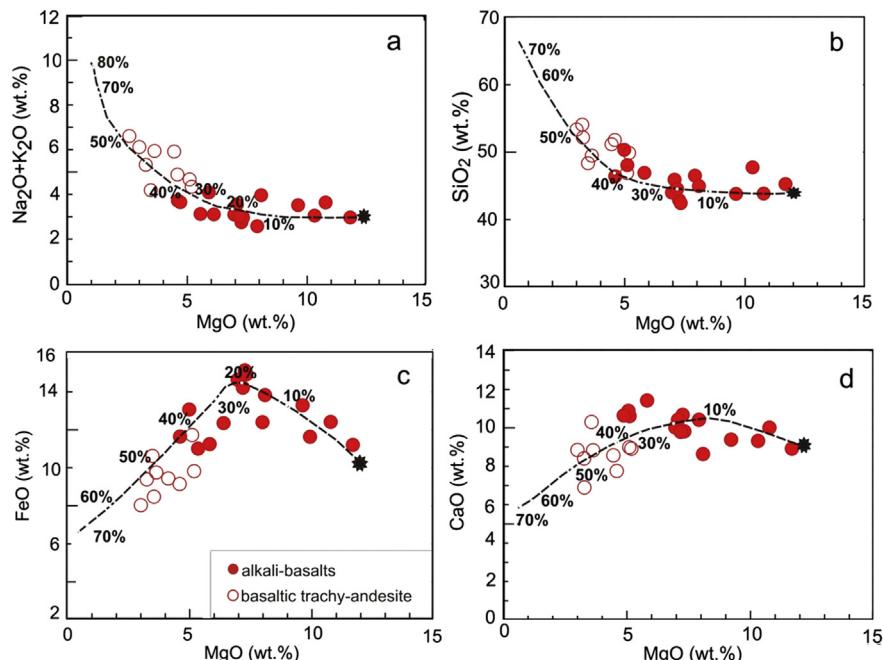


Figure 11. (a) MgO versus $\text{Na}_2\text{O} + \text{K}_2\text{O}$, (b) MgO versus SiO_2 , (c) MgO versus FeO , (d) MgO versus CaO in south Marzanabad volcanic rocks. Lines represent the results of fractional crystallization calculation by MELTS software with numbers that indicate the mass crystallized as wt.%.

and McLennan (1985) K/Nb in the Earth's crust is higher than mantle. More values of K/Nb in basaltic trachy-andesites could show the occurrence of some minor crustal contamination (Jung et al., 2006).

In primitive mantle normalized diagram, basaltic trachy-andesites show less depletion of K compared to alkali-basalts which may reflect the effect of minor crustal contamination or fractional crystallization. Zr/Nb varies from 4.27 to 5.59 in alkali-basalts which are less evolved; however in more evolved basaltic trachy-andesites, Zr/Nb ranges from 6.43 to 9, which could remark the effect of minor crustal contamination (Wörner, 1999).

The HFSE/LREE ratio can indicate the characters of source magma. According to Smith et al. (1999) ratios more than unity represent the asthenospheric origin for magma, whereas ratios less than unity represent the lithospheric origin. In south Marzanabad volcanic rocks Nb/La, Zr/La and Zr/Ce ratios are 1.15, 4.51 and 2.23 respectively, which suggest the melt origination from an asthenospheric mantle. Based on rare earth elements and isotopic composition, the studied rocks show the similar source characteristics and derive from the same magmatic source in depleted mantle with regard to bulk earth composition (Fig. 12).

8.3. Crystallization condition

Low Ti/Al ratios in Cpx composition specify the crystallization of Cpx in low pressures (e.g., Wilkinson, 1974; Dobosi et al., 1991; Włodyka, 2002; Ali and Ntaflos, 2011). The Ti/Al ratio varies from

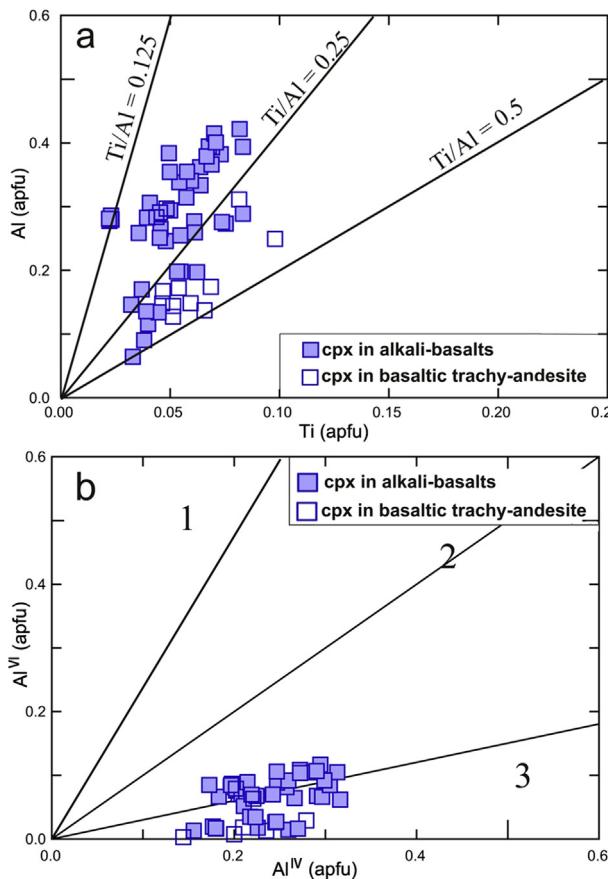


Figure 13. (a) Ti versus Al diagram in the pyroxenes of south Marzanabad volcanic rocks. (b) Plot of Al^{VI} versus Al^{IV} in the pyroxenes of south Marzanabad volcanic rocks (calculation is based on 6 oxygens), fields are: 1—eclogites, 2—granulites and inclusions in basalts, 3—igneous rocks (Aoki and Shiba, 1973).

0.077 to 0.513 in alkali-basalts and from 0.154 to 0.537 in basaltic trachy-andesites (Fig. 13a). The average of this ratio in alkali-basalts and basaltic trachy-andesites respectively is 0.20 and 0.34. Concerning the lower Ti/Al ratio in Cpx of alkali-basalts, they could be crystallized in rather higher pressures than Cpx of basaltic trachy-andesites. On the other hand, increasing of Al^{VI}/Al^{IV} ratio in Cpx, shows the pressure increment during crystallization (Green and Ringwood, 1968; Wass, 1979). Al^{VI}/Al^{IV} ratio varies from 0.00 to 0.49 in alkali-basalts with an average of 0.24 and from 0.00 to 0.39 in basaltic trachy-andesites with an average of 0.07 (Fig. 13b). As a result, the crystallization of Cpx in alkali-basalts took place in slightly higher pressures than Cpx in basaltic trachy-andesites.

8.4. Partial melting of source magma and mantle potential temperature

The differentiation degree of LREE relative to HREE is defined by the (La/Yb)_N ratio that varies from 7.00 to 20.93 with an average of 13.53 in south Marzanabad volcanic rocks. High (La/Yb)_N in studied rocks suggests the low degree of partial melting in their mantle source (Jung et al., 2006).

The HREE enrichment degree is represented by the (Dy/Yb)_N ratio. The partial melts generated from garnet-lherzolite source, have (Dy/Yb)_N > 1.06 (Blundy et al., 1998; Peters et al., 2008). (Dy/Yb)_N ratio ranges from 1.29 to 2.80 with an average of 1.81 in Marzanabad volcanic rocks, which signifies the garnet was a residual phase during the partial melting.

We used PRIMELT2.XLS software from Herzberg and Asimow (2008) to calculate primary magma composition and mantle potential temperature. The four most primitive rock samples of south Marzanabad volcanic rocks plot above the dashed-line in Fig. 14 that characterizes olivine-fractionated primary magma from peridotitic sources. For calculating the most proper primary magma composition for Marzanabad volcanic rocks, we used the most primitive RD7 sample with Fe₂O₃/TiO₂ = 1.0 to estimate Fe₂O₃ (Herzberg and Asimow, 2008). The calculated primary magma composition have 44.75 SiO₂, 2.13 TiO₂, 15.89 Al₂O₃, 2.13 Fe₂O₃, 9.75 FeO, 11.95 MgO and 9.78 CaO (all in wt.%).

The calculated mantle potential temperature for Marzanabad volcanic rocks is 1370 °C. The ambient mantle potential temperatures that make MORB is about 1300–1454 °C (Herzberg et al., 2007; Putirka et al., 2007), showing that the Marzanabad volcanic rocks come up from an ambient mantle.

For modeling the partial melting of common upper mantle source, REE systematics like the plot of La/Yb vs. Dy/Yb (Fig. 15) can

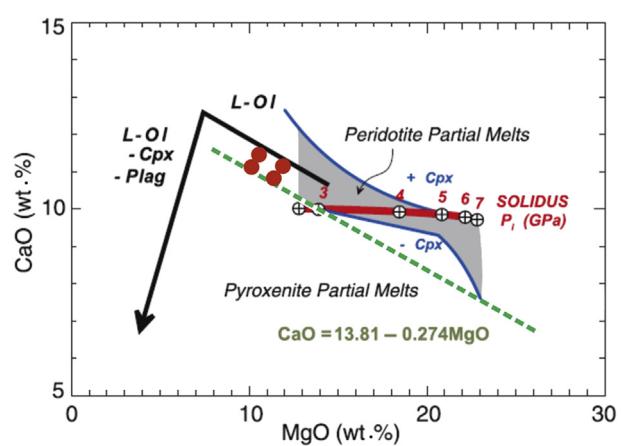


Figure 14. CaO and MgO diagram of the south Marzanabad volcanic rocks compared with partial melts of pyroxenite and peridotite (Herzberg and Asimow, 2008).

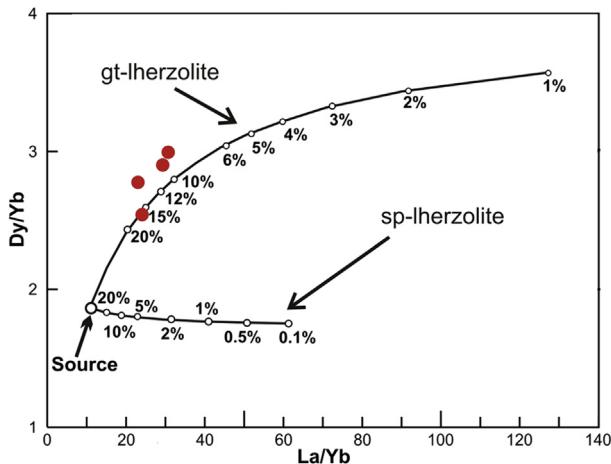


Figure 15. La/Yb vs. Dy/Yb covariation for the south Marzanabad volcanic rocks. Partial melting curves were calculated using a non-modal, fractional melting model (Shaw, 1970).

be useful (Thirlwall et al., 1994; Baker et al., 1997; Jung et al., 2006). This plot is easily recognizes between melting in the garnet peridotite stability field and spinel peridotite stability field owing to the strong fractionation of HREE by garnet.

Partial melting in the source magma of south Marzanabad volcanic rocks was reconstructed (Fig. 15). Partial melting curves were calculated using a modal, fractionated melting model (Shaw, 1970). Mineralogical composition of primitive mantle (66% olivine, 14% orthopyroxene, 14% clinopyroxene, 6% garnet) and chemical composition of primitive mantle are taken from Mertz et al. (2001). Mineral-melt distribution coefficients are taken from McKenzie and O’Nions (1991, 1995) and normalizing values are from McDonough and Sun (1995). In both diagrams of Figs. 15 and 16, the most primitive RD7 sample, reveals nearly 15% of partial melting in the garnet peridotite stability field, which produces the primary magma of south Marzanabad volcanic rocks.

8.5. Melt generation

The alkali basalt magmatism in central Alborz is related to deep mantle melting or mantle plume associated with crustal contamination in local extensional system or deep faults system (Haghnazari et al., 2009; Jaffarian et al., 2009). Alborz zone reveals

an elevation 3000–5000 m above sea level. According to geophysical studies, mantle lithosphere is almost absent underneath the Alborz Mountains (e.g., Sodoudi et al., 2009; Mirnejad et al., 2010). The crustal thickness under the central Alborz is estimated about 35–45 km (Dehghani and Makris, 1984; Amjadi et al., 2012). However insufficient crustal base and the relatively thin lithosphere underneath the Alborz can predicate that the asthenospheric mantle is backing the high elevation (Sodoudi et al., 2009). Ansari et al. (2011) believed that alkaline magmatism in central Alborz zone was generated by the deep mantle melting and the delamination of sub-continental lithosphere.

The calculated mantle potential temperature shows that south Marzanabad volcanic rocks generated from the mantle at ambient temperature which is against the mantle plume hypothesis (Hastie and Kerr, 2010).

In summary, in south Marzanabad area in north of central Alborz, like many other parts of Iran, an extensional rift basin was originated during Cretaceous. Based on isotopic and REE data of south Marzanabad volcanic rocks, alkaline volcanism was occurred by the extension and partial melting of asthenospheric mantle in the rift basin without major affecting of contribution subcontinental lithospheric mantle or crustal contamination. Erupted alkaline volcanic rocks overlapped the upper Jurassic Shemshak sediments with the angular unconformity. Existence of magmatic chambers en route of ascending magma could cause crystal fractionation. Finally in late Cretaceous, the Laramide orogenic phase terminated the rifting extensional system and volcanism.

9. Conclusion

South Marzanabad volcanic rocks underwent fractional crystallization that is confirmed by petrographical and geochemical studies. Basaltic trachy-andesites formed from alkali-basalts due to fractional crystallization. Based on the presented fractionation model of south Marzanabad volcanic rocks, the composition of more evolved samples could be explained by up to almost 50% of fractional crystallization.

The source magma of south Marzanabad rocks was generated by the low degree of partial melting from the garnet bearing mantle source with regard to high values of $(\text{La}/\text{Yb})_{\text{N}}$ and $(\text{Dy}/\text{Yb})_{\text{N}}$. Based on low $^{87}\text{Sr}/^{86}\text{Sr}$, high $^{143}\text{Nd}/^{144}\text{Nd}$ and HFSE/REE ratios, south Marzanabad volcanic rocks show the similar source characteristics and derivation from the depleted mantle source with regard to bulk earth composition.

The calculated mantle potential temperature shows that south Marzanabad volcanic rocks generated from the mantle at ambient temperature which is contrary to mantle plume hypothesis. In Cretaceous, alkaline volcanism was occurred by upwelling and decompressional melting of an asthenospheric mantle in the rift basin without major involvement of subcontinental lithospheric mantle or crustal contamination.

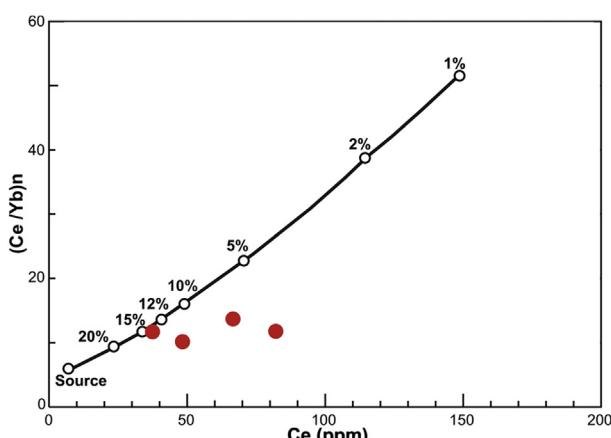
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Figure 16. Chondrite normalized Ce/Yb vs. Ce for south Marzanabad volcanic rocks. Normalization values are from McDonough and Sun (1995).



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