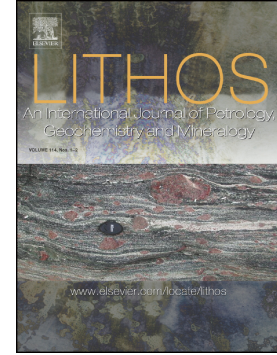


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Crust-mantle interaction inferred from the petrology and Sr-Nd-Pb isotope geochemistry of Eocene arc lavas from the Kahrizak Mountains, north-central Iran

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**Abstract**

The Eocene volcanic rocks from the Kahrizak Mountains in north-central Iran are part of the Urumieh-Dokhtar magmatic arc, which runs parallel to the Main Zagros Thrust as the Neo-Tethys suture. These volcanic rocks, similar to those from eastern Pontides and northern Anatolia, Turkey, were mainly produced during the Eocene magmatic flare-up associated with the Arabia-Eurasia convergence. The rock suite includes basalt, trachyandesite/andesite and trachydacite/rhyolite lavas and pyroclastic deposits that evolved compositionally from calc-alkalic to shoshonitic. Their normalized trace element concentration patterns are moderately enriched in light rare earth element and depleted in high field-strength elements (HFSE; e.g., Nb, Ta, Ti). They have narrow ranges of initial Pb isotopic ratios and  $^{143}\text{Nd}/^{144}\text{Nd}_i$ , but highly variable  $^{87}\text{Sr}/^{86}\text{Sr}_i$ . The new analyses indicate that the parental magmas of the volcanic rocks were derived from a mantle source that had been enriched by fluids released from a subducted oceanic slab. The fluids introduced significant amounts of large ion lithophile elements, but negligible HFSE to the source. The parental magmas underwent fractional crystallization and assimilation of upper crustal materials to produce the range of volcanic rocks. Integration of new analyses with regional data suggests that the Eocene volcanic rocks from north-central Iran, together with ~coeval volcanic rocks in eastern Pontides and northern Anatolia, were most probably derived from

a lithospheric mantle source that had been previously metasomatized by fluids derived from a subducted slab before and during the Arabia-Eurasia collision.

Index terms: Iran, Urumieh-Dokhtar zone, Kahrizak geochemistry, Eocene magmatic flare-up, lithospheric mantle.

## 1. Introduction

Magma genesis in subduction zones is generally more complex than in divergent plate boundaries and intraplate settings due to the wide range of igneous, sedimentary, metamorphic, and aqueous fluid components that potentially participate during the partial melting, magma mixing and assimilation processes that generate arc magma. The complexity is multiplied in continental margins because the ascent of magma through thick and compositionally heterogeneous overriding plate, combined with prolonged magma storage, bring about additional compositional modifications observed in continental arc magmas. Over the last two decades extensive investigations on crust-mantle interactions have been carried out for some of the major Cenozoic continental arc systems such as the Andes (e.g., Mattioli et al., 2006; Schiano et al., 2010), Indonesia (Gertisser and Keller; 2003) and Turkey (Temizel et al. 2012); however, modern detailed geochemical methods have seldom been applied to the Eocene arc of Iran.

In this paper we deal with a poorly investigated Eocene arc volcanic suite from the Kahrizak Mountains in north-central Iran in order to better constrain its tectono-magmatic evolution in the context of coeval magmatism in the region (Fig. 1). Eocene arc volcanism represents the most voluminous magmatic event in Iran, so much so that it has been ranked with famous flare-ups worldwide (Verdel et al., 2011). The volcanism is ultimately associated with the Neo-Tethyan plate

subduction during the Mesozoic and Cenozoic (e.g., Omrani, 2008; Verdel et al., 2011; Deevsalar et al., 2017; Zhang et al., 2018). The Eocene arc volcanic suite from north-central Iran is compositionally diverse and includes mantle-derived mafic to crustal-contaminated felsic end-members, although some intermediate varieties are also present. In this contribution we focus on crust-mantle interactions by employing the petrography, mineral compositions, major and trace element chemistry, and Sr, Nd and Pb radiogenic isotopes of the Eocene arc volcanic suite.

## **2. Geological setting and field relations**

The substrate of the Iranian Paleogene arc consists of a mosaic of continental basement blocks that was consolidated during the Neoproterozoic-Early Cambrian Cadomian orogeny along the northern margin of Gondwana (Ramezani and Tucker, 2003; Hassanzadeh et al., 2008; Rahmati-Ilkhchi et al., 2011). The opening of Neo-Tethys Ocean in the Permian time left these basement blocks situated between a new ocean that was expanding and an older one that was being consumed (Berberian and King, 1981). The subduction initiation along the northern margin of the Neo-Tethys, beneath the Sanadaj-Sirjan zone of Iran, followed the collision between northern Iran and Eurasia during closing of the Paleo-Tethys (Berberian and Berberian, 1981; Horton et al., 2008; Hassanzadeh and Wernicke, 2016). In the Cretaceous-Paleocene time, arc magmatism

diachronously shifted to the northeast and formed the Urumieh-Dokhtar zone in central Iran (Verdell et al., 2011; Hosseini et al., 2017).

The Kahrizak Mountains are located twelve miles south of Tehran in north-central Iran (Fig. 1). They are part of the Urumieh–Dokhtar arc (UDA) in the northern sector of the Central Iran tectonic province, which is bounded by the Sanandaj–Sirjan zone in the west-southwest, and by the Alborz Mountains in the north. To the east, the UDA includes the Great Kavir Depression, which is underlain by one of the aforementioned Gondwana basement blocks exposed in several tectonic windows, with igneous protoliths ranging in age from ca. 580 to 520 Ma (e.g., Ramezani and Tucker, 2003; Hassanzadeh et al., 2008; Rahmati-Ilkhchi et al., 2011). Sedimentary cover of the Great Kavir Block includes thick ( $\leq 9$  km) Paleozoic-to-Cenozoic sequences that are largely comparable to those overlying the other basement blocks. Geochronological data obtained from the few outcrops of the Great Kavir Block additionally indicate granite-tonalite plutonic emplacement at ca. 215 Ma in the Saghand area (Ramezani and Tucker, 2003) and a medium-pressure amphibolite-facies metamorphism at ca. 166 Ma in the Rezveh area (Rahmati-Ilkhchi et al., 2011) to the southeast and east, respectively, of the Kahrizak Mountains.

Eocene calc-alkaline volcanic rocks occur almost throughout Central Iran and partly in neighboring provinces (Fig. 1; e.g., Verdell et al., 2011; Moghadam et

al., 2016). Radiometric ages (U-Pb and  $^{40}\text{Ar}/^{39}\text{Ar}$ ) of lava and pyroclastic samples from UDA and Alborz Mountains as well as mafic and silicic plutonic rocks from the Sabzevar, Khorram Darreh and Saghand areas in Central Iran and Sanandaj–Sirjan zone range from 55 to 37 Ma (Ramezani and Tucker, 2003; Hassanzadehe et al., 2008; Verdell et al., 2011; Moghadam et al., 2016). This magmatic flare-up occurred supposedly while the Neo-Tethyan slab began a slow roll-back after a subhorizontal subduction in the Late Cretaceous-Paleocene times (Verdel et al., 2011; Zhang et al., 2018; and references therein). The Eocene volcanic rocks were largely erupted in shallow marine environment, but concomitant basin and range tectonism also generated subaerial volcanic eruptions and rapid accumulations of thick sediments, volcano-sedimentary deposits, pyroclastics, and lava flows in the subsiding basins (Amidi et al., 1984). Volcanism diminished and the region began uplifting at the closure of the Neo-Tethys Ocean, and this was followed by the Arabia-Eurasia collision that is estimated to be in late Eocene (ca. 35 Ma - Bottrill et al., 2012), or late Oligocene (ca. 27 Ma, Pirouz et al., 2017). Deposition of detrital and shallow marine sediments with sparse volcanism ensued during the late Oligocene-early Miocene time and resulted the conspicuous trio of Lower Red, Qom and Upper Red Formations indicating broad subsidence and uplift, most likely due to the dynamic topography of the overlying plate during continental collision (e.g., Bottrill et al., 2012). The Qom Formation is carbonate-dominant



and the thickest (500 – 1000 m) in central Iran, but it extends northwest Iran, and into the southern Turkey with varying thicknesses.

The Eocene volcanic rocks exposed in the northern and eastern parts of the Kahrizak Mountains mainly consist of about 300 m of variegated rhyolites and dacitic to trachytic tuffs. These are overlain by ignimbrites that are mainly trachydacite in composition and are easily distinguished by their massive outcrops and steep slopes. Widespread distribution of acidic tuffs and ignimbrites suggests that first volcanic activity in the area was explosive. The explosive phase was followed by lava eruptions. In some parts of the study area, the pyroclastic rocks are overlain by a lesser volume of rhyolitic lavas. A ~130 m thick sequence of basalt and pyroxene-bearing basaltic andesite, basaltic trachyandesite and trachyandesite also occurs in the Kahrizak Mountains. Zeolites are common secondary minerals. The strata are dissected by several faults which are N- and NW-trending.

### **3. Analytical methods**

More than 200 thin-sections of volcano-sedimentary rocks from various stratigraphic levels of the Kahrizak Mountains were analyzed under the microscope to identify their mineralogy and petrographic characteristics, and to select the least altered samples – i.e., those without veinlets and amygdales as well

as secondary calcite, quartz, chlorite, zeolites, and/or clay minerals. Chemical compositions of minerals were determined using the JEOL 8200 electron microprobe at the California Institute of Technology using a focused electron beam with an accelerating voltage of 15 kV and a beam current of 25 nA. Standards for analysis were anorthite ( $\text{SiK}\alpha$ ,  $\text{AlK}\alpha$ ,  $\text{CaK}\alpha$ ); albite ( $\text{NaK}\alpha$ ); fayalite ( $\text{FeK}\alpha$ ); forsterite ( $\text{MgK}\alpha$ );  $\text{Mn}_2\text{SiO}_4$  ( $\text{MnK}\alpha$ );  $\text{TiO}_2$  ( $\text{TiK}\alpha$ );  $\text{Cr}_2\text{O}_3$  ( $\text{CrK}\alpha$ ); and microcline ( $\text{KK}\alpha$ ). A precision approaching <1% relative error and accuracy as good as 1-2% were obtained. Quantitative elemental microanalyses were processed with the CITZAF correction procedure (Armstrong, 1995), and analytical results are given in supplementary Table S1.

Through petrographic inspections, 14 representative samples were selected as the least altered for determining whole rock compositions at the ACME Analytical Laboratories (Vancouver, Canada). Major element compositions were analyzed by inductively-coupled plasma-atomic emission spectrometry (ICP-AES). Trace elements, including rare earth elements (REE), were determined by inductively-coupled plasma-mass spectrometry (ICP-MS). The major and trace element compositions of two samples (K.91.31 and K.91.14) were analyzed at the Lab West Mineral Analysis LTD- Australia, and Zarazma Mineral Studies Co, Iran. The major and trace element analyses including their precisions are presented in Table 1 and supplementary Table S2.

Strontium, Nd and Pb isotope ratios for 9 of the samples were determined using a 9-collector, Micromass Sector 54 thermal ionization mass spectrometer (TIMS) at the Scripps Institution of Oceanography, University of California, San Diego. The sample preparation procedure used is similar to that described in Tian et al. (2008). Chips of whole rock samples were hand-picked to avoid fracture- and vug-filled portions as well as weathered crust, ultrasonicated in dilute HNO<sub>3</sub> acid for about 30 minutes, rinsed with ultrapure H<sub>2</sub>O, and dried in an oven at 105 °C overnight prior to powdering in an alumina ceramic grinder. About 35 mg powder of each sample were digested with a double-distilled, 2:1 mixture of concentrated HF:HNO<sub>3</sub> acid in a clean Teflon beaker. Lead was first separated by re-dissolving the dried samples in 1N HBr and then passing the solutions through a small ion exchange column in an HBr medium. Strontium and REE were separated from the residual solutions in an ion exchange column using HCl as the eluent. Finally, Nd was separated from the rest of the REE in an ion exchange column using alpha hydroxyisobutyric acid as the eluent. The isotopic analyses are presented in Table 2, which also includes the details of the TIMS procedure as footnotes.

## **4. Results**

### **4.1. Petrography and mineral chemistry**

The bulk compositions of the Eocene lava flows and pyroclastic rocks from the Kahrizak Mountains range from basalt to rhyolite. Based on their mineralogy and major element composition, these rocks generally belong to two main groups, calc-alkaline basaltic and alkaline trachydacite-rhyolite groups (Fig. 2). The pyroclastic rocks include tuff and ignimbrite are widespread in the area. Below we present the results of our investigation of each compositional group.

#### 4.1.1. The calc-alkaline basaltic group

This group is commonly observed closely together in the field and includes basalt and intermediate rocks including basaltic trachyandesite, trachyandesite and andesite. The basalts show a wide range of textures, from phyrlic, microphyric, hyalophyric, glomerophyric to rarely intersertal, intergranular, and trachytic. Their typical mineral assemblages include plagioclase, olivine, clinopyroxene, orthopyroxene, and Fe-Ti oxide (Table S1 and Figs. 3A-F). Plagioclase ( $\text{Na}_{0.10-0.51}\text{Ca}_{0.29-0.92}\text{Si}_{0.20-3.03}\text{Al}_{0.77-1.95}\text{O}_8$ ) ranges from labradorite to bytownite in composition ( $\text{An}_{55-90}$ ; Figs. 4A, B). Plagioclase phenocrysts generally exhibit oscillatory zoning and sieve texture (Figs. 3A-C); some have reaction rims and are rounded. Those with oscillatory zoning generally have bytownite to anorthite cores and albite ( $\text{An}_{25.9}$ ) and andesine ( $\text{An}_{35.7-45.9}$ ) rims (Fig. 4C). Some basalts have

phenocrysts with andesine rim and oligoclase ( $\text{An}_{26.4-32.4}$ ) core; others show crystals of Na-sanidine ( $\text{K}_{0.22}\text{Na}_{0.24}\text{Al}_{0.53}\text{Si}_{1.46}\text{O}_8$ ) in the cores of oligoclase.

Olivine occurs as euhedral to subhedral crystal, range from chrysolite ( $\text{Fe}_{0.19-0.20}\text{Mg}_{0.46-0.51}\text{Si}_{1.18-1.23}\text{O}_4$ ) – hyalosiderite ( $\text{Fe}_{0.19-0.29}\text{Mg}_{0.44-0.49}\text{Si}_{1.17-1.75}\text{O}_4$ ) in composition (Fig. 4E) and appears to be zoned. Clinopyroxene ( $\text{Wo}_{43-49}\text{En}_{37-43}\text{Fs}_{13-16}$ ) occurs as euhedral to subhedral grains and is diopside-augite in composition ( $\text{Na}_{0.03-0.63}\text{Ca}_{0.26-1.17}\text{Fe}_{0.04-0.04}\text{Mg}_{0.03-1.12}\text{Al}_{1.13-1.18}\text{Ti}_{0.01-0.04}\text{Si}_{2.41-2.58}\text{O}_6$ ) (Fig. 4D). Clinopyroxene also displays zonation and reaction textures with more sodic ( $\text{Wo}_{79}\text{En}_{10}\text{Fs}_{11}$ ) rim composition (Figs. 4d, F). The Fe-Ti oxides are mainly magnetite-titanomagnetite ( $\text{Ti}_{0.33-0.37}\text{Fe}_{2.37-2.57}\text{O}_4$ ) (Table S1).

The trachyandesites and basaltic trachyandesites consist mainly of plagioclase, clinopyroxene, hornblende, biotite, and Fe-Ti oxide. Plagioclase ( $\text{An}_{36-59}$ ) (Fig. 4A and Table S1) occurs as euhedral to subhedral crystals and typically shows sieve texture and normal zoning. Hornblende occurs as subhedral grains and is classified (Hawthorne, 1983) as magnesio-hastingsite and magnesian hastingsite ( $\text{Na}_{0.68-0.72}\text{Ca}_{1.71-1.73}\text{Fe}_{1.51-1.54}\text{Si}_{5.86-5.95}\text{Al}_{1.94-2.0}\text{O}_2(\text{OH})_2$ ; magnesian number (Mg#) = 0.65-0.70 (Fig. 4F and Table S1).

The basaltic rocks were strongly affected by fluid/rock interaction and, thus, contain secondary minerals due to alteration. The paragenetic sequence of alteration began with precipitation of phyllosilicates on walls of veins followed by

crystallization of Na-zeolite minerals (analcime, tetranatrolite and natrolite) and Ca-zeolites (thomsonite/natrolite, gonnardite, chabazite, stilbite, scolecite/mesolite, and heulandite) from the fluid. The latter is a mixture of sea and magmatic waters (Yazdani et al., 2014). The major-trace element and isotopic composition of zeolites and host rocks suggests that the alteration occurred penecontemporaneously with volcanic eruptions in a submarine environment.

#### **4.1.2. The alkaline trachydacite-rhyolite group**

This group includes rhyolite lavas, trachydacitic ignimbrites and tuffs. The mineral assemblage in rhyolites includes plagioclase, quartz and biotite in a glassy, trachytic groundmass. There are two varieties of glassy groundmass - dark brown and yellow. The rhyolites generally exhibit hyalo-microlitic porphyritic and spherulitic textures with perlitic cracks. Some quartz and many biotite grains have corroded rims. Ignimbrites vary in texture from vitrophyric to eutaxitic; the fiamme-bearing ignimbrites show vitroclastic and fluidal structures, in which the lenticular flames and phenocrysts are oriented along the flow direction. Tuffs of rhyolitic to dacitic composition consist of vitric, crystal and lithic tuffs.

## **4.2. Major and trace element compositions**

The samples analyzed are mainly lava flows and two samples of pyroclastic rocks - an ignimbrite and a tuff. Notably, the analyses were done on samples that are petrographically the least altered. Hence, the chemical and isotope analytical results are considered a reflection of the primary composition of the volcanic rocks in the following discussion, unless noted otherwise.

The Kahrizak volcanic rocks display a large range of composition, with  $\text{SiO}_2$  contents ranging from 48 to 73 wt.% and Mg# of the basaltic rocks ranging from 30 to 43 (Tables 1 and S2). They range from basalt, basaltic trachyandesite, trachyandesite, to rhyolite (Fig. 2; Le Bas et al., 1986). The basaltic and andesitic rocks tend to be calc-alkaline to high-K calc-alkaline whereas the rhyolite and trachydacitic rocks tend to be high-K calc-alkaline to shoshonitic (Pecerillo and Taylor, 1976). According to the classification of Draper and Johnson (1992), the basalts are high-alumina and low magnesian ( $\text{SiO}_2 = 48\text{-}51$  wt%;  $\text{Al}_2\text{O}_3 = 17\text{-}21$  wt%;  $\text{MgO} = 4\text{-}7$  wt%;  $\text{CaO} = 7.5\text{-}11$  wt%). The samples generally exhibit negative CaO, MgO,  $\text{Al}_2\text{O}_3$ ,  $\text{FeO}_t$ ,  $\text{TiO}_2$ , Ni, Sr, and Co correlations, but positive  $\text{K}_2\text{O}$ , Ba, and Th correlations with  $\text{SiO}_2$  (Fig. 5). All these variations are consistent with fractionation of the observed mineral phases olivine + clinopyroxene + plagioclase  $\pm$  hornblende  $\pm$  magnetic  $\pm$  apatite in the volcanic rocks.

Relative to normal mid-ocean ridge basalts (MORB; Sun and McDonough, 1989), the samples show selective enrichments in large-ion lithophile elements

(LILE; e.g., Sr, K, Rb, Ba, Th) and to a lesser extent in La and Ce, but depletions in some high field-strength elements (HFSE) such as Ta, Nb, Zr, and Hf (Fig. 6A). In general, the Kahrizak Mountains volcanic rocks and some Eocene arc volcanic rocks from the UDA (Omran et al., 2008), eastern Pontides and northern Anatolia show similar trace element concentration patterns.

The volcanic rocks are moderately enriched in light REE (Fig. 6B). That is, the samples show moderately fractionated, subparallel REE concentration patterns ( $\text{La}/\text{Lu}_{\text{Ch}} = 4-11$ ), indicating they may have come from a common, or at least, similar sources. Their REE concentrations increase with increasing differentiation from basalts to rhyolites. Moreover, the rhyolitic rocks display moderate negative Eu-anomalies (mean  $\text{Eu}_{\text{Ch}}/\text{Eu}^* = 0.512-0.726$ ), suggesting plagioclase fractionation.

### 4.3. Pb, Nd and Sr isotopic ratios

The samples analyzed show a limited range of Pb isotopic values ( $^{206}\text{Pb}/^{204}\text{Pb}_i = 18.431-18.688$ ,  $^{207}\text{Pb}/^{204}\text{Pb}_i = 15.533-15.616$ , and  $^{208}\text{Pb}/^{204}\text{Pb}_i = 37.647-38.380$ ; Table 2; Figs. 7A, B). In general, they appear to define linear arrays in Pb-Pb isotopic diagrams that either overlap and/or are parallel with the linear Pb isotopic fields of other Eocene lavas from Iran and of Tethyan basalts. The samples also have a narrow range of  $^{143}\text{Nd}/^{144}\text{Nd}_i$  (0.51257 – 0.51264) but their  $^{87}\text{Sr}/^{86}\text{Sr}_i$  (0.69265-0.70729) are heterogeneous and include usually low



values, clearly indicating seawater alteration (Rossel et al., 2014) and/or upper crustal assimilation (Alagna et al., 2010). Thus, a number of what were selected as the least altered samples may have also been variably affected by either alteration or assimilation, or both. In general, however, the  $^{87}\text{Sr}/^{86}\text{Sr}_i$  and  $^{143}\text{Nd}/^{144}\text{Nd}_i$  of the bulk of basaltic and intermediate samples are lower than the majority of Eocene lavas from Iran and Turkey, but overlap with the Eocene high-Ti gabbros from the Sanandaj–Sirjan zone in western Iran that also have low  $^{143}\text{Nd}/^{144}\text{Nd}_i$  and highly variable  $^{87}\text{Sr}/^{86}\text{Sr}_i$  (Fig. 7C).

Another significant feature of Kahrizak volcanic rocks is their measured Pb isotopic ratios, best exemplified by  $^{207}\text{Pb}/^{204}\text{Pb}$ , generally increase with increasing  $\text{SiO}_2$ , i.e., from basalt to trachyandesite and pyroclastic rocks (Fig. 8A). Their measured  $^{143}\text{Nd}/^{144}\text{Nd}$  and  $^{87}\text{Sr}/^{86}\text{Sr}$  also increase with increasing  $\text{SiO}_2$  (Figs. 8B, C).

## 5. Discussion

The Kahrizak volcanic rocks lack primitive rocks with  $\text{Mg}\# > 70$  and high compatible element abundances, e.g.,  $\text{Ni} > 200$  ppm,  $\text{Cr} > 400$  ppm, that can be considered to represent magmas derived directly from peridotitic mantle (e.g., Tatsumi and Eggins, 1995). However, the Kahrizak volcanic rocks have similar initial Pb and Nd isotopic ratios, clearly indicating their cogenetic nature. Thus,

before discussing the magma genesis and source composition of these volcanic rocks, we first examine whether they are related to one another by fractional crystallization, crustal assimilation, or a combination of both (AFC), and magma mixing.

### 5.1. Assimilation-fractional crystallization (AFC)

As noted earlier, major and trace element variations of the analyzed volcanic rocks are consistent with crystal fractionation from a common or very similar parental magmas. Specifically, the observed major and trace element variations (Fig. 5) are consistent with fractionation of olivine + clinopyroxene + plagioclase ± hornblende ± titanomagnetite in the basalts, plagioclase + hornblende + clinopyroxene ± titanomagnetite in the andesites, and plagioclase + quartz + biotite ± hornblende ± sanidine ± titanomagnetite ± apatite in the rhyolites and trachydacites. In rhyolitic samples, negative Eu anomalies are indicative of feldspar removal during fractionation. However, such an anomaly is not present in the basalts. This observation, together with high Al<sub>2</sub>O<sub>3</sub> and Ba contents and low La/Sr ratio (0.02-0.04), rule out any significant plagioclase fractionation in the basalts (Mattioli et al., 2006). On the other hand, in addition to their negative Eu anomalies (Fig. 6B), the Sr contents of the rhyolitic and pyroclastic rocks decrease with increasing SiO<sub>2</sub> (Fig. 5) and <sup>87</sup>Sr/<sup>86</sup>Sr (not shown). These compositional

features indicate the importance of plagioclase fractionation in the magmatic evolution of rhyolitic samples as the plagioclase partition coefficient for Sr is 1.8 in felsic rocks, but 0.1 for other trace elements (James, 1982). Moreover, the rhyolitic rocks are significantly low in  $P_2O_5$  and  $TiO_2$  relative to the basaltic rocks, indicating the potential roles of apatite and Fe–Ti oxide fractionation.

Significantly, the increase in REE concentrations of Kahrizak volcanic rocks from basalts to rhyolites (Fig. 6B) is accompanied by a decrease in  $Dy/Dy^*$  values, which are a measure of the concavity of a REE pattern (Davidson et al., 2013). All samples have  $Dy/Dy^* < 1.0$ , or have concave up REE patterns;  $Dy/Dy^*$  together with  $Dy/Yb$  decrease with differentiation (not shown), indicating that amphibole is a dominant fractionating phase as well (Davidson et al., 2013). This is significant because the Sr/Y ratios of the Kahrizak volcanic rocks are as high as 76 in the basalts. Although a Sr/Y of 76 is still low compared to those of ‘adakites’ that are purported to be melts from subducted oceanic basalts at high pressure where garnet is stable (e.g., Castillo, 2012), they overlap with adakitic lavas from Iran and Turkey, some of which have indeed been proposed as partial melts of subducted basaltic crust (e.g., Ghorbani et al., 2011). Combined with their relatively low  $La/Yb_N$  values ( $\leq 10$ ), the relatively high Sr/Y of Kahrizak basalts is most probably not due to partial melting of subducted basalt, but is also an indication of

hornblende fractionation from parental hydrous magmas (e.g., Castillo et al., 1999; Moghadam et al., 2016).

Many studies on the evolution of arc magmas have shown that assimilation of crustal material is an important process to modify the trace element and isotopic composition of mantle-derived magmas (e.g., Schiano et al., 2010). For example, contamination of primary magmas by mature and thickened paleo-arc crust was proposed to be an important factor in the collisional and post-collisional Tertiary magma evolution in north-central Turkey (Temizel et al., 2012). Magma mixing is also important. For instance, Amidi et al., (1984) have argued that mixing of basaltic and paligenetic felsic magmas was responsible for producing the calc-alkaline Eocene volcanic rocks in Central Iran. Therefore, it is essential to evaluate the effects of crustal contamination on mantle-derived magmas, e.g., through mixing of mantle- and crustal-derived melts, on the composition of the Kahrizak volcanic rocks.

It is noteworthy that the trace element composition of bulk continental crust is quite similar to the composition of typical arc magmas. Thus, some of the major compositional features of crustal contamination are similar to, or hard to distinguish from, those of arc magmas derived from a mantle source metasomatized by aqueous fluids and/or melts derived from a subducted basaltic crust and overlying sediments (Verdel et al., 2011). As mentioned earlier, however,

Kahrizak volcanic rocks show evidence of mineral-melt disequilibria including (1) Ca-rich plagioclase phenocrysts that exhibit oscillatory zoning and are rimmed with highly sodic plagioclase, sieve texture, corroded and rounded crystals, and early crystallization of Na-sanidine in the cores of oligoclase (Figs. 3A-C); (2) zoning and reaction rims in olivine phenocrysts; and (3) partial melting and reaction rim in clinopyroxenes (Figs. 3D-F). These textures are usually interpreted as a result of magma mixing (Tsuchiyama, 1985) although they may also occur due to rapid decompression, where heat loss is minor relative to ascent rate (Nelson and Montana, 1992). Disequilibrium textures in zoned pyroxene phenocrysts are results of either assimilation of continental crust, rapid fall of  $H_2O$  pressure or magma mixing (Sakuyama, 1979).

Many volcanic systems most probably experience rapid decompression and reduction in pressure is a simpler mechanism to produce the above-mentioned disequilibrium textures as it requires no addition of heat or mass. Rapid decompression may also operate in conjunction with magma mixing because magmas generally decompress as they ascend. Thus, some of the observed variations in incompatible elements (e.g., enrichments of K, Rb, and Ba) may have been the result of crustal assimilation and/or mixing (Fig. 5). Note that although most of these elements are fluid-mobile and, thus, their enrichments could be attributed to recent low temperature alteration, we reiterate that petrographic and

zeolite data indicate the alteration and volcanic eruptions were penecontemporaneously in a submarine environment. Moreover, Rb/Sr and Rb generally increase with measured  $^{87}\text{Sr}/^{86}\text{Sr}$  from basalt to rhyolite and trachydacite (Figs. 8D, E). As noted earlier, the Kahrizak volcanic rocks also display roughly positive correlations between measured radiogenic isotopes and  $\text{SiO}_2$  (Figs. 8A-C). As these ca. 40 Myr old volcanic rocks are neither temporally nor spatially stratified, and can be traced back to a common or very similar sources, the observed isotopic and trace element versus isotopic relationships of the bulk of the analyses most likely are not the result of alteration. Instead, these relationships indicate that crustal assimilation, in addition to fractional crystallization, was involved in the petrogenesis of the Kahrizak volcanic rocks.

One way to illustrate the possible effects of the AFC process in the Kahrizak volcanic rocks is shown in Figure 8E. The  $^{87}\text{Sr}/^{86}\text{Sr}$  ratio versus Rb relationship of the Kahrizak basaltic and intermediate rocks defines a trend that is close to or parallels the straight line that links the composition of an enriched mantle source and a continental crust end-member (Scharer, 1991). Along this line, an AFC model is constructed using a value of  $r = 0.4$  and  $K_{d(\text{Rb})} = 0.15$  for the mineral phases present in the basalts. This model indicates that the basalts are magmas coming from an enriched mantle and experience ca. 40% AFC with continental crust. Note that although some samples (one basalt and two rhyolites) plot far from

the line, indicating their Rb and  $^{87}\text{Sr}/^{86}\text{Sr}$  compositions are most probably altered, the overall trend of Kahrizak volcanic rocks still provides useful information on their petrogenesis.

In summary, geochemical and isotopic data indicating that fractional crystallization and assimilation of upper crust material have played an important role in the petrogenesis of the Kahrizak volcanic rocks.

## 5.2. Partial melting of the source

In this section, the concentrations of some fluid-immobile, incompatible trace elements (i.e., Ta, Th, Yb, Ti, and Nb) are used to constrain the relative degree of partial melting of the mantle source to produce the primary Kahrizak basaltic rocks (e.g., Gribble et al., 1996). In general, the moderately enriched REE patterns (Fig. 6B) and relatively low  $\text{La}/\text{Lu}_{\text{Ch}}$  (4-11) ratio of the Kahrizak basalts discount the role of garnet in their mantle source. The same is true for the Eocene arc volcanic rocks from northern Anatolia (Keskin et al., 2008), but not for those from the eastern Pontides (Temizel et al., 2012). Specifically, the Yb and  $\text{TiO}_2$  compositions of Kahrizak basaltic rocks are consistent with the presence of spinel in their mantle source (Fig. 9A). Model calculations indicate that the Kahrizak basalts can be produced by about 10% to 30% partial melting of a spinel lherzolite, similar to that of the Mariana Trough basalts. The Th/Ta and Nb/Ta ratios of

Kahrizak basalts also suggest a relatively high degree of partial melting at shallow depth (Fig. 9B). In detail, the Nb/Ta ratios, as proxy for degree of melting, of Kahrizak basalts are displaced toward high values similar to the rift basalts from the Mariana Trough at near-constant Th/Ta ratios ( $Nb/Ta = 12-19$ ). Thus, alteration-resistant, incompatible trace element data suggest that the Kahrizak volcanics were produced by high degree of partial melting in the spinel stability field.

### 5.3. Nature of the magma source

As discussed above, major and trace elements and Nd and Pb isotope data suggest a common source for the pyroclastics and lava flows in the Kahrizak Mountains. In  $^{206}Pb/^{204}Pb_i$  versus  $^{208}Pb/^{204}Pb_i$  and  $^{207}Pb/^{204}Pb_i$  plots (Figs. 7A, B), the Kahrizak volcanic rocks generally plot above the northern hemisphere reference line (NHRL) and toward the proposed enriched mantle II (EMII) mantle end-member (Zindler and Hart, 1986) as well as overlap with the field for global pelagic sediments (Ferguson and Klein, 1993). On the other hand, their depletion of Nb and Ta relative to LILEs can be attributed primarily to two processes: (1) addition of a LILE-enriched, but Nb and Ta poor subduction component to their mantle source or (2) assimilation of continental crust (Keskin et al., 2008). As also discussed above, assimilation of continental crust combined with fractional



crystallization (AFC), plus alteration, may have played a role in generating the compositional variation of the Kahrizak volcanics, but these are clearly evident mainly in the differentiated rocks.

In general, many geochemical features of Kahrizak volcanic rocks, particularly the relatively more primitive basalts, indicate that they are associated with subduction. For example, higher Th/Ce, Nb/Zr and Th/Nb ratios and lower Pb/Nd ratios in high-Al arc magmas relative to MORB are commonly interpreted as the result of sediment input into their mantle source (e.g., Gertisser and Keller, 2003). The Kahrizak volcanic rocks indeed display relatively high Al<sub>2</sub>O<sub>3</sub> contents (14.18-21.5 wt.%), Th/Ce (0.02-0.57), Nb/Zr (0.06–0.22) and Th/Nb (0.05-1.64) ratios, and low Pb/Nd (0.09–0.71) ratios, which indeed suggest the involvement of subducted sedimentary materials in their magma source, as is proposed for eastern Pontides (Temizel et al., 2012; Aydinçakir and Sen, 2013). The elevated P content of Kahrizak basalts (P<sub>2</sub>O<sub>5</sub> = 0.3-1.0%) can also be explained by the presence of sediment in their source (Raos and Crawford, 2004).

The source enrichment features in the Kahrizak volcanic rocks can be constrained through a plot of Th/Yb versus Ta/Yb (Fig. 10), as these elemental ratios can be effectively used to display source variation and crustal contamination (Pearce et al., 1990). In the plot, the Kahrizak volcanic rocks have high Th/Yb for given Ta/Yb ratios, suggesting they were derived from a mantle source containing

a subduction component. Significantly, the trachydacites and rhyolites have Ta/Yb values that indeed plot along lines for AFC and mixing with continental crust, consistent with the earlier discussion. In comparison, volcanic rocks from eastern Pontides, northern Anatolia and UDA exhibit elongated fields that are subparallel to the mantle metasomatism array. Thus, the geochemical characteristics of Kahrizak basaltic rocks were likely inherited from a mantle source that had been metasomatized by a subduction component. Correlatively, the low Nb/La and Ce/Pb ratios of the Kahrizak volcanics (Figs. 11A, B) indicate the influence of fluids dehydrated from subducted slab. Altogether these data suggest that the Kahrizak primary magmas were derived by partial melting of a mantle that had been metasomatized by fluids enriched with subduction component. However, this brings the question regarding the true or original nature of the mantle that was metasomatized by the subduction fluids.

Figure 11B also clearly shows that although the Th/La ratio of Kahrizak rhyolitic lavas have a bulk upper crust affinity, the basaltic lavas have a Th/La ratio that is higher than that of MORB (Plank, 2005). Moreover, the Kahrizak volcanic rocks have a limited and low range of  $^{143}\text{Nd}/^{144}\text{Nd}_i$  and most of the basaltic rocks have sub-bulk silicate Earth  $^{86}\text{Sr}/^{87}\text{Sr}_i$  (Fig. 7C). These values are within the field of sublithospheric mantle (e.g., Farmer, 1988; Arndt, 2013). In addition, Eocene volcanic rocks that occur in almost all of Central Iran and adjacent Urumieh-

Dokhtar and Sanandaj-Srjan provinces have a compositional range that is highly consistent with a lithospheric mantle source (Temizel et al., 2012; Keskin et al., 2008; Pearce et al., 1990). Consequently, the parental magmas of the Kahrizak volcanic rocks were most probably derived from a lithospheric mantle source that was subsequently enriched by fluids produced during earlier subduction events; the magmas further experienced AFC during ascent to the surface. The parental magmas were produced by moderately high degree of partial melting from such enriched lithospheric mantle source at a shallower depth (probably temperature less than 900 °C according to Figs. 4A, B) and possess the isotopic and elemental signatures that are commonly observed in north-central Iran and surrounding regions.

#### **5.4. Implications for the geodynamic evolution of north-central Iran and surrounding areas**

The Kahrizak Mountain volcanism is part of the Eocene magmatic flare-up in Central Iran, eastern Pontides and northern Anatolia. A variety of mechanisms have been proposed to explain the origin of the flare-up. These include volcanism related to subduction (Berberian and Berberian, 1981; Shahabpour 2007), slab-melting (Ghorbani et al., 2011), rifting and back-arc magmatism (Kazmin et al., 1986; Emami, 1981), slab break-off (e.g., Jahangiri 2007; Hassanzadeh et al.,

2009), and, lately, extensional arc flare-up due to slab roll-back following a period of flat-slab subduction (Verdel et al., 2011; Zhang et al., 2018). Among these proposals, we find that the slab roll-back scenario of Verdel et al. (2011) is most consistent with our data and literature data on Eocene magmatism, as summarized below. (1) The magmatism covers a wide area and was initiated prior to the Arabian-Eurasia continental collision. (2) It was temporally restricted to a ca. 17 Myr period. (3) Despite variations, the magmas overall have arc-like geochemical characteristics. (4) There is a trend in some localities for the magma composition to be initially calc-alkaline and, then, change to alkaline and shoshonitic.

Verdel et al.'s proposed mechanism is a four-stage model that started with a pre-conditioning of the arc lithosphere due to steep slab descent, followed by the spread of the pre-conditioning due to the Cretaceous flattening of the slab in the second stage. It was during the third stage that the Eocene volcanic rocks having major and trace element characteristics that are typical of continental arc magmatism were erupted. The magmas came from the decompression melting of the subduction-preconditioned arc lithospheric mantle or subcontinental lithospheric mantle (SCLM) due to lithospheric extension and crustal thinning accompanying slab rollback. Extension continued during the fourth stage in late Oligocene to Miocene, and the flare-up ended when the supply of preconditioned

SCLM was exhausted, and replaced by asthenosphere-derived, OIB-type volcanism.

We also concur that one of the key aspects of the appropriate mechanism is extensive preconditioning of the SCLM, but we want to emphasize that the preconditioning process most likely predated the 50 Myr flat-slab subduction (Verdel et al., 2011). The region experienced the subduction of the Neo-Tethys slab from at least the Early Jurassic until late Oligocene. The long-term subduction underneath Central Iran greatly affected the composition of the region's SCLM and asthenosphere, portions of which were trapped or wedged between subducting slabs and overlying plates. Fluids from the subducting slabs metasomatized the SCLM and the mantle wedges, which, as is typical along convergent margins, were the sources of arc magmas erupted in various parts of the region. As the Neo-Tethys closed, subduction slowed down and this may have increased the effectiveness of mantle fertilization (Verdel et al., 2011).

The uppermost asthenosphere and particularly the SCLM may have indeed partially melted due to decompression during lithospheric extension and crustal thinning (Verdel et al., 2011). However, the SCLM also could have been thermally perturbed because of the partial removal of its lithospheric root either by delamination or detachment of subducted slab (Pearce et al., 1990; Temizel et al., 2012; Bottrill et al., 2012).-In general, magmas formed during the flare-up were

dominated by basaltic melts from the metasomatized SCLM. Some of these magmas, however, assimilated crustal mineral while fractional crystallization was taking place to produce the rhyolitic high K-calc-alkaline to shoshonitic lavas. The assimilation process became more pronounced towards the end of the flare-up when magmatic input from the SCLM decreased and asthenosphere-derived, OIB-type magmas were added (Verdel et al., 2011). The end of the flare-up occurred during the 'soft' Arabia-Eurasia collision that is commonly estimated to be in late Eocene (Bottrill et al., 2012), although arc and associated extensional magmatism continued until late Oligocene to Miocene (Verdel et al., 2011; Zhang et al., 2018). Thus, partial melting of metasomatized lithospheric mantle accompanied by assimilation of the crust generated the Eocene calc-alkaline to high-k calc-alkaline and shoshonitic basaltic to rhyolitic rocks in north-central Iran and surrounding areas (see also, e.g., Temizel and Arslan, 2008; 2009).

## **6. Conclusions**

(1) The Kahrizak Mountains in north-central Iran generally consists of Eocene basaltic to rhyolitic lavas interbedded with pyroclastic deposits formed in a shallow marine environment. The lavas include basalt, basaltic trachyandesite/andesite, trachydacite and rhyolite that can be classified into calc-alkaline basaltic group and

high K-calc-alkaline to shoshonitic trachydacite-rhyolite group. The petrography of the volcanic rocks shows evidence of mineral-melt disequilibria.

(2) Despite some evidence of alteration, major and trace element variations indicate a general increase in incompatible trace elements with differentiation, consistent with the effects of fractional crystallization. The disequilibrium textures and complex compositional variations of minerals and other geochemical data are consistent with contamination or assimilation of crustal rocks. Overall, trace element variations suggest that the volcanic rocks in north-central Iran were derived from a relatively enriched lithospheric mantle source and had assimilated continental crust materials.

(3) The distinct enrichment in LILE and to a lesser extent in light REE, but depletion in HFSE of the lavas are similar to those of Paleogene volcanic rocks in eastern Pontides (northeastern Turkey) and northern Anatolia (north-central Turkey). Moreover, the rhyolitic rocks have compositional similarities with the upper crust.

(4) The basaltic rocks have a narrow range of Pb and Nd isotopic composition that indicates a common, fairly homogeneous source. However, the measured Sr, Pb and, to a certain extent, Nd isotopic ratios correlate with degree of differentiation; these suggest an additional enriched component was involved in the petrogenesis of the volcanic rocks.

(5) Data indicate that the original source of the basaltic magmas was the lithospheric mantle that was previously metasomatized by fluids derived from the subducted oceanic slabs. Partial melting of such a source produced the bulk of the calc-alkaline basaltic group. Assimilation of continental crust and mixing with the basaltic magmas produced the intermediate to rhyolitic group.

(6) The Eocene volcanic rocks in north-central Iran and surrounding areas were derived through partial melting of the subduction-preconditioned subcontinental lithospheric mantle and asthenosphere during extension and thinning of the lithosphere at the waning stages of the closing of the Neo-Tethys Ocean.

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Fig. 1. (A) Simplified geological map of the Iran and Turkey region showing major sutures and continental blocks (modified after Kheirkhah et al., 2009; Whitechurch et al., 2013). The main Paleo- and Neo-Tethyan sutures are represented by heavy lines with filled triangles and the Eocene volcanic rocks in Iran by green areas (after Agard et al., 2011). Upper Cretaceous-Eocene volcanic rocks and Eocene volcanic-sedimentary successions along the Izmir-Ankara suture (Keskin et al., 2008) across Turkey are also shown. The location of study area is marked by blue square south of Tehran. (B) Simplified geological map of the Kahrizak area (after Afaghi et al., 1986).

Fig. 2. Total alkali versus silica classification diagram (Le Bas et al., 1986) for the Kahrizak (KH) volcanic rocks. The volcanic rocks generally plot along the alkaline-subalkaline divide of Irvine and Baragar (1971). Also plotted for reference are Eocene volcanic rocks from Urumieh-Dokhtar (UD) (Omrani et al., 2008; Verdel et al., 2011), Eocene volcanic rocks from Sabzevar (SZ) (Moghadam et al., 2016), eastern Pontides (Aydincakir and Sen, 2013; Temizel et al., 2012) and northern Anatolia (Keskin et al., 2008).

Fig. 3. Photomicrographs of the Kahrizak volcanic rocks displaying some of their textures and mineralogy. (A) Zoning of plagioclase in basalt; (B) and (C) plagioclase sieve texture in andesite; (D), (E), and (F) zoning and reaction rim (marked with arrows) in clinopyroxenes in andesitic-basalt and basalt, respectively. Abbreviations: Cpx: clinopyroxene; Plg: plagioclase; Zeo: zeolite; Opa: opaque; Chl: chlorite.

Fig. 4. (A, B) An-Ab-Or triangular plots showing the composition of feldspars in the Kahrizak volcanic rocks. The temperature curves are from Elkins and Grove (1990). Note that some of the plagioclases plot on curves that partially melt as high as 900°C. (C) Anorthite content of the plagioclases plotted against distance from the core. (D) Clinopyroxene (Morimoto, 1988), olivine (E) and hornblende (Hawthorne, 1983) (F) classification diagrams for the Kahrizak volcanic rocks. In Figure 4E: 1= edenitic hornblende; 2= ferro-edenitic hornblende; 3= magnesian hastingsitic hornblende; 4= hastingsitic hornblende.

Fig.5. SiO<sub>2</sub> (wt%) versus major oxide (wt%) and select trace elements (ppm) variation plots for the Kahrizak volcanic rocks. See text for discussion.

Fig. 6. (A) Normal-MORB normalized multi-element and (B) chondrite-normalized REE spider diagrams for the Kahrizak volcanic rocks (normalizing values are from Sun and McDonough, 1989). The upper continental crust (Rudnick and Gao, 2003) is plotted for comparison. Eocene volcanic rocks from eastern Pontides and northern Anatolia are from Temizel et al. (2012) and Keskin et al. (2008), respectively.

Fig. 7. (A)  $^{206}\text{Pb}/^{204}\text{Pb}_i$  versus  $^{207}\text{Pb}/^{204}\text{Pb}_i$  and (B)  $^{208}\text{Pb}/^{204}\text{Pb}_i$  for the Kahrizak volcanic rocks. Data for Urumieh-Dokhtar are from Mirnejad et al. (2011), Eocene volcanic rocks for Sabzevar (SZ) are from Moghadam et al. (2016), Tethyan basalts and global pelagic sediments are from Ferguson and Klein (1993) and Pearce et al. (1995), Mariana Trough are from Straub and Zellmer (2012), and upper continental crust (UCC) and lower continental crust are from Hofmann (1997). (C)  $^{87}\text{Sr}/^{86}\text{Sr}_i$  versus  $^{143}\text{Nd}/^{144}\text{Nd}_i$  plot for the Kahrizak volcanic rocks. Eocene lavas from Iran (Urumieh-Dokhtar from Omrani et al., 2008; also gabbro in Sanandaj-Sirjan from Deevsalar et al., 2017), and from Turkey (Temizel et al., 2012; Aydincakir and Sen, 2013; Kaygusuz, 2009; Keskin et al., 2006; Pearce et al., 1990; Kurt et al., 2008; Varol et al., 2007; Pamić et al., 1995) are shown for comparison. Fields for MORB and mantle array are from Wilson (1989) and McCulloch et al. (1994); proposed EMI, EMII, HIMU, and DM mantle end-

members and bulk Earth compositions are from Zindler and Hart (1986), the Northern Hemisphere Reference Line (NHRL) is from Hart (1984), and the seawater alteration trend is from McCulloch et al. (1994). CAV=calc-alkaline volcanics. Two Kahrizak samples with very low  $^{87}\text{Sr}/^{86}\text{Sr}_i$  were not plotted.

Fig. 8. (A) Measured  $^{207}\text{Pb}/^{204}\text{Pb}$ , (B)  $^{143}\text{Nd}/^{144}\text{Nd}$  and (C)  $^{87}\text{Sr}/^{86}\text{Sr}$  versus  $\text{SiO}_2$  (wt.%) plots for the Kahrizak volcanic rocks showing possible fractional crystallization (FC) and assimilation-fractional crystallization (AFC) trends (SCM = subcontinental mantle). (D) measured  $^{87}\text{Sr}/^{86}\text{Sr}$  versus  $\text{Rb}/\text{Sr}$  and (E)  $\text{Rb}$  plots for the Kahrizak volcanic rocks. Solid line represents mixing between enriched SCLM composition and continental crust from Scharer (1991). Numbers next to the mixing line correspond to % contribution from sediments. Bulk rock Rb partition coefficient ( $K_d$ ) value used in the AFC model was calculated from individual mineral partition coefficient values from the GERM Partition Coefficient ( $K_d$ ) Database (<https://earthref.org/KDD/>)

Fig. 9. Plots to estimate the degrees of partial melting that generated the Kahrizak basalts. (A) Ti versus Yb with a fixed primitive upper mantle (PUM) composition (Gribble et al. 1996). This plot assumes that the Mariana Trough backarc basin basalts (BABB) were produced by 6% to 24% partial melting in the spinel lherzolite stability field, with the highest value of 34% representing an arc-like basalt (Pearce and Stern, 2006). (B) Th/Ta (a proxy for the deep subduction component) versus Nb/Ta (a proxy for degree of melting and mantle depletion). Central and southern Mariana arcs = CIP and SSP, respectively and northern Mariana arc = NSP. See text for discussion.

Fig. 10. Th/Yb versus Ta/Yb diagram (after Pearce et al., 1990) for the Kahrizak volcanic rocks. Average continental crust = av. CC from Taylor and McLennan (1985). Data for Urumieh-Dokhatr are from Omrani et al. (2008) and Verdel et al. (2011); those from Turkey are from Aydinçakir and Sen (2013), Temizel et al. (2012) and Keskin et al. (2008). See text for discussion.

Fig. 11 (A) Nb/La versus Ba/Rb (N-MORB and E-MORB values are from Wang et al., 2004) and (B) Th/La versus Ce/Pb (reference ratio for Th/La from Plank, 2005; Ce/Pb reference ratio from Wang et al., 2004) diagrams for the Kahrizak volcanic rocks. See text for discussion.

Table 1. Major and trace element contents of representative volcanic rocks from the Kahrizak Mountains, north-central Iran.

Sample	K.91 .14	K.91 .31	K. B.3	K.E. 30	K.E L.1	K.E T.5	K.S. 18"	K.SH .21	K.Y. 10	S0- 19	* +/-
SiO <sub>2</sub>	68.3	67.4	50.	57.0	70.0	50.4	58.6		72.5	61.1	0.8
(wt%)	5	0	11	3	0	5	5	49.50	6	3	8
TiO <sub>2</sub>	0.47	0.33	1.0 6	1.44	0.78	1.13	0.90	1.23	0.39	0.69	0.0 1
Al <sub>2</sub> O <sub>3</sub>	15.4 1	16.1 8	17. 29	16.3 9	14.8 4	18.4 7	18.4 3	19.23	14.1 8	13.9 5	0.2 8
Fe <sub>2</sub> O <sub>3</sub>	2.14	1.40	3.9 5	3.96	2.18	4.00	3.15	4.11	1.33	7.47	0.1 6
MnO	0.08	0.08	0.1 5	0.10	0.13	0.15	0.07	0.14	0.05	0.13	0.0 1
MgO	0.56	0.20	6.7 3	2.60	0.09	4.62	2.27	4.21	0.63	2.88	0.0 8



CaO	1.36	3.42	9.7 2	6.17	0.34	10.2 3	4.65	10.23	2.36	6.00	0.1 6
Na <sub>2</sub> O	3.58	4.78	2.5 8	3.54	1.85	2.89	4.71	2.88	3.75	4.11	0.2 3
K <sub>2</sub> O	6.49	5.06	1.3 3	3.07	8.07	1.18	3.41	1.36	3.52	1.29	0.0 7
P <sub>2</sub> O <sub>5</sub>	0.11	0.15	0.3 3	0.82	0.15	0.34	0.59	0.35	0.09	0.32	0.0 1
LOI	2.32	2.58	2.9 0	2.50	4.40	2.50	3.10	3.10	4.40	1.94	
Total	100. 05	100. 02	99. 99	99.9 8	99.9 5	99.9 9	99.9 9	99.96	100. 00	99.9 1	
Sc (ppm)	5.00	7.00	29. 00	20.0 0	11.0 0	30.0 0	10.0 0	29.00	6.00	27.0 0	1.1 0
V	36.0	31.0	264	181	19.0	284	64.0	307	29.0	165	19. 0
Co	4.10	3.70	48. 10	27.3 0	22.5 0	43.2 0	20.0 0	43.20	17.9 0	24.0 0	2.0 0
Ni	3.00	2.00	32. 90	1.50	0.30	13.0 0	0.30	15.30	0.50	470. 00	44. 00

Rb	161	141	26. 4	89.5	194	21.9	78.3	30.2	143	19.5	1.8 0
Sr	68.4	198	603	559	77.8	660	587	690	407	317	30. 0
Zr	60.0	95.0	70. 3	197	252	87.3	199	85.3	191	112	9.2
Nb	12.7	13.3	5.5	15.2	19.0		15.9		11.9	69.0	8.0
	0	0	0	0	0	6.60	0	7.70	0	0	0
Cs	2.60	2.20	1.0 0	1.80	1.20	0.30	1.00	0.30		48.0 4.50	0.3 0
Ba	887	759	382	694	675	338	788	320	697	486	20
La	39.8	32.4	15.	37.8	35.1	16.7	37.3		26.9	71.3	5.8
	0	0	70	0	0	0	0	16.70	0	0	0
Ce	70.7	55.3	32.	81.2	76.9	36.2	77.5		50.8	161.	8.4
	0	0	40	0	0	0	0	36.00	0	00	0
Pr	7.79	6.22	3.9 2	9.39	8.46	4.32	8.82	4.50	5.23		19.4 1.4 0 0
Nd	26.8	22.7	16.	39.4	31.9	17.9	35.1		19.4	75.7	6.4
	0	0	80	0	0	0	0	19.80	0	0	0
Sm	5.39	4.26	4.5	8.70	6.90	4.10	7.50	4.70	3.70	13.7	1.0

			0							0	0
Eu	0.87	1.02	1.3 2	2.25	1.37	1.33	1.99	1.54	0.74	3.81	0.2 4
Gd	4.94	4.32	4.1 6	8.55	5.94	4.43	6.67	4.50	3.37	10.5 3	0.7 8
Tb	0.65	0.65	0.6 8	1.26	0.98	0.61	0.94	0.69	0.60	1.41	0.0 8
Dy	3.60	3.73	3.8 4	7.37	5.99	4.09	5.57	4.20	3.27	7.50	0.3 8
Ho	0.76	0.81	0.7 4	1.36	1.24	0.77	1.07	78.00	0.66	1.39	0.1 7
Er	2.23	2.42	2.0 6	3.94	3.69	2.21	3.30	2.34	2.02	3.78	0.4 8
Tm	0.45	0.41	0.3 0	0.60	0.57	0.31	0.48	0.32	0.33	0.55	0.0 5
Yb	2.74	2.75	1.9 2	3.83	4.05	2.15	3.25	2.34	2.37	3.55	0.1 7
Lu	0.46	0.43	0.2 8	0.59	0.62	0.32	0.54	0.34	0.40	0.53	0.0 4
Hf	1.86	2.51	1.9	5.50	8.10	2.60	5.00	2.40	5.40	3.10	0.1

			0								3
			0.4								0.6
Ta	0.85	0.75		1.10	1.40	0.40	0.90	0.50	1.00	4.90	0
			0								0
		16.1	1.6		13.4						
Pb	5.80			3.70		2.20	5.80	3.00	6.80		
		0	0		0					-	-
		15.3	2.0		15.0				12.8	13.0	1.1
Th	20.8			8.50		2.40	5.40	1.90			
		0	0		0				0	0	0
			0.6							19.4	2.4
U	3.60	3.34		2.40	4.00	0.70	1.60	0.60	4.80		
			0							0	0

\*at 95% confidence interval. Additional information on accuracy and precision can be found at [www.acmelab.com](http://www.acmelab.com).

Table 2. Sr, Nd and Pb isotopic ratios of volcanic rocks from the Kahrizak Mountains, north-central Iran.

Samp	$^{87}\text{Sr}$	$(^{87}\text{Sr})$	$^{143}\text{Nd}$	$(^{143}\text{Nd})$	$\epsilon$	$^{206}\text{Pb}$	$(^{206}\text{Pb})$	$^{207}\text{Pb}$	$(^{207}\text{Pb})$	$^{208}\text{Pb}$	$(^{208}\text{Pb})$
le	$^{86}\text{S}$	$^{86}\text{Sr}$	$d/^{144}$	$/^{144}\text{Nd}$	N	$b/^{204}$	$/^{204}\text{Pb}$	$b/^{204}$	$/^{204}\text{Pb}$	$b/^{204}$	$/^{204}\text{Pb}$
name	r	)i	Nd	)i	d	Pb	)i	Pb	)i	Pb	)i
K.ET.	0.7	0.70	0.51	0.512	-	18.7	18.63	15.6	15.60	38.7	38.36

5	074	728	2689	580	0.	58	6	23	5	81	7
	49	5			1						
					2						
					-						
	0.7										
K.SH	0.70	0.51			0.	18.6	18.52	15.5	15.57	38.5	38.27
	051										
.21	488	2687	0.512	2	03	6	89	8	93	7	
	03										
	7		574	4							
					-						
	0.7										
K.E.3	0.70	0.51	0.512	0.	18.6	18.43	15.6	15.56	38.7	37.83	
	053										
0	455	2671	566	4	80	1	05	9	04	2	
	46										
	7			1							
					-						
	0.7										
K.EL.	0.69	0.51		0.	18.5	18.43	15.5	15.53	38.4	38.06	
	049										
1	264	2686	0.512	0	51	6	50	3	89	4	
	70										
	8		583	6							
	0.7	0.70		0.							
K.91.		0.51			18.7	18.63	15.6	15.61	38.7	38.38	
	066	310	0.512	8							
31			2720		13	3	28	6	41	0	
	18	6		631	7						
	0.7	0.69	0.51	0.512	0.	18.9	18.68	15.6	15.61	39.0	37.64
K.91.											
14	079	633	2718	623	7	26	8	50	5	09	7

	45	7			0						
	0.7	0.70			0.						
K.S.1			0.51	0.512		18.6	18.50	15.6	15.59	38.6	38.29
8"	052	455			0						
			2692	591		06	0	10	5	47	3
	15	7			8						
					-						
	0.7										
		0.70	0.51	0.512	0.	18.6	18.47	15.6	15.58	38.6	38.20
K.B.3	049										
		470	2699	572	2	16	2	10	9	81	6
	16										
		0			9						
	0.7	0.70			1.						
K.Y.1			0.51	0.512		18.7	18.46	15.6	15.59	38.7	38.06
0	061	446			0						
			2729	639		33	2	33	3	84	9
	95	3			1						

Strontium, Nd and Pb isotope ratios were analyzed using a 9-collector, Micromass Sector 54 thermal ionization mass spectrometer (TIMS). Total procedural blanks are 35 pg for Sr, 10 pg for Nd and 60 pg for Pb. Strontium isotopic ratios were fractionation-corrected to  $^{86}\text{Sr}/^{88}\text{Sr} = 0.1194$  and are reported relative to  $^{87}\text{Sr}/^{86}\text{Sr} = 0.710254 \pm 0.000018$  (n=22) for NBS 987 during the period of analysis. Neodymium isotopic ratios were measured in oxide form, fractionation corrected to  $^{146}\text{NdO}/^{144}\text{NdO} = 0.72225$  ( $^{146}\text{Nd}/^{144}\text{Nd} = 0.7219$ ) and are reported relative to  $^{143}\text{Nd}/^{144}\text{Nd} = 0.511856 \pm 0.000016$  (n = 19) for the La Jolla Nd Standard during

the period of analysis. Lead isotopic ratios were analyzed using the double-spike method to correct for mass fractionation during analyses; separate measurements of spiked and unspiked samples were made on different aliquots from the same dissolution. The SBL-74  $^{207}\text{Pb}$ – $^{204}\text{Pb}$  double-spike from the University of Southampton was used. During the analysis period, the method produced the following results for NBS981:  $^{206}\text{Pb}/^{204}\text{Pb} = 16.930 \pm 0.002$ ,  $^{207}\text{Pb}/^{204}\text{Pb} = 15.490 \pm 0.003$  and  $^{208}\text{Pb}/^{204}\text{Pb} = 36.700 \pm 0.009$  ( $n = 11$ ).  $2\sigma$  precisions for individual runs are better than these.

## HIGHLIGHTS

- The Kahrizak volcanic rocks are part of the Eocene magmatic flare-up in Iran
- The Kahrizak volcanic rocks have a lithospheric source but arc-like geochemistry
- The flare-up initiated prior to Arabian-Eurasia collision and lasted for ca. 17 Myr
- Subduction fluid pre-conditioning of the lithospheric mantle preceded the flare-up
- Flare-up due to melting of pre-conditioned mantle during lithospheric extension



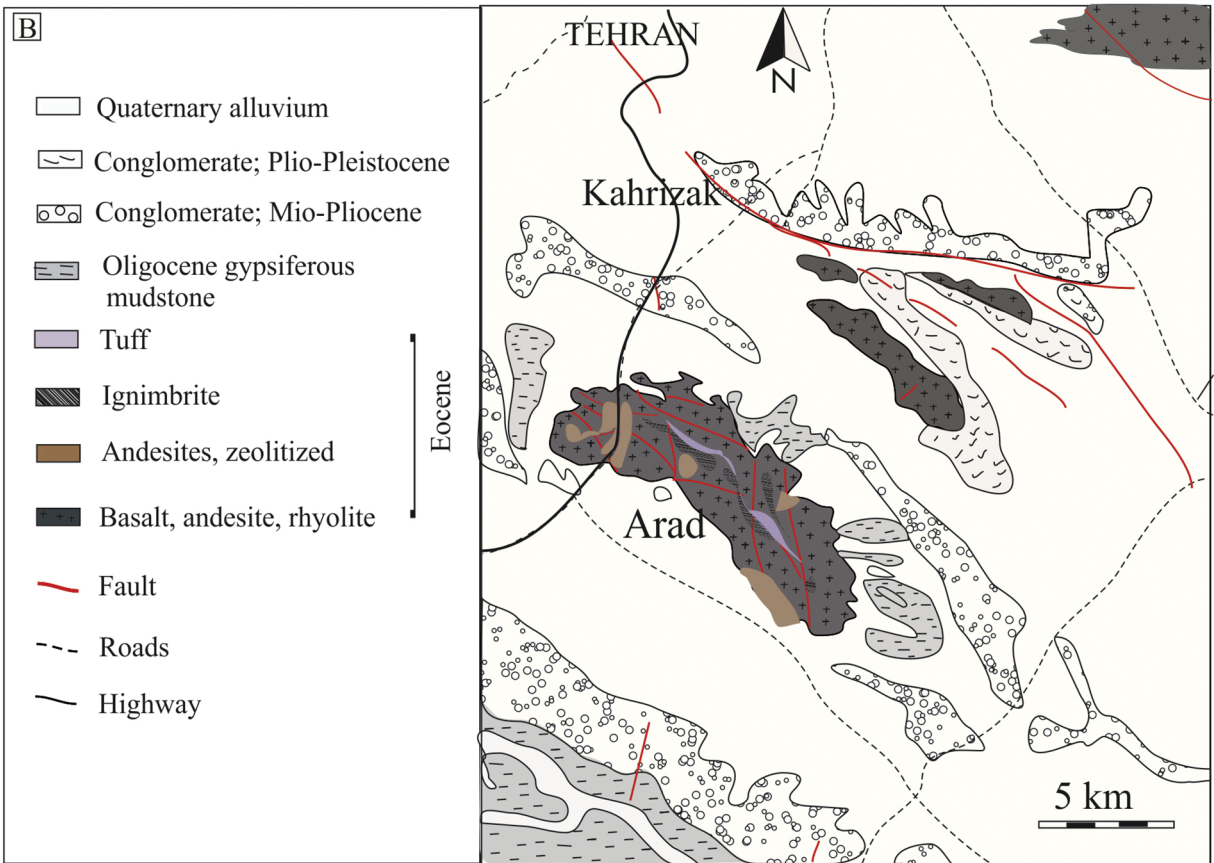
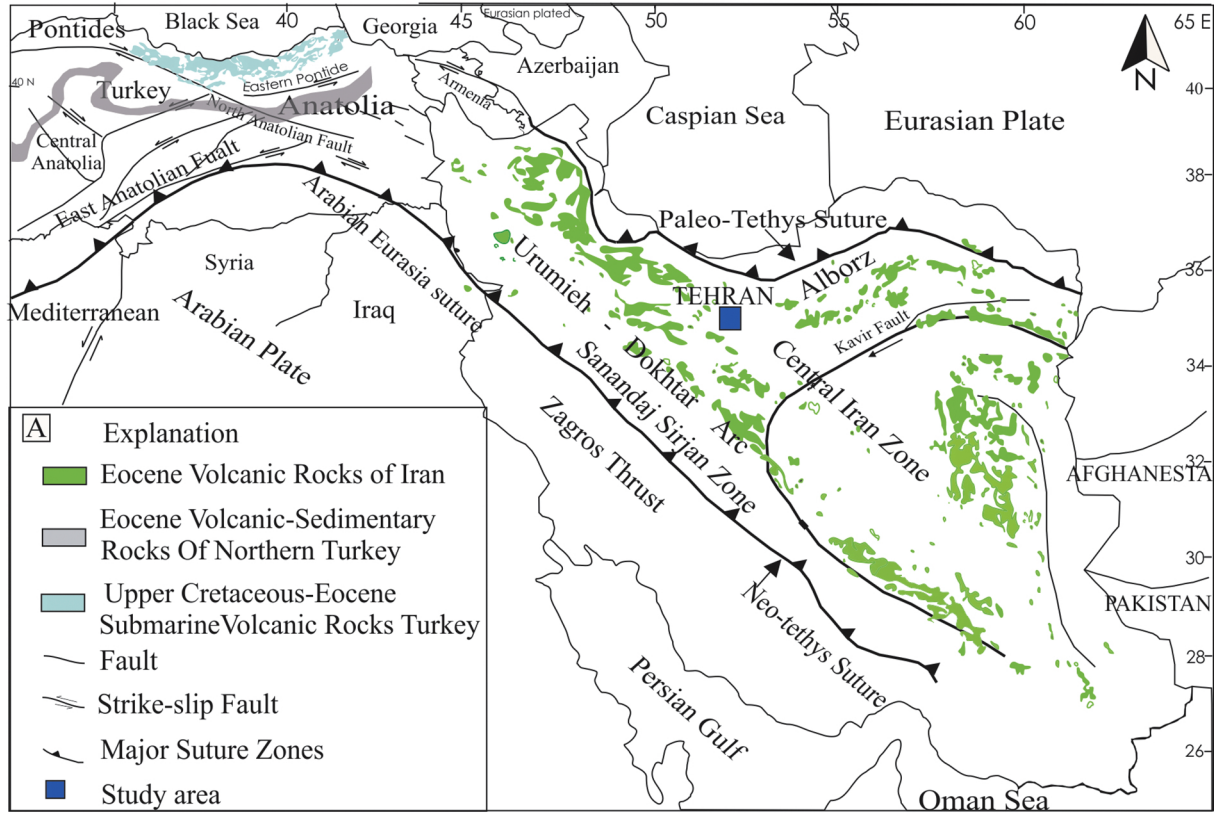


Figure 1

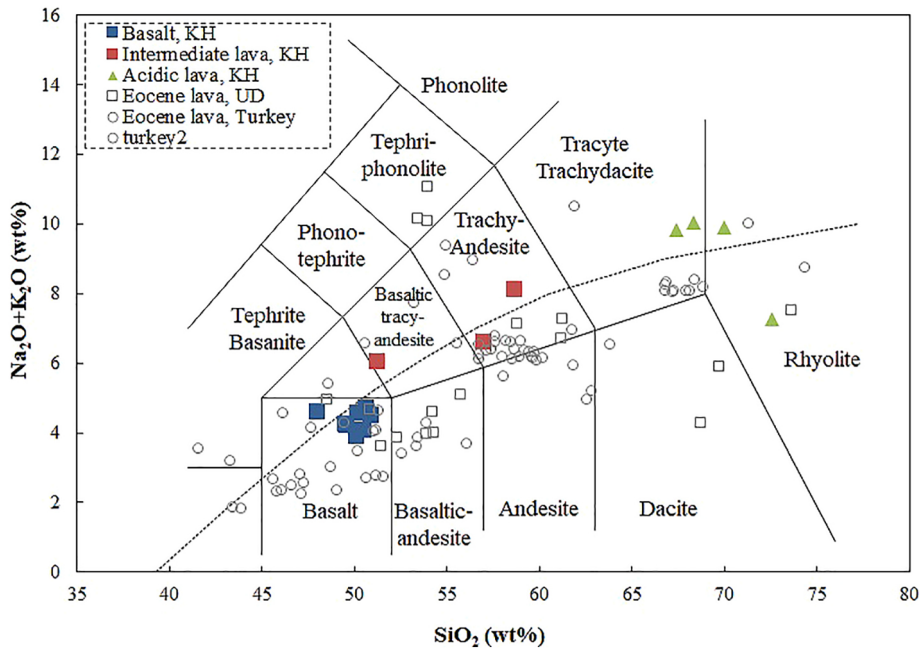


Figure 2

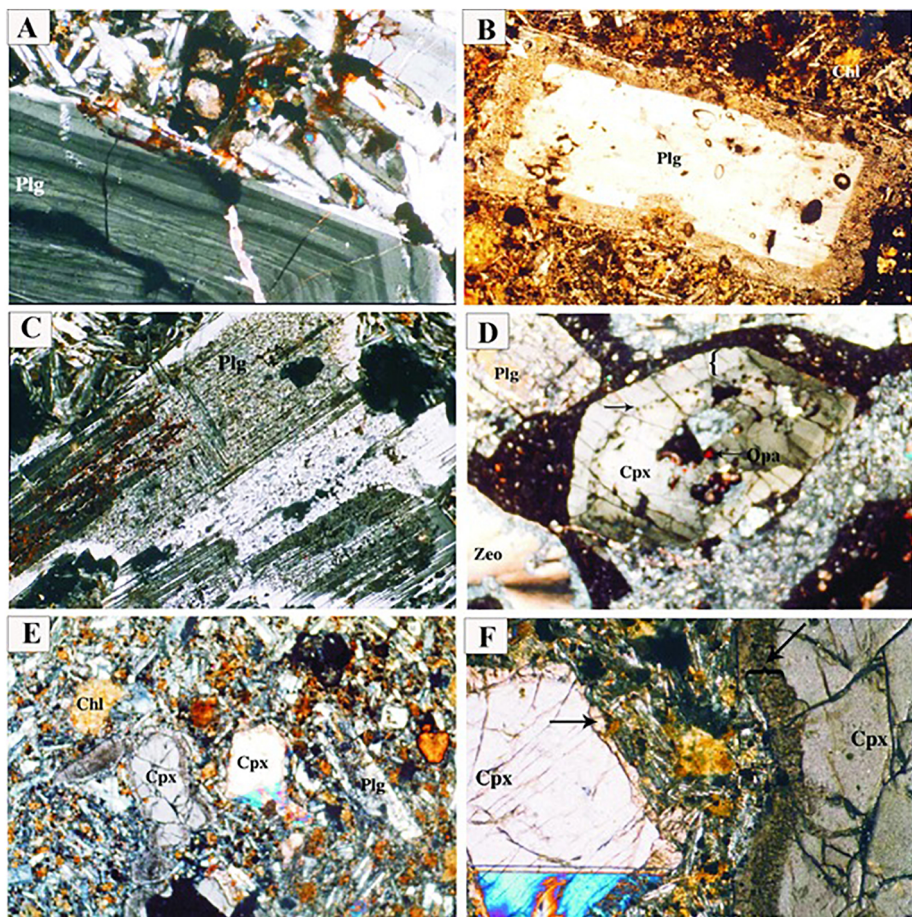
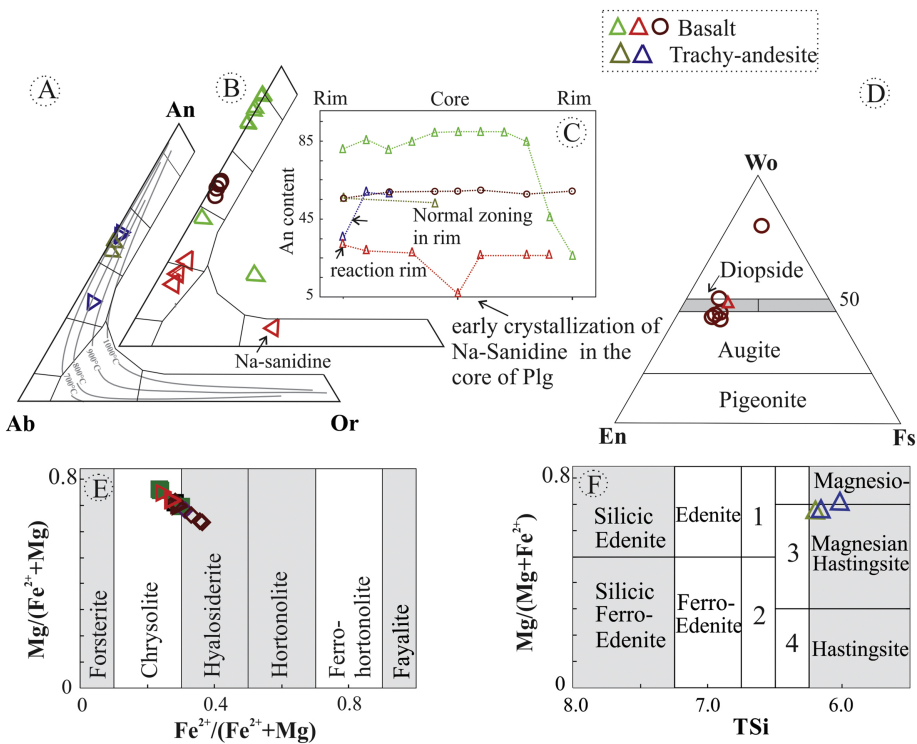


Figure 3



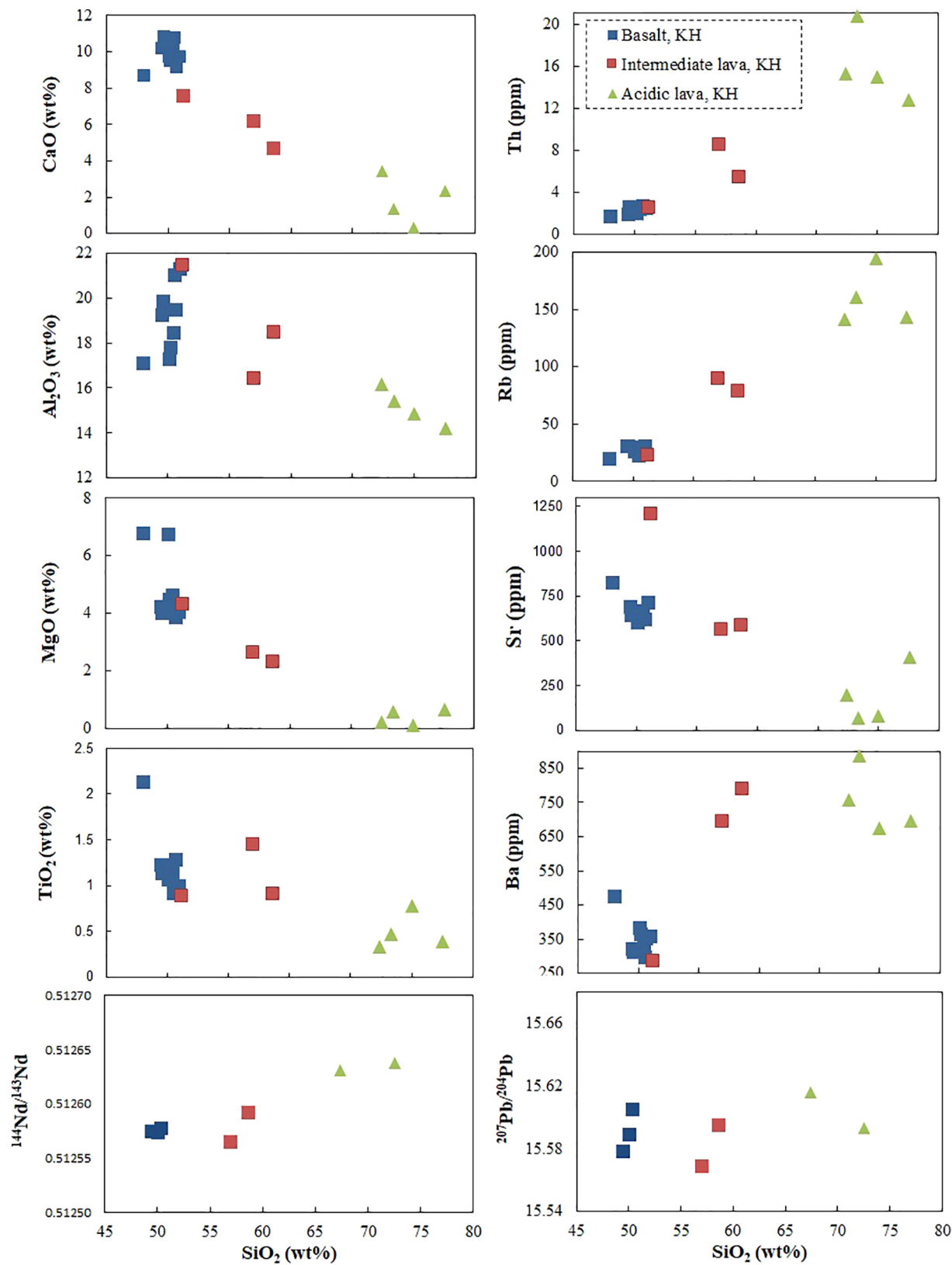


Figure 5

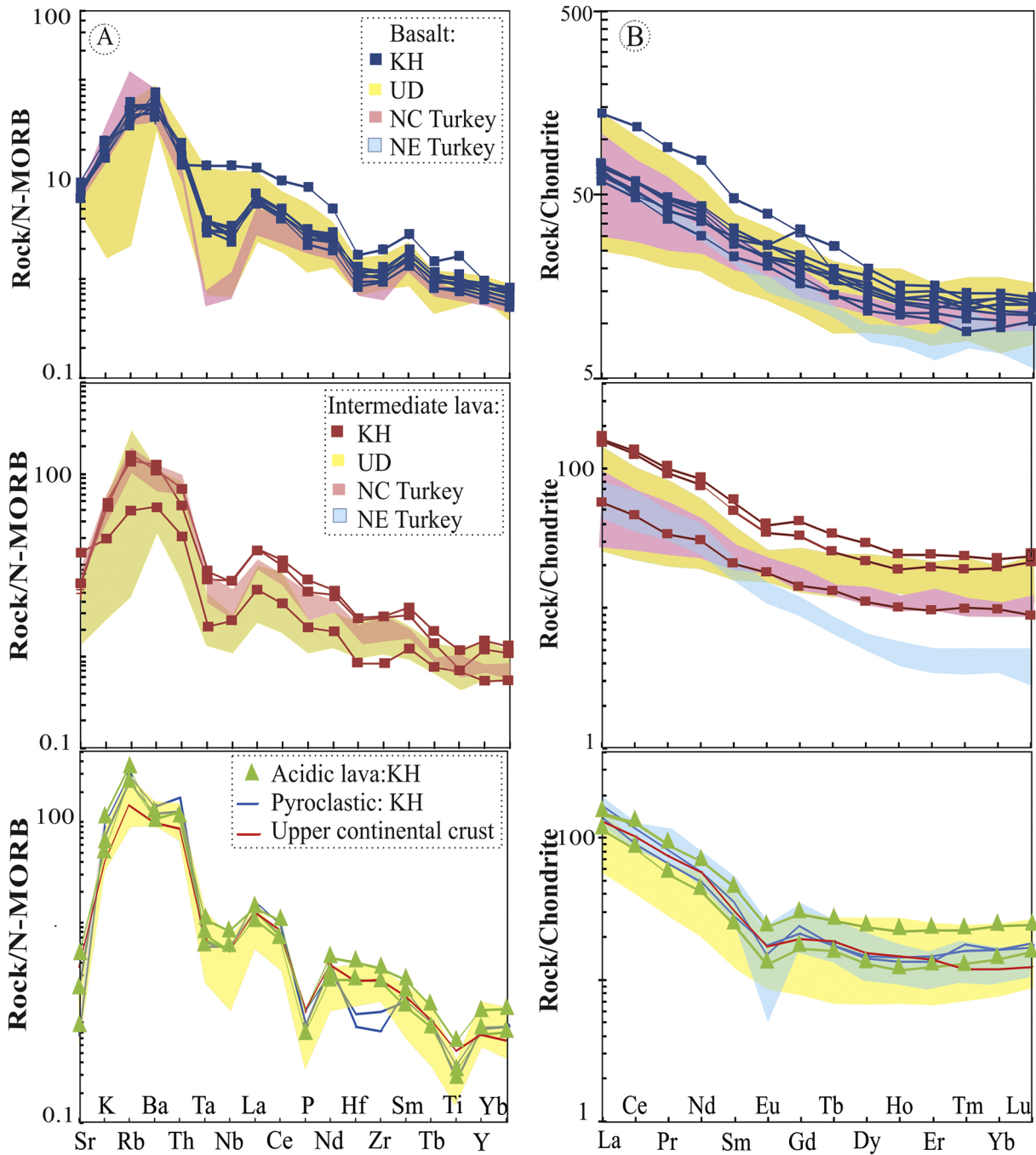


Figure 6

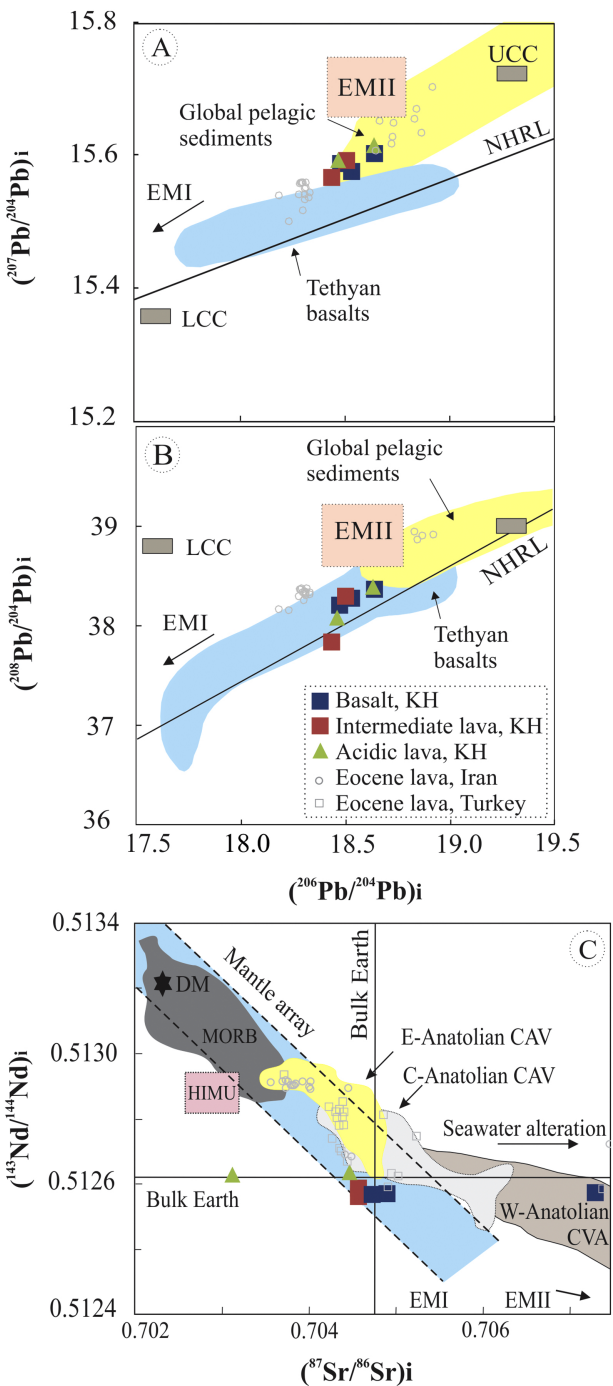


Figure 7

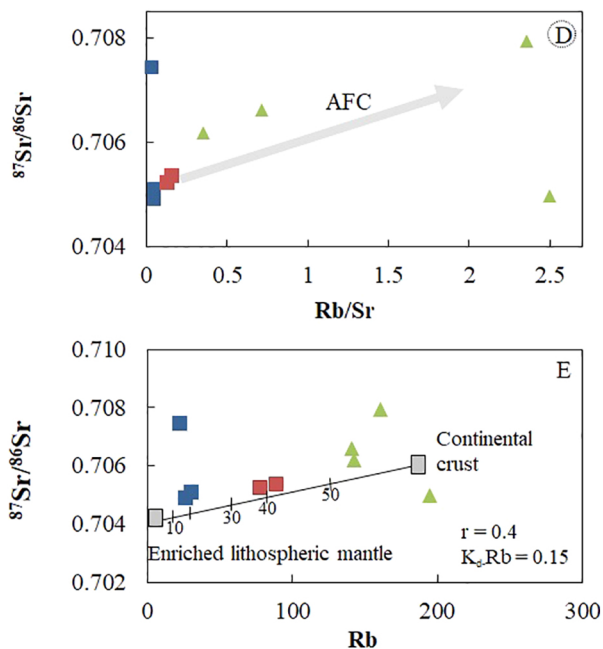
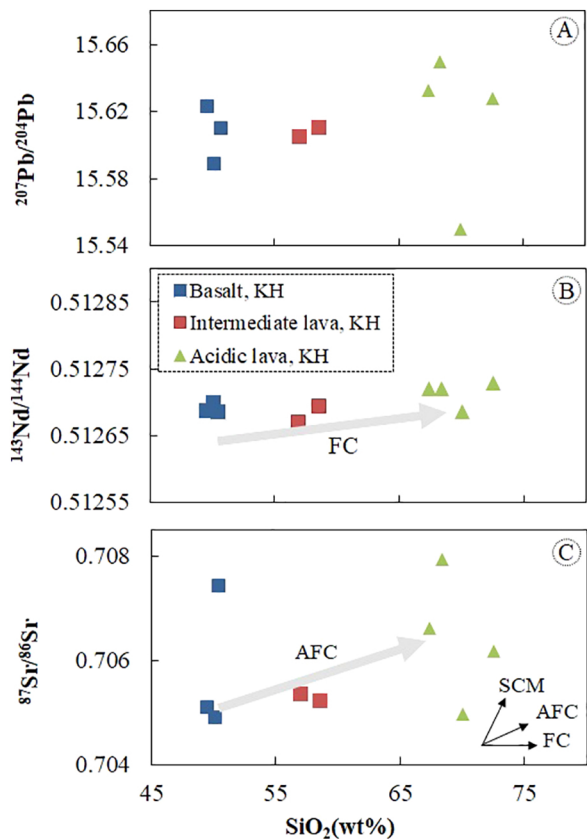


Figure 8



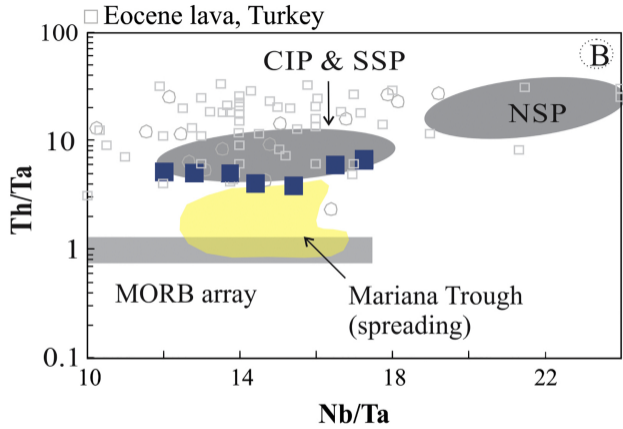
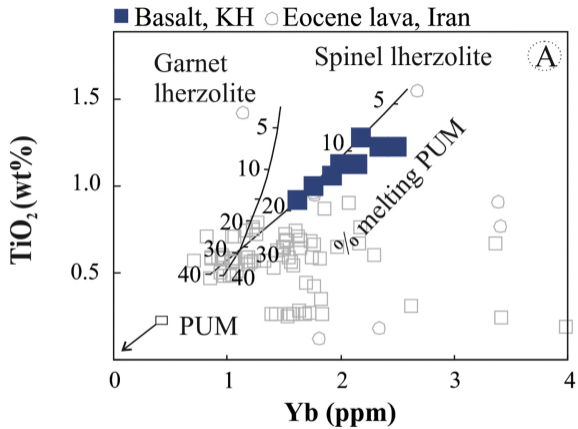


Figure 9

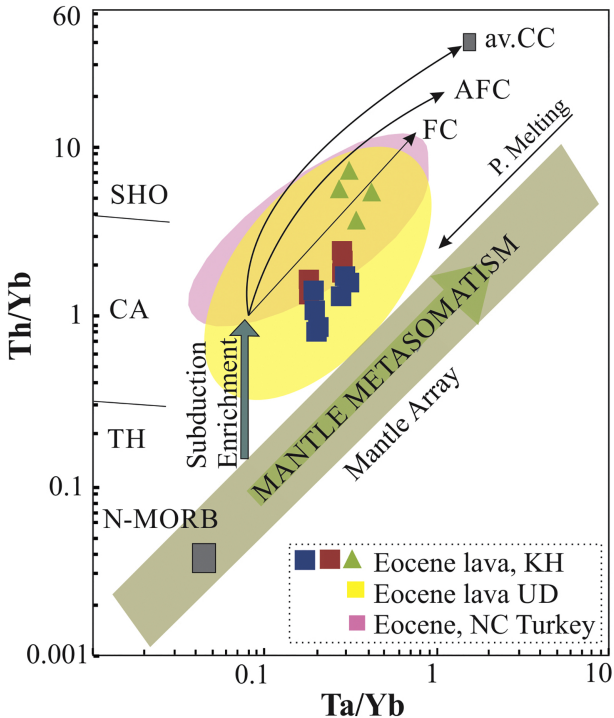


Figure 10

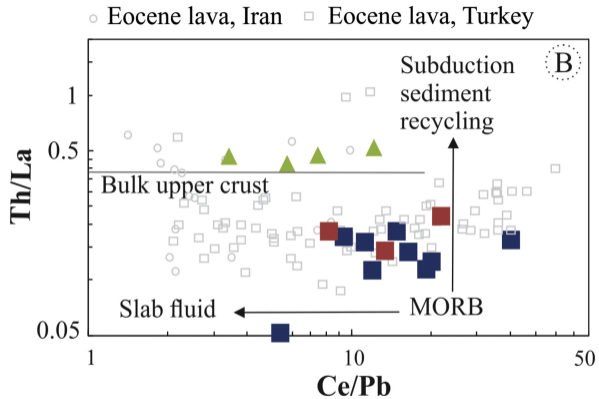
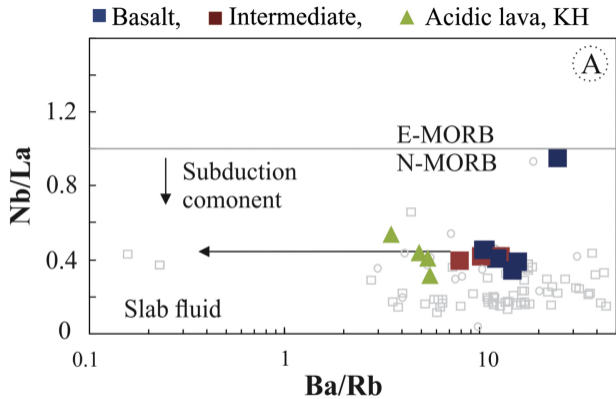


Figure 11