Geochemistry of skarn and porphyry deposits in relation to epithermal mineralization in the Arasbaran metallogenic zone, NE Tabriz, Iran

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The Arasbaran metallogenic zone in northern Iran is part of the Alborz–Azerbaijan magmatic zone, which developed along the southern margin of Eurasia during the Early Mesozoic–Late Cenozoic. This region hosts precious and base metal mineralization, including porphyry, skarn, and epithermal copper, molybdenum, and gold deposits. Rare earth element variations across all the deposits are similar, indicating a similar source for these elements. The north-west trending belt comprising the Nabiyan to the Sonajih deposits consistently shows chiefly alkaline conditions of formation. Fluid inclusion studies indicate that both high and low temperature hydrothermal fluids participated in the formation of all of the deposits. The mineralization age decreases from north to south and east to west and, although metal zonation is complex, the Cu–Au association post-dated the Cu–Mo mineralization reflecting that the ore fluid evolved in terms of both cooling and chemical changes due to fluid–fluid and fluid–rock interactions. In this region most deposits record a concentric zonation, with the centres preserving porphyry and skarn deposits and deposits becoming progressively epithermal toward the outer parts of the mineralizing system. According to this, the mineralization age decreases from the porphyry and skarn deposits to the epithermal deposits. The homogenization temperature and salinity both decrease from the centre to the outer zone. The pattern of homogenization temperature zonation, which is concordant with salinity zonation, suggests that fluids migrated up-dip and towards the margins of the zonation system.

Key words: Arasbaran metallogenic zone, porphyry deposits, skarn deposits, epithermal deposits, fluid inclusions, Iran.

INTRODUCTION

The east-west trending Alborz Magmatic Belt (AMB) in northern Iran is divided into western and eastern parts. The western part, referred to here as the Alborz–Azerbaijan Magmatic Belt (AAMB), is subdivided into the Arasbaran metallogenic zone (AMZ) in the north and the Tarom–Hashtjin metallogenic province (THMP) in the south (Fig. 1; Azizi and Jahangiri, 2008; Azizi and Moinevaziri, 2009). The eastern part consists of basic and felsic tufts and lavas with alkaline to shoshonitic affinities, whereas the western part consists of andesitic to dacitic lavas and many calc-alkaline to shoshonitic granitoid bodies (Radmard et al., 2019).

This region in the Tethyan realm is an important component of a regional metallogenic zone extending from the Pontide belt in Turkey to the Himalayas. Large Cu–Mo porphyry deposits, Cu-skarn occurrences (Mollai, 1993) and Cu–Mo–Au epithermal-vein deposits (Radmard et al., 2019) in this area indicate the economic value and potential of mineralization in this magmatic belt and hence requirement of more systematic studies of metallogenesis and resource exploration.

Epithermal, base metal and porphyry deposits in Iran are mainly related to magmatic belts, including the NW–SE trending Urmieh–Dokhtar and E–W-trending Alborz which are part of the Alpine–Himalayan orogenic system (Yang et al., 2009). The most important porphyry deposits in Iran include the (Fig. 1): Sungun (Calagari, 2003), Dali (Zarasvandi et al., 2015), Aliabad (Zarasvandi et al., 2005), Sarcheshmeh (Aftabi and Atapour, 2010), Meiduk (Aftabi et al., 2008) and Shadan (Karimpour et al., 2014) deposits. Some of the recognized epithermal Au deposits of Iran are located in the AMZ (Fig. 2): Masjeddaghi, Mazrae-e-Shadi, Zaglic, Safikhanlo (Alirezaei et al., 2011).

Demonstrating the origin of hydrothermal fluids responsible for the formation of epithermal deposits is one of the essential steps in the study of this type of deposits, and the concentrations of rare earth elements (REEs), La to Lu in the hydrothermal fluids, may yield useful data regarding the origin of ore-forming elements including base metals (Kato, 1999). REEs behave coherently during most geological processes due to similar chemical and physical properties (e.g., Taylor and McLennan, 1985). In spite of this similarity, REEs can be fractionated during geochemical processes. Thus, REE patterns normalized to standard material have been widely used

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as a tool to reveal various geochemical processes. However, there are only a few geochemical studies on REEs in the deposits located in the AMZ.

Fluid inclusions within hydrothermal veins have been recognized as records of mineralizing fluids and the processes by which mineral deposits were formed (Wilkinson, 2001). One of the essential aims of a fluid inclusion study is to establish a geochemical model through collecting data about the components and physico-chemistry of the palaeofluids (Williams-Jones and Heinrich, 2005). Fluid inclusion studies form an important tool for constraining the physico-chemical features of the hydrothermal fluids responsible for pervasive alteration and mineralization processes (Nash and Theodore, 1971; Nash, 1976; Batchelder, 1977; Chivas and Wilkins, 1977; Elminan, 1977; Roedder, 1984).

Studies on the mineralization in the AMZ have been limited and provide little insight into the origin of the deposits located in this zone (e.g., Mazraeh-e-Shadi, Safikhano, Sungun, Mazraeh, Anjerd, Masjed袋ghl, Zaglig). In this paper, we describe the geological setting and geochemical characteristics of the deposits in the AMZ, including REEs and fluid inclusions, which help determine the relationships between the deposits and act as guidelines for future exploration in the AAMB. Our results extend knowledge of the epithermal mineralization processes in the AAMB, and provide exploration criteria for similar epithermal ores in this area and in other parts of the AMB in northwestern Iran.

**GEOLOGICAL SETTING**

**Regional tectonic setting.** There are five major tectonic zones in northwestern Iran related to the geodynamic evolution of the Tethyan realm between the Arabian and Eurasian plates during the Early Mesozoic–Late Cenozoic. These zones include, from south to north, the Zagros Suture and Fold-Thrust Belt, the Sanandaj–Sirjan Zone (SSZ), the Urmieh–Dokhtar Magmatic Belt (UDMB), the Central Iran blocks, and the AAMB (Alavi, 1994), shown in Figure 1.

The UDMB represents a continental arc that formed as a result of the subduction of Neotethyan oceanic crust under the SSZ in the late Mesozoic (Alavi, 1994). North of the UDMB, discrete fragments of the Central Iran blocks crop out from beneath the Tertiary volcanic cover. Prior to the Triassic continental rifting episode, the Central Iran, South Armenian, and Tauride blocks formed a microcontinent that was a part of the northern edge of Gondwana (Sosson et al., 2005; Azizi and Moinevaziri, 2009; Mollai et al., 2009; Dilek et al., 2010). All three blocks include a Proterozoic crystalline basement overlain by Paleozoic–Mesozoic sedimentary sequences (Sosson.

![Fig. 1. A simplified geotectonic map of Iran (Nabavi, 1976) with location of the Arasbaran metallogenic zone (AMZ) encompassing the deposits studied](image-url)
The calc-alkaline and shoshonitic magmatic activity in the UDMB started in the Eocene and continued into the Quaternary (Daliran and Neubaer, unpublished data). The youngest volcanic activity in the UDMB is mainly calc-alkaline in nature and is associated with tectono-magmatic processes related to post-collisional, intra-continental rifting events (Richards, 2003). The south of the UDMB is dominated by basic to intermediate igneous rocks with calc-alkaline to alkaline affinities and host the Sarcheshmeh and Meiduk porphyry copper deposits (Fig. 1). No porphyry copper mineralization has yet been reported from the UDMB zone to the north.

The AMZ in the western zone is separated from the UDMB by a narrow fragment of the Central Iran microcontinent. To the north-west, the AAMB extends into the Lesser Caucasus in Armenia and into the Eastern Pontides in Turkey. The AMZ developed along the southern margin of Eurasia. Upper Jurassic–Cretaceous flysch deposits and platform carbonates deposited adjacent to this margin are overlain by Eocene volcanic rocks, and are intruded by Oligo-Miocene, shallow-crustal to hypabyssal plutons that collectively make up the AMZ (Fig. 2). A geological map of deposits in this region is shown in Figure 3.

Throughout the Paleogene and Neogene, the AMZ was bordered to the north and north-east by the Moghan basin, in which Eocene to Miocene clastic rocks accumulated, and in the south and south-west by a fluvi-lacustrine terrestrial basin, in which Miocene red beds including clastic sediments and evaporites were deposited. During the Late Miocene to Quaternary, the Lesser Caucasus and the AMZ underwent regional contraction, shortening first in a N–NW direction and subsequently in a NNE direction. The NNE-oriented crustal shortening was accompanied by NNW stretching and extension, and associated intensive alkaline magmatism in a broad zone of dextral transtension in the hinterland of the Arabia–Eurasia collision front (Mohajel and Fergusson, 2000; Sosson et al., 2005; Dilek et al., 2010).

**Regional geology.** The magmatic rocks of the AMZ are high-K calc-alkaline to alkaline (Aghazadeh et al., 2011; Aslabanha and Foden, 2012; Castro et al., 2013; Nabatian et al., 2016), and are tectonically linked to subduction of the Neo-Tethys oceanic crust beneath the Iranian plate, and subsequent collision of Arabia with Eurasia during the Alpine–Hi-
malayan orogeny that caused the formation of different orogenic belts, including the AAMB, in which occur such deposits as the Mazraeh-e-Shadi, Sungun, Safikhanlo, Sarilar, and others (Alavi, 1994).

Cenozoic rocks of the AMZ on a Na$_2$O + K$_2$O versus SiO$_2$ discrimination diagram, can be compositionally classified as subalkaline basalt, andesite/basaltic andesite, and rhyodacite/dacite, and they plot mainly in the subalkaline field, though some of them fall into the alkaline field (Fig. 4).

Samples with 50 to ~68 wt.% SiO$_2$ are calc-alkalic to alkalic (Fig. 5A), whereas samples with >68 wt.% SiO$_2$ are generally calcic (Frost et al., 2001); a limited subset of the volcanic rocks associated with the Mazraeh-e-Shadi deposit are particularly calcic. Cenozoic plutonic rocks of the AMZ are meta-to peraluminous (Fig. 5B).

On the Rb versus Y + Nb tectonic discrimination diagram of Pearce et al. (1984), most of the igneous rocks associated with the deposits fall in the volcanic arc granite field, while some of the younger (Miocene) quartz-monzonitic and monzonitic plutons straddle the boundary with, and fall into the field of, within-plate granites (Fig. 6). On an Nb versus Y tectonic discrimination diagram, most of the igneous rocks are placed near the top of the volcanic arc granitoids (VAG field), but also span the compositional fields plotting in the third VAG, syn-COLG and WPG fields (Fig. 6).

**Metallogenic belts.** Mineralization and formation of ore deposits in orogenic belts commonly accompany igneous and associated hydrothermal activities (Sillitoe, 1997). Although some of this magmatism is related to active subduction zone processes as in volcanic arcs, the distribution and genesis of many important mineral deposits and occurrences in young orogenic systems appear to be spatially and temporally related to tectonic events that are either late in the collisional history or post-collisional in origin (Laffitte, 1984; Mitchell, 1996; Sillitoe, 1997).
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**Fig. 3C – Mazraeh-e-Shadi (Radmard et al., 2017)**
Fig. 3D, E – Safikhanlo and Zaglic (Alirezaei et al., 2011); F – Sonajil (Hosseinizadeh, 2008); G – Mazraeh (Mollai, 1993)
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Fig. 4. Plots of representative rocks from the deposits on a total alkali vs. SiO$_2$ diagram (Le Maitre, 2002)

Fig. 5A – Na$_2$O+K$_2$O-CaO versus SiO$_2$ diagram of igneous rocks associated with the deposits (Frost et al., 2001); B – a Shand’s index diagram for the Arasbaran granitoids

Discrimination fields for different types of granitoids after Maniar and Piccoli (1989) and Shand (1927); symbols as in Figure 4
The origin of heat flux, melt, and fluids, that collectively play a major role in metal transport and mineralization in post-collisional settings, raises fundamental questions in evaluating the relationships between metallogeny, magmatism, and regional tectonics in such orogenic systems.

The AMZ hosts precious and base metal mineralization, including copper, molybdenum, and gold within porphyry, skarn, and epithermal deposits (Tables 1 and 2). According to the distribution of the ore deposit types, three metallogenic zones are distinguished within the AMZ (Daliran et al., 2007), as shown in Figure 7:

b1. Zone A contains Cu ± Mo ± Au porphyry and skarn deposits, and stockwork Cu–Mo–Au mineralization types.

![Fig. 6. Trace-element, tectonic setting-discrimination variation diagram showing the composition of igneous rocks associated with the deposits](image)

![Fig. 7. Distribution of metallogenic zones and types of mineralization in the AMZ exposed in northwestern Iran (Jamalia et al., 2010)](image)
<table>
<thead>
<tr>
<th>Metallogenic zone</th>
<th>Deposit</th>
<th>Location</th>
<th>Metals</th>
<th>Magmatism/mineralization age [Ma]</th>
<th>Mineralization type</th>
<th>Geology host/country rocks</th>
<th>Economic potential</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>Sungun</td>
<td>38°41'53&quot;N 46°42'21&quot;E</td>
<td>Cu–Mo</td>
<td>22.9 ±0.2 and 21.7 ±0.2 Ma</td>
<td>porphyry, skarn</td>
<td>U. Cretaceous limestone intruded by granodiorite-monzonite plutons</td>
<td>800 Mt at 0.6% Cu; open pit mining</td>
<td>Simmonds et al. (2017)</td>
</tr>
<tr>
<td></td>
<td>Nabijan</td>
<td>38°48'58&quot;N 46°48'24&quot;E</td>
<td>Cu–Au</td>
<td>U. Eocene to Oligocene</td>
<td>porphyry, skarn</td>
<td>U. Cretaceous volcano-sedimentary rocks intruded by monzonitic plutons</td>
<td>0.4 ppm Au; 0.1% Cu</td>
<td>Baniadam (2002), Shokohi (2007)</td>
</tr>
<tr>
<td></td>
<td>Anjerd</td>
<td>38°39'17&quot;N 47°03'53&quot;E</td>
<td>Cu–(Fe)</td>
<td>Miocene</td>
<td>skarn</td>
<td>U. Cretaceous volcano-sedimentary rocks intruded by the Sheivar granodiorite-monzonite pluton</td>
<td>400,000 t at 1.5% Cu</td>
<td>Mollai (1993), Mollai et al. (2009), Atalu (2006)</td>
</tr>
<tr>
<td></td>
<td>Mazraeh</td>
<td>38°39'17&quot;N 47°03'53&quot;E</td>
<td>Cu/Au</td>
<td>10 Ma</td>
<td>skarn</td>
<td>U. Cretaceous volcano-sedimentary rocks intruded by the Sheivar granodiorite-monzonite pluton</td>
<td>400,000 t at 1.5% Cu</td>
<td>Hassanpour (2010)</td>
</tr>
<tr>
<td></td>
<td>Haftches hmem</td>
<td>38°45'42&quot;N 46°39'13&quot;E</td>
<td>Cu–Mo</td>
<td>27.05 ±0.37 Ma</td>
<td>porphyry</td>
<td>U. Cretaceous limestone intruded by granodiorite-monzonite plutons</td>
<td>1–2% Mo; 2 ppm Au; open pit mining</td>
<td>Aghazadeh et al. (2015)</td>
</tr>
<tr>
<td></td>
<td>Aniq</td>
<td>38°49'08&quot;N 46°22'13&quot;E</td>
<td>Cu–Mo–Au</td>
<td>L. Miocene</td>
<td>stockwork, intrusion-related, porphyry</td>
<td>medium- to high-K, calc-alkaline granodiorite, monzonite and diorite-gabbro plutons</td>
<td>1–10 ppm Au; in prospection</td>
<td>Jamali and Mehrabi (2006)</td>
</tr>
<tr>
<td></td>
<td>Gharehchilar</td>
<td>38°50'49&quot;N 46°22'13&quot;E</td>
<td>Cu–Mo–Au</td>
<td>25.19 ±0.19 to 31.22 ±0.28 Ma</td>
<td>stockwork, intrusion-related, porphyry</td>
<td>medium- to high-K, calc-alkaline granodiorite, monzonite and diorite-gabbro plutons</td>
<td>1–10 ppm Au; in prospection</td>
<td>Simmonds and Moazzeh (2015)</td>
</tr>
<tr>
<td></td>
<td>Zaglic</td>
<td>38°26'44&quot;N 47°21'06&quot;E</td>
<td>Au</td>
<td>Oligocene</td>
<td>low-sulphidation epithermal</td>
<td>Eocene dacitic and andesitic rocks intruded by calc-alkaline plutons</td>
<td>1–10 ppm Au; in prospection</td>
<td>Jamali and Mehrabi (2015)</td>
</tr>
<tr>
<td></td>
<td>Sonajil</td>
<td>38°11'34&quot;N 47°16'49&quot;E</td>
<td>Cu±Au</td>
<td>Oligocene</td>
<td>porphyry</td>
<td>Middle to u. Eocene andesitic rocks intruded by medium- to high-K calc-alkaline subvolcanic rocks</td>
<td>0.2% Cu; in prospection</td>
<td>Jamali and Mehrabi (2015)</td>
</tr>
<tr>
<td></td>
<td>Vali lu</td>
<td>38°20'36&quot;N 46°52'33&quot;E</td>
<td>As</td>
<td>Miocene–Pliocene</td>
<td>epithermal</td>
<td>Miocene</td>
<td>Abandoned mine</td>
<td>Jamali and Mehrabi (2015)</td>
</tr>
<tr>
<td></td>
<td>Mivehrud</td>
<td>38°32'59&quot;N 46°15'31&quot;E</td>
<td>Cu–Au</td>
<td>9.12 ±0.19 Ma</td>
<td>porphyry, epithermal</td>
<td>U. Cretaceous to Palaeocene flysch deposits intruded by medium- to high-K calc-alkaline subvolcanic rocks</td>
<td>1 ppm Au, 0.1% Cu, low grade; 3 ppm Au, Sb in veins; in prospection</td>
<td>Alirezaei et al. (2016)</td>
</tr>
<tr>
<td></td>
<td>Noldareh</td>
<td>38°47'23&quot;N 46°03'10&quot;E</td>
<td>As</td>
<td>Miocene–Pliocene</td>
<td>epithermal</td>
<td>Miocene</td>
<td>Abandoned mine</td>
<td>Jamali and Mehrabi (2015)</td>
</tr>
<tr>
<td></td>
<td>Masjedda gh</td>
<td>38°52'38&quot;N 46°56'19&quot;E</td>
<td>Cu–Au</td>
<td>20.46 ±3.55 Ma</td>
<td>porphyry, high-sulphidation epithermal</td>
<td>Eocene andesitic rocks intruded by calc-alkaline subvolcanic rocks</td>
<td>0.33% Cu, 2 ppm Au</td>
<td>Agahzadeh et al. (2015)</td>
</tr>
</tbody>
</table>

Table 1

Characteristics of the ore deposits associated with AMZ
The Sungun, Hafcheshmeh, Nabijan, and Mazraeh mines are located in this zone (Fig. 7).

2. Zone B comprises several epithermal gold deposits and occurrences (Zaglic, Safikhanlo, Sarilar, and Mazraeh-e-Shadi mines) (Fig. 7). Prominent examples of this type of mineralization are hosted by Eocene volcanic rocks.

3. Zone C contains deposits occurring in the mountain range from the Sabalan volcano in the south-east to the Jofia region in the north-west. The main examples include the Mivehrud, Masjedjaghi, and Sonajil deposits.

Widespread, WNW-trending alteration zones that are parallel to the regional faults occur mainly within Eocene volcanic rocks in Zone B. Considering the alteration assemblages, vein minerals, and fluid inclusion data, Ebrahimli et al. (2009) recognized various types of epithermal system in the AMZ, as follows (Fig. 2): (1) low-sulphidation (Safikhanlo and Zaglic); (2) intermediate (e.g. Mazraeh-e-Shadi); (3) high-sulphidation (e.g. Masjedjaghi).

METHODS

In the present study, numerous field samples were taken and selected for different purposes. Polished and thin sections were studied by optical microscopy at Tabriz University. Most samples were taken from across the skarn, host rocks and quartz-sulfide breccia veins (related to epithermal deposits). Quantitative values of major and minor elements, trace elements, and REEs were determined by inductively coupled plasma-atomic emission spectrometry and inductively coupled plasma-mass spectrometry analysing methods (ICP-MS) in the Amdel Mineral Laboratory, Australia. Approximate detection limits for the elements are: Au = 0.005 ppm, Ag = 0.2 ppm, Pb, Zn and Cu = 0.5 ppm. Microthermometric studies were conducted on 100–150 µm-thick polished slabs of quartz from silicified and mineralized zones. Sub-surface samples containing quartz veinlets obtained from diamond drill holes and surface samples were selected for thermometric analyses. Microthermometric studies were carried out at the Lorestan University Geological Department—Fluid Inclusion Laboratory using a Linkam THMSG600 freezing-heating stage mounted on an Olympus microscope. The freezing-heating stage has temperature range that varies from –196–600°C. Calibrations were performed using cesium nitrate (melting point of +414°C), n-hexane (freezing point of 1–94.3°C) and synthetic fluid inclusion standards. The heating rate was 5–10°C/min at higher temperatures (>100°C), with a reproducibility of ±1°C, but was reduced to 0.1–0.5°C/min near phase transformation, with a reproducibility of ±0.1°C. Salinities of liquid-rich fluid inclusions were calculated from measured ice-melting temperatures using the equation of Bodnar (1993). Interpolation of data and creation of longitudinal sections for the veins was done using the Kriging interpolation function built in the Surfer software, version 9, Excel, Auto CAD, Fluids (Bakker, 1999), Clathrates (Bakker, 1997) and Flincore (Brown, 1989). The basic statistics, minimum, maximum, mean and standard deviation were calculated for every vein by SPSS software, version 16.

**PORPHYRY DEPOSITS**

**Sungun.** The largest known porphyry Cu–Mo deposit in north-west Iran is the Sungun porphyry copper deposit (PCD), which is also the second largest PCD in Iran (after Sarcheshmeh PCD). The Sungun deposit, located in the NW part of Iran ~100 km NE of Tabriz, the most prominent porphyry Cu–Mo in Zone A with ore reserves of ~796 Mt at 0.6% Cu, and probable reserves >1 Gt (Simmonds and Moazzen, 2015), has been mined within the enriched, sericitic shallow mineralization zone.

The Sungun mineralization, represented by a chalcopyrite-pyrite-bornite assemblage, is associated with Oligo-Miocene granodiorite-monzonite and diorite-granodiorite plutons (Hezarkhani, 2006). The parental magma of these plutons was medium- to high-K calc-alkaline in composition and was emplaced into the country rock at shallow crustal depths of 2 km and at temperatures of ~670–780°C (Hezarkhani, 2006). The existence of primary hornblende in the fresh diorite-granodiorite rocks suggests that the parental magma was hydrous. It is inferred that the exsolution of fluids from wet magma at shallow depths during the early crystallization stage of the stocks and plutons caused the precipitation of Mo and Cu minerals in the intrusions (Hezarkhani and Williams-Jones, 1998).

Four alteration types are developed in Sungun deposit: potassic, phyllic, propylitic and argillic. The earliest alteration is represented by potassic mineral assemblages (dominated by K feldspar) developed pervasively and as halos around veins in the deep and central parts of the Sungun stock (Hezarkhani and Williams-Jones, 1998). The potassic alteration displays a close spatial association with copper and molybdenum mineralization, as much as 60% of the copper and all of the molybdenum was possibly emplaced during this alteration episode (Hezarkhani and Williams-Jones, 1998). Three types of fluid inclusion are typically observed in quartz veinlets at Sungun: (1) vapour-rich, two-phase, (2) liquid-rich two-phase (Fig. 8A) and (3) multi-phase. However, occasionally mono-phase solid (minute halite crystal) inclusions are also observed in some samples. Multi-phase inclusions include those containing liquid, vapour, and solid phases. There may be only one (principally halite) or several solid phases (e.g., halite, sylvite, and an unknown transparent solid). The homogenization temperature varies between 192–584°C and salinity varies between 1–68 wt.% NaCl equivalents. The microthermometric data and calculated parameters such as salinities, homogenization temperatures and salinities vs. homogenization temperatures binary diagram for all deposits is graphically illustrated in Figure 9.

**Sonajil.** Rocks of the Sonajil region include Eocene andesite lava, porphyry microdiorite, Incheh granitoid and Ouzzadagli Plio-Quaternary volcanic rocks. Porphyry microdiorite is the main host rock of porphyry type Cu–Mo mineralization in the Sonajil. Activity of hydrothermal fluids caused formation of sulphide, sulphosalt and oxide (magnetite, hematite) minerals in the form of dissemination, veins and veinlets.

**Table 2**

<table>
<thead>
<tr>
<th>Deposit name</th>
<th>Homogenization temperature [°C]</th>
<th>Salinity (wt.% NaCl equivalent)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sungun</td>
<td>336.35</td>
<td>31.47</td>
</tr>
<tr>
<td>Sonajil</td>
<td>400.25</td>
<td>22.96</td>
</tr>
<tr>
<td>Mazraeh</td>
<td>356.11</td>
<td>23.92</td>
</tr>
<tr>
<td>Nabijan</td>
<td>195</td>
<td>5.7</td>
</tr>
<tr>
<td>Mazraeh-e-Shadi</td>
<td>227.94</td>
<td>1.94</td>
</tr>
<tr>
<td>Zaglic</td>
<td>200.9</td>
<td>4.47</td>
</tr>
<tr>
<td>Safikhanlo</td>
<td>267.5</td>
<td>1.77</td>
</tr>
<tr>
<td>Anjerd</td>
<td>356.11</td>
<td>23.92</td>
</tr>
</tbody>
</table>
These veins/veinlets contain sulfides (pyrite, chalcopyrite, molybdenite, bornite, galena, tetrahedrite, tennantite and enargite), hydroxides and oxides (magnetite and specularite) and carbonate (malachite and azurite) minerals. Activity of supergene solutions caused the formation of a predominantly goethite-enriched cap containing copper oxides and carbonates. Low pyrite abundance, outcropping of the potassic zone, rapid uplift, a basic composition of the host stock and low hypogene grade are the reasons for poor secondary chalcocite enrichment in a form of poorly developed blanket.

Various types of fluid inclusion including mono-phase vapour, two-phase liquid and vapour (Fig. 8B), and multiphase liquid-vapour-solid of primary origin are present within quartz-sulfide veinlets. $T_H$ (L-V) for halite-bearing inclusions homogenizing by the disappearance of halite and of vapour are 260–565°C and 320–520°C, respectively and salinity is 35.3–69% NaCl. $T_H$ (L-V) for two-phase inclusion homogenization temperatures are 180–565°C, and salinity is 0.7–15.17% NaCl. In a bivariate plot of $T_H$-salinity, two distinct populations of high and low salinity fluids are recognizable and most of the data points relating to the high-salinity plot above the halite saturation curve. The coexistence of vapour-rich two-phase and halite-bearing inclusions having similar $T_H$ ranges can indicate boiling in the Sonajil porphyry deposit.

**SKARN DEPOSITS**

The skarn deposits of the Ahar region can be classified petrologically into endoskarn, exoskarn and ore skarn. Each of these can be further subdivided on the basis of the predominant mineral assemblage. Skarn mineralogy can be complex (Meinert et al., 2005). The dominant skarn minerals are garnet, calcite, pyroxene, actinolite and epidote which are accompanied by quartz, feldspar, minor vesuvianite and hornblende.

Copper-iron skarn deposits occur prominently in the AMZ along the intrusive contacts of the Oligo-Miocene granodiorite-monzonite plutons with the Cretaceous limestones. The most important skarn deposits in this zone are at Mazraeh, Anjerd, Javanshaykh and Gavdel. In the Sungun district, copper-skarn deposits are found close to the porphyry copper at the contact between a hypabyssal monzodiorite intrusion and a Cretaceous limestone. The Sungun skarn was previously mined in the 1920s by the Soviet Union. Porphyry stock emplacement led to the formation of contact metamorphism and skarn-type mineralization along its border with carbonates at the north-east of the Sungun open pit (Calagari and Hosseinzadeh, 2006). The associated contact metasomatic alteration and skarn mineralization are best developed in places where the fracture density in the carbonate rocks is relatively high. Metasomatic effects are well pronounced at the contact of the porphyry stock with carbonate rocks, and diminish gradually outwards up to 55 m from the contact.

Einaudi (1982) distinguished between reaction skarns and ore skarns. The former, of limited extent, are formed along shale-limestone contacts during metamorphism. The latter, as the name implies, are the skarns that contain mineralization, and are formed as a result of infiltration of fluids derived from igneous intrusions. In the Ahar region the processes that lead to the formation of skarn deposits include three stages as follows: (1) Emplacement of plutonic magma which leads to isochemical contact metamorphism (calcite, pyroxene, plagioclase); (2) prograde metasomatic skarn formation as the pluton cools and an ore fluid develops, (epidote, tremolite/actinolite, chlorite) and; (3) retrograde alteration of earlier formed mineral assemblages, leading to the formation of hydrosilicate minerals along with ore deposition (Mollai et al., 2014). In general, skarns are zoned from endoskarns in the causative pluton, to proximal garnet and distal pyroxene, a pattern which also indicates the overall oxidation state of the skarn system (Meinert et al., 2005).

Weak skarn mineralization occurs also at the contact of the Mivehrud subvolcanic body with Upper Cretaceous marly limestone in Zone C. The mineral assemblage consists of chalcopy-

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*Fig. 8. Photomicrographs of primary fluid inclusions at: A – Sungun, B – Sonajil, C – Mazraeh, D – Nabijan, E – Mazraeh-e-Shadi, F – Zaglic*

L – liquid, V – vapour
rite-pyrite-magnetite with minor galena, sphalerite, and molybdenite, as well as andradite, grossular, epidote, and pyroxene (Mollai, 1993; Atalu, 2006).

Textural examination shows that at least three types of garnet are distinguishable. From oldest to youngest these are as follows: (1) fine-to-medium-grained, anisotropic garnets which are mainly present in the outer parts of the skarn zone. (2) Medium- to coarse-grained, subhedral to euhedral, dark brown garnets. They occur as patchy aggregates with conspicuous growth zoning, and are isotropic in their central zones, and anisotropic at their margins. (3) Veinlets of medium- to coarse-grained, isotropic and anisotropic garnets cross-cut type 2 garnets and partially replace them (Calagari and Hosseinzadeh, 2006). Optical anisotropy in garnet is due to their departure from cubic symmetry, as a result of partial replacement of \((\text{SiO}_2)\) by \((\text{OH})\) to form hydrogarnet. Rose and Burt (1979) suggested that anisotropic garnet will form due to fluctuation in fluid composition resulting from variable admixture with meteoric water. Zoned garnets from skarn deposits of the Ahar region do not show systematic compositional variation from the core to the rim of the crystal.

**Mazraeh.** Based on Mollai (1993), among skarn deposits along the northern margin of the Ahar from west to east, the Mazraeh Cu–Fe skarn deposit is the most typical. It is located 20 km north of Ahar town (Fig. 7). The origin and development of the skarn can be related to a granitic intrusion of Miocene age, which has intruded a sequence of calcareous rocks. On the basis of petrological observations, the skarn can be subdivided into exoskarn, endoskarn and ore skarn (Mollai, 1993; Mollai et al., 2009). The main mineral constituents of the skarns are garnet, magnetite, calcite, chalcopyrite, epidote, hematite and pyroxene, accompanied by quartz, pyrite, bornite, coevalite, chalcocite, plagioclase and chlorite. The bulk chemistry and spatial variation indicate that the endoskarn was the result of interaction between the Mazraeh granodiorite with crystalline limestone and metasomatic alteration by hydrothermal fluids enriched in Mg, Fe, Cu, P, Ag, Zn, Pb, Cd, Mo, Mn. These elements point towards a magmatic source, and have been contributed to the system from the magma as well as the host rock. The crystalline limestone was the source for Ca and Mg in the case of the endoskarn. The transformation of granodiorite into endoskarn was accomplished by addition of 1.4 to 15% CaO along with 7.17% of total iron into the granodiorite, accompanied by ~15.5% depletion in \((\text{SiO}_2)\). The ore-stage is dominated by liquid-rich fluid inclusions (Fig. 8C); vapour-rich inclusions are rare. The homogenization temperature varies between 435 and 232°C and salinity ranges from 1.15 to 49 wt.% NaCl equivalents (Mollai, 1993).

**Nabijan.** The Nabijan skarn deposit is located 40 km west of Kaleibar city, East Azerbaijan province in the Alborz–Azerbaijan structural zone (Fig. 2). Geological rock units outcropping in the area consist of volcano-sedimentary rocks (trachyandesite, andesite, shale and limestone) of Cretaceous age, intruded by Oligo-Miocene monzogranite, monzogranodiorite and diorite. These intrusions and related hydrothermal activities caused the Au mineralization of the Cretaceous host rocks, as well as of the monzogranites. The mineralization in the monzogranite occurred as a silicic stockwork and sheeted veins with pyrite and minor chalcopyrite, sphalerite and galena. The other type of mineralization is skarn, which is present as sparse irregular veins with

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**Fig. 9.** Histogram of homogenization temperatures, salinity and plots of homogenization temperatures vs. salinity (wt.% NaCl equivalent) for all fluid inclusions showing two populations

Each box shows the median, quartiles, and extreme values within a category: A – Sungun, B – Sonajil, C – Mazraeh, D – Nabijan, E – Mazraeh-e-Shadi, F – Zaglic, G – Safikhanloo
small amounts of Au in the carbonates. Preliminary estimation of ore reserve amounts to 320 kt with average Au grade ~1.37 g/ton. Surface and drilling data enabled determination of two important anomalies. One is in the contact zone of intrusive rocks with the Cretaceous volcanics and carbonates, representing the skarn zone. In this anomaly, the correlation coefficient index between Au and Cu, Pb, Zn, Ag, and As is low, whereas in the second one which coincides with the stockwork mineralization within the monzogranite, it is high. The homogenization temperature varies between 168–224°C and salinity varies between 3.27–8.51 wt.% NaCl equivalent (Fig. 8D). The occurrence of hydrothermal breccias, bladed calcite, adularia, and rare coexisting vapour- and liquid-dominant inclusions suggest that boiling occurred during evolution of the ore fluids. The large variations in Th and the salinity values can be explained by boiling and/or mixing.

EPITHERMAL MINERALIZATION

Widespread, WNW-trending alteration zones that are parallel to the regional faults occur mainly within the Eocene volcanic rocks (e.g., Safikhano and Mazraeh-e-Shadi) of Zone B. The mineralization is restricted to silica, silica-carbonate, and silica-barite veins, although stockwork and sheeted veinlets occur locally. Pyrite is the main sulphide mineral and is associated with minor chalcopyrite, sphalerite, and galena. Gold is found as fine particles in silica and in sulphides. Some sericitic alteration occurs at deeper structural levels, where it is associated with quartz veins and veinlets that consist of transparent and milky quartz with crustiform and comb-like textures, typical of epithermal mineralization environments. Fluid inclusion studies show a wide range of homogenization temperatures (120–350°C) and salinity (1–15 wt.% NaCl equivalent). A positive correlation exists between the higher salinities and the high base metal contents in the veins (Ebrahimi et al., 2009).

Zaglik and Safikhano. The Zaglik and Safikhano epithermal gold prospects are located to the west of the Cenozoic AAMB in NW Iran. These prospects are characterized by extensive outcrops of Cretaceous flysch type strata and Cenozoic volcanic and plutonic rocks. They are only 4 km apart, and lie in an area covered by Eocene–Miocene volcanic, pyroclastic and intrusive rocks. The oldest rocks include dark grey to green, porphyritic andesite, basaltic andesite, tachy-andesite to latite-andesite of Upper Eocene age. The rocks consist mainly of plagioclase (andesine–oligoclase), hornblende, clinopyroxene and subordinate biotite and quartz. The dominantly andesite lavas are overlain by light to dark green, fine-grained tuff of intermediate composition, with scattered crystals of plagioclase, hornblende, pyroxene and biotite, as well as andesitic tuff breccias and andesitic breccias. The volcanic–pyroclastic succession in Safikhano was intruded by a hypabyssal intrusion composed dominantly of syenite associated with subordinate granite and monzogranite. The intrusive rocks consist mainly of plagioclase, alkali feldspars, quartz, hornblende, and biotite. Apatite and Fe–Ti oxides are common accessory minerals. An Oligocene age is proposed for the intrusion by analogy with similar intrusions in the Ahar quadrangle. Numerous acidic to intermediate dykes intruded the Upper Eocene volcanic and pyroclastic rocks and the Oligocene intrusive rocks. The dykes vary in length and width between 50–500 and 1–20 m, respectively.

Mineralization is mainly restricted to quartz and quartz-carbonate veins and veinlets. Pyrite is the main sulphide, associated with subordinate chalcopyrite and bornite. Gold occurs as microscopic and submicroscopic grains in quartz and pyrite.

Host volcanic rocks from both Safikhano and Zaglik areas are medium- to high-K calc-alkaline and alkaline rocks and display fractionated REE patterns, with LREE significantly enriched relative to the HREE. On primitive mantle-normalized plots, they display depletions in Nb, Ti and P, and enrichments in Nb, which are common characteristics of arc-related magmas worldwide. Hydrothermal alteration minerals developed in the wall rocks include quartz, calcite, pyrite, kaolinite, montmorillonite, illite, chlorite, and epidote. Minor alunite occurs at Safikhano. Gold is locally enriched in the altered rocks immediately adjacent to the veins. The ore-stage quartz from both prospects is dominated by liquid-rich fluid inclusions (Fig. 8E); vapour-rich inclusions are rare. The homogenization temperature varies between 176–224 and 199–317°C and salinity varies between 1.4–9.6 and from <1 to 6.7 wt.% NaCl equivalent, for Safikhano and Zaglik, respectively. The occurrence of hydrothermal breccias, bladed calcite, adularia, and vapour- and liquid-dominant inclusions suggests that boiling resulted from the evolution of ore fluids. The large variations in Th and the salinity values can be explained by boiling and/or mixing. A lack of sulphate minerals in the veins suggests that sulphides and gold precipitated from a reduced, H_2S-dominant fluid. The δ^34S values for the ore fluid vary between ~4.6 and ~9.3‰. Sulphur could have been derived directly from magmatic sources, or leached from the volcanic and plutonic country rocks (Alirezaei et al., 2011). Ore formation in the Zaglik and Safikhano areas occurred in response to mixing, boiling, and interactions with wall rocks. Considering the intermediate-argillic alteration, low contents of base metal sulphides, and the overall low salinities, the Zaglik and Safikhano occurrences can be classified as low-sulphidation epithermal systems.

Mazraeh-e-Shadi. The Mazraeh-e-Shadi epithermal deposit is located ~130 km to the north-east of Tabriz in the magmatic rocks of the central Iranian geostructural zone (Fig. 2). More than 10 Cu–Mo–Au porphyry deposits and epithermal gold systems have been identified in the Arasbaran metallogenic zone, e.g. Sungun (Calagari, 1997), Mazraeh, Sungun and Anjerd (Mollai, 1993; Calagari and Hosseinzadeh, 2006; Karimzadeh Somarin and Moayyed, 2002), Masjedaghi (Akbarpur, 2007), Zaglik (Heidarzadeh, 2006), Safikhano (Safikhano, 2002), Sharafabad (Purmik, 2006; Ebrahimi et al., 2009).

In the Mazraeh-e-Shadi area three types of lithologies are represented (Radmand et al., 2017), as follows: Eocene pyroclastic rocks, Oligocene–Pliocene volcanic rocks, and Quaternary volcanicogenic conglomerate (agglomerate) and aluvial units. The Eocene pyroclastic, volcanic and agglomeratic rocks (usually of an andesitic composition) are overlain by dacitic tuff, dacite, hornblende andesite, trachyandesite and andesite volcanic rocks of Oligocene and Pliocene age. Numerous dacite to andesitic dykes intruded into the volcanic and pyroclastic rocks are of Upper Eocene age. The youngest rock units in the area are Quaternary alluvial plain and river deposits. The Quaternary volcanicogenic conglomerate (agglomerate) crops out in the vicinity of the Mazraeh-e-Shadi deposit in the southwestern part of the study area. Subvolcanic rocks such as quartz monzodiorite-diorite porphyry intruded the Eocene volcanic and volcano-sediementary units and caused mineralization and alteration. Gold mineralization in the Mazraeh-e-Shadi is related to Eocene volcanic and intrusive rocks. Outcrops of subvolcanic
rocks extend over a distance of ~1,650 m and their width ranges from 300 to 450 m in the area.

Three main types of fluid inclusions were identified in the samples collected from silicified and mineralized zones; Type 1 – liquid-rich fluid inclusions (LV) consists of liquid + vapour with the liquid phase volumetrically dominant, and lack daughter crystals (Fig. 8F). The diameters of these fluid inclusions ranges from 4 to 92 µm. These fluid inclusions are common in all the mineralized quartz veins. They occur in almost all samples and made up the largest numbers of the inclusions studied. The first ice-melting temperatures ($T_{fm}$) of these inclusions range from –19 to –29°C and the last ice-melting temperature ($T_{m\text{ ice}}$) is between –3.2 and 2.2°C. These inclusions yield homogenization temperatures from 160 to 308°C and salinities are 0.17–5.2 wt.% NaCl equivalent. Type 2 – two-phase vapour-rich (VL) fluid inclusions are common in exposure quartz vein samples. The diameters of these fluid inclusions ranges from 6 to 17 µm. These inclusions homogenized to a gas phase. The homogenization temperatures, ranging from between 173 and 324°C, are the highest homogenization temperatures obtained from quartz in the study area and salinities are 0.16–1.8 wt.% NaCl equivalent. The gas bubbles in the (VL) inclusions occasionally occupy >80% of the inclusion volume. Type 3 – vapour monophase (V) inclusions have sizes from 10 to 31 µm (Radmard et al., 2017). Quartz veins containing liquid-rich and vapour-rich inclusions indicate boiling during hydrothermal evolution, resulting in relatively higher gold grades (up to 813 ppb Au).

GEOCHEMISTRY OF RARE EARTH ELEMENTS

The study of REE behaviour in rocks and minerals is of note for the following reasons. First, REEs induced special behaviours during various geochemical processes, which are used as detectors in recognition of ores and rock origin as well as of geological and tectonic settings. Secondly, environmental issues are highly significant to radioactive wastewater management and REEs are known as fissionable plutonium and uranium products in nuclear power plants (Brooking, 1984; Rard, 1988). Thirdly, economically speaking, they have several applications of extraordinary value. REEs are conveniently divided into three subgroups: HREE, MREE and LREE (Rollinson, 1993). The fractionation degree between REEs can be determined by the LREE/HREE ratio. It has been pointed out that replacement of REEs by calcium is one of their significant features (Knarchenko and Pokrovsky, 1995; Pollard, 1995).

Geochemical considerations indicate that the concentration values of rare earth elements in the deposits range from 55 to 234 ppm (Appendix 1*).

To evaluate the fractionation degree between REEs, various ratios of REEs, such as (La/Yb)n, (La/Sm)n and (Gd/Yb)n were used. (La/Yb)n determines the degree of fractionation of LREEs from HREEs during geochemical processes (Aubert et al., 2001; Yusoff et al., 2013), while the two other ratios establish the degree of fractionation between LREEs and MREEs and between MREEs and HREEs, respectively (Yusoff et al., 2013).

The values of these ratios range for (La/Yb)n from 8.88 to 88.77, for (La/Sm)n from 3.36 to 28.41 and for (Gd/Yb)n from 1.57 to 5.17 (Appendix 2). The greatest fractionation occurred

* Supplementary data associated with this article can be found, in the online version, at doi: 10.7306/gq.1520
between LREEs and HREEs (up to 89 times) with a relatively minor fractionation between MREEs and HREE. These ratios reflect fractionations as follows: LREE/HREE > LREE/MREE > MREE/HREE. The highest values of the three ratios have been observed in the Mazraeh-e-Shadi and Nabijan deposits (Appendix 1): (La/Yb)n = 89, (La/Sm)n = 28.41 and (Gd/Yb)n = 5.17 that represent large fractionation of REE in the Mazraeh-e-Shadi deposit not only because of high LREE but also on account of low MREE and HREE. Since the distribution pattern of REEs is governed by mainly geochemical conditions, the La/Y ratios render useful data for determination of pH in the environment of ore formation. Values of La/Y >1 show alkaline conditions, while values of La/Y <1 indicate acidic conditions (Crinci and Jurkowic, 1990). This ratio in this region ranges from 1.32 to 29.17 (Appendix 2 and Fig. 13).

Calculations of Eu and Ce anomalies are defined by the following equations (Taylor and McLennan, 1985): Ce/Ce* = (2(2Ce/Ce chondrite)/(La/La chondrite + Pr/Pr chondrite)) and Eu/Eu* = (2(Eu/Eu chondrite)/(Sm/Sm chondrite + Gd/Gd chondrite)). The Eu and Ce anomaly values across the lithological units range from 0.60–1.29 and 0.61–0.98, respectively (Appendix 1 and Fig. 14). Values >1 and <1 are called positive and negative anomalies, respectively. Plotting the Ce/Ce* values of all samples on a Ce/Ce* vs. ΣREEs binary diagram represents one population including <1. This situation is also repeated for Eu/Eu* and (Pr/Yb)n (Fig. 15).
**DISCUSSION**

**REEs.** Rare earth element variations across the all deposits are similar indicating a similar source for the REE. All the deposits along the profile from Nabijan to the Sonajil show chiefly alkaline character. As the shear zones formed under relatively similar PT conditions, it is likely that the stability of the REE-bearing minerals was controlled by the characteristics of the fluids, i.e., pH, oxygen fugacity ($f_{O_2}$) and concentrations of potential REE ligands ($SO_4^{2-}$, $CO_3^{2-}$, $OH^-$, $F^-$, $PO_4^{3-}$). These REE ligands may precipitate into REE minerals or REE-bearing minerals in response to changes in fluid composition, because of fractional crystallization, fluid-rock reaction, or fluid-mixing processes (Rolland et al., 2003). Precipitation and dissolution of various REE minerals at different stages of shear zone formation illustrate that changes in fluid chemistry occurred in individual shear zones (Rolland et al., 2003). The Nabijan deposit has the highest La/Y values (29.17), indicating basic conditions of hydrothermal alteration during mineralization (Fig. 13). The basic conditions of hydrothermal alteration decrease from epithermal deposits to the porphyry and skarn deposits, from the north to the south and west to east due to the relationships between these deposits (Fig. 13). This relationship indicates that porphyry and skarn deposits have an early age and epithermal deposits are later in origin.

All the deposits, based on the REE values and Ce anomalies, can be grouped into one category with an alkaline pH, negative Ce anomalies and high REE values. This is in fact an enrichment zone. The positive anomalies of Ce indicate that Ce was fixed as $Ce^{4+}$ under acidic and oxidising conditions. The only logical and acceptable factor for the negative anomalies for Ce in this region is the coupled complexation of $Ce^{4+}$ with carbonate ligands and its leaching during the mineralization processes (Karadag et al., 2009).

Since the Eu carriers are plagioclases, Eu concentrations might be related to the degree of plagioclase alteration. The Eu positive anomaly strongly indicates early stages of rock alteration in acidic environments (Kikawada, 2001). The positive anomalies of Eu and ($La/Lu$) imply acidic fluids at high temperature (Wood, 1990). Considering the three parameters Ce/Ce*,

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**Fig. 14. Variations of Ce/Ce* and Eu/Eu* values**

Abbreviations are given in Figure 10

**Fig. 15A – relationship between Ce/Ce* vs. $\Sigma$REE; B – $(Pr/Yb)_{cn}$ vs. $\Sigma$REE; C – Eu/Eu* vs. $\Sigma$REE in the Sungun, Zaglic, Safikhano, Nabijan, Mazraeh, Mazraeh-e-Shadi and Sonajil deposits**
Eu/Eu* and (Pr/Yb)n in all the deposits indicates alkaline and redox conditions of hydrothermal fluids enriched in REE with negative anomalies of Eu and Ce (Fig. 15). Geochemically, these deposits show similar major and trace element characteristics and overlapping inter-element ratios, indicating common melt sources; their enrichment in light lithophile element and high incompatible element abundances suggests derivation of their magmas from a subduction-metasomatized, subcontinental lithospheric mantle source.

Fluid inclusions and the mineralization age. Plots of salinity versus homogenization temperatures for fluid inclusions (Fig. 9) indicate that two types of hydrothermal fluids participated in the formation of all deposits: (a) high temperature and (b) low temperature fluids. At Zaglic, Safikhanlo and Nabijan the two types of hydrothermal fluids are unclear due to insufficient microthermometric data.

Projection of microthermometry data of fluid inclusions on the Th versus salinity diagram (Fig. 16A) indicates characteristics of a typical epithermal, skarn and porphyry deposit. Figure 15B shows that mineralization has taken place in a temperature range of 150 to 585°C. The depth of mineralization (Fig. 17) decreases from porphyry and skarn deposits (Sungun, Sonajil and Mazraeh) to the epithermal deposits (Mazraeh-e-Shadi, Zaglic and Safikhanlo).

In this region most deposits show concentric zonation, with the centres preserving porphyry and skarn deposits (i.e. Sungun, Sonajil and Mazraeh), and outer deposit types becoming progressively epithermal (i.e. Mazraeh-e-Shadi, Zaglic and Safikhanlo). As follows from Table 1, mineralization is characterized by a pronounced regional zonation in the age and distribution of metals. The mineralization age decreases from the north to the south and east to west, and from the porphyry and skarn deposits to the epithermal deposits (Fig. 18). Although metal zonation is complex, the metal content changes from a Cu–Mo to Cu–Au association with time, reflecting an ore fluid that evolved in terms of both cooling and chemical changes due to fluid–fluid and fluid–rock interactions.

Fig. 16. Plots of salinity versus homogenization temperatures of primary fluid inclusions
Geochemistry of skarn and porphyry deposits in relation to epithermal mineralization in the Arasbaran metallogenic zone...

Fig. 17. Plots of homogenization temperatures vs. depth for all fluid inclusions in all deposits (Hass, 1971)

Fig. 18. Variation of the magmatism/mineralization age in the AMZ and its extension northwards to the Lesser Caucasus

The mineralization age decreases from the north-east to the south-west
Fig. 19. The homogenization temperature (°C) variation

Fig. 20. The salinity (wt.% NaCl equivalent) variation
Geochemistry of skarn and porphyry deposits in relation to epithermal mineralization in the Arasbaran metallogenic zone...

Fig. 21. The homogenization temperature (°C) variation in porphyry and skarn deposits (from Sungun to Sonajil)

Fig. 22. The salinity (wt.% NaCl equivalent) variation in porphyry and skarn deposits (from Sungun to Sonajil)
he homogenization temperature (°C) zonation and salinity (wt.% NaCl equivalent) zoning (Figs. 19 and 20) are recognized from the centres (Sungun, Sonajil and Mazraeh) towards the margins of the mineralizing system (Mazraeh-e-Shadi, Zaglic, Safikhano and Valili). These zonation patterns (Figs. 18–20) indicate that three mineralization centres represented by porphyry and skarn deposits (Sungun, Sonajil and Mazraeh) were formed in this region, surrounded by epithermal deposits. There is a distinct zonation from Sungun to Sonajil, with marked decrease in salinity (wt.% NaCl equivalent) and increase in homogenization temperature (°C; Figs. 21 and 22). Summarizing, the homogenization temperature and salinity decreases from the centre to the outer zone, suggesting that fluids migrated up-dip in the central parts of the mineralizing system. Thus, high- and low-sulfidation epithermal deposits reflect end-members in a continuum of magmatic-hydrothermal processes that progressively incorporated more non-magmatic waters as the volcanic system wand, or as one moves away from the volcanic centre.

CONCLUSIONS

Cenozoic rocks of the Arasbaran metallogenic zone compositionally can be classified as subalkaline basalt, andesite/basaltic andesite, and rhyodacite/dacite and they plot mainly in the subalkaline field. Most of the igneous rocks associated with the deposits fall in the volcanic arc granites field, while some of the younger (Miocene) quartz-monzonitic and monzonitic plutons straddle the boundary with, and fall into the sub-alkaline field. Most of the igneous rocks as so site/basaltic andesite, and rhyodacite/dacite and they plot compositionally can be classified as sub-alkaline basalt, andesite and rhyodacite/dacite and they plot compositionally can be classified as sub-alkaline basalt, andesite and rhyodacite/dacite.

In this region most deposits record a concentric zonation on a regional scale, suggesting that the heat was generated in the centre of the mineralizing system, where the porphyry and skarn deposits were formed (i.e. in the Sungun and Sonajil areas). The relationship between the deposits indicates that the epithermal deposits post-dated the porphyry and skarn mineralization. This is supported by the homogenization temperatures and by salinities recorded in fluid inclusion data, which demonstrate that the fluids migrated up-dip and were spreading towards the margins of the zonation pattern. This is shown by the zonation from Sungun to Sonajil, with marked decreases in salinity and increases in homogenization temperature.

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