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Deciphering the magma storage conditions and preeruptive processes at Cumaçay: a Plio-Quaternary volcanic eruption center in Eastern Anatolia, Turkey

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Abstract: We investigate the geology and petrology of Cumaçay, one of the Plio-Quaternary eruption centers of postcollisional volcanism in Eastern Turkey, using a combination of geochronology, bulk-rock geochemistry, thermobarometry, and thermodynamic simulations. Our new K-Ar ages reveal an age of 3.5–0.97 Ma for the eruptive products, which spread around an area of approximately 1000 km². Mineral-melt equilibria for olivine, orthopyroxene, clinopyroxene, and plagioclase allow estimation of crystallization pressures, and temperatures indicate two separate crustal storage of crystallization at 14–28 km and 5–15 km. The temperature estimation from diverse methods ranges from 954 °C to 1224 °C. Thermodynamic modelings using Magma Chamber Simulator (MCS) at a representative pressure (5 kbar) imply that the assimilation and crustal contamination (AFC) resulted in evolved melts with various SiO₂ contents at mid to lower crustal depths. Calculated wide range of temperatures, compositional variety from evolved members, and MCS recharge simulation at a representative pressure (2.5 kbar) reveal that the magma mixing between contrasted melts characterizes the final chemical dispersal of the erupted lavas.

Key words: Eastern Anatolia, magma plumbing system, thermobarometry, mineral chemistry

1. Introduction
Reconstructing the magma storage conditions at deep and shallow crustal depths and their petrological evolution is one of the aims of igneous petrology. It is critical to understanding and monitoring future eruption scenarios (e.g., Mclnt, 2005; White and McCausland, 2016; Cashman et al., 2017). Besides geophysical investigations of fossil and active magma storage regions, petrologic studies of erupted products provide valuable information about magma storage and ascent dynamics. For example, minerals can monitor the whole range of crystallization temperature, pressure, and composition of magma in which they developed during storage and ascent (e.g., Eskandari et al., 2018). This approach has been applied to various intracontinental volcanic regions. Eastern Anatolia is represented by basaltic to rhyolitic lava flows, pyroclastic deposits of postcollisional magmatism extending from the Middle Miocene to historical times (e.g., Keskin et al., 1998; Keskin, 2003; Özdemir et al., 2016, 2019, 2022; Açlan and Altun, 2018; Oyan, 2018). The voluminous postcollisional volcanic activity that erupted from various eruption centers has covered the region and formed extensive basaltic plateaus and active-dormant volcanoes (Pearce et al., 1990; Yilmaz et al., 1998; Keskin, 2003, 2007; Karaoğlu et al., 2005; Özdemir et al., 2006, 2019, 2022; Lebedev et al., 2010, 2016; Özdemir and Güleç, 2014; Oyan et al., 2016, 2017; Açlan et al., 2020). The volume and rock type of the volcanics show variations in time. The Middle Miocene and Quaternary are dominated by intermediate to acidic eruptions; however, the Late Miocene, Pliocene, and Early Quaternary are rich in basaltic products (Özdemir et al., 2022). With the exceptions of limited studies on the magma storage conditions of individual volcanic centers (e.g., Özdemir et al., 2011, 2022; Macdonald et al., 2015; Karaoğlu et al., 2017) previous works on the eastern Anatolian volcanics are mainly concerned with the mantle source region, time-space distribution, and connection with the previous subduction and collision events. Our study focuses on Cumaçay, one of eastern Anatolia’s poorly known eruption centers. We report detailed investigations of various ranges of lava types with new K-Ar ages. A combination of whole-rock, mineral chemistry, and thermodynamic simulations is used to reveal magma storage conditions and chemical variations of the Cumaçay.
2. Analytical methods
The whole rock geochemistry of the Cumaçay volcanics was determined by ICP-ES and ICP-MS following lithium metaborate/tetraborate fusion and dilute nitric acid digestion at Acme Analytical Laboratories in Canada. Mineral major element compositions were obtained using a CAMECA SX-100 five-wavelength spectrometer (WDS) instrument at the University of Bristol. Minerals were analyzed using a 10 nA beam current, 5 µm beam diameter, and a 20 kV accelerating voltage. Sr and Nd isotopic compositions were conducted at the Pacific Centre for Isotopic and Geochemical Research (PCIGR) at the University of British Columbia. The results of Sr and Nd isotopes were determined on a Thermo-Finnigan Triton TIMS and a Nu Plasma MC-ICP-MS, respectively, using the procedure of Weis et al. (2006). K-Ar age determinations were carried out at the Isotope laboratory of IGEM-RAS, Moscow. The radiogenic $^{40}$Ar analysis was obtained by the isotope dilution method with mono-isotope $^{38}$Ar as a spike on the mass spectrometer MI-1201 IG (SELMI). The precision of the results was controlled by measurements of isotopic compositions of atmospheric argon. FBA-01 spectrometer was used to acquire potassium content in the samples.

3. Results
3.1. Geology and age of the Cumaçay volcanics
Cumaçay volcanics are one of the constituents of eastern Anatolian postcollisional volcanism, exposed 50 km northwest of Ağrı city, covering an area of more than 1000 km$^2$ (Figure 1). Active right-lateral strike-slip Balık Göl fault cuts across the products of volcanism from the northwest toward the southeast (Sağlam-Selçuk et al., 2021). They cover the Lower Paleozoic metamorphic rocks (Topuz et al., 2017, 2021), allochthonous Late Cretaceous Neotethyan ophiolites/mélanges (Kağızman Complex), Paleocene to Miocene sedimentary rocks and Upper Miocene pyroclastics and intermediate volcanics (Keskin, 1992). The Cumaçay volcanics were first named by Kral and Çağlayan (1980) and are mainly composed of basalt to rhyolitic rocks and intermediate to acidic pyroclastic rocks. They are exposed in extensive areas; however, the best outcrops are primarily located around the Cumaçay village (Ergen and Sümengen, 2018). The current study mainly focused on the basaltic, intermediate, and acidic lava members exposed around the Cumaçay village. Our new K-Ar isotopic ages indicate that the Cumaçay volcanics erupted between Upper Pliocene to Pleistocene (Supplementary File 1). The volcanism started with basaltic lava flow, evolved to intermediate and acidic members, and ended with eruption of younger basaltic lava flows. The observed initial product of Cumaçay is a porphyritic basaltic lava flow, exposed northern parts of the study area, flows over the basement strata, and mostly covered by younger basaltic and intermediate members of the volcanism. A single K-Ar age yields 3.5 ± 0.18 Ma. Following the basaltic lava flow, various lava compositions including basaltic trachyandesites, basaltic andesites,
trachyandesites, andesites, and dacites have erupted. The basaltic trachyandesites and basaltic andesites display porphyritic and glassy textures. K-Ar ages of those lavas are 2.4 ± 0.1 and 2.4 ± 0.12 Ma. Andesites (including trachyandesites) are the most widespread lava flows within the Cumaçay volcanics. Obtained K-Ar ages from the three samples are between 2.9 ± 0.2 and 2.16 ± 0.16 Ma. They are generally porphyritic, occupy high topographies, and mostly overlie the older basaltic and basaltic trachyandesitic lavas. Dacitic lavas have an aphyric texture and cover the andesitic lavas in the southern and northern parts of the study area. The obtained single K-Ar age is 2.1 ± 0.15 Ma. Cumaçay volcanism has ended up with the eruption of two basaltic lava flows. One of them is exposed mainly to northern parts as thick layers, forms columnar jointing, and covers the intermediate and acidic members of the Cumaçay. We obtained a single K-Ar isotopic age of 0.97 ± 0.1 Ma (Supplementary File 1).

3.2. Whole rock and isotope geochemistry

Seventy-two rock samples from Cumaçay volcanics were analyzed for major and trace element compositions. The majority of the samples have low LOI values (<1 wt%), suggesting that the rocks are fresh (Supplementary File 1). Major element compositions (on a hydrous basis) range between 49.25 and 64.62 wt% for SiO₂ and 1.43–7.43 wt% for MgO. Cumaçay volcanics show a compositional spectrum from basalts to dacites (Figure 2). Volcanism started with the mildly alkaline basaltic lava flows, continued with subalkaline intermediate to acidic members, and ended up with transitional and subalkaline basaltic flows. The transitional and subalkaline members are calc-alkaline (Figure 2), and the high Na₂O/K₂O indicates their sodic nature. On conventional Harker diagrams (Figure 3), CaO/Al₂O₃, Fe₂O₃, TiO₂, MgO, Sc, Ni, and Co display negative, but K₂O and total alkalis define negative correlations with SiO₂. The good correlations of selected major and trace element contents would appear to reflect the continuous effect of the chemical differentiation at Cumaçay volcanism. MORB-normalized trace element and chondrite-normalized REE distributions of selected Cumaçay volcanics are given in Figure 4. Large ion lithophile elements (LILE) are enriched in all samples compared to the high field strength elements (HFSE). The element enrichment level seems to increase from basalts to dacites. Basaltic andesites, basaltic tracyandesites, andesites, and dacites have negative Pb, P, and Ti contents. All rock groups display depletion of Nb and Ta, indicating inherited subduction components in mantle source or crustal contamination from country rocks en route to the surface. The Cumaçay volcanics show variable enrichment of light rare earth elements (LREE) over heavy rare earth elements (HREE) with weak negative Eu anomalies increasing andesites to dacites. The extent of enrichment rises slightly with increasing whole-rock SiO₂ contents (e.g., La/Yb₁/₂; 7.3–7.6 for basalts and 7.6–9 for dacites). The medium rare earth element (MREE) to HREE remains mostly flat over this interval displaying slight fractionation in andesites and dacites (e.g., Sm/Yb₁/₂; 1.9–3.2 for andesites and 2.3–3.4 for dacites). ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd isotopic ratios of the volcanics vary in the range of 0.703616–0.704762 and 0.512878–0.512757 (Supplementary File 1). The basaltic samples display the lowest Sr and highest Nd isotopic compositions; in contrast, the more evolved lavas have more radiogenic values.

3.3. Mineralogy and petrography

The majority of the Cumaçay volcanics are rich in crystals and porphyritic with hyalopilitic groundmass;
Figure 3. Selected major and trace element variation diagrams of Cumaçay volcanics.
however, basalts are entirely crystalline. All basaltic lavas contain varying amounts of phenocrysts (>0.5 mm) and microphenocrysts (0.1–0.5 mm) composed of olivine, plagioclase, clinopyroxene, and Fe-Ti oxides. Olivines are in the form of phenocrysts and microphenocrysts with a maximum length of 1.5 mm. They have iddingsite along their rims and display skeletal texture in subalkaline members (Figure 5a). Clinopyroxene is ubiquitous microphenocrysts in all basaltic lava flows. Clinopyroxene and plagioclase display subophitic and glomeroporphyritic textures. Plagioclase phenocrysts are the most common mineral in all basalts, reach up to 1.5 mm in size, and are unzoned or slightly oscillatory zoned. They include melt, pyroxene, and olivine inclusions. Basaltic andesites and basaltic trachyandesites are weakly and highly porphyritic within a fine-grained groundmass. The pyroxenes are common as microphenocrysts. The size of both olivine and plagioclase phenocrysts reaches up to 1.5 mm. Sieve-textured plagioclase crystals have abundant melt and mineral inclusions (e.g., pyroxenes, Figure 5b). The groundmass is crystalline and contains pyroxene, plagioclase, olivine, Fe-Ti oxide microcrystals, and glass. Andesites are texturally highly variable; some are aphyric with feldspar microlites and glassy groundmass, whereas porphyritic ones contain plagioclase, clinopyroxene, orthopyroxene, and Fe-Ti oxides. The size of the plagioclases ranges up to 2 mm; some of them have corroded rims. Orthopyroxenes are more abundant than the clinopyroxenes compared to mafic members of the Cumaçay. Dacites are the most evolved members of the Cumaçay volcanism. Some of them are highly vesicular and aphyric with glassy groundmass. The porphyritic examples contain plagioclase, clinopyroxene and orthopyroxene phenocrysts (Figure 5c). The orthopyroxene is the most abundant mafic phase. The amount of clinopyroxene is reduced compared to the intermediate and mafic members. The groundmass contains plagioclase, pyroxene Fe-Ti oxides, and glass. Andesites and dacites are rich in microcrystalline enclaves and crystal clots made of pyroxene, plagioclase, and Fe-Ti oxides (Figure 5d).

3.4. Mineral chemistry
To show the evolution of mineral chemistry with changing the host rock SiO$_2$ contents, we present analyses as a series of stacked histograms ranked downwards in order of increasing SiO$_2$ (Figure 6). Mineral chemistry data including one sample each from basalt (Cum175), basaltic trachyandesite (Cum158), andesite (Cum160), and dacite (Cum241), and four samples from basaltic andesites (Cum164;179;181;199) are given in Supplementary File 2.

Olivine
Olivine crystals within different rock groups are normally zoned with forsterite (Fo) rich cores. Fo contents of olivine phenocrysts and microcrystals in basalts range between 81 and 84 (Figure 6a). The Fo of four different basaltic andesites ranges between 68 and 86 mol%. Phenocrysts Fo contents of basaltic andesites decrease with increasing the host rock SiO$_2$. Olivine microcrystals have the lowest Fo compared with the phenocrysts within the same rock. Some of the olivines with high MgO wt% overlap with the olivine compositions in basalt.

Orthopyroxene
Orthopyroxene is present in basaltic andesite to dacites with increasing abundance. There are scarce pigeonite crystals in these rocks. Orthopyroxenes have a composition range of En$_{57-79}$ and can be classified as enstatite with a 2–4 mol% wollastonite component. As for clinopyroxenes, orthopyroxenes in the basaltic andesites and dacites display relatively limited ranges in Mg# (78–82 and 56–71, respectively), whereas andesite ranges widely between these two extremes (66–81) (Figure 6b). Normally and reversely zoned crystals coexist in andesitic lava.

Clinopyroxene
Clinopyroxenes are Mg-rich augites (En$_{32-53}$, Fs$_{9-23}$, Wo$_{32-45}$) and present in basalts to dacites with decreasing
abundance. They occur as microcrystals in basalts, basaltic trachyandesites, and basaltic andesites; however, they form phenocrysts in andesite and dacites. Unzoned, normally, and reversely zoned crystals coexist in the same sample. The Mg# (molar Mg/(Mg+Fe) of clinopyroxenes shows a decrease from mafic to acidic members (Figure 6c). The Mg# of clinopyroxenes in basalt, basaltic andesites, and basaltic trachyandesite is in the range of 69–76, 66–85, and 63–76, respectively. However, Mg# of clinopyroxenes in andesite has a wide compositional range (63–81; Figure 6c). A single clinopyroxene in a dacitic sample yields an Mg# of 63 (not shown).

Plagioclase

Plagioclase is the most abundant phenocryst and occurs in almost every rock sample either as phenocrysts or in the groundmass. The composition of plagioclases ranges from bytownite to andesine. Overall, there is a general decrease in anorthite (An) content with increasing host rock SiO₂. Some of the plagioclase phenocrysts in basalt typically have slightly more An-rich rims than cores. Composition of the plagioclases in basalt is in the range of An₆₉-₆₀ (Figure 6d). Plagioclase in basaltic andesites records the highest anorthite contents (An₄₈-₇₉). The lower An contents are typically from the rim or microcrystal compositions. Plagioclases in basaltic trachyandesite have a lower content of An (An₃₂-₄₃) compared to basalts and basaltic andesites (Figure 6d). Normal zoning is dominant with a rimward decrease of only 1–2 mol% An. The An contents of phenocrysts display a wide range in andesite (An₃₁-₇₀) and dacite (An₃₇-₆₄), which is more prominent in andesite.

Fe-Ti oxides

Ilmenite and titanomagnetite are observed in most of the Cumaçay volcanics. They occur as groundmass crystals, inclusions in other phenocrysts, and rarely as phenocrysts. The ulvospinel component in the cores of titanomagnetites ranges from 24 to 66 mol%. Ilmenite contents in ilmenite crystals range from 59 to 96 mol%.

3.5. Conditions of magma storage and eruption

In this section, we report and discuss our results using the various multiple minerals and mineral/liquid methods available in the literature. Intensive magmatic parameters, such as temperature (T), pressure (P), oxygen fugacity (fO₂), and melt water content (H₂O) at preeruptive conditions, could be presented through these models. The results of the different approaches are given in Table 1.
3.5.1. Initial estimates by independent variables
One of the main difficulties of employing the geothermometers is the requirement of some parameters, such as P, H₂O, and T. It is, therefore, crucial that independent methods accurately determine one or more variables (e.g., Polo et al., 2017). For this reason, we first applied Fe-Ti oxide and clinopyroxene-liquid to calculate the temperature and pressures.

Coexisting magnetite-ilmenite pairs are rare in the Cumaçay volcanics, most of which are from groundmass microcrystals. Due to the rarity of the touching oxide pairs, we also used individual crystals within the same thin section for temperature, and oxygen fugacity (fO₂) estimates. Each magnetite was checked against each ilmenite analysis for the same sample to test the Mg/Mn equilibrium (Bacon and Hirschmann, 1988). Only data that passed this test were used for the calculations (Table 1). Suitable pairs were then processed using the method of Anderson and Lindsley (1985), with the recalculation scheme of Stormer (1983) for fO₂ and T estimates (Figure 7a). In basalt, Fe-Ti oxide temperatures range between 1009 and 1090 °C. Temperatures from basaltic andesites

Figure 6. Frequency distribution diagrams for olivine (a), orthopyroxene (b), clinopyroxene (c), and plagioclase (d) grouped based on the host rock composition.
Table 1. Temperature, pressure, and melt H$_2$O estimated for Cumaçay volcanics (B: basalt, BA: basaltic andesite, BTA: basaltic trachyandesite, A: andesite, D: dacite).

<table>
<thead>
<tr>
<th>Sample no.</th>
<th>Temperature (°C)</th>
<th>Pressure (kbar)</th>
<th>H$_2$O melt (wt%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cum175 (B)</td>
<td>1009–1090</td>
<td>1166–1187</td>
<td>1117–1153</td>
</tr>
<tr>
<td>Cum164(BA)</td>
<td>885–987</td>
<td>1121–1132</td>
<td>1094</td>
</tr>
<tr>
<td>Cum179(BA)</td>
<td>999</td>
<td>1140–1200</td>
<td>1153–1170</td>
</tr>
<tr>
<td>Cum181(BA)</td>
<td>842–888</td>
<td>1168–1186</td>
<td>1146–1161</td>
</tr>
<tr>
<td>Cum199(BA)</td>
<td>893–1019</td>
<td>1152–1171</td>
<td>1138–1170</td>
</tr>
<tr>
<td>Cum158(BTA)</td>
<td>857–1016</td>
<td>1050–1116</td>
<td>1042–1111</td>
</tr>
<tr>
<td>Cum160(A)</td>
<td>1081–1115</td>
<td>1045</td>
<td>1072–1084</td>
</tr>
<tr>
<td>Cum241(D)</td>
<td>954–967</td>
<td>954–990</td>
<td>964–974</td>
</tr>
</tbody>
</table>
and basaltic trachyandesite yield lower temperatures than the basaltic sample, ranging from 842 to 1019 °C and 848 to 1016 °C, respectively. Fe-Ti oxide crystallization temperatures from the dacite narrowly range between 954 and 967 °C.

The second method applied is the clinopyroxene-liquid thermometer of Putirka et al. (2003), which requires a pressure input and not water content. Pressure input has minimal effect on calculating the crystallization temperatures. Two different equilibrium tests were carried out to apply the clinopyroxene-liquid thermometers; the exchange coefficient between melt and clinopyroxene $K_D^{cpx-lq} = 0.28 \pm 0.08$ (Putirka, 2008) and the comparison of the measured clinopyroxene geochemistry with the predicted ones obtained from regression analysis of clinopyroxene-melt pairs. (Figure 7b). The whole-rock composition was assumed to be nominal melt composition in both tests. Only the measured clinopyroxenes that pass both requirements for equilibrium were used for temperature estimates. Clinopyroxene-melt pairs of basaltic sample have crystallization temperatures of 1166–1186 °C. Temperatures from basaltic andesites lie from 1140 to 1200 °C. Basaltic trachyandesite and andesite have crystallization temperatures of 1050–1116 °C and 1081–1115 °C, respectively. The calculated pressures from this method yield 3.7–6.6 kbar for basalt and 0.9–7.1 kbar for basaltic andesites (Table 1).

**H₂O contents**

The plagioclase-liquid hygrometer of Waters and Lange (2015) was used to calculate the preruptive melt H₂O contents of Cumaçay volcanics. The method is based on the crystal-liquid exchange reaction between the albite and anorthite components and is temperature-dependent (pressure has a minor effect). Bulk compositions of the rocks are used as the nominal liquid, and selected plagioclase core (an-max) compositions, most likely to be in equilibrium with the bulk rock, are used for the melt water concentrations. Our calculations used pressures and average temperatures from clinopyroxene–liquid (except dacitic sample) methods as P and T inputs. The temperature for a dacitic sample is from the Fe-Ti oxide thermometer. Calculated H₂O contents range from 0.6 to 0.7 wt% for basalt, 0.8 to 1.6 wt% for basaltic andesites, 2.1 to 2.2 wt% for basaltic trachyandesites, 1.6 to 1.9 wt% for andesite, and 3.7 to 4 wt% for dacite (Table 1).

**Oxygen fugacity**

Oxygen fugacity was defined using the Fe-Ti oxide equilibria following the same method for Fe-Ti oxide thermometry. Each Fe-Ti oxide pair was processed by the method reported by Spencer and Lindsley (1981), with the recalculation scheme of Stormer (1983), and calculated as log units relative to the FMQ buffer (Figure 7a). ∆FMQ values range from –0.9 to 1.77. The lowest fO2 is recorded by basaltic andesite (∆FMQ: –0.05 to +0.06; Cum181). The highest fO2 is from basaltic trachyandesite (∆FMQ: 1.21–1.77; Cum158).

3.5.2. Magma temperatures and pressures

Once obtaining the first T-P-H₂O wt%, we can apply other accurate methods to calculate crystallization P and T for Cumaçay volcanics. A summary of the different approaches is given in Table 1. The olivine-liquid (Putirka, 2008), clinopyroxene-liquid (Putirka, 2008), plagioclase-liquid (Putirka, 2005), and orthopyroxene-liquid (Putirka, 2008) geothermometers were used to monitor the crystallization conditions of the Cumaçay volcanics.

**Clinopyroxene-liquid**

The equilibrium conditions between clinopyroxene and nominal liquid (whole rock composition) are the same as described above. Melt H₂O contents are from the method of Walters and Lange (2015) (Table 1). Equations of 33 and 30 from Putirka (2008) are used to estimate temperature and pressure. Clinopyroxene-melt pairs of basaltic sample produce temperatures of 1117–1153 °C and pressures of 1.3–5.2 kbar. T and P conditions of basaltic andesites lie from 1094 to 1170 °C and 2 to 7.6 kbar, respectively. The basaltic trachyandesite and andesite have relatively lower temperatures ranging from 1042 to 1111 °C. The crystallization pressure of basaltic trachyandesite is in the range of 4.3–4.8 kbar. A single clinopyroxene-melt pair yields a pressure of 2.6 kbar for the andesite.

**Orthopyroxene-liquid**

Orthopyroxene compositions were evaluated for equilibrium with whole rock compositions based on $K_D^{opx-lq} = 0.29 \pm 0.06$ (Putirka, 2008) (Figure 7c). Crystals not in equilibrium with nominal melt are excluded and not considered in interpretations. Equilibrium mineral-melt pairs, the input of melt H₂O contents, and preferred pressures are given in Table 1. Based on the equations of 28a and 29a from Putirka (2008), basaltic andesite, andesite, and dacite have temperatures and pressures of 1152–1156 °C and 4 kbar, 1056–1079 °C and 1–3 kbar, 954–990 °C and 1–4 kbar, respectively.

**Olivine-liquid**

Olivine-liquid thermometry (Putirka, 2008; Eq.22) was used to calculate the temperatures for basalts and basaltic andesites. The composition of the whole rock is used for the nominal liquid composition. The mineral-liquid equilibria were checked using the method of Roeder and Emslie (1970). The phenocrysts that passed the equilibrium test ($K_D^{ol-liq} = 0.30 \pm 0.03$; Figure 7d) are used for temperature estimates. Since the olivine liquid equilibria is sensitive to pressure and water contents, we used calculated melt water contents from Walter and Lange (2015) (Table 1) and maximum pressure (7.6 kbar) from clinopyroxene-liquid equilibria (Table 1). The temperatures obtained from the olivine-liquid thermometer...
are in the range of 1211–1224 °C for basalts and 1152–1170 °C for basaltic andesites.

**Plagioclase-Liquid**

Putirka (2008) proposed that the temperature, pressure, and melt H₂O content have little effect on the Ab-An exchange coefficient. Thus the K_{D_{pl-liq}} (Ab-An) exchange between crystal and melt 0.28 ± 0.11 for T > 1050 °C and 0.1 ± 0.05 < 1050 °C can be used for equilibrium (Putirka, 2008) (Figure 7e). Plagioclase-liquid equilibrium temperatures and pressures are estimated by carrying out equations 23 and 25a from Putirka (2005) using estimated melt H₂O content for each rock group (Table 1). Based on the available plagioclase compositions, water, and pressure data, the plagioclase liquid geothermometer yield a temperature and pressure range of 1203 to 1207 °C and 6.9–7.8 kbar for basalts, 1138–1172 °C and 3.7–6.7 kbar for basaltic andesites, 1081 to 1087 °C and 3.2–6.54 kbar for basaltic trachyandesite, 1072 to 1084 °C and 2.8–4.9 kbar for andesite, and 964 to 974 °C and 1–1.4 kbar for dacite.

**Summary of temperature and pressure**

Calculated temperatures from diverse methods are broadly consistent with fractionation from basalts (≤1224 °C) to dacites (954 °C) (Figure 8a). Temperatures generally decrease with increasing SiO₂ contents of host rock SiO₂. However, andesite (Cum160) has a nearly similar temperature range to that of the basaltic trachyandesitic

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**Figure 7.** a) Temperature versus oxygen fugacity, calculated from Fe-Ti equilibria using the ILMAT program (Lepage 2003). b) Clinopyroxene-liquid equilibrium test according to observed and predicted (Dihd) compositions (Putirka, 1999). Equilibrium test based on the Fe-Mg exchange reaction between orthopyroxene (c) or olivine (d) and liquid (Fe-Mg_{opx-liq} = 0.29 ± 0.06 and Fe-Mg_{ol-liq} = 0.30 ± 0.03; Putirka, 2008). c) Plagioclase-liquid equilibrium test for the Ab-An exchange reaction by the model of Putirka (2005, 2008). The stability field was drawn using a value for Ab-An_{pl-liq} = 0.28 ± 0.11, T > 1050 °C (continuous lines), and Ab-An_{pl-liq} = 0.1 ± 0.05, T < 1050 °C (dotted lines).
lava (Cum158). Olivine-liquid temperatures overlap or are slightly higher than the plagioclase liquid temperatures in basalt and basaltic trachyandesites. Clinopyroxene-liquid temperatures have the lowest values in basalts; however, they mostly match up with the plagioclase-liquid and orthopyroxene-liquid temperatures in basaltic andesites, basaltic trachyandesite, and andesite. Fe-Ti oxide temperatures from diverse rock groups are predominantly from groundmass microcrystals rather than phenocrysts and lower than the other methods (except for dacite) and probably reflect most closely the temperature shortly before the eruption due to their fast reequilibration (Venezky and Rutherford, 1999) or underwent reequilibration upon slow cooling of the lava flow (e.g., Özdemir et al., 2011).

The preeruptive pressures using different thermodynamic barometers vary between approximately 8 kbar and approximately 1 kbar and mostly overlap (Figure 8b). Crystallization pressures from diverse methods for basalt and basaltic andesites mainly clustered between 3.7 and 7.8 kbar. Basaltic trachyandesite displays a wide range of crystallization pressures from 3.2 to 6.5 kbar. Andesite and dacite yield pressures of 1–4.9 and 1–4 kbar, respectively.

4. Discussion

4.1. Assessing magma differentiation processes

Cumaçay volcanics have low ranges of Mg#, Co, and Ni (Supplementary File 1), suggesting they are not primary melts and have undergone fractional crystallization. The covariation diagrams of whole-rock major and trace elements show negative correlations in CaO/Al₂O₃, MgO,
Sc, Ni, and Co with increasing content of SiO₂ (Figure 3), indicating crystallization of olivine and pyroxene from their parental melts. Drops in contents of Fe₂O₃ and TiO₂ (Figure 3) with increasing degrees of differentiation are consistent with the fractionation of Fe-Ti oxides that are ubiquitous as microcrystals. The lack of strong negative Eu anomaly in basaltic and intermediate rock indicates an insignificant role of plagioclase. However, a decrease in Sr contents of the rocks with SiO₂ contents > 62 wt% and slightly negative Eu anomaly in REE patterns (Figure 4) suggest the role of plagioclase in dacitic members.

The crystallization temperatures obtained from various thermometers also indicate a progressive decrease from basaltic members of Cumaçay towards more fractionated lavas. However, the ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd, isotopic compositions of the Cumaçay volcanics vary systematically between 87 Sr/86 Sr compositions of the Cumaçay volcanics vary systematically between 87 Sr/86 Sr and 143 Nd/144 Nd, isotopic basaltic members of Cumaçay towards more fractionated thermometers also indicate a progressive decrease from basaltic members of Cumaçay towards more fractionated lavas. However, the ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd, isotopic compositions of the Cumaçay volcanics vary systematically towards more fractionated thermometers also indicate a progressive decrease from basaltic members of Cumaçay towards more fractionated lavas. However, the ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd, isotopic compositions of the Cumaçay volcanics vary systematically towards more fractionated thermometers also indicate a progressive decrease from basaltic members of Cumaçay towards more fractionated lavas. However, the ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd, isotopic compositions of the Cumaçay volcanics vary systematically.
evolved samples (Figure 9), which could result from contamination from the continental crust. Since FC models fail to reproduce evolved acidic members in Na$_2$O and Al$_2$O$_3$ space and linear geochemical trends observed in CaO, MgO, and FeO (approximately 54–65 %wt SiO$_2$) contents, we have performed a series of MCS models to test whether the assimilation of wall rock melt during fractional crystallization could overcome these problems. We chose the same basalt (Cum175) that we used in FC models as parental magma composition, and Paleozoic schist (CMK13) represents the continental crust (wall rock composition) of the region (Table 2). Modeled major element compositions are given in Figure 11. Total anatectic melt assimilated by magma range between 22

Table 2. Parental melt (M), wall rock (WR), and recharge (R) compositions and parameters used in FC, AFC, and RFC runs*.

<table>
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<th>FC and AFC</th>
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<th>RFC</th>
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<tr>
<td></td>
<td>M (Cum175)</td>
<td>WR (CMK13)</td>
<td>M (#Cum238)</td>
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<tr>
<td>SiO$_2$ (%)</td>
<td>49.90</td>
<td>54.99</td>
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<td>Recharge trigger T(°C)</td>
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*For AFC runs parent magma Fe$^{2+}$/Fe$^{3+}$ was calculated at FMQ, FMQ+1, and FMQ+2 at liquidus temperatures after adding 0.5–1 wt% H$_2$O; for WR Fe$^{2+}$/Fe$^{3+}$ was calculated at FMQ, FMQ+1, and FMQ+2 at 750 °C and at 5 kbar after adding 1 wt% H$_2$O; for RFC run parent magma Fe$^{2+}$/Fe$^{3+}$ was calculated at FMQ+2 at liquidus and at 2.5 kbar after adding 4 wt% H$_2$O.
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and 25 mass unit. Compared to FC runs, AFC runs predict lower crystallinity ranging between approximately 74% and 54%. Crystallization phase assemblage is similar to the FC runs and composed of orthopyroxene (approximately 10–21 wt%), clinopyroxene (17–26 wt%), pigeonite (3–16 wt%), plagioclase (approximately 42–52 wt%), and Fe-Ti oxide (approximately 4–13 wt%). Apatite could not be stabilized in AFC runs. All major oxide contents of the modeled LLDs indicate higher values compared to the FC runs in a given SiO$_2$. The higher water contents do not have much effect on most of the major oxides; however, predicted higher Al$_2$O$_3$ contents depend upon the late crystallization of the plagioclase. Modeled curves at FMQ+1 and FMQ+2 with 1 wt% H$_2$O contents reproduce most of the basaltic, intermediate, and acidic members of Cumaçay for CaO, FeO, Al$_2$O$_3$, and to a lesser extent, MgO. Majority of Na$_2$O+K$_2$O contents of the Cumaçay volcanics ranging from basalts to dacites, replicated by FMQ+1 AFC run, which implies that the crustal contamination at mid to lower crustal depths could be responsible for the major element diversity of Cumaçay volcanics.

4.1.3. Recharge and fractional crystallization (RFC) model

The textural and compositional diversity of phenocrysts, wide distribution of calculated pressures and temperatures, and the existence of microcrystalline enclaves and crystal clots in andesites and dacites suggest magma mixing/recharge may be responsible for the generation of some of the evolved members of Cumaçay volcanics. Moreover, the linear trend and the mismatch between MCS-AFC curves and the Cumaçay whole rock data in the range of approximately 58–65 SiO$_2$ wt% and approximately 2.5–4 wt% MgO (Figure 11) also support this idea. To test the viability of mixing processes, we performed the RFC

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**Figure 10.** Fractional crystallization (FC) thermodynamic modeling results from Magma Chamber Simulator (MCS) software (Bohrson et al., 2014;2020) using the major element compositions of Cumaçay volcanics. Major elements are recalculated on an anhydrous basis.
model using dacite (Cum238) for resident magma and a basaltic trachyandesite (Cum179) for recharge magma compositions (Table 2). The initial water concentrations of samples are in accordance with the plagioclase–liquid hygrometer for similar rock groups. The RFC model terminates 40 wt% of magma crystallization. The composition of solid cumulate is approximately 17 wt.% orthopyroxene, approximately 10 wt% clinopyroxene, approximately 61 wt% plagioclase, and approximately 12 wt% Fe-Ti oxide, which is similar to that FC and AFC runs. As demonstrated in Figure 11, the liquid line descent show suitable matches with the intermediate and acidic samples for MgO and the alkalis (Na₂O+K₂O) for comparable SiO₂ concentrations. In the MgO diagram, the andesitic and the dacidic samples that show linear tendency between approximately 58–65 SiO₂ wt% and approximately 2.5–4 wt% MgO contents correlate better with modeled liquid after the recharge event. Assimilation of country rocks may also incorporate shallow-level processes.

4.2. Magma plumbing system
Our observation from mineral thermobarometry and MCS runs let us have some limitations on the subvolcanic magma plumbing system under the Cumaçay. We suggest a two-stage petrogenetic model similar to that proposed for the Süphan volcano by Özdemir et al. (2011). In our model, mantle-derived basalt intrudes the lower to mid-crust (deep magma reservoir), where it disturbs the geothermal gradient forming a deep-crustal hot zone comparable to the model of Annen et al. (2006), which is suggested for arc environments. The surrounding crust undergoes partial melting when it exceeds the solidus and forms hybrid melt that includes remaining melts from the crystallization of the basalt and anatectic melt from the wall rock. AFC processes produce evolved melts that may contain various
differing SiO₂ content distributed across a wide depth range. It is challenging to constrain the precise depth range of the hot zone; however, thermobarometry results indicate a crystallization pressure range of approximately 4–8 kbar or 14–28 km. Additionally, geophysical studies suggest that the thickness of the crust in eastern Anatolia is in the range of 40–45 km (e.g., Özacar et al., 2010; Alkan et al., 2020) and have a low-velocity zone between 20 and 30 km (e.g., Angus et al., 2006; Karaoglou et al., 2017; Zhu, 2018; Alkan, 2022), representing a partial melt pocket in the middle crust under the Late Cenozoic volcanic centers. The generated evolved melts migrate to shallow crustal levels, where they produce a subvolcanic magma reservoir. The crystallization pressures suggest around approximately 1.5–4.5 kbar or 5–15 km depth. Similar shallow magma storage regions have been reported for Plio-Quaternary (e.g., Karaoglou et al., 2018) and Quaternary (e.g., Özdemir et al., 2011; Macdonald et al., 2015; Karaoglou et al., 2017) volcanic centers in eastern Anatolia. The shallow magma storage region is where various melts come across and interact with each other. In addition to the wide range of calculated temperature/pressures and compositional variety of phenocrysts, the presence of crystal clots in andesite and dacite supply other evidence for the magma mixing (replenishment) process in a shallow magma reservoir. The plagioclase, pyroxene, and Fe-Ti oxide-bearing crystal clots in andesites and dacites indicate that they likely originated from the highly crystalline part (e.g., crystal mush) of the shallow reservoir. Injection of hot hybrid (or primitive) mafic magma into such reservoirs caused thermal erosion on crystal mushes (e.g., Neave et al., 2014; Zhang et al., 2015; Feng and Zhu, 2018). Finally, crystal clots can be formed by the disaggregation of those mushy zones (e.g., Feng and Zhu, 2018). The final chemical composition of intermediate and acid magmas of Cumaçay volcanics would be the result of mixing between evolved (e.g., dacite) and basaltic (e.g., basalt; basaltic trachyandesite) melts.

5. Conclusion
Cumaçay consists of genetically related lava flows ranging in composition from basalt to dacite, representing one of the Plio-Quaternary eruption centers of postcollisional volcanism in Eastern Turkey. Our new K-Ar ages and whole rock geochemistry indicate that the volcano started with the mildly alkaline basaltic lava flows (3.5 Ma), continued with subalkaline intermediate to acidic members (2.9–2.1 Ma), and ended up with transitional and subalkaline basaltic flows (2–0.97 Ma). Deep (approximately 14–28 km) and shallow (approximately 5–15 km) storage reservoirs have been recognized based on thermobarometric calculations beneath the Cumaçay. Our thermodynamic modeling using MCS reveals that the mantle-derived primitive magma intruded into the lower to the middle crust, where fractionation and crustal assimilation resulted in evolved melts with various SiO₂ contents. The generated evolved melts then migrate to shallow crustal levels, where they produce a subvolcanic magma reservoir. Petrographical observations, combined with whole-rock major element modeling, reveal that the evolved rocks (andesite and dacite) may be produced via combined processes of fractional crystallization, the interaction of colder (e.g., dacite) and hotter magmas (e.g., basalt; basaltic trachyandesite) ascending from depth and recycling of early formed crystals at shallow magma storage region.

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Supplementary Data
Supplementary data can be accessed at the following link: https://aperta.ulakbim.gov.tr/record/252343#.ZDfLRXZByUk

References


