The nature of the proximal volcaniclastic materials from the 2021 eruption Fukutoku-1 Oka-no-Ba in the Izu-Bonin arc. 2 3 4 Kenta Yoshida\*, Yoshihiko Tamura, Tomoki Sato, Erika Tanaka, Noriko Tada, Morihisa Hamada, Takeshi Hanyu, Qing Chang, Shigeaki Ono 5 6 7 Research Institute for Marine Geodynamics, Japan Agency for Marine-Earth Science and 8 Technology, Natsushima-cho 2-15, Yokosuka, 237-0061 Japan. 9 10 Correspondence 11 Kenta Yoshida, Natsushima-cho 2-15, Yokosuka, 237-0061 Japan. 12 E-mail: yoshida ken@jamstec.go.jp 13 **Funding Information** 14 Japan Society for the Promotion of Science, KAKENHI, Grant Nos. JP19K14825 and 15 JP19H01999 to K.Y. and JP21H01195 to Y. T. 16 17 Conflict of interest disclosure 18 19 There are no entities or relationship, etc. presenting a potential conflict of interest requiring 20 disclosure in relation to this manuscript. 21 22 This is a non-peer review preprint that has been submitted to *Island Arc*; future versions may have 23 different content. 24 25

#### Abstract

Fukutoku-Oka-no-Ba is a submarine volcano located at 24°17.1′N/141°28.9′E in the Izu–Bonin–Mariana arc, and is one of the most active volcanoes in Japan. This volcano produced an explosive eruption in August 2021 that generated a large amount of volcaniclastic material, some of which drifted westward to Japan and the coastal area of East Asia as a pumice raft. The pumice clasts that drifted for >1000 km were mostly homogeneous and identical to those produced by past historical eruptions. The clasts have trachytic compositions (SiO<sub>2</sub> = 61-63 mass% and Na<sub>2</sub>O+K<sub>2</sub>O = 8.6-10.0 mass%) and contain augite, plagioclase, olivine (Mg#  $\sim$ 65), and magnetite, along with a small number of mafic enclaves containing diopside and high-Mg olivine (Mg#  $\sim$ 92).

We undertook a research cruise to investigate the proximal volcaniclastic materials by dredging. The proximal materials include pumice, weakly vesiculated lapilli, and volcanic rocks, which have trachytic composition (SiO<sub>2</sub> contents up to 64.5 mass%). The main minerals in the proximal material are similar to those in the drift pumice, although remnants of mafic magma do not occur in the SiO<sub>2</sub>-rich samples. The compositional variations of the SiO<sub>2</sub>-rich proximal samples are due to plagioclase-dominated fractional crystallization, and the higher-SiO<sub>2</sub> samples yielded lower temperature and pressure among the studied samples.

The petrographic and geochemical characteristics of the proximal and drift ejecta from Fukutoku-Oka-no-Ba suggest the magma reservoir was stratified into two parts. The lower part experienced magma mixing with a limited volume of mafic magma, whereas the upper part underwent plagioclase-dominated fractional crystallization. The accumulation of differentiated high-SiO<sub>2</sub> magma in the upper part of the magma reservoir might have increased the pressure in the magma conduit, which then exceeded the critical pressure required for an explosive eruption.

Keywords: Fukutoku-Oka-no-Ba, Izu-Bonin-Mariana arc, phreatomagmatic eruption, drift pumice

# 1. Introduction

A large volume of pumice clasts was generated by the 2021 eruption of Fukutoku-Oka-no-Ba (FOB) volcano, which is located at 24°17.1′N/141°28.9′E in the Izu–Bonin–Mariana arc (Fig. 1a). This is one of the most active volcanoes in Japan. The pumice clasts drifted and became distributed over a wide area around the islands of Japan and in many places in the East Asia, including Taiwan, Philippines, and Thailand (Yoshida et al., 2022a, b, c). The large amounts of drift pumice filled port areas and hindered vessel navigation, which led to significant economic losses (Maeno et al., 2022; Yoshida et al., 2022a, c; Asami and Takahashi, 2023). The drift pumice is geochemically homogeneous (Yoshida et al., 2022a) and similar to that generated during past eruptions in 1904, 1914, 1986, and 1992 (Tsuya,

1937; Yoshida et al., 1987; Kato, 1988; Nakano and Kawanabe, 1992; Sun et al., 1998), although the color and texture of the pumice are variable (Yoshida et al., 2022a). A notable characteristic of the pumice generated by FOB is the minor but common occurrence of black-colored pumice, which consists mainly of transparent brown glass with magnetite nanocrystals (i.e., nanolites) (Yoshida et al., 2022a). Although recent studies have focused on nanolite precipitation and the mechanism of large eruptions (Cáceres et al., 2020, 2021; Di Genova et al., 2020; Scarani et al., 2022), the actual processes that trigger eruptions remain unclear (Dubosq et al., 2022). The biotite-bearing hydrous mineral assemblage of nanolite and higher Fe<sup>3+</sup>/ΣFe ratio of the nanolite-bearing glass than the nanolite-free glass indicate that oxidation due to fluid infiltration induced nanolite-precipitation and subsequent bubble nucleation, which resulted in the explosive eruption (Yoshida et al., 2023).

The 2021 eruption of FOB was large enough to be observed by satellite imaging, and the volcanic column reached a height of ~16 km, corresponding to stratospheric level (Maeno et al., 2022; Yoshida et al., 2022a). Based on Himawari-8 satellite images, Maeno et al. (2022) provided a detailed timeseries of the phreatomagmatic eruption from FOB during 13–15 August 2021. The 2021 FOB eruption column consisted of a vapor-rich plume and a small amount of volcaniclastic materials. The pumice contains virtually no hematite nanolites, which indicates that most of the pumice clasts were ejected into water and subsequently floated due to their buoyancy (Maeno et al., 2022; Yoshida et al., 2023). Whether the pumice floated or sunk depended on its microtexture (Mitchell et al., 2021), and the microtexture of the FOB pumice varies widely due to the presence of nanolites (Yoshida et al., 2022a, 2023). The drift pumice sampled on-land only represents the pumice clasts continued to float over a long transport distance (up to 4000 km). Yoshida et al. (2022b, c) noted that the pumice rafts that drifted longer distances tended to comprise smaller clasts with lesser amounts of black pumice that contained larger vesicles.

To understand the nature of the 2021 eruption and magmatic system of FOB, it is necessary to investigate the full range of erupted volcaniclastic materials, including those that did not float. As such, we undertook a research cruise using the R/V *Yokosuka* to investigate these deposits. A dredge survey was undertaken inside and outside the submarine caldera to collect fresh rock samples, both with and without significant vesiculation. In this paper, we describe the petrographic and geochemical features of these samples. These new and previously reported data constrain the nature of the magmatic system of FOB, which can be used to predict and mitigate hazards from submarine volcanoes in the far-off sea.

## 2. Materials and Methods

### 2.1. Geological Background and Research Cruise

FOB is one of the most southern volcanoes in the Izu-Bonin arc. The volcano is located within a

large volcanic complex that extends approximately 15 and 30 km for E–W and N–S direction, respectively (Fig. 1b). The complex comprises the Kita-Fukutoku-Tai volcano, Kita-Fukutoku caldera, and Minami-Ioto volcano (from north to south), and FOB is the central cone of Kita-Fukutoku caldera, which is ~2 km in diameter (Ito et al., 2011). The summit of FOB before the 2021 eruption had an oval shape (elongate NE–SW) and was flat at a depth of ~30 m below sea level (Ito et al., 2011). Seismic and geomagnetic surveys indicate that a low-velocity zone exists to the north of FOB, which can be attributed to the partially molten region of the magma reservoir (Nishizawa et al., 2002; Onodera et al., 2003). The magmatic system of FOB and Kita-Fukutoku caldera are still poorly understood. Based on Nd and Pb isotopic compositions, Sun et al. (1998) suggested that the magma of FOB is distinct from those of the nearby volcanoes of Ioto, but similar to those of the Hiyoshi Volcanic Complex in the northern part of the Mariana arc.

The 2021 FOB eruption occurred in the morning of 13 August 2021 and continued to the morning of 16 August (Japan Meteorological Agency, 2021). Underwater sound and infrasound remote observations indicate that the eruption started at 5:55 AM (in Japan Standard Time; Maeno et al., 2022; Metz, 2022).

We undertook a geological survey and sampling around FOB during six dredging operations using the R/V *Yokosuka* during cruise YK22-15 from 14 to 27 August 2022. A double-towing dredging system was used (Supplementary Fig. 1). Figure 1b shows the locations of the dredging operations. Given that the cruise was carried out just 1 yr after the explosive eruption, we remained at least two nautical miles (~3.7 km) from the volcanic vent for safety reasons (green circle in Fig. 1b). We consider that the samples collected by the dredges represent the geology of the volcano, although there remains uncertainty as to whether the dredged samples are representative of their specific locations.

Three dredges (YK22-15 D03, D05, and D08) targeted the western side of the volcanic vent along the western slope of the caldera. YK22-15 D08 was a continued dredging operation of YK22-15 D03, because the D03 line was cancelled due to stacking of the dredger. YK22-15 D06 targeted the eastern slope of the outer rim of the crater and YK22-15 D07 targeted the outer slope of the topographic high from the rim of the crater. YK22-15 D04 targeted the southern slope of Kita-Fukutoku-Tai. Detailed locations of the dredges are described in the Supplementary Material.

2.2. Analytical Methods

Mineral and volcanic glass compositions were determined using a field emission gun electron microprobe (EMP) analyzer equipped with five wavelength-dispersive X-ray detectors (JEOL: JXA-8500F) at Japan Agency for Marine-Earth Science and Technology (JAMSTEC; Yokosuka, Japan). Natural and synthetic standards were used to calibrate the quantitative analyses following the procedure of Yoshida et al. (2022a). The analytical conditions were 15 kV and 10 nA for the accelerating voltage and beam current, respectively, except for analyses of olivine. For some olivine

grains, we used an accelerating voltage of 20 kV and beam current of 25 nA. The beam diameter was set to 3  $\mu$ m for minerals and 5  $\mu$ m for glass. Fe<sup>3+</sup>/ $\Sigma$ Fe values for clinopyroxene were calculated such that the total cations were four on a six-oxygen basis, whereas Fe<sup>3+</sup>/ $\Sigma$ Fe values for magnetite were calculated for Fe<sup>2+</sup> + Mg + Mn = 1 on a four-oxygen basis. All Fe in olivine was assumed to be ferrous. Anorthite, albite, and orthoclase contents of plagioclase were calculated as Ca/(Ca+Na+K)×100, Na/(Ca+Na+K)×100, and K/(Ca+Na+K)×100, respectively.

Cathodoluminescence analysis of silica minerals was conducted using a scanning electron microscope (ThermoScientific: Quanta FEG450) equipped with a cathodoluminescence system (Gatan: MonoCL 4) at JAMSTEC.

Raman spectra were obtained with a Raman spectrophotometer (RAMANtouch VIS-HP-MAST; Nanophoton) equipped with a 532 nm semiconductor green laser at JAMSTEC. The laser power on the sample surface was 1-2 mW, and data were acquired in  $2 \times 20$  s cycles to eliminate accidental cosmic rays. The spectrometer was calibrated to the 520.7 cm<sup>-1</sup> peak of a Si wafer.

Whole-rock major element compositions were determined following the methods of Tani et al. (2005) and Sato et al. (2020), by X-ray fluorescence (XRF) spectrometry (Rigaku ZSX Primus II). Prior to analysis, the samples were crushed to pebble size (5–10 mm) and desalinated using hot water (~40 °C) and by boiling in Milli-Q water. Desalinization was checked using a AgNO<sub>3</sub> solution and corresponding precipitation of AgCl, so that no precipitation occurs. The desalinized samples were then washed with Milli-Q water and acetone in an ultrasonic bath, and powdered in an agate mortar or with a Multi-beads Shocker pulverizer. Finally, a mixture of 0.4 g of sample powder and 4 g of Li<sub>2</sub>B<sub>4</sub>O<sub>7</sub> was fused and made into glass beads for XRF analysis. Accuracy and reproducibility of the major element data are better than  $\pm 1\%$  and  $\pm 2\%$  (relative standard deviations), respectively.

We also analyzed the whole-rock trace element compositions by solution inductively coupled plasma-mass spectrometry (ICP-MS; iCAP Qc; ThermoFisher Scientific). The rock powders were digested in HF, HClO4, and HNO3. We also analyzed a reference basalt (JB-2; Jochum et al., 2016), which yielded results that are in good agreement with certified values (Table S1).

## 3. Results

## 3.1. Sample descriptions

The recovered rock samples were initially described onboard. The samples include gray to brown pumice, black volcanic rock with variable degrees of vesiculation, and tuff. For convenience, we omit the cruise number (YK22-15) and describe the dredge number with the prefix D and the sample number with the prefix R. For example, D03R01 is the first rock sample from dredge number three.

The rock samples are generally fresh and without Mn oxide coatings. The pumice clasts are mostly rounded. In contrast, the black volcanic rock samples are angular to subangular, indicating subsequent

reworking was insignificant. The samples collected from the western lines (D03, D05, and D08) contain a considerable amount of pumice and angular volcanic rock clasts that are <10 cm in length (Fig. 2a-b). The most common pumice is gray in color, and is similar to those collected as drift pumice (Yoshida et al., 2022a), while some pumice is dark gray to black (Fig. 2c). Although the drift pumice contains black spots including mafic enclaves and plagioclase-dominated crystal clots with nanolitebearing glass, such features are rare in the dredged samples. Tuff breccia clasts up to 20 cm in size were also recovered. Several black lapilli clasts with a vitreous luster (i.e., obsidian) were also found (Fig. 2d). Based on the deep-sea camera observations, the seafloor around the D08 position was calm and covered with white volcanic ash (Fig. 1c). The dredge D06 was conducted on the southeastern side of the central vent, and sampled black volcanic rocks with variable degrees of vesiculation that were up to 10 cm in size (Fig. 2e). Smaller pumice clasts were also recovered. During dredge D07 on the northeastern slope of the outer caldera rim, an outcrop of black volcanic rocks was observed by the deep-sea camera (Fig. 1d). The volcanic rock sample D07R01 has a pillow-like structure (Fig. 2f), indicative of the subaqueous extrusion. Dredge D07 also recovered large pumice clasts up to 22 cm in length, which have a well-developed tubular texture (Fig. 2g). This texture is referred to as woody pumice, and is considered to form during a submarine eruption (Kato, 1987).

Nine representative samples, including volcanic rock and pumice clasts, were selected for petrographic and geochemical investigations.

# 3.2. Whole Rock Geochemistry

The nine samples have trachytic compositions regardless of their appearance (i.e., glassy volcanic rock/lapilli or pumice; Table 1; Fig. 3a), with higher  $SiO_2$  (63.7–64.4 mass% on an anhydrous basis) and total alkali ( $Na_2O+K_2O=10.2-10.4$  mass%) contents as compared with the drift pumice (Yoshida et al., 2022a, b), except for one sample (Fig. 3a). The glassy lapilli sample D06R02 was collected from the southeastern slope of the caldera and has a slightly  $SiO_2$ -poor composition that overlaps those of the drift pumice. The whole rock major element compositions of the dredged samples and drift pumice define linear trends in Harker diagrams (Figs. 3b–d).

Plots of SiO<sub>2</sub> versus selected trace elements are shown in Figures 3e–f. Three drift pumice compositions reported by Yoshida et al. (2022a) are also shown. Zr/Y and La/Sm ratios of the dredged samples are approximately 9.0 and 8.3, respectively, which are higher than those of the drift pumice, except for sample D06R02 (Figs. 3e–f) that has a similar composition as the drift pumice.

N-MORB-normalized trace element diagrams show that the studied samples have similar patterns to the drift pumice from the 1986 and 2021 eruptions (Fig. 3g)

## 3.3. Petrography and Mineral/Volcanic Glass Chemistry

Based on the occurrence of microlites that are visible under an optical microscope, the pumice and

202 volcanic rock samples respectively were subdivided into microlite-rich and microlite-free types (Table 203 1; Figs. 4a-b). Regardless of the appearance of the pumice or volcanic rock, and occurrence of 204 microlites, all the samples contain a similar phenocryst mineral assemblage of clinopyroxene (Cpx), 205 plagioclase (Pl), olivine (Ol), and magnetite (Mag) (Fig. 4b), with minor amounts of apatite occurring as inclusions and a groundmass phase (Fig. 4c). In a few samples, pyrrhotite occurs as inclusions in 206 207 clinopyroxene and plagioclase. The microlite-rich pumice clast D08R11 contains cristobalite (Crs) in 208 relatively large vesicles (Figs. 4e and 5d).

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The microlite-free glassy samples (D06R02 and D08R04) contain pale brown groundmass glass (Fig. 4a) that is weakly vesiculated. In contrast, the microlite-rich glassy samples (D03R01, D07R01, D07R05, and D06R01) contain dark brown groundmass glass (Fig. 4b) with abundant microlites of plagioclase and magnetite (Fig. 5a-b). Figure 4b shows that the abundance of microlites is nearly zero in the interstitial glass between the phenocryst minerals. Raman microscopy revealed that the dark brown glass in the microlite-rich samples has a Raman peak of magnetite nanolites at ~670 cm<sup>-1</sup>, while this peak was not recognized for the pale brown glass in the microlite-free samples (Fig. 4f).

The microlite-free pumice sample (D03R03) is gray and has a strongly vesiculated groundmass that consists of colorless glass. The glass surrounding the plagioclase phenocrysts is colorless and nanolitefree (Fig. 4c), while the plagioclase phenocrysts in the drift pumice is commonly associated with nanolite-bearing brown glass, even though the groundmass glass is generally colorless. Figure 4d shows a plagioclase phenocryst in a gray-type drift pumice from the FOB eruption that was collected in Thailand and described by Yoshida et al. (2022b). The nanolite-rich pumice sample (D08R11) has a dusty and highly vesiculated groundmass (Fig. 4e). Although most vesicles are very small (<50 μm), we identified large vesicles (>100 μm) that occasionally contain spherical cristobalite (Fig. 4e).

Mafic magma components such as mafic enclaves and Mg-rich olivine (Mg# ~90) were not recognized in the dredged samples.

Below, we describe the characteristics of respective minerals and volcanic glass. Representative EMP analyses are shown in Tables 2–4.

Plagioclase is the most abundant minerals in the studied samples, and is generally euhedral and up to 5mm in length. Plagioclase generally has a Ca-rich core with An<sub>40-49</sub>Ab<sub>54-48</sub>Or<sub>4-3</sub> and a Ca-poor rim with An<sub>30-38</sub>Ab<sub>56-62</sub>Or<sub>7-4</sub> (Figs. 5a and 6a), which is similar to the core-rim compositions reported for the drift pumice from FOB (Yoshida et al., 2022a). Brown-colored melt inclusions are common, some of which contain optically-visible microlites of magnetite.

Clinopyroxene in the studied samples has an almost homogeneous augite composition with Mg# ~80 (Fig. 6b). Samples D06R01 and D07R05 exhibit significant zonation in Fe-Mg ratios, with D07R05 having cores with Mg# = 93 and rims with Mg# = 78, and D06R02 having cores with Mg# = 72 and rims with Mg# = 80 (Figs. 5b & 6b).

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Olivine in the studied samples exhibits mostly homogeneous with Fo<sub>62-65</sub>, where olivine in sample

D07R01, D07R05, and D08R11 exhibits increase in Fe contents in the outermost rims (Fig. 5c).

Magnetite in the studied samples occurs both as inclusions and a groundmass mineral. The magnetite contains considerable amounts of TiO<sub>2</sub> (up to 11 mass%) and Al<sub>2</sub>O<sub>3</sub> (up to 3.2 mass%). The MgO content of magnetite is a useful proxy for temperature (Canil and Lacourse, 2020). The  $X_{Mg-mag}$  values (Mg/[Mg +  $\Sigma$ Fe]) vary among samples, but are mostly  $X_{Mg-mag}$  = 0.06 (Fig. 6c). D07R01 has the lowest  $X_{Mg-mag}$  values for fine-grained magnetite in the groundmass ( $X_{Mg-mag}$  = 0.044–0.048), whereas the magnetite in Ca-poor rims yielded higher  $X_{Mg-mag}$  values of 0.053-0.057 (Fig. 5a). The highest  $X_{Mg-mag}$  values were obtained for sample D06R02, which is a microlite-free volcanic rock, with  $X_{Mg-mag}$  = 0.063–0.072.

The presence of cristobalite in sample D08R11 was confirmed by Raman spectroscopy (Fig. 5d). EMP analysis showed the impurity of Al<sub>2</sub>O<sub>3</sub> and Na<sub>2</sub>O for 1.0 and 0.6 mass%, respectively (Table 4), corresponding to similar molar amounts (~0.0115 on a two-oxygen basis; Table 4). This is indicative of the incorporation of Al and Na as [AlO<sub>4</sub>/Na<sup>+</sup>]<sup>0</sup> (Schipper et al., 2020). Cathodoluminescence imaging of the cristobalite clearly revealed a radial zoning, which is characteristic of chemical vapor deposition (Schipper et al., 2020). The occurrence of cristobalite in sample D08R11 is similar to that of other deep-sea cristobalite occurrences, such as rhyolite in the Havre, Kermadec Arc (Ikegami et al., 2018), and Kikai Caldera, Japan (Hamada et al., 2023).

Glass chemistry was also determined by EMP analysis. For the microlite-rich samples, the sites of EMP analyses were carefully determined using backscattered electron images and the results showing the mineral signature were rejected as mixtures. The glass in the dredged samples has relatively high SiO<sub>2</sub> (>65 mass%) and total alkali (Na<sub>2</sub>O+K<sub>2</sub>O > 10 mass%) contents. The glassy lapilli sample (D06R02), with a whole-rock composition similar to those of the previously reported drift pumice data (Fig. 6d) also has a similar glass composition as the drift pumice (Maeno et al., 2022; Yoshida et al., 2022a). The microlite-rich pumice sample (D08R11) has a wide range of SiO<sub>2</sub> contents (64–73 mass%) and exhibits a negative correlation between SiO<sub>2</sub> and Na<sub>2</sub>O + K<sub>2</sub>O (Fig. 6d). The glass compositions for each sample vary with the whole-rock compositions except for sample D08R11. For example, D07R01 and D07R05 have the most SiO<sub>2</sub>-rich compositions (Fig. 6e). We did not find melt inclusions with low SiO<sub>2</sub> contents (<60 mass%) like those reported from the drift pumice.

The groundmass glass in sample D08R11 is heterogeneous with respect to halogen contents, and has a lower Cl content (<0.2 mass%) as compared with the other samples (0.3–0.4 mass%; Fig. 6e). The groundmass glass of the eastern dredge samples (D07R01, D07R05, and D06R01) has relatively higher Cl contents (up to 0.4 mass%), whereas samples from the western dredges and sample D06R02 have Cl contents of ~0.3 mass%.

#### 4. P-T Estimates

Since pressure estimation using the available mineral paragenesis is not easy, Yoshida et al. (2022a) used a machine-learning-based clinopyroxene single mineral geobarometer (Petrelli et al., 2020) for the drift pumice from FOB, and obtained a pressure of  $\sim$ 250 MPa for the magma reservoir. In the present study, we also used the same clinopyroxene geobarometer (Table 3). Most samples consistently yielded pressures range of 215–279 MPa, except for two samples that yielded higher (D03R01 = 388 MPa) and lower (D07R01 = 177 MPa) pressures. The cores of zoned clinopyroxene in samples D07R05 and D06R01 yielded higher pressures of 445 and 371 MPa, respectively.

Another pressure indicator is the Cl content of groundmass glass. Figure 6e shows that the western samples (D03R01, D03R03, and D08R04) and sample D06R02 have Cl contents of ~0.3 mass%, whereas the eastern samples (D07R01, D07R05, and D06R01) have Cl contents of up to 0.4 mass%. The glass in sample D08R11 has very low Cl contents down to mostly zero. Based on the experimental studies, the maximum Cl solubility in alkaline melts, such as trachyte, exhibits a negative correlation with increasing pressure and is mostly independent of for compositional differences (Signorelli and Carroll, 2002; Balcone-Boissard et al., 2016). A Cl content of 0.4 mass% corresponds to a pressure of <200 MPa, and thus the higher Cl contents of the eastern samples (except for D06R02) can indicate a lower pressure of their equilibration.

Temperature conditions can be determined using the magnetite geothermometer (Canil and Lacourse, 2020) where  $X_{Mg\text{-mag}}$  values can be transformed into temperatures. As shown in Figure 6c, the calculated temperatures for most samples are 920–940 °C, where sample D06R02 yielded higher temperatures of 940–960 °C. Rim compositions of the groundmass magnetite in some samples (D08R11, D07R01, F07R05, and F06R01) yielded temperatures of <900 °C (Fig. 6c). All of these low-temperature samples contain abundant microlites in the groundmass. In sample D07R01, magnetite inclusions in the plagioclase rims yielded  $X_{Mg\text{-mag}} \sim 0.06$ , corresponding to a temperature of ~925 °C whereas those in the groundmass yielded temperatures of 882–896 °C.

Other conventional geothermometers were applied to mineral-melt pairs carefully selected after thinsection observations. The olivine-melt geothermometer (Putirka et al., 2007) was applied to a melt inclusion and surrounding olivine in sample D07R05 and yielded a temperature of 940 °C. The olivine-melt pair in sample D03R01 yielded a higher temperature of 962 °C. The clinopyroxene-melt geothermometer (Neave and Putirka, 2017) was applied to a melt inclusion-clinopyroxene pair in sample D03R01 and yielded a temperature of 981 °C and pressure of 90 MPa. According to Perkins and Vielzeuf (1992), olivine and clinopyroxene compositions can exist in equilibrium, and thus olivine-melt and clinopyroxene-melt pairs might reflect similar conditions in the magma reservoir, although the pressure value obtained from clinopyroxene-melt pair (D03R01) is apparently lower than other estimates. The plagioclase-melt geothermometer (Putirka, 2008) was applied to the plagioclase

rims  $(An_{31}Ab_{61}Or_8)$  and groundmass glass in sample D07D01, and yielded a temperature of 908–964 °C under the assumption of 0–2 mass % of  $H_2O$  in melt.

In summary, the coarse-grained phenocryst minerals consistently yielded temperatures of 920–980 °C, whereas rims of coarse-grained magnetite and fine-grained magnetite recorded lower temperatures down to 882 °C.

#### 5. Discussion

## 5.1. The variation of magma composition

Bivariate compositional diagrams of the major elements (Figs. 3b–d) show we linear trends in the low- and high-SiO<sub>2</sub> ranges (61.0–63.5 and 63.5–64.5 mass%, respectively). The reported whole-rock compositions of the drift pumice, including past eruptions, are variable and scattered (Fig. 3a), and plot in the low-SiO<sub>2</sub> range. Selected trace element ratios also exhibit differences between low- and high-SiO<sub>2</sub> samples (Fig. 3e-f). We also show the trace element composition of mafic melt inclusions found in olivine and clinopyroxene (Yoshida et al., 2022a). In the magmatic system of FOB, remnants of primitive mafic magma are limitedly recognized, and include mafic melt inclusions (Maeno et al., 2022; Yoshida et al., 2022a), high-Mg olivine (Kato, 1988; Yoshida et al., 2022a, b), and calcic plagioclase (Yoshida et al., 2022a), most of which occur within or are closely associated with black pumice (Yoshida et al., 2022a, b). The lower Zr/Y and La/Sm ratios of the drift pumice (i.e., low-SiO<sub>2</sub> range) as compared with the dredged samples (i.e., high-SiO<sub>2</sub> range) can be explained by the effect of the mafic melt component, although the whole-rock compositions show little change. Relatively scattered compositions in the low-SiO<sub>2</sub> range (Fig. 3a) could reflect heterogeneity in the magma reservoir caused by partial mixing with a small amount of mafic component.

In contrast, the compositional trends in the higher-SiO<sub>2</sub> range are characterized by moderate changes in all elements and a CaO decrease with increase in SiO<sub>2</sub>. We modeled the compositional change from the medium-SiO<sub>2</sub> sample (D08R04) to the highest-SiO<sub>2</sub> sample (D07R05) by assuming fractional crystallization of the minerals in sample D08R04 (Tables 2-4). As a result, the difference between samples D08R04 and D07R05 can be best explained by extraction of 5.6% plagioclase, 0.3% clinopyroxene, 0.2% olivine, and 0.5% magnetite (i.e., fractionation path A in Figs. 3b–d).

The model calculations indicate that the FOB magma compositions reflect the occurrence of mixing and fractionating domains. The former domain is characterized by low-SiO<sub>2</sub> compositions partly due to the involvement of a small amount of mafic magma. This domain is heterogeneous which may be due to incomplete mixing. In contrast, the SiO<sub>2</sub>-rich samples represent a gradual change due to plagioclase-dominated fractionation. The SiO<sub>2</sub>-rich samples contain high-Cl glass (Fig. 6f), which is an indicative of low pressure, and this fractionating domain might represent the upper most part of the magma reservoir that is not mixing with the deeper part of the reservoir.

5.2. Implications for the FOB Magmatic System

We undertook dredge sampling around the area of FOB, including the outer slope of Kita-Fukutoku caldera (Fig. 1b). Although previous studies reported the drift pumice had a trachytic composition with  $SiO_2 <\sim 63$  mass%, our dredged samples included more  $SiO_2$ -rich samples with  $SiO_2 > 64$  mass%. Both the  $SiO_2$ -rich and -poor samples have high total alkali contents of up to 10.5 mass%, and thus we consider that all these samples represent the magma reservoir of FOB.

The crustal structure beneath FOB was investigated by acoustic and geomagnetic surveys of Nishizawa et al. (2002) and Onodera et al. (2003). The geomagnetic data indicate a low geomagnetic anomaly in the northwestern region of the volcanic vent of FOB. Onodera et al. (2003) interpreted this anomaly to represent partially molten rocks (i.e., the magma reservoir).

The dredge D07 was conducted on the outer slope of Kita-Fukutoku caldera and volcanic rock samples were collected from a seafloor outcrop (Fig. 1d). Although it is unclear whether these samples were generated by the recent volcanic activity of FOB or represent an older volcanic unit of Kita-Fukutoku caldera, a consistent trend including volcaniclastic materials collected from the inside of the caldera indicates that the chemical composition of the D07 samples reflects the endmember characteristics of the current FOB magma reservoir. The samples from D07 have the most SiO<sub>2</sub>-rich compositions and lowest temperatures of ~880 °C (Fig. 6c), which can be interpreted to reflect plagioclase-dominated fractional crystallization during cooling.

The pumice sample containing cristobalite (D08R11) possibly represents remnant volcaniclastic material deposited within the conduit, which was ejected during the early stage of the eruption. Cristobalite occurring in vacancies in volcanic rocks are common in submarine volcanoes (e.g., Ikegami et al., 2018; Hamada et al., 2023), as well as in on-land volcanoes (e.g., Schipper et al., 2017). Schipper et al. (2017, 2020) suggested that cristobalite crystallizes by chemical vapor deposition due to the degassing of halogen species (HF and HCl), and provides evidence of shallow and slow cooling typical of a volcanic plug. After the 2021 eruption, FOB exhibited weak degassing activity, which is evident from the bubbles reaching the sea surface above the summit (Fig. 7a). Samples such as D08R11 could have been generated during inter-eruption activity in a blocked volcanic vent.

Consequently, our results for the dredged samples can be summarized as follows (Fig. 7b). The magma reservoir of FOB is heterogeneous and consists of two domains. Most of the volcaniclastic materials ejected by the 2021 FOB eruption (i.e., pumice) were derived from the deeper low-SiO<sub>2</sub> domain. The deeper domain is characterized by the involvement of a small amount of mafic magma, as is evident from some trace elements (Figs. 3e–f), and the limited occurrence of mafic enclaves (Yoshida et al., 2022a) and nanolite-bearing black pumice (Yoshida et al., 2023). Based on the microto nano-scale petrographical observation, Yoshida et al. (2023) suggested that the intrusion of hydrous mafic magma induced oxidation and corresponding nanolite-precipitation in the limited volume of the

magma which enhanced heterogeneous bubble nucleation (Pistone et al., 2017; Di Genova et al., 2020). This bubble nucleation served magma convection in the deeper domain, resulting in incomplete mixing and considerable heterogeneity. The shallower part is characterized by higher-SiO<sub>2</sub> contents and lower pressures of <200 MPa, corresponding to a depth of <8 km assuming a crustal density of 2400 kg/m³. The shallower domain consists of more differentiated magma, indicating the gravity differentiation. The SiO<sub>2</sub>-rich less-dense magma may have avoided magma convection and mixing with the deeper domain and does not contain mafic magma components. As a result, the whole-rock and melt compositions became more SiO<sub>2</sub> rich, and plagioclase became more stable at lower temperatures. This SiO<sub>2</sub>-rich magma was either ejected with the pumice and sunk immediately after the explosive eruption, or was effusively erupted from a location other than central vent of FOB, such as dredge site D07. The storage of the differentiated high-SiO<sub>2</sub> magma in the upper part of the magma reservoir might have plugged the volcanic conduit and increased the pressure, which ultimately led to the high explosivity of the 2021 eruption.

# Summary

Proximal samples of volcaniclastic sediments and volcanic rocks of FOB exhibit differences as compared with drift pumice collected from coastal areas of Japan (i.e., >1000 km from the volcano). The proximal rocks include pumice and volcanic rocks, most of which have higher SiO<sub>2</sub> contents (>63.5 mass%) as compared with drift pumice from the 2021 and earlier eruptions (61–63 mass%), although the constituent minerals are the same. Compositional variations of the SiO<sub>2</sub>-rich proximal samples resulted from plagioclase-dominated fractional crystallization, and the higher-SiO<sub>2</sub> samples yielded lower temperatures and pressures. The petrographic and chemical characteristics of the proximal and drifted ejecta from FOB suggest the magma reservoir contains two distinct domains.

# Supplementary Materials

The detailed positions of the dredges, reference standard analysis for the whole-rock trace element composition, and the outline of the dredge procedure are provided in the Supplementary Material.

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## Figure Captions

Figure 1. (a-b) Bathymetric maps. (a) Wide-area map showing the distribution of the Izu-Bonin arc. Bathymetric data are from ETOPO1 (doi:10.7289/V5C8276M). (b) Close-up map around FOB that shows the locations of the dredge sites. Bathymetric data were collected during the YK22-15 cruise. The green circle indicates the two-nautical-mile area of prohibited entry due to safety reasons. (c) Sandy seafloor covered with volcanic ash, which was observed during dredge D08 on the western slope of FOB. (d) Outcrop of volcanic rock observed during dredge D07 on the northeastern outer slope of Kita-Fukutoku Caldera.

Figure 2. Photographs of representative hand specimens collected during the cruise. (a) Well-vesiculated pumice, (b) black-colored lapilli, and (c) black-colored pumice collected during dredges D03 and D08 on the western slope of FOB. (d) Lapilli clasts collected during dredge D05 on the northwest of FOB. (e) Weakly vesiculated trachyte clast collected from the southeastern slope of FOB. The white areas are mainly plagioclase. (f) Volcanic rock and (g) well-developed woody pumice collected during dredge D07 on the northeastern outer slope of Kita-Fukutoku caldera.

Figure 3. (a) Total alkalis versus silica diagram showing the compositions of the dredged samples and literature data. (b-d) Plots of major elements versus SiO<sub>2</sub>, which exhibit linear relationships. (e-f) Plots of selected trace element ratios versus SiO<sub>2</sub>. Data for pumice from previous eruptions and melt inclusion in the 2021 drift pumice are also shown. (g) Trace element patterns of dredged samples and drift pumice from the 2021 and 1986 eruptions. Compositions were normalized to N-MORB (Sun & McDonough, 1989). Abbreviations: Y22a, Yoshida et al. (2022a); Y22b, Yoshida et al. (2022b); M22, Maeno et al. (2022); K88, Kato (1988); S98, Sun et al. (1998); Y87, Yoshida et al. (1987).

Figure 4. (a) Photomicrograph of the microlite-free lapilli clast D08R04. The groundmass consists of weakly vesiculated pale brown glass. (b) Microlite-rich lava D07R01 containing magnetite nanolites in the groundmass glass, which was confirmed by Raman spectroscopy. The interstitial glass is generally microlite-free. (c) Cross-polarized light (XPL) and plane-polarized light (PPL) photomicrographs of the pumice clast D03R03 collected from the western slope of FOB. The glass adjacent to the plagioclase phenocryst is colorless and transparent. (d) Photomicrograph of a pumice clast that drifted from FOB to the Gulf of Thailand. The groundmass surrounding the plagioclase phenocryst has a brown color, which is due to magnetite nanolite crystallization. A

detailed petrographic description was provided by Yoshida et al. (2022b). (e) Photomicrograph of the weakly vesiculated pumice clast D08R11 that contains phenocrysts of olivine, plagioclase, and magnetite. Spherical cristobalite also occurs in the vesicles. (f) Representative Raman spectra of the colored glass. Although the groundmass glass in D06R02 and D08R04 is pale brown, it does not exhibit a magnetite nanolite peak at 670 cm<sup>-1</sup>. The brown glass in D07R01 exhibits a nanolite peak in its Raman spectrum.

Figure 5. (a) Backscattered electron (BSE) image of the plagioclase phenocryst in the microlite-rich volcanic rock clast D07R01. The temperatures calculated based on the X<sub>Mg-mag</sub> values of magnetite crystals included in the plagioclase rim are also shown. (b) Zoned clinopyroxene in the microlite-rich volcanic rock clast D07R05. The Mg# value of clinopyroxene in the core and rim, and temperatures calculated from magnetite are also shown. (c) Zoned olivine observed in sample D07R01. Representative Mg# values of the core and rim are also shown. (d) Raman spectrum of cristobalite in sample D08R11. A reference spectrum taken from the RRUFF database (Lafuente et al., 2016) is shown for comparison. (e) BSE image of the microlite-rich pumice clast D08R11, showing a mineral aggregate of plagioclase, clinopyroxene, and magnetite. Temperatures calculated for magnetite are also shown. (f) Cathodoluminescence image of the cristobalite in the boxed area of (e), which has a radial texture.

Figure 6. (a-b) Ternary compositional diagram for (a) plagioclase and (b) clinopyroxene in the studied samples. (c) X<sub>Mg-mag</sub> values of magnetite in the studied samples and the corresponding temperatures calculated following the method of Canil and Lacourse (2020). (d-f) Glass compositions of the studied samples shown in plots of (d) SiO<sub>2</sub>-total alkalis, (e) SiO<sub>2</sub>-CaO, and (f) halogen contents.

Figure 7. (a) Photographs of the sea surface above the volcanic vent of FOB taken on 22 August 2022. Floating bubbles were observed. Under the sea surface, the wall of the volcanic vent is visible. (b) Schematic diagram of the FOB magmatic system inferred from results of this study. See the main text for details.

**Tables** 602 603 Table 1. Whole-rock geochemical compositions of the dredged samples, showing the trace element compositions of selected samples. 604 605 Table 2. Representative compositions of plagioclase. 606 Footnote: \*Total iron as FeO. 607 608 609 Table 3. Representative compositions of clinopyroxene and olivine. 610 Footnote: \*Total iron as FeO. †Pressures for clinopyroxene were calculated using the method of on 611 Petrelli et al. (2020). 612 613 Table 4. Representative compositions of magnetite and silica mineral. 614 Footnote: \*Total iron as FeO.  $\dagger X_{\text{Mg-mag}}$  calculated based on Canil and Lacourse (2020) and 615  $\ddagger$ Temperature calculated using  $X_{\text{Mg-mag}}$ .

# Fig. 1

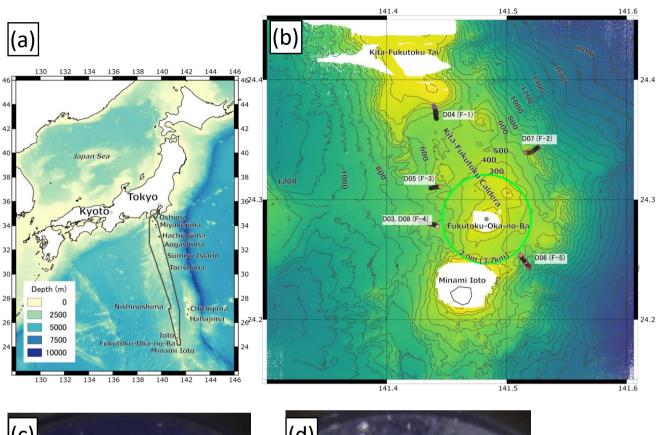






Fig. 2

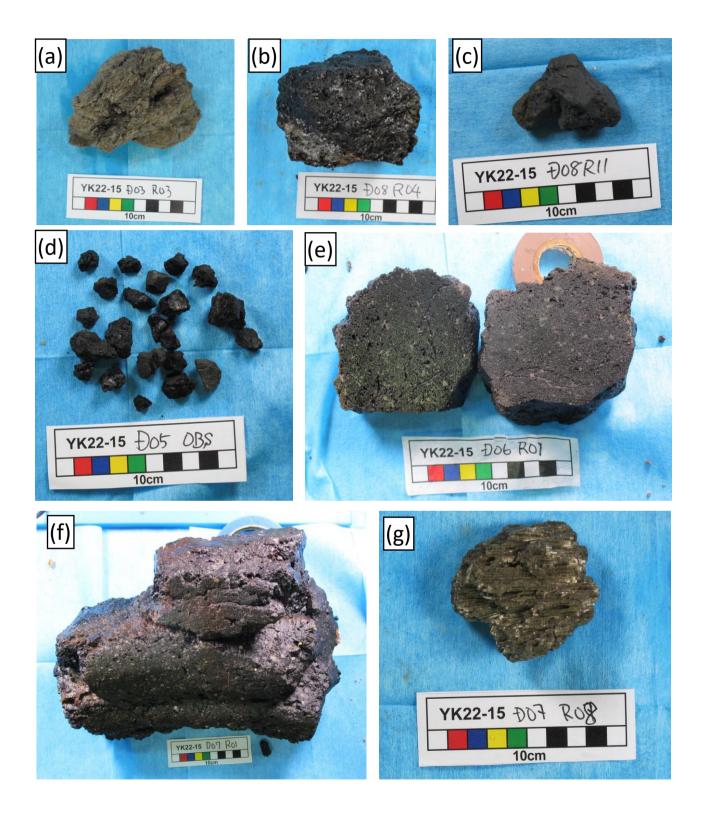


Fig. 3

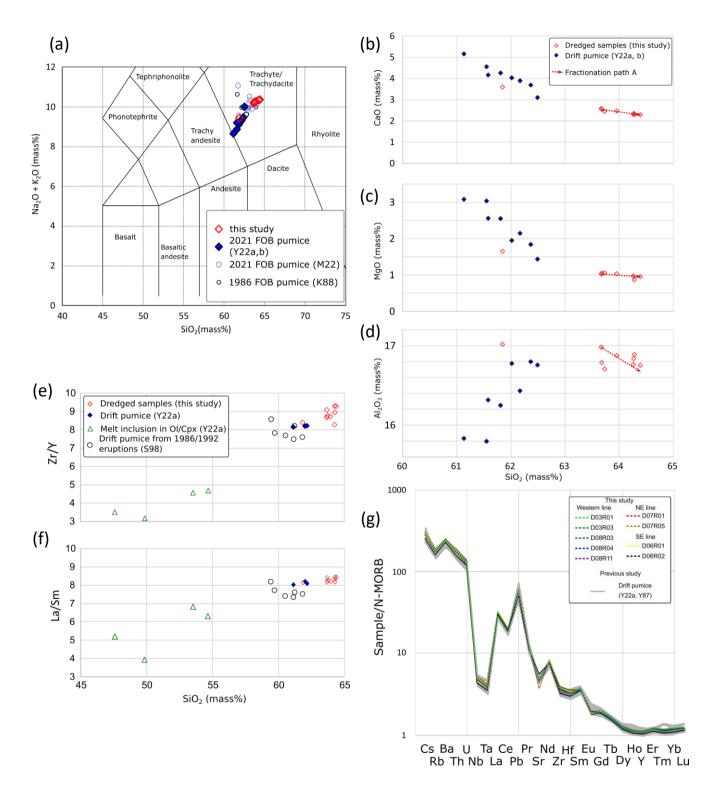


Fig. 4

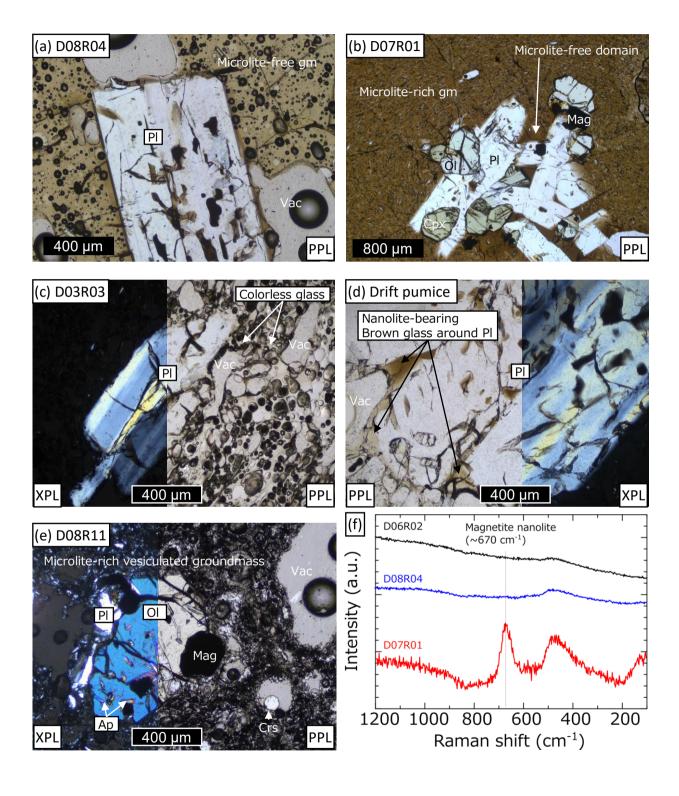


Fig. 5

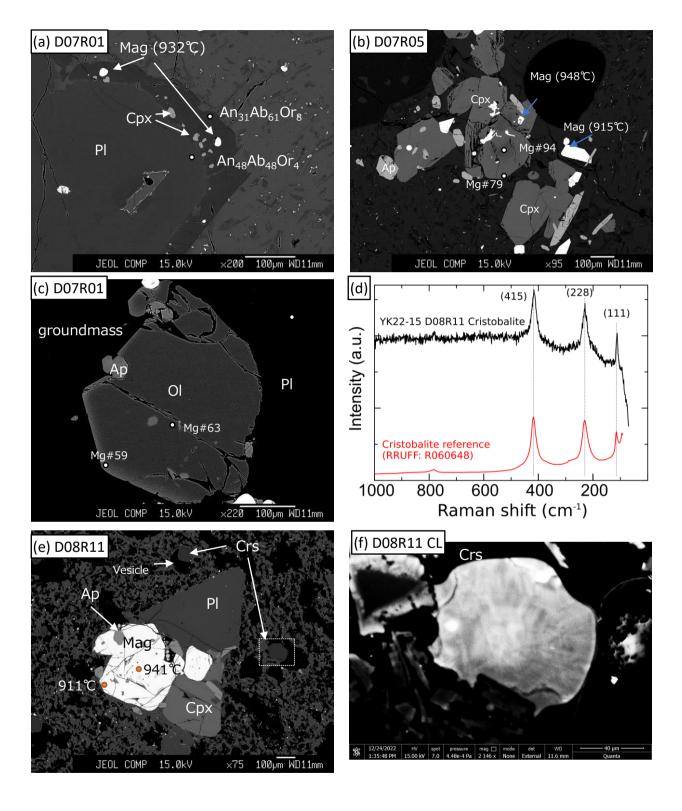


Fig. 6

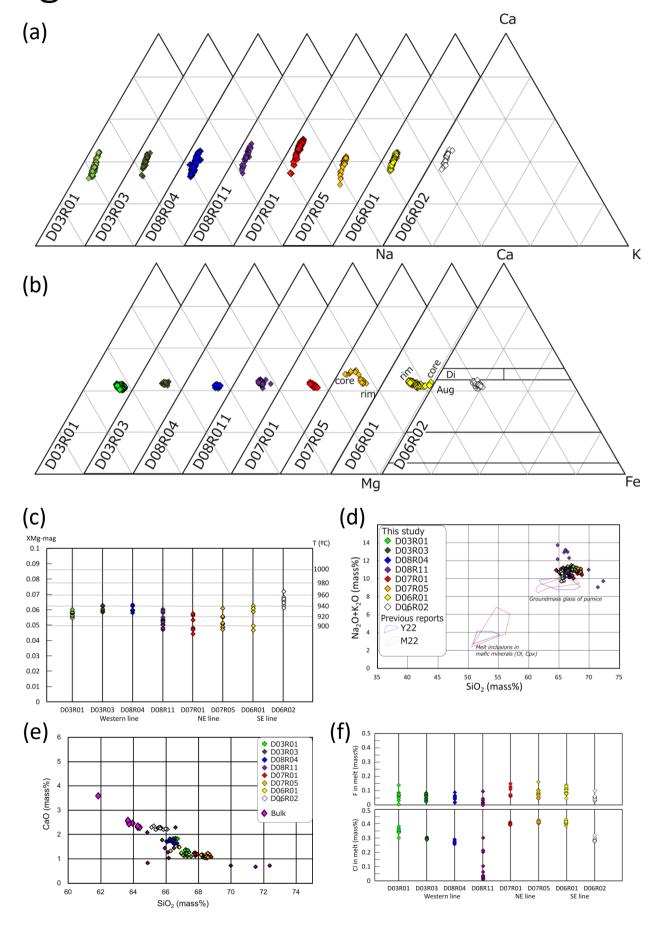
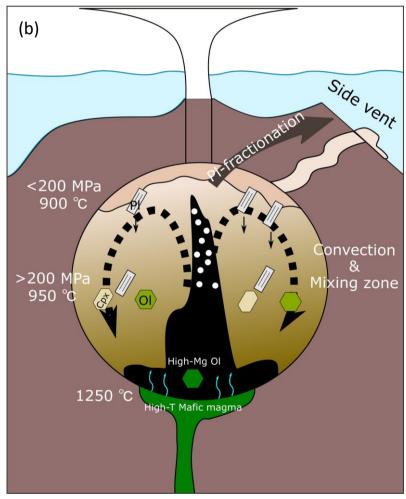


Fig. 7





	Western I	ine						
Sample No.	D03R01	D03R02	D03R03	D03R04	D04R01	D08R03	D08R04	D08R11
type	lapilli	pumice	pumice	pumice	pumice	lapilli	lapilli	pumice
microlite	rich	-	free	-	-	-	free	rich
in mass%	)							
SiO <sub>2</sub>	63.06	61.33	62.74	61.04	60.43	62.52	62.47	63.02
$TiO_2$	0.57	0.59	0.58	0.59	0.65	0.59	0.58	0.57
$Al_2O_3$	16.57	16.43	16.56	16.17	16.74	16.39	16.66	16.51
Fe <sub>2</sub> O <sub>3</sub>	4.68	5.30	4.74	5.46	6.03	5.17	4.91	4.60
MnO	0.17	0.17	0.17	0.17	0.19	0.17	0.17	0.17
MgO	0.85	1.50	1.02	1.86	1.42	1.04	1.01	0.94
CaO	2.26	3.15	2.42	3.59	3.36	2.40	2.54	2.31
Na <sub>2</sub> O	5.04	4.94	5.06	4.76	4.82	4.91	4.95	5.05
$K_2O$	5.11	4.65	5.02	4.63	4.56	5.14	5.04	5.09
$P_2O_5$	0.22	0.24	0.23	0.23	0.28	0.24	0.24	0.22
Total	98.52	98.30	98.53	98.48	98.48	98.57	98.57	98.47
LOI	0.82	1.25	1.60	1.47	1.87	0.44	0.47	1.75
in ppm								
Sc	5.765		4.607			3.48	5.078	2.703
Co	5.105		5.44			5.911	6.039	5.104
Ni	0.604		0.85			0.69		0.532
Cu	15.668		13.926			17.532		12.898
Rb	102.895		97.786			101.468		97.351
Sr	381.091		401.854			385.298		380.95
Υ	31.473		30.462			31.749		30.792
Zr	282.1		266.22			278.422		255.013
Nb	11.603		11.139			11.44		11.041
Cs	2.044		1.963			2.018		1.944
Ва	1628.653		1557.937				1571.212	
La	80.889		77.939			81.187		78.154
Ce	151.691 16.309		145.449 15.722			151.712 16.416		146.606 15.904
Pr Nd	58.724		56.303			59.309		57.213
Nd Sm	9.68		9.476			9.809		9.545
Eu	1.953		1.909			1.982		9.545 1.961
∟u	1.900	•	1.509			1.302	1.331	1.501

Gd	7.058	6.818	7.292	7.028	6.928
Tb	1.086	1.047	1.098	1.061	1.061
Dy	5.753	5.632	5.912	5.698	5.698
Но	1.165	1.122	1.179	1.155	1.153
Er	3.585	3.522	3.654	3.602	3.601
Tm	0.539	0.513	0.541	0.517	0.525
Yb	3.684	3.554	3.724	3.594	3.586
Lu	0.568	0.549	0.575	0.557	0.554
Hf	6.868	6.538	6.959	6.678	6.531
Ta	0.525	0.512	0.533	0.515	0.513
TI	0.237	0.262	0.232	0.22	0.079
Pb	17.641	17.386	17.578	17.279	12.601
Th	21.248	20.221	21.474	20.853	20.767
U	6.108	5.829	6.219	5.97	6.169

NE line			SE line	
D07R01	D07R05	D07R09	D06R01	D06R02
lava	lava	woody pumice	lapilli	lapilli
rich	rich	-	rich	free
63.13	63.27	62.33	62.58	60.70
0.55	0.55	0.55	0.59	0.60
16.47	16.46	16.74	16.50	16.70
4.77	4.67	4.67	5.07	5.85
0.16	0.16	0.16	0.16	0.17
0.95	0.94	1.04	1.04	1.62
2.28	2.26	2.68	2.51	3.53
4.84	4.83	4.90	4.92	4.68
5.31	5.35	5.05	5.13	4.58
0.20	0.20	0.21	0.24	0.26
98.67	98.68	98.34	98.73	98.69
1.03	1.02	2.70	0.63	0.69
3.765	3.171		5.942	5.946
5.951	5.678		6.242	9.864
1.315	1.284		0.745	4.824
22.445	19.442		20.727	34.807
107.589	107.074		103.452	89.679
335.262	340.913		393.279	500.262
31.204	31.177		31.28	29.066
290.57	290.074		284.739	244.3
12.01	12.086		11.762	10.175
2.121	2.108		2.039	1.788
1557.2	1558.452		1584.703	1451.749
81.591	81.026		82.04	75.035
152.237	153.946		154.28	139.477
16.384	16.261		16.438	15.138
58.124	58.206		59.568	55.082
9.644	9.598		9.741	9.201
1.839	1.836		1.947	1.963

6.924	6.942	7.104	6.716
1.067	1.065	1.065	1.017
5.753	5.676	5.743	5.456
1.151	1.151	1.16	1.076
3.641	3.673	3.64	3.37
0.531	0.538	0.531	0.488
3.661	3.704	3.618	3.352
0.576	0.574	0.569	0.523
7.19	7.238	6.956	6.081
0.559	0.552	0.542	0.467
0.279	0.287	0.22	0.212
18.213	18.305	17.404	15.525
22.493	22.578	21.507	18.934
6.494	6.522	6.284	5.538

Sample N	N-D03R01		D03R03		D08R04		D08R11	
Note	core	rim	core	rim	core	rim	core	rim
SiO <sub>2</sub>	56.75	59.50	58.32	59.68	57.83	59.62	56.30	60.19
TiO <sub>2</sub>	0.01	0.07	0.07	0.06	0.00	0.02	0.06	0.00
$Al_2O_3$	26.37	24.83	26.23	24.52	25.58	24.54	26.79	24.25
$Cr_2O_3$	0.00	0.13	0.00	0.01	0.01	0.00	0.04	0.00
FeO*	0.56	0.61	0.49	0.55	0.48	0.42	0.55	0.54
MnO	0.00	0.00	0.02	0.00	0.02	0.00	0.01	0.08
MgO	0.05	0.05	0.06	0.03	0.03	0.07	0.02	0.02
CaO	8.96	7.23	8.82	7.01	8.58	6.89	10.03	7.08
$Na_2O$	5.82	6.72	6.17	6.97	6.14	6.78	5.59	6.74
$K_2O$	0.69	0.97	0.79	1.11	0.75	1.00	0.59	1.17
total								
Ο	8	8	8	8	8	8	8	8
Si	2.57	2.66	2.60	2.68	2.61	2.68	2.54	2.70
Ti	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Al	1.41	1.31	1.38	1.30	1.36	1.30	1.43	1.28
Cr	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Fe	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02
Mn	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Mg	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Ca	0.44	0.35	0.42	0.34	0.42	0.33	0.49	0.34
Na	0.51	0.58	0.53	0.61	0.54	0.59	0.49	0.59
K	0.04	0.06	0.04	0.06	0.04	0.06	0.03	0.07
An	44	35	42	33	42	34	48	34
Ab	52	59	53	60	54	60	49	59
Or	4	6	4	6	4	6	3	7

D07R01		D07R05		D06R01		D06R02	
core	rim	core	rim	core	rim	core	rim
56.48	60.24	58.32	60.40	57.86	57.99	56.52	58.64
0.05	0.08	0.04	0.04	0.00	0.07	0.06	0.04
26.72	24.32	25.45	24.48	26.40	25.11	25.85	24.31
0.05	0.00	0.00	0.00	0.00	0.00	0.03	0.04
0.58	0.60	0.48	0.45	0.48	0.48	0.70	0.62
0.00	0.05	0.04	0.00	0.01	0.09	0.05	0.00
0.04	0.06	0.06	0.01	0.08	0.04	0.02	0.04
9.88	6.33	8.20	6.64	9.09	7.92	9.47	7.52
5.47	6.88	6.35	7.11	5.95	6.56	6.06	6.69
0.53	1.25	0.88	1.35	0.81	0.92	0.78	1.02
8	8	8	8	8	8	8	8
2.55	2.70	2.63	2.69	2.59	2.63	2.56	2.66
0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
1.42	1.28	1.35	1.29	1.39	1.34	1.38	1.30
0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
0.02	0.02	0.02	0.02	0.02	0.02	0.03	0.02
0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
0.00	0.00	0.00	0.00	0.01	0.00	0.00	0.00
0.48	0.30	0.40	0.32	0.44	0.38	0.46	0.37
0.48	0.60	0.55	0.61	0.52	0.58	0.53	0.59
0.03	0.07	0.05	0.08	0.05	0.05	0.05	0.06
48	31	40	31	44	38	44	36
48	61	55	61	52	57	51	58
3	7	5	8	5	5	4	6

Sample No.	D03R01	D03R03	D08R04	D08R11	D07R01	D07R05	
Note						core	rim
SiO <sub>2</sub>	52.66	52.58	52.65	52.65	52.42	53.39	53.00
TiO <sub>2</sub>	0.38	0.39	0.36	0.34	0.40	0.19	0.38
$Al_2O_3$	0.64	1.48	0.67	1.82	0.85	1.68	1.22
$Cr_2O_3$	0.02	0.10	0.00	0.00	0.05	0.09	0.07
FeO*	9.43	9.09	9.61	8.79	9.10	4.13	9.62
MnO	0.19	0.99	0.11	0.68	0.13	0.12	0.79
MgO	15.36	15.35	15.21	15.44	15.45	17.50	15.26
CaO	19.88	20.22	20.16	20.53	20.19	23.32	20.59
Na <sub>2</sub> O	0.45	0.38	0.48	0.38	0.41	0.22	0.45
K <sub>2</sub> O	0.01	0.01	0.00	0.00	0.00	0.00	0.00
total	99.03	100.59	99.23	100.63	99.00	100.63	101.37
	0	6	0		6	0	6
0	6	6	1.07	6			6
Si	1.97	1.94	1.97				1.94
Ti	0.01	0.01	0.01	0.01			0.01
Al	0.03		0.03				0.05
Cr	0.00	0.00	0.00	0.00			0.00
Fe <sup>3+</sup>	0.04		0.05	0.06			0.07
Fe <sup>2+</sup>	0.25	0.22	0.25	0.21	0.24	0.06	0.22
Mn	0.01	0.03	0.00	0.02	0.00	0.00	0.02
Mg	0.86	0.84	0.85	0.85	0.86	0.94	0.83
Ca	0.80	0.80	0.81	0.81	0.81	0.90	0.81
Na	0.03	0.03	0.03	0.03	0.03	0.02	0.03
K	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Mg#	77	79	77	80	78	94	79
P (MPa, P20)†	388	261	254	236	177	445	276

D06R01		D06R02		D03R01	D03R03	D08R04	D07R01
core	rim						core
53.42	52.62	52.85	SiO <sub>2</sub>	37.41	37.12	37.64	36.44
0.18	0.41	0.39	$TiO_2$	0.05	0.02	0.00	0.02
0.93	1.69	1.33	$Al_2O_3$	0.00	0.01	0.02	0.00
0.02	0.00	0.02	$\mathrm{Cr_2O_3}$	0.00	0.01	0.00	0.00
11.45	8.87	9.78	FeO*	31.03	30.51	30.31	32.50
1.07	0.65	0.69	MnO	1.57	1.54	1.52	1.68
14.19	15.05	15.11	MgO	31.64	32.53	32.09	30.49
19.88	3 20.58	20.12	CaO	0.15	0.17	0.18	0.15
0.46	0.43	0.38	Na <sub>2</sub> O	0.03	0.00	0.01	0.01
0.00	0.03	0.02	$K_2O$	0.00	0.00	0.01	0.01
101.59	100.33	100.69	NiO	0	0	0	0
			total	101.88	101.91	101.77	101.30
6	6	6	Ο	4	4	4	4
1.97	1.94	1.95	Si	1.00	0.99	1.00	0.99
0.00	0.01	0.01	Ti	0.00	0.00	0.00	0.00
0.04	0.07	0.06	Al	0.00	0.00	0.00	0.00
0.00	0.00	0.00	Cr	0.00	0.00	0.00	0.00
0.04	0.05	0.05	Fe	0.69	0.68	0.68	0.74
0.31	0.23	0.26	Mn	0.04	0.03	0.03	0.04
0.03	0.02	0.02	Mg	1.26	1.29	1.28	1.24
0.78	0.83	0.83	Ca	0.00	0.00	0.01	0.00
0.79	0.81	0.80	Na	0.00	0.00	0.00	0.00
0.03	0.03	0.03	K	0.00	0.00	0.00	0.00
0.00	0.00	0.00					
72	2 78	76		64	66	65	63
371	279	215					

	D07505		D00501	D00500
	R07R05		D06R01	D06R02
rim	core	rim		
36.36	36.51	37.15	37.10	37.38
0.00	0.01	0.01	0.00	0.00
0.02	0.01	0.69	0.01	0.02
0.00	0.00	0.02	0.00	0.01
34.63	32.52	35.46	31.03	29.18
1.72	1.65	1.79	1.41	1.23
28.87	30.20	26.52	31.94	33.40
0.15	0.15	0.19	0.16	0.17
0.01	0.04	0.18	0.00	0.01
0.02	0.00	0.16	0.02	0.00
0.001	0	0	0	0.002
101.77	101.09	102.17	101.67	101.39
4	4	4	4	4
0.99	0.99	1.01	0.99	1.00
0.00	0.00	0.00	0.00	0.00
0.00	0.00	0.02	0.00	0.00
0.00	0.00	0.00	0.00	0.00
0.79	0.74	0.81	0.70	0.65
0.04	0.04	0.04	0.03	0.03
1.18	1.23	1.08	1.28	1.33
0.00	0.00	0.01	0.00	0.00
0.00	0.00	0.01	0.00	0.00
0.00	0.00	0.01	0.00	0.00
60	62	57	65	67

Sample No.	D03R01	D03R03	D08R04	D08R11		D07R01	
Note				in Cpx ancgm		in Pl rim	gm
SiO <sub>2</sub>	0.10	0.09	0.08	0.09	0.10	0.16	0.13
TiO <sub>2</sub>	10.93	10.53	10.38	11.26	10.95	11.27	12.26
$Al_2O_3$	2.81	2.83	2.95	2.61	2.70	2.59	2.07
$Cr_2O_3$	0.06	0.00	0.02	0.12	0.00	0.09	0.13
FeO*	78.09	77.56	78.01	77.77	78.35	75.66	75.54
MnO	1.05	0.96	1.02	0.94	1.12	0.17	1.10
MgO	2.81	2.70	2.91	2.68	2.17	2.58	2.08
CaO	0.03	0.00	0.03	0.03	0.01	0.11	0.03
Na <sub>2</sub> O	0.01	0.00	0.03	0.00	0.05	0.00	0.00
K <sub>2</sub> O	0.04	0.01	0.00	0.02	0.00	0.01	0.10
total	95.92	94.68	95.44	95.52	95.45	92.64	93.44
Ο	4	4	4	4	4	4	4
Si	0.00	0.00	0.00	0.00	0.00	0.01	0.00
Ti	0.29	0.28	0.28	0.30	0.29	0.31	0.34
Al	0.12	0.12	0.12	0.11	0.11	0.11	0.09
Cr	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Fe <sup>3+</sup>	1.49	1.50	1.50	1.48	1.49	1.46	1.45
Fe <sup>2+</sup>	0.82	0.83	0.82	0.83	0.85	0.85	0.85
Mn	0.03	0.03	0.03	0.03	0.03	0.01	0.03
Mg	0.15	0.14	0.15	0.14	0.12	0.14	0.11
Ca	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Na	0.00	0.00	0.00	0.00	0.00	0.00	0.00
K	0.00	0.00	0.00	0.00	0.00	0.00	0.00
XMg-mag†	0.060	0.058	0.062	0.058	0.047	0.057	0.047
T_XMg (° C) ‡	934	929	940	928	893	926	892

D07R05	D06R01		D06R02	D08R11
2011103	core	rim	2001102	Crs
0.09		0.07	0.10	98.21
11.90		10.22	9.33	0.143
2.59		2.87	3.19	0.973
0.06		0.06	0.05	0.02
77.23		78.43	77.51	0.066
1.14		0.94	0.84	0.037
2.32		2.31	3.12	0.006
0.01	0.00	0.00	0.00	0.014
0.00		0.05	0.00	0.597
0.01	0.02	0.04	0.00	0.005
95.34		94.99	94.14	
4	4	4	4	2
0.00	0.00	0.00	0.00	0.99
0.32	0.28	0.28	0.25	0.00
0.11	0.12	0.12	0.13	0.0115
0.00	0.00	0.00	0.00	0.00
1.46	1.50	1.50	1.52	0.00
0.84	0.82	0.85	0.81	0.00
0.03	0.03	0.03	0.03	0.00
0.12		0.12	0.17	0.00
0.00	0.00	0.00	0.00	0.00
0.00		0.00	0.00	0.0116
0.00		0.00	0.00	0.00
0.051	0.063	0.050	0.067	
905	941	902	953	

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Supplementary Material 1

Dredge assembly used during the cruise YK22-15

During the cruise YK22-15, we used a Satsuki type dredger and depressor-towing body. This assembly is employed to protect the opto-electrical composite cable from any damaging during dredging (Fig. S1). The Satsuki type dredger was connected to the depressor-towing body by a lead wire with 200 m length. The position of the dredger was monitored by a transponder attached to the depressor-towing body (Fig. S2a). The Satsuki type dredger has a size of 800 mm  $\times$  740 mm  $\times$  120 mm (width, height, and length, respectively) and  $\sim$ 240 kg.

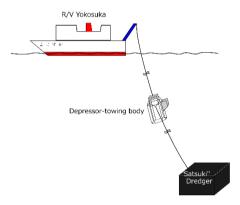


Fig. S1. Schematic image of the dredge assembly.

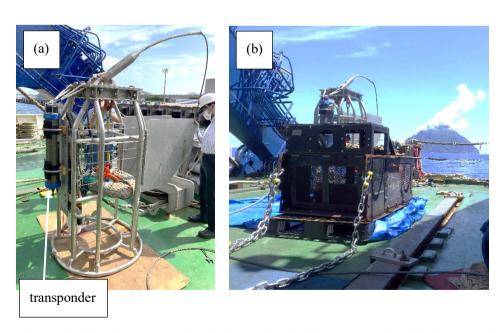


Figure S2. (a) A depressor-towing body equipped with a transponder. (b) Photograph of Satsuki type dredger which is ready for operation.

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Table S1. Details of the dredge positions

	Hit bottom		Off bottom		Depth (m)*1		Tension	Survey
Dredge number							max.	time
							(kN)	(hh:mm)
	Lat.	Lon.	Lat.	Lon.	On the	Off the		
	$(SOQ^{*1})$	$(SOQ^{*1})$	$(SOQ^{*1})$	$(SOQ^{*1})$	bottom	Bottom		
D02	24-	141-	24-	141-	446	252	41.6	0:36
D03	16.8737N	25.6601E	16.8458N	25.8852E	446	353		
D.0.1	24-	141-	24-	141-	324   299	200	13.0	0:59
D04	22.0755N	26.4721E	22.5723N	26.4072E		299		
505	24-	141-	24-	141-	2.50	100	18.1	1:00
D05	18.5790N	26.2145E	18.6416N	26.6848E	369	182		
- 0 -	24-	141-	24-	141-		649 361	16.6	1:12
D06	14.6284N	31.1048E	15.1767N	30.6928E	649			
	24-	141-	24-	141-		256	28.1	1:17
D07	20.6611N	31.6256E	20.3546N	31.0876E	525			
D08	24-	141-	24-	141-	202	294	17.3	1:02
	16.8088N	26.0734E	16.6746N	26.5212E	383			

<sup>\*</sup>SOQ = SGK Transponder's position

<sup>\*</sup>SOJ = Ship's position

Table S2. Trace element analysis of secondary standard (JB-2) and the reference value Sample JB-2, mea: JB-2, reference\* ( $\mu$ g/g)

Sample	JD-Z, mea: Ji	D-Z, reference	$(\mu g)$
Sc	51.728	54.08	
Co	34.64	37.57	
Ni	12.417	14.77	
Cu	209.443	222.1	
Rb	5.732	6.4	
Sr	175.628	178.2	
Υ	21.115	23.56	
Zr	43.894	48.25	
Nb	0.439	0.565	
Cs	0.754	0.8	
Ва	208.065	218.1	
La	2.131	2.281	
Ce	6.297	6.552	
Pr	1.096	1.129	
Nd	6.168	6.392	
Sm	2.208	2.266	
Eu	0.792	0.836	
Gd	3.15	3.123	
Tb	0.572	0.5863	
Dy	3.915	3.868	
Но	0.854	0.863	
Er	2.584	2.537	
Tm	0.376	0.393	
Yb	2.525	2.529	
Lu	0.382	0.3894	
Hf	1.49	1.487	
Та	0.032	0.0396	
TI	0.036	0.034	
Pb	4.963	5.25	
Th	0.252	0.2576	
U	0.155	0.1528	

<sup>\*</sup>Reference of JB-2 is taken from GeoReM database (Jochum et al. Geostand. Geoanal. Res. 40, 333-350, 2016)