Article

Origin of Pumice in Sediments from the Middle Okinawa Trough: Constraints from Whole-Rock Geochemical Compositions and Sr-Nd-Pb Isotopes

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Abstract: Frequent volcanic activity has occurred in the Okinawa Trough (OT) during the late Quaternary, which attracted much attention to the origin of volcanic rocks. Pumice collected from the seafloor has been extensively investigated, whereas few studies paid attention to the pumice in the sediment. The geochemical compositions of pumice preserved in sediments generally provide insight into past volcanic activity and regional magmatism. Here, we present major and trace element compositions and Sr-Nd-Pb isotope data, together with the established age framework for pumice samples recovered from sediment core S9 in the middle OT (MOT) to investigate their possible formation. Compositionally, the S9 pumice samples are dacite and are characterized by relatively higher Sr ($^{87}\text{Sr}/^{86}\text{Sr} = 0.70480–0.70502$) and Pb ($^{206}\text{Pb}/^{204}\text{Pb} = 18.321–18.436$, $^{207}\text{Pb}/^{204}\text{Pb} = 15.622–15.624$, and $^{208}\text{Pb}/^{204}\text{Pb} = 38.52–38.63$) and lower Nd ($^{143}\text{Nd}/^{144}\text{Nd} = 0.51272–0.51274$) isotope compositions than basalts from the MOT. The geochemical compositions of pumice clasts from different layers of core S9 display no temporal variation trends and vary within narrow ranges. On the basis of the geochemical characteristics of S9 pumice samples, we infer that the parent magma of these samples might generate from hybrid magma through an extensive fractional crystallization process. The Indian Ocean MORB-type mantle was first metasomatized by the subducted Philippine Sea sediments to form the primitive magma; then, followed by assimilation of a small amount of lower crustal component occurred in the lower crust. The long-term magmatism and relatively consistent isotopic compositions indicate that a magma chamber might have existed in the lower crust of the MOT between 11.22 and 12.96 cal. ka BP.

Keywords: pumice in sediments; Sr-Nd-Pb isotope; hybrid magma; fractionation crystallization; origin of pumice; middle Okinawa Trough

1. Introduction

The subduction zone, as a crucial interface of material exchange between the Earth’s crust and mantle, is one of the critical areas for plate subduction and consumption as well as for the generation of magmatism [1–3]. The back-arc basin is an essential part of trench-arc-basin subduction system. Achievements of the Ocean Drilling Program (ODP) have revealed that back-arc basins...
play an important role in the study of geodynamics and provide valuable information about the origin of mantle-derived magma, interactions between the crust and mantle, crustal accretion and oceanic-continental transitions [4–6]. In recent decades, the studies of back-arc basin magmatism have gained increased attention [7–9].

The Okinawa Trough (OT) located behind the Ryukyu arc-trench system is an active marginal sea basin that is still in the early stage of expansion [10–12]. Extensive volcanic activity has given rise to numerous volcanic rocks, making the OT an ideal area for studying the magmatism associated with back-arc basin initiation [13,14]. Volcanic rocks can provide information about magma evolution and geodynamic mechanisms in the OT [15]. Pumice is the most widely distributed volcanic rock in the OT [16,17]. Zhai [18] reported that pumice was mainly distributed in the central and northern part of the OT, which can be observed both on the seafloor and in sediment interlayers. The geochemical and mineralogical characteristics of pumice were used to investigate the source magma compositions [19–21], magmatic evolution [14], and implications for magma chambers [22] in the OT. The origin of pumice in the OT; however, remains to be clarified despite much research effort [19,23–25]. Most of the studies suggest that pumice were derived from basaltic magma through fractional crystallization [26,27], while a few studies suggest the partial remelting of andesites [23]. The divergence of these interpretations can be ascribed to the lack of an age framework and the poor recognition of pumice types.

The OT is characterized by complex magmatic and tectonic activity, and has different geological backgrounds in three sections divided by the Tokara Strait (~29.3°N) and the Kerama Gap (~26°N) [12,28,29]. Due to the difficulties in dating young geological material [7,10,25,30,31], a majority of studies fail to establish the age model when analyzing the magma sources and evolutions of volcanic rocks in the OT [19,20,23]. Because the magmatic components and tectonic conditions in the study area have likely changed in this rapidly developing region during different eruption times, the lack of age framework may result in problematic interpretation of the results. The volcanic material (e.g., pumice clasts) also contributed significantly to the sedimentation in the OT since the late Quaternary, which are well documented by previous studies [18,30,32]. Pumice in sediments can provide pumice samples beneath the seafloor for geochemical study, and their chronological framework can be established using radiocarbon dating. Nonetheless, previous studies were mainly focused on pumice that recovered from the surface of the seafloor with a television (TV) grab sampler or trawl sampler [19,20,23], and few studies have focused on pumice recovered from marine sediments. We collected a sediment core from a slope setting in the middle OT (MOT) to investigate the origin of the pumice in sediment of the study area for the first time.

Here, we present the geochemical data and eruption age of pumice clasts in sediment core S9 from the MOT. The objectives of this study are (1) to characterize the geochemical compositions of the pumice, (2) to identify the potential magma sources of the pumice, and (3) to investigate the possible petrogenesis of the pumice in the sediment from the MOT.

2. Geological Setting and Samples

The OT is a newborn back-arc basin that is located in the eastern part of the East China Sea (ECS) and formed between the ECS shelf and the Ryukyu volcanic arc, which is related to the subduction of the Philippine Sea Plate beneath the Eurasian Plate (Figure 1) [33–35]. The total length of the OT is approximately 1300 km; it starts from the Kyushu Island in the north and ends at Taiwan Province in the south [8,12]. The OT is characterized by a complex geological background with obviously thin crust compared with the ECS shelf and the Ryukyu arc and anomalously high heat flow and gravity values [11,34,36]. Generally, the OT can be divided into three segments by the Tokara Strait and the Kerama Gap: the northern OT (NOT), the MOT, and the southern OT (SOT) [37]. Magmatism in the OT is bimodal, showing dominantly basaltic and rhyolitic rocks with small amounts of intermediate volcanic rocks [11,38]. Pumice is widely distributed in the three segments of the OT, whereas basalts mainly occur in the middle and southern segments [26]. Although volcanic rocks are widespread in the OT, few studies have reported on the eruption age of volcanic rocks collected from the OT due to
difficulty in dating young volcanic rocks [7,17]. According to previous studies, the earliest eruption age of volcanic rocks is approximately 4 Ma [31], and most of the volcanic rocks collected from the OT are younger than 1 Ma [25,30]. Additionally, the OT is located in a semi-abyssal area and represents a transitional environment between the shelf sedimentary environment and the abyssal sedimentary environment [10]. The OT is covered by large quantities of semi-abyssal sediments [33,39], especially in the MOT with 2–3 km-thick sediment layers [40]. These sediments are mainly composed of terrigenous sediments from around large rivers, as well as abundant biodetritus and volcanic material [32], which preserves valuable information about the magmatism and make the OT an ideal area for investigating back-arc magmatic activity.

**Figure 1.** Simplified tectonic map of the Okinawa Trough (OT), showing sampling locations of the studied pumice and volcanic rocks used in this study. Yellow star represents the sampling location of core S9. Blue inverted triangles and green circles represent andesites [26] and basalts [16,17,24,27] from the middle OT (MOT), respectively. Red triangle represents a Quaternary volcano named Kuro-shima [37]. Two black dashed lines represent the Tokara Fault and Kerama Gap [12]. Solid arrows indicate plate motion vectors of the Philippine Sea Plate relative to the Eurasian Plate [35].

All pumice reported in this study was recovered from piston core S9, which was collected from the MOT at a water depth of 1062 m (Figure 1) with a piston corer. The total length of core S9 was 251 cm. Volcanic materials in S9 were mainly composed of pumice clasts that were found between 226 and 237 cm with sizes of 0.5–15 mm and a grayish white color (Figure 2). All of the pumice samples were
glassy, fresh, and slightly porphyritic. The main phenocryst minerals are orthopyroxene, amphibole, and Fe-Ti oxides. We sampled the pumice clasts at two-centimeter intervals and numbered them F1-6 for geochemical analysis as shown in Table 1.

![Figure 2](image_url). (a) Close-up image of core S9 containing centimeter-thick pumice clasts layers. (b) Hand specimen photographs of pumice recovered from core S9. (c-f) Representative scanning electron microscope (SEM) images of S9 pumice. The eruption ages of S9 pumice samples were calculated with Bayesian analysis.

| Table 1. Major and trace elements and Sr-Nd-Pb isotopic compositions of S9 pumice samples. |
|---------------------------------|----------------|----------------|----------------|----------------|----------------|----------------|
| Sample | F1 | F2 | F3 | F4 | F5 | F6 |
| Major elements (wt.%) | | | | | | |
| SiO₂ | 68.57 | 68.1 | 67.86 | 68.88 | 69.37 | 69.18 |
| TiO₂ | 0.63 | 0.69 | 0.68 | 0.65 | 0.67 | 0.63 |
| Al₂O₃ | 13.17 | 13.21 | 13.29 | 13.08 | 13.29 | 12.92 |
| Fe₂O₃T | 3.53 | 3.81 | 3.67 | 3.53 | 3.6 | 3.31 |
| MnO | 0.09 | 0.1 | 0.1 | 0.1 | 0.1 | 0.09 |
| MgO | 0.78 | 0.86 | 0.84 | 0.74 | 0.78 | 0.64 |
| CaO | 2.57 | 2.73 | 2.61 | 2.51 | 2.55 | 2.28 |
| Na₂O | 4.02 | 3.99 | 3.98 | 4.05 | 4.06 | 4.06 |
| K₂O | 2.59 | 2.53 | 2.48 | 2.59 | 2.59 | 2.7 |
| P₂O₅ | 0.1 | 0.11 | 0.1 | 0.1 | 0.1 | 0.08 |
| LOI | 3.78 | 3.45 | 4.17 | 3.67 | 2.75 | 3.78 |
| Total | 99.82 | 99.58 | 99.77 | 99.9 | 99.85 | 99.68 |
Table 1. Cont.

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87Sr/86Sr = 0.704857 0.704799 0.704896 0.704941 0.705015
143Nd/144Nd = 0.512732 0.512732 0.512743 0.512736
206Pb/204Pb = 0.48 0.48 0.41 0.43 0.49 0.47

a εNd = [(143Nd/144Nd)sample/(143Nd/144Nd)CHUR-1]*10000; (143Nd/144Nd)CHUR = 0.512638 cite from Jacobsen and Wasserburg [41]; CHUR=chondrite uniform reservoir; Fe2O3T is total Fe.

3. Analytical Methods

3.1. Pretreatment of Pumice Samples

The analyzed pumice clasts in this study were selected from the sediment of core S9. Because pumice vesicles are connected and easily filled by fine-grained sediments, the selected pumice must be effectively cleaned before chemical analyses. The sediments were first placed in a beaker and treated with 5% H2O2 at room temperature for at least 12 h to effectively remove the organic matter. The loose sediments were repeatedly washed with deionized water and dried in an oven at 50 °C for 24 h in preparation for selection. After that pumice clasts were selected from the residues with a binocular microscope. To effectively remove the remaining filler in the pumice vesicles, we followed the methods of Huang et al. [25] and Zhang et al. [21] for further washing. The selected pumice clasts were first washed in an ultrasonic cleaner with deionized water, and the water was changed every 20 min until the remaining water was sufficiently clear. Then, pumice samples were treated with 0.5 mol/L dilute hydrochloric acid and bathed for two hours at 60 °C to remove any secondary carbonates and Fe-Mn.
oxides. To completely remove the Cl ions, pumice clasts were then bathed in hot deionized water for several days, and the water was changed every day until no precipitation formed when AgNO$_3$ was added to the supernatant. After this step, pumice clasts were dried and ground with an agate mortar to a size of 200 mesh for further analysis.

3.2. Major and Trace Element Analysis

In this study, the selected six samples of pumice clasts from core S9 were measured for major and trace element contents, and five of them were measured for their Sr-Nd-Pb isotope ratios.

The major element compositions of the pumice samples were determined by X-ray fluorescence (XRF; Phillips PW2404) at the State Key Laboratory of Biogeology and Environmental Geology of China University of Geosciences (Wuhan), China. Aliquots of 0.5000 ± 0.0005 g of powdered sample were accurately weighed and mixed with 5 g of Li$_2$B$_4$O$_7$ to produce fused glass disks at 1050 °C in an auto-melting prototyping machine for analysis. Loss on ignition (LOI) was determined by the weight difference before and after calcinations at 1000 °C. The certified reference material GSR-3 was also measured many times for data quality control. The analytical precision and accuracy for major elements were generally better than 3%.

All trace element data were obtained by inductively coupled plasma mass spectrometry (ICP-MS; Perkin Elmer Elan 9000) at the ALS Laboratory in Guangzhou, China. Approximately 40 mg of sample powder was weighed and completely digested with concentrated HF-HNO$_3$-HClO$_4$ in an airtight Teflon container. The solution was diluted to approximately 40 g with ultrapure water for testing. Certified reference standards GBM908-10, MRGeo08, OGEoo8, OREAS-45e, OREAS 120, OREAS 460, OREAS-100a, and SY-4 were measured repeatedly with the samples for assessing the analytical results. The precision and accuracy of trace element analysis were generally better than 5%.

3.3. Sr-Nd-Pb Isotope Analysis

Sr-Nd-Pb isotope compositions were measured on a thermal ionization mass spectrometer MAT-262 (Finnigan Company, Germany) at the Radiogenic Isotope Geochemistry Laboratory, University of Science and Technology of China. Approximately 100 mg of powdered samples was placed in a 15 mL Teflon container. Then, 8–10 drops of purified HClO$_4$ and 2–3 mL of purified HF were added, followed by airtight heating for approximately one week to fully dissolve the samples. The separation and purification of Sr-Nd-Pb isotopes are different. In detail, Sr isotopes were obtained from a quartz exchange column with 5 mL AG50W-X12 exchange resin (200–400 mesh), Nd isotopes were obtained from a quartz exchange column with 1.7 mL Teflon powder, and Pb isotopes were obtained from an AG1-X8 anion exchange resin. Additionally, Sr isotope ratios were measured by the Ta metal band and Ta emitter, Nd isotope ratios were measured by the Re metal band, Pb isotope ratios were measured by the Re metal band with silica gel as emitter. The values of $^{86}$Sr/$^{88}$Sr = 0.1194 and $^{146}$Nd/$^{144}$Nd = 0.7219 were used for mass fractionation calibration of the measured Sr and Nd isotopic ratios. The long-term measurement results of standard solution NBS987 was $^{87}$Sr/$^{86}$Sr = 0.710249 ± 0.000012 (2σ, n = 38) and that of standard solution La Jolla was $^{143}$Nd/$^{144}$Nd = 0.511869 ± 0.000006 (2σ, n = 25). The measured results of Pb standard solution NBS981 was used for mass fractionation calibration of Pb isotopes. Multiple measurement results showed that the fractionation coefficient of Pb isotopes was 0.1% per atomic mass unit at 1300 °C. The accuracies of Sr and Nd isotope ratios were better than 0.003%, and those of Pb isotope ratios were better than 0.01%. During the whole process, the background values of Sr, Nd and Pb isotope analyses of the bulk sample were < 200 pg, < 100 pg, and < 200 pg, respectively. The analytical details were described by Chen et al. [42]. For convenience, $^{143}$Nd/$^{144}$Nd ratios were converted to εNd with the chondritic uniform reservoir value of 0.512638 [41].

3.4. Radiocarbon Dating

To constrain the eruption age of S9 pumice samples, planktonic foraminifera from three key layers (220–221, 229–230, and 243–244 cm; including two previous data values from Jiang et al. [43]) were
selected for radiocarbon dating by using accelerator mass spectrometers (AMS) at the American Beta Analytical Laboratory. AMS $^{14}C$ age data were calibrated to calendar years before AD 1950 with the CALIB 7.0.4 program, the calibration data set Marine13.14c [44] and Delta R = 39 ± 18 [45]. The Bacon software package was used to establish the age-depth model for core S9 [46]. In this study, we define the sedimentary ages of S9 pumice samples as their eruption age.

4. Geochemical Results

The measured major element, trace element, and Sr-Nd-Pb isotope compositions of pumice clasts are shown in Table 1.

4.1. Major and Trace Elements

All six pumice clasts plot in the field of dacite in the total alkali versus silica (TAS) diagram [47] with SiO$_2$ contents ranging from 67.86–69.37 wt.% (Figure 3a), relatively lower than most reported pumice data in the MOT [19,23,26]. The K$_2$O versus SiO$_2$ diagram shows that these pumice belong to medium-K calc-alkaline series (Figure 3b) [48]. Compared with MOT basalts and andesites, S9 pumice clasts have lower CaO (2.28–2.73 wt.%), MgO (0.64–0.86 wt.%), Al$_2$O$_3$ (12.92–13.29 wt.%), Fe$_2$O$_3^{T}$ (3.31–3.81 wt.%), and TiO$_2$ (0.63–0.69 wt.%) concentrations and higher Na$_2$O (3.98–4.06 wt.%) and K$_2$O (2.48–2.70 wt.%) contents (Figure 4).

The normal mid-ocean ridge basalt (N-MORB) normalized incompatible element patterns of S9 pumice samples are shown in Figure 5, which for comparison, also includes basalts from the MOT [16,27] and basalts from the northern and middle Ryukyu volcanic front [37]. Consistent with previously published pumice data, S9 pumice samples show pronounced enrichments in large ion lithophile elements (LILEs), Th and U, and depletions in high field strength elements (HFSEs), which suggests characteristics of subduction-related magma [49]. Additionally, S9 pumice samples exhibit significant positive K and Pb anomalies, negative Nb and Ta anomalies, and clearly negative P and Ti anomalies. Furthermore, highly incompatible trace elements (e.g., Rb and Ba; Figure 4) exhibit excellent linear correlations among the MOT basalt, andesite, and S9 pumice, whereas compatible elements (e.g., Co, Sc and V; Figure 4) decrease with increasing SiO$_2$ content from basalt to S9 pumice.

Figure 3. Classification diagrams of S9 pumice and volcanic rocks from the MOT. (a) Total alkali versus silica (TAS) diagram [47]; (b) K$_2$O versus SiO$_2$ diagram, the boundaries of different series were drawn according to [48]. MOT = middle Okinawa Trough. Data sources are from Zhang et al. [23]; Hoang and Uto [27]; Zhai and Gan [16]; Shinjo and Kato [26]; Guo et al. [24]; Guo et al. [19]; Zhai et al. [9]; Huang et al. [25]; Shinjo et al. [17].
Figure 4. Harker diagrams of S9 pumice samples, basalts, andesites, dacites, and other types of pumice from the MOT. The data sources are the same as in Figure 3.

Figure 5. Normal-MORB normalized trace element concentration patterns (a,b), and chondrite-normalized REE patterns (c,d) for the studied pumice compared with basalts from the MOT [16,17,24,27], and basalts from the northern and central Ryukyu volcanic front [37,50–53]. The normalization constants are from Sun and McDonough [54]. MOT = middle Okinawa Trough; NRVF = northern Ryukyu volcanic front; MRVF = middle Ryukyu volcanic front.
As shown on the chondrite-normalized rare earth elements (REEs) diagram (Figure 5c,d), the S9 pumice samples are enriched in light REEs (LREEs) relative to heavy REEs (HREEs) ([La<sub>N</sub>/Yb<sub>N</sub>] = 3.35–3.53), showing convex downward patterns. The bulk REE values (ΣREE = 110.06–113.62 ppm) of S9 pumice are relatively high compared with those of the associated basalts, almost twice as much as basalt ΣREE values, and the HREE concentrations are also slightly higher than those of basalt. Additionally, REE patterns show pronounced negative Eu anomalies with δEu (δEu = 2*Eu/(Sm+Gd)) ranging from 0.6 to 0.69.

4.2. Sr-Nd-Pb Isotopes

The isotope data of S9 pumice samples are presented in Table 1. These representative data have also been plotted on various covariation diagrams to allow comparison with rocks from known geological settings [55,56].

On the Sr-Nd isotope diagram (Figure 6), our pumice samples record nearly constant isotopic ratios and plot differently from those of the previously reported volcanic rocks in the MOT. In detail, S9 pumice samples have relatively higher 87Sr/86Sr ratios (0.70480–0.70502) and lower 143Nd/144Nd ratios (0.51272–0.51274; εNd = 1.67–2.05) compared with the MOT basalt [57], andesite and type I rhyolite [26] (Table 1). Additionally, these pumice samples have Sr-Nd isotope ratios similar to those of the central Ryukyu basalts [50,53] and plot within the overlapping fields of Indian Ocean MORB and Philippine Sea sediment [57–60] in the Sr-Nd isotope diagrams (Figure 6).

Figure 6. 87Sr/86Sr versus 143Nd/144Nd diagram for the studied pumice compared to volcanic rocks from the MOT [17,24,26,27,57], basalts from the northern West Philippine Basin [55,56], basalts from the northern Ryukyu volcanic front [37,50,51], basalt from the central Ryukyu arc [37]. Indian MORB, Pacific MORB and Philippine Sea sediments (PetDB database; http://www.earthchem.org/petdb) are also shown for comparison. EMI and EMII data are from Zindler and Hart [61].

Figure 7 illustrates the Pb isotope compositions of the samples. Although S9 pumice samples have 206Pb/204Pb (18.321–18.436) values similar to those of associated basalts, they display different trends on the Pb-Pb isotope diagrams. In detail, S9 pumice samples have nearly constant 207Pb/204Pb values ranging from 15.622 to 15.624. The values of 208Pb/204Pb are ranging from 38.52 to 38.63, which are higher than those of MOT basalts and northern Ryukyu volcanic front basalts [37,51,52] (Figure 7). Additionally,
all these pumice samples lie distinctly above the Northern Hemisphere reference line (NHRL) [62] and are compositionally within the field of southern Kyushu and Indian Ocean MORB (Figure 7).

Figure 7. Plots of (a–b) 207Pb/204Pb versus 206Pb/204Pb and (c,d) 208Pb/204Pb versus 206Pb/204Pb diagrams of S9 pumice samples and basalts from the MOT [17,57] and its adjacent regions. The Northern Hemisphere reference line (NHRL) [62] and representative fields (DM, EMI, EMII, PREMA, and BSE) [61] are shown for reference. Yellow inverted triangles represent basalts from the southern segment of northern Ryukyu volcanic front [37]. Data of Indian MORB and Pacific MORB are from PetDB database (http://www.earthchem.org/petdb), data of Philippine Sea sediments are from Hauff et al. [58], Plank and Langmuir [59], and Shu et al. [57]. Letters A-D in (b) and (d) represent different components discussed in the text. MOT = middle Okinawa Trough. NRVF = northern Ryukyu volcanic front.

5. Discussion

5.1. Constraints on Eruption Age

Reliable chronological framework form a basis for precise and quantitative analyses of geological processes [63]; however, eruption ages for volcanic rocks in the OT are deficient due to the difficulties in dating young geological material [64–66]. The OT is a young and active back-arc basin with volcanic activity younger than 4 Ma [31]. Chen et al. [30] and Huang et al. [25] used the U-series disequilibrium dating method to determine the ages of felsic volcanic rocks in the OT, and both imply a widespread felsic volcanic activity in the late Pleistocene.

As mentioned above, pumice clasts used in this study were selected from sediment core S9, all samples are fresh with sharp edges and corners (Figure 2b). Previous studies have suggested that all the vesicles in pumice are connected [67] and their density are lower than seawater [19]. Nevertheless, most pumice clasts sink immediately upon immersion in water after explosive volcanism [68], which means pumice is unlikely to float far and the buoyancy of freshly erupted submarine pumice is transient, especially when the pumice is hot and has erupted from the deep seafloor (e.g., depths >~200 m, temperatures > 300 °C) [68,69]. In addition, studies on pumice associated with the OT have all regarded the rocks as products of back-arc volcanism [19,23,26]. These results suggest that pumice...
samples deposited in core S9 were close to the eruption center with short-distance transportation. Therefore, we are confident that the eruption ages of pumice samples used in this study can be well constrained with the radiocarbon dating data of planktonic foraminifera.

To establish a reliable and probabilistic age framework, Bayesian accumulation histories for deposits are the best choice to constrain the sedimentation history of the investigated layers [70]. In this study, Bayesian statistics were performed by using the Bacon R software (v2.3.5) [46] to reconstruct an age-depth model for S9 by combining radiocarbon dates with prior information. Detailed analytical procedures were given in Blaauw and Christen [46]. The age-depth model was developed on the basis of three $^{14}$C ages of foraminifera selected from layers at 220–221, 229–230, and 243–244 cm (Table 2), and the results have a 95% confidence interval. As shown in Table 3, the ages of S9 pumice samples are between 11.22 cal. ka BP and 12.96 cal. ka BP, corresponding well with felsic volcanic activity that occurred during the late Pleistocene and early Holocene [25,30]. Thus, we reasonably infer that an approximately 1.74-kyr eruption record is well preserved in core S9.

Table 2. Radiocarbon dating data for core S9.

<table>
<thead>
<tr>
<th>Depth (cm)</th>
<th>Material</th>
<th>AMS $^{14}$C Age (yr BP)</th>
<th>±1σ</th>
<th>Calendar Age (cal yr BP)</th>
<th>±2σ</th>
</tr>
</thead>
<tbody>
<tr>
<td>220–221</td>
<td>Planktonic foraminifera</td>
<td>9050 a</td>
<td>30</td>
<td>9675</td>
<td>142.5</td>
</tr>
<tr>
<td>229–230</td>
<td>Planktonic foraminifera</td>
<td>10750</td>
<td>40</td>
<td>11972</td>
<td>236</td>
</tr>
<tr>
<td>243–244</td>
<td>Planktonic foraminifera</td>
<td>12437 a</td>
<td>30</td>
<td>13877</td>
<td>128.5</td>
</tr>
</tbody>
</table>

* data cited from Jiang et al. [43].

Table 3. The calculated eruption ages of S9 pumice samples.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Depth (cm)</th>
<th>Eruption Ages (Cal. yr BP)</th>
<th>±2σ</th>
</tr>
</thead>
<tbody>
<tr>
<td>F1</td>
<td>226–227</td>
<td>11219</td>
<td>214</td>
</tr>
<tr>
<td>F2</td>
<td>228–229</td>
<td>11718</td>
<td>235</td>
</tr>
<tr>
<td>F3</td>
<td>230–231</td>
<td>12220</td>
<td>189.5</td>
</tr>
<tr>
<td>F4</td>
<td>232–233</td>
<td>12470</td>
<td>198</td>
</tr>
<tr>
<td>F5</td>
<td>234–235</td>
<td>12708</td>
<td>142.5</td>
</tr>
<tr>
<td>F6</td>
<td>236–237</td>
<td>12964</td>
<td>149.5</td>
</tr>
</tbody>
</table>

5.2. The origin of S9 Pumice

The formation of volcanic rocks in the OT is complex due to the variable Quaternary magmatic activity in the three different segments [11,12,16,17]. Although grayish white pumice is widely distributed in the OT, its origin remains controversial due to the lack of available isotope data and an age framework. Previous studies suggest that there are different types of pumice with various compositions in the OT, and different models were carried out to interpret their formations: (1) most of the studies accept that rhyolitic magmas are derived from basaltic magmas through fractional crystallization or assimilation and fractional crystallization (AFC) rather than through partial melting, as they have similar Sr-Nd isotope compositions [9,19]; (2) Zhang et al. [23] proposed that the partial melting of cognate mafic to intermediate igneous rocks could also generate rhyolites with similar Sr-Nd isotope compositions.

5.2.1. Fractional Crystallization

Major element concentrations of S9 pumice samples (e.g., MgO and CaO) display a linear trend with the MOT basalts and andesites (Figure 4). The concentrations of incompatible trace elements (e.g., Ba and La) gradually increase while the concentrations of compatible trace elements (e.g., Co and V) decrease with increasing SiO$_2$ contents from MOT basaltic rocks to andesites to S9 pumice samples (Figure 4). These results indicate a magmatic evolution process from basaltic to felsic volcanic rocks. In addition, evidence from other representative elements (e.g., P, Ti, and Eu) also suggests a process of fractional crystallization.
In the early stage of magmatic crystallization, Ti and P can enter mineral phases and form ilmenite, titanomagnetite, and apatite, resulting in depletion of these elements in the residual melt during the later stage of magma evolution [19]. In the N-MORB-normalized trace element patterns, strong depletions in P and Ti in S9 pumice samples compared with associated basalts likely suggest fractionation of apatite and Fe-Ti oxides (Figure 5). Similar to volatiles, P generally shows enrichment during the late stage of felsic magmatic evolution [54]. However, P in this study shows evident depletion, further indicating that the S9 pumice are not derived from felsic magma. The K/Rb versus SiO$_2$ diagram among MOT basalt, andesite, and S9 pumice display a decreasing trend (Figure 4), which may be attributed to fractional crystallization of hornblende and this is consistent with our observation that small amount of the hornblende can be observed in S9 pumice samples. During the process of magmatic fractional crystallization, the crystallization of plagioclase generally leads to a significant negative Eu anomaly in the residual melt [15,19]. The chondrite-normalized REE patterns of S9 pumice displays a trend subparallel to that of the MOT basalts with obvious negative Eu anomalies ($\delta$Eu = 0.60–0.69; Figure 5), which suggests that the magma experienced the crystallization of plagioclase. All of these findings imply that the parent magma of S9 pumice clasts experienced a crystal fractionation process.

5.2.2. Magma Sources of S9 Pumice

The SiO$_2$ contents of pumice clasts in this study (67.86–69.37 wt.%) are relatively lower than those of previously reported pumice in the MOT (generally > 70 wt.%) [19,23,26], suggesting that S9 pumice may have different magma sources.

Studies about the magma sources of volcanic rocks in the OT were carried out since the early 1990s [7,17]. These studies have proposed that isotopic compositions provide valuable information about regional magmatic activity and evolution. However, available isotopic data are relatively few (especially Pb isotope compositions), and the source type of magma remains controversial despite many previous studies [23,26,38]. In this study, we use trace elements and Sr-Nd-Pb isotope compositions and characteristics to perform reliable analysis for the magma sources of S9 pumice. This study further confirms the well-recognized concept that magma of the back-arc basin have a complex multisource origin [12,71].

The OT is a back-arc basin located behind the Ryukyu arc-trench system that has developed above the subduction zone [35,72]. The mantle wedge beneath the Ryukyu volcanic arc is affected by various subduction components (e.g., sediments, fluids and altered oceanic crust) [37]. Therefore, the nature of the mantle wedge and the subduction components are key factors in analyzing the origins of volcanic rocks from the OT. There are two main viewpoints regarding the nature of the mantle wedge in the study area: Indian MORB-type mantle and the Pacific MORB-type mantle [12,27]. Figure 7 shows variations in $^{207}$Pb/$^{204}$Pb versus $^{206}$Pb/$^{204}$Pb and $^{208}$Pb/$^{204}$Pb versus $^{206}$Pb/$^{204}$Pb, along with the values of basalts from the MOT and the southern segment of northern Ryukyu volcanic front. These samples apparently define three distinctly separate trends and reflect the mixing of subduction components and the mantle, although components A, B, C, and D may not represent the exact Pb isotopic compositions of these end members.

Figure 7b,d show that component A is characterized by high $^{206}$Pb/$^{204}$Pb, $^{207}$Pb/$^{204}$Pb, and $^{208}$Pb/$^{204}$Pb concentrations, which is consistent with the upper continent crust. This feature can be explained by the sediment derived from the subducting Philippine Sea plate [37,57], which is widely considered to be a potential source of volcanic rocks in the OT [19,27,57,71]. Shinjo et al. [37] suggests that component B is most likely the Pacific Ocean MORB-type mantle. The effect of Pacific Ocean MORB-type mantle on OT magma is limited to the northeastern end of the NOT, and the boundary between the Indian Ocean MORB-type and the Pacific Ocean MORB-type possibly lies between Kuro-Shima and the northern segment of the north Ryukyu volcanic front (Figure 1) [12,27,37]. Please note that core S9 is located at the southwestern side of the boundary (Figure 1), and the Pb isotope characteristics of MOT basalts are convergent along the mixing line between component A and C (Figure 7b,d). Evidence from Pb isotope composition have indicated that the eastern Asian-western Pacific asthenosphere resembles
Indian Ocean MORB [73]. Findings from Guo et al. [19] and Chen et al. [71] have also suggested that the mantle wedge beneath in the middle and southern part of the OT is Indian Ocean MORB-type. Sr, Nd, and Pb isotope compositions have also shown that MOT basalt, andesite, type I rhyolite and S9 pumice are well within the Indian MORB-type field (Figures 6 and 7) [54,74]. Consequently, we infer that component C is most likely the Indian Ocean MORB-type mantle.

S9 pumice samples have relatively high $^{207}$Sr/$^{206}$Pb and $^{208}$Pb/$^{204}$Pb ratios and clearly plot leftward, trending towards an enriched mantle I (EMI)-like endmember (Figure 7a,b). This trend is different from that of basalts and seems difficult to be explained by mixing Indian Ocean MORB-type mantle and subducted sediment. This indicates that an isotopically distinct component may be responsible for this phenomenon, i.e., a component with low $^{206}$Pb/$^{204}$Pb and relatively high $^{207}$Pb/$^{204}$Pb ratios compared with depleted mantle. Similar characteristics are shown in $^{87}$Sr/$^{86}$Sr versus $^{143}$Nd/$^{144}$Nd diagram (Figure 6), S9 pumice samples have lower $^{143}$Nd/$^{144}$Nd and higher $^{87}$Sr/$^{86}$Sr than the MOT basalts and plots closer to EMI-like mantle material [61]. This further suggests that an EMI-like component might have contributed to the magmatic compositions of S9 pumice. This finding is in agreement with previous studies from the OT [17,19,27]. Generally, crustal contamination may lead to increases in SiO$_2$ and incompatible element contents and decreases in most compatible element contents [13,37].

In Figure 8, our geochemical data are compared with available data of basalts and andesites from the MOT. Figure 8 shows that Sr-Nd isotope compositions have linear correlations with SiO$_2$ and MgO contents among the MOT basalts, andesites, and S9 pumice samples. SiO$_2$ correlate positively with $^{87}$Sr/$^{86}$Sr and negatively with $^{143}$Nd/$^{144}$Nd (Figure 8a,d), while MgO correlate negatively with $^{87}$Sr/$^{86}$Sr and positively with $^{143}$Nd/$^{144}$Nd (Figure 8b,e), indicating a crustal assimilation process. Additionally, $^{87}$Sr/$^{86}$Sr gradually increases, while $^{143}$Nd/$^{144}$Nd decreases with increasing Th/La ratios among the MOT basalts, andesites, and S9 pumice (Figure 8c,f). These results suggest that the variations in the chemical and isotopic compositions of S9 pumice samples may reflect the process of crustal contamination in the MOT. Accordingly, this enriched component and crustal assimilation revealed by the geochemical characteristics of S9 pumice may most likely indicate the contribution of the lower crust, which has higher $^{87}$Sr/$^{86}$Sr and much lower $^{143}$Nd/$^{144}$Nd ratios than S9 pumice. Some lower crustal xenoliths from North China and lower crustal granodiorite from southern Kyushu generally yield higher $^{87}$Sr/$^{86}$Sr ratios and much lower $^{143}$Nd/$^{144}$Nd ratios than S9 pumice [75–78]. Letouzey and Kimura [72] suggested that the middle and northern parts of the OT were an uplifted land area constructed from Cretaceous granite and older host rocks before the late Miocene. In addition, Cretaceous granite fragments were obtained by dredging and coring in the OT [79,80]. Kizaki [28] suggested that the geological characteristics beneath the central grabens of the OT are similar to that of the Inner Zone of southwest Japan, which is featured by abundant exposures of Cretaceous to Paleogene felsic volcanic and plutonic rocks. The Sr-Nd isotope compositions of granites from the Kyushu Island display the signature of the lower crust [77,78], which can be used as representative of the lower crustal endmember.

To further verify our hypothesis, two endmember isotopic models are used to perform calculations of magmatic compositions for S9 pumice (Figures 9 and 10, Table 4) and establish their potential links with MOT basalts. The isotopic ratios and element concentrations of the endmembers used in this study are listed in Table 4. In the Sr-Nd isotope mixing model (Figure 9), variations in $^{87}$Sr/$^{86}$Sr and $^{143}$Nd/$^{144}$Nd ratios of S9 pumice samples, basalts, andesites, and type I rhyolite from the MOT indicate genetic difference among these volcanic rocks (i.e., they have different magmatic compositions). The isotopic characteristics of S9 pumice samples have suggested that their magmatic compositions may contribute from the Indian Ocean MORB-type mantle, the Philippine Sea sediments, and EMI-like component (Figures 6 and 7). The geochemical compositions of S9 pumice have also suggested the occurrence of crustal assimilation during the magmatic ascent process (Figure 8).

Therefore, upper continent crust (UCC) [81–83] is considered to be another endmember to evaluate the effect of contamination and deduce the magmatic composition of S9 pumice.
Figure 8. Plots of (a–c) $^{87}\text{Sr}/^{86}\text{Sr}$ versus SiO$_2$, MgO, and Th/La, and (d–f) $^{143}\text{Nd}/^{144}\text{Nd}$ versus SiO$_2$, MgO, and Th/La diagrams for the studied pumice and volcanic rocks from the MOT. The source data of volcanic rocks from the MOT are the same as in Figure 6. MOT = middle Okinawa Trough.

Figure 9. Calculation diagram of $^{87}\text{Sr}/^{86}\text{Sr}$ versus $^{143}\text{Nd}/^{144}\text{Nd}$ for S9 pumice samples and volcanic rocks from the MOT. Igneous rocks from the southern Kyushu, which exhibit a lower crustal provenance are also shown for comparison [13,77,78]. The black lines represent mixing between two endmembers (UCC = upper continental crust; PSs = Philippine Sea sediments; SK = southern Kyushu). The compositions of endmembers used for the calculations are listed in Table 4. MOT = middle Okinawa Trough. The source data of volcanic rocks in the MOT are the same as in Figure 6.
Figure 10. Calculation diagram of $^{143}$Nd/$^{144}$Nd versus $^{206}$Pb/$^{204}$Pb for S9 pumice and volcanic rocks from the MOT. The endmembers used for calculations in this figure are the same as in Figure 9. The source data of basalts in the MOT are the same as in Figure 6. UCC = upper continental crust; PSs = Philippine Sea sediments; LC = lower crust.

Table 4. Endmember compositions used in isotopic modeling.

<table>
<thead>
<tr>
<th>Elements</th>
<th>Indian MORB $^a$</th>
<th>UCC $^b$</th>
<th>LC $^c$</th>
<th>PSs $^d$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sr (µg/g)</td>
<td>7.664</td>
<td>333</td>
<td>250</td>
<td>177.222</td>
</tr>
<tr>
<td>Nd (µg/g)</td>
<td>0.581</td>
<td>27</td>
<td>19.5</td>
<td>33.423</td>
</tr>
<tr>
<td>Pb (µg/g)</td>
<td>0.018</td>
<td>25.46</td>
<td>7.55</td>
<td>43.545</td>
</tr>
<tr>
<td>$^{87}$Sr/$^{86}$Sr</td>
<td>0.702987</td>
<td>0.72045</td>
<td>0.70625</td>
<td>0.7122</td>
</tr>
<tr>
<td>$^{143}$Nd/$^{144}$Nd</td>
<td>0.513048</td>
<td>0.511999</td>
<td>0.5124</td>
<td>0.5123</td>
</tr>
<tr>
<td>$^{206}$Pb/$^{204}$Pb</td>
<td>18.1047</td>
<td>18.637</td>
<td>17.98</td>
<td>18.462</td>
</tr>
</tbody>
</table>

$^a$ Sr-Nd-Pb isotope data are average composition of Indian Ocean MORB from the PetDB database (http://www.earthchem.org/petdb). Trace element concentrations are from Workman and Hart [74] and Sun and McDonough [54].

$^b$ The composition of UCC is cited from Meng et al. [81], Zhang et al. [82], and Wedepohl [83].

$^c$ LC = lower crust, Sr-Nd isotope data are from references [77,78]. Pb isotope composition is from references [77,78,84].

$^d$ Philippine Sea sediment (PSs) is a composition of sediment at DSDP Sites 442/443/444. Trace and isotope compositions are from Shu et al. [57] and Mahoney [60].

To constrain the magmatic compositions of S9 pumice samples in this study, $^{87}$Sr/$^{86}$Sr, $^{143}$Nd/$^{144}$Nd, and $^{206}$Pb/$^{204}$Pb of the pumices, as well as potential sources, the Indian Ocean MORB-type mantle, the Philippine Sea sediments, lower crustal component, and UCC, together with previously reported volcanic rocks from the MOT were plotted against each other (Figures 9 and 10). S9 pumice samples mainly fall along the mixing line between the Indian Ocean MORB-type mantle and the Philippine Sea sediments, and tend to have a trend towards the mixing line between the Indian Ocean MORB-type mantle and lower crustal component [77,78,84], but far away from the mixing line between the Indian Ocean MORB-type mantle and the UCC (Figure 9). These results suggest that the magmatic composition of S9 pumice is a two-endmembers mixture of the Indian Ocean MORB-type mantle and the Philippine Sea sediments. The UCC did not appear to have significant influence on the magmatic composition.
of S9 pumice samples. In addition, it might be assumed that the parent magma for S9 pumice was slightly contaminated by a lower crustal component, which is relatively evident in the $^{143}\text{Nd}/^{144}\text{Nd}$ versus $^{206}\text{Pb}/^{204}\text{Pb}$ diagram (Figure 10).

The $^{143}\text{Nd}/^{144}\text{Nd}$ versus $^{206}\text{Pb}/^{204}\text{Pb}$ diagram was further employed to evaluate the contribution of lower crustal component on the magmatic composition of S9 pumice (Figure 10). The mixture of Indian Ocean MORB-type mantle and Philippine Sea sediments was inadequate to explain the Pb isotope composition of magma that S9 pumice derived from. Lower crustal component is considered to be an endmember for calculation, as expected, the magmatic compositions of S9 pumice may originate from the mixture of Indian Ocean MORB-type mantle, subducted Philippine Sea sediments, and lower crustal component. In addition, we found that the Sr-Nd-Pb isotope composition of S9 pumice samples, which were erupted in different periods (12.96 cal. ka BP to 11.22 cal. ka BP), displaying no temporal variation trend and varying in narrow ranges (Tables 1 and 3). According to the above discussion, we infer that S9 pumice was generated from a hybrid magma that was mainly composed of primitive mantle-derived magma with slight assimilation of a lower crustal source, while the primitive mantle-derived magma was originated from Indian Ocean MORB-type mantle metasomatized by subducted Philippine Sea sediments. The magma sources of S9 pumice clasts have remained relatively stable during the eruption history.

To investigate the magma evolution process of the S9 pumice samples, quantitative modeling of fractional crystallization based on major element contents are carried out according to Rayleigh fractionation calculations using constraints of least-squares modeling (Supplementary Table S1). Shinjo and Kato [26] proposed that the observed REE patterns of andesite, dacite, and rhyolite can be reproduced using a three-step fractional crystallization model (i.e., basalt-andesite-dacite-rhyolite), suggesting that dacite can be produced through a two-step fractional crystallization of the associated basalt. According to previous studies [17,26,27], the basalt and andesite in the MOT tend to have similar Sr-Nd isotope compositions (Figure 8), indicating that the S9 pumice samples might be slightly contaminated by the lower crustal material when the magma evolved from andesite to dacite. Therefore, we mainly focus on the fractional crystallization from andesite to dacite (i.e., S9 pumice) in this study. We used the mixed andesite (i.e., 99.5–98.5% 287-2A + 0.5–1.5% lower crust; 287-2A from Shinjo and Kato [26]) to calculate the REE compositions of S9 pumice samples (Table 5; Figure 11). Because S9 pumice samples have similar geochemical compositions, their normalized formulas of the crystalline phases are nearly consistent. Approximately 50% fractionation of plagioclase, clinopyroxene, amphibole, magnetite, and apatite can account for the compositional variation from the mixed andesite to S9 pumice clasts. Therefore, we infer that the fractional crystallization of associated basalt can generate the observed compositions of S9 pumice clasts.

### Table 5. Partition coefficients ($K_d$ values) used in modeling.

<table>
<thead>
<tr>
<th>Element</th>
<th>Pl</th>
<th>Opx</th>
<th>Amph</th>
<th>Mt</th>
<th>Ap</th>
<th>Bulk D</th>
</tr>
</thead>
<tbody>
<tr>
<td>La</td>
<td>0.19</td>
<td>0.031</td>
<td>1</td>
<td>1.5</td>
<td>14.5</td>
<td>0.707</td>
</tr>
<tr>
<td>Ce</td>
<td>0.111</td>
<td>0.028</td>
<td>0.899</td>
<td>1.3</td>
<td>21.1</td>
<td>0.697</td>
</tr>
<tr>
<td>Nd</td>
<td>0.09</td>
<td>0.47</td>
<td>1.6</td>
<td>1</td>
<td>32.8</td>
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<td>Sm</td>
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<td>0.28</td>
<td>2</td>
<td>1</td>
<td>46</td>
<td>1.318</td>
</tr>
<tr>
<td>Eu</td>
<td>0.376</td>
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<td>3.2</td>
<td>0.6</td>
<td>30.4</td>
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</tr>
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<td>Gd</td>
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<td>0.039</td>
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<td>0.3</td>
<td>43.9</td>
<td>1.12</td>
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<td>Dy</td>
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<td>1.223</td>
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<td>15.4</td>
<td>0.966</td>
</tr>
<tr>
<td>Lu</td>
<td>0.056</td>
<td>0.71</td>
<td>2.1</td>
<td>1</td>
<td>13.8</td>
<td>0.981</td>
</tr>
</tbody>
</table>
The variations in geochemical and mineralogical compositions of volcanic rocks from the OT suggest a convecting magma chamber for continuous supply of magma during this period. Geochemical studies and assimilated by a small amount of lower crustal component in the magma chamber, followed by extensive fractional crystallization to generate the parent magma. On the basis of our radiocarbon dating data, we conclude that core S9 records felsic volcanic activity in the MOT, which occurred during 12.96–11.22 cal. ka BP.

### 5.3. Implications for the Existence of a Millennial-Scale Magma Chamber

The existence of a magma chamber is generally confirmed by geophysical characteristics (e.g., seismic tomography and thermal considerations) [89–91] or large-volume volcanic eruptions [92,93]. The variations in geochemical and mineralogical compositions of volcanic rocks from the OT suggest a complex magmatic system beneath the OT [20,22,71].

Through a comprehensive study of the Sr, Nd, and Pb isotope and trace element compositions of S9 pumice samples, we infer that a magma chamber existed in the lower crust of the MOT. This hypothesis can be supported by many evidence. In this study, the felsic magma was erupted from 12.96 to 11.22 cal. ka BP and lasted for at least 1.74 ka. The felsic pumice studied by Chen et al. [30] and Huang et al. [25] from the MOT erupted in 12.66 cal. ka BP and 17.7 cal. ka BP. This indicate that felsic volcanic activities were widespread in the MOT during the late Pleistocene and early Holocene. The similarities (small variations) in the trace element abundances,REE patterns, and Sr-Nd-Pb isotope compositions (Table 1) among these pumice samples in S9 suggest the existence of a long-term, relatively stable and convecting magma chamber for continuous supply of magma during this period. Geochemical studies
by Guo et al. [19] and Zhang et al. [23] inferred that the white pumice magma chamber was probably located in the lower crust, and mineralogical study by Zhai et al. [9] implied that the crystallization of plagioclase and pyroxene occurred approximately 30 km below the seafloor. Therefore, we infer that the location of magma chamber of the felsic pumice in S9 is in the lower crust, even near the Moho. The Indian Ocean MORB-type mantle and a small amount of lower crustal component and subducted Philippine Sea sediments were mixed, assimilated and homogenized within a shallow magma chamber with a homogeneous isotopic signature (Table 1). Then the magma would further crystallize and fractionate into dacitic magma, after that the melt moved upward and finally erupted from the seafloor. This study supports the existence of a magma chamber beneath the MOT, which played an important role in the creation of the large volumes of silicic magma with nearly identical isotopic compositions from 12.96 to 11.22 cal. ka BP.

6. Conclusions

In this study, we presented new major and trace element concentrations and Sr, Nd, and Pb isotope data, as well as eruption ages for pumice samples recovered from sediment core S9 from the MOT. The geochemical compositions of S9 pumice samples erupted during 12.96 and 11.22 cal. ka BP display no temporal variation trend and vary in narrow ranges. All pumice clasts are dacite, showing characteristics of subduction-related magma and recording relatively higher Sr ($^{87}$Sr/$^{86}$Sr = 0.70480–0.70502) and Pb ($^{206}$Pb/$^{204}$Pb = 18.321–18.436, $^{207}$Pb/$^{204}$Pb = 15.622–15.624, and $^{208}$Pb/$^{204}$Pb = 38.52–38.63) and lower Nd ($^{143}$Nd/$^{144}$Nd = 0.51272–0.51274) isotope compositions than basalts from the MOT. The parent magma of S9 pumice was most likely generated from a hybrid magma through an extensive fractional crystallization process. The hybrid magma was mainly formed from the mantle-derived magma (i.e., Indian Ocean MORB-type mantle metasomatized by subducted Philippine Sea sediments) assimilated by a small amount of lower crustal component. The data suggest that a magma chamber existed in the lower crust of the MOT during 11.22–12.96 cal. ka BP and supplied felsic magma for at least 1.74 ka.

Supplementary Materials: The following are available online at http://www.mdpi.com/2077-1312/7/12/462/s1, Supplementary Table S1 shows the results of the Rayleigh fractionation calculations based on the constraints of least-squares.


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Conflicts of Interest: The authors declare no conflict of interest.

References


42. Chen, P.; Siebel, W.; Satir, M.; Terzioğlu, M.; Saka, K. Geochronology of the Karadere basement (NW Turkey) and implications for the geological evolution of the Istanbul zone. *Int. J. Earth Sci.* **2002**, *91*, 469–481. [CrossRef]


90. Detrick, R.S.; Buhle, P.; Vera, E.; Mutter, J.; Orcutt, J.; Madsen, J.; Brocher, T. Multi-channel seismic imaging of a crustal magma chamber along the East Pacific Rise. Nature 1987, 336, 35–41. [CrossRef]

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