20. Rb-Sr ISOTOPE SYSTEMATICS AND Sr/Ca-Ba/Ca RATIOS OF NAURU BASIN BASALTS, DEEP SEA DRILLING PROJECT LEG 89¹

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ABSTRACT

Four samples of Nauru Basin basalts (Cores 94 to 109 of Hole 462A, sub-bottom depth 1077-1209 m) have 87 Sr/ 86 Sr ratios in the range 0.7037 to 0.7038, which is distinctly higher than the ratios of N-type MORB. The Rb contents of the samples are depleted in comparison with those of MORB and ocean-island basalts. These chemical and isotopic characteristics are identical to those of the basalts previously drilled during Leg 61 (Cores 75 to 90 of Hole 462A), and are explained in terms of inhomogeneity of the source region in the mantle or later alteration effects.

Sr/Ca-Ba/Ca systematics of 15 samples from Cores 462A-94 to 462A-109 and 14 samples from Cores 462A-75 to 462A-90 suggest that the Nauru Basin basalts are derived from a mantle peridotite by 20 to 30% partial melting with subsequent plagioclase crystallization.

INTRODUCTION

The Nauru Basin basalts cored in Hole 462A during DSDP Leg 61 have elemental abundances similar to those of N-type MORB, except that they have lower K and Rb concentrations and distinctly higher ⁸⁷Sr/⁸⁶Sr ratios than N-type MORB (Fujii et al., 1981). Trace-element characteristics are intermediate between those of N- and Ttypes of MORB (Batiza et al., 1980). Therefore, Tokuyama and Batiza (1981) proposed naming these peculiar basalts "ocean-plateau tholeiite," as a new type of basalt produced by intraplate volcanism.

One of the most interesting problems of the Nauru Basin basalts is their age. DSDP Site 462 lies on the magnetic anomaly between M26 and M27, which corresponds to about 155 Ma (Larson and Schlanger, 1981), but the age of the basalts was determined to be Aptian (~110 Ma) from the fossils in the interbedded sediments (Site 462 report, this volume) and to be 110 \pm 3 Ma by means of ⁴⁰Ar-³⁹Ar dating of one sample, 462A-32-1, 46-49 cm (Ozima et al., 1981). This age discrepancy casts doubt on the validity of the magnetic-anomaly age, unless the argument of Larson and Schlanger (1981) is accepted that anomalies M26 and M27 originate in the Pacific Plate basalts that presumably underlie the Aptian basalts.

On Leg 89, the 1068.5 m of Hole 462A cored during Leg 61 was deepened by 140.5 m, to 1209 m sub-bottom (Leg 89 Scientific Party, 1983). The newly drilled section also contained basalt sheet-flows, which represent earlier stages of volcanism of the lower flows cored during Leg 61. We characterize here the Rb-Sr and Sr/Ca-Ba/ Ca systematics of volcanic rocks in the Nauru Basin, using the newly drilled basalts, as a continuation of work on Leg 61 basalts (Fujii et al., 1981).

EXPERIMENTAL METHODS

Each sample was pulverized, using an agate mortar, to powder finer than 150 mesh. Ba, Sr, and Ca were determined by inductively coupled plasma-optical emission spectroscopy (Onuma et al., 1981).

For two samples, 462A-102-4, 3-5 cm and 462A-109-1, 106-108 cm, we removed the extremely fine grains ("fine whole rock") by settling in acetone. We then carried out a further density separation of the powder by centrifugation in mixed methylene iodide and acetone (densities of 2.62, 2.90, and 3.30 g/cm³). Plagioclase is concentrated in the 2.62 to 2.90-g/cm³ fraction, and clinopyroxene is concentrated in the >3.30-g/cm³ fraction. Rb and Sr determinations were done using isotope dilution. For each sample, Rb–Sr and ⁸⁷Sr/⁸⁶Sr analyses were done in 5 fractions: "whole rock," "fine whole rock," 2.62 < ρ < 2.90, 2.90 < ρ < 3.30, and ρ > 3.30 g/cm³. Details of strontium-isotope analyses were described by Notsu (1983). The measured range of ⁸⁷Sr/⁸⁶Sr ratios of NBS 987 standard was 0.71028 to 0.71033, and the mean value was 0.71030 during the period of this study. Blanks for Rb and Sr throughout a whole procedure were approximately 0.3 ng and 2.5 ng, respectively.

RESULTS AND DISCUSSION

Bulk Rb-Sr Characteristics of the Nauru Basin Basalts

Table 1 shows the results of Rb–Sr and ⁸⁷Sr/⁸⁶Sr analyses. The ⁸⁷Sr/⁸⁶Sr ratios of the Leg 89 samples are 0.70372 to 0.70379, identical to those of the lower sill basalts of the Nauru Basin (0.70358–0.70403; mean 0.70370) (Fujii et al., 1981). The Rb and Sr contents of the Leg 89 samples are also similar to those of the lower sill basalts. This is reasonable because the Leg 89 samples represent the continuation downward of the lower sills drilled during Leg 61. We emphasize that the ⁸⁷Sr/ ⁸⁶Sr ratios of the Nauru Basin basalts are distinctly higher than those of N-type MORB and similar to those of oceanic island basalts. We also confirm that the Rb contents are depleted relative to those in MORB and oceanic island basalts. This means either that the source re-

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Table 1. Rb and Sr contents and ⁸⁷Sr/⁸⁶Sr ratios of Hole 462A basalts.

Sample	Rb (ppm) ^a	Sr (ppm) ^b	⁸⁷ Rb/ ⁸⁶ Sr	87 _{Sr} /86 _{Sr}
94-6, 38-40				
Whole rock	0.85 ± 7	93	$0.0246~\pm~20$	0.70372
102-4, 3-5				
Whole rock	0.40 ± 2	97	0.0120 ± 6	0.70375
Fine whole rock	0.81 ± 2	120	0.0196 ± 5	0.70372
$2.62 < \rho < 2.90$	0.18 ± 1	185	0.0028 ± 2	0.70372
$2.90 < \rho < 3.30$	0.25 ± 2	42	0.0174 ± 14	0.70370
$\rho > 3.30$	0.16 ± 2	6.3	0.075 ± 9	0.70390
105-2, 7-9				
Whole rock	0.31 ± 2	99	$0.0092~\pm~6$	0.70373
109-1, 106-108				
Whole rock	0.43 ± 1	102	0.0122 ± 3	0.70379
Fine whole rock	0.78 ± 2	124	0.0182 ± 4	0.70374
$2.62 < \rho < 2.90$	0.22 ± 2	198	0.0032 ± 3	0.70366
$2.90 < \rho < 3.30$	0.21 ± 2	92	0.0067 ± 6	0.70370
$\rho > 3.30$	0.16 ± 2	9.5	$0.048~\pm~6$	0.70390

Note: Density (ρ) is in g/cm³.

^a Errors are 2σ mean.

^b 2σ mean errors are less than 2%.

^c 2σ mean error of each analysis is less than 0.00002, but the reproducibility of the same sample is less than 0.00005. So the uncertainty associated with each ratio is estimated to be ± 0.00005 .

gion of the Naura Basin basalts has chemical and isotopic characteristics different from those of MORB or oceanic island basalts or that the later alteration effects have changed the original chemistry of the Nauru Basin basalts.

The Attempt to Obtain Rb-Sr Internal Isochron Age

The Rb-Sr results of separated fractions of two samples show that for both samples Rb is concentrated in the "fine whole-rock" fraction and Sr is concentrated in the 2.62 $< \rho < 2.90$ -g/cm³ fraction. According to the petrographic data for the thin section prepared from 462A-102-4, 7-8 cm (which is adjacent to our analyzed Sample 462A-102-4, 3-5 cm in the same rock piece) and for the thin section prepared from 462A-109-1, 100-103 cm (which is near Sample 462A-109-1, 106-108 cm), both samples have undergone alteration. Sample 462A-109-1, 100-103 cm is more abundant in clay minerals than Sample 462A-102-4, 7-8 cm (Site 462 report, this volume). Because clay minerals are more readily crushed finer, they are concentrated in the "fine whole-rock" fraction as a result of grain-size distinction in our separation procedure. Therefore, Rb may be enriched in this fraction. On the other hand, Sr is concentrated in plagioclase.

Figures 1 and 2 show the relationships between the ${}^{87}\text{Sr}/{}^{86}\text{Sr}$ and ${}^{87}\text{Rb}/{}^{86}\text{Sr}$ ratios for separated fractions of Samples 462A-102-4, 3-5 cm and 462A-109-1, 106-108 cm, respectively. For both cases, the fractions with $\rho > 3.30 \text{ g/cm}^3$ have higher ${}^{87}\text{Rb}/{}^{86}\text{Sr}$ and ${}^{87}\text{Sr}/{}^{86}\text{Sr}$ ratios than the other four fractions, but it is difficult to determine precise ages at such low ${}^{87}\text{Rb}/{}^{86}\text{Sr}$ values (<0.08) and in such young material. For the purpose of discus-



Figure 1. 87 Sr/ 86 Sr ratios versus 87 Rb/ 86 Sr ratios of Sample 462A-102-4, 3-5 cm basalt. W: whole rock. F: fine whole rock. L: 2.62 $< \rho < 2.90$ g/cm³. M: 2.90 $< \rho < 3.30$ g/cm³. H: $\rho > 3.30$ g/ cm³. 100 Ma and 200 Ma reference lines are drawn to indicate a scale of age.



Figure 2. ⁸⁷Sr/⁸⁶Sr ratios versus ⁸⁷Rb/⁸⁶Sr ratios of Sample 462A-109-1, 106-108 cm basalt. W: whole rock. F: fine whole rock. L: $2.62 < \rho < 2.90 \text{ g/cm}^3$. M: $2.90 < \rho < 3.30 \text{ g/cm}^3$. H: $\rho > 3.30 \text{ g/cm}^3$. 100 Ma and 200 Ma reference lines are drawn to indicate a scale of age.

sion, 100-Ma and 200-Ma reference lines are drawn through the "whole-rock" samples in both figures. If we assume that alteration occurred soon after cooling, we can only say that the Rb-Sr data indicate that Samples 462A-102-4, 3-5 cm and 462A-109-1, 106-108 cm are more than several tens Ma but less than a few hundreds Ma. According to the Site 462 report (this volume), lower Aptian or older sediment was drilled at the sub-bottom depth of 1123 m. Our estimate of Rb-Sr age is consistent with the sediment age, but also includes the Jurassic age of the magnetic anomaly (Larson and Schlanger, 1981). A more precise absolute age of the Nauru Basin basalts is therefore a prerequisite for discussion of the nature of the primary magma generation.

A possibility remains, however, that, because of alteration, the Nauru Basin basalts may not give a meaningful Rb-Sr age. During alteration, the original distribution of Rb and Sr in a rock is disturbed. By basaltseawater interaction at high temperature, Rb is leached from the rock and enriched in clay minerals (Menzies and Seyfried, 1979). As mentioned before, the two samples analyzed in this work contain clay minerals. It is not known whether such alteration disturbs the Rb-Sr isochron in these samples.

Sr/Ca-Ba/Ca Diagram for Elucidation of Magma Genesis

Figure 3 shows a Sr/Ca-Ba/Ca diagram (SB diagram) (Onuma et al., 1983) for analysis of the partial melting process in the mantle and the fractional crystallization process in the magma chamber.

In mantle peridotite (for example, a garnet peridotite with chondritic Sr/Ca and Ba/Ca ratios), Sr^{2+} and Ba^{2+} , with larger ionic radii, should be accommodated in accessory mineral phases such as apatite and phlogopite, whereas Ca^{2+} , with smaller ionic radius, should be taken up in major mineral phases such as garnet and clinopyroxene, as well as in the accessory phases. Orthopyroxene and olivine do not accept Ca^{2+} , Sr^{2+} , or Ba^{2+} .

A primary magma generated by small degrees of partial melting should have larger Sr/Ca and Ba/Ca ratios, since the accessory minerals are the first to melt. A primary magma with a larger degree of partial melting should have smaller Sr/Ca and Ba/Ca ratios because of addition of Ca^{2+} to the melt by decomposition of garnet and clinopyroxene. Melting and/or crystallization of olivine does not change Sr/Ca and Ba/Ca ratios of the melt.

If a series of primary magmas derived from a common mantle peridotite is available, we could expect to get a partial-melting line with a slope of about 45° passing the mantle peridotite in the SB diagram. Primitive basalts from an island-arc region fall on a partial-melting line through a mantle peridotite with chondritic Sr/ Ca and Ba/Ca ratios. The slope of about 45° means that the partial melting process is operating at nearly constant Sr/Ba ratio with variable Ca supply. The degrees of partial melting marked on the line are calculated numbers based on a melting experiment of a garnet peridotite by Mysen and Kushiro (1977). A series of primary magmas derived from a Ba-enriched source ("fertile mantle") or from a Ba-depleted source ("depleted mantle") with the same Sr contents should make different partial melting lines, as shown in Figure 3.

Plagioclase and clinopyroxene fractionation of a primary magma can be studied using Sr/Ca-Ba/Ca systematics. Olivine and magnetite crystallization cannot be visualized in the SB diagram, since these minerals do not accept Ca^{2+} , Sr^{2+} , or Ba^{2+} . Orthopyroxene crystallization might increase Sr/Ca and Ba/Ca ratios of the primary magmas in negligibly small amount, because orthopyroxene accepts small amounts of Ca^{2+} but excludes Sr^{2+} and Ba^{2+} . On the other hand, clinopyroxene, garnet, amphibole, and plagioclase crystallization changes Sr/Ca and Ba/Ca ratios of the primary magma drastically under the crystal-structure control of the Ca, Sr, and Ba partition between the magma and these minerals. As shown in Figure 3, a primary magma is evolved toward the upper right by clinopyroxene crystallization,



Figure 3. Schematic presentation of the principle of the Sr/Ca-Ba/Ca diagram for elucidation of magma genesis. (Cpx = clinopyroxene; Pl = plagioclase.)

whereas the magma is evolved toward the right by plagioclase crystallization.

Nauru Basin Basalts on the Sr/Ca-Ba/Ca Diagram

Figure 4 shows an SB diagram for Nauru Basin basalts (Hole 462A, lower unit) along with the Reykjanes Ridge basalts (Hole 409). Data are taken from Table 2 and Fujii et al. (1981) for the Nauru Basin basalts and from Tarney et al. (1979) for the Reykjanes Ridge basalts.

As already pointed out by Fujii et al. (1981), Nauru Basin basalts are very similar to N-type MORB in their Ca, Sr, and Ba abundances. The Nauru Basin basalts and N-type MORB are both depleted in Ba compared with the Reykjanes Ridge basalts. In other words, the Reykjanes Ridge basalts are more primitive than the socalled N-type MORB.

The Reykjanes Ridge basalts make a triangular region, whereas the Nauru Basin basalts make a horizontal band, on the SB diagram. The triangular region defined by the Reykjanes Ridge basalts suggests that (1) the source material is slightly depleted in Ba compared with a chondritic mantle, (2) most of the basalts are generated by 20 to 30% partial melting of the source material, with subsequent plagioclase crystallization, and (3) several basalts are derived from the source material by 7 to 15% partial melting without plagioclase crystallization. The horizontal band defined by the Nauru Basin basalts suggests, on the other hand, that (1) the source material is strongly Ba-depleted compared with the chon-

Table 2. Ca, Sr, and Ba contents (ppm) of Hole 462A basalts.

Sample (interval in cm)	Ca	Sr	Ba	Sr/Ca	Ba/Ca
94-1, 123	77600	95	10	1.22×10^{-3}	1.3×10^{-4}
94-3, 117	79500	101	9	1.27×10^{-3}	1.1×10^{-4}
94-6, 38-40	77600	92	9	1.19×10^{-3}	1.2×10^{-4}
96-3, 118-120	77800	95	9	1.22×10^{-3}	1.2×10^{-4}
98-7, 70-72	80100	96	6	1.20×10^{-3}	0.7×10^{-4}
99-1, 143-145	78400	92	6	1.17×10^{-3}	0.8×10^{-4}
100-2, 74	77200	94	6	1.22×10^{-3}	0.8×10^{-4}
100-2, 133-135	77900	95	7	1.22×10^{-3}	0.9×10^{-4}
102-4, 3-5	77800	96	11	1.23×10^{-3}	1.4×10^{-4}
103-1, 76-78	79400	98	5	1.23×10^{-3}	0.6×10^{-4}
104-1, 88-90	78900	98	6	1.24×10^{-3}	0.8×10^{-4}
105-2, 7-9	78600	102	6	1.30×10^{-3}	0.8×10^{-4}
106-1, 24	78000	98	5	1.26×10^{-3}	0.6×10^{-4}
108-2, 36-38	76400	98	7	1.28×10^{-3}	0.9×10^{-4}
109-1, 106-108	79600	105	6	1.32×10^{-3}	0.8×10^{-4}

dritic mantle, and (2) the basalts are derived from the source material by 20 to 30% partial melting with subsequent plagioclase crystallization, if we assume that Sr is fixed but Ba is mobile in the mantle.

Thus, the difference in Ba content between the Nauru Basin basalts and the Reykjanes Ridge basalts may be explained by different source materials. The source material for the Nauru Basin basalts corresponds to a "Badepleted" or "barren" mantle; the source material for the Reykjanes Ridge basalts corresponds to a "Ba-normal" or "normal" mantle.

There is an alternative explanation, however, for the difference between them. Ca and Sr are accommodated



Figure 4. Comparison, based on Sr/Ca-Ba/Ca systematics, of the Nauru Basin basalts and the Reykjanes Ridge basalts.

in clinopyroxene and plagioclase in the basalts, but Ba is originally situated in the groundmass or glass, which is easily altered. Therefore, the possibility that Ba in the specimens had been leached out during the subsequent alteration process cannot be ruled out.

Whatever results follow, the SB diagram suggests that the Nauru Basin basalts are derived from a mantle peridotite by 20 to 30% partial melting with subsequent plagioclase crystallization.

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