3-1986

Gabbroic Xenoliths and Host Ferrobasalt from the Southern Juan de Fuca Ridge

Jacqueline Eaby Dixon
California Institute of Technology, jdixon@usf.edu

David A. Clague
U.S. Geological Survey

Jean-Philippe Eissen
Universite Louis Pasteur

Follow this and additional works at: https://digitalcommons.usf.edu/msc_facpub

Part of the [Life Sciences Commons](#)

Scholar Commons Citation

Dixon, Jacqueline Eaby; Clague, David A.; and Eissen, Jean-Philippe, "Gabbroic Xenoliths and Host Ferrobasalt from the Southern Juan de Fuca Ridge" (1986). *Marine Science Faculty Publications*. 1322.
https://digitalcommons.usf.edu/msc_facpub/1322

This Article is brought to you for free and open access by the College of Marine Science at Digital Commons @ University of South Florida. It has been accepted for inclusion in Marine Science Faculty Publications by an authorized administrator of Digital Commons @ University of South Florida. For more information, please contact digitalcommons@usf.edu.

Gabbroic Xenoliths and Host Ferrobasalt From the Southern Juan de Fuca Ridge

JACQUELINE EABY DIXON¹ AND DAVID A. CLAGUE

U.S. Geological Survey, Menlo Park, California

JEAN-PHILIPPE EISSEN²

Universite Louis Pasteur, Laboratoire de Petrographie, Strasbourg, France

Rare isotropic gabbroic xenoliths occur in sheet and lobate flow fragments of nearly aphyric ferrobasalt collected along a 12-km section of the southern Juan de Fuca Ridge. Xenoliths comprise $\ll 1\%$ of the dredge contents and range in size from 1 cm³ (glomerocryst) to 240 cm³. The xenoliths have ophitic to intersertal texture with 5–50% interstitial glass of ferrobasaltic composition more evolved than the host lava. On the basis of texture and mineralogy, the xenoliths have been subdivided into three types: type I, plagioclase + olivine + glass; type II, plagioclase + augite + glass $\pm$ olivine; and type III, plagioclase + augite + olivine + glass $\pm$ pigeonite (partially inverted) + Fe-Ti oxides. Mineral and glass inclusion compositions suggest a sequence of evolution for the three xenolith types in which type I is the least evolved and type III is the most evolved. Application of a graphical pyroxene geothermometer to augite in xenolith types II and III yields crystallization temperatures of 1100°–1200°C and to host-lamellae pairs in inverted pigeonite yields subsolidus equilibrium temperatures of 1100°–1150°C. Coexisting titanomagnetite-ilmenite pairs in type III xenoliths yield temperature estimates of 1000°–1070°C and log $f_{O_2} = -9.7$ to -10.8 . We infer that the xenoliths represent the partially crystalline “mush” boundary zone of a magma chamber based on the abundance of interstitial glass, zonation of mineral grains in the most crystalline samples, and coherence of chemical trends between interstitial glass, glass inclusions, and mineral phases. The evolved composition of the xenoliths provides evidence for the presence of melts more fractionated than the host ferrobasalt in the magma chamber. The erupted ferrobasalt is a hybrid lava formed by mixing these highly evolved melts with more primitive melts.

INTRODUCTION

Recent studies of oceanic crust suggest that size and longevity of axial magma chambers beneath oceanic spreading centers and the petrology of the erupted igneous rocks are largely a function of spreading rate. Magma chambers beneath slow spreading centers are thought to be small and in some places transitory [Stakes *et al.*, 1984], while beneath intermediate to fast spreading centers they are steady state [Macdonald, 1982]. A natural consequence of steady state magma chambers undergoing continuous fractionation is mixing between batches of primitive magma and residual differentiates. Mixing is now recognized as an important petrologic process in the generation of mid-oceanic ridge basalt (MORB) and has been invoked by various authors to explain (1) the eruption of homogeneous basalt compositions over long time periods [Usselman and Hodge, 1978], (2) the presence of anomalous phenocrysts and melt inclusions [Dungan and Rhodes, 1978], (3) the concentrations of incompatible elements in some moderately evolved compositions in excess of that predicted by simple fractional crystallization models [O'Hara, 1977; Bryan and Moore, 1977; Bryan *et al.*, 1979; Stakes *et al.*, 1984], (4) a reversal in mineral crystallization sequences [Walker *et al.*, 1979], (5) magma with chemical and mass balance characteristics of plagioclase accumulation even though the lavas contain few or no plagioclase phenocrysts [Flower, 1982], and (6) the homogeneity of isotopic compositions along moderate to fast spreading centers [Cohen and O'Nions, 1982; Allegre *et al.*, 1983; Batiza, 1984].

The role of mixing in the production of ferrobasalt, how-

ever, is less certain. Clague and Bunch [1976] used linear least squares mixing models of major elements to show that ferrobasalt can be produced by shallow level fractionation of plagioclase, clinopyroxene, and minor olivine in the average proportions of 9.3:7.7:1 with up to 74% of the parental magma fractionally crystallizing. Matthey and Muir [1980], on the other hand, found that a model for mixing of periodically injected batches of primitive magma with residual differentiates could more closely predict variations in major and trace elements in ferrobasalt from Deep Sea Drilling Project (DSDP) sites 424 and 425 near the Galapagos spreading center 86°W.

Natland [1983] found no geochemical evidence for mixing between olivine tholeiite and ferrobasalt at the East Pacific Rise near the Siqueiros Fracture Zone. He argued that ferrobasalt in this area formed due to crystal fractionation in small isolated magma bodies such as dikes or shallow intrusions above larger magma chambers. Sinton *et al.* [1983] proposed that ferrobasalt is generated in small isolated magma bodies behind propagating rift tips. Farther behind the propagating rift tip, extreme differentiation becomes less likely as the steady state thermal configuration of a normal ridge is approached. These studies suggest that the extent of differentiation of MORB may be controlled by a delicate balance between cooling and magma supply rates.

The effects of differentiation and mixing are recorded in oceanic plutonic rocks, but at the present time, sampling of the plutonic rocks is severely biased toward the Atlantic and Indian oceans, where slow spreading rates result in large displacements along transform faults and rift valley boundaries [Engel and Fisher, 1975; Hodges and Papike, 1976; CAY-TROUGH, 1979; Fox and Stroup, 1981]. Data for gabbros from intermediate to fast spreading centers are less abundant [Vanko and Batiza, 1982; Hebert *et al.*, 1983]. Moreover, interpretation of magmatic conditions is impeded by deformation, metamorphism, or brecciation of most oceanic gabbroic and ultramafic rocks.

¹Now at Division of Geological and Planetary Sciences, California Institute of Technology, Pasadena, California.

²Now at Office de la Recherche Scientifique et Technique Outre-Mer, Noumea, New Caledonia.

Fig. 1. Map showing location of mid-ocean ridge segments in the northeast Pacific. Location of U.S. Geological Survey study area is shown by the solid square.

Rare glass-bearing gabbroic xenoliths occur in nearly aphyric ferrobassalt collected from the southern Juan de Fuca Ridge. In this report we present major and trace element data for the host ferrobassalt and mineral phase chemistry for these unaltered gabbroic xenoliths. Detailed discussion of lava chemistry provides a framework for comparison with the gabbroic xenoliths. The data are used to evaluate the role of mixing in the formation of ferrobassalt and the longevity of subaxial magma chambers at the Juan de Fuca Ridge.

GEOLOGIC SETTING

The Juan de Fuca Ridge extends 500 km from the Blanco Fracture Zone to the Sovanco Fracture Zone and is a moderate-rate spreading center (30 mm yr⁻¹ half rate) separating the Pacific and Juan de Fuca plates (Figure 1). Previous petrologic studies of basalt from the Juan de Fuca Ridge show that the southern two thirds of the ridge (south of the Cobb offset) is characterized by strongly fractionated mid-oceanic ridge basalt (MORB) with high FeO, TiO₂, MnO, P₂O₅, Na₂O, and K₂O and low MgO, CaO, and Al₂O₃ [Kay et al., 1970; Scheidegger, 1973; Detrick and Lynn, 1975; Vogt and Byerly, 1976; Wakeham, 1977; Delaney et al., 1982]. Ferrobassalt (>10% FeO*, total iron calculated as FeO) is particularly abundant near the Juan de Fuca Ridge–Blanco Fracture Zone intersection and 20–30 km south of the Cobb offset. The maximum iron and titanium enrichment in the lavas occurs behind zones identified as propagating rifts in both regions [Wilson et al., 1984].

High-amplitude magnetic anomalies (600–1200 gammas) are spatially associated with ferrobassalt at the Juan de Fuca Ridge–Blanco Fracture Zone intersection [Vogt and Byerly, 1976]. Vogt and Byerly postulated that the iron and titanium enrichment could result from fractional crystallization in a

subaxial conduit transporting magma longitudinally away from the Cobb hotspot and damming of the flow at fracture zones.

Incompatible element ratios are highly variable, both along the ridge and within single dredge hauls; most are intermediate between normal and enriched MORB [Lias and Rhodes, 1982]. The rare earth element (REE) patterns range from light REE depleted to light REE enriched with a dominance of fairly flat REE patterns [Schilling et al., 1982]. Sr isotopic ratios of basalt along the Juan de Fuca Ridge have a small range of values; the average ⁸⁷Sr/⁸⁶Sr ratio is 0.70249 ± 0.00014. Though subtle variations exist along strike, the Sr isotopic data do not show any systematic variation relative to the Cobb hotspot [Eaby et al., 1984].

STUDY AREA

The study area is a 12-km segment of the axial valley approximately 15 km north of the Juan de Fuca Ridge–Blanco Fracture Zone intersection (Figure 1). This area was selected in part due to its relatively simple tectonic setting. The ridge segment has a linear symmetrical axial rift valley and is located away from hotspots and actively propagating rifts. Preliminary descriptions of dredged lavas, dredged sulfides, and the geologic setting of the ridge segment are published elsewhere [Eaby and Clague, 1982; Koski et al., 1982; Normark et al., 1982; Morton et al., 1983]. Dredge locations and bathymetry of the study area are shown in Figure 2. The axial valley is extremely flat floored and approximately 1 km wide and 100 m deep.

Seismic reflection studies in the study area [Morton, 1984] show a weak reflector at approximately 2.3 km (0.8–1.0 s) depth interpreted as a shallow level magma reservoir. The low amplitude of the reflection suggests that the velocity contrast across the interface is small, gradational (i.e., partially crystalline mush), or both.

Deep-tow photographic surveys of the study area [Normark et al., 1982; Lichtman et al., 1983] indicate that lava in the axial valley is dominantly sheet and lobate flow forms, characteristic of high effusion rates (Figures 3a–3c), with rare pillows (Figure 3d), characteristic of low effusion rates. Most lava displays brilliant glassy reflections. The axial floor is smooth and unfissured outside a nearby continuous linear depression 5–25 m deep and 50–100 m wide that bisects the valley. Sediment cover within the axial valley is least (<25%) within the central zone of collapse features and increases toward the valley margin (>25%). Zn-rich massive sulfide deposits are aligned along the central bathymetric depression. Valley walls are dominated by pillow flows lacking glassy reflections and having 40–80% sediment cover. The inner valley walls, including a low terrace about 30 m above the valley floor, are formed by steep normal faults.

HOST LAVA

Petrography and Distribution

The recovered basalt samples from the axial rift valley are nearly aphyric with <1% phenocrysts; plagioclase (1–5 mm) >> augite (1–2 mm) > olivine (1–2 mm). The samples are all nonvesicular (<0.5%) glassy to microcrystalline basalt. Basalt morphology is dominantly sheet, lobate, and hollow lobate (blisters) flow fragments (90%) with minor amounts of pillow fragments (10%). The sheet and lobate flow fragments are commonly 4–8 cm thick and have thick glassy selvages on upper and lower surfaces. The upper glassy selvages have a maximum thickness of 13 mm. Palagonite thicknesses range from 0 to 2 μm, suggesting an age of <1000 years [Hekinian and Hoffert, 1975].

Fig. 2. Scientists aboard the U.S. Geological Survey research vessel *S. P. Lee* conducted acoustic transponder navigated dredging, deep-toward camera surveying, and water sampling during September 1981 (cruise L11-81-WF). Locations of dredge stations are shown with arrows indicating direction of movement of the dredge. Dredges 3, 8, and 16 sampled older material from the axial walls. Contour interval is 25 m.

Lava samples from the axial valley walls are dominantly pillow fragments texturally similar to samples from the axial valley. The outer glassy selvage has commonly spalled off leaving only a few millimeters of glass; palagonite thicknesses on original glass surfaces range from 5 to 16 μm , suggesting an age of 2000–5000 years. Three small porphyritic glass fragments recovered in dredge 8 are probably from the valley walls and are the only porphyritic samples collected in the study area. These flat glass chips contain 10% total phenocrysts with plagioclase $\gg$ augite $>$ olivine. The porphyritic glass chips have the thickest palagonite and Mn-oxide coatings of all the samples. Where the dredge path could have sampled both the walls and valley, our assignment of location of the samples is based on lava age and morphology of the sampled fragments.

Chemical Composition

The host lavas for the xenoliths are mid-ocean ridge ferrobasalt that exhibits a narrow range of compositional variation. Average microprobe analyses of glassy margins of samples from each dredge are listed in Table 1. A representative group of 16 X ray fluorescence whole-rock major and trace element analyses were selected from a larger set of 30 analyses and are listed in Table 2. The lavas from the axial valley are enriched in iron and titanium ($\text{FeO}^* = 11.6\text{--}12.2$ wt %, $\text{TiO}_2 = 1.8\text{--}1.9$ wt %, Mg number = 52.5–54.7) compared to normal MORB ($\text{FeO}^* < 10.0$ wt %, $\text{TiO}_2 < 1.5$ wt %). Basalt from the axial valley walls is also ferrobasalt but exhibits a slightly wider compositional range; the most (Mg number = 50.3) and least (Mg number = 60.4) differentiated lavas are from the valley walls.

Normative compositions of the nearly aphyric lavas range from slightly (up to 1.5%) olivine normative to slightly (up to 1.7%) quartz normative with the vast majority of the samples being slightly quartz normative. The porphyritic sample 8-2 is the least differentiated lava from the area and contains 3.7% normative olivine. There is a general trend of decreasing normative olivine and increasing normative quartz with increasing FeO/MgO ratios, but higher Na_2O values also force the norm toward higher normative olivine.

In general, increasing FeO^* , Na_2O , K_2O , TiO_2 , P_2O_5 , MnO , and SO_3 are coupled with decreasing MgO , Al_2O_3 , and CaO . SiO_2 concentrations show no systematic variation with increasing Mg number. Plots of TiO_2 versus FeO^*/MgO and CaO versus MgO illustrate these trends (Figures 4a and 4b); data from the Galapagos spreading center at approximate longitudes of 85° and 95°W [Byerly et al., 1976; Clague et al., 1981; Fornari et al., 1983] are plotted for comparison. The most striking aspect of the data set is the chemical homogeneity of samples from the axial valley. The 24 whole-rock X ray fluorescence analyses of lavas from the valley floor have an average analysis and standard deviation of $\text{SiO}_2 = 49.89 \pm 0.18$, $\text{Al}_2\text{O}_3 = 13.65 \pm 0.08$, $\text{Fe}_2\text{O}_3^* = 13.27 \pm 0.19$, $\text{MgO} = 6.87 \pm 0.11$, $\text{CaO} = 11.10 \pm 0.10$, $\text{Na}_2\text{O} = 2.53 \pm 0.12$, $\text{K}_2\text{O} = 0.18 \pm 0.02$, $\text{TiO}_2 = 1.86 \pm 0.04$, $\text{P}_2\text{O}_5 = 0.18 \pm 0.01$, and $\text{MnO} = 0.21 \pm 0.01$, which is statistically identical to the analytical precision.

Trace element abundances are listed with major element analyses in Table 2 and shown on MgO variation diagrams in Figure 5. The range in trace element concentrations (V 308–378, Cr 331–112, Ni 99.3–49.0, Zn 100–131, Ga 17.7–20.3, Rb 0.9–2.4, Sr 104.4–109.3, Y 32.1–53.3, Zr 96.9–187, Nb 2.9–6.4

Fig. 3. Deep-tow photographs of typical basalt morphologies found in the axial valley. (a) Low-relief sheet flows with light sediment cover which forms the dominant basalt morphology in the axial valley. (b) Collapsed hollow lobate sheet flow. (c) Lobate basalt flow. (d) Pillowed basalt flow.

ppm) are similar to those found in ferrobalt with comparable FeO^*/MgO from the Galapagos spreading center [Perfit *et al.*, 1983]. Increasing FeO^* and TiO_2 are accompanied by increasing concentrations of incompatible elements (V, Zr, Y, Zn, Nb) and decreasing concentrations of compatible elements (Ni, Cr).

Ratios of Zr/Nb for the 24 samples from the valley average 23.4 ± 1.1 and are toward the less depleted range for normal-MORB (N-MORB). Similarly, the valley samples have an average Zr/Y ratio of 3.24 ± 0.07 , an average Y/Nb ratio of 7.21 ± 0.4 , and an average $\text{P}_2\text{O}_5/\text{K}_2\text{O}$ ratio of 1.02 ± 0.05 (Figure 4c). These trace and minor element ratios are all within the ranges found for N-MORB elsewhere and are consistent with the low $^{87}\text{Sr}/^{86}\text{Sr}$ ratios (0.7026) reported for three samples from the valley floor of this segment of the Juan de Fuca Ridge [Eaby *et al.*, 1984]. The six analyzed samples from the valley walls, while more variable in major element composition than the 24 valley floor samples, have consistently higher $\text{P}_2\text{O}_5/\text{K}_2\text{O}$ ratios (1.16 ± 0.09), Zr/Nb ratios (28.4 ± 2.9), and Y/Nb ratio (8.9 ± 1.3).

Phenocrysts and Microphenocrysts

Plagioclase compositions are given in Tables 3a and 3b. Plagioclase phenocrysts and microphenocrysts are weakly zoned, skeletal to euhedral laths in form, and up to 5 mm in length. Plagioclase phenocrysts in sample 8-2 range in composition from An_{70} to An_{77} . The average composition of 46 microphenocryst analyses is $\text{An}_{74.5}$ with a range of An_{70} to An_{84} . Analyses with maximum and minimum An contents for several samples are listed in Table 3a.

Olivine occurs as euhedral, hopper, and skeletal microphenocrysts. Representative olivine microphenocryst compositions are given in Table 4a and shown in Figure 6. Olivine microphenocryst compositions range from Fo_{80} to Fo_{83} . The most magnesian olivine (Fo_{83}) occurs in sample 8-2.

Representative pyroxene analyses are given in Table 5 and shown on the pyroxene quadrilateral in Figure 6. Pyroxene compositions referenced in the text are expressed in terms of percent Wo, Fs, and En, where Wo is the atomic proportion $100\text{Ca}/(\text{Ca} + \text{Fe} + \text{Mg})$, $\text{Fs} = 100\text{Fe}/(\text{Ca} + \text{Fe} + \text{Mg})$, and $\text{En} = 100\text{Mg}/(\text{Ca} + \text{Fe} + \text{Mg})$. Quadrilateral components were calculated using the method of Lindsley and Andersen [1983] for comparison to Lindsley's [1983] graphical pyroxene geothermometer and are listed in Table 5. Phenocrysts of augite in sample 8-2 range in composition from $\text{Wo}_{40.1}\text{Fs}_{9.1}\text{En}_{50.8}$ to $\text{Wo}_{39.4}\text{Fs}_{10.0}\text{En}_{50.6}$. The Cr_2O_3 contents range from 0.60 to 1.27 wt % and are about a factor of 3 higher than Cr_2O_3 contents of pyroxene in the xenoliths. The phenocrysts have the most refractory compositions of all analyzed pyroxene. Composition of augite microphenocrysts in samples 22-34 and 26-13B range from $\text{Wo}_{40.5}\text{Fs}_{10.4}\text{En}_{49.1}$ to $\text{Wo}_{36.0}\text{Fs}_{11.5}\text{En}_{52.5}$. Compositions of phenocrysts and microphenocrysts in the host lava are similar to phenocryst compositions in Galapagos spreading center lavas [Perfit and Fornari, 1983] and Nazca plate lavas [Mazzullo and Bence, 1976].

Fractionation and Source Characteristics

Variations in major and trace element composition of ferrobalt from the southern Juan de Fuca Ridge are similar to the variations in lavas with comparable FeO^*/MgO ratio

TABLE 1. Averaged Major Element Microprobe Analyses of Basalt Glasses

	Dredge													
	3-6	8	11	12	4	3	23	5	17	22	26	16	16-6	13
Number of samples	(1)	(1)	(1)	(1)	(7)	(3)	(1)	(7)	(8)	(8)	(7)	(3)	(1)	(1)
Location	wall	wall	valley?	valley	valley	wall	valley	valley	valley	valley	valley	valley	wall?	wall
SiO ₂	50.8	50.6	50.8	50.8	50.6 ± 0.1	50.7 ± 0.6	51.2	50.5 ± 0.2	50.5 ± 0.3	50.7 ± 0.4	50.5 ± 0.4	51.0 ± 0.3	50.7	50.7
Al ₂ O ₃	14.4	14.3	14.1	14.3	14.2 ± 0.2	14.1 ± 0.2	14.1	14.1 ± 0.2	14.1 ± 0.2	14.0 ± 0.1	14.1 ± 0.1	14.0 ± 0.1	14.1	14.0
FeO*	11.6	11.3	12.1	12.1	12.1 ± 0.2	12.0 ± 0.1	11.9	12.1 ± 0.2	12.2 ± 0.1	12.3 ± 0.1	12.3 ± 0.1	12.4 ± 0.2	13.1	12.9
MgO	7.41	7.19	7.10	6.98	6.95 ± 0.08	6.85 ± 0.06	6.78	6.81 ± 0.16	6.84 ± 0.10	6.75 ± 0.10	6.76 ± 0.06	6.78 ± 0.05	6.62	6.26
CaO	11.9	12.4	11.4	11.7	11.3 ± 0.3	11.8 ± 0.3	11.1	11.3 ± 0.3	11.2 ± 0.3	11.2 ± 0.3	11.3 ± 0.2	11.4 ± 0.2	11.5	11.0
Na ₂ O	2.53	2.58	2.61	2.45	2.59 ± 0.08	2.48 ± 0.11	2.71	2.64 ± 0.08	2.58 ± 0.07	2.66 ± 0.04	2.65 ± 0.05	2.50 ± 0.05	2.52	2.64
K ₂ O	0.09	0.10	0.15	0.14	0.15 ± 0.01	0.16 ± 0.01	0.18	0.15 ± 0.01	0.16 ± 0.01	0.16 ± 0.01	0.15 ± 0.01	0.16 ± 0.00	0.14	0.19
TiO ₂	1.78	1.77	1.92	1.90	1.88 ± 0.05	1.89 ± 0.04	1.88	1.89 ± 0.05	1.90 ± 0.02	1.92 ± 0.01	1.92 ± 0.02	1.98 ± 0.04	2.14	2.14
P ₂ O ₅	0.14	0.17	0.18	0.16	0.17 ± 0.01	0.18 ± 0.01	0.22	0.18 ± 0.01	0.17 ± 0.01	0.18 ± 0.02	0.17 ± 0.01	0.18 ± 0.02	0.18	0.27
MnO	0.17	0.17	0.19	0.18	0.18 ± 0.01	0.18 ± 0.01	0.18	0.18 ± 0.01	0.18 ± 0.01	0.18 ± 0.01	0.19 ± 0.01	0.19 ± 0.01	0.19	0.19
SO ₃	0.14	0.14	0.16	0.16	0.15 ± 0.01	0.15 ± 0.01	0.14	0.15 ± 0.01	0.15 ± 0.01	0.15 ± 0.01	0.15 ± 0.01	0.15 ± 0.01	0.18	0.15
Total	101.0	100.7	100.7	100.9	100.2	100.5	100.4	100.0	100.0	100.2	100.2	100.7	101.4	100.4
Mg #†	55.90	55.67	53.86	53.37	53.25	53.05	52.96	52.66	52.70	52.11	52.08	52.10	50.02	49.07
FeO/MgO‡	1.40	1.42	1.53	1.56	1.57	1.58	1.58	1.60	1.60	1.64	1.64	1.64	1.78	1.85

Glasses were analyzed for major and minor elements with a three-channel ARL-EMX electron microprobe using U.S. Geological Survey standards of A-99 for Ti, Na, Si, and Fe; VG-2 and Indian Ocean glass 113716 for Al, Ca, and Mg; K-spar 2 for K; apatite for P; pyrrhotite for S; and fayalite for Mn. Sample current was 15 nA and counting time was 60 s. Six or more were analyzed and averaged to give a representative value for the sample. Matrix corrections were made using a Fram72 program [Beeson, 1967].

†Mg # is the atomic proportion (Mg/Mg + Fe).

‡FeO = 0.9 × FeO*.

TABLE 2. Representative XRF Major and Trace Element Analyses of Host Lava

	Sample															
	8-2	3-6	12-3	4-28	11-0	3-3	4-23	16-8	5-7	17-1	26-1	23-1	22-17	5-67	16-6	13-0
Location	wall	wall	valley	valley	valley?	wall	valley	valley	valley	valley	valley	valley	valley	valley	wall?	wall
SiO ₂	49.4	49.5	49.5	49.9	49.6	50.1	49.9	49.7	50.1	49.7	50.0	50.0	50.0	49.7	49.7	49.9
Al ₂ O ₃	14.5	13.9	13.7	13.7	13.7	13.6	13.6	13.5	13.6	13.6	13.7	13.7	13.6	13.7	13.4	13.5
Fe ₂ O ₃	11.6	12.5	12.9	12.9	13.1	13.2	13.3	13.3	13.5	13.3	13.4	13.5	13.4	13.4	14.0	13.8
MgO	8.01	7.45	7.09	7.00	6.98	6.89	6.94	6.80	6.87	6.78	6.78	6.80	6.77	6.74	6.58	6.32
CaO	11.9	11.7	11.3	11.2	11.1	11.2	11.1	11.1	11.0	11.0	11.0	11.1	11.0	11.0	11.1	10.7
Na ₂ O	2.35	2.31	2.38	2.50	2.47	2.51	2.47	2.36	2.71	2.60	2.53	2.58	2.57	2.69	2.44	2.56
K ₂ O	0.11	0.12	0.17	0.17	0.17	0.18	0.18	0.18	0.18	0.17	0.18	0.18	0.18	0.18	0.17	0.21
TiO ₂	1.50	1.69	1.84	1.79	1.84	1.84	1.87	1.92	1.88	1.87	1.88	1.88	1.88	1.89	2.06	2.11
P ₂ O ₅	0.13	0.15	0.17	0.18	0.17	0.18	0.18	0.18	0.18	0.18	0.17	0.19	0.19	0.18	0.20	0.26
MnO	0.21	0.20	0.21	0.21	0.22	0.21	0.21	0.20	0.22	0.21	0.21	0.22	0.21	0.21	0.22	0.23
Total	99.7	99.5	99.3	99.6	99.4	99.9	99.8	99.2	100.2	99.4	99.9	100.2	99.8	99.7	99.9	99.6
Mg #†	60.4	56.7	54.8	54.5	53.9	53.5	53.5	53.0	52.9	52.8	52.7	52.5	52.7	52.5	50.8	50.3
calculated FeO‡	9.36	10.2	10.4	10.4	10.6	10.7	10.8	10.8	10.9	10.8	10.8	10.9	10.9	10.9	11.4	11.2
FeO/MgO	1.17	1.37	1.47	1.49	1.52	1.55	1.56	1.59	1.59	1.59	1.59	1.60	1.61	1.62	1.73	1.77
V	308.	345.	373.	368.	375.	387.	365.	399.	361.	372.	369.	365.	364.	368.	398.	378.
Cr	331.	209.	169.	100.	97.	124.	94.8	108.	92.3	90.1	86.2	88.5	86.0	89.4	95.7	112.
Ni	99.	69.	60.	57.	56.	53.	56.	54.	49.	52.	50.	54.	50.	50.	51.	49.
Zn	100.	117.	115.	128.	119.	132.	118.	121.	119.	119.	120.	117.	119.	121.	127.	131.
Ga	17.7	18.9	19.0	18.4	18.6	18.8	18.4	18.8	20.5	18.6	18.5	19.5	19.6	18.9	19.1	20.3
Rb	0.9	0.8	1.6	1.7	1.6	1.4	2.0	1.9	1.4	1.8	1.4	1.2	1.7	2.1	1.3	2.4
Sr	104.4	87.4	103.5	112.4	109.9	109.8	109.8	114.0	115.5	114.1	113.7	114.4	114.2	114.3	99.2	109.3
Y	32.1	36.5	39.9	38.6	39.3	39.1	40.6	40.8	40.2	39.3	39.8	39.9	40.7	40.8	44.7	53.3
Zr	96.9	111.	125.	123.	128.	130.	133.	134.	131.	128.	130.	128.	133.	132.	139.	187.
Nb	2.9	4.1	5.9	5.3	5.3	5.0	5.5	6.2	5.5	5.7	5.5	5.4	5.2	5.5	4.8	6.4
Zr/Nb	33	27	21	24	24	26	24	22	24	23	24	24	26.	24.	29.	29.
Zr/Y	3.02	3.04	3.13	3.26	3.26	3.32	3.28	3.28	3.26	3.26	3.27	3.21	3.27	3.24	3.11	3.51
Y/Nb	11.1	8.90	6.76	7.28	7.42	7.82	7.38	6.58	7.31	6.89	7.24	7.39	7.83	7.42	9.31	8.33

Major element data of fresh microcrystalline samples were obtained by X ray fluorescence analysis (XRF) of fused glass discs, prepared by fusing the sample with a lanthanum-bearing lithium borate fusion mixture at the University of Massachusetts, Ronald B. Gilmore Memorial Laboratory. Trace elements (TiO₂, V, Cr, Ni, Zn, Ga, Rb, Sr, Y, Zr, and Nb) were determined by XRF analyses on pressed powder pellets. Detection limits and precision (in ppm) for the trace element analyses are 0.7 and 0.5 for Nb, 0.7 and 1 for Zr, 0.5 and 2 for Sr, 3 and 1 for Zn, 3 and 3 for Ni, 3 and 5 for Cr, 4 and 5 for V, and 0.005 and 0.02 (wt %) for TiO₂.

†Mg # = atomic proportion Mg/(Mg + Fe).

‡Calculated FeO = (0.9 FeO*).

Fig. 4. Variation diagrams for major and minor element compositions of host lava from the southern Juan de Fuca Ridge. Data from the Galapagos spreading center (GSC) are included for comparison. (a) Plot of TiO_2 versus FeO^*/MgO shows the narrow range in composition at the lower end of the ferrobalt field of the Juan de Fuca Ridge basalt data from this study superposed on the wide range in compositions from the Galapagos spreading center differentiation suite [Byerly *et al.*, 1976; Fornari *et al.*, 1983]. (b) Plot of CaO versus MgO shows the trend of decreasing CaO with decreasing MgO due to crystallization of olivine, clinopyroxene, and plagioclase. Juan de Fuca Ridge data from this study exhibit the same trends as the Galapagos spreading center data [Clague *et al.*, 1981; Fornari *et al.*, 1983] but are more limited in extent of differentiation. (c) Plot of P_2O_5 versus K_2O for Juan de Fuca Ridge data from this study compared to Galapagos spreading center data [Clague *et al.*, 1981]. The ratio of two incompatible elements should be indicative of the source regions. Data from Galapagos spreading center at 85°W, Galapagos spreading center at 95°W, and Juan de Fuca Ridge (this study) define linear but distinct trends suggesting differences in the source regions. Data from the axial valley form a tight cluster in Figures 4b and 4c. Data from the valley walls are more scattered.

from the Galapagos spreading center. Galapagos spreading center lavas from near 95°W and 86°W exhibit decreasing P_2O_5 , TiO_2 , V, and Zn contents at very high FeO^*/MgO ratios; these trends are consistent with late stage fractionation of apatite and titanomagnetite. Lavas from the southern Juan

de Fuca Ridge are more fractionated than normal MORB but show a more restricted compositional range than lavas from the Galapagos spreading center and do not appear to have reached saturation with titanomagnetite or apatite.

These major element trends for the Juan de Fuca samples can be modeled by fractionation of plagioclase, clinopyroxene, and lesser olivine as described by Clague and Bunch [1976]. Fractionation, however, may not be a unique solution. Mixing of relatively primitive magmas with extremely evolved magmas residing in the subaxial chamber can produce nearly identical major element geochemical trends. The role of mixing will be developed later in the paper.

The Sr isotopic data [Eaby *et al.*, 1984] and the trace and minor element ratios discussed above indicate that these lavas from the southern Juan de Fuca Ridge are derived from a relatively depleted source region. Several other samples from this section of the ridge have rare earth patterns that are depleted in light rare earth elements, as one might expect based on the other trace element and isotopic data. These samples have chondrite-normalized La/Sm of 0.67–0.75 [Schilling *et al.*, 1982], similar to the 95°W Galapagos spreading center source which has chondrite-normalized $\text{La}/\text{Sm} = 0.67$ [Clague *et al.*, 1981] and $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of approximately 0.7027–0.7028 [Verma and Schilling, 1982]. In contrast, the source region beneath 85°W on the Galapagos spreading center has chondrite-normalized $\text{La}/\text{Sm} = 0.46$ and $^{87}\text{Sr}/^{86}\text{Sr} = 0.7024$ –0.7025. We conclude that the source region beneath this section of the southern Juan de Fuca Ridge is comparably depleted to that at 95°W but is far less

Fig. 4c

Fig. 5. Trace element content as a function of MgO content for lava from the southern Juan de Fuca Ridge compared to data from the Galapagos spreading center [Clague *et al.*, 1981; Perfit *et al.*, 1983]. Lines are drawn by eye. Trace element contents of samples from the axial valley form a tight cluster, while trace element contents of samples from the axial walls define the extremes of the enclosed fields.

depleted than that at 85°W on the Galapagos spreading center. The source regions underlying the Juan de Fuca Ridge and near 95°W on the Galapagos spreading center, although comparably depleted in terms of La/Sm and $^{87}\text{Sr}/^{86}\text{Sr}$, are not

identical in that Juan de Fuca Ridge source has higher abundances of Sr and Na.

Differences in trace element ratios [Clague *et al.*, 1981; LeRoex *et al.*, 1983] for the axial valley and axial valley wall

samples indicate that the older valley wall lavas were derived from a more depleted source than the younger valley floor lavas. This observation suggests that the source regions for Juan de Fuca MORB may be heterogeneous on a time scale of thousands to tens of thousands of years.

XENOLITHS

Xenoliths comprise <1% of the dredge contents. They range in size from 1 cm³ (glomerocryst) to 240 cm³ (Figures 7a and 7b). Xenoliths commonly occur near the upper glassy selvage and along fractures of the sheet and lobate flow fragments. The common occurrence of xenoliths on fracture surfaces suggests that a mechanism such as differential contraction during cooling between the host lava and the xenolith created strain that induced fracturing.

On the basis of texture and mineralogy, xenoliths have been subdivided into three types: type I: medium-grained; plagioclase + olivine + glass; type II: medium-grained; plagioclase + augite ± olivine + glass; and type III: medium- to coarse-grained; plagioclase + augite + olivine ± inverted pigeonite ± Fe-Ti oxides + glass. All xenoliths have small volumes (<1%) of sulfide globules.

The following sections describe the textures and phase chemistry of each xenolith type. Compositions of plagioclase, olivine, pyroxene, oxides, and interstitial glass are given in Tables 3, 4, 5, 6, and 7 and shown in Figure 6, 8, and 10. Glass inclusion data (Tables 8 and 9 and Figure 11) will be discussed separately. Modal compositions of xenoliths are shown in Table 10.

Type I Xenoliths

Three small (<1 cm³) olivine plus plagioclase xenoliths were found. Xenoliths 5-17, 17-7, and 17-12 are open-textured, medium-grained hypidiomorphic interstitial gabbros. Type I xenoliths are a loose cluster of crystals with olivine (up to 3 mm in length) subophitically enclosing or interstitial to euhedral plagioclase (up to 3 mm). Mineral grains are fresh and have not reacted with the surrounding melt. A rim of variolitic crystallization formed upon quenching around plagioclase.

Plagioclase occurs as weakly zoned laths having the same compositional range as the microphenocrysts in the lavas. The compositional range in sample 17-12 is An₆₈ to An₇₃ and in sample 5-17 is An₆₅ to An₇₃. The average composition of 24 analyses is An_{72.8}, slightly more sodic than the average microphenocryst composition.

Olivine compositions within each sample are homogeneous; the range of olivine compositions in sample 17-7 is Fo_{78.1} to Fo_{78.8} and in sample 17-12 is Fo_{78.5} to Fo_{78.9}. These compositions are intermediate between compositions of microphenocrysts and olivine in xenolith types II and III. The range of compositions in sample 5-17 is Fo_{71.5} to Fo_{72.5}, which are more like those for olivine in type II xenoliths. NiO contents in type I olivine were not analyzed.

Interstitial glass (25–30 modal %) is compositionally similar to the surrounding host glass. FeO/MgO (using FeO = FeO* × 0.9) is 1.62 for the interstitial glass similar to 1.60 for the host lava. Interstitial glass contains 1.92 wt % TiO₂, 2.58 wt % Na₂O, and 0.18 wt % P₂O₅.

Type II Xenoliths

Type II xenoliths are composed dominantly of plagioclase, augite, and glass. Minor volumes of olivine are present. Samples 17-1 and 5-0 (Figure 7a) are representative of type II xenoliths and are described in detail below.

Sample 17-1, the largest open-textured xenolith (240 cm³), forms a toe of a sheet flow fragment and is texturally similar

to glassy, highly porphyritic lava (Figure 7c). Euhedral to subhedral augite, and rarely olivine, poikilitically enclose the plagioclase laths to form subophitic clusters. Abundant glass inclusions indicate rapid crystal growth. Glass adjacent to plagioclase and augite is opaque brown due to variolitic crystallization. Interstitial glass comprises 50 modal % of the xenolith and is pale translucent brown away from the crystal grain boundaries. Flow of melt around crystals is indicated by variations in color and incipient crystallization (Figure 7d). The absence of embayed crystals or mineral zonation suggests that glass and minerals have not reacted.

Sample 5-0 is a medium to coarse grained hypidiomorphic granular gabbro (Figure 7e). Euhedral plagioclase laths are similar in size and morphology to those in 17-1 but exhibit slight zoning. Augite oikocrysts are up to 1 cm across. Interstitial glass (22 modal %) is opaque due to incipient crystallization. There is no evidence of interaction between xenolith minerals and the surrounding lava; mineral-glass interfaces are sharp with euhedral crystal forms.

The majority of the type II xenoliths consist of subophitic to interstitial clusters of plagioclase + augite + glass, smaller than (<1 cm³) but similar to xenolith 5-0. Rare inclusions of olivine in augite and vice versa occur, but generally these phases are not intergrown.

Plagioclase laths in type II xenoliths range in composition from An_{61.5} to An_{74.0} and have an average composition of An_{68.6}. Plagioclase in open-textured xenoliths (i.e., 17-1) is weakly zoned and shows little variation within a single xenolith. Plagioclase compositions from more crystalline xenoliths have slight normal oscillatory zoning. Early formed plagioclase laths, which are poikilitically enclosed by augite, tend to be more Ca-rich than small tabular grains growing adjacent to interstitial glass or at the xenolith-host lava boundary.

Olivine in type II xenoliths occurs as small discrete grains at the edge of the xenolith in samples 5-0 and 5-77 or as subhedral to anhedral grains subophitically enclosing small euhedral plagioclase laths in sample 5-1. These olivine grains are unzoned and range in composition from Fo_{72.8} to Fo_{73.2}.

Pyroxene in type II xenoliths occurs as homogeneous oikocrysts and more rarely as small discrete grains in the interstitial glass. The dominant compositional trend is one of decreasing Mg, Ca, and Cr and increasing Ti with increasing Fe content. Oikocrysts range in composition from Wo_{41.5}Fs_{8.8}En_{49.7} (sample 4-7) to Wo_{40.0}Fs_{12.3}En_{47.7} (sample 17-65). Analyses 11 and 14 in Table 5 are typical of most of the oikocrysts. An Fe-enrichment trend is also observed between oikocrysts and small augite grains in the interstitial glass or near the xenolith-host lava boundary. Analyses of pyroxenes from sample 5-1 (analyses 12 and 13 in Table 5) show that the oikocryst has 100 × Fe/(Fe + Mg) ratio of 15.9, while a small grain in the interstitial glass has a ratio of 21.5. The more differentiated composition of late forming pyroxene is also observed in samples 5-0 and 5-13.

Interstitial glass compositions in type II xenoliths define Fe-enrichment trends with increased crystallinity of the xenoliths. Compositions range from similar to the host lava (FeO/MgO (using FeO = FeO* × 0.9) ratio is 1.67) to more differentiated than the host lava (FeO/MgO ratio is 2.59). TiO₂ content ranges from 1.90 to 2.65 wt %. Na₂O content ranges from 2.67 to 3.28 wt %. P₂O₅ content ranges from 0.20 to 0.33 wt %. K₂O content ranges from 0.18 to 0.33 wt %. Chemical variation trends defined by the interstitial glass are similar to trends observed in the host glass but extend to more evolved compositions. Figures 8a and 8b show increasing TiO₂ with decreasing Mg number and increasing P₂O₅ and K₂O at essentially the same P₂O₅/K₂O ratio.

Type III Xenoliths

Type III xenoliths are generally coarser-grained, more crystalline, and mineralogically more evolved than types I and II. The most crystalline type III xenoliths are characterized by zoning in plagioclase and augite, a wide range of grain sizes, intricate intergrowth textures, and random mineral grain orientation. Type III xenoliths have reacted with the host melt more than the other xenolith types. Samples 26-1A, 5-7A, 5-7B, and 5-8B are typical of type III xenoliths (Figures 7b, 7f, 7g, and 7h).

Sample 26-1A is a medium-grained hypidiomorphic granular intersertal gabbro. Grain sizes range from 0.1 to 6 mm for plagioclase, 0.3 to 3 mm for olivine, and 0.2 to 5 mm for augite. The texture is more equigranular than other type III xenoliths. Mineral grains have no preferred orientation. Plagioclase is subhedral to euhedral. Larger grains exhibit slight optical zonation. Although normal zonation is most

common, rare oscillatory zonation is also present. Olivine and augite appear to have crystallized simultaneously. Crystal-mesostasis interfaces are sharp.

Samples 5-7A and 5-8B are coarse-grained hypidiomorphic granular olivine gabbro. In sample 5-7A olivine is the main oikocryst phase and occurs as large (5 mm across), optically continuous subhedral grains. Olivine and augite subophitically enclose plagioclase but do not enclose each other. Larger plagioclase grains are subhedral because of impingement of co-crystallizing olivine. Only 1 modal % interstitial glass remains. In sample 5-8B, augite is the main oikocryst phase which subophitically encloses plagioclase (up to 1 cm in length). Augite and plagioclase exhibit intricate intergrowth textures (Figure 9a), indicating simultaneous growth. Normal and oscillatory zoning occurs in the larger plagioclase grains. In 5-7A and 5-8B the larger augite grains exhibit optical zoning. Interstitial glass comprises 3 modal %. Augite has no exsolution lamellae of orthopyroxene, but minute exsolution

TABLE 3a. Plagioclase Analyses

	Microphenocrysts					Type I		Type II						
	1	2	3	4	5	6	7	8	9	10	11	12	13	14
Sample	4-35	4-35	8-2	8-2	5-7A	17-12	17-12	5-0	5-0	5-0	5-0	5-0	5-13	5-13
Comment†	max	min	max	min		max	min	core	midway	rim	small in gm	small in aug	small in gm	small in aug
SiO ₂	47.6	51.0	49.4	51.9	50.0	47.4	51.1	50.8	50.8	51.0	51.5	50.4	49.4	50.2
Al ₂ O ₃	32.7	30.6	31.6	29.0	31.3	32.8	30.4	31.1	30.9	31.2	30.4	31.1	30.7	30.5
FeO	0.38	0.51	0.43	0.48	0.52	0.44	0.49	0.41	0.52	0.65	0.54	0.53	0.75	0.61
CaO	17.4	14.7	16.0	14.5	14.9	17.9	13.9	14.2	14.0	14.3	13.6	14.4	14.5	13.9
NaO	1.79	3.27	2.57	3.43	3.11	1.60	3.68	3.55	3.66	3.56	3.91	3.41	3.24	3.40
K ₂ O	...	...	...	...	0.04	...	...	0.03	0.02	0.04	0.03	0.04	0.02	0.04
Total	100.9	100.2	100.0	100.3	99.9	100.2	99.6	100.1	99.9	100.8	100.0	99.9	98.6	98.6
Cations per Formula Unit														
Si	2.192	2.326	2.263	2.382	2.290	2.180	2.340	2.316	2.321	2.312	2.348	2.302	2.293	2.322
Al	1.777	1.644	1.707	1.568	1.687	1.779	1.639	1.670	1.661	1.667	1.631	1.677	1.679	1.661
Fe	0.014	0.019	0.017	0.019	0.020	0.017	0.019	0.016	0.020	0.025	0.021	0.020	0.029	0.024
Ca	0.858	0.718	0.785	0.714	0.731	0.882	0.682	0.691	0.684	0.693	0.664	0.707	0.719	0.687
Na	0.160	0.290	0.226	0.303	0.276	0.144	0.324	0.313	0.324	0.313	0.346	0.303	0.292	0.305
K	...	...	...	...	0.002	...	...	0.001	0.001	0.002	0.002	0.002	0.001	0.002
Total	5.001	4.997	4.998	4.986	5.006	5.002	5.004	5.007	5.011	5.012	5.012	5.011	5.013	5.001
An %	84.3	71.2	77.6	70.2	72.5	86.0	67.8	68.8	67.8	68.8	65.6	69.9	71.0	69.1
Type II														
	15	16	17	18	19	20	21	22	23	24	25	26	27	28
Sample	5-77	5-77	17-1	17-1	17-1	17-1	17-4	17-4	17-4	17-65	17-65	17-65	17-66	17-66
Comment†	large in gm	small in gm	large in gl	small in gl	large in aug	small in gl	core in aug	small in aug	small in aug	small in aug	small in gm	small in aug	small in gl	small in gl
SiO ₂	50.7	50.4	50.6	50.1	50.4	49.9	50.1	49.5	49.1	51.3	52.3	51.5	51.2	50.5
Al ₂ O ₃	29.8	30.0	29.9	32.1	30.2	30.4	30.9	30.7	31.0	29.6	28.9	29.5	30.6	30.6
FeO	0.60	0.68	0.58	0.60	0.62	0.53	0.55	0.60	0.60	0.56	0.60	0.57	0.59	0.56
CaO	14.3	13.7	13.4	14.6	13.7	14.1	14.1	14.9	14.9	13.0	12.7	12.9	13.6	14.3
Na ₂ O	3.25	3.57	3.67	3.19	3.65	3.38	3.35	2.98	2.87	3.95	4.34	4.14	3.58	3.30
K ₂ O	0.04	0.04	0.07	0.03	0.02	0.03	0.03	0.03	0.03	0.05	0.05	0.05	0.03	0.04
Total	98.6	98.4	98.2	100.6	98.6	98.4	99.0	98.7	98.5	98.5	98.9	98.6	99.6	99.3
Cations per Formula Unit														
Si	2.344	2.338	2.348	2.273	2.332	2.316	2.307	2.294	2.281	2.369	2.405	2.374	2.340	2.319
Al	1.622	1.639	1.632	1.719	1.647	1.662	1.679	1.678	1.696	1.611	1.565	1.603	1.647	1.657
Fe	0.023	0.027	0.023	0.023	0.024	0.020	0.021	0.023	0.023	0.022	0.023	0.022	0.022	0.022
Ca	0.708	0.677	0.666	0.711	0.677	0.701	0.696	0.738	0.740	0.645	0.624	0.639	0.668	0.706
Na	0.291	0.320	0.330	0.281	0.327	0.304	0.299	0.268	0.258	0.354	0.387	0.370	0.317	0.294
K	0.002	0.002	0.004	0.002	0.001	0.002	0.002	0.002	0.002	0.003	0.003	0.003	0.002	0.002
Total	4.990	5.003	5.003	5.009	5.008	5.005	5.004	5.003	5.000	5.004	5.007	5.007	4.996	5.000
An %	70.7	67.8	66.6	71.5	67.4	69.6	69.8	73.2	74.0	64.4	61.5	63.1	67.7	70.5

TABLE 3a. (continued)

	Type III												
	29	30	31	32	33	34	35	36	37	38	39	40	41
Sample Comment†	17-66 small in gl	17-69 lg in gm	17-69 small in gm	5-7A core large	5-7A small center	5-7A small edge	5-7A core edge	5-7A core center	5-8B core center	5-8B rim center	16-7A small in gl	26-1A small in gl	26-1A small in aug
SiO ₂	51.1	50.5	50.3	50.8	50.5	52.9	49.7	49.9	51.1	52.9	55.3	50.4	49.0
Al ₂ O ₃	30.5	30.2	30.5	30.9	31.2	28.1	30.6	30.4	30.8	29.1	27.5	29.9	31.1
FeO	0.57	0.54	0.57	0.63	0.43	0.67	0.62	0.64	0.60	0.68	0.66	0.66	0.65
CaO	13.9	13.9	14.2	14.3	14.4	11.8	14.1	14.0	13.9	11.9	12.1	13.2	14.4
Na ₂ O	3.48	3.53	3.27	3.53	3.51	4.62	3.27	3.46	3.61	4.60	4.78	3.93	3.05
K ₂ O	0.04	0.05	0.04	0.03	0.03	0.17	0.06	0.04	0.04	0.09	0.08	0.05	0.07
Total	99.6	98.7	98.9	100.2	100.1	98.3	98.4	98.4	100.0	99.3	100.4	98.1	98.3
	Cations per Formula Unit												
Si	2.337	2.332	2.322	2.317	2.303	2.441	2.307	2.317	2.328	2.416	2.495	2.341	2.280
Al	1.644	1.645	1.658	1.658	1.679	1.531	1.675	1.658	1.655	1.568	1.461	1.639	1.706
Fe	0.022	0.021	0.022	0.024	0.016	0.026	0.024	0.024	0.023	0.026	0.025	0.025	0.025
Ca	0.683	0.688	0.700	0.699	0.702	0.585	0.701	0.699	0.679	0.583	0.583	0.655	0.716
Na	0.308	0.316	0.293	0.312	0.311	0.413	0.295	0.312	0.319	0.408	0.418	0.354	0.275
K	0.002	0.003	0.002	0.002	0.002	0.010	0.003	0.002	0.002	0.005	0.005	0.003	0.004
Total	4.996	5.005	4.997	5.012	5.013	5.006	5.005	5.012	5.006	5.006	4.987	5.017	5.006
An %	68.8	68.3	70.5	69.0	69.2	58.0	70.2	69.0	67.9	58.5	58.0	64.7	72.0

†max, min refer to maximum and minimum An content; core, rim refer to location within large zoned grains; center, edge refer to location within xenolith; gm = groundmass; aug = augite; gl = glass; large = >4 mm; small = <2 mm. Microprobe data for plagioclase were obtained using an ARL-EMX three-channel microprobe and reduced using the matrix correction program of Fram 72 [Beeson, 1967]. U.S. Geological Survey standards used in analysis of plagioclase were Amelia Albite for Na; Crystal Bay Bytownite for Al, Si, Ca; K-spar 2 for K; and Fayalite for Fe. Sample current was 15 nA and counting time was 60 s.

lamellae of oxides occur along grain boundaries. Partially inverted pigeonite occurs as an interstitial phase; remnant cores of pigeonite are surrounded by coalesced blebs of augite in a host of orthopyroxene. Figures 9b, 9c, and 9d shows the blebby exsolution texture. Anhedral titanomagnetite and ilmenite occur in the interstitial glass and as isolated interstitial phases (Figures 9e and 9f). Fe-Ti oxides do not show optical exsolution.

The presence of subtle planar extinction discontinuities in olivine and melt-filled fractures that crosscut mineral grains indicates that these xenoliths have been deformed (Figures 9g, 9h, and 9i). The mesostasis-xenolith interface, along both the xenolith edge and melt-filled fractures, is more jagged indicating reaction with the melt. Plagioclase appears to have reacted with the host melt more than augite or olivine and commonly has a sawtooth appearance caused by fingers of melt intruding the plagioclase (Figure 9h).

TABLE 3b. Summary of Plagioclase Compositions

	Micro- phenocrysts	Xenoliths		
		I	II	III
Number analyzed	(46)	(24)	(24)	(12)
SiO ₂	49.7	50.0	50.6	51.3
Al ₂ O ₃	31.1	30.9	30.5	30.0
FeO	0.53	0.55	0.58	0.62
CaO	15.6	15.0	14.0	13.4
Na ₂ O	2.92	3.10	3.51	3.84
K ₂ O	...	...	0.04	0.07
Total	99.9	99.6	99.2	99.2
Average An %	74.5	72.8	68.6	65.7
Range An %	70-84	68-86	61-74	58-72

Samples 5-7B and 16-7A consist dominantly of plagioclase and augite in a loose cluster of crystals. Pigeonite, found only during microprobe analysis, has not inverted and is optically continuous with adjacent augite. Minerals in these two samples have moderately differentiated compositions and exhibit zoning and evidence for reaction with the host melt. Zones of abundant glass inclusions occur between augite with different extinction angles and were probably trapped between intergrowing crystals.

Olivine in type III xenoliths is coarser-grained and more iron-rich than microphenocrysts in the host lava and olivine in xenolith types I and II. The average composition is Fo_{69.6}. Olivine in samples 26-1A occurs as small (<1 mm) discrete grains with an average composition of Fo_{71.7}. Olivine in sample 17-3 occurs as two small grains at the edge of the xenolith with composition of Fo_{69.9}. Olivine in sample 5-7A occurs as large (5 mm) optically continuous subhedral grains and has the widest range in composition. Compositions of the cores of the large olivine grains range from Fo_{74.3} to Fo_{74.8}. One olivine has a slight zonation from core (Fo_{72.3}) to rim (Fo_{71.0}). The most iron-rich olivine (Fo_{63.7}) occurs in sample 16-7B, a xenolith similar to 5-7A. CaO-contents range from 0.23 to 0.33 wt % and are lower than CaO contents of microphenocrysts and olivine in type II xenoliths.

Plagioclase in type III xenoliths is coarser-grained and more complexly zoned than plagioclase in type I and II xenoliths. There is broad compositional overlap between type II and III plagioclase. The average composition of 10 plagioclase analyses is An_{65.7}. In general, increasing iron enrichment in the interstitial glass is accompanied by decreasing An content in the plagioclase.

Plagioclase in sample 26-1A ranges in composition from An_{64.8} to An_{72.0}. Two large plagioclase grains enclosed in olivine in sample 5-7A have An contents of 69.0 and 70.2%.

TABLE 4a. Representative Olivine Analyses

	Microphenocrysts				Type I			Type II			Type III				
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
Sample Number analyzed†	5-7 (3)	8-2 (3)	17-1 (1)	26-1 (1)	5-17 (4)	17-7 (3)	17-12 (2)	5-1 (3)	5-77 (2)	5-0 (3)	5-7A (1)	5-7A (1)	17-3 (3)	26-1A (3)	16-7B (3)
SiO ₂	39.5	39.8	40.1	39.7	39.8	39.3	40.3	37.2	37.0	36.5	37.5	38.9	36.7	37.1	37.6
FeO	18.6	15.7	17.5	17.5	24.5	19.8	19.2	24.9	24.9	24.5	23.2	26.3	27.4	25.9	31.0
MnO	0.30	0.26	0.29	0.28	0.37	0.30	0.28	0.39	0.39	0.38	0.36	0.40	0.44	0.40	0.44
MgO	41.8	43.6	42.0	41.6	35.3	40.4	39.8	37.3	37.5	37.6	38.7	36.2	35.5	36.5	30.5
CaO	0.44	0.46	0.55	0.52	...	...	...	0.35	0.34	0.36	0.33	0.24	0.33	0.33	...
NiO	0.16	0.16	0.14	0.13	...	...	...	0.15	0.17	0.17	0.19	0.19	0.18	0.19	0.08
Total	100.8	100.0	100.6	99.7	100.1	99.9	99.6	100.2	100.3	99.5	100.3	100.2	100.6	100.4	99.6
Cations per Formula Unit															
Si	1.002	1.005	1.013	1.012	1.040	1.010	1.031	0.980	0.977	0.971	0.980	0.980	0.978	0.981	1.023
Fe	0.394	0.331	0.370	0.373	0.536	0.426	0.411	0.549	0.548	0.545	0.508	0.585	0.608	0.571	0.704
Mn	0.006	0.006	0.006	0.006	0.008	0.007	0.006	0.009	0.009	0.009	0.008	0.009	0.010	0.009	0.010
Mg	1.580	1.638	1.580	1.580	1.375	1.545	1.519	1.468	1.478	1.492	1.511	1.435	1.413	1.446	1.237
Ca	0.012	0.013	0.015	0.014	...	...	...	0.010	0.010	0.010	0.009	0.007	0.009	0.009	...
Ni	0.003	0.003	0.003	0.003	...	...	...	0.004	0.004	0.004	0.004	0.004	0.004	0.004	0.002
Total	2.997	2.996	2.987	2.988	2.961	3.010	2.969	3.020	3.026	3.031	3.020	3.020	3.022	3.020	2.976
Fo%	80.0	83.2	81.0	80.9	72.0	78.4	78.7	72.8	72.9	73.2	74.8	71.0	69.9	71.7	63.7
Olivine-Melt Equilibria Calculations [Roeder, 1974]‡															
(MgO/FeO) ¹ *	0.67	0.83	0.72	0.71	0.43	0.61	0.62	0.45	...	0.46	0.50	0.41	0.39	0.43	0.30
D	0.25	0.25	0.24	0.23	...	0.27	0.26	0.27	...	0.28	...	...	0.26	0.29	...
FeO/FeO*	0.84	0.76	0.78	0.77	...	0.90	0.89	0.89	...	0.93	...	...	0.87	0.98	...

Microphenocrysts from host lava; Type I, II, and III refer to xenoliths. Olivine data were obtained using an ARL-SEM automated nine-channel microprobe and reduced using the matrix correction program of Bence and Albee [1968]. Standards used in the analysis of olivine were San Carlos olivine for Mg and Si; fayalite for Fe and Mn; and a synthetic Ni-doped diopside. Sample current was 20 nA. Backgrounds for NiO were calculated by hand using a mean atomic number method based on synthetic Ni-free fayalite and Fo_{97.5} from marble.

†Averaged analyses vary less than 0.5 Fo% (1σ).

‡Calculations use host glass compositions with microphenocrysts and intersertal glass compositions with xenoliths.

Plagioclase in sample 5-8B shows complex zoning; a large, normally zoned plagioclase has a core composition of An_{67.9} and rim composition of An_{58.5}. Small plagioclase grains in the interstitial glass in samples 5-7A and 16-7A have the most sodic compositions (An_{58.0}) and reflect the more differentiated composition of the residual liquid. Intergrowths of plagioclase grains commonly have zones of abundant glass inclusions at the grain boundaries.

Pyroxene in type III xenoliths occurs as augite, magnesian pigeonite, and orthopyroxene. Sample 26-1a contains only augite that is fairly equigranular and compositionally homogeneous. Compositions range from Wo_{40.2}Fs_{10.3}En_{49.5} to Wo_{39.5}Fs_{11.8}En_{48.7}. Large augite oikocrysts in samples 5-7A

and 5-8B become more iron-rich toward the margins. Cores of augite oikocrysts have a compositional range of Wo_{40.5}Fs_{9.7}En_{49.7} to Wo_{39.2}Fs_{11.9}En_{49.0} with augite in 5-8B slightly more iron-rich than augite in 5-7A. Core compositions are similar to augite compositions in 26-1A. Points near the edge of an augite oikocryst in sample 5-7A are Wo_{37.2}Fs_{13.3}En_{49.5} and in sample 5-8B are Wo_{38.5}Fs_{17.4}En_{44.1}.

Augite in sample 5-7B is iron-rich with a core composition of Wo_{35.0}Fs_{16.8}En_{48.2} and a margin composition of Wo_{33.8}Fs_{17.6}En_{48.6} (analysis 7 in Table 5). A small grain of pigeonite in sample 5-7b has a composition of Wo_{8.5}Fs_{26.0}En_{65.5} and is optically continuous with the adjacent augite. Augite and pigeonite compositions in sample 16-7A are similar to those in sample 5-7B. A similar occurrence of augite and pigeonite as single crystals and as alternating sectors in complexly zoned phenocrysts has been reported by Byerly [1980] in evolved basalt from the Galapagos spreading center near 95°W.

Primary pigeonite, partially inverted to host-lamellae pairs of orthopyroxene and augite, occurs as an interstitial phase in 5-7A and 5-8B. Remnant cores of pigeonite have a compositional range of Wo_{6.1}Fs_{23.7}En_{70.2} to Wo_{9.1}Fs_{25.2}En_{65.7}. These compositions are less iron-rich than pigeonite in sample 5-7B. Surrounding these cores of pigeonite are areas of host orthopyroxene with exsolved augite forming blebs microns to tens of microns in width. Augite blebs have compositions of Wo_{39.0}Fs_{12.8}En_{48.2} to Wo_{37.2}Fs_{16.3}En_{46.4} in sample 5-7A and Wo_{36.9}Fs_{14.9}En_{48.2} to Wo_{37.6}Fs_{18.7}En_{43.6} (analysis 19 in Table 5) in sample 5-8B. The exsolved augite has iron contents similar to the margins of the large, zoned augite oikocrysts.

Host orthopyroxene commonly has patchy extinction.

TABLE 4b. Summary of Olivine Compositions

	Micro-phenocrysts	Xenoliths		
		I	II	III
Number analyzed	(17)	(3)	(3)	(4)
SiO ₂	39.8	39.8	36.9	37.3
FeO	17.5	21.2	24.8	27.2
MnO	0.28	0.32	0.39	0.42
MgO	42.5	38.5	37.5	35.0
CaO	0.49	...	0.35	0.31
NiO	0.15	...	0.16	0.19
Total	100.7	99.8	100.1	100.4
Average Fo %	81.2	76.4	73.0	69.6
Range Fo %	79-83	72-79	72-74	64-75

Fig. 6. Pyroxene quadrilateral and histograms of Fo content in olivine and An content in plagioclase showing the sympathetic compositional variation of coexisting mineral phases in the host lava and xenoliths. The pyroxene quadrilateral on the left shows the large variation of CaO content at relatively constant FeO/MgO ratios for the phenocrysts (sample 8-2) and microphenocrysts related to rapid cooling. The pyroxene quadrilateral on the right shows the iron enrichment trend found in types II and III xenoliths. Magnesium pigeonite and orthopyroxene occur only in type III xenoliths. Tie lines connect pyroxene analyses from inverted pigeonite in samples 5-7A, 5-7B, and 5-8A. Histogram of Fo content in olivine shows the iron enrichment trend in olivine from host lava and xenoliths. Plagioclase compositions show much more overlap than pyroxene or olivine. In general, mineral compositions in types II and III xenoliths overlap, but minerals in type III xenoliths have a wider range in composition, extending to more differentiated compositions.

Ranges in composition in sample 5-7A are $Wo_{3.9}Fs_{20.7}En_{75.4}$ (analysis 22 in Table 5) to $Wo_{4.6}Fs_{27.2}En_{68.2}$ (analysis 24 in Table 5) and in sample 5-8B are $Wo_{4.0}Fs_{24.1}En_{71.9}$ to $Wo_{4.3}Fs_{32.4}En_{63.3}$ (analysis 26 in Table 5). One discrete grain of orthopyroxene ($Wo_{4.4}Fs_{26.9}En_{68.7}$) occurs in sample 5-7A. This orthopyroxene grain is adjacent to augite and probably represents complete migration of exsolved augite to the margin of an inverted pigeonite. Much of the scatter in the host-lamellae pair data occurs in the direction of the tie lines and probably reflects overlap of the electron beam on fine exsolution lamellae.

Minor element distribution in host-lamellae pairs are similar to contents in other terrestrial and lunar inverted pigeonites [DeVore, 1955, 1957; Brown, 1957; Carstens, 1958; Ghose et al., 1973; Walker et al., 1973]. Ti, Al, and Cr are partitioned into the high-Ca phase, while Mn is partitioned into the low-Ca phase (Figure 10).

Representative analyses of Fe-Ti oxides are given in Table 6. Table 6 includes calculation methods of Stormer [1983], Anderson [1968], Carmichael [1967], and Lindsley and Spencer [1982] for ulvöspinel and ilmenite components, temperature, and oxygen fugacity to allow direct comparison of these data to data from other studies. Due to the high minor element contents (Mn, Mg, Al, and V) in the xenolith oxides, the recalculation method of Stormer is used in the text.

Irregular grains of Fe-Ti oxides occur within interstitial glass and in interstitial locations in samples 5-7A and 5-8B. Titanomagnetite and ilmenite occur as discrete or composite grains. Titanomagnetite compositions are fairly homogenous; ulvöspinel component varies from 59.8 to 63.0%. Con-

centrations of minor elements (Al_2O_3 , 2.51–3.36 wt %; V_2O_5 , 0.88–1.30 wt %; MnO, 0.38–0.47 wt %; MgO, 2.82–4.37 wt %) are similar to minor element contents in titanomagnetite phenocrysts in ferrobalt and andesite from the Galapagos Rift [Perfit and Fornari, 1983].

Ilmenite is more abundant than titanomagnetite and exhibits a wider range in composition. MgO content in ilmenite is higher than in coexisting titanomagnetite. Ilmenite compositions have been subdivided into ilmenite and high-Mg ilmenite. Ilmenite (<7.0 wt % MgO) occurs as isolated grains or as composite grains with titanomagnetite. Ilmenite component varies from 87.8 to 89.4%, MgO contents range from 3.67 to 6.89 wt %, and Al_2O_3 ranges from 0.32 to 0.56 wt %. These compositions are similar to, but with slightly higher MgO contents than, ilmenite from Galapagos Rift lavas [Perfit and Fornari, 1983]. High-Mg ilmenite has MgO contents of 7.82–8.44 wt %. High MgO content is accompanied by higher TiO_2 and lower Fe_2O_3 and FeO contents. Ilmenite component ranges from 89.8 to 93.8%. High-Mg ilmenite occurs only as isolated crystals, but there is no apparent correlation between size of grain; location of grain within the xenolith; surrounding silicate phases; presence of adjacent vapor bubble, sulphide phase, or glass; on ilmenite composition.

Interstitial glasses are more evolved and have a wider compositional range than interstitial glasses in types I or II. FeO/MgO ratios (with $FeO = FeO^* \times 0.9$) range from 2.14 to 3.38 (see Table 7). Ranges in minor element contents are $TiO_2 = 2.31$ –4.71 wt %, $Na_2O = 2.90$ –3.91 wt %, $K_2O = 0.21$ –0.69 wt %, and $P_2O_5 = 0.24$ –1.03 wt %. CaO/Al_2O_3 ratios vary from 0.62 to 0.76. Interstitial glass compositions

for sample 5-7A are not reported because of beam overlap on incipiently crystallizing titanomagnetite and apatite.

Glass Inclusions

Glass inclusions are present in nearly every mineral phase and range in size from less than a micron across to more than a hundred microns in length. They usually exhibit negative-crystal shapes relative to the host crystal showing that they represent small volumes of magma entrapped during the growth of the crystal [Sobolev and Kostyuk, 1975; Roedder, 1979].

Compositions of glass inclusions are presented in Table 8 and are plotted on Figure 11a. The wide scatter in the glass inclusion data is related to the type of xenolith, the composition of the host crystal, and the relative degree of crystallization of the inclusion. Glass inclusion compositions define a broad trend of increasing differentiation from type I to type III xenoliths. Analyses of glass inclusions and interstitial glass in each xenolith type have similar ranges in elemental ratios, with the exception of sample 5-17, which compositionally resembles type II xenoliths. Figure 11b shows the parallel trends, on a plot of FeO^*/CaO and FeO^*/MgO , of data from glass inclusions, interstitial glass, and host lava. Superimposed on this trend is the effect of the growth of the host crystal. Crystallization results in a decrease of CaO when the inclusion is trapped in plagioclase, a decrease of MgO when trapped in

olivine, or a decrease in CaO and MgO when trapped in augite. The distance of the xenolith from the glassy rim of the host lava controls the degree of crystallization of the host crystal [Clocchiatti, 1980]. Thus the xenoliths located farthest from the glassy rim have glass inclusion compositions most displaced from the MORB fractionation trend.

The composition of the original trapped liquid may be calculated, however, by the intersection of fractionation lines of the coexisting host minerals on a triangular oxide plot [Watson, 1976; Clocchiatti, 1977]. Each Juan de Fuca xenolith type contains at least two different coexisting minerals; therefore glass inclusions can be used to estimate the magma composition. These estimated initial compositions are presented in Table 9.

Crystallization Sequence

In general, the crystallization sequence is coherent for the entire xenolith suite. The presence of euhedral plagioclase grains and its occurrence as inclusions in olivine and augite indicate that plagioclase was the liquidus phase for liquids from which the xenoliths crystallized. Textural evidence suggests that olivine and augite began crystallizing essentially simultaneously; rare intergrowths of olivine in augite and vice versa occur, but olivine and augite more commonly are mutually exclusive. Compositional evidence suggests that olivine may have begun crystallizing slightly earlier than augite be-

TABLE 5. Representative Pyroxene Analyses

	Phenocrysts					Microphenocrysts					Type II			
	1	2	3	4	5	6	7	8	9	10	11	12	13	14
Sample Type	8-2 aug	8-2 aug	8-2 aug	8-2 aug	8-2 aug	22-34 aug	26-13B aug	26-13B aug	26-13B aug	26-13B aug	5-0 aug	5-1 aug	5-1 aug	17-1 aug
SiO_2	51.6	51.3	50.5	53.4	51.2	51.8	51.7	52.0	51.6	52.7	52.5	51.9	52.3	53.2
Al_2O_3	3.63	3.54	6.45	1.80	3.83	2.54	2.27	3.51	2.80	1.29	2.40	3.08	2.02	2.49
FeO^*	5.63	5.67	5.75	5.77	6.15	7.37	6.65	6.91	7.34	7.45	6.09	5.99	8.56	6.19
MgO	16.4	17.1	16.0	18.1	17.5	18.3	17.6	17.7	18.8	19.0	17.4	17.4	17.2	17.9
CaO	21.1	20.0	19.3	19.9	19.0	19.9	20.2	19.2	17.9	18.1	20.3	20.4	18.9	20.2
Na_2O	0.25	0.23	0.24	0.20	0.24	...	...	...	...	...	0.23	0.25	0.25	0.24
TiO_2	0.79	0.68	0.72	0.52	0.73	...	...	...	...	...	0.53	0.55	0.65	0.51
Cr_2O_3	1.15	1.00	1.07	0.60	0.96	...	...	...	...	...	0.26	0.41	0.04	0.31
MnO	0.17	0.20	0.21	0.20	0.20	...	...	...	...	...	0.20	0.23	0.31	0.23
Total	100.7	99.7	100.2	100.5	99.8	99.9	98.3	99.2	98.4	98.5	99.9	100.2	100.4	101.2
Cations per Formula Unit														
Si	1.880	1.882	1.842	1.941	1.875	1.891	1.918	1.911	1.907	1.949	1.927	1.901	1.927	1.925
Al	0.156	0.153	0.277	0.077	0.165	0.109	0.099	0.152	0.122	0.056	0.104	0.133	0.088	0.106
Fe	0.172	0.174	0.175	0.175	0.188	0.225	0.206	0.212	0.227	0.230	0.187	0.183	0.264	0.187
Mg	0.891	0.935	0.870	0.980	0.955	0.996	0.973	0.969	1.036	1.047	0.951	0.952	0.946	0.964
Ca	0.824	0.786	0.754	0.775	0.745	0.778	0.803	0.756	0.709	0.717	0.798	0.798	0.748	0.783
Na	0.018	0.016	0.017	0.014	0.017	...	...	...	...	...	0.016	0.018	0.018	0.017
Ti	0.022	0.019	0.020	0.014	0.020	...	...	...	...	...	0.015	0.015	0.018	0.014
Cr	0.033	0.029	0.031	0.017	0.028	...	...	...	...	...	0.008	0.012	0.001	0.009
Mn	0.005	0.006	0.006	0.006	0.006	...	...	...	...	...	0.006	0.007	0.010	0.007
Total	4.001	4.000	3.992	3.999	3.999	3.999	3.999	3.999	4.001	3.999	4.012	4.019	4.020	4.012
Mineral Norm [Lindsley and Andersen, 1983]														
Wo	38.3	36.1	31.4	37.7	33.9	33.5	36.1	33.3	30.8	33.3	38.1	...	35.8	37.2
En	53.0	55.5	57.1	53.3	56.9	59.5	55.8	56.0	59.8	56.7	53.3	...	52.6	54.4
Fs	8.7	8.4	11.5	9.0	9.2	7.0	8.1	10.7	9.4	10.0	8.6	...	11.6	8.4
Others	12.9	12.8	17.6	7.0	13.5	10.9	8.2	9.0	9.3	5.1	9.1	12.2	9.4	9.5
Calculated Fe_2O_3	0.90	1.21	0.00	0.37	1.24	3.94	2.31	0.97	2.31	1.65	1.25	2.19	1.98	1.38
Mineral Norm*														
Wo	43.7	41.5	41.9	40.1	39.4	39.0	40.5	39.0	36.0	36.0	41.2	41.3	38.2	40.5
En	47.2	49.3	48.6	50.8	50.6	50.0	49.1	50.0	52.5	52.5	49.1	49.2	48.3	49.8
Fs	9.1	9.2	9.5	9.1	10.0	11.0	10.4	10.9	11.5	11.5	9.7	9.5	13.5	9.7

TABLE 5. (continued)

Sample Type	Type III											
	15	16	17	18	19	20	21	22	23	24	25	26
	26-1A aug	5-7A aug	5-7B aug	5-7A aug-ex	5-8B aug-ex	5-7A pig	5-7B pig	5-7A opx	5-7A opx	5-7A opx	5-8B opx	5-8B opx
SiO ₂	52.5	53.1	51.8	51.9	51.8	53.6	53.4	54.0	53.7	53.5	53.2	53.0
Al ₂ O ₃	2.18	1.83	2.04	2.16	2.02	1.00	0.86	1.20	0.82	0.76	0.96	0.88
FeO*	6.69	6.85	11.1	9.49	11.7	15.6	17.0	13.7	16.3	17.8	18.7	20.8
MgO	17.5	17.9	17.3	17.7	15.3	24.7	24.1	28.0	26.0	25.0	24.2	22.8
CaO	19.7	19.6	16.7	17.3	18.3	4.33	4.30	2.03	2.07	2.34	2.26	2.16
Na ₂ O	0.22	0.22	0.29	0.23	0.30	0.07	0.08	0.02	0.03	0.03	0.03	0.02
TiO ₂	0.51	0.45	0.78	0.75	1.01	0.41	0.35	0.61	0.33	0.48	0.54	0.50
Cr ₂ O ₃	0.09	0.09	0.09	0.06	0.02	0.03	0.05	0.04	0.04	0.03	0.03	0.03
MnO	0.23	0.26	0.38	0.33	0.40	0.47	0.51	0.40	0.44	0.53	0.53	0.57
Total	99.7	100.3	100.5	100.0	100.8	100.2	100.6	100.0	99.7	100.4	100.5	100.9
Cations per Formula Unit												
Si	1.934	1.941	1.918	1.921	1.925	1.954	1.953	1.943	1.961	1.956	1.951	1.955
Al	0.094	0.079	0.089	0.094	0.089	0.043	0.037	0.051	0.035	0.033	0.042	0.038
Fe	0.206	0.209	0.345	0.293	0.363	0.474	0.519	0.412	0.498	0.543	0.574	0.642
Mg	0.959	0.978	0.954	0.977	0.845	1.343	1.311	1.500	1.412	1.360	1.323	1.256
Ca	0.779	0.769	0.662	0.685	0.729	0.169	0.168	0.078	0.081	0.092	0.089	0.085
Na	0.016	0.015	0.021	0.016	0.022	0.005	0.006	0.001	0.002	0.002	0.002	0.001
Ti	0.014	0.012	0.022	0.021	0.028	0.011	0.010	0.016	0.009	0.013	0.015	0.014
Cr	0.003	0.003	0.003	0.002	0.001	0.001	0.002	0.001	0.001	0.001	0.001	0.001
Mn	0.007	0.008	0.012	0.010	0.012	0.015	0.016	0.012	0.014	0.016	0.016	0.018
Total	4.012	4.014	4.026	4.019	4.014	4.015	4.022	4.014	4.013	4.016	4.013	4.010
Mineral Norm [Lindsley and Andersen, 1983]												
Wo	37.3	37.1	...	...	36.0	9.8	9.0	4.0	4.2	4.7	4.6	4.4
En	53.2	53.5	...	...	46.4	67.5	66.0	76.0	71.6	68.4	67.0	63.6
Fs	9.5	9.4	...	...	17.6	22.7	25.0	20.0	24.3	27.0	28.0	32.0
Others	8.4	7.8	10.4	10.0	9.7	5.0	4.6	5.0	4.0	3.6	4.4	4.2
Calculated Fe ₂ O ₃	1.26	1.39	2.29	2.07	1.51	0.90	0.80	0.67	0.66	0.28	0.47	0.40
Mineral Norm*												
Wo	40.1	39.3	33.8	35.0	37.6	8.5	8.4	3.9	4.1	4.6	4.5	4.3
En	49.3	50.0	48.6	50.0	43.6	67.6	65.6	75.4	70.9	68.2	66.6	63.3
Fs	10.6	10.7	17.6	15.0	18.7	23.9	26.0	20.7	25.0	27.2	28.9	32.4

Analysis 12 from core or oikocryst; 13 from small grain at edge of xenolith; analyses 18,23 and 19,25 are host-lamellae pairs. Pyroxene data were obtained using a ARL-SEMQ automated nine-channel microprobe and reduced using the matrix correction program of Bence and Albee [1968]. Standards used in analysis of pyroxene were synthetic Di-Jd for Si, Ca, Mg, Na, and Al; TiO₂ for Ti; Cr₂O₃ for Cr; Mn₂O₃ for Mn; and fayalite for Fe. Sample current was 20 nA, and counting time was 60 s.

*Wo = Ca/(Ca + Mg + Fe); En = Mg/(Ca + Mg + Fe); Fs = Fe/(Ca + Mg + Fe).

†aug = augite; aug-ex = exsolved augite; pig = pigeonite; opx = orthopyroxene.

cause the least evolved compositions are from type I (olivine + plagioclase) xenoliths. Primary pigeonite occupies interstitial positions and is a late crystallizing phase. Fe-Ti oxides were the last phases to crystallize and occur in the interstitial glass or along grain boundaries.

These observations suggest the following paragenetic sequence for the xenoliths: initial crystallization of plagioclase followed by plagioclase + olivine + augite, plagioclase + olivine + augite + pigeonite, and plagioclase + olivine + augite + inversion of pigeonite + Fe-Ti oxides. After crystallization of pigeonite and Fe-Ti oxides the xenoliths underwent ductile deformation, indicated by subtle kink banding in olivine, and brittle fracture, indicated by melt-filled fractures crosscutting the most crystalline xenoliths. Quenching of the xenoliths in the host lava preserved the different stages in the crystallization history.

The crystallization sequence defined by the xenoliths is similar to the sequence found in lavas from the Galapagos spreading center at approximately 85°W. Perfit and Fornari [1983] found the following crystallization sequence in Gala-

pagos lavas: initial crystallization of plagioclase + olivine followed by plagioclase + augite + olivine, plagioclase + augite + pigeonite, and plagioclase + augite + pigeonite + Fe-Ti oxides. In the Galapagos lavas, olivine ceases to crystallize in ferrobalt containing small microphenocrysts of pigeonite. The coexistence of olivine and pigeonite in the most crystalline xenoliths reflects the metastable assemblage caused by continued differentiation of the residual melt during in situ crystallization. The crystallization sequence found in the Juan de Fuca Ridge xenoliths and Galapagos spreading center lavas differs from results of crystallization experiments done on a differentiated sample from the Oceanographer Fracture Zone [Walker et al., 1979] in the lack of chromian spinel as a near-liquidus phase, the occurrence of olivine before augite, and the occurrence of magnetite.

General Mineral Trends

Figure 6 is a compilation of pyroxene, olivine, and plagioclase data for phenocrysts, microphenocrysts, and xenoliths. Figure 8 shows the compositional trends of hot and interstitial

Fig. 7. Photographs and photomicrographs of xenoliths in basalt from the Juan de Fuca Ridge. (a) Sample 5-0; medium-grained gabbroic xenolith enclosed in aphyric host lava. Black near end of scale bar is interstitial glass. (b) Sample 5-7A; type III xenolith exposed on fracture surface of sheet flow fragment. Note sharp textural contrast between aphyric host lava and xenolith. (c) Sample 17-1A; open textured, glassy xenolith. Plagioclase laths (white) are subophitically surrounded by augite. Dark grey is glass. Augite and plagioclase contain abundant glass inclusions. Plane light. (d) Sample 17-1A; close-up of flowage texture in glass around augite crystal. Plane light. (e) Sample 5-0; photomicrograph of section cut through sample shown in Figure 7a. Ophitic texture with augite (light grey with well-developed cleavage) surrounding plagioclase laths (white). Sprays of plagioclase laths extend into interstitial glass (black, center of photo). Crossed nicols. (f) Sample 26-1A; medium-grained, fairly equigranular xenolith. Black material in upper left of photo is host lava; black material within xenolith is interstitial glass. Augite (grey) and plagioclase (white and light grey) are dominant minerals. Crossed nicols. (g) Sample 5-7A; coarse-grained type III xenolith. Olivine (light grey with irregular fractures) and plagioclase are the dominant minerals. Pyroxene is a darker grey. An inverted pigeonite can be seen just above the left end of the scale bar; Figure 9b is a close-up of this pigeonite. Plane light. (h) Sample 5-8B; coarsened-grained type III xenolith. Large plagioclase in middle of photo. Augite is medium grey in lower half of xenolith. Olivine is light grey with irregular fractures in the central upper portion of the photo. Close-up of plagioclase and pyroxene intergrowth texture found in upper right edge of photo appears in Figure 9a.

Fig. 8. Major and minor element variation diagrams showing evolved interstitial glass compositions relative to host lava compositions. (a) TiO_2 versus FeO^*/MgO plot shows the increasing TiO_2 and FeO^* contents with increasing differentiation. (b) P_2O_5 versus K_2O plot shows enrichment of incompatible elements in the interstitial melts during differentiation.

glasses. The transition from type I to type III xenoliths is marked by sympathetic compositional variation of coexisting augite, plagioclase, olivine, and glass. Minerals in the xenoliths are more evolved than phenocrysts and microphenocrysts in the host lavas. The Fe-enrichment trends defined by analyses of mafic phases are similar to trends defined by phenocrysts in lavas from the Galapagos spreading center [Byerly, 1980; Perfit and Fornari, 1983], oceanic gabbros [Miyashiro and Shido, 1980; Tiezzi and Scott, 1980; Vanko and Batiza, 1982; Hebert et al., 1983], and other tholeiitic intrusions such as the Skaergaard [Wager and Deer, 1939], the Stillwater Complex [Hess, 1960], and the Bushveld [Truter, 1955].

Plagioclase shows a less systematic variation between xenolith types. Plagioclase in type II and type III xenoliths show similar ranges in composition but are more sodic than those in type I xenoliths or the microphenocrysts.

Olivine analyses provide the most conclusive evidence that the xenoliths formed in equilibrium with melts more evolved than the host lavas because olivine-liquid Fe/Mg partitioning is independent of temperature [Roeder, 1974]. Olivine-melt equilibrium calculations for microphenocrysts and host lava using $D_{\text{ol-Liq}} = (\text{MgO}/\text{FeO})_{\text{Liq}} / (\text{FeO}/\text{MgO})_{\text{ol}}$ [Roeder, 1974] indicate that the distribution coefficient ($D_{\text{ol-Liq}}$) is approximately 0.24 (assuming that all iron is FeO). An assumed distribution coefficient of 0.30 results in calculated FeO/FeO^* ratios of 0.74–0.84. Perfit and Fornari [1983] arrived at a similar conclusion for olivine phenocrysts from the Galapagos spreading center.

Olivine shows and Fe-enrichment trend from type I to type III (Figure 12). In contrast to the host lava-microphenocryst equilibria, however, the xenolith olivine-interstitial glass equilibria calculations are consistent with the melt having FeO/FeO^* ratio of 0.9 and olivine-melt distribution coefficient of approximately 0.30. The higher FeO/FeO^* ratios imply that

the conditions of formation of the gabbros are more reduced than the conditions of formation of the microphenocrysts.

Pyroxene shows a systematic decrease in Cr_2O_3 , increase in TiO_2 , and slight decrease in Al_2O_3 with increasing $\text{Fe}/(\text{Fe} + \text{Mg})$ ratio (Figure 10). These data suggest that the xenoliths are comagmatic. Augite compositions of type II and type III xenoliths overlap, but the most Fe-enriched augite occurs in a type III xenolith with partially inverted pigeonite. Compositions of augite phenocrysts and microphenocrysts do not follow the Fe-enrichment trend defined by the different xenolith types but instead are depleted in Ca (Wo_{37-40}) and enriched in Al at relatively constant Fe/Mg ratios. Similar trends have been noted in Ca-pyroxenes from DSDP hole 425 [Mattey and Muir, 1980] and hole 319 [Mazzullo and Bence, 1976]. The Ca depletion trend has been related to kinetic effects upon quenching [Natland, 1980].

Partitioning of FeO and MgO between coexisting clinopyroxene and glass is shown on Figure 13. Calculated $K_D = (\text{FeO}^*/\text{MgO})_{\text{Cpx}} / (\text{FeO}^*/\text{MgO})_{\text{Liq}}$ are 0.21 for phenocrysts-host glass pairs and 0.15–0.20 for xenolith pyroxene-interstitial glass pairs. Lower K_D calculated for the xenoliths imply that they formed under more reduced conditions than the phenocrysts, consistent with the olivine data. These values are lower than the 0.26 average value reported for the FAMOUS area [Bryan, 1979].

The occurrence of high Mg contents in ilmenite and the existence of such wide ranges of compositions within single xenoliths are unusual in an oceanic environment. MgO contents in oceanic samples rarely exceed 5.5 wt % [Haggerty, 1976; Byerly, 1980; Perfit and Fornari, 1983]. The most influential factor controlling ilmenite chemistry is the composition of the magma from which it crystallized. Magnesian ilmenite is characteristic of kimberlites, where it forms in equilibrium

with extremely high MgO melts [Haggerty, 1976]. If melt composition were the only factor, then ilmenite crystallizing from extremely differentiated melt, as is indicated by the silicate mineralogy in the Juan de Fuca Ridge xenoliths, should have low MgO contents. Pinckney and Lindsley [1976] showed that the preference of Mg for the rhombohedral phase increases with decreasing temperature. Ilmenite grains with vastly different MgO contents occur within millimeters of one another; therefore it is unlikely that temperature differences caused a factor of 2 difference in Mg content. The origin and significance of the high-Mg ilmenite is unknown.

Geothermometry and Cooling Rate

Plagioclase-melt equilibria calculations [Drake, 1976] for samples 5-7A and 8-2 indicate that the phenocrysts and microphenocrysts in the host lava are in equilibrium with a dry melt in the temperature range of 1200°–1250°C. The presence of water in the melt increases the An content of plagioclase [Johannes, 1978] and results in calculation of anomalously high temperatures; hence the plagioclase-melt equilibria temperatures for a dry melt are maximum temperature estimates, although the low volatile content estimates in normal MORB [Delaney et al., 1978] and ferrobassalt [Byers et al., 1983, 1984] suggest that water in the melt should not have significantly affected plagioclase equilibrium.

Use of the Lindsley [1983] graphical geothermometer for augite in type II xenoliths yields estimates of crystallization temperatures of 1100°–1200°C. The temperature range for phenocrysts and microphenocrysts is 1100°–1300°C; these estimates are not reliable because of quenching control of CaO content. Subsolidus equilibrium temperatures can be estimated by using coexisting orthopyroxene-augite-pigeonite assemblages found in inverted pigeonite. Compositions of host-lamellae pairs in sample 5-8B and 5-7A yield temperature estimates of 1100°–1150°C and 1130°C, respectively. Noninverted pigeonite in sample 5-7B also yields a temperature of 1130°C. High-Fe augite in sample 5-72 gives a temperature of 1090°C.

The presence of exsolved Ca-rich pyroxene in pigeonite indicates slow cooling. The character of exsolution lamellae in pyroxene reflect bulk composition, maximum temperature, and cooling rate [Robinson, 1980]. Inverted pigeonite with blebby augite, similar to inverted pigeonite in the Juan de Fuca xenoliths, has been called "Kintoki-San type" [Kuno,

1966] and has been found in plutonic [Brown, 1957] and hypabyssal [Ishii, 1973] environments. Ishii and Takeda [1974] suggest that the blebby texture is produced when pigeonite undergoes eutectoidal decomposition into orthopyroxene and augite at the temperature of the pigeonite eutectoid reaction point.

Minimum stability temperatures for pigeonite in the xenoliths can be derived graphically from a plot of minimum stability for pure Ca-Mg-Fe pigeonite, relative to augite and orthopyroxene, as a function of $X = \text{Fe}^{2+}/(\text{Fe}^{2+} + \text{Mg})$ [Lindsley, 1983] and give an estimate of $1150 \pm 30^\circ\text{C}$. The use of minimum temperature of stability curves at low pressures of natural pigeonite, instead of pure Ca-Mg-Fe pigeonite [Ross and Huebner, 1979], results in a slightly higher estimate of 1180°C. A minimum temperature of stability of approximately 1150°C is consistent with subsolidus temperatures of 1100°–1150°C for orthopyroxene-augite pairs in inverted pigeonite.

Inverted pigeonite with blebby exsolution of augite occurs in a two-pyroxene gabbro recovered at DSDP site 334. Pigeonite from site 334 gabbro has complex exsolution patterns because equilibration to low temperatures resulted in multiple sets of exsolution lamellae [Hodges and Papike, 1976]. Pigeonite in Juan de Fuca gabbro xenoliths has simple exsolution patterns reflecting the arrested equilibrium at higher temperatures. Partly inverted pigeonite has been observed in terrestrial hypabyssal rocks, such as the Palisades Sill [Walker et al., 1973] and the Kintoki-San dike in Hakone Volcano [Ishii and Takeda, 1974], and in lunar and meteoric pyroxenes [Papike and Bence, 1972; Ghose et al., 1973; Harlow et al., 1979]. We assume that inversion of pigeonite and other subsolidus equilibration reactions in the xenoliths would have proceeded to completion had eruption and quenching not abruptly changed the rate of cooling. Therefore the similarity in texture between pigeonite from the xenoliths and that from the Palisades Sill implies that the rate of cooling of the Palisades Sill was greater than that of the magma chamber source of the xenoliths.

Estimates of temperature and log f_{O_2} using the Stormer calculation method for composite Fe-Ti oxides are 1027°C and -10.47 in sample 5-8b (Pair 1 + 3) and 1054°C and -9.92 in sample 5-7a (pair 6 + 9). The composite grains should be the most reliable T and f_{O_2} indicators. Calculations based on Carmichael [1967] and Lindsley and Spencer [1982] yield T estimates consistently lower by 40°–60°C and log f_{O_2} lower by up to 1.30. Scatter in the estimates increases with increasing minor element contents. The use of high-Mg ilmenite in the magnetite-ilmenite mineral pair results in lower temperature estimates (859°–919°C) and lower log f_{O_2} estimates (-12.5 to -14.0). Maximum and minimum temperature and f_{O_2} estimates are shown in Table 6. Lack of optically resolvable Fe-Ti oxide exsolution indicates quenching without significant slow cooling after oxides formed.

Discussion

Textural evidence from the Juan de Fuca Ridge xenoliths suggests that the xenoliths formed by in situ crystallization near the margin of a magma chamber. Xenoliths from Juan de Fuca are characterized by random orientation of grains, wide range of grain sizes, zoning deep within the crystals in the more crystalline xenoliths, and abundant intergrowth textures. Textures characteristic of cumulate gabbros, such as homogeneous cumulate phases, limited grain size variation, and edge zoning, are not present. The Juan de Fuca Ridge xenoliths have textures similar to, except for the abundance of glass, those of high-level isotropic gabbro from the Samail ophiolite [Pallister and Hopson, 1981]. The sympathetic variation of interstitial glass, glass inclusion, and mineral compositions

Fig. 9. (opposite) Photomicrographs of textures found in type III xenoliths. (a) Sample 5-8B; intergrowth texture of augite and plagioclase. Crossed nicols. (b) Sample 5-7A; blebby exsolution texture in inverted pigeonite. Plagioclase is medium grey with albite twinning in the upper part of the photo. Orthopyroxene is dark grey in the center of the pyroxene. Augite is light grey in the central right of the photo. Crossed nicols. (c) Sample 5-8B; blebby exsolution texture in inverted pigeonite. Augite is light grey and orthopyroxene is dark grey. Exsolution more aligned along crystallographic planes than in other pigeonites. Crossed nicols. (d) Sample 5-7A; blebby exsolution texture in inverted pigeonite. This sample shows the great variability in bleb width. A core of pigeonite remains in the upper central part of the photo. Crossed nicols. (e) and (f) Samples 5-7A and 5-8B; anhedral oxides (white) in interstitial glass (grey). Photomicrographs were taken with half reflected and half transmitted light to show position of oxides within interstitial glass. (g) Sample 5-8B; melt-filled fracture through olivine crystal indicating a stage of brittle deformation. Olivine has subtle extinction discontinuity which runs approximately through the center of the photo. Olivine below the discontinuity is slightly darker grey than above. Box defines area of Figure 9i. Crossed nicols. (h) Sample 5-8B; melt-filled fracture through augite and plagioclase. Augite and plagioclase have started to react with the melt. Crossed nicols. (i) Sample 5-8B; close-up of fracture in Figure 9g. Reaction between plagioclase, olivine, and melt can be seen by the ragged crystal-melt boundaries. Crossed nicols.

Fig. 10. Minor element variation diagram for pyroxene in host lava and xenoliths. Augite from the xenoliths (solid circles) exhibits a systematic decrease in Cr_2O_3 and Al_2O_3 and increase in TiO_2 with increasing $\text{Fe}/(\text{Fe} + \text{Mg})$ and can be related to a single fractionation trend. Scatter in TiO_2 and Al_2O_3 contents in phenocrysts and microphenocrysts is a result of disequilibrium crystallization during rapid cooling.

support the interpretation of in situ crystallization and document the changing composition of the interstitial liquid.

Several lines of evidence indicate that the Juan de Fuca xenoliths formed by quenching of a not yet holocrystalline mush and not by partial melting of a holocrystalline gabbro. Examination of the entire textural suite shows that for the more open-textured xenoliths, euhedral grains are set in a matrix of glass. The grain size and crystallinity increase at the expense of interstitial glass. An interstitial origin for the glass in the more crystalline xenoliths is indicated by the location of glass in interstitial areas at grain contacts, homogeneity of glass within a given xenolith regardless of adjacent mineral phases, and absence of reaction textures at interstitial glass-mineral contacts.

In contrast to the sharp interstitial glass-mineral boundaries, evidence of disequilibrium between melt and xenolith minerals occurs at host lava-xenolith mineral and injected melt-xenolith mineral boundaries for the most differentiated type III xenoliths. Thus some reaction between xenoliths and host lava occurred between the time of entrainment and the time of quenching on the seafloor, but evidence of partial melting to produce the interstitial glass is not present.

The response of the xenoliths to stress is a function of their texture and temperature at the time of deformation. Xenolith 17-1 has swirls of melt around the crystals, while xenoliths 5-7A and 5-8B are crosscut by melt-filled fractures. Based on

the previous discussion of geothermometry, these type III xenoliths equilibrated to temperatures of approximately 1000°C but no lower before they were quenched at the surface. The confining pressure during fracturing is difficult to constrain because fracturing could have occurred in the magma chamber as the fragment was ripped from the wall rock, during transport to the surface, or during eruption. Nonetheless, the gabbro xenoliths were brittle enough to be fractured. Fracturing has also been documented in Cayman Trough gabbros, where Malcolm [1981] interpreted veins of clinopyroxene and brown hornblende as residual magmatic fluid injected into fractures in the gabbros near their solidus temperature.

The most striking features of the xenoliths are their evolved composition with respect to the host lava and the sharp textural contrast between the xenoliths and the nearly aphyric host lava. Crystal clots less evolved than the host lava have been found in Iceland [Larsen, 1979] and International Program of Ocean Drilling (IPOD) leg 46 site 396B [Sato *et al.*, 1978] and are interpreted to be cognate xenoliths. Gabbro xenoliths from Iceland are most common in porphyritic basalt and can be explained by a single episode of crystallization. A large gabbroic xenolith with composition more evolved than the host melt occurs in basalt from DSDP site 504B near the Costa Rica Rift [Natland, 1983]. This single xenolith was interpreted to be a fragment removed from the walls of a

TABLE 6. Fe-Ti Oxide Analyses

Type Number of points analyzed	Sample 5-8B					Sample 5-7A							
	1	2	3	4	5	6	7	8	9	10	11	12	13
	mag (4)	mag (1)	ilm (3)	ilm (1)	hi-Mg ilm (1)	mag (2)	mag (3)	mag (2)	ilm (2)	ilm (3)	ilm (3)	hi-Mg ilm (2)	hi-Mg ilm (1)
<i>Recalculated Analyses [Carmichael, 1967]</i>													
SiO ₂	0.01	0.03	0.00	0.00	0.00	0.03	0.01	0.01	0.00	0.00	0.01	0.00	0.00
TiO ₂	20.6	20.1	48.0	48.4	51.6	20.2	21.2	20.2	48.9	48.7	49.1	50.5	51.6
Al ₂ O ₃	2.53	2.51	0.32	0.45	0.41	2.67	2.61	3.36	0.56	0.41	0.47	0.45	0.38
V ₂ O ₃	1.12	1.06	0.43	0.37	0.84	1.25	0.88	1.30	0.45	0.74	0.86	0.95	0.69
Cr ₂ O ₃	0.05	0.09	0.01	0.00	0.06	0.02	0.01	0.05	0.00	0.00	0.02	0.03	0.16
Fe ₂ O ₃	26.8	27.8	10.6	10.2	7.31	27.3	26.0	27.3	11.5	10.3	10.1	9.26	5.72
FeO	45.8	44.4	36.2	34.5	31.0	45.1	45.8	43.5	31.4	33.8	32.2	29.9	32.1
MnO	0.47	0.38	0.46	0.38	0.36	0.44	0.46	0.39	0.36	0.40	0.37	0.34	0.42
MgO	2.82	3.51	3.67	4.86	8.44	3.20	3.32	4.37	6.89	5.38	6.54	8.52	7.82
Total	100.2	99.9	99.7	99.2	100.0	100.2	100.3	100.5	100.1	99.7	99.7	100.0	98.9
<i>Cations Per Formula Unit (Sp. = 3, Rh. = 2)</i>													
Si	0.000	0.001	0.000	0.000	0.000	0.001	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Ti	0.564	0.548	0.893	0.895	0.921	0.551	0.577	0.543	0.884	0.892	0.893	0.902	0.935
Al	0.109	0.108	0.009	0.013	0.012	0.114	0.111	0.141	0.016	0.012	0.013	0.013	0.011
V	0.033	0.031	0.009	0.007	0.016	0.036	0.026	0.037	0.009	0.015	0.017	0.018	0.013
Cr	0.001	0.003	0.000	0.000	0.001	0.001	0.000	0.001	0.000	0.000	0.000	0.001	0.003
Fe ⁺³	0.729	0.760	0.197	0.189	0.130	0.745	0.709	0.733	0.208	0.189	0.184	0.165	0.104
Fe ⁺²	1.401	1.348	0.748	0.709	0.615	1.366	1.384	1.299	0.630	0.688	0.650	0.593	0.646
Mn	0.014	0.012	0.010	0.008	0.007	0.014	0.014	0.012	0.008	0.008	0.008	0.007	0.009
Mg	0.149	0.190	0.135	0.178	0.298	0.173	0.179	0.233	0.247	0.195	0.236	0.302	0.281
Total	3.000	3.001	2.001	1.999	2.000	3.001	3.000	2.999	2.002	1.999	2.001	2.001	2.002
<i>Recalculated Mole Fractions</i>													
	usp	usp	ilm	ilm	ilm	usp	usp	usp	ilm	ilm	ilm	ilm	ilm
Storner [1983]	0.624	0.598	0.892	0.894	0.920	0.610	0.630	0.609	0.878	0.892	0.892	0.898	0.938
Anderson [1968]	0.582	0.554	0.884	0.882	0.904	0.566	0.588	0.553	0.859	0.879	0.876	0.878	0.926
Carmichael [1967]	0.564	0.549	0.893	0.895	0.921	0.552	0.577	0.544	0.884	0.894	0.893	0.902	0.935
Lindsley and Spencer [1982]	0.604	0.588	0.900	0.904	0.934	0.594	0.617	0.595	0.895	0.904	0.907	0.916	0.947
<i>Mineral Pair Calculations (mag + ilm)</i>													
	(1 + 3)	(2 + 4)	(2 + 5)	(6 + 9)	(7 + 9)	(8 + 10)	(8 + 13)						
	T	f _{O₂}	T	f _{O₂}	T	f _{O₂}	T	f _{O₂}	T	f _{O₂}	T	f _{O₂}	T
Storner [1983]	1027	-10.47	1001	-10.82	919	-12.51	1054	-9.92	1072	-9.70	1015	-10.61	859
Anderson [1968]	1015	-10.47	996	-10.68	938	-11.84	1058	-9.60	1078	-9.39	1002	-10.55	869
Carmichael [1967]	978	-11.08	959	-11.38	884	-13.03	990	-10.78	1010	-10.53	963	-11.27	830
Lindsley and Spencer [1982]	988	-11.10	963	-11.51	861	-13.81	994	-10.92	1013	-10.68	970	-11.41	806

Analyses 1 and 3 are from a composite grain, as are 6 and 9. Fe-Ti oxide data were obtained using an ARL-SEMQ automated nine-channel microprobe and reduced using the matrix correction program of Bence and Albee [1968]. Standards used in the analysis of Fe-Ti oxides were synthetic V-doped diopside for Si and V; Fe₂O₃ Fe; TiO₂ for Ti; synthetic spinel for Mg and Al; Mn₂O₃ for Mn and Tiesbaghi chromite for Cr. Sample current was 20 nA. The vanadium intensity (k-alpha) was calculated by subtracting the Ti_{k-beta} peak from the measured total intensity.

magma chamber or shallow conduit system that originally contained more fractionated melt. The abundance of glass and a broad spectrum of texture and compositions in the Juan de Fuca xenolith suite suggest that formation of the gabbros is intimately involved with the magma chamber and not the product of crystallization in a small isolated magma pocket. The evolved compositions suggest the existence of melt more evolved than the host melt in the magma chamber.

HAWAIIAN LAVA LAKES: AN ANALOGY

A puzzling aspect of the xenoliths is the large variability in mineral modes (Table 10). Estimated bulk xenolith compositions do not, in general, reflect lava compositions. Also the dominance of augite + plagioclase xenoliths is contrary to

what one would expect from simple crystallization along a liquid line of descent, which requires olivine in addition to augite + plagioclase. The problem may be one of scale. The large-scale view is one of in situ crystallization at the top of a cooling magma body, while in detail the actual processes may be extremely complex. Drill core studies of Hawaiian lava lakes; such as Kilauea Iki [Richter and Moore, 1966; Helz, 1980], Makaopuhi [Wright and Okamura, 1977], and Alae [Wright and Peck, 1978], document the small-scale complexities of crystallization in a cooling magma body.

A feature common to all the studied lava lakes in Hawaii is the presence of zones of relatively coarse-grained, glassy, vesicular segregation veins that appear to fill fractures and have high differentiated compositions [Wright and Okamura, 1977]. These segregation veins form by physical segregation of differ-

TABLE 7. Average Interstitial Glass Analyses

	Type I		Type II				Type III				
	17-7	17-69	17-1A	5-0	5-1	5-72	5-7B	26-1A	16-7A	17-3	5-8B
Analyses average	(3)	(2)	(4)	(4)	(3)	(5)	(2)	(4)	(2)	(2)	(3)
SiO ₂	50.6	50.6	51.4	50.8	51.6	51.1	51.3	51.7	51.4	51.3	52.7
Al ₂ O ₃	13.7	14.1	13.7	13.7	14.0	14.5	13.4	13.9	13.6	13.2	13.1
FeO*	12.2	12.3	12.9	13.5	14.0	14.1	13.5	12.9	14.2	15.0	13.2
MgO	6.78	6.65	6.21	5.77	5.65	4.90	6.25	5.42	5.17	5.04	3.51
CaO	10.9	10.8	10.3	10.1	9.62	10.0	10.2	9.83	9.25	9.33	8.05
Na ₂ O	2.58	2.70	2.86	3.02	3.17	3.06	2.81	3.05	3.04	3.03	3.69
K ₂ O	0.18	0.18	0.22	0.22	0.23	0.21	0.29	0.30	0.30	0.27	0.64
TiO ₂	1.92	1.90	2.10	2.36	2.41	2.44	2.46	2.55	2.93	2.97	2.96
P ₂ O ₅	0.18	0.20	0.22	0.26	0.27	0.23	0.29	0.30	0.37	0.33	0.84
MnO	0.19	0.19	0.20	0.22	0.21	0.21	0.22	0.20	0.22	0.23	0.23
SO ₃	0.16	0.15	0.17	0.17	0.19	0.20	0.15	0.17	0.19	0.20	0.11
Total	99.4	99.8	100.3	100.1	101.4	101.0	100.9	100.3	100.7	100.9	99.0
Mg #	51.0	50.3	47.5	44.6	43.1	39.5	46.5	44.1	40.5	38.7	33.2
FeO/MgO†	1.62	1.67	1.87	2.11	2.24	2.59	1.94	2.14	2.48	2.68	3.38

Analytical procedure for intersertal glasses was the same as the procedure for glassy rims, except that counting time was 120 s.

FeO = total Fe calculated as FeO.

†FeO/MgO calculated using $0.9 \times \text{FeO}^*$.

entiated melt by filter pressing from a rigid crystal framework. Melt, as defined by drilling, is a crystal-liquid mush too fluid to support the weight of the drill string, a condition reached when the magma contains approximately 45% glass and 55% crystals [Helz, 1980]. Wright and Peck [1978] also note that the composition of the liquid will vary in composition according to the efficiency of the filter press in separating liquid from crystals. Differences in the ratios of minerals brought in will produce a differentiate that does not lie on a "liquid line of descent." Helz [1980] documents that the total thickness of segregation veins in Kilauea Iki, the largest and most complex of the Hawaiian lava lakes, is 3–6 m in the central part of the lake, corresponding to 6–11% of the upper crust. Formation of segregation veins in the crystallizing zone above the southern Juan de Fuca Ridge magma chamber can explain the

highly variable bulk compositions of the xenoliths and the deviation from a simple liquid line of descent.

Segregation veins need fractures to form. Fracturing in lava lakes, which are static bodies, may be associated with degassing of lava and partially molten crust. Additional large horizontal fractures may form when the upper crust becomes more or less supported by the walls of the crater and fails to track the lens of melt as it cools and solidifies [Wright and Okumura, 1977]. Mid-ocean ridge magma chambers differ from Hawaiian lava lakes in that volatiles are kept in solution by high hydrostatic and lithostatic loads. The absence of volatile release as a fracturing mechanism, however, is more than compensated for by active tectonics at mid-ocean ridges. Lava lake studies show that fracturing can be sustained down to the crust-melt interface. Similarly, rifting at the ridge crest in the

TABLE 8. Average Glass Inclusion Analyses

	Type I		Type II		Type III				Type I		Type III		Type II		Type III			
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18
Sample	17-12	5-17	5-15	5-11	26-1A	16-7A	16-7B	17-3	17-12	5-17	26-1A	16-7B	5-11	5-15	16-7A	17-3	26-1A	16-7B
Host Mineral	plag	plag	plag	plag	plag	plag	plag	plag	ol	ol	ol	ol	aug	aug	aug	aug	aug	aug
Number of analyses	(2)	(3)	(1)	(3)	(3)	(3)	(6)	(3)	(2)	(3)	(1)	(2)	(3)	(4)	(2)	(3)	(2)	(1)
SiO ₂	50.6	47.5	48.1	51.3	47.4	50.8	44.6	51.1	50.2	51.2	51.2	53.2	51.3	52.4	50.5	50.9	54.9	52.0
Al ₂ O ₃	8.34	7.76	4.93	13.8	6.68	12.3	3.93	12.9	14.6	13.7	14.2	13.9	12.7	16.9	10.6	14.6	17.1	16.6
FeO*	16.5	20.9	19.9	12.6	21.3	14.2	27.5	15.3	11.7	13.8	14.2	14.3	13.5	13.2	15.1	13.8	10.6	13.1
MgO	9.08	9.02	10.9	5.71	9.25	5.55	9.28	4.28	3.95	3.53	3.22	1.71	5.50	2.54	7.10	3.96	2.15	1.09
CaO	10.3	9.50	10.1	10.3	9.04	9.38	8.56	8.69	12.7	10.7	11.3	9.54	10.3	7.37	9.76	8.68	6.77	6.10
Na ₂ O	1.15	1.04	0.81	2.26	1.19	2.62	0.67	3.02	2.67	2.75	2.82	3.38	2.61	3.36	2.28	3.23	4.33	4.28
TiO ₂	2.04	2.96	3.51	2.25	3.24	2.32	3.81	2.82	2.09	2.59	2.16	2.85	2.51	2.33	2.82	2.84	2.40	3.77
P ₂ O ₅	0.15	0.28	0.28	0.23	0.24	0.30	0.24	0.29	0.14	0.22	0.21	0.22	0.26	0.19	0.34	0.26	0.37	0.40
MnO	0.25	0.39	0.37	0.20	0.32	0.26	0.44	0.24	0.21	0.21	0.18	0.21	0.18	0.20	0.27	0.19	0.16	0.21
Total	98.4	99.4	98.9	98.7	98.7	97.7	99.0	98.6	99.3	98.7	99.5	99.3	98.9	98.5	98.8	98.4	98.8	97.6
FeO*/CaO	1.60	2.20	1.97	1.22	2.36	1.51	3.21	1.76	0.92	1.29	1.26	1.50	1.31	1.79	1.55	1.59	1.57	2.15
FeO*/MgO	1.82	2.31	1.83	2.21	2.30	2.56	2.96	3.57	2.96	3.91	4.41	8.36	2.45	5.20	2.13	3.48	4.93	12.0
CaO/MgO	1.13	1.05	0.93	1.80	0.98	1.69	0.92	2.03	3.22	3.03	3.51	5.58	1.87	2.90	1.37	2.19	3.15	5.60

Fig. 11a. Compositions of glass inclusions trapped in xenolith minerals plotted on MgO, CaO, FeO* triangular plot. Compositions of the host rock glasses (small dots) are plotted for comparison and fall on a MORB fractionation line. Symbols used are type I xenoliths: sample 5-7, solid triangle; sample 5-17, triangle with dot; sample 17-12, open triangle. Type II xenoliths: sample 5-11, solid circle; sample 5-15, circled dot; sample 16-7A, open circle; sample 17-2, horizontally half solid circle; sample 17-3, vertically half solid circle. Type III xenoliths: sample 16-7B, cross; sample 26-1A, plus. The short dashed lines link the inclusions in the different crystals of two type II xenoliths. Long dashed lines divide the field by host minerals.

oceanic environment may cause fractures that extend to this boundary. Rifting of the oceanic crust, therefore, may cause percolation of differentiated liquids into fractures at or near the ridge crest at mid-ocean ridges.

MAGMA MIXING AND FORMATION OF XENOLITHS

We favor a model of repetitive magma chamber replenishment to account for the presence of xenoliths within aphyric lava. Replenishment occurs before the magma chamber has completely solidified and results in mixing, rejuvenated convection, and superliquidus temperatures [Walker et al., 1979]. The boundary for rejuvenated convection would be the brittle-ductile boundary, approximately 45% liquid, as defined by the Hawaiian lava lake studies. The host lava is a hybrid formed by mixing highly evolved residual melt with a new batch of primitive melt [Clague et al., 1983]. Mixing is supported by the homogeneous lava composition within the axial valley and is probably controlled by density differences. Dense picritic magma may pond at the base of the magma chamber after injection as modeled by Huppert and Turner [1981]. Sparks et al. [1980] calculated that the density of picritic liquids decreases with decreasing temperature due to removal of olivine. The density of ferrobalt, on the other hand, increases with decreasing temperature due to removal of dominantly plagioclase. Convective mixing will occur when the density of the overlying ferrobalt is greater than or equal to the density of

Fig. 11b. Plot of $(\text{FeO}^*/\text{CaO})_{\text{liq}}$ versus $(\text{FeO}^*/\text{MgO})_{\text{liq}}$, showing the evolved compositions of interstitial glass and glass inclusions relative to host lava compositions. Lines join data from a single xenolith but from two different minerals. FeO^*/CaO and FeO^*/MgO ratios calculated in glass inclusions of olivine are systematically 10–15% higher than the same ratios calculated in coexisting plagioclase. Glass inclusion data from sample 5-17, a type I xenolith, are compositionally similar to interstitial glass from type II and III xenoliths.

the underlying olivine tholeiite to picrite. Eruption may be triggered by injection of the next batch of primitive magma. The time between injections of melt must be long enough to allow mixing and resorption of phenocrysts to occur but not long enough to allow the hybrid melt to cool to the liquidus temperature when new phenocrysts would form.

The linear rift valley, abundance of sheet flows, absence of fissuring, and active hydrothermal venting in the U.S. Geological Survey (USGS) study area suggest that the magma chamber has been replenished within the last few hundred years. Initial conduit formation for the eruption fractured the crystallizing boundary zone of the magma chamber. High effusion rate eruptions of superliquidus magma, responsible for the formation of extensive aphyric sheet flows, carried the gabbro fragments to the surface.

CONCLUSIONS

Basalt from the southern Juan de Fuca Ridge is ferrobalt derived from a depleted MORB source. Subtle differences in trace element ratios between basalt from the axial walls (2000–5000 years old) and axial valley floor (age < 1000 years) indicate that mantle source regions for Juan de Fuca MORB are variable on a time scale of thousands to tens of thousands of years.

Rare glassy gabbroic xenoliths found in this aphyric ferrobalt are divided into three types based on mineralogy and texture; two types are more evolved than the host lava. The coarsest-grained xenoliths equilibrated to temperatures of

TABLE 9. Estimated Initial Compositions of Glass Inclusions

Host Glass	Type I	Type II	Type II	Type III	Type I	Type III	Type III	Type III
Sample	17-12	5-15	5-11	16-7a	5-17	26-1A	16-7B	17-3
				<i>Inclusive in Olivine</i>				
FeO*/CaO	1.07 ± 0.02	1.04 ± 0.01	...	...	1.39 ± 0.01	1.43 ± 0.01	1.73 ± 0.03	...
FeO*/MgO	1.77 ± 0.05	1.74 ± 0.06	...	...	2.67 ± 0.06	2.72 ± 0.06	3.62 ± 0.26	...
CaO/MgO	1.65 ± 0.02	1.68 ± 0.04	...	...	1.89 ± 0.04	1.93 ± 0.04	2.17 ± 0.14	...
				<i>Inclusive in Plagioclase</i>				
FeO*/CaO	...	1.05 ± 0.01	1.07	1.19 ± 0.02	1.22 ± 0.15	1.32 ± 0.04	1.33 ± 0.04	1.63 ± 0.08
FeO*/MgO	...	1.74 ± 0.04	1.79	2.07 ± 0.08	2.12 ± 0.75	2.42 ± 0.19	2.40 ± 0.20	3.06 ± 0.45
CaO/MgO	...	1.66 ± 0.03	1.67	1.74 ± 0.06	1.74 ± 0.54	1.83 ± 0.13	1.81 ± 0.13	1.82 ± 0.27
				2.04 ± 0.26

Fig. 12. Plot of $(\text{FeO}^*/\text{MgO})_{\text{ol}}$ versus $(\text{FeO}^*/\text{MgO})_{\text{liq}}$ showing olivine-melt equilibria based on the method of Roeder [1974]. Arrow extends from analyzed $(\text{FeO}^*/\text{MgO})_{\text{liq}}$ (all iron as FeO) to $(\text{FeO}^*/\text{MgO})_{\text{liq}}$ using $\text{FeO} = 0.9 \times \text{FeO}^*$. Olivine-melt pairs are composed of phenocrysts (open square) and microphenocrysts (open circle) with host glass, and xenolith olivine with coexisting interstitial glass. Calculated K_D for phenocrysts/microphenocryst-host glass pairs are lower than K_D calculated for xenolith olivine-interstitial glass pairs.

1100°–1150°C based on two-pyroxene geothermometry and 1030°–1050°C based on Fe-Ti oxide geothermometry. These coarsest-grained xenoliths were able to sustain brittle fracture. Gabbroic xenoliths represent partially crystalline wallrock formed by in situ crystallization at the top of the magma chamber. The formation of segregation veins of differentiated material by filter pressing, as documented in Hawaiian lava lake studies, may be an important process in in situ crystallization at mid-ocean ridges. Crystallization of some xenoliths from these segregation veins may explain the great variability in mineral modes and the abundance of augite + plagioclase xenoliths.

We propose that the magma chamber under the southern Juan de Fuca Ridge has undergone recent replenishment. Ferrobasalt from the southern Juan de Fuca Ridge is a hybrid melt formed by mixing of residual differentiate with more primitive melt introduced during replenishment. Total resorption of phenocrysts due to superliquidus temperatures and rejuvenated convection may be responsible for the aphyric texture of the ferrobasalt. The gabbros crystallized from evolved liquids along the cooling boundary zone of a magma chamber prior to replenishment. The xenoliths were probably carried to the surface by high-rate eruptions of superliquidus magma that resulted in extensive aphyric sheet flows.

Fig. 13. Plot of $(\text{FeO}^*/\text{MgO})_{\text{Cpx}}$ against $(\text{FeO}^*/\text{MgO})_{\text{liq}}$ showing pyroxene melt equilibria. Pyroxene compositions are more evolved and variable than phenocrysts/microphenocrysts compositions. Bryan [1979] calculated a K_D of 0.26 for pyroxene in basalt from the FAMOUS area. The one type III pyroxene with high calculated K_D may reflect disequilibrium between pyroxene and interstitial glass due to crystallization of Fe-Ti oxides, thereby lowering the FeO^*/MgO ratios in the glass.

TABLE 10. Xenolith Modal Mineralogy

Sample	Interstitial Glass	Plag	Ol	Aug	Pig	Fe-Ti Oxides	Counts
<i>Type I</i>							
17-7	28.5	52.1	19.4	...	...	...	1000
17-12	25.0	39.0	36.0	...	...	...	1000
Average*	26.7	45.6	27.7	...	...	...	
<i>Type II</i>							
5-0	21.9	38.6	trace	39.5	...	...	1000
5-1	16.2	45.4	18.2	20.2	...	...	1000
5-10	48.7	31.7	...	19.6	...	...	1000
5-11	20.9	39.3	...	39.8	...	...	1000
5-13	11.9	45.0	...	43.1	...	...	1000
16-5	25.0	48.0	...	27.0	...	...	100
16-7	35.1	27.1	...	37.8	...	...	1000
17-1A	51.0	27.5	1.0	20.5	...	...	1000
17-2	23.0	36.0	...	41.0	...	...	100
17-4	7.6	41.5	...	50.9	...	...	100
22-25	11.0	55.0	...	34.0	...	...	100
22-26	1.0	33.7	...	65.3	...	...	100
Average*	28.3	36.8	2.6	32.3	...	...	
<i>Type III</i>							
5-7A	1.2	52.4	38.7	3.7	4.0	trace	1000
5-7B	19.8	32.1	...	48.1	trace	...	1000
5-8B	3.0	47.5	6.1	40.4	2.0	1.0	1000
16-7	5.5	56.5	0.5	34.7	2.8	trace	1000
17-3	4.3	83.0	2.8	9.9	...	...	1000
26-1A	16.0	51.0	7.0	26.0	...	...	1000
Average*	8.3	53.7	9.2	27.1	1.5	0.2	

*Weighted by the number of points counted. Type I, type II, and type III xenoliths represent 2, 80, and 18% of the xenoliths present.

Acknowledgments. We thank the captain and crew of the *S. P. Lee* for their support during fair and more often foul weather. The first author would like to thank Mike Rhodes and Joel Sparks at the University of Massachusetts, Ronald B. Gilmore Memorial Laboratory for their guidance in XRF analysis and Lew Calk for assistance in operation of the USGS microprobe. We would also like to thank Kay McDaniel for typing the tables. Thoughtful reviews by Charlie Bacon, Rosalind Helz, John Sinton, and Mike Dungan improved the manuscript. The third author thanks the French Ministère des Relations Extérieures for providing a grant in support of a year of work at the U.S. Geological Survey.

REFERENCES

- Allegre, C. J., B. Dupre, and B. Hamelin, Geochemistry of oceanic ridge basalts explained by blob mantle mixing, *Eos Trans. AGU*, 64, 324, 1983.
- Anderson, A. T., Oxidation of the LaBlache Lake titaniferous magnetite deposit, Quebec, *J. Geol.*, 76, 528–547, 1968.
- Batiza, R., Inverse relationship between Sr isotope diversity and rate of oceanic volcanism has implications for mantle heterogeneity, *Nature*, 309, 440–441, 1984.
- Beeson, M. H., A computer program for processing electron microprobe data, *U.S. Geol. Surv. Open File Rep.*, 40 pp., 1967.
- Bence, A. E., and A. L. Albee, Empirical correction factors for the electron microanalysis of silicates and oxides, *J. Geol.*, 76, 382–403, 1968.
- Brown, G. M., Pyroxenes from the early and middle stages of fractionation of the Skaergaard intrusion, East Greenland, *Mineral. Mag.*, 31, 511–543, 1957.
- Bryan, W. B., Regional variation and petrogenesis of basalt glasses from the FAMOUS area, Mid-Atlantic Ridge, *J. Petrol.*, 20, 293–325, 1979.
- Bryan, W. B., and J. G. Moore, Compositional variations of young basalts in the Mid-Atlantic Ridge rift valley near lat 36°49'N, *Geol. Soc. Am. Bull.*, 88, 556–570, 1977.
- Bryan, W. B., G. Thompson, and P. J. Michael, Compositional variation in a steady-state zoned magma chamber: Mid-Atlantic Ridge at 36°50'N, *Tectonophysics*, 55, 63–85, 1979.
- Byerly, G., The nature of differentiation trends in some volcanic rocks from the Galapagos spreading center, *J. Geophys. Res.*, 85, 3797–3810, 1980.

- Byerly, G. R., W. G. Melson, and P. R. Vogt, Rhyodacites, andesites, ferrobasalts and ocean tholeiites from the Galapagos spreading center, *Earth Planet. Sci. Lett.*, **30**, 215–221, 1976.
- Byers, C. D., D. W. Muenow, and M. O. Garcia, Volatiles in basalts and andesites from the Galapagos Spreading Center, 85° to 86°W, *Geochim. Cosmochim. Acta*, **47**, 1551–1558, 1983.
- Byers, C. D., D. M. Christie, D. W. Muenow, and J. M. Sinton, Volatile contents and ferric-ferrous ratios of basalt, ferrobasalt, andesite and rhyodacite glasses from the Galapagos 95.5°W propagating rift, *Geochim. Cosmochim. Acta*, **48**, 2239–2245, 1984.
- Carmichael, I. S. E., The iron-titanium oxides of salic volcanic rocks and their associated ferromagnesian silicates, *Contrib. Mineral. Petrol.*, **14**, 36–64, 1967.
- Carstens, H., Note on the distribution of some minor elements in coexisting ortho- and clinopyroxene, *Nor. Geol. Tidsskr.*, **38**, 257–260, 1958.
- CAYTROUGH, Geological and geophysical investigation of the Mid-Cayman Rise spreading center: Initial results and observations, in *Deep Drilling Results in the Atlantic Ocean: Ocean Crust Maurice Ewing Ser.*, vol. 2. Edited by M. Talwani, C. E. Harrison, and D. E. Hayes, pp. 66–95, AGU, Washington, D. C., 1979.
- Clague, D. A., and T. E. Bunch, Formation of ferrobasalt at East Pacific midocean spreading centers, *J. Geophys. Res.*, **81**, 4247–4256, 1976.
- Clague, D. A., F. A. Frey, G. Thompson, and S. Rindge, Minor and trace element geochemistry of volcanic rocks dredged from the Galapagos spreading center: Role of crystal fractionation and mantle heterogeneity, *J. Geophys. Res.*, **86**, 9469–9482, 1981.
- Clague, D. A., J. S. Eaby, and J. R. Delaney, Petrologic evidence for long-lived axial magma chambers on the Juan de Fuca Ridge, paper presented at the Fifth Annual Geodynamics Research Symposium on Oceanic Lithosphere: Origin, Structure, and Dynamics, Tex. A&M Univ., College Station, Tex., 1983.
- Clocchiatti, R., Les liquidus silicates pieges dans lse cristaux d'olivine, de plagioclase et de pyroxene: prise d'essai du magma. Application a un basalte a affinite tholeitique de la ride oceanique emergee d'Asal, *C. R. Hebd. Seances Acad. Sci.*, **284**, 2203–2206, 1977.
- Clocchiatti, R., Glassy inclusions in plagioclase and pyroxene phenocrysts in the chilled margin of a pillow lava from hole 417D, Deep Sea Drilling Project, *Initial Rep. Deep Sea Drill. Proj.*, 51–53, part 2, 1063–1067, 1980.
- Cohen, R. S., and R. K. O'Nions, The lead, neodymium and strontium isotopic structure of ocean ridge basalts, *J. Petrol.*, **23**, 299–324, 1982.
- Delaney, J. R., D. W. Muenow, and D. G. Graham, Abundance and distributions of water, carbon and sulfur in the glassy rims of submarine pillow basalts, *Geochim. Cosmochim. Acta*, **42**, 581–594, 1978.
- Delaney, J. R., J. L. Karsten, and D. A. Clague, Geochemistry of Juan de Fuca Ridge basaltic glasses, *Eos Trans. AGU*, **63**, 1147, 1982.
- Detrick, R. S., and W. S. Lynn, The origin of high-amplitude magnetic anomalies at the intersection of the Juan de Fuca Ridge and Blanco Fracture Zone, *Earth Planet. Sci. Lett.*, **26**, 105–113, 1975.
- DeVore, G. W., Crystal growth and the distribution of elements, *J. Geol.*, **63**, 471–494, 1955.
- DeVore, G. W., The association of strongly polarizing cations with weakly polarizing cations as a major influence in element distribution, mineral composition, and crystal growth, *J. Geol.*, **65**, 178–195, 1957.
- Drake, M. J., Plagioclase-melt equilibria, *Geochim. Cosmochim. Acta*, **40**, 457–465, 1976.
- Dungan, M. A., and J. M. Rhodes, Residual glasses and melt inclusions in basalts from DSDP legs 45 and 46: Evidence for magma mixing, *Contrib. Mineral. Petrol.*, **67**, 417–431, 1978.
- Eaby, J. S., and D. A. Clague, Preliminary description of basalt from the southern Juan de Fuca Ridge, *U.S. Geol. Surv. Open File Rep.*, **82-200C**, 17 pp., 1982.
- Eaby, J., D. A. Clague, and J. R. Delaney, Sr isotopic variations along the Juan de Fuca Ridge, *J. Geophysical Res.*, **89**, 7883–7890, 1984.
- Engel, C. G., and R. L. Fisher, Granitic to ultramafic rock complexes of the Indian Ocean ridge system, western Indian Ocean, *Geol. Soc. Am. Bull.*, **86**, 1553–1578, 1975.
- Flower, M. F. J., Cryptocumulate tholeiite as evidence for magma mixing at an intermediate-rate spreading axis, *Nature*, **229**, 542–545, 1982.
- Fornari, D. J., M. R. Perfit, A. Malahoff, and R. Embley, Geochemical studies of abyssal lavas recovered by DSRV Alvin from eastern Galapagos rift, Inca transform, and Ecuador rift, 1, Major element variations in natural glasses and spacial distribution of lavas, *J. Geophys. Res.*, **88**, 10,519–10,529, 1983.
- Fox, P. J., and J. B. Stroup, The plutonic foundation of the oceanic crust, in *The Sea*, vol. 7, *The Oceanic Lithosphere*, edited by C. Emiliani, pp. 119–218, Wiley-Interscience, New York, 1981.
- Ghose, S., I. S. McCallum, and E. Tidy, Luna 20 pyroxenes: Exsolution and phase transformation as indicators of petrologic history, *Geochim. Cosmochim. Acta*, **37**, 831–839, 1973.
- Haggerty, S. E., Opaque mineral oxides in terrestrial igneous rocks, *Short Course Notes Mineral. Soc. Am.*, **3**, H-101–H-169, 1976.
- Harlow, G. E., C. E. Nehru, M. Prinz, G. J. Taylor, and K. Keil, Pyroxenes in Serra de Mage: Cooling history in comparison with Moama and Moore County, *Earth Planet. Sci. Lett.*, **43**, 173–181, 1979.
- Hebert, R., D. Bibeau, and R. Hekinian, Ultramafic and mafic rocks from the Garret Transform fault near 13°30'S on the East Pacific Rise: igneous petrology, *Earth Planet. Sci. Lett.*, **65**, 107–125, 1983.
- Hekinian, R., and M. Hoffert, Rate of palagonitization and manganese coating on basaltic rocks from the rift valley in the Atlantic ocean near 36°50' north, *Mar. Geol.*, **19**, 91–109, 1975.
- Helz, R. T., Crystallization history of Kilauea Iki lava lake as seen in drill core recovered in 1967–1979, *Bull. Volcanol.*, **43**, 675–701, 1980.
- Hess, H. H., Stillwater igneous complex, Montana: A quantitative mineralogical study, *Mem. Geol. Soc. Am.*, **80**, 230 pp., 1960.
- Hodges, F. N., and J. J. Papike, DSDP site 334: Magmatic cumulates from oceanic layer 3, *J. Geophys. Res.*, **81**, 4135–4151, 1976.
- Huppert, H. E., and J. S. Turner, A laboratory model of a replenished magma chamber, *Earth Planet. Sci. Lett.*, **54**, 144–152, 1981.
- Ishii, T., Zoning, exsolution and phase equilibria in the Kintoki-san pyroxenes, Hakone volcano, paper presented at 80th Annual Meeting, Geol. Soc. of Jpn., Sendai, 1973.
- Ishii, T., and H. Takeda, Inversion, decomposition and exsolution phenomena of terrestrial and extraterrestrial pigeonites, *Geol. Soc. Jpn. Mem.*, **11**, 19–36, 1974.
- Johannes, W., Melting of plagioclase in the systems Ab-An-H₂O and Qz-Ab-An-H₂O at PH₂O = 5 kbars: An equilibrium problem, *Contrib. Mineral. Petrol.*, **66**, 295–303, 1978.
- Kay, R., N. J. Hubbard, and P. W. Gast, Chemical characteristics and origin of oceanic ridge volcanic rocks, *J. Geophys. Res.*, **75**, 1585–1613, 1970.
- Koski, R. A., R. Goodfellow, and R. M. Bouse, Preliminary description of massive sulfide samples from the Juan de Fuca Ridge, *U.S. Geol. Surv. Open File Rep.*, **82-200B**, 21 pp., 1982.
- Kuno, H., Review of pyroxene relations in terrestrial rocks in the light of recent experimental works, *Mineral. J.*, **5**, 21–43, 1966.
- Larsen, J. G., Glass-bearing gabbro inclusions in hyaloclastites from Tindfjallajökull, Iceland, *Lithos*, **12**, 289–302, 1979.
- LeRoex, A. P., H. J. B. Dick, A. J. Erlank, A. M. Reid, F. A. Frey, and S. R. Hart, Geochemistry, mineralogy and petrogenesis of lavas erupted along the southwest Indian ridge between the Bouvet triple junction and 11 degrees east, *J. Petrol.*, **24**, 267–318, 1983.
- Lichtman, G. S., W. R. Normark, J. R. Delaney, J. L. Morton, H. P. Johnson, and J. L. Karsten, Photogeology and evolution of the Juan de Fuca Ridge, *U.S. Geol. Surv. Open File Rep.*, **83-464**, 14 pp., 1983.
- Liias, R. A., and J. M. Rhodes, Evolution of basaltic magma along the Juan de Fuca Ridge, *Eos Trans. AGU*, **63**, 1147, 1982.
- Lindsley, D. H., Pyroxene thermometry, *Am. Mineral.*, **68**, 477–493, 1983.
- Lindsley, D. H., and D. J. Andersen, A two-pyroxene thermometer, *Proc. Lunar Planet. Sci. Conf. 13th*, Part 2, *J. Geophys. Res.*, **88**, suppl., A887–A906, 1983.
- Lindsley, D. H., and K. J. Spencer, Fe-Ti oxide geothermometry: Reducing analyses of coexisting Ti-magnetite (Mt) and ilmenite (Ilm), *Eos Trans. AGU*, **63**, 471, 1982.
- Macdonald, K. C., Mid-ocean ridges: Fine scale tectonic, volcanic and hydrothermal processes within the plate boundary zone, *Annu. Rev. Earth Planet. Sci.*, **10**, 155–190, 1982.
- Malcolm, F. L., Microstructures of the Cayman Trough gabbros, *J. Geol.*, **89**, 675–688, 1981.
- Mattey, D. P., and I. D. Muir, Geochemistry and mineralogy of basalts from the Galapagos spreading center, Deep Sea Drilling Project leg 54, *Initial Rep. Deep Sea Drill. Proj.*, **54**, 755–772, 1980.
- Mazzullo, L. J., and A. E. Bence, Abyssal tholeiites from DSDP leg 34: The Nazca plate, *J. Geophys. Res.*, **81**, 4327–4351, 1976.
- Miyashiro, A., and F. Shido, Differentiation of gabbros in the Mid-Atlantic Ridge near 24°N, *Geochem. J.*, **14**, 145–154, 1980.

- Morton, J. L., Oceanic spreading centers: Axial magma chambers, thermal structure, and small scale ridge jumps, Ph. D. thesis, 101 pp., Stanford Univ., Stanford, Calif., 1984.
- Morton, J. L., J. S. Eaby, W. R. Normark, J.-P. Eissen, M. Golan-Bac, J. B. Rapp, and J. M. Been, Basalt samples and water-column samples from the southern Juan de Fuca Ridge: Preliminary results of cruise L13-82-WF, *U.S. Geol. Surv. Open File Rep.*, 83-827, 15 pp., 1983.
- Natland, J. H., Effect of axial magma chambers beneath spreading centers on the compositions of basaltic rocks, *Initial Rep. Deep Sea Drill. Proj.*, 54, 833-850, 1980.
- Natland, J. H., A compositionally nearly steady-state magma chamber at the Costa Rica Rift: Evidence from basalt glass and mineral data, Deep Sea Drilling Project sites 501, 504, 505, *Initial Rep. Deep Sea Drill. Proj.*, 69, 811-858, 1983.
- Normark, W. R., J. E. Lupton, J. W. Murray, R. A. Koski, D. A. Clague, J. L. Morton, J. R. Delaney, and H. P. Johnson, Poly-metallic sulfide deposits and water-column tracers of active hydrothermal vents on the southern Juan de Fuca Ridge, *Mar. Technol. Soc. J.*, 16, 46-53, 1982.
- O'Hara, M. J., Geochemical evolution during fractional crystallization of a periodically refilled magma chamber, *Nature*, 266, 503-508, 1977.
- Pallister, J. S., and C. A. Hopson, Samail ophiolite plutonic suite: Field relations, phase variation, cryptic variation and layering, and a model of a spreading ridge magma chamber, *J. Geophys. Res.*, 86, 2593-2644, 1981.
- Papike, J. J., and A. E. Bence, Apollo 14 inverted pigeonites: Possible samples of lunar plutonic rocks, *Earth Planet. Sci. Lett.*, 14, 176-182, 1972.
- Perfit, M. R., and D. J. Fornari, Geochemical studies of abyssal lavas recovered by DSRV *Alvin* from eastern Galapagos rift, Inca transform, and Ecuador rift, 2, Phase chemistry and crystallization history, *J. Geophys. Res.*, 88, 10,530-10,550, 1983.
- Perfit, M. R., D. J. Fornari, A. Malahoff, and R. W. Embley, Geochemical studies of abyssal lavas recovered by DSRV *Alvin* from eastern Galapagos rift, Inca transform, and Ecuador rift, 3, Trace element abundances and petrogenesis, *J. Geophys. Res.*, 88, 10,551-10,572, 1983.
- Pinckney, L. R., and D. H. Lindsley, Effects of magnesium on iron-titanium oxides, *Geol. Soc. Am. Abstr. Programs*, 8 1051, 1976.
- Richter, D. H., and J. G. Moore, Petrology of the Kilauea Iki lava lake, Hawaii, *U.S. Geol. Surv. Prof. Pap.*, 537-B, 26 pp., 1966.
- Robinson, P., The composition space of terrestrial pyroxenes—Internal and external limits, in *Pyroxenes*, *Rev. Mineral.*, vol. 7, edited by C. T. Prewitt, pp. 419-475, Mineralogical Society of America, Washington, D. C., 1980.
- Roedder, E., Origin and significance of magmatic inclusions, *Bull. Mineral.*, 102, 487-520, 1979.
- Roeder, P. L., Activity of iron and olivine solubility in basaltic liquids, *Earth Planet. Sci. Lett.*, 23, 397-410, 1974.
- Ross, M., and J. S. Huebner, Temperature-composition relationships between naturally occurring augite, pigeonite and orthopyroxene at one bar pressure, *Am. Mineral.*, 64, 1133-1155, 1979.
- Sato, H., K. Aoki, K. Odamoto, and B. Fujita, Petrology and chemistry of basaltic rocks from hole 396B, IPOD/DSDP leg 46, *Initial Rep. Deep Sea Drill. Proj.*, 46, 115-141, 1978.
- Scheidegger, K. F., Temperatures and compositions of magmas ascending along midocean ridges, *J. Geophys. Res.*, 78, 3340-3355, 1973.
- Schilling, J.-G., G. Cousens, R. L. Chase, and J. R. Delaney, Juan de Fuca-Explorer Ridge: Rare earth contents, *Eos Trans. AGU*, 63, 1154, 1982.
- Sinton, J. M., D. S. Wilson, D. M. Christie, R. N. Hey, and J. R. Delaney, Petrologic consequences of rift propagation on oceanic spreading ridges, *Earth Planet. Sci. Lett.*, 62, 193-207, 1983.
- Sobolev, V. S., and V. P. Kostyuck, Magmatic crystallisation based on a study of melt inclusions (in Russian), Nauka Press, Novosibirsk, 1975. (Translated in part in *Fluid Inclusion Res.*, 9, 182-253, 1975.)
- Sparks, R. S. J., P. Meyer, and H. Sigurdsson, Density variation amongst mid-ocean ridge basalts: Implications for magma mixing and the scarcity of primitive lavas, *Earth Planet. Sci. Lett.*, 46, 419-430, 1980.
- Stakes, D. S., J. W. Shervais, and C. A. Hopson, The volcanic-tectonic cycle of the FAMOUS and AMAR valleys, Mid-Atlantic Ridge (36°47'N): Evidence from basalt glass and phenocrysts compositional variations for a steady state magma chamber beneath the valley midsections, AMAR 3, *J. Geophys. Res.*, 89, 6995-7028, 1984.
- Stormer, J. C., Jr., The effects of recalculation on estimates of temperature and oxygen fugacity from analyses of multicomponent iron-titanium oxides, *Am. Mineral.*, 68, 586-594, 1983.
- Tiezi, L. J., and R. B. Scott, Crystal Fractionation in a cumulate gabbro, Mid-Atlantic Ridge, 26°N, *J. Geophys. Res.*, 85, 5438-5454, 1980.
- Truter, F. C., Modern concepts of the Bushveld Igneous Complex, *Rep. 1*, pp. 77-92, Comm. for Tech. Coop. in Afr., South. Reg. Comm. of Geol., Salisbury, 1955.
- Usselman, T. M., and D. S. Hodge, Thermal control of low-pressure fractionation processes, *J. Volcanol. Geotherm. Res.*, 4, 265-281, 1978.
- van Andel, T. H., and R. D. Ballard, The Galapagos Rift at 86°W, 2, Volcanism, structure, and evolution of the rift valley, *J. Geophys. Res.*, 84, 5390-5406, 1979.
- Vanko, D. A., and R. Batiza, Gabbroic rocks from the Mathematician Ridge failed rift, *Nature*, 300, 742-744, 1982.
- Verma, S. P., and J.-G. Schilling, Galapagos hot spot-spreading center system, 2, ⁸⁷Sr/⁸⁶Sr and large ion lithophile element variations (85°W-101°W), *J. Geophys. Res.*, 87, 10,838-10,856, 1982.
- Vogt, P. R., and G. R. Byerly, Magnetic anomalies and basalt composition in the Juan de Fuca-Gorda Ridge Area, *Earth Planet. Sci. Lett.*, 33, 185-207, 1976.
- Wager, L. R., and W. A. Deer, Geological investigations in East Greenland, III, The petrology of the Skaergaard intrusion, Kangerdlugssuaq, East Greenland, *Medd. Grøn.*, 105, 1-352, 1939.
- Wakeham, S. E., Petrochemical patterns in young pillow basalts dredged from Juan de Fuca and Gorda ridges, M. S. thesis, Oregon State Univ., Corvallis, 1977.
- Walker, D., T. Shibata, and S. E. DeLong, Abyssal tholeiites from the Oceanographer Fracture Zone, II, Phase equilibria and mixing, *Contrib. Mineral. Petrol.*, 70, 111-125, 1979.
- Walker, K. R., N. G. Ware, and J. F. Lovering, Compositional variations in the pyroxenes of differentiated Palisades Sill, New Jersey, *Geol. Soc. Am. Bull.*, 84, 89-110, 1973.
- Watson, E. B., Glass inclusions as samples of early magmatic liquid: Determinative method and application to a South Atlantic basalt, *J. Volcanol. Geotherm. Res.*, 1, 73-84, 1976.
- Wilson, D. S., R. N. Hey, and C. Nishimura, Propagation as a mechanism of reorientation of the Juan de Fuca Ridge, *J. Geophys. Res.*, 89, 9215-9225, 1984.
- Wright, T. L., and R. T. Okamura, Cooling and crystallization of tholeiitic basalt, 1965 Makopuhi lava lake, Hawaii, *U.S. Geol. Surv. Prof. Pap.*, 1004, 78 pp., 1977.
- Wright, T. L., and D. L. Peck, Crystallization and differentiation of the Alae magma, Alae lava lake, Hawaii, *U.S. Geol. Surv. Prof. Pap.*, 935-C, 20 pp., 1978.

D. A. Clague, U.S. Geological Survey, 345 Middlefield Road, MS/999, Menlo Park, CA 94025.

J. E. Dixon, Division of Geological and Planetary Sciences, 170-25, California Institute of Technology, Pasadena, CA 91125.

J.-P. Eissen, ORSTOM, B.P. A5, Noumea, New Caledonia.

(Received July 22, 1985;
revised October 15, 1985;
accepted November 1, 1985.)