40. CHEMICAL STRATIGRAPHY OF BOREHOLES IN THE IZU-BONIN ARC FROM IN SITU NUCLEAR MEASUREMENTS¹

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ABSTRACT

In situ nuclear measurements were made in open-hole sections of Ocean Drilling Program Holes 792E and 793B. They provide quantitative estimates of a limited range of formation elements. A comparison of core and geochemical log data indicates reasonable agreements between the two. Limited analyses of a wide range of core samples indicate that although the chalk and nannofossil-rich silts are distinct, the admixture of sands, clays, and volcanogenic debris provides a range of homogeneous sediment compositions. The chemical homogeneity does not allow detailed distinct zoning on a simple chemical basis. Statistical analysis of the log data, however, reveals distinct and recognizable patterns that enable a chemostratigraphy to be derived for both sites. This chemostratigraphy is in agreement with the shipboard-derived lithologic units. It reveals considerable detail that may act as the focus for further evaluation.

INTRODUCTION

Site 792 is located on the western half of the Izu-Bonin forearc sedimentary basin, about 60 km east of the arc volcano Aoga Shima and 170 km west of the axis of the Izu-Bonin Trench. The location is upslope from a fork in the Aoga Shima Canyon, where the forearc sediments lap onto the edge of a basement high. The site was drilled five times in succession, achieving a total penetration of 885.9 mbsf; however, as a consequence of a stuck pipe, the downhole geochemistry is only in open hole from approximately this depth up to 286 mbsf. Six lithologic units were described based on the material recovered at Site 792; these are described in Table 1 (note that Unit I is not considered here as it does not occur in the open section of Hole 792E).

Site 793 is located in the center of the Izu-Bonin forearc sedimentary basin, 125 km west of the axis of the Izu-Bonin Trench; it lies in an interchannel area on the southern side of the broad Sumisu Jima Canyon. The site was drilled twice; Hole 793B was spudded with a reentry cone and penetrated to a total depth of 1682.0 mbsf. Downhole geochemical data were obtained in open hole between 1531.0 and 1034.0 mbsf. The interval between 99.7 and 586.5 mbsf was not cored; consequently, seven lithologic units were described from core obtained outside this region. These are described in Table 2, although only the lowermost Units V, VI, and VII correspond with the interval in which *in situ* geochemistry was measured in open hole.

In situ chemistry is measured through the deployment of a series of nuclear logging tools developed by Schlumberger. Full details of the suite of tools are given elsewhere (Hertzog et al., 1987; Anderson et al., 1990; Lovell and Anderson, 1988), but they basically consist of four measurement devices that yield intensity data for uranium, thorium, and potassium (natural gamma-spectroscopy tool); aluminum (aluminum activation tool); silicon, carbon, iron, sulfur, titanium, gadolinium, hydrogen, and chlorine (gamma spectroscopy tool). Measurements are made at 15-cm intervals downhole and are processed into weight percent oxides (except for U, Th, and Gd which are measured in ppm). During processing the data are smoothed such that the individual measurements relate to a volume of rock approximately two-thirds of a meter in diameter. The calibration of these comparatively novel geochemical logs has been the subject of considerable discussion, not least because of the difficulty of comparing different measurement techniques, sample bias in terms of volume and content, and the inherent problem created by the hostile downhole environment (see, for example, Harvey and Lovell, 1989). The chemistry of the formation is, however, important in addition to these technical difficulties. Although the log measurements have now been successfully calibrated in many sedimentary and basement formations (see, for example, Anderson et al., 1990), their application in regions of volcaniclastic sediments is hitherto not documented.

CORE-LOG COMPARISON

Table 3 presents a comparison of core and log geochemistry for a limited number of sample positions for which core was available in open-hole sections and for which geochemical logging tool (GLT) data existed. Figure 1 shows plots of core vs. GLT data for four of the oxides used in this study. In total, only seven geochemical variables were used in this preliminary study; they were chosen according to previous experience with GLT data in terms of expected primary mineralogy. A direct comparison of core and log data is difficult because of the inherent differences in the two measurement techniques, together with the errors involved in depth location of recovered core samples. Overall, however, the data show tolerably good agreement for most elements; elsewhere, however, in the sections of both holes, negative values occur in the GLT-derived values for SiO₂, TiO₂, and CaO; this is a result of processing difficulties described in Pratson et al. (this volume). In particular, the large diameter hole, combined with high-porosity sediments, is detrimental to the in situ measurement, resulting in a high hydrogen signal as well as poor counting statistics for the remaining elements.

Individual comparisons show the following:

1. SiO₂ has a limited range (40%–60%) in the cores but a much larger range (30%–75%) and typically higher (10%–15%) values in the GLT data.

2. Al_2O_3 shows good agreement with most values, lying close to a 1:1 line in Figure 1. At Site 792 some GLT values appear higher than the core values.

3. Although average GLT values for TiO_2 are reasonable, high (6.8%) values in the Hole 792E GLT data have yet to be substantiated by the core data. Negative values in the GLT data are a particular problem.

 FeO shows insignificant correlation, but the GLT data show good agreement in the 1%–6% range. The lack of correlation appears

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Table 1. Lithostratigraphy of Site 792.

Unit	Depth (mbsf)	Description				
I	0–183.7	upper Pliocene to Holocene nannofossil-rich, vitric silty clay and a clayey silt,				
п	183.7–357.4	interbedded with vitric silts and sands and minor pumiceous and scoriaceous gravels. middle and upper Miocene sandy mudstone, muddy sandstone, and silty claystone (all with nannofossil-rich claystone and				
ш	357.4-429.3	nannofossil chalk, and rare crystal-vitric siltstone and sandstone. upper Oligocene to middle Miocene intensely bioturbated, nannofossil-rich claystone and nannofossil chalk and rare crystal-vitric				
IV	429.3–783.4	siltstone and sandstone. upper Oligocene vitric sandstone and volcanic sandy conglomerate containing claystone interclasts, with minor silty claystone and				
v	783.4-804.0	claystone with nannofossil-rich intervals. Altered volcanic sandstone with claystone intraclasts				
VI	804.0-885.9	Porphyritic andesite with minor basaltic andesite and dacite.				

Table 2. Lithostratigraphy of Site 793.

Unit or subunit	Depth (mbsf)	Description		
IA	0-32.5	Pleistocene		
IB	32.5-99.7	Sediments		
п	586.5-591	Olivine-clinopyroxene-orthopyroxene diabase		
ш	591-735.7	lower to middle Miocene nannofossil-rich silty		
IV	735.7–759.0	claystone, vitric siltstone, and sandstone nannofossil rich clayey siltstone. lower Miocene claystone, nannofossil, and		
		nannofossil-rich claystone, and vitric siltstone.		
v	759.0-1373.1	upper lower and upper Oligocene vitric		
		sandstone, purniceous sandstone, granule- to fine-pebble conglomerate, siltstone, clavey siltstone, and silty claystone.		
VI	1373.1-1403.9	upper lower Oligocene, very poorly suited volcanic breccia with sandy matrix and		
		mixed fresh to altered clasts of mainly plagioclase-rich andesite.		
VII	1403.9–1682	upper-lower Oligocene breccias and massive to pillowed flows of porphyritic		
		clinopyroxene-orthopyroxene and aphyric,		
		basaltic andesites and andesites.		

to be caused by a small number of samples, mostly from Hole 792E with very high FeO yields. These may be real, as they are in other ODP holes where they have been associated with fracture and brecciation zones.

5. CaO shows reasonable but questionable agreement as a high negative value for the minimum of the full GLT data set is present. This is related to only a few sample points and is probably a consequence of overestimating the high SiO_2 concentrations during the closure model.

 K₂O exhibits reasonable agreement between the GLT and core despite low concentrations.

Attempts to consider the geochemistry of individual sedimentary facies rely heavily on the results of shipboard analyses (Schlich, Wise, et al., 1990); these criteria are inevitably subject to modification following more extensive shore-based studies. Table 4 and Figure 2 show the results of a classification of the laboratory XRF data in terms of gross sedimentary description. The similarity between the majority of the units, excepting the chalk and nannofossil-rich horizons, is notable. This is accentuated when only the elements measurable downhole by the GLT are considered; the basement basaltic rock, for

instance, are only clearly distinguished in Table 4 by their higher MgO and lower Na₂O contents.

DOWNHOLE GEOCHEMISTRY

Given the apparent uniformity of the different sediments in terms of the downhole geochemistry, it is inevitable that the downhole logs show little character when compared with the main lithologic units (see Pratson et al., this volume). Figures 3 and 4 show the log responses for each hole. The main features are related respectively at each site to the location of a major fault zone and the transition into basement. In Hole 792E the dominant characteristics of the downhole geochemistry appear to be related to individual horizons that show peaks in K2O, Th, and U. An alternative approach to the variation in downhole geochemistry is to search for "natural" geochemical groups. The data have, therefore, been subjected to an iterative, nonhierarchical cluster analysis (INCA) using a Euclidean distance model on z-standardized measurements. Initially, 20 groups were fitted; a distinct break in the mean-squared deviation between groups occurs at Groups 11 (Hole 792E) and 12 (Hole 793B). Further details of the method are given in Lebart et al. (1984), Shepherd et al. (1987), and Harvey and Lovell (1989).

The results of these analyses are plotted in Figures 5 and 6 for Holes 792E and 793B, respectively. The group centroid compositions are given in Table 5 and 6. In each case, a chemostratigraphy is derived from the relative proportions of different groups. The chemostratigraphy is also plotted in the same figures alongside the major lithologic units.

At Site 792 the derived chemostratigraphy shows excellent agreement with the shipboard descriptions. In addition to picking out each individual unit, the chemostratigraphy highlights the fault zone and the transition to basement. Indeed, the INCA classification suggests far more detail exists than originally observed; integration of these results with further detailed core descriptions and high-resolution *in situ* FMS images may enhance our interpretation of the chemostratigraphy. At Site 793 the open-hole section only penetrates three separate units. Again the chemostratigraphy provides far more detail, which requires both a reappraisal of the core and a consideration of finer sedimentological aspects.

Further detailed analysis of the GLT data, including an appraisal of the errors likely in this type of environment, is necessary to quantify the significance of the derived chemostratigraphy. The detailed lithostratigraphy derived from the FMS images (see Hiscott et al., this volume) will be critical in this calibration exercise, in that it provides the missing boundaries to units otherwise poorly sampled by coring.

CONCLUSIONS

A comparison of the core and geochemical log data indicates reasonable agreement between the two within the limited range of samples available for the comparison, although the estimates for silica are probably too high in the log data, and some queries still remain about the processing of the GLT data.

Initial, but very limited, geochemical data on a wide range of core samples indicate that, although the chalk and nannofossil-rich silts are not unexpectedly very distinct, the admixture of sands, clays, and volcanogenic debris leads to a range of homogeneous sediment compositions. Considerably more core-based geochemical work will be necessary to characterize these unusual sediments fully.

As a consequence of the relative chemical homogeneity among the core samples, an initial examination of the geochemical logs for both sites shows no distinct or simple pattern relating to the geology of the sites except for the recognition of a prominent fault zone and the sediment/basement interface at Site 792.

Statistical analyses of the log data have revealed, however, that distinct and recognizable patterns in the GLT logs exist from which a chemostratigraphy has been derived for both sites. The derived chemostratigraphy, at both sites, is in excellent agreement with the shipboard-derived

	GLT			Core				
	Mean	SD	Min.	Max.	Mean	SD	Min.	Max.
Hole 792E:				New York	A RECEIPTION			-
SiO ₂	57.19	11.15	30.95	71.48	47.37	8.71	16.50	57.99
Al ₂ O ₃	18.10	4.41	8.12	23.60	15.19	3.71	3.99	19.50
TiO ₂	0.66	1.08	-2.32	2.48	0.63	0.19	0.13	0.87
FeO	10.49	4.56	6.17	26.26	7.84	2.12	2.19	10.63
CaO	10.24	8.35	0.83	34.43	10.39	8.54	2.40	40.93
K ₂ O	1.10	0.79	0.27	2.88	0.88	0.69	0.14	2.55
S	0.81	3.02	0.00	14.06	0.15	0.19	0.05	1.04
Hole 793B:								
SiO ₂	66.01	5.53	56.28	75.78	52.80	2.39	49.65	57.78
Al ₂ O ₃	15.67	1.97	12.54	18.36	14.89	1.68	11.73	17.48
TiO ₂	1.10	0.35	0.40	1.55	0.54	0.14	0.27	0.74
FeO	9.30	2.07	6.19	13.47	7.41	0.90	5.56	8.51
CaO	5.95	2.80	1.23	12.06	7.65	1.39	5.31	9.81
K ₂ O	0.90	0.27	0.55	1.32	0.78	0.32	0.33	0.27
S	1.08	1.30	0.00	5.28	0.05	0.03	0.02	0.12

Table 3. Comparison of core and geochemical logging tool (GLT) data, Holes 792E and 793B.

Notes: Mean for Hole 792E is N = 27 and for Hole 793B, N = 14. SD = standard deviation, Min. = minimum, and Max. = maximum. Geochemistry of core samples was determined by XRF at the University of Nottingham.



Figure 1. Comparison of core and geochemical logging tool (GLT) data. Open triangles = Hole 792E, and open squares = Hole 793B.

Table 4. Average of core sample groups from the major lithologic units.

		Nannofossil-rich claystone				
	Chalk	А	в	С	Clastics	Basalts
SiO ₂	28.30	44.16	51.06	56.28	50.20	54.18
Al ₂ O ₃	7.90	10.79	14.92	13.02	16.39	13.57
TiO ₂	0.33	0.45	0.67	0.64	0.66	0.37
Fe ₂ O ₃	4.36	6.18	8.29	6.80	8.34	8.94
MgO	2.63	2.51	4.77	3.77	5.56	8.81
CaO	28.35	17.83	7.72	7.26	7.10	7.48
Na ₂ O	2.34	2.08	2.74	2.51	3.74	1.96
K ₂ O	0.38	1.01	1.48	0.97	0.96	0.87
MnO	0.23	0.30	0.26	0.15	0.22	0.17
P2O5	0.13	0.12	0.12	0.11	0.11	0.01
LOI	25.14	14.55	7.74	8.27	6.56	3.42
N	3	5	10	5	20	4

N = number of core samples per group. A, B, and C distinguished by visual grain-size description ranging from fine (A) to coarse (C).

Table 5. Centroid composition determined from the INCA classification of GLT responses from Hole 792E, open section.

Group	Α	В	С	D	E	F
SiO ₂	55.72	44.66	45.46	66.32	69.48	44.40
Al2O3	19.64	14.47	14.97	11.68	16.58	14.05
TiO ₂	1.49	-0.67	2.17	1.61	0.94	2.24
FeO	10.96	13.47	16.27	11.42	14.56	
K ₂ O	0.85	1.50	0.86	0.63	0.62	1.07
S	0.17	1.34	0.63	0.37	0.05	11.83
N	558	106	136	314	830	77
Group	G	н	I	J	К	
SiO ₂	61.81	16.37	31.85	63.85	61.72	
Al2O3	22.42	17.88	15.61	16.07	19.08	
TiO ₂	1.12	2.84	-0.58	-0.05	1.17	
FeO	7.79	23.07	14.09	8.79	8.05	
CaO	5.20	12.60	24.21	9.16	6.37	
K ₂ O	0.69	2.08	1.92	0.89	2.06	
S	0.08	20.33	12.68	0.18	0.16	
N	894	17	47	366	415	

Notes: The INCA was run using a Euclidian model with z-standardization and using the GLT measurements for SiO₂, Al₂O₃, TiO₂, FeO, CaO, K₂O, and S. N = number of responses allocated to each group; number of responses classified = 3760.

Table 6. Centroid compositions determined from the INCA classification of GLT responses for Hole 793B, open section.

Group	Α	в	С	D	Е	F
SiO ₂	74.05	64.93	74.03	64.51	70.76	54.42
Al2O3	4.96	15.94	12.41	15.16	15.67	15.88
TiO ₂	0.48	0.84	0.41	1.12	0.97	1.05
FeO	9.49	8.13	8.89	8.77	6.93	13.56
CaO	8.90	8.12	2.80	4.98	4.24	5.95
K ₂ O	0.81	0.88	1.22	0.90	0.91	1.02
S	1.31	1.18	0.25	4.5	0.53	8.09
N	125	510	340	42	430	7
Group	G	Н	I	J	К	L
SiO ₂	64.77	59.46	68.05	57.51	61.44	66.83
Al2O3	18.56	16.42	14.00	4.06	22.79	14.46
TiO ₂	1.20	1.14	0.48	-2.36	1.54	0.62
FeO	9.07	11.42	10.68	17.21	9.17	9.66
CaO	4.95	9.41	5.29	19.44	0.41	5.70
K ₂ O	0.87	0.92	1.00	1.06	4.66	2.01
S	0.58	1.23	0.51	3.10	0.00	0.73
N	548	308	546	12	19	336

Notes: The INCA was run using a Euclidian model with z-standardization and using the GLT measurements for SiO₂, Al₂O₃, TiO₂, FeO, CaO, K₂O, and S. N = number of responses allocated to each group; number of responses classified = 3198.

lithologic units, but it is somewhat more detailed and provides a set of chemically derived units as well that will act as the focus for further evaluation of the core and other log measurements.

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Figure 2. Geochemical classification of core samples, Holes 792E and 793B. The four points trending toward the CaO position are the nannofossil-rich claystones and chalk samples.



Figure 3. Downhole geochemical logs, Hole 792E.



Figure 4. Downhole geochemical logs, Hole 793B.



Figure 5. Site 792 iterative, nonhierarchical cluster analysis (INCA) classifications. A through K are different geochemical groups; INCA zones are characteristic intervals defined by these groups.



Figure 6. Site 793 iterative, nonhierarchical cluster analysis (INCA) classifications. A through L are different geochemical groups; INCA zones are characteristic intervals defined by these groups.