21. MINERALOGY OF ALTERED BASAL VOLCANICLASTIC SEDIMENTS AT SITES 447 (WEST PHILIPPINE BASIN) AND 450 (PARECE VELA BASIN), DEEP SEA DRILLING PROJECT LEG 59

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ABSTRACT

The authigenic minerals contained in the altered basal intervals of volcaniclastic sediments from Sites 447 and 450 of Deep Sea Drilling Project Leg 59 are dioctahedral smectite (with variable crystallinity), phillipsite, and sanidine. Sanidine seems the most widespread and common product of basal alteration in the Philippine Sea marginal basins. The neomorphic mineral suites may have been produced by (1) halmyrolysis of the volcaniclastic sediments; (2) halmyrolysis of the underlying basalts; or (3) hydrothermalism associated with basaltic intrusions. At Site 450, other authigenic minerals occur (carbonates, analcime, clinoptilolite, Fe-Mn oxides), and the basal paragenesis is consistent with a hydrothermal origin. Such a process could have produced temperatures up to 200°C in the tuffs lying as much as 2 meters above the contact with a basaltic intrusion. Products of low-temperature alteration, however, are also present in the altered interval of this site.

INTRODUCTION

The basal sediments lying immediately above a series of basalt flows are commonly altered and often metal-rich, and can be produced either (1) by hydrothermalism, which may or may not be associated with former spreading centers (Bonatti, 1975), or (2) by sea-water alteration (halmyrolysis) of the basaltic basement (Hummel, 1922; Natland, 1973). Alteration of basal sediments has been studied in recent years utilizing numerous samples from DSDP sites, but pertinent data from marginal back-arc basins are relatively scarce (Bonatti et al., 1978).

This chapter studies the mineralogy of the altered intervals of volcaniclastic sequences cored during DSDP Leg 59 at Sites 447 and 450 in the Philippine Sea complex of marginal basins (Fig. 1). The purpose is to give a detailed account of the suites of neomorphic minerals and to discuss their possible genetic processes.

METHODS

We analyzed powders of bulk samples and oriented slides of clay minerals (<2-µm fraction) with a Philips X-ray diffractometer, using the CuKα and FeKα radiations and a goniometer speed of 1°/2θ per minute. Quantitative estimates of the amorphous and mineral fractions were performed following the procedures given by Cook et al. (1975), but our mineral percentages are calculated on the whole sample (i.e., amorphous matter + X-ray diffractive minerals = 100). The analytic results are reported in Table 1. All samples also have been studied by optical microscopy on smear slides, and some of them have been examined with the scanning electron microscope (SEM).

SITE 447

Hole 447A was drilled in the West Philippine Basin (Fig. 1). A total of 113 meters of sediments was cored before encountering the basaltic basement. The sedimentary column contains, from the mudline core downward:

1) 37.5 meters of a barren, dark brown pelagic clay (Cores 1–4);

2) 9.5 meters of middle-upper Oligocene calcareous pelagic clay (Core 5);

3) 38.0 meters of middle Oligocene carbonate-bearing ruditic units, comprising tuffaceous volcaniclastic breccias and polymeric breccias (Cores 6–9);

4) 2.1 meters of barren, altered, very fine tuff or mudstone, in variegated layers 8 to 40 cm thick with shades of gray, brown, olive, olive gray, and grayish green (Core 10, Section 1 and part of Section 2);

5) 25.9 meters of middle Oligocene tuffs and volcaniclastic breccias and/or conglomerates (Core 10, part of Section 2, through Core 13).

Beneath this unit, the basaltic basement has been cored down to a total sub-bottom depth of 296.5 meters (Cores 14–36).

We analyzed samples from the variegated, altered tuffs of the upper part of Core 10. Apart from the variable colors, these are rather monotonous deposits, which show only diffuse mottling in shades slightly lighter or darker than the background. Seven samples have been taken at each major color change. Figure 2 shows X-ray mineralogy by mineral graphs.

The amorphous fraction ranges from 68% to 84%. The X-ray mineral fraction contains dioctahedral smectite of the beidellite–montmorillonite group, phillipsite, and sanidine. Minor amounts of aragonite crystals and iron–manganese amorphous oxides and hydroxides have been detected on smear slides. Phillipsite appears only in the two lowest samples, close to the bottom of the altered zone. In phillipsite-bearing samples, the amorphous fraction is low and the smectite decreases in amount (from 5–12% to 1–2%) and in crystallinity (v/p ratio from >0.5 to <0.5). Sanidine approaches the theoretical composition (almost pure monoclinic K-feldspar) and is fairly constant (6–12%) throughout Core 10.

All these minerals seem to derive from the alteration of a previous unit rich in volcanic glass. However, our
Figure 1. Schematic location map of the marginal basins of the Philippine Sea with locations of the sites studied in this chapter.

Table 1. Quantitative X-ray mineralogy of the analyzed samples.

<table>
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<tr>
<th>Sample (intervals in cm)</th>
<th>Anorthoclase (%)</th>
<th>Smectite (&lt;vp)</th>
<th>Pyrite (%)</th>
<th>Calcite (%)</th>
<th>Plagioclase (%)</th>
<th>Sandstone (%)</th>
<th>Chlorite (%)</th>
<th>Totobiotite</th>
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</table>

Note: - = absence, + = presence, ++ = relative abundance.

The sample was confined only to the altered zone, so that we cannot exclude the possibility that the sanidine and smectite were at least in part already present as components of the volcaniclastic host rock.

SITE 450

Hole 450 was drilled in the eastern half of the Parece Vela Basin (Fig. 1). Its sedimentary column is represented, from the mudline core down to a sub-bottom depth of 333 meters, by:

1. 26.5 meters of Pleistocene to upper Miocene dark brown, ash-bearing pelagic clay (Sub-unit 1a, Cores 1-3);
2. 61.5 meters of upper to middle Miocene ash-rich pelagic clay with minor dark gray ash layers (Sub-unit 1b, Cores 4 through 10);
3. 236.5 meters of a middle Miocene turbiditic sequence of dark gray vitric and/or crystal tuffs, fine vitric tuffs, and tuffaceous volcaniclastic conglomerates, all altered in varying degrees to clay minerals and with nannofossil-bearing hemipelagic layers (Sub-unit 2a, Cores 10 through the top of 35);
4. 8.5 meters of barren, vitric tuff and fine tuff, altered to various shades of reddish brown (Sub-unit 2b, lowermost Core 35 and uppermost 36). Except for the color change, the depositional characteristics are exactly the same as those of Sub-unit 2a.
5. 0.01 meters of thermally and chemically metamorphosed pale yellowish white fine vitric tuff (Sub-unit 2c, part of Core 36).

This sequence rests on a pillowed intrusive plagioclase-olivine phryic basalt, drilled to a total sub-bottom depth of 340 meters.
Figure 2. X-ray mineral graphs for Core 10 of Hole 447A. (To the left of the lithologic column, sediment colors are indicated by using the standard shipboard color chart system. Smectite crystallinity [v/p] is measured according to Biscaye, 1965.)

We have analyzed a total of 22 samples from the lowermost portion of the unaltered Sub-unit 2a and from the whole Sub-unit 2b. The detailed X-ray mineralogy is shown by mineral graphs in Figure 3.

Sub-unit 2a (Samples 35-1, 106-109 cm to 35-3, 46-49 cm of Table 1)

The vitric tuffs of this unaltered interval contain 68 to 85% amorphous matter, and the X-ray diffractive fraction is given by scarce dioctahedral smectite (the same as Hole 447A) with fair to high crystallinity (v/p ratio 0.6-0.7), by abundant plagioclase (10-30%, An>60), and by occasional calcite. The calcite is biogenic (skeletons of nannofossils), low in magnesium, and is contained in hemipelagic layers that are sometimes burrowed and mottled. At places, layers of crystal vitric tuff occur in the sequence (e.g., Sample 35-3, 8-10 cm). These have lower amorphous and higher plagioclase contents than do vitric tuffs. Biogenic calcite and plagioclase represent primary components of these submarine turbiditic tuffs; the latter are crystals ejected with glass by subaerial and/or submarine explosions.
associated with arc volcanism. Smectite contents fluctuate with the amorphous fraction and derive from devitrification of glass.

Upper Portion of Sub-unit 2b (Samples 35-3, 51–54 cm to 35-3, 89–91 cm of Table 1)

Just below the major color change to red-brown shades, the amorphous fraction slightly increases to more than 85%. This is probably related to the addition of amorphous iron–manganese compounds over the volcanic glass background. Plagioclase significantly decreases (7–12%) and smectite crystallinity is slightly less than in Sub-unit 2a (v/p ratio 0.5–0.6). Todorokite is present in some layers, concentrated in black blebs, streaks, and mottles. Clinoptilolite occurs occasionally, amounting to about 2% in Sample 35-3, 59–63 cm. Aragonite, not detected by X-ray diffraction, occurs in some smear slides as platelets, twins, and radiating crystals. Most of these are euhedral, but some have small etching pits and scars. Calcite is virtually absent in this interval, even in hemipelagic layers, which suggests a possible leaching of biogenous carbonate remains.

Lower Portion of Sub-unit 2b (Samples 36-1, 5–8 cm to 36-2, 112–115 cm of Table 1)

The amorphous fraction is around 80% throughout this interval. A strong decrease in smectite crystallinity (v/p from 0.6 to 0.3–0.4) occurs below Sample 36-1, 70–76 cm. Phillipsite appears in correspondence with such a decrease and is inversely correlated with the amorphous fraction. A few euhedral to subhedral analcime crystals have been detected by SEM in the lowermost samples of this interval. Plagioclase further decreases (less than 5%) and eventually disappears completely below Sample 36-1, 70–76 cm. Euhedral to subhedral, almost pure sanidine appears from the top sample of this interval in concentrations of 5 to 10%. Biogenic calcite is absent, but limpid lumps and clusters of recrystallized low-Mg calcite are abundant from Sample 36-1, 70–76 cm to Sample 36-2, 5–8 cm. Rare aragonite (?) crystals have also been observed on smear slides. Hematite has been found by X-ray diffraction only in sample 36-1, 70–76 cm, although amorphous iron–manganese oxides and hydroxides are widespread in smear slides from the whole Sub-unit 2b.

DISCUSSION

The altered basal volcaniclastic sediments at Sites 447 and 450 contain authigenic smectite, phillipsite, and sanidine. At Site 450 only, other types of authigenic zeolites and carbonates occur. All these minerals represent common products of submarine alteration of tuffs, as recognized in numerous previously studied DSDP sites. Authigenic sanidine, however, seems widespread in the Leg 59 sites, whereas reports of this mineral from other oceanic areas are few (see references in Kastner and Stonecipher, 1978).

Submarine alteration of a given host rock depends on composition of interstitial water, age, and temperature (Hay, 1978). This means that different causes can produce the same diagenetic minerals. From a general view-point, the eventual products of alteration of basal volcaniclastic sediments can derive from (1) halmyrolisis of the volcaniclastic sediments, (2) halmyrolisis of the underlying basaltic basement, and (3) hydrothermalism (up to 300–400°C) associated with the emplacement of the basaltic basement.

Core evidence for the cause of the basal alteration in Hole 447A is rather poor. In Site 450, on the contrary, such causes can be assumed with some confidence; our discussion will therefore be confined to this site. Petrographic evidence (the basaltic basement is pillowowed and altered) and the intrusive contact still preserved in the baked tuffs of Sub-unit 2c show that the basalt was emplaced and cooled beneath a few-meters-thick sequence of previously deposited tuffs. Hajash (this volume) has found that the chemical composition of the altered tuffs is consistent with hydrothermal alteration, even if a low-temperature overprint is present. These tuffs thus would have experienced, at least during the initial stages of alteration, temperatures significantly higher than those of normal deep-sea environments. The host rock unaffected by hydrothermal alteration is represented by the unaltered tuffs of Sub-unit 2a. In fact, all of Unit 2 (Cores 10 through 36) shows great textural and depositional uniformity, and we can reasonably assume that the basal altered interval had the same composition as the overlying unaltered section prior to the basalt intrusion and associated hydrothermalism. The range of minerals found in the host rock and in the altered interval is schematically reported in Figure 4. We will discuss separately the possible significance of different mineral groups.

Plagioclase

Calcium-rich plagioclase, which is a primary crystalline component of the host rock, decreases across the
uppers boundary of the altered zone and eventually disappears about 200 cm above contact with the basalt. Plagioclase is unstable in hydrothermal conditions, the more calcium-rich varieties being the less resistant. Experimental work has demonstrated that plagioclase transforms to other minerals or is completely destroyed at temperatures of about 200°C (Deer et al., 1967). Its disappearance in Hole 450 might suggest that temperatures as high as 200°C affected the tuffs up to 2 meters above the intrusive contact (Fig. 3). Minerals that may derive from the alteration of plagioclase will be discussed later.

Carbonates
Biogenic, low-Mg calcite disappears in the altered zone, where recrystallized spar calcite and euhedral to subhedral aragonite occur. Aragonite-bearing layers overlie the calcite ones (Fig. 4), suggesting that different authigenic carbonates can form depending on temperatures or on concentration of Mg ions (Bishoff and Fyfe, 1968; Natland, 1973). Etching on aragonite crystals may indicate that this mineral is not stable under the present conditions. The recrystallized lumps of calcite should reflect high rates of precipitation for this mineral. They might derive from hydrothermal dissolution of previous skeletal carbonates (Castellari and Sartori, 1978) with subsequent reprecipitation, but they could come at least in part from the precipitation of the Ca released from glass and from anorthite-rich plagioclase. High amounts of carbonates in our altered interval suggest that under hydrothermal conditions neomorphic calcite may form (Bonatti, 1966) and replace previous volcaniclastic beds (Hein and Scholl, 1978).

Smectite
Dioctahedral smectite is present in all samples. Amounts and crystallinity do not vary significantly except for the lowermost 2 meters of the altered interval, where crystallinity is lower than that above (Fig. 3). In the same bottom interval phillipsite is present. Smectite derives from low-temperature devitrification of glass, and its decrease in crystallinity close to the intrusive contact might be explained by either of these two hypotheses:

Hydrothermalism
The tuffs already had a low-temperature smectite component when they underwent the hydrothermal process. Dioctahedral smectites are unstable in hydrothermal conditions and transform to other clay minerals and mixed layers above 300°C (Eberl et al., 1978). We can hypothesize that the hydrothermalism affected these phyllosilicates by decreasing their crystallinity. Lack of further transformation would suggest that the altered tuffs underwent temperatures significantly lower than 300°C. In conditions unfavorable for smectite formation, phillipsite authigenesis could have been enhanced (see the following).

Hyalmyrolisis
Smectite formed at Site 450 after the emplacement and cooling of the basalt. It found favorable conditions everywhere in the tuff sequence except for the basal 2 meters, where the peculiar physiochemical setting of basalt hyalmyrolisis favored the formation of phillipsite. According to Hay (1978), the activity of dissolved species such as H+, alkali ions, etc., determines whether a clay mineral or a zeolite is formed from alteration of tuffs.

Zeolites
From the intrusive contact upward in the altered interval, we observe the following sequence of zeolites: analcime (scarce) + phillipsite + phillipsite + clinoptilolite (Fig. 4). The trend is to a progressive hydration of the authigenic zeolites far from the heat source, which suggests that a thermal gradient might have been established in the intruded tuff sequence. In hydrothermal conditions, analcime starts forming at higher temperatures than phillipsite (250° and 200°C in Höller and Wirshing, 1974, 1978; Wirshing, 1975), and clinoptilolite forms at lower temperatures (about 140°C in Hawkins et al., 1978). Such temperatures are in good accordance with the paragenesis found in Sub-unit 2b, because plagioclase (which is destroyed at about 200°C, as is discussed earlier) coexists with clinoptilolite and disappears where phillipsite is present (Fig. 4). Plagioclase should be at least in part the precursor of zeolite. The relationship between phillipsite and low-crystallinity smectite at the bottom of Hole 450 suggests, however, that some of this zeolite might also derive directly from glass (in conditions unfavorable to smectite formation) or from previously formed smectite.

Sanidine
Authigenic sanidine is not necessarily a high-temperature mineral. Apart from data reported by Stewart et al. (1973), most of the known occurrences of this mineral are diagenetic in origin (see Kastner and Stonecipher, 1978). Moreover, authigenic sanidine has been experimentally formed at temperatures below 80°C by Flehmig (1977). In Site 450 (and also in Hole 447A), sanidine does not show great variation in concentration. Confined and extending throughout the altered zone, sanidine thus differs from phillipsite and low-crystallinity smectite (Fig. 4). It can also coexist with partly altered plagioclase. These observations suggest that sanidine might have formed in Site 450 at relatively low temperatures, deriving mostly from glass and subordinately from plagioclase. In fact, our samples of sanidine approach pure composition very closely, as is the case for authigenic K feldspars replacing former glass. Sanidine after plagioclase should be rich in Na according to Deer et al. (1967).

Todorokite and Hematite
These minerals, together with the amorphous ferromanganese compounds detected in smear slides, suggest a direct supply of Fe and Mn from alteration of the underlying basalt. Todorokite occurs higher in Sub-unit 2b than in hematite, suggesting that some fractionation processes have occurred between the Fe and the more mobile Mn (Krauskopf, 1957). The presence of diagenetic crystalline hematite has been observed in the basal
sediments of the West Philippine Basin by Bonatti et al. (1979). A diagenetic mobilization and recrystallization of hematite and todorokite in our samples would be indicated by their occurrence in blebs and mottles.

CONCLUSIONS

The authigenic minerals contained in the altered basal intervals of volcaniclastic sediments from Sites 447 and 450 of DSDP Leg 59 are dioctahedral smectite (with variable crystallinity), phillipsite, and sanidine. Sanidine seems the most widespread and common product of basal alteration in the marginal basins of the Philippine Sea.

The neomorphic mineral suites may have been produced by (1) halmyrolysis of the volcaniclastic sediments; (2) halmyrolysis of the underlying basalts; or (3) hydrothermalism associated with basaltic emplacement and cooling.

At Site 450, where other authigenic minerals occur (carbonates, analcime, clinoptilolite, Fe-Mn oxides), the basal paragenesis is consistent with a hydrothermal origin. Such a process could have produced temperatures up to 200°C in tuffs lying as high as 2 meters above the contact with a basaltic intrusion. Products of low-temperature alteration, however, are also present in the altered intervals of this site.

ACKNOWLEDGMENTS

E. Bonatti read parts of the manuscript and offered useful suggestions. N. Morandi discussed some aspects of alteration of volcanics. G. Busatti and W. Landuzzi made laboratory analyses, and G. Zini drew the illustrations.

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