The oxygen isotopic compositions (δ^{18}O) of basalts from DSDP Leg 46 range from 5.9 to 8.9‰ (SMOW), indicating 1 to 15 per cent weathering induced by cold sea water. Basalts from the bottom portion of the hole, including basaltic gravel, have lower δ^{18}O than basalts from the upper parts of the hole, consistent with the bottom portion of the hole being sealed off from direct sea-water circulation prior to drilling. The δ^{18}O (34.7‰) of vug-filling carbonate from the highly altered central portion of the 9-meter cooling unit in Core 15 indicates that the alteration is a result of cold sea-water reaction and is not deuteric alteration.

METHODS
Oxygen was extracted from the rock powders by the BrFs method (Clayton and Mayeda, 1963), reacted with carbon to form CO₂, and analyzed on an isotope ratio mass spectrometer. The isotopic composition of sample, x, is given:

\[ \delta_x = \frac{^{18}O}{^{16}O} \frac{x}{\text{standard}} - 1 \times 1000. \]

The analyses are reported relative to Standard Mean Ocean Water (Craig, 1961). The carbonates were analyzed by the H₃PO₄ method (McCrea, 1950).

OXYGEN ISOTOPE GEOCHEMISTRY
The values of δ^{18}O for ten basaltic and two carbonate samples for DSDP Hole 396B are presented in Table 1. The δ^{18}O values for basalts range from 5.9‰ for a pillow rind fragment from the lower part of the hole (26-1, 3-7 cm) to 8.9‰ for an iron oxide stained sample from the vuggy interior of the 9-meter cooling unit in Core 15 (15-3, 83-89 cm). Although water contents are not presently available for these samples, on the reasonable assumption that these samples fall on the same trend as other deep-sea samples, the measured δ^{18}O values represent from 1 to 15 per cent weathering, with an average value of approximately 9 per cent. The amount of weathering indicated for the Leg 46 basalts is quite comparable to that found for other young crustal samples such as those from Legs 34 and 37 (Muehlenbachs, 1976, 1977a) and considerably more weathered than samples from Legs 34, 37, and 46.

INTRODUCTION
Hole 396B of Leg 46 (located at 22°59.14'N, 43°30.90'W, approximately 160 km east of the Mid-Atlantic Ridge) penetrated 205.5 meters into basement. Basement at the site consists primarily of basaltic pillow lavas and fragmental basaltic material, and is overlain by 150.5 meters of nannofossil ooze, the basal part of which is Miocene (~13 m.y.). The purpose of this paper is to report on the oxygen isotope geochemistry of samples from Site 396B and to relate that chemistry to interaction between basaltic basement material and ocean water.

Several studies have shown that the δ^{18}O values of unaltered oceanic basalts are very uniform, approximately 5.7 ± 0.3‰ (Taylor, 1968; Garlick and Dymond, 1970; Muehlenbachs and Clayton, 1972; Pineau et al., 1976; Anderson and Lawrence, 1976; and Muehlenbachs, 1976, 1977a). The same studies have shown that most ocean basalts are a few per mil enriched in δ^{18}O because of partial reaction with cold sea water. Consideration of basaltic dredge samples and a wide range of DSDP samples indicates that there is a high positive correlation between the δ^{18}O of the basaltic samples and their total water content (Muehlenbachs and Clayton, 1972; Pineau et al., 1976; Muehlenbachs, 1977a,b). Thus, to a good approximation the samples may be considered as mixtures of normal-δ^{18}O (~5.7‰), fresh, anhydrous basalt and high-δ^{18}O (~27‰) smectites formed during weathering at low temperatures (~4°C), and the percentage of oxygen isotope exchange experienced by the sample is a direct measure of the degree of weathering the sample has undergone.

Samples recovered by DSDP Legs 34, 37 (Muehlenbachs, 1976, 1977a), and 46 (this study) are approximately 10 per cent weathered, more or less uniformly to a depth of at least 580 meters into the basaltic layer of the relatively young oceanic crust. Isotopic evidence indicates that both dredge samples of similar age (Muehlenbachs and Clayton, 1976) and the Cretaceous and Jurassic basalts recovered by DSDP Legs 11, 26, 27 (Muehlenbachs, 1977b), and 35 (Anderson and Lawrence, 1976) are considerably more weathered than samples from Legs 34, 37, and 46.
breccia near the top of the basalt sequence (5-1, 63-68 cm) yields a δ18O of 34.5‰ and calcite from vugs in cooling unit Sample 15-3, 83-89 cm yields a δ18O value of 34.7‰. If the carbonates are pure calcites in equilibrium with nor-
mal sea water, their isotopic composition implies a temperature of ~3°C (O'Neil et al. 1969), a value within the range of 7° to −10°C calculated for carbonates from Legs 34 (Seyfried et al., 1976), 37 (Muehlenbachs, 1977a), and for dredged basalts (Muehlenbachs and Clayton, 1972). The indicated sub-freezing temperature is probably a result of the presence of divalent cations other than calcium in the carbonates.

The basaltic samples in the upper part of the hole (Cores 5 to 22), with the exception of Sample 7-1, 50-57 cm, are more enriched in 18O than the three analyzed samples from the lower part of the hole (from Cores 26, 32, and 33). Sample 7-1, 50-57 cm, one of the Leg 46 internal chemical standards, evolved a considerable quantity of water during grinding in a ball mill (M. Dungan, written communication) and probably has not yielded a meaningful δ18O value. Ignoring this sample, the average 18O value for the upper six basaltic samples is 8.3‰ with a standard deviation of 0.4‰, while the average value for the lower three samples, including basaltic gravel Sample 33-1, 60-65 cm, is 6.5‰ with a standard deviation of 0.6. The lower 18O values in the bottom portion of the hole are indicative of less interaction with sea water and are consistent with heat flow evidence that the bottom portion of the section at Hole 396B was sealed off from direct sea-water circulation prior to drilling of the hole (Erickson, this volume).

Two samples from the 9-meter cooling unit in Core 15 were analyzed, one from the relatively fresh upper portion of the unit and one from the vuggy, iron-oxide stained central portion. Sample 15-3, 83-89 cm, from the central portion has a higher 818O (8.9‰) than Sample 15-2, 120-123 cm (7.8‰), consistent with the more altered appearance of the former. In addition, the 818O (34.7‰) of the vug-filling carbonate from Sample 15-3, 83-89 cm, is a strong indica-

### TABLE 1

**δ18O (SMOW) of Samples From DSDP Leg 46**

<table>
<thead>
<tr>
<th>Sample (Interval in cm)</th>
<th>Basalt δ18O</th>
<th>Carbonate δ18O</th>
<th>Carbonate δ13C (PDB)</th>
</tr>
</thead>
<tbody>
<tr>
<td>396B-5-1, 63-68</td>
<td>8.5</td>
<td>34.5</td>
<td>+1.9</td>
</tr>
<tr>
<td>396B-7-1, 50-57</td>
<td>5.5</td>
<td>34.5</td>
<td>+1.9</td>
</tr>
<tr>
<td>396B-11-1, 22-28</td>
<td>8.2</td>
<td>34.5</td>
<td>+1.9</td>
</tr>
<tr>
<td>396B-15-2, 120-123</td>
<td>7.8</td>
<td>34.5</td>
<td>+1.9</td>
</tr>
<tr>
<td>396B-15-3, 83-89</td>
<td>8.9</td>
<td>34.7</td>
<td>+5.4</td>
</tr>
<tr>
<td>396B-20-5, 12-20</td>
<td>8.5</td>
<td>34.7</td>
<td>+5.4</td>
</tr>
<tr>
<td>396B-22-3, 28-38</td>
<td>7.9</td>
<td>34.7</td>
<td>+5.4</td>
</tr>
<tr>
<td>396B-26-1, 3-7</td>
<td>5.9</td>
<td>34.7</td>
<td>+5.4</td>
</tr>
<tr>
<td>396B-32-1, 100-105</td>
<td>7.0</td>
<td>34.7</td>
<td>+5.4</td>
</tr>
<tr>
<td>396B-33-1, 60-65</td>
<td>6.7</td>
<td>34.7</td>
<td>+5.4</td>
</tr>
</tbody>
</table>


bLeg 46 internal chemical standard 1.

cLeg 46 internal chemical standard 2.

dLeg 46 internal chemical standard 3.

eBasaltic gravel.

### CONCLUSIONS

The major conclusions that may be drawn from this study are: (1) samples from DSDP Hole 396B exhibit δ18O values similar to those from other young crustal samples that have undergone low-temperature reaction with sea water; (2) the lower δ18O values in the bottom portion of the hole are consistent with the lower clastic sequence being sealed off from direct circulation of sea water prior to drilling; and (3) the δ18O values from vug filling carbonate in Sample 15-3, 83-89 cm, indicates that the more intense alteration in the central portion of the 9-meter cooling unit is a result of low-temperature reaction with sea water rather than deuteric alteration.

### REFERENCES


