2. DATA REPORT: ISOTOPIC COMPOSITION OF PORE FLUIDS, NEW JERSEY SHELF AND SLOPE¹

Mitchell J. Malone² and Jonathan B. Martin³

ABSTRACT

An investigation of the isotopic composition of the interstitial waters was conducted at Sites 1071, 1072, and 1073 on the New Jersey continental shelf and slope during Ocean Drilling Program Leg 174A. Sites 1071 and 1072 are closely spaced drill holes on the continental shelf located ~130 km from the shoreline in 88 and 98 m of water, respectively. Site 1073 is located on the continental slope in 640 m water and penetrated a total of 664 m of sediment of which ~520 m is Quaternary age. A total of 125 oxygen and hydrogen isotopic analyses of pore fluids are presented from all three sites. Twelve strontium isotopic ratios are reported from Site 1071.

INTRODUCTION

The majority of previous investigations of marine interstitial water chemistry during the Deep Sea Drilling Project and Ocean Drilling Program (ODP) have been conducted on fluids recovered from pelagic and hemipelagic sediments. Most interstitial water studies on continental shelves have been limited to the upper meters of the sediment column principally because of the difficulties of drilling in shallow waters. This has been particularly true on the New Jersey shelf because of potential shallow-gas hazards. One attempt at shelf drilling was initiated by the United States Geological Survey Atlantic Margin Coring Project (AM-COR) (Hathaway et al., 1979). In widely spaced, shallow holes across the New Jersey continental shelf, AMCOR researchers noted a fresh to ¹Malone, M.J., and Martin, J.B., 2000. Data report: Isotopic composition of pore fluids, New Jersey shelf and slope. *In* Christie-Blick, N., Austin, J.A., Jr., and Malone, M.J. (Eds.), *Proc. ODP, Sci. Results*, 174A, 1–11 [Online]. Available from World Wide Web: <http://wwwodp.tamu.edu/publications/174A_SR/ VOLUME/CHAPTERS/SR174A02.PDF>. [Cited YYYY-MM-DD] ²Ocean Drilling Program and Department of Geology and Geophysics, 1000 Discovery Dr., Texas A&M University, College Station TX 77845, USA. malone@odpemail.tamu.edu

malone@odpemail.tamu.edu ³Department of Geology, PO Box 112120, University of Florida, Gainesville FL 32611, USA.

Initial receipt: 15 November 1999 Acceptance: 14 June 2000 Web Publication: 2 October 2000 Ms 174ASR-156

brackish water plume (low Cl⁻) extending at least 100 km offshore. They believed this fluid to be relict Pleistocene water and suggested that its presence is an impediment to seawater infiltration of mainland freshwater aquifers (Hathaway et al., 1979; Kohout et al., 1988). The drilling of two closely spaced sites on the outer shelf ~130 km from the shoreline (Site 1071 in 88 m water depth and Site 1072 in 98 m water depth) (Austin, Christie-Blick, Malone, et al., 1998) provided the opportunity to further explore the origin and distribution of the low Cl-plume (Fig. F1).

Most notably, McDuff (1985), Schrag and DePaolo (1993), Schrag et al. (1996), and Burns and Maslin (1999) have observed an effect of sealevel fluctuations on pore-water chemistry. They observed positive excursions in Cl⁻ and δ^{18} O at shallow depths (~50 meters below seafloor [msbf]), which are consistent with increases in salinity and δ^{18} O of seawater during the last glacial maximum (LGM). However, the positive excursions in Cl⁻ and δ^{18} O have only been observed at shallow depths (i.e., for the LGM) because at greater depths any signal is dampened by diffusive smoothing. High-resolution interstitial water sampling was undertaken at Site 1073 (640 m water depth) on the New Jersey slope to examine such variability (Austin, Christie-Blick, Malone, et al., 1998) (Fig. F1). Postcruise stable isotopic analyses were performed to supplement shipboard data.

METHODS

Pore-fluid samples were collected using routine shipboard squeezing of whole-round sediment samples immediately after retrieval. Additional details on shipboard analyses and data are reported in Austin, Christie-Blick, Malone, et al. (1998). Oxygen and hydrogen isotopic analyses were performed in duplicate at Mountain Mass Spectrometry (Evergreen, Colorado, U.S.A.) using an automated preparation system. Oxygen isotopic ratios were determined using the CO₂-water equilibration method of Epstein and Mayeda (1953). Hydrogen isotopic ratios were determined using the equilibration method of Horita (1988, 1989), whereby the water sample is equilibrated with hydrogen gas with the aid of Hokka bead platinum catalysts. Replicate analyses of the standard through the course of the analyses yielded a standard deviation of $0.04\%_0$ for δ^{18} O and $0.4\%_0$ for δ D.

Sr isotope measurements were made on ~150 μ L of pore water, which was pipetted into Teflon containers and completely dried. The resulting salts were dissolved in 50 μ L of 3.5-N HNO₃. Strontium was separated from this solution with Sr-selective crown ether resin, following a technique modified from Pin and Bassin (1992). The Sr blank for the technique is 100 pg. The separated Sr was loaded onto tungsten filaments and analyzed in the Department of Geology at the University of Florida for ⁸⁷Sr/⁸⁶Sr isotope ratios using a VG Micromass 354 triple collector thermal ionization mass spectrometer in dynamic mode. Instrumental mass fractionation was corrected to a ⁸⁶Sr/⁸⁸Sr ratio of 0.1194. Numerous replicate measurements of the NIST-987 standard over the past several years have yielded a ⁸⁷Sr/⁸⁶Sr value of 0.710235, with an external precision of ±0.000023 (2 σ). This external precision represents the minimum uncertainty assigned to any individual sample.

F1. Location of Sites 1071–1073 drilled during Leg 174A, p. 5.



RESULTS

Results of isotopic analyses of interstitial waters are compiled for Sites 1071, 1072, and 1073 in Tables **T1**, **T2**, and **T3**, respectively. In addition to the data, ODP sample identifier and depth (in mbsf) of each sample are also tabulated. Data are depicted graphically vs. depth in Figures **F2**, **F3**, and **F4**.

ACKNOWLEDGMENTS

The analyses presented in this report were funded by a JOI/USSSP grant. We are indebted to the ODP Leg 174A technical and scientific parties, especially co-chief scientists James A. Austin, Jr. and Nicholas Christie-Blick. Comments from Rick Murray and James Austin are much appreciated. We acknowledge ODP for access to data and samples.

T1. Isotopic composition of interstitial waters, Site 1071, p. 9.

T2. Isotopic composition of interstitial waters, Site 1072, p. 10.

T3. Isotopic composition of interstitial waters, Site 1073, p. 11.

F2. Interstitial water isotopic data vs. depth, Site 1071, p. 6.



F3. Interstitial water isotopic data vs. depth, Site 1072, p. 7.



F4. Interstitial water isotopic data vs. depth, Site 1073, p. 8.



REFERENCES

- Austin, J.A., Jr., Christie-Blick, N., Malone, M.J., et al., 1998. *Proc. ODP, Init. Repts.*, 174A: College Station, TX (Ocean Drilling Program).
- Burns, S.J., and Maslin, M.A., 1999. Composition and circulation of bottom water in the western Atlantic Ocean during the last glacial, based on pore-water analyses from the Amazon Fan. *Geology*, 27:1011–1014.
- Epstein, S., and Mayeda, T., 1953. Variation of ¹⁸O content of waters from natural sources. *Geochim. Cosmochim. Acta*, 4:213–224.
- Hathaway, J.C., Poag, C.W., Valentine, P.C., Miller, R.E., Schultz, D.M., Manheim, F.T., Kohout, F.A., Bothner, M.H., and Sangrey, D.A., 1979. U.S. Geological Survey core drilling on the Atlantic Shelf. *Science*, 206:515–527.
- Horita, J., 1988. Hydrogen isotope analysis of natural waters using an H₂-water equilibration method: a special implication to brines. *Chem. Geol.*, 72:89–94.
 - _____, 1989. Analytical aspects of stable isotopes in brines. *Chem. Geol.*, 79:107–112.
- Kohout, F.A., Meisler, H., Meyer, F.W., Johnston, R.H., Leve, G.W., and Wait, R.L., 1988. Hydrogeology of the Atlantic continental margin. *In Sheridan*, R.E., and Grow, J.A. (Eds.), *The Geology of North America* (Vol. I-2): *The Atlantic Continental Margin*, U.S. Geol. Soc. Am., 463–480.
- McDuff, R.E., 1985. The chemistry of interstitial waters, Deep Sea Drilling Project Leg 86. *In* Heath, G.R., Burckle, L.H., et al., *Init. Repts. DSDP*, 86: Washington (U.S. Govt. Printing Office), 675–687.
- Pin, N.C., and Bassin, C., 1992. Evaluation of a strontium-specific extraction chromatographic method for isotopic analysis in geological materials. *Anal. Chim. Acta*, 269:249–255.
- Schrag, D.P., and DePaolo, D.J., 1993. Determination of δ^{18} O of seawater in the deep ocean during the last glacial maximum. *Paleoceanography*, 8:1–6.
- Schrag, D.P., Hampt, G., and Murray, D.W., 1996. Pore fluid constraints on the temperature and oxygen isotopic composition of the glacial ocean. *Science*, 272:1930–1932.

Figure F1. Location of Sites 1071–1073 drilled during Leg 174A. Bathymetry is given in meters. In addition, AMCOR sites drilled offshore New Jersey (Hathaway et al., 1979) are also shown (6009–6011 and 6020).







Figure F3. Interstitial water isotopic data from Site 1072 vs. depth. Pore-water Cl⁻ data vs. depth are also shown (Austin, Christie-Blick, Malone, et al., 1998). SMOW = standard mean ocean water.





Figure F4. Interstitial water isotopic data from Site 1073 vs. depth. SMOW = standard mean ocean water.

Table T1. Isotopic composition of interstitial waters, Site 1071.

Core, section, interval (cm)	Depth (mbsf)	δ ¹⁸ O SMOW	δD SMOW	⁸⁷ Sr/ ⁸⁶ Sr
174A-1071A-				
2H-1, 145-150	6.55	-0.65	8.15	
3H-2, 145-150	10.80	-0.92	8.23	0.709347
4H-2, 145-150	14.90	-0.98	5.92	0.709334
6H-1, 75-80	26.45	-1.22	1.23	0.709306
174A-1071B-				
2X-1, 140-150	41.20	-1.35	-1.74	
3X-CC, 0-50	49.64	-1.44	-1.07	
5X-2, 145-150	61.35	-1.32	0.50	
5X-3, 140-150	62.80	-1.26	-0.12	0.709230
174A-1071C-				
2X-2, 140-150	61.30	-1.32	0.84	
6X-CC, 21-29	97.09	-1.05	0.60	0.709382
7X-CC, 13-23	105.73	-1.13	-2.06	
8X-CC, 17-27	116.34	-0.92	-2.02	
9X-2, 140-150	127.40	-1.13	1.69	0.709701
Replicate	127.40	-1.13	1.69	0.709686
12X-2, 140-150	151.40	-1.06	-0.81	
15X-5, 140-150	169.80	-1.06	-0.89	0.709167
19X-CC, 0-16	201.16	-0.93	-1.69	
22X-1, 140-150	229.80	-1.02	2.46	
23X-1, 140-150	239.10	-1.07	0.76	
25X-2, 140-150	254.50	-1.13	3.36	0.709361
174A-1071D-				
1R-1, 52-62	0.52	-0.86	-0.88	0.709364
Replicate	0.52	-0.86	-0.88	0.709374
4R-CC, 6-12	28.36	-1.06	-4.00	
5R-1, 24-30	38.24	-1.40	-2.70	
6R-1, 97-107	48.67	-1.36	-0.57	
7R-1, 33-43	57.73	-1.33	-1.34	0.709181
8R-1, 17-27	62.57	-1.34	-1.76	0.709214
174A-1071F-				
1R-1, 140-150	253.40			
2R-1, 39-45	261.79	-1.14	-0.87	
6R-1, 118-128	321.38	-1.06	-1.69	
7R-1, 90-100	330.50	-0.88	-3.33	
8R-1, 0-10	339.00	-0.86	-1.08	
9R-2, 74-86	350.04	-0.74	-6.18	
10R-3, 140-150	362.30	-0.81	-2.36	0.709018
11R-2, 58-68	369.48	-0.85	-6.39	

Note: SMOW = standard mean ocean water.

Table T2. Isotopic composition of interstitial waters, Site 1072.

Core, section,	Depth	$\delta^{18}O$	δD
interval (cm)	(mbsf)	SMOW	SMOW
174A-1072A-			
1R-1, 134-144	1.34	-0.38	6.53
2R-1, 140-150	8.90	-0.52	4.99
3R-1, 140-150	18.40	-0.53	5.71
4R-2, 140-150	29.60	-0.59	0.19
7R-1, 140-150	52.40	-0.81	0.60
9R-2, 140-150	63.90	-0.81	-0.84
15R-1, 33-43	90.03	-0.81	3.32
16R-1, 104-114	95.04	-0.86	3.86
19R-2, 140-150	111.30	-0.64	2.50
22R-2, 140-150	130.30	-0.53	2.77
29R-1, 140-150	161.70	-0.57	2.97
33R-1, 140-150	182.20	-1.07	2.14
39R-3, 140-150	211.70	-1.05	1.14
43R-1, 140-150	230.50	-0.82	1.55

Note: SMOW = standard mean ocean water.

δ¹⁸Ο Core, section, Depth δ¹⁸O δD Core, section, Depth δD interval (cm) (mbsf) SMOW SMOW (mbsf) SMOW SMOW interval (cm) 174A-1073A-32X-3, 135-150 287.2 -0.56 -6.86 1H-1, 140-150 1.4 0.89 3.64 33X-5, 135-150 298.0 -0.60 -6.70 1.02 1H-3, 140-150 4.4 3.58 304.5 -0.67 -6.70 34X-3, 135-150 9.5 2H-2, 140-150 1.13 3.33 35X-4, 135-150 315.4 -0.64 -6.62 2H-3, 140-150 11.0 1.08 3.70 174A-1073A-2H-5, 140-150 14.0 1.01 3.56 36X-3, 135-150 322.8 -0.77 -7.38 3H-1, 130-140 17.4 0.89 3.24 37X-3, 135-150 332.2 -0.80 -7.79 3H-3, 130-140 20.2 0.97 2.80 38X-5, 135-150 344.6 -0.78 -7.67 3H-5, 130-140 0.99 2.77 23.0 39X-4, 135-150 352.5 -0.70 -6.87 4H-1, 130-140 26.9 1.06 1.88 40X-4, 135-150 361.9 -0.35 -5.40 29.8 1.01 4H-3, 130-140 2.68 41X-6, 135-150 374.3 -0.29 -5.92 4H-5, 130-140 32.7 1.00 2.81 42X-5, 135-150 382.2 -0.30-6.52 5H-6, 140-150 44.1 0.76 3.14 43X-5, 135-150 391.7 -0.45 -5.89 6H-7, 130-150 3.25 54.1 0.75 44X-3, 135-150 398.0 -0.38 -6.38 7H-6, 140-150 63.0 0.79 2.57 45X-5, 135-150 409.9 -0.51 -6.58 8H-6, 140-150 72.5 0.78 2.42 46X-5, 135-150 418.7 -0.56 -6.53 9H-6, 140-150 82.0 1.19 2.74 47X-6, 135-150 430.4 -0.50 -6.91 10H-6, 140-150 91.5 0.63 3.21 48X-5, 135-150 438.3 -0.67 -6.73 11H-6, 140-150 101.0 0.65 2.18 49X-5, 135-150 447.8 -0.52 -7.22 12H-4, 140-150 107.5 0.51 1.68 50X-5, 135-150 456.5 -0.58 -7.16 13H-5, 140-150 1.93 118.5 0.38 51X-5, 135-150 466.7 -0.63 -7.30 14H-5, 140-150 128.0 0.47 1.56 -7.94 52X-5, 135-150 475.6 -0.59 15H-6, 140-150 139.0 0.37 1.38 53X-5, 135-150 484.3 -0.64 -7.33 16H-2, 140-150 142.5 0.45 -0.17 54X-5, 135-150 494.6 -0.46 -7.13 16H-6, 140-150 148.5 0.47 0.58 55X-5, 135-150 504.0 -0.57 -6.86 17H-6, 140-150 158.0 0.60 -0.44 56X-5, 135-150 513.3 -0.72 -7.64 18H-5, 140-150 166.0 0.56 -1.50 57X-1, 135-150 516.7 -0.65 -8.37 19H-6, 140-150 176.9 0.41 -0.22 57X-4, 135-150 521.2 -0.58 -8.83 20H-4, 140-150 183.4 0.33 -1.25 58X-4, 135-150 530.7 -0.56 -7.29 21H-4, 140-150 190.9 0.29 -0.88 59X-6, 135-150 543.2 -0.41 -8.64 22H-2, 140-150 197.5 0.32 -1.2060X-5, 135-150 549.2 -0.41 -8.27 22H-3, 140-150 199.0 0.30 -1.57 61X-6, 135-150 -0.59 558.8 -8.45 22H-5, 140-150 202.1 0.24 -1.76 62X-5, 135-150 567.5 -0.76 -7.33 23H-5, 140-150 210.2 0.25 -2.68 63X-5, 135-150 -0.48576.8 -8.98 24H-1, 134-146 213.1 0.29 -4.16 64X-5, 135-150 586.2 -0.50 -7.72 25X-3, 135-150 220.1 0.19 -2.27 65X-5, 135-150 595.6 -0.87 -7.93 26X-3, 140-150 228.8 0.26 -2.08 66X-5, 135-150 605.2 -0.92 -8.72 27X-4, 135-150 239.8 -0.03 -3.24 67X-5, 135-150 614.6 -0.83 -6.92 28X-4, 135-150 249.4 0.04 -1.92 68X-5, 135-150 624.2 -0.94 -8.9829X-3, 135-150 257.4 0.01 -2.65 69X-5, 135-150 633.7 -0.99 -8.92 30X-3, 135-150 268.4 -0.26 -4.35 31X-4, 135-150 277.9 -0.61 -5.41

 Table T3. Isotopic composition of interstitial waters, Site 1073.