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Characterization of Fracture Surfaces in Dolomite Rock, Culebra Dolomite Member, Rustler Formation

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CHARACTERIZATION OF FRACTURE SURFACES IN DOLOMITE ROCK, CULEBRA DOLOMITE MEMBER, RUSTLER FORMATION*

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ABSTRACT

The Culebra Dolomite Member, Rustler Formation, southeastern New Mexico, is characterized by a high fracture porosity. Bedding plane fractures are predominant, but vertical and high-angle fractures are also common. The information presented in this report for horizontal fractures shows that: (1) horizontal water-bearing fractures in dolomite and calcite rock tend to occur in zones where clay and quartz are concentrated, particularly along clay seams, and (2) secondary minerals, primarily gypsum and some calcite, are precipitated from solution onto the fracture surfaces. No surfaces of vertical or high-angle fractures that were clearly identifiable as water-bearing were discovered in the cores examined.

Dolomite compositions on the fracture surfaces are no different than those in the bulk rock, indicating that aqueous alteration of dolomite did not occur to any significant extent. Where present, calcite in these samples, both in the bulk rock and fracture surfaces, is a product of recrystallization from dolomite caused by aqueous alteration, usually near surface.

It is probable that the vertical fracture surfaces are not as clay-rich as the horizontal ones, since accumulations of clay occur along horizontal planes due to sedimentation.

Our data argue that since the cation exchange capacity of clay minerals is so much higher than that of dolomite, calcite, or gypsum, and the clay minerals are a major component of the fracture surface mineralogy, the sorption of radionuclides due to the clay will far outweigh that of the other minerals. This fact should be taken into account in any study of the transport of radionuclides through the Culebra Dolomite.

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I. INTRODUCTION

The Culebra Dolomite Member of the Rustler Formation in southeastern New Mexico is characterized by a high fracture porosity. Bedding plane fractures are predominant, but vertical and high-angle fractures are also common. In the WIPP-19 drill core (Chapter III), the bedding plane fracture density varies from 3 to 8 per vertical foot below 764 feet depth, and 1 to 3 per vertical foot above 764 feet. Irregularly-curved vertical fractures occur with a frequency of 1 to 4 per vertical foot, and high angle fractures (60 to 70°) are spaced at about 1 to 5 per vertical foot (Ferrall and Gibbons, 1979). Water-bearing (open) fracture surfaces are easily recognized because they are darker than the bulk dolomite due to aqueous alteration.

The objective of this report is to describe the water-bearing fracture surfaces in detail, in terms of texture and mineralogy and to compare and contrast the composition and mineralogy of the surfaces with that of the bulk rocks. Analytical methods used in this study include x-ray diffraction (XRD) analysis, x-ray fluorescence (XRF) spectroscopic analysis, atomic absorption (AA) spectroscopy, electron microscope elemental analysis, and optical microscopy.

II. SITE GEOLOGY

The Upper Permian (Ochoan) Rustler Formation is a sequence of evaporite and clastic rock units deposited in the Delaware, Midland, Palo Duro, and Dalhart Basins in southeastern New Mexico and western Texas. In the Delaware Basin (Figure II-1), which is ringed by the Capitan Reef Complex, the Rustler Formation overlies the Salado Formation, which is composed mainly of thick halite beds and is in turn overlain by the Dewey Lake Red Bed Formation, which is composed almost entirely of mud/siltstone.

The Rustler Formation is divided into four recognizable units (Figure II-2). In ascending order, they are the lower (unnamed) member (argillaceous halite, mud/siltstone, and anhydrite), the Culebra Dolomite Member, the Tamarisk Member (halite, mudstone, anhydrite, and gypsum), the Magenta Dolomite Member, and the Forty-niner Member (anhydrite, gypsum, and mudstone).

In the boreholes from which samples were taken for this study, the Culebra Dolomite Member varies in thickness from about 20 feet to 30 feet. It is primarily composed of massive, laminated dolomite with some clay, quartz, and gypsum, although some cores show extensive brecciation and/or large vugs and voids. Clay laminae of varying thicknesses (2 mm to several centimeters) are common, and the entire unit is underlain by a 0.5 to 3 feet thick black shale (Sowards, Glenn, and Keil, 1991). In the ventilation and access shafts of the WIPP repository, the uppermost 6 to 12" of the black shale are deformed and tilted. This deformation is evidently due to collapse of the dolomite unit. There are no obvious depositional features in the lower 2.5 feet of the black shale. The black shale grades into a reddish/brown shale that is clearly a solution residue: there are no depositional features (laminae, etc.), and the texture is chaotic. This reddish/brown shale overlies the uppermost anhydrite unit in the lower member; the contact is very uneven, but fairly well defined. The anhydrite unit itself is relatively unfractured. It is clear that the dissolution of an argillaceous halite bed above the anhydrite unit and of several other halite units in the lower member and uppermost Salado Formation caused the collapse of the Culebra and overlying units. This collapse resulted in extensive fracturing in the dolomite units (Culebra and Magenta), particularly in the Culebra. The reason that the anhydrite and shale units are not as fractured as the dolomite units is that anhydrite and shale are able to deform in a more plastic manner than the harder dolomite.

In the areas where the dolomite unit lies near the surface, calcite is often present instead of dolomite. This calcite is the result of dedolomitization of dolomite.

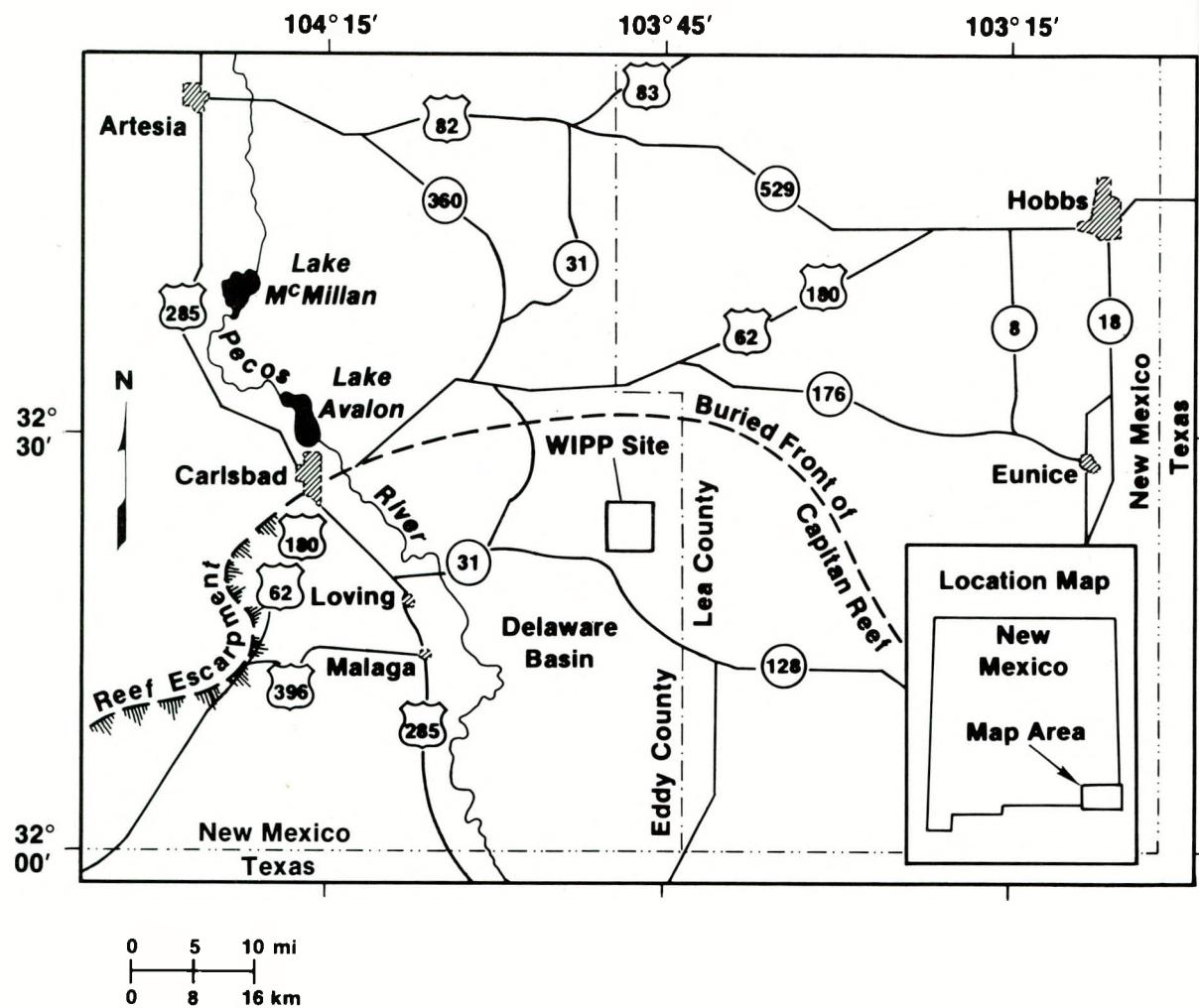
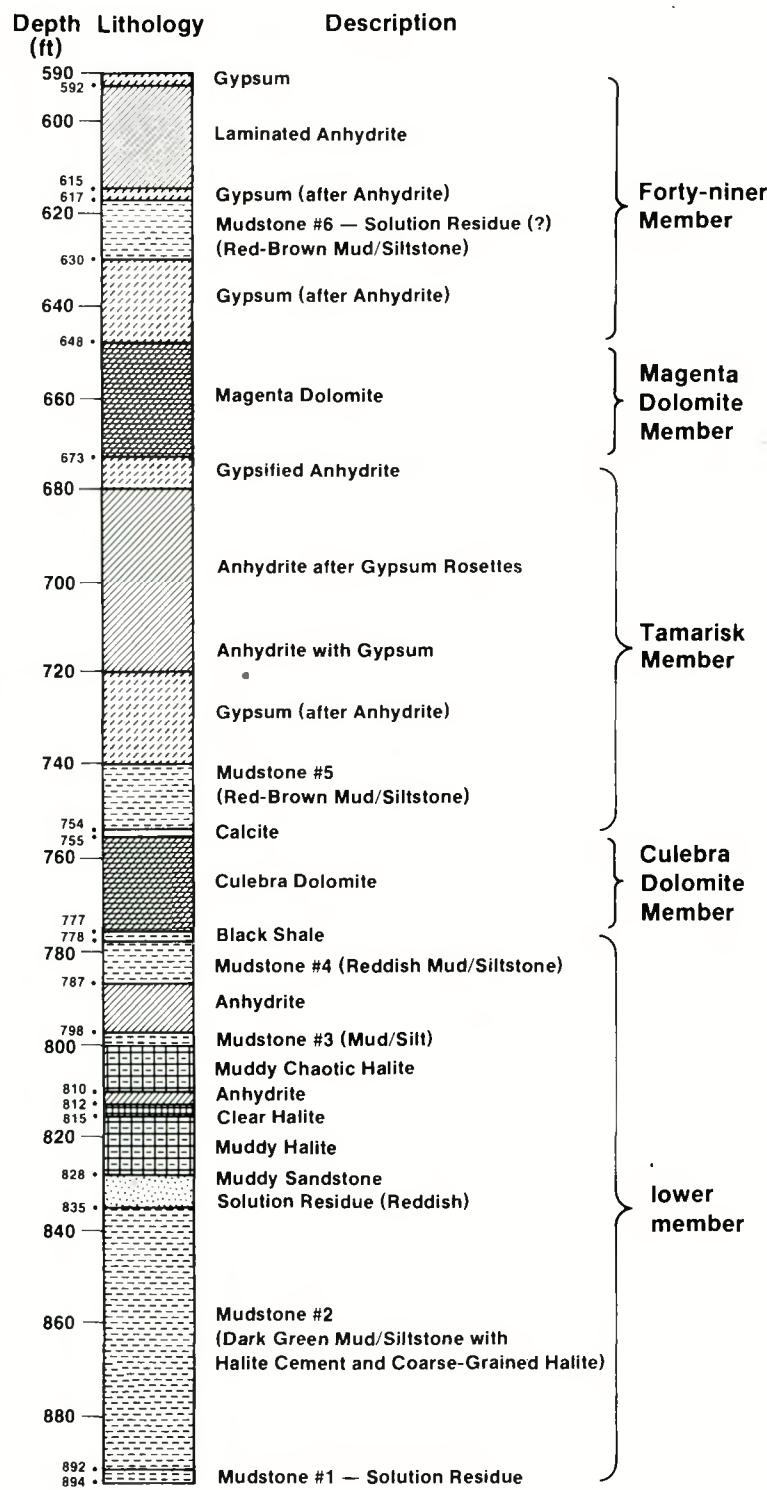


Figure II-1. Regional Setting, Delaware Basin, Southeastern New Mexico (from Borns et al., 1983).



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Figure II-2. Lithologic Log of WIPP-19 Core, Rustler Formation (modified from Ferrall and Gibbons, 1979).

III. PROJECT SAMPLES

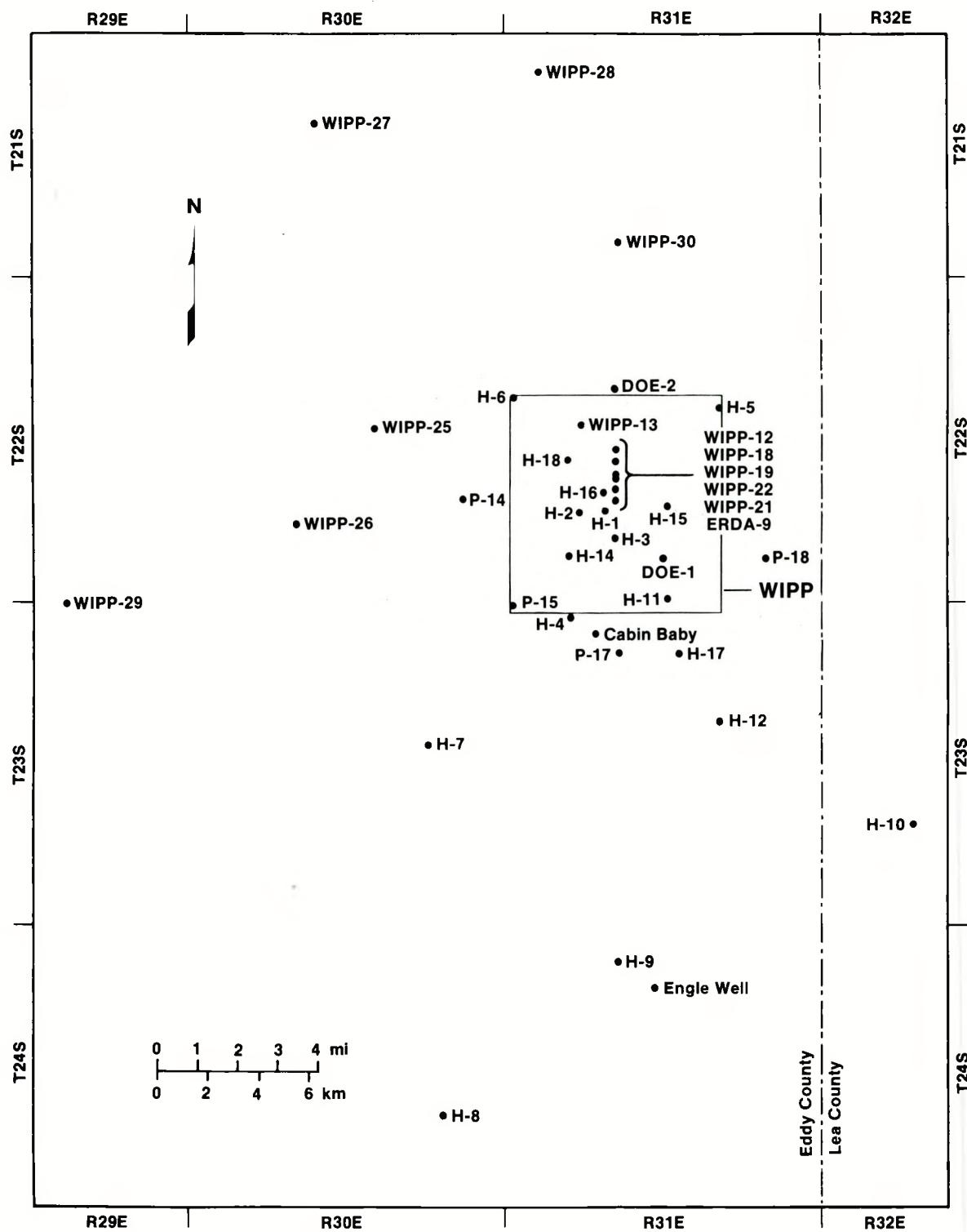
Eighteen samples were selected from various drill cores located in the vicinity of the Waste Isolation Pilot Plant (WIPP) site (Figure III-1).

Selection Criteria

Samples for this study were selected primarily on the basis of a well-defined, water-bearing fracture surface. Five samples were selected from the WIPP-29 and WIPP-32 cores located in the vicinity of Nash Draw (Figure III-1). In these cores, the Culebra Dolomite is very shallow, and the extra samples were chosen to better characterize the fractures where near surface alteration has occurred.

Core Locations, Sample Depths, and Descriptions

Table III-1 lists the well number, sample depths, and descriptions for the 18 samples selected for this study. Figure III-1 shows the locations of the various wells.



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Figure III-1. Location of Wells in the Vicinity of the WIPP Site.

Table III-1. Sample Locations and Descriptions

Sample ID	Well	Depth (feet)	Description
CS1	WIPP-12	838.5-838.7	Massive dolomite (vuggy, dark grey)
CS2	WIPP-13	712.1-712.4	Brecciated dolomite (fractured, dark grey)
CS3	WIPP-13	705.3	Massive dolomite (laminated, brown)
CS4	WIPP-13	714.0	Massive dolomite (light brown)
CS5	WIPP-26	187.0	Brecciated dolomite
CS6	WIPP-27	305.0	Brecciated dolomite (friable, light grey)
CS7	WIPP-28	447.5	Massive clay/dolomite (laminated, dark grey)
CS8	WIPP-29	27.0	Massive dolomite (light grey)
CS9	WIPP-30	633.5	Massive dolomite (contorted laminae, tan)
CS10	WIPP-30	639.0	Massive dolomite (vuggy, tan)
CS11	WIPP-30	635.0	Massive dolomite (brecciated, tan)
CS12	WIPP-32	57.0	Massive dolomite (vuggy)
CS13	WIPP-32	91.1	Claystone (grey)
CS14	WIPP-32	55.0	Massive limestone (brown)
CS15	WIPP-32	56.0	Massive limestone (red)
CS16	WIPP-32	62.0	Massive dolomite (light grey)
CS17	WIPP-33	57.0	Massive gypsum (brown)
CS18	WIPP-34	836.0	Massive dolomite (grey)

IV. FRACTURE SURFACE PETROGRAPHY

Hand Specimen Description

Water-bearing fracture surfaces are easily identified in hand specimen: they are much darker (dark grey/brown to black) than freshly broken surfaces (Figures IV-1 to IV-4). The horizontal fractures almost always occur along clay-rich seams, either layers of nearly pure clay or dolomite (and calcite) layers that are especially rich in clay minerals (Figures IV-1 and IV-4). The darkening can be attributed to two factors: (1) the presence of clay minerals (clay separates from Culebra Dolomite rocks are usually dark grey to black), and (2) oxidation of surface minerals due to contact with oxygen-rich waters. The zone of penetration of the darkened (altered) surface area is usually not deep, typically a few tens of microns, but in some samples where the rock near the fracture surface is friable, the darkened zone can penetrate a few millimeters. When the fracture occurs along a layer of pure clay, a dense system of horizontal fractures may develop, rather than a single fracture. Clay layers may be stripped away to reveal successive darkened (altered) surfaces, which are only a few tens or hundreds of microns apart. Thus, a clay seam can become a multilayer channel and present a very large surface area to the fluid moving in the fractures. Fractures that occur in rock that is primarily dolomite or calcite are single layered, and the altered zone is shallow.

Figure IV-3 shows a very pitted fracture surface. This surface is similar to the interior of large vugs that are frequently seen in Culebra Dolomite. Although the majority of the vugs are not interconnected and do not form a part of the transport network, some undoubtedly do, and this surface may be part of a large void in which water did flow.

Optical Microscopy

Polished thin sections perpendicular to the fracture surfaces were prepared from samples in which the surface was not too friable. Of the eight sections prepared, only four fracture surface rims were preserved: CS1, CS4, CS14, and CS18 (Figures IV-6 to IV-9).

Figure IV-5 shows a clay-rich area along the fracture surface rim. The clay area dominates the rim, and very little dolomite is directly on the surface. Areas like this one are not unusual, but the zone near the fracture rim shown in Figure IV-6 is a far more common feature. This area is quite clay-rich, but the rim itself appears to be composed of about 50% clay and 50% dolomite.

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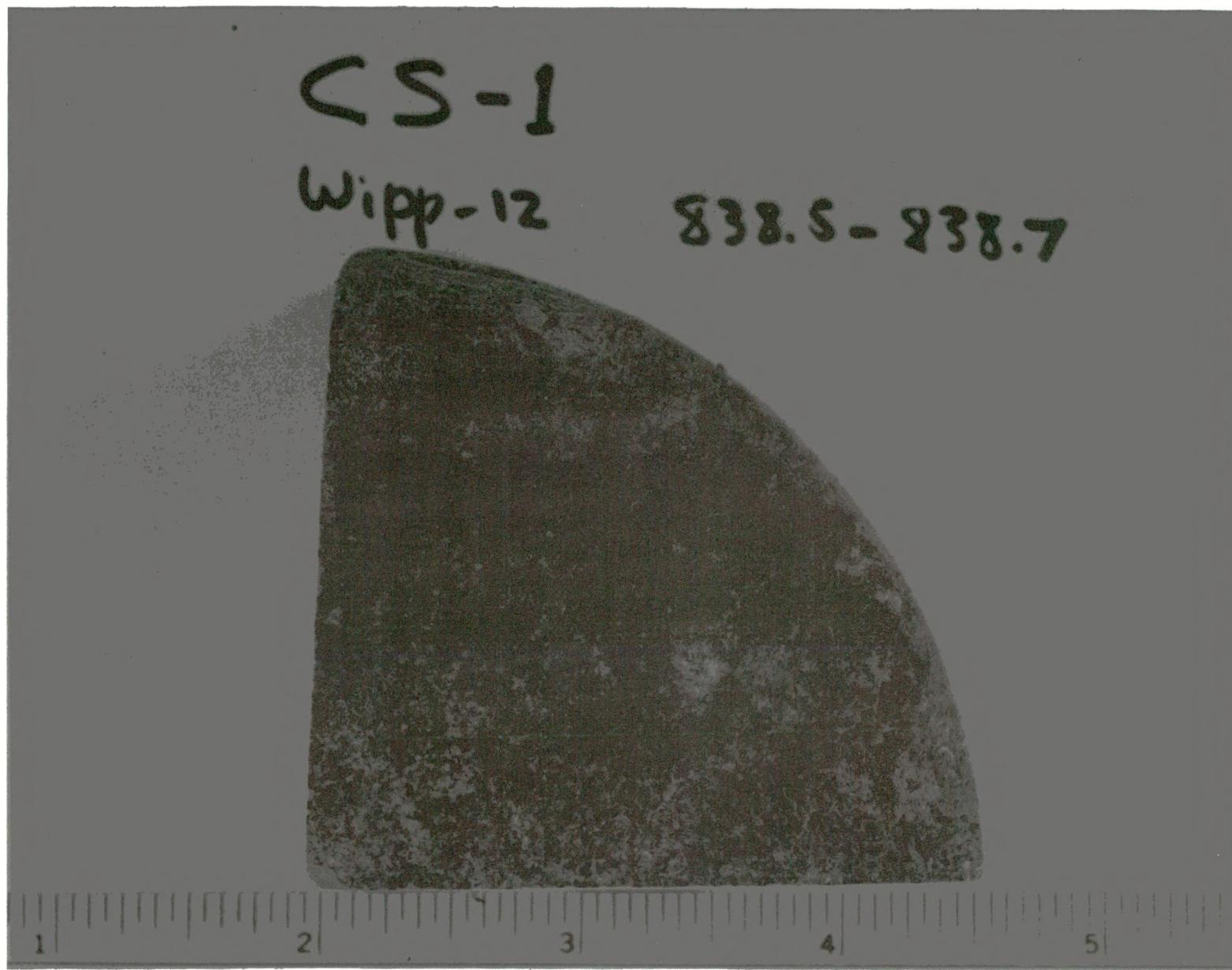


Figure IV-1. Fracture Surface of Sample CS1, Top View.

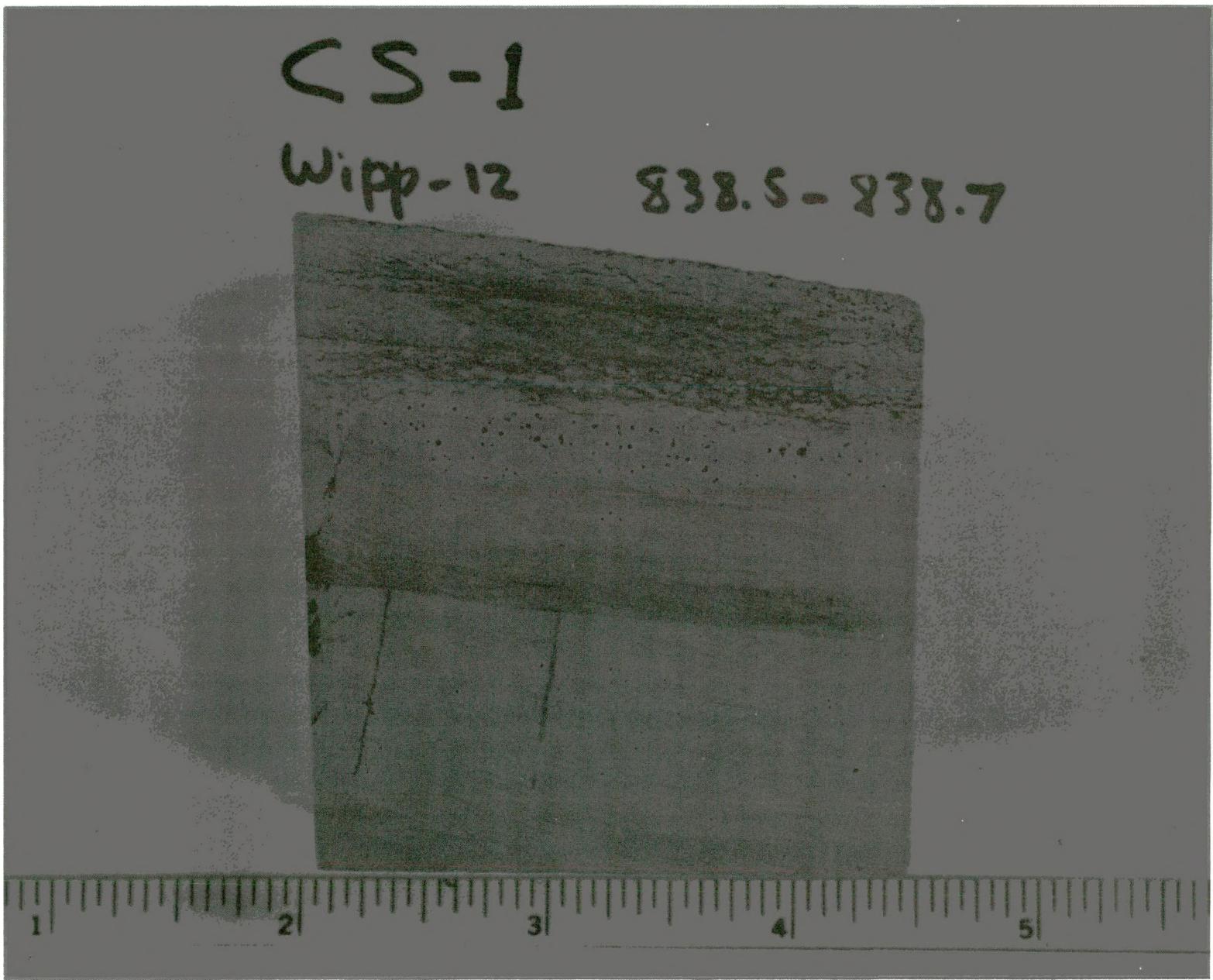


Figure IV-2. Fracture Surface of Sample CS1, Side View.

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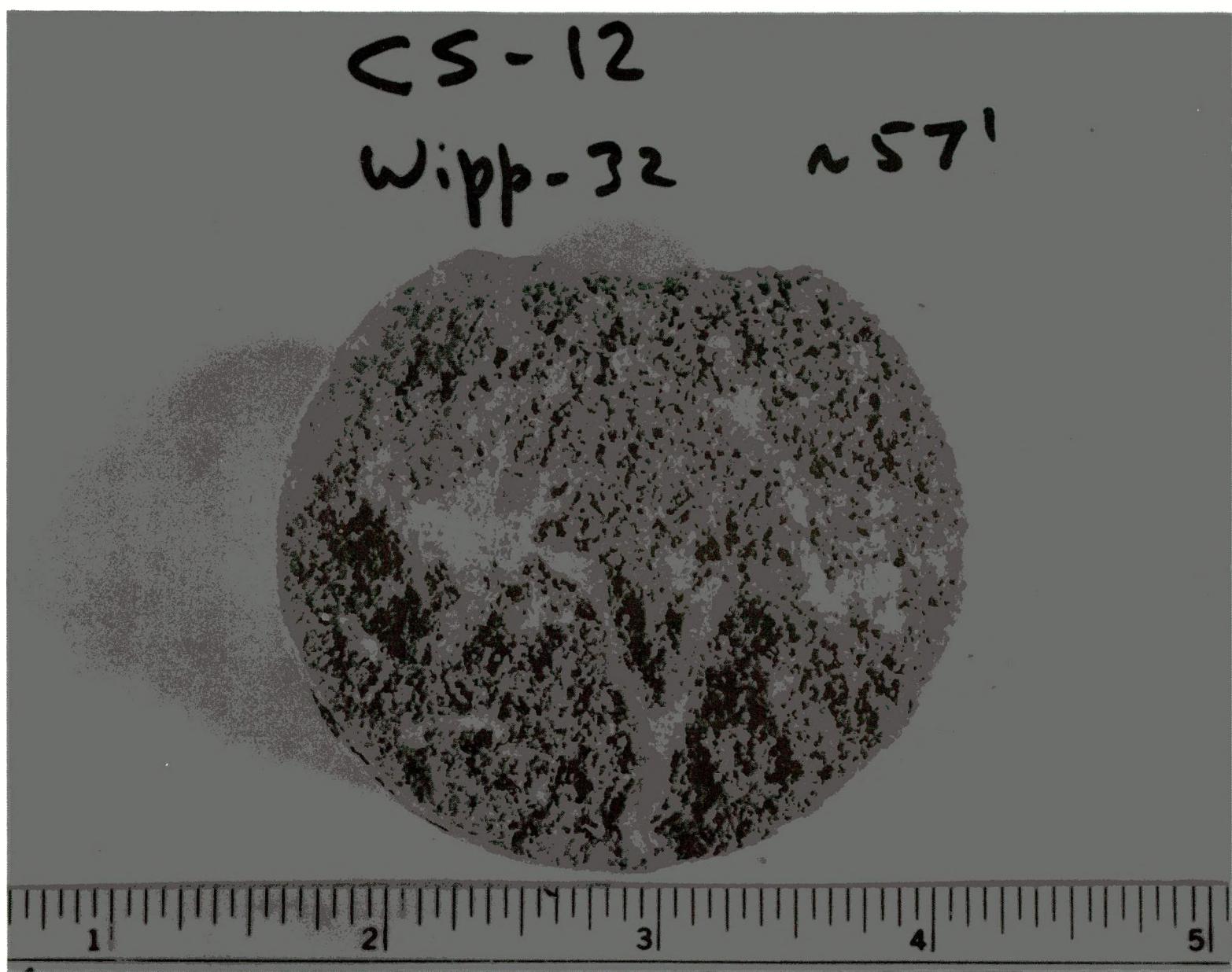


Figure IV-3. Fracture Surface of Sample CS12, Top View.



Figure IV-4. Fracture Surface of Sample CS18, Top View.

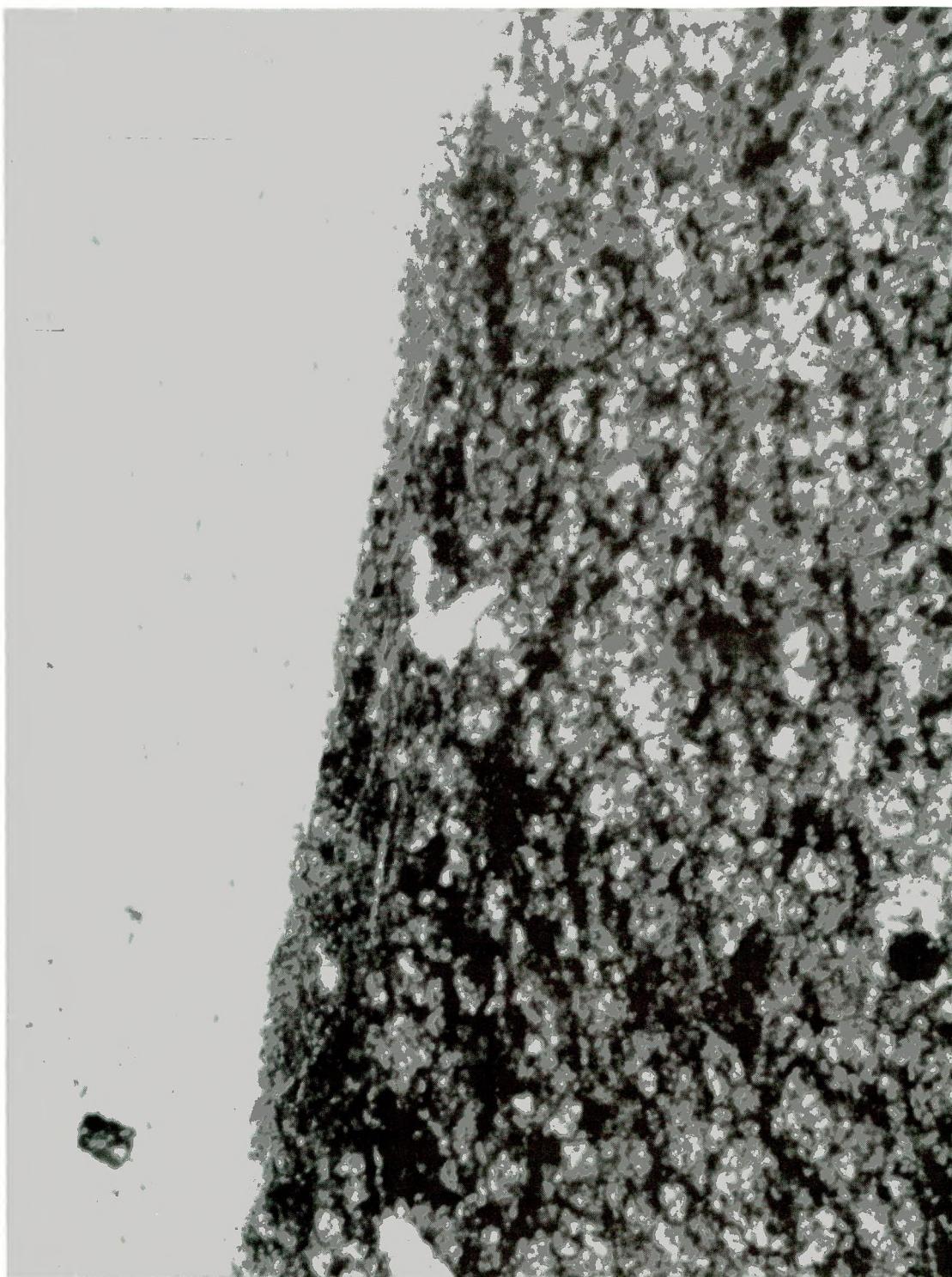


Figure IV-5. Thin-Section Photomicrograph of Fracture Surface Rim of Sample CS4. Plane polarized light, M=1000X. Note dense clay-rich area on surface rim.

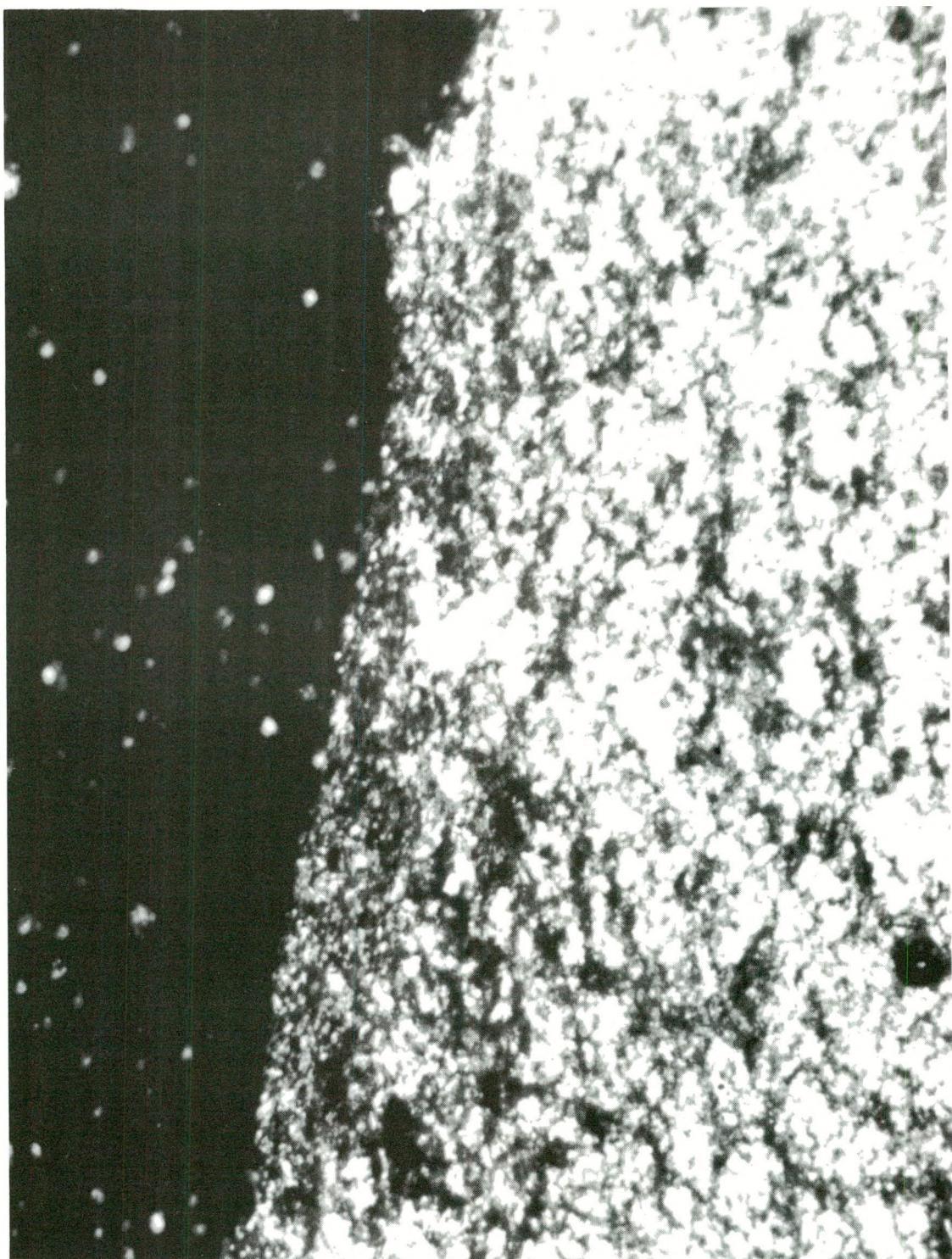


Figure IV-6. Thin-Section Photomicrograph of Fracture Surface Rim of Sample CS4. Crossed polars, $M = 1000X$. Note clay-rich area near surface rim.

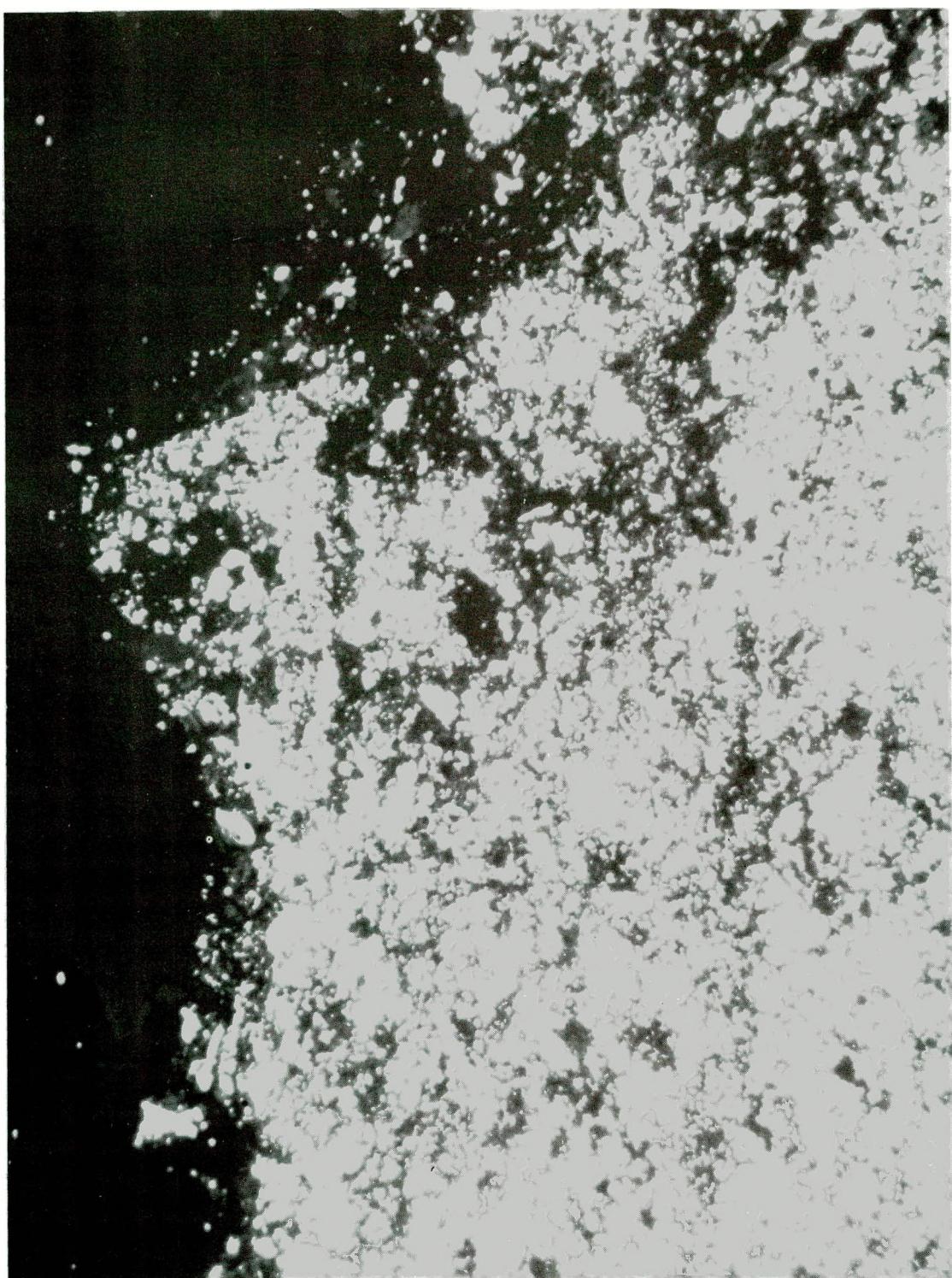


Figure IV-7. Thin-Section Photomicrograph of Fracture Surface Rim of Sample CS4. Crossed polars, M=1000X. Note gypsum lining on fracture rim.

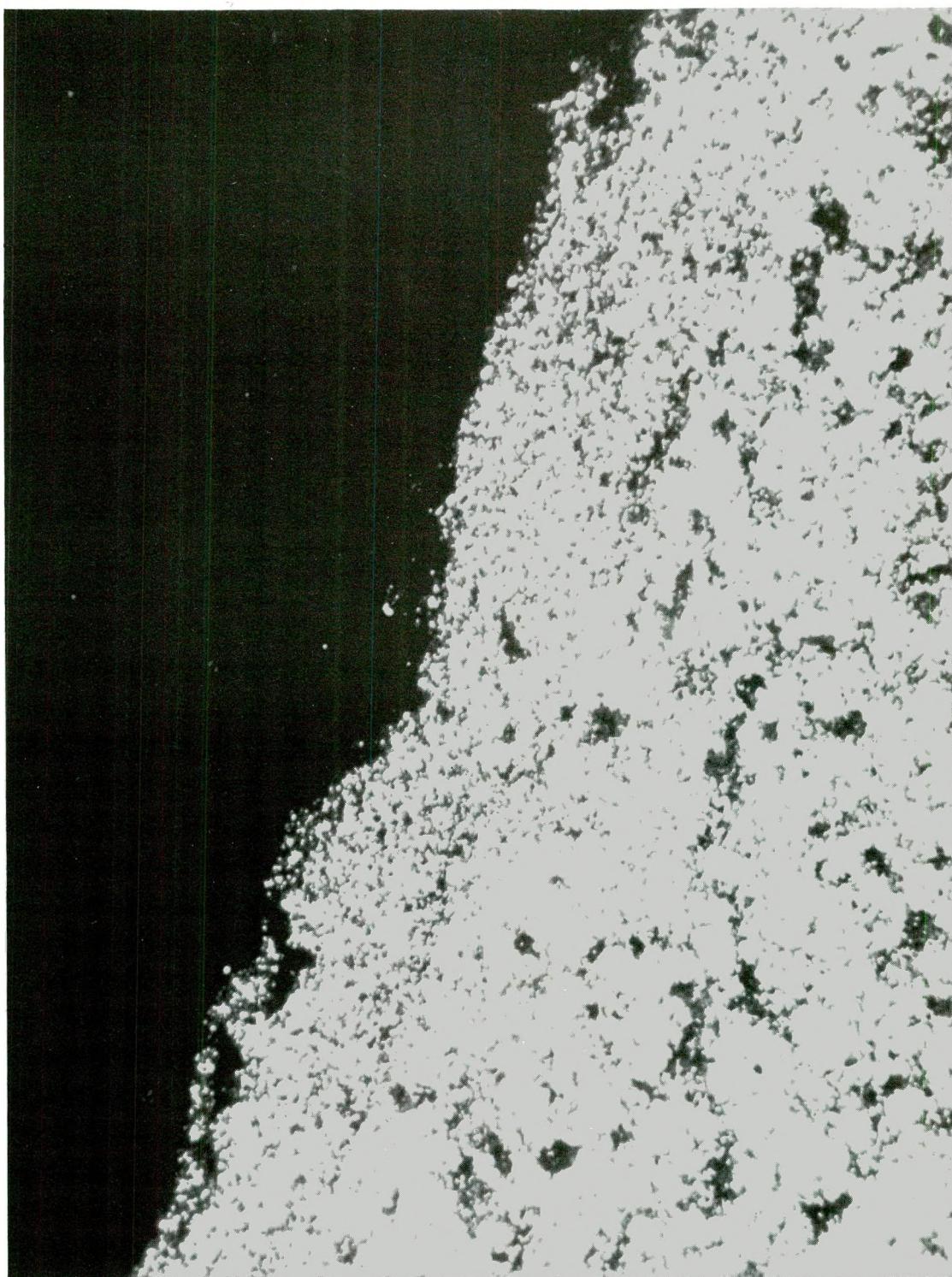


Figure IV-8. Thin-Section Photomicrograph of Fracture Surface Rim of Sample CS18. Crossed polars, M=1000X. Note fine-grained dolomite on fracture rim.



Figure IV-9. Thin-Section of Photomicrograph of Fracture Surface Rim of Sample CS15. Crossed polars, M=1000X.

Figure IV-7 shows a gypsum and clay-rich area near the fracture surface. Along the rim, a lining of gypsum shields the clay from the fracture surface, so there was probably little contact between the water moving in the fracture and the clay minerals near the surface.

In Figure IV-8 there is an area of extremely fine-grained dolomite near the fracture surface. This area appears to be free of clay. The average grain size in the normal dolomite away from the rim is about $7\mu\text{m}$, compared to about $0.6\mu\text{m}$ in the fine-grained area. This feature was not observed in the other thin sections or elsewhere in the thin section of sample CS4 and is probably fairly rare. It may be interpreted in one of two ways: (1) the fine-grained area is a zone of alteration in which the dolomite has recrystallized into "micromicritic" dolomite, or (2) the area is a primary feature, and the fracture traversed the fine-grained area because the rock was weaker there. Neither explanation is particularly satisfactory since, in the first case, dolomite that recrystallizes from calcite, for example, is invariably coarse-grained; in the second case, it would be a great coincidence for such an unusual feature to be directly in the path of the fracture.

Figure IV-9 shows coarse-grained calcite (about $25\mu\text{m}$ in diameter) near the fracture rim of sample CS15, a limestone. Although the area is clay-rich, calcite grains dominate the fracture rim itself. The texture and grain size of the calcite are typical for the limestones, both near the fracture surfaces and away from them, although larger grain sizes were observed, and twinned crystals are common. This coarse-grained subhedral calcite is quite different from that found at the top of the Culebra in the WIPP-19 core (754' depth, Sowards, Glenn, and Keil, 1991). In that case, the calcite was micritic (grain size about $4-5\mu\text{m}$), with anhedral grains, and very clay-rich. The difference in texture might be explained by assuming that the calcite in the WIPP-19 core is the result of primary deposition, and the coarse-grained calcite observed in these samples is the result of dedolomitization (calcitization) of dolomite due to near surface aqueous alteration.

V. WHOLE ROCK AND FRACTURE SURFACE COMPOSITIONS

The compositional data for the bulk rock and fracture surfaces, measured in component oxides, are listed in Tables V-1 and V-2. These tables are derived from Tables C-1 and C-2 in Appendix C by subtracting the halite component and renormalizing. The data for the bulk rock were obtained by x-ray fluorescence spectroscopic analysis (XRF), with the exception of the values for SO_3 , which were measured gravimetrically. For the fracture surface analyses, atomic absorption spectroscopy (AA) was used, with the exception of the sulfate measurements. Based on the totals, it is estimated that the XRF data for the bulk rock are accurate to about 10% of the amounts present. The fracture surface analyses were obtained from scrapings from the surfaces, and very little material was obtained (typically 50 mg). The AA data for such a small sample are probably not accurate to more than about 50% of the amount present.

Figures V-1 to V-8 show the composite bulk rock and fracture surface abundances of SiO_2 , Al_2O_3 , MgO , and CaO in bar-diagram form. In the bar-diagrams, it can be easily seen that SiO_2 is greatly concentrated on the fracture surfaces (Figures V-1 and V-2). In all samples, except CS7 and CS13, the fracture surface concentration is greater than the bulk rock concentration (in samples CS1 and CS3 the fracture surface concentrations were not measured due to insufficient sample). Similarly, Al_2O_3 is higher in concentration on the fracture surfaces in all samples (Figures V-3 and V-4), with the same two exceptions mentioned above, CS7 and CS13. The increase in SiO_2 and Al_2O_3 can be explained by the fact that fracture surfaces tend to occur along clay- and quartz-rich seams in the rock.

The fracture surface MgO concentrations are lower than those in the bulk rock in all samples except CS6, CS12, CS14, CS15, and CS17 (Figures V-5 and V-6). Since MgO occurs in both clay and dolomite, its behavior is not quite as easily explained as that of SiO_2 and Al_2O_3 , which occur only in the clastic components. An increase in clay and quartz on the fracture surfaces is accompanied by a decrease in dolomite, so if the sample contains primarily dolomite (as in samples CS1 to CS11), one would expect a slight decrease in MgO if both SiO_2 and Al_2O_3 increase. The increase in MgO on the fracture surfaces of samples CS14, CS15, and CS17 is due to the fact that CS14 and CS15 are limestones and contain primarily calcite, and CS17 is gypsum: since the fractures occur along clay-rich seams, and there is essentially no MgO in the major mineral components, MgO is consequently enriched on the surfaces.

The behavior of CaO is also somewhat complicated, since it is a component of dolomite, calcite, and gypsum. In dolomite rocks (CS1 through CS12, CS16 and CS18), one would expect CaO to be higher in the bulk rock than in the fracture

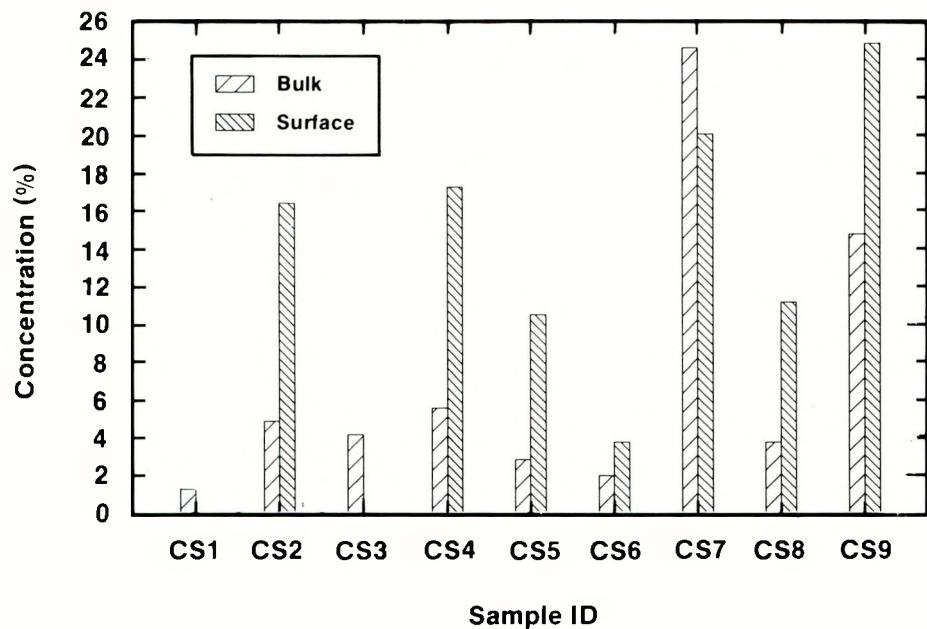
Table V-1. Bulk Rock Compositions

Sample ID	Well	Depth	SiO ₂	TiO ₂	Al ₂ O ₃	Fe ₂ O ₃	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	SO ₃	Total
CS1B	WIPP-12	838.60	1.27	0.02	0.24	0.34	0.03	20.22	29.50	0.07	0.07	0.04	0.16	51.94
CS2B	WIPP-13	712.30	4.87	0.08	1.15	0.60	0.02	19.50	27.40	0.04	0.37	0.04	0.07	54.13
CS3B	WIPP-13	705.30	4.14	0.06	0.98	0.47	0.03	19.27	27.80	0.06	0.31	0.04	0.07	53.23
CS4B	WIPP-13	714.00	5.56	0.05	1.14	0.56	0.02	19.30	27.60	0.05	0.37	0.04	0.04	54.72
CS5B	WIPP-26	187.50	2.82	0.04	0.65	0.37	0.04	19.65	27.80	0.07	0.29	0.04	0.05	51.81
CS6B	WIPP-27	305.00	1.95	0.03	0.41	0.29	0.02	20.05	27.90	0.25	0.24	0.05	0.09	51.28
CS7B	WIPP-28	447.50	24.02	0.29	5.51	0.78	0.03	19.02	18.89	1.29	1.19	0.03	0.10	71.15
CS8B	WIPP-29	27.00	3.73	0.06	0.84	0.44	0.02	19.80	27.50	0.22	0.34	0.05	0.06	53.05
CS9B	WIPP-30	633.50	14.27	0.17	2.81	0.09	0.01	19.15	26.14	1.75	1.12	0.13	0.19	65.83
CS10B	WIPP-30	639.00	2.70	0.04	0.55	0.34	0.02	20.40	29.90	0.05	0.18	0.10	0.05	54.32
CS11B	WIPP-30	635.00	1.35	0.02	0.33	0.30	0.02	20.05	29.80	0.30	0.13	0.10	0.07	52.46
CS12B	WIPP-32	57.00	0.35	0.01	0.10	0.08	0.01	11.40	43.60	0.05	0.02	0.18	0.12	55.92
CS13B	WIPP-32	91.10	53.36	0.74	12.96	4.32	0.01	16.15	0.95	0.05	2.32	0.21	0.03	91.10
CS14B	WIPP-32	55.00	2.20	0.02	0.38	0.15	0.02	0.84	54.40	0.04	0.13	0.25	0.13	58.56
CS15B	WIPP-32	56.00	2.95	0.03	0.50	0.22	0.02	0.84	52.60	0.04	0.17	0.25	0.12	57.73
CS16B	WIPP-32	62.00	1.55	0.02	0.32	0.14	0.01	17.40	32.50	0.05	0.12	0.12	0.07	52.30
CS17B	WIPP-33	570.00	1.77	0.02	0.24	0.10	0.00	0.35	32.20	0.10	0.04	N.A.	45.97	80.79
CS18B	WIPP-34	836.00	2.45	0.04	0.50	0.34	0.02	20.20	27.80	0.06	0.18	0.10	0.09	51.77

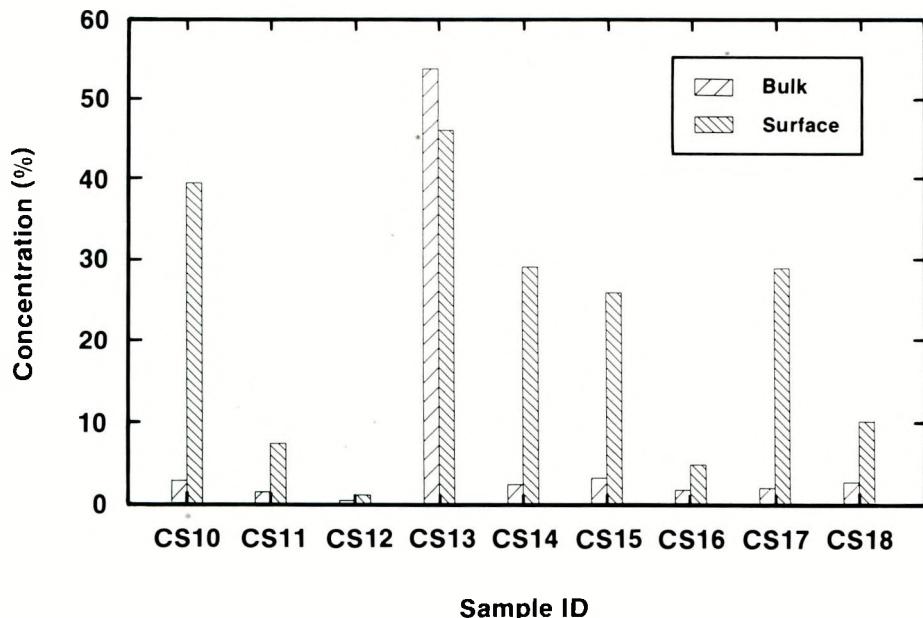
Table V-2. Fracture Surface Compositions (NaCl Data Subtracted)

Sample ID	Depth	SiO ₂	Al ₂ O ₃	Fe ₂ O ₃	MgO	CaO	K ₂ O	SO ₃	Total
CS1S	838.60	*	7.80	1.11	15.61	35.67	0.89	*	61.09
CS2S	712.30	16.42	4.54	1.32	17.64	21.27	1.23	*	62.43
CS3S	705.30	*	2.69	0.72	17.97	24.07	1.05	*	46.49
CS4S	714.00	17.28	4.61	1.66	18.28	20.38	0.13	1.48	63.83
CS5S	187.50	10.50	1.43	0.48	19.03	26.31	0.62	*	58.38
CS6S	305.00	3.74	0.65	0.28	20.41	28.51	0.32	*	53.09
CS7S	447.50	20.07	4.95	1.36	18.54	19.75	0.93	0.03	65.62
CS8S	27.00	11.16	1.82	1.48	10.19	24.53	1.25	*	50.43
CS9S	633.50	24.82	6.13	1.66	16.27	16.40	1.95	*	67.24
CS10S	639.00	39.37	8.12	2.06	11.83	10.55	2.16	*	74.10
CS11S	635.00	7.27	2.11	0.58	17.46	26.88	0.61	3.37	58.28
CS12S	57.00	0.98	0.14	0.12	13.26	40.57	0.06	*	55.13
CS13S	91.10	45.79	11.13	3.14	11.37	11.23	1.57	0.29	84.53
CS14S	55.00	28.92	6.88	2.10	9.82	24.04	1.83	0.01	73.60
CS15S	56.00	25.74	5.98	1.97	8.01	28.78	1.34	0.01	71.82
CS16S	62.00	4.56	0.71	0.33	16.91	34.03	0.24	0.61	57.40
CS17S	57.00	28.69	5.82	2.15	4.70	17.64	1.31	21.09	81.40
CS18S	836.00	9.91	2.55	4.21	16.37	28.69	0.66	*	62.40

* = Insufficient Sample

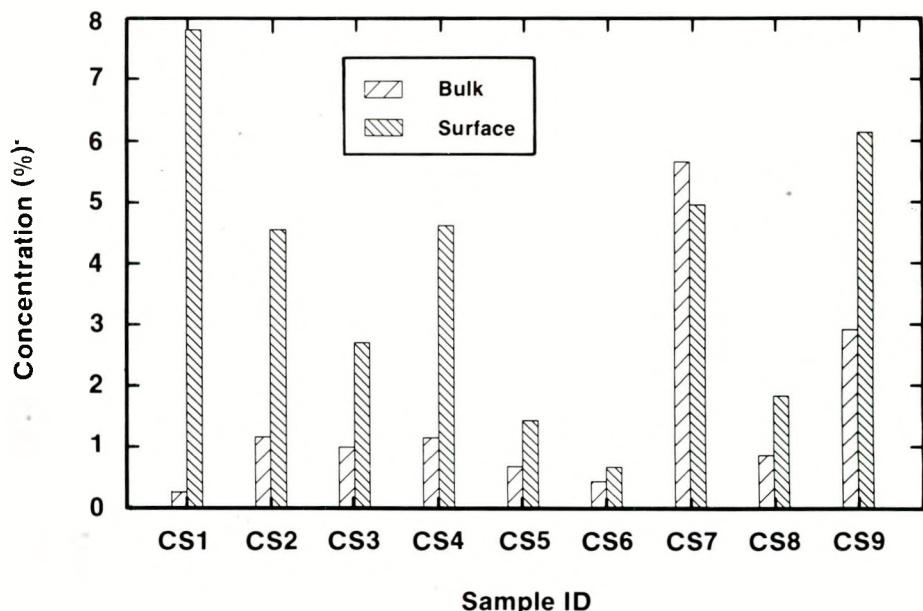


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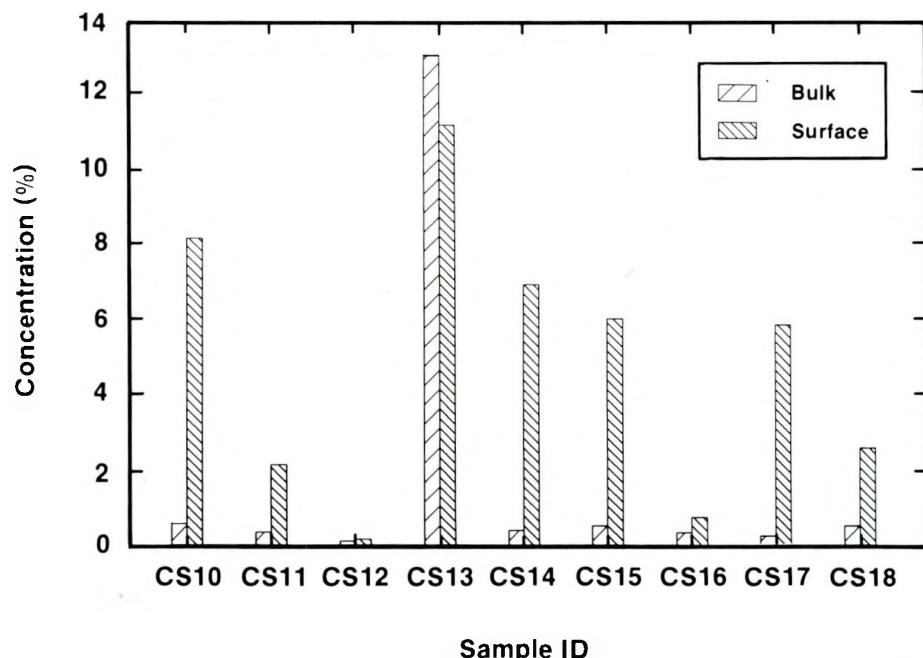
Figure V-1. SiO_2 Concentrations of CS1 through CS9.

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Figure V-2. SiO_2 Concentrations of CS10 through CS18.

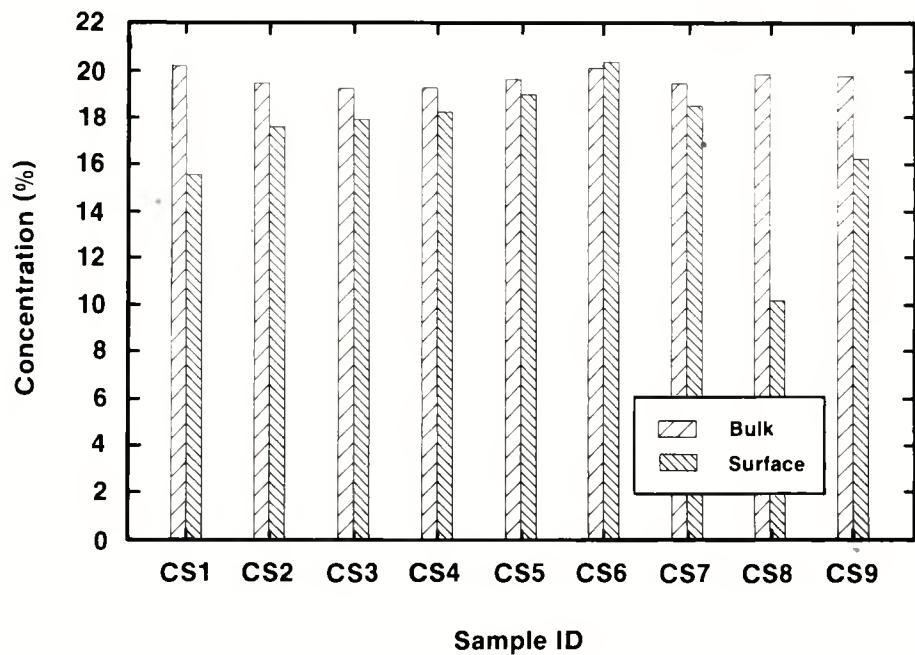


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Figure V-3. Al_2O_3 Concentrations of CS1 through CS9.

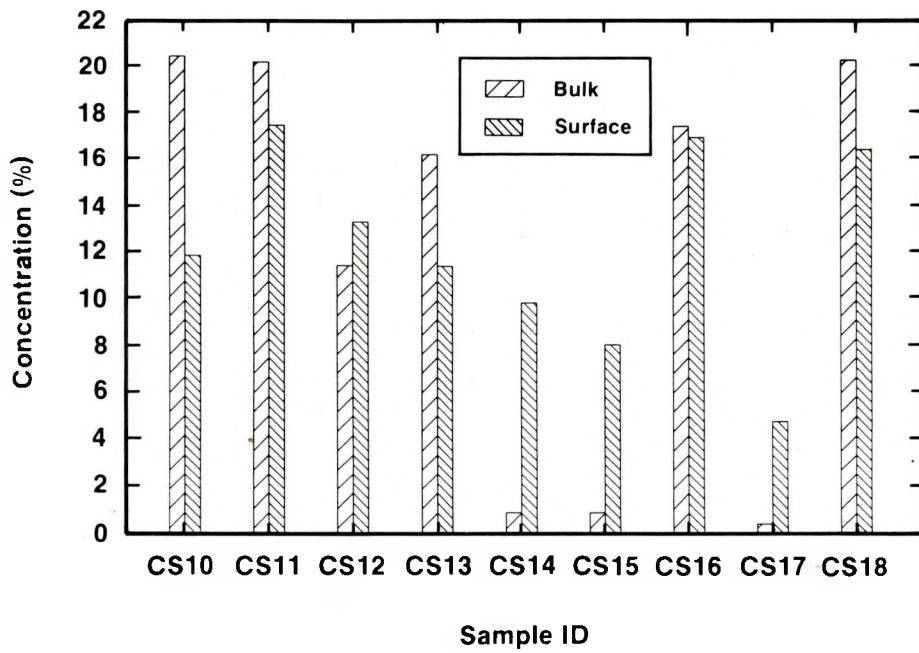
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Figure V-4. Al_2O_3 Concentrations of CS10 through CS18.



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Figure V-5. MgO Concentrations of CS1 through CS9.



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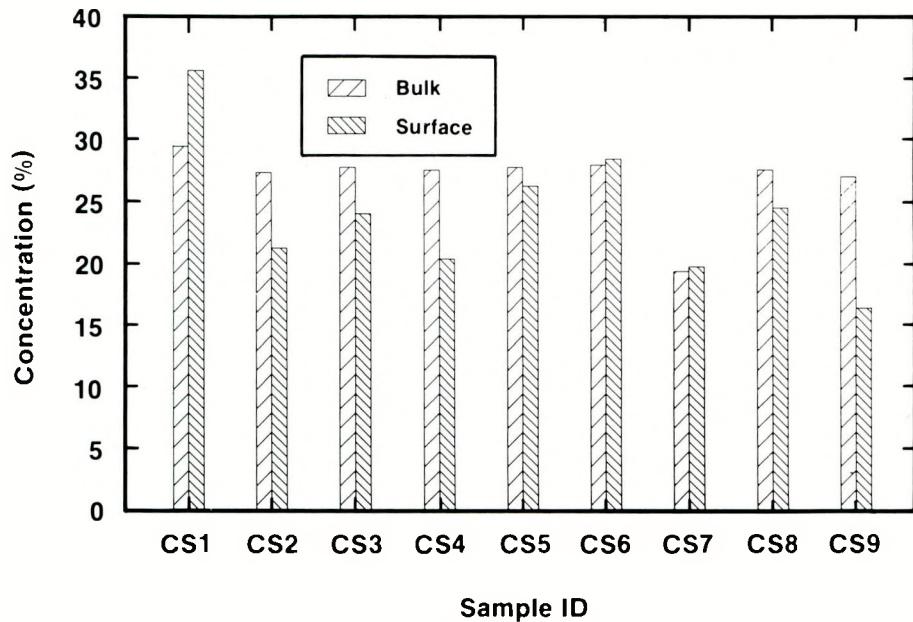
Figure V-6. MgO Concentrations of CS10 through CS18.

surface, and this is generally the case, with a few exceptions, in which the bulk and surface concentrations are similar (Figures V-7 and V-8). In the two limestones, CS14 and CS15, the surface CaO is much lower in concentration than that in the bulk rock, and in the claystone (CS13), the surface mode is greater.

Fe₂O₃ is concentrated on the fracture surfaces in all samples except CS6, where they are nearly equal, and CS13, where the bulk rock concentration is higher. This trend closely follows that of the clay mineral modes (Figures V-9 and V-10), so it may be assumed that the concentration of Fe₂O₃ on the surface is due to the Fe₂O₃ content of the clay minerals.

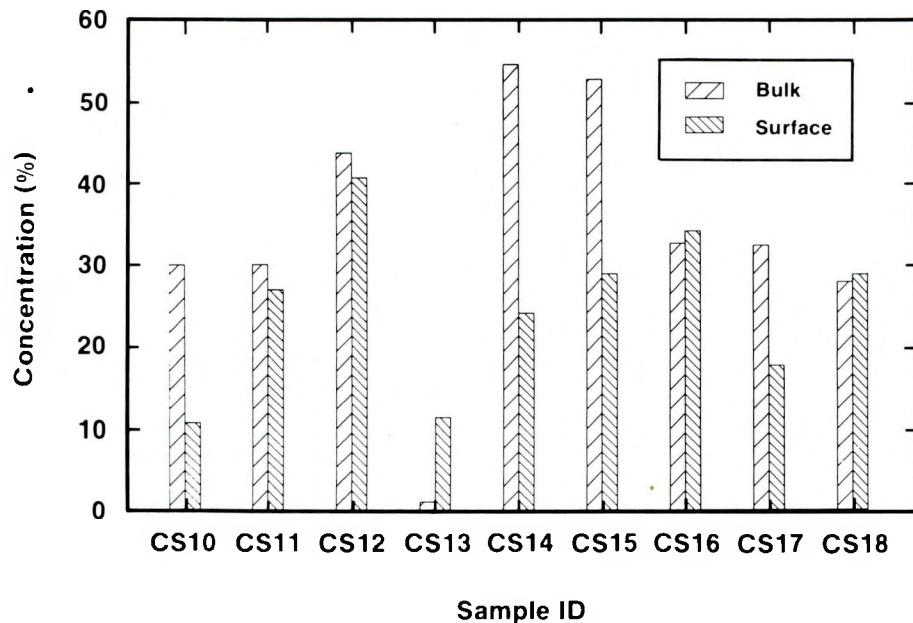
K₂O is also concentrated on the fracture surfaces (Tables V-1 and V-2). This is also due to the higher clay mode on the surfaces, since illite is one of the components of the clay mineral assemblage (Sewards, Williams, and Keil, 1991), and illite typically contains about 7-8% K₂O.

Since the variation in composition is intimately associated with the variation in mineralogy, the above discussion may be more easily understood when the individual mineral modes are discussed in the following section.



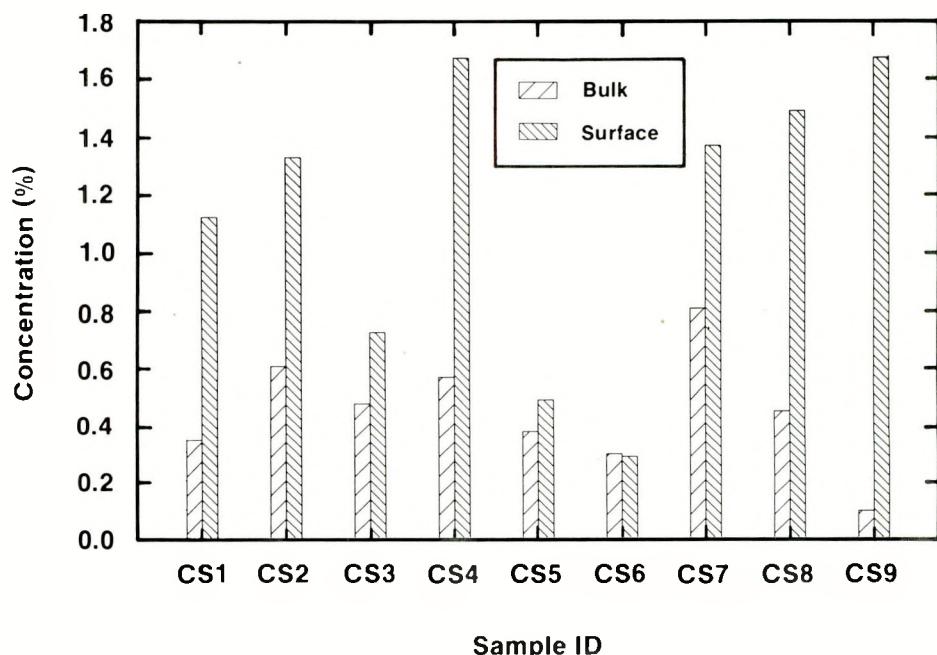
TRI-6342-459-0

Figure V-7. CaO Concentrations of CS1 through CS9.

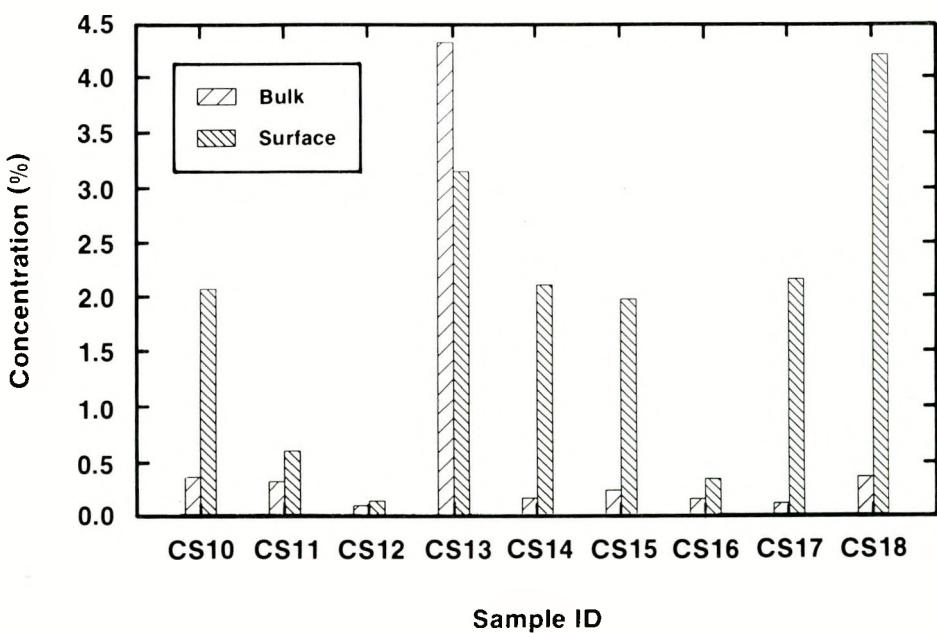


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Figure V-8. CaO Concentrations of CS10 through CS18.



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Figure V-9. Fe₂O₃ Concentrations of CS1 through CS9.

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Figure V-10. Fe₂O₃ Concentrations of CS10 through CS18.

VI. WHOLE ROCK AND FRACTURE SURFACE MINERALOGY

Table VI-1 lists the minerals that were identified in the bulk rock and fracture surfaces by XRD. These include dolomite, calcite, gypsum, halite, quartz, and clay. Halite in these samples is an artifact of the drilling process: a mixture of brine and mud was used to drill the wells; when the fluid evaporated, halite precipitated on the core surfaces.

The results of the whole rock and fracture surface mineral mode calculations, which are based on the compositional data included in the previous section, are listed in Tables VI-2 and VI-3. These tables are derived from Tables C-3 and C-4, listed in Appendix C, by subtracting the halite modes and normalizing to 100%. The mineral mode calculation method is discussed in Appendix B.

Clay

The clay modes for the bulk rock and fracture surface scrapings are displayed in bar diagram form in Figures VI-1 and VI-2. With the exception of two samples, CS7, a clay-rich dolomite, and CS13, a claystone, fracture surface clay contents are considerably higher than those of the bulk rock. The average bulk rock clay mode for these samples is 4.6% (excluding CS7 and CS13), whereas the average mode in the fracture surface scrapings is 18%, nearly four times the bulk rock average.

The reason the clay modes in the fracture surface scrapings are so much higher than the bulk rock modes is that the fractures occur primarily along clay-rich seams, since these are the weakest layers in the rock. In sample CS13, a claystone, the surface mode is less than that of the bulk mode. This is due to the fact that secondary calcite has been deposited on the fracture surface (Figure VI-7). Similarly, in sample CS7, the bulk clay mode is greater than the mode in the fracture surface scrapings because the fracture occurred along a quartz-rich layer (Figure VI-3).

Quartz

The quartz modes follow approximately the same trend as the clay modes: quartz modes in the fracture surface scrapings are substantially greater than those of the bulk rock (Figures VI-3 and VI-4). The zero values for the modes in the fracture surface scrapings in samples CS1 and CS3 in Figure VI-3 are due to the fact that there was insufficient sample to determine SiO_2 in these samples (Table VI-2).

Table VI-1. Semi-Quantitative Mineral Modes Determined by XRD

Sample ID	Well	Depth	Bulk				Surface								
			Evaporite Minerals			Clastic Minerals	Evaporite Minerals			Clastic Minerals					
			Dolomite	Calcite	Gypsum	Halite	Quartz	Clay	Dolomite	Calcite	Gypsum	Halite	Quartz	Clay	
CS1	WIPP-12	838.6	****				*	*	**		**	****	**	**	**
CS2	WIPP-13	712.3	****				*	*	****		***	*	**	**	**
CS3	WIPP-13	705.3	****				*	*	****		*	***	**	**	**
CS4	WIPP-13	714.0	****				*	*	****		**	***	**	**	**
CS5	WIPP-26	187.5	****				*	*	****		*	**	**	**	**
CS6	WIPP-27	305.0	****				*	*	****		**	*	*	*	*
CS7	WIPP-28	447.5	****				**	**	****		**	**	**	**	**
CS8	WIPP-29	27.0	****				*	*	***		****	****	*	*	*
CS9	WIPP-30	633.5	****			*	**	**	****		**	**	***	**	**
CS10	WIPP-30	639.0	****				*	*	**		**	*	****	****	****
CS11	WIPP-30	635.0	****				*	*	***	*	**	***	**	**	**
CS12	WIPP-32	57.0	***	***	**		*	*	****	****		*	*	*	*
CS13	WIPP-32	91.1					****	****		***		****	****	****	****
CS14	WIPP-32	55.0		****			*	*		****			**	**	**
CS15	WIPP-32	56.0		****						****			**	**	**
CS16	WIPP-32	62.0	****	**					****	***			**	**	**
CS17	WIPP-33	570.0			****		*	*			****		***	**	**
CS18	WIPP-34	836.0	****				*	*	**		**	****	*	*	*

**** = Very abundant
 *** = Abundant

** = Present
 * = Trace

Table VI-2. Bulk Rock Mineral Modes (Normalized to 100%)

Sample ID	Dolomite	Calcite	Gypsum	Halite	Clay	Quartz	Total
CS1B	97.85				1.61	0.54	100.00
CS2B	90.91				7.73	1.35	100.00
CS3B	92.26				6.59	1.14	100.00
CS4B	90.38				7.57	2.05	100.00
CS5B	94.66				4.49	0.86	100.00
CS6B	96.40				2.87	0.73	100.00
CS7B	58.62				34.66	6.72	100.00
CS8B	93.05				5.76	1.19	100.00
CS9B	77.26			0.83	16.83	5.08	100.00
CS10B	95.46				3.56	0.98	100.00
CS11B	97.48				2.19	0.34	100.00
CS12B	59.47	39.56	0.26		0.67	0.04	100.00
CS13B	3.03				83.77	13.20	100.00
CS14B	0.00	96.46			2.52	1.03	100.00
CS15B	0.00	95.19			3.38	1.44	100.00
CS16B	89.90	7.40			2.13	0.57	100.00
CS17B	0.00		97.40		1.58	1.02	100.00
CS18B	95.56				3.48	0.96	100.00

Table VI-3. Fracture Surface Mineral Modes (Normalized to 100%)

Sample ID	Dolomite	Calcite	Gypsum	Clay	Quartz	Total
CS1S	25.01		43.31	31.68	0.00	100.00
CS2S	60.18		0.86	26.45	12.51	100.00
CS3S	81.78		0.00	18.22	0.00	100.00
CS4S	59.45		2.64	25.39	12.53	100.00
CS5S	82.73		0.00	8.48	8.79	100.00
CS6S	93.06			3.86	3.08	100.00
CS7S	58.48			26.97	14.55	100.00
CS8S	43.72		33.95	12.08	10.25	100.00
CS9S	45.69		0.47	35.01	18.83	100.00
CS10S	27.89		0.00	43.49	28.62	100.00
CS11S	71.81	3.68	6.44	12.47	5.60	100.00
CS12S	65.19	33.04		0.90	0.87	100.00
CS13S		14.86		54.99	30.14	100.00
CS14S		37.46		40.06	22.48	100.00
CS15S		44.99		34.89	20.12	100.00
CS16S	78.41	13.32		4.37	3.90	100.00
CS17S			41.17	35.22	23.62	100.00
CS18S	78.56		0.00	14.17	7.27	100.00

Dolomite

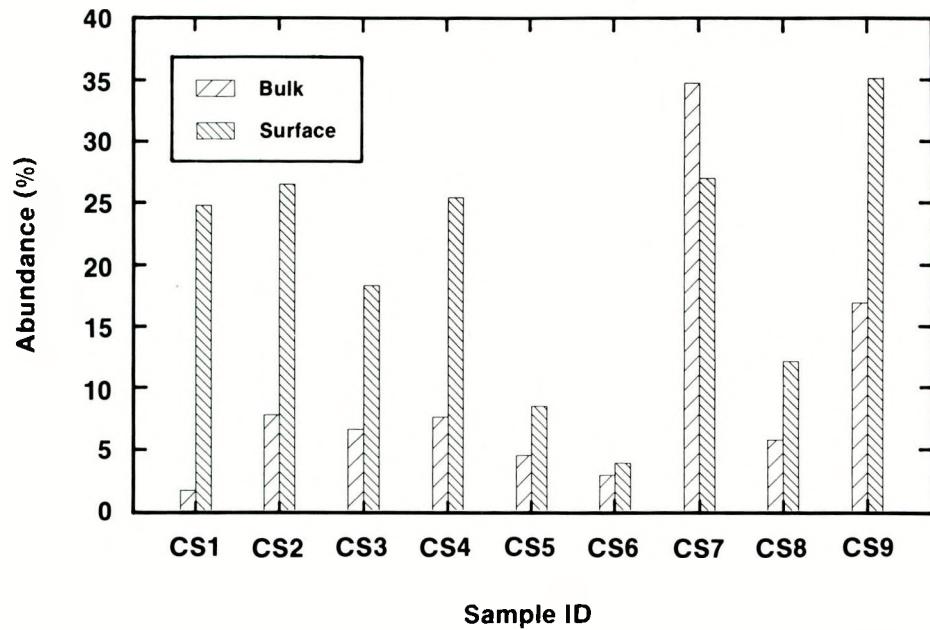
In the samples containing dolomite, the fracture surfaces have less dolomite than the bulk rock, except for sample CS12, where the fracture surface has slightly more (Figures VI-5 and VI-6). In sample CS1, the mode in the fractures surface scrapings is only about 1/3 of the bulk mode; this is due to two factors: the mode in the fracture surface scrapings in this sample is considerably greater than the bulk mode (about 25% vs. 1.6%), and there is about 40% gypsum on the fracture surface. This gypsum is obviously of secondary origin, since the bulk gypsum mode is zero. Sample CS8 is very similar: the dolomite mode in the fracture surface scrapings is about 1/2 of the bulk mode, while there is 12% clay and 34% gypsum on the fracture surface.

Calcite

Calcite only appears in samples CS11 through CS16 (Figure VI-7). With the exception of CS11 (depth 635'), all these samples are from shallow cores (55' to 92' depth). In samples CS11 and CS13, in which the bulk calcite mode is zero, calcite in the fracture surface scrapings is a secondary precipitate. In sample CS16, the mode in the surface scrapings is greater than the bulk mode, which would also indicate secondary precipitation. In the remaining three samples, calcite in the bulk rock appears to have recrystallized from dolomite (see Chapter IV), and the fracture surfaces contain less calcite than the bulk rock. As in the dolomite samples, this is due to the fact that the fractures occurred along clay- and quartz-rich seams.

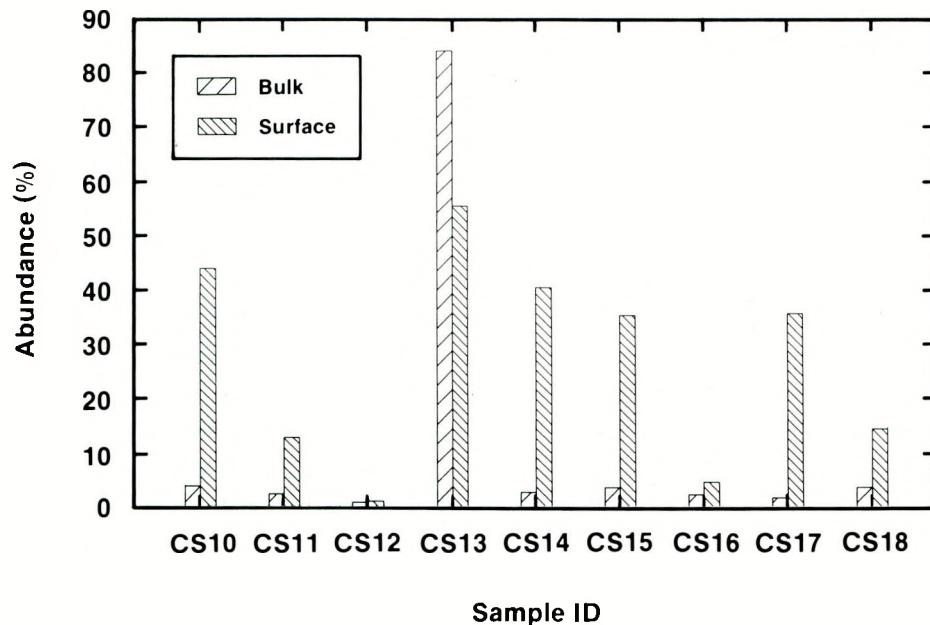
Gypsum

With the exception of sample CS17, which is massive gypsum, gypsum only appears in the fracture surface modes (samples CS1, CS2, CS4, CS8, CS9, and CS11) (Figures VI-8 and VI-9). Clearly, the gypsum precipitated from solution on the fracture surfaces in these samples. In sample CS17, the surface mode is much lower than the bulk mode. Again, it appears that the fracture surface occurred along a clay-rich vein, since the surface clay mode is 35%.



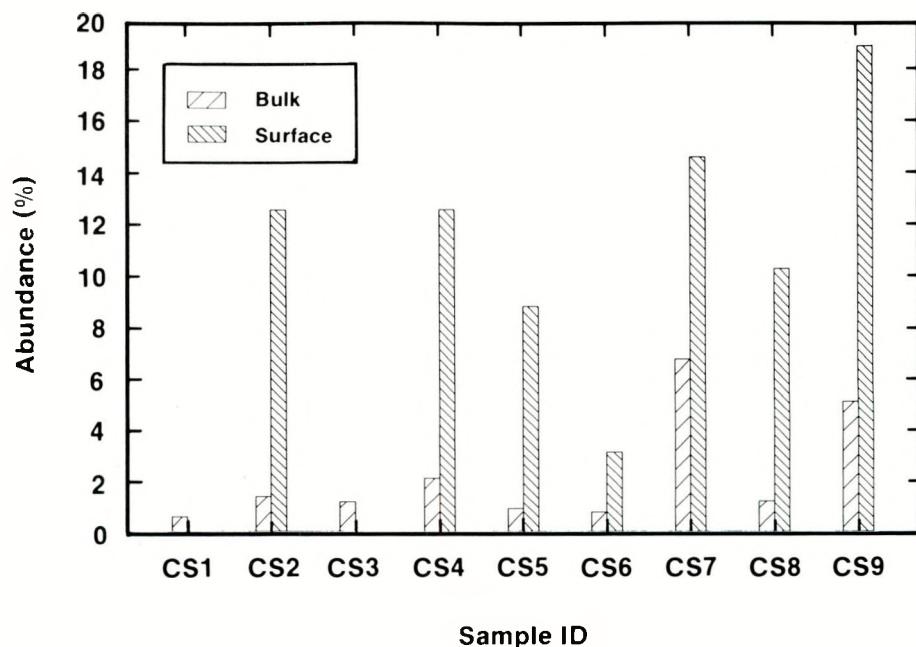
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Figure VI-1. Clay Modes of CS1 through CS9.



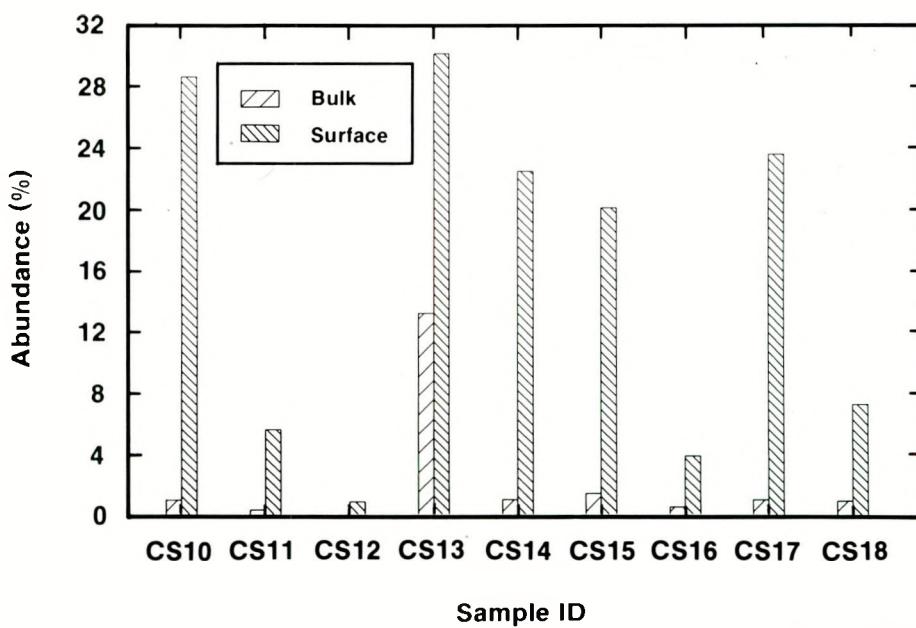
TRI-6342-464-0

Figure VI-2. Clay Modes of CS10 through CS18.



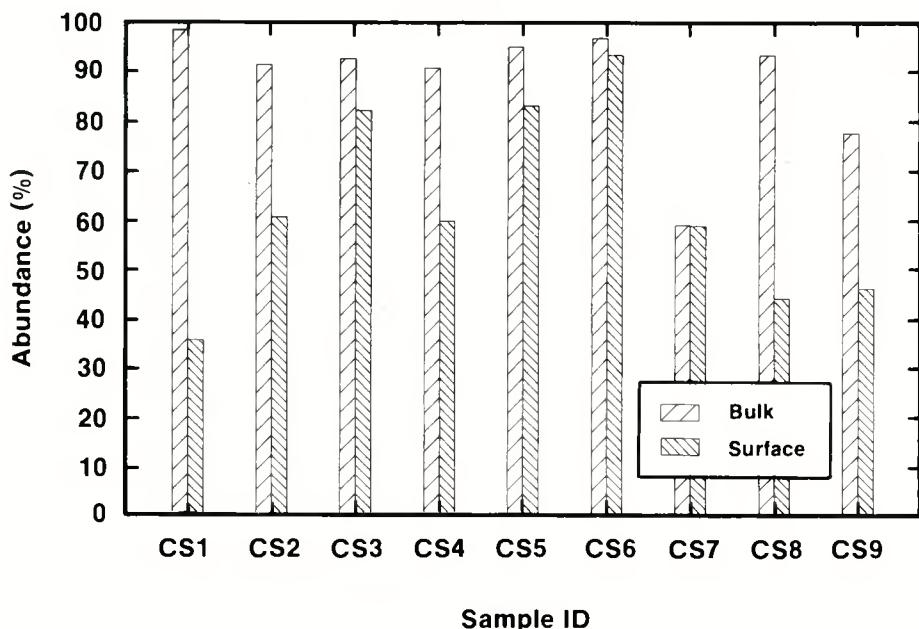
TRI-6342-465-0

Figure VI-3. Quartz Modes of CS1 through CS9.



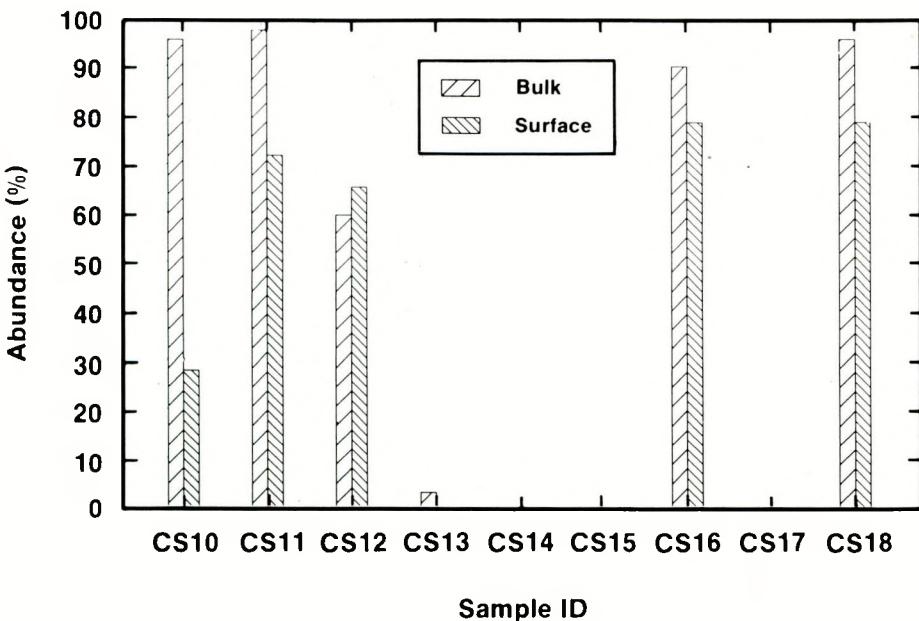
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Figure VI-4. Quartz Modes of CS10 through CS18.



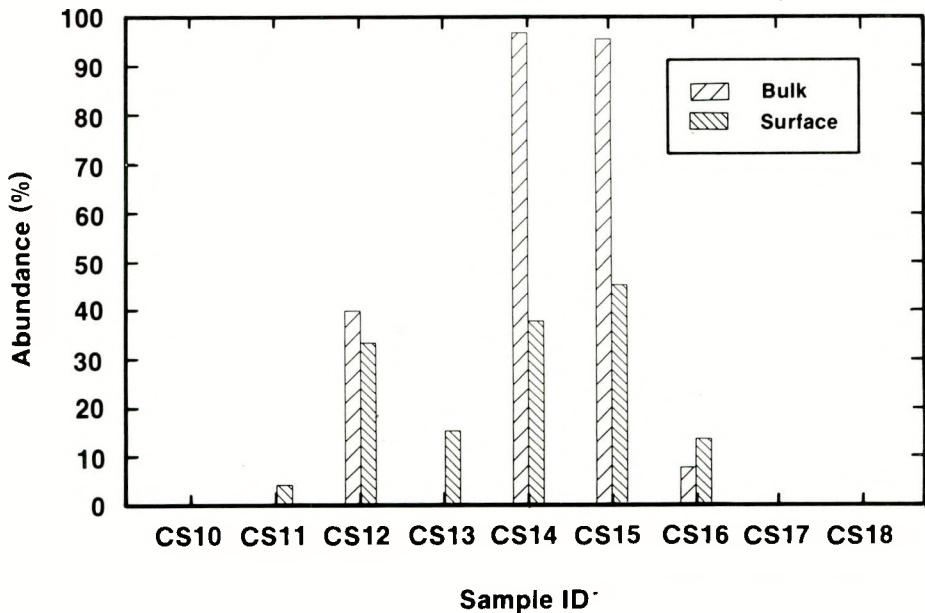
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Figure VI-5. Dolomite Modes of CS1 through CS9.



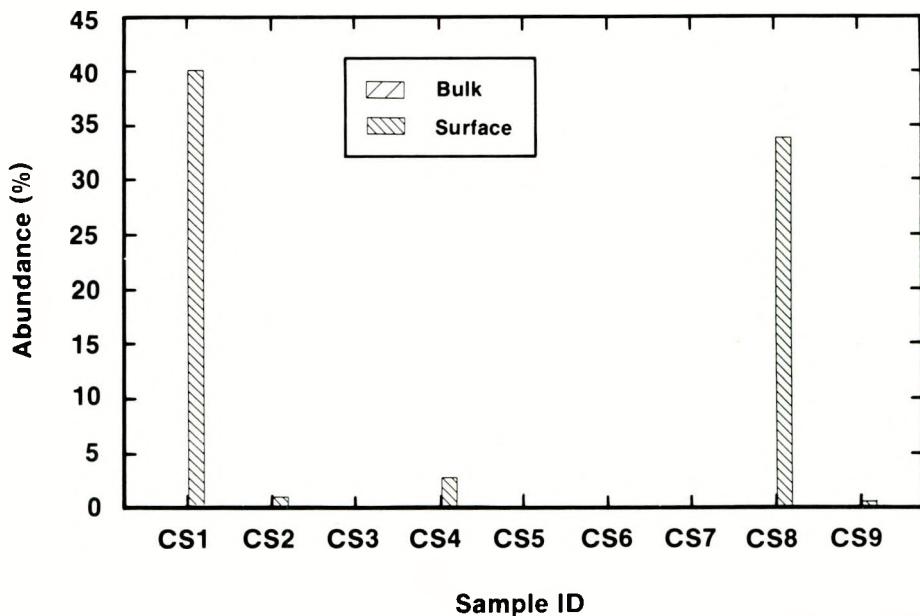
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Figure VI-6. Dolomite Modes of CS10 through CS18.



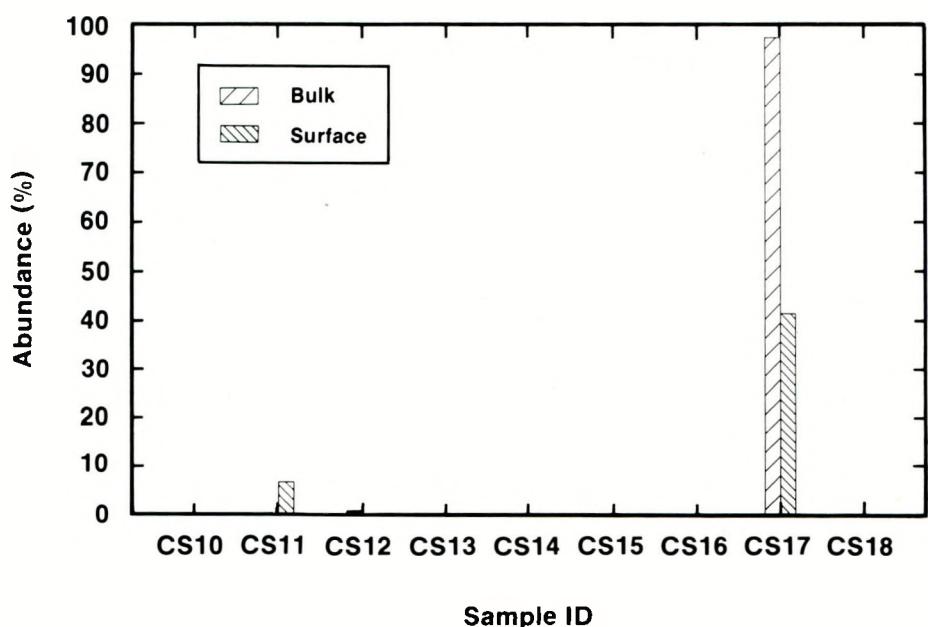
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Figure VI-7. Calcite Modes of CS10 through CS18.



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Figure VI-8. Gypsum Modes of CS1 through CS9.



TRI-6342-471-0

Figure VI-9. Gypsum Modes of CS10 through CS18.

VII. DOLOMITE COMPOSITIONS

Compositions of dolomite grains on a fracture surface were obtained on the electron microprobe for only one sample, CS4. Other thin sections, in which the fracture surface rim was not destroyed by the polishing process, did not yield good totals (51.1% for an ideal dolomite) because the dolomite grains were either too small or were intermixed with clay. Table VII-1 shows the results for sample CS4. The difference between the compositions of the dolomite grains on the fracture surface and those in the bulk rock is insignificant. They are also very similar to those of other Culebra Dolomite samples (Sewards, Williams, and Keil, 1991). Clearly, the compositions of the dolomite grains in this sample were not affected by fluid moving in the fractures.

Table VII-1. Dolomite Compositions, Sample CS4

#	Oxides				Bulk		Cations		
	CaO	FeO	MgO	Total	Ca	Fe	Mg	Total	
1	25.80	0.20	18.44	44.44	1.000	0.006	0.994	2.000	
2	30.89	0.10	20.75	51.74	1.032	0.003	0.965	2.000	
3	28.40	0.20	19.69	48.29	1.015	0.006	0.979	2.000	
4	29.78	0.15	21.53	51.46	0.995	0.004	1.001	2.000	
5	30.34	0.05	20.70	51.09	1.025	0.001	0.973	2.000	
6	28.92	0.00	20.14	49.06	1.016	0.000	0.984	2.000	
7	28.85	0.38	19.28	48.51	1.031	0.011	0.959	2.000	
8	30.99	0.00	20.49	51.48	1.042	0.000	0.958	2.000	
9	29.17	0.00	20.47	49.64	1.012	0.000	0.988	2.000	
10	28.38	0.19	20.62	49.19	0.992	0.005	1.003	2.000	
11	29.44	0.10	19.43	48.97	1.041	0.003	0.956	2.000	
12	29.99	0.00	18.99	48.98	1.063	0.000	0.937	2.000	
					1.022	0.003	0.975	2.000	

#	Oxides				Surface		Cations		
	CaO	FeO	MgO	Total	Ca	Fe	Mg	Total	
1	29.81	0.00	19.87	49.68	1.038	0.000	0.962	2.000	
2	27.41	0.00	18.87	46.28	1.021	0.000	0.979	2.000	
3	28.70	0.00	20.03	48.73	1.015	0.000	0.985	2.000	
4	27.36	0.00	17.52	44.88	1.058	0.000	0.942	2.000	
5	28.80	0.00	20.06	48.86	1.016	0.000	0.984	2.000	
6	29.75	0.00	20.81	50.56	1.013	0.000	0.987	2.000	
7	27.63	0.00	19.84	47.47	1.000	0.000	1.000	2.000	
8	27.54	0.43	19.53	47.50	1.000	0.012	0.987	2.000	
9	28.25	0.52	19.76	48.53	1.006	0.014	0.979	2.000	
10	29.49	0.34	19.68	49.51	1.032	0.009	0.959	2.000	
11	28.03	0.10	17.83	45.96	1.059	0.003	0.938	2.000	
12	30.20	0.15	20.89	51.24	1.017	0.004	0.979	2.000	
					1.023	0.004	0.973	2.000	

VIII. DISCUSSION

Two main conclusions can be drawn from the information presented above: (1) horizontal water-bearing fractures in dolomite and calcite rock tend to occur in zones where clay and quartz are concentrated, particularly along clay seams, and (2) secondary minerals, primarily gypsum and some calcite, are precipitated from solution onto the fracture surfaces. The clay modes in fracture surface scrapings in dolomite rock range from about 1% to 43%, with an average of 18%, whereas clay modes in the bulk rock in these samples range from less than 1% to 7%, with an average of 4.6%. Similarly, for the two limestone samples, the clay mode is much greater on the fracture surface than in the bulk rock (37% vs. 2%). Secondary gypsum is an important constituent of the fracture surface mineralogy in these samples. Secondary calcite is present in only one sample from a deep core and all five shallow cores.

Dolomite compositions on the fracture surfaces are no different from those in the bulk rock, indicating that aqueous alteration of dolomite did not occur to any significant extent (with the possible exception of some dissolution and recrystallization).

Where present, calcite in these samples, both in the bulk rock and fracture surfaces, is a product of recrystallization from dolomite caused by aqueous alteration, usually near surface.

It should be mentioned that the surfaces examined in this study are all due to horizontal fractures; no surfaces of vertical or high-angle fractures that were clearly identifiable as water-bearing were discovered in the cores examined. It is probable that the vertical and near-vertical fracture surfaces are not as clay-rich as the horizontal ones, since accumulations of clay occur along horizontal planes due to sedimentation.

The implications of these results for the WIPP repository are obvious: since the cation exchange capacity of clay minerals is so much higher than that of dolomite, calcite, or gypsum, and the clay minerals are a major component of the fracture surface mineralogy, the sorption of radionuclides due to the clay will far outweigh that of the other minerals. This fact should be taken into account in any study of the transport of radionuclides through the Culebra Dolomite.

IX. REFERENCES

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APPENDIX A: ANALYTICAL PROCEDURES

X-Ray Diffraction Analysis

Small portions of the ground and sieved whole rock and fracture surface powders were placed in Plexiglas containers (2.5 x 2.5 x 0.4 cm), which have a 1 mm deep hollowed-out compartment. The surface of the powder was then scraped off level with the top surface of the Plexiglas container. The container was placed in the sample holder of a Scintag PAD-V automated diffractometer and analyzed from 2° 2-θ to 60° 2-θ at a scanning rate of 3° per minute using a 0.03° chopper increment.

X-Ray Fluorescence Analysis

Whole rock samples were ground with a mortar and pestle and then passed through a 100 mesh sieve. Fused glass disks were prepared according to standard procedures (Norrish and Chappell, 1967) and analyzed on a Rigaku 3064M x-ray fluorescence spectrometer for 10 elements: SiO₂, Al₂O₃, TiO₂, Fe₂O₃, MnO, MgO, CaO, Na₂O, K₂O, and P₂O₅. Four standards were used: (1) NBS-88b (National Bureau of Standards - Dolomitic Limestone); (2) BCS-CRM-393 (British Chemical Standard - Chemical Reference Material); (3) Dol-1 (Echantillon-type de Calcite), and (4) AM-PAD44 (Amostra Pedrao 44).

Atomic Absorption Spectroscopy

Fracture surface samples were ground and sieved, dissolved in hydrofluoric and perchloric acid mixture, and analyzed on a Perkin-Elmer 303 atomic absorption spectrophotometer for seven elements: SiO₂, Al₂O₃, Fe₂O₃, MgO, CaO, Na₂O, and K₂O (McLaughlin, 1967).

Electron Microprobe Elemental Analysis

Polished thin sections of the samples were prepared using no water and maintaining a temperature below 60°C. The sections were coated with carbon in a vacuum evaporator. Mineral grains were analyzed with a JEOL 733 electron microprobe using an acceleration potential of 15KV, a beam current of 2 nanoamperes, a beam diameter of 1.5 microns, for a period of 80 seconds per analysis. Analyses were corrected according to standard Bence-Albee procedures.

Appendix A References

McLaughlin, R.J.W. 1967. "Atomic Absorption Spectroscopy." *Physical Methods in Determinative Mineralogy*, J. Zussman, ed. Academic Press, 514p.

Norrish, K., and B.W. Chappell. 1967. "X-ray Fluorescence Spectrography." *Physical Methods in Determinative Mineralogy*, J. Zussman, ed. Academic Press, 514p.

APPENDIX B: MODAL MINERALOGICAL CALCULATIONS

Modes for the minerals identified by x-ray diffraction (XRD) were determined from the compositional data obtained by x-ray fluorescence (XRF) and atomic absorption (AA). Based on the compositions of the individual minerals, a particular element, when present in only one mineral, was used to determine the mode of that mineral. For example, the only phase containing aluminum in these samples is clay, and electron microprobe analyses of clay samples from the Culebra Dolomite (Sowards, Williams, and Keil, 1991) show that the clay aggregates contain an average of 15% Al_2O_3 ; thus, the weight percent of clay is calculated according to the formula:

$$\text{Clay(wt\%)} = \text{Al}_2\text{O}_3 / 0.15$$

Quartz, since it contains only SiO_2 , is determined by:

$$\text{Quartz(wt\%)} = \text{SiO}_2 - \text{Clay(wt\%)} \times 0.46$$

since the average SiO_2 content of the clay fraction is 46%. The remaining mineral modes are determined using the following formulae:

$$\text{Dolomite(wt\%)} = (\text{MgO} - \text{Clay} \times 0.15) / 0.19$$

$$\text{or: Dolomite(wt\%)} = (\text{CaO} - \text{Gypsum} \times 0.336) / 0.304$$

$$\text{Gypsum(wt\%)} = \text{SO}_3 / 0.465$$

$$\text{or: Gypsum(wt\%)} = (\text{CaO} - \text{Dolomite(wt\%)} \times 0.304) / 0.326$$

$$\text{Calcite(wt\%)} = (\text{CaO} - \text{Dolomite} \times 0.304) / 0.56$$

Appendix B Reference

Sowards, T., M. Williams, and K. Keil. 1991. *Mineralogy of the Culebra Dolomite Member of the Rustler Formation*. SAND90-7008. Albuquerque, NM: Sandia National Laboratories.

APPENDIX C: DATA TABLES

Tables C-1 and C-2 are the raw data from the bulk rock and fracture surface compositional analyses. Tables V-1 and V-2 are derived from these by converting the Na₂O data to NaCl, subtracting this, and normalizing. Similarly, Tables C-3 and C-4 are the results of the mineral mode calculations from Tables C-1 and C-2. Tables VI-2 and VI-3 are derived from these by removing the halite modes and normalizing to 100%.

Table C-1. Bulk Rock Compositions

Sample ID	Well	Depth	SiO ₂	TiO ₂	Al ₂ O ₃	Fe ₂ O ₃	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	SO ₃	Total
CS1B	WIPP-12	838.60	1.27	0.02	0.24	0.34	0.03	20.22	29.50	0.07	0.07	0.04	0.16	51.94
CS2B	WIPP-13	712.30	4.87	0.08	1.15	0.60	0.02	19.50	27.40	0.04	0.37	0.04	0.07	54.13
CS3B	WIPP-13	705.30	4.14	0.06	0.98	0.47	0.03	19.27	27.80	0.06	0.31	0.04	0.07	53.23
CS4B	WIPP-13	714.00	5.56	0.05	1.14	0.56	0.02	19.30	27.60	0.05	0.37	0.04	0.04	54.72
CS5B	WIPP-26	187.50	2.82	0.04	0.65	0.37	0.04	19.65	27.80	0.07	0.29	0.04	0.05	51.81
CS6B	WIPP-27	305.00	1.95	0.03	0.41	0.29	0.02	20.05	27.90	0.25	0.24	0.05	0.09	51.28
CS7B	WIPP-28	447.50	24.02	0.29	5.51	0.78	0.03	19.02	18.89	1.29	1.19	0.03	0.10	71.15
CS8B	WIPP-29	27.00	3.73	0.06	0.84	0.44	0.02	19.80	27.50	0.22	0.34	0.05	0.06	53.05
CS9B	WIPP-30	633.50	14.27	0.17	2.81	0.09	0.01	19.15	26.14	1.75	1.12	0.13	0.19	65.83
CS10B	WIPP-30	639.00	2.70	0.04	0.55	0.34	0.02	20.40	29.90	0.05	0.18	0.10	0.05	54.32
CS11B	WIPP-30	635.00	1.35	0.02	0.33	0.30	0.02	20.05	29.80	0.30	0.13	0.10	0.07	52.46
CS12B	WIPP-32	57.00	0.35	0.01	0.10	0.08	0.01	11.40	43.60	0.05	0.02	0.18	0.12	55.92
CS13B	WIPP-32	91.10	53.36	0.74	12.96	4.32	0.01	16.15	0.95	0.05	2.32	0.21	0.03	91.10
CS14B	WIPP-32	55.00	2.20	0.02	0.38	0.15	0.02	0.84	54.40	0.04	0.13	0.25	0.13	58.56
CS15B	WIPP-32	56.00	2.95	0.03	0.50	0.22	0.02	0.84	52.60	0.04	0.17	0.25	0.12	57.73
CS16B	WIPP-32	62.00	1.55	0.02	0.32	0.14	0.01	17.40	32.50	0.05	0.12	0.12	0.07	52.30
CS17B	WIPP-33	570.00	1.77	0.02	0.24	0.10	0.00	0.35	32.20	0.10	0.04	N.A.	45.97	80.79
CS18B	WIPP-34	836.00	2.45	0.04	0.50	0.34	0.02	20.20	27.80	0.06	0.18	0.10	0.09	51.77

Table C-2. Fracture Surface Compositions

Sample ID	Well	Depth	SiO ₂	Al ₂ O ₃	Fe ₂ O ₃	MgO	CaO	Na ₂ O	K ₂ O	SO ₃	Total
CS1S	WIPP-12	838.60	*	2.80	0.40	5.60	12.80	34.00	0.32	*	55.52
CS2S	WIPP-13	712.30	15.90	4.40	1.28	17.08	20.60	1.68	1.19	*	60.85
CS3S	WIPP-13	705.30	*	2.10	0.56	14.04	18.80	11.60	0.82	*	47.36
CS4S	WIPP-13	714.00	15.60	4.16	1.50	16.50	18.40	5.16	0.12	1.34	61.28
CS5S	WIPP-26	187.50	9.58	1.30	0.44	17.36	24.00	4.66	0.57	*	57.47
CS6S	WIPP-27	305.00	3.46	0.60	0.26	18.90	26.40	3.92	0.29	*	53.57
CS7S	WIPP-28	447.50	18.90	4.66	1.28	17.46	18.60	3.08	0.87	0.03	63.60
CS8S	WIPP-29	27.00	7.10	1.16	0.94	6.48	15.60	19.30	0.79	*	50.43
CS9S	WIPP-30	633.50	23.30	5.76	1.56	15.28	15.40	3.24	1.83	*	64.81
CS10S	WIPP-30	639.00	38.20	7.88	2.00	11.48	10.24	1.58	2.09	*	71.47
CS11S	WIPP-30	635.00	4.49	1.30	0.36	10.78	16.60	20.28	0.38	2.08	55.91
CS12S	WIPP-32	57.00	0.96	0.14	0.12	12.94	39.60	1.26	0.06	0.02	54.98
CS13S	WIPP-32	91.10	45.50	11.06	3.12	11.30	11.16	0.34	1.56	0.29	81.21
CS14S	WIPP-32	55.00	28.40	6.76	2.06	9.64	23.60	0.96	1.80	0.01	71.17
CS15S	WIPP-32	56.00	25.40	5.90	1.94	7.90	28.40	0.70	1.33	0.01	69.64
CS16S	WIPP-32	62.00	4.48	0.70	0.32	16.60	33.40	0.98	0.24	0.60	57.00
CS17S	WIPP-33	570.00	28.30	5.74	2.12	4.64	17.40	0.72	1.29	20.80	78.89
CS18S	WIPP-34	836.00	2.64	0.68	1.12	4.36	7.64	38.90	0.18	*	54.40

* = Insufficient Sample

Table C-3. Bulk Rock Mineral Modes (Unnormalized)

Sample ID	Dolomite	Calcite	Gypsum	Halite	Clay	Quartz	Total
CS1B	97.04				1.60	0.53	99.17
CS2B	90.13				7.67	1.34	99.14
CS3B	91.45				6.53	1.13	99.12
CS4B	90.79				7.60	2.06	100.45
CS5B	91.45				4.33	0.83	96.61
CS6B	91.78				2.73	0.69	95.20
CS7B	62.14				36.73	7.12	105.99
CS8B	90.46				5.60	1.15	97.21
CS9B	85.99			0.93	18.73	5.65	111.30
CS10B	98.36				3.67	1.01	103.04
CS11B	98.03				2.20	0.34	100.56
CS12B	59.47	39.56	0.26		0.67	0.04	100.00
CS13B	3.13				86.40	13.62	103.14
CS14B		97.14			2.53	1.03	100.71
CS15B		93.93			3.33	1.42	98.68
CS16B	89.89	7.40			2.13	0.57	100.00
CS17B			98.86		1.60	1.03	101.49
CS18B	91.45				3.33	0.92	95.70

Table C-4. Fracture Surface Mineral Modes (Unnormalized)

Sample ID	Dolomite	Calcite	Gypsum	Halite	Clay	Quartz	Total
CS1S	14.74		25.52	64.09	18.67	0.00	123.01
CS2S	66.74		0.96	3.17	29.33	13.88	114.07
CS3S	62.84		0.00	21.87	14.00	0.00	98.71
CS4S	64.95		2.88	9.73	27.73	13.69	118.98
CS5S	84.53		0.00	8.78	8.67	8.98	110.96
CS6S	96.32			7.39	4.00	3.18	110.89
CS7S	67.37			5.81	31.07	16.76	121.00
CS8S	28.00		21.74	36.38	7.73	6.57	100.42
CS9S	50.11		0.52	6.11	38.40	20.65	115.78
CS10S	33.68		0.00	2.98	52.53	34.58	123.77
CS11S	49.89	2.56	4.47	38.23	8.67	3.89	107.71
CS12S	67.37	34.14		2.38	0.93	0.90	105.72
CS13S		19.93		0.64	73.73	40.41	134.72
CS14S		42.14		1.81	45.07	25.29	114.31
CS15S		50.71		1.32	39.33	22.69	114.05
CS16S	83.68	14.21		1.85	4.67	4.16	108.57
CS17S			44.73	1.36	38.27	25.66	110.02
CS18S	25.13		0.00	73.33	4.53	2.33	105.32

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