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Dames & Moore

FINAL REPORT

Phase II Investigation
Underground Storage Tank Area

CARRIER CORPORATION
Syracuse, New York

January 16, 1987

Job No. 14750-002

Dames & Moore



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BUREAU OF
HAZARDOUS WASTE TECHNOLOGY
DIVISION OF SOLID AND
HAZARDOUS WASTE

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1.0 Introduction

This report details the Phase II investigation in which Dames & Moore was contracted by the Carrier Corporation to install two additional monitoring wells, and perform sampling and analysis on groundwater samples from seven wells, at Carrier's manufacturing facility off Thompson Road in Syracuse, New York. This Phase II investigation follows a Phase I investigation which was performed at the site by Dames & Moore in late 1985 and early 1986.

The site location is shown on Figure 1. The area of primary concern, and which is the focus of this investigation, is an underground storage tank area which includes four tanks; two 20,000 gallon cement tanks and two 8,000 gallon steel tanks. The tanks contain a variety of different materials including acids and caustics, waste coolants and waste oil. In the past, trichloroethene still bottoms were also stored in the tanks. Closure of the tanks is currently being planned by Carrier.

This report includes a description of the underground tanks and their history, a description of the work performed, and an analysis of the data. Recommendations for additional studies are also provided.

1.1 Site Location

The location of the Carrier Corporation's facility in Syracuse, New York is shown on Figure 1. The facility is in the northeast portion of Syracuse, approximately one mile south of the New York State Thruway. The site is bounded by Sanders Creek to the north, Thompson Road to the west, Kinne Street to the east and residential roads and fields to the south.

The site slopes slightly to the north and northwest towards Sanders Creek. The elevation difference between the north and south

boundaries is about 10 to 20 feet, based on the USGS topographic map of the site.

This study investigated a group of below-grade tanks located west of the center of the site. The location of the tanks is shown in Figure 2. The land surrounding the tanks, like the rest of the site, has a slight slope to the north. The elevation in the area of the tanks is about 405 feet above mean sea level (MSL).

1.2 Site Background

This section provides a summary of a history of the four tanks which was provided by Carrier Corporation to Dames & Moore.

The underground tank area includes two 20,000 gallon underground cement tanks. One contains acids and caustics, with some chromic acid, and the other contains water soluble coolants (cutting oils). The other two tanks are 8,000 gallon underground steel tanks. One contains waste oil and the other previously contained trichloroethene still bottoms. It is now empty. (Note: Trichloroethene can also be spelled trichloroethylene. Carrier uses the latter spelling while Galson Technical Services, the laboratory for this project, uses the former. Since the laboratory results used the former spelling, that spelling is used in this report for consistency.)

The two concrete tanks were installed around 1945 by General Electric (G.E.), former owner of the site. They were used primarily as central settling and neutralization basins for their storm, process, and sanitary wastewater. The wastewater was then discharged into the sewer system.

In the 1950's, Carrier purchased the property from G.E. and use of the tanks for wastewater was discontinued. They remained unused and in 1969 they were filled in with stone. In 1978, Carrier rebuilt these concrete tanks for use in storing spent process wastes prior to their

disposal. Wastes temporarily stored at this time were: oils, acids, caustics, trichloroethene still bottoms and coolants.

In 1980, Carrier installed the two steel 8,000 gallon storage tanks. Addition of these tanks allowed segregation of the various process wastes. One steel tank was reserved for waste oils and the other for waste trichloroethene still bottoms. One concrete tank was used for acids and alkalis, and the other for waste coolants.

Around 1982, the use of the one steel tank to store waste trichloroethene still bottoms was discontinued. The material was completely pumped out. Since that time, still bottom waste has been put into drums for disposal offsite. The tank remains in place.

1.3 Previous Investigation

On March 24, 1986, Dames & Moore produced a final report which described the Phase I investigation at this Carrier Corporation facility. The Phase I study included the installation of five monitoring wells; one well upgradient and four wells downgradient of the waste tanks being investigated. These wells were sampled and analyzed for pH, total organic carbon (TOC), total organic halogens (TOX), specific conductance, volatile organic compounds (VOCs), oil and grease, PCBs and selected metals. Five subsurface soil samples were also collected and analyzed for chromium, trichloroethene and oil and grease. Groundwater contamination and flow directions were assessed.

The results of the Phase I investigation indicated that VOCs were present in the groundwater samples, and were probably attributable to the tank area. Wells MW-2 and MW-4 contained the highest measured concentrations of VOCs. These are the two wells closest to the underground tanks. Contamination of the groundwater with metals was not found to be significant. Analysis of the soil samples showed detectable amounts of trichloroethene at MW-4 and oil and grease in soil samples from borings MW-2, 3, and 4.

Based on these findings, this Phase II investigation was initiated to further define the extent of the groundwater contamination.

1.4 Phase II Investigation

The scope of work for this Phase II investigation consisted of three tasks. Task 1 consisted of the installation of two shallow overburden monitoring wells, located downgradient of the contaminated groundwater found in the Phase I investigation. In Task 2, the wells were developed for one hour each, surveyed, and sampled for laboratory analyses, consisting of: pH, specific conductance, TOX, TOC, and VOC(s). Task 3 was report preparation. The field work for this Phase II investigation began on October 13, 1986.

1.5 Organization of this Report

This report consists of five sections. This introduction (Section 1.0) summarizes the site and history of investigations. Section 2.0 describes the procedures employed in performing the tasks for this Phase II investigation. Section 3.0 details the data analysis results in terms of subsurface stratigraphy, groundwater flow and chemical analyses, and provides a concise summary of the findings of this Phase II investigation within the context of the Phase I results. Section 4.0 presents our conclusions and recommendations regarding the site conditions. Appendix A contains the laboratory analytical results and Appendix B contains the boring logs for all seven on-site wells.

2.0 Site Investigation Procedures

The scope of this investigation included the installation of two additional monitoring wells and sampling of these two wells and the five existing wells installed during the Phase I study in December, 1985. This section describes the procedures used in performing these activities.

2.1 Soil Borings and Monitoring Well Installations

The location of the monitoring wells in this Phase II investigation were selected after discussions with Ed Besaw of Carrier Corporation. The locations were to be far enough downgradient of the waste tanks previously mentioned, so that the wells were beyond the contaminant plume identified during the Phase I work. Figure 2 shows the location of the two wells installed (MW-5 and MW-6) in relation to the five Phase I wells.

The soil borings were performed with a truck mounted drilling rig, which utilized continuous flight augers to advance the bore holes. No drilling fluid is used in this process. Representative soil samples were obtained using a split-spoon sampler in general accordance with ASTM Specification D-1586. Using this method, a 2-inch nominal O.D. split-spoon sampler is driven into the soil by a 140-pound hammer with a free fall of 30 inches. Representative portions of the soil samples were placed in jars and retained by Dames & Moore. Boring logs were prepared by Dames & Moore, based on visual observations of the samples.

During drilling, the samples were scanned with an HNU photo-ionization meter to detect releases of volatile organic compounds (VOCs). The samples were again scanned after being placed in jars and brought to room temperature (head space test). No significant releases of VOCs were measured in either test.

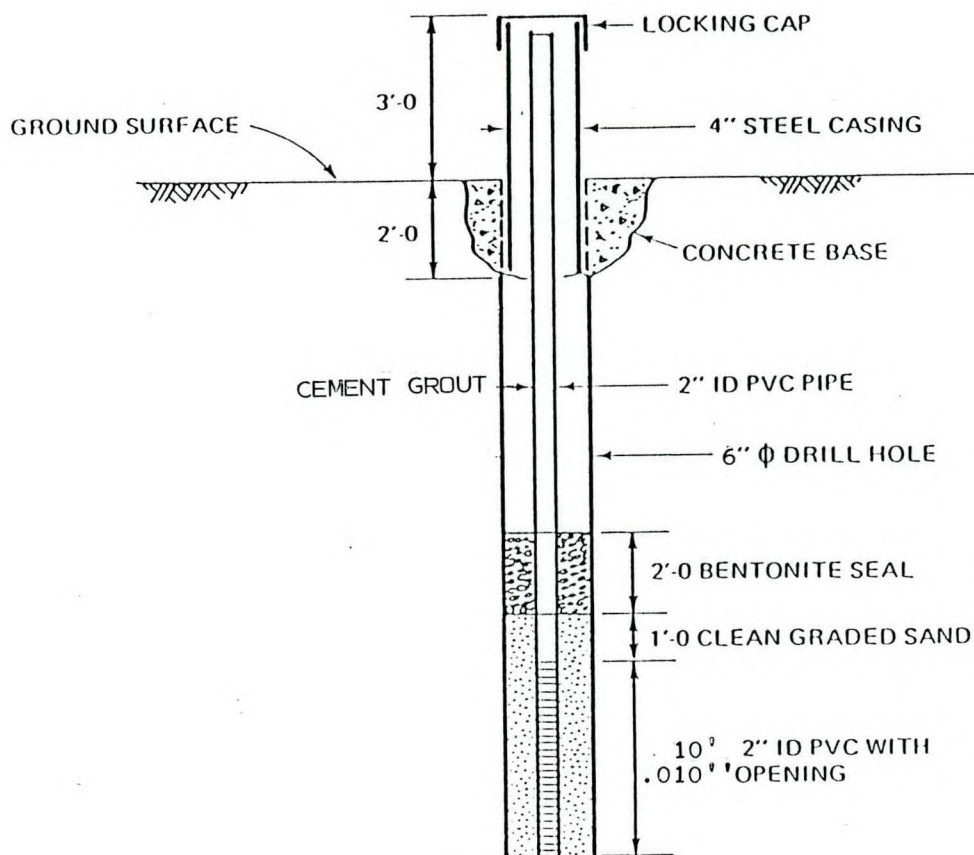
The soils were sampled at intervals of at least every five feet, to a final depth of 15.0 feet. The wells were then installed. The well materials consist of 2-inch diameter PVC with .010-inch slotted screens. In both wells 10-foot screens were used. A typical well schematic is presented as Figure 3. Where practicable, the well screens were installed to intersect the water table. This would allow those compounds which are lighter than water to enter the well. The two wells installed as part of the Phase II work (MW-5 and MW-6) monitor the upper part of the overburden aquifer.

After the wells were installed, 4-inch diameter protective casings were grouted in place over the well pipe. The following day, the wells were developed by evacuation for one hour each to remove fine sediments, and provide better quality groundwater samples. The wells were installed on October 13, 1986 and developed on October 14, 1986. The new wells were surveyed in relation to the existing wells and tied into an on-site benchmark on October 23, 1986.

2.2 Sampling and Analysis

On October 17, 1986, samples from all seven wells installed under the Phase I and II work were collected and sent to a laboratory for analysis of priority pollutant volatile organic compounds (VOCs), TOX, TOC, pH and specific conductance utilizing EPA-approved protocols and procedures. Additional samples from each well were field tested for pH, specific conductance and temperature.

Prior to collecting the sample to be analyzed, two to three well volumes of water were removed from each well. This process provides a more representative sample of the water in the aquifer. A stainless steel bailer was used for evacuation and sampling of groundwater. Between wells, the bailer was rinsed with successive washes of hexane, methanol and distilled, deionized water to limit possible cross-contamination. The samples were contained in laboratory-prepared bottles and packed in an ice-filled cooler. The seven groundwater



TYPICAL OBSERVATION WELL

samples were collected and delivered to the laboratory, Galson Technical Services in East Syracuse, on the same day.

3.0 Data Analysis Results

This section details the results of the data analysis which was conducted as part of the Phase II investigation. These results incorporate pertinent information from the Phase I investigation as well. This section presents the data analysis results in three subsections; soil stratigraphy, groundwater flow and groundwater quality, as investigated within the scope of work previously outlined.

3.1 Soil Stratigraphy

The soils in the vicinity of the underground waste tanks consist of medium to coarse textured lake sediments which are dominated by fine sand and silt, are relatively gravel-free and moderately permeable (see boring logs for MW-2, 3S, 3D and 4 in Appendix B). Downgradient of the tanks, in the vicinity of MW-6, the soils are predominantly glacial till deposits. The soil samples taken from MW-6 indicate the soils there are stiff, slowly permeable and contain 10 to 20% gravel. Below a depth of about 8.5 feet, the soil becomes increasingly moist, sandy and relatively gravel-free. Farther downgradient, at MW-5, the soils consist of fill to at least a depth of 7.0 feet. Underlying the fill are organic-rich lake sediments or possibly alluvial deposits associated with Sanders Creek. Lake sediments were encountered at the bottom of MW-5, a soft silty-clay soil.

Based on the limited soil characterization provided by the few borings performed, it would appear that the underground tanks are located in relatively coarse, moderately permeable sediments. Downgradient groundwater flow toward MW-6 may be impeded by the dense, slowly permeable soils encountered there. However, due to the presence of foundations, utility conduits, etc. at the site, defining the path of contaminant movement is difficult. Impermeable or slowly-permeable materials can allow increased contaminant movement if they are breeched by man-made subsurface drainage features. As a result, predictions of

present and future migration pathways based upon soil stratigraphy alone may be misleading.

3.2 Groundwater Flow

Table 1 contains information regarding monitoring well construction and water levels taken at the time of sampling on October 17, 1986. The overburden units which occur within the screened portion of the wells are listed with accompanying elevation information. Those units described on the boring logs as having less than 20 percent silt and clay are marked with an asterisk. These are the zones which, based on the boring log descriptions, would be the most permeable zones in the boring. If contaminants are present in the groundwater, they would probably be transmitted through these zones most easily.

The northward groundwater flow direction is confirmed by the water level information in Table 1. The elevation of the groundwater surface in the overburden aquifer, as monitored by the wells, is highest at MW-1. The groundwater surface is slightly lower in elevation at MW-2 and MW-4, and continues to lower toward MW-3, MW-6 and MW-5. The water levels from MW-3D and MW-3S indicate a slight downward gradient is present at that location.

3.3 Groundwater Quality

3.3.1 Volatile Organic Compounds

Tests to detect volatile organic compounds (VOCs) were performed on a sample from each of the seven wells installed during the Phase I and II work. The results are presented in Table 2. Only those compounds which were detected are listed. The analytical data are presented in their entirety in Appendix A. The concentrations are compared to applicable New York State groundwater standards or guidance values. Also listed are the results from a lab blank which is a quality assurance measure to determine if bottle preparation or shipment and

TABLE 1

Monitoring Well Details

<u>WELL NUMBER</u>	<u>UNIT SCREENED</u>	<u>SCREEN ELEVATION</u>	<u>WATER ELEVATION (10/17/86)</u>
MW-1	Silty Sand Lacustrine	405.2 - 395.2	402.4
MW-2	Silty Lacustrine	399.5 - 395.5	399.3
	Sandy Lacustrine*	395.5 - 391.5	
	Silty Lacustrine	391.5 - 389.5	
MW-3S	Silty Lacustrine	399.6 - 397.6	398.1
	Sandy Lacustrine*	397.6 - 393.6	
	Silty Lacustrine	393.6 - 389.6	
MW-3D	Silty-Sand Lacustrine	381.5 - 379.0	397.5
	Sandy Till*	379.0 - 376.5	
MW-4	Gravel Fill*	399.5 - 391.5	400.8
	Silty Lacustrine	391.5 - 389.5	
MW-5	Silty Fill	390.2 - 388.2	394.2
	Silty Alluvium	388.2 - 380.2	
MW-6	Silty Till	399.4 - 389.4	395.7

*Permeable Zones

TABLE 2

Volatile Organic Compounds Detected
in Groundwater Samples in (ug/l)

	SAMPLE ID:	MW-1	MW-2	MW-3D	MW-3S	MW-4	MW-5	MW-6	LAB BLANK	LOQ	NYSDEC STANDARD OR CRITERIA
BENZENE		ND	< 4.4	ND	ND	< 4.4	ND	ND	ND	4.4	
CHLOROFORM		ND	ND	ND	ND	2	ND	ND	ND	1.6	100.0A
1,1-DICHLOROETHANE		ND	380-	ND	240+	500-	< 4.7	ND	ND	4.7	50.0B
1,2-DICHLOROETHANE		ND	40-	ND	11+	26+	ND	ND	ND	2.8	1.0B
1,1-DICHLOROETHENE		ND	880+	ND	70+	290+	ND	ND	ND	2.8	0.9B
TRANS-1,2-DICHLOROETHENE		ND	610-	ND-	60-	60-	ND	ND	ND	1.6	50.0B
ETHYLBENZENE		ND	ND	ND	ND	< 7.2	ND	ND	ND	7.2	
METHYLENE CHLORIDE		3	19	4	7	15	7	3	8	2.8	10.0B
TETRACHLOROETHENE		ND	ND	ND	ND	< 4.1	ND	ND	ND	4.1	
1,1,1-TRICHLOROETHANE		ND	620+	ND	< 3.8	250-	ND	ND	ND	3.8	50.0B
1,1,2-TRICHLOROETHANE		ND	180+	ND	< 5.0	13-	ND	ND	ND	5.0	0.5B
TRICHLOROETHENE		ND	240+	ND	ND	6-	< 1.9	ND	ND	1.9	10.0A 5.0B
TOLUENE		ND	< 6.0	ND	ND	64	< 6.0	< 6.0	ND	6.0	
VINYL CHLORIDE		ND	6600	ND	77	7000	ND	ND	ND	5.0	5.0A

A - New York State Department of Environmental Conservation, NYCRR 703.5 - Standards for Class 6A groundwaters.

B - New York State Department of Environmental Conservation "Ambient Water Quality Criteria" memo dated August 5, 1983. Guidance Values.

LOQ - Limit of Quantitation

(<) - Less than Limit of Quantitation

ND - Not Detected

(+) - Indicates increase in concentration since Phase I study

(-) - Indicates decrease in concentration since Phase I study

storage have caused contamination. It appears from the results that methylene chloride concentrations detected are the result of contamination from either bottle preparation, shipping or extraction processes which are part of the analytical procedure. Methylene chloride is commonly used by laboratories in these extraction processes. As a result of methylene chloride being detected in the lab blank, it is probably attributable to some form of external contamination previously mentioned.

No other VOCs were detected in MW-1, the upgradient well. As expected, the highest concentrations of VOCs were detected at MW-2 and MW-4, the locations nearest the tanks. At these locations, the applicable standards or guidance values were exceeded for 1,1-dichloroethane, 1,2-dichloroethane, 1,1-dichloroethene, 1,1,1-trichloroethane, 1,1,2-trichloroethane, trichloroethane and vinyl chloride. For these and other detected compounds, it generally appears that the concentrations are less in MW-4, and higher in MW-2, as compared to the results from the Phase I sampling.

Vinyl chloride concentrations were the highest of those detected. This could be attributable to degradation of the PVC well screen and casing by the contaminants in the groundwater. For instance, trichloroethene is listed as having a severe effect on PVC (Cole Parmer, Inc., 1984). The high concentrations of trichloroethene and other associated solvents is probably deteriorating the PVC wells over time.

No VOCs, other than methylene chloride, were detected in MW-3D. MW-3S contained slightly higher levels of several VOCs as compared to the Phase I sampling. Standards or guidance values were exceeded in MW-3S for 1,1-dichloroethane, 1,2-dichloroethane, 1,1-dichloroethene, trans-1,2-dichloroethene and vinyl chloride.

At MW-5, near Sanders Creek, trichloroethene, 1,1-dichloroethane and toluene were reported as below the limit of quantitation. The limit of quantitation is, in essence, the detection limit for the analytical

instrument. It is the smallest amount of a particular compound that can be accurately quantified. Results which are reported to be below the limit of quantitation are estimates, and can be the result of interferences in the analysis. Because these three compounds were reported as below the limit of quantitation, as was toluene at MW-6, there is not conclusive evidence that these compounds are present in wells MW-5 and MW-6.

The only VOC reportedly stored in the waste tanks was trichloroethene. However, other products, primarily various types of dichloroethenes and dichloroethanes were also found in the groundwater samples. In groundwater systems where trichloroethene has been released, it is quite common to also find the dichloro type compounds. They are decomposition products of trichloroethene.

An example of a decomposition chain of a chloroethane compound is: 1,1,1-trichloroethane $\longrightarrow$ 1,1-dichloroethane $\longrightarrow$ chloroethane.

Other chloroethene and chloroethane compounds exhibit similar transformations. These transformations probably account for the wide variety of compounds found in the groundwater. The fact that these compounds were detected indicates that the VOCs being found are due to past leakage or spillage. However, the rates of decomposition are not well established and cannot be used to determine the time a leak or spill occurred.

The following subsections summarize the results of the other analyses performed. Table 3 presents those results.

3.3.2 Total Organic Halogens

Total organic halogens (TOX) were detected in very small amounts (<0.1 mg/l) at MW-1, MW-3D, MW-5 and MW-6. The highest levels were at

TABLE 3

Summary of Other Analytical Results

<u>WELL ID</u>	<u>TOX</u> <u>mg/l</u>	<u>TOC</u> <u>mg/l</u>	<u>SP. COND.</u> <u>umhos/cm</u>	<u>pH</u>
MW-1	< 0.1	20-	1550+	7.0-
MW-2	10+	35+	2740+	6.9
MW-3S	2.6-	11+	1790-	7.2
MW-3D	< 0.01-	8+	2180+	7.4+
MW-4	3.6-	82-	1230+	6.8-
MW-5	0.02	16	2130	6.8
MW-6	< 0.01	6	570	7.8

(+) - Indicates increase since Phase I study

(-) - Indicates decrease since Phase I study

MW-2, MW-4 and MW-3S. These levels were generally slightly less than the Phase I results.

3.3.3 Total Organic Carbon

Total organic carbon (TOC) was measured in each well, and found to be highest at MW-2 and MW-4. Interestingly, the levels for MW-1 and MW-5 were above that of MW-3S. Generally, all values were slightly below those found in the Phase I work.

3.3.4 Specific Conductance

The conductivity of groundwater can serve as an indication of the quantity of dissolved ions present. An increase in the conductivity indicates more (or different valence), ions are entering the groundwater between two monitoring points. As was the case during the Phase I work, the highest conductivity value was at MW-2. MW-3D, MW-5 and MW-3S were also in the upper range. The lowest values were at MW-6 and MW-4.

3.3.5 pH

The pH in all wells varied between 6.8 and 7.8 which is within the normal, expected range.

3.4 Summary of Results

Two additional monitoring wells were installed. MW-5 occurs in a downgradient location within alluvial sediments. MW-6 is also downgradient of the tanks and occurs within glacial till.

All wells were sampled and analyzed for VOCs, TOX, TOC, pH and specific conductance. Wells MW-1 and MW-3D were generally free of VOCs. The highest concentrations of VOCs were detected in MW-2 and MW-4, with lower concentrations at MW-3S.

Trichloroethene, 1,1-dichloroethane and toluene were reported as below the limit of quantitation at MW-5, and toluene was reported as below the limit of quantitation at MW-6. As a result, it cannot be shown conclusively that the compounds were present in the samples from these wells.

Overall, the types of compounds detected in the five existing wells (MW-1, 2, 3S, 3D, 4) were generally similar. In about one-half of the cases where similar compounds were detected, the concentrations were lower than those reported during the Phase I investigation.

TOX, TOC, pH and specific conductance all were in general agreement with the previous sampling results.

4.0 Recommendations

The high concentration of vinyl chloride in the groundwater samples from MW-2, 3S and 4 indicate degradation of the PVC well casing. If monitoring is to continue on a long-term basis, stainless steel wells could be installed. This would lessen the interaction between the contaminants and the well casing materials.

The closure of the tanks will most likely result in removal of wells MW-2 and MW-4, due to their proximity to the tanks. If wells MW-3S and MW-3D are to remain as future monitoring points, it is suggested that these wells be replaced with stainless steel well screens and casings.

APPENDIX A

LABORATORY ANALYTICAL RESULTS



Galson

Technical Services, Inc.

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LABORATORY ANALYSIS REPORT

Client: DAMES & MOORE

Task Number: 86101707

Location: CARRIER

Job Number: G6274

Date Sampled: 17-OCT-1986

TOTAL ORGANIC HALOGENS

Sample ID	Lab ID	MG/L
MW-1	D24394	<0.1
MW-2	D24395	10
MW-3D	D24396	<0.01
MW-3S	D24397	2.6
MW-4	D24398	3.6
MW-5	D24399	0.02
MW-6	D24400	<0.01

($<$) - Less Than LOQ
($>$) - Greater Than
NA - Not Applicable
ND - Not detectable
NS - Not specified
MG - Milligrams
 $\mu\text{g/l}$ - Micrograms Per Liter
LOQ - Limit of Quantitation

Method(s): TOX ANALYZER

Footnotes:

Submitted by: *E.H.*

Approved by: *E.H. Galson*

Date: 4-NOV-1986



Galson

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LABORATORY ANALYSIS REPORT

Client: DAMES & MOORE

Task Number: 86101707

Location: CARRIER

Job Number: G6274

Date Sampled: 17-OCT-1986

TOTAL ORGANIC CARBON

Sample ID	Lab ID	MG/L
MW-1	D24387	20
MW-2	D24388	35
MW-3D	D24389	8
MW-3S	D24390	11
MW-4	D24391	82
MW-5	D24392	16
MW-6	D24393	6

($<$) - Less Than LOQ
($>$) - Greater Than
NA - Not Applicable
ND - Not detectable
NS - Not specified
MG - Milligrams
 $\mu\text{g/l}$ - Micrograms Per Liter
LOQ - Limit of Quantitation

Method(s): EPA 600/4-79-020

Footnotes:

Submitted by: *Alex*

Approved by: *E. Galson*

Date: 4-NOV-1986



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LABORATORY ANALYSIS REPORT

Client: DAMES & MOORE

Job Number: G6274

Task Number: 86101707

Location: CARRIER

Date Sampled: 17-OCT-1986

Sample ID	Lab ID	pH ELECTRODE	SPECIFIC CONDUCTANCE μ HMS/CM
MW-1	D24380	7.0	1550
MW-2	D24381	6.9	2740
MW-3D	D24382	7.4	2180
MW-3S	D24383	7.2	1790
MW-4	D24384	6.8	1230
MW-5	D24385	6.8	2130
MW-6	D24386	7.8	570

(<) - Less Than LOQ
(>) - Greater Than
NA - Not Applicable
ND - Not detectable
NS - Not specified
MG - Milligrams
 μ g/l - Micrograms Per Liter
LOQ - Limit of Quantitation

Method(s): EPA 600/4-79-020

Footnotes:

Submitted by: *D. L. Fager*

Approved by: *Eric Holme*

Date: 4-NOV-1986



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LABORATORY ANALYSIS REPORT

Client: DAMES & MOORE
Task Number: 86101707
Location: CARRIER

Job Number: G6274

Date Sampled: 17-OCT-1986

	Lab ID:	D24401	D24402	D24403	D24404	D24405	D24406	D24407	IN-HOUSE	
	Client ID:	MW-1	MW-2	MW-3D	MW-3S	MW-4	MW-5	MW-6	BLANK	LOQ
BENZENE	µg/l	ND	<4.4	ND	ND	<4.4	ND	ND	ND	4.4
BROMOMETHANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
BROMODICHLOROMETHANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	2.2
BROMOFORM	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	4.7
CARBON TETRACHLORIDE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	2.8
CHLOROBENZENE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	6.0
CHLOROETHANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
2-CHLOROETHYL VINYL ETHER	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	10.0
CHLOROFORM	µg/l	ND	ND	ND	ND	2	ND	ND	ND	1.6
CHLOROMETHANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
DIBROMOCHLOROMETHANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	3.1
1,3-DICHLOROBENZENE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
1,2-DICHLOROBENZENE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
1,4-DICHLOROBENZENE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
1,1-DICHLOROETHANE	µg/l	ND	380	ND	240	500	<4.7	ND	ND	4.7
1,2-DICHLOROETHANE	µg/l	ND	40	ND	11	26	ND	ND	ND	2.8
1,1-DICHLOROETHENE	µg/l	ND	880	ND	70	290	ND	ND	ND	2.8
TRANS-1,2-DICHLOROETHENE	µg/l	ND	610	ND	60	60	ND	ND	ND	1.6
1,2-DICHLOROPROPANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	6.0
CIS-1,3-DICHLOROPROPENE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
TRANS-1,3-DICHLOROPROPENE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
ETHYLBENZENE	µg/l	ND	ND	ND	ND	<7.2	ND	ND	ND	7.2
METHYLENE CHLORIDE	µg/l	3	19	4	7	15	7	3	8	2.8
1,1,2,2-TETRACHLOROETHANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	6.9
TETRACHLOROETHENE	µg/l	ND	ND	ND	ND	<4.1	ND	ND	ND	4.1
1,1,1-TRICHLOROETHANE	µg/l	ND	620	ND	<3.8	250	ND	ND	ND	3.8
1,1,2-TRICHLOROETHANE	µg/l	ND	180	ND	<5.0	13	ND	ND	ND	5.0
TRICHLOROETHENE	µg/l	ND	240	ND	ND	6	<1.9	ND	ND	1.9
TRICHLOROFLUOROMETHANE	µg/l	ND	ND	ND	ND	ND	ND	ND	ND	5.0
TOLUENE	µg/l	ND	<6.0	ND	ND	64	<6.0	<6.0	ND	6.0
VINYL CHLORIDE	µg/l	ND	6600	ND	77	7000	ND	ND	ND	5.0

(<) - Less Than LOQ
(>) - Greater Than
NA - Not Applicable
ND - Not detectable
NS - Not specified
MG - Milligrams
µg/l - Micrograms Per Liter
LOQ - Limit of Quantitation

Method(s): EPA 624

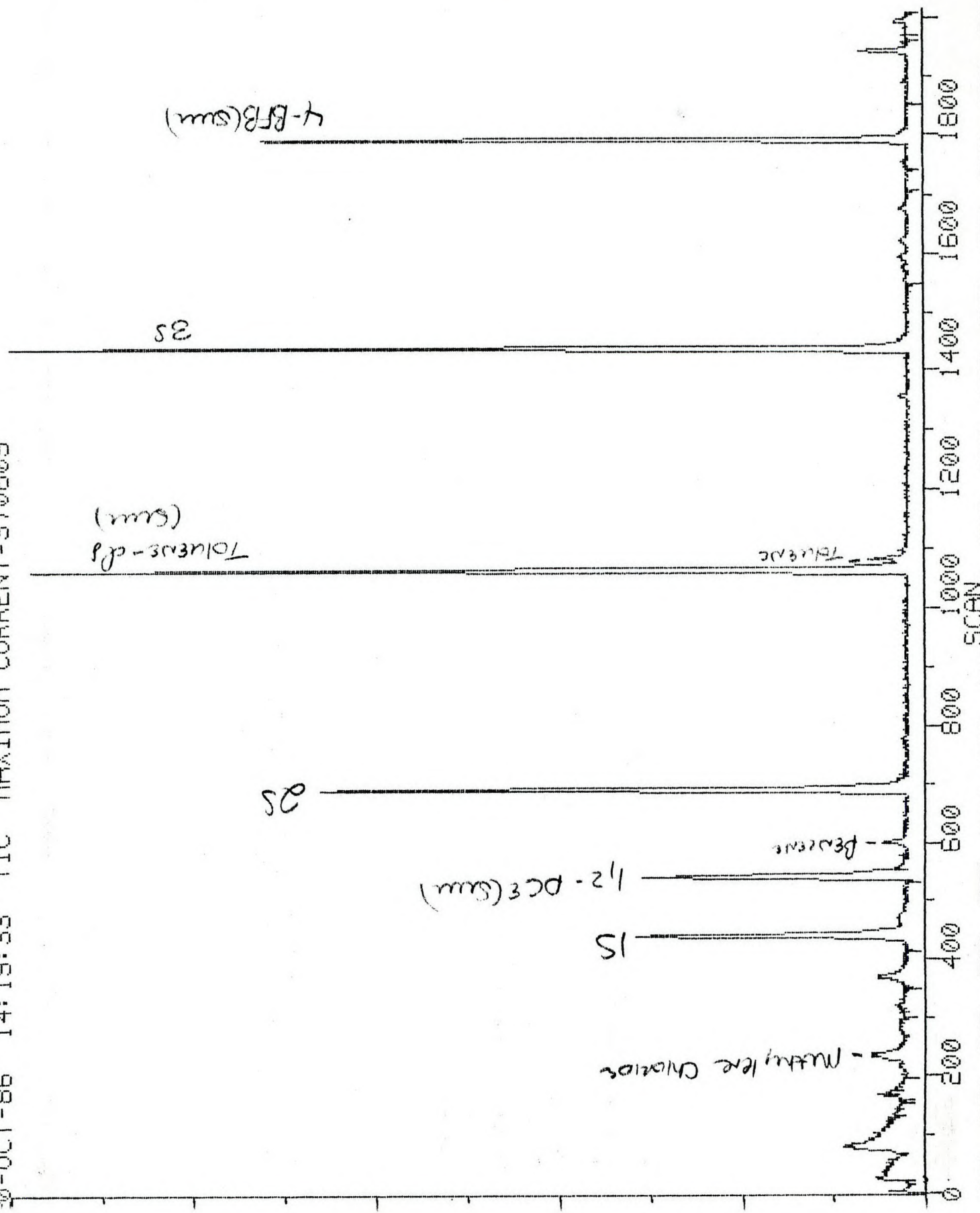
Footnotes:

Submitted by: *J. C. ...*

Approved by: *Eric ...*

Date: 4-NOV-1986

124401 DAMES&MOORE MW-1 +1.0UL INT STD, SURR EMV 2.85
0-OCT-86 14:19:33 TIC MAXIMUM CURRENT=370609



EXTREL Quantitation Report

Name: _____ W.D. _____
Analyst: _____ Date: _____
Reviewer: _____ Date: _____
Comments: _____

Library used: SY:VOASUR
Data file name: SY:D24401
Injection time: 20-OCT-86 14:19:33
Comments:
DAMES&MOORE MW-1 +1.0UL INT STD,SURR EMV 2.85
Dilution factor: 1.00

Library entries as follows:

Standards:

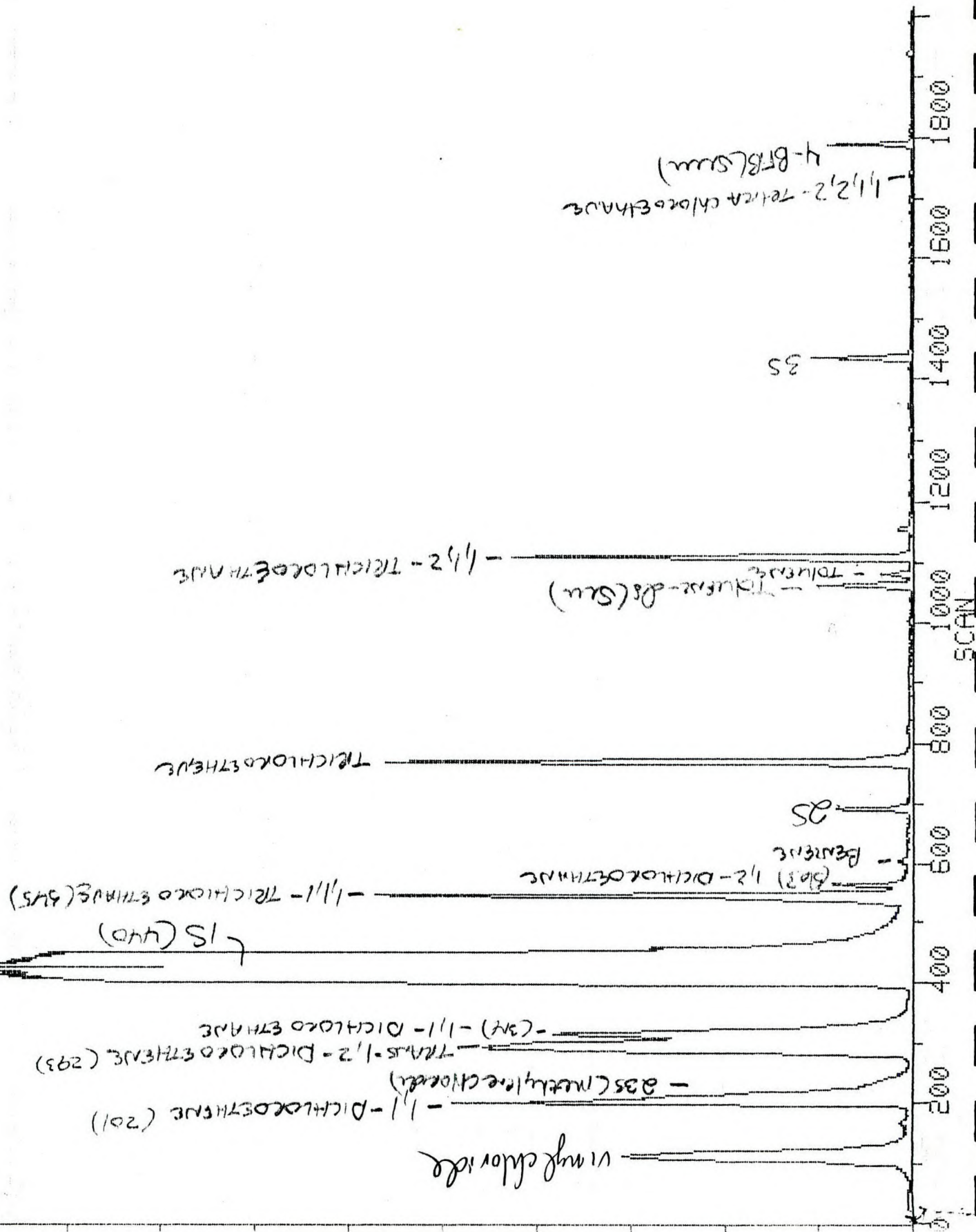
1S BROMOCHLOROMETHANE
2S 1,4-DIFLUOROBENZENE
3S CHLOROBENZENE-D5

Targets:

1T 1,2-DICHLOROETHANE-D4
2T TOLUENEN-D8
3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	5.12	441			STD	0.79	100.0	%
2S	7.43	689			STD	0.73	100.0	%
3S	14.43	1438			STD	0.80	100.0	%
1T	6.07	542	65. / 128.	323136. / 122376.	1	0.77	100.7	%
2T	10.92	1062	98. / 117.	710720. / 579792.	3	0.74	102.0	%
3T	17.72	1790	95. / 117.	301696. / 579792.	3	0.54	95.9	%

24402 5.0ML MW-2 DAMES&MOORE +1.0 UL INT STD, SURR EMU2.85
0-OCT-86 15:00:31 TIC MAXIMUM CURRENT=3664832



EXTREL Quantitation Report

Name: _____ W.O. _____
Analyst: _____ Date: _____
Reviewer: _____ Date: _____
Comments: _____

Library used: SY:VOASUR
Data file name: SY:D24402
Injection time: 20-OCT-86 15:00:31
Comments:
5.0ML MW-2 DAMES&MOORE +1.0 UL INT STD,SURR EMV2.85
Dilution factor: 1.00

Library entries as follows:

Standards:

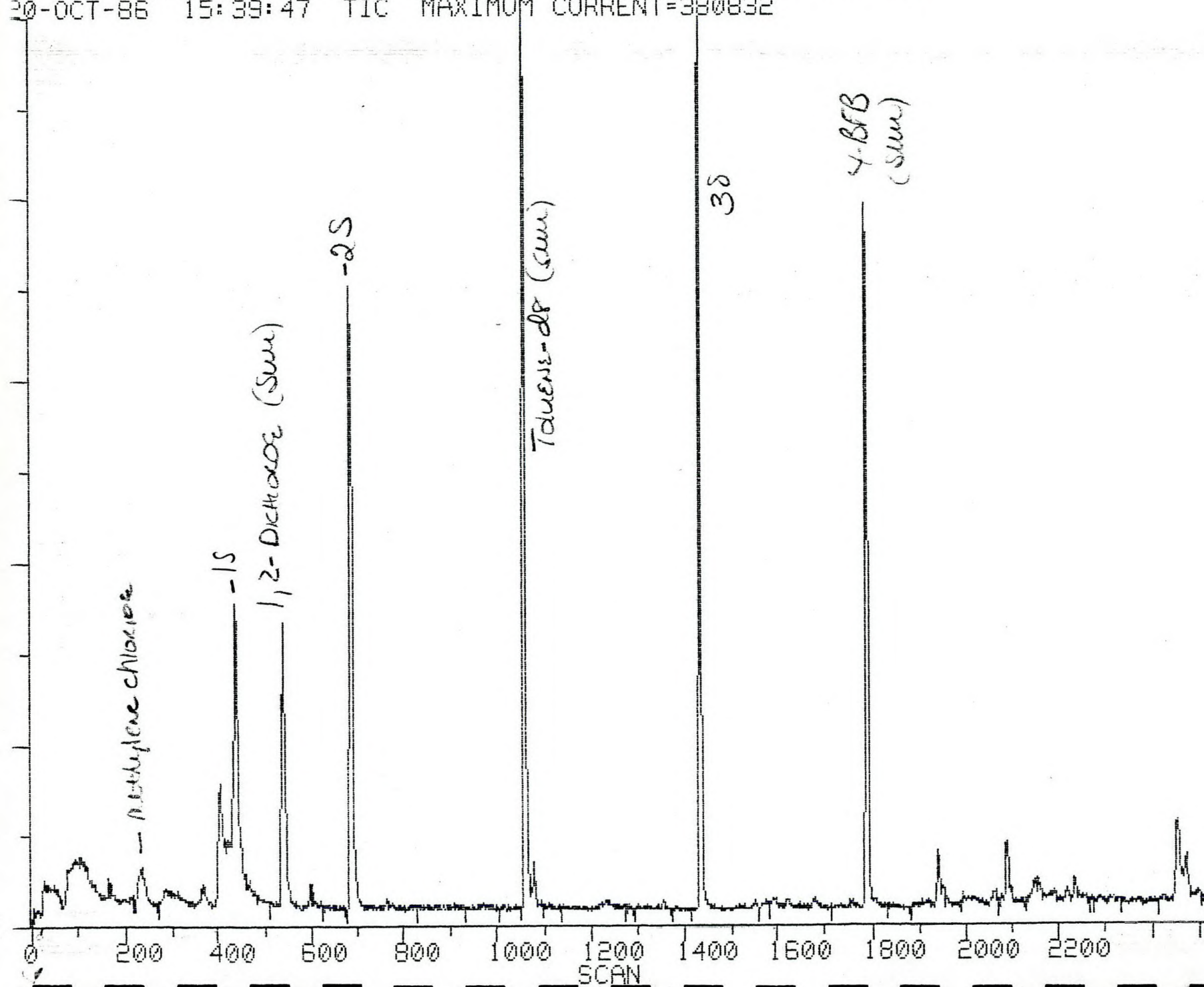
1S BROMOCHLOROMETHANE
2S 1,4-DIFLUOROBENZENE
3S CHLOROBENZENE-D5

Targets:

1T 1,2-DICHLOROETHANE-D4
2T TOLUENEN-D8
3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	5.10	440✓			STD	0.23	100.0	%
2S	7.45	691✓			STD	0.73	100.0	%
3S	14.42	1436✓			STD	0.76	100.0	%
1T			Not Found					
2T	10.92	1062✓	98./ 117.	752728./ 640084.	3	0.74	97.8	%
3T	17.72	1789✓	95./ 117.	359200./ 640084.	3	0.70	103.4	%

024403 5.0ML MW-3D DAMES8MOORE +1.0UL INT STD, SURR EMV2.85
20-OCT-86 15:39:47 TIC MAXIMUM CURRENT=380832



EXTREL Quantitation Report

Name: _____ W.O. _____
Analyst: _____ Date: _____
Reviewer: _____ Date: _____
Comments: _____

Library used: SY:VDASUR
Data file name: SY:D24403
Injection time: 20-OCT-86 15:39:47
Comments:
5.0ML MW-3D DAMES&MOORE +1.0UL INT STD,SURR EMV2.85
Dilution factor: 1.00

Library entries as follows:

Standards:

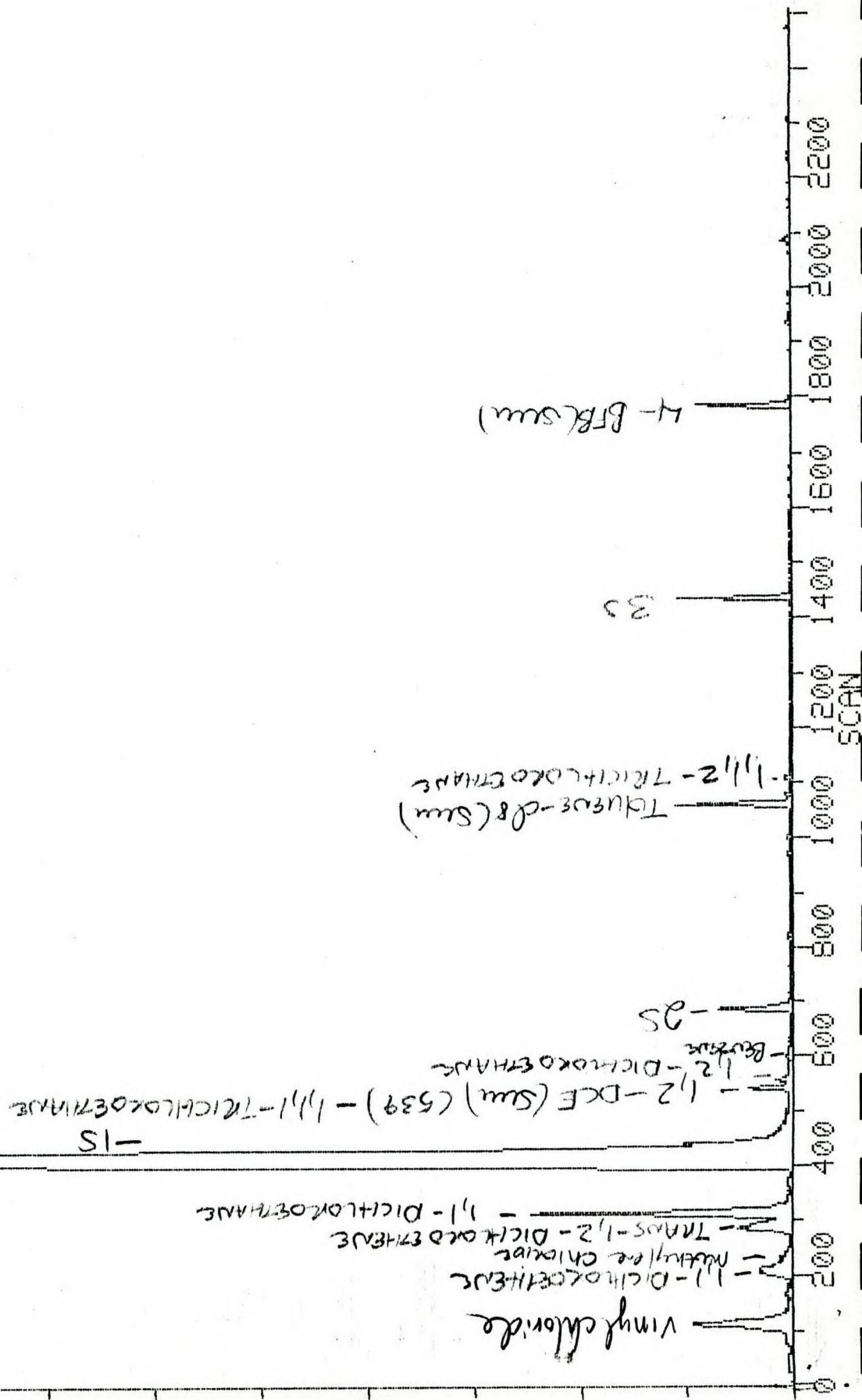
1S BROMOCHLOROMETHANE
2S 1,4-DIFLUOROBENZENE
3S CHLOROBENZENE-D5

Targets:

1T 1,2-DICHLOROETHANE-D4
2T TOLUENEN-D8
3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	5.07	436			STD	0.79	100.0	%
2S	7.40	686			STD	0.73	100.0	%
3S	14.42	1435			STD	0.76	100.0	%
1T	6.02	537	65. / 128.	337472. / 127976.	1	0.84	100.6	%
2T	10.90	1060	98. / 117.	758912. / 617040.	3	0.81	102.3	%
3T	17.70	1788	95. / 117.	322936. / 617040.	3	0.57	96.4	%

24404 MW-35 DAMES & MOORE EMU2.85
0-OCT-86 16:27:35 TIC MAXIMUM CURRENT=3605210



EXTREL Quantitation Report

Name: _____ W.D. _____
Analyst: _____ Date: _____
Reviewer: _____ Date: _____
Comments: _____

Library used: SY:VDASUR
Data file name: SY:D24404
Injection time: 20-OCT-86 16:27:35
Comments:
MW-3S DAMES \$ MOORE EMV2.85
Dilution factor: 1.00

Library entries as follows:

Standards:

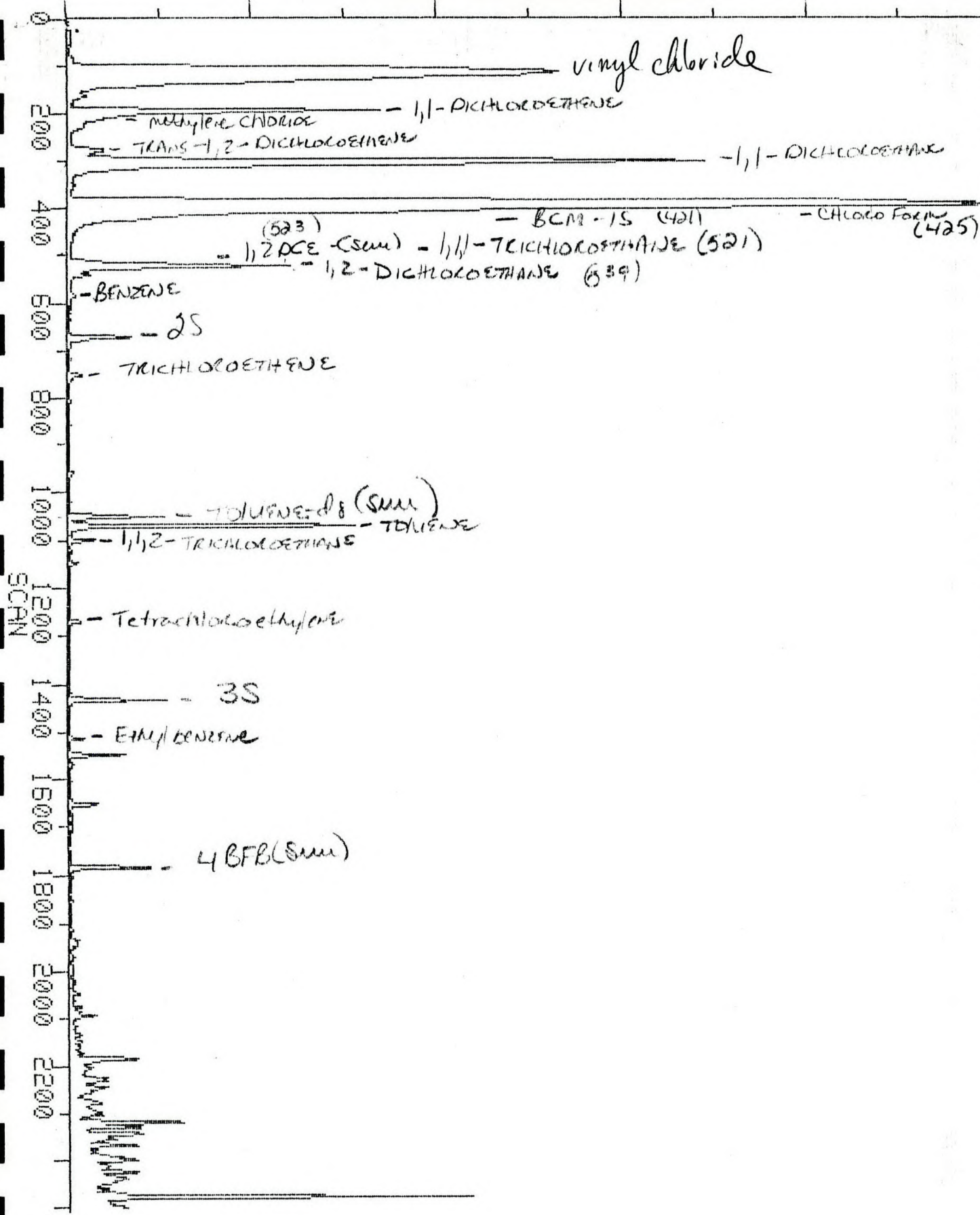
1S BROMOCHLOROMETHANE
2S 1,4-DIFLUOROBENZENE
3S CHLOROBENZENE-D5

Targets:

1T 1,2-DICHLOROETHANE-D4
2T TOLUENEN-D8
3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	5.10	440			STD	0.71	100.0	%
2S	7.42	687			STD	0.73	100.0	%
3S	14.42	1436			STD	0.80	100.0	%
1T	6.03	539	65. / 128.	354566. / 122926.	1	0.79	110.0	%
2T	10.90	1060	98. / 117.	781248. / 638256.	3	0.81	101.8	%
3T	17.67	1784	95. / 117.	339994. / 638256.	3	0.57	98.1	%

0224405 5.0ML DAME58MOORE + 1.0UL INT STD/SURR EMU2.85
20-OCT-86 17:13:11 TIC MAXIMUM CURRENT=3638992



EXTREL Quantitation Report

Name: _____ W.D. _____
Analyst: _____ Date: _____
Reviewer: _____ Date: _____
Comments: _____

Library used: SY:VOASUR
Data file name: SY:D24405
Injection time: 20-OCT-86 17:13:11
Comments:
5.0ML DAMES&MOORE + 1.0UL INT STD/SURR EMV2.85
Dilution factor: 1.00

Library entries as follows:

Standards:

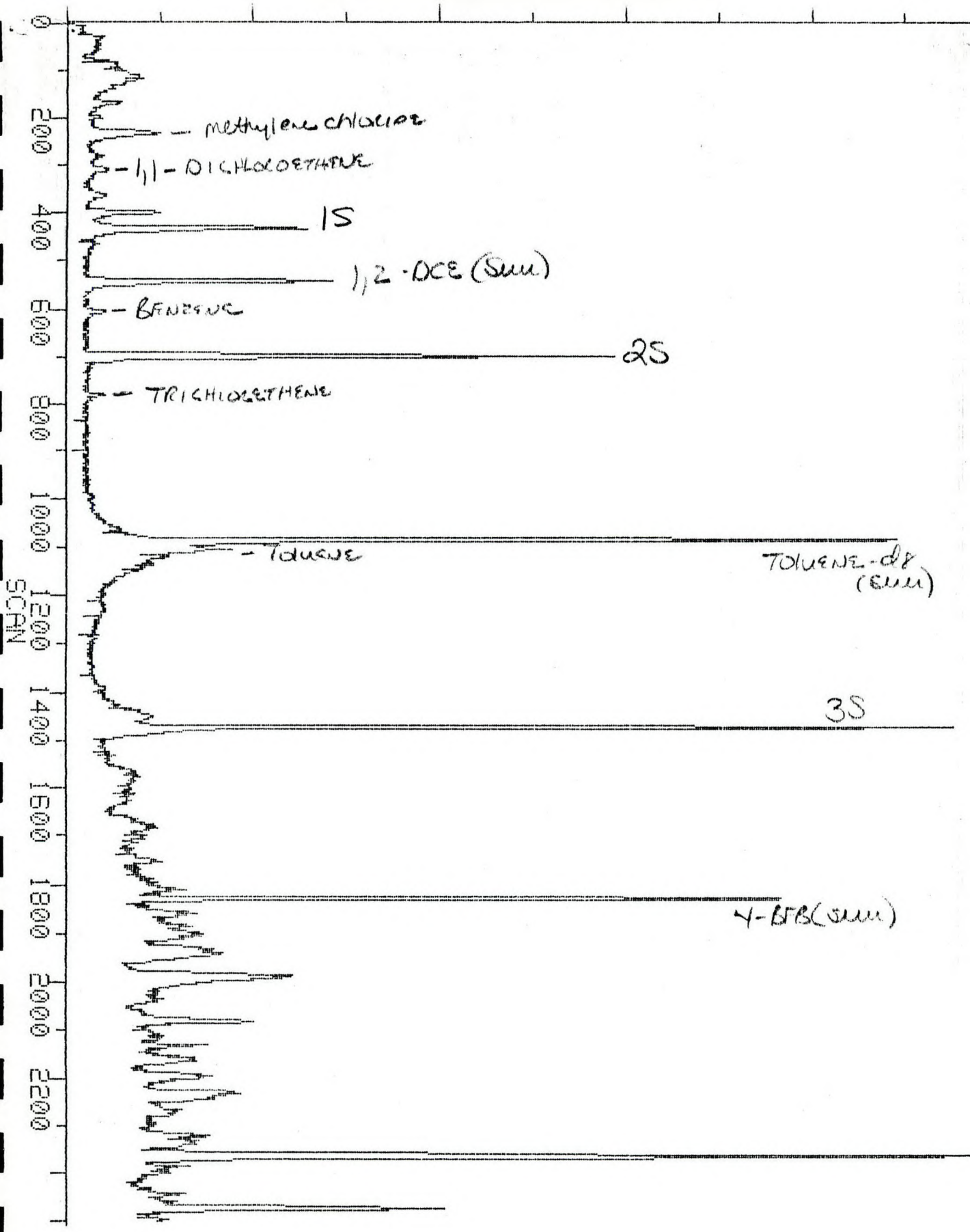
1S BROMOCHLOROMETHANE
2S 1,4-DIFLUOROBENZENE
3S CHLOROBENZENE-D5

Targets:

1T 1,2-DICHLOROETHANE-D4
2T TOLUENEN-D8
3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	4.93	421 ✓			STD	0.67	100.0	%
2S	7.27	671 ✓			STD	0.73	100.0	%
3S	14.37	1431 ✓			STD	0.84	100.0	%
1T	5.88	523 ✓	65. / 128.	314784. / 90672.	1	0.54	132.4	%
2T	10.82	1050 ✓	98. / 117.	832544. / 665904.	3	0.70	104.0	%
3T	17.67	1784 ✓	95. / 117.	349488. / 665904.	3	0.57	96.7	%

024406 5.0ML DAMESBMOORE MA-5 + 1.0UL INT STD/ SURR EMU 2.85
20-OCT-86 17:44:01 TIC MAXIMUM CURRENT=441631



EXTREL Quantitation Report

Name: _____ W.O. _____
Analyst: _____ Date: _____
Reviewer: _____ Date: _____
Comments: _____

Library used: SY:VDASUR
Data file name: SY:D24406
Injection time: 20-OCT-86 17:44:01
Comments:
5.0ML DAMES&MOORE MW-5 + 1.0UL INT STD/ SURR EMV 2.85
Dilution factor: 1.00

Library entries as follows:

Standards:

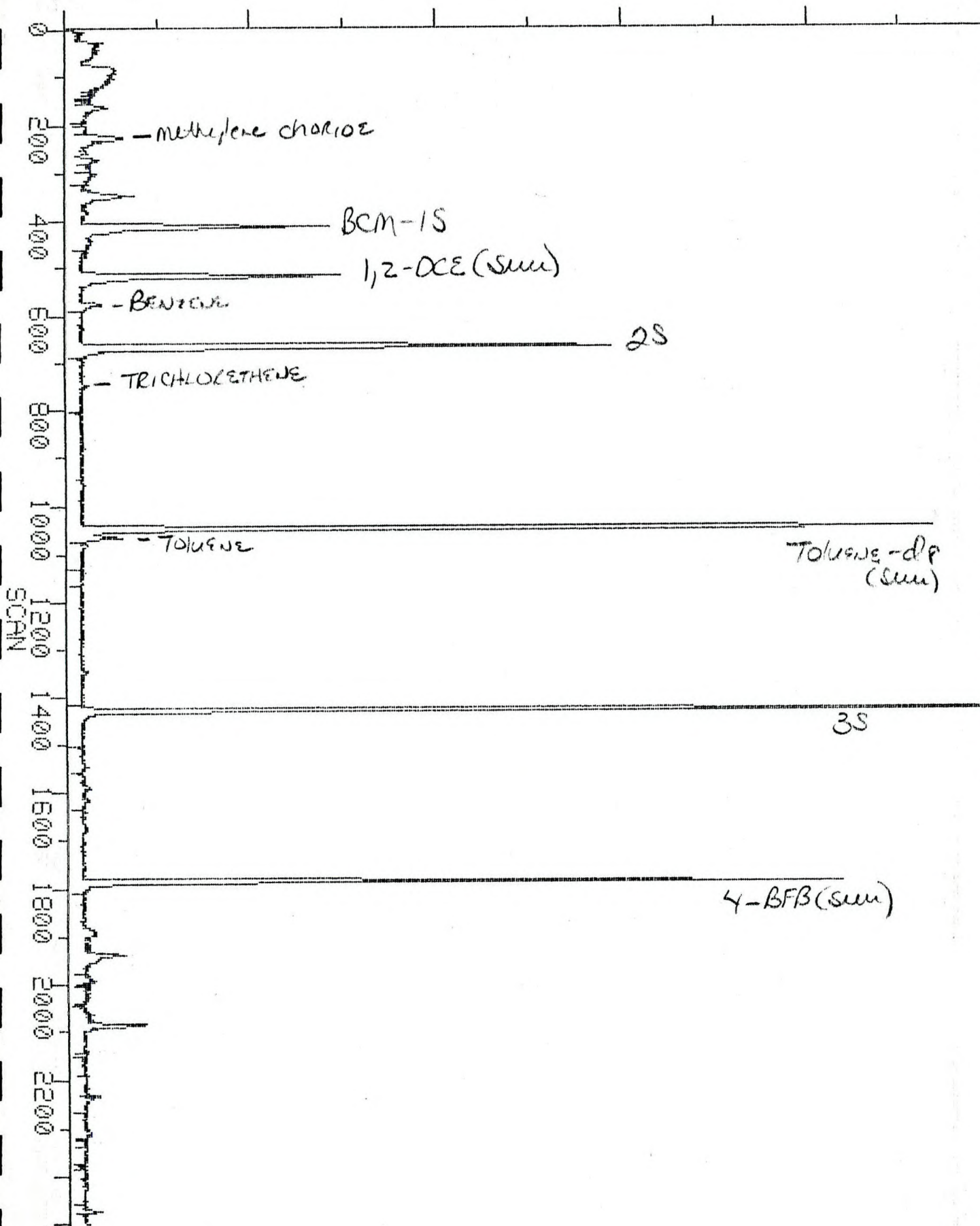
1S BROMOCHLOROMETHANE
2S 1,4-DIFLUOROBENZENE
3S CHLOROBENZENE-D5

Targets:

1T 1,2-DICHLOROETHANE-D4
2T TOLUENEN-D8
3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	5.03	432			STD	0.79	100.0	%
2S	7.50	696			STD	0.79	100.0	%
3S	14.73	1470			STD	0.76	100.0	%
1T	6.05	540	65./ 128.	347970./ 133502.	1	0.74	99.4	%
2T	11.13	1085	98./ 117.	795456./ 653968.	3	0.81	101.2	%
3T	18.08	1828	95./ 117.	346056./ 653968.	3	0.54	97.5	%

D24407 5.0ML DAMESMOORE MM-6 + 1UL INT STD/SURR 2.85EMV
20-OCT-86 22:00:15 TIC MAXIMUM CURRENT=450941



EXTREL Quantitation Report

Name: _____ W.O. _____
 Analyst: _____ Date: _____
 Reviewer: _____ Date: _____
 Comments: _____

Library used: SY:VDASUR
 Data file name: SY:D24407
 Injection time: 20-OCT-86 22:00:15
 Comments:
 5.0ML DAMES&MOORE MW-6 + 1UL INT STD/SURR 2.85EMV
 Dilution factor: 1.00

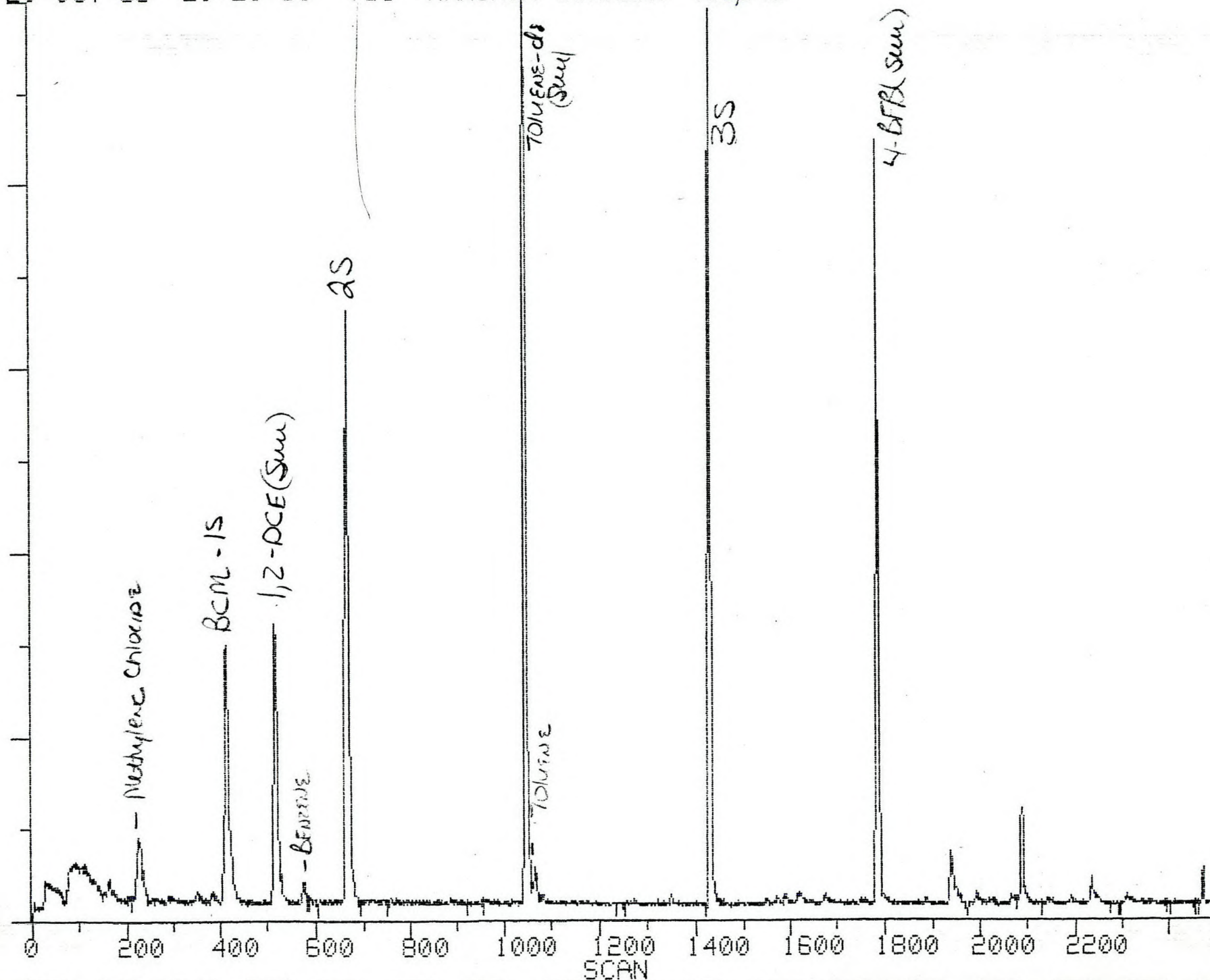
Library entries as follows:

Standards:
 1S BROMOCHLOROMETHANE
 2S 1,4-DIFLUOROBENZENE
 3S CHLOROBENZENE-D5

Targets:
 1T 1,2-DICHLOROETHANE-D4
 2T TOLUENEN-D8
 3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	4.88	416			STD	0.87	100.0	%
2S	7.23	667			STD	0.73	100.0	%
3S	14.35	1429			STD	0.76	100.0	%
1T	5.80	514	65./ 128.	395076./ 145375.	1	0.74	103.7	%
2T	10.78	1047	98./ 117.	802824./ 752384.	3	0.72	88.7	%
3T	17.68	1785	95./ 117.	408928./ 752384.	3	0.70	100.1	%

DMBLK 5.0ML BLANK DAMES&MOORE + INTSTD/SURR EMU2.85
20-OCT-86 20:20:30 TIC MAXIMUM CURRENT=419542



EXTREL Quantitation Report

Name: _____ W.O. _____
Analyst: _____ Date: _____
Reviewer: _____ Date: _____
Comments: _____

Library used: SY:VDASUR
Data file name: SY:DMBLK
Injection time: 20-OCT-86 20:20:30
Comments:
5.OML BLANK DAMES&MOORE + INTSTD/SURR EMV2.85
Dilution factor: 1.00

Library entries as follows:

Standards:

1S BROMOCHLOROMETHANE
2S 1,4-DIFLUOROBENZENE
3S CHLOROBENZENE-D5

Targets:

1T 1,2-DICHLOROETHANE-D4
2T TOLUENEN-D8
3T 4-BFB

No.	Time	Scan	Tmass/Smass	Tarea/Sarea	Ref	Fit	Conc	Units
1S	4.88	416			STD	0.83	100.0	%
2S	7.23	667			STD	0.73	100.0	%
3S	14.37	1430			STD	0.84	100.0	%
1T	5.82	515	65./ 128.	390792./ 154686.	1	0.84	96.4	%
2T	10.78	1047	98./ 117.	896432./ 735152.	3	0.72	101.4	%
3T	17.68	1787	95./ 117.	396616./ 735152.	3	0.57	99.4	%

APPENDIX B

BORING LOGS

**DAMES & MOORE
BORING LOG**

CLIENT: Carrier
LOCATION: Syracuse, NY

BORING NO.: MW-1
SURFACE ELEV: 409.16'

DRILLING METHOD: Hollow-Stem Auger

SAMPLING METHOD: Split Spoon

DATE STARTED: 12/19/85

DATE FINISHED: 12/19/85

SAMPLE NO.	BLOWS/FT	SAMPLE TYPE	DEPTH IN FT.	SOIL GRAPH	MATERIAL DESCRIPTION
			0	ML	Silt, little fine sand, (fill); brown
1	30	SS	1	FILL	
			2		Silt, little fine sand, brown
2	42	SS	3	ML	
			4		Silty fine sand; trace clay, brown
3	16	SS	5		Top of screen at 4 feet
			6		
4	12	SS	7		
			8		
			9	SM	
			10		
5	13	SS	11		
			12		
			13		
			14		Bottom of screen at 14 feet
6	17	SS	15	CL ML	Clay, little silt
			16		Boring terminated at a depth of 16.0 feet on 12/19/85.

EXPLANATION:



SCREENED AREA

**DAMES & MOORE
BORING LOG**

CLIENT: Carrrier
LOCATION: Syracuse, NY

BORING NO.: MW-2
SURFACE ELEV: 404.46'

DRILLING METHOD: Hollow-Stem Auger

SAMPLING METHOD: Split Spoon

DATE STARTED: 12/19/85

DATE FINISHED: 12/19/85

SAMPLE NO.	BLOWS/FT	SAMPLE TYPE	DEPTH IN FT.	SOIL GRAPH	MATERIAL DESCRIPTION
			0	----	Asphalt, 1"
			1	GP	Gravel, little fine sand and silt; brown
			2		
1	23	SS	3		Silt, little fine sand grading to some fine sand, brown
			4	ML	
2	16	SS	5		Top of screen at 5 feet
			6		
3	14	SS	7		Fine sand, little silt; brown
			8		
			9	SM	
			10		
4	23	SS	11		
			12		
			13	ML	Silt, little clay, trace fine sand; brown
5	7	SS	14	CL	
			15		Bottom of screen at 15 feet
			16		
			17		
			18		
			19		
			20		Boring terminated at a depth of 20.0 feet on 12/19/85.

EXPLANATION:



SCREENED AREA

**DAMES & MOORE
BORING LOG**

CLIENT: Carrier
LOCATION: Syracuse, NY

BORING NO.: MW-3S
SURFACE ELEV: 403.56'

DRILLING METHOD: Hollow-Stem Auger

SAMPLING METHOD: Split Spoon

DATE STARTED: 12/18/85

DATE FINISHED: 12/19/85

SAMPLE NO.	BLOWS/FT	SAMPLE TYPE	DEPTH IN FT.	SOIL GRAPH	MATERIAL DESCRIPTION
			0	----	1" Asphalt
			1		Silt, little fine sand, some fine sand layers; brown
			2		
			3	ML	
			4		Top of screen at 4 feet
			5		
			6		Fine sand, little silt; brown
			7		
			8	SM	
			9		
			10		Silt, some clay, trace fine sand; brown
			11		
			12	ML	
			13		
			14		Bottom of screen at 14 feet Boring terminated at a depth of 14.0 feet on 12/19/85. No samples taken. Boring was adjacent to MW-3D and assumed to have the same stratigraphy.

EXPLANATION:



SCREENED AREA

**DAMES & MOORE
BORING LOG**

CLIENT: Carrier
LOCATION: Syracuse, NY

BORING NO.: MW-3D
SURFACE ELEV: 403.46'

DRILLING METHOD: Hollow-Stem Auger

SAMPLING METHOD: Split Spoon

DATE STARTED: 12/18/85

DATE FINISHED: 12/18/85

SAMPLE NO.	BLOWS/FT	SAMPLE TYPE	DEPTH IN FT.	SOIL GRAPH	MATERIAL DESCRIPTION
			0	----	1" Asphalt
1	12	SS	1		Silt, little fine sand, some fine sand layers; brown
			2		
			3	ML	
			4		
2	18	SS	5		
			6		Fine sand, little silt; brown
3	34	SS	7	SM	
			8		
4	23	SS	9		
			10		Silt, some clay, trace fine sand; brown
5	4	SS	11		
			12		
6	7	SS	13		
			14		
			15	ML	
7	6	SS	16		
			17		
			18		
			19		
			20		Fine sand and silt; brown
8	8	SS	21	SM	
			22	ML	Top of screen at 22 feet
			23		
			24		
			25		
			26	SM	Sand, little fine to medium gravel and silt; brown (this layer correspondsto till noted on other on-site boring logs)
			27		Bottom of screen at 27 feet Boring terminated at a depth of 27.0 feet on 12/18/85.

EXPLANATION:



SCREENED AREA

**DAMES & MOORE
BORING LOG**

CLIENT: Carrier
LOCATION: Syracuse, NY

BORING NO.: MW-4
SURFACE ELEV: 404.46'

DRILLING METHOD: Hollow-Stem Auger

SAMPLING METHOD: Split Spoon

DATE STARTED: 12/19/85

DATE FINISHED: 12/19/85

SAMPLE NO.	BLOWS/FT	SAMPLE TYPE	DEPTH IN FT.	SOIL GRAPH	MATERIAL DESCRIPTION
			0		Gravel, little fine sand and silt; brown (fill)
			1		
			2		
1	20	SS	3	GW	Top of screen at 5 feet
			4	GM	
2	4	SS	5	FILL	
			6		
3	2	SS	7		
			8		
			9		
			10		
4	10	SS	11		
			12		
			13		
5	4	SS	14	ML	
			15		Bottom of screen at 15 feet Boring terminated at a depth of 15 feet on 12/19/85.

EXPLANATION:



SCREENED AREA

**DAMES & MOORE
BORING LOG**

CLIENT: Carrier
LOCATION: Syracuse, NY

BORING NO.: MW-5
SURFACE ELEV: 395.46'

DRILLING METHOD: 4 1/4" Hollow Stem Auger

SAMPLING METHOD: Split Spoon

DATE STARTED: 10/13/86

DATE FINISHED: 10/13/86

SAMPLE NO.	BLOWS/FT	SAMPLE TYPE	DEPTH IN FT.	SOIL GRAPH	MATERIAL DESCRIPTION
			0	ML	Extremely moist, black sand-silt-clay fill with little gravel and wood fragments
1	13	SS	1	FILL	
			2	SM	Extremely moist to wet, reddish-brown fine to coarse sand, some silt, trace fine fine gravel, loose
			3		
2	10	SS	4	FILL	
			5	ML	Top of screen at 5 feet Extremely moist, gray sand-silt-clay with little fine gravel, medium stiff
3	5	SS	6		Extremely moist, black organic rich silt and fine sand, numerous roots and wood fragments
			7	ML	
4	4	SS	8		Extremely moist, black silt and fine to medium sand with organic debris, roots, wood chips, some bedding apparent
			9		
5		SS	10		grading to fine sand, some silt, little organic debris, occasional dark stained beds, loose, slight tendency to liquify when disturbed
			11	SM	
6	4	SS	12		Wet, grayish-brown silty clay Bottom of screen at 15 feet
			13		
7	2	SS	14	CL	Boring terminated at a depth of 15.0 feet on 10/13/86.
			15		

EXPLANATION:



SCREENED AREA

**DAMES & MOORE
BORING LOG**

CLIENT: Carrier
LOCATION: Syracuse, NY

BORING NO.: MW-6
SURFACE ELEV: 404.68'

DRILLING METHOD: 4 1/4" Hollow Stem Auger

SAMPLING METHOD: Split Spoon

DATE STARTED: 10/13/86

DATE FINISHED: 10/13/86

SAMPLE NO.	BLOWS/FT	SAMPLE TYPE	DEPTH IN FT.	SOIL GRAPH	MATERIAL DESCRIPTION
			0		Auger to 1' and began sampling at 1'
			1		Moist, reddish-brown silt with little fine gravel and occasional thin (<1" thick) sand lens, stiff, becoming very stiff below 3.0 feet
1	10	SS	2		
			3	ML	
2	24	SS	4		Top of screen at 5 feet
			5		
3	24	SS	6		
			7		Moist, reddish-brown, fine to medium sand, some silt, little to trace fine gravel, very dense in place
4	51	SS	8		
			9		
5	19	SS	10		grades at 9.0 feet to extremely moist with a slight tendency to liquify when disturbed, medium dense
			11	SM	
6	12	SS	12		
			13		grades to fine to coarse sand, some silt, occasional thin (<2") silt beds, medium dense
7	20	SS	14		
			15		

Bottom of screen at 15 feet
Boring terminated at a depth of 15.0 feet on 10/13/86.

EXPLANATION:



SCREENED AREA