1

PHASE II REMEDIAL INVESTIGATION REPORT FORMER GRIESS-PFLEGER TANNERY SITE WAUKEGAN, ILLINOIS

RECEIVED

NOV 30 1995 IEPAVDLPC

Prepared for:
Commonwealth Edison Company
One First National Plaza
Chicago, Illinois
and
Beggs and Cobb Corporation
Peabody, Massachusets

Prepared by: Metcalf & Eddy, Inc. One Pierce Place, Suite 1400W Itasca, Illinois

ORIGINAL



TABLE OF CONTENTS

	1 1.0 INTI	RODUCTION	·	
1.			COPE OF REMEDIAL INVESTIGATION	
1.	2 REFO	ORT ORGAN	IZATION	
ECTIO	1 2.0 ENV	VIRONMENT	TAL INVESTIGATION	
2.	1 GEOI	LOGIC/HYD	ROGEOLOGIC INVESTIGATIONS	
2.	2 ENVI	RONMENT	AL SAMPLING PROCEDURE	
	2.2.1	Exploratory	Soil Boring	
		2.2.1.1	Purpose	
		2.2.1.2	Location and Rationale	
		2.2.1.2	Exploratory Borehole Soil Sampling	
	222		Well Installation - Shallow and Deep	
	2.2.2	2.2.2.1		
		2.2.2.2	Purpose	
		2.2.2.3	Delling Mothed	
			Drilling Method	
	0.00	2.2.2.4	Monitoring Well Borehole Soil Sampling	
			Well Installation/Construction Details	
			Well Development Procedures	
	2.2.5		T Level Measurements	
		Groundwate	er Sampling	
	2.2.7		Subsurface Soil Sampling	
		2.2.7.1	Purpose	
			7.1(a) Site Classification	:
			7.1(b) Parameter Specific Soil Sampling	
		2.2.7.2	Locations and Rationale	- :
		2.2.7.3	Surface and Subsurface Soil Sampling Methods	
			and Equipment	
	2.2.8		ter and Sediment Sampling	,
		2.2.8.1	Purpose	- 1
		2.2.8.2	Locations	
		2.2.8.3	Sampling Methods and Equipment	- 2
			me Estimation	- 1
	2.2.10	O Surveying	Methodology	- 2
	2.2.1	1 Decontami	nation Procedures	- 1
2.	3 SAM	PLE PACKA	GING AND SHIPMENT	- 1
2.	4 INVE	STIGATIVE	DERIVED WASTE	-
.0 R	SULTS A	ND DISCUS	SION OF ENVIRONMENTAL INVESTIGATION	
3.			****************************	
3.		HYDROGEC	DLOGY	
3.	מינות נ	OTTAINE	N OF CHROMIUM WASTE EXCLUSION	

STED OF STREET

LIST OF TABLES AS AN ATTACHMENT (Continued)

4-2 Soil Analytical Results

Pesticide/PCB Compounds

Semi-Volatile Organic Compounds

Inorganic Compounds

TCLP Inorganic Compounds

4-3 Soil Analytical Results

PCDD/PCDF Compounds

4-4 Asbestos Analytical Results

4-5 Soil Analytical Results

Inorganic Compounds - Arsenic Only

TCLP Inorganic Compounds - Arsenic Only

4-6 Wetland Analytical Results

Soil Analytical Results - Semi-Volatile Organic Compounds

Surface Water Analytical Results - Semi-Volatile Organic Compounds

Soil Analytical Results - Pesticide/PCB Compounds

Surface Water Analytical Results - Pesticide/PCB Compounds

Soil Analytical Results - Inorganic Compounds

Surface Water Analytical Results - Inorganic Compounds

4-7 Wetland Analytical Results

AVS and SEM Analytical Results

Soil Chemistry - Total Organic Carbon

Wet Chemistry - Total Organic Carbon

LIST OF FIGURES AS AN ATTACHMENT

- 2-1 Monitoring Well, Soil Boring, Sediment Sample, Production Waste Location Map
- 2-2 Monitoring Well Locations on the Waukegan Generating Station Property
- 3-1 North South Geologic Cross Section
- 3-2 East West Geologic Cross Section
- 3-3 Shallow Groundwater Monitoring Well Contour Map
- 3-4 Deep Groundwater Monitoring Well Contour Map
- 4-1 Surficial Total Pesticide Concentration Map
- 4-2 Subsurficial Total Pesticide Concentration Map
- 4-3 Surficial Total Polychlorinated Biphenyls Concentration Map
- 4-4 Subsurficial Total Polychlorinated Biphenyls Concentration Map
- 4-5 Surficial Total Polynuclear Aromatic Hydrocarbons Concentration Map
- 4-6 Subsurficial Total Polynuclear Aromatic Hydrocarbons Concentration Map
- 4-7 Surficial Chromium Concentration Map
- 4-8 Subsurficial Chromium Concentration Map
- 4-9 Surficial Lead Concentration Map
- 4-10 Surficial Arsenic Concentration Map
- 4-11 Subsurficial Arsenic Concentration Map

EXECUTIVE SUMMARY

The former Griess-Pfleger Tannery is located on the northeast corner of Sand (Pershing) and Dahringer Road. The property consists of approximately 38 acres and is bordered by Dahringer road to the south, Elgin Joliet and Eastern Railroad on the east, a spur of the Chicago Northwestern railroad, which conjoins with the Elgin Joliet and Eastern Railroad (EJ&E) to the north, and Sand (Pershing) Road to the west. The property is currently owned by Commonwealth Edison Company.

The former Griess-Pfleger Tannery was established in 1917 and operated as a leather tanning facility until 1973. As evidenced from the analytical data, chrome tanning processes are believed to have been utilized. Chrome tanning, as generally practiced, consisted of nine basic steps and utilized a number of chemicals in the tanning process. The tanning process produced waste by-products in the form of gaseous reaction products, wastewater, wastewater sludge, and solid waste. Further research is being conducted to determine whether arsenic was part of the tanning process.

In January 1989, the United States Environmental Protection Agency's (USEPA) Field Investigation Team (FIT) performed a preliminary site investigation. Laboratory analyses indicated elevated levels of chromium and lead in the soil.

In June 1992, Commonwealth Edison contracted Metcalf & Eddy, Inc. (M&E) to conduct a Remedial Investigation (RI). The RI conducted at the former tannery was performed voluntarily in cooperation with the Illinois Environmental Protection Agency.

Phase I Remedial Investigation

In May and June 1993, M&E performed various field activities. These activities included installation of seven shallow groundwater monitoring wells; collection of: eleven sediment samples, sixty-eight soil samples (forty-three subsurfical, fifteen surfical and ten monitoring well soil boring samples), two geotechnical samples, nine production waste samples, one cistern water sample, and seven groundwater samples; and performance of a magnetometer survey.

The soil and sediment samples were collected and analyzed for Target Compound List (TCL) volatile and semi-volatile organic compounds, TCL pesticides/PCBs, and Target Analyte List (TAL) inorganic analytes. The production waste samples were analyzed for TCL volatile and semi-volatile organic compounds, TCL pesticides/PCBs, TAL inorganic analytes, and Toxicity Characteristic Leaching Procedure (TCLP) Compounds. The groundwater and cistern water samples were collected for the TCL volatile and semi-volatile organic compounds, TCL pesticides/PCBs, and TAL inorganic analytes in addition to Total Dissolved Solids (TDS). In summary, the Phase I RI investigation results indicated the following:

Numerous inorganic analytes were identified. Elevated levels of chromium and lead were detected consistently in the surface and subsurface soil throughout Areas II and III of the site. Pesticides, PCBs, and base-neutral acid extractable compounds (BNAs), predominantly polynuclear aromatic hydrocarbons (PNAs), were detected in the soil sporadically throughout the site. The concentration of these compounds decreased with depth.

Groundwater samples were collected from all seven shallow groundwater monitoring wells. The laboratory analyses indicated that the base neutral and acid extractable compounds, pesticides/PCBs, and volatile organic compounds were not in any of the groundwater samples collected. Inorganic analytes, arsenic and lead, were detected in one monitoring well above the Illinois Class I Standards (IAC Title 35, Subpart D, Section 620.410(a)). Iron and manganese were also detected above the Illinois Class I Standards. Analytical results for the background sample also indicated levels of iron and manganese above the Class I Standard. Total Dissolved Solids were present above the Illinois Class I Standard in all seven monitoring wells. These levels are presumed to be naturally occurring.

Three of nine production waste samples collected exceeded the TCLP regulatory limits, two for chromium and one for lead.

More detail regarding the Phase I investigation is provided in the March 1994 Remedial Investigation Report - Phase I for the former Griess-Pfleger Tannery Site, Waukegan, Illinois. This report was made final on July 21, 1994 with minor revisions.

Phase II Remedial Investigation

From January to September of 1995, M&E performed additional field activities as part of the Phase II RI. The objectives of the Phase II activities were to define the magnitude and extent of surface and subsurface impact, characterize the type of constituents in the subsurface, and determine if off site areas have been impacted by former tannery operations. The Phase II data, when used in conjunction with the Phase I investigation data, will allow a Baseline Risk Assessment/Ecological Risk Assessment to be completed.

To accomplish the above mentioned objectives, Metcalf & Eddy, Inc. performed various media sampling. This included performing: chromium speciation, waste volume estimation, and installation of one exploratory soil boring and five additional monitoring wells (two shallow and three deep); and collecting: eighty-one surface/subsurface soil samples, two production waste samples, three wetland sediment/surface water samples, two dioxin/furan samples, two asbestos samples, and five groundwater samples.

Based upon the results obtained from the Phase II investigation, four additional monitoring wells were installed and sampled during Phase IIA and IIB of the investigation. In addition, as part of the Phase IIB investigation, 14 groundwater samples were collected through the use of a Geoprobe^R sampling unit.

Geologic Characterization

Both the shallow and deep groundwater monitoring wells installed at the former tannery indicate that groundwater is flowing under unconfined conditions. Deeper soil borings indicated that a clay lens is apparent and discontinuous at the site. Saturated soil was found approximately three to five feet below grade. Static water conditions indicate that the shallow groundwater flow direction is to the east under a gradient of 0.0014 feet per foot. The deeper groundwater flow direction is generally to the east under a hydraulic gradient of 0.0017 feet per foot.

Based on the results of the Phase II site investigation, the geology of the site consists of a well graded to poorly graded sand to silty sand with fill and peat. Clay was apparent in two soil borings but not in significant quantities to justify describing it as a confining layer.

Analytical Characterization

Samples collected were analyzed for either Level III or Level IV IEPA QA/QC. The level of QA/QC was determined based upon the purpose of the sample.

Groundwater

Groundwater analytical results indicated that volatile organic compounds, semi-volatile organic compounds, and pesticide/PCBs, were not detected. However, several inorganic analytes were detected in concentrations exceeding IEPA regulatory limits, including arsenic (total and dissolved), chromium (total), lead (total), and mercury (total). Analytical results for the inorganic analytes indicate that the constituents generally adhere to soil particles whose size is greater than 0.45 micrometers in diameter. Overall, it appears that the inorganic constituents are adsorbed to the soil and not part of the groundwater matrix, with the exception of arsenic.

The Phase II investigation indicated that groundwater containing arsenic exceeding the Illinois Class I Standard has migrated to the eastern edge of the tannery property boundary. Subsequently, two additional monitoring wells were installed as part of the Phase IIA investigation at Commonwealth Edison's Waukegan Generating Station. Analytical results from these two monitoring wells confirmed that groundwater containing arsenic had migrated under the El&E railroad tracks and onto Commonwealth Edison's Waukegan Generating Station property. A Phase IIB investigation was undertaken that utilized a Geoprobe[®] unit to collect groundwater samples to define the extent of the arsenic plume. Using the Geoprobe[®] collected data, two permanent monitoring wells were installed to confirm the arsenic plume delineation. At the present time, this delineation indicates that the arsenic plume has migrated 400 feet from the former Tannery.

Other inorganic analytes which were detected and exceeded the IEPA regulatory limit include: cadmium (total and dissolved), iron (total), and manganese (total and dissolved). Analytical results for the background sample indicated elevated levels of iron and manganese. Elevated

and a more than the state of th

130 D00742

levels of these constituents were also apparent during Phase I sampling activities. Therefore, these constituents can be considered indicative of naturally occurring constituents.

Total Dissolved Solids were detected at or above the Illinois Class I Standard of 1,200,000 μ g/L in eleven of the twelve monitoring wells. These elevated levels are considered naturally occurring.

Surface and Subsurface Soils

Soil analytical results indicate that asbestos was not detected in either of the two soil samples collected.

Elevated levels of semivolatile organic compounds, consisting mainly of polynuclear aromatic hydrocarbons were identified at various locations throughout the site.

Polychlorinated biphenyls were identified at elevated levels in the vicinity or the production waste disposal area.

Significantly elevated levels of pesticides were not identified during the Phase II investigation, indicating that the elevated pesticide concentration identified at one location during the Phase I investigation was an isolated occurrence.

Dioxins and furans were identified in soil samples collected during the Phase II investigation. However, the levels of dioxins/furans identified were not above the USEPA screening levels for protection of human health and the environment.

Elevated levels of chromium, lead, and to a lesser extent arsenic were identified throughout Areas II and III of the site.

Seventeen soil samples exhibiting elevated levels of chromium were analyzed for TCLP metals. The amount of leachable chromium in all of these samples was found to be below regulatory limits. Additionally, these samples were analyzed for hexavalent chromium. It was determined that the chromium exists in the soil as greater than 99.97% trivalent chromium, the more stable, benign, and less mobile form of the two. Leachable metals were not identified in any of the samples above the TCLP regulatory limit, with the exception of mercury at one sampling location.

SECTION 1.0 INTRODUCTION

1.1 PURPOSE AND SCOPE OF REMEDIAL INVESTIGATION

The purpose of the Phase II Remedial Investigation (RI) is to provide adequate information to further delineate the magnitude and extent of the surface and subsurface impacted areas of the former tannery previously depicted during the Phase I RI. Specifically, the Phase II RI aided in the determination of the volume of the environmental media and helped determine whether off-site areas have been impacted by former tannery operations. These activities were accomplished by performing the following activities:

- Exploratory Soil Boring/Monitoring Well Installation One exploratory soil boring was
 installed. The objective of this boring was to determine whether a uniform and
 continuous confining layer was present at depth prior to installing the deeper monitoring
 wells.
- Three deep monitoring wells and two shallow monitoring wells were installed to supplement the seven shallow monitoring wells that previously existed on site. The purpose of the deeper monitoring wells was to determine if deeper stratigraphic units or groundwater bearing zones have been impacted. The shallow monitoring wells were installed to assess the extent of groundwater impact and in addition to determine whether constituents were migrating to or from off-site locations.
- Four additional shallow monitoring wells were installed as part of the Phase IIA and IIB of the RI at Commonwealth Edison's Waukegan Generating Station to determine the extent of the arsenic plume migration. In addition, 14 shallow groundwater samples were collected as part of the Phase IIB investigation utilizing a Geoprobe^R groundwater sampling unit.
- Waste Volume Estimation The objective of this was to determine the gross volume of tannery waste material (tannery hide material like production waste sample PW-5) located in the southern edge of Area III's southwestern section.
- Chromium Waste Exclusion The purpose of this is to determine if on-site wastes qualify
 for the chromium hazardous waste exclusion as stated in 35 IAC 721.104 (b)(6) as it
 pertains to tannery waste streams.
- Wetland Sediment Soil and Surface/Subsurface Soil Sampling The objective of soil sample collection was to collect soil samples in areas not previously sampled and in areas suspected of containing elevated levels of constituents.

- Surface soil samples were collected and analyzed for select inorganic TCLP parameters, specifically chromium, lead, mercury, and arsenic, throughout the site to determine the applicable regulatory framework.
- The speciation of chromium; trivalent chromium (Cr^{*}) or hexavalent chromium (Cr^{*}), was determined.
- Soil samples were collected in and around select areas in which elevated levels of specific constituents were identified during Phase I activities to more fully define the lateral extent of impact.
- Soil samples were collected and analyzed for asbestos to determine if it was present in dredge sand used as fill at the site.
- Soil samples were collected where PCBs were detected during Phase I activities.

 Two samples were analyzed for dioxin/furans.
- Off-site wetland surface water and sediment samples were collected from the wetland area located south of the southeastern section of Area III.
- Sampling and Analysis Surface/subsurface soil, wetland surface water and sediment, and groundwater media samples were collected and analyzed. The investigation analyte list was reduced for both soil and groundwater to those parameters which were detected in elevated levels during Phase I of the RI. However, those samples submitted for analysis from newly installed monitoring wells or from the wetland area were analyzed for Level IV TCL/TAL parameters. Media samples collected during the Phase II RI investigation were analyzed for the following parameters:
 - △ Groundwater/Wetland Surface Water Environmental Sampling (Phase II):

Monitoring Wells Installed During Phase I Activities (7):

Level III TCL SVOCs, TCL Pesticides/ PCBs (P/PCBs), Total and Dissolved RCRA Metals, Hexavalent Chromium, Total Suspended Solids (TSS), Total Dissolved Solids (TDS), Turbidity, Cyanide.

Monitoring Wells Installed During Phase II Activities (5):

Level IV CLP Target Analyte List (TAL, Total and Dissolved), TCL VOCs, TCL SVOCs, TCL P/PCBs, Hexavalent Chromium, TSS, TDS, Turbidity, Cyanide.

THE RESIDENCE OF THE PROPERTY OF THE PROPERTY

Wetland Surface Water (3):

Level IV CLP TCL SVOCs, TCL P/PCBs,

TOC, Total and Dissolved RCRA Metals.

Monitoring Wells Installed During Phase IIA (2):

Level III Total and Dissolved RCRA

Metals.

Geoprobe^R Groundwater Samples During Phase IIB (14):

Level III Total and Dissolved Arsenic.

Monitoring Wells Installed During Phase IIB (2):

Level III Total and Dissolved RCRA

Metals.

Soil Environmental Sampling (Phase II):

Phase II Monitoring Well

Soil Borings:

Level IV CLP TAL TCL SVOCs, TCL

P/PCBs.

Wetland Sediment Sampling:

Level IV CLP TCL SVOCs, TCL P/PCBs, RCRA Metals, TOC, Acid Volatile Sulfide and Selected Simultaneously Extractable Metals (AVS/SEM), Hexavalent Chromium.

Surface/Subsurface Soil Sampling: TCLP and

TCLP and RCRA Metals; Specific Parameters: Hexavalent Chromium, P/PCBs, Dioxin/Furan, Polynuclear Aromatic Hydrocarbons (PNAs), Asbestos, and Arsenic; Level III TCL SVOC and TCL

P/PCBs.

Two deviations from the Phase II RI work plan activities occurred:

In-situ hydraulic conductivity tests were not conducted in any of the newly installed monitoring wells as stated in M&E's work plan. The purpose of performing the hydraulic conductivity tests would be to determine whether hydraulic communication existed between the shallow and the deep aquifers if a competent clay confining layer was present in the subsurface. Because this confining layer was not apparent and the strata in the newly installed monitoring wells were similar to that encountered during Phase I activities, hydraulic conductivities were assumed to be similar.

A Shelby tube sample was not collected from the subsurface because no clay layer was encountered. The purpose of the Shelby Tube would be to gather geotechnical data to ensure that the clay layer was competent and continuous with no discernable sand layers. The soil was to be analyzed for the following parameters using appropriate ASTM standards:

•	D 2216		Moisture Content	
٠	D 2434-68	•	Permeability of Granular Soil via Flexible Wall Permeameter	
•	D 422-63/ D 1140-54	-	Particle Size Analysis of Soils (Combined Analysis)	
	D 2937-83		Density of Soil	
•	D 4318-84	-	Liquid Limit, Plastic Limit, Plasticity Index of Soil	
•	D 2248	-	Visual Soil Classification	

All data gathered from both phases of the investigation were completed in anticipation of performing a Baseline Risk Assessment/Ecological Risk Assessment (BRA/ERA). Conducting a BRA will determine the risk to human health and the environment. Additionally, the ERA is conducted as part of the BRA to evaluate the possibility of adverse ecological effects occurring as a result of exposure. The performance of these activities will be evaluated after all Phase II data have been gathered, reduced, and integrated with Phase I data.

To perform the above referenced activities, M&E utilized existing IEPA-approved planning documents for the proposed sample collection and analysis. The documents that were utilized included the Field Sampling Plan (FSP), dated February 1993, and the Quality Assurance Project Plan (QAPP), dated December 1992. Both of these documents were prepared for the Phase I RI at the Former Griess-Pfleger Tannery Site and were deemed fully applicable for the Phase II RI.

The former Griess-Pfleger Tannery is in the Illinois Pre-Notice Site Cleanup Program (formerly known as the Illinois Voluntary Cleanup Program). M&E and its laboratory subcontractor abided by the data quality objectives set forth in the Analytical Quality Assurance Plan (AQAP) for the IEPA Bureau of Land Pre-Notice Site Cleanup Program. The levels of Data Quality applied during the Phase II RI included Level III, Engineering and Level IV, Confirmational. Details of the level of QA/QC are provided in Section 2 of this report.

1.2 REPORT ORGANIZATION

This RI Report is divided into five sections. These sections include:

 Section 1.0, Introduction. This section provides a brief overview describing the site activities and work objectives. فيأمل مونتانيه الموارينين ماياراتها المتنائد المانكانية الكامايل والمايتان المتنازية المانين

- Section 2.0, Environmental Investigation. This section details the methods and techniques of the environmental investigation. Monitoring well, exploratory soil boring, wetland surface water/sediment, and surface and subsurface soil sampling methods and decontamination methods are explained.
- Section 3.0, Results and Discussion of Environmental Investigations. This section details
 site specific geology and hydrogeology. Most information gathered during Phase I RI
 activities has not deviated. New information, not submitted as part of the Phase I
 Report, is presented in this section.
- Section 4.0, Nature and Extent of Contamination. This section describes the nature, extent, and magnitude, of contamination in the soil and groundwater.
- Section 5.0, Summary, Conclusions, and Recommendations. This section summarizes the findings and presents recommendations.
- Section 6.0, References.

The following sections, which were part of the Phase I Report, will not be a part of the Phase II Report as explained below.

- Site Background. The Phase I Report (Section 2.0, Site Background) described the
 detailed site history, background, and provided a summary of previous investigations.
 This information was not reiterated in its entirety but it is briefly provided as part of the
 Phase II Executive Summary.
- Environmental Setting. The Phase I Report (Section 3.0, Environmental Setting), provides information regarding land use and surrounding population, climatology, surface water, soils, topography, regional geology and stratigraphy, regional hydrogeology, and regional groundwater use. This information has not deviated since Phase I RI Report preparation. Therefore, this information is not provided in this report submission.

2.2.1 Exploratory Soil Boring

2.2.1.1 Purpose

According to the Illinois State Water Survey, available well logs indicated that a clay layer had been documented to exist :9 to 22 feet below grade. The exploratory soil boring (EB-1) was drilled on-site to determine whether a uniform and continuous subsurface confining layer existed for the installation of deeper monitoring wells. Existing on-site monitoring wells (MW-1, MW-2, MW-3, MW-4, MW-5, MW-6, and MW-7) extend to a depth of approximately thirteen feet below grade. Only poorly graded to well graded sands or gravelly sands were encountered to the base of each borehole. A confining layer was not encountered.

2.2.1.2 Location and Rationale

One exploratory soil boring was drilled to 30 feet below grade. The location of this boring was based upon ease of accessibility. The exploratory boring was located in the southwestern section of Area II.

2.2.1.3 Exploratory Borehole Soil Sampling

Soil samples were collected continuously to 30 feet below grade during drilling. A two foot long split-spoon sampler (ASTM D1586) was driven by a 140-pound hammer free-falling 30 inches. Upon retrieval, the split spoon was opened and the soil sample was screened using a photoionization detector (PID). A stainless steel knife was used to part the sample in order to survey the interior portion of the soil sample for logging purposes. The stainless steel knife was decontaminated between sample intervals in accordance with the procedures described in Section 2.2.11. An aliquot of soil was collected from the split spoon and placed into sealable plastic bags. After the soil gasses were allowed to equilibrate, a PID reading was taken. Photoionization detection readings are illustrated on the geologic logs located in Appendix A.

The soil samples were described using ASTM Method D2488 (Description of Soils) and classified in the field using the Unified Soil Classification System (USCS). Descriptions and classifications were recorded onto geologic logs which are included in Appendix A.

It was anticipated that the exploratory borehole would penetrate a competent and continuous (one without discernable sand) confining layer approximately 25 feet below grade. Field personnel pursued a greater depth (30 feet) to determine if the confining layer was further below grade. Because a confining layer was not encountered, soil samples were not collected for geotechnical analysis.

2.2.2 Monitoring Well Installation - Shallow and Deep

2.2.2.1 Purpose

Five monitoring wells, two shallow (MW-8, MW-9) and three deep (MW-1A, MW-5A, and MW-7A) were installed during field activities. The purpose of installing the shallow monitoring wells was to better characterize the site to determine if the arsenic impacted the groundwater east of SB-06 and to determine if additional off-site influences could be affecting the site. The deep monitoring wells were installed to determine: whether deeper stratigraphic units were affected by former tannery operations, the groundwater flow direction for the deeper stratigraphic unit(s), and whether contamination is migrating at depth.

Subsequent to installation of the five monitoring wells, Phase IIA and IIB investigations were undertaken. These investigations consisted of installation of four additional shallow monitoring wells (MW-10, MW-11, MW-12, and MW-13) and collection of 14 groundwater samples utilizing a Geoprobe^R groundwater sampling unit. These additional monitoring wells were installed to determine the extent of the arsenic plume that had migrated to the eastern edge of the site boundary, as shown by the arsenic concentration in MW-8.

2.2.2.2 Location and Rationale

Two water table monitoring wells (screens intersecting the water table) were installed as part of the field activities. One monitoring well (MW-8) was installed adjacent to the eastern property fence line. The other monitoring well (MW-9) was installed approximately 215 feet west of the eastern property fence line in the northern portion of the site.

Three deep monitoring wells were installed as part of the field activities. Monitoring well MW-7A is located off-site and in the Illinois Department of Transportation's (IDOT's) right-of-way approximately 65 feet east of the soft shoulder of Amstuz Expressway and approximately one-third mile south of Greenwood Avenue. This location was amended with respect to an Illinois Environmental Protection Agency (IEPA) November 30, 1994 request regarding improving the triangulation between the proposed three deep monitoring wells. Two other deep monitoring wells, MW-1A and MW-5A, are located adjacent to downgradient monitoring wells MW-1 and MW-5, respectively. All three monitoring wells will help determine: a) whether the deeper stratigraphic units have been affected by former tannery operations, b) the groundwater flow direction for the deeper stratigraphic unit(s), and c) whether contamination is migrating off-site at depth.

The four additional shallow monitoring wells were installed at Commonwealth Edison's Waukegan Generating Station. One monitoring well (MW-10) was installed approximately 300 feet east of MW-1, and MW-11 was installed approximately 200 feet east of MW-8. Monitoring well MW-12 was installed 300 feet east of MW-10 and MW-13 was installed 450 feet east of MW-11. The locations of the monitoring wells at the Waukegan Generating Station are shown in Figure 2-2.

ni manganakan sah

2.2.2.3 Drilling Method

Due to the shallow nature of the groundwater (approximately three feet below grade) and the general sponginess of the overlying soil, an all-terrain vehicle (ATV) was utilized.

Prior to drilling, the drill rig and related equipment were decontaminated. All drilling was initiated using 4.25-inch hollow stem augers (HSA). Small amounts of potable water were added into the hollow stem augers to alleviate heaving sands. Potable water was obtained from Commonwealth Edison's Waukegan Generating Station which acquires its water from the Waukegan Public Water Supply (The station's water was analyzed prior to inception of Phase I activities). Deep soil borings MW-5A and MW-1A were straight drilled (without sampling) to approximately ten feet below grade because the upper stratigraphy was previously defined during the Phase I RI by the adjacent shallow monitoring wells. Continuous split spoon samples were collected from 10 feet below grade to total depth in MW-1A and MW-5A and from the ground surface to total depth in MW-7A because the distance between MW-7 and MW-7A was too great to correlate the surfical stratigraphy. The stratigraphy was determined by using a split spoon sampler (ASTM D 1586) driven by a 140-pound hammer free-falling 30 inches. The samples were field screened utilizing a photoionization detector (PID). All soil samples were described in the field using ASTM Method D 2488 and classified using the Unified Soil Classification System (USCS). Geologic logs are included as Appendix A.

All drill cutting were containerized in 55 -gallon drums and staged at a central location at the site, with the exception of the four monitoring wells installed at the Generating Station. With approval from IEPA, the cuttings from these monitoring wells was thin spread around the well location. For the soil that was containerized, each drum was labelled with the well number, date, and drum contents.

2.2.2.4 Monitoring Well Borehole Soil Sampling

Upon split spoon sampler retrieval, the sampling device was opened and screened using a PID. A stainless steel knife was used during the screening process to part the sample in order to survey portions of the sample. Portions of the sample were quickly collected and placed into soil sampling jars. Containers for volatile organic compound analysis were filled first, SVOCs second, and P/PCBs and metals last. The stainless steel knife was cleaned between sample intervals. One soil sample was collected from the bottom of each deep monitoring well borehole: MW-1A, 23 - 25 feet below grade; MW-5A, 22 - 24 feet below grade; and MW-7A, 23 - 25 feet below grade.

2.2.3 Monitoring Well Installation/Construction Details

All monitoring wells were installed through the HSAs. All monitoring wells were constructed of 2-inch diameter Type 304 stainless steel with flush joints. The well screens consisted of Type 304 continuous wire-wrap stainless steel with 0.010-inch openings. The well screens were 10 feet in length for both shallow and deep monitoring wells.

Deep Monitoring Wells - MW-1A, MW-5A, and MW-7A

Due to the presence of heaving sands, borings for both MW-5A and MW-7A were over drilled one foot. Therefore, prior to placing the well string in the borehole, 1.0 foot of clean 20/40 sized silica sand was place into the bottom of these boreholes. Heaving sands were not a problem for monitoring well MW-1A. Therefore, this well string was placed at the bottom of the borehole. The silica sand filter pack was placed in the annular space around the well screens to minimize intrusion of fine-grained sediment into the monitoring well. The filter pack was added to the deep monitoring well's annular space to a level approximately two feet above the screened interval.

While slowly adding the filter pack, the HSAs were incrementally withdrawn to allow the filter pack material to drop out of the bottom of the HSA and to prevent the formation from collapsing around the well screen. An eleven to twelve foot thick bentonite chip seal was placed above the filter pack and hydrated with distilled water. Due to the relative shallowness of the groundwater, grout was not necessary to complete the monitoring wells. The bentonite seal was allowed to hydrate a minimum of four hours prior to placing the concrete apron around each monitoring well. The exact dimensions of the annular space materials were adjusted in the field on a well by well basis.

Above grade protective covers (4 inch diameter by 7 feet long) were provided for monitoring wells MW-1A and MW-5A. Monitoring well MW-7A was completed with a flush mounted aluminum cover per IDOT's requirements. Each deep monitoring well was provided with an expandable locking cap. Keyed alike Masterlock padlocks were provided for each monitoring well. A magnet was placed inside MW-7A's above grade annular space for ease of locating the flush mounted monitoring well during the winter time.

Shallow Monitoring Wells

Heaving sands were not a problem for shallow monitoring wells MW-8, MW-9, and MW-11. Therefore, the well strings were placed at the bottom of the drilled borehole. The silica sand filter pack was placed in the annular space to minimize intrusion of fine-grained sediment into the monitoring well.

While slowly adding the filter pack, the HSAs were incrementally withdrawn to allow the filter pack material to drop out of the bottom of the HSA and to prevent the formation from collapsing around the well screen. The filter pack was added to the shallow monitoring well's annular space to a height approximately one-half foot above the screened interval. Due to the shallow nature of the groundwater, it was more important to have an effective surface seal to impede the infiltration of surface water. A 1.5 foot thick bentonite seal was placed above the filter pack and hydrated with distilled water. Due to the relative shallowness of the monitoring wells, grout was not necessary to complete the monitoring wells. The bentonite seal was allowed to hydrate a minimum of four hours prior to placing the concrete apron around each monitoring well. The

exact dimensions of the annular space materials were adjusted in the field on a well by well basis.

Above grade protective covers were provided for the shallow monitoring wells. For MW-8 and MW-9, the standard 4 inch diameter by 7 feet long steel protective covers were cut down to approximately 4 feet in length so as not to interrupt the shallow groundwater flow into the well screens. The remaining monitoring wells installed at the Waukegan Generating Station were completed with flush mount well boxes that were cemented in place.

Organic clay soil cuttings from monitoring well MW-9 adhered to the interior of the HSAs. To avoid bridging of annular space materials, the drillers placed approximately 50 gallons of potable water inside the HSAs to wash out the organic clay.

Heaving sands were encountered during the installation of MW-10, MW-12, and MW-13. Minimal amounts of potable water were added to each of these wells during both drilling and installation. The approximate volumes of water added during installation was 5 gallons, 20 gallons, and 50 gallons, respectively.

Monitoring well construction diagrams are included in Appendix B. The following information was recorded on the geologic/monitoring well construction diagrams.

- General information including the drilling contractor, well number, well site, date
 the well construction was initiated and finished, and the name of the driller and
 supervising geologist or engineer.
- Specific information including the drilling method, borehole diameter, type and diameter of the protective casing, riser pipe and well screen, type of annular backfill, annular seal and filter pack, and depths to the top of the annular seal and filter pack and total well and boring depth.

Typical well construction details of the above depth measurements were made in the field using a weighted tape. Measurements are accurate to within 0.01 feet.

2.2.4 Monitoring Well Development Procedures

Well development allows for the free flow of water through the disturbed formation into the filter pack and well screen. Prior to well development and purging, all equipment were decontaminated in accordance to the procedures outlined in the Field Sampling Plan, Section 2.2.9. Well development and purging was accomplished by manually bailing using a three-foot long disposable Teflon™ bailer, or a five foot long PVC bailer. Development was not conducted until the bentonite pellet seal in each well was allowed to set for a minimum of 24 hours. Immediately before sampling, the wells were purged a minimum of three well volumes. Purging the monitoring well allows for a representative sample to be collected from the aquifer.

Each well volume was determined by measuring the static water level in the well with an electronic interface probe (IP) to the nearest 0.01 foot. The static water level from the top of the well casing was subtracted from the total depth of the well from the top of the well casing to determine the height of the water column in the well. The height of the column multiplied by the area of the well equalled one well volume.

Monitoring wells were considered developed after development water was relatively sediment free, and field parameters (pH, temperature, and specific conductance) stabilized to within 10 percent. Calibration, operation, and maintenance procedures for the pH, specific conductivity, and temperature meter is detailed in the approved QAPP. Field parameter readings were collected after each well volume was removed. The parameters were considered stabilized when three successive readings were within 10 percent. Typically, three to five well volumes were removed from each monitoring well during development. Well development/well purging field parameter data tables are illustrated in Appendix C.

Monitoring well MW-9 was developed by using a Grundfos submersible pump. The submersible pump was raised and lowered within the monitoring well and utilized like a surge block. Approximately 50 gallons of water was removed from this monitoring well. This amount equalled the total volume of potable water placed in the monitoring well by the drilling subcontractor to remove the organic clay from the inside of the hollow stem augers. For the monitoring wells that encountered heaving sands during well installation, the amount of water added was removed prior to initiation of the development process. An additional three to five well volumes of water were then removed as part of the well development process.

All development water was containerized in 55-gallon drums.

2.2.5 Static Water Level Measurements

The static water levels were measured in all of the monitoring wells installed at the site. Water levels were measured with an electronic interface probe (IP). Measurements were collected by lowering the probe into the well until the instrument emitted an audible tone. Pepth to water from the top of the stainless steel well riser was measured to the nearest 0.01 ft. Total depth of each monitoring well was determined by lowering the IP to the bottom of the well and sounding for total depth. All water level measurements, date and time, instrument used, and field personnel were recorded in a bound field logbook. The electronic interface probe was decontaminated prior to and after a reading was collected as described in Section 2.2.11. Static water level measurements are included in Appendix D.

2.2.6 Groundwater Sampling

In order to assess groundwater quality at the former tannery site, a second round of groundwater samples was collected from the seven monitoring wells installed by M&E during Phase I of the RI. The analytical analyte list was reduced to those parameters which were detected during

Phase I activities. These wells were sampled for Level III TCL SVOCs, TCL P/PCBs, total and dissolved RCRA Metals, hexavalent chromium, TSS, TDS, and Turbidity.

An initial round of groundwater samples was collected from the five monitoring wells installed during Phase II activities. These samples were analyzed for Level IV CLP total and dissolved TAL parameters, TCL VOCs, TCL SVOCs, TCL P/PCBs, hexavalent chromium, TSS, TDS, and Turbidity in accordance with the QAPP and the Illinois Pre-Notice Program Analytical Quality Assurance Plan. Analysis for TSS, TDS, and turbidity were performed in accordance with the methodologies described in Methods for Chemical Analysis of Water and Wasies, EPA 600/4-79-020: TDS - 160.1, TSS -160.2, and Turbidity - 180.1.

Table 2-2 illustrates the sample identification number and its corresponding laboratory analysis.

The four monitoring wells installed at the Waukegan Generating Station were sampled for Level III total and dissolved RCRA metals. In addition, MW-11 and MW-12 were analyzed for TSS, TDS, and TOC.

The monitoring wells were purged by manual bailing using a disposable Teflon™ bailer. During purging, the pH, specific conductivity, temperature, color, odor, and relative turbidity of the groundwater were recorded. Measurements were made after each well volume was purged and were recorded in the field log book. Groundwater samples were collected with disposable Teflon™ bailers.

Sample containers were filled directly from the bailer. Sample containers for VOCs were filled first, followed by containers for semi-volatile organic compounds, pesticides and PCBs, and metals analyses. The samples collected for dissolved metal analysis were first collected in a one-liter plastic bottle. The sample was then filtered at the well site using a disposable 0.45 micron positive pressure filter. Subsequent to filtering, the water was transferred to other sample bottles for dissolved metal analysis. The sample bottles were pre-preserved by the laboratory. The metal samples were preserved with nitric acid (HNO_i) to a pH < 2. The aliquot for cyanide analysis was preserved with sodium hydroxide (NaOH) to a pH of > 12. After sample collection, all samples were placed directly into a sample cooler where a temperature of 4°C was maintained. Chain of custody procedures for the samples are discussed in Section 2.3.

2.2.7 Surface and Subsurface Soil Sampling

2.2.7.1 Purpose

In order to assess the presence, extent, and magnitude of subsurface impact, additional sample collection occurred. Eighty-one soil samples were collected and analyzed from 51 sample locations. From the results of the Phase I investigation, the trends identified indicate that the highest constituent concentrations were located at the surface (0 - 2 feet below grade) and decreased with depth.

	Table 2-2 Soll/Sediment/Groundwater Sample Identification Number and Corresponding Analysis			
Surface Soil Samples (51):	SB-44A, SB-55A, SB-46A, SB-47A, SB-41A, SB-49A, SB-50A, SB-51A, SB-52A, SB-53A, SB-54A, SB-55A, SB-56A, SB-57A, SB-54A SB-59A, SB-60A, SB-61A, SB-62A, SB-61A, SB-64A/CA, SB-65A, SB-66A/CA, SB-67A/CA, SB-69A/CA, SB-70A, SB-71A, SB-72A SB-73A, SB-74A, SB-75A, SB-76A, SB-77A, SB-78A, SB-79A, SB-80A, SB-81A, SB-82A, SB-83A, SB-84A, SB-83A, SB-86A, SB-100 SB-101, SB-102, SB-103, SB-104, SB-105, SB-106, SB-107			
Subserface Soil Samples (30):	SB-44B, SB-45B, SB-46B, SB-47B, SB-44B, SB-49B, SB-50B, SB-51B, SB-51B, SB-59B, SB-60B, SB-62B, SB-64CB, SB-67B/CB, SB-69B/CB, SB-71B, SB-73B, SB-77B, SB-77B, SB-79B, SB-80B, SB-81B, SB-82B, SB-83B, SB-84B, SB-83B, SB-86B, MW-1A, MW-5A MW-7A.			
Production Waste Samples (2):	PW-5. PW-9.			
Groundwater Samples (16):	MW-1, MW-1A, MW-2, MW-3, MW-4, MW-5, MW-5A, MW-6, MW-7, MW-7A, MW-8, MW-9, MW-10, MW-11, MW-12, MW-13.			
Wetland Samples (6):	WL-1, WL-2, WL-3, WL-5W-1, WL-5W-2, WL-5W-3,			
Perticide Samples (4):	Surface, Subsurface, and Sediment SB-52A, SB-53A, SB-53B, SB-54A.			
PCB Samples (15):	SB-35A, SB-36A, SB-57A, SB-58A, SB-38B, SB-59A, SB-60A, SB-100A, SB-101A, SB-102A, SB-103A, SB-104A, SB-105A, SB-105A, SB-106A			
PNA Samples (4):	SB-56A, SB-38A, SB-58B, SB-60A.			
Diozin/Puran Samples (2):	SB-55A, SB-59A.			
Asbestos samples (2):	AS-1, AS-2.			
Arsenic Specific Samples (36):	SB-44A, SB-44B, SB-45A, SB-45B, SB-46A, SB-46B, SB-47A, SB-47B, SB-48A, SB-48B, SB-49A, SB-49B, SB-50A, SB-50B, SB-51A, SB-51B, SB-77A, SB-77B, SB-77A, SB-78B, SB-77A, SB-79B, SB-79B, SB-80A, SB-80B, SB-81A, SB-81B, SB-82A, SB-82B, SB-82B, SB-83A, SB-83B, SB-84A, SB-84B, SB-85B, SB-85B, SB-86A, SB-86B, SB-86B			
AVS/SEM Samples - (3):	WL-1, WL-2, WL-3.			
TAL Metale (3):	MW-1A, MW-5A, MW-7A.			
TCL SVOC (18):	MW-1A, MW-5A, MW-7A, WL-1, WL-2, WL-J, SB-61A, SB-62A, SB-62B, SB-61A, SB-64A/CA, SB-65A, SB-66A/CA, SB-67A/CA, SB-67B/CB, SB-68A, SB-69A/CA, SB-69B/CB.			
TCL P/PCBs (IB):	MW-1A, MW-7A, MW-7A, WL-1, WL-2, WL-3, SB-61A, SB-62A, SB-62B, SB-63A, SB-64A/CA, SB-63A, SB-66A/CA, SB-67A/CA, SB-67B/CB, SB-64A, SB-69A/CA, SB-69B/CB.			
RCRA Meials (28):	WL-1, WL-2, WL-3, SB-60B. SB-61A, SB-62A, SB-62B, SB-63A, SB-64A/CA, SB-64CB, SB-65A, SB-66A/CA, SB-67A/CA, SB- 67B/CB, SB-63A, SB-69A/CA, SB-69B/CB, SB-70A, SB-71A, SB-71B, SB-72A, SB-73A, SB-73B, SB-74A, SB-73A, SB-76A, PW-S			

Soil/Sediment/Groundwater Sample Identification Number and Corresponding Analysis

Hexavelest Chromium (22): WL-1, WL-2, WL-3, 58-608, S8-64A/CA, S8-64CB, S8-66A/CA, S8-67B/CA, S8-67B/CB, S8-69A/CA, 58-69B/CB, S8-70A, S8-

71A, SB-71B, SB-72A, SB-73A, SB-73B, SB-74A, SB-75A, SB-76A, PW-5, PW-9.

SB-60B, SB-64A/CA, SB-64CB, SB-66A/CA, SB-67A/CA, 67B/CB, SB-69A/CA, SB-69B/CB, SB-70A, SB-71A, SB-71B, SB-72A, SB-TCLP Metals (19):

73A, 58-73B, 58-74A, 58-75A, 58-76A, PW-5, PW-9.

Surface Water/Groundwater

TSS (14): MW-1, MW-1A, MW-2, MW-3, MW-4, MW-5, MW-5A, MW-6, MW-7, MW-7A, MW-8, MW-9, MW-10, MW-11.

WL-1, WL-2, WL-3, WL-SW-1, WL-SW-2, WL-SW-3, MW-10, MW-11,

TDS (14): MW-1, MW-1A, MW-2, MW-3, MW-4, MW-5, MW-5A, MW-6, MW-7, MW-7A, MW-8, MW-9, MW-10, MW-11.

Turbidity (12): MW-1, MW-1A, MW-2, MW-3, MW-4, MW-5, MW-5A, MW-6, MW-7, MW-7A, MW-8, MW-9.

Cyanide (12): MW-1, MW-1A, MW-2, MW-3, MW-4, MW-5, MW-5A, MW-6, MW-7, MW-7A, MW-8, MW-9.

RCRA Metals (14); MW-1, MW-2, MW-3, MW-4, MW-5, MW-6, MW-7, MW-10, MW-11, MW-12, MW-13, WL-SW-1, WL-SW-2, WL-SW-3.

SVOC (12): MW-1, MW-2, MW-3, MW-4, MW-5, MW-6, MW-7, MW-8, MW-9, WLSW-1, WLSW-2, WLSW-3, MW-1A, MW-5A, MW-7A. Level IV SVOC (3):

P/PCB (12):

MW-1, MW-2, MW-3, MW-4, MW-5, MW-6, MW-7, MW-8, MW-9, WL-SW-1, WL-SW-2, WL-SW-3, MW-1A, MW-5A, MW-7A. Level IV P/PCB (3):

Level IV VOCs (5): MW-1A, MW-5A, MW-7A, MW-8, MW-9.

Level IV TAL (5): MW-1A, MW-5A, MW-7A, MW-8, MW-9.

Hexavalent Chromium (12): MW-1, MW-1A, MW-2, MW-3, MW-4, MW-5, MW-5A, MW-6, MW-7, MW-7A, MW-8, MW-9.

TOC (t):

re collected on February 2, 7, and 8, Isan 29, and August 31, 1995. ing samples were collected on January 31, and February 1, 1995. all buring samples were collected on January 12, 17, 18, and 31, 1995 and February 24, 1995. all well-must samples were collected on April 10, 1995.

Parameter specific soil sampling occurred in select areas in which elevated levels of specific constituents were identified in Phase I, to more fully define the lateral extent of impact. The following lists specific activities and their respective purpose.

Table 2-2 illustrates the sample identification number and its corresponding laboratory analysis.

2.2.7.1 (a) Site Classification

In order to determine the regulatory status of the site, both surface and subsurface soil samples were collected and analyzed for RCRA metals, TCLP metals and hexavalent chromium (Cr⁺). These analyses were intended to determine the status of the chromium at the former tannery for both regulatory and Risk Assessment purposes. Soil samples were collected from areas shown to have elevated chromium based upon Phase I results. Production waste sample locations PW-5 and PW-9 were analyzed for the same parameters.

2.2.7.1 (b) Parameter Specific Soil Sampling

In order to more accurately assess the presence and to define the vertical and lateral extent of certain constituents, parameter-specific and area-specific sampling was conducted.

Pesticides

In the vicinity of SB-39, a total of four surface soil samples were collected and analyzed for pesticides to determine the extent of pesticide impact in the area.

Polychlorinated Biphenyls, Polynuclear Aromatic Hydrocarbons, and Dioxin/Furans

In the southeastern section of Area III, elevated levels of PCBs were identified during Phase I activities. A total of 15 surface soil samples were collected. Additionally, of those soil samples, four samples were collected and analyzed for polynuclear aromatic hydrocarbons (PNAs) and two samples were collected and analyzed for dioxin/furans. These soil samples were analyzed for dioxin/furans due to concerns raised by the Community Advisory Group (CAG) members regarding alleged former on-site burning of PCBs.

Asbestos Sampling

To assess the presence of asbestos in dredge sand used as fill at the site, two soil samples were collected and analyzed for asbestos using Polarized Light Microscopy (PLM). These samples were collected due to concerns raised at the CAG meeting regarding asbestos deposited into Lake Michigan and being dredged with the sand and subsequently stored temporarily on site.

Arsenic

In the vicinity of MW-1 and SB-06, surface and subsurface soils were analyzed for arsenic only. The purpose of this sampling was to identify the possible source and/or the extent of arsenic in the subsurface. A total of 36 surface and subsurface soil samples were collected.

2.2.7.2 Locations and Rationale

Thirty subsurface samples were obtained from the unsaturated zone immediately above the Static Water Level (SWL) (located approximately 3 feet below grade) and 51 surface soil samples were obtained from the 51 soil boring locations. These surface and subsurface samples were collected in an effort to assess the presence and magnitude of impact to the surface and subsurface from former tannery operations. The surface samples were collected from the surface to a depth of approximately one foot below grade.

2.2.7.3 Surface and Subsurface Soil Sampling Methods and Equipment

Split spoon soil samples were collected from the shallow and deep monitoring wells. Soil samples were collected by using a split-spoon sampler (ASTM D1586) driven by a 140-pound hammer free-falling 30 inches. Once retrieved, the split spoon was opened and screened using a PID. A stainless steel knife was used during the screening process to part the sample in order to survey portions of the sample which were not disturbed by the split spoon sampler. The stainless steel knife was decontaminated between sample intervals as described in Section 2.2.11. Soil collected from the split-spoon sampler was transferred directly to the sample containers.

Due to the shallow nature of the groundwater table at the site, accessibility of the drill rig, and number of soil samples to collect, stainless steel hand augers were used to collect surface and subsurface soil samples. Soil samples collected with a hand auger were composite. A stainless steel spoon was used to transfer the soil from the hand auger into a stainless steel compositing bowl. Soil collected for VOC samples were immediately packed into the sample container and headspace was minimized. All samples were capped as quickly as feasible.

2.2.8 Surface Water and Sediment Sampling

2.2.8.1 Purpose

Wetland surface water and sediment samples were collected and analyzed to provide data for the Ecological Risk Assessment (ERA) and aid in determining if constituents have migrated off-site. In addition to the Level IV CLP TCL SVOC, TCL P/PCB, TOC and RCRA Metal analysis performed on both the surface water and sediment samples, the sediment samples were analyzed for both Hexavalent Chromium and Acid Volatile Sulfide and Selected Simultaneously

Extractable Metals (AVS/SEM)¹. The TOC Methodology will aid in the determination of the bioavailability of organic material. The SEM/AVS Methodology will determine the persistence of metal constituents in the wetland sediment. The surface water samples were also analyzed for total and dissolved RCRA metals.

2.2.8.2 Locations

A total of three sediment and three surface water samples were collected during Phase II RI activities. The samples were collected in areas where aquatic biota systems were established and thriving. Samples were not collected from an infertile ditch or an intermittent pond or stream. The three samples were collected in the wetland area immediately south of the southeastern site boundary.

2.2.8.3 Sampling Methods and Equipment

The following information presents the sampling method for collecting wetland surface water and sediment samples from the wetland area.

Due to the relatively shallow nature of the water in the wetland area, field personnel traversed the wetland to collect sediment samples. Upon reaching the desired location, field personnel were careful so as not to disturbed the sediment. Prior to collecting the sediment samples, surface water samples were collected by lowering the entire sampling bottle (in an upright position) below the water surface. The mouth of the sample container was approximately four to six inches below the water surface. Samples for TCL SVOC analyses were collected first, followed by TCL P/PCBs, TOC, and total and dissolved RCRA metals.

The sediment samples were collected by using a stainless steel hand auger. Each sample was collected from the upper 12 inches of the wetland sediment. Cattail root matter was encountered during boring. Soil samples collected with a hand auger were composited. A stainless steel spoon was used to transfer the soil from the hand auger into a stainless steel compositing tray. Samples collected for non-volatile organic compound analysis were transferred from the sampling device to a compositing tray then to their respective sample containers. All samples were capped as quickly as feasible and placed into iced coolers to preserve them at 4°C.

2.2.9 Waste Volume Estimation

To determine the gross volume of tannery waste material (tannery hides) present at the site, field personnel traversed the southwestern section of Area III and tested several areas with a hand auger to map the areas anticipated to contain tannery waste material, much like that of PW-5

AVS and SEM are operational definitions in the analysis of sulfide and associated metals (cadmium, copper, lead, moreoury, sichal, and zinc) in equatic sediments. The SEM/AVS ratio has been useful in predicting if a sediment is not scutely toxic (SEM/AVS ratio is less than one) or shows only potential for toxicity (SEM/AVS ratio is greater than one). This is because other binding phases, such as metal oxides and organic metter, may also bind metals thereby reducing their binavallability.

(bluish grey shavings). A site walk-through revealed that material is confined to an area approximately 15 feet long by four feet wide and three feet deep at the location of PW-5. The total volume of this material is estimated to be 180 cubic feet (6.66 cubic yards). No other location in the southwestern section of the site consisted of the production waste material.

2.2.10 Surveying Methodology

M&E conducted surveying at the former Griess-Pfleger Tannery to establish horizontal and vertical control. The horizontal survey located monitoring wells and soil borings. The vertical survey established top of riser (TOR) elevations of the monitoring wells and ground surface elevations.

The horizontal survey was conducted using a Global Positioning System (GPS) with an accuracy of \pm 5 - 10 millimeters. It was determined that due to the many obstructions at the site blocking the path between points (trees, shrubs), a GPS survey would be more efficient. The type of GPS survey method used was Rapid Static. For the Rapid Static survey, a temporary reference station was situated inside the entrance gate. Field personnel used roving receivers to record the locations of the sampling points. Computer software was used to process the data collected from the receivers and assigned North - East coordinates to the unknown points. The coordinates were used to create a map showing monitoring wells, soil borings, and sediment samples.

The vertical survey of the monitoring well riser locations was established with an automatic level. A reference benchmark was established on MW-4 whose elevations was known from Phase I activities. The riser elevations were taken on the north side of the risers.

2.2.11 Decontamination Procedures

Potable water for decontamination was obtained from Commonwealth Edison's Waukegan Station. This facility obtains its water from the City of Waukegan. Analytical data for the City of Waukegan potable water supply source was obtained during Phase I activities.

All drill rigs, drilling equipment, split spoons, sampling spoons, hand augers, well casing and screen, well development equipment, and water level measurement equipment were decontaminated upon arrival to the site, between each monitoring well or soil boring location and prior to departure. A portable wash tub was used to collect decontamination rinsate water. The rinse water was transferred from the wash tub to labeled 55-gallon drums and staged in a central area. The drums were labelled as to the contents of each drum and date.

Decontamination of the drill rig, downhole tools, well screen and casing, and well development equipment consisted of high pressure steam washing. Any visible residue after steam cleaning was scrubbed with a brush and a solution of phosphate-free laboratory grade detergent (Alconox) and potable water followed by a final steam cleaning. All decontaminated equipment was allowed to air dry prior to use. Well materials were wrapped in new plastic for transport to the well location.

Split spoons were decontaminated between each new well location following the procedure described above. However, during drilling, the split spoons were decontaminated between each sampling interval with a brush and a solution of phosphate-free laboratory grade detergent (Alconox) and potable water followed by a final rinse with distilled water.

New disposable Teflon^{to} bailers were used to develop and collect groundwater samples from each of the monitoring wells. Therefore, decontamination of this equipment was not needed.

The electronic water level indicator, pH, specific conductivity, and temperature probes were decontaminated at the well site between measurements. The submersible pump was decontaminated with a series of rinses with potable water, dilute nitric acid (10%) solution, and a final double rinse with distilled water. The equipment was allowed to air dry prior to being wrapped in new plastic bags for transport.

2.3 SAMPLE PACKAGING AND SHIPMENT

After collection, all samples were labelled, sealed, packaged, and delivered in person to IEA Laboratory, Inc. in Schaumburg, Illinois. Level III QA/QC samples were analyzed at the Schaumburg location whereas the Level IV CLP QA/QC samples were shipped by IEA Schaumburg to IEA's laboratory in Cary, North Carolina, where they were analyzed. The samples were shipped via next day courier. Temperature in the coolers was maintained at 4°C through the use of ice sealed in plastic bags.

A chain of custody record was completed and accompanied samples during shipment to the laboratory. The chain of custody record was sealed in plastic for protection and taped inside the lid of the cooler. A copy of the chain of custody record was retained by the sampling team. All records included: (1) sample numbers; (2) date and time of collection; (3) locations where samples were collected; (4) type of sample: grab or composite; (5) analytical parameters requested; (6) names and signatures of samplers; and (7) names of persons involved in the chain of possession from time of collection to receipt at the laboratory.

Because each cooler was hand-delivered to the laboratory, custody seals were not used on the coolers.

2.4 INVESTIGATIVE DERIVED WASTE

The fate of drummed investigative derived waste material will be based upon the proper characterization of materials (decontamination/purge/well development water and soil cuttings) currently stored in drums on site. Material will be transported off-site under Commonwealth Edison's direction.

SECTION 3.0 RESULTS AND DISCUSSION OF ENVIRONMENTAL INVESTIGATION

3.1 SITE GEOLOGY

The site geology is characterized as consisting of surfical unconsolidated material, made-land soils. The subsurface soils, typically described, ranged from pale yellow to black, moist to wet, loose to dense, poorly graded to well graded silty sand to sand. The silty sands are found as surfical deposits but grade to sands at depth. Trace amounts of fine subrounded gravels are present throughout the strata. The sand grain particles range from fine to coarse. Clay was present in the deeper soil borings (EB-1 and MW-7A). The thickness of the clay lens ranged from 4 - 9 inches and is discontinuous. Some boring/monitoring wells (MW-6, MW-4, MW-7A, and MW-9) have spongy organic matter (peat) ranging from the surface to 13 feet below grade. Admixed with the sands are coal, tannery waste material (animal hair), organic material (roots, vegetation), gravel, slag, and asphalt. Tannery wastes can be found up to a depth of 10 feet but, the asphalt and gravel only extend to a depth of five feet below grade. The sand extends to the base of all the borings. The geologic logs of the soil boring/monitoring wells are included in Appendix A. Figures 3-1 and 3-2 illustrate the north-south and west-east cross sections of the site, respectively.

3.2 SITE HYDROGEOLOGY

Groundwater at this site is considered a Class I Potable Resource Groundwater by definition of Title 35 of the Illinois Administrative Code (IAC), Subtitle C, Part 620, Subpart B, Groundwater Classification (e.g. groundwater which is presently being used or has the potential for being put to conventional use).

Both the shallow and the deep monitoring wells installed at the former tannery indicate that groundwater is flowing under unconfined conditions. Deeper soil borings indicate the presence of a clay lens but it was not thick enough (at least 3 to 5 feet thick) nor continuous across the site to segregate the water bearing unit into two separate units.

Using the most recent groundwater elevation data (February 24, 1995) the static water level (SWL) ranged from approximately 93.24 feet (MW-5) to 95.62 feet (MW-7A). The tabulated water elevation data is illustrated in Appendix D. The SWL measurements were collected from the ten on-site monitoring wells and two off-site monitoring wells, all of which are screened in the unconsolidated sediments. The saturated thickness is determined to be at least twenty-three feet. It is anticipated that the saturated thickness of the aquifer is greater because a confining layer was not encountered.

Shallow groundwater flow is toward the east under a gradient of 0.0014 feet/foot. This calculation is identical to the shallow gradient calculated for Phase I activities. The deeper groundwater flow directions is toward the east under a hydraulic gradient of 0.0017 feet/foot.

eardioveried

The hydraulic conductivity testing for the newly installed monitoring wells was not done because material encountered during Phase II activities is similar to that which was discovered during Phase I activities. Because no thick and continuous clay layer was present, it can be interred that hydraulic communication exists between the upper and lower water bearing units.

Figures 3-3 and 3-4 are groundwater contour maps illustrating the flow direction from the shallow and deep groundwater monitoring wells, respectively. These data were collected on February 24, 1995 and confirm that the shallow and deep water is one aquifer.

3.3 DETERMINATION OF CHROMIUM WASTE EXCLUSION

Based on 35 IAC 721.104 (Exclusion from the Definition of Hazardous Wastes), specific solid chromium wastes are not hazardous if certain conditions are met. These conditions, as shown below, are met by applicable site wastes (PW-5 and PW-9).

A) The waste fails the toxicity characteristic test because chromium is present and does not fail the test for any other constituents.

During the Phase 1 investigation, two samples of production waste (PW-5 and PW-9) had constituent concentrations that exceeded the Toxicity Characteristics Leaching Procedure (TCLP) regulatory standards of 5.0 mg/l. Samples PW-5 and PW-9 had a chromium concentrations of 8.06 mg/l and 24.2 mg/l, respectively. The remaining constituents were below the TCLP regulatory limits. As a result, these samples meet the requirements of this subpart of the exclusion

During Phase II investigation, production waste samples PW-5 and PW-9 were resampled. The respective chromium concentrations were 0.6 mg/l and 14 mg/l. The remaining constituents were below the TCLP regulatory limits. As a result, these samples meet the requirements of this subpart of the exclusion.

B) The chromium in the waste is exclusively (or nearly exclusively) trivalent chromium.

Phase II chromium speciation results indicate that chromium used during the tanning process consists of 99.9% trivalent chromium.

C) The waste is generated from an industrial process which uses trivalent chromium exclusively (or nearly exclusively) and the process does not generate hexavalent chromium.

According the U.S. EPA, chromium in the tanning process must be in the trivalent form (and in an acid medium) to perform the tanning (U.S. EPA 1974).

According to available documents and analytical data, trivalent chromium was used in the tanning process. A typical chrome tanning process consists of nine separate steps.

The decidibilities are after a secure of the first of a second

In this process, sulfuric acid and sodium chloride are used to pickle the hides. The pickled hides are immersed in a acidified solution of sodium dichromate. Therefore the chrome was present in an acid medium.

D) The waste is typically and frequently managed in non-oxidizing environments.

In soils, hexavalent chromium may be quickly converted to trivalent chromium which is relatively immobile and the more benign of the two species. Trivalent chromium may be oxidized to hexavalent chromium, but this appears to requires the presence of manganese oxides (U.S. EPA 1980).

A 5-year U.S. EPA study analyzed the land application of trivalent chromium-containing tannery process sludges and found that the chromium remained primarily in the topsoil. Trace amounts of chromium were found in surface water runoff from the sludge-loaded soil, but the chromium transport was associated with movement of soil particles. Oxidation of the chromium to hexavalent chromium was not detected (U.S. EPA 1986).

The tannery waste deposits (PW-5 and PW-9) at the site contained trivalent chromium and no detectable hexavalent chromium.

- E) Specific examples of chromium wastes that match these characteristics are given. However, if the water is one of these types, it must still pass TCLP toxicity analysis for constituents beside chromium (must be below the regulatory limits) and must pass hazardous waste characteristic analysis (ignitability, corrosivity, etc.) Among the specific wastes listed are:
 - Chrome (blue) trimmings generated by hair pulp/chrome tan/retan/wet finish or hair save/chrome tan/retan/wet finish or retan/wet finish or no beamhouse or through-the-blue or shearling tanning processes.
 - ii) Chrome (blue) shavings generated by pulp/chrome tan/retan/wet finish or hair save/chrome tan/retan/wet finish or retan/wet finish or no beamhouse or throughthe-blue or shearling tanning processes.
 - iii) Buffing dust generated by hair pulp/chrome tan/retan/wet finish or hair save/chrome tan/retan/wet finish or retan/wet finish or not beamhouse or throughthe-blue tanning processes.
 - iv) Sewer screening generated by hair pulp/chrome tan/retan/wet finish or hair save/chrome tan/retan/wet finish or retan/wet finish or not beamhouse or throughthe-blue tanning processes.

CANAL PROPERTY LANG.

99130,000767

- v) Wastewater treatment sludges generated by hair pulp/chrome tan/retan/wet finish or hair save/chrome tan/retan/wet finish or retan/wet finish or not beamhouse or through-the-blue tanning processes.
- vi) Wastewater treatment sludges generated by hair pulp/chrome tan/retan/wet finish or hair save/chrome tan/retan/wet finish or through-the-blue tanning processes.
- vii) Waste scrap leather from the leather tanning, shoe manufacturing, and other leather product manufacturing industries.
- viii) Wastewater treatment sludges from production of titanium dioxide pigment using chromium-bearing ores by the chloride process.

The examples provided in the above subsections include applicable site wastes (PW-5 and PW-9).

SECTION 4.0 NATURE AND EXTENT OF CONTAMINATION

4.1 GROUNDWATER

In addition to the seven existing shallow monitoring wells, five additional monitoring wells were installed as part of the Phase II investigation, two shallow and three deep. All twelve of the wells were sampled in February, 1995.

The five new wells were sampled for full TAL and TCL, with analysis conducted under IEPA Level IV QA/QC. The seven existing wells were sampled for TCL SVOC, TCL P/PGBs and the RCRA metals. These analyses were conducted under IEPA Level III QA/QC. All metals analyses were conducted on both filtered and unfiltered samples. All samples collected were analyzed by IEA Laboratories. The Level III samples were analyzed at IEA's Schaumburg, Illinois facility. The Level IV samples were analyzed at IEA's Cary, North Carolina facility.

In addition to the parameters identified above, all of the monitoring wells were sampled for hexavalent chromium, total suspended solids (TSS), total dissolved solids (TDS), and turbidity. These analyses were all conducted under Level III QA/QC.

The determination that elevated levels of chemical constituents were present in the groundwater was made by comparing the analytical results obtained with the constituent concentration limits found for Class I Groundwater in IAC Title 35, Subpart D, Section 620.410.

Analytical results from the Phase II investigation indicated that groundwater containing arsenic exceeding the Illinois Class I Standard has migrated to the eastern edge of the tannery property boundary. Subsequently, two additional monitoring wells were installed as part of the Phase IIA investigation at Commonwealth Edison's Waukegan Generating Station. Analytical results from these two monitoring wells confirmed that groundwater containing arsenic had migrated under the EI&E railroad tracks and onto Commonwealth Edison's Waukegan Generating Station property. A Phase IIB investigation was undertaken that utilized a Geoprobe^R unit to collect groundwater samples to define the extent of the arsenic plume. Using the Geoprobe^R collected data, two additional permanent monitoring wells were installed to confirm the arsenic plume delineation. At the present time, the delineation indicates that the arsenic plume has migrated 400 feet from the former Tannery.

The groundwater samples collected from the Phase IIA and IIB monitoring wells were analyzed the total and dissolved RCRA metals under Level III QA/QC. The 14 Geoprobe^R samples were analyzed for arsenic under Level III QA/QC.

4.1.1 Volatile Organic Compounds

Volatile organic compounds were not detected in any of the newly installed Phase II monitoring wells, MW-1A, MW-5A, MW-7A, MW-8, and MW-9. Samples for VOCs were not collected

from the existing monitoring wells because the Phase I investigation showed a lack of VOCs in the groundwater.

4.1.2 Base-Neutral/Acid Extractable Compounds

Two SVOCs were detected in the groundwater samples collected, di-n-butyl phthalate and bis(2-ethylhexyl) phthalate. Both of these compounds were detected in five of the twelve samples collected. All of the di-n-butyl phthalate were qualified "JB", indicating that the laboratory method blank also contained this analyte, which indicates that this compound is present due to laboratory contamination.

The bis(2-ethylhexyl) phthalate concentrations ranged from an estimated concentration of 2 ppb in MW-1A and MW-7A to an estimated concentration of 5 ppb in MW-5A. Bis(2-ethylhexyl) phthalate, as well as most of the phthalate family, are common laboratory contaminants. The presence of both of these compounds is likely due to laboratory contamination.

4.1.3 Pesticides/PCBs

Pesticides and PCBs (P/PCBs) were not detected in any of the groundwater samples collected.

4.1.4 Inorganics

As expected, inorganic analytes were detected in all samples collected. Many of these analytes are considered naturally occurring in groundwater at varying levels. Additionally, a significant decrease in analyte concentrations is observed when comparing the total (unfiltered) versus dissolved (filtered) concentrations. This concentration decrease from the total to the dissolved value is a result of the acid preservative dissolving the constituents from the suspended solids in the sample containers.

The Class I Groundwater Standard for chromium, as found in IAC Title 35, Subpart D, Section 620.410(a), is 100 μ g/l. All of the filtered groundwater samples collected showed dissolved chromium concentrations at less than 40 percent of the Class I standard. The dissolved chromium concentrations ranged from 1 μ g/l in MW-4 to 40 μ g/l in MW-2.

Chromium was detected above Class I standards in the unfiltered samples from 8 of the twelve wells sampled. Monitoring wells MW-4, MW-5, MW-7, and MW-7A had unfiltered total chromium concentrations below the 100 μ g/l standard. The highest total concentration identified was 6,670 μ g/l in MW-8. The corresponding dissolved concentration was 10.1 μ g/l, a reduction of greater than 600 times the total amount. This data shows that the majority of the chromium is bound to soil particles greater than 0.45 μ m in size, and not readily desorbed into the aquifer. While reductions of this magnitude were not always realized, this reduction is indicative of general trends throughout the site. The upgradient wells, MW-7 and MW-7A, contained total chromium at concentrations of 19 and 11 μ g/l and dissolved chromium at concentrations of 8.2 and 1 μ g/l, respectively.

The filtered samples collected during Phase II show chromium results similar to those for the unfiltered samples collected during Phase I of the investigation. The reason for the increase in the chromium concentration in the Phase II unfiltered samples is unknown, but may be due to the geologic conditions encountered, and the locations of the new monitoring wells.

All of the Phase I and Phase II unfiltered monitoring well samples were analyzed for hexavalent chromium. The hexavalent chromium values were all less than 10 μ g/l, with the exception of MW-4 which contained 41 μ g/l. The total chromium concentration in this sample was 53 μ g/l. The dissolved chromium concentration in this sample was less than 10 μ g/l. The total chromium concentration for both the filtered and unfiltered samples are both below the Class I standard of 100 μ g/l.

Total arsenic concentrations exceeded the Class I standard in four of the monitoring wells; MW-1, MW-2, MW-8, MW-9, MW-10, and MW-11. The highest observed concentration was 2,100 μ g/l in MW-1.

Dissolved arsenic concentrations exceeded the Class I standard of 50 μ g/l in five monitoring wells; MW-1, MW-8, MW-9, MW-10, and MW-11. The arsenic concentrations in these five wells ranged from 83 μ g/l in MW-10 to 1,100 μ g/l in MW-1. It should be noted that MW-2 contained 49 μ g/l, very close to the Class I standard of 50 μ g/l. The arsenic concentration in MW-1 has decreased since the last Phase I concentration of 6,470 μ g/l, while the concentration in MW-2 has increased from 28 μ g/l.

It has not been confirmed that arsenic usage occurred at this site. However, research is being conducted to verify the type of tanning processes that were utilized on-site.

Total cadmium was identified in four wells above the Class I standard of 5 μ g/I; MW-1, MW-3, MW-4, and MW-5. The maximum total cadmium concentration was 18 μ g/I. Two of these four wells, MW-4 and MW-5, exhibited dissolved cadmium concentrations either at or above the Class I standard at 5 and 15 μ g/I, respectively. Total cadmium was not identified in any of the groundwater samples collected during the Phase I investigation.

Total mercury concentrations in MW-2 and MW-8 were found to be above the Class I standard of 2 μ g/I, at 3 and 4.6 μ g/I, respectively. All dissolved mercury values were below the Class I standard. All total mercury values from the Phase I samples were below the Class I standard.

Total lead values exceeded the Class I standard of 7.5 μ g/l in all monitoring wells with the exception of MW-1, MW-3, MW-4, MW-5, MW-6, and MW-12. Concentrations ranged from 12.4 μ g/l in MW-5A to 168 μ g/l in MW-8. All dissolved lead concentrations were below the Class I standard.

For all of the Phase II wells, both the filtered and unfiltered samples exceeded the Class I standard of 150 μ g/l for manganese. The highest concentration was found in MW-5A at 879 μ g/l. These results are consistent with the Phase I groundwater samples which all exceeded the

Class I standard. Additionally, MW-7A, the upgradient well, contained one of the higher manganese concentrations.

Total iron concentrations exceeded the Class I standard of 5,000 μ g/l in all of the newly installed wells. The highest concentration was found in MW-5A at 19,100 μ g/l. Dissolved iron exceeded the Class I standard in only one well, MW-5A, at 2 concentration of 15,100 μ g/l. Four of the seven Phase I monitoring wells exceeded the Class I standard during Phase I sampling activities. These elevated concentrations are the result of the high TDS values which are considered naturally occurring.

All monitoring wells, with the exception of MW-7A, exceeded the total dissolved solids (TDS) Class I Standard of 1,200,000 μ g/l. The TDS result for MW-7A was 930,000 μ g/l. These results correspond with the high TDS values obtained during the Phase I investigation. These results show that the high TDS values are naturally occurring.

Overall, the total chromium concentrations from the Phase I sampling event are comparable to those obtained for the filtered Phase II samples. The arsenic concentration in MW-1 has decreased by a factor of three from Phase I to Phase II. The arsenic concentration in MW-2 showed an increase over the Phase I result. At the present time, the arsenic plume, as shown by the results obtained for MW-9, has reached the eastern edge of the site boundary. The Phase IIA monitoring wells, MW-10, and MW-11 show that the arsenic plume has migrated under the EJ&E railroad tracks and onto Commonwealth Edison's Waukegan Generating Station property. Based upon the results of the Phase IIB investigation, the arsenic plume appears to have migrated a distance of approximately 400 feet to the east of the site. Total lead, cadmium and mercury levels have shown an increase in concentration between Phase I and Phase II to above the Class I standards. Phase II dissolved concentration for lead and mercury were below the Class I standard. The reason for the increase in these constituents is unknown, but may be attributable to accumulation of very fine soil particles (less than $0.45\mu m$ in diameter) in the wells.

Analytical results for all groundwater samples are summarized in Table 4-1.

4.2 SURFACE AND SUBSURFACE SOILS

Several areas of concern were identified during the Phase I investigation. The concerns associated with the different areas were caused by either a specific compound, for example arsenic in surface/subsurface soils, or a specific class of compounds, such as PCBs. As a result, additional sampling was conducted in these areas of concern for the specific compound or class of compound. This Section summarizes the results of the investigations into the specific areas of concern.

In order to supplement the Phase I investigation, 51 surface soil, 30 subsurface soil, three sediment and two production waste samples were collected. The analytical parameters varied dependant upon the purpose of the sample. Samples were collected to more fully define the extent of certain constituents, characterize the nature of the constituents present, collect sufficient

on the reserve the second

data to conduct both a Baseline Risk Assessment (BRA) and Ecological Risk Assessment (ERA), and to determine if constituents had migrated beyond the site boundary.

In order to combine the Phase I and Phase II investigations, the analytical data summaries presented below incorporate results from both phases of the investigation. Non-parameter specific data summaries are presented in Table 4-2. All of the Figures presented incorporate data collected during both the Phase I and Phase II investigations.

4.2.1 Pesticides at Location SB-39

During the Phase I investigation, elevated levels of DDT, DDD, and DDE were identified in the surface soils at sampling location SB-39. The total pesticide concentration at this location was $48,000 \,\mu\text{g/kg}$. In order to determine the extent of the surfical pesticides in this area, three surface and one subsurface samples were collected. Analytical results for these four samples indicate that there is not a significant amount of pesticides in these samples, which shows that the elevated pesticide level in the surface soil at this location is an isolated occurrence.

Analytical results for these samples are summarized in Table 4-2. Surfical and subsurfical total pesticide distribution maps are contained in Figures 4-1 and 4-2, respectively.

4.2.2 Polychlorinated Biphenyls and Polynuclear Aromatic Hydrocarbons

During the Phase I investigation, elevated levels of PCBs and PNAs were observed in Area III. In order to more fully define the extent of the PCBs and PNAs, fifteen additional Phase II soil samples were collected, 14 surfical and one subsurfical. All 15 samples were analyzed for PCBs. In addition, four of the samples were analyzed for PNAs. PCBs were detected in all fifteen of the Phase II samples collected. The PCB concentrations ranged from 220 μ g/kg at SB-100A to 116,800 μ g/kg at SB-56A.

Similar to the Phase I investigation, the predominant Aroclor identified was 1248 coupled with lesser quantities of Aroclor 1254. Small amounts of Aroclor 1260 were found in several of the samples collected. The source of the PCBs at these locations is unknown.

Analytical results for the samples collected during the Phase II investigation are summarized in Table 4-2. Surfical and subsurfical total PCB distribution maps are contained in Figures 4-3 and 4-4, respectively.

4.2.3 Polychlorinated Dibenzo Dioxins/Furans

In response to requests from the Community Action Group (CAG), two of the soil samples collected were analyzed for polychlorinated dibenzo dioxins/furans (PCDD/PCDF). One sample was located near the production waste disposal area and one located in an area exhibiting elevated Phase I PCB concentrations. All of the PCDD and PCDF compounds were identified in SB-55A, and 14 out of the 17 compounds were identified in sample SB-59A. Concentrations

the above more about the second limited

of the various compounds ranged from 5.7 parts per trillion (ppt) of 2,3,4,7,8-hexachlorodibenzofuran in sample SB-59A to 140,000 ppt 1,2,3,4,6,7,8-heptachlorodibenzodioxin in sample SB-59A.

The U.S. EPA has a method for calculating the relative risk to humans of the various PCDD/PCDF compounds. These relative risks are based upon the toxicity of 2,3,7,8-tetrachlorodibenzodioxin (2,3,7,8-TCDD), believed to be the most toxic PCDD/PCDF. With the relative risk factors applied, both of these samples are below the 1 part per billion (ppb) 2,3,7,8-TCDD risk based cleanup threshold. Analytical results for the dioxin/furans are summarized in Table 4-3.

4.2.4 Asbestos

Based upon information supplied by the CAG and others, sand dredged from Lake Michigan and temporarily stored at the former tannery may have been impacted with asbestos. In order to determine the presence of asbestos fibers, two sand samples were submitted for asbestos fiber analysis by polarized light microscopy (PLM). Asbestos fibers were not detected in either of the samples. Analytical results for the asbestos samples are summarized in Table 4-4.

4.2.5 Arsenic Specific Sampling

In order to identify the possible source of the arsenic detected in MW-1 during the Phase I investigation, 36 soil samples; 18 surfical and 18 subsurfical, were collected and analyzed for arsenic only. Slightly elevated arsenic levels were identified in the surface and subsurface soils in the vicinity of MW-1. The arsenic levels identified ranged from 1.2 mg/kg in SB-45B to 10,000 mg/kg in SB-50B. While no distinct arsenic source was identified, the elevated arsenic levels are present throughout Area II. In addition, the two samples exhibiting the highest arsenic concentrations, SB-44A at 4,600 mg/kg and SB-50B at 10,000 mg/kg, were tested for leachable arsenic. The results for both of these samples did not exceed the 5 mg/l regulatory limit. Analytical results from these arsenic samples are summarized in Table 4-5.

4.2.6 Chromlum Speciation

In order to more fully characterize site conditions, the valence state of the chromium was determined. Chromium generally occurs in two valance states, the more benign and less mobile trivalent state (Cr⁺³) and the more toxic, more mobile hexavalent state (Cr⁺⁶). A total of 22 soil and sediment samples were collected and analyzed for Cr⁺³ and Cr⁺⁶. The highest Cr⁺⁶ concentration in any of the samples collected was 16 mg/kg in sample SB-64A/CA. The corresponding Cr⁺³ concentration was 49,000 mg/kg. This data shows that the chromium present at the site is almost exclusively in the more benign, less mobile trivalent state. The percentages of trivalent chromium range from 99.97% Cr⁺³ in sample SB-64A/CA up to 99.992% Cr⁺³ in sample 67A/CA. Hexavalent chromium analytical results are summarized in Table 4-2.

A STATE OF THE PARTY OF THE PAR

4.2.7 Site Classification/Characterization

In order to determine if the waste material exhibits the toxicity characteristic as defined in 40 CFR 261.24, a total of 12 surface and five subsurface soil samples were collected and analyzed for RCRA and TCLP metals. In addition, two of the production waste samples were re-sampled for RCRA and TCLP metals.

None of the soil samples collected contained leachable chromium over the 5 mg/l regulatory limit. Total chromium concentrations ranged from 230 mg/kg in SB-69A/CA to 54,000 mg/kg in SB-73B.

Samples SB-69A/CA and SB-69B/CB both exceeded the regulatory limit of 0.2 mg/l for mercury, at 0.57 and 0.58 mg/l, respectively. This sample was taken from the production waste area, and based upon the relatively low total chromium values (230 mg/kg and 1600 mg/kg), this sample does not reflect overall site conditions, only this localized area.

Production waste sample PW-9 exceeded the TCLP regulatory limit of 5 mg/l for chromium at a concentration of 14 mg/l. This result is similar to the TCLP chromium concentration of 24.2 mg/l that was obtained for this sample during the Phase I investigation. Unlike the Phase I investigation, PW-5 did not exceed the TCLP lead 'limit.

All other samples reported TCLP concentration below the regulatory limits.

4.3 CONSTITUENT DELINEATION

Upon completion of the Phase I investigation, it was determined that there were certain areas in which insufficient analytical data had been collected. As a result, additional data was collected in these areas. This sampling was in response to data gaps identified during preparation of the Phase I RI, not in response to a specific area of concern. This section summarizes the results obtained during this portion of the Phase II investigation.

In order to fill existing data gaps, and to gather additional data for purposes of the BRA, ERA, and site characterization, a total of 15 samples were collected and analyzed for TCL SVOCs, TCL P/PCBs, and RCRA metals.

The samples collected to further delineate extent were SB-61A through SB-69A, SB-62B, SB-67B, SB-69B, and the three samples collected during monitoring well installation, MW-1A, MW-5A, and MW-7A.

Nine of the 15 samples collected were surface soils and 6 were subsurface soils. The three subsurface soil samples collected during monitoring well installation were analyzed for the Target Analyte List (TAL) inorganics.

- i. Le carrier d'année de la company de

The following sections summarize the findings of the Phase II investigation and incorporates data from both the Phase I and II investigations into the colored Figures showing constituent distribution. Where appropriate, pertinent Phase I investigation material is incorporated into this section to complete the site overview. Analytical data for these samples is summarized in Table 4-2.

4.3.1 Base-Neutral/Acid Extractables

The only SVOC detected in the soil samples collected during monitoring well installation was di-n-butyl phthalate. This compound was identified in all three samples at concentration ranging from 240 μ g/kg in MW-5A to 2,200 μ g/kg in MW-1A. All of these concentrations were qualified with a "B", which indicates that this compound is present in the laboratory blank. It is likely that di-n-butylphthalate is not present in the environment at the sampling locations.

The remaining 12 samples, which were collected to gather additional information and close data gaps, were found to contain only PNAs. The PNAs were identified in eight of the 12 samples collected. The concentrations identified ranged from 550 μ g/kg in SB-64A/CA to 36,000 in SB-62A. The highest PNA concentration identified during the Phase I investigation was 172,000 μ g/kg in SB-15A. The Phase II data shows the same trends as the Phase I data with regard to PNA distribution; decreasing concentrations with increasing depth. Surfical and subsurfical total PNA distribution maps are contained in Figures 4-5 and 4-6, respectively.

4.3.2 Pesticides/PCBs

Pesticides and PCBs were not detected in any of the soil samples collected during monitoring well installation.

As expected from the Phase I data, low levels of pesticides were found in eight of the nine surface soils collected. Only surface soil sample SB-69A/CA did not contain pesticides. None of the subsurface samples were found to contain pesticides. Of the eight surface soils that contained pesticides, the highest total pesticide concentration observed was 2,066 μ g/kg in sample SB-63A. The second highest total pesticide concentration was 626 μ g/kg in SB-61A. The remaining six samples all contained less than 125 μ g/kg total pesticides. Similar to the Phase I investigation, total pesticide concentrations tended to decrease with increasing depth.

Polychlorinated biphenyls were identified in five of the 12 surface and subsurface soil samples collected. Similar to the Phase I investigation, the PCBs were not found in the samples collected in Area I, only in the production waste area and in the southern portion of Area II. The total PCB concentrations ranged from $160 \mu g/kg$ in SB-65A to $1,080 \mu g/kg$ in SB-69B/CB.

4.3.3 Inorganics

Significant quantities of inorganics were not detected in any of the samples collected during monitoring well installation. The arsenic concentrations ranged from 1.2 mg/kg in MW-5A to 3.4 mg/kg in MW-1A.

The chromium concentration ranged from 11.8 mg/kg in MW-1A to 73.5 mg/kg in the upgradient monitoring well location, MW-7A. The lead concentrations ranged from 2.2 mg/kg in MW-1A to 9.0 mg/kg in MW-7A.

The goal of collecting these samples was to more fully define the areas impacted by chromium, lead, and to a lesser extent, arsenic and mercury. Analytical results vary among the samples collected during this portion of the investigation. For example, chromium concentration range from 5.8 mg/kg in SB-62B to 49,000 mg/kg in SB-64A/CA. The surfical and subsurfical chromium distribution maps are contained in Figures 4-7 and 4-8, respectively. The surfical lead distribution map is contained in Figure 4-9. The surfical and subsurfical arsenic distribution maps are contained in Figures 4-10 and 4-11, respectively.

4.4 WETLAND SURFACE WATER/SEDIMENT SAMPLING

In order to collect additional data to support the ERA, three surface water and sediment samples were collected from the wetland area south and east of Area III. These samples were analyzed for TCL SVOCs, TCL P/PCBs, RCRA Metals, Acid Volatile Sulfide (AVS), Simultaneous Extractable Metals (SEM), and Total Organic Carbon (TOC).

4.4.1 Base-Neutral/Acid Extractables

The only class of compounds identified in the wetland sediment samples was PNAs. They were identified in all three samples at relatively low concentrations. Total PNA concentrations ranged from 430 μ g/kg in WL-2 to 2,590 μ g/kg in WL-1.

The only SVOC detected in the wetland surface water samples was bis(2-ethylhexyl)phthalate in sample WL-2 at an estimated concentration of 3 μ g/l. Phthalates, as a class of compounds are common laboratory contaminants. The presence of bis(2-ethylhexyl)phthalate in the environment at this sampling location is questionable. Analytical results are summarized in Table 4-6.

4.4.2 Pesticides/PCBs

Only one pesticide was identified in the three sediment samples collected. Methoxychlor was identified in sample WL-1 at an estimated concentration of 3.9 μ g/kg.

Aroclor-1248 was identified in WL-2 and WL-3 at estimated concentrations of 81 μ g/kg and 120 μ g/kg, respectively.

A STATE OF THE STA

Pesticides and PCBs were not detected in any of the surface water samples collected

Analytical results are summarized in Table 4-6.

4.4.3 Inorganics

Seven of the eight RCRA metals were not present in the wetland sediments at significant concentrations.

Chromium concentrations in the wetland sediment ranged from 9,410 mg/kg in sample WL-2 to 14,100 mg/kg in WL-3. These values are lower than those found throughout Area II and parts of Area III.

All hexavalent chromium concentrations were below the laboratory Practical Quantitation Limit (PQL). The valence state of the chromium in the wetland sediments is greater than 99.999% Cr⁻¹.

For the surface water samples, both filtered and unfiltered samples were collected. The dissolved metals samples were filtered prior to preservation utilizing a $0.45\mu m$ filter.

Barium, chromium, and lead were the only inorganics identified in the surface water samples. These analytes were detected at very low concentrations below Illinois Class I groundwater standards. All barium and detected lead results were flagged with a "B", indicating laboratory induced contamination.

The total chromium concentration in WL-SW-1 was 37.3 μ g/l, and the dissolved concentration was 6.5 μ g/l. By filtration, it is shown that the majority of the chromium is adsorbed to suspended soil particles and not dissolved in the water. Additionally, all of the low level chromium detections were also flagged with a "B", indicating minor laboratory contamination.

All hexavalent chromium concentrations were below the laboratory PQL.

Analytical results are summarized in Table 4-6.

4.4.4 Acid Volatile Sulfide/Simultaneous Extractable Metals

Acid Volatile Sulfide (AVS) and SEM analysis was performed on three wetland sediment samples to determine their persistence in the aquatic sediments. Acid Volatile Sulfide analysis is determined in a deoxygenated environment whereby sulfide is liberated and trapped. The sulfide concentration is determined by using a spectrophotometer. The SEMs are determined by Inductively Coupled Plasma Spectrometry (ICP). The six metals that are considered in the analysis are cadmium, copper, lead, mercury, nickel, and zinc. Values may be less than those obtained by other methodologies because sample preparation does not require such a rigorous digestion.

The SEM/AVS ratio is useful in predicting the toxicity of aquatic sediments. If the SEM/AVS ratio is less than unity (one), no significant toxicity exists. This is because binding phases (metal oxides, organic matter) reduces their bioavailability. If the SEM/AVS ratio is greater than one, then a toxic potential exists in the aquatic sediment. The laboratory analysis for wetland sediment sample WL-1 indicates a value less than one indicating that metal toxicity would not occur. However, analysis for WL-2 and WL-3 indicate that a toxicity potential exits in the aquatic sediment. Therefore, these metals are available and could potentially cause an ecological risk.

In addition, the surface water and sediment samples were analyzed for total organic carbon (TOC). The purpose of the TOC analysis is to aid in the determination of the bioavailability of organic material, which will enhance completion of the ERA. The TOC content of the soil ranged from 140,000 mg/kg in WL-1 to 330,000 in WL-3. For the surface water samples, the TOC content ranged from 5.9 mg/l in WL-SW-3 to 12 mg/l in WL-SW-1.

Analytical results for these analyses are contained in Table 4-7.

the state of the s

SECTION 5.0 SUMMARY AND RECOMMENDATIONS

5.1 PURPOSE

The purpose of the Phase II investigation activities was to supplement the Phase I investigation results by collecting sufficient data to fully characterize the site in terms of the impact to on and off site media from chemical compounds. The Phase II activities, used in conjunction with the Phase I activities, will also serve as the primary database for a Baseline Risk Assessment (BRA) and Ecological Risk Assessment (ERA).

The principal soil contaminants at this site, in reference to lateral extent and concentration, are chromium and, to a lesser extent, arsenic, lead, and mercury. Polychlorinated biphenyls, PNAs, and pesticides are present in specific areas of the site. Impact to the groundwater stems primarily from dissolved arsenic.

Parameter specific analyses were conducted during Phase II activities to further determine the extent, concentration, toxicity, and trend of the above compounds. In addition, analyses for dioxins/furans and asbestos were performed on select samples due to community input.

5.2 SUMMARY OF RESULTS

Four samples were collected for pesticide analysis surrounding soil boring SB-39 due to elevated pesticide levels detected at this location during the Phase I investigation. The analytical results indicated that the pesticide concentrations were low and occur in an isolated area. Fifteen soil samples were collected for PCB analysis from Area III due to elevated PCBs in a localized area. Polychlorinated biphenyls were detected in all fifteen soil samples collected during Phase II sampling activities. The predominant Aroclor was 1248.

In response to requests from the CAG, two soil samples were analyzed for PCDD/PCDF due to their concern that PCBs were burned at the site. The analytical results indicated the presence of PDCC/PCDF compounds but, not above the 1 ppb 2,3,7,8-TCDD risk based cleanup threshold. Two additional samples were analyzed for asbestos. It was the concern of CAG members that sand, used as fill at the site, contained asbestos. Asbestos was not present in the two samples.

To supplement data gaps and to gather additional information, additional soil samples were collected and analyzed for TCL Pesticides/PCBs, TCL SVOCs, and RCRA metals Pesticides were detected in eight of the nine soil samples collected. Polychlorinated biphenyls were detected in five of the twelve soil samples collected. Similar to the Phase I investigation, pesticides and PCBs tended to decrease with increasing depth.

TO SECURITION OF THE SECOND

Polynuclear aromatic hydrocarbons were detected in eight of 12 soil samples collected during Phase II sampling activities.

Due to elevated arsenic levels detected in select locations of Area II during Phase I activities, thirty-six soil samples were collected and analyzed for arsenic to define the extent of impact within this area. Slightly elevated levels of arsenic were present throughout Area II. Two of the thirty-six soil samples that exhibited the highest arsenic values, SB-44A and SB-50B, were tested for leachable arsenic. Neither result exceeded the 5 μ g/l TCLP regulatory limit.

As part of the investigation activities, twenty two soil and sediment samples were collected and analyzed to determine the valence state of chromium; trivalent, the less mobile and less toxic form, or hexavalent, the more mobile and more toxic form. The data indicated that, at a minimum, 99.97% of the chromium exists in the trivalent state.

To determine whether the material on site might be classified as a characteristic waste, twelve surface and five subsurface samples were analyzed for TCLP and RCRA metals. Two production waste samples, PW-5 and PW-9, were re-sampled for RCRA and TCLP metals. Only two soil samples, SB-69A/CA and SB-69B/CB exceeded the TCLP regulatory limit in both cases for mercury. These samples were taken in the production waste area and reflect only an isolated area and not the entire site. Production waste sample PW-9 exceeded the chromium TCLP regulatory limit of 5 mg/l. Phase I analytical results for production waste sample PW-9 also indicated a chromium concentration exceeding the TCLP regulatory limit. Phase II analytical results for Production Waste sample PW-5 did not exceed any TCLP regulatory limits. However, PW-5 had exceeded the chromium TCLP limit during the Phase I investigation. Both PW-5 and PW-9 appear to meet the chromium waste exclusion determination.

Phase II soil samples further defined the areas identified during Phase I as being impacted by chromium and to a lesser extent; lead, arsenic, and mercury.

Groundwater analytica' results indicated levels established for Class Groundwater in Illinois Administrative Code (IAC) Title 35, Subpart D, Section 620.140, were not exceeded for VOCs, BNAs, and pesticides/PCBs. But, as expected, inorganic analytes were detected in all samples collected. Dissolved arsenic was detected above the Class I Groundwater Standard (50 µg/l) in five monitoring wells; MW-1, MW-8, MW-9, MW-10, and MW-11. One monitoring well, MW-2, was close to the arsenic Class I Standard but was not included as an exceedance. Two monitoring wells, MW-4 and MW-5, were at or above the dissolved cadmium Class I Groundwater Standard. Dissolved manganese, iron, and TDS exceeded their respective Class I Groundwater Standard in most monitoring wells, however, since these constituents were prevalent in elevated levels during Phase I sampling and in the upgradient monitoring wells during both phases of the investigation, these levels are considered naturally occurring. Dissolved concentrations (filtered samples) did not exceed the Class I Standard for mercury, chromium, and lead.

19 - 14 41 35

The Phase IIA and IIB investigation determined the extent of the arsenic plume in the groundwater. Results from these investigations indicate that the plume has migrated approximately 400 feet to the east of the former tannery, under the EI&E railroad tracks and onto Commonwealth Edison's Waukegan Generating Station.

The groundwater samples were analyzed for hexavalent chromium to determine the valence state of chromium in the groundwater. One monitoring well, MW-4, contained 41 μ g/l of total hexavalent chromium. Total chromium concentrations for this well was 53 μ g/l whereas the dissolved chromium was less than 10 μ g/l.

To support an ERA, three wetland surface water and sediment samples were collected and analyzed for TCL SVOCs, TCL P/PCBs, RCRA Metals, AVS/SEM, and TOC.

Of the SVOCs, only PNA compounds were detected in all three sediment samples. One SVOC, bis(2-ethylhexyl)phthalate, was detected at an estimated valued in the surface water. This phthalate is a common laboratory contaminant.

One pesticide, methoxychlor, was detected in sediment sample WL-1 at 3.9 μ g/kg. Aroclor 1248 was identified at estimated concentrations in sediment samples WL-2 and WL-3 at 81 μ g/kg and 120 μ g/kg, respectively. Surface water analytical results did not indicate the presence of any pesticides or PCBs.

Seven of the eight RCRA metals were not detected at significant levels in each sediment sample collected in the wetland area. Chromium exhibited the highest concentrations ranging from 9,410 mg/kg in WL-2 to 14,100 mg/kg in WL-3. Chromium speciation analysis indicated that 99.99% of the chromium exists as trivalent chromium.

Total and dissolved RCRA metals analysis was performed on the surface water samples. Dissolved lead was detected at concentrations below Illinois Class I groundwater standards. Dissolved barium and chromium were identified and were flagged with a "B" qualifier indicating laboratory induced contamination.

Acid Volatile Sulfide and Simultaneously Extractable Metals were performed on three sediment sample to determine their persistence in the aquatic sediments. The analyses indicate that a toxicity potential exist in sediment samples WL-2 and WL-3 which could pose as an ecological risk.

Total organic carbon analysis were performed on both the sediment and surface water samples to determine the bioavailability of organic material. These data will enhance the completions of the ERA.

At this point, data appears sufficient for performing a BRA/ERA, and further site investigation does not appear necessary. From the activities performed and the data collected, it appears that the next step for this site is to perform a BRA/ERA. The human health risk assessment process

99430.000782

provides a necessary point of definition for whether a site or particular environmental media, is in need of remediation or requires mitigation measures.

A BRA is an evaluation of the potential threat to human health and the environment in the absence of any remedial action. A BRA is key to the process of investigating and formulating appropriate responses to environmental releases or contamination. The risk assessment identifies potential receptors (both human and environmental) and evaluates the likelihood that contaminants present at a site will adversely affect these receptors. It provides information to help EPA determine whether remedial action is necessary at a site.

Ecological risk assessments help identify environmental problems, establish environmental investigation priorities, and provide a scientific basis for regulator actions. The ERA helps evaluate the likelihood that adverse ecological effects may occur or are occurring as a result of exposure. The ERA is a qualitative and/or quantitative appraisal of the actual or potential effects of a site on plants and animals.

SECTION 6.0 REFERENCES

- U.S. EPA, 1974, Leather Tanning and Finishing, EPA-440/1-74-016a, Office of Air and Water Programs, Washington, D.C.
- U.S. EPA, 1980, <u>Hazardous Waste Land Treatment</u>, SW-874, Office of Water and Waste Management, Washington, D. C.
- U.S. EPA, 1986, Project Summary Field Investigation and Evaluation of Land Treating Tannery Sludges, EPA/600/S2-86/033, Robert S. Kerr Environmental Research Laboratory, Ada, Oklahoma.

Definitions of Data Qualifiers and Terminology

- A This flag is utilized to indicate that a tentatively identified compound (TIC) is a suspected additional product formed during sample processing and caution should be applied in interpreting these results.
- This flag is used when the analyte is found in the associated blank as well as in the sample. It indicates possible and probable blank contamination and warms the data user to use caution when applying the results of this analytes.
- BQL Below quantitation limit indicates the compound was not detected in the samples above the practical quantitation limit.
- C Indicates that a pesticide identification has been confirmed utilizing GC/MS techniques.
- D indicates that sample extract was diluted by the factor listed due to the sample matrix and/or concentration levels. All method detection limits or practical quantitation limits for the particular samples are therefore increased by this dilution factor.
- E Indicates that the concentration of the specific compound exceeded the calibration range of the instrument for that particular analysis.
- J Indicates an estimated value. It indicates that the compound was analyzed for and determined to be present in the sample. This flag is used either when estimating a concentration for tentatively identified compounds where a 1:1 response is assumed, or when the mass spectral data indicate the presence of a compound that met the identification criteria but the result is less that the sample quantitation limit ut greater than zero.
- ND Indicates the compound or analyte was not detected in the sample above the method detection limit or the practical quantitation limit for the particular analysis.
- PQL The practical quantitation limit is the lowest level that can be reliably achieved within specified limits of precision and accuracy during routine operating conditions.
- U indicates the compound was analyzed for but not detected in the sample above the applicable quantitation limit.

TAULÉ 4-1 GROUNDWATER ANALYTICAL RESULTS FORMER GREISS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number	MW-1A	AE-WM	MW-7A	MW-0	E-WH	Regulator
IEA Sample IO Humber	950219504	950219501	950219509	950218502	950219503	Slandards
VOLATILE ORGANIC COMPOUNDS						
HLOROMETHANE	<10	<10	<10	<10	<10	
ROMOMETHANE	<10	<10	<10	<10	<10	
VININ CHLORIDE	<10	<10	<10	<10	<10	2
CHLOROETHANE	<10	<10	<10	<10	<10	
WETHYLENE CHLORIDE	<10	<10	<10	<10	<10	
CETONE	<10	<10	<10	<10	<10	
CARBON DISULFIDE	<10	<10	<10	<10	<10	
I_1-DICHLOROETHENE	<10	<10	<10	<10	<10	7
,1-DICHLOROETHANE	<10	<10	<10	<10	<10	
,2-DICHLOROETHENE (TOTAL)	<10	€10	<10	<10	<10	100
HLOROFORM	<10	<10	<10	<10	<10	
2-DEHLOROETHANE	<10	<10	<10	<10	<10	3
-BUTANONE	<10	<10	<10	<10	<10	
,1,1-TRICHLORGETHANE	<10	<10	<10	<10	<10	200
ARBON TETRACHLORIDE	<10	<10	<10	<10	<10	3
ROMODICHLOROMETHANE	<10	<10	<10	<10	<10	
1,2-DICHLOROPROPANE	<10	<10	<10	<10	<10	5
CIS-1,3-DICHLOROPROPENE	<10	<10	<10	<10	<10	
TRICHLOROETHENE	<10	<10	<10	<10	<10	5
DISROMOCHLOROMETHANE	<10	<10	<10	<10	<10	
1,1,2-TRICHLOROETHANE	<10	<10	<10	<10	<10	
BENZENE	<10	<10	<10	<10	<10	5
TRANS-1,3-DICHLOROPROPENE	<10	<10	<10	<10	<10	
BROMOFORM	<10	<10	<10	<10	<10	
-METHYL-2-PENTANONE	<10	<10	<10	<10	<10	
-HEXANONE	<10	<10	<10	<10	<10	
TETRACHLOROETHENE	<10	<10	<10	<10	<10	5
1,1,2,2-TETRACHLOROETHANE	<10	<10	<10	<10	<10	
TOLUENE	<10	<10	<10	<10	<10	1000
CHLOROSENZENE	<10	<10	<10	<10	<10	100
ETHYLBENZENE	<10	<10	<10	<10	<10	700
STYPENE	<10	<10	<10	<10	<10	100
XYLENES (TOTAL)	<10	<10	<10	<10	<10	10000

All units are in ug/L (ppb).

Sample ID Number EA Sample ID Number	MW-1 950290005	HW-1A 950219504	950301003	MW-3 850290003	MW-4 950301001	MW-5 950290001	84W-5A 850218501	MW-0
SEMI-VOLATILE ORG. COMPOUNDS						,		
PHENOL	<10	<10	<10	<10	<10	<10	<10	<10
BIS (2-CHLOROETHYLIETHER	<10	<10	<to-< td=""><td><10</td><td><10</td><td><10</td><td><10</td><td><10</td></to-<>	<10	<10	<10	<10	<10
2-CHLOROPHENOL	<10	<10	<10	<10	<10	<10	<10	<10
1,3-DICHLOROBERZENE	<10	<10	· <10	<10	<10	<10	<10	<10
1,4-DICHLOROBEKZENE	<10	<10	<10	<10	<10	<10	<10	<10
BLENZYL ALCOHOL	<10	<10	<10	<10	<10	<10	<10	<10
1,2-DICHLOROSENZENE	<10	<10	<10	<10	<10	<10	<10	<10
2-METHYLPHENOL	<10	<10	<10	<10	<10	<10	<10	<10
BIS (2-CHLOROISOPROPYL) ETHER	<10	<10	<10	<10	<10	<10	<10	<10
4-METHYLPHENOL	<10	<10	<10	<10	<10	<10	<10	<10
N-NITROSO-01-N-PROPYLAMINE	<10	<10	<10	<10	<10	<10	<10	<10
HEMACHLOROTHEANE	<10	<10	<10	<10	<10	<10	<10	<10
MITROBENZENE	<10	<10	<10	<10	<10	<10	<10	<10
ISOPHORONE	<10	<10	<10	<10	<10	<10	<10	<10
2-NITROPHENOL	<10	<10	<10	<10	<10	<10	<10	<10
2,4-DIMETHYLPHENOL	<10	<10	<10	<10	<10	<10	<10	<10
BENZOIC ACID	<10	<10	<10	<10	<10	<10	<10	<10
BIS (2-CHLOROETHOXY) METHANE	<10	<10	<10	<10	<16	<10	<10	< 10
2,4-DICHLORPHENOL	<10	<10	<10	<10	<10	<10	<10	<10
1,2,4-TRICHLOROBENZENE	<10	<10	<10	<10	<10	<10	<10	<10
NAPHTHALENE	<10	<10	<10	<10	<10	<10	<10	<10
4 CHLORGANILINE	<10	<10	<10	<10	<10	<10	<10	<10
HEIACHLOROBUTADENE	<10	<10	<10	<10	<10	<10	<13	<10
4 CHLORO-3-METHYLPHENOL	<10	<10	<10	<10	<10	<10	<10	<10
2METHYLNAPHTHALENE	<10	<10	<10	<10	<10	<10	<10	<10
HEXACHLOROCYCLOPENTADENE	<10	<10	<10	<10	<10	<10	<10	<10
2.4.6-TRICHLOROPHENOL	<10	<10	<10	<10	<1D	<10	<10	<10
2,4,5-TRICHLOROPHENOL	<50	<25	<50	<50	<50	<50	<25	<50
2-CHLORONAPHTHALENE	<10	<10	<10	<10	<10	<10	<10	<10
2-MITROANILINE	<50	< 25	<50	<30	<50	<50	<25	<50
DIMETHYLPHTHALATE	<10	<10	<10	<10	<10	<10	<10	<10
ACENAPHTHYLENE	<10	<10	<10	<10	<10	<10	<10	<10
2.6-DINETROTOLUENE	<10	<10 St	<10	<10	<10	<10	<10	<1D
3-NITROANLINE	<50	<25	<50	<50	<90	<50	<25	<80
ACENAPHTHENE	<10	<10	<10	<10	<10	<10	<10	<10

Units are in ug/L (ppb) NA – Analysis not performed.

BELONO. OSTAN

TABLE 4-1 (Continued) GROUNDWATER ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Semple 10 Number	HW-7	MW-7A	MW-8	NW-8
EA Sample D Number	050290007	950290013	850290010	950290011
BEM-YOLATLE ON COMPOUNDS				
PHENOL	<10	<10	<10	<10
BIS (2-CHLOROETHM)ETHER	<10	<10	<10	<10
2-CHLOADPHENOL	<10	<10	<10	<10
1,3-DICHLOROGENZERS	<10	<10	<10	<10
1.4-DICHLOROGENZEIE	<10	<10	<10	<10
BÉNZYL ALCIOHOL	<10	<10	<10	<10
1,2-DICHLOROGENZE	<10	<10	<10	<10
2-METHYLPHENOL	<10	<10	<10	<10
BIS (2-CHLOROISOPROPYL) ETHER	<10	<10	<10	<10
4-METHYLPHENOL	<10	<10	<10	<10
N-NTROSO-DI-H-PROPYLANINE	<10	<10	<10	<10
HEXACHLOROTHEANE	<10	<10	<10	<10
NTROBENZENE	<10	<10	<10	<10
ESOPHORONE	<10	<10	<10	<10
Z-HTROPHENOL	<10	<10	<10	<10
2,4-DIMETH NYLPHE NOL	<10	<10	<10	<10
BENZOIC ACID	<10	<10	<10	<10
BIB (2-CHLOROETHOXY) METHANE	<10	<10	<10	<10
2.4-DICHLORPHENOL	<10	<10	<10	_ <10
1,2,4-TRICHLOROBERZENE	<10	<10	<10	<10
HAPHTHALENE	<10	<10	<10	<10
4 CHLOROANILNE	<10	<10	<10	<10
HEXACHLOROBUTADENE	<10	<10	<10	<10
4 CHLORO~3-METHREHENOL	<10	<10	<10	<10
ZMETRYLNAPKIHALDE	€10	<10	<10	<10
NEXACHLORIOGYCLOPENTADIENE	<10	<10	<18	<10
2.4.4—TRICHLOROPHEROL	<10	<10	<10	<10
2,4,5- (NICHLOROPHE)(OL	<50	<25	<25	<25
Z-CHLORONAPHTHALENE	<10	<10	<10	<10
2-NITROANILINE	<50	<23	<25	<25
DMETHILPHTHALATE	<10	<10	<10	<10
ACENAPHTHYLENE	<10	<10	<10	<10
2,0-DHITACTOLUENE	<10	<10	<10	<10
3-MTMOLNE	<50	<25	<25	<25
ACENAPHTHENE	<10	<t0< td=""><td><10</td><td><10</td></t0<>	<10	<10

Units are in ug/L (ppb)

Bample ID Humber	MW-1	MW-1A	NW-z	MW-3	MW-4	MW-5 950290001	MW-5A	MW-4	MW-6
IEA Sample IO Number	950290005	950219504	950301003	950290003	620201001	950290001	930218301	930301003	#3030 IOU
EM-VOLATLE ORG. COMPOUNDS									<50
2,4-DINITROPHENOL	<50	<25	<50	<50	<50	<50	<25	<50 <50	<50
4-AETROPHENOL	<50	<25	<50	<50	<50	<60	<25		
DIBENZOFURAN	<10	<10	<10	<10	<10	<18	<10	<10	<10
4-DINTROTOLUENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
DIETHYLPHTHALATE	<10	<10	<10	<10	<10	<10	<10	<10	<10
L-CHLOROPHENYL PHENYL ETHER	<10	<10	<10	<10	<10	<10	<10	<10	<10
PLUORENE	<10	<10_	<10	<10	<10	<10	<10	<10	<10
I-NITROANLINE	<50	<29	<50	<50	<\$0	<60	<25	<\$0	<50
I,S-DINTRO-2-METHYLPHENOL	<50	<25	<50	<50	<50	<60	<25	<50	<50
I-NTROSCOPHENTLAMNE (I)	<10	<10	<10	<10	<10	<10	<10	<16	<10
- BAOMOPHENYL PHENYL ENTER	<10	<10	<10	<10	<10	<10	<10	<10	<10
HEXACHLONOBENZENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
ENTACHLOROPHENOL	<50	<23	<50	<50	<50	<80	<25	<50	<\$0
PHENANTHRENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
WITHFLACENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
DI-N-BUTYUPHTHALATE	<10	2.8	<10	<10	<10	<10	2.53	<10	<10
FLUORANTHENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
TYRENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
BUTYLBENZYLPHTHALATE	<10	<10	<10	<10	<10	<10	<10	<10	<10
T-DICHLOROBENZIDINE	<10	<10	<10	<10	<10	<10	<10	<10	<10
SENZOKAJANTHFUCENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
CHRYSENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
BIS(2-ETHYLHEXYLIPHTHALATE	<10	2J	<10	<10	<10	<10	_54	<10	<10
DI-N-OCTYLPHTHALATE	<10	<10	<10	<10	<10	<10	<10	<10	<10
SENZORIFLUORANTHENE	<10	<10	<10	01>	<10	<10	<10	<10	<10
SENZOGOFLUORANTHENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
BENZOJAJPYRENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
NDENO(1,2,3-CD)PYRENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
DIBENZO/A HANTHRACENE	<10	<10	<10	<10	<10	<10	<10	<10	<10
BENZOIG HAPERYLENE	<10	<10	<10	<10	<10	<10	<1B	<10	<10

NA -- Analysis not performed Units are in ugit. (ppb)

MWG13-15_46647

684000 OETER

Sample ID Number	MW-7	MW-7A	MW-a	MW-B
IEA Sample ID Humber	950290007	950290013	950290010	950290011
SEMI-VOLATILE ORG. COMPOUNDS				
2,4-DINTROPHENOL	<50	<25	<25	<25
4-NITROPHENOL	<50	<25	<21	<25
DBENZOFURAN	<10	<10	<10	<10
2,4-DINITROTOLUENE	<10	<10	<10	<10
DETHILLATE	< 10	<10	<10	<10
4-CHLOROPHENYL PHENYL ETHER	<10	< 10	<10	<10
FLUORENE	<10	<10	<10	<10
4-NITROANILINE	<50	<25	<23	<23
4.4-DINITRO-2-METHYLPHENOL	< 50	<25	<25	<25
K-NITROSCOPHENYLAMINE (I)	<10	<10	<10	<10
I-BROMOPHENYL PHENYL EHTER	<10	<10	<10	<10
HEXACHILOROBENZEHE	<10	<10	<10	<10
PENTACHLOROPHENOL	<50	<25	<25	<25
PHENANTHRENE	< 10	<10	<10	<10
NITHPLACENE	<10	<10	<10	<10
D-N-BUTYLPHTHALATE	<10	2.10	2.8	2.IB
TLUORANTHENE	< 10	<10	<10	<10
PYRENE	<10	<10	<10	<10
RUTYLBENZYLPHTHALATE	<10	<10	<10	<10
J-DICHLOROBENZIONE	<10	<10	<10	<10
BENZO(A)ANTHRACENE	<10	<10	<10	<1D
CHRYSENE	<10	<10	<10	<10
HOZ-ETHYLHENYLPHTHALATE	<10	2,1	43	3,1
M-H-OCTYLPHTHALATE	<10	<10	<10	<10
ENZORIFLUORANTHENE	<10	<10	<10	<10
ENZOPOTLUGRANTHENE	<10	<10	<10	<10
ENZOJAJPYRENE	<10	<10	<10	<10
NDENO(1,2,3-CD)PYRENE	<10	<10	<10	<10
DIBENZOVAJNANTHRACENE	<10	<10	<10	<10
ENZOIG, HAPENYLENE	<10	<10	<10	<10

MA — Analysis not performed Units are in ug/L (ppb)

Sample ID Number		. MW_1A	MW-2	MW-s	MW-4	MW-5	MW-5A	MW-6
IEA Sample ID Number	950290005	950219504	950301003	950290003	950301001	950290001	950219501	950301005
PEST/PCB COMPOUNDS	L							
ALPHA-BHC	< 0.050	<0.050	< 0.050	<0.050	<0.050	< 0.050	<0.050	< 0.050
BETA-BHC	<0.050	< 0.050	< 0.050	<0.050	<0.050	< 0.050	<0.050	< 0.050
DELTA-BHC	< 0.050	<0.050	< 0.050	< 0.050	< 0.050	<0.050	<0.050	<0.050
GAMMA-BHC (LINDANE)	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050
HEPTACHLOR	<0.050	<0.050	< 0.050	< 0.050	<0.050	<0.050	< 0.050	< 0.050
ALDRIN	< 0.050	<0.050	< 0.050	< 0.050	< 0.050	<0.050	< 0.050	< 0.050
HEPTACHLOR EPOXIDE	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050	<0.050_	<0.050	< 0.050
ENDOSULFAN I	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050	<0.050	< 0.050	< 0.050
DIELDRIN	<0.10	< 0.10	< 0.10	<0.10	<0.10	<0.10	<0.10	<0.10
4,4'+-ODE	< 0.10	< 0.10	< 0.10	<0.10	<0.10	<0.10	<0.10	< 0.10
ENDRIN	< 0.10	< 0.10	< 0.10	<0.10	<0.10	<0.10_	<0.10	<0.10
ENDOSULFAN II	<0.10	<0.10	<0.10	<0,10	<0,10_	<0.10	<0.10	<0.10
4,4'-000	< 0.10	< 0.10	<0.10	<0.10	< 0.10	<0,10	<0.10	<0,10
ENDOSULFAN SULFATE	<0.10	<0.10	<0.10	< 0.10	<0.10	<0.10	<0.10	<0.10
4,4'-DDT	< 0.10	<0.10	< 0.10	<0.10	<0.10	<0.10	<0.10	<0.10
METHOXYCHLOR	< 0.50	< 0.50	< 0.50	<0.50	< 0.50	< 0.50	<0.50	< 0.50
ENDRIN KETONE	< 0.10	< 0.10	<0.10	<0.10	<0.10	<0.10	<0.10	<0,10
ENORIN ALDEHYDE	< 0.10	< 0.10	<0.10	<0.10	< 0.10	<0.10	<0.10	<0.10
ALPHA-CHLORDANE	<0.050	<0.050	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050	<0.050
GAMMA-CHLORDANE	< 0.050	< 0.050	< 0.050	< 0.050	< 0.050	<0.050	< 0.050	<0.050
TOXAPHENE	<5,0	<5.0	<5.0	<5.0	<5.0	<5.0	<5.0	< 5.0
AROCLOR - 1016	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
AROCLOR - 1221	<2.0	<2.0	<2.0	<2.0	< 2.0	<2.0	<2.0	<2.0
AROCLOR 1232	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
AROCLOR - 1242	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
AROCLOR 1248	<1.0	· <1.0	<1.0	<1.0	<1.0	<1.0	<1.0	< 1.0
AROCLOR - 1254	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
AROCLOR - 1260	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0

Units are in ug/L (ppb).

<u>}</u>

TABLE 4-1 (Continued)
GROUNDWATER ANALYTICAL RESULTS
FORMER GRIESS-PFLEGER TANNERY
COMMONWEALTH EDISON COMPANY
WAUKEGAN, ILLINOIS

Sample ID Number IEA Sample ID Number	MW-7 950290007	MW-7A 950219505	MW-8 950219502	MW-9 950219503
PEST/PCB COMPOUNDS	30023000)	B30213303	930213302	830219303
ALPHA-BHC	< 0.050	< 0.050	< 0.050	< 0.050
BETA-BHC	< 0.050	< 0.050	< 0.050	< 0.050
DELTA-BHC	< 0.050	< 0.050	< 0.050	< 0.050
GAMMA-BHC (UNDANE)	< 0.050	< 0.050	< 0.050	< 0.050
HEPTACHLOR	< 0.050	< 0.050	< 0.050	< 0.050
ALDRIN	< 0.050	< 0.050	< 0.050	<0.050
HEPTACHLOR EPOXIDE	< 0.050	< 0.050	< 0.050	< 0.050
ENDOSULFAN I	< 0.050	< 0.050	< 0.050	< 0.050
DIELDRIN	<0.10	< 0.10	< 0.10	< 0.10
4,4'-DDE	< 0.10	< 0.10	< 0.10	< 0.10
ENDRIN	<0.10	< 0.10	<0.10	< 0.10
ENDOSULFAN II	<0.10	< 0.10	<0.10	< 0.10
4,4'-DDD	<0.10	< 0.10	<0.10	< 0.10
ENDOSULFAN SULFATE	< 0.10	<0.10	<0.10	< 0.10
4,4'-DDT	<0.10	<0.10	<0.10	< 0.10
METHOXYCHLOR	<0.50	< 0.50	<0.50	< 0.50
ENDRIN KETONE	< 0.10	< 0.10	<0.10	< 0.10
ENDRIN ALDEHYDE	<0.10	<0.10	<0.10	< 0.10
ALPHA-CHLORDANE	< 0.050	< 0.050	< 0.050	< 0,050
GAMMA-CHLORDANE	< 0.050	< 0.050	< 0.050	< 0.050
TOXAPHENE	<5.0	<5.0	<5.0	<5.0
AROCLOR - 1018	<1.0	<1.0	<1.0	<1.0
AROCLOR - 1221	<2.0	<2.0	<2.0	<2.0
AROCLOR - 1232	<1.0	<1.0	<1.0	<1.0
AROCLOR 1242	<1.0	<1.0	<1.0	<1,0
AROCLOR - 1248	<1.0	<1.0	< 1.0	<1.0
AROCLOR - 1254	<1.0	<1.0	<1.0	<1.0
AROCLOR - 1250	<1.0	<1.0	<1.0	<1.0

Units are in ug/L (ppb).

MWG13-15_46650

\$3130 JOSE 74.

Sample ID Number	MW-1	MW-1 (Filtered)	MW-1A	MW-1A (Filtered)	MW-2	MW-2 (Filtered)	MW-3	MW-3 (Fithered)	Regulatory
IEA Sample ID Number	850290005	950290008	950219504	950219504F	950301003	950301004	950290003	950290004	Limit
NORGANIC COMPOUNDS			, r						
LUMMUM	NA	NA .	1770	33.0B	NA .	NA	NA.	NA NA	
ANTIMONY	NA	NA	4.98	<2.0	NA	NA	NA.	NA	
ARSENIC	J. C. 103 (25)	BERNIE FOR	36.9	6.18	正是对对环境	49.0	3.5	<2.0	50 - tL
SARIVIA	59,0	<50.0	28,CB	13.78	120.0	70.0	1200	<\$0,0	2000 - IL
STYLLUM	NA	NA NA	<1.0	1.0	NA	NA NA	NA	NA	
ADMUM	STREET, SECTION	<1.0	<1.0	1,18	< 5.0	<5.0	ST 157 1374	<5.0	6 – #L
CAUCIUM	NA	NA.	234000	221000	NA NA	NA	NA	NA	
CHROMUM	CALCULATE.	25.0	ALCOHOL: U.S.	3.88	12.00	40.0	沙罗西州亚东	13.0	100 - IL
EXAVALENT CHROMUM	< 0.01	NA .	< 0.01	NA	< 0.01	NA	<0.01	NA	
COBALT	NA	NA	4.08	1,58	NA	_ NA	NA.	NA.	1000 IL
COPPER	l NA	NA	8.68	2.19	NA _	NA	NA.	NA	650 - IL
RON	NA.	NA	1. 1.7 also \$2.1	1460	NA	NA _	NA NA	NA	5000 - IL
EAD	<3.9	<3,0	是57世海	1,0	CART TOP	<3.0	7,2	<3.0	7.5 - IL
MAGNEBIUM	NA	NA NA	62300	77600	NA	NA	NA	NA	
MANGANESE	NA	NA ·	151-2	TANK TINGE	NA	NA	NA NA	NA NA	150 - IL
MERCURY	<0.20	0.51	0.24	1.0	经数据的	<0,20	<0.20	<0.20	2 - JL
ACKEL	NA	NA	22,1	14,18	NA	NA_	1 NA	NA	100 - IL
MUREATO	NA.	NA	5800	5730	NA NA	NA .	NA	NA.	
SELEWUM	<3.0	<3.0	<3.0	9.4	<3.0	<3.0	<3.0	<3,0	50 - IL_
SLVER	<10.0	<10.0	<1.0	<1.0	<10.0	<10.0	<18.0	<10.0	60 IL
SODIUM	NA NA	NA	45500	47100	NA.	NA	NA NA	NA NA	
HALLIUM	NA.	NA	<3.0	<3.0	NA	NA NA	NA	NA NA	
VANADIUM	NA	NA	10.10	<1.0	NA	NA.	NA.	NA	
TING	MA	NA.	68.6	3.98	NA	NA.	NA.	NA.	5000 - IL
TOTAL CYANDE	<0.005(A)	NA.	<10.0	HA	<0.005(A)	NA.	<0.006(A)	NA NA	200 - IL

Unite ere in ug/L (ppb) RA -- Hot embysed. ** -- Action level under review, (A) -- Units ere in reg/L

	4.5					4		4.4	41 - 1670
		FC	OUNDWATE PAMER GRIE PAMONWEA	4-1 (Continued) R ANALYTICAL RI 88-PFLEGER TA LTH EDIBON COI EGAN, ILLINOIS	NNERY				K
mole (D Humber	MW-4	NW-4 (Fillated)	MW-S	MW-6 (Fithered)	MW-5A	MW-SA (Filtered)	36W-5	I sau a see . a I	m
ample ID Humber	950301001	950301002	950290001	830290003	850218301	850219501F	950301005	950001008	Regulator
HEC COMPOUNDS						634216-221	930301023	830001000	Land
	NA	NA	NA	. NA	2220	48.9	NA.	NA I	
	NA	NA NA	NA	NA.	< 2.0	<2.0	NA.	NA	
	19.0	2.5	<2.0	<2.0	<3.0	10.2	3.5	<2.0	50 - H.
	120.0	<50.0	<50.0	< 50.0	63.55	44.55	<50.0	<50.0	2000 - H
	NA.	NA .	RA	NA NA	<1.0	<1.0	NA.	NA.	
	27-37-38 to	Hellie Harris	PERSENT	William X of the	1,08	<1.0	<6.0	<5.0	5-1
	NA	MA	NA	NA	233000	240000	NA.	NA NA	
	53.0	<10.0	80.0	18.0	Bereit Teres	1,03)	41 - 32-52-52	10.0	100 - IL
CHRONINIK	41.0	NA.	<0.01	NA	<1.0	NA.	<0.01	NA	
	NA	NA.	NA	NA NA	8.78	3.68	NA	NA NA	1000 - E.
	NA	NA,	NA	NA.	15,18	<1.0	NA.	NA.	850 - IL
	NA	NA	NA	NA	ALC: TOWN	经验证证证证	NA	NA NA	5000 - IL
	<3.0	<3.0	<3.0	< 3.0	表表示。	2,38	8.7	<3.0	7.6 - IL
		NA .	NA	NA	67000	89400	NA	NA NA	
	HA		NA	, NA	1557 TYPES	KENGKY 1 (58-39)	NA	NA.	150 - IL
	NA NA	NA.				<0.2	0,28	0.26	2 - K
		KA <0.20	< 0.20	<0,20	< 0,20				
	NA			<0,20 NA	<0.20 9,68			NA I	100 - "
	NA <0.20	<0.20	<0.20			4.28	NA	NA NA	100 - IL
	NA <0.20 NA	<0 20 NA	<0.20 NA	NA	21800	4.28 23700	NA NA	NA NA	
	NA <0.20 NA NA	<0 20 NA NA	<0.20 NA NA	NA NA	9,08 21800 <3.0	4.28 23700 20.5	NA NA <3,0	NA €3,5	50 - IL
	NA	<0 20 NA NA <3.0 <10.0	<0.20 NA NA 5.0 <10.0	NA NA 8,9 <10.0	9,98 21800 <3.0 1,25	4.28 23790 20.8 <1.0	NA <3,0	NA <3,5 <10.0	50 - IL 50 - IL
	NA <0.20 NA NA <3.0 <10.0	<0 20 NA NA <3.0	<0.20 NA NA 8.0 <10.0	NA NA 8,9 <10.0	9,68 21800 <3.0 1.25 136000	4.28 23700 20,6 <1,0 149000	NA NA <3,0 13,0 NA	NA <3,0 <10.0 NA	50 - IL 50 - IL
	NA <0.20 NA NA <3.0 <10.0 NA	<0 20 NA NA NA <3.0 <10.0	<0.20 NA NA 8.0 <10.0 NA NA	NA NA 8,9 <10.0 NA	9,68 21800 <3.0 1,28 136000 <3.0	4.28 23700 20.6 <1.0 149000 4.08	NA NA <3,0 13,0 NA NA	NA <3.0 <10.0 NA NA	50 - IL 50 - IL
	NA <0.20 NA NA <3.0 <10.0 NA NA	<0 20 NA NA NA <3.0 <18.0 NA NA	<0.20 NA NA 8.0 <10.0	NA NA 8,9 <10.0	9,68 21800 <3.0 1.25 136000	4.28 23700 20,6 <1,0 149000	NA NA <3,0 13,0 NA	NA <3,0 <10.0 NA	50 - IL 50 - IL

\$5"V

Sample 10 Humber	MW-7	MW-7 (Filtered)	MW-7A	WW-7A (Filtered)	MW-B	MW-8 (Filtered)	MW-0	MW-8 (Filtered)	Regulatory
IEA Sample ID Number	950290007	950290008	950219505	950219305F	950219502	850219502F	950219303	950219303F	Limit
NORGANIC COMPOUNDS	3								
ALUMINUM	NA NA	NA.	1600	35.79	2390	41.58	1170	25.00	
ANTRIAONY	NA	_ NA	<2.0	<2.0	33.66	5.7B	8.4B	<2.0	
ARSENC	0.0	<2.0	5.6B	6.69	AMETITE SEE		第23年27年22章	TO THE PARTY OF TH	50 - 11
DANUM	190.0	58.0	00.7B	70.98	1028	31.28	121B	90.00	2000 - IL
BERYLLIUM	NA NA	NA	<1.0	<1.0	<1.0	<1.0	<1,0	<1.0	
CADIMUM	<8.0	<5.0	<1.0	<1.0	1.78	≪1.0	<1.0	<1.0	5 - IL
CALCIUM	NA NA	NA	148000	112000	250000	200000	191000	158000	
CHROMUM	19.0	11.0	8.2B	1.0	EN CONTRACTOR	10.1	建筑工程等	1.68	100 - IL
HEXAVALENT CHROMIUM	<0.01	<0.01	<0.01	NA NA	<0.01	NA	< 0.01	NA _	
COBALT	NA.	NA NA	4,68	E3.1	4,18	2.78	80.9	<1.0	1000 - IL
COPPER	1 NA	NA NA	12.48	<1,0	24.68	1.78	13.69	<1.9	680 - IL
RON	NA NA	NA NA	PASSES TO FREE	2360	THE TOTAL	2770	が他日日開発器	2090	5000 - IL
LEAD	A 10 (10 (10)	<1.0	HAS DEED	<1.0	Adda Hard	2,48		1.48	7.5 - IL
MAGNESIUM	NA	NA	49100	34500	125000	127000	32400	29500	
MANGANESE	NA .	NA NA	THE STEEL	"林"是"我"	100	REPORTED AND A	音級把打扮發	207	150 - IL
MERCURY	0.39	<0,20	< 0.20	<0.20	国語文文馆 理	< 0.20	0.40	<0.20	2 - IL
HCKEL_	NA	NA .	9.28	4.08	10.69	3.18	22.28	4.08	100 - IL
MURRATO	NA	NA NA	138000	14100	5990	6120	54800	58700	
BELENIUM	<3,0	<3.0	<3,0	11,1	<3.0	27.9	43.0	12.9	50 - IL
SILVER	<10.0	<10.0	<1.0	<1.0	<1.0	<1.0	<10	<1.0	50 - IL
SOCIUM	NA NA	NA	144000	148000	64800	90400	152000	180000	
THALLIUM	NA.	NA.	<3.0	<3.0	<3.0	4.58	<3.0	<3.0	
VANADIUM	NA	NA	10.29	<1.0	13.68	<1,0	5.15	<1.9	
ZINC	NA.	NA.	52.4	1.28	143	4.1B	181	1.78	5000 - IL.
TOTAL CYANDE	<0.505 (A)	NA	<10.0	NA.	<10.0	NA	<10.0	NA NA	200 - IL

Rample ID Humber	MW-10	MW-10 (Filtered)	MW-11	MW-11 (Filtered)	MW-12	MW-12 (Fithcod)	MW-13	MW-13 (FViered)	Regulatory
IEA Sample IO Humber	851187051	951197002	85*187003	951197004	951587002	851567001	931547004	851547000	timit
INCREMIC COMPOUNDS						STATE OF THE PARTY			100000
ALUMPUN	NA	NA	NA	NA NA	NA	NA I	NA.	NA I	
AHTIMONY	NA.	NA NA	NA .	NA NA	NA.	NA.	NA.	_ NA	
ARSENC	ALC: NO PERSON	(大三)次例 计中间程序	STATE AND SEC	在地域的企业的经验	3.6	2	10	8.6	60 - A.
BANKIM	120	68	200	<50.0	50	<50,0	130	79	2000 HL
DERYLLRIM	NA.	NA.	NA	NA.	NA	NA.	NA	NA NA	
CADMUM	<5,0	经验的基础的证据	新疆五百里	逐級分下 跨南外	<6.0	3377 ST 57538	<5.0	<3.0	8-L
CALDUM	NA	MA	_ NA	NA .	HA	NA	NA	NA	
CHROMINA	21	<10.0	80	<10,0	<10.0	<100	11	<10.0	100 - IL
HEXAVALENT CHROMIUM	NA	NA.	NA.	NA.	NA	NA.	HA	NA NA	
COBALT	NA	HA	NA	NA.	NA.	NA NA	HA	NA NA	1000 - AL
COPPER	NA.	MA	NA.	NA NA	HA	NA.	NA	NA NA	660 - IL
PION	NA.	MA	NA	NA.	HA	NA.	NA	NA	5000 - IL
FAD	BUILDING STORY	<3.0	17	<3.0	<3.0	9.1	1911		7.5 - FL
MUNESUM	NA	NA	NA	NA I	NA	NA.	NA	NA I	
MANGANESE	NA.	NA	FLA	NA.	NA	NA.	MA	NA I	150 - IL
MERCURY	<0.2	<0.2	< 0.2	<0.2	<0.2	<0.2	<0.2	<0.2	2 - L
NCKE.	NA.	NA	NA	NA I	NA	NA.	NA.	NA I	100 - IL
POTASSRIM	NA.	NA	NA	NA I	NA	NA.	NA	NA	
SELEWAN	<3,0	<3.0	<3.9	<3.0	<30.0	<30.0	<30.0	C20.0	50 - HL
SLVER	<10.0	<10.0	<10.0	<10.0	<10.0	<10.0	<10.0	<10.0	50 - fL
SODANI	NA	NA	NA	NA I	NA	NA.	KA	NA I	
THALLIUM	NA NA	NA.	NA	NA .	MA	NA.	NA	HA	
MUGAHAN	NA.	NA.	NA	NA.	NA	HA	NA	NA.	
ONC	NA	NA NA	NA	NA I	HA	NA.	NA	NA I	5000 - IL
TOTAL CYANIDE	NA	NA.	NA	NA.	MA	NA NA	NA	NA I	200 - R.

Unite are in ugit. (ppb) NA = Not embysed. ** — Auther land under review, (A) — Unite ere in mg/l.

Sample ID Number IEA Sample ID Number WET CHEMISTRY	MW-1	MW-1A	MW-2	.MW-3	MW-4	MW-5	MW-SA	MW-6	Regulatory
	95029005	9502205504	950301003	95029000	950301001	950290001	950220501	950301005	Limit
TURBIDITY TOTAL DISSOLED SOLIDS TOTAL SUSPENDED SOLIDS	30000 160000 72000	100000 1300000 180000	47000 1700000 160000	420000 1800000 220000	4000000 1200000 150000	100000 2700000 50000	280000 1700000 160000	110000 1500000 54000	1200000

Sample ID Number IEA Sample ID Number	MW-7 95029007	MW-7A 950220505	MW-8 950220502	MW-9 950220503	Regulatory Limit
WET CHEMISTRY					
TURBIDITY	310000_	600000	430000	170000	
TOTAL DISSOLED SOLIDS	1200000	930000	1500000	1200000	1200000
TOTAL SUSPENDED SOLIDS	720000	630000	510000	190000	

NA - Not analyzed.

94130.000797

Sample ID Number	SB-52A	SB-53A	SB-538	SB-54A	SB-55A	SB-56A	SB~57A
IEA Sample ID Number	950153017	950153018	950153019	950153020	950182001	950182002	
PESTICIDE/PCB COMPOUNDS							
ALPHA-BHC	<8.0	<8.0	<8.0	<8.0	NA	NA.	NA
BETA-BHC	<8.0	< 8.0	<8.0	<8.0	NA	NA.	NA
DELTA-BHC	<8.0	<8.0	<8.0	<8.0	NA	NA	NA
GAMMA-BHC (LINDANE)	<8.0	<8.0	<8.0	<8.0	NA.	NA	NA
HEPTACHLOR	<8.0	<8.0	<8.0	<8.0	NA	NA NA	NA
ALDRIN	<8.0	<8.0	<8.0	<8.0	NA	NA	NA
HEPTACHLOR EPOXIDE	<8.0	<8.0	<8.0	<8.0	NA	NA.	NA
ENDOSULFANI	<8.0	<8.0	<8.0	<8.0	NA	NA.	NA
DIELDRIN	<16.0	<16.0	<16.0	u	NA	NA.	NA.
4,4'-DOE	410	20	<15.0	39	NA	NA.	NA
ENDRIN	<18.0	<16.0	<16.0	<16.0	NA	NA	NA
ENDOSULFAN II	<16.0	<16.0	<16.0	<10.0	NA	NA NA	NA.
4,4°-000	190	<18.0	<16,0	<18.0	NA	NA	NA.
ENDOSULFAN SULFATE	<16.0	<16.0	<16.0	<16.0	NA.	NA	NA.
4,4°-00T	590	40	33		NA.	NA	NA.
METHOXYCHLOR	<80.0	<80.0	<80.0	<60.0	NA	NA	NA
ENDRIN ALDEHYDE	<16.0	<16.0	<10.0	<16.0	NA	NA	NA
ALPHA-CHLORDANE	<18.0	<16.0	<18.0	<16.0	NA	- NA	NA
GAMMA-CHLORDANE	<80.0	<80,0	<60.0	<80.0	NA	NA	NA
TOXAPHENE	<160.0	<160.0	<150.0	<160.0	NA	NA.	NA
TOTAL PESTICIDES	1190	60	93	222	0		
AROCLOR - 1015	NA NA	NA .	NA	NA.	<80.0	U	U
AROCLOR - 1221	NA NA	NA.	NA.	NA	<80.0	U	Ŭ
AROCLOR - 1232	NA	NA	NA.	NA.	<80.0	U	ŭ.
AROCLOR - 1242	NA NA	NA	NA.	NA NA	<80.0	Ü	Ü
AROCLOR - 1248	NA.	NA	NA.	NA NA	<80.0	91000	4600
AROCLOR - 1254	NA	NA.	NA	NA NA	580		910
ARDCLOR - 1260	NA NA	NA NA	NA	NA NA	<160,0	5800	310
TOTAL AROCLORS Inits are in ug/kg.	0	0	0	0	580		5820

In public prompted on 1

Sample ID Number	S8-58A	SB-588	58-58A	SB-60A	SB - 61A	SB - 62A	SB-52B	SB-63A
IEA Sample ID Number	950182004	950182005	950182006	950182007	950182008	950162009	950182010	950182011
PESTICIDE/PCB COMPOUNDS		•						
ALPHA-BHC	NA.	" NA	NA	NA.	<1.7	<1.7	<1.7	<1.7
BETA-BHC	NA NA	NA	NA	NA NA	<1.7	<1.7	<1.7	<1.7
DELTA-BHC	NA NA	NA NA	NA	NA NA	<1.7	<1.7	<1.7	<1.7
GAMMA-BHC (LINDANE)	NA	NA	NA	NA.	<1.7	<1.7	<1.7	<1.7
HEPTACHLOR	NA	NA	NA	NA NA	<1,7	<1.7	<1.7	<1.7
ALDRIN	NA NA	NA	NA	NA.	<1.7	<1.7	<1.7	<1.7
HEPTACHLOR EPOXIOE	NA	NA	NA	NA	<1.7	<1.7	<1.7	<1.7
ENDOSULFAN I	NA	NA.	NA	- NA	<1.7	<1.7	<1.7	<1.7
DIELDAIN	· NA	NA	NA	NA	<1.7	<1,7	<1.7	<1.7
4,4'-DDE	NA NA	NA	NA	NA	380	25	<1.7	870
ENDRIN	NA NA	NA .	NA.	N/A	<3.3	<3.3	<3.3	<3.3
ENDOSULFAN II	NA NA	NA.	NA.	NA NA	<3.3	<3.3	<3.3	<3.3
4,4'DDD	. NA	NA NA	NA .	NA	36	<3.3	<3.3	220
ENDOSULFAN SULFATE	NA NA	NA	NA.	NA NA	<3.3	<3,3	<3.3	<3,3
4,4'-DDT	NA	NA	NA.	NA NA	210	30	<3.3	890
METHOXYCHLOR	NA NA	NA	NA.	NA	<17	<17	<17	<17
ENDRIN ALDEHYDE	NA.	NA NA	NA.	NA	<3.3	<3.3	<3.3	<3.3
ALPHA-CHLORDANE	NA NA	NA	NA	NA NA	<3.3	<3.3	<3,3	4
GAMMA-CHLORDANE	NA.	NA.	NA	NA	<1.7	<1.7	<1.7	30
TOXAPHENE	NA NA	NA.	NA	NA NA	<1.7	<1.7	<1.7	<1.7
TOTAL PESTICIDES	0	D	0	0	526	55	0	208
AROCLOR ~ 1016	<80.0	<80.0	<60.0	<80.0	<33	<33	<33	<33
AROCLOR - 1221	<60.0	<80.0	<80,0	<80.0	<87	<67	<67	<67
AROCLOR - 1232	<60.0	<60.0	<80.0	<80.0	<33	<33	<33	<33
AROCLOR - 1242	<80.0	<80.0	<80.0	<60.0	<33	<33	<33	<33
AROCLOR - 1248	1100	<80.0	26000	<80.0	<33	<33	<33	<33
AROCLOR 1254	680	1900	9500	1200	<33	<33	<33	<33
AROCLOR - 1260	210	200			<33	<33	<33	<33
TOTAL AROCLORS	1990	2100	37600	1450	0	0	0	

hits are in ug/kg.

Sample ID Number	SB-MA/CA				SB-67B/CB	SB-58A	SB~69A/CA	\$8-69B/C8
IEA Sample ID Number	950182012	950182014	950182015	950182016	950182017	950182018	950182019	950182020
PESTICIDE/PCB COMPOUNDS								
ALPHA-BHC	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7
BETA-BHC	<1,7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7
DELTA-BHC	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7
GAMMA-BHC (LINDANE)	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1,7	<1.7
HEPTACHLOR	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7
ALDRIN	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7
HEPTACHLOR EPOXOE	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7
ENDOSUFANI	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7	<1.7
DIELDRIN	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3
4.4-DDE	18	23	<3.3	73	<3.3	14	<3.3	<3.3
ENDRIN	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3
ENDÖSULFANII	43	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3
4,4-D00	13	6.2	<3.3	9	<3.3	12	<3.3	<3.3
ENDOSULFAN SULFATE	43	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3
4,4-DDT	9.3	13	13	36	<3.3	<3.3	<3.3	<3.3
METHOXYCHLOR	<17	<17	<17	<17	<17	<17	<17	<17
ENDRIN ALDEHYDE	<3.9	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3	<3.3
ALPHA-CHLORDANE	35	14	<1.7	<1.7	<1.7	14	<1.7	<1.7
GAMMA-CHLORDANE	25	. 11	<1.7	<1.7	<1.7	17	<1.7	<1.7
TOXAPHENE	<170	<170	<170	<170	<170	<170	<170	<170
TOTAL PESTICIDES	100.3	87.2	13	118	0	57	. 0	
AROCLOR 1016	<33	<33	<33	<33	<33	<33	<33	<33
AROCLOR - 1221	<57	<67	<67	<67	<67	<67	<67	<67
AROCLOR - 1232	<33	<33	<33	<33	<33	<33	<33	<33
AROCLOR - 1242	<33	<33	<33	<33	<33	<33	<33	<33
AROCLOR - 1248	<33	<33	<33	<33	<33	<33	<33	<33
AROCLOR 1254	<33	<33	<33	<33	<33	<33	410	820
AROCLOR - 1280	<33	160	170	220	<33	<33	180	200
TOTAL AROCLORS Units are in ug/kg.	0	160	170	220	0	0		1080

Sample ID Number	SB-100A	SB-101A	SB-102A	SB-103A	SB-104A	SB-105A	8B-106A	SB-107A
IEA Sample ID Number	950424005	950424001	950424004	950424003	950424006	950424002	950424008	950182007
PESTICIDE/PCBS COMPOUNDS								
ALPHA-BHC	NA	NA NA	NA .	NA	NA	NA _	NA	NA NA
BETA-BHC	NA NA	NA NA	NA NA	NA	NA	NA _	NA V	NA
DELTA-BHC	NA NA	NA	NA	NA.	NA _	NA	NA .	NA
GAMMA-BHC (LINDANE)	NA	NA	NA	NA	NA_	NA .	NA	NA_
HEPTACHLOR	NA	NA.	NA	NA	NA -	NA	NA	NA
ALDRIN	NA .	NA	NA .	NA NA	NA	NA .	NA	NA .
HEPTACHLOR EPOXIDE	NA	NA	NA	NA	NA .	NA NA	NA .	NA
ENDOSULFANI	NA	NA.	NA NA	NA.	NA	NA _	NA NA	NA
DIELDRIN	NA	NA .	NA.	NA	NA .	NA _	NA	NA NA
4.4-DDE	NA	NA.	NA.	NA	NA	NA .	NA_	NA
ENDRIN	NA.	NA	NA NA	NA	NA	NA _	NA NA	NA
ENDOSULFAN II	NA	NA	NA NA	NA	NA NA	NA .	NA .	NA NA
4.4°-DDD	NA	NA	NA NA	NA	NA	NA	NA	NA
ENDOSULFAN SULFATE	NA	NA.	NA.	NA	NA	NA	NA.	NA
4.4'-DDT	NA NA	NA NA	NA	NA	NA	NA	NA	NA.
METHOXYCHLOR	NA.	NA	NA	NA	NA	NA .	NA.	NA NA
ENDAIN ALDEHYDE	NA	NA.	NA	NA	NA	NA _	NA NA	NA
ALPHA-CHLORDANE	NA.	NA.	NA	NA	NA	NA _	NA	NA
GAMMA-CHLORDANE	NA.	NA	NA.	NA NA	NA NA	NA	NA .	NA
TOXAPHENE	NA	NA	NA	NA NA	NA.	NA.	NA.	NA.
TOTAL PESTICIDES	0	0	0	0	0	. 0	G	
AROCLOR 1016	<80.0	<60.0	<60.0	<80.0	<60.0	<80.0	<80.0	<33
AROCLOR - 1221	<80.0	<80.0	<80.0	<60.0	<80.0	<80.0	<80.0	<67
AROCLOR - 1232	<80.0	<80.0	<80.D	<80.0	<80.0	<80.0	<80.0	<33
AROCLOR - 1242	<80.0	<80.0	<80.0	<80.0	<80.0	<80.0	<60.0	<33
AROCLOR - 1248	<60.0	240	<80.0	<80.0	210	36000	<80.0	670
AROCLOR - 1254	220	460	290	2500	250	19000	240	<160.0
AROCLOR - 1260	<160.0	260			<160	27000	<160.0	<160.0
TOTAL ARDCLORS	220							670

Units are in ug/kg.

Sample ID Number	MW-1A	MW-5A	MW-7A
IEA Semple ID Number	950211802	950211803	950211801
PESTICIDE/PCBS COMPOUNDS			
ALPHA-BHC	<2.0	<2.0	<2.0
BETA-BHC	<2.0	<2.0	<2.0
DELTA-BHC	<2.0	<2.0	<2.0
GANNA-BHC (UNDANE)	<2.0	<2.0	<2.0
HEPTACHLOR	<2.0	<2.0	<2.0
ALDRIN	<2.0	<20	<2.0
HEPTACHLOR EPOXIDE	<20	<2.0	<2.0
ENDOSULFANT	<2.0	<2.0	<2.0
DIELDRIN	<4.0	<4.0	<4.0
4,4'-DDE	<4.0	<4.0	<4.0
ENDRIN	<4.0	<4.0	<4.0
ENDOSULFAN II	<4.0	<4.0	<4.0
4,4'-000	<4.0	<4.0	<4.0
ENDOSULFAN SULFATE	<4.0	<4.0	<4.0
4,4'-DOT	<4.0	<4.0	<4.0
METHOXYCHLOR	<20	<20	<20
ENDRIN ALDEHYDE	<4.0	<4.0	<4.0
ALPHA-CHLORDANE	<2.0	<2.0	<2.0
GAMMA-CHLORDANE	<2.0	<2.0	<2.0
TOXAPHENE	<200	<200	<200
TOTAL PESTICIDES	0	0	*
AROCLOR - 1015	<40	<40	<40
AROCLOR - 1221	<81	<81	<81
AROCLOR - 1232	<40	<40	<40
AROCLOR - 1242	<40	<40	<40
AROCLOR - 1248	<40	<40	<40
AROCLOR - 1254	<40	<40	<40
AROCLOR - 1260	<40	<40	<40
TOTAL AROCLORS	0	0	

Jnits are in ug/kg

MWG13-15_46660

208000 05T

										and the state of the state of
•			TAR	LE 4-2 (Co	(beunitne					
				NALYTICAL						
						EDM				
			FORMER G							
					SON COMP	ANY				
			WA	UKEGAN, II	LLINOIS					
Sample ID Humber	SB-S4A	3B-38A	58-58B	88-80A	18-61A	8B-62A	8B-62B	AE9-88	58-63A	EB-64A/CA
IEA Sample 10 Humber	950102002	950162004	950182005	950102007	950182008	\$50182009	950182010	950182011	850182011	950182012
EMI-VOLATILE ORG. COMPOUNDS								`		
HENOL	NA	NA NA	NA.	NA	<330	<310	<330	<330	<330	<330
IS (2-CHLOROETHYLLETHER	NA	NA .	NA	NA NA	<330	<330	<330	<330	<330	<330
-CHLOROPHENOL	NA	NA	NA.	NA NA	<330	<330	<330	<330	<330	<330
,3-DICHLOROBENZENE	NA .	NA	NA	NA .	<330	<330	<330	<330	<330	<330
4-DICHLOROBENZENE	NA	NA.	NA	NA	<330	<336	<330	<330	<338	<330
ENZYL ALCOHOL	, NA	NA	NA.	NA.	<330	<330	<330	<330	<330	<330
7-DICHLOROBERZENE	NA NA	NA	NA	NA	<330	<350	<330	<330	<338	<330
- METHYLPHENOL	NA NA	NA	NA .	NA NA	<330	<330	<330	<230	<330	<330
IS 12-CHLOROISOPROPYLI ETHER	NA	NA	NA.	NA	<330	<330	<330	<330	<338	<330
-METHYLPHENOL -NETROSO-DI-N-PROPYLANINE	NA NA	NA NA	NA NA	NA NA	<330 <330	<330	<330 <330	<330	<330 <330	<330 <330
EXACHLOROTHEANE	NA NA	NA NA	NA NA	NA.		<330	<330	<330	<330	<330
TROBENZENE	NA.	NA.		NA	<330	<330	<330	<338	<330	<330
SOPHORONE	- NA	NA -	NA NA	NA	<330	<330	<330	<330	<330	<330
- NITROPHENOL	NA.	NA NA	NA NA	NA	<330	<330	<330	<330	<330	<330
4-DIMETHYLPHENOL	NA.	NA NA	NA NA	NA NA	<330	<330	<330	<330	<330	<330
ENZOIC ACID	NA	NA	NA	NA.	<330	<330	<330	<330	<330	<330
ES (2-CHLOROETHOXY) METHANE	NA	NA.	NA.	NA NA	<330	<330	<330	<330	<330	<330
4-DICHLORPHENOL	NA	NA	NA	NA.	<330	<330	<330	<330	<330	<330
2.4-TRICHLOROBENZENE	NA .	NA NA	NA	NA.	<330	<330	<330	<330	<330	<330
APHTHALENE	<880	<950	<\$80	<660	1600	<330	<330	<330	<330	<330
CHLOROANLINE	NA	NA.	NA -	NA	<330	<330	<330	<330	<330	<330
EXACHLOROBUTADIENE	NA	NA	NA .	NA NA	<330	<330	<330	<330	<330	<330
CHLORD-3-METRYLPHENOL	NA.	NA.	NA.	NA.	<660	<860	<680	<680	<680	<680
METHYLHAPHTHALENE	NA	HA	NA	HA	<335	<330	<330	<330	<330	<330
EXACHLOROCYCLOPENTADIENE	NA	NA .	NA	NA	<330	<330	<330	<330	<330	c330
4.4-TRICHLOROPHENOL	NA	NA NA	NA	NA.	<330	<330	<330	<330	<330	<330
4,5-TRICHLDROPHENOL	NA	NA	NA	NA	<1600	<1600	<1600	<1600	<1800	<1900
-CHLORONAPHTHALENE	NA	NA NA	NA .	NA	<330	<330	<330	<330	<330	<230
- NITROANILINE	NA .	NA NA	NA_	NA.	< 1600	<1000	<1800	<1000	<1800	<1800
METHYLPHTHALATE	NA	NA NA	NA	NA NA	<330	<330	<330	<330	<330	<330
CENAPHTHYLENE	<660	<160	<660	< 850	<330	<330	<330	<330	<130	<330
#-DINTROTOLUENE	NA .	NA NA	NA .	NA NA	<330	<330	<330	<330	<330	<330
ARBAZOLE	NA	NA	NA.	NA.	870	<330	<330	<338	<330	<330
TOTAL PNAs Inter are in ug/lig (ppb)			0		1900			0	10	

Sample ID Number	88-65A		SB-STAICA	38- 678/CB	88-88A	SB-SPA/CA	88-ee8/C8	MW-1A	MW-SA	MW-TA
IEA Sample ID Humber	950162014	950182015	950162016	950182017	950182018	950142019	95018202C	950211802	P50211803	05021160
SEMI-VOLATILE ORG. COMPOUNDS										
PHENOL	<330	<330	<330	<330	<330	<330	<330	<400	<390	<300
BIS (2-CHLOROETHYLJETHER	<330	<330	<330	< 330	<330	<330	<330	<400	<390	<300
2-CHLOROPHENOL	<330	<330	<330	<330	<330	<330	<330	<400	<390	<300
1,1-DICHLOROSENZENE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<300
1,4-DICHLOROBENZENE	<330	<330	<330	<330	<330	<530	<330	<400	<390	<300
DENZYL ALDOHOL	<330	<330	<330	<330	<330	<330	<330	<400	<390	<300
1,2-DICHLOROBENZENE	<330	<330	<330	<330	<330	<330	<330	<400	<380	<290
2-METHYLPHENOL	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
BIS IZ-CHLOROISOPROPYL) ETHER	<330	<330	<330	<330	<330	<330	<330	<400	<380	<390
I - METHYLPHENOL	<330	<330	<330	<330	<330	<330	<330	<400	<380	<390
N-NITROSO-DI-N-PAOPYLAMINE	<330	<530	<330	<330	<330	<330	<330	<400	<300	<300
HEXACHLOROTHEANE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
NITROBENZENE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
SOPHORONE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
-NITROPHENOL	<330	<330	<330	<330	<330	<330	<330	<400	<390	<290
2.4-DIMETHYLPHENOL	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
BÉNZOIC ACID	<330	<330	<333	<330	<330	<330	<330	<400	<390	<390
BIS (2-CHLOROETHOXY) METHANE	<330	<330	<330	<330	<330	<330	<330	<400	<590	<390
2.4-DICHLORPHENOL	<330	<330	<339	<330	<330	<330	<330	<400	<3₽0	<390
1.2.4-TRICHLOROBENZENE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
NAPHTHALENE	<330	<330	<330	<330	<300	€330	<330	<400	<390	<380
4 CHLOROANLINE	<330	<330	<33C	<330	<330	<330	<330	<400	<390	<390
HEXACHLOROBUTADIENE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
4 CHLORD -3 - METHYLPHENOL	<880	<680	<8G0	<660	<840	<600	<860	<400	<390	<390
METHYLNAPHTHALENE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
HEXACHLOROCYCLOPENTADIENE	<330	<330	<330	<230	<330	<330	<330	<400	<390	<390
2.4.6 - TRICHLOROPHENOL	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
24.5-TRICHLOROPHENOL	<1500	<1800	<1600	¢1800	<1600	< 1800	<1800	<970	<850	<840
-CHLORONAPHTHALENE	<330	<330	<330	<330	<330	<330	<330	<400	<380	<300
-NITROANLINE	<1800	<1600	<1800	< 1800	<1800	<1600	<1600	<870	<950	<940
DIMETHYLPHTHALATE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<300
ACEMAPHTHYLENE	<330	<330	<330	<330	<330	<230	<330	<400	<390	<380
2.4- DWITROTOLLENE	<330	<330	<330	<330	<330	<330	€330	<970	<850	<940
CARBAZOLE	<330	<330	<330	<330	<330	<330	<330	<400	<380	<300
TOTAL PNAs	0				- 100				0	

MA — Analysis not performed Units are in ug/lig (ppb)

Sample ID Humber	30-56A	A12-E2	89-546	A08-88	SB-GIA	58-62A	80-620	AC3 -E3	8B-63A	88-94A/C/
EA Sample ID Humber	950182002	9501B2004	950182005	950182007	950182008	950102009	950182010	950182011	950182011	950102012
SEMI-VOLATILE ORG, COMPOUNDS										
3-NITROANLINE	NA.	NA _	NA	NA NA	<1800	<1600	<1800	<1800	<1600	<1600
ACENAPHTHENE	<1200	<1200	<1200	<1200	1800	<330	<330	330	<330	<330
2,4-DINITROPHENOL	NA	NA .	NA.	NA	<1800	<1600	<1600	<1800	<1800	<1800
4-NITROPHENOL	NA .	NA	NA NA	NA NA	<1600	<1600	<1500	<1800	<1600	<1800
DIBENZOFURAN	NA	NA	NA	NA	1400	<330	<330	<330	<330	<330
2,4-DINTROTOLUENE	NA .	NA	NA .	NA	<330	<330	<530	<330	<330	<330
DIETHYLPHTHALATE	NA	NA	NA	NA NA	<330	<130	<330	<336	<330	<310
4-CHLOROPHENYL PHENYL ETHER	NA	NA	NA NA	NA NA	<330	<330	<330	<330	<330	<330
FLUORENE	200	<146	<140	<140	2200	<330	<238	<330	<330	<330
4-NITROANILINE	NA .	NA.	NA	NA NA	<1600	<1800	<1800	<1800	<1800	<1800
4.4-DINTRO-2-METHYLPHENOL	NA .	NA.	NA	NA	<1600	<1800	<1600	<1600	<1600	<1800
H-NTROSODIPHENYLAMINE (I)	NA .	NA .	NA	NA .	<330	<330	<330	<330	<330	<330
4-BROMOPHENYL PHENYL EKTER	NA NA	NA NA	NA	NA.	<330	<330	<330	<330	<330	<330
HEXACHLOROBENZENE	NA	NA	NA	NA .	<330	<330	<330	<330	<330	<330
PENTACHLOROPHENOL	NA	NA .	NA NA	NA.	<1000	<1800	<330	<330	<330	<330
PHENANTHRENE	820	<680	<880	<660	13000	4200	<330	2600	2800	<330
ANTHRACENE	<880	<680	<680	<860	<330	<230	<330	410	560	<330
DI-N-BUTYLPHTHALATE	NA	NA	NA	NA	<330	<330	<330	<330	<330	<330
FLUORANTHENE	870	< 580	<860	700	13000	4000	<330	_ 2400	2190	<330
PYRENE	1600	<180	<150	250	26000	9800	<330	6500	8430	85
BUTYLBENZYLPHTHALATE	NA.	NA	NA.	NA	<330	<330	<330	<330	<330	<330
3,5-DICHLOROSENZIDINE	NA NA	NA	NA.	NA NA	<680	<660	<660	<660	<640	<680
BENZO(A)ANTHRACENE	200	20	3000	510	9290	2900	<330	1500	1500	<330
CHRYSENE	260	<100	<100	780	8900	3000	<330	1700	1700	<330
BISIZ-ETHYLHEXYLIPHTHALATE	NA.	NA	NA	NA	<330	<330	<330	480	730	<330
DI-N-OCTYLPHTHALATE	NA	NA	NA	·NA	<330	<330	<330	<330	<330	<330
BENZOIB)FLUGRANTHENE	450	38	42	1490	8000	2700	<330	1700	1900	<330
BENZOK)FLUORANTHENE	170	<11		750	3300	1800	<330	870	1500	<330
BENZO/A)PYRENE	390	37		1200	4800	1900	<330	1400	1700	<330
INDENO(1,2,3-CD)PYRENS	270		<29	1200	8400	2000	<330	1300	<330	<330
DIBENZOJA, HJANTHRACENE	35	<20	<20	190	1900	0001	<330	<330	<330	<330
BENZOIG.H.MPERYLENE	250	51	81	1600	8130	2300	<330	1300	<330	<330
TOTAL PNA	8055	187		8540		34000	0		19780	

NA - Analysis not pertoened

MWG13-15_46663

SOBOGO GETS

Sample ID Number IEA Sample ID Number	38-05A 950182014	88-60A/CA 950152015	SB-67A/CA 950182018	58-67B/CB 650182017	38-66A 950182018	88-89A/CA 850182019	88-898/CB 850182020	MW-1A 950211802	MW-5A 950211903	MW-7A
SEMI-VOLATRE ORG. COMPOUNDS				-	030100010	***************************************	930107040	130411002	830211903	1 03021190
3-NITROANLINE	<1000	<1600	<1800	<1800	<1800	<1600	<1800	<270	<850	<940
ACENAPHTHENE	<330	<330	<330	<330	<330	<330	<330	<400	<300	<590
2,4-DINTROPHENOL	<1600	<1600	<1600	<1400	<1800	<1800	<1800	<970	<950	<940
4-NITROPHENOL	<1800	<1800	<1800	<1600	<1800	<1600	<1600	< 070	<850	<840
DIBENZOFURAN	<330	<330	<330	<330	<330	<330	<330	<400	<386	<390
2,4-DINITROTOLUENE	<330	<330	<330	<330	<330	<330	<338	<400	<300	<300
DIETHYLPHIHALATE	<330	<330	<330	<330	<330	<330	<330	<400	<300	<390
4-CHLOROPHENYL PHENYL ETHER	<330	<330	<330	<330	<330	<330	<330	<400	<380	<390
FLUORENE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<390
4 - NITROANILINE	<1800	<1500	<1800	<1600	<1800	<1500	<1600	<970	<950	< 840
4.6-DINITRO-2-METHYLPHENOL	<1800	<1600	<1600	<1800	<1800	<1600	<1800	<970	<850	<940
N-NTROSODPHENYLAMINE (1)	<330	<130	<330	<330	<330	<330	<330	<400	<390	<380
4-BROMOPHENYL PHENYL EHTER	<330	<330	<330	<339	<330	<330	<330	<400	<300	<390
HEXACHLOROBENZENE	<330	<330	<330	<330	<230	<330	<330	<400	<390	<390
PENTACHLOROPHENOL	<330	<130	<330	<330	- <330	<230	<330	<970	<950	C340
PHENANTHRENE	<330	480	<330	<330	<330	<330	<330	<400	<390	<290
ANTHRACENE	<330	<130	<330	<330		<330	<330 <330	<400	<390	<390
DI-N-BUTYLPHTHALATE	<330	<330	<330	<330	<330	<330	€330	2220JB	240.8	200/B
FLUORANTHENE	<330	500	<330	<330	<330	<330	€330	<400		
PYRENE	870	1500	400	<330	730				<390	<390
BUTYLBENZYLPHTHALATE	<330	<330				<330	<330	<400	<390	<380
3.T-DICH ORDBENZIONE	<440	<660	<330	<330	<330	<330	<330	<400	<300	<390
BENZO(AIANTHRACENE	340		<860	<660	<660	<860	<880	<400	<380	<390
CHRYSENE		370	<330	<330	<330	<330	<330	<400	<390	<390
	<330	510	<330	<330	<330	<330	<338	<400	<390	<390
BIS(2-ETHYLHEXYLIPHTHALATE	<330	<330	700	<330	1400	450	490	<400	<390	<390
	<330	<330	<330	<330	<330	<330	<330	<400	<380	<390
BENZOIS)FLUORANTHENE	<330	<330	<330	<330	<330	<330	<330	<400	<300	<390
BENZOOGFLUORANTHENE BENZOIAIPYRENE	<350	<500	<330	<330	<330	<330	<330	<400	<300	<390
	<330	<130	<330	<330	<330	<330	<330	<400	<380	<390
NDENO(1,2,3-CD)PYRENE	<530	<330	<330	<330	<330	<330	<330	<400	<300	<290
DIBENZO(A,MANTHRACENE	<330	<330	<330	<330	<330	<330	<330	<400	<390	<300
BENZOIGH APENYLENE TOTAL PNA	<220	<330	<330	<330	<330 730	<330	<330	<400	<300	<380

NA - Analysis not performe



				(E)						
			SOIL FORMER G COMMON	BLE 4-2 (Con ANALYTICAL I RIESS-PFLEI NEALTH EDISI AUKEGAN, ILL	RESULTS BER TANNER ON COMPAN					
Sample ID Number	58-50B	\$8-51A	88-02A	8B-62B	88-63A	88-84A/CA	BB-64CB		BB-BSA/CA	
IEA Sample ID Number	950188014	950182008	950182009	950182010	950182011	950182012	950182013	950182014	950182015	950182916
INDRBANIC COMPOUNDS										
LUUINUM	NA NA	NA	NA NA	NA NA	NA.	NA	NA NA	NA	NA NA	- NA
NTMONY	NA	<u>N</u> A	NA	NA	NA.	NA	NA	NA.	NA	NA
RSENIC	7.1	6.0	5.0	1.9		9.4	140**	63**	380**	78**
ARIUM	740	520	180	<4.0	140	180	- 300	120	250	320_
ERYLUUM	NA	NA	NA	NA.	NA .	NA	NA	NA	NA	NA.
ADMINIM	45	4.7	1,4	< 0.40	22	81	4,4	2,8	4.9	2.5
CALCIUM	NA	NA	NA NA	NA	NA .	NA NA	NA	NA NA	NA NA	NA NA
HROMIUM	5000	1100	180	5.8	36000	49000	27000	32000	31000	36000
EXAVALENT CHROMIUM	<1.4	NA	NA	NA	NA.	16	12	NA	<2.7	2,5
OBALT	NA	NA.	NA.	NA	NA NA	NA NA	NA	NA	NA_	NA.
OPPER _	NA NA	NA .	NA NA	NA NA	NA.	NA	NA	NA	NA NA	NA
RON	NA NA	NA	NA NA	NA	NA NA	NA.	NA NA	NA	NA	NA
EAD	1000	220	170	1,8*	480	930	470	720	550	_410
IANGANESE	NA NA	NA	NA	NA	NA	NA NA	NA	NA	NA	NA_
IERCURY	3.2	0.96	0.092	< 0.094	5.0	4.5	0.54	5,5	0,39	0.25
HCKEL	NA NA	NA NA	NA	NA	NA	NA NA	NA	NA.	NA	NA
POTASSIUM	NA.	NA NA	NA NA	NA NA	NA	NA NA	NA.	NA.	NA	NA NA
ELENIUM	<3.7	<1.5	51.Z	1.2	<2.5	<3.4	<3.7	<3,5	<2.7	<2.8
ILVER	260	< 0.91	<1.1	<0.08	<20	<2.2	<2.4	<2.0	<2.4	2,3
MUIDOS	NA	NA .	NA	NA NA	NA	NA	NA NA	MA	NA	NA.
HALLIUM	NA	NA NA	NA	NA	NA _	NA	NA	NA.	NA	NA NA
/ANADIUM	NA	NA .	NA -	NA	NA.	NA NA	NA_	NA	NA	NA.
ZINC	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA NA
CYANIDE	NA.	NA.	NA	NA.	NA	NA NA	NA	NA	I NA	NA

Urite are in rights (ppm)
NA - Not analyzed.
* - Analysis by 7421 on 2/3/85 with PQAL of 0.3 mg/kg
** - PQL varied with samples weight and percent solids.

			TAP	ILE 4-2 (Cont	frauni					
				ANALYTICAL F						
				RIESS-PFLEC		Y				
			COMMONY	VEALTH EDISC	N COMPAN	Y				
			WA	UKEGAN, ILLI	NOIS					
Sample ID Number	88-608	88-01A	8B-02A	89-628	88-63A	SB-E4A/CA	88-64CB	88-65A	BB-66A/CA	BB-STAICA
IEA Sample ID Number	950188914	950182008	980182809	950182010	950182011	950182012	950102013	950182014	950182015	950182018
MORGANIC COMPOUNDS										
ALUMENUM	NA	NA .	, NA	NA	NA	NA	NA	NA .	NA .	NA
WHITH WITH	NA	NA	, NA	NA	NA	NA	NA.	NA	_ NA	_ NA
ARBENIC	7.1	6.0	5.0	1,9	111	9.4	140**	63**	380**	78**
ARIUM	740	520	180	<4.0	140	180	300	120	250	320
SERVILIUM	NA	NA NA	_ NA	NA.	NA	NA NA	NA	NA.	NA NA	NA
DADMIUM	45	4.7	1.4	<0.40	22	81	4.4	2.0	4.9	2.5
CALCIUM	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
CHROMIUM	8000	1100	180	5.8	36000	49000	27000	32000	51000	56000
HEXAVALENT CHROMIUM	<1.4	NA .	NA _	NA	NA	16	12	NA	<2.7	2.5
COBALT	NA	NA	NA	NA_	NA	NA	NA	NA	NA	NA
COPPER	NA	NA	NA	NA_	ŅA	NA	NA	NA.	NA	NA
RON	NA	NA	NA .	NA	NA	NA	NA	NA	NA	NA
EAO	1000	220	170	1.8*	480	830	470	720	560	410
MANGANESE	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
MERCURY	3.2	0,98	0.092	< 0.094	5.0	4.5	0.54	5.5	0.39	0.25
HICKEL	NA.	NA_	NA .	NA	NA	NA	NA	NA	NA	NA
POTABBIUM	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
BELENIUM	<3.7	<1.5	<1.7	1.2	<2.5	< 3,4	<3.7	<3.5	<2.7	<2,8
BILVER	260	<0.01	<1.1	< 9.08	<20	<2.2	<2.4	<2.0	<2.4	2.3
SODIUM	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
THALLIUM	NA	NA.	NA	NA	_NA	NA.	NA	NA .	NA .	NA NA
MUIGANAY	NA	NA	NA	NA	NA	NA	NA NA	NA	. NA	NA
ZINC	NA NA	NA	NA	NA	NA	NA	,NA	NA	NA	NA
CYANIDE	NA	NA	NA.	NA	NA	NA	NA	NA	NA.	NA

Units are in mg/kg (ppm)

NA - Not analyzed.

"- Analyzes by 7421 on 2/3/95 with PQAL of 0.3 mg/kg

"- PQL varied with samples weight and percent solids.

Table 4–2 (Considued) Soil analytical results Former Griebs-Pfleger Tannery Commonwealth Edison Company Waukegan, Illinois

Sample 10 Number	88-478	88-68A	88-59A/CA	68-698/C8	88-70A	98-71A	58-71B	89-72A	5B-73A	88-788
IEA Sample ID Number	950182017	950182018	950182019	950162020	950188091	950188002	950186003	950138004	950188006	25016800
MORGANIC COMPOUNDS										
ALUMNUM	NA	NA NA	NA	NA	NA	NA	NA	NA	NA	NA
ANTIMONY	NA NA	NA	NA	NA	NA	NA	NA	NA .	NA	NA
ARBENIC	370**	16	4.0	5.4	<3.6	38**	_20	14	11	98**
BARIUM	130	140	250	280	120	150	120	150	68	330
BERYLLIUM	NA	NA NA	NA	NA NA	NA	NA	NA	NA	NA	NA
CADMIUM	3.2 1	3.7	9.5	21	40	6,5	3.3	14	<1.2	6.2
CALCIUM	NA	NA	NA NA	NA	NA.	NA	NA	NA	NA.	NA
CHROMIUM	3600	44500	230	1600	820	27000	37000	49000	42000	54000
HEXAVALENT CHROMIUM	< 1.6	NA	<1.8	<1.9	<1.5	<1.75	<2.1	<2.3	<2.4	<3.1
COBALT	NA	NA.	NA .	NA	NA	NA	NA	NA_	NA	NA
COPPER	NA	NA.	NA	NA	NA	NA	NA	NA _	NA	NA
IRON	NA.	NA	NA	NA	NA	NA_	NA .	NA.	NA	, NA
LEAD	100	970_	180	700	360	680	920	1200	1000	\$30
MANGANESE	NA	NA	NA.	NA.	NA.	NA	NA NA	NA	NA	NA
MERCURY	0.55	9.5	16	35	3.9	6,8	11	16	11	4,3
NICKEL	NA	NA NA	NA	NA NA	NA	NA.	NA	NA	NA	NA
POTASSIUM	NA	NA .	NA	NA						
BELENIUM	<1.9	<3.7	<1.8	<2.4	<3.0	<4.1	<6.3	<6.3	<6.1	<9,1
SILVER	<1.3	<2.0	92	230	89	<1.4	<2.0	2.7	<2.3	< 3.0
SODIUM	NA .	NA	NA	NA	NA	NA.	NA	NA.	NA	NA
THALLIUM	NA	NA	NA	NA NA	NA	NA	NA	NA.	NA.	NA
VANADIUM	NA	NA	NA	NA	NA	. NA	NA	NA _	HA	NA
ZNC	NA	NA NA								
CYANIDE	NA NA	NA .	NA NA	NA .	NA	NA NA	NA	NA.	NA	NA.

NA - Not ensigned.

- Analysis by 7421 on 2/3/85 with PQAL of 0.3 mg/kg

- PQL varied with samples weight and percent solids.

Sample ID Number	6B-74A	8B-75A	8B-76A	PW-5	PW-B	MW-1A	MW-BA	MW-7A	MW-10	MW-11
IEA Sample ID Number	9501688008	9501688009	9501888010	9501888005	850188801 t	950188802	950188803	#501#4801	951167901	
MORGANIC COMPOUNDS										00110100
ALUMINUM	NA NA	NA	NA I	NA	NA .	988	1280	1880	NA .	MA
ANTIMONY	NA NA	NA	NA	NA	NA.	< 0.47	< 0.38	<0.39	NA	NA
ARSENIC	16	18	12	5.1	0.68	3.4	1.28	3,1	6.3	5.0
BARRUSE	390	220	460	39	< 0.4	3.1B	4.39	17.3B	18	34
BERYLUUM	NA	NA	NA	NA	NA	< 0.24	<0.15	0.24B	NA	MA
CADMIUM	<1.6	42	3,6	1.7	<1.4	< 0.24	0.198	0.21B	< 0.58	0.76
CALCIUM	NA	. NA	NA	NA	NA	38800	25700	29700	NA	NA
CHROMIUM	40000	41000	37000	21000	23000	11.8	20.0	73.8	7.4	5.9
HEXAVALENT CHROMIUM	<5.0	3.5	< 3.03	<2.7	<2.5	NA	NA	NA	NA	NA
COBALT	NA	. NA	NA	NA	NA.	1.68	68.1	2.08	NA	NA
COPPER	NA NA	_ NA	NA	NA	NA	1.7B	3.1B	4.49	NA	NA
IRON	NA.	NA NA	NA	NA	NA	2680	3840	4350	NA	NA
LEAD	490	1400	191	16	16	2.2	3.0	9.0	1.2	1.6
MAGNESIUM	NA .	NA	NA	NA	NA	18400	12300	1114200	NA	NA
MANGANESE	NA NA	NA	NA	NA	NA	141	1116	130	NA	NA
MERCURY	1.4	0.75	0.24	2.5	< 0.2	<0.12	<0.11	<0.11	<0.12	< 0.13
NICKEL .	NA .	_ NA	NA	NA	NA	2.58	2.98	7.58	NA	NA
POTABBIUM	NA	_ NA	NA	NA	NA	237B	344B	358B	NA	NA
BELEHIUM	<11	<7.7	<7.8	<7.7	<7.1	< 0.71	< 0.53	< 0.58	<1.5	<1.6
BILVER	<3.2	<2.2	<2.9	<2.1	<2.6	<0.24	<0.18	< 0.19	<1.1	<1.2
SODIUM	. NA	NA	NA	NA	NA	76.38	126	1138	NA	NA
THALLIUM	NA	NA	NA	NA	NA	0.728	< 0.53	<0.58	NA	NA
WINDHAM	NA	NA	NA	NA	NA	7.8B	9.9	11.3	NA	NA
ZING	NA	NA	NA	NA	NA	20.2	22.1	20.B	NA	NA.
CYANIDE	NA.	NA	NA	NA	NA.	< 0.68	<0.54	<0.64	NA	NA

NA — Not analyzed.

* — Analysis by 7421 on 2/3/95 with PQAL of 0.3 mg/kg

** — PGL varied with samples weight and percent solids.

A STATE OF THE STA

TABLE 4-3 SOIL ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number	88-55A	SB-59A
PACE Sample ID Number	SAM0213061	SAM0213071
PCDD/PCDF COMPOUNDS		
2378-TCDD	74.17	ND
12378-PeCDD	209.09	ND
123478-HxCDD	269.05	58.44
123578-HxCDD	470.68	422.87
123789HxCDD	464.46	167,09
1234678-HpCDD	7406,49	13993.85
OCDD	51187,16	138807.5
2378-TCDF	74.24	37.12
12378~PeCDF	263.73	233.57
23478-HxCDF	20.67	5.7
123478-HxCDF	58,33	86,96
123678-HxCDF	25.57	40.25
123769HxCDF	17.02	29.92
234678-HxCDF	48.17	70.76
1234678 - HpCDF	611.14	1923.63
1234789-HpCDF	61.31	
OCDF	850.32	_7785.78
Total TCDD	1125.17	3,81
Total PeCDD	1485.03	<u>N</u> D
Total HxCDD	4851.65	2429.83
Total HpCDD	15510.15	24455,53
Total TCDF	610,67	544.83
Total PeCDF	702.25	1016.67
Total HxCDF	2997.99	6245.72
Total HpCDF	4426,37	15924.28

Units are pg/g - parts per trillion (ppt)

ASBESTOS ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWFALTH EDISON COMPANY WAUKEGAN, ILLINOS

Sample ID Number	AS-1	AS-2
Asbestos Analytical Results		
Asbestos	None	None
% Non-Asbestos Fiber - Cellulose	30	40
% Non-Fiber Matter - Mineral Grains	70	60

Bulk asbestos analysis by PLM-DS

Sample ID Number	SB-44A	SB-44B 950153002	ı
EA Sample ID Number	950153001	950153002	٤
INORG, COMPD - Arsenic Only			
Arsenic	4600	5,0	L
Units are in mg/kg (ppm)			
Sample ID Number	SB-44A	Regueltory	ł
IEA Sample ID Number	950153010	Umit	ı
INORG, COMPD - Amenic Only			l
TCLP Arsenic	0.4	5.0	Ł
	9.7		

Sample ID Number	SB-49A	SB-49B	SB-50A	SB-50B	SB-51A	SB-51B	SB-77A	SB-77B	SB-78A	
IEA Sample ID Number	950153010	950153011	950153012	950153014	950153015	950153016	950248015	950248016	950248017	950248018
INORG, COMPD - Arsunic Only					'					
Arsenic	16	9.5	3,5	10000	16	25	8.5	8.5	13	7.8

Units are in mg/kg (ppm)

Sample ID Number	\$8-508	Regualtory
EA Sample ID Number	950153011	Limit
INORG, COMPD - Arsenic Only		
TCLP Arsenic	2.2	5.0

Units are in mg/L (ppm)

Sample ID Number IEA Sample ID Number	SB-79A 950248013	SB-799 950245014	SB-80A 950248019	SB-608 950248020	SB-81A 950153001	SB-81B 950153002	SB-82A 950153003	SB-82B 950153004	SB-83A 950153005	SB-838 950153006
INORG. COMPD - Arsenic Only						_				
Arsenic	23	780	1600	710	49	55	180	240	12	22
Units are in mo/kg (ppm)						13.				

	TT-28-12-17	100 000	1 00 000 1	in and	50 000 I	Loop Toop
Sample ID Number	58-84A	58-84B	ZR-R2V	20-628	38-00A	20-000
Sample ID Number IEA Sample ID Number	950153007	950153008	850153009	850153010	950153011	950153012
INORG, COMPD - Arsenic Only						
Acconic	7.1	10	210	70	21	4800

Units are in mg/kg (ppm)

TABLE 4-8 WETLAND SOIL ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number	WL-1	WL-2	WL-3
IEA Sample ID Number	9504269-04A	8504269-05A	9504269-06/
BEMI-VOL ORG. COMPOUNDS			
PHENOL	<1900	<1500	<2200
BIS (2-CHLOROETHYL)ETHER	<1000	<1600	<2200
2-CHLOROPHENOL	< 1900	<1600	<2200
1,3-DICHLOROBENZENE	<1900	<1600	<2200
1.4-DICHLOROBENZENE	< 1900	<1600	<2200
BENZYL ALCOHOL	< 1900	< 1800	<2200
1,2-DICHLOROSENZENE	<1900	<1800	<2200
2-METHYLPHENOL	< 1900	<1800	<2200
BIS (2-CHLOROISOPROPYL) ETHER.	< 1900	<1800	<2200
4-METHYLPHENOL	<1900	<1800	<2200
N-NITROSO-DI-N-PROPYLAMINE	< 1900	<1600	<2200
HEXACHLOROTHEANE	<1900	<1600	<2200
MITROBENZENE	<1900	<1000	<2200
SHOROHE	<1900	<(600	<2200
2-NITROPHENOL	< 1900	<1800	<2200
2,4-DIMETHYLPHENOL	<1900	<1800	<2200
BENZOIC ACID	< 1900	<1600	<2200
BIS (2-CHLOROETHOXY) METHANE	< 1900	<1600	<2200
2,4-DICHLORPHENOL	< 1900	<1800	<2200
1,2,4-TRICHLOROBENZENE	<1900	<1000	<2200
NAPHTHALENE	<1900	<1600	<2200
4 CHLOROANILINE	< 1900	<1800	<2200
HEXACHLOROBUTADIENE	< 1900	<1800	<2200
4 CHLORO-3-METHYLPHENOL	<1900	<1600	<2200
ZMETHYLNAPHTHALEHE	<1900	<1800	<2200
HEXACHLOROCYCLOPENTADIENE	<1900	<1800	<2200
2,4,E-TRICHLOROPHENOL	<1900	<1800	<2200
2,4,5-TRICHLOROPHENOL	<4700	<400G	<5300
-CHLORONAPHTHALENE	<1900	<1800	<2200
2-NITROANLINE	<4700	<4000	<£300
DIMETHYLPHTHALATE	< 1900	<1800	<2200
ACENAPHTHYLENE	< 1900	<1000	<5300
2,4-DINTROTOLUENE	< 1900	<1600	<2290
3-NITROANLINE	<4700	< 4000	<5300
ACENAPHTHENE	<1900	<1800	<2200

Units are in ug/kg (ppb)

TABLE 4-6 (Continued) WETLAND SOIL ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Semple ID Humber IEA Sample ID Humber	WL-1 9504289-04A	WL-2 9504289-05A	WL-3 8504209-05A
SEMI-VOL, ORG. COMPOUNDS			
2.4 - DINITROPHENOL	<4700	<4000	<5300
4-NITROPHENOL	<4700	<4000	<5300
DIBENZOFURAN	< 1800	<1600	<2200
2,4 - DINITROTOLUENE	<1900	<1600	<2200
DETHYLPHTHALATE	<1900	<1600	<5500
4-CHLOROPHENYL PHENYL ETHER	<1900	<1800	<2200_
FLUORENE	<1900	<1600	<2200
4-NITROANILINE	<4700	<4000	<5300
4,6-DINITRO-2-METHYLPHENOL	<4700	<4000	<5300
N-NITROSODIPHENYLAMINE (1)	<1900	<1000	<2200
4-BROMOPHENYL PHENYL EHTER	<1900	<1800	<2200
HEXACHLOROBENZENE	<1800	<1800	<2200
PENTACHLOROPHENOL	<4700	<4000	<8300
PHENANTHRENE	400J	<1600	300J
ANTHRACENE	<1900	<1600	<2200
CARBAZOLE	< 1900	<1800	<2200
DI-N-BUTYLPHTHALATE	< 1900	<1800	<2290
FLUORANTHÈNÈ	350J	230J	2501
PYRENE	450J	5007	420J
BUTYLBENZYLPHTHALATE	<1900	<1800	<2200
3,3' -DICHLOROBENZIDINE	<1900	<1600	<2200
BENZO(A)ANTHRACENE	2103	<1609	<2200
CHRYSENE	290J	<1600	270J
BIS(Z-ETHYLHEXYLIPHTHALATE	<1900	<1600	<2200
DI-N-OCTYLPHTHALATE	<1900	<1600	<5500
BENZO(B)FLUORANTHENE	250J	<1600	250J
BENZONGFLUORANTHENE	210J	<1800	<2200
SENZOJAJPYRENE	230	<1800	<2200
INDENO(1,2,3-CD)PYRENE	<1900	<1600	<2200
DIBENZO(A, H) ANTHRACENE	<1900	<1600	<2200
BENZOIG, H. & PERYLENE	<1900	<1600	<2200

NA - Analysis not performed

TABLE 4-6 (Continued) WETLAND SURFACE WATER ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number IEA Sample ID Number	WL-1 950428901	WL-2	WL-3 950426903
SEMI-VOL ORG. COMPOUNDS	100075001	830426902	930428903
PHENOL	<10	<10	<10
BIS 12-CHLOROETHYLETHER	<10	<10	<10
Z-CHLOROPHENOL	<10	<10	<10
I.3-DICHLOROBENZENE	<10	<10	<10
1.4-DICHLOROBENZENE	<10	<10	<10
BENZYL ALCOHOL	< 10	<10	<10
1.2-DICHLOROBENZENE	<10	<10	<10
Z-METHOLPHENOL	<10	<10	
813 (2-CHLOROISOPROPYL) ETHER	<10	<10	<10
4-METHYLPHENOL			<10
N-NITROSO-DI-N-PROPYLAMINE	<10	<10	<10
	مهرست والمساكد	<10	<10
HEXACHLOROTHEANE	<u><10</u>	<10	<10
NITROBENZENE	< <u>!D</u>	<10	<10
ISOPHORONE	<10	<u><1D</u>	<10
2-NITROPHENOL	<10	<10	<10
2,4-DIMETHYLPHENOL	<10	<10	<10
BENZOIC ACID	<10	<10	<10_
BIS (2-CHLOROETHOXY) METHANE	<10	<10	<10
2,4-DICHLORPHENOL	<10	<10	<10
1,2,4-TRICHLOROSENZENE	<10	<10	<10
NAPHTHALENE	<10	<10	<10
4 CHLOROANUNE	<10	<10	<10
HEXACHLOROBUTADIENE	<10	<10	<10
4 CHLORO -3-METHYLPHENOL	<10	<10	<10
ZMETHYLNAPHTHALENE	<10	<10	<10
HEXACHLOROCYCLOPENTADIENE	<10	<10	<10
2,4,6 - TRICHLOROPHENOL	<10	<10	<10
2,4,5-TRICHLOROPHENOL	<25	<25	<25
2-CHLOROHAPHTHALENE	<10	<10	<10
2-NITROANLINE	<23	<25	<23
DIMETHYLPHTHALATE	<10	<10	<10
ACENAPHTHYLENE	<10	<10	<10
2.0-DINITROTOLUENE	<10	<10	<10
3-NITROANS,INE	<25	<29	<25
ACENAPHTHENE	<10	<10	<10

Units are in ug/L (ppb)

TABLE 4-6 (Continued) WETLAND SURFACE WATER ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number	WL-1	WL-2	WL-3
IEA Sample IO Number	9504269~04A	9504289-05A	9504259-05
SEMI-VOL ORG COMPOUNDS			
2,4-DINITROPHENOL	<25	<25	<25
4-NITROPHENOL	<25	<25	<25
DIBENZOFURAN	<10	<10	< 10
2,4-DINTROTOLUENE	<10	<10	<10
DIETHYLPHTHALATE	< 10	<10	<10
4-CHLOROPHENYL PHENYL ETHER	<10	<10	< 10
FLUORENE	<10	<10	<10
4-NITROANILINE	<25	<25	<25
4,6-DINETRO-2-METHYLPHENOL	<25	<25	<25
N-NITROSCOPHENYLAMINE (1)	<10	<10	<10
4-BROMOPHENYL PHENYL EHTER	<10	<10	<10
HEXACHLOROBENZENE	<10	<10	<10
PENTACHLOROPHENOL	<25	<25	<25
PHENANTHRENE	<10	<10	<10
ANTHRACENE	<10	<10	<10
CARBAZOLE	<10	<10	<10
DI-N-BUTYLPHTHALATE	<10	<10	<10
FLUORANTHERE	<10	<10	<10
PYRENE	<10	<10	<10
BUTYLBENZYLPHTHALATE	<10	<10	<10
3.3'-DICHLOROBENZIONE	<10	<10	<10
BENZO(A)ANTHRACENE	<10	<10	<10
CHRYSENE	<10	<10	<10
BISIZ-ETHYLHEXYLIPHTHALATE	<10 '	3.7	< 10
DI-N-OCTYLPHTHALATE	<10	<10	<10
BENZO(B)FLUORANTIÆNE	<10	<10	<10
BENZO(K)FLUORANTHENE	<10	<10	<18
SENZO(A)PYRENE	<10	<10	<10
INDENO(1,2,3-CD)PYRENE	<10	<10	<10
DIBENZOIA HANTHRACENE	<10	<10	<10
BEHZO(G.H.SPERYLENE	<10	<10	<10

Units are in ug/L

TABLE 4-6 (Continued) WETLAND SOIL ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number	WL-1	WL-2	WL-3
IEA Sample ID Number	950469-04A	950469-05A	
PEST/PCBS COMPOUNDS			
ALPHA-BHC	<10	<8.5	<11
BETA-BHC	<10	< 8.5	<11
DELTA-BHC	<10	< 0.5	<11
GAMMA-BHC (LINDANE)	<10	<8.5	<11
HEPTACHLOR	<10	< 8.5	<11
ALDRIN	<10	<8.5	<11
HEPTACHLOR EPOXIDE	<10	<8.5	<11
ENDOSULFAN I	<10	< 8.5	<11
DIELDRIN	<19	<16	<22
4,4'-DDE	<19	<16	<12
ENDRIN	<19	<16	<22
ENDOSULFAN II	<19	<16	<22
4,4'-DDD	<19	<16	<21
ENDOSULFAN SULFATE	<19	<16	<22
4,4'-D0T	<19	<16	<22
METHOXYCHLOR	3.9JP	<85	<110
ENDRIN KETONE	<19	<16	<22
ENDRIN ALDEHYDE	<19	<16	<22
ALPHA-CHLORDANE	<10	<8.5	<11
GAMMA-CHLORDANE	<10	< 8.5	<11
TOXAPHENE	<1000	<850	<1100
AROCLOR - 1016	<190	<160	<220
AROCLOR - 1221	<390	<340	<440
AROCLOR - 1232	<190	<160	<220
AROCLOR - 1242	<190	<160	<220
AROCLOR - 1248	<190	81J	120JP
AROCLOR - 1254	<190	<160	<220
AROCLOR - 1260	< 190	<160	<220

Units are in ug/kg.

TABLE 4-6 (Conlinued) WETLAND SURFACE WATER ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number	WL-SW-1	WL-SW-2	WL-SW-3
IEA Sample ID Number	9504269-01A	9504269-02A	9504269~03A
PEST/PCBS COMPOUNDS			
ALPHA-BHC	< 0.050	< 0.050	< 0.050
BETA-BHC	< 0.050	< 0.050	<0.050
DELTA-BHC	< 0.050	< 0.050	< 0.050
GAMMA-BHC (LINDANE)	< 0.050	< 0.050	<0.050
HEPTACHLOR	< 0.050	< 0.050	< 0.050
ALDRIN	< 0.050	< 0.050	< 0.050
HEPTACHLOR EPOXIDE	< 0.050	< 0.050	< 0.050
ENDOSULFAN I	< 0.050	< 0.050	<0.050
DIELDRIN	< 0.10	<0.10	<0.10
4,4'-DDE	<0.10	<0.10	<0.10
ENDRIN	<0.10	< 0.10	<0.10
ENDOSULFAN II	< 0.10	<0.10	<0.10
4,4'-000	< 0.10	< 0.10	<0.10
ENDOSULFAN SULFATE	< 0.10	< 0.10	< 0.10
4,4'-DDT	< 0.10	< 0.10	< 0.10
METHOXYCHLOR	< 0.50	<0.50	< 0.50
ENDRIN KETONE	< 0.10	< 0.10	<0.10
ENDRIN ALDEHYDE	< 0.10	< 0.10	< 0.10
ALPHA-CHLORDANE	< 0.050	< 0.050	< 0.050
GAMMA-CHLORDANE	< 0.050	< 0.050	<0.050
TOXAPHENE	<5.0	<5.0	<5.0
AROCLOR - 1016	<1.0	<1.0	<1.0
AROCLOR - 1221	<2.0	<2.0	<2.0
AROCLOR - 1232	<1.0	<1.0	<1.0
AROCLOR - 1242	<1.0	<1.0	<1.0
AROCLOR - 1248	<1.0	<1.0	<1.0
AROCLOR 1254	<1.0	< 1.0	<1.0
AROCLOR - 1260	<1.0	₹1.0	<1.0

Units are in ug/L

TABLE 4-6 (Conlinued) WETLAND SOIL ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number	WL-1	WL-2	WL-3
IEA Sample ID Number	950426904	950426905	950426906
INORGANIC COMPOUNDS			
Arsenic	41,3	34.6	43,3
Barken	126	92.38	265
Cadmium	3.5	1.58	3.3
Chromium	11400	9410	14100
Hexavalent Chromium	<0.18	<0.17	<0.18
Lead	143	128	154
Mercury	2.6	0.77	0.81
Selenium	0.0	2.08	2.18
Silver	3.68	2.68	2.1B

Units are in mg/kg

MWG13-15_46678

DEBUGO. DETAG

TABLE 4-5 (Conlinued) WETLAND SURFACE WATER ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID Number IEA Sample ID Number	WL-SW-1 950426901	WL-SW-1F	WL-SW-2	WL-SW-2F 950426902F	WL-SW-3 950426903	WL-SW-31
INORGANIC COMPOUNDS	950426901	8304208016	830420802	330420802F	830420803	1204208031
Arsenic	<3.0	<3.0	<3.0	<3.0	<3.0	<3.0
Barlum	45.48	42.88	42.9B	39.48	40.1B	38.18
Cadmium	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
Chromium	37.3	6.5B	3.68	1.3B	2,98	<1.0
Hexavalent Chromlum	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
Lead	2.38	2.6B	<2.0	<2.0	<2.0	<2.0
Mercury	< 0.20	<0.80	< 0.20	< 0.40	<0.20	< 0.40
Selenium	<3.0	<3.0	<3.0	<9,0	<3.0	<3.0
Silver	<1.0	<1.0	<1.0	<1.0	<1,0	<1.0

Units are in ug/L

TABLE 4-7 AVS and SEM ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOIS

Sample ID	Cadmium	Copper	Lead	Mercury	Nickel	Zinc
WL1#1	4.473	54.00	209.4	0,89	90.74	740.2
WL-1#2	4.068	27.21	139.9	0.00	38.54	502.9
WL-2#1	3.598	56.03	205.1	0.20	13.72	272.2
WL-2#2	3.966	44.68	183.9	0.19	12.79	_303,B
WL-3#1	2.506	31.41	72.2	0.12	10.24	128,4
WL-3#2	2.378	29.02	72.0	0.05	25.05	149.7

Units are in mg/kg

Sample ID	Cadmium	Copper	Lead	Mercury	Nickel	Zinc	SEM
WL-1#1	0.04	0.850	1.011	0.004	0.524	11.32	13.75
WL-1#2	0.036	0.428	0.675	0.000	0.857	9.22	11,02
WL-2#1	0.032	0.913	0.990	0.001	0.234	4.18	6.33
WL-2#2	0.035	0.703	0.868	0.001	0.218	4.65	6.49
WL-3#1	0.022	0.494	0.348	0.001	0.174	1.96	3.00
WL-3#2	0.021	0.457	0.347	0.000	0.427	2.29	3.54

SEM = Cd+Cu+Pb+Hg+Ni+Zn Units are in micromoles per gram

1	Sample ID	SEM 1	SEM 2	Ave. SEM
i	WL-1	13.75	11.02	12,39
	WL-2	6,33	6.49	6.41
	WL-3	3.00	3.54	3.27

Units are in micromoles per gram

Sample ID	AV8 1	AVS 2	Ava. AVS
WL-1	23.2	19.35	21.28
WL-2	1.41	1.36	1.39
WI _3	2.89	3.07	6.04

Units are in micromoles per gram

Semple ID	W dry/ W wel	Ave. AVS		Ave. SEM/ Ave. AVS
WL-1	0.18	21.28	12.39	0.58
WL-2	0.18	1.39	8.41	4.61
WL-3	0.25	2,98	3.27	1.10

Units are in micromoles per gram

MWG13-15_46680

PS45010008ea

TABLE 4-7 (Continued) WETLAND SOIL ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONWEALTH EDISON COMPANY WAUKEGAN, ILLINOS

Sample ID Number IEA Sample ID Number	WL-1	WL-2	WL-3 NA
SOIL CHEMISTRY	-		
TOC	140000	270000	330000

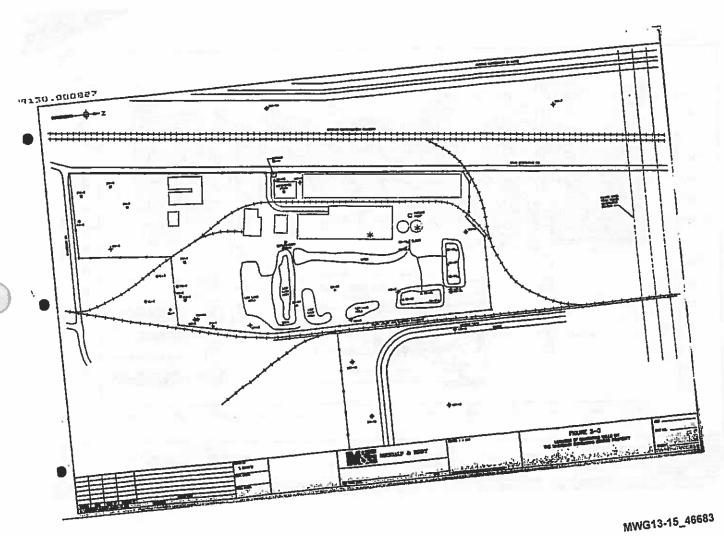
Units are in mg/L (opm)

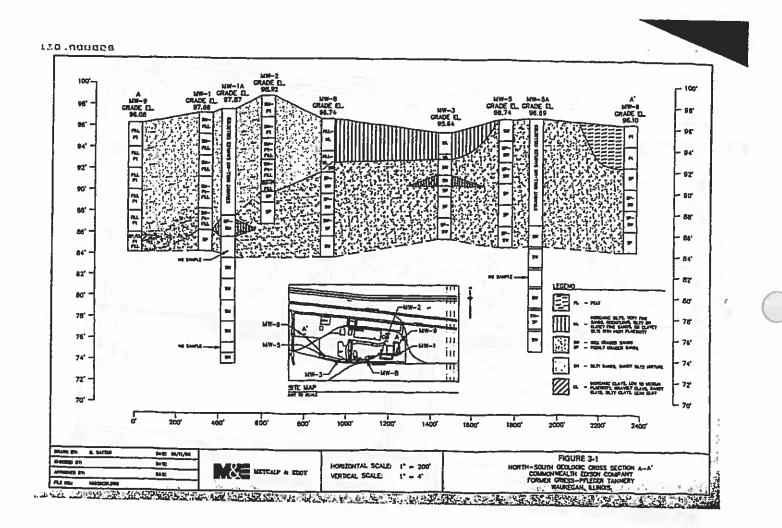


TABLE 4-7 (Continued) WETLAND SURFACE WATER ANALYTICAL RESULTS FORMER GRIESS-PFLEGER TANNERY COMMONVEALTH EDISON COMPANY WAUKEGAN, ILLINOS

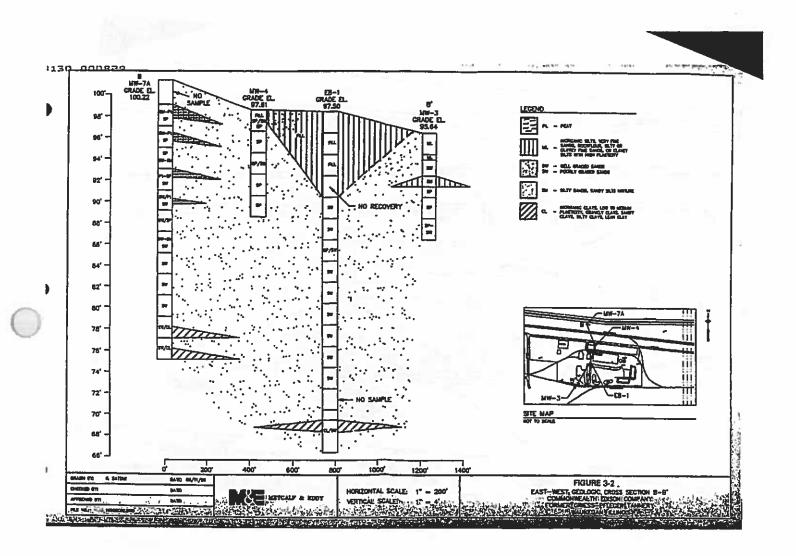
Sample ID Number	WL-SW-1	WL-8W-2	WL-SW-3
IEA Sample ID Number	NA	NA	NA
WET CHEMISTRY			
TOC	12	9,2	5.9
14.4 . 1 . 4 . 1			

Units are in mg/L (ppm)

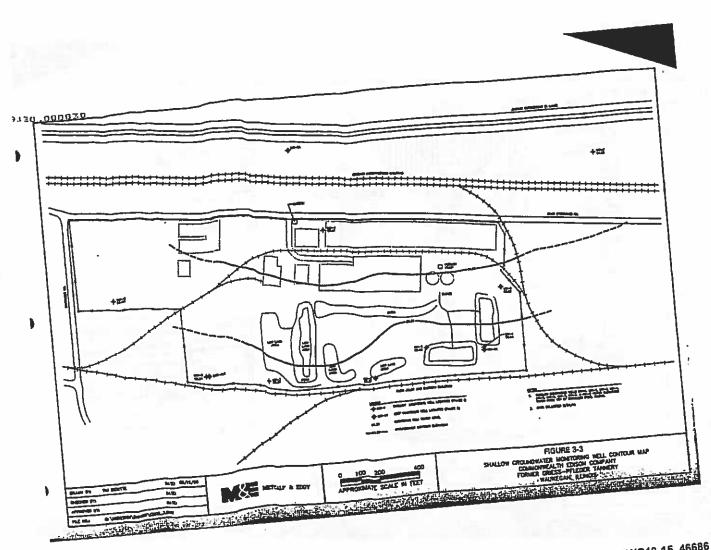




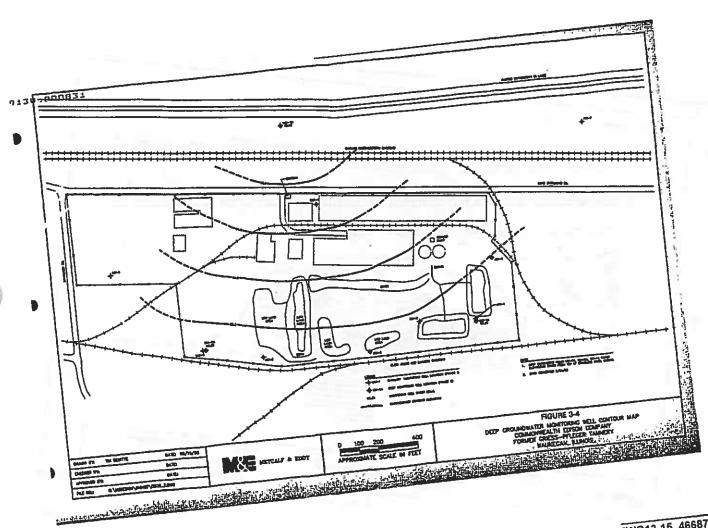
MWG13-15_46684



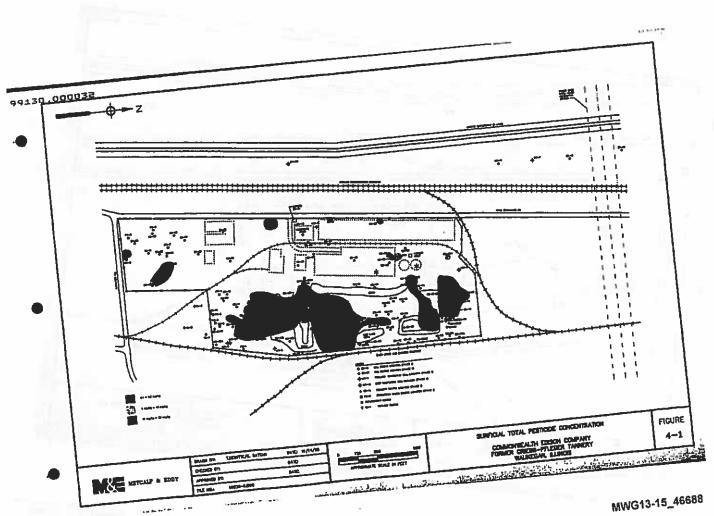
MWG13-15_46685

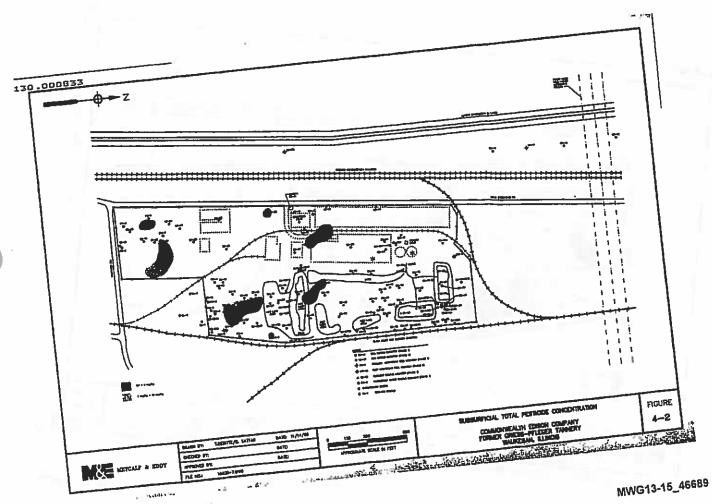


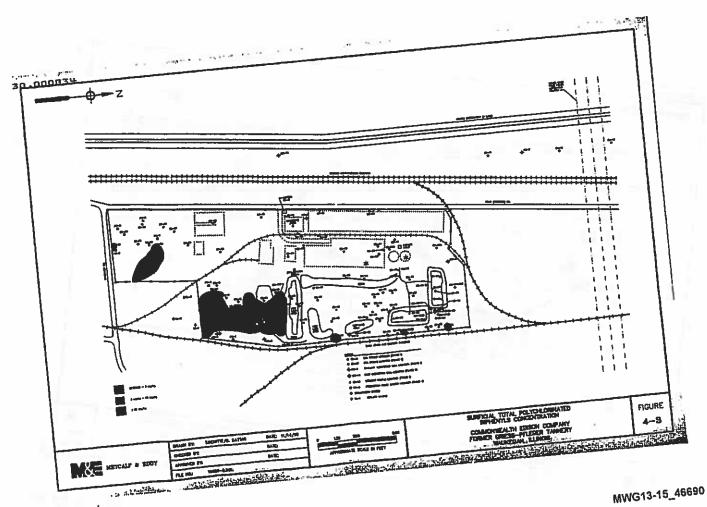
MWG13-15_46686

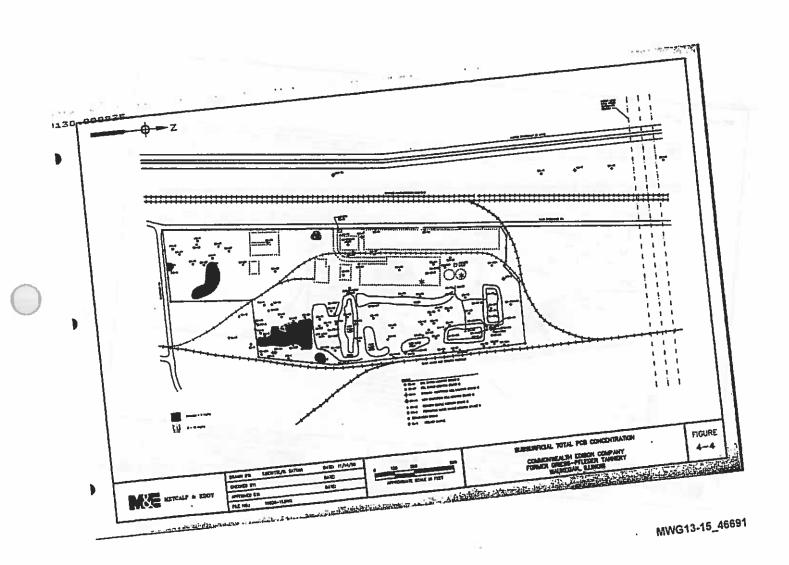


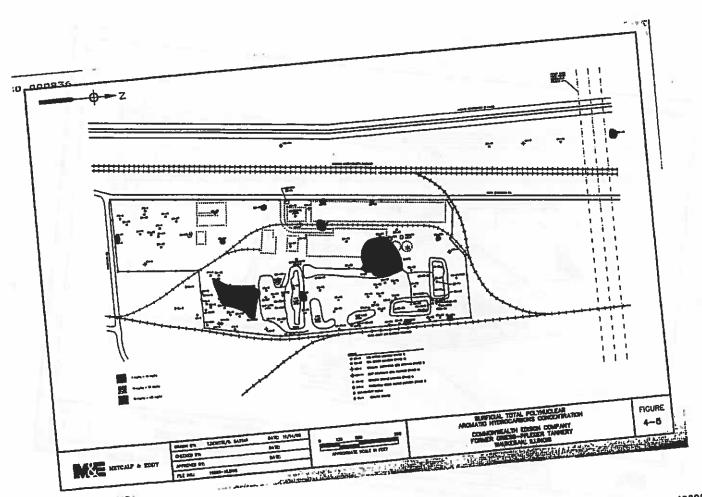
MWG13-15_46687



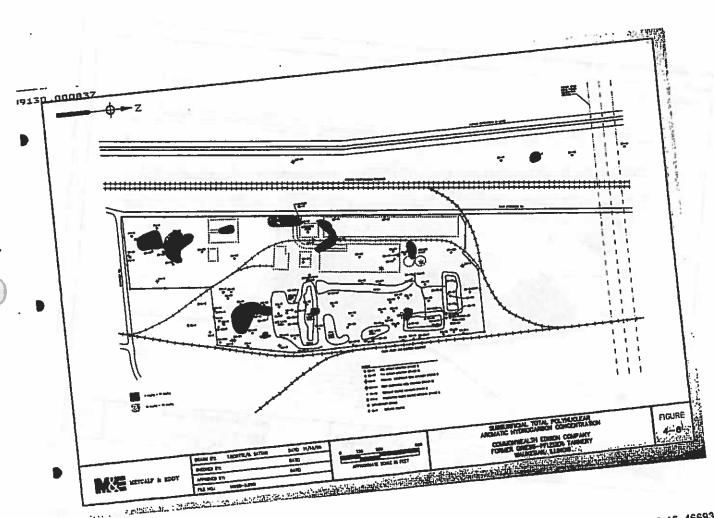




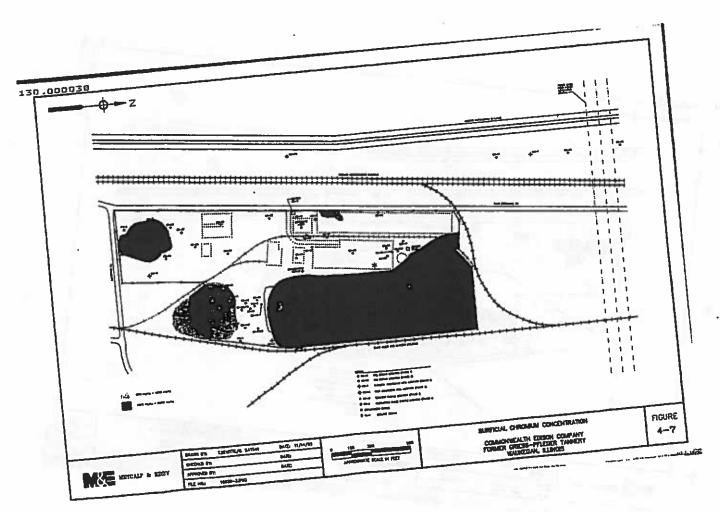




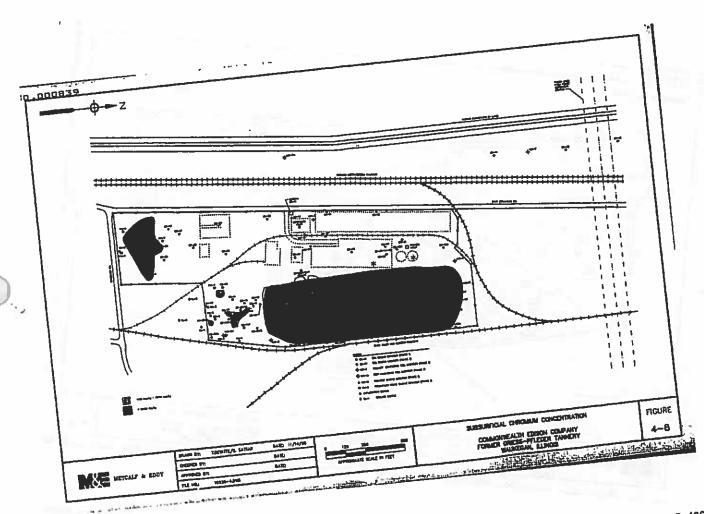
MWG13-15_46692



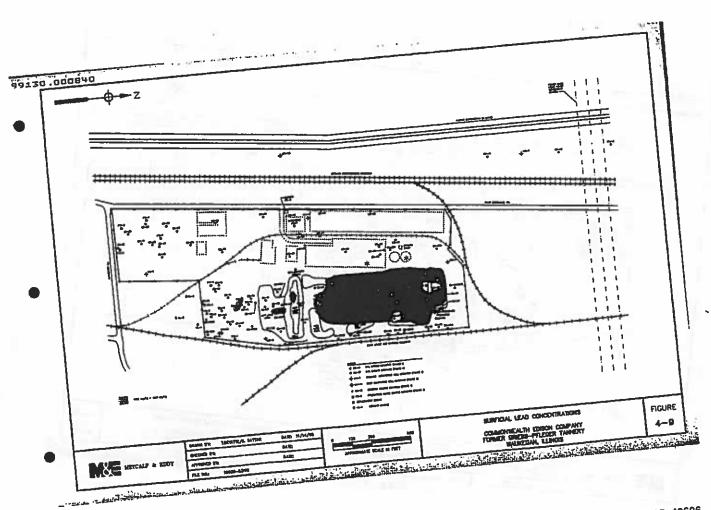
MWG13-15_46693



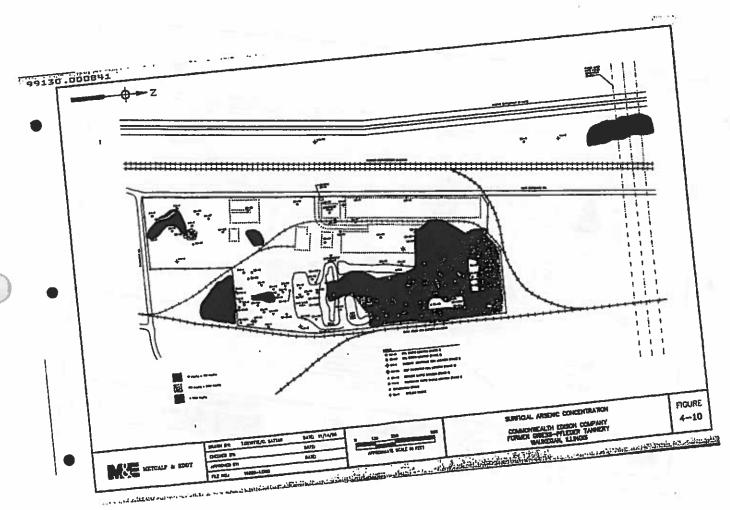
MWG13-15_46694



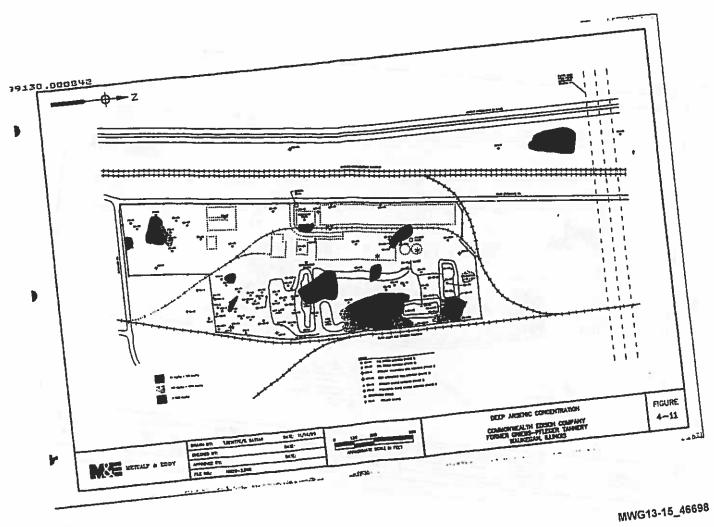
MWG13-15_46695



MWG13-15_46696



MWG13-15_46697



	F	CATIO	N				UOB NO. 016920		488				
4	D/	HRING	ER I	ROAD:	WES	OF	OF EJAE R.R. WAUKEGON -						
	U (CNTR	ACTO	R WT	D-80	ART	LONGYEAR ENG/GEO: STORY BEGUN: 1-3	95					
		IG:						130	One.				
	4.2		**.*	LLOW				MO	20' F				
DEPTH (C.T.)	SALPIE TYPE/NO.	SAMPLE	SALPLE RECOVERY	BLOW COUNT (per 8 inches)	RECOVERY OR ROD.	0 (med	SAMPLE DESCRIPTION	Caupalic					
٦	1	1-3	24	10/16/	100		BROWN TO BLACK, SATURATED LOOSE, GRANULAR SLAG-LIKE						
	SS						MATERIAL (SATURATED 2.5-3' B.G.)						
	2	3-5	12	7/7/10	50	-	BLACK, SATURATED LOOSE GRANULAR SLAG-LIKE MATERIAL WITH SOME SAND FILL		.FJĞ				
1	55	5-7	0	29/9/	o		NO RECOVERY						
-	3 SS	5-/	0	29/9/	-								
-	4	7-9	10	15/%/	42	-	GREY, SATURATED, LOOSE, FINE TO MEDIUM SAND WITH TRACE						
1	SS		- 3				COARSE SAND, SUBROUNDED, POORLY SORTED						
	5 SS	9-11	12	15/1/1	50	-	GREY TO BLACK, SATURATED, LOOSE, FINE TO MEDIUM SAND (SUBROUNDED) WITH SOME C. GRAVEL AND SOME PEBBLES. POORLY SORTED						
+	6 S\$	11-13	24	15/7/10	100	-	TOP 13"-DARK GREY, SATURATED, LOOSE, F. SAND W TRACE C. SAND & TRACE PERRIES (MODERATELY WELL SORTED) MEDICLE 4"-GRAY, SATURATED LOOSE, SUBMANGULAR GRAVEL AND SLAG, PEICE OF LEATHER (2" ONA & 1/4" THICK), TOP 4"-GRAY, SATURATED, LOOSE, F. SAND W/ SOME C. SAND.		SP/S				
	7 SS	13–15	14	17/25 5003	58	-	GREY, SATURATEDM SLIGHTL DENSE, F-M SAND, WITH TRACE C. SAND, POORLY SORTED, SUBROUNDED		SW				
1	_	15–17	16	4/21 6/17	67	-	GREY, SATURATED, MODERATELY DENSE, F-M SAND, W/ TRACE		SW				
1	SS						SAND, TRACE PEBBLES, POORLY SORTED, SUBROUNDED.		J.,				
	g SS	17–19	24	15/25 32/60	100	10	GREY, SATURATED, MODERATELY DENSE, F-M SAND, W/ TRACE C. SAND & F. GRAVEL, POORLY SORTED, SUBROUNDED.		5W				
-	10 SS	19-21	24	9/19 70/55	100	-	SAME AS ABOVE		sw				
	11	21-23	24	36/50	100	Ξ	DARK GREY IN TOP 12" & LIGHT GREY IN BOTTOM 12", SATURATED MODERATE. DIDISE, F-M SAND, W/ TRACE C. SAND AND F. GRAVEL, POORLY SORTED, SUBROUNDED.		SW				
	SS 12	23-25	24	11/30 5003	100	-	GREY, SATURATED, MODERATELY DENSE, F-M SAND, W/ TRACE C. SAND & F. GRAVEL, POORLY SORTED, SUBROUNDED.		SW				
	55							-	-				
		_					STRAIGHT DRILL, NO SAMPLES COLLECTED						
	13 S\$	26-30	24	12/16/ 21/56	100	Ξ	GRAY, DENSE, WET, SLIGHTLY PLASTIC SILTY CLAY (TOP 4") GREY SATURATED, MOD. DENSE, F-M SAND W/ TRACE F. GRAVELL, POORLY SORTED.		a/s				
1	1				200			1	2.0				

		et. C		WW.	T.TH	EDISC	N			SHEET	ORING	
31		OCATIO						JOB NO.	016920	1	MW.	
i	N D	OR THE	GER	CORN ROAD	ER O	F SA T OF	ND AND EJ&E R.R.	LOCATION:	EGON		OTAL	
18							LONGYEAR	ENG/GEO:	STORY	97.87 BEGUN: 2-	25	2.0
	ILL.			BILE				DRILLER:	ERIC		2-95 2-95	₹1
IC		SIZE:		LING.			GRND.	WATER (DE				
	4.2	!5 T	HC	MOTIK			GER 6.49	(TOC) / 9	3.30 (TOC)	CLEAR, COLD,	MID.	7 -
DEPTH (FT.)	SAMPLE TYPE/NO.	Junes Junes	SALPLE	(per 6 inches)	M RECOVERY CM RGD.	(W-4)	STORES	SAM DESCR			CRAPHIC	CLASSFICATION
							STRAIGHT DRILL Y	WITH HOLLOW STE	M AUGER TO 10 ROWN ADJACENT	FT. B.G. MONITORING		
					-	1/2	WELL MW-1A				1	
			1371									
-			- 27									
											71	
		111										
			H V									
-	1	10-12	18	3/10/	75	4	5"-MED. GREY, W SORTED W/ SILT,	ET, SUBROUNDED, BOTTOM 13"-GRE	VERY DENSE, M- YISH BROWN, WET	C SAND, FAIRLY WELL , POORLY SORTED, SOME SILT		SP-SI
4	SS	12.14		35/		0.2		·		SOME SILT		3-34
ŀ	2 SS	12-14	0	5064	0	0.3	NO SAMPLE, SAMP	TEL WASSIEL FROM	35/11 25:00M			
ŀ	3	14-16	24	7/17	100	0.9	TOP 10"-MED. ORI TRACE C-VC SAN	EY, WET. V. DENS	E, SUBROUNDED	M. SAND W/		
1	SS		-	20/40			TRACE C-VC SAN SAME AS ABOVE, TO V. FINE SANO	D. SUBANGULAR Y COLOR CHANGE Y	V/ SOME SILT. BK ELLOWISH BROWN	TTOM 14"— FINING DOWN		SM
1	4	16-18	24	¥%4	100	0.3	MED. CREY CRADIN	IG TO YELLOW BR	DWN, WET, MODE	RATELY DENSE, V. FINE		
I	SS			-,,,,			SARU W/ F-H SA	MU, SUBMQUNDED,	MODERATELY WE	IL SORTED, SOME SILT.		SM
	5	18-20	24	3/4/7	100	1.0	OLIVE GREY TO BE C. SAND W/ SILT.	TOWNISH GREY, LC SUBROUNDED. TR	OSE TO MEDIUM	DENSITY, V. FINE TO EVBANGULAR TRACE		6
	55			177			HAIR, ROOTS.					SM
-		20-22	24	8/11	100	_	IV. FINE TO M. SUE	SROUNDED SAND!	SH BROWN, WET, N/ SILT, TRACE I	LOOSE TO MEDUM DENSI	ח	SM
-	SS						SULFUR ORGANIC S	See CIT				
ŀ	7	23-25	24	3/2/17	100	_	OUVE BROWN TO B	ROWN, DARK GRE	Y, DENSE FINE 1	O V. FINE SUBANGULAR	-	
+	SS		-7	12/17	100		OR SAND W/ SILT.	GRADING TO A V	. FINE SAND TON	ARD BOTTOM.		SM
ŀ											_	-
İ												1.1
	1			_								
L												
L										7- 7/4-1		
H	_				_							
Į.	491 F	TYPE	·e -				NOTES:				<u> </u>	

130 0000H7

GEOLOGIC LOG

_	E LO	T: CO CATIC ORTHE AHRING	N: AST	CORN	ER O	F SAI	N ND AND EJ&E R.R.	ND LOCATION: GRND. ELEV. TOTAL				RING MW- TAL (SA SEPTH	
DR		ONTR	ACTO	R: WT	D-BC	ART	LONGYEAR	ENG/GEO:	STORY	BEGUN:	1-31	-95		
	ILL I			BILE			TV	DRILLER:	ERIC		1-31	-95		
HO	LE S	IZE:		LING					EPTH/ELEV.):	WEATHER:				
	4.2	5	HO	LLOW	STEN	A AUG			93.26 (TOC)		DY.	MID.	20° F	
SAUPLE	SAMPLE TIME	SALPLE	SAMPLE	BLOW COUNT (per 6 inches)	X RECOVERY OR ROD.	QL (wdd)		SAMPLE DESCRIPTION						
							STRAIGHT DRII GEOLOGIY ALR WELL MW-5.	L WITH HOL EADY DETER	OW STEM AUGE MINED FROMM A	R TO 10 FT. B.G DJACENT MONITO	RING			
	1 SS	1012 125-145		47/70/ 27/30 -/7 32/50		0	GREY SATURA	Y SORTED, S	UBROUNDED.	C. SAND, MODER M SAND W/ TRA Y POORLY SORTE	CE		SW SW	
	3 55	15–17	10	10/16 21/27	42	0	SAME AS ABO	VE						
	4	17.5-19.5	12	7511/8	50	-	GREY, SATURA F. GRAVEL. PO	TED. MODERA	ATELY DENSE, F D, SUBROUNDED	-C SAND, TRACE	OF		SW	
	5 SS	20-22	24	2/12/ 10/14	87.5	-	SUBROUNDED, BOT	TOM 4" - CRE	, SATURATED, MODI	/ Trace F. Gravel, Erately Dense F. S MATELY POOPLY SORT	CHA		SW-SP	
	_	225-245	20	15/17/ 21/30	83	-	GREY, SATURA SAND, SUBROU	TED, MODER	ATELY DENSE, P	OORLY SORTED.	FC		5W	
	SS		_											
S/	MPL SS=	E TYP	ES SPOC	N, S1	=SH(LBY	NOTES TUBE DRILLE	D TO 26' B.	o. (1 FOOT OVE	R 25') BECAUSE INSTALLED TO	OF .			

Vetcalf & Sale

	PROJECT: COMMONWEALTH EDISON SITE LOCATION: JOB NO. 016920											RING MW-	
Ş				CODM	ED O	E SAI	ND AND	JOB NO.	016920	GRND. ELEV.		AL I	
							EJ&E R.R.		KEGON	100.22		25	
DR		CONTR	ACTO	R: WT	D-80	ART	LONGYEAR	ENG/GEO:	STORY		-31		
_	DRILL CONTRACTOR: WTD-BOART DRILL RIG: MOBILE DRILL - A						\TV	DRILLER:	ERIC	FINISHED: 2	-1-	95	
HC	LE 9	SIZE:		LLOW					PTH/ELEV.): 95.62 (TOC)	WEATHER: CLE			
DEPTH (FT.)	SAUPLE TYPE/NO.	SAUPLE	SAMPLE	BLOW COUNT (per 6 hohes)	X RECOVERY OR ROD.			SAN	IPLE RIPTION			DBAPHIC LOG	
	1	1-3	21	3/4/6/	87.5	0	TOP 17"-GREY TO	D SLACK, DAMP.	SOFT SILTY SAND	W/ TRACE PEBBLES, V	"/_		-
	SS			7	0,.0	-	FIBROUS ORGANIC MOIST WELL SORT	: Matter (Peat). Ted, M-Sahd, 50	BOTTOM 4"-TAN	TO DARK BROWN, SUG	HILY		2M-
	2	3-5	24	1/3/4/	100	0	TOP 9"-GREY &	BLACK, DAMP SIL	TY SAND, SOFT, W	ORGANIC MATERIAL			
	SS			-			W/TRACE C. SAN SUBROUNDED.	D. BOTTOM 9"-BI	JK, MOIST, SOFT, Y	RLY SORTED F-M SAN ELL SORTED ML SAND			SM-
	3	5-7	24	3/3/4/	100	0	TOP 5"-TAN, MO	ST, SOFT, WELL !	SORTED, MEDIUM S.	AND, SUBROUNDED.			9
	SS						SORTED. GRADE 1			COENAIELT POORLT			570
	4	7-9	10	4/10/	42	0	TOP 4"-BLK, MOIS (SUBROUNDED). BO	T, SPONGY ORGAN OTTOM 6"-TAN, S	HIC MATTER (PEAT, ATURATED, LOOSE,	W/ F. SAND F-C SAND W/ F. GRA	AVEL	ш	PT-
*	SS 5	9~11	20	3/%	83	0	TAN. SATURATED.	LOOSE, SOFT, F-	C SAND, TRACE P	EBBLES, POORLY SORT T), TAN, SATURATED !	TED.		SW,
	SS	11-13	24	3/9/7	100	0				F-C SAND & F-M GR			S
	SS	11-13	2.7	11/17	100	-	BOTTOM 12"-BUK SORTED, SUNROU	TO BRN. SATURA	ITED, MODERATELY	DENSE, F. SAND, WELL	-		2M
	7	13-15	19	11/17/	24	0	GREY SATURATED	POORLY SORTED	, F-C SAND W/ F	CRAVEL 1" MOIST,			SW/
	SS		,				LOOSE F-V C. SA	ND & F-M CRAV	EL SUBROUNDED.	TED, POORLY SONTED,			Ś
	8	15-17	24	19/33	100	0	PEBBLES, GRADINI	G TO BUK-GRY, F	NSE, POORLY SORT	ED. F-C SAND GRAVE E F. GRAVEL, MODERA	TELY		Š
	SS			14 22			POORLY SORTED.		COOTER C A A	3 W / 10 A 6 ***	1		Ľ
	9	17–19	24	15/18 22/30	100	0	SUBROUNDED.	, SOFT, POORLY	auries, r-G SAM	D W/ TRACE F. GRAVE			s
	10	10-21	22	7/3/	92	0	GREY TO TAN. SA	TURATED F-M S	UND W/ TRACE C.	SAND & F. GRAVEL S	UÐ	-	H
	SS			1/3/			RUNDED POORLY	SORTED, GRADE 1	no coarser matei	HALS. GREY TO TAN, -C SAND W/ GRAVEL			s
		21-23	24	12/18 20/27	100	0	TRACE PEBBLES. GREY, SATURATED	, MODERATELY D	ENSE, POORLY SOR	TED, F-C SAND W/		-	٢
	SS			20/2/			F. GRAVEL SUGH	L BOTTOM 4" V. TLY TO NON - P	SIPP, MUIST, SILT LASTIC.	LAT W/ IHAGE			SW
	12	23-25	20	4/5/8/	83	0	GREY, SATURATED	MODERATELY OF	ENSE, F-C SAND Y	/ TRACE F. GRAVEL.	ST.		5W
	SS				11.00		SILTY CLAY W/ TI						3"
·												Ţ	

SAMPLE TYPES SS=SPLIT SPOON, ST=SHELBY TUBE

Metoalf & Bady

NOTES: MONITORING WELL INSTALLED AT 24' B.G. FLUSH MOUNT DUE TO IDOT R.O.W. SPECIFICATIONS.

PROJECT: COMMONWEALTH EDISON SITE LOCATION: NORTHEAST CORNER OF SAND AND DAHRINGER ROAD, WEST OF EJ&E R.R.									JOB NO. 016920 LOCATION: WAUKEGON			GRN			ORING N MW-8 OTAL DEI		
R	u c	ONTR	ACTO	R: WT	D-80	ART	LONGYE	AR	ENG/GE	_	TORY	BEG	UN:	2-1-			
_	LL F			BILE					DRILLER		RIC		SHED:	2-1-			
10	4.2			LING LLOW					WATER (TOC)		H/ELEV.		D. TO U			F	
DEVIN (F1.)	TYPE/NO.	SAMPLE	SAMPLE	BLOW COUNT (per 6 inches)	A RECOVERY OR ROD.			SAMPLE DESCRIPTION							OBAPHEC LOG	CLASSFICATION	
	1 SS	1-3	3	YX.	13	0.6	BROWN-OU LEATHER LO	VE, M COKING	XST, V. STIFF) W/ SALT, E	FIBROL	IS MATERIAL UND 1.7 ppm	(ALMOST	LEATHER	r (WORN		FILL	
	2 SS	3-5	18	16%	75	0.4			TO BROWN TO BOTTOM 10" - D, F. GRAVEL							FILL ML, SW	
	3 SS	5-7	15	1/3/4	62.5	0.4	F. GRAVEL	SOME								SP-	
$\frac{1}{1}$	4 SS	7-9	22	10/24 30/25	92	1.4	PAT 8.5 H.C.		TURATED, MC Y POORLY S), MODERATE							SP-	
	5 SS	9-11	19	16/24	79	-	CREY, SATU SUBROUNDED (LESS COAR	SE MA	MODERATEL SAND. BECC TERIAL) TURATED, MC							sw-	
-	SS	11-13	24	15/17 23/30	100	.8	F-C SAND,	TRACE	OF GRAVEL	TRACE	OF PEDBLES					SW	
	·											H					
-			-					-								-	
\dashv				-												+-	

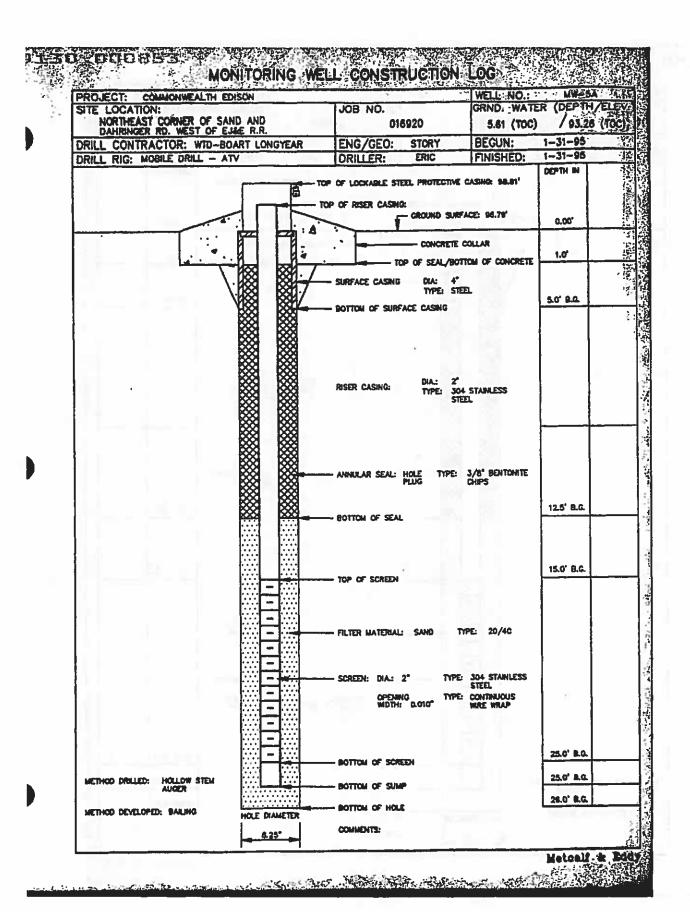
SAMPLE TYPES SS=SPLIT SPOON, ST=SHELBY TUBE

NOTES:
END OF BORING 13'. MONITORING WELL INSTALLED
TO 13' B.G.

Metoalf & Ridge

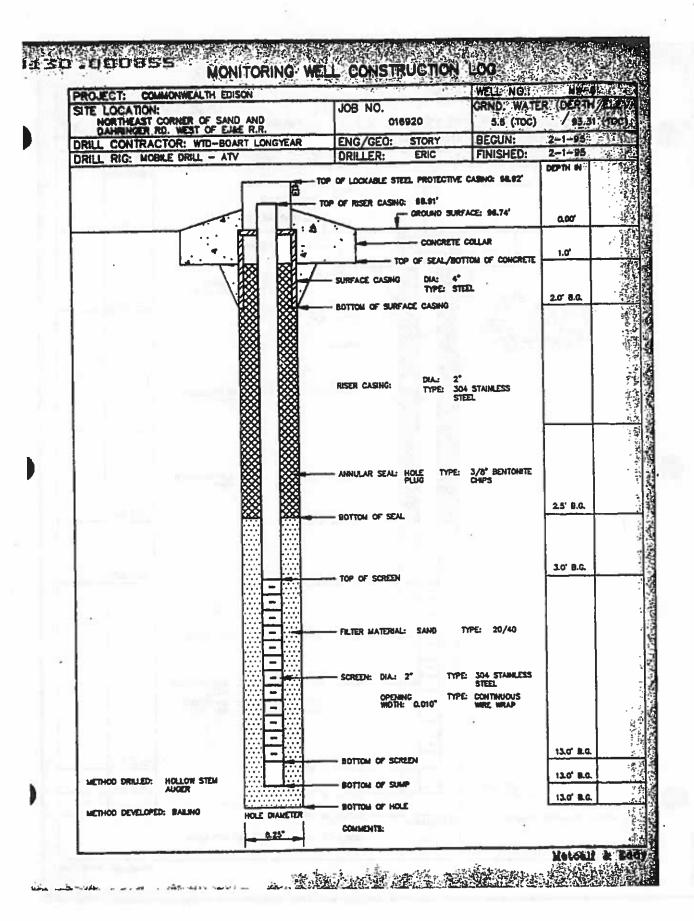
		T: CC		SHEET	BORI								
SIT	•	CATIC				Turi.		JOB NO.	016920	1	TOTA	W-	- 111
	NORTHEAST CORNER OF SAND AND							LOCATION:	(F00)	GRND. ELEV.			EF.
	DAHRINGER ROAD, WEST OF EJ&E RILL CONTRACTOR: WTD-BOART LONGY								KEGON	_		13'	
								ENG/GEO:	STORY		2-2-9 2-2-9		-
	ILL F	-		BILE				DRILLER:	ERIC				V
110	HOLE SIZE: DRILLING METHOD: 4.25 HOLLOW STEM AUG							93.86 (TOC)					
DEPTH (FT.)	SALPLE TYPE/NO.	SAMPLE	ANSAOOSH STARTYS	BLOW COUNT (per 8 hiches)	X RECOVERY OR ROD.	(web) Old	Harry		APLE RIPTION		CRUDHEC	991	
										CT 11 CT 1100TO			
	1	1-3	17	,	71	0	PLASTIC SELTY CI	LAY W/ FIBROUS	(HIDE) MATERIAL	iet, v. stiff, modera (sewer-like/petroli	EUM		FI
	SS			·			ODOR)	* U P/51/64	- CH APPA CITY	NAV W ~ AV W/			
	2	3-5	4	15/3/	17	0	ROOT MATTER.	r, v. Dense, Non	-rladiig SLIT (CLAY TO CLAY W/			FI
	SS			100									
	3	5-7	24	13%	100	0	BLK, WET, V. SOF ODOR), GRADE TO	FT, SILTY CLAY W CREY//BLK. V.	/ HAIR & HIDE M SOFT. WET, SLIGH	aterial (Sever-Swai Itly Plastic, Silty C	LAY		FI
	SS			6/12	100								-
	4	7-9	24	8/12 26/28	100	0	BOTTOM 8 -GRE	Y, SOFT, SUGHT	DERATELY DENSE	CLAY W/ HAIR MATE , MODERATELY WELL	BOAL.		n
	SS 5	9~11	24	12/12	100	Ò							H
	SS	9~11	24	12/12 24/28	100	0	(SEWER-SWAMP (ODOR) 2"-TAN S. RACE COARSE SA	ATURATED, MOD. 1	TH HAIR MATERIAL DENSE, POORLY SORTE DARK GREY, SATURAT	D. ED	Į.	FI
				8/18	455		WELL SORTED, SL	JBROUNDED F. SA	NO. DENSE WELL	SORTED F. SAND. S	UB-	_	H
	5 5 5	11-13	24	6/18 28/35	100	0	ROUNDED, 5"-BL	k, saturated, p R-Swamp Goor). ED F. Sand Graf	LASTIC, V. SOFT,	SILTY CLAY W/ HAR LY, SATURATED MOD. I TEXTURE SECUENC —	DENSE.		0
			1 =										
							-						
													H
	—		-										
	_	-		-		-	7/						
							1						
				-									r
						-							
				_			1						
						*							
S	MPL SS=S	E TYP SPLIT	ES SPOC	N, S1	r=she	LBY	TUBE THE INST	DE OF THE AUGE	RS FROM THE STI	r was lost in the i cky silty clay so it , not bridge in the	HAT THE	10	a

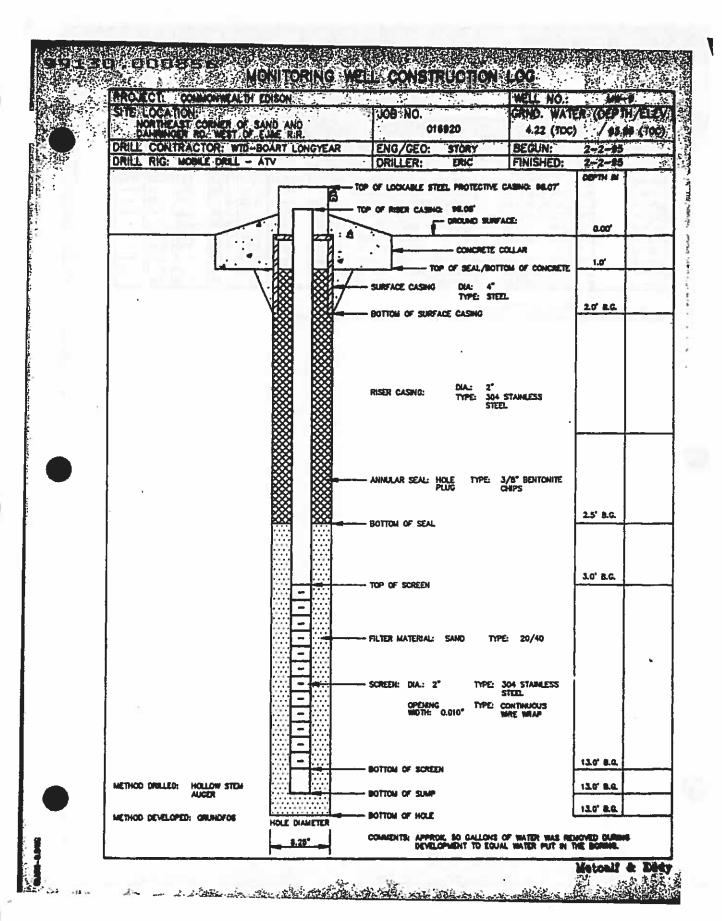
Marketine in the second residence in the contract of the second s 130.000852 MONITORING WELL CONSTRUCTION LOG MW-1A PROJECT: COMMONWEALTH EDISON WELL NO .: GRND. WATER (DEPTH/ELEV) SITE LOCATION: JOB NO. NORTHEAST CORNER OF SAND AND DAHRINGER RD. WEST OF EJ&E R.R. 016920 6.49 (TOC) /93.30 (TOC) 2-2-95 BEGUN: DRILL CONTRACTOR: WTD-BOART LONGYEAR ENG/GEO: STORY 2-2-95 DRILL RIG: MOBILE DRILL - ATV FINISHED: DRILLER: DEPTH IN TOP OF LOCKABLE STEEL PROTECTIVE CASING: 99.82" TOP OF RISER CASING: 99.78° CROUND SURFACE: 87.87 0.00 CONCRETE COLLAR 1.0 TOP OF SEAL/BOTTOM OF CONCRETE SURFACE CASING DIA: DIA: 4" TYPE: STEEL 5.0' 8.0. BOTTOM OF SURFACE CASING 2° 304 STANLESS RISER CASING: TYPE: STEEL ANNULAR SEAL: HOLE PLUG 13.0° B.G. ROTTOM OF SEAL 15.0" B.C. TOP OF SCREEN TYPE: 20/40 FILTER MATERIAL: SAND SCREEN: DIA: 2" TYPE: 304 STAINLESS STEEL OPENING WOTH: 0.010° TYPE: CONTRIBOUS 25.0° Q.C. BOTTOM OF SCREEN 25.0' B.C. METHOD DRILLED: HOLLOW STEM AUGER BOTTOM OF SUMP 25.0° B.C. BOTTOM OF HOLE METHOD DEVELOPED: BAILING HOLE DIAMETER COMMENTS: Metcalf & Eddy



9430.000854 MONITORING WELL CONSTRUCTION LOG MM-7A PROJECT: COMMONWEALTH EDISON WELL NO .: SITE LOCATION:
NORTHEAST CORNER OF SAND AND
DAHRINGER RD. WEST OF EJIE R.R. GRND. WATER (DEPTH/ELEY) JOB NO. 016920 ·/ 95.62 (TOG) 4.81 (TOC) STORY 2-1-95 DRILL CONTRACTOR: WTD-BOART LONGYEAR BEGUN: ENG/GEO: DRILL RIG: MOBILE DRILL - ATV 2-1-95 ERIC FINISHED: DRILLER: DEPTH IN DOWNDARLE LOCKING CAP . TOP OF SURFACE CASING: 100.67 TOP OF RISER CASING: 100.43" GROUND SURFACE: 100.22" 0.00 CONCRETE PAD 8" ALUMINUM FLUSH MOUNT SURFACE CASING TYPE 6° OF SAND PLACED ON TOP——
OF PELLETS TO HELP PREVENT
FROST HEAVE OF THE CEMENT
COLLAR. A MAGNET WAS PLACED
RISDE AND ON TOP OF THE SAND
FOR EASE OF LOCATION IN CASE
IT WAS COVERED WITH SNOW. 1.0' B.G. BOTTOM OF SURFACE CASING TOP OF SEAL BACKFILL TYPE: DIA.: TYPE: 2" 304 STAINLESS STEEL RISER CASING: TYPE: 3/8" BENTONITE CHPS ANNULAR SEAL: HOLE PLUG 120' B.C. BOTTOM OF SEAL 14.0' B.G. TOP OF SCREEN FILTER MATERIAL: SAND TYPE: 20/40 SCREEN: DIA: 2" 304 STAINLESS STEEL TYPE: CONTINUOUS OPENING WOTH: 0.010" 24.0" B.G. BOTTOM OF SCREEN METHOD DRILLED: HOLLOW STEM 24.0' B.Q. BOTTOM OF SUMP BOTTOM OF HOLE METHOD DEVELOPED: BAILING HOLE DIAMETER COMMENTS: FLUSH MOUNTED PER IDOT R.O.W. 0.25 Metoelf & Eddy

and setting the second section to the second





The second second					*	
L. West Street	6,48 (6,57,16,64 (6,	72 1150.00	1060.00 1000.0	1150.00		2.00 8.00 7-
MW-1A	7.11 6.90 6.65 6.	81 900.00	850.00 900.0	00.000	1.50 2.00	2.00 1.50
MW-2	7.42 7.84 7.57 7.	41 7.31 1534.00	1569.00 1563.0	1606.00 1617.00	7.10 4.44	9.00 9.00
MW-3	8.45 5.37 8.40 -	1300,00	1350.00 1300.0	b	3.50 4.00	4.00
MW-4	6.75 6.97 8.98 7.	.02 1260.00	1220.00 1240.0	0 1240.00		4.44 4.50
MW-5	6.52 6.58 6.67 -	1400.00	1160.00 1650.0	0		2.00
MW-5A	5.85 6.45 6.41 6.	.00 1200.00	0.008 00.0011	0 1300.00	2.90 0.00 -	0.50 3.90
MW-6	7.30 7.29 7.27 7.	.30 1549.00	1565.00 1574.0	0 1590.00	6.44 7.30	7.28 7.89
MW-7		• 500,00	1100.00 1100.0	0 1000.00	_,	7.00 5.50
MW-7A	7.19 7.02 6.74 -	700.00	700.00 700.0	0 650.00		5.00 3.00 :
MW-8	6.72 6.65 6.73 6.	.87 6.86 1100.00	1200.00 1250.0	0 1250.00 1250.00	1.00 1.00	1.00 1.00 1.00
MW-9	6.60 6.30 6.36 6.	.49 1000.00	1000.00 1000.0	0 1100.00	2,50 1,00	1.00 4.00

pH meter not working appropriately.
 Measurements were taken with the Hydac digital conductivity/temperature/pH Tester.

GOMNONWEALTH EDISON ORMER GRIESS - PFLEGER VANNERY SITE WAUKEGAN ILLINOIS GROUNDWATER ELEVATIONS

	20.0		7.7		
		TOR	GRD:	D7 *.	CORR.
		ELEV,	ELEV.	* *	AT ELEV
MELT, ID.	DATE	(FEET)	(FEET)		(FEET)
MW=1	5/24/94	99.44	97,66	51, 7	93.76
	6/4/94	99.44	97.66	5.84	83.62
	2/24/95	99.44	97.68	6.10	93.34
MW-1A	2/24/95	99.79	97.87	6.49	93.30
MW-2	5/24/94	101.60	98.92	7.67	93.93
	6/4/94	101.60	98.92	7.91	93.69
	2/24/95	101.60	98.92	8.08	93.52
MW-3	5/24/94	97.89	95.64	4.63	93.26
	6/4/94	97.89	95,64	4.78	93.11
	2/24/95	97.89	95.64	4.61	93.28
MW-4	5/24/94	99,69	97.61	5.60	94.09
	6/4/94	99.69	97.61	5.79	93,90
	2/24/95	99.69	97.61	5,28	94.41
MW-5	5/24/94	98.53	96.71	5,39	93.14
	6/4/94	98.53	96.71	5,50	93.03
	2/24/95	98.53	96.71	5.29	93.24
MW-5A	2/24/95	98,87	96.79	5.61	93.26
MW-6	5/24/94	98.44	96.10	4.87	93.57
	6/4/94	98.44	96.10	5.03	93.41
	2/24/95	98.44	96.10	4.84	93.60
MW-7	5/24/94	104.11	101.96	8.43	95,68
	6/4/94	104.11	101.96	8,53	95.58
	2/24/95	104.11	101.96	9,52	94.59
MW-7A	2/24/95	100.43	100,22	4,81	95.62
MW-8.	2/24/95	98,91	96.74	5.60	93,31
MW-9	2/24/95	98.08		4.22	93.86

Groundwater Elevations are taken from top of stainless steel casing.

