

SITE INVESTIGATION/REMEDIAL ALTERNATIVES REPORT WORKPLAN Former Provan Ford Facility 146-172 Mill Street Newburgh, New York

Prepared for:

City of Newburgh 83 Broadway Newburgh, New York 12550

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July 2000

Project No. PROVA001

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INTRODUCTION

First Environment Inc., (First Environment) was retained by the City of Newburgh to conduct site investigation and remediation activities at the Former Provan Ford facility located at 146-172 Mill Street Newburgh, New York. The activities outlined in this workplan are based on a site investigation conducted in October 1999 by First Environment as well as proposed investigation and remediation activities resulting from a site meeting held on February 8, 2000 where First Environment met with members of the New York Department State of Environmental Conservation (NYSDEC) and McGoey Hauser and Edsall. In attendance were Mssrs. David Camp and Larry Ricci of the NYSDEC, Mr. William Hauser of McGoey, Hauser and Edsall and Mssrs. Thomas Bambrick and Scott Green of First Environment.

The City of Newburgh is conducting the activities under the Municipal Assistance Environmental Restoration Projects "Brownfields Program". The following workplan details the site investigation and remedial investigation activities either conducted or proposed for the Former Provan Ford site. The activities include: the disposal of stockpiled soil; excavation and disposal of impacted soil related to USTs; the sampling, excavation and disposal of surface soil impacted by petroleum materials; the characterization and disposal to stored drums; the collection and analysis of soil samples; the installation, sampling and analysis of groundwater monitoring wells; the closure of underground storage tanks and piping; sealing interior drain connections; and the preparation of a Site Investigation/Remedial Alternatives Report (SI/RAR). All work will be performed in accordance with applicable state and local requirements.

ENVIRONMENTAL SETTING

GENERAL INFORMATION

The Former Provan Ford Facility is located at 146-172 Mill Street, Newburgh, New York. The adjoining properties east of the subject property and across Mill Street are residential. The adjacent to and properties across Route 9-W (Robinson Street) include several commercial companies, including Marjam Building Supply, Levinson Used Car Dealer and a car wash. The adjacent property south of the Former Provan facility is occupied by the Orange County Plumbing Supply Company. A Dairy Cone Ice Cream Store and an old warehouse/commercial building now occupied by a Limousine Service are located adjacent to and northwest of the subject property. The adjoining property north of the Former Provan Ford facility across Dickson Street is a vacant brick warehouse/light manufacturing building which appears to be several decades old.

The subject property, comprises approximately 3.5 acres, is located on the northwest corner of Mill Street and Deyo Place in Newburgh, New York. The land on which the facility lies is covered approximately 20% by the facility building. The remaining portion of the property is covered with gravel and/or concrete. Some portions of the northeast and west areas of the property are covered with earthen fill and vegetation. Two concrete pads, where trucks were formerly washed are also present immediately west of the facility building. The property is located in a mixed commercial and residential use area.

The Former Provan Ford building consists of one slab-on-grade concrete structure. This building encompasses approximately 18,000 square feet and is used primarily for garage and storage space. The building is segregated in two halves; the northern half, which was built in the 1940s and is now used exclusively for garbage storage; and the southern half, which was added to the original structure in the 1950s and is used for petroleum and parts storage as well on-going truck repair operations. According to the

current site occupant, the site building had been occupied by Former Provan Ford since its construction. Prior to the construction of the facility, the site was undeveloped. The boundaries of the site are delineated by Route 9-W (Robinson Street) on the west, The Orange County Plumbing Supply Company on the south, Dickson Street on the north and Mill Street on the east. The location of the facility is illustrated on the Newburgh Quadrangle United States Geological Survey (USGS) 7.5 Minute Topographic Map (revised 1977) as Figure 1.

SITE HISTORY

Gary's Heavy Truck repair was the occupant from March 1998 to March 2000. The City of Newburgh became the owner in March 1998 after it foreclosed on the former Provanowned property and has subsequently evicted Gary's Heavy Truck Repair (March 2000). At the time of its operation, Gary's Heavy Truck repair activities included the service and maintenance of heavy trailer-trucks in the southern garage area. The facility also includes a large amount of petroleum and refuse storage on-site, however, these were not included in the Truck repair activities.

Prior to 1998, Former Provan Ford had occupied the property for approximately fifty years. Former Provan's on-site operations included the washing, service and maintenance of Ford tanker trucks.

The Former Provan facility consisted of two attached garage maintenance areas, a small office area and an external wash rack which was located on a concrete pad and attached to the western wall of the garage. Empty tanker trucks would arrive at the site, where they would be cleaned in the wash rack, then serviced and fueled before they were sent back on the road for petroleum distribution.

TOPOGRAPHY AND DRAINAGE

The USGS 7.5 Minute Topographic Quadrangle Map for the Site is included in Figure 1 and shows the area in which the Former Provan Ford Facility is located. The Facility is approximately 500 feet above mean sea level and the surrounding topography slopes to the southeast toward the Quassaick Creek approximately 1/8 mile south of the site area.

WATER/WASTEWATER AND STORMWATER

Only sanitary wastewater is discharged from the subject property. Stormwater on the subject site flows overland with the topographical gradient (east/southeast) and discharges to either Mill Street or one of several storm drains located on the paved portions of the subject property.

GEOLOGY AND HYDROGEOLOGY

The Former Provan property lies within the Valley and Ridge Physiographic Province. The predominant features associated with this province are narrow valleys and ridges formed as a result of differential erosion of the underlying sandstone and shale formations. Specifically, the Newburgh area is characterized by alluvial deposits underlain by meta-sedimentary and sedimentary bedrock formations.

Alluvium in the area is comprised of flood plain sediments (sand, silt and clay) associated with the Hudson River. Glacial deposition in the area consists primarily of till and unsorted outwash. The outwash is predominantly a mixture of gravel, sand, silt and clay. The bedrock underlying this region consists of middle Ordovician Taconic Sequence, primarily shales and graywackes.

Soil borings conducted at the property indicate the upper 2 to 3 feet is primarily composed of fill material. A sand and gravel layer extends from approximately 2 to 8 feet below grade and is underlain by a lower permeability layer of silt. The above-

mentioned unconsolidated soil overlies weathered bedrock at approximately 15 feet below grade. The water table varies between 8 to 14 feet below grade. Groundwater is believed to flow to the east/southeast in the direction of the topographic gradient.

PREVIOUS WORK SUMMARY

An initial investigation was initiated by First Environment on October 19 to 22, 1999. During the investigation, soil borings and soil samples were collected for analysis; in addition, the installation sampling and analysis of groundwater monitoring wells was conducted to provide a baseline for soil and groundwater quality at the site.

In total, First Environment installed 27 geoprobe soil borings, of which, soil samples were collected at 15 locations for laboratory analysis. Using a geoprobe, soil borings were installed at or immediately adjacent to each of the six known USTs, the exterior Oil Water-Separator and associated pipeline, the Truck Wash Rack and associated drains as shown in Figure 2. These areas present the potential for the greatest environmental concern due to the former contents, associated handling practices, and age of the above-referenced tanks and pipelines. Geoprobe soil analytical results adjacent to USTs and the oil/water separator are summarized in Table 1. These results indicate 12 of 15 soil samples exceed the NYSDEC TAGM recommended soil clean up objectives for various volatile organic compounds and/or semi-volatile organic compounds.

Five surface soil samples were collected from 0 to 6-inches below grade adjacent to the Wash Rack Area, Oil/Water Separator, the UST tank islands, as well as the former Truck Parking Area as shown in Figure 3. Soil collection was biased toward areas that contain visible petroleum stained soil. Each soil sample exceeds the NYSDEC TAGM recommended soil clean up objective for various volatile organic compounds and/or semi-volatile organic compounds as shown in Table 2.

A total of seven temporary one-inch piezometers (PZ-1 to PZ-7) were installed using a Geoprobe to assess the water quality at each of the former UST locations and the

oil/water separator as illustrated in Figure 4. The water table was encountered from approximately 8 to 14 feet below grade and petroleum product was detected on the water table at five of the seven wells. Groundwater samples were collected at each piezometer for laboratory analysis in attempt to establish a baseline for VOC concentrations at each UST location and at the oil water separator. Six of seven piezometer locations indicate the groundwater quality has been impacted above the NYSDEC groundwater standard for volatile organic compounds. The groundwater analytical results are summarized in Table 3. Attachment A includes the well construction logs for piezometers (PZ-1 to PZ-7). All drilling, well installation, soil and groundwater sampling procedures are included in the Quality Assurance/Quality Control Plan (QA/QC) provided in Attachment B.

In addition, at the request of the NYSDEC approximately 60.8 tons of surface soil was stockpiled for disposal. This soil was sampled by First Environment on December 15, 1999 for waste characterization. Stockpiled soil was identified as non-hazardous and disposed of by TPS technologies, Inc. on February 8, 2000.

Also, during the February 8, 2000 site walk between First Environment and the NYSDEC, a limited number of drums were identified that pose a threat to the environment. Therefore, on February 24, 2000, a total of 9 drums were overpacked by CODE Environmental prior to pending off-site disposal.

PROPOSED SCOPE OF WORK FOR AREAS OF ENVIRONMENTAL CONCERN

UST REMOVAL, SOIL SAMPLING AND GROUNDWATER ACTIVITIES

Six USTs have been identified for removal at the Former Provan Ford Facility, including: 1) 20,000-gallon diesel tank, 2) 10,000-gallon diesel tank, 3) 8,000-gallon heating oil tank, 4) 4,000-gallon gasoline tank, 5) 3,000-gallon waste oil tank, and 6) 550-gallon heating oil tank. The NYSDEC case manager will be notified at least one (1) week in advance to the removal of the USTs. In addition, the NYSDEC will inspect all tank excavations prior to backfilling.

During the previous investigation, First Environment discovered each UST has impacted soil immediately adjacent to the tank. Furthermore, floating product was detected in wells adjacent to all but one UST. In order to better evaluate potential soil excavation limits and water quality, First Environment will install 9 additional monitoring wells prior to removing the USTs. Eight of the nine wells will be installed downgradient of the existing USTs and one well will be installed upgradient of the existing USTs and operations building. The upgradient well will establish background water quality levels.

The UST/piping excavation, removal and disposal activities will include pumping residual petroleum material, excavation of significantly impacted soil, and removal of existing concrete pads that overlie the USTs. Prior to removing the USTs and associated piping, each tank and piping run will be inerted, pumped, and cleaned of any residual petroleum materials as described in First Environment UST activities and procedures included as Attachment C. First Environment estimates approximately 5,000 gallons of residual petroleum and water will be pumped from the USTs prior to removal. After the USTs have been removed, any significant accumulation of product in the excavation will be removed from the excavation using a VAC truck prior to backfilling. Once the USTs have been cleaned, they will be removed and examined for

holes and then taken to a scrap-metal facility for disposal. During the removal, First Environment estimates between 15 to 30 cubic yards of contaminated soil per tank will be excavated and stockpiled for waste classification and disposal. As part of the excavation process, excavated soil will be field screened and visually inspected to determine potential re-use. Once field screened, the soil will be segregated into piles consisting of a petroleum contaminated and non-contaminated soil. At which time, waste characterization samples will be collected for the impacted soil pile for purpose of classifying the soil for disposal. In addition, a total of 70 cubic yards of concrete pad will be removed and disposed.

Subsequent to excavating any highly contaminated soil, post excavation soil samples will be collected at each of the UST excavations in accordance with NYSDEC STARS Memo #1 and SPOTs Memo #14. A total of six post-excavation soil samples will be collected at each UST excavation base, consisting of at least one composite sample from each of the four sidewalls and two from the excavation base. At tank area 5 & 6, First Environment will combine the post excavation sampling due to their close proximity to one another. If visually impacted soil remains subsequent to excavating soil, a grab sample will be collected pursuant to SPOTs Memo #14. Also, one sample per 20 linear feet of piping run will be collected and analyzed. Upon completion of soil excavation and post-excavation sampling, clean distinguishable soil will be backfilled in each excavation to grade. Figure 5 illustrates the approximate area of each UST excavation and the proposed location of post excavation soil samples. The Health and Safety Plan for these activities is provided in Attachment D.

In accordance with the STARS document, soil impacted by gasoline compounds, will be analyzed for VOCs and MTBE using Method 8021. For soil impacted by fuel oil, samples will be analyzed for both VOCs and base/neutral compounds using Methods 8021 and 8270, respectively. In areas impacted by waste oil, soil will be analyzed for VOCs plus TICs (Method 8260) and SVOCs plus TICs (Method 8270) as well as PCBs using Method 8082 plus sulfuric acid clean up and TAL metals.

A total of nine monitoring wells are proposed by First Environment. Eight 2-inch PVC flush mounted monitoring wells will be used to delineate the extent of groundwater contamination associated with the USTs as well as the installation of one upgradient well to provide background water quality data. The proposed well locations are illustrated in Figure 6. As per Section 5.5.3.2 – "Permanent Monitoring Wells," of the NYSDEC Sampling Guidelines and Protocols, the 2-inch wells will installed using hollow stem augers or continuous flight augers. Casings with well screens shall be installed in unconsolidated and semi-consolidated soils to prevent soil and other foreign material from entering the well during pumping. The annular space surrounding the screen will be backfilled with sand and filter pack and the remainder of the annular space surrounding the casing above the screen will be backfilled with bentonite, and/or cement bentonite grout, depending on the conditions of the site. A typical well construction diagram for unconsolidated material has been provided as Figure 7.

Once the wells have been installed, they will be developed to a sediment free discharge. Following a two-week period to allow the wells to stabilize, an initial round of groundwater samples will be collected at all 16 wells that exist at the site.

All 16 wells onsite will be surveyed and then sampled for chemical constituents consistent with NYSDEC STARS guidance document. These include for gasoline compounds, groundwater analysis for VOCs and MTBE using Method 8021. For groundwater impacted by fuel oil, STARS recommends analysis for both VOCs and base/neutral compounds using Methods 8021 and 8270, respectively. At the former waste oil tank, groundwater will be analyzed for VOCs using Method 8260 plus TICs and SVOCs using Method 8270 plus TICs as well as PCBs (Method 8082) and TAL metals.

First Environment will conduct rising and falling head slug tests at each of the 16 wells in an effort to determine the permeability of the water-bearing zone. This data will be used to characterize the Hydrogeology at the site and will be included in the SI/RAR.

SURFACE SOIL SAMPLING

Soil staining has been observed at five areas on the property. First Environment has identified that surfical soil concentrations, exceed NYSDEC recommended soil clean up objectives for PAHs, metals and/or VOCs at each of the five areas. As a result, First Environment recommends additional investigation at each of the five areas.

At each area, four additional sample locations will be sampled at 0-6 inches and 6 to 12-inches below grade to horizontally and vertically delineate the extent of contamination. In addition, two background samples will be collected and analyzed from 0 to 6 inches at "Clean Areas" away from Former Provan Operations. Surficial soil sample locations are illustrated in Figure 8. Soil samples will be collected and analyzed for base neutrals (Method 8270), PCBs, and metals, since the petroleum stained soil is likely the result of waste oil leaks associated with heavy truck transport. Samples collected at 6 to 12-inches below grade will also be analyzed for VOCs (Method 8021). Based on the results, the need to excavate soil for remedial purposes will be evaluated. The QA/QC Plan summarizing the soil sampling procedures is provided in Attachment B. The Health and Safety Plan for these activities is provided in Attachment D.

DRUM CHARACTERIZATION & DISPOSAL

The Former Provan facility does not currently generate any hazardous waste or petroleum oil waste which requires disposal. However, 81 drums (56 outside; 25 inside) of waste oil, hydraulic oil and other miscellaneous petroleum products are currently staged on at the Former Provan property. The 56 drums, which are located outside, are on the western portion of the property beneath the truck cleansing rack. The drums were staged at this location as part of NYSDEC's initial stabilization of the property after drums of spilled oil were discovered in June, 1998. According to the site representative, the interior 25 drums have been undisturbed since Former Provan vacated the property.

During the February 8, 2000 site walk between First Environment and the NYSDEC, a number of dented and rusted drums were observed as well as one drum that was leaking petroleum material. Drums that were identified as a potential threat to the environment were overpacked. On February 24, 2000, a total of 9 drums were overpacked by CODE Environmental.

Based on initial screening, the drums at Provan site contain petroleum products and waste oils. Drums will be characterized by a field chemist based on using physical methods with confirmation samples taken whenever necessary. The physical method includes: identifying drum markings, visual inspection of drum contents utilizing a glass drum sleeve, and obtaining MSDS's when possible. An OVA will also be utilized to monitor organic vapors. The field chemist will perform field screening for pH using pH paper and will perform field flash point tests in the field. Lastly, the chemist will perform field solubility and reactivity screening to determine solubility and/or reactivity of the petroleum contents.

Confirmation composite sampling for PCB's, Total Organic Halides, and Flashpoint will be conducted on drums that contain waste oil and petroleum contaminated solids. The analytical methods used for drum characterization will be as follows: total organic halides using EPA SW-846 method 3540A/9020A; Ignitability using SW-846 methods 1020; PCBs using SW-846 methods 3540A/9020A. A composite sample will be taken from no more than 10 drums for the analyses. Any drums failing the field screening for field ignitability will be analyzed to confirm the flash point by certified laboratory. We have selected Integrated Analytical Laboratories, a New York State Department of Health & Environmental Laboratory Approval Program (NYSDOH ELAP) certified laboratory to perform lab analyses.

Prior to drum disposal, each drum will be characterized by a field chemist. Based on preliminary testing, we expect 10% of the drum to be hazardous and the other 90%, non-hazardous.

PIPE AND DRAIN ASSESSMENT

Based on historical maps (provided by the City of Newburgh or site owners and operators) and site observations, First Environment will develop a map that illustrates the on-site pipeline and drains.

OILWATER SEPARATOR PUMP-OUT, CLEANING AND DISPOSAL

First Environment will pump-out, clean, and dispose of the contents of both the interior and exterior oil/water separators. First Environment estimates a total of 2,500 gallons of petroleum material and sludge will be pumped out of the oil/water separators and disposed. The interior separator will be cleaned and disposed at a scrap-metal yard along with its associated piping. All inside building drains that lead to the exterior separator will be sealed with concrete. Prior to sealing, the drains each will be cleaned. Once the exterior separator has been cleaned, it will be visually inspected to determine its integrity. Also, a sample of the groundwater from the well immediately downgradient of the separator will be collected and analyzed to determine if there has been any adverse impacts resulting from past use of the separator. Since the separator is relatively new, if the integrity is sound and there are no identifiable impacts from past use, then the separator will be left in place, as is, for potential future use. The exterior drains will also be cleaned of petroleum materials along with exterior oil/water separator.

SITE INVESTIGATION/REMEDIAL ALTERNATIVE REPORT

First Environment will provide an SI/RAR that documents the findings of the site investigation and proposed remedial alternative activities, including results of the UST excavation, removal and subsequent post-excavation soil sampling activities, soil disposal activities, groundwater monitoring results and recommendations consistent with 6NYCRR Part 375-1.10(c)(1-7); in addition to, remediation recommendations in accordance with guidelines specified in STARS for the management of excavated

contaminated soil as well as active groundwater remediation alternatives, including but not limited to, Risk Based Corrective Action Measures.

PROJECT SCHEDULE

A project schedule illustrating the major tasks is provided on Figure 9. The completion of these tasks are subject to receiving a fully executed contract.

WELL CONSTRUCTION LOGS

First Environment's well construction logs are included as Attachment A.

QUALITY ASSURANCE/QUALITY CONTROL PLAN

First Environment's Quality Assurance/Quality Control Plan is included as Attachment B.

UST ACTIVITIES AND PROCEDURES

First Environment UST Removal Procedures are included as Attachment C.

HEALTH AND SAFETY PLAN

First Environment's site-specific health and safety plan for the activities to be performed at the Provan Ford site is included as Attachment D.

CITIZEN PARTICIPATION PLAN

First Environment's Citizen Participation Plan is included as Attachment E.

COST SUMMARY & QUOTATIONS

First Environment's Cost Summary is included as Attachment F.

SOIL BORING ANALYTICAL RESULTS PROVAN FORD, NEWBURGH NEW YORK TABLE 1

Polychlorinated Biphenyls (PCBs)	TOTAL BN's & TIC's	TOTAL TICS	benzo[g,h,i]perylene	Dibenz[a,h]anthracene	Indeno[1,2,3-cd]pyrene	Benza[a]pyrene	Benzo[k]fluoranthene	Benzo[b]fluoranthene	Distance of the state of the st	Corysene	(Benzo[a]anthracene	Butylbenzylphthalate	Pyrene	Fluoranthene	Di-n-butylphthalate	Anthracene	Phenanthrene	N-Nitrosodiphenylamine	Fluorene	Diethylphthalate	Dibenzofuran	Acenaphthene	Acenaphthylene	2-Methylnaphthalene	Naphthalene	Semivolatile Organic Compounds	Naphthalene	n-Butylbenzene	4-isopropyitoluene	sec-Butylbenzene	1,2,4-Trimethylbenzene	tert-Butylbenzene	1,3,5-Trimethylbenzene	n-Propylbenzene	Isoprovibenzene	Total Xylenes	Ethylhenzene	Tolliene	Benzene	Volatile Organic Compounds - 8021 List	SAMPLE DATE	SAMPLE LOCATION	
10,000	n/a	n/a	50,000	14	3,200	61	1,100	1,100	50,000	5000	224	50,000	50,000	50,000	B, 100	50,000	50,000	n/a	50,000	7,100	6,200	50,000	41,000	36,400	13,000		13,000	n/a	n/a	. n/a	n/a	n/a	n/a	n/a	⊓/a	1,200	5.500	500	3		OBJECTIVE	RECOMMENDED	
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1	215,259	192,400	116	c	r 06	18	r 08	177	= 6	= 0	400	, C	432	420	_	211	1,220	c	254	=	184 J	137	108 J	9,450	9,420		992	c	c	c	1,550		769	256 J	128 J	2130	561	= 6	=		10/19/99	GS-4	
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ı	308426 J	190,940	117486 J	: c	: c	c	c	c	C	94 48	c (= 0	: c	: c	2,4/0	567 J	2,330	2,050	0 00	3 -		000	n 0 c	1,000	2,930		3,820	3 -	: c	: c	26,500	5	7,900	2,990	P 056	17.400	3,080	1030 J	c		10/22/99	G-26	חל חל
ı	622160 J	375,700	246460 J	: c	. c	c	c	c	1,230	169,000	c (- 100	13 20	755	725 1	1,200	6,800	6,530	1,0/0			1,200	3 -	01,000	12,700		Ω.Φ.Ψ	-	4,230	3 -	106,000*	, c	31,400	15,200	6,260		16,200	9 120	c		10/22/99	G-27	מנים

NOTES:

BN = Semivolatile Compound
TIC = Tentatively Identified Compound
= Not analyzed
J = The concentration was detected at a value below the MDL
u = Analyzed for but not detected at the MDL
SHADED = Analyte detected in excess of recommended soil cleanup objective
* = Results from diluted analysis
All compounds measured in parts per billion (ppb)

TABLE 2 SURFACE SOIL ANALYTICAL RESULTS PROVAN FORD, NEWBURGH NEW YORK

SAMPLE ID	RECOMMENDED	SS-1	SS-2	SS-3	SS-4	SS-5
SAMPLE LOCATION	SOIL	SS-1	SS-2	SS-3	SS-4	\$ \$-5
SAMPLE DEPTH (feet below grade)	CLEANUP	0-6"	0-6"	0-6"	0-6"	0-6"
SAMPLE DATE	OBJECTIVE	10/22/99	10/22/99	10/22/99	10/22/99	10/22/99
					İ	
Volatile Organic Compounds - 8021 List						
Total Xylenes	1,200	1,950	-	••	_	-
1,3,5-Trimethylbenzene	n/a	1,610	_	-	_	-
1,2,4-Trimethylbenzene	n/a	3,510	_		_	-
Naphthalene -	13,000	1,190	. –	_	-	-
Semivolatile Organic Compounds						
Naphthalene	13,000	372	409 J .	u	u	u
2-Methylnapthalene	36,400	2,160	501 J	u	น	u
Acenaphthylene	41,000	147 J	2,590	422	1,920	4,660
Acenaphthene	50,000	291	2,190	u	2,770	u
Dibenzofuran	6,200	372 J	1,410	u	ប	u
Fluorene	50,000	523	3,220	u	2,400	u
N-Nitrosodiphenylamine	n/a	1,640	6,030	u	u .	u
Phenanthrene	50,000	2,440	37,800	309	54,200	2,720
Anthracene	50,000	406	7,340	345	8,630	3,130
Carbazole	n/a	υ	2,900	u	5,650	u
Fluoranthene	50,000	137 J	92,000	1,690	167,000	24,600
Pyrene	50,000	224	92,300	1,730	110,000	42,800
Benzo[a]anthracene	224	U	30,400	1,370	47,200	16,600
Chrysene	400	u	33,100	1,260	50,900	18,000
bis[2-Ethylhexyl]phthalate	50,000	u	85,100	317	4,540	2,570
Benzo[b]fluoranthene	1,100	u	59,100	2,830	102,000	47,800
Benzo[k]fluoranthene	1,100	u	15,800	1,050	37,100	17,800
Benzo[a]pyrene	61	u	24,900	1,300	48,700	36,000
Indeno[1,2,3,-cd]pyrene	3,200	u	6,230	683	14,700	13,700
Dibenz(a,h,)perylene	14	u	2,040	299	4,720	4,410
Benzo[g,h,i]perylene	50,000	u	6,290	658	13,300	15,200
TOTAL BN's	n/a	8712 J	511650 J	14,263	675.730	249,930
TOTAL TIC's	n/a	94,700	3,565,700	6,728	280,650	139,620
TOTAL BN's & TIC's	n/a	103412 J	4077350 J	20,991	956,380	389,550
Metals						
Arsenic	7,500	3,630	4,860	20,800	4,300	6,960
Barium	300,000	42,300	58,100	43,200	88,200	124,000
Cadmium	1,000	u	956	438	5,420	2,720
Chromium	10,000	17,800	17,000	16,500	22,500	21,100
Lead	350,000	11,000	80,500	43,300	120,000	240,000
Mercury	100	23	36	66	76	61
Selenium	2,000	u	u	บ	u	u
Silver	N/A	u	u	u	u	u
Polychlorinated Biphenyls (PCBs)	10,000	_	-		u	

NOTES:

BN = Semivolatile Compound TIC = Tentatively Identified Compound

The later of the state of

* = Results from diluted analysis

All compounds measured in parts per billion (ppb)

TABLE 3

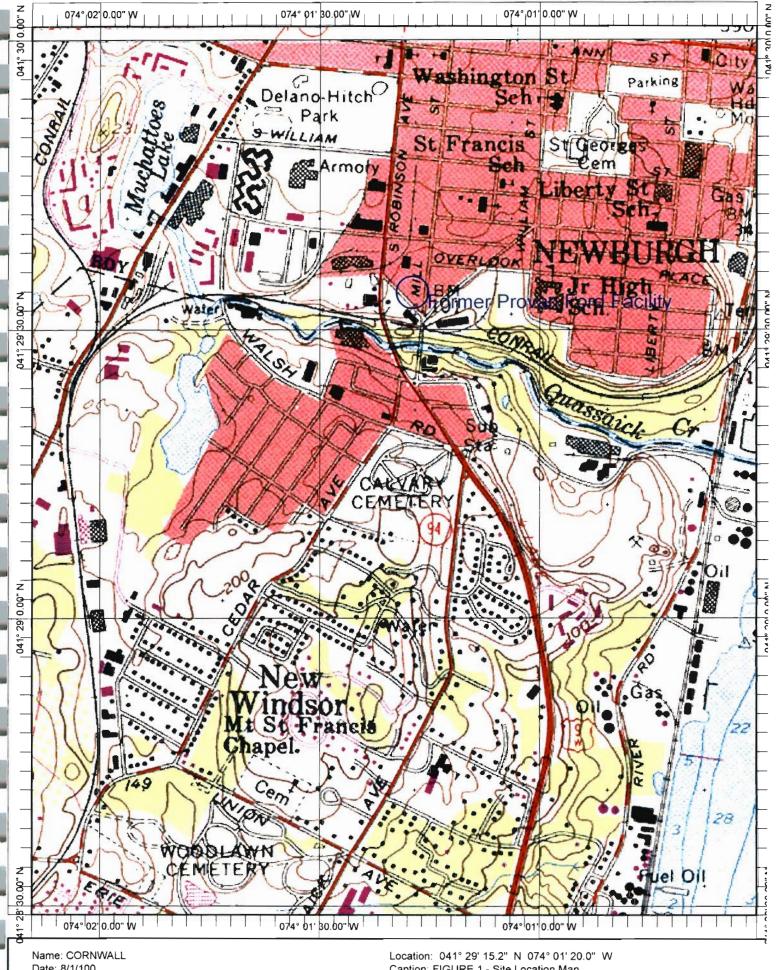
GROUNDWATER ANALYTICAL RESULTS PROVAN FORD, NEWBURGH NEW YORK

WELL ID/SAMPLE ID	RECOMMENDED GROUNDWATER	PZ-1	PZ-2	PZ-3	PZ-4	PZ-5	9-Z4	PZ-7
SAMPLE DATE	CLEANUP OBJECTIVE	10/21/99	10/21/99	10/21/99	10/21/99	10/21/99	10/22/99	10/22/99
Volatile Organic Compounds								
Vinyl Chloride 1,1-Dichloroethene	27 52	5,230	מ כ	. n	ם ם	ככ	88 u	1,240
Methylene Chioride Methyl-t-Butyl Ether (MTBE)	5	406	= =	□ ←	n 2 070	5 5	= =	n 2697
1,1-Dichloroethane	s	568		-		, ,	145	2,610
Chloroform	7	123	n	ח	ח	ח	ם .	n
1,1,1-1 (ICI)forbetinane 11,2-Dichloroethane(EDC)	വറ	7,000 U	3 2	л 38	.	3 5	. u	16.400
Benzene	0.7	160	10,200	- 01	13,100 *		73	374
Trichloroethene	'n	13,3001	ח	11	ח	ם	n	ח
Toluene	Ŋ	597	28,000*	2	45,000 *	ם	478	2,050
Tetrachloroethene	2	1,290	ח	4	ח	ם	ח	ם
Ethylbenzene	2	260	2,530	4	4,860	5	958	663
Total Xylenes	Ω.	317	14,200	4	19,600	ָ י	3,250	2,340
TOTAL VO's	n/a	25,322	54,930	89	89,630	ם	5,233	34,065
TOTAL TIC's	ח/מ	34,500	48,020	109	43,140	D	22,200	45,910
TOTAL VO'C & TIC's	n/a	59,822	102,950	197	132,770	כ	27,433	79,975
Metals								
Lead	n/a	10	611	ł	206	;	1	ı

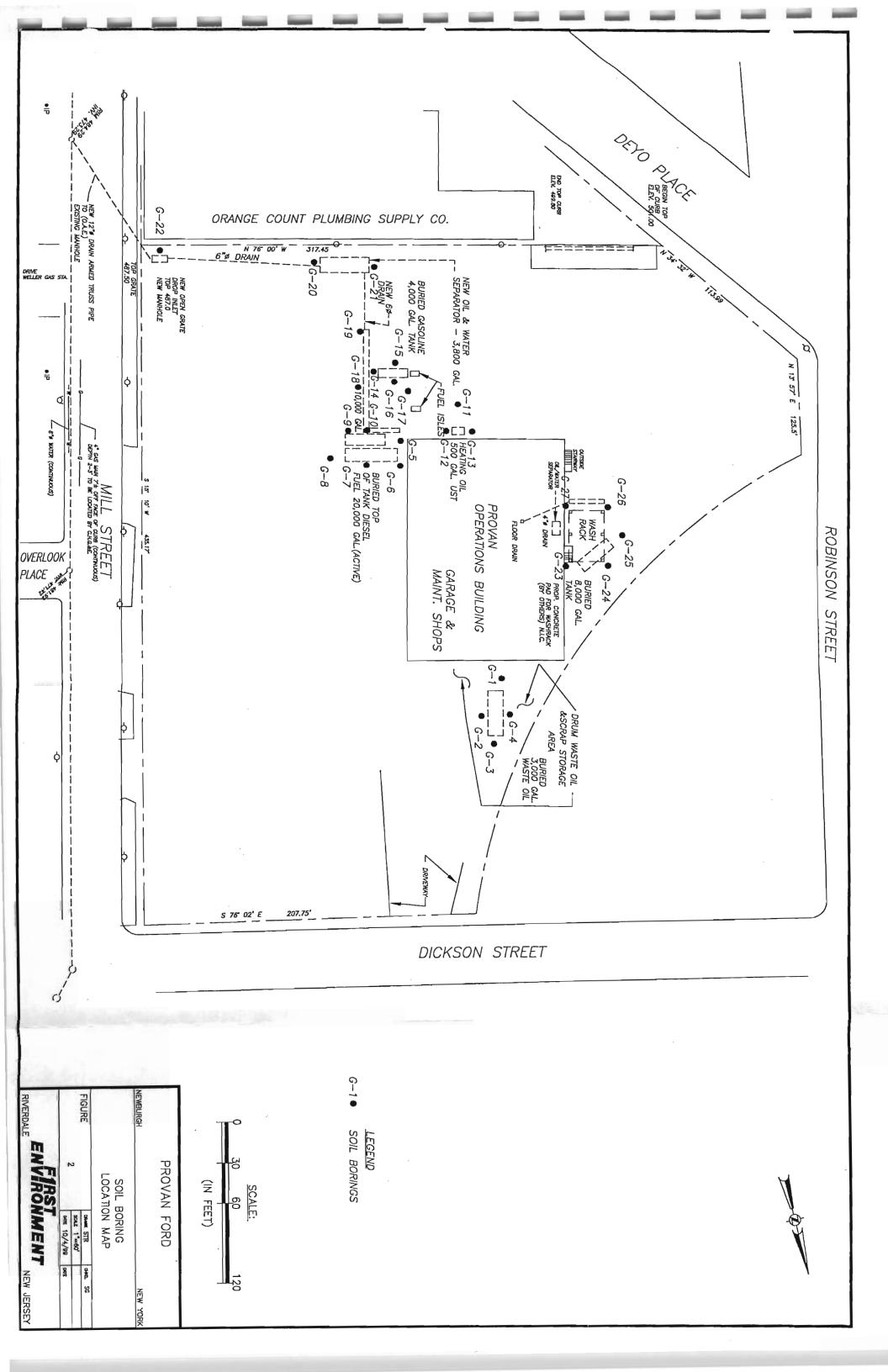
NOTES:

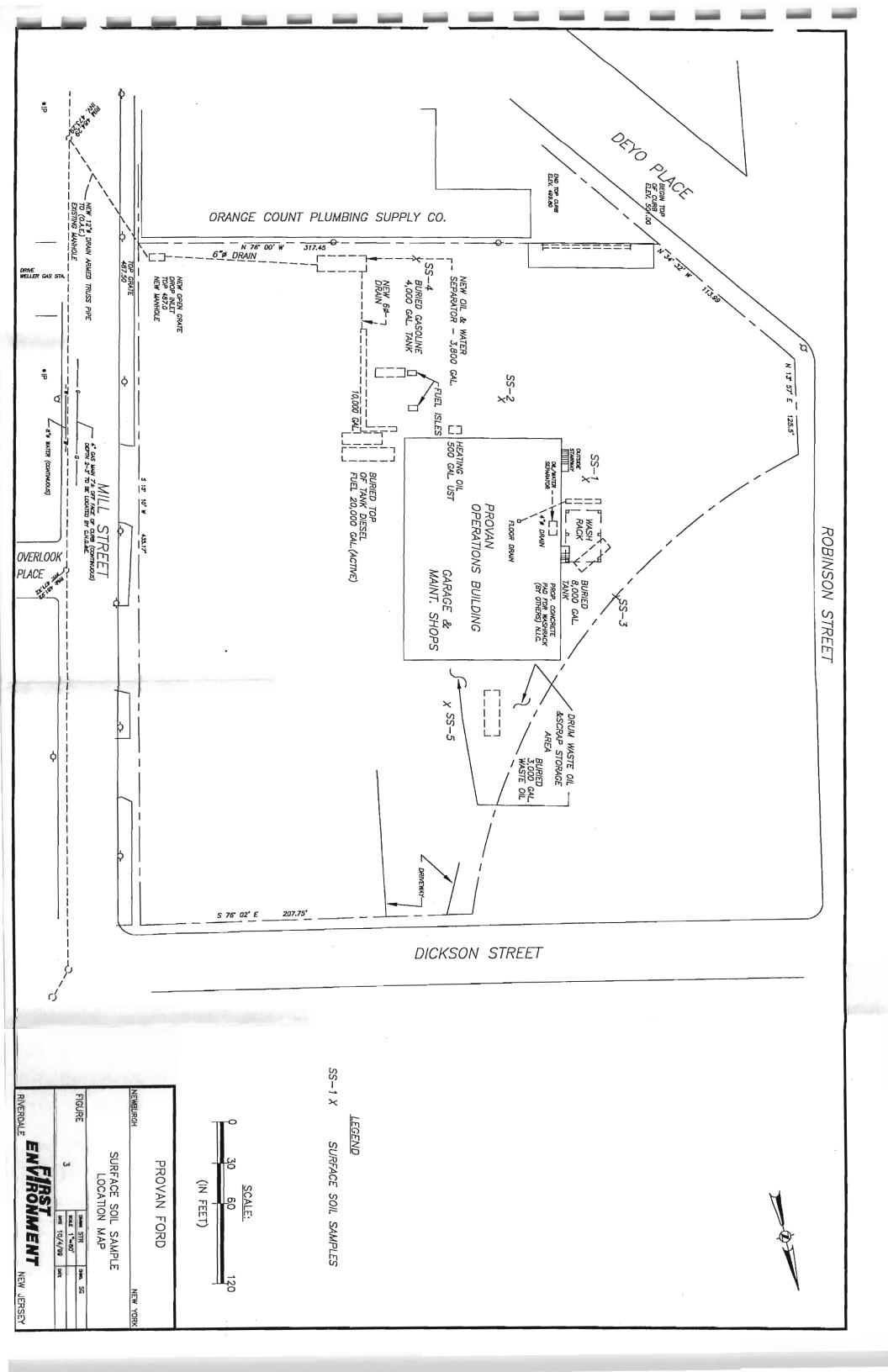
VO = Volatile Compound TIC = Tentatively Identified Compound

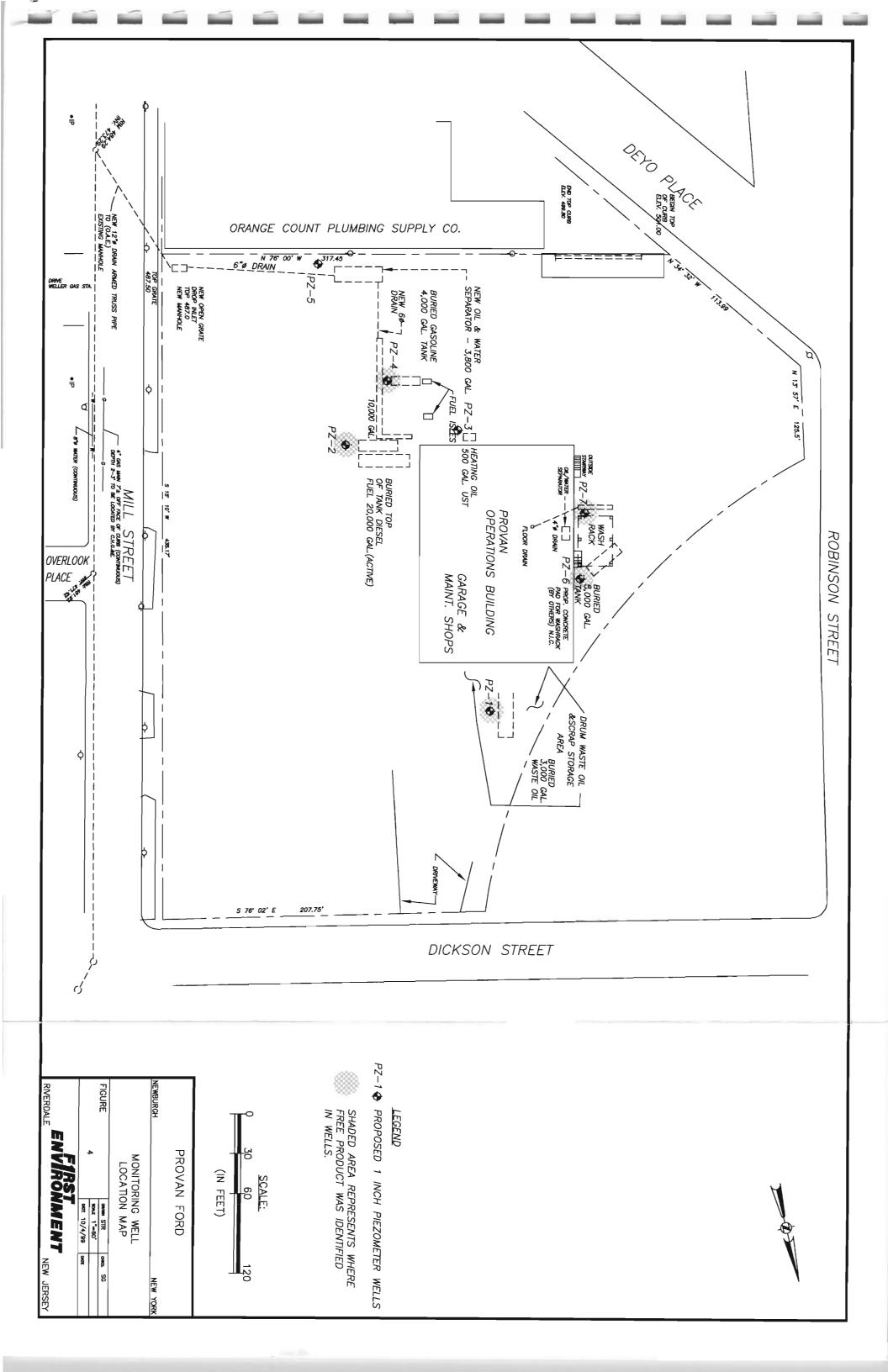
-- = Not analyzed
J = The concentration was detected at a value below the MDL
u = Analyzed for but not detected at the MDL
SHADED = Analyte detected in excess of recommended soil cleanup objective
* = Results from diluted analysis
All compounds measured in parts per billion (ppb)

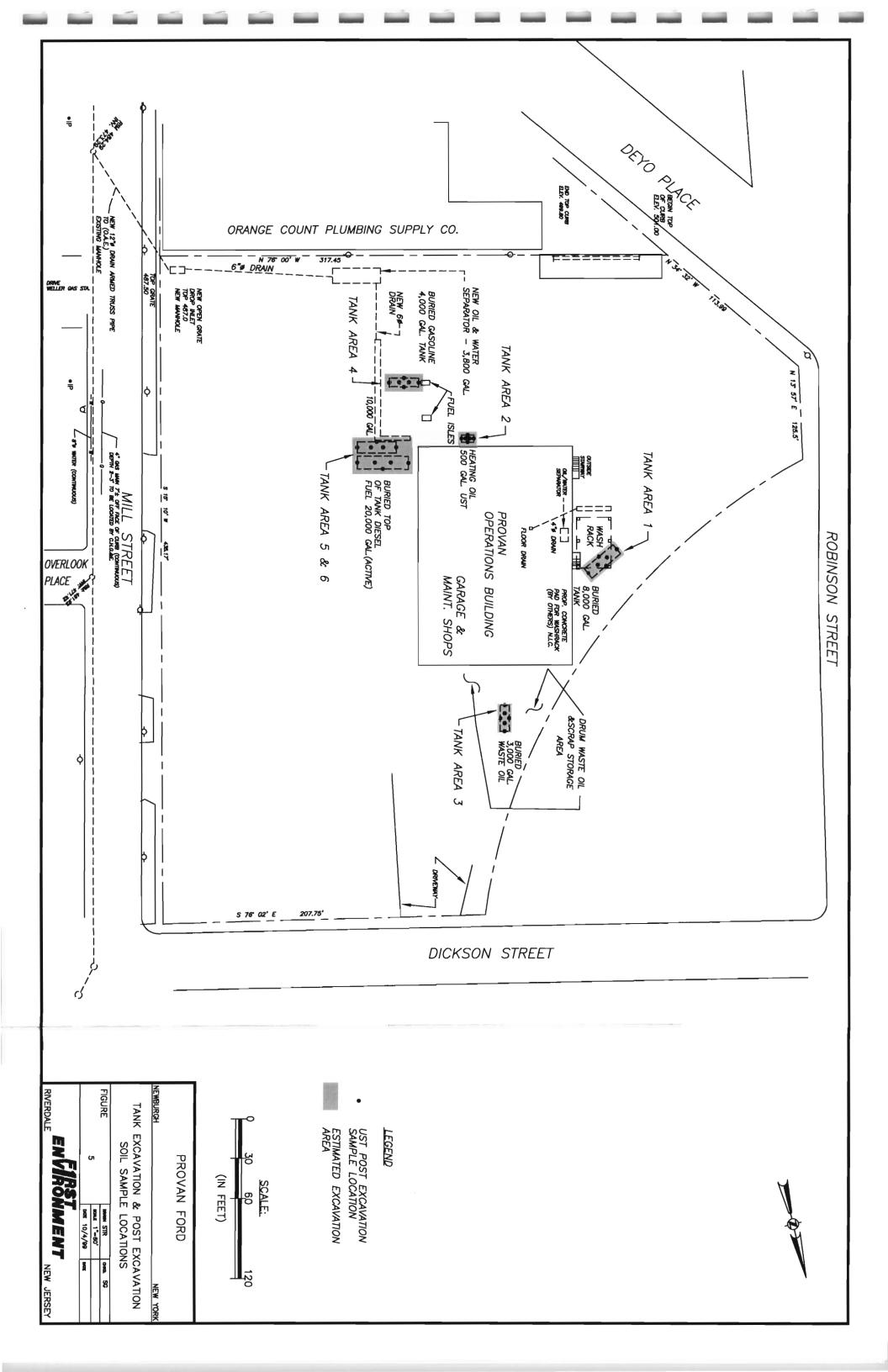


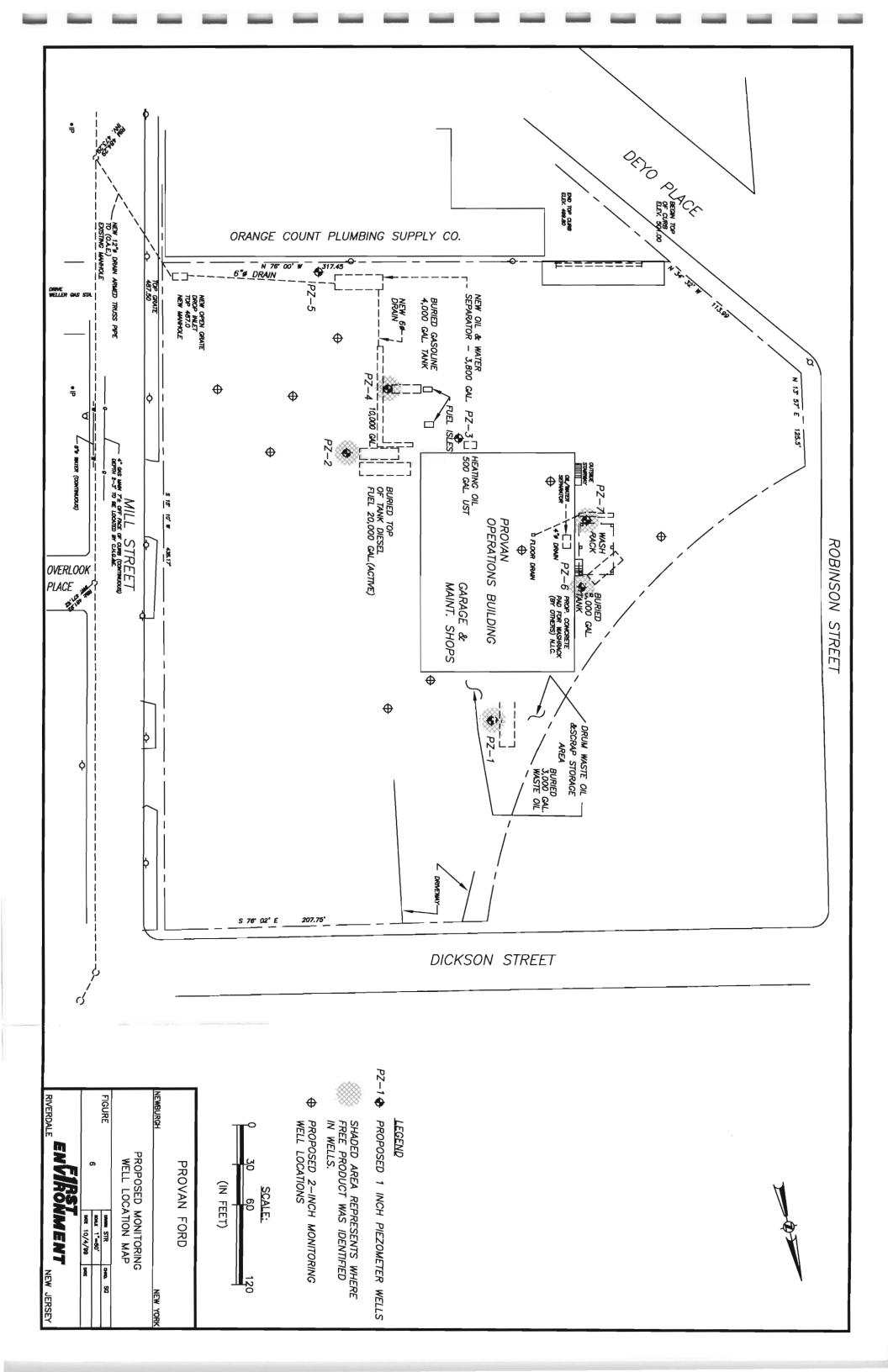
Date: 8/1/100 Scale: 1 inch equals 1000 feet Location: 041° 29' 15.2" N 074° 01' 20.0" W
Caption: FIGURE 1 - Site Location Map
Former Provan Ford Facility
Newburgh, NY

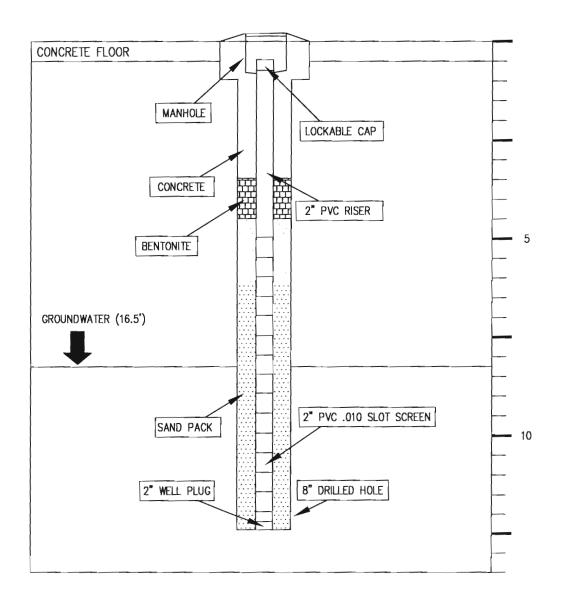




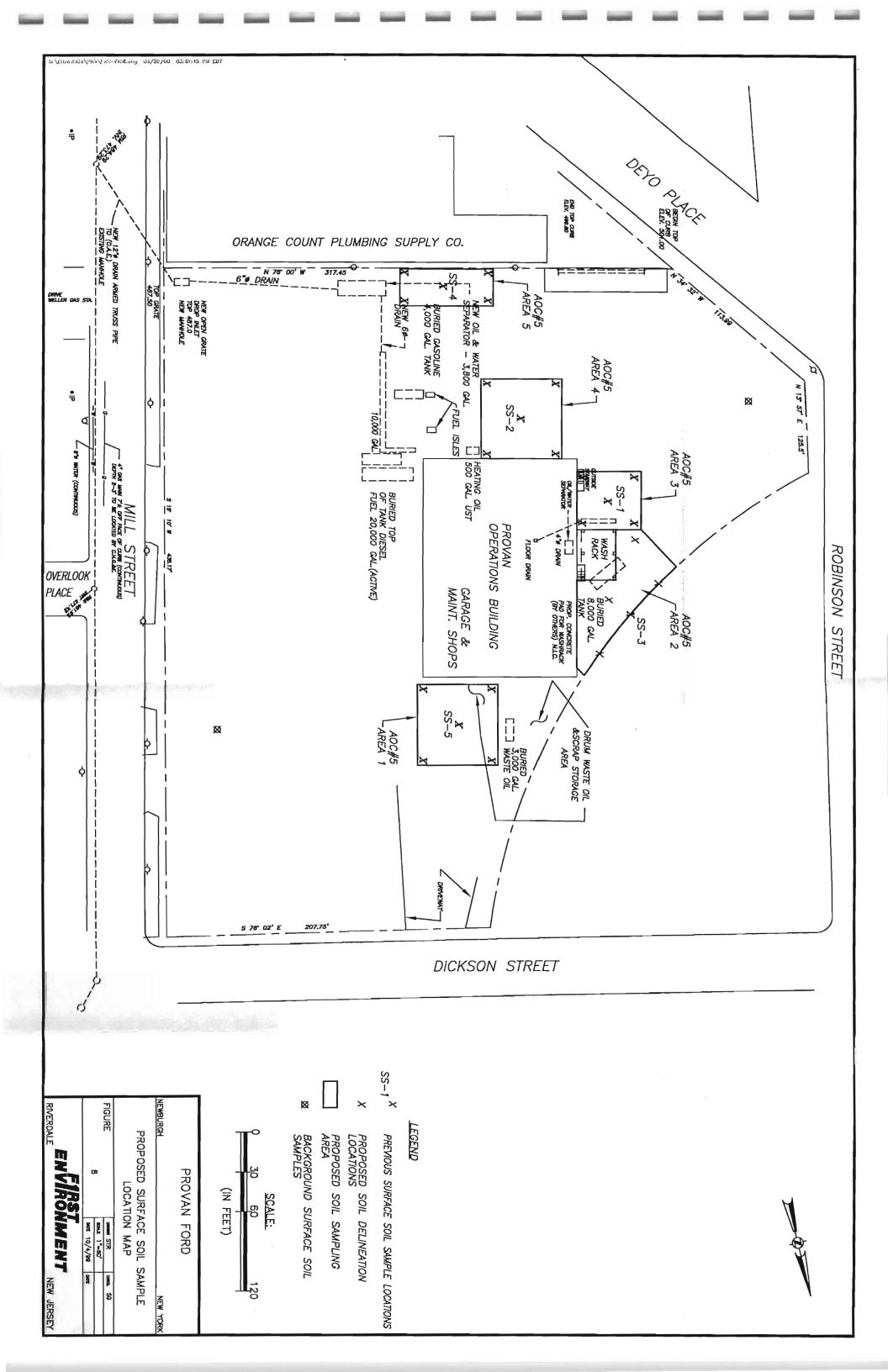


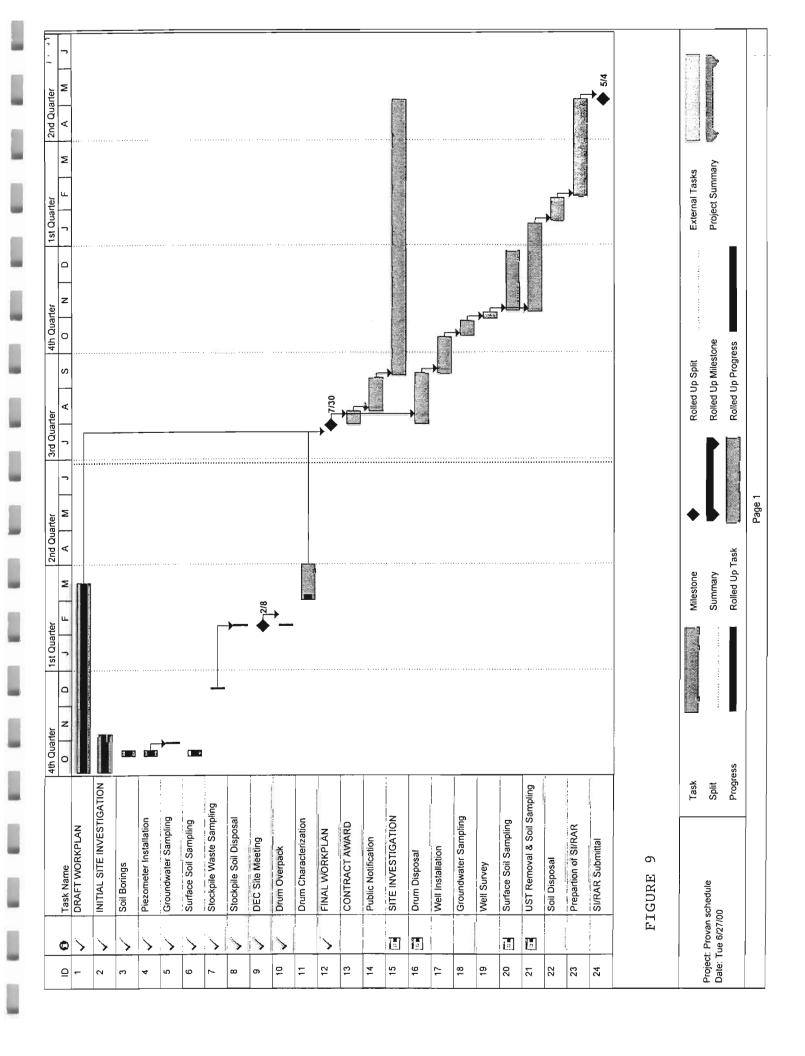






	PROVAN	FORD	
NEWBURGH			NEW YORK
	WELL CONSTRU	CTION DIA	GRAM
FIGURE		ORMAN CSN/STR	crean, TB
, ,,,,,,,,	7	SONLE NTS	
	,	DATE 8/12/99	DALE
RIVERDAL	F1RS1 ENVIRON	IMENT	NEW JERSEY





FIRST ENVIRONMENT DRILLING LOG

Boring No. ___PZ-1_____ Permit No. _n/a____ Project No. Prova001___ t PROVAN Sheet _1_ of _1_ Newburgh, NY te Started ___10/19/99_____ __evation (TC) ____ Hole Diameter __2"____ Date Completed ___10/19/99_ (GS) Boring Method __Direct-Push____ Development Method __P____ Water Level (Data) Strata Column Instrument Reading Sample No. Moisture Content Blows on Recovery Sample Sampler Well Construction Description 0" 6" 12" 18" 6" 12" 18" 24" hole Concrete fill, asphalt lock-28 50 0-4 able Brown Sand, little Silt cap concrete Little Gravel PVC benriser tonite 600 36 4-8 Х Green Sand and Gravel 1" PVC Y .010 slot screen sand pack 500 8-12 41 Silt and Clay, tr. F Sand 2" dia. hole well plug TD at 14.65 ft below grade Product Detected on Water Table **Boring Method Development Method** Sampler Type HSA - Hollow Stem Auger AL - Air Lift Driven Split Spoon Pressed Shelby Tube or Macro P - Pumping CFA - Continuous Flight Auger -DC - Driving Casing Rock Core J - Jetting SB - Surge Block MD - Mud Drilling NR. No Recovery

FIRST ENVIRONMENT DRILLING LOG

Boring No. __PZ-2____ Project No. Prova001_____ Permit No. _n/a_____ Sheet _1__ of _1__ nt PROVAN_ Newburgh, NY_ ate Started ___10/21/99 _____evation (TC) _____ Date Completed 10/21/99 (GS) Boring Method __Direct-Push ____ Development Method __P____ Hole Diameter __2"__ Water Level (Data) Strata Column Instrument Reading Sample No. Moisture Content Blows on Recovery Sampler Well Construction Description -0"-6"-12"-18" 6" 12" 18" 24" manhole Concrete fill, asphalt 42 40 0-4 able some Brown Sand, little clay cap concrete PVC benriser tonite 4-8 36 765 Green Sand, little Gravel 1" PVC .010 slot screen sand pack 28 1000 8-12 X Gr./green Silt and Clay 2" dia. hole Y well Weather Rock TD at 15.15 ft below grade Product Detected on Water Table **Boring Method** Sampler Type Development Method HSA - Hollow Stem Auger AL - Air Lift Driven Split Spoon CFA - Continuous Flight Auger -Pressed Shelby Tube or Macro P - Pumping DC - Driving Casing Rock Core J - Jetting MD - Mud Drilling

SB - Surge Block

No Recovery

Boring No. __PZ-3____ Permit No. _n/a____ Sheet _1__ of _1__ Project No. Prova001___ nt PROVAN Newburgh, NY Date Completed 10/21/99 te Started 10/21/99 (GS) evation (TC)___ Boring Method __Direct-Push____ Development Method __P____ Hole Diameter __2"____ Water Level (Data) Strata Column Instrument Reading Sample No. Blows on Moisture Content Recovery Sample Depth (ft.) Sampler Well Construction Description 0" 6" 12" 18" 6" 12" 18" 24" manhole Cinder of asphalt, Rock fill lock-20 24 0-4 able Some Sand & Silt cap concrete benriser tonite 75 48 Green Sand, little Gravel and some fill Х 1" PVC .010 slot screen sand pack 46 0 8-12 Green Silt and Clay layer 2" dia. Y hole well plug Weather Rock TD at 14.75 ft below grade Development Method Boring Method Sampler Type HSA - Hollow Stem Auger AL - Air Lift Driven Split Spoon CFA - Continuous Flight Auger > Pressed Shelby Tube or Macro P - Pumping DC - Driving Casing Rock Core J - Jetting MD - Mud Drilling SB - Surge Block No Recovery

Boring No. __PZ-4____ Project No. Prova001_____ Permit No. _n/a____ nt PROVAN_ Sheet _1__ of _1__ Newburgh, NY__ Date Completed __10/21/99____ ate Started ____10/21/99_____ levation (TC) (GS) Hole Diameter __2"___ Boring Method __Direct-Push____ Development Method __P___ Strata Column Water Level (Data) Instrument Reading Sample No. Moisture Content Blows on Recovery Sample Depth (ft.) Sampler Well Construction Description 0" 6" 12" 18" 6" 12" 18" 24" manhole Cinder of asphalt, Rock fill lock-40 575 0-4 able Some Sand, little silt,tr gravel cap concrete benriser tonite 1418 34 Green Sand, some gravel, little clay 4-8 .010 slot screen sand pack Y 36 1500 8-12 Green Silt, Rock frags, and small cobbles 2" dia. hole well plug Weather Rock TD at 14.6 ft below grade Product was detected on Water table Boring Method Development Method Sampler Type HSA - Hollow Stem Auger AL - Air Lift Driven Split Spoon CFA - Continuous Flight Auger-Pressed Shelby Tube or Macro P - Pumping DC - Driving Casing Rock Core J - Jetting SB - Surge Block MD - Mud Drilling No Recovery

Boring No. __PZ-5____ Project No. Prova001____ Permit No. _n/a____ nt PROVAN Sheet _1__ of _1__ Newburgh, NY___ te Started 10/20/99 revation (TC) Hole Diameter 2" Date Completed ___10/20/99___ Boring Method __Direct-Push____ Development Method __P____ Water Level (Data) Strata Column Instrument Reading Sample No. Blows on Moisture Content Recovery Sample Depth (ft. Sampler Well Construction Description 0" 6" 12" 18" 6" 12" 18" 24" manhole Cinders, stained soil 15 0-4 48 able F. Sand and Gravel сар concrete benriser tonite 4-8 24 20 Green Sand/silt,little clay little Gravel 1" PVC .010 slot screen sand pack 38 141 8-12 Х Green Silt layer, Rock frags Y 2" dia. hole well plug TD at 11.55 ft below grade **Boring Method** Sampler Type Development Method HSA - Hollow Stem Auger AL - Air Lift Driven Split Spoon CFA - Continuous Flight Auger -Pressed Shelby Tube or Macro P - Pumping Rock Core DC - Driving Casing J - Jetting MD - Mud Drilling SB - Surge Block No Recovery

Boring No. __PZ-6____ Permit No. _n/a____ Sheet _1__ of _1__ Project No. Prova001____ int PROVAN __Newburgh, NY______ate Started ___10/22/99_____ Date Date Completed __10/22/99 levation (TC) _____ (GS) Boring Method Direct-Push Development Method P Strata Column Water Level (Data) Instrument Reading Sample No. Blows on Moisture Content Recovery Depth (ft.) Sample Sampler Well Construction Description 0" 6" 12" 18" 6" 12" 18" 24" manhole Cinders, heavy stained soil 2500 0-4 40 F. Sand and Gravel able сар concrete PVC riser tonite 48 3700 F-M Sand and Gravel 4-8 sheen petr. Odor 1" PVC .010 slot screen sand pack 4200 8-12 36 Green/silt, stained soil х Y 2" dia. hole well plug TD at 14.36 ft below grade Product on Water Table **Boring Method Development Method** Sampler Type HSA - Hollow Stem Auger Driven Split Spoon AL - Air Lift CFA - Continuous Flight Auger-Pressed Shelby Tube or Macro P - Pumping DC - Driving Casing J - Jetting Rock Core

MD - Mud Drilling

SB - Surge Block

No Recovery

Boring No. __PZ-7____

Permit No. _n/a____ Sheet _1__ of _1__ Project No. Prova001____ nt PROVAN Newburgh, NY Date Completed __10/22/99 ite Started ____10/22/99___ revation (TC) (GS) Boring Method __Direct-Push____ Development Method __P____ Hole Diameter __2"____ Strata Column Water Level (Data) Instrument Reading Sample No. Blows on Moisture Content Recovery Sample Depth (ft.) Sampler Well Construction Description 0" 6" 12" 18" 6" 12" 18" 24" manhole Cinders, heavy stained soil lock-39 302 0-4 Gravel & Sand able сар concrete benriser tonite 1200 48 F-M Sand and Gravel 4-8 little clay 1" PVC Y .010 slot screen sa.ıd pack 1500 48 8-12 Х Green Silt and Clay, little gravel product encountered 2" dia. hole well plug TD at 12.2 ft below grade Product on Water Table Boring Method Development Method Sampler Type HSA - Hollow Stem Auger Driven Split Spoon AL - Air Lift \boxtimes CFA - Continuous Flight Auger-Pressed Shelby Tube or Macro P - Pumping DC - Driving Casing J - Jetting Rock Core SB - Surge Block MD - Mud Drilling No Recovery

QUALITY ASSURANCE/QUALITY CONTROL (QA/QC) PLAN

The soil, groundwater, surface water and sediment sampling techniques to be employed at the Provan facility in Newburgh, New York, are detailed below. All efforts will be made to eliminate sample contamination and maximize the reliability to the analytical results. These efforts include proper use and cleaning of sampling equipment and sample containers to eliminate sample contamination, use of a quality assurance program to maximize accuracy and precision of the analytical results, proper installation of groundwater monitoring wells and the use of chain-of-custody procedures to track the samples from source to analysis and minimize the opportunity for tampering.

SAMPLING EQUIPMENT AND PROCEDURES

SAMPLING EQUIPMENT AND CLEANING PROCEDURES

The sample containers, glass jars with Teflon™ lined-plastic screw-on lids, will be provided by a New York State Department of Health (NYSDOH) Environmental Laboratory Approval Program (ELAP) Certified Laboratory. Containers used to collect soil and water samples will be specifically designed for that purpose. The containers will be cleaned prior to shipment by the laboratory, using standard, in-house procedures.

Soil samples will be collected with either a hand auger, split-spoon sampler or Geoprobe Macrocore device. If a split-spoon sampler or Macrocore device is used, a drill rig will be used to drive the sampler to the required depth and obtain soil samples. The split-spoon sampler will be 24 inches in length and in accordance with ASTM D1586-67. The Macrocore will be 48 inches in length. The Macrocore sampler will be equipped with a sample retention device and acetate liner to ensure sample quality.

All soil sampling equipment will be cleaned with a wire or bristle brush to remove any clinging soils or materials. This will be followed by a washing with a phosphate-free detergent and water. The equipment will then be rinsed with clean water, distilled water, methanol (used to avoid contaminating soils with acetone), and finally, distilled water. The drilling tools will be cleaned with a steam cleaner prior to use and between work on individual boreholes.

Prior to sampling groundwater, wells will be purged by bailing or pumping, depending on well productivity. If bailing is used, one PVC bailer will be dedicated to each well for use in purging and one stainless steel, Polyethylene, or Teflon™ bailer will be dedicated to each well for use in sampling.

All purge/sampling equipment will be cleaned before transporting to the field. Bailers will be cleaned by the certified laboratory using standard in-house procedures. After drying, the bailers will be wrapped in aluminum foil for transportation to the field. Pumps used to purge wells prior to sampling will be cleaned by rinsing with detergent, potable water, distilled water, methanol and distilled water.

USE OF QUALITY ASSURANCE SAMPLES (BLANKS, DUPLICATES)

Travel blank samples, duplicates and blank samples generated in the field will serve as an independent check on the laboratory and field sampling techniques. These samples will be coded to minimize the chance of laboratory identification.

The following QA/QC samples will be collected:

- Duplicate samples will be collected for each media sampled. The number of duplicates will, at a minimum, equal five percent of the samples collected for that media.
- One travel/trip blank, consisting of distilled water as prepared by the laboratory will be and analyzed for VOCs for each two days of sampling.

 One field blank for each media sampled that day will be submitted for analysis for volatile organics.

SAMPLING METHODOLOGY

All subsurface work will be conducted in a manner that produces reliable information of subsurface conditions and representative soil, sediment, surface water and groundwater samples for analysis. A First Environment degreed hydrogeologist, geologist, engineer or equivalent will supervise all drilling and sampling procedures.

Soil Sample Collection Methodology

Soil Samples.

The location of each borehole will be referenced by a grid system or some other survey control. A drill rig/Geoprobe or hand auger will be used to produce boreholes at the proper depths at each predetermined sampling location. Sample depths will be measured to ensure they are correct. Samples will be taken by driving a split-spoon sampler, Macrocore or hand auger into the undisturbed material below the bottom of the borehole. The sampler will be equipped with a sample retention device.

Prior to sample collection, sample depths will be screened with Photoionzation Detector (PID) to insure that collected samples are representative of actual soil conditions. This will be accomplished through head space analysis. The PID will be calibrated to 100 ppm isobutylene prior to each days activities.

A soil sample will be collected from the appropriate sampler or hand auger. To prevent contamination of sample bottle by windblown soils, each bottle will remain sealed until sample collection. Upon soil collection, the sample will be split in two bottles for headspace analysis and laboratory analysis. The sample containing the highest

headspace PID reading at each boring location will be submitted to the laboratory for analysis. This sample will bottle have the following information recorded on it:

- Job Name and Location
- Sample Location
- Time and Date of Sampling
- Depth of Sampling
- Analysis
- Boring Number

The jar will then be placed in a cooler and kept at 4°C until transported to the laboratory. This procedure will be repeated at each sample location and for successive samples at the same location.

Each sample device will be examined in the field to aid in evaluation of site stratigraphy. If boreholes are located at or near the water table, they will be sealed with a cement bentonite grout acceptable for use in monitoring wells.

Monitoring Well Installation Technique and Design

To ensure that representative samples of the groundwater are obtained, monitoring wells will be installed in accordance with NYSDEC monitoring well installation and design specifications for unconsolidated material as presented in Section 5.5.3.2 of the NYSDEC Sampling Guidelines and Protocols. All wells will be constructed of threaded, flush joint, schedule 40 PVC wells materials, supplied by the drilling contractor and will be installed using auger drilling techniques.

Upon completion of the borings, an appropriate length of 2-inch PVC, 0.010-inch slotted well screen will be installed through the hollow stem augers from approximately 10 feet

below to 5 feet above the water table. Should the water table be encountered within 7 feet or less of the ground surface, the well screen will extend to within 2 feet of the ground surface. The remainder of the well will consist of 2-inch PVC casing which will extend over the ground surface. Filter sand will be placed in the annulus between the screen and the borehole to a level of at least 6 inches above the top of the screen. A bentonite pellet seal will be placed on top of the filter sand. The remainder of the annulus will be grouted with a cement bentonite ground acceptable for use in monitoring wells. The surface protection will consist of a lockable steel casing, extending approximately 2 feet above the ground surface and anchored in cement. In areas accessible to vehicular traffic, road boxes may be installed.

To complete the monitoring well installation, each well will be developed by pumping, bailing or an equivalent method. This will remove fines generated during the installation and ensure that hydraulic continuity is established between the well and the aquifer.

Groundwater Flow Direction

A New York registered land surveyor will survey the reference elevation of the top of the PVC monitoring well casings. Water level measurements will be recorded to within 0.01 feet, using an electric water-level indicator. This information will be used to determine groundwater flow direction and construct groundwater contour maps.

Groundwater Sample Collection Methodology

Water samples will be collected no sooner than two weeks after development of the monitoring wells. Prior to sample collection, a minimum of three well volumes will be evacuated using a pump of dedicated bailer depending on well production. After purging, a bailer will be submerged beneath the water column in the well, filled and raised to the surface. The sample collection jar will be filled directly from the bailer. Bailing will continue until each sample jar is filled and closed. Care will be taken to ensure that samples tested for volatile organics have no air space. Sample jars will be

kept closed until the time of collection, to prevent airborne contamination of the sample container. Specific conductance and pH will be measured in the field.

After closing the sample jar, the following information will be recorded on the sample container:

- Job Name
- Sample Media
- Sample Location
- Time and Date of Sampling
- Analysis

The sample will then be placed in a cooler and kept at 4°C until transported to the laboratory for analysis. This process will be repeated for each well.

A field log will also be kept and the following information recorded for each sample:

- Time and Date of Sampling
- Weather
- Name of Sampler
- Water Level Prior to Purge
- Total Well Depth
- Volume Purged
- Purging Method
- Sampler Type
- Presence and Description of any Free Product
- pH and Specific Conductance
- Other Characteristics (odor, color, etc.)
- Turbidity

CHAIN-OF-CUSTODY PROCEDURES

Each sample will be recorded separately on the chain-of-custody manifest as part of the sampling procedure. The information obtained for each sample will include the following:

- Sample Identification
- Sampler's Name
- Time and Date of Sampling
- Sample Laboratory Number
- Analysis to be Performed
- Laboratory Name

Chain-of-Custody procedures will include the following:

- · All samples will be listed on a chain-of-custody manifest.
- All personnel responsible for sampling, transporting and receiving samples will sign the chain-of-custody manifest.
- Analyst's name and laboratory will be recorded on the chain-of-custody manifest.
- Samples will be transported in a secured container with the chain-of-custody manifest attached.
- Samples will be kept in a locked vehicle or within sight of a custodian until received by the laboratory.

PROVISIONS FOR SPLIT SAMPLES

Approximately one week prior to soil and groundwater sampling, the department will be notified of the sampling times and dates. It will then have the option of observing or obtaining split samples.

ANALYTICAL LABORATORY AND METHODS

ANALYTICAL LABORATORY

INTEGRATED ANALYTICAL LABORATORIES (IAL)

Randolph, New Jersey 07869

ANALYTICAL METHODS

For soil impacted by gasoline compounds, the STARS guidance document specifies analysis for VOCs and MTBE using Method 8021. For soil impacted by fuel oil, STARS recommends analysis for both VOCs and base/neutral compounds using Methods 8021 and 8270, respectively. In areas impacted by waste oil, soil will be analyzed for VOCs (method 8260) plus TICs and SVOCs (Method 8270) plus TICs as well as PCBs (method 8082) plus sulfuric acid clean up for PCBs and TAL metals. The analytical methods to be used for drum characterization are as follows: total organic halides - EPA SW- 846 method 3540A/9020A; Ignitability - SW-846 method 1020; and PCBs - SW-846 methods 3540A/9020A. The detection limits are attached to this Plan.

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TOTALE COMPOUNDS		Instrument Average Mod	Reported MDI	Reported MDL	pos s
Acctone	X	1.640	20	2500	
Benzene	X	0.450	5	625	
Aromochloromethane	x	0.540	5	625	-
Bromodichloromethane	X		5	625	- '
Bromoform	Z		5	625	-
Promomethane	X		5	625	- .
2-Butanonc(MEK)	T _x	0.700	20	2500	\dashv
arbon disulfide	X	4	5	625	-
arbon tetrachloride	X	0.540	5	625	-
Chlorobenzenc	X	0,200	5	625	- ·
hloroethane	X	0.370	5	625	-
hloroform	X	0.390	5	625	-
Chloromethane	X	0.730	5	625	_
is-1,2-Dichloroethene	X	0,390	5	625	-
is-1,3-Dichloropropene	Z	0.370	5	625	-
Cyclohexane	x	0.570	-	023	x MDL not available
2-Dibromo-3-chioropropane	X	0:650	5.1	625	NIDE HOLAVAIIADIE
ibromochloromethane	x	0.230	5	625	- ·
1,2-Dibromocthano/Ethylene dibromide(EDB)	_	0.230	5	625	-
2-Dichlorobenzene	X	0.480	5	625	-
3-Dichlorobenzene	X	0.480	5	625	-
1,4-Dichlorobenzene	X	0.370	5	625	
Chlorodifluoromethane	X	0.620	5	625	-
I-Dichlorocthane	x	0.450	5	625	┥.
1,2-Dichloroethane(EDC)	X	0.420	5	625	
1 - Dichlorocthene	×	0.280	5	625	-
2-Dichloropropane	r r	0.680	5	625	-
Ethylbenzene	x	0.370	5	625	
2-Hexanone	x	0.620	20	2500	388
opropylbonzene **(ica noto)	- x	0.370	-5	625	-
Methyl acetate	$\frac{1}{x}$	0.070		023	x MDL not available
Methylcyclohexane	$\frac{1}{x}$				x MDL not available
Methyl-2-pentanone/Methyl Isobutyl Ketone (MIB	$\frac{\hat{x}}{x}$	0.420	20	2500	X WIDE HOLEVAIIADIO
Methylene Chloride	x	0.420	5	625	1
Methyl-tertiary-butyl ether(MTBE)		1.270	5	625	1
rene	X	0.200	5	625	1
1.1,2,2-Tetrachloroethane	X	0.390	5	625	2
Terrachloroethene (PERC)	X	0.310	5	625	(1)
luene	X	0.510	5	625	
tal Xylenes	X	1.160	5	625	
trans-1,2-Dichloroethene	x	0.540	5	625	
ns-1,3-Dichloropropene	x	0.310	5	625	
4-Trichlorobenzene	×	0.390	5	625	
1,1,1-Trichlorocthane	X	0.730	5	625	
,2-Trichloroethane	x	0.450	5	625	- C 1
chloroethene	×	0.340	5	625	
Trichlorofluoromethane					1
Themoreties of the trial te	X	0.480	5	625	
.2-Trichlor-1,2,2-trifluorethane	X	0.480	5	625	x MDL not available

TCL from EPA Internet site 6/8/99. TCL changed 1/1/00, RR confirmed.

	_ /	/ ±	_/	1
SAMMUVALL	/ / /	ر القبل الم	g/ g ([2]
	3 /.,		3 / 2 / 5 / 5 / 5 / 5 / 5 / 5 / 5 / 5 / 5	Ĕ /
		A PER	Reported MDL's,	3 /
Acenaphthene	x	0.111	33.3	· ·
Acenaphthylene	x	0.176	33.3	
Acetophenone		0.170	33.3	X - MDL Not Available
	X	0.182	33.3	A MDL Not Available
Anthracene	X	0.162	33.3	V MDI Net Asseileble
Atrazine Benzaldehyde	X	<u> </u>	-	X - MDL Not Available X - MDL Not Available
		0.249	23.2	X - MDL Not Available
Benzo[a]anthracene	X		33.3	-
Benzo[a]pyrene	X	0.335	33.3	- ·
Benzo[b]fluoranthene	X	0.545	33.3	4 .
Benzo[g,h,i]perylene	х	0.546	33.3	4
Benzo[k]fluoranthene	. X	0.690	33.3	<u> </u>
1,1-Biphenyl	x			X - MDL Not Available
Bis(2-chloroethoxy)methane	X	0.161	33.3	<u></u>] ·
Bis(2-chloroethyl)ether	. х	0.167	33.3	
Bis(2-chloroisopropyl)ether **	X	0.548	33.3	** Compound also known as 2.2-oxybis (1-Chloropropane
Bis(2-ethylhexyl)phthalate	x	0.732	33.3]
4-Bromophenyl-phenylether	x	0.447	33.3].
Butylbenzylphihalate	x	0:570	-33:3	
Caprolactam	х			X - MDL Not Available
Carbazole "	X	0.289	33.3	Ţ.
4-Chloro-3-methylphenol	х	0.432	33.3	1
4-Chloroaniline	x	0.563	33.3	1
2-Chloronaphthalene	х	0.295	33.3	1
2-Chlorophenol	x	0.167	33.3	· .
4-Chlorophenyl-phenylether	x	0,442	. 33.3	1
Chrysene	x	0.402	33.3	1
Dibenz[a,h]anthracene	X	0.421	33.3	1
Dibenzofuran	Z	0.168	. 33,3	1 ·
3,3'-Dichlorobenzidine	z.	0.337	33.3	1
2,4-Dichlorophenol	X	0.516	33.3	· ·
Diethylphthalate	x	0.297	33.3	1
Dimethylphthalate	x	0.248	33.3	· ·
P,4-Dimethylphenol	Z	0.260	33.3	· ·
Di-n-butylphthalate	x	0.478	33.3	1
4,6-Dinitro-2-methylphenol	Z Z	0.887	33:3	·
2,4-Dinitrophenol	x	0.405	33.3	1
2,4-Dinitrotoluene	$\frac{1}{x}$	0.864	33.3	1
2,6-Dinitrotoluene		0,444	33.3	1 ·
Di-n-octylphthalate	x	0.629	33.3	1
luoranthene	х	0.409	33.3	1
Fluorene	x	0.372	33.3	
Hexachlorobenzene	x	0.502	· 33.3	1
lexachlorobutadiene	х	0.418	33.3	
Hexachlorocyclopentadiene	X	0.332	33.3	
Hexachloroethane	x	0.415	33.3] '
ndeno[1,2,3-cd]pyrene	x	0.623	33.3	
sophorone	Z	0.259	33.3	
2-Methylnaphthalene	2	0.184	33.3	

SEMINORME	TCL	Instrument MDL, C (Aqueon	Reported MDL's, Soil (p.ct.)
 Methylphenol (o-Cresol)	x	0.252	33.3
4-Methylphenol (p-Cresol)	X	0.285	33.3
apthalene	X	0.132	33.3
Nitroaniline	X	0.510	33.3
3-Nitroaniline	X	0.696	33.3
Nitroaniline	x	0.929	33.3
trobenzene	X	0.414	33.3
2-Nitrophenol	X	0.819	33.3
4-Nitrophenol	X	0.687	33.3
Nitroso-di-n-propylamine	X	0.245	33.3
Nitrosodiphenylamine	X	0.338	.33.3
Pentachlorophenoi	, x	0.751	33.3
enanthrene	x	0.146	33.3
enol	x	0.269	33.3·
Pyrene	X	0.262	33.3
1,5-Trichlorophenol	X	0.445	33.3
,6-Trichlorophenol	X	0.583	33.3

COMPOUNDS	⁷ CL 8082	Reported Aq (ppb) MDL's	Reported Soli (PPb) MDL's	
Aroclor 1016	Х	0.2	6.68	
Aroclor 1221	Х	0.2	6.68	1
Aroclor 1232	х	0.2	6.68	
Aroclor 1242	Х	0.2	6.68	
Aroclor 1248	X	0.2	6,68	
Aroclor 1254	X	0.2	6.68	
Aroclor 1260	X	0.2	6.68	

PSTIBILES

COMPOUNDS	TCL 8081.4	Reported Aq (Ppb) MDL's	Reported Soil (Ppb) MDL's
4,4'-DDD	X	0.005	0.167
4,4'-DDE	X	0.005	0.167
4,4'-DDT	X	0.005	0.167
Aldrin	X ·	0.005	0.167
alpha-BHC	X	0.005	0.167
alpha-Chlordane	Х	0.005	0.167
beta-BHC	X	0.005	0.167
delta-BHC	X	0.005	0.167
Dieldrin	X	0.005	0.167
Endosulfan I	X	0.005	0.167
Endosulfan II	X	0.005	0.167
Endosulfan sulfate	X	0.005	0.167
Endrin	X	0.005	0.167
Endrin aldehyde	X	0.005	0.167
Endrin Ketone	X	0.005	0.167
gamma-BHC (Lindane)	X	0.005	0.167
gamma-Chlordane	X	0.005	0.167
Heptachlor	X	0.005	0.167
Heptachlor Epoxide	X	0.005	0.167
Methoxychlor	X	0.005	0.167
Toxaphene	X	0.025	0.835

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COMPOUNDS	Aqueous - MDL (Ppm) - ICP 6010	Monitoring Well - MDL (PPM) - ICP/MS	Soil MDL (PPM)- ICP/MS 6020
Aluminum	0.1	0.1	2.0
Antimony	0.1	0.008	2.0
Arsenic	0.1	0.004	0,2
Barium	0.01	0.02	5.0
Beryllium	0.02	0.004	0.2
Cadmium	0.005	0.0006	Ó.2
Calcium	0.4	0.8 .	20.0
Chromium	0.01	0.02	0.6
Cobalt	0.02	0.04	0.4
Copper	0.02	0.04	0.4
Iron	0.05	0.10	3.0
Lead	0.004/0.029	0.004/0:029	2.0
Magnesium	0.10		· 20.0
Manganese	0.005	0.01	0.4
Метситу	· E .	Ø	<u>E</u>
Nickel	0.01	0.02	0.6
Potassium	0.1		20.0
Selenium ·	0.1	800.0	2.0
Silver	0.02	0.0004	
Sodium	0.1		. 20.0
Thallium	0.004	0.008	0,08
Vanadium	0.015	0.03	0.3
Zinc	0.01	0.02	1.0

☑ = MDL for TCLP, Wastewater & Monitoring Well is 0.0005
by ColdVapor and 0.0125 for Soil by Cold Vapor.

-					
	SOIL - Reported	AQUEOUS - Reported			
COMPOUNDS	MDL	MDL	Aqueous Method	Soil Method	Holding Times
Cyanide, Total (ppm)	1.00	50'0	335.2	0106	14 days

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UST ACTIVITIES & PROCEDURES

The following activities will be conducted as part of the underground storage tank and piping removal process.

- All work will be performed in accordance with the appropriate industry standard, American Petroleum Institute publication API 1604; ANSI, NFPA and all other applicable government regulations;
- Acquisition of all required permits including notification for the appropriate oversight authority;
- Overseeing the activities required to ensure proper closure of the tank including excavation, cleaning, safety monitoring, and backfilling to grade;
- Soil sampling the tank excavation (in accordance with NYSDEC STARs and SPOTs Memos)
- Preparing the required closure documentation.

Upon receipt of the required state and local permits, the following tank decommissioning procedures will be employed at the site.

TANK DECOMMISSIONING PROCEDURES

Upon receipt of all permits and approvals, the underground storage tanks and associated piping will be scheduled for removal. Tank decommissioning will be implemented following the appropriate industry standards, as stated above. The tank decommissioning procedures are described below.

TANK PREPARATION

Prior to excavation, the following activities will be performed:

- Underground utilities will be identified by knowledgeable persons.
- Before initiating work in the tank areas, a monitoring instrument will be used to assess vapor and oxygen concentrations in the work area.
- All material transfer piping will be drained, cut and plugged with an impermeable material, or capped to avoid any spillage.
- Residual product found in the tanks will be pumped out and containerized for proper off-site disposal or recycling. (A unit price for disposal is provided in the cost estimate below).

TANK & IMPACTED SOIL REMOVAL PROCEDURES

The following tank removal procedures will be conducted:

- After removing or disconnecting the transfer piping, the soils around the tank will be excavated.
- The excavation will conform to the applicable trenching and shoring requirements.
- Upon removal, the tanks will be placed on plastic and secured prior to inspection and cleaning.
- The tanks will be labeled "do not re-use" after removal from the ground and before removal from the site.

- The tank will be removed from the site as promptly as possible.
- All transfer piping will be removed and properly disposed of.
- Excavated soil will be segregated into piles consisting of a pile petroleum contaminated and non-contaminated soil. Soils will be segregated as potentially clean and contaminated based in visual, olefactory and PID criteria. Soil samples for waste characterization will be collected prior to offsite disposal of impacted soil in accordance with methods described by STARS and SPOTs. Once the soil has been characterized, the soil will disposed of at an appropriate facility.

SITE ASSESSMENT

Site assessment activities, required by the NYSDEC, will be performed to determine if the previous operation of the underground storage tanks or piping has impacted the underlying soils or groundwater at the site. The site assessment activities will include visually inspecting the tanks, excavating soil and collecting post excavation soil samples for certified laboratory analysis as required by the NYSDEC.

SAMPLING AND ANALYSIS

First Environment will field screen all excavated soil with a portable photoionization detector to evaluate potential soil contamination. In accordance with NYSDEC STARS & SPOTS Guidance Documents, one post excavation soil sample will be collected at each side-wall and two composite samples at the tank base. One soil sample will be collected six-inches below the transfer piping for every 20 feet of transfer piping. If visually impacted soil remains subsequent to excavating soil, a grab sample will be collected pursuant to SPOTs Memo #14. Soil samples will be collected by First

Environment and submitted to a NYSDOH ELAP certified laboratory for analysis per STARS and SPOTS requirements depending on the tank contents.

SITE HEALTH AND SAFETY PLAN FIRST ENVIRONMENT, INC.

SITE NAME Provan Ford		
SITE ADDRESS Route 9W and Deyo Place	CLIENT City of Newburgh	
Newburgh, New York 12550	PROJECT # PROVA001	
SITE PHONE	REVISION # DATE	TE6/26/2000
CONTACT Bill Hauser (City of Newburgh)	CONTACT PHONE (914)-562-8642	
OBJECTIVES: (Summarize below)		
Removal of Diesel, qasoline & waste oil underground storage tanks, the installation of soil borings and groundwater monitoring wells, waste drum removal, and excavation, stockpiling and disposal of contaminated soil.	 the installation of soil borings and contaminated soil. 	groundwater monitoring wells,
TYPE OF SITE: (Check as many as applicable)		
()Active (X)Secure ()Enclosed space ()Uncontrolled	()Recovery (X)Remediation	tion ()Unknown
(X)Inactive ()Unsecure ()Landfill ()Industrial	()Well Field ()Other: (specify)	(specify)
DESCRIPTION AND FEATURES: Summarize below. Include principal open	Include principal operations and unusual features (size, containers, buildings,	containers, buildings, dikes,
power line, terrain, etc.).		
Site is currently occupied by Gary's Trucking Repair Service.		
SURROUNDING AREA/DIRECTION:(X)Residential ()Industrial (X)Commercial	ercial ()Rural ()Urban	()Other:
DISTANCE TO NEAREST ENVIRONMENTALLY SENSITIVE RECEPTORS:		
Water Body 1/8 Mile Wetlands Potable Well	SchoolRe	Residence
Building	Depth to Groundwater	
Water Body (Quassaick Creek); Hospital (St. Luke's Hospital)		

Page 2 of 2

THIS PAGE RESERVED FOR MAP & EVACUATION ROUTE

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		Total Control
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or known exposures Former tanker truck cleaning & filling station for commercial trucks. Site has approx. 75 drums containing In addition to history, include complaints from public, previous agency actions, 50% of drums have been overpacked. (Summarize below. injuries, etc.) primarily oils. HISTORY:

Other: (specify)	
()Other:	
() Unknown	
()Gas	
(X) Sludge	
(x) Soil	
(x) Solid	
(X) Liquid	
WASTE TYPES: (X) Liquid	

WASTE CHARACTERISTICS: (Check as many as applicable.)

() Radioactive	
(X) Flammable	
() Corrosive	

() Reactive	() Other: (specify)
(x) Volatile	() Unknown
() Toxic	(X) Inert

Potential for unknown contaminants during soil excavation

PRINCIPAL DISPOSAL METHODS AND PRACTICES:

Tank contents will be pumped out and transported off site for disposal by a licensed waste hauler. Drums and excavated soil will Be transported off site by a licensed waste hauler and disposed of off site. All waste will be disposed of at NYSDEC licensed Facilities

HAZARDS OF CONCERN:

x	Heat Stress (attach guidelines)	×	(X) Noise	() Biological (animals,insects,plants)
×	Cold Stress (attach guidelines)	$\hat{}$) Inorganic Chemicals	(X) 'Physical (terrain, water bodies)
(x)	(X) Explosive/Flammable	(X	(X) Organic Chemicals	(x) Electrical
(x)	(X) Oxygen Deficient	$\hat{}$	() Radiological	<pre>(X) Mechanical (machinery/euipment)</pre>
$\hat{}$	Other: (specify)			

Heat Stress/Cold Stress hazards depend on current temperatures during work activities

HAZARDOUS MATERIAL SUMMARY: (Circle waste type and estimate amounts by category.)

PCBs	Metals	Halogens	Dyes/Inks	Cyanides	Caustics	Acids	CHEMICALS: Amount/Units:
Other	Paint Chips	Non-Ferrous Smelter	Milling/Mine Tailings	Flyash	Ferrous Smelter	Asbestos	SOLIDS: Amount/Units:
Other	Tank Bottoms	Refinery	POTW Sludge	Paint Pigments	Metals	Aluminum	SLUDGES: Amount/Units:
			(abcort).	Other	Non-Halogenated Solvents	Halogenated Solvents	SOLVENTS: Amount/Units:
			ממיני לדדי מדמת מי	Wasta Otl/Sludge	Other	Oily Wastes	OILS: Amount/Units:
Stockpiled Soil		Other	Radiological	Municipal	Laboratory	Hospital	OTHER: Amount/Units:

Pest/Herbicide (specify): (specify):

Phenols

Pickling Liquors

Other (specify): Tank Bottoms

JUSTIFICATION: OVERALL HAZARD EVALUATION: () High (× Medium () Low () Unknown (Where tasks have different hazards, evaluate each.)

Typical from petroleum based UST's rarely are hazardous so an interim medium hazard evaluation has been designated There are unknown contaminant concentrations within the tank bottoms and contaminated soil to be excavated. Contaminant concentrations

FIRE/EXPLOSION POTENTIAL: () High (x) Medium () Low () Unknown

Fire/explosion hazards do exist when removing qasoline USTs. The gasoline USTs to be removed will be inerted prior to removal.

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KNOWN	HIGHEST OBSERVED CONCENTRATION (specify units and media)	PEL/TLV ppm or mg/m³ (specify)	IDLH ppm or mg/m³ (specify)		SYMPTOMS/EFFECTS OF ACUTE EXPOSURE
(VOLATILES)		-			
Benzene (MSDS attached) Toluene (MSDS attached) Ethylbene (MSDS attached) Xylenes (MSDS attached)		1 ppm 100 ppm 100 ppm 100 ppm	1,000 ppm 2,000 ppm 2,000 ppm 1,000 ppm		See MSDS See MSDS See MSDS See MSDS
(SEMI-VOLATILES)					
Acenapthene (MSDS attached) Benzo(a)pyrene (MSDS attached) Bromodichloromethane (MSDS attached) Methyl Ethyl Ketone (MSDS attached) Dichlorodifluoromethane (MSDS attached) 1,2-Dichloroethylene (MSDS attached) Dichlorobenzene (MSDS attached) Dichlorobenzene (MSDS attached) Dichlorobenzene (MSDS attached) Naphthalene (MSDS attached) Phenanthrene (MSDS attached) Pyrene (MSDS attached)	ached) (ched) hattached) hed) ached)				
NA NE U	A = Not Available E = None Established = Unknown	S = Soil GW = Groundwater SW = Surface Wat	A = F = SI =	Air Flyash Sludge	D = Drums TK = Tanks L = Lagoon

Soil Excavation/Stockpiling/Disposal TASK 4 TASK DESCRIPTION/SPECIFIC TECHNIQUE/STANDARD OPERATING PROCEDURES/SITE LOCATION (Attach additional sheets as necessary.) Drum Removal TASK 3 Soil Borings/Well Installations TASK 2 UST Removal TASK 1

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FIELD ACTIVITIES COVERED UNDER THIS PLAN

TYPE

LEVEL OF PROTECTION PRIMARY CONTINGEN

CONTINGENCY

SCHEDULE

08/09/00 3:19 PM

PROTECTIVE CLOTHING: (X) Not Needed	() Encapsulating Suit:	() Splash Suit:	() Apron:	(XX) Tyvek Coverall:	() Saranex Coverall:	() Coverall:	() Other:	GLOVES: () Not Needed	() Undergloves:	(X) Gloves: (XX) - Nitrile and/or latex	() Overgloves:	OTHER: (specify)			•
RESPIRATORY: (X) Not Needed	() SCBA, Airline:	() APR:	(XX) Cartridge:	() Escape Mask:	() Other:	HEAD AND EYE: () Not Needed	(X) Safety Glasses: (XX)	() Face Shield:	() Goggles:	(X) Hard Hat: (XX)	(X) Ear Protection: (XX)	() Other:	BOOTS: () Not needed	Boots: (X), (XX) - Steel Toe	Overboots:

(XX) Contingency

(X) Primary

1234

TASKS: LEVEL:

A B C D Modified

PROTECTIVE EQUIPMENT: (Specify by task.) UST REMOVAL

TASKS: LEVEL: BOOTS: () Not needed Overboots: Boots: (X), () Other: () Goggles: HEAD AND EYE: (X) Ear Protection: (XX) (X) Hard Hat: (XX) () Face Shield: (X) Safety Glasses: (XX) () Other: () Escape Mask: () APR: () SCBA, Airline: RESPIRATORY: A B C D Modified 1234 (XX) Cartridge: (XX) - steel toe (X) Primary () Not Needed (X) Not Needed (X) Contingency OTHER: (specify) GLOVES: () Not Needed () Undergloves: () Coverall: PROTECTIVE CLOTHING: (X) Not Needed () Overgloves: (X) Gloves: (XX) - nitrile and/or latex () Other: () Saranex Coverall: () Apron: () Splash Suit: () Encapsulating Suit: (XX) Tyvek Coverall:

PROTECTIVE EQUIPMENT: (Specify by task.) SOIL BORING/WELL INSTALLATION

PROTECTIVE EQUIPMENT: (Specify by task.) DRUM REMOVAL

																LEVEL:	TASKS:
Overboots:	Boots: (X), (XX)	BOOTS: () Not needed	() Other:	(X) Ear Protection: (XX)	(X) Hard Hat: (XX)	() Goggles:	() Face Shield:	(X) Safety Glasses: (XX)	HEAD AND EYE: () Not Needed	() Other:	() Escape Mask:	(XX) Cartridge:	() APR:	() SCBA, Airline:	RESPIRATORY: (X) Not Needed	A B C D Modified	1 2 3 4 (X) Primary (XX) Contingency
			OTHER: (specify)	() Overgloves:	(X) Gloves: (XX) - nitrile and/or latex	() Undergloves:	GLOVES: () Not Needed	() Other:	() Coverall:	() Saranex Coverall:	(XX) Tyvek Coverall:	() Apron:	() Splash Suit:	() Encapsulating Suit:	PROTECTIVE CLOTHING: (X) Not Needed		

PROTECTIVE EQUIPMENT: (Specify by task.)

SOIL EXCAVATION/STOCKPILING/DISPOSAL

task.)
ζq
(Specify
EQUIPMENT:
MONITORING

INSTRUMENT	TASK	ACTION GUIDELINES	COMMENTS	
Combustible Gas Indicator	1234	<10% LEL 10-20% LEL >20% LEL 21.0% 0 <21.0% 0 <19.5% 0	Continue monitoring. Potential explosion hazard, notify SHSC Explosion hazard; interrupt task/evacuate Oxygen normal Oxygen Deficient, notify SHSC Interrupt task/evacuate	() Not Needed
Radiation Survey Meter	1234	3 x Background: >2 mr/hr Note: Annual ex	id: Notify SHSC Interrupt task/evacuate exposure not to exceed 100 mrem/yr or 50 .rem/hr average	() Not Needed
Photoionization Detector	1234	Specify:		() Not needed
()11.7 ev (X)10.6 e	ev ()10.2 ev	()9.8 ev ()ev	
Type:				
Flame Ionization Detector	1234	Specify:		() Not Needed
Type:				
Personal Gas Detector	1234	Specify:		() Not Needed
Type:				
Dust Monitor	1234	Specify:		() Not Needed
Type:				
Other, Specify:	1234	Specify:		
Type:				
Discharge Control Measures:	ıres:			
Drainage Structures	1234		Hay bails(if needed)	() Not Needed
Surface Water Groundwater	4 c		Booms(if necessary)	
Soil	n m	Specify:		(x) Not Needed
Dust	1234	Specify:		Not
Vapors	2	Specify:		(X) Not Needed
,				

DECONTAMINATION PROCEDURES:

All equipment that comes in contact with contaminated soil/material will be decontaminated on-site prior to removal from the site. If necessary, decontamination water will be drummed on-site for future disposal.

PERSONNEL DECONTAMINATION:

Soils/materials generated on-site. All PPE that comes in contact with contaminated soil/materials will be removed on-site and disposed of along with the contaminated

Containment and Disposal Method:

() Not Needed

SAMPLING EQUIPMENT DECONTAMINATION:

All sampling equipment used will be disposable (one time use only) and will be disposed of following use. When spli During soil borings and well installation, the spoon will be decontaminated with DI water and alconox detergeant. decontamination water will be drummed on-site for future disposal. When split spoons are used, tergeant. If necessary, the

Containment and Disposal Method:

() Not Needed

HEAVY EQUIPMENT DECONTAMINATION:

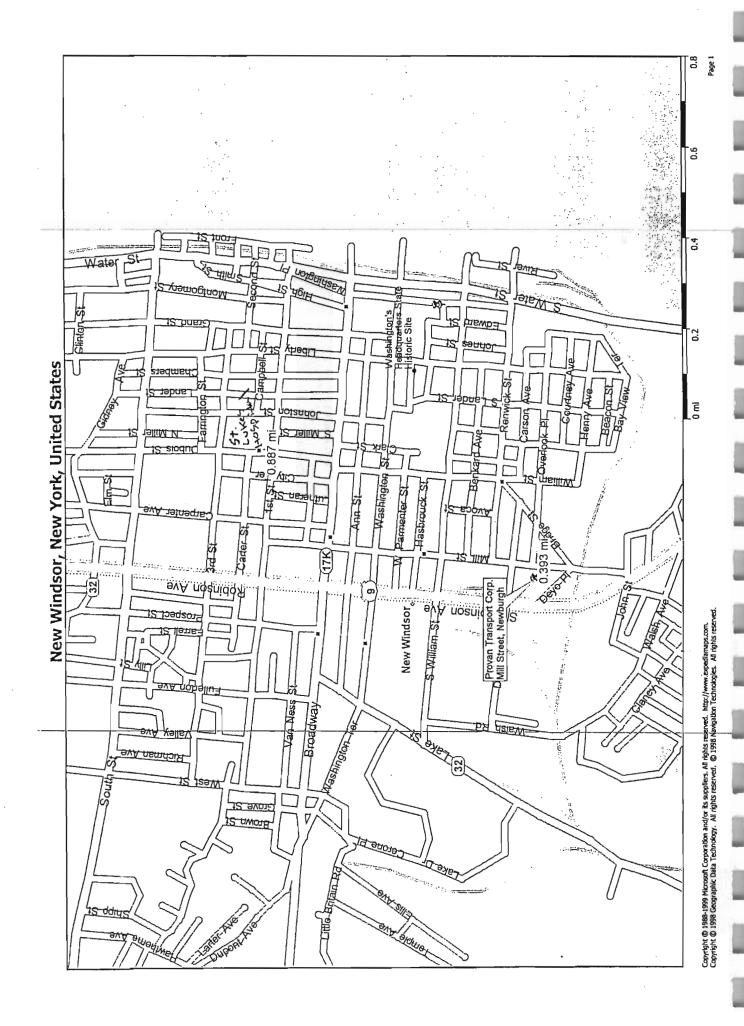
All equipment that comes in contact with contaminated soil/materials will be decontaminated on-site prior to removal from the site. If necessary, decontamination water will be drummed on-site for future disposal.

Containment and Disposal Method:

() Not Needed

EMERG	EMERGENCY RESPONSE EQUIPMENT: (check all that apply) /	(attach map indicating location)
(XX)	20 lb ABC Fire Extinguisher ()	Shower
(XX)	First Aid Kit (XX)	PPE
<u> </u>	Stretcher or Blanket ()	SCBA
(XX)	Potable Water (XX)	Spill Control Equipment
<u> </u>	Eye Wash (XX)	Radio/Phone
^ _	Other	

Hd		973-616-9700	973-616-9700	973-616-9700	973-616-9700	973-616-9700				800-457-7362	:	914-562-1212	914-562-1133		914-632-3701	800-462-6642			Maries of the state of the stat	7 10 10 10 10 10 10 10 10 10 10 10 10 10	914-561-4400			
		B. Tod Delaney	Corey Nachshen	Corey Nachshen	Scott Green	BettyJo Armstrong	and the second s	Vacant								Apple and the second se			TARE TO PASSE AND DESCRIPTION OF THE REAL PROPERTY OF THE PASSE OF THE		St. Luke's Hosp.	70 Dubois St.		1.0 mile
L'IL RGENC. CONTRAIN.	24-Hour Emergency Line:	Health and Safety Manager:	Site Supervisor:	Site H & S Coordinator:	Project Manager:	Personnel Manager	Site Owner:	Site Operator:	EPA Contact:	State Environmental Agency:	State Spill Contractor:	Fire Department:	Police Department:	State Police:	Health Department:	Control Center:	MEDICAL EMERGENCY	Ambulance:	Paramedic:	Medical Consultant:	Hospital Name:	Hospital Address:	Name of Contact at Hospital:	Distance to Hospital: (Attach map and description of
, ONE							202-366-4488	800-424-8802	800-262-8200	800-424-9300				ry Response Plan.	written for the use	lance. The Poison	i, and objectives stated it to exist at the time of	realis to respectively		Date 2-28.2000	Date 3-6-60	Jate 3-13-00	×)
EMENGENCY CONTROL	Water Supply:	Telephone:	Gas:	Electric:	Sewer Authority	Dept of Public Works:	DOT Hazardous Materials Information:	National Response Center Hotline:	CMA Chemical Referral Center:	CHEMTREC:	Other:		CONTINGENCY PLANS:	See First Environment, Inc. Emergency Response Pl	NOTE: This Health and Safety Plan has been written for the	for specific trained personnel who are under medical surveillance.	plan is applicable for the specific purposes, and objectives stated and is representative of carditions believed to exist at the time of the content of the	for its use by others.	HEALTH AND SAFETY PLAN APPROYALS	Prepared by	Reviewed by	HSM Signature Dr Od. Mello		



PERSONNEL & RESPONSIBILITIES: (Attach additional sheets as necessary.)

NAME	FIRM	HEALTH/TRAINING CLEARANCE	RESPONSIBILITIES	SITE TASK	SIGNATURE	DATE
1. Scott Green	First Environment	YES	Project Manager	(j) (2) (3) (4)	(ed)	3-8-90
2. Corey Nachshen	First Environment	Yes	Site Supervisor	1234	110	
3.				1 2 3 4		
4.				1 2 3 4		
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UTILITY MARK-OUT:	Date notified:	Confirm	Confirmation Number:		Initials:	: : : : : : : : : : : : : : : : : : : :

All First Environment personnel assigned to work on-site have attended a 40-hour hazardous materials site operations and safety training course or have met the equivalent experience and training provision of 29 CFR 1910.120, and have been certified medically fit by a qualified occupational physician to work on hazardous sites and to wear a respirator.

By signing above, employees and subcontractors acknowledge that they have read and understand this Site Health and Safety Plan. All site workers under the direction of First Environment must sign this sheet before commencing the site tasks to which they have been assigned for this

Cold stress normally occurs in temperatures at or below freezing, or under certain circumstances, in temperatures of 40°F. Extreme cold for a short time may cause severe injury to exposed body surfaces or result in profound generalized cooling, causing death. Areas of the body which have high surface area-to-volume ratio, such as fingers, toes, and ears, are the most susceptible. Two factors influence the development of a cold weather injury: ambient temperature and the velocity of the wind. For instance, 10°F with a wind of 15 miles per hour (mph) is equivalent in chilling effect to still air at -18°F. An equivalent chill temperature chart relating the actual dry bulb temperature and wind velocity is presented in Table 4-2.

TABLE 4-2
WIND CHILL TEMPERATURE CHART

Estimated Wind					Actual	Temper	rature Re	eading (°	F)				
Speed (in mph)	50	40	30	.20	10.	0:	-10.	-20	-30	-40	-50	-60	
					Equiva	lent Chi	ii Tempe	erature (°	F)				
calm	50	40	30	20	10 -	0	· -1 0	-20	-30	-40	-50	-60	
5	48	37	27	16	6 :.	-5	-15	-26	-36	-47	-57	-68	
10	40	28	16	4	-9	-24	-33	-46	-58	-70	-83	-95	
15	36	22	9	-5	-18	-32	-45	-58	-72	-85	-99	-112	
20	32	18	4	-10	-25	-39	-63	-67	-82	-96	-110	-121	
25	30	16	0	-15	-29	-44	-59	-74	-88	-104	-118	-133	
30	28	.13	-2	-18	-33	-48	-63	-79	-94	-109	-125	-140	
. 35	27	11	4	-20	-35	-51	-67	-82	-98	-113	-129	-145	
40	26	10	-6	-21	-37_	-53	-69	-85_	-100	-116	-132	-148	
(Wind speeds greater than 40 mph have little additional effect.)	Maxim	E DANG lum dar of secu	iger of	false	of expo		sh	GREAT DANGER Flesh may freeze within 30 seconds.					
		Tr	ench fo	oot and	Immersi	on foot i		ur at any		n this ch	art.		

[This chart was developed by the U.S. Army Research Institute of Environmental Medicine, Natick, MA (Source: ACGIH Threshold Lin Values for Chemical Substances and Physical Agents)].

Local injury resulting from cold is included in the generic term frostbite. There are several degrees of tissue damage associated with frostbite. Frostbite of the extremities can be categorized into:

- . Frost Nip or Incipient Frostbite characterized by suddenly blanching or whitening of skin.
- Superficial Frostbite skin has a waxy or white appearance and is firm to the touch, but tissue beneath is
 resilient.
- Deep Frostbite tissues are cold, pale, and solid; extremely serious injury.

Systemic hypothermia is caused by exposure to freezing or rapidly dropping temperature. It can be fatal. symptoms are usually exhibited in five stages: 1) shivering; 2) apathy, listlessness, sleepiness, and (sometimes) rapid cooling of the body to less than 95°F; 3) unconsciousness, glassy stare, slow pulse, and slow respiratory rate; 4) freezing of the extremities; and 5) death. Trauma sustained in freezing or sub-zero conditions requires spec attention because an injured worker is predisposed to secondary cold injury. Special provisions must be made apprevent hypothermia and secondary freezing of damaged tissues in addition to providing for first aid treatment. To avoid cold stress, personnel must wear protective clothing appropriate for the level of cold and physical activity addition to protective clothing, preventive safe work practices, additional training, and warming regimens may be utilized to prevent cold stress.

Safety Precautions for Cold Stress Prevention

- For air temperature of 0°F or less, the hands should be protected by mittens. For exposed skin, continuous exposure should not be permitted when air speed and temperature results in a wind chill temperature of -25°F.
- At air temperatures of 36°F or less, field personnel who become immersed in water or whose clothing becomes
 wet must be immediately provided with a change of clothing and be treated for hypothermia.
- If work is done at normal temperature or in a hot environment before entering the cold, the field personnel must ensure that their clothing is not wet as a consequence of sweating. If wet, field personnel must change into dry clothes prior to entering the cold area.
- If the available clothing does not give adequate protection to prevent hypothermia or frostbite, work must be modified or suspended until adequate clothing is made available or until weather conditions improve.
- Field personnel handling evaporative liquid (e.g., gasoline, alcohol, or cleaning fluids) at air temperatures below 40°F must take special precaution to avoid soaking of clothing or gloves with the liquids because of the added danger of cold injury due to evaporative cooling.

Safe Work Practices

- Direct contact between bare skin and cold surfaces (≤ 20°F) should be avoided. Metal tool handles and/or equipment controls should be covered by thermal insulating material.
- For work performed in a wind chill temperature at or below 10°F, workers should be under constant protective
 observation (buddy system). The work rate should be established to prevent heavy sweating that will result in
 wet clothing. For heavy work, rest periods must be taken in heated shelters and workers should be provided with
 an opportunity to change into dry clothing if needed.
- Field personnel should be provided the opportunity to become accustomed to cold-weather working conditions and required protective clothing.
- Work should be arranged in such a way that sitting or standing still for long periods is minimized.

During the warming regimen (rest period), field personnel should be encouraged to remove outer clothing to permit sweat evaporation or to change into dry work clothing. Dehydration, or loss of body fluids, occurs insidiously in the cold environment and may increase susceptibility to cold injury due to a significant change in blood flow to the extremities. Fluid replacement with warm, sweet drinks and soups is recommended. The intake of coffee should be limited because of diuretic and circulatory effects.

Heat stress is caused by a number of interacting factors, including environmental conditions, clothing, workload, etc., as well as the physical and conditioning characteristics of the individual. Since heat stress is one of the most common illnesses associated with heavy outdoor work conducted with direct solar load and, in particular, because wearing PPE can increase the risk of developing heat stress, workers must be capable of recognizing the signs and symptoms of heat-related illnesses. Personnel must be aware of the types and causes of heat-related illnesses and be able to recognize the signs and symptoms of these illnesses in both themselves and their co-workers.

Heat rashes are the one of the most common problems in hot work environments. Commonly known as prickly heat, a heat rash is manifested as red papules and usually appears in areas where the clothing is restrictive. As sweating increases, these papules give rise to a prickling sensation. Prickly heat occurs in skin that is persistently wetted by unevaporated sweat, and heat rash papules may become infected if they are not treated. In most cases, heat rashes will disappear when the affected individual returns to a cool environment.

Heat cramps are usually caused by performing hard physical labor in a hot environment. These cramps have been attributed to an electrolyte imbalance caused by sweating. It is important to understand that cramps can be caused both by too much and too little salt.

Cramps appear to be caused by the lack of water replenishment. Because sweat is a hypotonic solution (plus or minus 0.3% NaCl), excess salt can build up in the body if the water lost through sweating is not replaced. Thirst cannot be relied on as a guide to the need for water; instead, water must be taken every 15 to 20 minutes in hot environments.

Under extreme conditions, such as working for 6 to 8 hours in heavy protective gear, a loss of sodium may occur. Drinking commercially available carbohydrate-electrolyte replacement liquids is effective in minimizing physiological disturbances during recovery.

Heat exhaustion occurs from increased stress on various body organs due to inadequate blood circulation, cardiovascular insufficiency, or dehydration. Signs and symptoms include pale, cool, moist skin; heavy sweating; dizziness; nausea; headache, vertigo, weakness, thirst, and giddiness. Fortunately, this condition responds readily to prompt treatment.

Heat exhaustion should not be dismissed lightly, however, for several reasons. One is that the fainting associated with heat exhaustion can be dangerous because the victim may be operating machinery or controlling an operation that should not be left unattended; moreover, the victim may be injured when he or she faints. Also, the signs and symptoms seen in heat exhaustion are similar to those of heat stroke, which is a medical emergency.

Workers suffering from heat exhaustion should be removed from the hot environment, be given fluid replacement, and be encouraged to get adequate rest.

Heat stroke is the most serious form of heat stress. Heat stroke occurs when the body's system of temperature regulation fails and the body's temperature rises to critical levels. This condition is caused by a combination of highly variable factors, and its occurrence is difficult to predict.

Heat stroke is a medical emergency. The primary signs and symptoms of heat stroke are confusion; irrational behavior; loss of consciousness; convulsions; a lack of sweating (usually); hot, dry skin; and an abnormally high body temperature, e.g., a rectal temperature of 41°C (105.8°F). If body temperature is too high, it causes death. The elevated metabolic temperatures caused by a combination of work load and environmental heat load, both of which contribute to heat stroke, are also highly variable and difficult to predict.

If a worker shows signs of possible heat stroke, professional medical treatment should be obtained immediately. The worker should be placed in a shady area and the outer clothing should be removed. The worker's skin should be wetted and air movement around the worker should be increased to improve evaporative cooling until professional methods of cooling are initiated and the seriousness of the condition can be assessed. Fluids should be replaced as soon as possible. The medical outcome of an episode of heat stroke depends on the victim's physical flaters and the timing and effectiveness of first aid treatment.

Regardless of the worker's protestations, no employee suspected of being ill from heat stroke should not be sent home or left unattended unless a physician has specifically approved such an order.

Proper training and preventive measures will help avert serious illness and loss of work productivity. Preventing heat stress is particularly important because once someone suffers from heat stroke or exhaustion, that person may be predisposed to additional heat injuries.

Heat Stress Safety Precautions

Heat stress monitoring and work rest cycle implementation should commence when the ambient adjusted temperature exceeds 72°F. A minimum work rest regimen and procedures for calculating ambient adjusted temperature are described in the Table 4-1. Additional personnel must be provided to allow appropriate rest to prevent heat-related illness, as necessary to maintain required production and job continuity.

TABLE 4-1 WORK/REST SCHEDULE

Adjusted Temperature	Work-Rest Regimen Normal Work Ensemble	Work-Rest Regimen Impermeable Ensemble
90°F (32.2°C) or above	After each 45 minutes of work	After each 15 minutes of work
87.5°-90°F (30.8°-32.2°C)	After each 60 minutes of work	After each 30 minutes of work
82.5°-87.5°F (28.1°-30.8°C)	After each 90 minutes of work	After each 60 minutes of work
77.5°-82.5°F (25.3°-28.1°C)	After each 120 minutes of work	After each 90 minutes of work
72.5°-77.5°F (30.8°-32.2°C)	After each 150 minutes of work	After each 120 minutes of work

- For work levels of 250 kilocalories/hour (Light-Moderate Type of Work)
- ^b Calculate the adjusted air temperature (ta adj) by using this equation: ta adj °F = ta °F + (13 x % sunshine). Measure air temperature (ta) with a standard mercury-in-glass thermometer, with the bulb shielded from radiant heat. Estimate percent sunshine by judging what percent time the sun is not covered by clouds that are thick enough to produce a shadow. (100 percent sunshine = no cloud cover and a sharp, distinct shadow; 0 percent sunshine = no shadows.)
- A normal work ensemble consists of cotton coveralls or other cotton clothing with long sleeves and pants.
- The information presented above was generated using the information provided in the American Conference of Governmental Industrial Hygienists (ACGIH) Threshold Limit Values (TLV) Handbook.

In order to determine if the work rest cycles are adequate for the personnel and specific project site conditions additional monitoring of individuals heart rates will be conducted during the rest cycle. To check the heart rate, count the radial pulse for 30 seconds at the beginning of the rest period. If the heart rate exceeds 110 beats per minute, shorten the next work period by one-third and maintain the same rest period

Additional one or more of the following control measures can be used to help control heat stress and are mandatory if any worker has a heart rate (measure immediately prior to rest period) exceeding of 115 beats per minute:

- Workers will be encouraged to drink plenty of water and electrolyte replacement fluids throughout the day.
- On-site drinking water will be kept cool (50 to 60°F).
- A work regimen that will provide adequate rest periods for cooling down will be established, as required.

- All personnel will be advised of the dangers and symptoms of heat stroke, heat exhaustion, and heat cramps.
- Cooling devices, such as vortex tubes or cooling vests, should be used when personnel must wear impermeable clothing in conditions of extreme heat.
- Personnel should be instructed to monitor themselves and co-workers for signs of heat stress and to take additional breaks as necessary.
- A shaded rest area must be provided. All breaks should take place in the shaded rest area.
- Personnel must not be assigned to other tasks during breaks.
- Personnel must remove impermeable garments during rest periods. This includes white Tyvek-type garments.
- All personnel must be informed of the importance of adequate rest, acclimation, and proper diet in the
 prevention of heat stress disorders.

Material Safety Data Sheet

From Genium's Reference Collection Genium Publishing Corporation 1145 Catalyn Street Schenectady, NY 12303-1836 USA (518) 377-8855

Material Name: BENZENE



No. 316

BENZENE (Revision D)

Issued: November 1978 Revised: April 1988

SECTION 1. MATERIAL IDENTIFICATION

25

Description (Origin/Uses): Used in the manufacture of medicinal chemicals, dycs, linoleum, airplane dopes, varnishes, and lacquers; and as a solvent for waxes, resins, and oils.

Other Designations: Benzol; Phene; Phenylhydride; C.H.; NIOSH RTECS No. CY1400000; CAS No. 0071-43-2

Manufacturer: Contact your supplier or distributor. Consult the latest edition of the Chemicalweek Buyers' Guide (Genium ref. 73) for a list of suppliers.

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*See sect. 8 K 4

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SECTION 2. INGREDIENTS AND HAZARDS

Benzene, CAS No. 0071-43-2

Ca 100

SHA PEL

8-Hr TWA: 1 ppm
15-Min Ceiling: 5 ppm

*See NIOSH, RTECS, for additional data with references to irritative, mutagenic, tumorigenic, and reproductive effects.

Action Level: 0.5 ppm

ACGIH TLV, 1987-88

TLV-TWA: 10 ppm, 30 mg/m³

Toxicity Data*

Human, Inhalation, LC₄: 2000 ppm/5 Min

Human, Oral, TD₄: 130 mg/kg _

Human, Inhalation, TC₄: 210 ppm

SECTION 3. PHYSICAL DATA

Boiling Point: 176'F (80°C)
Melting Point: 42°F (5.5°C)

Vapor Pressure: 75 Torrs at 68°F (20°C)

Vapor Density (Air = 1): >1

Water Solubility (%): Slight % Volatile by Volume: 100 Molecular Weight: 78 Grams/Mole

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Molecular Weight: /8 Grams/Mole

Specific Gravity (H,O = 1): 0.87865 at 68°F (20°C)

Appearance and Odor: A colorless liquid; characteristic aromatic odor.

SECTION 4 FIRE	AND EXPLOSION DA	TA	LOWER	UPPER
Flash Point and Method	Autoignition Temperature	Flammability Limits in Air		
12°F (-11.1°C) CC	928°F (498°C)	% by Volume	1.3%	7.1%

Extinguishing Media: Use dry chemical, foam, or carbon dioxide to put out benzene fires. Water may be ineffective as an extinguishing agent because it can scatter and spread the fire. Use water to cool fire-exposed containers, flush spills away from exposures, disperse benzene vapor, and protect personnel attempting to stop an unignited benzene leak.

Unusual Fire or Explosion Hazards: Benzene vapor is heavier than air and can collect in low-lying areas such as sumps or wells. Eliminate all sources of ignition there to prevent a dangerous flashback to the original liquid benzene. Danger: Explosive and flammable benzene vapor-air mixtures can easily form at room temperature; always use this material in a way that minimizes dispersion of its vapor into general work areas.

Special Fire-fighting Procedures: Wear a self-contained breathing apparatus (SCBA) with a full facepiece operated in the pressure-demand or positive-pressure mode.

SECTION 5. REACTIVITY DATA

Benzene is stable in closed containers during routine operations. It does not undergo hazardous polymerization.

Chemical Incompatibilities: Hazardous chemical reactions involving benzene and the following materials are reported in Genium reference 84: bromine pentafluoride, chlorine, chlorine trifluoride, chromic anhydride, nitryl perchlorate, oxygen, ozone, perchlorates, perchloryl fluoride and aluminum chloride, permanganates and sulfuric acid, potassium peroxide, silver perchlorate, and sodium peroxide.

Conditions to Avoid: Avoid all exposure to sources of ignition and to incompatible chemicals.

Hazardous Products of Decomposition: Toxic gases like carbon monoxide (CO) may be produced during benzene fires.



One Genium Plaza Schenectady, NY 12304-4690 USA (518) 377-8854

Material Safety Data Sheets Collection: Sheet No. 317

Toluene

Revision: E, 9/92 Errata: 2/9 Issued: 8/79

Section 1. Material Identification

Toluene (C.H.CH1) Description: Derived from petroleum i.e., dehydrogenation of cycloparaffin fractions followed by the aromatization of saturated aromatic hydrocarbons or by fractional distillation of coal-tar light oil and purified by rectification. Used widely as a solvent (replacing benzene in many cases) for oils, resins, adhesives, natural rubber, coal tar, asphalt, pitch, acetyl celluloses, cellulose paints and varnishes; a diluent for photogravure inks, raw material for organic synthesis (benzoyl & benzilidene chlorides, saccharine, TNT, toluene diisocyanate, and many dyestuffs), in aviation and high octane automobile gasoline, as a nonclinical thermometer liquid and suspension solution for navigational instruments. Other Designations: CAS No. 108-88-3, Methacide, methylbenzene, methylbenzol, phenylmethane, toluol, Tolu-sol. Manufacturer: Contact your supplier or distributor. Consult latest Chemical Week Buyers' Guide(73) for a suppliers list.

Cautions: Toluene is an eye, skin, and respiratory tract imtant becoming narcotic at high centrations. Liver and kidney damage has occurred. Pregnant women chronically exposed to toluene have shown teratogenic effects. Toluene is highly flammable.

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Section 2. Ingredients and Occupational Exposure Limits

Toluene. < 100%: may contain a small amount of benzene (- 1%), xylene, and nonaromatic hydrocarbons.

1991 OSHA PELS

8-hr TWA: 100 ppm (375 mg/m³) 15-min STEL: 150 ppm (560 mg/m³)

1990 IDLH Level 2000 ppm

1990 NIOSH RELS TWA: 100 ppm (375 mg/m³)

STEL: 150 ppm (560 mg/m³)

1992-93 ACGIH TLV (Skin) TWA: 50 ppm (188 mg/m³)

1990 DFG (Germany) MAK* TWA: 100 ppm (380 mg/m³) Half-life: 2 hr to end of shift Category II: Substances with systemic effects

Peak Exposure Limit: 500 ppm, 30 min average value, 2/shift

1985-86 Toxicity Data†

Man, inhalation, TCL: 100 ppm caused hallucinations, and changes in motor activity and changes in psychophysiological tests.

Human, oral, LD, : 50 mg/kg; toxic effects not yet reviewed

Human, eye: 300 ppm caused irritation.

Rat, oral, LD s: 5000 mg/kg

Rat, liver: 30 µmol/L caused DNA damage.

* Available information suggests damage to the developing fetus is probable. tSee NIOSH, RTECS (XSS250000), for additional irritation, mutation, reproductive, and toxicity data.

Section 3. Physical Data

Refraction Index: 1.4967 at 20 °C/D

Bolling Point: 232 °F (110.6 °C) Melting Point: -139 °F (-95 °C) Molecular Weight: 92.15 Density: 0.866 at 68 °F (20/4 °C) Surface Tension: 29 dyne/cm at 68 °F (20 °C) Viscosity: 0.59 cP at 68 'F (20 °C)

Water Solubility: Very slightly soluble, 0.6 mg/L at 68 °F (20 °C) Other Solubilities: Soluble in acetone, alcohol, ether, benzene, chloroform, glacial acetic acid, petroleum ether, and carbon disulfide. Vapor Pressure: 22 mm Hg at 68 'F (20 'C); 36.7 mm Hg at 86 'F (30 'C) Saturated Vapor Density (Air = 0.075 lb/ft^3 or 1.2 kg/m^3): 0.0797 lb/ft^3 or 1.2755 kg/m^3

Odor Threshold (range of all referenced values): 0.021 to 69 ppm

Appearance and Odor: Colorless liquid with a sickly sweet odor.

Section 4. Fire and Explosion Data

Autoignition Temperature: 896 °F (480 °C) LEL: 1.27% v/v UEL: 7.0% v/v Flash Point: 40 °F (4.4 °C) CC

Extinguishing Media: Toluene is a Class 1B flammable liquid. To fight fire, use dry chemical carbon dioxide, or 'alcohol-resistant' foam. Water spray may be ineffective as toluene floats on water and may actually spread fire. Unusual Fire or Explosion Hazards: Concentrated vapors are heavier than air and may travel to an ignition source and flash back. Container may explode in heat of fire. Toluenes' burning rate = 5.7 mm/min and its flame speed = 37 cm/sec. Vapor poses an explosion hazard indoors, outdoors, and in sewers. May accumulate static electricity. Special Fire-fighting Procedures: Because fire may produce toxic thermal decomposition products, wear a self-contained breathing apparatus (SCBA) with a full facepiece operated in pressure-demand or positive-pressure mode. Structural firefighter's protective clothing provides only limited protection. Apply cooling water to sides of tanks until well after fire is out. Stay away from ends of tanks. For massive fire in cargo area, use monitor nozzles or unmanned hose holders; if impossible, withdraw from fire and let burn. Withdraw immediately if you hear a rising sound from venting safety device or notice any tank discoloration due to fire because a BLEVE (boiling liquid expanding vapor explosion) may be imminent. Do not release runoff from fire control methods to sewers or waterways.

Section 5. Reactivity Data

Stability/Polymerization: Toluene is stable at room temperature in closed containers under normal storage and handling conditions. Hazardous polymerization can't occur. Chemical Incompatibilities: Strong oxidizers, concentrated nitric acid, nitric acid, elifuric acid, dinitrogen tetroxide rilver perchlorate, bromine trifluoride, tetranitromethane, and 1,3-dichloro-5,5-dimethyl-2,4-imidazolididione. Conditions to Avold: Contact with heat, ignition sources, or incompatibles. Hazardous Products of Decomposition: Thermal oxidative decomposition of toluene can produce carbon dioxide, and acrid, imitating smoke.

Section 6. Health Hazard Data

Carcinogenicity: The IARC, (164) NTP, (169) and OSHA(164) do not list toluene as a carcinogen. Summary of Risks: Toluene is irritating to the eyes, nose, and respiratory tract. Inhalation of high concentrations produces a narcotic effect sometimes leading to come as well as liver and kidney damage. 93% of inhaled toluene is retained in the body of which 80% is metabolized to benzoic acid, then to hippuric acid and excreted in urine. The remainder is metabolized to o-cresol and excreted or exhaled unchanged. Toluene metabolism is inhibited by alcohol ingestion and is synergistic with benzene, asphalt fumes, or chlorinated hydrocarbons (i.e. perchloroethylene). Toluene is readily absorbed through the skin at 14 to 23 mg/ cm2/hr. Toluene is absorbed quicker during exercise than at rest and appears to be retained longer in obese versus thin victims; presumably due to lipid solubility. There is inconsistent data on toluene's ability to damage bone marrow; chronic poisoning has resulted in anemia and leucopenia w biopsy showing bone marrow hypo-plasia. These reports are few and some authorities argue that the effects may have been due to benzene contampants. Chronic inhalation during pregnancy has been associated with teratogenic effects on the fetus including microcephaly, CNS dysfunction, attentional deficits, developmental delay + language impairment, growth retardation, and physical defects including a small midface, short palpebrai fissures, with deep-set eyes, low-set ears, flat nasal bridge with a small nose, micrognathia, and blunt fingertips. There is some evidence that toluc causes an autoimmune illness in which the body produces antibodies that cause inflammation of its own kidney. Continue on next page



One Genium Plaza Schenectady, NY 12304-4690 USA (518) 377-8854

Material Safety Data Sheets Collection:

Sheet No. 385 Ethylbenzene

Issued: 8/78

Revision: B, 9/92

Section L. Material Identification

Ethylbenzene (C6H2C2H3) Description: Derived by heating benzene and ethylene in presence of aluminum chloride with subsequent distillation, by fractionation directly from the mixed xylene stream in petroleum refining, or dehydrogenation of naphthenes. Used as a solvent, an antiknock agent in gasoline; and as an intermediate in production of synthetic rubber, styrene, cellulose acetate, diethylbenzene, acetophenone, ethyl anthraquinone, propyl oxide, and α-methylbenzol alcohol. Other Designations: CAS No. 100-41-4, ethylbenzol, EB, phenylethane, NCI-C56393. Manufacturer: Contact your supplier or distributor. Consult latest Chemical Week Buyers' Guide(73) for a suppliers list.

3 2* * Skin

absorption.

Cautions: Ethylbenzene is a skin and mucous membrane irritant considered the most irritating of the benzene series. Inhalation causes acute and chronic central nervous system (CNS) effects. It is highly flammable and forms explosive mixtures with air.

0 R PPE - Sec. 8 † Chronic effects

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Section 2. Ingredients and Occupational Exposure Limits

Ethylbenzene, ca >99.0%. Impurities include - 0.1% meta & para xylene, - 0.1% cumene, and - 0.1% toluene.

1991 OSHA PELS

8-hr TWA: 100 ppm (435 mg/m³) 15-min STEL: 125 ppm (545 mg/m³) Action Level: 50 ppm (217 mg/m³)

1990 IDLH Level 2000 ppm

1990 NIOSH REL TWA: 100 ppm (435.mg/m3) STEL: 125 ppm (545 mg/m³) 1992-93 ACGIH TLVs TWA: 100 ppm (434 mg/m³) STEL: 125 ppm (545 mg/m³) 1990 DFG (Germany) MAK TWA: 100 ppm (440 mg/m³) Category 1: local irritants

Peak Exposure Limit: 200 ppm, 5 min momentary value, max of 8/shift Danger of cutaneous absorption

1985-86 Toxicity Data*

Human, inhalation, TC, : 100 ppm/8 hr caused eye effects, sleep, and respiratory changes.

Human, lymphocyte: 1 mmol/L induced sister chromatid

Rat, oral, LD : 3500 mg/kg; toxic effects not yet reviewed Rat (female), inhalation, TCL: 1000 ppm/7 hr/day, 5 days/ wk, for 3 wk prior to mating and daily for 19 days of gestation produced pups with high incidence of extra ribs. (179)

* See NIOSH, RTECS (DA0700000), for additional irritation, mutation, reproductive, and toxicity data.

Section 3. Physical Data

Boiling Point: 277 'F (136 'C) Melting Point: -139 °F (-95 °C) Surface Tension: 31.5 dyne/cm Ionization Potential: 8.76 eV Viscosity: 0.64 cP at 77 'F (25 'C) Refraction Index: 1.4959 at 68 °F (20 °C) Relative Evaporation Rate (ether = 1): 0.0106 Bulk Density: 7.21 lb/Gal at 77 °F (25 °C) Critical Temperature: 651 'F (343.9 'C) Critical Pressure: 35.6 atm

Molecular Weight: 106.16 Density: 0.863 at 77 'F (25 'C)

Water Solubility: Slightly, 14 mg/100 mL at 59 °F (15 °C)

Other Solubilities: Miscible in alcohol, ether; soluble in carbon tetrachloride, benzene, sulfur dioxide, and many organic solvents; insoluble in ammonia

Odor Threshold: 2.3 ppm

Vapor Pressure: 7.1 mm Hg at 68 °F (20 °C); 10 mmHg at 78.62 °F (25.9 °C); 100 mm Hg

165.38 'F (74.1 'C)

Saturated Vapor Density (Air = 0.075 lb/ft3 or 1.2 kg/m3): 0.0768 lb/ft3 or 1.2298 kg/m3

Appearance and Odor: Colorless, flammable liquid with a pungent odor.

Section 4. Fire and Explosion Data

Flash Point: 64 °F (18 °C) CC

Autolgnition Temperature: 810 'F (432 'C)

LEL: 1.0% v/v

UEL: 6.7% v/v

Extinguishing Media: Class 1B Flammable liquid. For small fires, use dry chemical, carbon dioxide, or 'alcohol-resistant' foam. For large fires, use fog or 'alcohol-resistant' foam. Use water only if other agents are unavailable: EB floats on water and may travel to an ignition source and spread fire. Unusual Fire or Explosion Hazards: Burning rate = 5.8 mm/min. Vapors may travel to an ignition source and flash back. Container may explode in heat of fire. EB poses a vapor explosion hazard indoors, outdoors, and in sewers. Special Fire-fighting Procedures: Because fire may produce toxic thermal decomposition products, wear a self-contained breathing apparatus (SCBA) with a full facepiece operated in pressure-demand or positive-pressure mode. Cool container sides with water until well after fire is out. Stay away from ends of tanks. For massive fire in cargo area, use monitor nozzles or unmanned hose holders; if impossible, withdraw from area and let fire burn. Withdraw immediately if you hear rising sound from venting safety device or notice any tank discoloration due to fire. Do not release runoff from fire control methods to sewers or waterways.

Section 5. Reactivity Data

Stability/Polymerization: Ethylbenzene is stable at room temperature in closed containers under normal storage and handling conditions. Hazardous polymerization cannot occur.

Chemical Incompatibilities: Reacts vigorously with oxidizers.

Conditions to Avoid: Exposure to heat and oxidizers.

Hazardous Products of Decomposition: Thermal oxidative decomposition of EB can produce acrid smoke and irritating furnes.

Section 6. Health Hazard Data

Carcinogenicity: The IARC, (164) NTP, (169) and OSHA(164) do not list EB as a carcinogen. Summary of Risks: Occupational exposure to EB alone is rare since it is usually present together with other solvents. EB is irritating to the eyes, skin, and respiratory tract. Vapor inhalation produces varying degrees of CNS effects depending on concentration. The liquid is absorbed through the skin but vapors are not. 56 to 64% of inhaled ethylbenzene is retained and metabolized. Urinary metabolites following exposure to 23 to 85 ppm for 8 hr are mandelic acid (64%), phenylglyoxylic acid (25%), and methylphenylcarbinol/1-phenyl ethanol (5%). Concurrent exposure to xylene and ethylbenzene causes slower excretion of EB metabolites. Based on the rat LD50, one manufacturer gives 3 to 4 oz. as the lethal dose for a 100 lb person. Continue on next page



One Genium Plaza Schenectady, NY 12304-4690 USA (518) 377-8854

Material Safety Data Sheets Collection:

Sheet No. 318 Xylene (Mixed Isomers)

Issued: 11/80 Revision: E, 9/92 Errata: 12/94

NFPA

HMIS

PPE ± + Chronic

Effects \$ Sec. 8

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Section 1. Material Identification

Xylene (Mixed Isomers) (C2H10) Description: The commercial product is a blend of the three isomers [ortho-(o-), meta-(m-), para-(p-)] with the largest proportion being m- xylene. Xylene is obtained from coal tar, toluene by transalkylation, and pseudocumene. Used in the manufacture of dyes, resins, paints, varnishes, and other organics; as a general solvent for adhesives, a cleaning agent in microscope technique; as a solvent for Canada balsam microscopy; as a fuel component; in aviation gasoline, protective coatings, sterilizing catgut, hydrogen peroxide, perfumes, insect repellants, pharmaceuticals, and the leather industry; in the production of phthalic anhydride, isophthalic, and terephthalic acids and their dimethyl esters which are used in the manufacture of polyester fibers; and as an indirect food additive as a component of adhesives. Around the home, xylene is found as vehicles in paints, paint removers, degreasing cleaners, lacquers, glues and cements and as solvent/vehicles for pesticides.

Other Designations: CAS No. 1330-20-7 [95-47-6; 108-38-3; 106-42-3 (o-, m-, p-isomers)], dimethylbenzene, methyltoluene, NCI-C55232, Violet 3, xylol.

Manufacturer: Contact your supplier or distributor. Consult latest Chemical Week Buyers' Guide(13) for a suppliers list.

Cautions: Xylene is an eye, skin, and mucous membrane irritant and may be narcotic in high concentrations. It is a dangerous fire hazard.

Section 2. Ingredients and Occupational Exposure Limits

Xylene (mixed isomers): the commercial product generally contains - 40% m-xylene; 20% each of o-xylene, p-xylene, and ethylbenzene; and small quantities of toluene. Unpurified xylene may contain pseudocumene.

1991 OSHA PELS 8-hr TWA: 100 ppm (435 mg/m³)

15-min STEL: 150 ppm (655 mg/m³)

1990 IDLH Level 1000 ppm

1990 NIOSH RELS TWA: 100 ppm (435 mg/m³) STEL: 150 ppm (655 mg/m³) 1992-93 ACGIH TLVs

TWA: $100 \text{ ppm} (434 \text{ mg/m}^3)$ STEL: 150 ppm (651 mg/m³)

BEI (Biological Exposure Index): Methylhippuric acids in urine at end of shift: 1.5 g/g creatinine

1990 DFG (Germany) MAK TWA: 100 ppm (440 mg/m³)

Category II: Substances with systemic effects

Half-life: < 2 hr

Peak Exposure: 200 ppm, 30 min, average value, 4 peaks per shift

1985-86 Toxicity Data*

Human, inhalation, TC_{Lo}: 200 ppm produced olfaction effects, conjunctiva irritation, and other changes involving the lungs, thorax, or respiration. Man, inhalation, LCto: 10000 ppm/6 hr; toxic effects not yet reviewed.

Human, oral, LD_{Lo}: 50 mg/kg; no toxic effect noted. Rat, oral, LD₅₀: 4300 mg/kg; toxic effect not yet reviewed.

Rat, inhalation, LC50: 5000 ppm/4 hr; toxic effects not yet reviewed.

See NIOSH, RTECS (XE2100000), for additional toxicity data.

Section 3. Physical Data

Boiling Point Range: 279 to 284 'F (137 to 140 'C)*
Boiling Point: ortho: 291 'F (144 'C); meta: 281.8 'F (138.8 'C);

para: 281.3 °F (138.5 °C)

Freezing Point/Melting Point: ortho: -13 °F (-25 °C); meta: -53.3 °F (-47.4 °C); para: 55 to 57 °F (13 to 14 °C) Vapor Pressure: 6.72 mm Hg at 70 °F (21 °C)

Saturated Vapor Density (Air = 1.2 kg/m³): 1.23 kg/m³, 0.077 lbs/ft³

Appearance and Odor: Clear, sweet-smelling liquid.

* Materials with wider and narrower boiling ranges are commercially available.

Molecular Weight: 106.16

Specific Gravity: 0.864 at 20 °C/4 °C Water Solubility: Practically insoluble

Other Solubilities: Miscible with absolute alcohol, ether, and many other organic liquids.

Octanol/Water Partition Coefficient: logKow = 3.12-3.20

Odor Threshold: 1 ppm Viscosity: <32.6 SUS

Section 4. Fire and Explosion Data

Flash Point: 63 to 77 °F (17 to 25 °C) CC Autoignition Temperature: 982 °F (527 °C) (m-) LEL: 1.1 (m-, p-); 0.9 (o-) UEL: 7.0 (m-, p-); 6.7 (o-,

Extinguishing Media: For small fires, use dry chemical, carbon dioxide (CO₂), water spray or regular foam. For large fires, use water spray, fog or regular foam. Water may be ineffective. Use water spray to cool fire-exposed containers. Unusual Fire or Explosion Hazards: Xylene vapors or liquid (which floats on water) may travel to an ignition source and flash back. The heat of fire may cause containers to explode and/or produce irritating or poisonous decomposition products. Xylene may present a vapor explosion hazard indoors, outdoors, or in sewers. Accumulated static electricity may occur from vapor or liquid flow sufficient to cause ignition. Special Fire-fighting Procedures: Because fire may produce toxic thermal decomposition products, wear a self-contained breathing apparatus (SCBA) with a full facepiece operated in pressure-demand or positive-pressure mode. Structural firefighter's protective clothing will provide limited protection. If feasible and without risk, move containers from fire area. Otherwise, cool fire-exposed containers until well after fire is extinguished. Stay clear of tank ends. Use unmanned hose holder or monitor nozzles for massive cargo fires. If impossible, withdraw from area and let fire burn. Withdraw immediately in case of any tank discoloration or rising sound from venting safety device. Do not release runoff from fire control methods to sewers or waterways.

Section 5. Reactivity Data

Stability/Polymerization: Xylene is stable at room temperature in closed containers under normal storage and handling conditions. Hazardous polymerization cannot occur. Xylene is easily chlorinated, sulfonated, or nitrated. Chemical Incompatibilities: Incompatibilities include strong acids and oxidizers and 1,3-dichloro-5,5-dimethyl-2,4-imidazolidindione (dichlorohydrantoin). Xylene attacks some forms of plastics, rubber, and coatings. Conditions to Avoid: Avoid heat and ignition sources and incompatibles. Hazardous Products of Decomposition: Thermal oxidative decomposition of xylene can produce carbon dioxide, carbon monoxide, and various hydrocarbon products.

Section 6. Health Hazard Data

arcinogenicity: The IARC, (164) NTP, (169) and OSHA(164) do not list xylene as a carcinogen. Summary of Risks: Xylene is an eye, mucous inembrane, and respiratory tract irritant. Irritation starts at 200 ppm; severe breathing difficulties which may be delayed in onset can occur at high concentrations. It is a central nervous system (CNS) depressant and at high concentrations can cause coma. Kidney and liver damage can occur with xylene exposure. With prolonged or repeated cutaneous exposure, xylene produces a defatting dermatitis. Chronic toxicity is not well defined, but it is less toxic than benzene. Prior to the 1950s, benzene was often found as a contaminant of xylene and the effects attributed to xylene such as blood dyscrasias are questionable. Since the late 1950s, xylenes have been virtually benzene-free and blood dyscrasias have not been associated with xylenes. Chronic exposure to high concentrations of xylene in animal studies have demonstrated mild reversible decrease in red and white cell counts as well as increases in platelet counts. Continue on next page



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Material Safety Data Sheet Collection

Acenaphthene

MSDS No. 975

Date of Preparation: 10/95

Section 1 - Chemical Product and Company Identification

Product/Chemical Name: Acenaphthene

Chemical Formula: C10H6(CH2)2

CAS Number: 83-32-9

Synonyms: 1,2-dihydroacenaphthylene; 1,8-dihydroacenaphthalene; 1,8-ethylenenaphthalene; ethylenenaphthalene;

naphthyleneethylene; periethylenenaphthalene

Derivation: By passing ethylene and benzene or naphthalene through a red hot tube; by heating tetrahydroacenaphthene with sulfur to 356 °F (180 °C); or by reacting acenaphthenone or acenaphthenequinone by high-pressure hydrogenation in decalin with nickel at 356 to 464 °F (180 to 240 °C). Occurs as a by-product in coal tar production during the high-temperature carbonization or coking of coal.

General Use: Used as an intermediate for dyes, pharmaceuticals, insecticides, fungicides, and plastics.

Vendors: Consult the latest Chemical Week Buyers' Guide. (73)

Section 2 - Composition / Information on Ingredients

Acenaphthene, ca 98 %wt

OSHA PEL

Nuisance Particulates

8-hr TWA: 15 mg/m³ (total dust), 5

mg/m³ (respirable fraction)

ACGIH TLV*

Nuisance Particulates TWA: 10 mg/m³

NIOSH REL

None established

DFG (Germany) MAK

None established

* Notice of intended change to 10 mg/m³ (inhalable), 3 mg/m³ (respirable).

Section 3 - Hazards Identification

ἀሴሴሴሴ Emergency Overview ፌሴሴሴ

Acenaphthene exists as white, needle-like crystals. There is very limited information on the toxicity of acenaphthene. It is irritating to the skin, eyes, and mucous membranes and may cause vomiting if large amounts are ingested. Animal studies indicate possible mutagenic activity. It is combustible.

Potential Health Effects

Primary Entry Routes: Inhalation, skin and eye contact.

Target Organs: Eyes, skin, respiratory tract.

Acute Effects

Inhalation: Irritation of the respiratory tract may occur.

Eye: Irritation may occur. Skin: Irritation may occur.

Ingestion: Ingestion of large amounts may cause vomiting. Irritation of the gastrointestinal tract may occur.

Carcinogenicity: IARC, NTP, and OSHA do not list acenaphthene as a carcinogen.

Medical Conditions Aggravated by Long-Term Exposure: Pre-existing skin disorders.

Chronic Effects: None reported.

HMIS H

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PPE* Sec. 8

Section 4 - First Aid Measures

Inhalation: Remove exposed person to fresh air and support breathing as needed.

Eye Contact: Do not allow victim to rub or keep eyes tightly shut. Gently lift eyelids and flush immediately and continuously with flooding amounts of water. Consult a physician or ophthalmologist if pain or irritation persist.

Skin Contact: Quickly remove contaminated clothing. Rinse with flooding amounts of water followed by a thorough soap and

Ingestion: Never give anything by mouth to an unconscious or convulsing person. Contact a poison control center. Unless the poison control center advises otherwise, have the conscious and alert person drink 1 to 2 glasses of water to dilute. Vomiting may be spontaneuos if large amounts are ingested.

After first aid, get appropriate in-plant, paramedic, or community medical support.

Note to Physicians: Treatment is symptomatic and supportive.

Wilson Risk

Scale R 1 I 2

S K

Section 10 - Stability and Reactivity

Stability: Acenaphthene is stable at room temperature in closed containers under normal storage and handling conditions. Polymerization: Hazardous polymerization does not occur.

Chemical Incompatibilities: Acenaphthene reacts with molecular oxygen in the presence of alkali-earth metal bromides to form acenaphthequinone; reacts with ozone in the presence of alkali-earth metal hydroxides to form 1,8-naphthaldehyde carboxylic acid; and is oxidized to aromatic alcohols and ketones by reaction with transition metal catalysts.

Conditions to Avoid: Exposure to heat, ignition sources, and incompatibles.

Hazardous Decomposition Products: Thermal oxidative decomposition of acenaphthene can produce carbon oxide(s) and thick, acrid smoke.

Section 11 - Toxicological Information

Toxicity Data:*

Acute Effects:

Rat, intraperitoneal, LD50: 600 mg/kg

Mutagenicity:

Microorganisms (species unspecified): 3 mg (-S9) caused mutation.

Section 12 - Ecological Information

Ecotoxicity: Pimephales promelas (fathead minnow), $LC_{50} = 1700 \,\mu\text{g/L/72}$ hr, 1600 $\mu\text{g/L/96}$ hr; Salmo gairdneri (rainbow trout), $LC_{50} = 1570 \,\mu\text{g/L/24}$ hr, 1130 $\mu\text{g/V48}$ hr, 800 $\mu\text{g/L/72}$ hr, 670 $\mu\text{g/L/96}$ hr.

Environmental Fate: In soil, acenaphthene will biodegrade under aerobic conditions with a half-life of 10 to 60 days. A soil absorption coefficient of 2065 to 3230 indicates slight mobility. In water, biodegradation will occur under aerobic conditions with a half-life of 1 to 25 days, as well as photolysis in direct sunlight. Volatilization is another means of removal with half-lives of 11 hr from a model river and 39 days from a model pond which considers the effect of adsorption. In air, acenaphthene reacts with photochemically-produced hydroxyl radicals with a half-life of 7.2 hr.

Section 13 - Disposal Considerations

Disposal: Acenaphthene is a good candidate for rotary-kiln incineration. Contact your supplier or a licensed contractor for detailed recommendations. Follow applicable Federal, state, and local regulations.

Section 14 - Transport Information

DOT Transportation Data (49 CFR 172.101):

Shipping Name:

Environmentally Hazardous Substances, solid, n.o.s. Shipping Symbols: –

Hazard Class: 9
ID No.: UN3077

Packing Group: III Label: Class 9

Special Provisions (172.102): 8,

B54, N50

Packaging Authorizations
a) Exceptions: 173.155

b) Non-bulk Packaging: 173.213

c) Bulk Packaging: 173.240

Quantity Limitations

a) Passenger, Aircraft, or Railcar: None

b) Cargo Aircraft Only: None

Vessel Stowage Requirements

a) Vessel Stowage: A

b) Other: -

Section 15 - Regulatory Information

EPA Regulations:

RCRA Hazardous Waste (40 CFR 261.33): Not listed

CERCLA Hazardous Substance (40 CFR 302.4) listed per CWA. Sec. 307(a)

CERCLA Reportable Quantity (RQ), 100 lb (45.4 kg)

SARA Toxic Chemical (40 CFR 372.65): Not listed

SARA EHS (Extremely Hazardous Substance) (40 CFR 355): Not listed

OSHA Regulations:

Air Contaminant (29 CFR 1910.1000, Table Z-1, Z-1-A): Not listed*

^{*} See NIOSH, RTECS (AB1000000), for additional toxicity data.



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Benzo(a)pyrene

MSDS No. 164

Date of Preparation: 2/94

Section 1 - Chemical Product and Company Identification

Product/Chemical Name: Benzo(a)pyrene

Chemical Formula: C₂₀H₁₂; a polynuclear aromatic hydrocarbon

CAS No.: 50-32-8

Synonyms: BaP; 3,4-benz(a)pyrene; BP; 3,4-benzopyrene; 3,4-benzpyrene. Formerly called 1,2-benzpyrene.

Derivation: Synthesized from pyrene and succinic anhydride.

General Use: Benzo(a)pyrene is no longer used or produced commercially in the US. In its pure form, benzo(a)pyrene may be used as a research laboratory reagent. It also occurs in combustion products of coal, oil, petroleum, wood and other biological matter, in motor vehicle and other gasoline and diesel engine exhaust; in charcoal-broiled foods; in cigarette smoke and general soot and smoke of industrial, municipal, and domestic origin. It occurs naturally in crude oils, shale oils, coal tars, gases and fly ash from active volcances and forest fires. Vendors: Consult the latest Chemical Week Buyers' Guide. (73)

Section 2 - Composition / Information on Ingredients

Benzo(a)pyrene, ca 100 %wt; except in laboratories, benzo(a)pyrene is usually mixed with other coal tar pitch chemicals.

Consider exposure limits for coal tar pitch volatiles as a guideline. However, because benzo(a)pyrene is considered a probable carcinogen to humans, it is recommended that exposures to carcinogens be limited to the lowest feasible concentration.

OSHA PELS

Coal tar pitch volatiles 8-hr TWA: 0.2 mg/m³

ACGIHTLY's

A2: Suspected Human Carcinogen

NIOSH REL

10-hr TWA: 0.1 mg/m³

Carcinogen; coal tar pitch volatile, cyclohexane extractable fraction.

DFG (Germany) MAK None established IDLH Level

700 mg/m³
Coal tar pitch volatiles (benzene soluble

fraction)

Section 3 - Hazards Identification

ፌሴሴሴሴ Emergency Overview ልሴሴሴሴ

Benzo(a)pyrene is a pale yellow, crystalline solid or powder that is irritating to the skin, eyes, and respiratory tract. It is a carcinogen and mutagen. Handle with extreme caution!

Potential Health Effects

Primary Entry Routes: Inhalation, ingestion. Target Organs: Respiratory system, bladder, kidneys, skin.

Acute Effects: Inhalation: Respiratory tract irritation. Eye: Irritation and/or burns on contact. Skin: Irritation with burning sensation, rash, and redness; dermatitis on prolonged exposure. Sunlight enhances effects (photosensitization). Ingestion: None reported.

Carcinogenicity: IARC, NTP, NIOSH, ACGIH, EPA, and MAK list benzo(a)pyrene as: an IARC 2A (probably carcinogenic to humans: limited human evidence, sufficient evidence in experimental animals), an NTP-2 (reasonably anticipated to be a carcinogen: limited evidence from studies in humans or sufficient evidence from studies in experimental animals), a NIOSH-X (carcinogen defined with no further categorization); an ACGIH TLV-A2 (suspected human carcinogen: carcinogenic in experimental animals, but available epidemiological studies are conflicting or insufficient to confirm an increased risk of cancer in exposed humans); an EPA-B2 (sufficient evidence from animal studies, inadequate evidence or no data from epidemiological studies); and an MAK-A1 (capable of inducing malignant tumors as shown by experience with humans) carcinogen, respectively.

Medical Conditions Aggravated by Long-Term Exposure: Respiratory system, bladder, kidney, and skin disorders. Chronic Effects: Inhalation: Cough and bronchitis. Eye: Photosensitivity and irritation. Skin: Skin changes such as thickening, darkening, pimples, loss of color, reddish areas, thinning of the skin, and warts. Sunlight enhances effects (photosensitization). Other: Gastrointestinal (GI) effects include leukoplakia (a pre-cancerous condition characterized by thickened white patches of epithelium on mucous membranes, especially of the mouth). Cancer of the lung, skin, kidneys, bladder, or GI tract is also possible. Smoking in combination with exposure to benzo(a)pyrene increases the chances of developing lung cancer. Persons with a high degree of inducibility of the enzyme aryl hydrocarbon hydroxylase may be a high risk population.

Comments: Pregnant women may be especially susceptible to exposure effects of benzo(a)pyrene; exposure may damage the fetus. In general, polyaromatic hydrocarbons such as benzo(a)pyrene tend to localize primarily in body fat and fatty tissues (for ex. breasts) and are excreted in breast milk. Benzo(a)pyrene may also affect the male reproductive system (testes and sperm).

Section 4 - First Aid Measures

Inhalation: Remove exposed person to fresh air and support breathing as needed.

Wilson Risk Scale R 1 I 4 S 4 K 1

HMIS
H 2*
F I
R 0
*Chronic
Effects
PPE†
†Sec. 8

annual basis for employees 45 yr of age or older or with 10 or more years of exposure to coal tar pitch volatiles. Train workers about the hazards of benzo(a)pyrene and the necessary protective measures to prevent exposure. Periodically inspect lab atmospheres, surfaces such as walls, floors, and benches, and interior of fume hoods and air ducts for contamination. Post appropriate signs and labels on doors leading into areas where benzo(a)pyrene is used.

Respiratory Protection: Seek professional advice prior to respirator selection and use. Follow OSHA respirator regulations (29 CFR 1910.134) and, if necessary, wear a MSHA/NIOSH-approved respirator. The following respirator recommendations are for coal tar pitch volatiles. For any unknown concentration, wear any SCBA with a full facepiece and operated in a pressure-demand or other positive pressure mode, or any supplied-air respirator with a full facepiece and operated in a pressure-demand or other positive pressure mode in combination with an auxiliary SCBA operated in pressure-demand or other positive pressure mode. For escape, wear any air-purifying full facepiece respirator (gas mask) with a chin-style or front- or back-mounted organic vapor canister having a high-efficiency particulate filter, or any appropriate escape-type SCBA. Select respirator based on its suitability to provide adequate worker protection for given working conditions, level of airborne contamination, and presence of sufficient oxygen. For emergency or nonroutine operations (cleaning spills, reactor vessels, or storage tanks), wear an SCBA. Warning! Air-purifying respirators do not protect workers in oxygen-deficient atmospheres. If respirators are used, OSHA requires a written respiratory protection program that includes at least: medical certification, training, fit-testing, periodic environmental monitoring, maintenance, inspection, cleaning, and convenient, sanitary storage areas.

Protective Clothing/Equipment: Wear chemically protective gloves, boots, aprons, and gauntlets to prevent prolonged or repeated skin contact. In animal laboratories, wear protective suits (disposable, one-piece and close-fitting at ankles and wrists), gloves, hair covering, and overshoes. In chemical laboratories, wear gloves and gowns. Wear protective eyeglasses or chemical safety, gas-proof goggles, per OSHA eye- and face-protection regulations (29 CFR 1910.133). Because contact lens use in industry is controversial, establish your own policy.

Safety Stations: Make available in the work area emergency eyewash stations, safety/quick-drench showers, and washing facilities.

Contaminated Equipment: Shower and change clothes after exposure or at the end of the workshift. Separate contaminated work clothes from street clothes. Launder before reuse. Remove benzo(a)pyrene from your shoes and clean personal protective equipment. Use procedures to ensure laundry personnel are not exposed.

Comments: Never eat, drink, or smoke in work areas. Practice good personal hygiene after using this material, especially before eating, drinking, smoking, using the toilet, or applying cosmetics.

Section 9 - Physical and Chemical Properties

Physical State: Solid

Appearance and Odor: Pale yellow monoclinic needles

with a faint, aromatic odor.

Vapor Pressure: >1 mm Hg at 68 'F (20 'C)

Formula Weight: 252,30

Specific Gravity (H2O=1, at 4 °C): 1.351

Water Solubility: Insoluble; 0.0038 mg (+/- 0.00031 mg)

in 1 L at 77 °F (25 °C)

Other Solubilities: Ether, benzene, toluene, xylene, concentrated hydrosulfuric acid; sparingly soluble in alcohol, methanol.

Boiling Point: >680 °F (>360 °C); 540 °F (310 °C) at 10 mm

Melting Point: 354 °F (179 °C)

Octanol/Water Partition Coefficient: log Kow= 6.04

Section 10 - Stability and Reactivity

Stability: Benzo(a)pyrene is stable at room temperature in closed containers under normal storage and handling conditions. It undergoes photo-oxidation when exposed to sunlight or light in organic solvents and is also oxidized by chromic acid and orone.

Polymerization: Hazardous polymerization cannot occur.

Chemical Incompatibilities: Strong oxidizers (chlorine, bromine, fluorine) and oxidizing chemicals (chlorates, perchlorates, permanganates, and nitrates).

Conditions to Avoid: Avoid heat and ignition sources and incompatibles.

Hazardous Decomposition Products: Thermal oxidative decomposition of benzo(a)pyrene can produce carbon monoxide and carbon dioxide.

Section 11 Toxicological Information

Toxicity Data:*

Tumorigenic Effects:

Rat, oral: 15 mg/kg produced gastrointestinal and musculoskeletal tumors.

Mouse, inhalation: 200 ng/m³/6 hr administered intermittently over 13 weeks produced tumors of the lungs

Rabbit, skin: 17 mg/kg administered intermittently over 57 weeks produced tumors of the skin and appendages.

Teratogenicity:

Rat, oral: 2 g/kg administered 28 days prior to mating and 1-22 days of pregnancy produced a stillbirth.

Rat, oral: 40 mg/kg on the 14th day of pregnancy caused changes in the extra embryonic structures.

Mouse, oral: 75 mg/kg administered to the female during the 12-14 day of pregnancy produced biochemical and metabolic effects on the newborn.



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Bromodichloromethane

MSDS No. 973

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Date of Preparation: 10/95

Section 1 - Chemical Product and Company Identification

Product/Chemical Name: Bromodichloromethane

Chemical Formula: CHBrCl2

CAS Number: 75-27-4

Synonyms: BDCM; dichlorobromomethane; methane, bromodichloro-; monobromodichloromethane

Derivation: Produced by treating the chlorine analog with hydrogen bromide in the presence of an aluminum halide catalyst.

Bromodichloromethane is found in treated (chlorinated) water at levels significantly higher than untreated water.

General Use: Used in research as a laboratory reagent or for chemical synthesis. Formerly used as a flame retardant, in fire

extinguishers, as a solvent for waxes, fats, and resins, and to separate minerals from salts.

Vendors: No longer produced or used commercially in the US.

Section 2 - Compositión / Information on Ingredients

Bromodichloromethane, ca 98+ % vol

OSHA PEL

None established

ACGIH TLV

None established

NIOSH REL

None established

DFG (Germany) MAK

None established

Section 3 - Hazards Identification

ជជជជជ Emergency Overview ជជជជជ

Bromodichloromethane exists as a colorless, volatile liquid. It is toxic to the central nervous system, resulting in sleep disturbances and fatigue. Liver and kidney damage may also occur. Bromodichloromethane is considered to be a potential human carcinogen based on animal studies. It is nonflammable as indicated by its former use as a flame retardant.

Potential Health Effects

Primary Entry Routes: Inhalation

Target Organs: Central nervous system (CNS), blood, liver, kidneys.

Acute Effects

Inhalation: CNS effects including fatigue, sleep disturbances, and incoordination. Methemoglobinemia (lack of oxygenated blood in body tissues), evident by a bluish tint to skin, lips, and fingernails, is also likely.

Eye: No effects reported.

Skin: No effects reported.

Ingestion: CNS effect may occur as via inhalation.

Carcinogenicity: IARC (Group 2B, possible human carcinogen with limited human evidence in the absence of animal evidence) and NTP (Group 2, probable human carcinogen with sufficient animal evidence and insufficient human evidence).

Medical Conditions Aggravated by Long-Term Exposure: None reported.

Chronic Effects: Repeated exposure to bromodichloromethane has resulted in liver and kidney carcinogenicity in animals. It is not known whether or not chronic exposure will produce cancer in humans. Human mutation data has been reported (Sec. 11). Other: Bromodichloromethane is metabolized to carbon monoxide in the liver. Therefore, toxic effects to the nervous system are due to carbon monoxide formation. This also explains why the liver can be adversely affected. Exposure to acetone and chlordecone potentiates liver toxicity.

Section 4 - First Aid Measures

Inhalation: Remove exposed person to fresh air and support breathing as needed.

Eye Contact: Do not allow victim to rub or keep eyes tightly shut. Gently lift eyelids and flush immediately and continuously with flooding amounts of water for at least 15 min.

Skin Contact: Quickly remove contaminated clothing. Rinse with plenty of water followed by a thorough soap and water wash. Ingestion: Never give anything by mouth to an unconscious or convulsing person. Contact a poison control center. Unless the poison control center advises otherwise, have the conscious and alert person drink 1 to 2 glasses of water to dilute. Inducement of vomiting should not be necessary except in cases of large ingestions.

After first aid, get appropriate in-plant, paramedic, or community medical support.

Note to Physicians: Treatment is symptomatic and supportive.

Wilson Risk Scale

R 1 I 3 S 1

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HMIS H 2* F 0 R 0

*Chronic Effects PPE†

PPE†

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Section 10 - Stability and Reactivity

Stability: Bromodichloromethane is stable at room temp. in closed containers under normal storage and handling conditions.

Polymerization: Hazardous polymerization does not occur. Chemical Incompatibilities: None reported.

Conditions to Avoid: Exposure to elevated temperatures.

Hazardous Decomposition Products: Thermal oxidative decomposition of bromodichloromethane can produce chlorine, bromine, phosgene, and carbon oxide(s) gases.

Section 11 - Toxicological Information

Toxicity Data:*

Acute Oral Effects:

Rat, oral, LD₅₀: 430 mg/kg caused somnolence, tremor, and liver changes.

Mouse, oral, LD₅₀: 450 mg/kg caused changes in circulation, fatty liver degeneration, and hemorrhage. Mutagenicity:

Human, lymphocyte: 400 μmol/L caused sister chromatid exchange.

Mouse, lymphocyte: 180 mg/L (+S9) caused mutation.

Multiple Dose Toxicity Data:

Rat, oral: 5670 mg/kg administered continuously for 4 weeks resulted in changes in liver weight and serum composition, and weight loss or decreased weight gain.

Tumorigenicity:

Rat, oral: 25.5 g/kg administered continuously for 2 years produced gastrointestinal and kidney tumors.

Mouse, oral: 38.25 g/kg administered continuously for 2 years produced liver tumors.

Section 12 - Ecological Information

Ecotoxicity: Data not found.

Environmental Fate: If released to soil, bromodichloromethane is expected to volatilize due to its high vapor pressure. Because it is highly mobile in soil, leaching will occur and biodegradation will occur under anaerobic conditions. In water, volatilization is expected to be relatively rapid with a typical half-life from streams and rivers of 35 hr. Bioconcentration is not expected to be significant (BCF = 0.72 to 1.37). In air, bromodichloromethane is expected to exist almost entirely in the vapor phase and react with hydroxyl radicals in the troposphere (half-life = 6.65 months). This persistent half-life indicates that long-range global transport is possible. Any bromodichloromethane that does not undergo reaction with hydroxyl radicals can reach the stratosphere, where it will be destroyed via photolysis.

Section 13 - Disposal Considerations

Disposal: Contact your supplier or a licensed contractor for detailed recommendations. Follow applicable Federal, state, and local regulations.

Section 14 - Transport Information

DOT Transportation Data (49 CFR 172.101):

Shipping Name: Environmentally Hazardous Substance, liquid, n.o.s.

Shipping Symbols: -

Hazard Class: 9 ID No.: UN3082 Packing Group: III Label: Class 9

Special Provisions (172.102): 8,

N50, T1

Packaging Authorizations a) Exceptions: 173.155

b) Non-bulk Packaging: 173.203

c) Bulk Packaging: 173.241

Quantity Limitations

a) Passenger, Aircraft, or Railcar: None

b) Cargo Aircraft Only: None

Vessel Stowage Requirements

a) Vessel Stowage: A

b) Other: -

Section 15 - Regulatory Information

EPA Regulations:

RCRA Hazardous Waste (40 CFR 261.33): Not listed

Listed as a CERCLA Hazardous Substance (40 CFR 302.4) per CWA, Sec. 307(a)

CERCLA Reportable Quantity (RQ), 5000 lb (2270 kg)

Listed as a SARA Toxic Chemical (40 CFR 372.65)

SARA EHS (Extremely Hazardous Substance) (40 CFR 355): Not listed

OSHA Regulations:

Air Contaminant (29 CFR 1910.1000, Table Z-1, Z-1-A): Not listed

^{*} See NIOSH, RTECS (PA5310000), for additional toxicity data.



One Genium Plaza Schenectady, NY 12304-4690 USA (518) 377-8854

Material Safety Data Sheets Collection:

Sheet No. 303 Methyl Ethyl Ketone

Issued: 9/79

Revision: D, 5/93

tion 1. Material Identification : 41 M hyl Ethyl Ketone (CH3CH2COCH3) Description: Derived by dehydrogenation or selective oxidation of sec-butyl 1 -NFPA alcohol; from mixed n-butylenes and sulfuric acid, followed by distillation to separate sec-butyl alcohol and then dehydro-2.7 (3)genation; or by controlled oxidation of butane. Used as a solvent for printing inks and cellulose compounds (nitrocellulose S 2* in micular), constituent of dewaxing compositions; in the manufacture of acrylic and vinyl surface coatings, paint • Skin re wers, cements and adhesives, artificial leather, cosmetics, lubricating oils, pharmaceuticals, smokeless powder and absorption explosives, and cleaning fluids. HMIS Other Designations: CAS No. 78-93-3, 2-butanone, ethyl methyl ketone, methylacetone, 2-oxobutane. 2 3 M ufacturer: Contact your supplier or distributor. Consult latest Chemical Week Buyers' Guide(73) for a suppliers list. 0 Candons: Methyl ethyl ketone is a flammable liquid. Vapors are irritating to the eyes and respiratory tract. It potentiates the PPE† neurotoxic potential of other chemicals and some findings suggest it may be neurotoxic itself. t Sec. 8

Setion 2. Ingredients and Occupational Exposure Limits

M yl ethyl ketone, ca 100% 1992 OSHA PELs

Tr-sitional Limit

8 TWA: 200 ppm (590 mg/m³) First Rule Limits

8-hr TWA: 200 ppm (590 mg/m³) 15 nin STEL: 300 ppm (885 mg/m³)

19 NIOSH RELs TWA: 200 ppm (590 mg/m³) STFL: 300 ppm (885 mg/m³) 1993-94 ACGIH TLVs

TWA: 200 ppm (590 mg/m³) STEL: 300 ppm (885 mg/m³)

1991 DFG (Germany) MAK TWA: 200 ppm (590 mg/m³)

Half-Life: < 2 hr

Category II: Substances with systemic effects Peak Exposure Limit: 400 ppm, 30 min.

average value, 4/shift

1992 Toxicity Data*

Rabbit, skin: 500 mg/24 hr caused severe irritation.

Rat, oral, LD50: 2737 mg/kg

Rat, inhalation, TC_{Lo}: 3000 ppm/7 hr from 6 to 15 days of pregnancy caused craniofacial abnormalities (including nose and tongue) as well as developmental abnormalities of the urogenital system and homeostasis. Human, inhalation, TC_{Lo}: 100 ppm/5 min caused eye

100

Action Section 1

irritation and respiratory changes.

1990 IDLH Level

3000 ppm

* See NIOSH, RTECS (EL6475000), for additional irritation, mutation, reproductive, and toxicity data.

Se tion 3. Physical Data

mang Point: 176 °F (80 °C)
Melting Point: -122.6 °F (-85.9 °C)

Varor Pressure: 71.2 mm Hg at 68 °F (20 °C)
Sai rated Vapor Density (Air = 0.075 lb/ft³ or

1. kg/m³): 0.085 lb/ft³ or 1.368 kg/m³

Bulk Density: 6.71 lb/gal

Remotion Index: 1.379 at 68 °F (20 °C)

Cr :al Temperature: 504 °F (262 °C)

Critical Pressure: 41 atm

Molecular Weight: 72.1

Density: 0.8045 g/mL at 68 °F (20 °C)

Water Solublity: MEK in water = 28%; water in MEK = 12.5% Other Solublities: Soluble in alcohol, benzene, ether, and fixed oils.

Surface Tension: 24.6 dyne/cm at 68 °F (20 °C)

Ionization Potential: 9.54 eV

Viscosity: 0.4 cP at 77 'F (25 'C)

Relative Evaporation Rate (ether = 1): 2.7

Octanol/Water Partition Coefficient: log Kow = 0.26 to 0.29

Appearance and Odor: Colorless, volatile, liquid with a sweet mint or acetone-like odor. The odor threshold is 25 ppm.

Se ion 4. Fire and Explosion Data

Flash Point: 16 'F (-9 °C) Autoignition Temperature: 759 'F (404 °C) | LEL: 1.4% at 200 "F (93 °C) | UEL: 11.4% at 200 °F (93 °C)

Ext guishing Media: A Class 1B flammable liquid. For small fires, use dry chemical, carbon dioxide, water spray, or alcohol-resistant foam. For

larg lires, use water spray, fog, or alcohol-resistant foam.

Unusual Fire/Explosion Hazards: Vapors may travel to ignition source and flash back. Container may explode in fire. Burning rate = 4.1 mm/min.

Special Fire-fighting Procedures: Because fire may produce toxic thermal decomposition products, wear a self-contained breathing apparatus (SC A) with a full facepiece operated in pressure-demand or positive-pressure mode. Structural firefighters' protective clothing provides only liming a protection. If possible without risk, move container from fire area. Apply cooling water to container sides until well after fire is out. Stay away from ends of tanks. For massive fire in cargo area, use monitor nozzles or unmanned hose holders; if impossible, withdraw and let fire burn. Withdraw immediately if you hear a rising sound from venting safety device or notice any tank discoloration due to fire. Do not release runoff from international to sewers or waterways; dike for proper disposal.

Section 5. Reactivity Data

Stability/Polymerization: Methyl ethyl ketone is stable at room temperature in closed containers under normal storage and handling conditions.

Haz dous polymerization cannot occur.

The ical Incompatibilities: Include chlorosulfonic acid, oleum (furning sulfuric acid), potassium-t-butoxide, hydrogen peroxide + nitric acid, 2-panol (forms explosive peroxides), chloroform + alkali, amines, ammonia, inorganic acids, caustics, copper, isocyanates, pyridines, and strong irres. MEK will soften or dissolve some plastics.

dons to Avoid: Exposure to heat, ignition sources, and incompatibles.

Lazar dous Products of Decomposition: Thermal oxidative decomposition of MEK can produce carbon dioxide gas and acrid smoke.

Section 6. Health Hazard Data

ar togenicity: The LARC. (183) NTP. (169) and OSHA(183) do not list methyl ethyl ketone as a carcinogen.

fummary of Risks: MEK vapors are irritating to the eyes and respiratory tract. Inhalation causes varying degrees of central nervous system



One Genium Plaza Schenectady, NY 12304-4690 USA (518) 377-8854

Material Safety Data Sheets Collection:

Sheet No. 308 Dichlorodifluoromethane

Issued: 11/77

Revision: D, 9/92

Section 1. Material Identification Dichlorodifluoromethane (CCl₂F₂) Description: Derived by reacting carbon tetrachloride and hydrogen fluoride in the Genlum 1.3 presence of an antimony halide catalyst, or by high temperature chlorination of vinylidene fluoride. Formerly used as an serosol propellant but due to it's role in ozone depletion, this use was banned by the USEPA on 12/15/78. Still used as a S refrigerant, leak detection agent, blowing agent for polymeric foams, a foaming agent in fire extinguishers, a solvent or diluent in fumigants for food and medical equipment sterilization; in the manufacture of glass bottles, preparing frozen tissue sections, in paint and varnish removers, water purification, and in thermal expansion valves. Other Designations: CAS No. 75-71-8, Algofrene type 2, Arcton, difluorodichloromethane, Eskimon-12, FC-12, fluorocarbon-12, Freon-12, Genetron-12, Halon, Isotron-12, propellant-12, refrigerant-12, Ucon-12. Manufacturer: Contact your supplier or distributor. Consult latest Chemical Week Buyers' Guide(73) for a suppliers list. PPE*

Cautions: Dichlorodiffuoromethane is considered low in toxicity but is a simple asphyxiant and produces weak irritative and narcotic effects in high concentrations. It may also induce serious cardiac arrythmias at high concentrations and in susceptible individuals. Contact with the liquid can cause frostbite.

Section 2. Ingredients and Occupational Exposure Limits

Dichlorodifluoromethane, ca 99.9%

1991 OSHA PEL

8-hr TWA: 1000 ppm (4950 mg/m³) 1990 IDLH Level

50,000 ppm 1990 NIOSH REL

10-hr TWA: 1000 ppm (4950 mg/m³)

1992-93 ACGIH TLV

TWA: 1000 ppm (4950 mg/m³) 1990 DFG (Germany) MAK TWA: 1000 ppm (5000 mg/m³) Category IV: Substances eliciting

very weak effects.

Peak Exposure Limit: 2000 ppm, 60 min momentary value, * 3/shift 1985-86 Toxicity Data†

Human, inhalation, TCLo: 200,000 ppm/30 min caused conjunctival irritation, fibrosing aveolitis, and changes in the liver.

Rat, inhalation, LC50: 80 pph/30 min; toxic effects not yet reviewed

Rabbit, inhalation, LCL: 80 pph/30 min; toxic effects not yet reviewed

* The momentary value is the level which should never be exceeded.

† See NIOSH, RTECS (PA8200000), for additional toxicity data.

Section 3. Physical Data

Boiling Point: -22 'F (-29.8 'C) Freezing Point: -252 °F (-158 °C) Vapor Pressure: 5.7 atm at 68 °F (20 °C)

Vapor Density (Air = 1): 4.16

Surface Tension: 9 dyne/cm Viscosity: 0.262 cP at 70 °F (21 °C)

Ionization Potential: 11.75 eV

Molecular Weight: 120.9

Density (liquid): 1.486 at -22 °F (-29.8 °C)

Water Solubility: Nearly insoluble, 0.03% at 77 °F (25 °C)

Other Solubilities: Soluble in most organic solvents, such as alcohol, ether, benzene, amyl chloride, bromobenzene, bromoform, n-butyl alcohol, butyl butyrate, carbon tetrachloride, and chloroform.

Critical Temperature: 232.7 °F (111.5 °C)

Critical Pressure: 43.2 atm

Appearance and Odor: Colorless, practically odorless (smells like ether at 20% in air) gas.

Section 4. Fire and Explosion Data

Flash Point: Nonflammable Autoignition Temperature: Nonflammable

LEL: None reported

UEL: None reported

Extinguishing Media: Use extinguishing media suitable for surrounding fire. Unusual Fire or Explosion Hazards: Cylinder may explode in heat of fire.

Special Fire-fighting Procedures: Because fire may produce toxic thermal decomposition products, wear a self-contained breathing apparatus (SCBA) with a full facepiece operated in pressure-demand or positive-pressure mode. Structural firefighter's protective clothing provides only limited protection. If possible without risk, move container from fire area. Apply cooling water to sides of containers until well after fire is out. Stay away from ends of tanks. Withdraw immediately if you hear a rising sound from venting safety device or notice any tank discoloration due to fire. Do not release runoff from fire control methods to sewers or waterways.

Section 5. Reactivity Data

Salamin : Stability/Polymerization: Dichlorodifluoromethane is stable until 1022 °F (550 °C) above which it begins to decompose. Hazardous polymerization cannot occur.

Chemical Incompatibilities: Chemically active metals such as sodium, potassium, calcium and powdered aluminum, zinc, or magnesium. Conditions to Avoid: Exposure to excessive heat and contact with incompatibles.

Hazardous Products of Decomposition: Thermal oxidative decomposition of dichlorodifluoromethane can produce earbon dioxide (CO2), and toxic phosgene, chlorine, hydrogen chloride, hydrogen fluoride, and fluorine gases.

Section 6. Health Hazard Data

Carcinogenicity: The IARC, (164) NTP, (169) and OSHA(164) do not list dichlorodifluoromethane as a carcinogen.

Summary of Risks: Dichloredifluoromethane is relatively low in toxicity but is a simple asphyxiant when oxygen levels decline to 15 to 16%, and produces unconsciousness at 6 to 8% oxygen. It also produces weak narcotic-like effects (when 10% mixture is inhaled) and eye irritation at high concentrations. Bursts of compressed gas or liquid can cause frostbite.

Medical Conditions Aggravated by Long-Term Exposure: Pre-existing ventricular ectopy.

Target Organs: Cardiovascular system and the central and peripheral nervous systems.

Continue on next page

Material Safety Data Sheet

From Genium's Reference Collection Genium Publishing Corporation 1145 Catalyn Street Schenectady, NY 12303-1836 USA (518) 377-8855



No. 359 ETHYLENE DICHLORIDE (Formerly 1,2-Dichloroethane) (Revision C)

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Issued: November 1978 Revised: August 1987

SECTION MATERIA MIDENTIFICATION

CHEMICAL NAME: ETHYLENE DICHLORIDE (Changed to reflect common industrial practice)

DESCRIPTION (Origin/Uses): Made from acetylene and HCl. Used as a degreaser, a scavenger in leaded gasoline, as an intermediate in the manufacture of vinyl chloride, in paint removers, in wetting and penetration agents, in ore flotation processes, as a fumigant, and as a solvent for fats, oils, waxes, and gums.

OTHER_DESIGNATIONS: 1,2-Dichloroethane; sym-Dichloroethane; Dutch Liquid; Dutch Oil; EDC;

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thane Dichloride; Ethylene Chloride; 1,2-Ethylene Dichloride; Glycol Dichloride; C2H4Cl2;	HMIS
NOSH RTECS KI0525000; CAS #0107-06-2	H 1
MANUFACTURERS/SUPPLIERS: Available from several suppliers, including:	F 3
low Chemical USA, 2020 Dow Center, Midland, MI 48640; Telephone: (517) 636-1000	R 0
	PPE* .
OMMENTS: Ethylene dichloride is a flammable, toxic liquid.	* See Sect. 8
PECTION OF INCREDIENTS AND THE AND THE	Property and the second second

SECTION 2. INGREDIENTS AND HAZARDS	%	HAZARD DATA
Ethylene Dichloride, CAS #0107-06-2; NIOSH RTECS #KI0525000 H H Cl - C - C - Cl	100	ACGIH Values 1987-88 TLV-TWA: 10 ppm, 40 mg/m ³ OSHA PEL* 1986-87 8-Hr TWA: 50 ppm; Ceiling: 100 ppm (15 Min.) NIOSH REL 1986-87 10-Hr TWA: 1 ppm Ceiling: 2 ppm (15 Min.) Toxicity Data
*The maximum allowable peak concentration (above the ceiling level value) of ethylene dichloride is 200 ppm for 5 minutes in any 3-hour period. COMMENTS: Additional data concerning toxic doses and tumorigenic, reproductive, and mutagenic effects is listed (with references) in the NIOSH RTECS 1983-84 supplement, pages 865-66.		Man, Inhalation, TC _{LO} : 4000 ppm/1 Hr Human, Oral, TD _{LO} : 428 mg/kg Man, Oral, TD _{LO} : 892 mg/kg Man, Oral, LD _{LO} : 714 mg/kg Rat, Oral, LD _{SO} : 670 mg/kg

SECTION 3. PHYSICAL DATA

Boiling Point ... 182.3'F (83.5'C)

Vapor Pressure ... 87 Torr at 77°F (25°C)

Water Solubility ... Soluble in about 120 Parts Water

Vapor Density (Air = 1) ... 3.4

Evaporation Rate (n-BuAc = 1) ... Not Listed Specific Gravity ... 1.2569 at 69°F (20°C) Freezing Point ... -31.9°F (-35.5°C) Molecular Weight ... 98.96 Grams/Mole

Appearance and odor: Colorless, clear liquid. Sweet, chloroformlike odor is typical of chlorinated hydrocarbons. The recognition threshold (100% of test panel) for ethylene dichloride is 40 ppm. Odor detection probably indicates an excessive exposure to vapor. High volatility and flammability, coupled with its toxicity and carcinogenic potential, make this material a major health hazard.

COMMENTS: Ethylene dichloride is miscible with alcohol, chloroform, and ether.

SECTION 4. FIRE A	ND EXPLOSION DATA		LOWER	UPPER
Flash Point and Method	Autoignition Temperature	Flammability Limits in Air		
See Below	775°F (413°C)	% by Volume	6.2	15.9

EXTINGUISHING MEDIA: Use chemical, carbon dioxide, alcohol foam, water spray/fog, or dry sand to fight fires involving ethylene dichloride. Direct water sprays may be ineffective extinguishing agents, but they may be successfully used to cool fire-exposed containers. Use a smothering effect to extinguish fires involving this material. UNUSUAL FIRE/EXPLOSION HAZARDS: Ethylene dichloride is a dangerous fire and explosion hazard when exposed to sources of ignition such as heat, open flames, sparks, etc. Its vapors are heavier than air and can flow along surfaces to distant, low-lying sources of ignition and flash back. If it is safe to do so, remove this material from the fire area. Ethylene dichloride burns with a smoky flame. SPECIAL FIRE-FIGHTING PROCEDURES: Wear a self-contained breathing apparatus with a full facepiece operated in a pressure-demand or another positive-pressure mode.

COMMENTS: Flash Point and Method: 56°F (13°C) CC; 65°F (18°C) OC.

OSHA Flammability Class (29 CFR 1910.106): IB. DOT Flammability Class (49 CFR 173.115): Flammable Liquid

SECTION 5. REACTIVITY DATA

Ethylene dichloride is stable. Hazardous polymerization cannot occur.

CHEMICAL INCOMPATIBILITIES include strong oxidizing agents. Explosions have occurred with mixtures of this material and liquid ammonia or dimethylaminopropylamine. Finely divided aluminum or magnesium metal may be hazardous in contact with ethylene dichloride.

CONDITIONS TO AVOID: Eliminate sources of ignition such as excessive heat, open flames, or electrical sparks, particularly in low-lying areas, because the explosive, heavier-than-air vapors will concentrate there.

PRODUCTS OF HAZARDOUS DECOMPOSITION can include vinyl chloride, chloride fumes, and phosgene. Phosgene is an extremely poisonous gas. Products of thermal-oxidative degradation (i.e., fire conditions) must be treated with appropriate



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Material Safety Data Sheets Collection:

Sheet No. 703 1,2-Dichloroethylene

Issued: 4/90

Section L. Material Identification

1,2-Dichloroethylene Description: An industrial solvent composed of 60% cis- and 40% trans-isomers. Both isomers, cis and trans, are made by partial chlorination of acetylene. Used as a general solvent for organic materials, lacquers, dye extraction, thermoplastics, organic synthesis, and perfumes. The trans-isomer is more widely used in industry than either the cis-isomer or the mixture. Toxicity also varies between the two isomers.

Other Designations: CAS No. 0540-59-0; C,H,Cl,; acetylene dichloride; cis-1,2-dichloroethylene; sym-dichloroethylene; trans-1,2-dichloroethylene, dioform.

Manufacturer: Contact your supplier or distributor. Consult the latest Chemicalweek Buyers' Guide(77) for a suppliers list.



Section 2. Ingredients and Occupational Exposure Limits

1,2-Dichloroethylene, ca 100%

OSHA PEL

8-hr TWA: 790 mg/m3, 200 ppm

ACGIH TLV, 1989-90

TLV-TWA: 790 mg/m3, 200 ppm

NIOSH REL, 1987 790 mg/m³, 200 ppm Toxicity Data*

Rat, oral, LD,: 770 mg/kg; toxic effects not yet reviewed Frog, inhalation, TC,: 117 mg/m³ inhaled for 1 hr affects the peripheral nerve and sensation (flaccid paralysis without anesthesia); behavior (excitement); lungs, thorax, or respiration (respiratory depression)

* See NIOSH, RTECS (KV9360000), for additional toxicity data.

Section 3. Physical Data

Boiling Point: 119 °F/48 °C Melting Point: -56 to -115 °F/-49 to -82 °C/

Vapor Pressure: 180 to 264 torr at 68 °F/20 °C

Vapor Density (Air = 1): 3.4

Molecular Weight: 96.95 g/mol Specific Gravity (H,O = 1 at 39 °F/4 °C): 1.27 at 77 °F/25 °C

Water Solubility: Insoluble

Appearance and Odor: A colorless, low-boiling liquid with a pleasant odor.

Section 4. Fire and Explosion Data

Flash Point: 37 F/2.8 °C, CC

Autoignition Temperature: 860 °F/460 °C

UEL: 12.8% v/v

Extinguishing Media: Use dry chemical, CO,, halon, water spray, or standard foam. Water may be ineffective unless used to blanket the fire. Unusual Fire or Explosion Hazards: This material's vapors are a dangerous fire hazard and moderate explosion hazard when exposed to any heat or ignition source or oxidizer.

Special Fire-fighting Procedures: Since fire may produce toxic fumes, wear a self-contained breathing apparatus (SCBA) with a full facepia operated in the pressure-demand or positive-pressure mode and a fully encapsulating suit. Vapors may travel to heat or ignition sources and flash back. Stay up wind and out of low areas. Be aware of runoff from fire control methods. Do not release to sewers or waterways.

Section 5. Reactivity Data

Stability/Polymerization: This material is stable at room temperature in closed containers under normal storage and handling conditions. Hazardous polymerization cannot occur.

Chemical Incompatibilities: This material is incompatible with alkalies, nitrogen tetraoxide, difluoromethylene, strong oxidizers, and dihyposluorite. When in contact with copper or copper alloys or by reaction with potassium hydroxide, explosive chloroacetylene may be released. Conditions to Avoid: Addition of hot liquid to cold 1,2-dichloroethylene may cause sudden emission of vapor that could flash back to an ignition

Hazardous Products of Decomposition: Thermal oxidative decomposition of 1,2-dichloroethylene can produce highly toxic fumes of chlori-(CI-).



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Material Safety Data Sheets Collection:

Sheet No. 514 p-Dichlorobenzene

Issued: 10/83

Revision: A, 11/90

Section 1. Material Identification

p-Dichlorobenzene (C,H,CL) Description: Derived by chlorinating monochlorobenzene. Used as a general insecticide; a moth repellent; a germicide; a chemical intermediate in the production of polyphenylene sulfide; a plastic used in the electrical and electronics industries; a space deodorant in products such as room deodorizers, urinal and toilet bowl blocks. and diaper pail deodorizers; and in producing 1,2,4 trichlorobenzene.

Other Designations: CAS No. 0106-46-7, 1,4-dichlorobenzene, dichlorocide, Evola, NCI-c 54955, Paracide, Paracrystals, Paradi, paradichlorobenzol, Paramoth, p-chlorophenyl chloride, PDB, Santochlor.

Manufacturer: Contact your supplier or distributor. Consult the latest Chemicalweek Buyers' Guide(71) for a suppliers list.

Cautions: p-Dichlorobenzene vapor is an eye and upper respiratory tract irritant. It is toxic to the liver. Prolonged exposure to high concentrations may cause weakness, dizziness, and weight loss. Flammable when exposed to heat, flame, or oxidizers.

Sec. 8

Section 2. Ingredients and Occupational Exposure Limits

p-Dichlorobenzene, ca 100%

1989 OSHA PELS

1990-91 ACGIH TLVs

1988 NIOSH REL

1985-86 Toxicity Data*

8-hr TWA: 75 ppm, 450 mg/m³ 15-min STEL: 110 ppm, 675 mg/m³ STEL: 110 ppm, 661 mg/m³

TWA: 75 ppm, 451 mg/m⁵

None established

Human, oral, TD, : 300 mg/kg produced sense organs and special senses (other eye effects); lungs, thorax, or respiration (other changes); and gastrointestinal

(hypermotility, diarrhea) effects

Human, eye: 80 ppm

1987 IDLH Level

1000 ppm

* See NIOSH, RTECS (CZ4550000), for additional irritative, mutative, reproductive, and toxicity data.

Section 3. Physical Data

Boiling Point: 345 'F (174 'C) at 760 mm Hg

Melting Point: 127.6 'F (53.1 'C)

Vapor Pressure: 10 mm Hg at 130.6 °F (54.8 °C)

Vapor Density (Air = 1): 5.08

Molecular Weight: 147.01

Specific Gravity: 1.248 at 131 °F (55 °C)

Water Solubility: Insoluble

Appearance and Odor: Volatile, white crystals with a distinctive mothball-like odor that becomes very strong at concentrations between 30 and 60 ppm. At concentrations of 80 to 160 ppm, vapors are painful to the eyes and nose. Odors and irritating effects are good warnings against overexposure to p-dichlorobenzene; however, individuals may develop tolerance to high concentrations.

Section 4. Fire and Explosion Data

Flash Point: 150 °F (66 °C), CC

Autolgnition Temperature: None reported

LEL: 1.7% V/V

UEL: None reported

Extinguishing Media: Use dry chemical, carbon dioxide, alcohol foam, or water spray. Use water spray to cool fire-exposed container, to disperse vapors, or to blanket a pool fire.

Unusual Fire or Explosion Hazards: Explosive and toxic mixtures may form in air when this material is heated, such as in a fire. Special Fire-fighting Procedures: Since fire may produce toxic fumes, wear a self-contained breathing apparatus (SCBA) with a full facepiece operated in the pressure-demand or positive-pressure mode and full protective clothing. Thoroughly decontaminate firefighting equipment after use. Be aware of runoff-from fire control methods. Do not release to sewers or waterways.

Section 5. Reactivity Data

Stability/Polymerization: p-Dichlorobenzene is stable at room temperature in closed containers under normal storage and handling conditions. Hazardous polymerization cannot occur.

Chemical Incompatibilities: p-Dichlorobenzene is incompatible with strong oxidizers and oxidizing agents.

Conditions to Avoid: Avoid incompatibilities and heat or ignition sources.

Hazardous Products of Decomposition: Thermal oxidative decomposition of p-dichlorobenzene includes carbon monoxide, chlorides, and chlorine.

Material Safety Data Sheet

From Genium's Reference Collection Genium Publishing Corporation 1145 Catalyn Street Schenectady, NY 12303-1836 USA (518) 377-8855



%

ca 100

No. 624

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NAPHTHALENE

Issued: November 1987

SECTION 1. MATERIAL IDENTIFICATION

Material Name: NAPHTHALENE

Description (Origin/Uses): Used as a moth repellant and in many industrial processes.

Other Designations: Naphthalin; Naphthene; Tar Camphor; C₁₀H₄; NIOSH RTECS No. QI0525000; CAS No. 0091-20-3

Manufacturer: Contact your supplier or distributor. Consult the latest edition of the Chemicalweek Buyer's Guide (Genium ref. 73) for a list of suppliers.

	/		/
H	AIS	V	
H	2		
F	2	R	1
R	0	I	4
PP	G*	S	1
*See sect. 8		V	2

SECTION 2 MINGREDIENTS AND HAZARDS

Naphthalene, CAS No. 0091-20-3

*Immediately dangerous to life and health

**See NIOSH RTECS for additional data with references to irritative, mutagenic, reproductive, and tumorigenic effects.

EXPOSURE LIMITS
IDLH* Level: 500 ppm

ACGIH TLVs, 1987-88
TLV-TWA: 10 ppm, 50 mg/m³
OSHA PEL
8-Hr TWA: 10 ppm, 50 mg/m³
Toxicity Data**
Child, Oral, LD_L: 100 mg/kg
Man, Unknown, LD_L: 74 mg/kg
Rat, Oral, LD_m: 1250 mg/kg

SECTION 3. PHYSICAL DATA

Boiling Point: 424 F (218 °C) Vapor Density (Air = 1): 4.4

Vapor Pressure: 0.087 Torr at 77°F (25°C)

Water Solubility: Insoluble

Specific Gravity (H,0 = 1): 1.162 at 68 F (20 °C)

Company of the Company of the Company

Melting Point: 176°F (80°C)

Molecular Weight: 128 Grams/Mole % Volatile by Volume: ca 100

Appearance and Odor: White crystalline flakes; strong coal tar odor.

SECTION 4. FIRE	AND EXPLOSION DA	TA	LOWER	UPPER
Flash Point and Method	Autoignition Temperature	Flammability Limits in Air	1 A 1000 Ta	F PO DE ALIXA
174°F (79°C) OC; 190°F (88°C) CC	979°F (526°C)	% by Volume	0.9	5.9

Extinguishing Media: Use water spray, dry chemical, or carbon dioxide to fight fires involving naphthalene. Caution: Foam or direct water spray applied to molten naphthalene may cause extensive foaming.

Unusual Fire or Explosion Hazards: Naphthalene is a volatile solid that gives off flammable vapor when heated (as in fire situations). This vapor is much denser than air and will collect in enclosed or low-lying areas like sumps. In these areas an explosive air-vapor mixture may form, and extra caution is required to prevent any ignition sources from starting an explosion or fire.

Special Fire-fighting Procedures: Wear a self-contained breathing apparatus (SCBA) with a full facepiece operated in the pressure-demand or positive-pressure mode.

SECTION 5. REACTIVITY DATA

Naphthalene is stable in closed containers at room temperature under normal storage and handling conditions. It does not undergo hazardous polymerization.

Chemical Incompatibilities: Naphthalene is incompatible with strong oxidizing agents, chromic anhydride, and mixtures of aluminum trichloride and benzoyl chloride.

Conditions to Avoid: Ignition sources like open flame, unprotected heaters, excessive heat, lighted tohacco products, and electric sparks must not occur in work areas where naphthalene vapor may become concentrated.

Hazardous Products of Decomposition: Toxic gases like carbon monoxide are produced during fire conditions. Irritating, flammable vapor forms below the melting point because even solid naphthalene has a significant vapor pressure.



Genium Publishing Corp.

One Genium Plaza Schenectady, NY 12304-4690 (518) 377-8854

Material Safety Data Sheet Collection

Phenanthrene

MSDS No. 905

(....).

Date of Preparation: 6/94

Section 1 - Chemical Product and Company Identification

Product/Chemical Name: Phenanthrene

Chemical Formula: (C6H4CH)2

CAS No.: 85-01-8 Synonyms: Phenantrin

Derivation: A polynuclear aromatic hydrocarbon found as a component of coal tar pitch volatiles (products of bituminous coal distillation). Produced from toluene, bibenzil, 9-methyl fluorene or stilbene by passage through red hot tubes or by diene

synthesis of 1-vinyl naphthalene and maleic anhydride.

General Use: Used in the manufacture of dyestuffs and explosives; in biological research or drug synthesis.

Vendors: Consult the latest Chemical Week Buyers' Guide. (73)

Section 2 - Composition / Information on Ingredients

Phenanthrene, ca 100 % wt

OSHA PEL*

8-hr TWA: 0.2 mg/m³

ACGIH TLV* TWA: 0.2 mg/m³ NIOSH REL*

10-hr TWA: 0.1 mg/m³, cyclohexane extractable fraction

DFG (Germany) MAK

None established

*Coal tar pitch volatiles (benzene soluble)

Section 3 - Hazards Identification

ជាជាជាជា Emergency Overview ជាជាជាជាជា

Phenanthrene exists as shiny crystals with a faint, aromatic odor. It can cause photosensitization of the skin. Phenanthrene is combustible and reacts dangerously with oxidizers.

Potential Health Effects

Primary Entry Routes: Skin contact.

Target Organs: Skin.

Acute Effects

Inhalation: Effects not reported.

Eye: Effects not reported.

Skin: Can cause photosensitization of the skin.

Ingestion: Effects not reported.

Carcinogenicity: Although it has produced skin cancer in experimental animals, the results were not statistically significant and IARC has assigned phenanthrene a Class 3 (unclassifiable as to carcinogenicity) designation. The NTP and OSHA do not list phenanthrene as a carcinogen.

Medical Conditions Aggravated by Long-Term Exposure: Skin disorders.

Chronic Effects: None reported.

Section 4 - First Aid Measures

Inhalation: Remove exposed person to fresh air and support breathing as needed.

Eye Contact: Do not allow victim to rub or keep eyes tightly shut. Gently lift eyelids and flush immediately and continuously with flooding amounts of water until transported to an emergency medical facility. Consult a physician immediately.

Skin Contact: Quickly remove contaminated clothing. Rinse exposed area with flooding amounts of water to remove loose material and then move quickly to a soap and water wash. For reddened or blistered skin, consult a physician.

Ingestion: Never give anything by mouth to an unconscious or convulsing person. Contact a poison control center. Unless the poison control center advises otherwise, have the conscious and alert person drink 1 to 2 glasses of water, then induce vomiting.

After first aid, get appropriate in-plant, paramedic, or community medical support.

Vote to Physicians: Treatment is symptomatic and supportive.

Wilson Risk Scale

R 1 I 3

S 3 K 1

HMIS H 1 F 1

PPE*

R

Section 10 - Stability and Reactivity

Stability: Phenanthrene is stable at room temperature in closed containers under normal storage and handling conditions. Polymerization: Hazardous polymerization does not occur.

Chemical Incompatibilities: Strong oxidizers.

Conditions to Avoid: Phenanthrene dust generation and exposure to heat ignition sources, or oxidizers.

Hazardous Decomposition Products: Thermal oxidative decomposition of phenanthrene can produce carbon oxide(s).

Section 11- Toxicological Information

Toxicity Data: *

Acute Oral Effects:

Mouse, oral, LD50: 700 mg/kg

Mutagenicity:

Rat, liver cell: 3 mmol/L caused DNA damage Human, lymphocyte: 100 µmol/L caused mutation

Carcinogenicity:

Mouse, skin: 71 mg/kg produced tumors at site of application.

See NIOSH, RTECS (SF7175000), for additional toxicity data.

Section 12 - Ecological Information

Ecotoxicity: Neanhes arenaceodentata, TLm = 0.6 ppm/96 hr, sea water at 71.6 °F (22 °C)

Environmental Degradation: If released to soil, some phenanthrene may biodegrade but the majority will bind to the soil without much leaching to groundwater. Volatilization is not expected to be significant. In water, it will adhere to particulates and sediment. Photolysis may occur near the surface producing toxic substances. Photolysis/photooxidation half-life = 8.4 hr. In the air, it will react with photochemically generated hydroxyl radicals (half-life = 1.67 days).

Soil Absorption/Mobility: Phenanthrene absorbs strongly to soil and sediment in water.

Section 13 - Disposal Considerations

Disposal: For treatment of phenanthrene contaminated water, the particulate bound portion can be removed by sedimentation, flocculation, and filtration. Chlorination is not recommended as it has been shown to produce mutagenic substances. The dissolved portion requires oxidation for partial removal. Contact your supplier or a licensed contractor for detailed recommendations. Follow applicable Federal, state, and local regulations.

Section 14 - Transport Information

DOT Transportation Data (49 CFR 172.101):

Packaging Authorizations

c) Bulk Packaging: 173.240

b) Non-bulk Packaging: 173.213

a) Exceptions: 173.155

Shipping Name:

Environmentally hazardous substances, solid, n.o.s.*

Shipping Symbols: ---

Hazard Class: 9 ID No.: UN3077 Packing Group: III

B54, N50

Label: Class 9 Special Provisions (172.102): 8,

Quantity Limitations

a) Passenger, Aircraft, or Railcar: None

b) Cargo Aircraft Only: None

Vessel Stowage Requirements

a) Vessel Stowage: A

b) Other: ---

* Classified as a hazardous substance when phenanthrene is in a quantity, in one package, which equals or exceeds the RQ of 5000 lb (2270 kg)

Section 15 - Regulatory Information

EPA Regulations:

RCRA Hazardous Waste Number: Not listed

Listed as a CERCLA Hazardous Substance (40 CFR 302.4) per CWA, Sec. 307(a)

CERCLA Reportable Quantity (RQ), 5000 lb (2270 kg)

SARA 311/312 Codes: 1

SARA Toxic Chemical (40 CFR 372.65): Not listed

SARA EHS (Extremely Hazardous Substance) (40 CFR 355): Not listed

OSHA Regulations:

Listed (coal tar pitch volatiles) as an Air Contaminant (29 CFR 1910.1000, Table Z-1, Z-1-A)

Section 16 - Other Information

References: 23, 73, 103, 124, 132, 133, 136, 139, 159, 164, 187, 190

Prepared By...... M Gannon, BA Industrial Hygiene ReviewPA Roy, MPH, CIH



1145 Catalyn Street Schenectady, NY 12303-1836 USA (518) 377-8854

Material Safety Data Sheets Collection:

Sheet No. 711 Pyrene

Issued: 4/90

Section 1. Material Identification

yrene Description: A condensed ring, polyaromatic hydrocarbon compound derived from coal tar. Also synthesized rom 0,0'-ditolyl. Used in biochemical research and as starting material for synthesizing benzo(a)pyrene. An ingredient of moked and broiled meat, tobacco smoke, and air pollution.

Other Designations: CAS No. 0129-00-0; C₁₆H₁₀; beta-pyrene; benzo(d,e,f)phenanthrene; benzo(d,e,f)phenathrene.

Manufacturer: Contact your supplier or distributor. Consult the latest Chemicalweek Buyers' Guide⁽⁷³⁾ for a suppliers list.

R 1 I 3 S 2* K - * Skin absorption

> R 0 PPG† † Sec. 8

Section 2. Ingredients and Occupational Exposure Limits

Pyrene, ca 100%

)SHA PEL 8-hr TWA: 0.2 mg/m³ ACGIH TLV, 1989-90

None established None established

NIOSH REL, 1987

Toxicity Data*

Rat, oral, LD_{so}: 2700 mg/kg ingested produces conjunctiva irritation, excitement, and muscle contraction

Rat, inhalation, LC₃₀: 170 mg/m³ inhaled produces conjunctiva

irritation, excitement, and muscle contraction

Gene mutation in mammalian cells; human cell types: 12 µmol/l

* See NIOSH, RTECS (UR2450000), for additional mutative, tumorigenic, and toxicity data.

Section 3. Physical Data

Jolling Point: 759 F/404 °C

Melting Point: 313 °F/156 °C

Vapor Pressure: 6.85 x 107 torr at 68 F/20 °C

Molecular Weight: 202.26 g/mol

Specific Gravity (H,O = 1 at 39 *F/4 *C): 1.271 at 73 *F/23 *C

Water Solubility: Insoluble (0.135 mg/l)

Appearance and Odor: Coloriess solid or a slight blue florescent solution. Tetracene impurities give pyrene a yellow color.

Section 4. Fire and Explosion Data

lash Point: None reported

Autoignition Temperature: None reported

LEL: None reported .

UEL: None reported

atinguishing Media: Use foam, dry chemical, and CO, to extinguish fire.

Unusual Fire or Explosion Hazards: Pyrene is a flammable and combustible material that heat and ignition sources may ignite. It burns rapidly with a flare-like effect.

pecial Fire-fighting Procedures: Since fire may produce toxic fumes, wear a self-contained breathing apparatus (SCBA) with a full facepiece perated in the pressure-demand or positive-pressure mode. Avoid skin contact. Be aware of runoff from fire control methods. Do not release to were or waterways.

ection 5. Reactivity Data

stability/Polymerization: Pyrene is stable at room temperature in closed containers under normal storage and handling conditions. Hazardous polymerization cannot occur.

lazardous Products of Decomposition: Thermal oxidative decomposition of pyrene can emit irritating fumes and acrid smoke.



Citizen Participation Plan for the Site Investigation/Remedial Alternatives Report Workplan Former Provan Ford Facility Environmental Protection Project B-00127-3

Prepared for City of Newburgh 83 Broadway Newburgh, NY 12550

July 2000

Project No. PROVA001

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INTRODUCTION

Citizen participation is an important and integral component of remedial programs in New York State. Integration of individuals and organizations into the remedial program helps ensure outcomes that account for both technical and human concerns for protecting public health and the environment. In order for this integration to be accomplished, a project-specific plan is needed to inform and involve community residents, public and private leaders, and other stakeholders. This Citizen Participation Plan (CPP) addresses this need by documenting the planned project-specific public outreach activities and resources organized for the remedial program associated with the Former Provan Ford Site in Newburgh, NY (Provan Site).

The primary purpose of this CPP is to outline a variety of communication methods that, based on applicable New York State law and New York State Department of Environmental Conservation (NYSDEC) guidance, provide for constructive communication of program activities among the stakeholders and other interested parties. This CPP includes methods intended to inform interested parties of program developments, elicit responses and public involvement and provide a central point of contact for inquiries regarding the remedial program for the Provan Site. Given this context, this CPP presents the planned communication and outreach activities, describes how interested individuals and groups can participate in the remedial program, and provides a variety of reference materials to facilitate gaining access to project-specific information and management personnel.

BASIC SITE INFORMATION

Site Description/Location/Brief History

The City of Newburgh took ownership of the Provan Ford Site in 1998 after it foreclosed on this 3.5 acre property. The City has leased the site for the past two years but has requested that the current occupant leave the site in the first quarter of the year 2000.

The Provan Ford Site is located at 146-172 Mill Street Avenue in Newburgh, NY. The approximate boundaries of the site include Route 9-W (Robinson Street) on the West, the Orange County Plumbing Supply Company on the South, Dickson Street on the North and Mill Street on the East. This is a mixed use area where the primary uses are commercial but some residential properties are located across Mill Street to the East.

Prior to 1998, Provan Ford had occupied the property for approximately fifty years. Prior to the construction of Provan Ford, the site was undeveloped. Provan's on-site operations included the washing, service and maintenance of Ford tanker trucks. This was a large truck dealership/repair facility and a petroleum/industrial tanker cleaning facility. It consisted of two attached garage maintenance areas, a small office area and an external wash rack which was located on a concrete pad and attached to the western wall of the garage. Empty tanker trucks would arrive at the site, where they would be cleaned out in the wash rack, then serviced and fueled before they were sent back on the road for petroleum distribution.

During Provan Ford's ownership of this site, the City of Newburgh's Building, Public Works and Fire Department's had on several occasions responded to the facility for various incidents involving discharges of petroleum/other substances, to the City's municipal sewer system from various portions of the site. The on-site drainage system is connected to the City of Newburgh's combined sewer system which ultimately discharges to the Hudson River. The neighborhood surrounding this property is on the City's Water and Sewer service.

An existing oil/water separator is located on the southeast portion of the site. The existing oil/water separator has not been properly maintained since it's installation and is currently full of sediments and petroleum products. The facility has in the past violated the City of Newburgh's Pretreatment Ordinance for oil and grease discharge of potentially flammable products to the City of Newburgh's sewer collection system. Several years ago several large explosions and fires were reputedly attributed to discharges from the project site. A structure immediately adjacent to the facility was destroyed by fire after a series of explosions within the City's sewer system. The New York State Department of Environmental Conservation Police have also responded to the facility on several occasions in the past. The combination of surface discharge of petroleum products along with the discharge to the City's storm sewer collection system impacted the property, surrounding areas and the receiving waters for many years.

Status of Current Site Investigations

In October of 1999 First Environment, Inc., an environmental engineering firm, was hired by the City of Newburgh to investigate the environmental problems at the site and completed an initial investigation in October 1999. This site investigation was continued in order to evaluate potential areas of environmental concern identified in the initial investigation. A second round of soil and groundwater sampling identified four areas of environmental concern: (1) the areas where the six underground storage tanks are located; (2) the area of the exterior oil/water separator and associated pipelines, (3) the truck wash rack area/drum storage and adjacent drains (4) and several areas of potential surface soil contamination. These areas of concerns are discussed more fully in the next section of this report as well as in the Site Investigation/Remedial Alternatives Report Workplan. A site location map (Figure 1) for Provan Ford Site as well as a site map which locates the areas of environmental concern (Figure 2) are included in this CPP. See Figures following page 16.

Petroleum contamination was identified in the second round of sampling. Additional sampling will be performed over the next twelve months as part of the Site Investigation

to further define the extent of both soil and groundwater contamination so that the remedial action plan developed will specifically address the concentration and movements of contamination identified for the site.

These site investigation activities will be completed as part of the Remedial Alternatives Workplan which this CPP is part of. Full reports on all data collected will be discussed in associated public meetings and documents.

The City of Newburgh is conducting these site investigations and remedial activities under the Municipal Assistance Environmental Restoration Projects or "Brownfield Program", included in the 1996 Clean Water/Clean Air Bond Act.

REMEDIAL PROGRAM DESCRIPTION

New York State Remedial Program Overview

The Municipal Assistance Environmental Restoration Projects "Brownfield Program" follows the typical process for investigating and remediating hazardous waste sites in New York State. This process begins when the Phase I and Phase II Preliminary Site Assessments (PSAs) are performed. The Phase I PSA generally involves a review of historical records and information that may serve to document any past uses of a site. The follow up Phase II PSA typically goes a step further by including some form of site sampling and investigation to characterize the present condition of a site, including an early determination of any potential hazard posed by materials or conditions encountered. The former Provan Ford Site Project is at this stage in the remedial program process. A Site Investigation/Remedial Alternatives Report (SI/RAR) will be completed based on the findings and the results of the Site Investigation/Remedial Alternatives Report Workplan of which this CPP is a part.

Description of the Completed and Proposed Project Investigations

An initial Site Investigation is complete and the results of this investigation are briefly outlined below:

- 27 soil borings were installed adjacent to the areas of environmental concern i.e., the six underground tank sites, the site of exterior oil/water separator and associated pipelines, and the site of the truck wash rack and associated drains. 16 soil samples were sent to an analytical laboratory and 12 of these samples showed petroleum contamination that exceeds NYSDEC allowable limits for volatile organic compounds and/or semi-volatile organic compounds.
- 5 soil surface samples were collected from 0-6 inches below grade at the same areas of concern and where soil staining was visible. Each of these samples exceeded NYSDEC samples for semi-volatile organic compounds and/or volatile organic compounds.
- 7 temporary monitoring wells were installed: Again, these were located adjacent to the six underground storage tanks and the exterior oil/water separator. The water table was encountered from 8 feet to 14 feet below grade and petroleum was detected on the water table at five of the seven wells. Laboratory analysis of samples from six of the seven wells show that the groundwater has been contaminated.
- Prior to the initial investigation activities at this site, NYSDEC had
 requested that 60 tons of soil be stockpiled for disposal. As part of the
 initial investigation, this stockpiled soil was sampled for waste
 characterization. The sampling results showed the stockpiled soil was
 not hazardous and it was properly disposed of by TPS Technologies on
 February 8, 2000.

On February 24, 2000, 9 drums were overpacked by the firm Code
 Environmental prior to pending off-site disposal. These nine drums are
 part of the drums stored in a fenced area on-site. The nine were dented
 and rusted and had been identified as being a potential threat to the
 environment.

Additional investigation work as well as remediation work is planned for the next 12 months. The objectives of this work are to thoroughly investigate and clearly define the extent, concentrations and movement of the contamination in the groundwater and soil at this site and the impacts of this contamination on the environment. Additional objectives of this site investigation include the definition of the extent to which contamination may have migrated or can be expected to migrate in the future both on and off-site and the identification of population and environmental receptors at risk. At the same time, remediation activities which are clearly required to reduce the already identified contamination will be carried out.

These proposed site investigation and remediation activities are briefly outlined below:

- The six underground storage tanks will be cleaned out and removed. Additional
 contaminated soil will be sampled, classified and removed. Additional soil
 samples will be collected and sent for analysis.
- Eight additional monitoring wells will be installed downgradient of the removed underground storage tanks. All 16 wells (9 new: 7 old) will be sampled for chemical constituents as required by NYSDEC.
- The initial Investigation had identified five areas on the property where surface soil may be contaminated. Additional sampling for poly-nuclear aromatic hydrocarbons (PAHs), metals and VOC is recommended for all five areas.

- 81 drums containing miscellaneous petroleum products (56 outside: 25 inside) are stored on this site. These drums will be properly disposed of off-site and prior to disposal each drum will be characterized by a field chemist.
- A map will be developed that illustrates historic and current on-site pipelines and drains.
- Both existing oil/water separators and drains will pumped out, cleaned, and petroleum products disposed of. All building drains inside the building will be sealed.

A Site Investigation/Remedial Alternatives Report will be prepared that will document the additional findings from this investigation and will also propose additional remediation activities. These results and recommendations will be discussed and presented at a public meeting. The Report itself will be available in the document repository.

The key decision points (milestones) and Project Schedule for the Provan Ford Site Investigation/Remedial Alternatives Report Workplan is available. See Provan Project Schedule following page 16.

CITIZEN PARTICIPATION ACTIVITIES

This section presents the specific citizen participation and outreach activities planned for implementation during the remedial program. Operating under overarching project-specific citizen participation goals, clearly defined objectives will be achieved by implementing a range of communication tools and methods. The planned activities are geared toward making project-specific information (e.g., work plans, technical reports, data, information sheet summaries) available to the public; facilitating communication among stakeholders (e.g., contact list, public meetings, comment periods); and notifying the public of document availability, public meetings, comment periods, and major program milestones.

Building locations where project-related documents will be available are provided in Appendix B. The community contact list is provided in Appendix C, while the Provan Ford Project management contacts (NYSDEC, New York State Department of Health [NYSDOH], and City of Newburgh representatives and Project officials are provided in Appendix D.

Goals and Objectives

The central goal of this CPP is to achieve effective, open communication between the City of Newburgh (owner of the former Provan Ford site), the citizens of Newburgh, the NYSDEC, and other stakeholders and interested parties. Corollary goals include:

- Communicate program goals and major milestones, actions, and outcomes;
- Inform citizens and others of on-going project activities, status, and progress,
- Provide citizens (and all stakeholders) a forum for input and comment;
- Engender a public understanding of the constituents of interest (chemicals observed in the project area), their potential risks to human health and the environment, and appropriate responses to mitigate those risks; and
- Inform the community about the duration and complexity of the remedial program.

In order to accomplish these goals, the following specific objectives will be pursued through the implementation of this CPP:

- Consistently communicate goals, accomplishments, and status of the project to the contact list (including community leaders, public officials, and the wider community, as necessary) through appropriate means;
- Maintain, update, and utilize the contact list,
- Educate the community, in lay terms, about the nature and magnitude of potential project area risks, including instructions for mitigating risk (if

- appropriate) and assurances that the environment and worker/public health and safety are protected;
- Provide interested parties the opportunity to review and comment on technical reports generated through the remedial program (e.g., public comment periods and document repository as required);
- Provide interested parties the opportunity to present opinions and ideas during the remedial program (e.g., conduct a public meeting/comment period and availability session as required);
- Provide response to public review and comment (e.g., prepare a responsiveness summary as required);
- Provide the news media with interviews or press releases, as available, to ensure accurate coverage of remedial program activities; and
- Periodically review the effectiveness of the citizen participation and outreach activities during the remedial program and make adjustments in this CPP's methods and/or activities, if necessary.

Tools and Methods

There are many ways to reach and communicate with the community of Newburgh as this CPP is implemented over the course of the remedial program. A variety of outreach tools and methods will be used because "The Community" is not a single entity, but comprised of various organizations, public an business leaders, and a diverse assemblage of individuals of all ages, education backgrounds, and cultures.

Interested parties will be informed and invited to participate in the planned citizen participation activities through appropriate means such as mailings to the contact list, press releases, or information sheets and other documents made available in the document repository.

The following specific public participation activities will be implemented to facilitate public involvement in the Provan Ford Project. Depending upon the nature of work

ongoing at the site, the NYSDEC and City of Newburgh may decide to use any or all of the following citizen participation tools and methods.

Contact List

The contact fist includes the names, addresses, and/or telephone numbers of individuals, groups, organizations, and/or media interested in or affected by the Provan Ford Project The contact list is periodically updated with the contact information of any individual, group, organization, or business that has an interest in the project and wishes to receive public mailings or announcements published regarding project activities. Names are added to the contact list as they are received from public meeting sign-in sheets or from any public inquiry requesting to be included on the list. When specific citizen participation activities are scheduled, the contact list is used to notify interested parties as to how and where they can participate in project decision-making. For example, when the NYSDEC prepares its PRAP, the contact list will be used to inform interested parties of the availability of the PRAP for public review and comment, and of the date, time, and location of the PRAP public meeting and public comment The first activity proposed for the Provan Project will be to mail an availability of the Site Investigation/Remedial announcement concerning the Alternatives Report Workplan to the contact list.

The contact list is provided as Appendix C of this CPP.

Document Repository

Document repositories will be established in publicly accessible locations that have the accommodations necessary to house and make project related documents available for community reference and review. The use of document repositories is intended to maximize public access to site information while minimizing abuse, destruction, of project documentation. The document repositories for the Provan Ford Project are:

NYSDEC Region 3 Office

21 South Putt Comers Road New Paltz, NY 12561-1696 Phone: (845) 256-3154 Hours: 9:00 a.m. to 4:30 p.m.

Newburgh Free Library

Head of Reference Department 124 Grand Street Newburgh, NY 12550 Phone: (845) 561-1985

Hours: 9:00 a.m. to 9:00 p.m. Monday through Thursday 9:00 am. to 5:00 p.m. Friday and

Saturday

1:00 p.m. to 5:00 p.m. Sunday

The following documents, as **available**, will be placed in the repositories and made available for public reference and use:

- Site Investigation/Remedial Alternatives Workplan;
- Quality Assurance/Quality Control Plan;
- Health and Safety Plan (HASP);
- Citizen Participation Plan (CPP);
- Site Investigation/Remedial Alternatives Report (SI/RAR);
- Proposed Remedial Action Plan (PRAP);
- Record of Decision and Responsiveness Summary; Remedial Design Documents (including design Plans and Specifications), and
- Other Materials (e.g., Information Sheets, Notices, etc.).

Public Meeting and Comment Period

After completion of the SI/RAR, the preferred remedy for the site will be presented in a Proposed Remedial Action Plan (PRAP) and be subject to public review and comment.

A public meeting and comment period will be held at a convenient time and location to gather public comments on the PRAP. A 45-day comment period will also be established to gather additional comments and questions from interested parties. During the public meeting, NYSDEC and City of Newburgh project managers will be on hand to summarize the findings of the SI/RAR and answer questions regarding project area characterization, data, activities, remedial alternatives, the preferred remedy presented in the PRAP, costs, implementation schedules, and criteria used in evaluating the preferred remedy. Note that representatives from the NYSDOH, City of Newburgh, or other public entities also may be available to answer questions at the public meeting.

Following discussion of the project's status, citizens and others have the opportunity to ask questions and provide comment on the RI/RAR findings and the PRAP. After the public meeting and comment period end, NYSDEC will review all public comments and, where applicable, incorporate the comments into the final remedial action decisions embodied in the Record of Decision (ROD). The ROD describes the remedial action to be taken at the project area, the rationale used to select a particular remedial alternative, and a summary of public comments with corresponding NYSDEC and/or NYSDOH responses.

Responsiveness Summary

Public questions, comments, and concerns voiced during the public meeting and collected during the comment period will be addressed by the NYSDEC and published in the ROD's Responsiveness Summary. Agency responses are to address both the broad general concerns and the significant specific questions communicated by the interested parties.

Availability Sessions

After the ROD is issued, implementation of the selected remedial action is planned and formally designed. Upon completion of the design documents that specify the exact

details of the remedial action, a fact sheet is prepared and an availability session is typically held. Availability sessions are less formal settings intended to provide an opportunity for NYSDEC, City of Newburgh, citizens, and other interested parties to discuss the remedial design and its implementation. For instance, before construction begins (if some form of construction is part of the design), citizens can learn of planned health and safety measures and other site controls intended to minimize such things as noise, traffic, and potential risk to human health and the environment.

During the availability session, project officials typically do not make formal presentations. However, by having project managers directly accessible to the community, the public participants have an opportunity to speak with project personnel on a near one-to-one basis. Availability sessions can be held on an as-needed basis throughout the project.

Fact Sheets

Fact sheets will be prepared and distributed to the contact list in order to announce major project milestones and accomplishments throughout the remedial program (e.g., start of construction, major report completion, etc.). Written in lay terms, information sheets will describe and summarize the project area's history, review the goals of the project, update the status of project plans and/or results, list opportunities for citizen involvement, list key contacts, and list the location of the document repositories.

SUMMARY

Guided by the goals and objectives of this CPP, implementation of the planned public outreach and citizen participation activities will ensure the timely communication of important program information of interest to the stakeholders of the Provan Ford Project Site and the City of Newburgh. Moreover, citizen involvement and interaction in the remedial program will be facilitated through specific opportunities such as public meetings, public comment periods, availability sessions, and use of the document repositories. Throughout the remedial program, this CPP and its specific outreach tools and methods will be monitored and, as required and agreed by the NYSDEC and City of Newburgh, will be adjusted to improve its effectiveness in responding to community needs.

FIGURES

- 1 Provan Project Site Location Map (Figure 1)
- 2 Provan Project Site Map Identifying Locations of Environmental Areas of Concern (Figure 2)
- Proposed Project Schedule for the implementation of the Provan Ford Site Investigation/Remedial Alternatives Report Workplan and the production of the Site Investigation/Remedial Alternatives Report

APPENDICES

Α	Glossary of Key	Citizen	Participation	Terms and	Major Program	Elements
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- Identification of Document Repositories and List of Available Documents Identification of Interested Parties (Contact List)
 Identification of Project Management Contacts В
- С
- D

Appendix A

Hazardous Waste Site Program Glossary and Acronyms

GLOSSARY

This glossary defines terms associated with New York's hazardous waste site citizen participation program, and important elements of the hazardous waste site remedial program. Words in **bold** in the definitions are defined elsewhere in the glossary. A list of acronyms often used in the remedial program begins on page D-7.

Administrative	9
Record	

Part of a site's **Record of Decision** which lists and defines documents used in the development of NYSDEC's decision about selection of a remedial action.

Availability Session

A scheduled gathering of program staff and members of the public in a casual setting, without a formal presentation or agenda but usually focusing on a specific aspect of a site's remedial process.

Citizen Participation

A program of planning and activities to encourage communication among people affected by or interested in hazardous waste sites and the government agencies responsible for investigating and remediating them.

Citizen Participation Plan

A document which must be developed at a site's **Remedial Investigation** stage. A CP plan describes the citizen participation activities that will be conducted during a site's remedial process.

Citizen Participation Record

A document prepared at a major remedial stage which describes the citizen participation activities required at that stage. A CP Record also directs a scoping process to determine if additional citizen participation activities are appropriate and feasible.

Citizen Participation Specialist

A staff member from an NYSDEC central office or regional office who has specialized training and experience to assist a **project manager** and to plan, conduct and site-specific citizen participation program.

Classification

A process to place a hazardous waste site within a category which defines its hazardous waste status and its threat or potential threat to public health and the environment. Sites are listed along with their classifications in the Registry of Inactive Hazardous Waste Disposal Sites.

Class 1 causing or representing an imminent danger of causing irreversible or irreparable damage to public health or environment -- immediate action required.

Class 2 - significant threat to public health or environment -- action required.

Class 2a - temporary classification assigned to a site for which there is inadequate or insufficient data for inclusion in any other classification.

Class 3 - does not present a significant threat to public health or environment -- action may be deferred.

Class 4 - site properly closed -- requires continued management.

Class 5 - site properly closed -- no further action required.

Delisted - site no longer considered an inactive hazardous waste disposal site.

Comment Period

A time period for the public to review and comment about various documents and DER actions. For example, a 45-day comment period is provided when DER issues a **Proposed Remedial Action Plan** (PRAP), and when DER proposes to Delist a site from the **Registry of Inactive Hazardous Waste Disposal Sites**.

Consent Order

A legal and enforceable agreement negotiated between NYSDEC and a **responsible party**. The order sets forth agreed upon terms by which a responsible party will undertake site investigation and/or cleanup, or pay for the costs of those activities. The order includes a description of the remedial actions to be taken by the responsible party with NYSDEC oversight, and a schedule for implementation.

Contact List

Names, addresses and/or telephone numbers of individuals, groups organizations, government officials and media affected by or interested in a particular

hazardous waste site. The size of a contact list and the categories included are influenced by population density, degree of interest in a site, the stage of the remedial process and other factors. It is an important tool needed to conduct outreach activities.

Delist

Action by which DER removes a hazardous waste site from the Registry of Inactive Hazardous Waste Disposal Sites upon determination that the site contains inconsequential amounts of hazardous waste; or that a remediated site no longer requires Operation and Maintenance; or that a remediated site does not require Operation and Maintenance. A proposal to delist a site triggers a public notification and comment period process.

Division of Environmental Enforcement (DEE) A unit within the New York State Department of Environmental Conservation which works with the **Division of Environmental Remediation** and others to negotiate with **responsible parties** to achieve agreements for the investigation and remediation of hazardous waste sites. A negotiated agreement is contained in a **consent order**.

Division of Environmental Remediation

Formerly the Division of Hazardous Waste Remediation, a major program unit within the New York State Department of Environmental Conservation created to manage the hazardous waste site remedial program from site discovery through Operation and Maintenance activities. Staff include: engineers, geologists, chemists, attorneys, participation specialists. citizen environmental program specialists and support staff.

Division of Hazardous Waste Remediation

(See Division of Environmental Remediation.)

Document Repository

A file of documents pertaining to a site's remedial and citizen participation programs which is made available for public review. The file generally is maintained in a public building near the hazardous waste site to provide access at times and a location convenient to the public.

Enforcement

NYSDEC's efforts, through legal action if necessary, to compel a **responsible party** to perform or pay for

site remedial activities. NYSDEC may perform this effort by itself or in concert with other agencies.

Environmental Quality Bond Act (EQBA) The 1986 Environmental Quality Bond Act which gives New York State bonding authority of up to \$1.2 billion to fund the State's share of the total cost of remediating hazardous waste sites in New York State.

Fact Sheet

A written discussion about part or all of a site's remedial process, prepared and provided by DER to the public. A fact sheet may focus on: a particular element of the site's remedial program; opportunities for public involvement; availability of a report or other information, or announcement of a public meeting or comment period.

Interim Remedial Measure (IRM) A discrete action which can be conducted at a site relatively quickly to reduce the risk to people's health and the environment from a well-defined hazardous waste problem. An IRM can involve removing contaminated soil and drums, providing alternative water supplies or securing a site to prevent access.

National Priorities List The U.S. Environmental Protection Agency's list of the most serious uncontrolled or abandoned hazardous waste sites identified for possible long-term remedial response using money from a special trust fund.

New York State Department of Health Agency within the executive branch of New York State government which: performs health-related inspections at suspected hazardouswaste sites; conducts health assessments to determine potential risk from environmental exposure; reviews Risk Assessments prepared during the Remedial Investigation and Feasibility Study; conducts health-related community outreach around sites; and reviews remedial actions to assure that public health concernsare adequately addressed.

New York State Department of Law Agency within the executive branch of New York State government which takes the lead on hazardous waste sites requiring civil enforcement through court action. Litigation can involve negotiations and court action with **responsible** parties to clean upsites; natural resource damage claims, and recovery of remedialcosts.

New York State Registry of Inactive Hazardous Waste Disposal Sites The "Registry." A document which NYSDEC is directed by law to maintain and which lists and provides information about everyhazardous waste site in New York State which meets criteriaestablished through a definition of hazardous waste and a classification system.

Operable Unit

A discrete part of an entire site that produces a release, threat of release, or pathway of exposure. An Operable Unit can receive specific investigation, and a particular remedy may be proposed. A **Record of Decision** is prepared for each Operable Unit.

Operation and Maintenance A period in which remedial action may be conducted following construction at a site (for example, operation of a "pump and treat" system), or which is performed after a remedial action to assure its continued effectiveness and protection of people's health and the environment. Activities can include siteinspections, well monitoring and other sampling.

Preliminary Site Assessment (PSA)

A PSA is DER's first investigation of a site. A PSA is performed to determine if a site meets New York State's definition of an inactive hazardous waste disposal site by confirming the presence of hazardous waste and determining if the site poses a significant threat to public health or the environment.

Project Manager

An NYSDEC staff member within the **Division of Environmental Remediation** (usually an engineer, geologist or hydrogeologist) responsible for the day-to-day administration or remedial activities at, and ultimate disposition of, a hazardous waste site. The Project Manager works with legal, health, **citizen participation** and other staff to accomplish site-related goals and objectives.

Proposed Remedial Action Plan

An analysis by DER of each alternative considered for the remediation of a hazardous waste site and a rationale for selection of the alternative it recommends. The PRAP is created based oninformation developed during the site's **Remedial** **Investigation and Feasibility Study.** The PRAP is reviewed by the public andother state agencies.

Public Meeting

A scheduled gathering of **Division of Environmental Remediation** staff with the affected/interested public to give and receive information, ask questions and discuss concerns about a site's remedial program. Staff from other NYSDEC divisions, legal and health staff, and staff from consultants and a responsible party often also attend. A public meeting, unlike an **availability session**, generally features a formal presentation and a detailed agenda.

Reclassification

A process by which DER redefines the threat posed by a hazardous waste site to public health and the environment by developing and assessing site information and, based on findings and conclusions, assigning a new **classification** code.

Record of Decision

A document which provides definitive record of the (ROD) cleanup alternative that will be used to remediate a hazardous waste site. The ROD is based on information and analyses developed during the Remedial Investigation/Feasibility Study and public comment.

Remedial Alternatives Report (RAR)

A report that contains an evaluation of options for the remediation of any contamination in, on, or under, or emanating from, a property that includes an analysis of data and other informationconcerning the nature and extent of that property's contamination and is generally performed concurrently, and in an interactive fashion, with the site investigation.

Remedial Construction The physical development, assembly and implementation of the remedial alternative selected to remediate a site. Construction follows the **Remedial Design** stage of a site's remedial program.

Remedial Design

The process following finalization of a **Record of Decision** in which plans and specifications are developed for the **Remedial Construction** of the alternative selected to remediate a site.

Remedial Investigation/ Feasibility Study (RI/FS) RI fully defines and characterizes the type and extent of hazardous waste contamination at the site. The FS, which may be which may be conducted during or after the RI, uses information developed during the RI to develop alternative remedial actions to eliminate or reduce the threat of hazardous waste contamination to public health and the environment.

Responsible Party

An individual or business who: currently owns or operates a hazardous waste site; or historically owned or operated a site when hazardous waste was disposed; or generated hazardous waste at a site; or transported hazardous waste to a site.

Responsiveness

A written summary of major oral and written comments received by DER during a comment period about key elelments of a site's remedial program, such as a Proposed Remedial Action Plan, and DER's response to those comments.

Site Investigation

A process undertaken to determine the nature and extent of contamination in, on, and under, and emanating from a property. The SI includes the gathering of sufficient information to determine the necessity for, and the selection of the appropriate method of, remediation of contamination in, on, or under, or emanating from a property.

Site Issues and Community Profile Scoping Sheet

A document prepared to support each Citizen
Participation Record. Each Scoping Sheet identifies issues and information important to DER and the community and information that needs needs to be exchanged at a particular remedial stage. The Scoping Sheet also summarizes information about the surrounding community, including demographics, special needs, etc.

Superfund

The common name for the Federal program established by The Comprehensive Environmental Response, Compensation and Liability Act of 1980, as amended in 1986. The Superfund law authorizes the U.S. Environmental Protection Agency to investigate and cleanup sites nominated to the **National Priorities List.**

Title 3 Project Remediation of a municipally owned site through the

State Superfund Title 3 Program whereby New York State pays 75 percent of eligible costs for remediation

and the municipality pays 25 percent.

Toll-Free "800" An information line maintained by the Division of

Number Environmental Remediation to provide convenier

Environmental Remediation to provide convenient access for people who have questions, concerns or information about hazardous waste sites and their

remedial programs.

ACRONYMS

AG -- New York State Attorney General's Office

ARAR -- Applicable, Relevant and Appropriate Requirement

C&D -- Construction and Debris

CERCLA -- Comprehensive Environmental Response, Compensation

and Liability Act of 1980

CO -- Consent Order
CP -- Citizen Participation
CPP -- Citizen Participation Plan
CPS -- Citizen Participation Specialist

CQC/CQA -- Construction Quality Control/Construction Quality Assurance

DEE -- Division of Environmental Enforcement

DER -- Division of Environmental Remediation, formerly the Division

of Hazardous Waste Remediation

DHWR -- Division of Hazardous Waste Remediation, now the Division

of Environmental Remediation

DOD -- Department of Defense
DOL -- Department of Law
DOW -- Division of Water

ENB -- Environmental Notice Bulletin

EQBA -- 1986 Environmental Quality Bond Act
EPA -- Environmental Protection Agency
F&W -- Division of Fish and Wildlife

FDA -- Food and Drug Administration

FSF -- Federal Superfund

FOIL -- Freedom of Information Law

FS -- Feasibility Study
FY -- Fiscal Year

GPM -- Gallons Per Minute
HeLP -- Health Liaison Program
IRM -- Interim Remedial Measure
mg/kg -- milligrams per kilogram
NAPL -- Non-Aqueous Phase Liquid

NPL -- National Priorities List

NYSDEC -- New York State Department of Environmental Conservation

NYSDOH -- New York State Department of Health

O&M -- Operation and Maintenance

OSHA -- Occupational Safety and Health Administration

OU -- Operable Unit

PAH -- Poly-Aromatic Hydrocarbon PCB -- Poly-Chlorinated Biphenyl

PM -- Project Manager

ppm/ppb/ppt -- parts per million/parts per billion/parts per trillion

PRAP -- Proposed Remedial Action Plan
PRP -- Potentially Responsible Party
PRS -- Priority Ranking System
PSA -- Preliminary Site Assessment
QA/QC -- Quality Assurance/Quality Control

RA -- Remedial Action

RCRA -- Resource Conservation and Recovery Act

RD -- Remedial Design
RFP -- Request for Proposals

RHWRE -- Regional Hazardous Waste Remediation Engineer

RI -- Remedial Investigation

RI/FS -- Remedial Investigation/Feasibility Study

ROD -- Record of Decision
RP -- Responsible Party
SSF -- State Superfund

TAGM -- Technical and Administrative Guidance Memorandum

TCLP -- Toxicity Characteristic Leaching Procedure
TSDF -- Treatment, Storage and Disposal Facility

ug/l -- micrograms per liter
USGS -- U.S. Geological Service
VCP -- Voluntary Cleanup Program
VOC -- Volatile Organic Compound

Appendix B – Identification of Document Repositories and List of Available Documents

Two document repositories have been established to provide the public with access to important project documents and other information. This information will include reports, data and other information gathered and developed during the course of the Site Investigation/Remedial Alternatives Report as well as fact sheets, public meeting announcements, the Proposed Remedial Action Plan, and Record of Decision.

NYSDEC Region 3 Office 21 South Putt Corners Road New Paltz, New York 12561 (845) 256-3154 Hours: Mon - Fri 8:30 am - 4:45 pm

Contact: Mike Knipfing, Citizen

Participation Specialist

Newburgh Free Library
Head of Reference Department
124 Grand Street
Newburgh, NY 12550
(845) 561-1985
Hours: Mon - Thu 9:00 am - 9:00 pm
Fri, Sat 9:00 am - 5:00 pm

Sun 1:00 pm - 5:00 pm

AVAILABLE DOCUMENTS

At the start of the Site Investigation, copies of the following documents were placed in the repositories:

Site Investigation/Remedial Alternatives Report Workplan Quality Assurance/Quality Control Plan Health and Safety Plan (HASP) Citizen Participation Plan (CPP)

When additional documents are developed during the Remedial Program, they will be made available in the repositories.

Appendix C – Identification of Interested Parties (Contact List)

The following contact list has been developed to help NYSDEC to keep the community informed about and involved in the SI/RAR process for the Provan Ford Site. The list includes adjacent/nearby property owners; local regional and state officials; local media; civic business and environmental organizations and others. The contact list will be reviewed periodically and updated as appropriate.

Provan Ford Contact List

Local Officials, Committees, and Boards

Mary L. Crabb, Mayor City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

Stephen J. Rockafellow Councilperson City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

Nicholas Valentine Councilperson City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

Marlene A. Koisch Councilperson City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

Regina M. Angelo Councilperson City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550 Harold J. Porr, III City Manager City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

Nancy D'Addio City Clerk, Registrar City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

James Monell Planning Board Chair City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

William F. Ketcham Corporation Counsel City of Newburgh 83 Broadway, City Hall Newburgh, NY 12550

Head of Reference Dept. Newburgh Free Library 124 Grand Street Newburgh, NY 12550 (914) 561-1985

County/Regional Officials, Boards and Organizations

Joseph G. Rampe Orange County Executive 255-275 Main Street Goshen, NY 10924

Donna L. Benson Orange County Clerk 255-275 Main Street Goshen, NY 10924

Comissioner Peter Garrison Orange County Department of Planning 124 Main Street Goshen, NY 10924 (914) 291-2318

Commisioner Maxcy J. Smith, MD Orange County Dept. of Health 124 Main Street Goshen, NY 10924 (914) 291-2332

Richard L. Hansen, Sr. Orange County Legislator P.O. Box 201, Rt. 284 Unionville, NY 10988 Vincent Poloniak, Director Community Development 223 Main Street Goshen, NY 10924

Harvey J. Burger Orange County Legislator P.O. Box 208 252 Montgomery Street Newburgh, NY 12550

Patrick J. Berardinelli, Sr. Orange County Legislator 17 Hillside Court Newburgh, NY 10950

Dino J. Sciamanna Orange County Legislator 15th District 255-275 Main Street Goshen, NY 10924

State and Federal Officials and Agencies

The Honorable William J. Larkin, Jr. New York State Senator 1093 Little Britain Road New Windsor, NY 12553

The Honorable Thomas J. Kirwan New York State Assembly 190 South Plank Road Newburgh, NY 12550

The Honorable Daniel P. Moynihan United States Senator 189 Main Street Oneonta, NY 13820

The Honorable Charles Schumer United States Senator 26 Federal Plaza, Ste. 31-100 New York, NY 10728

The Honorable Maurice D. Hinchey United States Representative 291 Wall Street Kingston, NY 12401

Mark VanDeusen NYS Department of Health Flanigan Square 547 River Street Troy, New York 12180 Ellen Stoutenburgh, Public Affairs Officer NYSDEC 21 South Putt Corners Road New Paltz, NY 12561

Michael J. Knipfing Citizen Participation Specialist NYSDEC 21 South Putt Corners Road New Paltz, NY 12561

Richard Morse Legislative Program Council's Staff New York State Assembly, Room 520 State Capitol Albany, NY 12224

Marc Moran, Director Region 3, NYSDEC 21 South Putt Corners Road New Paltz, New York 12561

Al Klauss NYSDEC 21 South Putt Corners Road New Paltz, New York 12561

David Camp NYSDEC 50 Wolf Road Albany, New York 12233-7010

Local/Regional Media:

City Editor El Clarin 48 Broadway Haverstraw, NY 10927

City Editor Ellenville Press P.O.Box 31 Ellenville, NY 12428

City Editor Mid Hudson Times P.O. Box 434 Walden, NY 12586

City Editor Times Herald Record 126 Main Street New Paltz, NY 12561

City Editor Tri-State Gazette 84-88 Fowler Street Port Jervis, NY 12771

City Editor, Greenwood Lake &West Milford News P.O.Box 1117 Greenwood Lake, NY 10925

City Editor Hudson Valley Black Press P.O.Box 2160 Newburgh, NY 12550

City Editor Hudson Valley Business Journal P.O.Box 339 Pine Island, NY 10969 City Editor Independent Republican P.O. Drawer A Goshen, NY 10924-0628

City Editor News of the Highlands P.O.Box 278 Highland Falls, NY 10928

City Editor
Orange County Post
P.O.Box 500
Washingtonville, NY 10992

City Editor Ottaway Newspapers, Inc. P.O.Box 401 Campbell Hall, NY 10916

City Editor Photo News 45 Gilbert Street Monroe, NY 10950

City Editor Strauss Newspapers P.O.Box 190 Warwick, NY 10990

City Editor
The Sentinel
P.O.Box 406
Vails Gate, NY 12584

City Editor
The Cornwall Local
P.O.Box B
Cornwall, NY 12518

City Editor
The Warwick Valley Dispatch
P.O.Box 594
Warwick, NY 10990

City Editor Times Herald Record 233 Broadway Newburgh, NY 12550

City Editor
Times Herald Record
40 Mulberry Street
Middletown, NY 10940

City Editor Warwick Advertiser P.O.Box 190 Warwick, NY 10990

News Director WALL/WKOJ 1 Broadcast Plaza Middletown, NY 10940

News Director WDLC/WTSX P.O.Box 920 Port Jervis, NY 12771

News Director WDST 118 Tinker Street Woodstock, NY 12498

News Director WELV/WTHN 12 Tucker Drive Poughkeepsie, NY 12601

News Director WGNY P.O.Box 2307 Newburgh, NY 12550 News Director WHUD/WLNA Radio Terrace Peekskill, NY 10566

News Director WTBQ 62 N. Main Street Florida, NY 10921

News Director WZAD P.O.Box 978 Wurstburo, NY 12790

News Director Access 8 TV P.O.Box 321 Port Jervis, NY 12780

News Director Cable 6 TV Industrial Drive Middletown, NY 10940

News Director Time Warner Cablevision P.O.Box 887 Middletown, NY 10940

News Director TKR Cable Co. 19 South Street Warwick, NY 10990

News Director WRNN TV 721 Broadway Kingston, NY 12401

Adjacent/Nearby Residents, Businesses, Property Owners, and Other Interested Parties

<u>Property</u>	Owner's Name & Address
145-147 S. Robinson Ave.	Visconti, Frederick J. Jr. c/o Visconti Limosine Inc. P.O. Box 2666 Newburgh, NY 12550
149-155 S. Robinson Ave.	Same as above
157-161 S. Robinson Ave.	Gehler, John C. 3 Forshee St. Monroe, NY 10950
17-23 Dickson St.	Visconti, Frederick J. Jr. 17-23 Dickson St. Newburgh, NY 12550
Dickson St.	New York Central Lines LLC 500 Water St. (J-910) Jacksonville, FL 32202
20-32 Deyo Pl. & 184 Mill St.	M.J.S. Orange County Associates 34 Cottage St. Middletown, NY 10940
146-158 S. Robinson Ave.	Baltsas Stauros & Pitsinos Pano N. 160 S. Robinson Ave. Newburgh, NY 12550
160 S. Robinson Ave.	Same as above
162-170 S. Robinson Ave.	The Buller Limited Partnership 20 Rewe Street Brooklyn, NY 11211
170-172 S. Robinson Ave.	Levinson Enterprises Inc. P.O. Box 615 Cornwall, NY 12518
176-184 S. Robinson Ave.	Davel Properties, Inc. 91 N. 12 th St. Brooklyn, NY 11211

Property Address	Owner's Name & Address
29-37 Deyo Place	O'Shea, Edward A. 430 Pearl St. Kingston, NY 12401
S. Robinson Ave.	Cannon, Robert J. 2689 Route 208 Walden, NY 12586
19-37 Deyo Place	M.J.S. Associates 270 Rte. 17S Mahwah, NJ 07430
Deyo Place	Same as above
139-143 S. Robinson Ave.	Feroli, Leonard P. & Ronald 224 Leslie Ave. Newburgh, NY 12550
133-137 S. Robinson Ave.	R & F Associates Inc. 135 S. Robinson Ave. Newburgh, NY 12550
Dickson St.	20 Dickson Street Corp. c/o Carlos Scheer 38 Dogwood Hills Rd. Newburgh, NY 12550
Carson Ave.	New York Central Lines LLC 500 Water St. (J-910) Jacksonville, FL 32202
136-144 Mill St.	Greenblatt, Charles & Samuel 110 Mill St. Newburgh, NY 12550
159 Mill St.	Weller, Fred 169 Mill St. Newburgh, NY 12550
157 Mill St.	Same as above
155 Mill St.	Same as above
153 Mill St.	Same as above

Property Address	Owner's Name & Address
145-151 Mill St.	Cohen, Kathleen V. 215 Carson Ave. Newburgh, NY 12550
137-143 Mill St.	Bermo, Gloria 137 Mill St. Newburgh, NY 12550
187-193 Mill St.	Fracasse, Richard E. c/o New Windsor Refrigeration & Air Cond. 187 Mill St. Newburgh, NY 12250
183-185 Mill St.	Smith, William & Edwards B. 183 Mill St. Newburgh, NY 12550
86-92 Bridge St.	Coen, Nancy 111 Larter Ave. Newburgh, NY 12550
169-175 Mill St.	Weller, Fred R. 169-175 Mill St. Newburgh, NY 12550
161-167 Mill St.	Woinicki, Mark L. & Maria Z. 167 Mill St. Newburgh, NY 12550

Environmental Groups

Scenic Hudson 9 Vassar Street Poughkeepsie, NY 12601 (914) 473-440

Clearwater, Inc. 112 Market Street Poughkeepsie, NY 12601 (914) 454-7673

Greenway Conservancy Capitol Building Capitol Station, Room 254 Albany, NY 12224 (518) 473-3835

The Nature Conservancy P.O. Box 601 Pawling, NY 12564

Orange Environment P.O. Box 25 Goshen, NY 10924 (914) 294-5852 Hudson River Pilots Assoc. 75 Alexander Street Yonkers, NY 10701

Karl Coplan, Esq. Pace/Riverkeeper 78 North Broadway White Plains, NY 10603

Environmental Citizens Coalition 33 Central Ave. Albany, NY 12210

Laura Haight NY PIRG 107 Washington Avenue Albany, NY 12210

Orange County Land Trust Executive Director John Gebhards P.O. Box 2442 Middletown, NY 10940

Appendix D - Identification of Project Management Contacts

For additional information about the program to investigate the Provan/Ford Site, the public is encouraged to contact any of the following project staff:

New York State Department of Environmental Conservation (NYSDEC):

NYSDEC Inactive Hazardous Waste Site Toll-Free Information Number: 1-800-342-9296

(Calls recorded 24 hours a day. Calls are returned during business hours)

David Camp, P.E. Project Engineer NYSDEC 50 Wolf Road Albany, New York 12233-7010 (518) 457-7924 Mike Knipfing Citizen Participation Specialist NYSDEC Region 3 21 South Putt Corners Road New Paltz, New York 12561 (914) 256-3154

New York State Department of Health (NYSDOH):

Mark VanDeusen
Outreach Unit
NYS Department of Health
Flanigan Square
547 River Street
Troy, New York 12180
1-800-458-1158 ext. 27530

Teresa Foster
Health Specialist
NYS Department of Health
Flanigan Square
547 River Street
Troy, New York 12180
1-800-458-1158 ext. 27880

City of Newburgh/First Environment, Inc. PROVAN Project Staff

Scott Green, P.G. Senior Hydrogeologist First Environment, Inc. 90 Riverdale Road Riverdale, NJ 07457 1-800-486-5869 Tom Bambrick Geophysicist First Environment, Inc. 90 Riverdale Road Riverdale, NJ 07457 1-800-486-5869

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\$120.00	\$100.00	\$80.00	\$60.00
Proj. Sci./Eng.	Senior Scientist	Scientist	Technical/Drafting

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Scientist Technical/Drafting	PHASE I	Site Visit Report Writing

Total Spent \$500.00 \$2,500.00 \$3,000.00

Subtotal:

WORKPLAN/HASP/CP/PREPARATION

Site Visit with DEC Report Writing

\$500.00	\$5,500.00	
	Subtotal:	
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	Days/or	Well &		Cost Per		
Personnel	Units	Sam.#	Materials	Day	Sample Cost	Cost
PHASE II (work conducted to date)						
Geoprobe (27 Borings & 7 Piezometers)	က	4	\$5	\$890	\$0	\$2,688
Mobilization	က	0	\$0	\$100	\$0	\$300
Piezometer Well Materials (7 piezometers)	0	7	\$450	\$0	\$0	\$3,150
Labor & Oversight & Sampling	2 5	0	\$0	\$800	\$0	\$8,000
VO STARS	0	23	\$0	\$0	\$115	\$2,645
SVOC STARS	0	15	\$0	\$0	\$185	\$2,775
RCRA Metals	0	5	\$0	\$0	\$80	\$400
PCBs	0	9	\$0	\$0	\$65	\$390
Lead	0	က	\$0	\$0	\$65	\$195
Field Equipment	က	0	\$0	\$1,000	\$0	\$3,000
Travel & Food Expenses	2	0	\$0	\$50	\$0	\$250
					Subtotal:	\$23,793

Subcontractor costs in italics.

Cost		\$450	\$2,128	\$304	\$850	\$800	\$850	\$300	\$1,200	\$2,000	\$100	\$9,582			\$700	\$200	\$150	\$1,320	\$1,600	\$2,000	\$12,045	\$2,400	\$2,000	\$250	\$22,665		\$890	\$100	\$1,600	\$7,400	\$3,200	\$2,600	\$1,000	\$100	\$16.890
Sample Cost		\$450	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	Subtotal:			\$35	\$25	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	Subtotal:		\$0	\$0	\$0	\$185	\$80	\$65	\$0	\$0	Subtotal:
Cost Per Day/Drum		\$0	\$35	\$5	\$850	\$800	\$850	\$100	\$800	\$1,000	\$50				\$0	\$0	\$75	\$660	\$800	\$250	\$165	\$800	\$1,000	\$20			\$890	\$100	\$800	\$0	\$0	\$0	\$1,000	\$50	
Materials		\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0				\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	0\$			\$5	\$0	\$0	\$0	20	\$0	\$0	0\$	
Well & Sam.#		1	0	0	0	0	0	0	0	0	0				0	0	0	0	0	0	0	0	0	0			20	0	0	40	40	40	0	0	
Days/or Units		0	8.09	8.09	1	~	1	6	1.5	2	2				20	80	2	2	2	∞	73	က	7	2				-	~	0	0	0	1	~	
Personnel		0	0	0	0	~	0	0	~	0	0				0	0	0	0	0	0	0	0	0	0				0	2	0	0	0	0	2	
	Soil Removal & Drum Overpack	Soil Sampling Stockpile (ID 27 samp)	Soil Disposal per ton cost	Soil Loading per ton	Soil Trucking day per ton	FEI Labor & Oversight	Drum Overpack Chemist	Over Pack Cost per drum	FEI Labor & Oversight	Field Equipment	Travel & Food Expenses		PHASE III (work to be conducted)	Drum Characterization & Disposal	Composite Samples	Drum Profile Cost	Transportation	Chemist Labor	Oversight Labor	Drum Disposal Cost (Haz)	Drum Disposal Cost (non-Haz)	Labor & Oversight	Field Equipment	Travel & Food Expense		Surface Soil Delineation Sampling	Geoprobe (20 surface soil borings)	Mobilization	Labor & Oversight & Sampling	SVOC STARS	RCRA Metals	PCBs	Field Equipment	Travel & Food Expense	-

Subcontractor cost in italics

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Cost		\$1,700	\$200	\$1,350	\$1,600	\$10,500	\$1,500	\$1,700	\$1,600	\$2,000	\$200	\$22,350		\$50,635	\$8,000	\$2,550	\$805	\$5,550	\$400	\$325	\$175	\$8,400	\$1,200	\$1,700	\$1,350	\$1,600	\$4,000	\$600	\$87,290		\$2,670	\$300	\$3,104	\$2,250	\$3,200	\$2,400
Sample Cost		\$0	\$0	\$450	\$0	\$0	\$0	\$0	\$0	\$0	\$0	Subtotal:		\$0	\$0	\$85	\$115	\$185	\$80	\$65	\$25	\$0	\$0	\$0	\$450	\$0	\$0	\$0	Subtotal:		\$0	\$0	\$0	\$0	\$0	\$0
Cost Per Day	•	\$850	\$100	\$0	\$800	\$35	\$5	\$850	\$800	\$1,000	\$50			\$5,000	\$800	\$0	\$0	\$0	\$0	\$0	\$0	\$35	\$5	\$850	\$0	\$800	\$1,000	\$50			\$890	\$100	\$0	\$0	\$800	\$800
Materials		\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0			\$635	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0			\$0	\$0	\$388	\$0	\$0	\$0
Well & Sam. #		0	0	က	0	0	0	0	0	0	0			0	0	30	7	30	3	5	7	0	0	0	E	0	0	0			0	0	0	0	0	0
Days/or Units		7	2	0	2	300	300	2	2	2	4		osal	10	10	0	0	0	0	0	0	240	240	7	0	2	4	12			က	ო	∞	15	2	က
Personnel		0	0	0	1	0	0	0	-	0	0		ing & Soil Dispo	0	_	0	0	0	0	0	0	0	0	0	0	-	0	0			0	0	0	0	2	~
	Surface Soil Disposal	Excavation	Mobilization/Demob	Soil Stockpile (ID 27 sample)	Labor & Oversight & Sampling	Soil Disposal per ton cost	Soil Loading per ton	Soil Trucking day per ton	FEI Labor & Oversight	Field Equipment	Travel & Food Expense		UST Excavation, Backfill, Sampling & Soil Disp	UST Removal/Restoration	Labor & Oversight & Sampling	VO STARS (method 8021)	VOC (method 8240)	SVOC STARS (8027)	RCRA Metals	PCBs	Lead	Soil Disposal per ton cost	Soil Loading per ton	Soil Trucking day per ton	Soil Stockpile (ID 27 sample)	Labor & Oversight	Field Equipment	Travel & Food Expense		Well Installation & Sampling	Drilling Rig	Mobilization	Well Materials 8 wells	Well Survey 15 wells	Slug Tests & Interpretation	Labor & Oversight & Sampling

Subcontractor cost in italics

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Cost	\$850 \$575 \$2.775	\$1,200	\$3.200	\$1,600	\$350 \$26,330	6 0 0	\$1,600	\$1,200	\$100	\$4,500		0	\$2,225	\$1,800	\$2,100	\$1,000	\$900	\$8,125		\$6,000	\$6,300	\$227 R25	620,1220
Sample Cost	\$85 \$115 \$185	\$80	\$25 80	0\$	\$0 Subtotal:	é	09 99 99	\$0	\$0	Subfotal:	Cost Per Gal	2	5	57	20	0	0\$ \$	Subtotal:		0\$ \$	Subtotal:	TOTAL	
Cost Per. Day	\$ 8 \$0 \$0	0\$ 80	\$000	\$1,000	\$50	000	8800	\$600	\$20			000	\$800	\$800	\$800	0000	\$800	?		\$1,000	3		
Materials	\$ \$ \$ \$0\$	\$0 \$0	0\$	0\$	\$0	ç	09	\$0	\$0			ć	0,4	04	\$500	0000	\$100	•		\$100) - -		
Well & Sam. #	10 5 15	4	∞ ⊂	0	0	c	0	0	0			•	0	0 (0	5	0 0	Þ		00)		
Days/or Units	000	04	ro c	၊က	7	•		2	_			0017	1500	0001	Ν,	_	r "	-		0 г			
Personnel		00	0 0	10	0	c	4 62	~	2		aning &	•	>	5	۲ (o	<i>-</i> c	1		← C)		
Molland & Samiland	Well Installation & Sampling (cont'd) GW VO STARS (method 8021) GW VO (method 8240) GW SVOC STARS (8027)	GW RCRA Metals GW PCBs	GW Lead	Field Equipment	Travel & Food Expense	Pipe Draing & Assessment	Pipeline Survey	Map Production	Travel & Food Expense		Water Separator Pump-out & Cleaning & Disnosal		Pump Exterior Separator	Fump Interior Separator	Cleaning Separator	Separator Disposal	Cleaning Drain to Separator	ומיכו מ ו סכם באף כו סכ	RI/RAW Report	Labor Materials			

Subcontractor cost in italics

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