# REMEDIAL INVESTIGATION WORK PLAN

1022 Old Country Road Plainview, New York 11803 Site Number 130201

Prepared for

MORTON VILLAGE REALTY COMPANY, INC. % John Curran Sive Paget and Riesel P.C. 460 Park Avenue New York, New York 10022

## **ROUX ASSOCIATES, INC.**

Environmental Consulting & Management



209 Shafter Street, Islandia, New York 11749 ♦ 631-232-2600

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#### **CERTIFICATION**

I, Joseph Duminuco, certify that I am currently a Qualified Environmental Professional as defined in 6 NYCRR Part 375 and that this Remedial Investigation Work Plan was prepared in accordance with all applicable statutes and regulations and in substantial conformance with the DER Technical Guidance for Site Investigation and Remediation (DER-10).

Joseph D. Duminuco

Name

September 14, 2015

Date

Signature

## LIST OF ACRONYMS

LIST OF ACRONYMS
μg/kgMicrograms per Kilogram
μg/LMicrograms per Liter
1,2 DCEcis-1,2-dichloroethene
AOCsAreas of Concern
ASPAnalytical Services Protocol (NYSDEC)
AWQSGVsAmbient Water Quality Standards and Guidance Values
bgsBelow Ground Surface
CAMPCommunity Air Monitoring Plan
CERCLAComprehensive Environmental Response, Compensation, and Liability Act
CLPContract Laboratory Protocol
CVOCsChlorinated Volatile Organic Compounds
DER-10NYSDEC DER-10 Technical Guidance for Site Investigation and Remediation
DODissolved Oxygen
DUSRData Usability Summary Report
EAExposure Assessment
ELAPEnvironmental Laboratory Approval Program
ESAEnvironmental Site Assessment
FSPField Sampling Plan
HASPHealth and Safety Plan
mg/LMilligrams per liter
MWMonitoring Well
NTUsNephelometric Turbidity Units
NYCRRNew York Codes, Rules, and Regulations
NYSDECNew York State Department of Environmental Conservation
NYSDOHNew York State Department of Health
ORPOxidation – Reduction Potential
PIDPhoto Ionization Detector
PPEPersonal Protective Equipment
PVCPolyvinyl Chloride
QAQuality Assurance
QAPPQuality Assurance Project Plan
QCQuality Control

#### **LIST OF ACRONYMS (Continued)**

RWP.....Remedial Work Plan

RI.....Remedial Investigation

RIWP.....Remedial Investigation Work Plan

SB.....Soil Boring

SCOs .....Soil Cleanup Objectives

SRI .....Supplemental Remedial Investigation

SVOCs .....Semivolatile Organic Compounds

TAL.....Target Analyte List

TCL.....Target Compound List

TWA .....Total Weighted Average

USEPA......United States Environmental Protection Agency

USGS .....United States Geological Survey

VOCs.....Volatile Organic Compounds

#### 1.0 INTRODUCTION

Roux Associates, Inc. (Roux Associates), on behalf of Morton Village Realty Company, Inc., (Morton Village), has prepared this revised Remedial Investigation Work Plan (RIWP) for the Morton Village Plaza property located at 998-1064 Old Country Road, Plainview, Nassau County, New York, 11803 (Site). This revised RIWP addresses comments provided by the New York State Department of Environmental Conservation (NYSDEC) in the Disapproval of the February 2015 Remedial Investigation Work Plan comment letter dated July 24, 2015. The Site consists of four buildings that are currently occupied by various retail stores. The focus of this RIWP is on the former Morton Village Cleaners, a/k/a Classic French Cleaners, (former Cleaners) tenant space (1022 Old Country Road – currently occupied by a Subway restaurant) and on an area (asphalt parking lot) immediately north of the former Cleaners tenant space. The former Cleaners tenant space and parking lot area are collectively referred to as, Area of Concern 1 (AOC-1). The Site is currently listed in the Registry of Inactive Hazardous Waste Disposal Sites in New York State as Site Number 130201 with a Classification "2" pursuant to Environmental Conservation Law (ECL) 27-1305. The Site location is shown on Figure 1. A detailed description of the Site is provided in Section 2 of this revised RIWP.

This revised RIWP is being submitted for NYSDEC review and approval. This revised RIWP has been developed in accordance with the DER-10 Technical Guidance for Site Investigation and Remediation (May 2010) issued by the NYSDEC. The purpose of the RI is to further determine the nature and extent of impacted soil and groundwater within AOC-1, locate the historical septic sanitary system associated with AOC-1 (potentially a continuing source area), evaluate soil vapor conditions, and generate sufficient data necessary to support the development of a Feasibility Study Work Plan (FSWP).

#### 1.1 RI Work Plan Organization

This revised RIWP contains a background section (Section 2) describing the Site, its history, and results of previous environmental investigations; a section defining the objectives and scope of the RI (Section 3); and Sections 4 and 5 that describe various project operations plans (e.g., Quality Assurance/Quality Control Plan, Health and Safety Plan). Reporting requirements and the project schedule are discussed in Section 6. Additionally, tables are provided that summarize all

previously collected environmental quality data. Previous sampling locations are presented on Figure 2. A map of the proposed sampling locations is presented on Figure 3.

#### 1.2 Project Team Contact Information

Roux Associates' Principal-In-Charge for this Site will be Joseph Duminuco, Principal Hydrogeologist and Vice President. Mr. Duminuco is based at Roux Associates' Islandia, New York headquarters office and can be reached at (631) 232-2600. Jeff Wills will be the Project Manager, and will be responsible for day-to-day management of the project, including preparation of work plans, and scoping and directing field activities.

The contact for the Property Owner is:

John Patrick Curran, Esq. Sive Paget & Riesel P.C. 460 Park Avenue New York, New York 10022

At this time, the following subcontractors have been selected for this project:

- <u>Drilling</u>: Aquifer Drilling and Testing, Inc., Mineola, New York
- Analytical: Alpha Analytical, Westborough, Massachusetts

Other subcontracted services have not yet been selected. This information will be provided to NYSDEC following contractor selection.

### 2.0 BACKGROUND

This section provides pertinent background information, including a description of the Site and its setting, the known history of the Site, and the results of previous environmental investigation work conducted at the Site.

## 2.1 Site Description and Setting

Property Location							
Property Name:	Morton Village Plaza						
Property Description:	The property is occupied by Morton Village Shopping Center, which consists of four buildings situated on four adjacent lots (Lots 10, 86, 88 and 89). The on-Site buildings are currently occupied by various professional businesses, retail stores, restaurants and a grocery store. The property is bordered by Knowles Street to the north, Old Country Road to the south, Lester Place to the east and Rex Place to the west.						
Property Address:	998-1064 Old Country Road						
Property Town, County, State:	Plainview, Nassau County, New York						
Property Tax Identification:	Block 555 Lots 10, 86, 88 and 89						
Property Topographic Quadrangle:	USGS Huntington Quadrangle, New York (1979)						
Nearest Intersection:	Rex Place and Old Country Road						
Area Description:	The area surrounding the Site is used mainly for residential purposes. Surrounding properties to the north, east and west are all residential properties. To the south of the Site, there are both residential properties as well as the Plainview-Old Bethpage Public Library.						
Current Site Zoning:	Commercial-Use, 452.14 - Area/Neighborhood Shopping Center						
Property Acreage:	9.936 acres						
Property Shape:	Rectangular						
Property Use:	The property is currently occupied by various professional businesses, retail stores, restaurants and a grocery store.						

#### 2.1.1 Site Operations

The Site is currently occupied by various professional businesses, retail stores, restaurants and a grocery store. The former Cleaners tenant space within AOC-1 is currently occupied by a Subway restaurant.

#### 2.1.2 Utilities

The following companies and municipalities currently provide utility services to the Site:

Utility	Provider
Electricity	PSEG
Natural Gas	PSEG
Sanitary/Storm Sewerage	Nassau County
Potable Water	Plainview Water District

#### 2.1.3 Topography/Hydrogeology

The property location is shown on the 1979 USGS Topographic Map of Huntington, New York. The surface elevation of the property is approximately 145 feet above mean sea level. Topography of the property slopes slightly to the south.

Groundwater was encountered at approximately 70 feet below ground surface (ft-bgs) during previous environmental investigations conducted by Leggette, Brashears & Graham, Inc. (LBG) and HRP Associates, Inc. (HRP). Based on previous environmental investigations conducted by LBG and HRP, groundwater beneath the Site flows to the south.

#### 2.1.4 Wetland Areas and Surface Water Bodies

There are no identified wetlands on or adjacent to the Site. There are no surface water bodies on or adjacent to the Site. The Long Island Sound is approximately 7 miles north of the Site.

#### **2.1.5 Soils**

According to the USGS, soils in the area of the Site are classified as Urban Land (Ug). Ug consists of areas where at least 80-85 percent of the surface is covered by asphalt, concrete, or other impervious building materials. Based on Soil Conservation Service STATSGO data, native soils beneath the Site and the surrounding area are listed as: Surficial Soils (fine sandy loam, loam,

loamy sand, silt loam and sand); Shallow Soils (silt loam); and Deeper Soils (sandy loam, gravelly – coarse sand, and coarse sand). Based on the previous investigations, described below, the soil beneath the Site consists primarily of fine to medium sand with varying amounts of silt and trace gravel.

#### **2.1.6 Underlying Formation**

The Site is located on the southeastern portion of Nassau County, New York. Based on previous investigations, the soil beneath the Site consists of sand with varying amounts of silt and gravel. The rock stratigraphic unit beneath the Site consists of Cenozoic, Quaternary rock and is categorized as a Stratified Sequence (USGS, 1994).

#### 2.1.7 Neighboring Properties

A review of neighboring properties from the Site and from public thoroughfares, and research of available information regarding the neighboring properties, was performed to identify evidence of environmental concerns that could adversely impact the Site. The Site is located in a mixed-use commercial and residential area of Plainview, Nassau County, New York. Based on the Environmental Site Assessment Phase I report prepared by LBG, a Gas Station historically occupied the property (Section 12, Block 555, Lot 6) adjacent to the southeast corner of the Site.

Direction	Operations
North	Residential Properties
East	Residential Properties
West	Residential Properties
South	Residential Properties and Plainview-Old Bethpage Public Library

#### 2.2 Site History

The following summary of Site history was based on a Phase I Environmental Site Assessment performed by LBG in February 2009.

The property was previously used for agricultural purposes until developed as a shopping center in 1956. The property has been improved with commercial developments and past commercial occupants included a dry cleaning facility (i.e., AOC-1) from the late 1950's through 2007.

Previous investigations (soil, groundwater, and vapor sampling) performed at the Site from 2006 to 2011 identified petroleum-related compounds and chlorinated compounds in the soil and soil vapor and chlorinated compounds in groundwater in the vicinity of AOC-1. The petroleum-related compounds were associated with a former underground storage tank (UST) that was identified and removed during Site characterization work in 2007 conducted by LBG, as described in Section 2.3.3. Operations at the former Cleaners have resulted in contamination of the soil around a sump located at the northern edge of the Site building within the basement of the former Cleaners, as well as the groundwater in the vicinity of the former Cleaners. Prior to the 1970's, there were cesspools and leaching fields (potentially a continuing source area) installed throughout the Site. The Site's sanitary system was not connected to the Municipal sanitary sewer line until the 1970's.

#### 2.3 Previous Environmental Investigations

This section provides an overview of the results of previous environmental investigations at the Site. The following environmental reports were reviewed by Roux Associates:

- Subsurface Investigation Letter Report Dry Cleaning Operation Morton Village Plaza prepared by Galdun Frankel Environmental dated October 2006 on behalf of Morton Village Realty Co., Inc.
- Environmental Site Assessment Phase II Report prepared by Leggette, Brashears & Graham, Inc. (LBG) dated September 2007 on behalf of Morton Village Realty Co., Inc.
- UST Closure and Remedial Summary Report Former Classic French Cleaners Morton Village Shopping Center prepared by LBG dated September 2008 on behalf of Morton Village Realty Co., Inc.
- Phase I Environmental Assessment Morton Village Plaza prepared by LBG dated February 2009 on behalf of Morton Village Realty Co., Inc.
- Site Characterization Report Former Morton Village Cleaners prepared by HRP Associates, Inc. dated August 2011 on behalf of the NYSDEC.

Results of these reports are discussed in the subsections below.

#### 2.3.1 Galdun Frankel Environmental, Subsurface Investigation Report

In September 2006, Galdun Frankel Environmental (GFE) completed a subsurface investigation at the Site to determine if soil or groundwater contamination has resulted from the operation of the former Cleaners. The GFE investigation included:

- Installation four soil boring (B1, B2, B3 and B4);
- Field screening of soil from each soil boring using a photoionization detector (PID) and visual observations; and
- Collection of six soil samples (B1-20', B2-3', B2-13', B2-25', B3-7' and B4-7'), one sediment sample (SUMP SEDIMENT) and two groundwater samples (B1GW and B2GW).

Soil boring B1 was installed on the southern side of the former Cleaners, and B2, B3 and B4 were installed on the northern side of the former Cleaners. B1 and B2 were advanced to a target depth of 85 ft-bgs; however, soil screening was not conducted from 50 to 85 ft-bgs at these locations. B3 and B4 were advanced to a target depth of 25 ft-bgs. Groundwater was observed at 75 ft-bgs. Groundwater samples were collected from B-1 and B-2 at a sample interval of 75 to 85 ft-bgs.

GFE inspected a sump located in the basement floor, on the north side of the former Cleaners. The sump received blow-down water from a boiler that was used in the dry cleaning process. The blow-down water was pumped to an overhead sewer line, which extended in an east to west direction, into adjoining commercial spaces. The sewer main is located to the north of the shopping center. GFE identified a void between the bottom of the basement floor slab and the top edges of the sump. A sediment sample was collected from inside the void. All soil, sediment and groundwater samples were sent to AmeriSci Laboratories and analyzed for volatile organic compounds (VOCs).

#### **Summary of Subsurface Investigation Results**

Field Screening

Elevated PID readings were detected in the first 13 feet of soil at B2. A PID reading of 1000 parts per million (ppm) was detected in soil from the 1 to 7 ft-bgs interval. Based on field screening activities, little to no evidence of contamination was observed from B1, B3 and B4.

Soil

Tetrachloroethylene (PCE) was detected in sample B2-3' at a concentration of 234 ppm.

#### Sediment

PCE was detected in SUMP SEDIMENT at a concentration of 3.74 ppm. Trichloroethylene (TCE) was detected in sample SUMP SEDIMENT at a concentration of 0.0147 ppm.

#### Groundwater

PCE was detected in samples B1GW and B2GW at a concentration of 11.9 parts per billion (ppb) and 546 ppb, respectively.

TCE was detected in sample B2GW at a concentration of 9.97 ppb. Cis-1,2-Dichloroethylene was detected in sample B2GW at a concentration of 5.09 ppb.

#### 2.3.2 Leggette, Brashears & Graham, Inc. Environmental Site Assessment Phase II

In May 2007, Leggette, Brashears & Graham, Inc. (LBG) completed a subsurface investigation at the Site to determine the vertical and horizontal extent of soil and groundwater contamination in the vicinity of the former Cleaners. LBG's investigation included:

- Sewer and septic investigation;
- Field screening of soil from six soil borings (GP-1, GP-2, GP-3, MW-1, MW-2 and MW-3);
- Installation of three groundwater monitoring wells (MW-1 through MW-3); and
- Collection of four soil samples (GP-1 (5-7), MW-2 (7-9), GP-3 (9-11) and MW-3 (11-13), three groundwater samples (MW-1 through MW-3) and one sediment sample (Sump Sediment). All samples were sent to AMRO Environmental Laboratories, Corp and analyzed for VOCs including methyl tertiary-butyl ether (MTBE).

Soil borings GP-1, GP-2, GP-3, MW-2 and MW-3 were installed on the northern side of the former Cleaners; MW-1 was installed on the southern side of the former Cleaners.

During drilling activities at soil boring GP-2, a 6-inch diameter pipe was encountered at 6 ft-bgs. The pipe was identified as an active sanitary sewer connected to the former Cleaners sanitary

facilities. Soil above and below the pipe was screened with a PID; no elevated PID readings were detected.

As per LBG, Building Department data did not reveal a septic tank in the area.

#### **Summary of Subsurface Investigation Results**

Field Screening

The highest PID reading was detected from the 5 to 7 ft-bgs interval at GP-1 at a concentration of 18 ppm. There were no elevated PID readings in soils screened from soil borings GP-2 and MW-1.

Soil

PCE was detected in samples GP-1 (5-7) and GP-3 (9-11) at a concentration of 410 ppb and 85 ppb, respectively.

Sediment

PCE was detected in sample Sump Sediment (0-1), collected from the bottom of the sump, at a concentration of 830 ppb.

Groundwater

PCE was detected in MW-1, MW-2 and MW-3 at a concentration of 120 ppb, 72 ppb and 93 ppb, respectively.

TCE was detected in MW-1, MW-2 and MW-3 at a concentration of 16 ppb, 6.9 ppb and 9.4 ppb, respectively.

Cis-1,2-Dichloroethylene was detected in MW-1, MW-2 and MW-3 at a concentration of 23 ppb, 4.8 ppb and 7.4 ppb, respectively.

#### 2.3.3 LBG UST Closure and Remedial Summary Report

In April 2008, LBG conducted remedial excavation activities at the Site, which included:

• Screening of soil during excavation activities with a PID;

- Removal of a sump structure located in the basement of the Site and excavation of soils beneath the former sump location;
- Location, excavation and removal of a suspected vent pipe located on the exterior of the north side of the Site;
- Removal of one 1,000-gallon underground storage tank (UST) encountered during exterior vent pipe excavation activities;
- Excavation of contaminated soils encountered in the vicinity of the former unknown UST;
- Collection of endpoint confirmation soil samples from the excavations for laboratory analysis; and
- Treatment of soil within the interior and exterior excavation areas with a remedial chemical oxidation compound in order to remediate any residual contamination.

The sump located in the basement of the former Cleaners measured two feet wide by two feet long by one foot deep. A 5.5 foot by 3.5 foot area was saw cut around the sump, and the entire structure was removed. Following removal of the sump structure, the area was excavated to 8 feet below the basement floor slab and a sump bottom soil endpoint sample was collected (Sump Bottom). Sump Bottom was submitted to Environmental Testing Laboratories, Inc., for analysis of VOCs, including MTBE.

During excavation activities of a vent pipe that was believed to have been associated with the heating oil above ground storage tank (AST) located in the former Cleaner's basement, a previously unknown 1,000-gallon single-wall steel UST was encountered. Approximately 50 gallons of oily water and sludge were removed from the interior of the UST by Metro Environmental. Upon removal, the UST was observed to be in poor condition with extensive surface corrosion and holes. Soil beneath the UST was impacted with petroleum. The New York State Department of Environmental Conservation was notified and spill number 08-00596 was assigned to the Site.

A total of seven endpoint soil samples (CL-N-SW, Bottom Main EX, East SW, West SW, Bottom F/O, South F/O and North SW F/O) were collected under the excavated UST. An eighth sample (South SW Base) was collected from the base of the building footing (former Cleaners).

All samples were submitted to Environmental Testing Laboratories, Inc., for analysis of VOCs including MTBE and semivolatile organic compounds (SVOCs).

During excavation activities, a total of 250.31 tons of soil was removed from the Site for disposal.

Following all excavation activities, a total of 360 pounds of the chemical oxidant RegenOx<sup>TM</sup> was mixed with water, according to manufacturer specifications, and applied directly to the southern portion of the UST excavation area along the foundation footings of the building, and 60 pounds of RegenOx<sup>TM</sup> mixed with water was applied directly to the bottom of the sump excavation. Prior to backfilling, a 4-inch slotted PVC sump was installed approximately 12.5 ft-bgs within the southern portion of the excavation, which was intended to serve as an additional precautionary measure to allow access to the subsurface for additional chemical oxidation applications and/or high-vacuum soil vapor extraction events, if necessary.

#### **Post-Excavation Sample Results**

There were no VOC or SVOC concentrations exceeding Technical and Administrative Guidance Memorandum (TAGM) 4046 from any of the endpoint samples collected within the UST excavation.

There were no VOC concentrations exceeding TAGM 4046 from the endpoint soil sample collected from the sump excavation.

Groundwater samples were not collected during UST closure and remedial activities.

#### 2.3.4 LBG Phase I Environmental Assessment

A Phase I ESA was conducted in 2009 by LBG on behalf of Morton Village, to inspect the Site and review historical and current land usage at and in the vicinity of the Site to evaluate if any RECs are present at or affecting the Site. The scope of work included a Site reconnaissance, review of Federal, State and local databases, research of historical documentation and review of available past environmental reports for the Site and Adjacent properties. In addition, the scope of work included the regulatory database investigation of all properties within ASTM standard search distances from the Site.

#### **Phase I Findings**

The Blue Angel Diner had corrosive cleaners stored in the basement as well as a refrigerant system for the two walk-in refrigerator/freezers. A floor drain was located directly beneath the refrigerant system equipment.

Numerous chemicals were observed in the basement of Hi Tech 1-Hour Photo in the area of a chemical mixing station, two photograph printing units and a chemical recovery system. Significant staining was observed on the floor surrounding the photograph development equipment as well as the chemical recovery system.

The main sanitary sewer line for Building B runs along the north side of the buildings and connects to the sewer main located on Rex Place. However, cesspools and leaching fields were installed on the Site prior to connection to the sewer and the locations of these former structures are unknown.

Stormwater is collected in catch-basins and is discharged through drywells/leaching pools.

#### 2.3.5 HRP Associates, Inc. Site Characterization Report

During November 2010, a Site characterization was completed by HRP Associates, Inc. (HRP) on behalf of the NYSDEC to investigate on-Site media (soil, soil vapor and groundwater) potentially impacted by operations of the former Cleaners. The HRP Site characterization included:

- Installation of five soil borings (SB-01 through SB-05) to a target depth of 120 ft-bgs;
- Collection of 26 soil samples from intervals described below:
  - Samples were collected at SB-1, SB-3, SB-4 and SB-5 from intervals: 79-81 ft-bgs, 89-91 ft-bgs, 99-101 ft-bgs, 109-111 ft-bgs, and 119-121 ft-bgs;
  - Samples were collected at SB-2 from intervals: 79-81 ft-bgs, 89-101 ft-bgs, 99-101 ft-bgs, and 114-116 ft-bgs;
- Collection of 22 discrete groundwater samples from the five soil borings (SB-1 through SB-5). The groundwater samples collected from SB-1 through SB-5 were identified as GW-1 through GW-5 and were collected from intervals described below:
  - Samples were collected at GW-1, GW-4, and GW-5 from intervals: 91-91.5 ft-bgs, 101-101.5 ft-bgs, 111-111.5 ft-bgs, and 121-121.5 ft-bgs;

- Samples were collected at GW-2 from intervals: 81-81.5 ft-bgs, 91-91.5 ft-bgs, 101-101.5 ft-bgs, 111-111.5 ft-bgs, and 118-120 ft-bgs;
- Samples were collected at GW-3 from intervals: 81-81.5 ft-bgs, 91-91.5 ft-bgs, 101-101.5 ft-bgs, 111-111.5 ft-bgs, and 121-121.5 ft-bgs;
- Development and collection of three groundwater samples from existing monitoring wells (MW-1 through MW-3); and
- Installation and collection of six soil vapor samples (SV-1 through SV-6) and one outdoor ambient air sample (AA-1). All soil vapor points were installed at a depth of 8 ft-bgs. SV-1 through SV-4 were installed in the parking lot behind (northern side) of the former Cleaners, and SV-5 and SV-6 were installed in the side walk in front (southern side) of the former Cleaner.

All soil and groundwater samples were sent to Environmental Laboratories Testing, Inc and all soil vapor samples were sent to Centek Laboratories. All soil samples were analyzed for VOCs. A total of five soil samples (one per boring) were also analyzed for SVOCs, pesticides, polychlorinated biphenyls (PCBs), and 8 RCRA Metals.

All groundwater samples were analyzed for VOCs, SVOCs and 8 RCRA Metals (Total and Dissolved).

All soil vapor samples were analyzed for VOCs (TO-15).

#### **Summary of Results**

Soil

There were no VOC or SVOC detections above Part 375 Unrestricted Use soil cleanup objectives (Unrestricted SCOs) or Commercial Use SCOs in any of the soil samples.

Chromium was detected in all five soil samples (SB-1 (99-101), SB-2 (79-81), SB-3 (89-91), SB-4 (119-121) and SB-5 (99-101) above Part 375 Unrestricted Use Soil Cleanup Objectives (Unrestricted SCOs) at concentrations ranging from 1.38 mg/kg to 2.02 mg/kg. There were no metal detections above Commercial SCOs.

#### Groundwater

Multiple VOCs including, PCE, TCE, cis-1,2-Dichloroethylene, and methyl tert-butyl ether (MTBE) were detected above NYSDEC Class GA Criteria in multiple samples collected from discrete groundwater samples (GW-1 through GW-5) as well as the existing monitoring wells samples, with the highest concentrations detected in GW-3 and MW-3. GW-3 and MW-3 are both located immediately south of the former Cleaners.

PCE was detected at concentrations ranging from 0.95 micrograms per liter ( $\mu$ g/L) to 88  $\mu$ g/L, as detected in samples GW-1 (121-121.5) and MW-3, respectively.

TCE was detected at concentrations of  $46 \,\mu\text{g/L}$ ,  $25 \,\mu\text{g/L}$  and  $43 \,\mu\text{g/L}$ , as in GW-3 (81-81.5), GW-3 (91-91.5) and MW-3, respectively.

No metals, pesticides or PCBs were detected above NYSDEC Class GA Criteria in any groundwater samples.

Soil Vapor

Multiple VOCs including, chlorinated VOCs and petroleum related VOCs were detected in all of the soil vapor samples and ambient air sample.

PCE was detected in soil vapor samples at a concentration ranging from 110 ppbv to 3,900 ppbv. PCE was detected in the outdoor ambient air sample at a concentration of 15 ppbv.

TCE was detected in soil vapor samples at a concentration ranging from 0.87 ppbv to 580 ppbv. TCE was not detected in the outdoor ambient air sample.

#### 2.4 Data Usability

Previous groundwater and soil analytical data developed by GFE, LBG and HRP were evaluated by Roux Associates and assumed reliable. All previous soil and groundwater analytical data are provided in Tables 1 and 2, respectively.

#### 3.0 RI WORK PLAN OBJECTIVES, SCOPE, AND RATIONALE

This section provides a description of the RI objectives, scope of work and rationale.

#### 3.1 Objectives and Rationale to Develop Remedial Investigation Work Plan

The previous environmental investigations have documented the following area of concern (AOC-1):

- Former Morton Village Cleaners, a/k/a Classic French Cleaners, (including the area [asphalt lot] immediately north of the former Cleaners):
  - Chlorinated VOCs (CVOCs) in soil and groundwater.

Based on the available environmental reports for the Site and known data gaps, including no source area being identified, the following objectives have been identified for the revised RIWP:

- Evaluate soil, groundwater and soil vapor quality associated with AOC-1;
- Evaluate the potential for impacts from the suspected historical septic system (potential source of CVOCs in soil/groundwater) that was associated with the former Cleaners; and
- Delineate the nature and extent of previously-documented CVOC impacts to soil and groundwater.

In order to address these objectives, this investigation will be completed in a phased approach (Phase A and Phase B).

Phase A of the RI will include the collection of soil samples and the installation of groundwater monitoring wells and collection of groundwater samples using a Geoprobe<sup>®</sup>. The purpose of this phase of the RI is to locate a suspected historical septic sanitary system and to collect sufficient sub-surface samples (soil and groundwater) to generate analytical data so that, together with the historical data generated by others, including groundwater and soil sampling, the appropriate sample locations for Phase B of the RI can be determined, as described below.

Phase B of the RI will include the installation and sampling of an upgradient groundwater monitoring well (RMW-3). The location of RMW-3 will be dependent on Phase A findings (i.e., groundwater flow direction). In addition, Phase B will include the completion of a soil vapor investigation at the Site. The soil vapor investigation will include the collection of soil vapor samples from locations specific to historical data and Phase A findings as well as the collection of

sub-slab and indoor air samples within the on-Site building (currently a Subway restaurant). The purpose of this phase of the RI is to collect sufficient sub-surface samples to generate analytical data so that, together with the historical data generated by others and data generated by Phase A of this RI, including groundwater and soil sampling, the Site will be sufficiently characterized to support the development of the FSWP.

All Phase B sample locations will be discussed with and approved by NYSDEC and New York State Department of Health (NYSDOH) prior to completion.

If a source area of the CVOC impacted groundwater (i.e., historical septic sanitary system) is not located, the environmental data collected during Phase A and B of the RI will be used in conjunction with the previous data to develop the scope of work for a supplemental RI. However, if a source area is identified, the environmental data will be used to develop the information necessary to support the development of a FSWP and identify the remediation track to be targeted.

#### 3.2 Phase A RI Scope

The scope of Phase A of the RI is designed to locate a suspected historical septic sanitary system and to collect sufficient sub-surface samples to generate analytical data so that, together with the historical data generated by others, including groundwater and soil sampling, the appropriate sample locations for Phase B of the investigation can be determined in order to sufficiently characterize the Site to support the development of the FSWP. To accomplish this, Phase A of the RI will focus on the following:

- Attempt to locate a potential source area (historical septic sanitary system); and
- The collection of soil and groundwater data sufficient to define the nature and extent of impacted media.

The scope of each component of Phase A of the RI is discussed in the following subsections. The proposed scope of work will be as follows:

• Site's records on both the town (Oyster Bay) and county (Nassau) level will be reviewed to determine whether a historic sanitary system (i.e. septic tank, leaching pools) exists beneath the Site that may have received discharges from the former Cleaners and may be acting as a continuing source of VOCs to groundwater beneath the Site.

- Conduct a geophysical survey including but not limited to, ground penetrating radar (GPR) to identify a suspected historical septic system in the vicinity of the former Cleaners space;
- Install six soil borings in order to delineate extent of impacts to soil, and address existing data gaps;
- Install one soil boring on the northeast side of the former UST excavation area, as requested by the NYSDEC in their letter dated July 24, 2015, to confirm current soil conditions in this area;
- Install up to two monitoring wells and collect groundwater samples to further assess groundwater quality; and
- Collect one groundwater grab sample (water table sample) in the vicinity of previously installed soil boring B-1, as requested by the NYSDEC in their July 24, 2015 comment letter, to confirm the groundwater quality at this location.

Detailed field sampling procedures are provided in the FSP (Appendix A). The proposed sampling locations are shown on Figure 3. Tables 3 and 4 summarize the approximate location and rationale for the proposed sampling locations.

#### 3.2.1 Historical Septic System Investigation

Prior to any invasive activity, Roux Associates will use a subcontractor to conduct a geophysical survey in an attempt to identify historical sanitary systems and underground utility lines (i.e., septic, sewer, water, electric, gas, etc.). The geophysical survey will utilize electromagnetic resonance and GPR technology. The survey will be performed in the back of the shopping plaza, behind the building that includes the former Cleaners (AOC-1).

#### 3.2.2 Soil Characterization

Six soil borings (RSB-1 through RSB-6) will be installed at locations biased toward the location (determined or assumed) of the historical septic sanitary system. The proposed soil boring locations are shown on Figure 3. Actual locations may vary based on field conditions (i.e., agency file reviews, results of the geophysical survey, access constraints, subsurface obstructions, and/or utilities). An additional soil boring (RSB-7) will be installed on the northeast side of the former UST excavation area (see Figure 3). All soil boring locations will be cleared to a minimum of five ft-bgs using hand tools and/or vacuum excavator to identify any potential subsurface utilities. Following utility clearance activities, the soil borings will be advanced using a Geoprobe® or

hollow stem auger (HSA) drill rig. Soil samples will be collected continuously, and will be visually inspected and screened for organic vapors with a photoionization detector (PID), until below the assumed bottom depth of the historical sanitary system (approximately 30 ft-bgs), with the exception of RSB-7. At RSB-7, soil samples will be collected continuously to two feet below the assumed UST bottom excavation depth of 7 ft-bgs (LBG, 2008). Soil lithology will be recorded according to the Unified Soils Classification System (USCS). If evidence of contamination (organic vapors, staining or odors) is present at 30 ft-bgs at RSB-1 through RSB-6 and 9 ft-bgs at RSB-7, the drilling will continue at the boring location until the evidence is no longer detected. A soil sample will be collected from the interval exhibiting the greatest evidence of impacts such as elevated PID detections, odors or staining. A second soil sample will be collected from the terminal depth of the boring (to be determined during field activities). If no evidence of impacts is observed during borehole advancement, the soil sample will be obtained from 28-30 ft-bgs interval at RSB-1 through RSB-6 and from 7-9 ft-bgs interval at RSB-7 and submitted for laboratory analysis.

Attempts will be made to reach the desired termination depths using a Geoprobe<sup>TM</sup> drill rig. If the Geoprobe<sup>TM</sup> drill rig is consistently unable to reach the termination depths, the sampling program may be modified to include the HSA drill rig.

All soil samples will be analyzed for the full Target Compound List (TCL) VOCs via United States Environmental Protection Agency (USEPA) Method 8260. All soil samples will be analyzed by Alpha Analytical of Westborough, Massachusetts, which is a New York State Department of Health Environmental Laboratory Approval Program-certified laboratory. Samples will be analyzed on a standard turnaround time and will be reported as Category B data deliverables, which will be used to generate a data usability summary report (DUSR).

All proposed soil borings not converted to monitoring wells will be backfilled with soil cuttings, if no evidence of impacted soil, and finished at grade with an asphalt patch. If evidence of impacted soil is observed at a soil boring, the soil will be containerized in 55 gallon DOT-approved drums and the soil boring will be backfilled with clean sand. Excess soil cuttings as well as the aforementioned impacted soil generated during soil boring activities will be containerized in 55 gallon DOT-approved drums, labeled, and stored on-Site pending laboratory results and proper off-Site disposal.

#### 3.2.3 Groundwater Investigation

If contamination is observed, the two groundwater monitoring wells (RMW-1 and RMW-2) are proposed to be installed in the two soil borings with the greatest degree of evidence of contamination. RMW-1 and RMW-2 will be installed at approximately the same depths (i.e., 70-90 ft-bgs) as existing monitoring wells MW-1 through MW-3 in order to verify previous results at that depth. Each monitoring well will be constructed of 2-inch diameter Schedule 40 polyvinyl chloride (PVC) with a 10-slot screen installed to span the water table. A gravel pack consisting of #2 Morie Sand or equivalent will be placed around the screen and up to two-feet above the top of the screened interval followed by a 1 to 2 foot layer of bentonite pellets. The bentonite pellets will be given time to hydrate before filling the remainder of the well annulus with bentonite grout using the tremie method. All monitoring wells will be completed with a flush-mounted manhole protective curb box installed at grade. The proposed monitoring well locations are shown on Figure 3. If evidence of contamination is not detected in any of the six soil borings, we propose to install one monitoring well in this area for additional Site characterization.

Each well will be developed to ensure proper hydraulic connection with the aquifer and to reduce/eliminate turbidity. The wells will be developed using a submersible pump, which will be surged periodically until well yield is consistent and has turbidity below 50 nephelometric turbidity units (NTUs), or the equivalent of 10 well casing volumes have been purged. Detailed procedures regarding well development are found in the FSP (Appendix A). Excess soil cutting and groundwater generated during well installation (including development and purging prior to sample collection) will be containerized in 55 gallon DOT-approved drums, labeled, and stored on-Site pending laboratory results and proper off-Site disposal.

To characterize groundwater flow and quality conditions following well installation and development, the new monitoring wells will be gauged and sampled. As part of the gauging round, water-level measurements will be recorded from the newly installed wells (RMW-1 and RMW-2) as well as existing monitoring wells (MW-1 through MW-3) to further define groundwater flow patterns beneath the Site.

Following the groundwater gauging event and waiting one week after well development, a groundwater sampling event will be completed. To ensure groundwater samples collected are representative of the conditions in the surrounding aquifer, monitoring wells will be purged prior to sample collection using low flow procedures as outlined in USEPA document titled "Low Stress (Low Flow) Purging and Sampling Procedures for the Collection of Groundwater Samples From Monitoring Wells" (USEPA, 2010). Additional information regarding groundwater sampling procedures is in the FSP (Appendix A).

Groundwater samples will be collected and analyzed for:

TCL VOCs via USEPA Method 8260

Field parameters (dissolved oxygen, pH, conductivity, turbidity, temperature and oxidation-reduction potential) will also be collected during well sampling activities.

In addition, a groundwater grab sample will be collected at the water table from a boring (RB-1) installed adjacent to previously completed soil boring B-1 (installed by HRP), as requested by the NYSDEC in their comment letter dated July 24, 2015. RB-1 will be installed using a Geoprobe® or HSA drill rig to establish current groundwater conditions at this location.

During groundwater grab sampling activities, a two foot long retractable stainless steel screen will be advanced by the Geoprobe® to two feet below the water table (approximately 70 ft-bgs). Upon reaching sampling depth (approximately 72 ft-bgs), the drill string will be retracted by two feet to expose the screen. A groundwater grab sample will be collected using the same sampling procedures as describe above.

If the Geoprobe® is unable to advance to sampling depth (approximately 72 ft-bgs), RB-1 will be completed with a HSA rig. Upon reaching sampling depth with the HSA, a temporary 2-inch diameter PVC well will be installed within the drill string (i.e., augers) and a groundwater grab sample will be collected using the sample sampling procedures describe above.

Groundwater grab sample will be collected and analyzed for:

• TCL VOCs via USEPA Method 8260

Field parameters (dissolved oxygen, pH, conductivity, turbidity, temperature and oxidation-reduction potential) will also be collected during groundwater grab sampling activities.

All groundwater samples will be analyzed by Alpha Analytical of Westborough, Massachusetts, which is a New York State Department of Health Environmental Laboratory Approval Program-certified laboratory. Samples will be analyzed on a standard turnaround time and will be reported as Category B data deliverables.

More information on sample analyses is provided in the QAPP (Appendix B).

#### 3.3 Phase B RI Scope

The scope of Phase B of the RI is designed to collect sufficient sub-surface samples to generate analytical data so that, together with the historical data generated by others and data generated by Phase A of this RI, including groundwater and soil sampling, the Site will be sufficiently characterized to support the development of the FSWP. To accomplish this, Phase B of the RI will focus on the following:

• The collection of an additional groundwater sample (i.e., upgradient monitoring well), and soil vapor, sub-slab and indoor air data sufficient to define the nature and extent of impacted media.

The scope of each component of Phase B of the RI is discussed in the following subsections. The proposed scope of work will be as follows:

- Installation and sampling of an upgradient monitoring well (location will be based on Phase A groundwater flow findings) in order to delineate extent of impacts to groundwater and address existing data gaps; and
- The installation and sampling of soil vapor monitoring points based on historical data and Phase A findings, including the collection and sampling of sub-slab and indoor air samples within the on-Site building (currently Subway restaurant).

All Phase B sample locations will be discussed with and approved by the NYSDEC/NYSDOH prior to completion.

#### 3.3.1 Groundwater Investigation

Based on Phase A groundwater flow findings, one upgradient monitoring well (RMW-3) will be installed to access upgradient groundwater conditions at the Site. RMW-3 will be installed and sampled using the same field procedures and laboratory methods used in Phase A of this RI. In addition, RMW-3 will be gauged and sampled following the procedures described in Section 3.2.3.

Following installation activities, an additional gauging event will be conducted and will include the collection of water-level measurements from all newly installed wells (RMW-1, RMW-2 and RMW-3) as well as existing monitoring wells (MW-1 through MW-3) to further define groundwater flow patterns beneath the Site.

#### 3.3.2 Soil Vapor Investigation

Based on historical data and Phase A findings, a soil vapor investigation will be conducted to assess both on-Site and off-Site soil vapor conditions. The data collected will be used to complete a qualitative exposure assessment. Soil vapor sampling will be conducted throughout the Site, based on historical data and Phase A findings. In addition, sub-slab and indoor air samples will be collected within the on-Site building to assess the current conditions within the building. Soil vapor samples will be collected with a Summa canister over a two-hour period using a regulator; the sub-slab and indoor air samples will be collected with a Summa canister over an eight-hour period using a regulator. All soil vapor, sub-slab and indoor are samples will be analyzed for VOCs via USEPA Method TO-15.

#### 3.4 Qualitative Exposure Assessment

A qualitative exposure assessment (EA) will be performed following the collection of all Phase A and Phase B and potential supplemental RI data. The EA will be performed in accordance with Section 3.3(c)4 of DER-10 and the NYSDOH guidance for performing a qualitative EA (NYSDEC DER-10; Technical Guidance for Site Investigation and Remediation; (Appendix 3 B). The results of the qualitative EA will be provided in the RI report.

#### 3.5 Fish and Wildlife Resource Impact Analysis

A Fish and Wildlife Resource Impact Analysis is not necessary due to the Site's location.

## 4.0 QUALITY ASSURANCE/QUALITY CONTROL (QA/QC) PROTOCOLS

The goal of QA/QC is to ensure that suitable and verifiable data results from sampling and analysis performed. To accomplish this, a Quality Assurance Project Plan (QAPP) has been prepared and is provided as Appendix B.

#### 5.0 HEALTH AND SAFETY

A site-specific Health and Safety Plan (HASP) and Community Air Monitoring Plan (CAMP) have been prepared for the Site and are provided in Appendix C. Decontamination plans and details can be found in Section 12.0 of the HASP (Appendix C).

The Property Owner and associated parties preparing the remedial documents submitted to the State, and parties performing this work are responsible for the safe performance of all invasive work, and for the integrity and safety of structures that may be affected by the intrusive activities (such as buildings, foundations and bridge footings). HAZWOPER training to on-Site workers and personnel will be provided as required for remedial investigation activities as appropriate. Copies of the 40 Hour OSHA course certificates for all on-Site personnel will be submitted to the NYSDEC and NYSDOH.

#### 6.0 REPORTING AND SCHEDULE

The following will be provided to the NYSDEC during the course of the RI work.

#### Progress Reports

Progress report submittals to be provided to NYSDEC will include the following:

- Daily Reports will be provided to the NYSDEC Project Manager during all periods of
  major investigative activity on remedial projects. These reports will include a summary of
  daily activities. These reports will also include a summary of substantive findings and
  other pertinent information including any complaints received from the public.
- Identification or lack of identification of the historical septic system during RI activities will be promptly communicated to the NYSDEC Project Manager.
- Monthly progress reports will be submitted to the NYSDEC Project Manager until the Certificate of Completion is issued.
- A Site map will be provided to identify locations discussed in progress reports provided to the NYSDEC.

#### Feasibility Study Work Plan (FSWP)

Following the completion of these phases of the RI and if a supplement RI is not warranted, a FSWP will be prepared and will include alternative remedies that will be evaluated to eliminate the Site's threat to public health or the environment. The FSWP will be submitted to the NYSDEC concurrently with the RI Report (described below) within two months of receipt of all final RI analytical data.

#### RI Report

Following the completion of RI activities a RI Report will be prepared within 2 months of receipt of all final RI analytical data and will include a description of the procedures followed and the results, including data summary tables and maps showing the extent of impacts to soil, groundwater and soil vapor. The RI Report will include all data developed during all RI activities, and will meet the technical requirements of NYSDEC DER-10; Technical Guidance for Site Investigation and Remediation. All RI analyses will be performed in accordance with the NYSDEC ASP, using USEPA SW-846 methods. The laboratory selected to analyze the field samples collected during the RI shall maintain a NYSDOH Environmental Laboratory Approval Program (ELAP) certification for each of the analyses listed in Section 3.0.

All laboratory data are to be reported in NYSDEC ASP Category B deliverables and will be delivered to NYSDEC in electronic data deliverable (EDD) format as described on NYSDEC's website (http://www.dec.ny.gov/chemical/62440.html). A Data Usability Report will be prepared meeting the requirements in Section 2.2(a)1.ii and Appendix 2B of DER-10 for all data packages generated for the RI.

Hazardous wastes and other contaminated media generated during RI phase will be stored, transported, and disposed in full compliance with applicable local, state, and federal regulations.

A table of the construction details for monitoring wells that have been built on-Site will be submitted. This will include date of construction, geologic interval screened, and current status (e.g., available for sampling, destroyed, intact, needs development, etc.).

The RI Report will include a conceptual site model that explains the occurrence of contaminant sources and their fate and transport at the Site in the context of the local Site stratigraphy and hydrogeology. The conceptual model will utilize both plan and cross-sectional views of the Site.

Copies of the transport manifests of hazardous and nonhazardous investigative waste will be provided to the NYSDEC Project Manager.

Following completion of the FSWP, a FS Report (FSR) will be prepared. The FSR will provide the basis for selection of a remedy that effectively eliminates the threat posed by contaminants at the Site, which will be used to prepare a Remedial Work Plan (RWP) for the Site.

Following completion of the FSR, a RWP will be prepared. The RWP will provide a detailed description of the remedial action and the remedial technology to be conducted for each area of concern.

SCHEDULE	
Reports/Work Plans	Date
Revised Remedial Investigation Work Plan Submittal	September 2015
Remedial Investigation Field Work	October 2015
Feasibility Study Work Plan Submittal	December 2015

SCHEDULE	
Reports/Work Plans	Date
Remedial Investigation Report Submittal	December 2015
Feasibility Study Report Submittal	February 2016
Remedial Work Plan Submittal	April 2016
Remedial Action	June 2016
Remedial Action Report Submittal	August 2016
Certificate of Completion Issuance	October 2016

## Remedial Investigation Work Plan 1022 Old Country Road, Plainview, New York 11803

**TABLES** 

- 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP
- 2. Summary of Volatile Organic Compounds in Groundwater Samples Collected by GFE, LBG and HRP
- 3. Proposed Soil Sampling Location Rationale
- 4. Proposed Groundwater Sample Location Rationale

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in µg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):		CL-N-SW <sup>3</sup> 4/16/2008 10		East SW <sup>3</sup> 4/16/2008 10	West SW <sup>3</sup> 4/16/2008 10	Bottom F/O <sup>3</sup> 4/16/2008 11	South SW F/O <sup>3</sup> 4/16/2008 10
1,1,1-Trichloroethane	680	500000	680		0.57 U	0.54 U	0.53 U	0.54 U	0.53 U	0.55 U	0.54 U
1,1,2,2-Tetrachloroethane					0.57 U 0.66 U	0.54 U	0.53 U 0.61 U	0.54 U	0.55 U 0.61 U	0.63 U	0.54 U
1.1.2-Trichloroethane					0.69 U	0.66 U	0.64 U	0.66 U	0.64 U	0.66 U	0.62 U
1.1-Dichloroethane	270	240000	270		0.63 U	0.59 U	0.54 U	0.59 U	0.54 U	0.60 U	0.49 U
1.1-Dichloroethene	330	500000	330		0.41 U	0.38 U	0.38 U	0.38 U	0.38 U	0.39 U	0.38 U
1.2.3-Trichlorobenzene					0.53 U	0.50 U	0.49 U	0.50 U	0.49 U	0.50 U	0.50 U
1.2.4-Trichlorobenzene					0.37 U	0.35 U	0.35 U	0.35 U	0.35 U	0.36 U	0.35 U
1.2-Dibromoethane					0.65 U	0.61 U	0.60 U	0.61 U	0.60 U	0.62 U	0.61 U
1,2-Dichlorobenzene	1100	500000	1100		0.52 U	0.49 U	0.48 U	0.49 U	0.48 U	0.49 U	0.49 U
1,2-Dichloroethane	20	30000	20		0.64 U	0.60 U	0.59 U	0.60 U	0.59 U	0.61 U	0.60 U
1,2-Dichloropropane					0.65 U	0.61 U	0.60 U	0.61 U	0.60 U	0.60 U	0.61 U
1,3-Dichlorobenzene	2400	280000	2400		0.58 U	0.55 U	0.54 U	0.55 U	0.54 U	0.56 U	0.55 U
1,4-Dichlorobenzene	1800	130000	1800		0.53 U	0.50 U	0.49 U	0.50 U	0.48 U	0.50 U	0.50 U
2-Butanone (MEK)	120	500000	120		2.44 U	2.31 U	2.26 U	2.31 U	2.26 U	2.33 U	2.31 U
2-Hexanone					2.18 U	2.06 U	2.02 U	2.06 U	2.02 U	2.08 U	2.06 U
4-Methyl-2-pentanone (MIBK)					2.37 U	2.24 U	2.19 U	2.24 U	2.19 U	2.26U	2.24 U
Acetone	50	500000	50		2.86 U	2.7 U	2.65 U	2.70 U	2.65 U	2.73U	2.70 U
Benzene	60	4400	60		0.58 U	0.55 U	0.54 U	0.55 U	0.54 U	0.56 U	0.55 U
Bromochloromethane					0.64 U	0.60 U	0.59 U	0.60 U	0.59 U	0.61 U	0.60 U
Bromodichloromethane					0.52 U	0.49 U	0.48 U	0.49 U	0.48 U	0.49 U	0.49 U
Bromoform					0.53 U	0.50 U	0.49 U	0.50 U	0.49 U	0.50 U	0.50 U
Bromomethane					0.54 U	0.51 U	0.50 U	0.51 U	0.50 U	0.51 U	0.51 U
Carbon disulfide					0.52 U	0.49 U	0.48 U	0.49 U	0.48 U	0.49 U	0.49 U
Carbon tetrachloride	760	2200	760		0.62 U	0.58 U	0.57 U	0.58 U	0.57 U	0.59 U	0.58 U
Chlorobenzene	1100	500000	1100		0.67 U	0.63 U	0.62 U	0.63 U	0.62 U	0.64 U	0.63 U
Chloroethane					0.77 U	0.73 U	0.71 U	0.73 U	0.71 U	0.74 U	0.73 U
Chloroform	370	350000	370		0.65 U	0.61 U	0.60 U	0.61 U	0.60 U	0.62 U	0.61 U
Chloromethane					0.55 U	0.52 U	0.51 U	0.52 U	0.51 U	0.52 U	0.52 U
cis-1,2-Dichloroethene	250	500000	250		0.50 U	0.47 U	0.46 U	0.47 U	0.46 U	0.47 U	0.47 U
cis-1,3-Dichloropropene					0.56 U	0.53 U	0.52 U	0.53 U	0.52 U	0.54 U	0.53 U
Cyclohexane					NA	NA	NA	NA	NA	NA	NA
Dibromochloromethane					0.51 U	0.48 U	0.47 U	0.48 U	0.47 U	0.48 U	0.48 U

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in μg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):		CL-N-SW <sup>3</sup> 4/16/2008 10		East SW <sup>3</sup> 4/16/2008 10	West SW <sup>3</sup> 4/16/2008 10	Bottom F/O <sup>3</sup> 4/16/2008 11	South SW F/O <sup>3</sup> 4/16/2008 10
Dibromochloropropane					NA	NA	NA	NA	NA	NA	NA
Dichlorodifluoromethane					0.41U	0.38 U	0.38 U	0.38 U	0.38 U	0.39 U	0.38 U
Ethylbenzene	1000	390000	1000		0.57 U	0.54 U	0.53 U	0.54 U	0.53 U	0.55 U	0.54 U
Freon 113					NA	NA	NA	NA	NA	NA	NA
Isopropylbenzene					0.48 U	0.46 U	0.45 U	0.46 U	0.45 U	0.46 U	0.46 U
m+p-Xylene					0.99 U	0.94 U	0.92 U	0.94 U	0.92 U	0.94 U	0.94 U
Methyl acetate					NA	NA	NA	NA	NA	NA	NA
Methylcyclohexane					NA	NA	NA	NA	NA	NA	NA
Methylene chloride	50	500000	50		1.03 U	0.98 U	0.96 U	0.98 U	0.96 U	0.99 U	0.98 U
MTBE	930	500000	930		0.57 U	0.54 U	0.53 U	0.54 U	0.53 U	0.55 U	0.54 U
o-Xylene					0.43 U	0.41 U	0.40 U	0.41 U	0.40 U	0.41 U	0.41 U
Styrene					0.47 U	0.45 U	0.44 U	0.45 U	0.44 U	0.45 U	0.45 U
Tetrachloroethene	1300	150000	1300		0.50 U	0.47 U	20.9	0.47 U	0.46 U	1.54 J	0.47 U
Toluene	700	500000	700		0.53 U	$0.70 \; J$	0.69 J	0.69 J	0.71 J	0.65 J	0.67 J
trans-1,2-Dichloroethene	190	500000	190		0.51 U	0.48 U	0.47 U	0.48 U	0.47 U	0.48 U	0.48 U
trans-1,3-Dichloropropene					0.46 U	0.44 U	0.43 U	0.44 U	0.43 U	0.44 U	0.44 U
Trichloroethene	470	200000	470		0.54 U	0.51 U	0.50 U	0.51 U	0.50 U	0.51 U	0.51 U
Trichlorofluoromethane					0.62 U	0.58 U	0.57 U	0.58 U	0.57 U	0.59 U	0.58 U
Vinyl chloride	20	1300	20		0.75 U	0.7 1U	0.69 U	0.71 U	0.69 U	0.71 U	0.71 U
Xylenes (total)	260	500000	1600		0.43 U	0.41 U	0.40 U	0.41 U	0.40 U	0.41 U	0.41 U

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

μg/kg - Micrograms per kilogram

NYSDEC - New York State Department of Environmental Conservation

Bold data indicates that parameter was detected above the NYSDEC Part 375 Unrestricted Use Standards

Shaded data indicates that parameter was detected above the NYSDEC Part 375 Commercial Use Standards

Boxed data indicates that parameter was detected above the NYSDEC Part 375 Protection of Groundwater Standards

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from LBG 2008 UST Closure Report

<sup>&</sup>lt;sup>4</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated value

U - Indicates that the compound was analyzed for but not detected

ft bg - Feet below grade

<sup>\*</sup>Reporting Limit was not provided in the associated report

NA - Not available in associated report

<sup>--</sup> No NYSDEC Part 375 Standards available

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in µg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):	North SW F/O <sup>3</sup> 4/16/2008 10	(Base) <sup>3</sup>	SB-5 <sup>4</sup> 11/12/2010 79-81	SB-5 <sup>4</sup> 11/12/2010 89-91	SB-5 <sup>4</sup> 11/12/2010 99-101	SB-5 RE <sup>4</sup> 11/12/2010 99-101
1 1 1 Tricklandshan	<b>C</b> 00	500000	<b>C</b> 00		0.60 H	0.54.11	£ 0.11	6 2 H	( ) II	6 <b>2</b> H
1,1,1-Trichloroethane 1,1,2,2-Tetrachloroethane	680	500000	680		0.60 U 0.70 U	0.54 U 0.62 U	5.9 U 5.9 U	6.2 U 6.2 U	6.2 U 6.2 U	6.2 U 6.2 U
1.1.2-Trichloroethane					0.70 U 0.73 U	0.62 U 0.66 U	5.9 U	6.2 U	6.2 U	6.2 U
1,1-Dichloroethane	270	240000	270		0.73 U 0.66 U	0.66 U 0.59 U	5.9 U	6.2 U	6.2 U	6.2 U
1,1-Dichloroethene					0.66 U 0.43 U		5.9 U			6.2 U
	330	500000	330		0.43 U 0.56 U	0.38 U	5.9 U 5.9 U	6.2 U 6.2 U	6.2 U	
1,2,3-Trichlorobenzene					0.36 U 0.39 U	0.50 U	5.9 U 5.9 U	6.2 U 6.2 U	6.2 U	6.2 U 6.2 U
1,2,4-Trichlorobenzene						0.35 U			6.2 U	
1,2-Dibromoethane	1100	 500000	1100		0.68 U 0.55 U	0.61 U	5.9 U 5.9 U	6.2 U	6.2 U	6.2 U 6.2 U
1,2-Dichlorobenzene	1100	500000	1100			0.49 U		6.2 U	6.2 U	
1,2-Dichloroethane	20	30000	20		0.67 U	0.60 U	5.9 U	6.2 U	6.2 U	6.2 U
1,2-Dichloropropane	2400	200000	2400		0.68 U	0.61 U	5.9 U	6.2 U	6.2 U	6.2 U
1,3-Dichlorobenzene	2400	280000	2400		0.61 U	0.55 U	5.9 U	6.2 U	6.2 U	6.2 U
1,4-Dichlorobenzene	1800	130000	1800		0.56 U	0.50 U	5.9 U	6.2 U	6.2 U	6.2 U
2-Butanone (MEK)	120	500000	120		2.58 U	2.31 U	29 U	31 U	31 U	31 U
2-Hexanone					2.30 U	2.06 U	29 U	31 U	31 U	31 U
4-Methyl-2-pentanone (MIBK)					2.49 U	2.24 U	29 U	31 U	31 U	31 U
Acetone	50	500000	50		3.02 U	2.70 U	29 U	31 U	31 U	31 U
Benzene	60	4400	60		0.61 U	0.55 U	5.9 U	6.2 U	6.2 U	6.2 U
Bromochloromethane					0.67 U	0.60 U	5.9 U	6.2 U	6.2 U	6.2 U
Bromodichloromethane					0.55 U	0.49 U	5.9 U	6.2 U	6.2 U	6.2 U
Bromoform					0.56 U	0.50 U	5.9 U	6.2 U	6.2 U	6.2 U
Bromomethane					0.57 U	0.51 U	5.9 U	6.2 U	6.2 U	6.2 U
Carbon disulfide					0.55 U	0.49 U	5.9 U	6.2 U	6.2 U	6.2 U
Carbon tetrachloride	760	2200	760		0.65 U	0.58 U	5.9 U	6.2 U	6.2 U	6.2 U
Chlorobenzene	1100	500000	1100		0.71 U	0.63 U	5.9 U	6.2 U	6.2 U	6.2 U
Chloroethane					0.81 U	0.73 U	5.9 U	6.2 U	6.2 U	6.2 U
Chloroform	370	350000	370		0.68 U	0.61 U	5.9 U	6.2 U	6.2 U	6.2 U
Chloromethane					0.58 U	0.52 U	5.9 U	6.2 U	6.2 U	6.2 U
cis-1,2-Dichloroethene	250	500000	250		0.52 U	0.47 U	5.9 U	6.2 U	6.2 U	6.2 U
cis-1,3-Dichloropropene					0.59 U	0.53 U	5.9 U	6.2 U	6.2 U	6.2 U
Cyclohexane					NA	NA	5.9 U	6.2 U	6.2 U	6.2 U
Dibromochloromethane					0.53 U	0.48 U	5.9 U	6.2 U	6.2 U	6.2 U

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in μg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):	4/16/2008	(Base) <sup>3</sup>	SB-5 <sup>4</sup> 11/12/2010 79-81	SB-5 <sup>4</sup> 11/12/2010 89-91	SB-5 <sup>4</sup> 11/12/2010 99-101	SB-5 RE <sup>4</sup> 11/12/2010 99-101
Dibromochloropropane					NA	NA	5.9 U	6.2 U	6.2 U	6.2 U
Dichlorodifluoromethane					0.43 U	0.38 U	5.9 U	6.2 U	6.2 U	6.2 U
Ethylbenzene	1000	390000	1000		0.60 U	0.54 U	5.9 U	6.2 U	6.2 U	6.2 U
Freon 113					NA	NA	NA	NA	NA	NA
Isopropylbenzene					0.51 U	0.46 U	5.9 U	6.2 U	6.2 U	6.2 U
m+p-Xylene					1.04 U	0.94 U	12 U	12 U	12 U	12 U
Methyl acetate					NA	NA	5.9 U	6.2 U	6.2 U	6.2 U
Methylcyclohexane					NA	NA	5.9 U	6.2 U	6.2 U	6.2 U
Methylene chloride	50	500000	50		1.09 U	0.98 U	1.8 J	6.2 U	2.4 J	6.2 U
MTBE	930	500000	930		0.60 U	0.54 U	5.9 U	6.2 U	6.2 U	6.2 U
o-Xylene					0.45 U	0.41 U	5.9 U	6.2 U	6.2 U	6.2 U
Styrene					0.50 U	0.45 U	5.9 U	6.2 U	6.2 U	6.2 U
Tetrachloroethene	1300	150000	1300		166	1.04 J	5.9 U	6.2 U	6.2 U	6.2 U
Toluene	700	500000	700		0.85 J	0.50 U	5.9 U	6.2 U	6.2 U	6.2 U
trans-1,2-Dichloroethene	190	500000	190		0.53 U	0.48 U	5.9 U	6.2 U	6.2 U	6.2 U
trans-1,3-Dichloropropene					0.49 U	0.55 U	5.9 U	6.2 U	6.2 U	6.2 U
Trichloroethene	470	200000	470		2.78 J	0.51 U	5.9 U	6.2 U	6.2 U	6.2 U
Trichlorofluoromethane					0.65 U	0.58 U	5.9 U	6.2 U	6.2 U	6.2 U
Vinyl chloride	20	1300	20		0.79 U	0.71 U	5.9 U	6.2 U	6.2 U	6.2 U
Xylenes (total)	260	500000	1600		0.45 U	0.41 U	5.9 U	6.2 U	6.2 U	6.2 U

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

μg/kg - Micrograms per kilogram

NYSDEC - New York State Department of Environmental Conservation

Bold data indicates that parameter was detected above the NYSDEC Part 375 Unrestricted Use Standards

Shaded data indicates that parameter was detected above the NYSDEC Part 375 Commercial Use Standards

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from LBG 2008 UST Closure Report

<sup>&</sup>lt;sup>4</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated value

U - Indicates that the compound was analyzed for but not detected

ft bg - Feet below grade

<sup>\*</sup>Reporting Limit was not provided in the associated report

NA - Not available in associated report

<sup>--</sup> No NYSDEC Part 375 Standards available

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in μg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):	SB-5 <sup>4</sup> 11/12/2010 109-111	SB-5 <sup>4</sup> 11/12/2010 119-121	SB-1 <sup>4</sup> 11/8/2010 79-81	SB-1 <sup>4</sup> 11/8/2010 89-91	SB-1 <sup>4</sup> 11/8/2010 99-101	SB-1 <sup>4</sup> 11/8/2010 109-111
1,1,1-Trichloroethane	680	500000	680		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,1,2,2-Tetrachloroethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,1,2-Trichloroethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,1-Dichloroethane	270	240000	270		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,1-Dichloroethene	330	500000	330		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,2,3-Trichlorobenzene					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,2,4-Trichlorobenzene					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,2-Dibromoethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,2-Dichlorobenzene	1100	500000	1100		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,2-Dichloroethane	20	30000	20		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,2-Dichloropropane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,3-Dichlorobenzene	2400	280000	2400		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
1,4-Dichlorobenzene	1800	130000	1800		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
2-Butanone (MEK)	120	500000	120		31 U	30 U	29 U	31 U	30 U	31 U
2-Hexanone					31 U	30 U	29 U	31 U	30 U	31 U
4-Methyl-2-pentanone (MIBK)					31 U	30 U	29 U	31 U	30 U	31 U
Acetone	50	500000	50		31 U	30 U	29 U	31 U	30 U	31 U
Benzene	60	4400	60		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Bromochloromethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Bromodichloromethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Bromoform					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Bromomethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Carbon disulfide					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Carbon tetrachloride	760	2200	760		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Chlorobenzene	1100	500000	1100		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Chloroethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Chloroform	370	350000	370		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Chloromethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
cis-1,2-Dichloroethene	250	500000	250		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
cis-1,3-Dichloropropene					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Cyclohexane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Dibromochloromethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in µg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):	SB-5 <sup>4</sup> 11/12/2010 109-111	SB-5 <sup>4</sup> 11/12/2010 119-121	SB-1 <sup>4</sup> 11/8/2010 79-81	SB-1 <sup>4</sup> 11/8/2010 89-91	SB-1 <sup>4</sup> 11/8/2010 99-101	SB-1 <sup>4</sup> 11/8/2010 109-111
Dibromochloropropane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Dichlorodifluoromethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Ethylbenzene	1000	390000	1000		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Freon 113					NA	NA	NA	NA	NA	NA
Isopropylbenzene					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
m+p-Xylene					12 U	12 U	12 U	12 U	12 U	12 U
Methyl acetate					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Methylcyclohexane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Methylene chloride	50	500000	50		6.1 U	2.2 J	3.2 J	4.1 J	6.0 U	4.0 J
MTBE	930	500000	930		4.5 J	5.6 J	5.9 U	6.1 U	6.0 U	6.1 U
o-Xylene					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Styrene					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Tetrachloroethene	1300	150000	1300		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Toluene	700	500000	700		1.4 J	1.1 J	1.2 J	1.8 J	1.7 J	6.1 U
trans-1,2-Dichloroethene	190	500000	190		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
trans-1,3-Dichloropropene					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Trichloroethene	470	200000	470		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Trichlorofluoromethane					6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Vinyl chloride	20	1300	20		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U
Xylenes (total)	260	500000	1600		6.1 U	6.0 U	5.9 U	6.1 U	6.0 U	6.1 U

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

μg/kg - Micrograms per kilogram

NYSDEC - New York State Department of Environmental Conservation

Bold data indicates that parameter was detected above the NYSDEC Part 375 Unrestricted Use Standards

Shaded data indicates that parameter was detected above the NYSDEC Part 375 Commercial Use Standards

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from LBG 2008 UST Closure Report

<sup>&</sup>lt;sup>4</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated value

U - Indicates that the compound was analyzed for but not detected

ft bg - Feet below grade

<sup>\*</sup>Reporting Limit was not provided in the associated report

NA - Not available in associated report

<sup>--</sup> No NYSDEC Part 375 Standards available

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in μg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):		SB-2 <sup>4</sup> 11/9/2010 79-81	SB-2 <sup>4</sup> 11/9/2010 89-91	SB-2 <sup>4</sup> 11/9/2010 99-101	SB-2 <sup>4</sup> 11/9/2010 114-116	SB-3 <sup>4</sup> 11/10/2010 79-81
1,1,1-Trichloroethane	680	500000	680		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,1,2,2-Tetrachloroethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1.1.2-Trichloroethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1.1-Dichloroethane	270	240000	270		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1.1-Dichloroethene	330	500000	330		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,2,3-Trichlorobenzene					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,2,4-Trichlorobenzene			<u></u>		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1.2-Dibromoethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,2-Dichlorobenzene	1100	500000	1100		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,2-Dichloroethane	20	30000	20		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,2-Dichloropropane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,3-Dichlorobenzene	2400	280000	2400		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
1,4-Dichlorobenzene	1800	130000	1800		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
2-Butanone (MEK)	120	500000	120		31 U	31 U	31 U	30 U	30 U	30 U
2-Hexanone					31 U	31 U	31 U	30 U	30 U	30 U
4-Methyl-2-pentanone (MIBK)					31 U	31 U	31 U	30 U	30 U	30 U
Acetone	50	500000	50		31 U	31 U	31 U	30 U	30 U	30 U
Benzene	60	4400	60		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Bromochloromethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Bromodichloromethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Bromoform					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Bromomethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Carbon disulfide					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Carbon tetrachloride	760	2200	760		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Chlorobenzene	1100	500000	1100		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Chloroethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Chloroform	370	350000	370		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Chloromethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
cis-1,2-Dichloroethene	250	500000	250		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
cis-1,3-Dichloropropene					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Cyclohexane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Dibromochloromethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

	NYSDEC	NYSDEC	NYSDEC					4		
	Part 375	Part 375	Part 375	Sample Designation:	$SB-1^4$	$SB-2^4$	$SB-2^4$	$SB-2^4$	$SB-2^4$	$SB-3^4$
Parameter	Unrestricted	Commercial	Protection of	Sample Date:	11/8/2010	11/9/2010	11/9/2010	11/9/2010	11/9/2010	11/10/2010
(Concentrations in µg/kg)	Use	Use	Groundwater	Sample Depth (ft bg):	119-121	79-81	89-91	99-101	114-116	79-81
										_
Dibromochloropropane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Dichlorodifluoromethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Ethylbenzene	1000	390000	1000		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Freon 113					NA	NA	NA	NA	NA	NA
Isopropylbenzene					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
m+p-Xylene					12 U					
Methyl acetate					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Methylcyclohexane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Methylene chloride	50	500000	50		3.1 J	6.1 U	2.2 J	2.4 J	1.9 J	6
MTBE	930	500000	930		6.1 U	6.1 U	6.2 U	6.0 U	2.7 J	5.9 U
o-Xylene					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Styrene					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Tetrachloroethene	1300	150000	1300		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	1.6 J
Toluene	700	500000	700		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
trans-1,2-Dichloroethene	190	500000	190		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
trans-1,3-Dichloropropene					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Trichloroethene	470	200000	470		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Trichlorofluoromethane					6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Vinyl chloride	20	1300	20		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
Xylenes (total)	260	500000	1600		6.1 U	6.1 U	6.2 U	6.0 U	6.0 U	5.9 U
/	200	223000			2.2 0	2.2 0	J 0	2.2 0	0	2.5

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

Bold data indicates that parameter was detected above the NYSDEC Part 375 Unrestricted Use Standards

Shaded data indicates that parameter was detected above the NYSDEC Part 375 Commercial Use Standards

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from LBG 2008 UST Closure Report

<sup>&</sup>lt;sup>4</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated value

U - Indicates that the compound was analyzed for but not detected

μg/kg - Micrograms per kilogram

ft bg - Feet below grade

<sup>\*</sup>Reporting Limit was not provided in the associated report

NA - Not available in associated report

<sup>--</sup> No NYSDEC Part 375 Standards available

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in μg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):	SB-3 <sup>4</sup> 11/10/2010 89-91	SB-3 <sup>4</sup> 11/10/2010 99-101	SB-3 <sup>4</sup> 11/10/2010 109-111	SB-3 <sup>4</sup> 11/10/2010 119-121	SB-4 <sup>4</sup> 11/11/2010 79-81	Duplicate <sup>4</sup> 11/11/2010
1,1,1-Trichloroethane	680	500000	680		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,1,2,2-Tetrachloroethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,1,2-Trichloroethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1.1-Dichloroethane	270	240000	270		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1.1-Dichloroethene	330	500000	330		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1.2.3-Trichlorobenzene					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,2,4-Trichlorobenzene					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1.2-Dibromoethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,2-Dichlorobenzene	1100	500000	1100		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,2-Dichloroethane	20	30000	20		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,2-Dichloropropane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,3-Dichlorobenzene	2400	280000	2400		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
1,4-Dichlorobenzene	1800	130000	1800		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
2-Butanone (MEK)	120	500000	120		31 U	30 U	30 U	30 U	30 U	30 U
2-Hexanone					31 U	30 U	30 U	30 U	30 U	30 U
4-Methyl-2-pentanone (MIBK)					31 U	30 U	30 U	30 U	30 U	30 U
Acetone	50	500000	50		31 U	30 U	30 U	30 U	30 U	30 U
Benzene	60	4400	60		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Bromochloromethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Bromodichloromethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Bromoform					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Bromomethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Carbon disulfide					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Carbon tetrachloride	760	2200	760		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Chlorobenzene	1100	500000	1100		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Chloroethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Chloroform	370	350000	370		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Chloromethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
cis-1,2-Dichloroethene	250	500000	250		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
cis-1,3-Dichloropropene					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Cyclohexane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Dibromochloromethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in μg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):	SB-3 <sup>4</sup> 11/10/2010 89-91	SB-3 <sup>4</sup> 11/10/2010 99-101	SB-3 <sup>4</sup> 11/10/2010 109-111	SB-3 <sup>4</sup> 11/10/2010 119-121	SB-4 <sup>4</sup> 11/11/2010 79-81	Duplicate <sup>4</sup> 11/11/2010
Dibromochloropropane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Dichlorodifluoromethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Ethylbenzene	1000	390000	1000		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Freon 113					NA	NA	NA	NA	NA	NA
Isopropylbenzene					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
m+p-Xylene					12 U	12 U	12 U	12 U	12 U	12 U
Methyl acetate					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Methylcyclohexane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Methylene chloride	50	500000	50		2 J	2.8 J	2.2 J	2.5 J	2.3 J	2.0 J
MTBE	930	500000	930		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
o-Xylene					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Styrene					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Tetrachloroethene	1300	150000	1300		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Toluene	700	500000	700		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
trans-1,2-Dichloroethene	190	500000	190		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
trans-1,3-Dichloropropene					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Trichloroethene	470	200000	470		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Trichlorofluoromethane					6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Vinyl chloride	20	1300	20		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U
Xylenes (total)	260	500000	1600		6.2 U	6.0 U	5.9 U	6.1 U	5.9 U	6.1 U

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

μg/kg - Micrograms per kilogram

NYSDEC - New York State Department of Environmental Conservation

Bold data indicates that parameter was detected above the NYSDEC Part 375 Unrestricted Use Standards

Shaded data indicates that parameter was detected above the NYSDEC Part 375 Commercial Use Standards

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from LBG 2008 UST Closure Report

<sup>&</sup>lt;sup>4</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated value

U - Indicates that the compound was analyzed for but not detected

ft bg - Feet below grade

<sup>\*</sup>Reporting Limit was not provided in the associated report

NA - Not available in associated report

<sup>--</sup> No NYSDEC Part 375 Standards available

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

Parameter (Concentrations in μg/kg)	NYSDEC Part 375 Unrestricted Use	NYSDEC Part 375 Commercial Use	NYSDEC Part 375 Protection of Groundwater	Sample Designation: Sample Date: Sample Depth (ft bg):	SB-4 <sup>4</sup> 11/11/2010 89-91	SB-4 <sup>4</sup> 11/11/2010 99-101	SB-4 <sup>4</sup> 11/11/2010 109-111	SB-4 <sup>4</sup> 11/11/2010 119-121
1,1,1-Trichloroethane	680	500000	680		6.3 U	6.4 U	6.2 U	6.1 U
1,1,2,2-Tetrachloroethane					6.3 U	6.4 U	6.2 U	6.1 U
1,1,2-Trichloroethane					6.3 U	6.4 U	6.2 U	6.1 U
1,1-Dichloroethane	270	240000	270		6.3 U	6.4 U	6.2 U	6.1 U
1,1-Dichloroethene	330	500000	330		6.3 U	6.4 U	6.2 U	6.1 U
1,2,3-Trichlorobenzene					6.3 U	6.4 U	6.2 U	6.1 U
1,2,4-Trichlorobenzene					6.3 U	6.4 U	6.2 U	6.1 U
1,2-Dibromoethane					6.3 U	6.4 U	6.2 U	6.1 U
1,2-Dichlorobenzene	1100	500000	1100		6.3 U	6.4 U	6.2 U	6.1 U
1,2-Dichloroethane	20	30000	20		6.3 U	6.4 U	6.2 U	6.1 U
1,2-Dichloropropane					6.3 U	6.4 U	6.2 U	6.1 U
1,3-Dichlorobenzene	2400	280000	2400		6.3 U	6.4 U	6.2 U	6.1 U
1,4-Dichlorobenzene	1800	130000	1800		6.3 U	6.4 U	6.2 U	6.1 U
2-Butanone (MEK)	120	500000	120		31 U	32 U	31 U	31 U
2-Hexanone					31 U	32 U	31 U	31 U
4-Methyl-2-pentanone (MIBK)					31 U	32 U	31 U	31 U
Acetone	50	500000	50		31 U	32 U	31 U	31 U
Benzene	60	4400	60		6.3 U	6.4 U	6.2 U	6.1 U
Bromochloromethane					6.3 U	6.4 U	6.2 U	6.1 U
Bromodichloromethane					6.3 U	6.4 U	6.2 U	6.1 U
Bromoform					6.3 U	6.4 U	6.2 U	6.1 U
Bromomethane					6.3 U	6.4 U	6.2 U	6.1 U
Carbon disulfide					6.3 U	6.4 U	6.2 U	6.1 U
Carbon tetrachloride	760	2200	760		6.3 U	6.4 U	6.2 U	6.1 U
Chlorobenzene	1100	500000	1100		6.3 U	6.4 U	6.2 U	6.1 U
Chloroethane					6.3 U	6.4 U	6.2 U	6.1 U
Chloroform	370	350000	370		6.3 U	6.4 U	6.2 U	6.1 U
Chloromethane					6.3 U	6.4 U	6.2 U	6.1 U
cis-1,2-Dichloroethene	250	500000	250		6.3 U	6.4 U	6.2 U	6.1 U
cis-1,3-Dichloropropene					6.3 U	6.4 U	6.2 U	6.1 U
Cyclohexane					6.3 U	6.4 U	6.2 U	6.1 U
Dibromochloromethane					6.3 U	6.4 U	6.2 U	6.1 U

Table 1. Summary of Volatile Organic Compounds in Soil Samples Collected by GFE, LBG and HRP, 1022 Old Country Road, Plainview, New York

	NYSDEC	NYSDEC	NYSDEC					
	Part 375	Part 375	Part 375	Sample Designation:	$SB-4^4$	$SB-4^4$	$SB-4^4$	$SB-4^4$
Parameter	Unrestricted	Commercial	Protection of	Sample Date:	11/11/2010	11/11/2010	11/11/2010	11/11/2010
(Concentrations in μg/kg)	Use	Use	Groundwater	Sample Depth (ft bg):	89-91	99-101	109-111	119-121
Dibromochloropropane					6.3 U	6.4 U	6.2 U	6.1 U
Dichlorodifluoromethane					6.3 U	6.4 U	6.2 U	6.1 U
Ethylbenzene	1000	390000	1000		6.3 U	6.4 U	6.2 U	6.1 U
Freon 113					NA	NA	NA	NA
Isopropylbenzene					6.3 U	6.4 U	6.2 U	6.1 U
m+p-Xylene					12 U	12 U	12 U	12 U
Methyl acetate					6.3 U	6.4 U	6.2 U	6.1 U
Methylcyclohexane					6.3 U	6.4 U	6.2 U	6.1 U
Methylene chloride	50	500000	50		2.6 J	3.2 J	3.1 J	3.1 J
MTBE	930	500000	930		6.3 U	6.4 U	6.2 U	6.1 U
o-Xylene					6.3 U	6.4 U	6.2 U	6.1 U
Styrene					6.3 U	6.4 U	6.2 U	6.1 U
Tetrachloroethene	1300	150000	1300		6.3 U	6.4 U	6.2 U	6.1 U
Toluene	700	500000	700		6.3 U	6.4 U	6.2 U	6.1 U
trans-1,2-Dichloroethene	190	500000	190		6.3 U	6.4 U	6.2 U	6.1 U
trans-1,3-Dichloropropene					6.3 U	6.4 U	6.2 U	6.1 U
Trichloroethene	470	200000	470		6.3 U	6.4 U	6.2 U	6.1 U
Trichlorofluoromethane					6.3 U	6.4 U	6.2 U	6.1 U
Vinyl chloride	20	1300	20		6.3 U	6.4 U	6.2 U	6.1 U
Xylenes (total)	260	500000	1600		6.3 U	6.4 U	6.2 U	6.1 U
•								

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

Bold data indicates that parameter was detected above the NYSDEC Part 375 Unrestricted Use Standards

Shaded data indicates that parameter was detected above the NYSDEC Part 375 Commercial Use Standards

Boxed data indicates that parameter was detected above the NYSDEC Part 375 Protection of Groundwater Standards

2517.0001Y.101/WKB

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from LBG 2008 UST Closure Report

<sup>&</sup>lt;sup>4</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated value

U - Indicates that the compound was analyzed for but not detected

μg/kg - Micrograms per kilogram

ft bg - Feet below grade

<sup>\*</sup>Reporting Limit was not provided in the associated report

NA - Not available in associated report

<sup>--</sup> No NYSDEC Part 375 Standards available

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

Parameter	NYSDEC AWQSGVs	Sample Designation: Sample Date:		B2GW <sup>1</sup>	MW-1 <sup>2</sup> 6/1/2007	MW-2 <sup>2</sup> 6/1/2007	MW-3 <sup>2</sup> 6/1/2007	MW-1 <sup>3</sup> 11/16/2010	MW-2 <sup>3</sup> 11/16/2010	MW-3 <sup>3</sup> 11/16/2010
(Concentrations in µg/L)	μg/L)	Sample Date.	<i>3</i> /20/2000	7/20/2000	0/1/2007	0/1/2007	0/1/2007	11/10/2010	11/10/2010	11/10/2010
1,1,1-Trichloroethane	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,1,2,2-Tetrachloroethane	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,1,2-Trichloroethane	1		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethane	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethene	5		NA	NA	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,3-Trichlorobenzene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,2,4-Trichlorobenzene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,2-Dibromoethane			NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichlorobenzene	3		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloroethane	0.6		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloropropane	1		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,3-Dichlorobenzene	3		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
1,4-Dichlorobenzene	3		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
2-Butanone (MEK)	50		NA	NA	10 U	10 U	11 U	5.0 U	5.0 U	5.0 U
2-Hexanone	50		NA	NA	2.0 U	2.0 U	2.0 U	5.0 U	5.0 U	5.0 U
4-Methyl-2-pentanone (MIBK)			NA	NA	10 U	10 U	10 U	5.0 U	5.0 U	5.0 U
Acetone	50		NA	NA	10 U	10 U	10 U	5.0 U	5.0 U	5.0 U
Benzene	1		NA	NA	1.0U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromochloromethane	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Bromodichloromethane	50		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Bromoform	50		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Bromomethane	5		NA	NA	NA	NA	NA	1.0 U	1.0 U	1.0 U
Carbon disulfide	60		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Carbon tetrachloride	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Chlorobenzene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Chloroethane	5		NA	NA	5.0 U	5.0 U	5.0 U	1.0 U	1.0 U	1.0 U
Chloroform	7		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Chloromethane			NA	NA	5.0 U	5.0 U	5.0 U	1.0 U	1.0 U	1.0 U
cis-1,2-Dichloroethene	5		5.09	ND*	23	4.8	7.4	1.3	0.95 J	43
cis-1,3-Dichloropropene	5		NA	NA	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Cyclohexane			NA	NA	NA	NA	NA	1.0 U	1.0 U	1.0 U
Dibromochloromethane	50		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Dibromochloropropane			NA	NA	5.0 U	5.0 U	5.0 U	NA	NA	NA

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

Parameter	NYSDEC AWQSGVs	Sample Designation: Sample Date:		B2GW <sup>1</sup> 9/26/2006	MW-1 <sup>2</sup> 6/1/2007	MW-2 <sup>2</sup> 6/1/2007	MW-3 <sup>2</sup> 6/1/2007	MW-1 <sup>3</sup> 11/16/2010	MW-2 <sup>3</sup> 11/16/2010	MW-3 <sup>3</sup> 11/16/2010
(Concentrations in µg/L)	$(\mu g/L)$									
Dichlorodifluoromethane	5		NA	NA	5.0 U	5.0 U	5.0 U	1.0 U	1.0 U	1.0 U
Ethylbenzene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Freon 113			NA	NA	NA	NA	NA	NA	NA	NA
Isopropylbenzene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
m+p-Xylene	5		NA	NA	2.0 U	2.0 U	2.0 U	2.0 U	2.0 U	2.0 U
Methyl acetate			NA	NA	NA	NA	NA	1.0 U	1.0 U	1.0 U
Methylcyclohexane			NA	NA	NA	NA	NA	1.0 U	1.0 U	1.0 U
Methylene chloride	5		NA	NA	5.0 U	5.0 U	5.0 U	1.0 U	1.0 U	1.0 U
MTBE	10		NA	NA	2.0 U	2.2	2. 0U	0.92 J	1.4	1.0 U
o-Xylene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Styrene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Tetrachloroethene	5		11.9	546	120	72	93	25	24	88
Toluene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
trans-1,2-Dichloroethene	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.7
trans-1,3-Dichloropropene			NA	NA	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Trichloroethene	5		ND*	9.97	16	6.9	9.4	2.3	2.4	26
Trichlorofluoromethane	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Vinyl chloride	2		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U
Xylenes (total)	5		NA	NA	2.0 U	2.0 U	2.0 U	1.0 U	1.0 U	1.0 U

AWQSGVs - Ambient Water-Quality Standards and Guidance Values

μg/L -Micrograms per liter

Bold data indicates that parameter was detected above the NYSDEC AWQSGVs

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated Value

U - Compound was analyzed for but not detected

<sup>\*</sup>Reporting Limit was not provided in the associated report

<sup>- -</sup> No NYSDEC AWQSGV available

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation:	GW-5 (91-91.5) <sup>3</sup>	GW-5 (101-101.5) <sup>3</sup>	GW-5 (111-111.5) <sup>3</sup>	<sup>3</sup> GW-5 (121-121.5) <sup>2</sup>	<sup>E</sup> GW-1 (91-91.5) <sup>3</sup>
Parameter	AWQSGVs	Sample Date:	11/12/2010	11/12/2010	11/12/2010	11/12/2010	11/8/2010
(Concentrations in µg/L)	(µg/L)						
1,1,1-Trichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2,2-Tetrachloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2-Trichloroethane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,3-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,4-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dibromoethane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloroethane	0.6		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloropropane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,3-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,4-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
2-Butanone (MEK)	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
2-Hexanone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
4-Methyl-2-pentanone (MIBK)			5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Acetone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Benzene	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromochloromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromodichloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromoform	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromomethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon disulfide	60		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon tetrachloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroethane	5		1.0 U	1.0 U	1.3	0.9 J	1.0 U
Chloroform	7		1.0 U	1.0 U	1.0 U	1.0 U	0.53 J
Chloromethane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
cis-1,2-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
cis-1,3-Dichloropropene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Cyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloropropane			NA	NA	NA	NA	NA

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: (	GW-5 (91-91.5) <sup>3</sup>	GW-5 (101-101.5) <sup>3</sup>	GW-5 (111-111.5) <sup>3</sup>	GW-5 (121-121.5) <sup>3</sup>	GW-1 (91-91.5) <sup>3</sup>
Parameter	AWQSGVs	Sample Date:	11/12/2010	11/12/2010	11/12/2010	11/12/2010	11/8/2010
(Concentrations in µg/L)	$(\mu g/L)$	_					
Dichlorodifluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Ethylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Freon 113			NA	NA	NA	NA	NA
Isopropylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
m+p-Xylene	5		2.0 U	2.0 U	2.0 U	2.0 U	2.0 U
Methyl acetate			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylcyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylene chloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
MTBE	10		1.0 U	1.0 U	1.0 U	1.0 U	6.2
o-Xylene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Styrene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Tetrachloroethene	5		2.7	3.3	0.96 J	1.0 U	2.1
Toluene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
trans-1,2-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
trans-1,3-Dichloropropene			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Trichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Trichlorofluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Vinyl chloride	2		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Xylenes (total)	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U

AWQSGVs - Ambient Water-Quality Standards and Guidance Values  $\mu g/L$  -Micrograms per liter

Bold data indicates that parameter was detected above the NYSDEC AWQSGVs

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated Value

U - Compound was analyzed for but not detected

<sup>\*</sup>Reporting Limit was not provided in the associated report

<sup>- -</sup> No NYSDEC AWQSGV available

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: G	W-1 (101-101.5) <sup>3</sup>	<sup>3</sup> GW-1 (111-111.5) <sup>3</sup>	GW-1 (121-121.5)	$^{3}$ GW-2 $(81-81.5)^{3}$	$GW-2 (91-91.5)^3$
Parameter	AWQSGVs	Sample Date:	11/8/2010	11/8/2010	11/8/2010	11/9/2010	11/9/2010
(Concentrations in µg/L)	(µg/L)						
1,1,1-Trichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2,2-Tetrachloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2-Trichloroethane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,3-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,4-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dibromoethane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloroethane	0.6		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloropropane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,3-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,4-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
2-Butanone (MEK)	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
2-Hexanone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
4-Methyl-2-pentanone (MIBK)			5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Acetone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Benzene	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromochloromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromodichloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromoform	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromomethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon disulfide	60		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon tetrachloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroform	7		1.0U	1.1	1.2	1.0 U	1.0 U
Chloromethane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
cis-1,2-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
cis-1,3-Dichloropropene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Cyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloropropane			NA	NA	NA	NA	NA

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: G	W-1 (101-101.5) <sup>3</sup>	GW-1 (111-111.5) <sup>3</sup>	GW-1 (121-121.5)	<sup>3</sup> GW-2 (81-81.5) <sup>3</sup>	$GW-2 (91-91.5)^3$
Parameter	AWQSGVs	Sample Date:	11/8/2010	11/8/2010	11/8/2010	11/9/2010	11/9/2010
(Concentrations in µg/L)	$(\mu g/L)$						
Dichlorodifluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Ethylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Freon 113			NA	NA	NA	NA	NA
Isopropylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
m+p-Xylene	5		2.0 U	2.0 U	2.0 U	2.0 U	2.0 U
Methyl acetate			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylcyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylene chloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
MTBE	10		6.6	2.9	1.0J	1.0U	3.2
o-Xylene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Styrene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Tetrachloroethene	5		2.1	3.1	0.95J	2.3	6.6
Toluene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
trans-1,2-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
trans-1,3-Dichloropropene			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Trichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	0.94 J
Trichlorofluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Vinyl chloride	2		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Xylenes (total)	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U

AWQSGVs - Ambient Water-Quality Standards and Guidance Values

Bold data indicates that parameter was detected above the NYSDEC AWQSGVs

μg/L -Micrograms per liter

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated Value

U - Compound was analyzed for but not detected

<sup>\*</sup>Reporting Limit was not provided in the associated report

<sup>- -</sup> No NYSDEC AWQSGV available

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: 3	W-2 (101-101.5)	GW-2 (111-111.5) (	GW-2RE (111-111.5)	$^{5}$ GW-2 $(118-120)^{3}$	GW-2RE (118-120
Parameter (Concentrations in μg/L)	AWQSGVs (μg/L)	Sample Date:	11/9/2010	11/9/2010	11/9/2010	11/9/2010	11/9/2010
1,1,1-Trichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2,2-Tetrachloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2-Trichloroethane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,3-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,4-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dibromoethane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloroethane	0.6		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloropropane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,3-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,4-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
2-Butanone (MEK)	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
2-Hexanone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
4-Methyl-2-pentanone (MIBK)			5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Acetone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Benzene	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromochloromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromodichloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromoform	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromomethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon disulfide	60		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon tetrachloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroform	7		0.52J	1.0U	1.0U	1.0U	1.0 U
Chloromethane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
cis-1,2-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
cis-1,3-Dichloropropene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Cyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloropropane			NA	NA	NA	NA	NA

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: 3	W-2 (101-101.5)	GW-2 (111-111.5) (	GW-2RE (111-111.5)	<sup>2</sup> GW-2 (118-120) <sup>3</sup>	GW-2RE (118-120)
Parameter	AWQSGVs	Sample Date:	11/9/2010	11/9/2010	11/9/2010	11/9/2010	11/9/2010
(Concentrations in $\mu g/L$ )	$(\mu g/L)$						
Dichlorodifluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Ethylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Freon 113			NA	NA	NA	NA	NA
Isopropylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
m+p-Xylene	5		2.0 U	2.0 U	2.0 U	2.0 U	2.0 U
Methyl acetate			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylcyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylene chloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
MTBE	10		1.0U	1.1	1.2	1.0 U	1.0 U
o-Xylene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Styrene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Tetrachloroethene	5		4.3	0.52J	0.57J	2.7	2.8
Toluene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
trans-1,2-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
trans-1,3-Dichloropropene			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Trichloroethene	5		0.51 J	1.0 U	1.0 U	1.0 U	1.0 U
Trichlorofluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Vinyl chloride	2		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Xylenes (total)	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U

AWQSGVs - Ambient Water-Quality Standards and Guidance Values

Bold data indicates that parameter was detected above the NYSDEC AWQSGVs

μg/L -Micrograms per liter

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated Value

U - Compound was analyzed for but not detected

<sup>\*</sup>Reporting Limit was not provided in the associated report

<sup>- -</sup> No NYSDEC AWQSGV available

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: (	GW-3 (81-81.5) <sup>3</sup>	GW-3RE (81-81.5) <sup>3</sup>	<sup>3</sup> GW-3 (91-91.5) <sup>3</sup>	GW-3 (101-101.5) <sup>3</sup>	GW-3 (111-111.5)
Parameter	AWQSGVs	Sample Date:	11/10/2010	11/10/2010	11/10/2010	11/10/2010	11/10/2010
(Concentrations in µg/L)	(µg/L)						
1,1,1-Trichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2,2-Tetrachloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2-Trichloroethane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,3-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,4-Trichlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dibromoethane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloroethane	0.6		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloropropane	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,3-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,4-Dichlorobenzene	3		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
2-Butanone (MEK)	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
2-Hexanone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
4-Methyl-2-pentanone (MIBK)			5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Acetone	50		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Benzene	1		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromochloromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromodichloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromoform	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromomethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon disulfide	60		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon tetrachloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chlorobenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroform	7		1.0U	1.0U	1.0U	1.0 U	1.0U
Chloromethane			1.0U	1.0U	1.0U	1.0 U	1.0U
cis-1,2-Dichloroethene	5		46	44	25	0.96 J	0.89J
cis-1,3-Dichloropropene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Cyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloromethane	50		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloropropane			NA	NA	NA	NA	NA

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: (	GW-3 (81-81.5) <sup>3</sup>	GW-3RE (81-81.5)	<sup>3</sup> GW-3 (91-91.5) <sup>3</sup> (	GW-3 (101-101.5) <sup>3</sup>	GW-3 (111-111.5
Parameter	AWQSGVs	Sample Date:	11/10/2010	11/10/2010	11/10/2010	11/10/2010	11/10/2010
(Concentrations in µg/L)	$(\mu g/L)$						
Dichlorodifluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Ethylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Freon 113			NA	NA	NA	NA	NA
Isopropylbenzene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
m+p-Xylene	5		2.0 U	2.0 U	2.0 U	2.0 U	2.0 U
Methyl acetate			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylcyclohexane			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Methylene chloride	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
MTBE	10		1.0U	1.0U	2	1	1.0 U
o-Xylene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Styrene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Tetrachloroethene	5		56	56	58	8.1	6.2
Toluene	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
trans-1,2-Dichloroethene	5		3	3.1	1.4	1.0U	1.0U
trans-1,3-Dichloropropene			1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Trichloroethene	5		16	16	17	1.3	0.94 J
Trichlorofluoromethane	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Vinyl chloride	2		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Xylenes (total)	5		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U

AWQSGVs - Ambient Water-Quality Standards and Guidance Values  $\mu g/L$  -Micrograms per liter

Bold data indicates that parameter was detected above the NYSDEC AWQSGVs

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated Value

U - Compound was analyzed for but not detected

<sup>\*</sup>Reporting Limit was not provided in the associated report

<sup>- -</sup> No NYSDEC AWQSGV available

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: GW-3 (121-121.5)	Duplicate <sup>3</sup>	GW-4 (91-91.5) <sup>3</sup>	GW-4 (101-101.5) <sup>3</sup>	GW-4 (111-111.5) <sup>3</sup>
Parameter	AWQSGVs	Sample Date: 11/10/2010	11/10/2010	11/10/2010	11/10/2010	11/10/2010
(Concentrations in µg/L)	(µg/L)	•				
	, C					
1,1,1-Trichloroethane	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2,2-Tetrachloroethane	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1,2-Trichloroethane	1	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethane	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,1-Dichloroethene	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,3-Trichlorobenzene	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2,4-Trichlorobenzene	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dibromoethane		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichlorobenzene	3	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloroethane	0.6	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,2-Dichloropropane	1	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,3-Dichlorobenzene	3	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
1,4-Dichlorobenzene	3	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
2-Butanone (MEK)	50	5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
2-Hexanone	50	5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
4-Methyl-2-pentanone (MIBK)		5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Acetone	50	5.0 U	5.0 U	5.0 U	5.0 U	5.0 U
Benzene	1	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromochloromethane	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromodichloromethane	50	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromoform	50	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Bromomethane	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon disulfide	60	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Carbon tetrachloride	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chlorobenzene	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroethane	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Chloroform	7	1.0U	1.0U	1.0U	1.0U	1.0U
Chloromethane		1.0U	1.0U	1.0U	1.0U	1.0U
cis-1,2-Dichloroethene	5	1.0U	28	1.0U	1.0U	1.0U
cis-1,3-Dichloropropene	5	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Cyclohexane		1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloromethane	50	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Dibromochloropropane		NA	NA	NA	NA	NA

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation: GW-3 (1	21-121.5) <sup>2</sup> Duplicate <sup>3</sup>	$GW-4 (91-91.5)^3$	$GW-4 (101-101.5)^3$	GW-4 (111-111.5) <sup>3</sup>
Parameter	AWQSGVs	Sample Date: 11/1	0/2010 11/10/2010	11/10/2010	11/10/2010	11/10/2010
(Concentrations in µg/L)	$(\mu g/L)$					
Dichlorodifluoromethane	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Ethylbenzene	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Freon 113		1	NA NA	NA	NA	NA
Isopropylbenzene	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
m+p-Xylene	5	2.	0 U 2.0 U	2.0 U	2.0 U	2.0 U
Methyl acetate		1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Methylcyclohexane		1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Methylene chloride	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
MTBE	10	1.	.0U 1.0U	30	22	11
o-Xylene	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Styrene	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Tetrachloroethene	5	0.	56 J <b>84</b>	5.7	4.6	3.6
Toluene	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
trans-1,2-Dichloroethene	5	1.	0 U 0.57 J	1.0U	1.0U	1.0U
trans-1,3-Dichloropropene		1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Trichloroethene	5	1.	0 U <b>19</b>	0.9J	0.62J	1.0U
Trichlorofluoromethane	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Vinyl chloride	2	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U
Xylenes (total)	5	1.	0 U 1.0 U	1.0 U	1.0 U	1.0 U

AWQSGVs - Ambient Water-Quality Standards and Guidance Values  $\mu g/L$  -Micrograms per liter

Bold data indicates that parameter was detected above the NYSDEC AWQSGVs

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated Value

U - Compound was analyzed for but not detected

<sup>\*</sup>Reporting Limit was not provided in the associated report

<sup>- -</sup> No NYSDEC AWQSGV available

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	MAGDEC	Cl- D ' '	CW 4 (101 101 5)3
Dawanatan	NYSDEC	Sample Designation:	GW-4 (121-121.5) <sup>3</sup> 11/10/2010
Parameter (Composition of in 1997)	AWQSGVs	Sample Date:	11/10/2010
(Concentrations in µg/L)	(µg/L)		
1,1,1-Trichloroethane	5		1.0 U
1,1,2,2-Tetrachloroethane	5		1.0 U
1,1,2-Trichloroethane	1		1.0 U
1,1-Dichloroethane	5		1.0 U
1,1-Dichloroethene	5		1.0 U
1,2,3-Trichlorobenzene	5		1.0 U
1,2,4-Trichlorobenzene	5		1.0 U
1,2-Dibromoethane			1.0 U
1,2-Dichlorobenzene	3		1.0 U
1,2-Dichloroethane	0.6		1.0 U
1,2-Dichloropropane	1		1.0 U
1,3-Dichlorobenzene	3		1.0 U
1,4-Dichlorobenzene	3		1.0 U
2-Butanone (MEK)	50		5.0 U
2-Hexanone	50		5.0 U
4-Methyl-2-pentanone (MIBK)			5.0 U
Acetone	50		5.0 U
Benzene	1		1.0 U
Bromochloromethane	5		1.0 U
Bromodichloromethane	50		1.0 U
Bromoform	50		1.0 U
Bromomethane	5		1.0 U
Carbon disulfide	60		1.0 U
Carbon tetrachloride	5		1.0 U
Chlorobenzene	5		1.0 U
Chloroethane	5		1.0 U
Chloroform	7		0.95J
Chloromethane			1.0U
cis-1,2-Dichloroethene	5		1.0U
cis-1,3-Dichloropropene	5		1.0 U
Cyclohexane			1.0 U
Dibromochloromethane	50		1.0 U
Dibromochloropropane			NA

Table 2. Summary of Volatile Orgainc Compounds in Groundwater Samples Collected by GFE, LBG and HRP 1022 Old Country Road, Plainview, New York

	NYSDEC	Sample Designation:	$GW-4 (121-121.5)^3$
Parameter	AWQSGVs	Sample Date:	11/10/2010
(Concentrations in µg/L)	$(\mu g/L)$		
Dichlorodifluoromethane	5		1.0 U
Ethylbenzene	5		1.0 U
Freon 113			NA
Isopropylbenzene	5		0.57 J
m+p-Xylene	5		2.0 U
Methyl acetate			1.0 U
Methylcyclohexane			1.0 U
Methylene chloride	5		1.0 U
MTBE	10		3.5
o-Xylene	5		1.0 U
Styrene	5		1.0 U
Tetrachloroethene	5		2.7
Toluene	5		1.0 U
trans-1,2-Dichloroethene	5		1.0U
trans-1,3-Dichloropropene			1.0 U
Trichloroethene	5		1.0U
Trichlorofluoromethane	5		1.0 U
Vinyl chloride	2		1.0 U
Xylenes (total)	5		1.0 U

AWQSGVs - Ambient Water-Quality Standards and Guidance Values

 $\mu g/L$  -Micrograms per liter

Bold data indicates that parameter was detected above the NYSDEC AWQSGVs

<sup>&</sup>lt;sup>1</sup>- Data was obtained from GFE 2006 Site Characterization Report

<sup>&</sup>lt;sup>2</sup>- Data was obtained from LBG 2007 Environmenal Site Assessment Phase II Report

<sup>&</sup>lt;sup>3</sup>- Data was obtained from HRP 2011 Site Charactization Report

J - Estimated Value

U - Compound was analyzed for but not detected

<sup>\*</sup>Reporting Limit was not provided in the associated report

<sup>- -</sup> No NYSDEC AWQSGV available

Table 3. Proposed Soil Sampling Locations, 1022 Old Country Road, Plainview, New York

Area of Concern (AOC)	Location	Matrix	Sample Depths*	Sample Parameters	Sampling Method**	Rationale
	RSB-1	Soil	Up to 2 samples: interval with the highest elevated PID reading, and the terminal depth of the boring (to be	TCL VOCs	SW-846 8260B	To delineate the nature and extent of contamination identified during previous investigations and identify the historic septic sanitary system
	RSB-2	Soil		TCL VOCs	SW-846 8260B	
	RSB-3	Soil		TCL VOCs	SW-846 8260B	
AOC-1	RSB-4	Soil		TCL VOCs	SW-846 8260B	
	RSB-5	Soil	determined during field activities)	TCL VOCs	SW-846 8260B	
	RSB-6	Soil		TCL VOCs	SW-846 8260B	
	RSB-7	Soil		TCL VOCs	SW-846 8260B	At the request of NYSDEC, to confirm that all impacted soil was removed during previous remedial actions (i.e., UST removal).

VOCs - Volatile Organic Compounds QA/QC samples will be collected as described in the QAPP (Appendix B)

<sup>\*</sup> Depths are in feet from grade; Additional intervals may be added based on field observations \*\* Laboratory will report to their minimum possible standards for each method (QAPP Table 2)

TCL - USEPA Contract Laboratory Program Target Compound List

Table 4. Proposed Groundwater Sampling Locations, 1022 Old Country Road, Plainview, New York

AOC-1	Area of Concern (AOC)	Monitoring Wells*	Matrix	Sample Depth**	Sample Parameters	Sampling Method***	Rationale
		RMW-1 and RMW-2	Groundwater	± 70-90	TCL VOCs	SW-846 8260B	To evaluate the nature and extent of groundwater impacts as indicated by previous investigations
	Groundwater	RWM-3	Groundwater	± 70-90	TCL VOCs	SW-846 8260B	At the request of NYSDEC, to evaluate upgradient groundwater
		RB-1	Groundwater	± 70-72	TCL VOCs	SW-846 8260B	At the request of NYSDEC, to evaluate the nature and extent of groundwater impacts as indicated by previous investigations

<sup>\*-</sup> Locations based on soil boring field screening results

TCL - USEPA Contract Laboratory Program Target Compound List

VOCs - Volatile Organic Compounds

QA/QC samples will be collected as described in the QAPP (Appendix C)

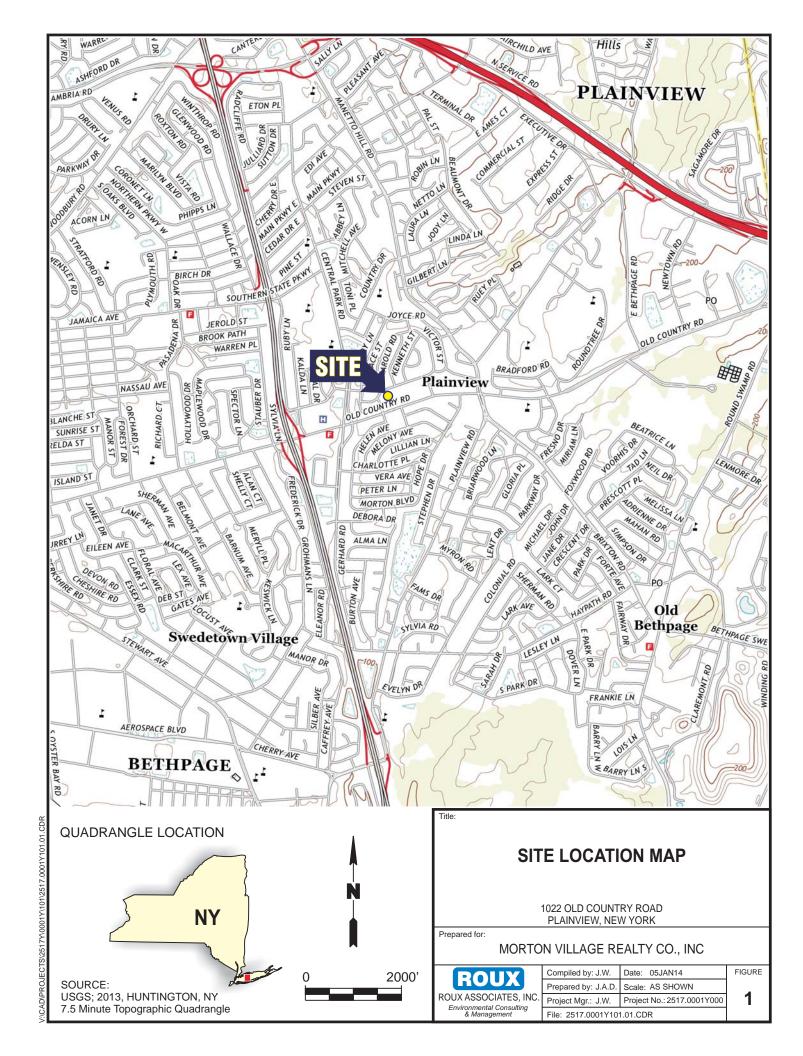
<sup>\*\*-</sup> Feet below grade, screened interval of well or grab sample

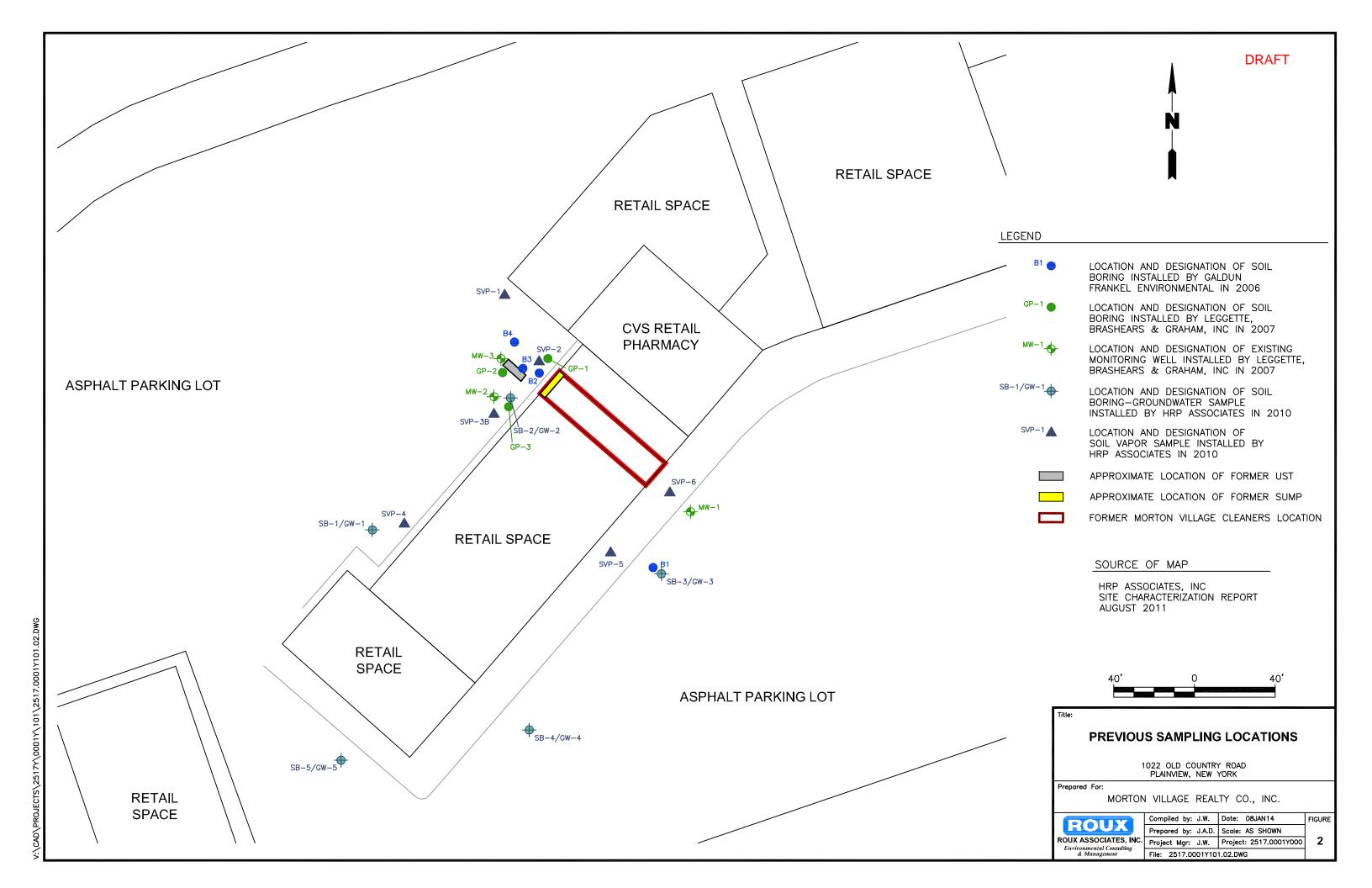
<sup>\*\*\*-</sup> Laboratory will report to their minimum possible standards for each method (QAPP Table 2)

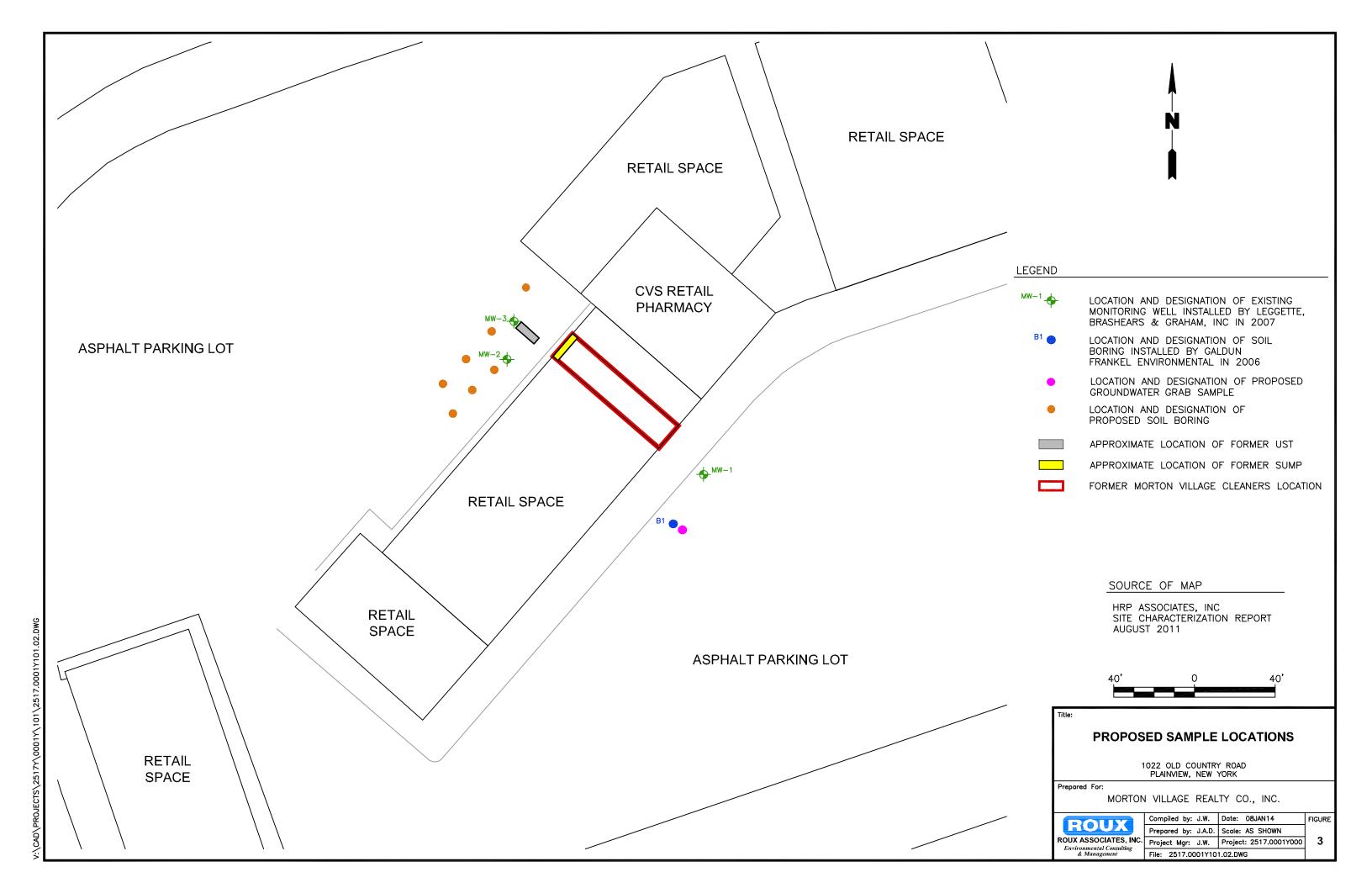
# Remedial Investigation Work Plan 1022 Old Country Road, Plainview, New York 11803

## **FIGURES**

- 1. Site Location Map
- 2. Previous Sampling Locations
- 3. Proposed Sample Locations







# Remedial Investigation Work Plan 1022 Old Country Road, Plainview, New York 11803

## **APPENDICES**

- A. Field Sampling Plan
- B. Quality Assurance Project Plan
- C. Site Health and Safety Plan

# Remedial Investigation Work Plan 1022 Old Country Road, Plainview, New York 11803

## **APPENDIX A**

Field Sampling Plan

# FIELD SAMPLING PLAN

1022 Old Country Road Plainview, New York 11803

Prepared for

MORTON VILLAGE REALTY COMPANY, INC. % John Curran Sive Paget and Riesel P.C. 460 Park Avenue New York, New York 10022

# **ROUX ASSOCIATES, INC.**

Environmental Consulting & Management



209 Shafter Street, Islandia, New York 11749 ♦ 631-232-2600

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### **TABLES**

- 1. Remedial Investigation Field and Quality Control Sampling Summary
- 2. Preservation, Holding Times and Sample Containers

### **ATTACHMENTS**

- 1. Roux Associates' Standard Operating Procedure for Tasks Described in this Field Sampling Plan
- 2. Chain of Custody Form

#### 1.0 INTRODUCTION

Roux Associates has developed this Field Sampling Plan (FSP) to describe in detail the field sampling methods to be used during performance of the Remedial Investigation (RI) for the site located at 1022 Old Country Road, Plainview, New York (Site). The focus of this RI Work Plan is on the former Morton Village Cleaners, a/k/a Classic French Cleaners, (former Cleaners) tenant space (1022 Old Country Road – currently occupied by a Subway restaurant) and on an area (asphalt parking lot) immediately north of the former Cleaners tenant space. The former Cleaners tenant space and parking lot area are collectively referred to as, Area of Concern 1 (AOC-1). The objective of the RI is to evaluate soil, groundwater and soil vapor (including sub-slab and indoor air) quality associated with AOC-1; evaluate the potential for impacts from suspected historical septic system (potential source of CVOCs in soil/groundwater) that was associated with the former Cleaners; and delineate the nature and extent of previously-documented CVOC impacts to soil, groundwater and soil vapor.

The FSP was prepared in accordance with directives provided in the DER-10 Technical Guidance for Site Investigation and Remediation (May 2010) issued by the New York State Department of Environmental Conservation (NYSDEC), and provides guidelines and procedures to be followed by field personnel during performance of the RI. Information contained in this FSP relates to sampling objectives, sampling locations, sampling frequencies, sample designations, sampling equipment, sample handling, sample analysis, and decontamination.

### 2.0 SAMPLING OBJECTIVES

Prior Site characterization work was conducted on the Site over the course of several years by various entities.

The objective of the proposed sampling is to determine the nature and extent of the known contamination on Site, to evaluate any additional AOCs (historic septic sanitary system) via geophysical survey and potential associated contamination, and to obtain a current representation of the environmental conditions at the Site.

The sampling procedures associated with characterization of soil, groundwater and soil vapor are discussed in detail in Section 4 of this FSP. A discussion of the data quality objectives (DQOs) is provided in the Quality Assurance Project Plan (QAPP) located in Appendix B of the RI Work Plan.

#### 3.0 SAMPLE MEDIA, LOCATIONS, ANALYTICAL SUITES, AND FREQUENCY

The media to be sampled during the RI include soil, groundwater and soil vapor. Sampling locations, analytical suites, and frequency vary by medium. A discussion of the sampling schedule for each medium is provided below, while the assumed number of field samples to be collected for each medium, including quality control (QC) samples, is shown in Table 1. Specifics regarding the collection of samples at each location and for each task are provided in Section 4 of this FSP.

#### 3.1 Soil Sampling

Soil samples underlying the Site will be collected at seven locations as shown in Figure 3 of the RI Work Plan. At each location, up to two soil samples will be collected: one soil sample will be collected from the interval that exhibits visual evidence of contamination or an elevated PID reading; and a second soil sample will be collected from the terminal depth of the boring (to be determined during field activities). All samples will be analyzed for Target Compound List (TCL) Volatile Organic Compounds (VOCs). The soil sampling rationale is summarized in Table 3 of the RI Work Plan.

Continuous soil samples will be collected using either the direst-push method via a Geoprobe<sup>®</sup> with a five-foot macrocore sampler or hollow stem auger (HSA). During soil boring activities, lithology will be recorded and soil will be field screened every two feet for VOCs using a PID.

As described in Section 3.2.2 of the RI Work Plan, soil borings locations may be adjusted based on findings from the geophysical survey.

#### 3.2 Groundwater Sampling

Groundwater samples will be collected from three groundwater monitoring wells that will be installed during the RI based on soil field screening results as well as groundwater flow direction. Following gauging activities, each well will be sampled for TCL VOCs. Field parameters including pH, turbidity, conductivity, dissolved oxygen, oxygen reducing potential and temperature will also be measured prior to sampling. The groundwater sampling rationale is summarized in Table 4 of the RI Work Plan.

## 3.3 Soil Vapor Sampling

A soil vapor investigation will be conducted as part of Phase B of this RI, which will include the collection of soil vapor, sub-slab and indoor air samples. The number and location of the samples will be determined following the completion of Phase A of this RI. All soil vapor, sub-slab and indoor air samples will be analyzed for VOCs via USEPA Method TO-15. All sample locations will be discussed with and approved by the NYSDEC and NYSDOH prior to completion.

#### 4.0 FIELD SAMPLING PROCEDURES

This section provides a detailed discussion of the field procedures to be used during sampling of the various media being evaluated as part of the RI (i.e., soil, groundwater and soil vapor). The soil and groundwater locations are shown on Figure 3 of the RI work plan and additional information including intervals to be sampled and sample rationale is provided in Tables 3 and 4 of the RI work plan.

### 4.1 Soil Sampling, Monitoring Well Installation/Sampling and Soil Vapor Sampling

Details for the collection of soil samples and the installation of monitoring wells, collection of groundwater samples and installation and sampling of soil vapor monitoring points (including subslab and indoor air sampling) are provided below.

#### 4.1.1 Soil Sampling

Borings will be advanced using a Geoprobe<sup>®</sup> direct-push or HSA drill rig. Samples of the soil profile will be collected in continuous five-foot increments using a 2-inch diameter macrocore sampler or two-foot increments using a 2-inch diameter split spoon to a maximum depth of approximately 70-75 feet below grade, which is the anticipated depth to the water table.

The soil from each five-foot or two-foot interval will be observed for lithology and evidence of contamination (e.g., staining, odors, and/or visible free product) and placed immediately thereafter into large Zip-loc<sup>TM</sup> bags for recording headspace. After a minimum of 15 minutes for equilibration with the headspace in the Zip-loc<sup>TM</sup> bag, each sample will be screened for organic vapors using a PID. Samples for VOC analysis will be placed in a laboratory-supplied jar prior to screening, due to the potential for loss of VOCs through volatilization. Soil samples from the soil borings will be collected according to Table 3 of the RI Work Plan. In all soil borings, soil samples will be collected from the interval that that exhibits visual evidence of contamination or an elevated PID reading. A second soil sample will be collected from the terminal depth of the boring (to be determined during field activities). These samples will be placed in the laboratory-supplied containers and shipped to the laboratory under chain of custody procedures in accordance with Roux Associates Standard Operating Procedures (SOPs). Upon completion, each boring will be allowed to collapse. Any borings remaining open will be backfilled with bentonite hole-plug,

which will be hydrated with potable water. The surface at each location will be restored with an asphalt patch.

Additional details regarding soil-sampling protocols are described in Roux Associates' SOP for the Collection of Soil Samples for Laboratory Analysis, which is provided in Attachment 1.

#### **4.1.2** Monitoring Well Installation

Following soil sampling activities, monitoring wells (RMW-1 and RMW-2) will be installed at the two soil borings with the greatest degree of evidence of contamination. Monitoring well RMW-3 will be installed as an upgradient well based on the groundwater flow direction observed during Phase A of this RI. RMW-1, RMW-2, and RMW-3 are proposed to be installed at a depth of approximately 70 to 90 feet below ground surface (ft-bgs). Monitoring wells will be constructed of 2-inch-inside-diameter, Schedule 40 polyvinyl chloride (PVC) casing and, 0.020-inch slot, machined screen. Well screens will be 20 feet long, and will be installed from approximately to 70 to 90 ft-bgs, depending upon the observed depth of the water table during drilling activities. A sand pack will be placed around the well screen, extending two feet above the top of the screened zone. Once the driller confirms the depth of the sand pack, a minimum two-foot-thick bentonite pellet seal will be placed above the sand pack. Once the pellets have been allowed to hydrate, a cement-bentonite grout will be pumped into the remaining annular space from the bottom up using a tremie pipe lowered to just above the bentonite seal. The wells will be completed using locking well plugs, and flush-mounted, bolt down, watertight, manhole covers cemented into place.

#### 4.1.3 Groundwater Sampling

Each well will be developed to remove any fine-grained material in the vicinity of the well screen and to promote hydraulic connection with the aquifer. The wells will be developed using a submersible pump, which will be surged periodically until well yield is consistent and has a turbidity below 50 nephelometric turbidity units (NTUs), if possible.

Groundwater samples will be collected no sooner than one week following development of the wells. Prior to sampling, depth to water will be measured at each well using an electronic water level indicator with an accuracy of +/-0.01 feet. All wells will then be purged and sampled using a bladder pump, or an alternative method, depending on the observed depth to groundwater and

logistical issues. Samples will be analyzed for TCL VOCs. Additional details for the collection of groundwater samples are included in the Roux Associates SOPs (Attachment 1).

All groundwater samples will be placed in the laboratory-supplied containers and shipped to the laboratory under chain of custody procedures in accordance with Roux Associates' SOPs.

#### 4.1.4 Soil Vapor and Sub-Slab Installations

Soil vapor points will be installed by hand or with a Geoprobe<sup>®</sup>; pending NYSDEC/NYSDOH approval of the sampling locations and depths, based on historical data and Phase A findings. Soil vapor points will be constructed with a six-inch long, stainless steel, mess sample screen attached to Teflon lined polyurethane tubing. At each soil vapor point, the sample screens will be installed at the bottom of the established sample interval and coarse sand will be added to the annulus between the sample screen and borehole, and up to six-inches above the top of the sample screen in order to create a one foot sample zone. Bentonite will be added to the top of the sand and hydrated with potable water to seal the sample zone. The surface will be completed with a flushmounted protective curb box.

Sub-slab soil vapor points will be installed hand tools. At each sub-slab soil vapor point, an approximate 0.5-inch hole will be drilled through the concrete slab to approximately six-inches below the slab, using a hand drill. A six-inch long, stainless steel, mess sample screen attached to Teflon lined polyurethane tubing will be installed at each sub-slab sample location. Coarse sand will be added to the annulus between the sample screen and borehole, and the surface will be sealed with bentonite or molding clay.

Soil vapor and sub-slab soil vapor will be collected no sooner than 24 hours following installation of the points.

#### 4.1.5 Soil Vapor Sampling

All soil vapor, sub-slab and indoor air samples will be collected in accordance with the October 2006 NYSDOH Final Guidance for Evaluating Soil Vapor Intrusion in the State of New York

(NYSDOH Guidance). Soil vapor and sub-slab samples will be collected utilizing the following procedural steps:

- 1. A clay seal will be placed around the sampling tubing in an effort to further minimize the infiltration of the atmospheric air present at land surface directly above the sampling point (ambient air).
- 2. The sampling tubing will be connected to a "T" connector three-way valve assembly, with one end of the "T" connector leading to a vacuum pump and the other end leading to a pre evacuated summa canister with a calibrated flow controller.
- 3. The soil vapor sample tubing and surrounding sand pack will be purged of approximately three volumes of air using a vacuum pump set at a rate of approximately 0.2 liters per minute.
- 4. A tracer gas (e.g., helium) will be used to enrich the atmosphere in the immediate vicinity of the sampling location where the sampling tubing intersected the ground surface in order to test the borehole seal and verify that ambient air is not inadvertently being drawn into the sample. A plastic container (i.e. bucket), with a soft seal, will be placed over the monitoring point and the helium will be injected into the bucket during purging of the monitoring point in an effort to enrich the interior of the bucket with the tracer gas. Note that the 3-way valve used to connect the sample tubing to the equipment will also be placed under a shroud (i.e., bucket enclosure) and included in the tracer gas verification. This will be done in an effort to ensure that the valve does not provide a means by which ambient air would enter the canister and possibly dilute the sample. Both the purge volume from the sample tubing (i.e., also the air that passed through the 3-way valve), and the helium enriched area within the bucket will be screened for the tracer gas. The gas will be measured with a MGD-2002 helium detector (by Dielectric) or equivalent meter, which measures the rate of helium leakage at the surface. If the screening results show that the rate of helium detected from the sampling tubing is greater than 10% of the helium detected in the enriched area (i.e., within the bucket), the seals around the sampling equipment will be reset and the sampling tubing will be re-purged until the tracer gas is no longer detected at levels greater than 10% of the enriched area.
- 5. Following the purging and tracer gas verification steps, the valve leading to the pump will be closed, the pump turned off, and the soil vapor directed to a 1-liter summa canister for sample collection. A laboratory-supplied, calibrated flow controller set for a 2-hour sample period (soil vapor samples) and an 8-hour sample period (sub-slab samples) will be used at each location, to restrict the sample collection rate to 0.2 liters per minute or less.

Indoor air samples will be collected using laboratory supplied 1-liter summa canisters and calibrated flow controller set for 8-hour sample period.

## 4.2 Geophysical Survey

A geophysical survey will be performed at the Site to attempt to locate the historic septic sanitary system and any additional previously unidentified AOC, as well as potential utility interference with the planned scope of work. Previous Site characterization work at the Site identified a former UST within the parking lot immediately north of the former Classic French Cleaners (AOC-1), which was removed in 2007. Roux Associates will target geophysical survey activities on areas that are previously uninvestigated within AOC-1 including the areas east and north of the former UST area.

#### 5.0 SAMPLE HANDLING AND ANALYSIS

To ensure quality data acquisition and collection of representative samples, there are selective procedures to minimize sample degradation or contamination. These include procedures for preservation of the samples as well as sample packaging and shipping procedures.

#### 5.1 Field Sample Handling

A detailed discussion of the number and types of samples to be collected during each task, as well as the analyses to be performed can be found in Section 3.0 of this FSP. The types of containers, volumes needed, and preservation techniques for the aforementioned testing parameters are presented in Table 2.

#### **5.2 Sample Custody Documentation**

The purpose of documenting sample custody is to confirm that the integrity and handling of the samples is not subject to question. Sample custody will be maintained from the point of sampling through the analysis. Specific procedures regarding sample tracking from the field to the laboratory are described in Roux Associates' SOP for Sample Handling (Attachment 1).

Each individual collecting samples is personally responsible for the care and custody of the samples. All sample labels should be pre-printed or filled out using waterproof ink. The technical staff will review all field activities with the Field Team Leader to determine whether proper custody procedures were followed during the fieldwork and to decide if additional samples are required.

All samples being shipped off-Site for analysis must be accompanied by a properly completed chain of custody form (Attachment D-2). The sample numbers will be listed on the chain of custody form. When transferring the possession of samples, individuals relinquishing and receiving will sign, date, and note the time on the record. This record documents transfer of custody of samples from the sampler to another person, to/from a secure storage area, and to the laboratory.

Samples will be packaged for laboratory pick up and/or shipment with a separate signed custody record enclosed in each sample box or cooler. Shipping containers will be locked and/or secured with strapping tape in at least two locations for shipment to the laboratory.

### **5.3** Sample Shipment

Laboratory courier services will be used for sample transport on this project. However, in the event that samples are shipped to the laboratory the following procedures will apply. Sample packaging and shipping procedures are based upon USEPA specifications, as well as U.S. Department of Transportation (DOT) regulations. The procedures vary according to potential sample analytes, concentration, and matrix, and are designed to provide optimum protection for the samples and the public. Sample packaging and shipment must be performed using the general outline described below. Additional information regarding sample handling is provided in Roux Associates' SOP for Sample Handling (Attachment 1).

All samples will be shipped within 12 hours of collection (when possible) and will be preserved appropriately from the time of sample collection. A description of the sample packing and shipping procedures is presented below:

- 1. Prepare cooler(s) for shipment.
  - Tape drain(s) of cooler shut;
  - Affix "This Side Up" arrow labels and "Fragile" labels on each cooler; and
  - Place mailing label with laboratory address on top of cooler(s).
- 2. Arrange sample containers in groups by sample number or analyte.
- 3. Ensure that all bottle labels are completed correctly. Place clear tape over bottle labels to prevent moisture accumulation from causing the label to peel off.
- 4. Arrange containers in front of assigned coolers.
- 5. Place packaging material at the bottom of the cooler to act as a cushion for the sample containers.
- 6. Arrange containers in the cooler so that they are not in contact with the cooler or other samples.
- 7. Fill remaining spaces with packaging material.
- 8. Ensure all containers are firmly packed with packaging material.

- 9. If ice is required to preserve the samples, ice cubes should be repackaged in double Zip-Lock<sup>TM</sup> bags, and placed on top of the packaging material.
- 10. Sign chain of custody form (or obtain signature) and indicate the time and date it was relinquished to Federal Express or other carrier, as appropriate.
- 11. Separate chain of custody forms. Seal proper copies within a large Zip-Lock<sup>TM</sup> bag and tape to cooler. Retain copies of all forms.
- 12. Close lid and latch.
- 13. Secure each cooler using custody seals.
- 14. Tape cooler shut on both ends.
- 15. Relinquish to Federal Express or other courier service as appropriate. Retain airbill receipt for project records. (Note: All samples will be shipped for "NEXT A.M." delivery).
- 16. Telephone laboratory contact and provide him/her with the following shipment information:
  - sampler's name;
  - project name;
  - number of samples sent according to matrix and concentration; and
  - airbill number.

#### 6.0 SITE CONTROL PROCEDURES

Site control procedures, including decontamination and waste handling and disposal, are discussed below.

#### **6.1 Decontamination**

In an attempt to avoid the spread of contamination, all drilling and sampling equipment must be decontaminated at a reasonable frequency in a properly designed and located decontamination area. Detailed procedures for the decontamination of field and sampling equipment are included in Roux Associates' SOPs for the Decontamination of Field Equipment, which is provided in Attachment D-1. The location of the decontamination area will be determined prior to the start of field operations. The decontamination area will be constructed to ensure that all wash water generated during decontamination can be collected and containerized for proper disposal.

#### **6.2** Waste Handling and Disposal

All waste materials (drill cuttings, decontamination water, etc.) generated during the RI will be consolidated and stored in appropriate bulk containers (drums, etc.), and temporarily staged at an investigation-derived-waste storage area on-Site. Roux Associates will then coordinate waste characterization and disposal by appropriate means.

## **TABLES**

- 1. Remedial Investigation Field and Quality Control Sampling Summary
- 2. Preservation, Holding Times and Sample Containers

Table 1. Remedial Investigation Field and Quality Control Sampling Summary

Sample Medium	Target Analytes	Field Samples	Replicates <sup>1</sup>	Trip Blanks <sup>2</sup>	Field Blanks <sup>3</sup>	Matrix Spikes <sup>1</sup>	Spike Duplicates <sup>1</sup>	Total No. of Samples
Soil	TCL VOCs	12	1	5	5	1	1	25
Groundwater	TCL VOCs	3	1	1	1	1	1	8
Soil Vapor <sup>4</sup>	VOCs (TO-15)	TBD	TBD	0	0	0	0	TBD

<sup>&</sup>lt;sup>1</sup>- Based on 1 per 20 samples or 1 per Sample Delivery Group (3 days max)

TCL - USEPA Contract Laboratory Program Target Compound List

VOCs - Volatile organic compounds

TBD - To be determined

<sup>&</sup>lt;sup>2</sup> - Based on 1 cooler per day

<sup>&</sup>lt;sup>3</sup> - Based on 1 per day

<sup>&</sup>lt;sup>4</sup> - Including sub-slab and indoor air

**Table 2. Preservation, Holding Times and Sample Containers** 

Analysis	Bottle Type	Preservation <sup>(a)</sup>	Holding Time <sup>(b)</sup>
<u>Target Compound List (TCL)</u> Soil			
TCL Volatile Organic Compounds (VOCs) SW-846 8260B	(3) 5 gram En Core Sampler	Cool to 4°C	48 hours from sample collection
Groundwater TCL VOCs 8260	(3) 40 mL Vials with telfon-lined cap	Cool to 4°C	14 days to extract, 40 days to analysis
Soil Vapor VOCs TO-15	(1) 1-liter Summa Canister	None	30 days from sample collection

<sup>(</sup>a) All soil and groundwater samples to be preserved in ice during collection and transport

<sup>(</sup>b) Days from date of sample collection.

## **ATTACHMENTS**

- 1. Roux Associates' Standard Operating Procedure for Tasks Described in this Field Sampling Plan
- 2. Chain of Custody Form

## **ATTACHMENT 1**

Roux Associates' Standard Operating Procedure for Tasks Described in this Field Sampling Plan

Date: May 5, 2000

#### 1.0 PURPOSE

The purpose of this standard operating procedure (SOP) is to establish guidelines for the sampling of ground-water monitoring wells for dissolved constituents. As part of the SOP for the sampling of ground-water monitoring wells, sample collection equipment and devices must be considered, and equipment decontamination and pre-sampling procedures (e.g., measuring water levels, sounding wells, and purging wells) must be implemented. Sampling objectives must be firmly established in the work plan before considering the above.

Valid water-chemistry data are integral to a hydrogeologic investigation that characterizes ground-water quality conditions. Water-quality data are used to evaluate both current and historic aquifer chemistry conditions, as well as to estimate future conditions (e.g., trends, migration pathways). Water-quality data can be used to construct ground-water quality maps to illustrate chemical conditions within the flow system, to generate water-quality plots to depict conditions with time and trends, and to perform statistical analyses to quantify data variability, trends, and cleanup levels.

#### 2.0 EQUIPMENT AND MATERIALS

- 2.1 In order to sample ground water from monitoring wells, specific equipment and materials are required. The equipment and materials list may include, but not necessarily be limited to, the following:
  - a. Bailers (Teflon<sup>TM</sup> or stainless steel).
  - b. Pumps (centrifugal, peristaltic, bladder, electric submersible, bilge, hand-operated diaphragm, etc.).
  - c. Gas-displacement device(s).
  - d. Air-lift device(s).
  - e. Teflon<sup>TM</sup> tape, electrical tape.
  - f. Appropriate discharge hose.
  - g. Appropriate discharge tubing (e.g., polypropylene, teflon, etc.) if using a peristaltic pump.
  - h. Appropriate compressed gas if using bladder-type or gas-displacement device.

- i. Portable generator and gasoline or alternate power supply if using an electric submersible pump.
- j. Non-absorbent cord (e.g., polypropylene, etc.).
- k. Plastic sheeting.
- 1. Tape measure (stainless steel, steel, fiberglass) with 0.01-foot measurement increments and chalk (blue carpenter's).
- m. Electronic water-level indicators (e.g., m-scope, etc.) or electric water-level/product level indicators.
- n. Non-phosphate, laboratory-grade detergent.
- o. Distilled/Deionized water.
- p. Potable water.
- q. Paper towels, clean rags.
- r. Roux Associates' field forms (e.g., daily log, well inspection checklist, sampling, etc.) and field notebook.
- s. Well location and site map.
- t. Well keys.
- u. Stop watch, digital watch with second increments, or watch with a second hand.
- v. Water Well Handbook.
- w. Calculator.
- x. Black pen and water-proof marker.
- y. Tools (e.g., pipe wrenches, screwdrivers, hammer, pliers, flashlight, pen knife, etc.).
- z. Appropriate health and safety equipment, as specified in the site health and safety plan (HASP).
- aa. pH meter(s) and buffers.
- bb. Conductivity meter(s) and standards.
- cc. Thermometer(s).

- dd. Extra batteries (meters, thermometers, flashlight).
- ee. Filtration apparatus, filters, pre-filters.
- ff. Plasticware (e.g., premeasured buckets, beakers, flasks, funnels).
- gg. Disposable gloves.
- hh. Water jugs.
- ii. Laboratory-supplied sample containers with labels.
- jj. Cooler(s).
- kk. Ice (wet, blue packs).
- ll. Masking, duct, and packing tape.
- mm. Chain-of-custody form(s) and custody seal(s).
- nn. Site sampling and analysis plan (SAP).
- oo. Site health and safety plan (HASP).
- pp. Packing material (e.g., bubble wrap)
- qq. "Zip-lock" plastic bags.
- rr. Overnight (express) mail forms.

#### 3.0 DECONTAMINATION

- 3.1 Make sure all equipment is decontaminated and cleaned before use (refer to the SOP for Decontamination of Field Equipment for detailed decontamination methods, summaries for bailers and pumps are provided below). Use new, clean materials when decontamination is not appropriate (e.g., non-absorbent cord, disposable gloves). Document, and initial and date the decontamination procedures on the appropriate field form and in the field notebook.
  - a. Decontaminate a bailer by: 1) wearing disposable gloves, 2) disassembling (if appropriate) and scrubbing in a non-phosphate, laboratory-grade detergent and distilled/deionized water solution, and 3) rinsing first with potable water and then distilled/deionized water.
  - b. Decontaminate a pump by: 1) wearing disposable gloves, 2) flushing the pump and discharge hose (if not disposable) first with a non-phosphate, laboratory-grade detergent and potable water solution in an appropriate

container (clean bucket, garbage can, or 55-gallon drum) and then with distilled/deionized water or potable water, and 3) wiping pump-related equipment (e.g., electrical lines, cables, discharge hose) first with a clean cloth and detergent solution and then rinsing or wiping with a clean cloth and distilled/deionized water or potable water.

3.2 Note that the decontamination procedures for bailers and pumps are the minimum that must be performed. Check the work plan to determine if chemicals specified by individual state regulatory agencies must also be used for decontamination procedures (e.g., hexane, nitric acid, acetone, isopropanol, etc.).

#### 4.0 CALIBRATION OF FIELD ANALYSIS EQUIPMENT

Calibrate field analysis equipment before use (e.g., thermometers, pH and conductivity meters, etc.). Refer to the specific SOP for field analysis for each respective piece of equipment. Document, and initial and date the calibration procedures on the appropriate field form, in the field notebook, and in the calibration log book.

#### 5.0 PROCEDURE

- 5.1 Document, and initial and date well identification, pre-sampling information, and problems encountered on the appropriate field form and in the field notebook as needed.
- 5.2 Inspect the protective casing of the well and the well casing, and note any items of concern such as a missing lock, or bent or damaged casing(s).
- 5.3 Place plastic sheeting around the well to protect sampling equipment from potential cross contamination.
- 5.4 Remove the well cap or plug and, if necessary, clean the top of the well off with a clean rag. Place the cap or plug on the plastic sheeting. If the well is not vented, allow several minutes for the water level in the well to equilibrate. If fumes or gases are present, then diagnose these with the proper safety equipment. Never inhale the vapors.
- 5.5 Measure the depth to water (DTW) from the measuring point (MP) on the well using a steel tape and chalk or an electronic sounding device (m-scope). Refer to the specific SOPs for details regarding the use of a steel tape or a m-scope for measuring water levels. Calculate the water-level elevation. Document, and initial and date the information on the appropriate field form and in the field notebook.
- 5.6 Measuring the total depth of the well from the MP with a weighted steel tape. Calculate and record the volume of standing water in the well casing on the appropriate field form and in the field notebook.

- 5.7 Decontaminate the equipment used to measure the water level and sound the well with a non-phosphate, laboratory-grade detergent solution followed by a distilled/deionized water rinse.
- 5.8 Purge the well prior to sampling (refer to the SOP for Purging a Well). The well should be pumped or bailed to remove the volume of water specified in the work plan. Usually three to five casing volumes are removed if the recharge rate is adequate to accomplish this within a reasonable amount of time.

If the formation cannot produce enough water to sustain purging, then one of two options must be followed. These include: 1) pumping or bailing the well dry, or 2) pumping or bailing the well to "near-dry" conditions (i.e., leaving some water in the well). The option employed must be specified in the work plan and be in accordance with regulatory requirements.

If the well is purged dry, then all the standing water has been removed and upon recovery the well is ready for sampling. However, depending on the rate of recovery and the time needed to complete the sampling round, one of the following procedures may have to be implemented: 1) the well may have to be sampled over a period of more than one day; 2) the well may not yield enough water to collect a complete suite of samples and only select (most important) samples will be collected; or 3) the well may not recover which will preclude sampling. Regardless of the option that must be followed, the sampling procedure must be fully documented. When preparing to conduct a sampling round, review drilling, development and previous sampling information (if available) to identify low-yielding wells in order to purge them first, and potentially allow time for the well to recover for sampling.

- 5.9 Record the physical appearance of the water (i.e., color, turbidity, odor, etc.) on the appropriate field form and in the field notebook, as it is purged. Note any changes that occur during purging.
- 5.10 If a bailer is used to collect the sample, then:
  - a. Flush the decontaminated bailer three times with distilled/deionized water.
  - b. Tie the non-absorbent cord (polypropylene) to the bailer with a secure knot and then tie the free end of the bailer cord to the protective casing or, if possible, some nearby structure to prevent losing the bailer and cord down the well.
  - c. Lower the bailer slowly down the well and into the water column to minimize disturbance of the water surface. If a bottom-filling bailer is used, then do not submerge the top of the bailer; however, if a top-filling bailer is used, then submerge the bailer several feet below the water surface.

- d. Remove and properly discard one bailer volume from the well to rinse the bailer with well water before sampling. Again, lower the bailer slowly down the well to the appropriate depth depending on the bailer type (as discussed above in 5.11 c). When removing the bailer from the well, do not allow the bailer cord to rest on the ground but coil it on the protective plastic sheeting placed around the well. Certain regulatory agencies require that the first bailer volume collected be utilized for the samples.
- 5.11 If a pump is used to collect the sample, then use the same pump used to purge the well and, if need be, reduce the discharge rate to facilitate filling sample containers and to avoid problems that can occur while filling sample containers (as listed in Number 5.14, below). Alternately, the purge pump may be removed and a thoroughly decontaminated bailer can be used to collect the sample.
- 5.12 Remove each appropriate container's cap only when ready to fill each with the water sample, and then replace and secure the cap immediately.
- 5.13 Fill each appropriate, pre-labeled sample container carefully and cautiously to prevent: 1) agitating or creating turbulence; 2) breaking the container; 3) entry of, or contact with, any other medium; and 4) spilling/splashing the sample and exposing the sampling team to contaminated water. Immediately place the filled sample container in a ice-filled (wet ice or blue pack) cooler for storage. If wet ice is used it is recommended that it be repackaged in zip-lock bags to help keep the cooler dry and the sample labels secure. Check the work plan as to whether wet ice or blue packs are specified for cooling the samples because certain regulatory agencies may specify the use of one and not the other.
- 5.14 "Top-off" containers for volatile organic compounds (VOCs) and tightly seal with Teflon<sup>TM</sup>-lined septums held in place by open-top screw caps to prevent volatilization. Ensure that there are no bubbles by turning the container upside down and tapping it gently.
- 5.15 Filter water samples (Procedure 4.6) collected for dissolved metals analysis prior to preservation to remove the suspended sediment from the sample. If water samples are to be collected for total metals analysis, then collect a second set of samples without field filtering.

In the event that the regulatory agency(ies) want unfiltered samples for metals analysis, a second set of filtered samples should also be collected. Because unfiltered samples are indications of total metals (dissolved and suspended) they are not representative of aquifer conditions because ground water does not transport sediment (except in some rare cases). Thus, the results for dissolved metals in ground water should be based on filtered samples even if both filtered and unfiltered sets are presented in a report.

- 5.16 Add any necessary preservative(s) to the appropriate container(s) prior to, or after (preferred), the collection of the sample, unless the appropriate preservative(s) have already been added by the laboratory before shipment.
- 5.17 Collect quality control (QC) samples as required in the work plan to monitor sampling and laboratory performance. Refer to the SOP for Collection of Quality Control Samples.
- 5.18 Conduct field analyses after sample collection is complete by measuring and recording the temperature, conductivity, pH, etc. (as called for in the work plan). Note and record the "final" physical appearance of the water (after purging and sampling) on an appropriate field form and in the field notebook.
- 5.19 Wipe the well cap with a clean rag, replace the well cap and protective cover (if present). Lock the protective cover.
- 5.20 Verify that each sample is placed in an individual "zip-lock" bag, wrapped with "bubble wrap," placed in the cooler, and that the cooler has sufficient ice (wet ice or blue packs) to preserve the samples for transportation to the analytical laboratory.
- 5.21 Decontaminate bailers, hoses, and pumps as discussed in the decontamination SOP. Wrap decontaminated equipment with a suitable material (e.g., clean plastic bag or aluminum foil). Discard cords, rags, gloves, etc. in a manner consistent with site conditions.
- 5.22 Complete all necessary field forms, field notebook entries, and the chain-of-custody forms. Retain one copy of each chain-of-custody form. Secure the cooler with sufficient packing tape and a custody seal.
- 5.23 Samples collected from Monday through Friday will be delivered within 24 hours of collection. If Saturday delivery is not available, samples collected on Friday must be delivered by Monday morning. Consult the work plan to determine if any of the analytes require a shorter delivery time.

**END OF PROCUDURE** 

## STANDARD OPERATING PROCEDURE 5.1 FOR COLLECTION OF SOIL SAMPLES FOR LABORATORY ANALYSIS

Date: May 5, 2000

#### 1.0 PURPOSE

The purpose of this Standard Operating Procedure (SOP) is to establish guidelines for the collection of soil samples for laboratory analysis. This SOP is applicable to soil samples collected from split-spoon samplers during drilling, hand auger samples, grab samples from stockpiled soils, surface samples, test pit samples, etc.

#### 2.0 CONSIDERATIONS

Soil samples may be collected in either a random or biased manner. Random samples can be based on a grid system or statistical methodology. Biased samples can be collected in areas of visible impact or suspected source areas. Soil samples can be collected at the surface, shallow subsurface, or at depth. When samples are collected at depth the water content should be noted, since generally "soil sampling" is restricted to the unsaturated zone. Equipment selection will be determined by the depth of the sample to be collected. A thorough description of the sampling locations and proposed methods of sample collection should be included in the work plan.

Commonly, surface sampling refers to the collection of samples at a 0 to 6 inch depth interval. Certain regulatory agencies may define the depth interval of a surface sample differently, and this must be defined in the work plan. Collection of surface soil samples is most efficiently accomplished with the use of a stainless steel trowel or scoop. For samples at greater depths a decontaminated bucket auger or power auger may be needed to advance the hole to the point of sample collection. Another clean bucket auger should then be used to collect the sample. To collect samples at depths of greater than approximately six feet the use of a drill rig and split spoon samples will usually be necessary. In some situations, sample locations are accessed with the use of a backhoe.

### 3.0 MATERIALS/EQUIPMENT

- a. A work plan which outlines soil sampling requirements.
- b. Field notebook, field form(s), maps, chain-of-custody forms, and custody seals.
- c. Decontamination supplies (including: non-phosphate, laboratory grade detergent, buckets, brushes, potable water, distilled water, regulatory-required reagents, aluminum foil, plastic sheeting, etc.).
- d. Sampling device (split-spoon sampler, stainless steel hand auger, stainless steel trowel, etc.).
- e. Stainless steel spoons or spatulas.
- f. Disposable sampling gloves.

## STANDARD OPERATING PROCEDURE 5.1 FOR COLLECTION OF SOIL SAMPLES FOR LABORATORY ANALYSIS

- g. Laboratory-supplied sample containers with labels.
- h. Cooler with blue or wet ice.
- i. Plastic sheeting.
- j. Black pen and indelible marker.
- k. Zip-lock bags and packing material.
- 1. Tape measure.
- m. Paper towels or clean rags.
- n. Masking and packing tape.
- o. Overnight (express) mail forms.

#### 4.0 DECONTAMINATION

All reusable sampling equipment will be thoroughly cleaned according to the decontamination SOP. Where possible, thoroughly pre-cleaned and wrapped sampling equipment should be used and dedicated to individual sampling locations. Disposable items such as sampling gloves, aluminum foil, and plastic sheeting will be changed after each use and discarded in an appropriate manner.

#### 5.0 PROCEDURE

- 5.1 Prior to collecting soil samples, ensure that all sampling equipment has been thoroughly cleaned according to the decontamination SOP. If samples are to be collected at depth, then the boring must be advanced with thoroughly cleaned equipment to the desired sampling horizon and a different thoroughly cleaned sampler must be used to collect the sample.
- 5.2 Using disposable gloves and a pre-cleaned, stainless steel spatula or spoon, extract the soil sample from the sampler, measure the recovery, and separate the wash from the true sample. Where allowed by regulatory agency(ies), disposable plastic spoons may be used.
- 5.3 Place the sample in a laboratory-supplied, pre-cleaned sample container. This should be done as quickly as possible and this is especially important when sampling for volatile organic compounds (VOCs). Samples to be analyzed for VOCs must be collected prior to other constituents.
- 5.4 The sample container will be labeled with appropriate information such as, client name, site location, sample identification (location, depth, etc.), date and time of collection, and sampler's initials.

## STANDARD OPERATING PROCEDURE 5.1 FOR COLLECTION OF SOIL SAMPLES FOR LABORATORY ANALYSIS

- 5.5 Using the remaining portion of soil from the sampler, log the sample in detail and record sediment characteristics (color, odor, moisture, texture, density, consistency, organic content, layering, grain size, etc.).
- 5.6 If soil samples are to be composited in the field, then equal portions from selected locations will be placed on a clean plastic sheet and homogenized. Alternately, several samples may be submitted to the laboratory for compositing by weight. The method used is dependent upon regulatory requirements. Specific compositing procedures shall be approved by the appropriate regulatory agency and described in the work plan. Samples to be analyzed for VOCs will not be composited unless required by a regulatory agency.
- 5.7 After the sample has been collected, labeled, and logged in detail, it is placed in a zip-lock bag and stored in a cooler at 4°C.
- 5.8 A chain-of-custody form is completed for all samples collected. One copy is retained and two are sent with the samples in a zip-lock bag to the laboratory. A custody seal is placed on the cooler prior to shipment.
- 5.9 Samples collected from Monday to Friday are to be delivered to the laboratory within 24 hours of collection. If Saturday delivery is unavailable, samples collected on Friday must be delivered by Monday morning. Check the work plan to determine if any analytes require a shorter delivery time.
- 5.10 The field notebook and appropriate forms should include, but not be limited to the following: client name, site location, sample location, sample depth, sample identification, date and time collected, sampler's name, method of sample collection, number and type of containers, geologic description of material, description of decontamination procedures, etc. A site map should be prepared with exact measurements to each sample location in case follow-up sampling is necessary.
- 5.11 All reusable sampling equipment must be thoroughly cleaned in accordance with the decontamination SOP. Following the final decontamination (after all samples are collected) the sampling equipment is wrapped in aluminum foil. Discard any gloves, foil, plastic, etc. in an appropriate manner that is consistent with site conditions.

**END OF PROCEDURE** 

Date: May 5, 2000

#### 1.0 PURPOSE

The purpose of this standard operating procedure (SOP) is to establish guidelines for sample handling which will allow consistent and accurate results. Valid chemistry data are integral to investigations that characterize media-quality conditions. Thus, this SOP is designed to ensure that once samples are collected, they are preserved, packed and delivered in a manner which will maintain sample integrity to as great an extent as possible. The procedures outlined are applicable to most sampling events and any required modifications must be clearly described in the work plan.

#### 2.0 CONSIDERATIONS

Sample containers, sampling equipment decontamination, quality assurance/quality control (QA/QC), sample preservation, and sample handling are all components of this SOP.

#### 2.1 Sample Containers

Prior to collection of a sample, considerations must be given to the type of container that will be used to store and transport the sample. The type and number of containers selected is usually based on factors such as sample matrix, potential contaminants to be encountered, analytical methods requested, and the laboratory's internal quality assurance requirements. In most cases, the overriding considerations will be the analytical methodology, or the state or federal regulatory requirements because these regulations generally encompass the other factors. The sample container selected is usually based on some combination of the following criteria:

## a. Reactivity of Container Material with Sample

Choosing the proper composition of sample containers will help to ensure that the chemical and physical integrity of the sample is maintained. For sampling potentially hazardous material, glass is the recommended container type because it is chemically inert to most substances. Plastic containers are not recommended for most hazardous wastes because the potential exists for contaminants to adsorb to the surface of the plastic or for the plasticizer to leach into the sample.

In some instances, however, the sample characteristics or analytes of interest may dictate that plastic containers be used instead of glass. Because some metals species will adhere to the sides of the glass containers in an aqueous matrix, plastic bottles (e.g., nalgene) must be used for samples collected for metals analysis. A separate, plastic

container should accompany glass containers if metals analysis is to be performed along with other analyses. Likewise, other sample characteristics may dictate that glass cannot be used. For example, in the case of a strong alkali waste or hydrofluoric solution, plastic containers may be more suitable because glass containers may be etched by these compounds and create adsorptive sites on the container's surface.

#### b. Volume of the Container

The volume of sample to be collected will be dictated by the analysis being performed and the sample matrix. The laboratory must supply bottles of sufficient volume to perform the required analysis. In most cases, the methodology dictates the volume of sample material required to complete the analysis. However, individual laboratories may provide larger volume containers for various analytes to ensure sufficient quantities for duplicates or other QC checks.

To facilitate transfer of the sample from the sampler into the container and to minimize spillage and sample disturbance, wide-mouth containers are recommended. Aqueous volatile organic samples must be placed into 40-milliliter (ml) glass vials with polytetrafluoroethylene (PTFE) (e.g., TeflonTM) septums. Non-aqueous volatile organic samples should be collected in the same type of vials or in 4-ounce (oz) wide-mouth jars provided by the laboratory. These jars should have PTFE-lined screw caps.

#### c. Color of Container

Whenever possible, amber glass containers should be used to prevent photodegradation of the sample, except when samples are being collected for metals analysis. If amber containers are not available, then containers holding samples should be protected from light (i.e., place in cooler with ice immediately after filling).

#### d. Container Closures

Container closures must screw on and off the containers and form a leak-proof seal. Container caps must not be removed until the container is ready to be filled with the sample, and the container cap must be replaced (securely) immediately after filling it. Closures should be constructed of a material which is inert with respect to the sampled material, such as PTFE (e.g., TeflonTM). Alternately, the closure may be separated from the sample by a closure liner that is inert to the sample material such as PTFE sheeting. If soil or sediment samples are being collected, the threads of the container must be wiped clean with a dedicated paper towel or cloth so the cap can be threaded properly.

#### e. Decontamination of Sample Containers

Sample containers must be laboratory cleaned by the laboratory performing the analysis. The cleaning procedure is dictated by the specific analysis to be performed on the sample. Sample containers must be carefully examined to ensure that all containers appear clean. Do not mistake the preservative as unwanted residue. The bottles should not be field cleaned. If there is any question regarding the integrity of the bottle, then the laboratory must be contacted immediately and the bottle(s) replaced.

## f. Sample Bottle Storage and Transport

No matter where the sample bottles are, whether at the laboratory waiting to be packed for shipment or in the field waiting to be filled with sample, care must be taken to avoid contamination. Sample shuttles or coolers, and sample bottles must be stored and transported in clean environments. Sample bottles and clean sampling equipment must never be stored near solvents, gasoline, or other equipment that is a potential source of crosscontamination. When under chain of custody, sample bottles must be secured in locked vehicles, and custody sealed in shuttles or in the presence of authorized personnel. Information which documents that proper storage and transport procedures have been followed must be included in the field notebook and on appropriate field forms.

## 2.2 Decontamination of Sampling Equipment

Proper decontamination of all re-usable sampling equipment is critical for all sampling episodes. The SOP for Decontamination of Field Equipment and SOPs for method-specific or instrument-specific tasks must also be referred to for guidance for decontamination of various types of equipment.

## 2.3 Quality Assurance/Quality Control Samples

QA/QC samples are intended to provide control over the proper collection and tracking of environmental measurements, and subsequent review, interpretation and validation of generated analytical data. The SOPs for Collection of Quality Control Samples, for Evaluation and Validation of Data, and for Field Record Keeping and Quality Assurance/Quality Control must be referred to for detailed guidance regarding these respective procedures. SOPs for method-specific or instrument-specific tasks must also be referred to for guidance for QA/QC procedures.

#### 2.4 Sample Preservation Requirements

Certain analytical methodologies for specific analytes require chemical additives in order to stabilize and maintain sample integrity. Generally, this is accomplished under the following two scenarios:

- a. Sample bottles are preserved at the laboratory prior to shipment into the field.
- b. Preservatives are added in the field immediately after the samples are collected.

Many laboratories provide pre-preserved bottles as a matter of convenience and to help ensure that samples will be preserved immediately upon collection. A problem associated with this method arises if not enough sample could be collected, resulting in too much preservative in the sample. More commonly encountered problems with this method include the possibility of insufficient preservative provided to achieve the desired pH level or the need for additional preservation due to chemical reactions caused by the addition of sample liquids to pre-preserved bottles. The use of pre-preserved bottles is acceptable; however, field sampling teams must always be prepared to add additional preservatives to samples if the aforementioned situations occur. Furthermore, care must be exercised not to overfill sample bottles containing preservatives to prevent the sample and preservative from spilling and therefore diluting the preservative (i.e., not having enough preservative for the volume of sample).

When samples are preserved after collection, special care must be taken. The transportation and handling of concentrated acids in the field requires additional preparation and adherence to appropriate preservation procedures. All preservation acids used in the field should be trace-metal or higher-grade.

### 2.5 Sample Handling

After the proper sample bottles have been received under chain-of-custody, properly decontaminated equipment has been used to collect the sample, and appropriate preservatives have been added to maintain sample integrity, the final step for the field personnel is checking the sample bottles prior to proper packing and delivery of the samples to the laboratory.

All samples should be organized and the labels checked for accuracy. The caps should be checked for tightness and any 40-ml volatile organic compound (VOC) bottles must be checked for bubbles. Each sample bottle must be placed in an individual "zip-lock" bag to protect the label, and placed on ice. The bottles must be carefully packed to prevent breakage during transport. When several bottles have been collected for an individual sample, they should not be placed adjacent to each other in the cooler to prevent possible breakage of all bottles for a given sample. If there are any samples which are known or suspected to be highly

contaminated, these should be placed in an individual cooler under separate chain-of-custody to prevent possible cross contamination. Sufficient ice (wet or blue packs) should be placed in the cooler to maintain the temperature at 4 degrees Celsius (°C) until delivery at the laboratory. Consult the work plan to determine if a particular ice is specified as the preservation for transportation (e.g., the United States Environmental Protection Agency does not like the use of blue packs because they claim that the samples will not hold at 4°C). If additional coolers are required, then they should be purchased. The chain-of-custody form should be properly completed, placed in a "zip-lock" bag, and placed in the cooler. One copy must be maintained for the project files. The cooler should be sealed with packing tape and a custody seal. The custody seal number should be noted in the field book. Samples collected from Monday through Friday will be delivered to the laboratory within 24 hours of collection. If Saturday delivery is not available, samples collected on Friday must be delivered by Monday morning. Check the work plan to determine if certain analytes require a shorter delivery time. If overnight mail is utilized, then the shipping bill must be maintained for the files and the laboratory must be called the following day to confirm receipt.

### 3.0 EQUIPMENT AND MATERIALS

- 3.1 General equipment and materials may include, but not necessarily be limited to, the following:
  - a. Sample bottles of proper size and type with labels.
  - b. Cooler with ice (wet or blue pack).
  - c. Field notebook, appropriate field form(s), chain-of-custody form(s), custody seals.
  - d. Black pen and indelible marker.
  - e. Packing tape, "bubble wrap," and "zip-lock" bags.
  - f. Overnight (express) mail forms and laboratory address.
  - g. Health and safety plan (HASP).
  - h. Work plan/scope of work.
  - i. Pertinent SOPs for specified tasks and their respective equipment and materials.
- Preservatives for specific samples/analytes as specified by the laboratory. Preservatives must be stored in secure, spillproof glass containers with their content, concentration, and date of preparation and expiration clearly labeled.

- 3.3 Miscellaneous equipment and materials including, but not necessarily limited to, the following:
  - a. Graduated pipettes.
  - b. Pipette bulbs.
  - c. Litmus paper.
  - d. Glass stirring rods.
  - e. Protective goggles.
  - f. Disposable gloves.
  - g. Lab apron.
  - h. First aid kit.
  - i. Portable eye wash station.
  - j. Water supply for immediate flushing of spillage, if appropriate.
  - k. Shovel and container for immediate containerization of spillage-impacted soils, if appropriate.

#### 4.0 PROCEDURE

- 4.1 Examine all bottles and verify that they are clean and of the proper type, number, and volume for the sampling to be conducted.
- 4.2 Label bottles carefully and clearly with project name and number, site location, sample identification, date, time, and the sampler's initials using an indelible marker.
- 4.3 Collect samples in the proper manner (refer to specific sampling SOPs).
- 4.4 Conduct preservation activities as required after each sample has been collected. Field preservation must be done immediately and must not be done later than 30 minutes after sample collection.
- 4.5 Conduct QC sampling, as required.
- 4.6 Seal each container carefully and place in an individual "zip lock" bag.
- 4.7 Organize and carefully pack all samples in the cooler immediately after collection (e.g., bubble wrap). Insulate samples so that breakage will not occur.

- 4.8 Complete and place the chain-of-custody form in the cooler after all samples have been collected. Maintain one copy for the project file. If the cooler is to be transferred several times prior to shipment or delivery to the laboratory, it may be easier to tape the chain-of-custody to the exterior of the sealed cooler. When exceptionally hazardous samples are known or suspected to be present, this should be identified on the chain-of-custody as a courtesy to the laboratory personnel.
- 4.9 Add additional ice as necessary to ensure that it will last until receipt by the laboratory.
- 4.10 Seal the cooler with packing tape and a custody seal. Record the number of the custody seal in the field notebook and on the field form. If there are any exceptionally hazardous samples, then shipping regulations should be examined to ensure that the sample containers and coolers are in compliance and properly labeled.
- 4.11 Samples collected from Monday through Friday will be delivered to the laboratory within 24 hours of collection. If Saturday delivery is not available, samples collected on Friday must be delivered by Monday morning. Check the work plan to determine if certain analytes require a shorter delivery time.
- 4.12 Maintain the shipping bill for the project files if overnight mail is utilized and call the laboratory the following day to confirm receipt.

END OF PROCEDURE

## STANDARD OPERATING PROCEDURE 9.1 FOR DECONTAMINATION OF FIELD EQUIPMENT

Date: May 5, 2000

#### 1.0 PURPOSE

The purpose for this standard operating procedure (SOP) is to establish the guidelines for decontamination of all field equipment potentially exposed to contamination during drilling, and soil and water sampling. The objective of decontamination is to ensure that all drilling, and soil-sampling and water-sampling equipment is decontaminated (free of potential contaminants): 1) prior to being brought onsite to avoid the introduction of potential contaminants to the site; 2) between drilling and sampling events/activities onsite to eliminate the potential for cross-contamination between boreholes and/or wells; and 3) prior to the removal of equipment from the site to prevent the transportation of potentially contaminated equipment offsite.

In considering decontamination procedures, state and federal regulatory agency requirements must be considered because of potential variability between state and federal requirements and because of variability in the requirements of individual states. Decontamination procedures must be in compliance with state and/or federal protocols in order that regulatory agency(ies) scrutiny of the procedures and data collected do not result in non acceptance (invalidation) of the work undertaken and data collected.

### 2.0 PROCEDURE FOR DRILLING EQUIPMENT

The following is a minimum decontamination procedure for drilling equipment. Drilling equipment decontamination procedures, especially any variation from the method itemized below, will be documented on an appropriate field form or in the field notebook.

- 2.1 The rig and all associated equipment should be properly decontaminated by the contractor before arriving at the test site.
- 2.2 The augers, drilling casings, rods, samplers, tools, rig, and any piece of equipment that can come in contact (directly or indirectly) with the soil, will be steam cleaned onsite prior to set up for drilling to ensure proper decontamination.
- 2.3 The same steam cleaning procedures will be followed between boreholes (at a fixed on-site location[s], if appropriate) and before leaving the site at the end of the study.
- 2.4 All on-site steam cleaning (decontamination) activities will be monitored and documented by a member(s) of the staff of Roux Associates, Inc.
- 2.5 If drilling activities are conducted in the presence of thick, sticky oils (e.g., PCBs) which coat drilling equipment, then special decontamination procedures may have to be utilized before steam cleaning (e.g., hexane scrub and wash).

## STANDARD OPERATING PROCEDURE 9.1 FOR DECONTAMINATION OF FIELD EQUIPMENT

2.6 Containment of decontamination fluids may be necessary (e.g., rinseate from steam cleaning) or will be required (e.g., hexane), and disposal must be in accordance with state and/or federal procedures.

#### 3.0 PROCEDURE FOR SOIL-SAMPLING EQUIPMENT

The following is a minimum decontamination procedure for soil-sampling equipment (e.g., split spoons, stainless-steel spatulas). Soil-sampling equipment decontamination procedures, especially any variation from the method itemized below, will be documented on an appropriate field form or in the field notebook.

- 3.1 Wear disposable gloves while cleaning equipment to avoid cross-contamination and change gloves as needed.
- 3.2 Steam clean the sampler or rinse with potable water. If soil-sampling activities are conducted in the presence of thick, sticky oils (e.g., PCBs) which coat sampling equipment, then special decontamination procedures may have to be utilized before steam cleaning and washing in detergent solution (e.g., hexane scrub and wash).
- 3.3 Prepare a non-phosphate, laboratory-grade detergent solution and distilled or potable water in a clean bucket.
- 3.4 Disassemble the sampler, as necessary and immerse all parts and other sampling equipment in the solution.
- 3.5 Scrub all equipment in the bucket with a brush to remove any adhering particles.
- 3.6 Rinse all equipment with copious amounts of potable water followed by distilled or deionized water.
- 3.7 Place clean equipment on a clean plastic sheet (e.g., polyethylene)
- 3.8 Reassemble the cleaned sampler, as necessary.
- 3.9 Transfer the sampler to the driller (or helper) making sure that this individual is also wearing clean gloves, or wrap the equipment with a suitable material (e.g., plastic bag, aluminum foil.

As part of the decontamination procedure for soil-sampling equipment, state and/or federal protocols must be considered. These may require procedures above those specified as minimum for Roux Associates, Inc., such as the use of nitric acid, acetone, etc. Furthermore, the containment and proper disposal of decontamination fluids must be considered with respect to regulatory agency(ies) requirements.

## STANDARD OPERATING PROCEDURE 9.1 FOR DECONTAMINATION OF FIELD EQUIPMENT

### 4.0 PROCEDURE FOR WATER-SAMPLING EQUIPMENT

The following is a decontamination procedure for water-sampling equipment (e.g., bailers, pumps). Water-sampling equipment decontamination procedures, especially any variation from the method itemized below, will be documented on an appropriate field form or in the field notebook.

#### 4.1 Decontamination procedures for bailers follow:

- a. Wear disposable gloves while cleaning bailer to avoid cross-contamination and change gloves as needed.
- b. Prepare a non-phosphate, laboratory-grade detergent solution and potable water in a bucket.
- c. Disassemble bailer (if applicable) and discard cord in an appropriate manner, and scrub each part of the bailer with a brush and solution.
- d. Rinse with potable water and reassemble bailer.
- e. Rinse with copious amounts of distilled or deionized water.
- f. Air dry.
- g. Wrap equipment with a suitable material (e.g., clean plastic bag, aluminum foil).
- h. Rinse bailer at least three times with distilled or deionized water before use.

#### 4.2 Decontamination procedures for pumps follow:

- a. Wear disposable gloves while cleaning pump to avoid cross-contamination and change gloves as needed.
- b. Prepare a non-phosphate, laboratory-grade detergent solution and potable water in a clean bucket, clean garbage can, or clean 55-gallon drum.
- c. Flush the pump and discharge hose (if not disposable) with the detergent solution, and discard disposable tubing and/or cord in an appropriate manner.
- d. Flush the pump and discharge hose (if not disposable) with potable water.
- e. Place the pump on clear plastic sheeting.
- f. Wipe any pump-related equipment (e.g., electrical lines, cables, discharge hose) that entered the well with a clean cloth and detergent solution, and rinse or wipe with a clean cloth and potable water.

# STANDARD OPERATING PROCEDURE 9.1 FOR DECONTAMINATION OF FIELD EQUIPMENT

- g. Air dry.
- h. Wrap equipment with a suitable material (e.g., clean plastic bag).

As part of the decontamination procedure for water-sampling equipment, state and/or federal protocols must be considered. These may require procedures above those specified as minimum for Roux Associates, Inc., such as the use of nitric acid, acetone, etc. Furthermore, the containment and proper disposal of decontamination fluids must be considered with respect to regulatory agency(ies) requirements.

## **ATTACHMENT 2**

Chain of Custody Form

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# **CHAIN OF CUSTODY**

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ANALYTICAL LABORATORY													

## Remedial Investigation Work Plan 1022 Old Country Road, Plainview, New York 11803

**APPENDIX B** 

Quality Assurance Project Plan

## QUALITY ASSURANCE PROJECT PLAN

1022 Old Country Road Plainview, New York 11803

Prepared for

MORTON VILLAGE REALTY COMPANY, INC. c/o John Curran Sive Paget and Riesel P.C. 460 Park Avenue New York, New York 10022

## **ROUX ASSOCIATES, INC.**

Environmental Consulting & Management



209 Shafter Street, Islandia, New York 11749 ♦ 631-232-2600

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### **TABLES**

- Field and Laboratory QC Summary
   Laboratory Reporting Limits for Soil, Water and Soil Vapor

#### 1.0 INTRODUCTION

This Quality Assurance Project Plan (QAPP) has been prepared to describe the measures that will be taken to ensure that the data generated during performance of the Remedial Investigation (RI) at 1022 Old Country Road, Plainview, New York (Site) are of quality sufficient to meet project-specific data quality objectives (DQOs). The QAPP was prepared in accordance with the guidance provided in New York State Department of Environmental Conservation (NYSDEC) Technical Guidance DER-10 (Technical Guidance for Site Investigation and Remediation) and the United States Environmental Protection Agency's (USEPA's) Guidance for the Data Quality Objectives Process (EPA QA/G-4).

#### 2.0 BACKGROUND, OBJECTIVES, AND SCOPE

In order to achieve project objectives, Roux Associates has developed a scope of work that includes sampling of soil and groundwater. A brief overview of each element of the RI scope of work is provided below. RI sampling locations are shown in Figure 3 of the RI Work Plan.

#### **2.1 Soil**

Samples of soil will be collected and analyzed at a minimum of seven locations for the following:

• Target Compound List (TCL) volatile organic compounds (VOCs)

#### 2.2 Groundwater

Groundwater samples will be collected from three new monitoring wells installed by Roux Associates during the RI. After gauging for water level measurements, each well will be sampled and analyzed for the following:

TCL VOCs

Field parameters, including temperature, pH, conductivity, redox potential, dissolved oxygen, and turbidity will also be measured.

#### 2.3 Soil Vapor

Soil vapor samples, include sub-slab and indoor air samples will be collected and analyzed during Phase B of this RI. Each soil vapor, sub-slab and indoor air sample will be collected and analyzed for the following:

• VOCs (TO15)

#### 3.0 PROJECT ORGANIZATION

The overall management structure and a general summary of the responsibilities of project team members are presented below.

#### Project Manager

Joseph Duminuco of Roux Associates will serve as Project Manager. The Project Manager is responsible for defining project objectives and bears ultimate responsibility for the successful completion of the investigation. This individual will provide overall management for the implementation of the scope of work and will coordinate all field activities. The Project Manager is also responsible for data review/interpretation and report preparation. Activities of the Project Manager are supported by the Project Quality Assurance Coordinator.

#### Field Team Leader

Jeff Wills of Roux Associates will serve as the Field Team Leader. The Field Team Leader bears the responsibility for the successful execution of the field program, as scoped in the RI Work Plan and the Field Sampling Plan (FSP). The Field Team Leader will direct the activities of all technical staff in the field as well all subcontractors. The Field Team Leader will also assist in the interpretation of data and in report preparation. The Field Team Leader reports to the Project Manager.

#### Laboratory Project Manager

Alpha Analytical (Alpha) of Westborough, Massachusetts, has been selected to analyze the field samples for this project and will be responsible for sample container preparation, sample custody in the laboratory, and completion of the required analysis through oversight of the laboratory staff. Alpha is an Environmental Laboratory Approval Program (ELAP) certified lab as described in Section 5. The Laboratory Project Manager will ensure that quality assurance procedures are followed and that an acceptable laboratory report is prepared and submitted. The Laboratory Project Manager reports to the Field Team Leader.

#### **Quality Assurance Officer**

Charles McGuckin, P.E. of Roux Associates/Remedial Engineering will serve as the Quality Assurance Officer (QAO) for this project. The QAO is responsible for conducting reviews,

inspections, and audits to ensure that the data collection is conducted in accordance with the FSP and QAPP. The QAO's responsibilities range from ensuring effective field equipment decontamination procedures and proper sample collection to the review of all laboratory analytical data for completeness and usefulness. The QAO reports to the Project Manager and makes independent recommendations to the Field Team Leader.

### 4.0 SAMPLING PROCEDURES

Detailed discussions of sampling, decontamination, and sample handling procedures are provided in the FSP (Appendix A of the RI Work Plan).

#### 5.0 QUALITY ASSURANCE/QUALITY CONTROL

The primary intended use for the RI data is to characterize Site conditions and determine if remediation needs to be undertaken at the Site. The primary DQO of the soil and groundwater sampling programs, therefore, is that data be accurate and precise, and hence representative of the actual Site conditions. Accuracy refers to the ability of the laboratory to obtain a true value (i.e., compared to a standard) and is assessed through the use of laboratory quality control (QC) samples, including laboratory control samples and matrix spike samples, as well as through the use of surrogates, which are compounds not typically found in the environment that are injected into the samples prior to analysis. Precision refers to the ability to replicate a value, and is assessed through both field and laboratory duplicate samples.

Sensitivity is also a critical issue in generating representative data. Laboratory equipment must be of sufficient sensitivity to detect target compounds and analytes at levels below NYSDEC standards and guidelines whenever possible. Equipment sensitivity can be decreased by field or laboratory contamination of samples, and by sample matrix effects. Assessment of instrument sensitivity is performed through the analysis of reagent blanks, near-detection-limit standards, and response factors. Potential field and/or laboratory contamination is assessed through use of trip blanks, method blanks, and equipment rinse blanks (also called "field blanks").

Table 1 lists the field and laboratory QC samples that will be analyzed to assess data accuracy and precision, as well as to determine if equipment sensitivity has been compromised. Table 2 shows the reporting limits and minimum detection limits achievable by the laboratory.

All RI "assessment" analyses (i.e., TCL VOCs) will be performed in accordance with the NYSDEC Analytical Services Protocol (ASP), using USEPA SW-846 methods. Alpha has been selected to analyze the field samples collected during the RI and shall maintain a New York State Department of Health (NYSDOH) ELAP Contract Laboratory Protocol (CLP) certification (Certification Number 10602) for each of the "assessment" analyses listed in Section 2.0.

All laboratory data are to be reported in NYSDEC ASP Category B deliverables and will be delivered to NYSDEC in electronic data deliverable (EDD) format as described on NYSDEC's website (http://www.dec.ny.gov/chemical/62440.html). A Data Usability Report will be prepared

by an independent party meeting DER-10 for all data packages gene		in Section	2.2(a)1.ii	and Appendix	2B	of
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## **TABLES**

- 1. Field and Laboratory QC Summary
- 2. Laboratory Reporting Limits for Soil and Water

Table 1. Field and Laboratory QC Summary 1022 Old Country Road, Plainview, New York

QC Check Type	Minimum Frequency	Use
Field QC		
Duplicate	1 per matrix per SDG*	Precision
Trip Blank	1 per VOC cooler	Sensitivity
Equipment Rinse Blank	1 per day	Sensitivity
Laboratory QC		
Laboratory Control Sample	1 per matrix per SDG	Accuracy
Matrix Spike/Matrix Spike Duplicate/Matrix Duplicate**	1 per matrix per SDG	Accuracy/Precision
Surrogate Spike	All organics samples	Accuracy
Laboratory Duplicate	1 per matrix per SDG	Precision
Method Blank	1 per matrix per SDG	Sensitivity

#### **Notes:**

<sup>\*</sup> SDG - Sample Delivery Group - up to 20 samples. Assumes a single extraction or preparation

<sup>\*\*</sup> Provided to lab by field sampling personnel

Table 2. Volatile Organic Compound Laboratory Reporting Limits for Soil, Groundwater and Soil Vapor Samples 1022 Old Country Road, Plainview, New York

Analysis Group Description	Method Description	Method Code			
Soil Analysis	Volatile Organic Compounds (GC/MS)	8260C			
,	Analyte	CAS#	RL	MDL	Units
	Methylene chloride	75-09-2	10	1.104	ug/kg
	1,1-Dichloroethane	75-34-3	1.5	0.0856	ug/kg
	Chloroform	67-66-3	1.5	0.37	ug/kg
	Carbon tetrachloride	56-23-5	1	0.21	ug/kg
	1,2-Dichloropropane	78-87-5	3.5	0.228	ug/kg
	Dibromochloromethane	124-48-1	1	0.1536	ug/kg
	1,1,2-Trichloroethane	79-00-5	1.5	0.304	ug/kg
	Tetrachloroethene	127-18-4	1	0.1402	ug/kg
	Chlorobenzene	108-90-7	1	0.348	ug/kg
	Trichlorofluoromethane	75-69-4	5	0.388	ug/kg
	1,2-Dichloroethane	107-06-2	1	0.1134	ug/kg
	1,1,1-Trichloroethane	71-55-6	1	0.1108	ug/kg
	Bromodichloromethane	75-27-4	1	0.1732	ug/kg
	trans-1,3-Dichloropropene	10061-02-6	1	0.1208	ug/kg
	cis-1,3-Dichloropropene	10061-01-5	1	0.1176	ug/kg
	1,1-Dichloropropene	563-58-6	5	0.1414	ug/kg
	Bromoform	75-25-2	4	0.236	ug/kg
	1,1,2,2-Tetrachloroethane	79-34-5	1	0.1008	ug/kg
	Benzene	71-43-2	1	0.118	ug/kg
	Toluene	108-88-3	1.5	0.1948	ug/kg
	Ethylbenzene	100-41-4	1	0.1274	ug/kg
	Chloromethane	74-87-3	5	0.294	ug/kg
	Bromomethane	74-83-9	2	0.338	ug/kg
	Vinyl chloride	75-01-4	2	0.1174	ug/kg
	Chloroethane	75-00-3	2	0.316	ug/kg
	1,1-Dichloroethene	75-35-4	1	0.262	ug/kg
	trans-1,2-Dichloroethene	156-60-5	1.5	0.212	ug/kg
	Trichloroethene	79-01-6	1	0.125	ug/kg
	1,2-Dichlorobenzene	95-50-1	5	0.1532	ug/kg
	1,3-Dichlorobenzene	541-73-1	5	0.135	ug/kg
	1,4-Dichlorobenzene	106-46-7	5	0.1384	ug/kg
	Methyl tert butyl ether	1634-04-4	2	0.0844	ug/kg
	p/m-Xylene	179601-23-1	2	0.1978	ug/kg
	o-Xylene	95-47-6	2	0.1718	ug/kg
	Xylene (Total)	1330-20-7	2	0.1718	ug/kg
	Xylene (Total)	1330-20-7	2	0.1718	ug/kg
	cis-1,2-Dichloroethene	156-59-2	1	0.1428	ug/kg
	Dibromomethane	74-95-3	10	0.1636	ug/kg
	Styrene	100-42-5	2	0.402	ug/kg
	Dichlorodifluoromethane	75-71-8	10	0.1908	ug/kg

Table 2. Volatile Organic Compound Laboratory Reporting Limits for Soil, Groundwater and Soil Vapor Samples 1022 Old Country Road, Plainview, New York

Analysis Group Description	Method Description	Method Code			
Soil Analysis	Volatile Organic Compounds (GC/MS)	8260C			
	Analyte Description	CAS Number	RL - Limit	MDL - Limit	Units
	Acetone	67-64-1	10	1.036	ug/kg
	Carbon disulfide	75-15-0	10	1.102	ug/kg
	Bromoform	75-25-2	5	0.61	ug/kg
	Isopropylbenzene	98-82-8	5	0.19	ug/kg
	1,1,2,2-Tetrachloroethane	79-34-5	5	0.52	ug/kg
	1,3-Dichlorobenzene	541-73-1	5	0.21	ug/kg
	1,4-Dichlorobenzene	106-46-7	5	0.67	ug/kg
	1,2-Dichlorobenzene	95-50-1	5	0.24	ug/kg
	1,2-Dibromo-3-Chloropropane	96-12-8	10	4.53	ug/kg
	1,2,4-Trichlorobenzene	120-82-1	5	0.75	ug/kg
	Dibromofluoromethane	1868-53-7	5		ug/kg
	2-Butanone	78-93-3	10	0.272	ug/kg
	Vinyl acetate	108-05-4	10	0.1322	ug/kg
	4-Methyl-2-pentanone	108-10-1	10	0.244	ug/kg
	1,2,3-Trichloropropane	96-18-4	10	0.1626	ug/kg
	2-Hexanone	591-78-6	10	0.666	ug/kg
	Bromochloromethane	74-97-5	5	0.276	ug/kg
	2,2-Dichloropropane	594-20-7	5	0.226	ug/kg
	1,2-Dibromoethane	106-93-4	4	0.1744	ug/kg
	1,3-Dichloropropane	142-28-9	5	0.1452	ug/kg
	1,1,1,2-Tetrachloroethane	630-20-6	1	0.318	ug/kg
	Bromobenzene	108-86-1	5	0.208	ug/kg
	n-Butylbenzene	104-51-8	1	0.1148	ug/kg
	sec-Butylbenzene	135-98-8	1	0.122	ug/kg
	tert-Butylbenzene	98-06-6	5	0.1354	ug/kg
	o-Chlorotoluene	95-49-8	5	0.1598	ug/kg
	p-Chlorotoluene	106-43-4	5	0.1328	ug/kg
	1,2-Dibromo-3-chloropropane	96-12-8	5	0.396	ug/kg
	Hexachlorobutadiene	87-68-3	5	0.228	ug/kg
	Isopropylbenzene	98-82-8	1	0.1038	ug/kg
	p-Isopropyltoluene	99-87-6	1	0.125	ug/kg
	Naphthalene	91-20-3	5	0.1384	ug/kg
	Acrylonitrile	107-13-1	10	0.514	ug/kg
	n-Propylbenzene	103-65-1	1	0.1092	ug/kg
	1,2,3-Trichlorobenzene	87-61-6	5	0.1476	ug/kg
	1,2,4-Trichlorobenzene	120-82-1	5	0.1818	ug/kg
	1,3,5-Trimethylbenzene	108-67-8	5	0.1434	ug/kg
	1,2,4-Trimethylbenzene	95-63-6	5	0.1414	ug/kg
	1,4-Dioxane	123-91-1	100	14.42	ug/kg
	1,4-Diethylbenzene	105-05-5	4	0.1598	ug/kg
	4-Ethyltoluene	622-96-8	4	0.124	ug/kg
	1,2,4,5-Tetramethylbenzene	95-93-2	4	0.1302	ug/kg
	Ethyl ether	60-29-7	5	0.26	ug/kg
	trans-1,4-Dichloro-2-butene	110-57-6	5	0.392	ug/kg

Table 2. Volatile Organic Compound Laboratory Reporting Limits for Soil, Groundwater and Soil Vapor Samples 1022 Old Country Road, Plainview, New York

Analysis Group Description	Method Description	Method Code			
Water Analysis	Volatile Organic Compounds (GC/MS)	8260C			
	Analyte Description	CAS Number	RL - Limit	MDL - Limit	Units
	Methylene chloride	75-09-2	2.5	0.7	ug/l
	1,1-Dichloroethane	75-34-3	2.5	0.7	ug/l
	Chloroform	67-66-3	2.5	0.7	ug/l
	Carbon tetrachloride	56-23-5	0.5	0.134	ug/l
	1,2-Dichloropropane	78-87-5	1	0.133	ug/l
	Dibromochloromethane	124-48-1	0.5	0.149	ug/l
	1,1,2-Trichloroethane	79-00-5	1.5	0.5	ug/l
	Tetrachloroethene	127-18-4	0.5	0.181	ug/l
	Chlorobenzene	108-90-7	2.5	0.7	ug/l
	Trichlorofluoromethane	75-69-4	2.5	0.7	ug/l
	1,2-Dichloroethane	107-06-2	0.5	0.132	ug/l
	1,1,1-Trichloroethane	71-55-6	2.5	0.7	ug/l
	Bromodichloromethane	75-27-4	0.5	0.192	ug/l
	trans-1,3-Dichloropropene	10061-02-6	0.5	0.164	ug/l
	cis-1,3-Dichloropropene	10061-01-5	0.5	0.144	ug/l
	1,1-Dichloropropene	563-58-6	2.5	0.7	ug/l
	Bromoform	75-25-2	2	0.65	ug/l
	1,1,2,2-Tetrachloroethane	79-34-5	0.5	0.144	ug/l
	Benzene	71-43-2	0.5	0.159	ug/l
	Toluene	108-88-3	2.5	0.7	ug/l
	Ethylbenzene	100-41-4	2.5	0.7	ug/l
	Chloromethane	74-87-3	2.5	0.7	ug/l
	Bromomethane	74-83-9	2.5	0.7	ug/l
	Vinyl chloride	75-01-4	1	0.33	ug/l
	Chloroethane	75-00-3	2.5	0.7	ug/l
	1,1-Dichloroethene	75-35-4	0.5	0.142	ug/l
	trans-1,2-Dichloroethene	156-60-5	2.5	0.7	ug/l
	Trichloroethene	79-01-6	0.5	0.175	ug/l
	1,2-Dichlorobenzene	95-50-1	2.5	0.7	ug/l
	1,3-Dichlorobenzene	541-73-1	2.5	0.7	ug/l
	1,4-Dichlorobenzene	106-46-7	2.5	0.7	ug/l
	Methyl tert butyl ether	1634-04-4	2.5	0.7	ug/l
	p/m-Xylene	179601-23-1	2.5	0.7	ug/l
	o-Xylene	95-47-6	2.5	0.7	ug/l
	Xylene (Total)	1330-20-7	2.5	0.7	ug/l
	Xylene (Total)	1330-20-7	2.5	0.7	ug/l
	cis-1,2-Dichloroethene	156-59-2	2.5	0.7	ug/l
	Dibromomethane	74-95-3	5	1	ug/l
	1,2,3-Trichloropropane	96-18-4	2.5	0.7	ug/l

Table 2. Volatile Organic Compound Laboratory Reporting Limits for Soil, Groundwater and Soil Vapor Samples 1022 Old Country Road, Plainview, New York

Analysis Group Description	Method Description	Method Code			
Water Analysis	Volatile Organic Compounds (GC/MS)	8260C			
-	Analyte Description	CAS Number	RL - Limit	MDL - Limit	Units
	Acrylonitrile	107-13-1	5	1.5	ug/l
	Styrene	100-42-5	2.5	0.7	ug/l
	Dichlorodifluoromethane	75-71-8	5	1	ug/l
	Acetone	67-64-1	5	1.46	ug/l
	Carbon disulfide	75-15-0	5	1	ug/l
	2-Butanone	78-93-3	5	1.94	ug/l
	Vinyl acetate	108-05-4	5	1	ug/l
	4-Methyl-2-pentanone	108-10-1	5	1	ug/l
	2-Hexanone	591-78-6	5	1	ug/l
	Bromochloromethane	74-97-5	2.5	0.7	ug/l
	2,2-Dichloropropane	594-20-7	2.5	0.7	ug/l
	1,2-Dibromoethane	106-93-4	2	0.65	ug/l
	1,3-Dichloropropane	142-28-9	2.5	0.7	ug/l
	1,1,1,2-Tetrachloroethane	630-20-6	2.5	0.7	ug/l
	Bromobenzene	108-86-1	2.5	0.7	ug/l
	n-Butylbenzene	104-51-8	2.5	0.7	ug/l
	sec-Butylbenzene	135-98-8	2.5	0.7	ug/l
	tert-Butylbenzene	98-06-6	2.5	0.7	ug/l
	o-Chlorotoluene	95-49-8	2.5	0.7	ug/l
	p-Chlorotoluene	106-43-4	2.5	0.7	ug/l
	1,2-Dibromo-3-chloropropane	96-12-8	2.5	0.7	ug/l
	Hexachlorobutadiene	87-68-3	2.5	0.7	ug/l
	Isopropylbenzene	98-82-8	2.5	0.7	ug/l
	p-Isopropyltoluene	99-87-6	2.5	0.7	ug/l
	Naphthalene	91-20-3	2.5	0.7	ug/l
	n-Propylbenzene	103-65-1	2.5	0.7	ug/l
	1,2,3-Trichlorobenzene	87-61-6	2.5	0.7	ug/l
	1,2,4-Trichlorobenzene	120-82-1	2.5	0.7	ug/l
	1,3,5-Trimethylbenzene	108-67-8	2.5	0.7	ug/l
	1,2,4-Trimethylbenzene	95-63-6	2.5	0.7	ug/l
	1,4-Dioxane	123-91-1	250	41.1	ug/l
	1,4-Diethylbenzene	105-05-5	2	0.7	ug/l
	4-Ethyltoluene	622-96-8	2	0.7	ug/l
	1,2,4,5-Tetramethylbenzene	95-93-2	2	0.65	ug/l
	Ethyl ether	60-29-7	2.5	0.63	ug/l
	trans-1,4-Dichloro-2-butene	110-57-6	2.5	0.7	ug/l

Table 2. Volatile Organic Compound Laboratory Reporting Limits for Soil, Groundwater and Soil Vapor Samples 1022 Old Country Road, Plainview, New York

Analysis Group Description	Method Description	Method Code			
Soil Vapor Analysis	Volatile Organic Compounds (GC/MS)	TO-15			
	Analyte Description	CAS Number	RL - Limit	MDL - Limit	Units
	Acetone	67-64-1	0.64	0.33	ug/m3
	1,3-Butadiene	106-99-0	0.60	0.10	ug/m3
	Benzene	71-43-2	0.86	0.11	ug/m3
	Bromodichloromethane	75-27-4	1.8	0.26	ug/m3
	Bromoform	75-25-2	2.8	0.49	ug/m3
	Bromomethane	74-83-9	1.0	0.17	ug/m3
	Bromoethene	593-60-2	1.2	0.20	ug/m3
	Benzyl Chloride	100-44-7	1.4	0.32	ug/m3
	Carbon disulfide	75-15-0	0.84	0.13	ug/m3
	Chlorobenzene	108-90-7	1.2	0.21	ug/m3
	Chloroethane	75-00-3	0.71	0.15	ug/m3
	Chloroform	67-66-3	1.3	0.16	ug/m3
	Chloromethane	74-87-3	0.56	0.21	ug/m3
	3-Chloropropene	107-05-1	0.85	0.16	ug/m3
	2-Chlorotoluene	95-49-8	1.4	0.22	ug/m3
	Carbon tetrachloride	56-23-5	1.7	0.21	ug/m3
	Cyclohexane	110-82-7	0.93	0.12	ug/m3
	1,1-Dichloroethane	75-34-3	1.1	0.15	ug/m3
	1,1-Dichloroethylene	75-35-4	1.1	0.27	ug/m3
	1,2-Dibromoethane	106-93-4	2.1	0.28	ug/m3
	1,2-Dichloroethane	107-06-2	1.1	0.12	ug/m3
	1,2-Dichloropropane	78-87-5	1.2	0.18	ug/m3
	1,4-Dioxane	123-91-1	0.97	0.58	ug/m3
	Dichlorodifluoromethane	75-71-8	1.3	0.20	ug/m3
	Dibromochloromethane	124-48-1	2.3	0.43	ug/m3
	trans-1,2-Dichloroethylene	156-60-5	1.1	0.37	ug/m3
	cis-1,2-Dichloroethylene	156-59-2	1.1	0.12	ug/m3
	cis-1,3-Dichloropropene	10061-01-5	1.2	0.15	ug/m3
	m-Dichlorobenzene	541-73-1	1.6	0.26	ug/m3
	o-Dichlorobenzene	95-50-1	1.6	0.23	ug/m3
	p-Dichlorobenzene	106-46-7	1.6	0.29	ug/m3
	trans-1,3-Dichloropropene	10061-02-6	1.2	0.15	ug/m3
	Ethanol	64-17-5	1.3	0.41	ug/m3
	Ethylbenzene	100-41-4	1.2	0.20	ug/m3
	Ethyl Acetate	141-78-6	0.97	0.29	ug/m3
	4-Ethyltoluene	622-96-8	1.3	0.21	ug/m3
	Freon 113	76-13-1	2.1	0.41	ug/m3

Table 2. Volatile Organic Compound Laboratory Reporting Limits for Soil, Groundwater and Soil Vapor Samples 1022 Old Country Road, Plainview, New York

Analysis Group Description	Method Description	Method Code			
Soil Vapor Analysis	Volatile Organic Compounds (GC/MS)	TO-15			
	Analyte Description	CAS Number	RL - Limit	MDL - Limit	Units
	Freon 114	76-14-2	1.9	0.29	ug/m3
	Heptane	142-82-5	1.1	0.12	ug/m3
	Hexachlorobutadiene	87-68-3	2.9	0.73	ug/m3
	Hexane	110-54-3	0.95	0.20	ug/m3
	2-Hexanone	591-78-6	1.1	0.35	ug/m3
	Isopropyl Alcohol	67-63-0	0.66	0.22	ug/m3
	Methylene chloride	75-09-2	0.94	0.63	ug/m3
	Methyl ethyl ketone	78-93-3	0.80	0.16	ug/m3
	Methyl Isobutyl Ketone	108-10-1	1.1	0.23	ug/m3
	Methyl Tert Butyl Ether	1634-04-4	0.97	0.19	ug/m3
	Methylmethacrylate	80-62-6	1.1	0.19	ug/m3
	Naphthalene	91-20-3	1.4	0.41	ug/m3
	Propylene	115-07-1	1.2	0.11	ug/m3
	Styrene	100-42-5	1.1	0.19	ug/m3
	1,1,1-Trichloroethane	71-55-6	1.5	0.18	ug/m3
	1,1,2,2-Tetrachloroethane	79-34-5	1.9	0.36	ug/m3
	1,1,2-Trichloroethane	79-00-5	1.5	0.26	ug/m3
	1,2,4-Trichlorobenzene	120-82-1	2.0	0.60	ug/m3
	1,2,4-Trimethylbenzene	95-63-6	1.3	0.19	ug/m3
	1,3,5-Trimethylbenzene	108-67-8	1.3	0.19	ug/m3
	2,2,4-Trimethylpentane	540-84-1	1.3	0.16	ug/m3
	Tertiary Butyl Alcohol	75-65-0	0.82	0.18	ug/m3
	Tetrachloroethylene	127-18-4	0.36	0.34	ug/m3
	Tetrahydrofuran	109-99-9	0.80	0.19	ug/m3
	Toluene	108-88-3	1.0	0.15	ug/m3
	Trichloroethylene	79-01-6	0.28	0.21	ug/m3
	Trichlorofluoromethane	75-69-4	1.5	0.21	ug/m3
	Vinyl chloride	75-01-4	0.69	0.10	ug/m3
	Vinyl Acetate	108-05-4	0.95	0.46	ug/m3
	m,p-Xylene	-	1.2	0.40	ug/m3
	o-Xylene	95-47-6	1.2	0.20	ug/m3
	Xylenes (total)	1330-20-7	1.2	0.20	ug/m3

## Remedial Investigation Work Plan 1022 Old Country Road, Plainview, New York 11803

**APPENDIX C** 

Site Health and Safety Plan

## **HEALTH AND SAFETY PLAN**

1022 Old Country Road Plainview, New York 11803

Prepared for

MORTON VILLAGE REALTY COMPANY, INC. % John Curran Sive Paget and Riesel P.C. 460 Park Avenue New York, New York 10022

## **ROUX ASSOCIATES, INC.**

Environmental Consulting & Management



209 Shafter Street, Islandia, New York 11749 ♦ 631-232-2600

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- 2. Hospital Route Map
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#### **APPENDICES**

- A. Job Safety and Health (OSHA) Poster
- B. Material Safety Data Sheets (MSDS)
- C. Job Safety Analysis
- D. Heat and Cold Stress Guidelines
- E. Health and Safety Briefing/Tailgate Meeting Form
- F. Medical Data Form
- G. Generic Community Air Monitoring Plan
- H. Accident Report and Investigation Form
- I. Acord Automobile Loss Form
- J. Near Loss Reporting Form
- K. OSHA Log of Occupational Injuries and Illnesses

### **APPROVALS**

By their signature, the undersigned certify that this Health and Safety Plan (HASP) is approved and will be utilized at the project site located at 1022 Old Country Road, Plainview, New York.

	_
Ray Fitzpatrick	Date
Office Health and Safety Manager	
Roux Associates, Inc.	
Joseph Duminuco	Date
Project Principal	
Roux Associates, Inc.	
Jeff Wills	Date
Project Manager/SHSO	
Roux Associates, Inc.	

#### 1.0 INTRODUCTION

This Site-specific Health and Safety Plan (HASP) has been prepared in accordance with 29 CFR 1910.120 Occupational Safety and Health Administration (OSHA) Hazardous Waste Operations and Emergency Response (HAZWOPER) and Roux Associates, Inc. (Roux Associates) Standard Operating Procedures (SOPs) and other OSHA requirements for job safety and health protection (Appendix A). It addresses all activities described below that are associated with the property located at 1022 Old Country Road, in the Hamlet of Plainview, New York (Site). The location of the Site is presented in Figure 1. The HASP will be implemented by the designated Site Health and Safety Officer (SSO) during work at the Site. The HASP attempts to identify all potential hazards at the Site; however, Site conditions are dynamic and new hazards may appear constantly. Personnel must remain alert to existing and potential hazards as Site conditions change and protect themselves accordingly.

Compliance with this HASP is required of all persons and subcontractors who perform fieldwork at the Site. The contents of this HASP may change or undergo revision based upon additional information made available to health and safety personnel, monitoring results, or changes in the technical scope of work. Any changes proposed must be reviewed and approved by the Office Health and Safety Manager (OHSM), with the SSO implementing the changes to the HASP.

Prior to performing work each task should be evaluated to determine the appropriate procedures that need to be followed.

#### 1.1 Scope of Work

In general, the tasks will include the following:

• Implementation of Remedial Investigation (RI) activities consisting of monitoring well installation, groundwater sampling, soil borings/sampling, installation of soil vapor and sub-slab monitoring points, and soil vapor sampling (including sub-slab and indoor air sampling).

Any change in scope will require a revision of this HASP to address any new hazards.

#### 2.0 EMERGENCY INFORMATION AND NOTIFICATION

Multiple emergency services may be obtained by calling 911. More specific numbers for local services are listed below.

Emergency Medical Service	911
Police: Nassau County Police Department	911
Fire: Plainview Fire Department	911
Hospital: North Shore University Hospital	516-719-3000
National Response Center	800-424-8802
Poison Control Center	800-222-1222
CHEMTREC	800-262-8200
Center for Disease Control	800-311-3435
USEPA (Region II)	212-637-5000
NYSDEC Emergency Spill Response	800-457-7362
First Care & Occupational Health Clinic	631-435-0110

Directions and maps to the Hospital and the Clinic are provided in Figures 2 and 3, respectively.

#### 2.1 Notification

As soon as first aid and/or emergency response needs have been met, the following parties are to be contacted by telephone: (Direct contact, no phone messages).

			Office:	<u>Cell</u> :
1.	Project Principal:	Joseph Duminuco	(631) 232-2600	(631) 921-6279
2.	Project Manager/SSO:	Jeff Wills	(631) 232-2600	(516) 637-0213
3.	Office Health and Safety Manager:	Ray Fitzpatrick	(631) 232-2600	(631) 484-1168
4.	Corporate Health & Safety Manager:	Joseph Gentile	(856) 423-8800	(610) 844-6911
5.	Office Manager:	Brian Morrissey	(631) 232-2600	(631) 921-6355

Accident reporting guidelines are outlined in section 13.5 of this HASP.

#### 3.0 HEALTH AND SAFETY PERSONNEL

This section briefly describes all Site personnel and their health and safety responsibilities for the RI work to be implemented at the Site. All personnel are responsible for ensuring compliance with the HASP.

#### **Project Principal (PP) – Joseph Duminuco – Roux Associates**

- Has the overall responsibility for the health and safety of Site personnel.
- Ensures that adequate resources are provided to the field health and safety staff to carry out their responsibilities as outlined below.

#### Office Health and Safety Manager (OHSM) – Ray Fitzpatrick – Roux Associates

- Implements the HASP.
- Performs or oversees site-specific training and approves revised or new safety protocols or field operations.
- Coordinates revisions of this HASP with Project Principal.
- Responsible for the development of new task safety protocols and procedures and resolution of any outstanding safety issues which may arise during the performance of site work.
- Review and approve all health and safety training and medical surveillance records for personnel and subcontractors.

#### Project Manager/Site Safety and Health Officer (PM/SSO) – Jeff Wills – Roux Associates

- Directs and coordinates health and safety monitoring activities.
- Ensures that field teams utilize proper personal protective equipment.
- Conducts initial onsite specific training prior to personnel and/or subcontractors commencing work.
- Conducts and documents daily pre-job safety briefings.
- Ensures that field team members comply with this HASP.
- Completes and maintains all accident investigation and reporting forms.
- Notifies PP and OHSM of all accidents/incidents.

- Notifies PP of daily field operations and work progress, who will then communicate at the end of the day to the designated representative the following:
  - 1. End of day tasks completed
  - 2. Next day's planned activities
  - 3. Third party issues
  - 4. Change of Plans approvals
- Determines upgrade or downgrade of personal protective equipment (PPE) based on Site conditions and/or real time monitoring results.
- Ensures that monitoring instruments are calibrated daily or as manufacturers suggested instructions determine.
- Submits and maintains health and safety field log books, daily safety logs, training logs, air monitoring result reports, weekly safety report.

#### **Field Personnel and Subcontractors**

- Report any unsafe or potentially hazardous conditions to the PM/SSO.
- Maintain knowledge of the information, instructions, and emergency response actions contained in the HASP.
- Comply with rules, regulations, and procedures as set forth in this HASP and any revisions, which are instituted.
- Prevent admittance to work Site by unauthorized personnel.

In the case that there is a change in any of the above personnel, all onsite personnel will be notified of the change. The new responsible party shall review and sign that they have been given a documented verbal full HASP review by Roux Associates and are aware of their responsibilities as outlined in this HASP.

#### 4.0 SITE LOCATION, DESCRIPTION, AND HISTORY

Descriptions of the Site and surrounding property usage are included in the following sections. The location of the Site is presented in Figure 1.

#### **4.1 Property Location and Description**

The Site is identified as Block 555, Lots 10, 86/89 and 88 on the Nassau County tax maps. The Site is located on the north side of Old Country Road between the corners of Rex Place and Lester Place. The Site consists of one two-story building and three one-story buildings and paved parking lot encompassing a total of 9.936 acres and is bordered by Knowles Street to the north, Old Country Road to the south, Lester Place to the east and Rex Place to the west. The surface elevation of the property is approximately 145 feet. Topography of the property slopes slightly to the south.

The surrounding properties to the north, east, and west are all residential properties. To the south of the Site, there are both residential properties as well as the Plainview-Old Bethpage Public Library.

The Site was previously used for agricultural use until developed as a shopping center in 1956. The Site has been improved with commercial developments since the late 1950s. Past commercial uses included a dry cleaning facility (former Morton Village Cleaners) from the late 1950s to 2007. Based on previous environmental investigations, the following Areas of Concern (AOCs) were identified at the Site:

- Former Morton Village Cleaners operations that were conducted between late 1950's and 2007.
- One historical Gas Station occupied the property adjacent to the southeast corner of the Site (Section 12, Block 555, Lot 6).
- Significant staining and standing oil observed in an elevator pit for the freight elevator in one of the existing Buildings (Building C) during a September 2008 site visit.

#### 5.0 WASTE DESCRIPTION/CHARACTERIZATION

#### 5.1 General

The following information is presented in order to identify the types of materials that may be encountered at the Site. The detailed information on these materials was obtained from:

- Sax's Dangerous Properties of Industrial Materials Lewis Eight Edition
- Chemical Hazards of the Workplace Proctor/Hughes
- Condensed Chemical Dictionary Hawley
- Rapid Guide to Hazardous Chemical in the Workplace Lewis 1990
- NIOSH Pocket Guide to Chemical Hazards 2005
- ACGIH TLV Values and Biological Exposure Indices
- OSHA 29 CFR 1910.1000

#### **5.2** Chemical Data Sheets

Several chemicals that may potentially be present in soils and groundwater at the Site, based on previous soil, soil vapor and groundwater sampling results and historic operations conducted at the Site that have been identified. The Summary of Toxicological Data is found in Table 1 and is provided for review of chemicals that may be encountered. The Summary of Toxicological Data Sheets provides information such as the chemicals characteristics, health hazards, protection, and exposure limits. Material Safety Data Sheets (MSDSs) for products that have been identified at the Site are available for review by project personnel (Appendix B).

#### **5.2.1** Contaminants of Concern

Soil and groundwater contaminants that may be encountered during drilling and sampling activities include both organic and inorganic compounds. Prior investigations at the Site have indicated detection of VOCs, Polycyclic Aromatic Hydrocarbons (PAHs) and metals in soil. Tetrachloroethene and trichloroethene were detected in groundwater and soil vapor samples. The toxicological, physical, and chemical properties of potential contaminants are presented in Table 1, and identified contaminants are presented in Appendix B.

#### 6.0 HAZARD ASSESSMENT

The potential to encounter chemical hazards is dependent upon the work activity performed (intrusive versus non-intrusive) and the duration and location of the work activity. Such hazards could include inhalation and/or skin contact with chemicals/gases that could cause: dermatitis, skin burns, being overcome by vapors or asphyxiation.

Physical hazards that may be encountered during Site work include heat and cold stress, being crushed, head injuries, punctures, cuts, falls, electrocution, bruises and other physical hazards due to motor vehicle operation, equipment use and power tools

Biological hazards may exist during Site activities. These hazards include exposure to insect bites/stings, animals and animal wastes, mold and blood borne pathogens.

Prior to the beginning of each new phase of work, a job safety analysis (JSA) (Appendix C) will be prepared by the PM/SSO with assistance from the OHSM. The analysis will address the hazards for each activity performed in the phase and will present the procedures and safeguards necessary to eliminate the hazards or reduce the risk. JSAs for each task will be reviewed with onsite personnel at each morning tailgate meeting and as tasks change throughout the day.

#### **6.1** Chemical Hazards

The potential for personnel and subcontractors to come in contact with chemical hazards may occur during the following tasks:

- Installation and sampling of soil borings, and
- Installation, gauging, purging and sampling of temporary monitoring wells.

For chronic and acute toxicity data, refer to Summary of Toxicological Data Sheets (Table 1) and MSDSs (Appendix B) for further details on compound characteristics.

#### **6.1.1 Exposure Pathways**

Exposure to these compounds during ongoing activities may occur through inhalation of contaminated dust particles, inhalation of VOCs and SVOCs, dermal absorption, and accidental ingestion of the contaminant by either direct or indirect cross-contamination activities.

#### **6.1.2 Operational Action Levels**

A decision-making protocol for an upgrade in levels of protection and/or withdrawal of personnel from an area based on exposure levels is outlined in Table 2.

#### **6.1.3 Additional Precautions**

Dermal absorption or skin contact with chemical compounds is possible during intrusive activities and while gauging, purging or sampling a monitoring well at the Site. The use of PPE in accordance with Section 8.2 and strict adherence to proper decontamination procedures should significantly reduce the risk of skin contact.

The potential for accidental ingestion of potentially hazardous chemicals is expected to be remote, when good hygiene practices are used.

#### **6.2 Physical Hazards**

A variety of physical hazards may be present during Site activities. These hazards are similar to those associated with any investigation-type project and include equipment operation and hazardous walking and working surfaces. The referenced hazards are not unique and are generally familiar to most hazardous waste site workers at environmental sites. Task-specific safety requirements for each phase will be covered during safety briefings.

#### **6.2.1 Heat Stress**

Heat stress is a significant potential hazard, associated with the use of protective equipment in a hot weather environment. The human body is designed to function at a certain internal temperature. When metabolism or external sources (fire or hot summer day) cause the body temperature to rise, the body seeks to protect itself by triggering cooling mechanisms. The PM/SSO will monitor the air temperature (as described later in this section) to determine potential adverse effects the weather can cause onsite personnel. Excess heat is dissipated by two means:

- Changes in blood flow to dissipate heat by convection, which can be seen as "flushing" or reddening of the skin in extreme cases.
- Perspiration, the release of water through skin and sweat glands. While working in hot environments, evaporation of perspiration is the primary cooling mechanism.

Protective clothing worn to guard against chemical contact effectively stops the evaporation of perspiration. Thus the use of protective clothing increases heat stress problems.

The major disorders due to heat stress are heat cramps, heat exhaustion, and heat stroke. Heat cramps are painful spasms, which occur in the skeletal muscles of workers who sweat profusely in the heat and drink large quantities of water, but fail to replace the bodies lost salts or electrolytes. Drinking water while continuing to lose salt tends to dilute the body's extracellular fluids. Soon water seeps by osmosis into active muscles and causes pain. Muscles fatigued from work are usually most susceptible to cramps.

Extreme weakness or fatigue, dizziness, nausea, and headache characterize heat exhaustion. In serious cases, a person may vomit or lose consciousness. The skin is clammy and moist, complexion pale or flushed, and body temperature normal or slightly higher than normal. Treatment is rest in a cool place and replacement of body water lost by perspiration. Mild cases may recover spontaneously with this treatment; severe cases may require care for several days. There are no permanent effects. As first aid treatment, the person shall be moved to a cool place. Body heat should be reduced artificially, but not too rapidly, by soaking the person's clothes in water and fanning them.

Heat stroke is considered a medical emergency and is caused by the breakdown of the body's regulating mechanisms. The skin is very dry and hot with red mottled or bluish appearance. Unconsciousness, mental confusion, or convulsions may occur. Without quick and adequate treatment, the result can be death or permanent brain damage.

Steps that can be taken to reduce heat stress are:

- Acclimate the body. Allow a period of adjustment to make further heat exposure endurable.
- Drink more liquids to replace the body water lost during sweating.
- Rest is necessary and should be conducted under the direction of the PM/SSO.
- Wear personal cooling devices. These are two basic designs; units with pockets for holding frozen packets and units that circulate fluid from a reservoir through tubes to different parts of the body. Both designs can be in the form of a vest, jacket, or coverall. Some circulating units also have a cap for cooling the head.

Heat stress is a significant hazard associated with using protective equipment in hot weather environments. Local weather conditions may produce conditions, which will require restricted work schedules in order to protect employees.

Appendix D contains procedures for heat stress; these will be used as a guideline and to provide additional information.

#### **6.2.2 Cold Stress**

Cold temperatures are a significant potential hazard. Examples of cold temperature hazards are frostbite and hypothermia.

Frostbite is the most common injury resulting from exposure to cold. The extremities of the body are most often affected. The signs of frostbite are:

- The skin turns white or grayish-yellow.
- Pain is sometimes felt early but subsides later. Often there is no pain.
- The affected parts feel intensely cold and numb.

Hypothermia is characterized by shivering, numbness, drowsiness, muscular weakness, and a low internal body temperature when the body feels extremely warm. This can lead to unconsciousness and death. With both frostbite and hypothermia, the affected areas need to be warmed quickly. Immersion in warm water is an effective means of warming the affected areas quickly. In such cases, medical assistance will be sought.

To prevent these effects from occurring, persons working in the cold shall wear adequate clothing and reduce the time spent in the cold area. The field PM/SSO is responsible for determining appropriate time personnel shall spend in adverse weather conditions and will monitor this.

Appendix D, which contains the Heat and Cold Stress Guidelines, provides additional information.

# **6.3 Biological Hazards**

The biological hazards, which have the potential to cause adverse health effects, are from exposure to domestic flies, mosquitoes, insects, animals and animal wastes, mold and bloodborne pathogens.

#### **6.3.1 Insect Stings**

Stings from insects are often painful, cause swelling and can be fatal if a severe allergic reaction such as anaphylactic shock occurs. If a sting occurs, the stinger should be scraped out of the skin, opposite of the sting direction. The area should be washed with soap and water followed by application of an ice pack.

If the victim has a history of allergic reaction, he shall be taken to the nearest medical facility. If the victim has medication to reverse the effects of the sting, it should be taken immediately.

If the victim experiences a severe reaction, a constricting band should be placed between the sting and the heart. The bitten area should be kept below the heart if possible. A physician shall be contacted immediately for further instructions.

#### **6.3.2** Animals and Animal Wastes

Due to the site location within a Suburban area, there lies the potential for various wildlife at the site, including, but not limited to, pigeons, bats, mice, rats, squirrels, raccoons, and feral cats. Certain animals can represent significant sources (vectors) of disease transmission. Precautions to avoid or minimize potential contact with (biting) animals (such as some of the above listed) or animal waste and/or deceased animals should be considered prior to all field activities. Rats, squirrels, raccoons, feral cats, and other wild animals can inflict painful bites which can also cause disease (as in the case of rabid animals). Site personnel should avoid contact with any of the above.

If contact occurs, be sure to clean the area thoroughly with soap and water as soon as possible. If a bite occurs, the area shall be cleaned thoroughly immediately with soap and water and medical attention shall be sought.

### **6.3.3 Bloodborne Pathogens**

The majority of the occupational tasks onsite will not involve a significant risk of exposure to blood, blood components, or body fluids. The highest risk of acquiring any bloodborne pathogen for onsite employees will be following an injury. When administering first aid care, there are potential hazards associated with bloodborne pathogens that cause diseases such as Human Immunodeficiency Virus (HIV), Hepatitis B (HBV), Hepatitis A (HAV), Hepatitis C (HCV), or the Herpes Simplex Virus (HSV). An employee who has not received the appropriate certification should never execute first aid and/or CPR.

In order to minimize any potential pathogen exposure, all employees should use the hand washing facilities on a regular basis. Additionally, the following universal precautions shall be followed to prevent further potential risk:

- Direct skin or mucous membrane contact with blood shall be avoided.
- Open skin cuts or sores shall be covered to prevent contamination from infectious agents.
- Body parts shall be washed immediately after contact with blood or body fluids that might contain blood, even when gloves or other barriers have been used.
- Gloves and disposable materials used to clean spilled blood shall be properly disposed of in an approved hazardous waste container.
- First aid responders shall wear latex or thin mil nitrile gloves when performing any procedure risking contact with blood or body substances.
- Safety glasses with attached side shields will be worn to protect the eyes from splashing or aerosolization of body fluids.
- A CPR mask will be worn when performing CPR to avoid mouth-to-mouth contact.
- Appropriate work gloves will be worn to minimize the risk of injury to the hands and fingers when working on all equipment with sharp or rough edges.
- Never pick up broken glass or possible contaminated material with your unprotected hands.
- Never handle wildlife (living or deceased) encountered onsite.

# **6.4 Hazard Assessment**

Task	Hazards	Risk of Exposure
Installation, gauging, purging and sampling of Groundwater	Inhalation/Skin Contact	Moderate/High
Monitoring Wells	Heat Stress/Cold Stress	Moderate
	Physical Injury	Moderate
Installation and sampling of Soil Borings	Inhalation/Skin Contact	Moderate/High
	Heat Stress/Cold Stress	Moderate
	Physical Injury	Moderate
Installation and sampling of soil vapor and sub-slab points and sampling of indoor air	Inhalation/Skin Contact	Moderate/High
	Heat Stress/Cold Stress	Moderate
	Physical Injury	Moderate

#### 7.0 TRAINING

# 7.1 General Health and Safety Training

In accordance with Roux Associates' corporate policies, and pursuant to 29 CFR 1910.120, hazardous waste site workers shall, at the time of the job assignment, have received a minimum of 40 hours of initial health and safety training for hazardous waste site operations. As a minimum, the training shall have consisted of instruction in the topics outlined in the above reference. Personnel who have not met the requirements for initial training will not be allowed to work in any Site activities in which they may be exposed to hazards (chemical or physical).

Completion of a 40-hour Health and Safety Training Course for Hazardous Waste Operations or an approved equivalent will fulfill the requirements of this section.

Roux Associates' PM/SSO has the responsibility of ensuring that personnel assigned to this project comply with these requirements.

# 7.2 Annual Eight-Hour Refresher Training

Current, annual 8-hour refresher training will be required of all hazardous waste site field personnel in order to maintain their qualifications for fieldwork. The following topics will be reviewed; toxicology, respiratory protection, medical surveillance, decontamination procedures, and personal protective clothing. In addition, topics deemed necessary by Roux Associates' Health and Safety Director may be added to the above list.

#### 7.3 Site-Specific Training

Site personnel will receive documented training that will specifically address the activities, procedures, monitoring and equipment for Site operations. It will include Site and facility layout, hazards, first aid equipment locations and emergency services at the Site, and will highlight all provisions contained within this HASP. This training will also allow field workers to clarify anything they do not understand and to reinforce their responsibilities regarding safety and operations for their particular activity.

# 7.4 Onsite Safety Meetings

Daily-documented (Appendix E) pre-work safety meetings will be presented each morning to discuss the scope-of-work for that day, potential safety concerns and control measures for those identified safety hazards as per the JSAs (Appendix C) for the upcoming activities.

The briefings will also provide a forum to facilitate conformance with safety requirements and to identify performance deficiencies related to safety during daily activities or as a result of safety audits by Roux Associates or other involved parties.

#### 7.5 First Aid and CPR

The PM/SSO will identify those individuals having first aid and CPR training in order to ensure that emergency medical treatment is available during field activities. The training will be consistent with the requirements of the American Red Cross Association and, as applicable, the American Heart Association. Certification and appropriate training documentation will be kept with the Site personnel records.

# **7.6 Additional Training / Procedures**

The OHSM may require additional or specialized training throughout the project. Such training shall be in the safe operation of heavy or power tool equipment or hazard communication training or other topic deemed Site appropriate.

#### 8.0 MEDICAL SURVEILLANCE PROCEDURES

#### 8.1 General

A Medical Surveillance Program has been established as part of this plan and is included in Appendix F. Roux Associates and subcontractor personnel performing field work at the Site are required to have passed a complete medical surveillance examination in accordance with 29 CFR 1910.120(f). A physician's medical release for work will be confirmed by the PM/SSO before an employee can begin Site activities. Such examinations shall include a statement as to the worker's present health status, the ability to work in a hazardous environment (including any required PPE, which may be used during temperature extremes), and the worker's ability to wear respiratory protection.

In the event that personal medical information is needed for emergency treatment, information will be made available to the treating health care professional through Roux Associates' Human Resources Department and the OHSM.

# 9.0 SITE CONTROL, PERSONAL PROTECTIVE EQUIPMENT, AND COMMUNICATIONS

A modified Site control approach may be utilized since activities will be limited to site inspection and groundwater sampling. If additional work is necessary, the following four-zone approach will be used in order to prevent the spread of contamination from the disturbed areas onsite.

#### 9.1 Site Control

If remedial activities are necessary, a four-zone approach will be employed. The four zones include: the Exclusion Zone (EZ), the Contamination Reduction Zone (CRZ), Contamination Reduction Corridor (CRC) and the Support Zone (SZ). A stepped remedial approach will be managed and the zones modified as the work progresses. Each of the areas will be defined through the use of control barricades and/or construction/hazard fencing. A clearly marked delineation between the SZ and the remaining three zones, the CRZ, CRC and EZ, will be maintained. The preferred method will utilize high visibility orange fencing and hand-driven metal posts, or orange cones. Signage will be posted to further identify and delineate these areas.

## 9.1.1 Support Zone

The Support Zone (SZ) is an uncontaminated area that will be the field support area for the Site operations. The SZ will contain the temporary project trailers and provides for field team communications and staging for emergency response. Appropriate sanitary facilities and safety equipment will be located in this zone. Potentially contaminated personnel or materials are not allowed in this zone. The only exception will be appropriately packaged/decontaminated and labeled samples. Meteorological conditions will be observed and noted from this zone, as well as those factors pertinent to heat and cold stress.

#### 9.1.2 Contamination Reduction Zone

A Contamination Reduction Zone (CRZ) is established between the exclusion zone and the support zone. The CRZ contains the Contamination Reduction Corridor (CRC) and provides an area for decontamination of personnel and equipment. The CRZ will be used for general Site entry and egress in addition to access for heavy equipment and emergency support services. Personnel are not allowed in the CRZ without:

- A buddy (co-worker);
- Appropriate PPE;

- Medical authorization;
- Training certification; and
- A need to be in the zone.

#### 9.1.3 Exclusion Zone

The area where contamination exists is considered to be the Exclusion Zone (EZ). All areas where excavation and handling of contaminated materials take place are considered the EZ. This zone will be clearly delineated by orange high visibility fencing. Safety tape may be used as a secondary delineation within the EZ. The zone delineation markings may be opened in areas for varying lengths of time to accommodate equipment operation or specific construction activities. The PM/SSO may establish more than one EZ where different levels of protection may be employed or where different hazards exist. Personnel are not allowed in the EZ without:

- A buddy (co-worker);
- Required minimum-level PPE;
- Medical authorization;
- Training certification; and
- A need to be in the zone.

#### 9.2 Personal Protective Equipment

The level of protection worn by field personnel will be enforced by the PM/SSO. Levels of protection for general operations are provided below and are defined in this section. Levels of protection may be upgraded at the discretion of the PM/SSO. All decisions on the level of protection will be based upon a conservative interpretation by the PM/SSO of the information provided by air monitoring results, environmental results and other appropriate information. Any changes in the level of protection shall be recorded in the health and safety field logbook.

### **9.2.1 Personal Protective Equipment Specifications**

The initial level of personal protective equipment is Level D. It is not anticipated that either Level B or Level C protection will be necessary.

The Minimum level of PPE for entry onto the Site is Level D PPE. The following equipment shall be used:

- Work uniform (long pants, sleeved shirt)
- Hard hat
- Steel toe work boots
- Safety glasses with attached side shields
- Boot covers (as needed)
- Hearing protection (as needed)
- High visibility clothing (shirt or vest)

Modified Level D PPE consists of the following:

- Regular Tyvek coveralls (Poly-coated Tyvek as required)
- Outer gloves: cut-resistant, leather, cotton (as required)
- Inner gloves: latex or nitrile (doubled) as required
- Chemical resistant boots over work boots (as required)
- Steel toe work boots
- Hard hat Safety glasses with attached side shields
- Hearing protection as needed

High visibility clothing (shirt or vest). Although not anticipated, any tasks requiring Level B personal protective equipment (PPE) will utilize the following equipment:

- Positive pressure, full facepiece, self-contained breathing apparatus (SCBA) or positive pressure, supplied air respirator with escape SCBA (NIOSH approved)
- Disposable coveralls (Tyvek, Poly-coated Tyvek, or Saranex)
- Gloves, inner: latex or nitrile
- Gloves, outer: cut-resistant
- Chemical resistant boots over the work boots
- Steel toe work boots

- Hard hat
- Hearing protection (as needed)
- Boot cover (as needed)

High visibility clothing (shirt or vest). For tasks requiring Level C PPE, the following equipment may be used in any combination:

- Full-face, air purifying, canister-equipped respirators (NIOSH approved) utilizing Organic Vapor/Acid Gas and P-100 filters (half-face if approved by PM/SSO)
- Disposable coveralls (Tyvek) as required
- Gloves, inner: latex or nitrile as required
- Gloves, outer: cut-resistant
- Chemical resistant boots over the work boots as required
- Steel toe work boots
- Hard hat
- Hearing protection (as needed)
- Safety glasses with attached side shields (if half-mask is utilized)
- Boot covers (as needed)
- High visibility clothing (shirt or vest)

#### 9.2.2 Site Specific Levels of Protection

Levels of protection for the proposed scope of work may be upgraded or downgraded depending on direct-reading instruments or personnel monitoring. The following are the initial levels of protection that shall be used for each planned field activity:

Activity	Initial Level of PPE
Installation, Gauging, Purging and Sampling of Temporary Monitoring Wells	D
Installation and sampling of Soil Borings	D

#### 9.3 Communications

Radios

If working in level C/B respiratory protection is required, personnel may find that communication becomes a more difficult task and process to accomplish. Distance and space further complicate this. In order to address this problem, electronic instruments, mechanical devices, or hand signals will be used as follows:

<u>Telephones</u> – Mobile telephones will be carried by designated personnel for communication with emergency support services/facilities.

 Two-way radios will be utilized onsite for communications between field personnel in areas where visual contact cannot be maintained and where hand signals cannot be employed.

Hand Signals – This communication method will be employed by members of the field team along with use of the buddy system. Signals become especially important when in the vicinity of heavy moving equipment and when using Level B respiratory equipment. The signals shall become familiar to the entire field team before Site operations commence, and will be reinforced and reviewed during site-specific training.

SignalMeaningHand gripping throatOut of air; can't breatheGrip partner's wristLeave area immediately; no debateHands on top of headNeed assistanceThumbs upOK; I'm all right; I understandThumbs downNo; unable to understand you, I'm not all right

#### 10.0 MONITORING PROCEDURES

#### 10.1 General

Monitoring will be performed as necessary to verify the adequacy of respiratory protection, to aid in Site layout, and to document worker exposure. If real-time breathing zone air monitoring in these areas indicates the presence of potentially hazardous materials in exceedances of the Action Levels for Worker Breathing Zone (Table 2), the OHSM will be contacted and a plan for implementing appropriate control measures will be developed. A documented safety briefing to communicate the new procedures to onsite personnel will be conducted. All monitoring instruments shall be operated by qualified personnel only and will be calibrated daily prior to use or, more often, as necessary. Additional monitoring may be required if exclusion zones are employed for specific site activities. General air monitoring will be performed in accordance with the Generic Community Air Monitoring Plan included in Appendix G during intrusive Site activities.

#### **10.2 Instrumentation**

The following monitoring instruments will be available for use during field operations as necessary. There will be a minimum of one of each piece of equipment on the Site at all times during intrusive activities:

- <u>Photoionization Detector</u> (PID) with 10.6 EV probe or Flame Ionization Detector (FID) or equivalent.
- Dust/Particulate Monitor (DM), MIE Miniram, or equivalent.

A PID will be used to monitor VOCs in active work areas during intrusive activities. VOCs shall also be measured upwind of the work areas to determine background concentrations.

A particulate monitor shall be used to measure concentrations of dust and particulate matter.

When deemed necessary, a CGI/O<sub>2</sub>/CO (or equivalent) meter shall be used to monitor for combustible gases, oxygen content and/ or carbon monoxide during confined space entry or when operating in areas with poor ventilation as the HSO deems necessary.

Calibration records shall be documented and recorded daily and included in the daily air monitoring report. This report will be specific to work area monitoring. All instruments shall be calibrated before and after each daily use in accordance with manufacturer's procedures.

#### 10.3 Action Levels

Action levels for the upgrading of PPE requirements in the HASP will apply to all Site work during investigation and remediation activities at the Site. Action levels are for known contaminants using direct reading instruments in the Breathing Zone (BZ) for VOCs and particulates, and at the source for combustible gases. The BZ will be determined by the PM/SSO, but is typically 4 to 5 feet above the work area surface or elevation. The action levels to be utilized for the Site are found in Table 2.

#### 11.0 SAFETY CONSIDERATIONS

#### 11.1 General

In addition to the specific requirements of this HASP, common sense should be used at all times. The following general safety rules and practices will be in effect at the site.

- Ignition sources within 35 feet of potentially flammable or contaminated material are strictly prohibited.
- Movement of vehicles and equipment, and other activities will be planned and performed with consideration for the location, height, and relative position of aboveground utilities and fixtures, including signs; lights; canopies; buildings and other structures and construction; and natural features such as trees, boulders, bodies of water, and terrain.
- Approved and appropriate safety equipment (as specified in this HASP), such as eye protection, hard hats, hand protection (nitrile, leather and/or cut resistant gloves as necessary), foot protection, and respirators, must be worn in areas where required.
- No eating, chewing tobacco, gum chewing or drinking will be allowed outside the SZ.
- Contaminated tools and hands must be kept away from the face.
- Personnel must use personal hygiene safe guards (washing up via hand towelettes or potable water) at the end of the shift.
- Each sample must be treated and handled as though it were contaminated.
- Persons with long hair and/or loose-fitting clothing that could become entangled in equipment (e.g., pumps, etc.) must take adequate precautions.
- Horseplay is prohibited in the work area.
- Work while under the influence of intoxicants, narcotics, or controlled substances is strictly prohibited.

#### 11.2 Sample Handling

Personnel responsible for handling of samples will wear the prescribed modified Level D protection. Samples are to be identified as to their hazard and packaged as to prevent spillage or breakage. Any unusual sample conditions shall be noted. Laboratory personnel and all field personnel shall be advised of sample hazard levels and the potential contaminants present. This can be accomplished by a phone call to the lab coordinator and/or including a written statement with the samples reviewing lab safety procedures in handling in order to assure that the practices are appropriate for the suspected contaminants in the sample.

#### 12.0 DECONTAMINATION AND DISPOSAL PROCEDURES

#### **12.1 Contamination Prevention**

Contamination prevention should minimize worker exposure and help ensure valid sample results by precluding cross-contamination. Procedures for contamination avoidance include:

# Personnel

- Do not walk through areas of obvious or known contamination.
- Do not directly handle or touch contaminated materials.
- Make sure that there are no cuts or tears on PPE.
- Fasten all closures in suits; cover with tape, if necessary.
- Particular care should be taken to protect any skin injuries.
- Stay upwind of airborne contaminants.
- Do not carry cigarettes, cosmetics, gum, etc., into contaminated areas.

# Sampling/Monitoring

• When required by the PM/SSO, cover instruments with clear plastic, leaving openings for sampling ports and air exhaust.

#### 12.2 Personnel Decontamination

If an exclusion zone (EZ) is employed at the Site, a field wash for equipment and PPE shall be set up and maintained for all persons exiting the EZ. The system will include a gross wash and rinse for all disposable clothing and boots worn in the EZ. As necessary, equipment and facilities will be available for personnel to wash their hands, arms, neck, and face.

## **12.3 Equipment Decontamination**

All potentially contaminated equipment used at the Site will be decontaminated to prevent contaminants from leaving the Site. The decontamination area will provide for the containment of all wastewater from the decontamination process. Respirators and any other PPE that comes in contact with contaminated materials shall pass through a field wash in the decontamination area, and a thorough decontamination at the end of the day. All decontamination rinse water will be collected and managed in accordance with all applicable regulations.

# 12.4 Decontamination during Medical Emergencies

If emergency life-saving first aid and/or medical treatment are required, normal decontamination procedures may need to be abbreviated or omitted. The Site PM/SSO or designee will accompany contaminated victims to the medical facility to advise on matters involving decontamination, when necessary. The outer garments can be removed if they do not cause delays, interfere with treatment, or aggravate the problem. Respiratory equipment must always be removed. Protective clothing can be cut away. If the outer contaminated garments cannot be safely removed, a plastic barrier between the individual and clean surfaces should be used to help prevent contaminating the inside of ambulances and/or medical personnel. Outer garments are then removed at the medical facility. Attempt to wash or rinse the victim if it is known that the individual has been contaminated with an extremely toxic or corrosive material, which could also cause severe injury or loss of life to emergency response personnel. For minor medical problems (ambulatory) or injuries, the normal decontamination procedures will be followed. Note that heat stroke requires prompt treatment to prevent irreversible damage or death. Protective clothing must be promptly removed. Less serious forms of heat stress also require prompt attention and removal of protective clothing immediately. Unless the victim is obviously contaminated, decontamination should be omitted or minimized, and treatment begun immediately.

#### 12.5 Disposal Procedures

A system of segregating all waste will be developed by the PM/SSO.

All discarded materials, waste materials, or other objects shall be handled in such a way as to preclude the potential for spreading contamination, creating a sanitary hazard, or causing litter to be left onsite. All potentially contaminated materials (e.g., clothing, gloves, etc.,) will be bagged or drummed as necessary, labeled and segregated for disposal. All non-contaminated materials shall be collected, bagged and labeled for appropriate disposal as domestic waste. All waste materials will be staged at the site.

#### 13.0 EMERGENCY PLAN

Should an emergency situation occur, the emergency plan, outlined in this section, shall be known by all onsite personnel prior to the start of work. The emergency plan will be available for use at all times during Site work. The plan provides the phone numbers for the fire, police, ambulance, hospital, poison control centers, and directions to the hospital from the Site. This information is to be found in Section 2 of this HASP.

Various individual Site characteristics will determine preliminary actions taken to assure that this emergency plan is successfully implemented in the event of a Site emergency. Careful consideration must be given to the proximity of neighborhood housing or places of employment, and to the relative possibility of Site release of vapors, which could affect the surrounding community.

The emergency coordinator shall implement the contingency plan whenever conditions at the Site warrant such action. The coordinator will be responsible for coordination of the evacuation, emergency treatment, and transport of Site personnel as necessary, and notification of emergency response units and the appropriate management staff.

In cases where the project principal or project manager is not available, the PM/SSO shall serve as the alternate emergency coordinator.

The PM/SSO during an emergency will perform air monitoring as needed, as well as lend assistance and provide health and safety information to responding emergency personnel.

Site Personnel will endeavor to keep non-essential personnel away from the incident until the appropriate emergency resources arrive. At that time the responders will take control of the Site. Site personnel may be asked to lend assistance to emergency personnel such as during evacuations, help with the injured, etc.

#### 13.1 Evacuation

Evacuation procedures will be discussed prior to the start of work and periodically during safety meetings. In the event of an emergency situation, such as fire, or explosion, an air horn,

automobile horn, or other appropriate device will be sounded for three (3) sharp blasts indicating the initiation of evacuation procedures. The emergency evacuation route shall be known by all site workers. Under no circumstances will incoming personnel or visitors be allowed to proceed into the area once the emergency signal has been given. The PM/SSO or project manager must ensure that access for emergency equipment is provided and that all combustion apparatuses have been shut down once the alarm has been sounded. All Site personnel will assemble in the designated nearest safe location. Once the safety of all personnel is established, the fire department and other emergency response groups will be notified by telephone of the emergency.

## **13.2** Personnel Injury

Emergency first aid shall be applied onsite as appropriate. For non-emergency situations, treatment should be sought, if needed, through the approved occupational health clinic. If necessary, the individual shall be decontaminated, if needed, and transported to the nearest hospital. The PM/SSO will contact the Human Resources Director and OHSM if medical information is needed.

The ambulance/rescue squad shall be contacted for transport as necessary in an emergency. However, since some situations may require transport of an injured party by other means, the injured person shall be escorted to the occupational health clinic or hospital. Maps to these facilities are shown in Figure 2.

#### 13.3 Accident/Incident Reporting

As soon as first aid and/or emergency response needs have been met, the following parties are to be contacted by telephone: (Direct contact, no phone messages).

			<u>Office</u> :	<u>Cell</u> :
1.	Corporate Health & Safety Manager:	Joseph Gentile	(856) 423-8800	(610) 844-6911
2.	Project Principal:	Joseph Duminuco	(631) 232-2600	(631) 921-6279
3.	Project Manager/SHSO:	Jeff Wills	(631) 232-2600	(516) 637-0213
4.	Office Health and Safety Manager:	Ray Fitzpatrick	(631)-232-2600	(631) 484-1168

Written confirmations of verbal reports are to be submitted within 24 hours. The report form entitled "Accident Report and Investigation Form" (Appendix H) is to be used for this purpose.

All representatives contacted by telephone are to receive a copy of this report. In addition to filling out the Accident Report and Investigation Form, if a Roux employee is involved in a motor vehicle accident, the employee must also complete the Acord form (Appendix I).

For reporting purposes, the term accident refers to fatalities, lost time injuries, spill or exposure to hazardous materials (radioactive materials, toxic materials, explosive or flammable materials), fire, explosion, property damage, or potential occurrence (i.e., near miss) of the above.

Any information released from the health care provider, which is not deemed confidential patient information, is to be attached to the appropriate form. Any medical information, which is released by patient consent, is to be filed in the individual's medical record and treated as confidential.

# 13.4 Personnel Exposure

Skin Contact: Use copious amounts of soap and water. Wash/rinse affected area

thoroughly, then provide appropriate medical attention. Eyes should be

rinsed for 15 minutes upon chemical contamination.

<u>Inhalation</u>: Move to fresh air and/or, if necessary, decontaminate/transport to hospital.

Ingestion: Decontamination and transport to emergency medical facility.

Puncture Wound

or Laceration: Decontamination and transport to emergency medical facility.

#### 13.5 Adverse Weather Conditions

In the event of adverse weather conditions, the PM/SSO or project manager will determine if work can continue without sacrificing the health and safety of all field workers. Some of the items to be considered prior to determining if work should continue are:

- Potential for heat stress and heat-related injuries.
- Potential for cold stress and cold-related injuries.
- Treacherous weather-related conditions.
- Limited visibility.
- Electrical storm potential.

Site activities will be limited to daylight hours and acceptable weather conditions. Inclement working conditions include heavy rain, fog, high winds, and lightning. Observe daily weather reports and evacuate if necessary in case of inclement weather conditions.

#### 13.5.1 Electrical Storm Guidelines

In the event that lightning and/or thunder are observed while working onsite, all onsite activities shall stop and personnel shall seek proper shelter (e.g., substantial building, enclosed vehicle, etc.). Work shall not resume until the threat of lighting has subsided and no lightning or thunder has been observed for 30 minutes. If the possibility of lightning is forecast for the day, advise the onsite personnel or the risks and proper procedure at the pre-work safety briefing. Continuously monitor for changing weather conditions and allow enough time to properly stop work if lightning if forecast.

# 14.0 LOGS, REPORTS AND RECORD KEEPING

The following is a summary of required health and safety logs, reports, and record keeping for this project.

## **14.1 Medical and Training Records**

The employer keeps medical and training records. The subcontractor employer must provide verification of training and medical qualifications to the PM/SSO. The PM/SSO will keep a log of personnel meeting appropriate training and medical qualifications for Site work. The log will be kept in the project file. Roux Associates will maintain medical records in accordance with 29 CFR 1910.20.

## 14.2 Onsite Log

The PM/SSO or project manager will keep a log of onsite personnel daily in the designated field book.

# **14.3 Exposure Records**

Applicable personal monitoring results, laboratory reports, calculations, and air sampling data sheets are part of an employee exposure record. These records will be kept by Roux Associates in accordance with 29 CFR 1910.20.

# **14.4 Lessons Learned Reports**

A lessons learned report must be completed following procedures given in Appendix J. The originals will be sent to Roux Associates for maintenance. Copies will be distributed as stated. A copy of the forms will be kept in the project file.

#### 14.5 Accident/Incident Reports

For any injury (OSHA Recordable or not), including "FYI" injuries (injuries where pain was felt, but not even first aid treatment was needed), and illnesses, all work on the activity where the injury/illness occurred will be stopped. An accident/incident report must be completed following procedures given in Appendix H. The originals will be sent to Roux Associates for maintenance. Copies will be distributed as stated. A copy of the forms will be kept in the project file.

# 14.6 OSHA Form 300

An OSHA Form 300 (Log of Occupational Injuries and Illnesses) (Appendix K) will be kept at the Site. All reportable injuries or illnesses will be recorded on this form. At the end of the project, the original will be sent to Roux Associates for maintenance.

# 14.7 Daily Health and Safety Briefing

The Daily Health and Safety Briefing form in Appendix E will be completed daily by the PM/SSO and submitted to the project manager.

# 15.0 FIELD TEAM REVIEW

Each Roux Associates employee or subcontractor shall sign this section after site-specific training is completed and before being permitted to work at the Site.

I have read and had Roux Associates verbally review this Health and Safety Plan prepared for this Site with me. I understand and will comply with the provisions contained therein.

Site/Project: 1022 Old Country Road, Plainview, New York

Date	Name	Signature	Company	

# **TABLES**

- 1. Toxicological, Physical and Chemical Properties of Compounds Potentially Present at the Site
- 2. Action Levels for Worker Breathing Zone

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
1,2,4-Trimethylbenzene	95-63-6	TWA 25 ppm (125	mg TWA 25 ppm (125 mg/m³)	None established	N.D.	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, nose, throat, respiratory system; bronchitis; hypochromic anemia; headache, drowsiness, fatigue, dizziness, nausea, incoordination; vomiting, confusion; chemical pneumonitis (aspiration liquid)	Eyes, skin, respiratory system, central nervous system, blood	Clear, colorless liquid with a distinctive, aromatic odor. BP: 337°F FI.P: 112°F UEL: 6.4% LEL: 0.9% Class II Flammable Liquid
1,2-Dichlorobenzene	95-50-1	TWA 25 ppm STEL 50 ppm	C 50 ppm (300 mg/m <sup>3</sup> )	C 50 ppm (300 mg/m <sup>3</sup> )	200 ppm	inhalation, skin absorption, ingestion, skin and/o eye contact	Irritation eyes, nose; liver, kidney damage; skin blisters r	Eyes, skin, respiratory system, liver, kidneys	Colorless to pale-yellow liquid with a pleasant, aromatic odor. [herbicide] BP: 357°F Fl.P: 151°F UEL: 9.2% LEL: 2.2% Class IIIA Combustible Liquid
1,2-Dichloroethane	107-06-2	TWA 10 ppm	Ca TWA 1 ppm (4 mg/m³) STEL 2 ppm (8 mg/m³)	TWA 50 ppm C 100 ppm 200 ppm [5-minute maximum peak in any 3 hours]	Ca [50 ppm]		Irritation eyes, corneal opacity; central nervous system depression; nausea, vomiting; dermatitis; liver, kidney, cardiovascular system damage; [potential occupational carcinogen]	Eyes, skin, kidneys, liver, central nervous system, cardiovascular system	Colorless liquid with a pleasant, chloroform-like odor. [Note: Decomposes slowly, becomes acidic & darkens in color.] BP: 182°F FI.P: 56°F UEL: 16% LEL: 6.2% Class IB Flammable Liquid
1,2-Dichloroethene (total)	540-59-0	TWA 200 ppm (79	0 m TWA 200 ppm (790 mg/m <sup>3</sup> )	TWA 200 ppm (790 mg/m³)	1000 ppm	inhalation, ingestion, skin and/or eye contact	Irritation eyes, respiratory system; central nervous system depression		Colorless liquid (usually a mixture of the cis & trans isomers) with a slightly acrid, chloroform-like odor BP: 118-140°F FI.P: 36-39°F UEL: 12.8% LEL: 5.6% Class IB Flammable Liquid
1,3,5-Trimethylbenzene	108-67-8	None established	TWA 25 ppm (125mg/m³)	None established	N.D.	Inhalation; ingestion; skin and/or eye contact	Eye, skin, nose, and throat, resp syst irritation; bronchitis; hypochromic anemia; headache, drowsiness, weakness, dizziness, nausea, incoordination, vomit, confusion; chemical pneumonitis	Eyes, skin, resp sys, CNS, blood	Clear, colorless liquid with a distinctive, aromatic odor BP: 329°F FL.P: 122°F Class II Flammable liquid
1,3,5-Trimethylbenzene	108-67-8	TWA 25 ppm (125	mg TWA 25 ppm (125 mg/m³)	None established	N.D	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, nose, throat, respiratory system; bronchitis; hypochromic anemia; headache, drowsiness, lassitude (weakness, exhaustion), dizziness, nausea, incoordination; vomiting, confusion; chemical pneumonitis (aspiration liquid)	Eyes, skin, respiratory system, central nervous system, blood	Clear, colorless liquid with a distinctive, aromatic odor. BP: 329°F Fl.P: 122°F Class II Flammable Liquid

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
1,4-Dichlorobenzene	106-46-7	TWA 10 ppm	Ca	TWA 75 ppm (450 mg/m³)	Ca [150 ppm]	inhalation, skin absorption, ingestion, skin and/or eye contact	Eye irritation, swelling periorbital (situated around the eye); profuse r rhinitis; headache, anorexia, nausea, vomiting; weight loss, jaundice, cirrhosis; in animals: liver, kidney injury; [potential occupational carcinogen]		Colorless or white crystalline solid with a mothball-like odor. [insecticide] BP: 345°F FI.P: 150°F LEL: 2.5% Combustible Solid
2,4-Dimethylphenol	105-67-9	None established	None established	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin, respiratory system, mouth, throat, stomach; r dizziness, weakness, fatigue, nausea, headache; systemic damage; moderate to severe eye injury.	Skin, CVS, eyes, CNS	Clear, colorless liquid with a faint ether or chloroform-like odor BP: 178°F
2-Butanone (MEK)	78-93-3	TWA 200 ppm (590 mg/m³) STEL 300 ppm (885 mg/m³)	TWA 200 ppm (590 mg/m³) STEL 300 ppm (885 mg/m³)	TWA 200 ppm (590 mg/m³)	3000 ppm	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, nose; headache; dizziness; vomiting; dermatitis	Eyes, skin, respiratory system, central nervous system	Colorless liquid with a moderately sharp, fragrant, mint- or acetone-like odor. BP: 175°F FI.P: 16°F UEL(200°F): 11.4% LEL(200°F): 1.4% Class IB Flammable Liquid
Acenaphthene	83-32-9	None established	None established	None established	None established	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, respiratory system	Eyes, skin, respiratory system	Brown solid
Acetone	67-64-1	TWA 500 ppm STEL 50 ppm	TWA 250 ppm (590 mg/m³)	TWA 1000 ppm (2400 mg/m³)	2500 ppm [10%LEL]	inhalation, ingestion, skin and/or eye contact	Irritation eyes, nose, throat; headache, dizziness, central nervous system depression; dermatitis	Eyes, skin, respiratory system, central nervous system	Colorless liquid with a fragrant, mint-like odor BP: 133°F FLP: 0°F UEL: 12.8% LEL: 2.5% Class IB Flammable Liquid
Anthracene	65996-93-2	TWA 0.2 mg/m <sup>3</sup>	Ca TWA 0.1 mg/m <sup>3</sup> (cyclohexane-extractable fraction)	TWA 0.2 mg/m <sup>3</sup> (benzene-soluble fraction)	Ca [80 mg/m <sup>3</sup> ]	inhalation, skin and/or eye contact	Dermatitis, bronchitis, [potential occupational carcinogen]	respiratory system, skin, bladder, kidneys	Black or dark-brown amorphous residue. Combustible Solids
Antimony	7440-36-0	TWA 0.5 mg/m <sup>3</sup>	TWA 0.5 mg/m <sup>3</sup>	$TWA~0.5~mg/m^3$	50 mg/m <sup>3</sup> (as S	yinhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, nose, throat, mouth; cough; dizziness; headache; nausea, vomiting, diarrhea; stomach cramps; insomnia; anorexia; unable to smell properly	Eyes, skin, respiratory system, cardiovascular system	Silver-white, lustrous, hard, brittle solid; scale-like crystals; or a dark- gray, lustrous powder. BP: 2975°F
Arsenic (inorganic)	7440-38-2 (metal)	TWA 0.01 mg/m3	Ca C 0.002 mg/m3 [15-min]	TWA 0.010 mg/m3	Ca [5 mg/m3 (as As)]	Inhalation; ingestion; skin absorption; skin and/or eye contact	Ulceration of nasal septum, dermatitis, GI disturbances, peripheral neuropathy, resp irritation, hyperpigmentation of skin, [potential occupational carcinogen]	Liver, kidneys, skin, lungs, lymphatic sys	Metal: sliver-gray or tin-white, brittle, odorless solid BP: sublimes
Asbestos	1332-21-4	TWA 0.1 f/cc	Ca 100,000 fibers/m3	TWA 0.1 fiber/cm3	Ca [IDLH value has not been determined]	E Inhalation; ingestion; skin and/or eye contact	Asbestosis (chronic exposure), dyspnea, interstitial fibrosis, restricted pulmonary function, finger clubbing, irritation eyes, [potential occupational carcinogen]	Respiratory system, eyes,	White or greenish (chrysotile), blue (crocidolite), or gray-green (amosite), fibrous, odorless solids. BP: decomposes

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
Asphalt fumes	8052-42-4	TWA 0.5 mg/m³(fumes)	Ca C 5 mg/m3 [15 min]	None established	Ca [IDLH value has not been determined]	e Skin absorption; inhalation; skin and/or eye contact	Irritation eyes, resp sys	Eyes, respiratory system	Black or dark brown cement-like substance Combustible solid
Barium	7440-39-3	TWA 0.5 mg/m3	None established	TWA 0.5 mg/m3	None established	Inhalation, ingestion, skin contact	Irritation skin, respiratory system,	(Skin, eyes, respiratory system	Yellow white powder BP: 1640 C
Benzene	71-43-2	TWA 0.5 ppm STEL 2.5 ppm	Ca TWA 0.1 ppm STEL 1 ppm	TWA 1 ppm STEL 5 ppm	Ca [500 ppm]	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin, nose, respiratory system; dizziness; r headache, nausea, staggered gait; anorexia, lassitude (weakness, exhaustion); dermatitis; bone marrow depression; [potential occupational carcinogen]	Eyes, skin, respiratory system, blood, central nervous system, bone marrow	Colorless to light yellow liquid with an aromatic odor [Note: Solid below 42 °F] BP: 176°F Fl.Pt = 12°F LEL:: 1.2% UEL: 7.8% Class B Flammable liquid
Benzo[a]anthracene	56-55-3	None established	None established	None established	None established	Inhalation; ingestion; skin absorption; skin and/or eye contact	; Irritation eyes, skin, respiratory system, CNS	Skin	Pale Yellow crystal, solid BP: 438 C
Benzo[a]pyrene	50-32-8	None established	TWA 0.1 mg/m3	TWA 0.2 mg/m3	None established		POISON. This material is an experimental carcinogen, mutagen, tumorigen, neoplastigen and teratogen. It is a probable carcinogen in humans and a known human mutagen. IARC Group 2A carcinogen. It is believed to cause bladder, skin and lung cancer. Exposure to it may damage the developing foetus. May cause reproductive damage. Skin, respiratory and eye irritant or burns.		Yellow crystals or powder [found in cigarette smoke, coal tar, fuel exhaust gas and in many other sources] BP: 495 C
Benzo[b]fluoranthene	205-99-2	None established	TWA 0.1 mg/m3	TWA 0.2 mg/m3	None established	Inhalation; ingestion; skin and/or eye contact	No data were identified on the toxicity of benzo[b]fluoranthene to humans. Based on results of studies in animals, IARC concluded that benzo[b]fluoranthene is possibly carcinogenic to humans	Respiratory system, skin, bladder, kidneys	Off-white to tan powder
Benzo[k]fluoranthene	207-08-9	None established	None established	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin, respiratory tract, gastrointestinal; fatal if r swallowed, inhaled, absorbed through the skin; vomiting, nausea, diarrhea	Lungs, respiratory system	Yellow crystals BP: 480 C

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Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
Beryllium	7440-41-7 (metal)	TWA 0.002 mg/m <sup>3</sup>	Ca C 0.0005 mg/m <sup>3</sup>	TWA 0.002 mg/m <sup>3</sup> C 0.005 mg/m <sup>3</sup> (30 minutes) with a maximum peak of 0.025 mg/m <sup>3</sup>	Ca [4 mg/m³ (as Be)]	inhalation, skin and/or eye contact	Berylliosis (chronic exposure): anorexia, weight loss, lassitude (weakness, exhaustion), chest pain, cough, clubbing of fingers, cyanosis, pulmonary insufficiency; irritation eyes; dermatitis; [potential occupational carcinogen]	Eyes, skin, respiratory system	Metal: A hard, brittle, gray-white solid. BP: 4532°F
Bis(2-ethylhexyl) phthalate	117-81-7	TWA 5 mg/m <sup>3</sup>	TWA 5 mg/m <sup>3</sup> STEL 10 mg/m <sup>3</sup> (do not exceed during andy 15-minute work period)	TWA 5 mg/m <sup>3</sup>	None established	inhalation, skin and/or eye contact	Irritation eyes, skin, nose, throat; affect the nervous system and liver; damage to male reproductive glands	Eyes, skin, nose, respiratory system, nervous system, reproductive system, liver	Colorless to light colored, thick liquid with slight odor
Butane	106-97-8	TWA 1000 ppm	TWA 800 ppm (1900 mg/m³)	None established	None established	inhalation, skin and/or eye contact (liquid)	Drowsiness, narcosis, asphyxia; liquid: frostbite	central nervous system	Colorless gas with a gasoline-like or natural gas odor. BP: 31°F UEL: 8.4% LEL: 1.6% Flammable Gas
Cadmium	7440-43-9 (metal)	TWA 0.01 mg/m <sup>3</sup>	Ca	TWA $0.005~\text{mg/m}^3$	Ca [9 mg/m³ (as Cd)]	inhalation, ingestion	Pulmonary edema, dyspnea (breathing difficulty), cough, chest tightness, substernal (occurring beneath the sternum) pain; headache; chills, muscle aches; nausea, vomiting, diarrhea; anosmia (loss of the sense of smell), emphysema, proteinuria, mild anemia; [potential occupational carcinogen]	respiratory system, kidneys, prostate, blood	Metal: Silver-white, blue-tinged lustrous, odorless solid. BP: 1409°F
Carbon Disulfide	75-15-0	TWA 1 ppm	TWA 1 ppm (3 mg/m³) STEL 10 ppm (30 mg/m³) [skin]	TWA 20 ppm C 30 ppm 100 ppm (30-minute maximum peak)	500 ppm	inhalation, skin absorption, ingestion, skin and/o eye contact	Dizziness, headache, poor sleep, lassitude (weakness, exhaustion), r anxiety, anorexia, weight loss; psychosis; polyneuropathy; Parkinson-like syndrome; ocular changes; coronary heart disease; gastritis; kidney, liver injury; eye, skin burns; dermatitis; reproductive effects	central nervous system, peripheral nervous system, cardiovascular system, eyes, kidneys, liver, skin, reproductive system	Colorless to faint-yellow liquid with a sweet ether-like odor. BP: 116°F FI.P: -22°F UEL: 50.0% LEL: 1.3% Class IB Flammable Liquid
Chlorobenzene	108-90-7	TWA 10 ppm	None established	TWA 75 ppm (350 mg/m³)	1000 ppm	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, nose; drowsiness, incoordination; central nervous system depression; in animals: liver, lung- kidney injury	Eyes, skin, respiratory system, central nervous system, liver	Colorless liquid with an almond- like odor BP: 270°F Fl.P: 82°F UEL: 9.6% LEL: 1.3%
Chloroethane	75-00-3	TWA 100ppm	Handle with caution in the workplace	TWA 1000 ppm (2600 mg/m³)	3800 ppm [10%LEL]	inhalation, skin absorption (liquid), ingestion (liquid), skin and/or eye contact	Incoordination, inebriation; abdominal cramps; cardiac arrhythmias, cardiac arrest; liver, kidney damage	Liver, kidneys, respiratory system, cardiovascular system, central nervous system	LEL: 1.3%  Colorless gas or liquid (below 54°F) with a pungent, ether-like odor.  BP: 54°F  Fl.P: NA (Gas) -58°F (Liquid)  UEL: 15.4%  LEL: 3.8%

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
Chloroform	67-66-3	TWA 10 ppm	Ca STEL 2 ppm (9.78 mg/m³) [60- minute]	C 50 ppm (240 mg/m <sup>3</sup> )	Ca [500 ppm]	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; dizziness, mental dullness, nausea, r confusion; headache, lassitude (weakness, exhaustion); anesthesia; enlarged liver; [potential occupational carcinogen]	Liver, kidneys, heart, eyes, skin, central nervous system	Colorless liquid with a pleasant odor BP: 143°F
Chromium	7440-47-3	TWA 0.5 mg/m³ (metal and Cr III compounds) TWA 0.05 mg/m³ (water-soluble Cr IV compounds) TWA 0.01 mg/m³ (insoluble Cr IV compounds)	TWA 0.5 mg/m <sup>3</sup>	TWA 1 mg/m <sup>3</sup>	250 mg/m³ (as Cr)	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin; lung fibrosis (histologic)	Eyes, skin, respiratory system	Blue-white to steel-gray, lustrous, brittle, hard, odorless solid. BP: 4788°F
Chrysene; Phenanthrene; Pyrene; Coal tar pitch volatiles	65996-93-2	TWA 0.2 mg/m3	Ca TWA 0.1 mg/m³ (cyclohexane- extractable fraction)	TWA 0.2 mg/m³ (benzene-soluble fraction)	Ca [80 mg/m <sup>3</sup> ]	Inhalation, skin and/or eye contact	Dermatitis, bronchitis, [potential occupational carcinogen]	Respiratory system, skin, bladder, kidneys	Black or dark-brown amorphous residue. Combustible Solids
cis-1,2-Dichloroethene	158-59-2	TWA 200 ppm	TWA 200 ppm	TWA 200 ppm	None established	inhalation, skin absorption, ingestion	Harmful if swallowed, inhaled, or absorbed through skin. Irritant. Narcotic. Suspected carcinogen	Skin	Colorless liquid BP: 60 C Fl.P: 4 C UEL: 12.8% LEL: 9.7 %
Copper	7440-50-8	TWA 0.2mg/m <sup>3</sup> (fume) 1 mg/m <sup>3</sup> (dusts and mists)	TWA 1 mg/m <sup>3</sup>	TWA 1 mg/m <sup>3</sup>	100 mg/m³ (as Cu)	Inhalation, ingestion, skin and/or eye contact	Irritation eyes, respiratory system; cough, dyspnea (breathing difficulty), wheezing	; Eyes, skin, respiratory system, liver, kidneys (increase(d) risk with Wilson's disease)	Noncombustible Solid in bulk form, but powdered form may ignite. BP: 4703°F
Dibenzo[a,h]anthracene	53-70-3	None established	None established	None established	None established	Inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin	Eyes, skin; skin photosensitization.	Colorless crystalline powder BP: 524°C
Diesel Fuel #2	68476-34-6	None established	None established	Designated as an OSHA Select Carcinogen	None established	ingestion, skin and/or eye contact	r Kidney damage; potential lung damage; suspected carcinogen; irritation of eyes, skin, respiratory tract; dizziness, headache, nausea; chemical pneumonitis (from aspiration of liquid); dry, red skin irritant contact dermatitis; eye redness, pain.		Clear yellow brown combustible liquid; floats on water; distinct diesel petroleum hydrocarbon odor.  BP: 356-716°F FI.P: 154.4-165.2°F LEL: 0.6% UEL: 7.0%
Ethylbenzene	100-41-4	TWA 100 ppm STEL 125 ppm	TWA 100 ppm (435 mg/m³) STEL 125 ppm (545 mg/m³)	TWA 100 ppm (435 mg/m³)	800 ppm [10%LEL]	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, mucous membrane; headache; dermatitis; narcosis, coma	Eyes, skin, respiratory system, central nervous system	Colorless liquid with an aromatic odor. BP: 277°F FI.P: 55°F UEL: 6.7% LEL: 0.8% Class IB Flammable Liquid
Fluoranthene	206-44-0	None established	None established	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; possible burns; heart and liver injury, r pulmonary edema, respiratory arrest, gastrointestinal disturbances.	Heart, liver, lungs.	Yellow needles.

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
Fluorene	86-73-7	None established	None established	None established	None established	inhalation, ingestion, skin and/or eye contact	Irritation skin, digestive tract	Skin	White crystals BP: 563°F
Fuel Oil #2	68476-30-2	TWA 100mg/m³ (aerosol and vapor, as total hydrocarbons)	None established s	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; CNS effects; nausea, vomiting, headache, r cramping, dizziness, weakness, loss of coordination,, drowsiness; kidney, liver damage	•	Clear or yellow to red oily liquid, kerosene-like odor BP: 347 - 689 °F UEL:5-6% LEL: 0.7-1.0%
Gasoline	8006-61-9	TWA 300 ppm STEL 500 ppm	Carcinogen	None established	Ca [IDLH value has not been determined]	e Skin absorption; inhalation; ingestion; skin and/or eye contact	Eyes and skin irritation, mucous membrane; dermatitis; headache; listlessness, blurred vision, dizziness, slurred speech, confusion, convulsions; chemical pneumonitis; possible liver, kidney damage [Potential occupational carcinogen]	Eyes, skin, respiratory system, CNS, Liver, Kidneys	Clear liquid with a characteristic odor, aromatic Fl.Pt = -45°F LEL = 1.4% UEL = 7.6% Classs 1B Flammable Liquid
Hexachlorobutadiene	87-68-3	TWA 0.02 ppm	Ca TWA 0.02 ppm (0.24 mg/m³/[skin]	None established	Ca [N.D.]	inhalation, skin absorption, ingestion, skin and/or eye contact	In animals: irritation eyes, skin, respiratory system; kidney r damage: [potential occupational carcinogen]	Eyes, skin, respiratory system, kidneys	Clear, colorless liquid with a mild, turpentine-like odor. BP: 419°F
Hydrogen Sulfide	7783-06-4	TWA (1 ppm) STEL (5 ppm) (adopted values for which changes are proposed in the NIC)	C 10 ppm (15 mg/m³) [10-minute]	C 20 ppm 50 ppm [10-minute maximum peak]	: 100 ppm	inhalation, skin and/or eye contact	Irritation eyes, respiratory system apnea, coma, convulsions; conjunctivitis, eye pain, lacrimation (discharge of tears), photophobia (abnormal visual intolerance to light), corneal vesiculation; dizziness, headache, lassitude (weakness, exhaustion), irritability, insomnia; gastrointestinal disturbance; liquid: frostbite	central nervous system	Colorless gas with a strong odor of rotten eggs. BP: -77°F UEL: 44.0% LEL: 4.0% Flammable Gas
Indeno[1,2,3-cd]pyrene	193-39-5	None established	None established	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; possible human carcinogen (skin); r weakness; affect liver, lung tissue renal tissue; impariment of blood forming tissue	Skin	Fluorescent green-yellow crystalline solid BP: 536 C
Indeno[1,2,3-cd]pyrene	193-39-5	None established	None established	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; possible human carcinogen (skin); r weakness; affect liver, lung tissue renal tissue; impariment of blood forming tissue	Skin	Yellowish crystal solid BP: 536 C
Isopropylbenzene	98-82-8	TWA 50 ppm	TWA 50 ppm (245 mg/m³) [skin]	TWA 50 ppm (245 mg/m³) [skin]	900 ppm [10%LEL]	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin, mucous membrane; dermatitis; headache, r narcosis, coma	Eyes, skin, respiratory system, central nervous system	Colorless liquid with a sharp, penetrating, aromatic odor. BP: 306°F Fl.P: 96°F UEL: 6.5% LEL: 0.9%

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
Lead	7439-92-1	$TWA~0.05~mg/m^3$	TWA (8-hour) 0.050 mg/m <sup>3</sup>	TWA 0.050 mg/m <sup>3</sup>	100 mg/m³ (as Pb)	inhalation, ingestion, skin and/or eye contact	Lassitude (weakness, exhaustion), insomnia; facial pallor; anorexia, weight loss, malnutrition; constipation, abdominal pain, colic; anemia; gingival lead line; tremor; paralysis wrist, ankles; encephalopathy; kidney disease; irritation eyes; hypertension	Eyes, gastrointestinal tract, central nervous system, kidneys, blood, gingival tissue	A heavy, ductile, soft, gray solid. BP: 3164°F Noncombustible Solid in bulk form
Manganese	7439-96-5 (metal)	TWA 0.2 mg/m <sup>3</sup>	TWA 1 mg/m <sup>3</sup> STEL 3 mg/m <sup>3</sup>	C 5 mg/m <sup>3</sup>	500 mg/m³ (as Mn)	inhalation, ingestion	Manganism; asthenia, insomnia, mental confusion; metal fume fever: dry throat, cough, chest tightness, dyspnea (breathing difficulty), rales, flu-like fever; low-back pain; vomiting; malaise (vague feeling of discomfort); lassitude (weakness, exhaustion); kidney damage	respiratory system, central nervous system, blood, kidneys	A lustrous, brittle, silvery solid. BP: 3564°F
Mercury (organo) alkyl compounds (as Hg)	7439-97-6	TWA 0.01 mg/m <sup>3</sup> STEL 0.03 mg/m <sup>3</sup> [skin]	TWA 0.01 mg/m <sup>3</sup> STEL 0.03 mg/m <sup>3</sup> [skin]	TWA 0.01 mg/m <sup>3</sup> C 0.04 mg/m <sup>3</sup>	2 mg/m <sup>3</sup> (as Hg)	inhalation, skin absorption, ingestion, skin and/or eye contact	Paresthesia; ataxia, dysarthria; vision, hearing disturbance; r spasticity, jerking limbs; dizziness; salivation; lacrimation (discharge of tears); nausea, vomiting, diarrhea, constipation; skin burns; emotional disturbance kidney injury; possible teratogenic effects		Appearance and odor vary depending upon the specific (organo) alkyl mercury compound
Mercury compounds [except (organo) alkyls] (as Hg) Mercury	7439-97-6	TWA 0.025 mg/m³ (elemental and inorganic forms)	Hg Vapor: TWA 0.05 mg/m <sup>3</sup> [skin] Other: C 0.1 mg/m3 [skin]	TWA 0.1 mg/m <sup>3</sup>	10 mg/m³ (as Hg)	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; cough, chest pain, dyspnea (breathing r difficulty), bronchitis, pneumonitis; tremor, insomnia, irritability, indecision, headache, lassitude (weakness, exhaustion); stomatitis, salivation; gastrointestinal disturbance, anorexia, weight loss; proteinuria	Eyes, skin, respiratory system, central nervous system, kidneys	Metal: Silver-white, heavy, odorless liquid. [Note: "Other" Hg compounds include all inorganic & aryl Hg compounds except (organo) alkyls.] BP: 674°F
Methyl tert-butyl ether (MTBE)	1634-04-4	TWA 50 ppm	No established REL	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, mucous membrane, respiratory; dizziness, r nausea, headache, intoxication	Eyes, skin, mucous membrane, respiratory system, central nervous system	Colorless liquid BP: 55.2 C
Methylene Chloride	75-09-2	TWA 50 ppm, A3 - suspected human carcinogen	Ca	TWA 25 ppm STEL 125 ppm	Ca [2300 ppm]	absorption,	Irritation eyes, skin; lassitude (weakness, exhaustion), r drowsiness, dizziness; numbness, tingle limbs; nausea; [potential occupational carcinogen]	Eyes, skin, cardiovascular system, central nervous system	Colorless liquid with a chloroform- like odor BP: 104°F UEL: 23% LEL: 13%

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
Naphthalene	91-20-3	TWA 10 ppm STEL 15 ppm	TWA 10 ppm (50 mg/m <sup>3</sup> ) STEL 15 ppm (75 mg/m <sup>3</sup> )	TWA 10 ppm (50 mg/m³)	250 ppm	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes; headache, confusion, excitement, malaise r (vague feeling of discomfort); nausea, vomiting, abdominal pain irritation bladder; profuse sweating; jaundice; hematuria (blood in the urine), renal shutdown; dermatitis, optical neuritis, corneal damage	Eyes, skin, blood, liver, kidneys, central nervous system ;	Colorless to brown solid with an odor of mothballs. BP: 424°F FI.P: 174°F UEL: 5.9% LEL: 0.9%
n-Butylbenzene	104-51-8	None established	None established	None established	None established	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; CNS depression, lung damage; nausea, r vomiting, headache, dizziness, weakness, loss of coordination, blured vision, drowsiness, confusion, disorientation	Eyes, skin,repiratory system, central nervous system	Colorless liquid with a sweet odor BP: 183 C FI.P: 59 C UEL: 5.8% LEL: 0.8%
Nickel	7440-02-0 (Metal)	TWA 1.5 mg/m³ (elemental) TWA 0.1 mg/m³ (soluble inorganic compounds) TWA 0.2 mg/m³ (insoluble inorganic compounds) TWA 0.1 mg/m³ (Nickle subsulfide)	Ca TWA 0.015 mg/m <sup>3</sup>	TWA 1 mg/m <sup>3</sup>	Ca [10 mg/m <sup>3</sup> (as Ni)]	inhalation, ingestion, skin and/or eye contact	Sensitization dermatitis, allergic asthma, pneumonitis; [potential occupational carcinogen]	Nasal cavities, lungs, skin	Metal: Lustrous, silvery, odorless solid. BP: 5139°F
Nitrobenzene	98-95-3	TWA 1 ppm	TWA 1 ppm (5 mg/m³) [skin]	TWA 1 ppm (5 mg/m³) [skin]	200 ppm	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, skin; anoxia; dermatitis; anemia; r methemoglobinemia; in animals: liver, kidney damage; testicular effects	Eyes, skin, blood, liver, kidneys, cardiovascular system, reproductive system	Yellow, oily liquid with a pungent odor like paste shoe polish. BP: 411°F Fl.P: 190°F LEL(200°F): 1.8%
n-Propylbenzene	103-65-1	None established	None established	None established	None established	inhalation, ingestion, skin and/or eye contact	Harmful if swallowed, Irritation eyes, skin, digestive tract, respiratory tract, central nervous system	Eyes, skin, central nervous system, respiratory system	colorless or light yellow liquid BP: 159 C Fl.P: 47 C UEL: 6% LEL: 0.8%
Petroleum hydrocarbons (Petroleum distillates)	8002-05-9	None established	TWA 350 mg/m <sup>3</sup> C 1800 mg/m <sup>3</sup> [15 min]	TWA 500 ppm (2000 mg/m³)	1,100 [10% LEL]	Inhalation; ingestion; skin and/or eye contact	Irritation eyes, skin, nose, throat; dizziness, drowsiness, headache, nausea; dried/cracked skin; chemical pneumonitis		Colorless liquid with a gasoline or kerosene-like odor BP: 86-460°F Fl. Pt = -40 to -86°F UEL: 5.9% LEL: 1.1% Flammable liquid
Phenol	108-95-2	TWA 5 ppm	TWA 5 ppm (19 mg/m <sup>3</sup> ) C 15.6 ppm (60 mg/m <sup>3</sup> ) [15-minute] [skin]	TWA 5 ppm (19 mg/m³) [skin]	250 ppm	inhalation, skin absorption, ingestion, skin and/or eye contact	Irritation eyes, nose, throat; anorexia, weight loss; lassitude r (weakness, exhaustion), muscle ache, pain; dark urine; cyanosis; liver, kidney damage; skin burns; dermatitis; ochronosis; tremor, convulsions, twitching	Eyes, skin, respiratory system, liver, kidneys	Colorless to light-pink, crystalline solid with a sweet, acrid odor. BP: 359°F UEL: 8.6% LEL: 1.8%

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
p-Isopropyltoluene	99-87-6	None established	None established	None established	None established	inhalation, skin absorption, eye contact	Irritation skin	CNS, skin	Colorless, clear liquid, sweetish aromatic odor BP: 350.8°F Class III Flammable liquid
sec-Butylbenzene	135-98-8	None established	None established	None established	None established	inhalation, skin absorption, ingestion, skin and/o eye contact	Irritation eyes, skin, upper airway central nervous system, headache, r dizziness; gastrointestinal disturbance		Colorless liquid BP: 344°F FI.P: 126 °F UEL: 6.9% LEL: 0.8% Combustible liquid
Selenium	7782-49-2	TWA 0.2 mg/m <sup>3</sup>	TWA 0.2 mg/m <sup>3</sup>	TWA 0.2 mg/m <sup>3</sup>	1 mg/m³ (as Se)	inhalation, ingestion, skin and/or eye contact	Irritation eyes, skin, nose, throat; visual disturbance; headache; chills, fever; dyspnea (breathing difficulty), bronchitis; metallic taste, garlic breath, gastrointestinal disturbance; dermatitis; eye, skin burns; in animals: anemia; liver necrosis, cirrhosis; kidney, spleen damage	Eyes, skin, respiratory system, liver, kidneys, blood, spleen	Amorphous or crystalline, red to gray solid. [Note: Occurs as an impurity in most sulfide ores.] BP: 1265°F
Silver	7440-22-4 (metal)	TWA 0.1 mg/m <sup>3</sup> (metal, dust, fumes) TWA 0.01 mg/m <sup>3</sup> (Soluble compounds, as Ag)	TWA 0.01 mg/m <sup>3</sup>	TWA 0.01 mg/m <sup>3</sup>	10 mg/m³ (as Ag)	inhalation, ingestion, skin and/or eye contact	Blue-gray eyes, nasal septum, throat, skin; irritation, ulceration skin; gastrointestinal disturbance	Nasal septum, skin, eyes	Metal: White, lustrous solid BP: 3632°F
tert-Butylbenzene	98-06-6	None established	None established	None established	None established	inhalation, skin absorption, ingestion,	Eye and respiratory irritant; CNS depression; liver or kidney damage	Respiratory system, central nervous system, eyes, liver, kidney	-
Tetrachloroethene	127-18-4	TWA 25 ppm STEL 100 ppm (STEL) listed as A3, animal carcinogen	Ca Minimize workplace exposure concentrations	TWA 100 ppm C 200 ppm (for 5 minutes in any 3-hour period), with a maximum peak of 300 ppm	Ca [150 ppm]	inhalation, skin absorption, ingestion, skin and/o eye contact	Irritation eyes, skin, nose, throat, respiratory system; nausea; flush r face, neck; dizziness, incoordination; headache, drowsiness; skin erythema (skin redness); liver damage; [potential occupational carcinogen]	Eyes, skin, respiratory system, liver, kidneys, central nervous system	Colorless liquid with a mild, chloroform-like odor. BP: 250°F Noncombustible Liquid
Toluene	108-88-3	TWA 20 ppm	TWA 100 ppm (375 mg/m³) STEL 150 ppm (560 mg/m³)	TWA 200 ppm C 300 ppm 500 ppm (10- minute maximum peak)	500 ppm	inhalation, skin absorption, ingestion, skin and/o eye contact	Irritation eyes, nose; lassitude (weakness, exhaustion), r confusion, euphoria, dizziness, headache; dilated pupils, lacrimation (discharge of tears); anxiety, muscle fatigue, insomnia paresthesia; dermatitis; liver, kidney damage	Eyes, skin, respiratory system, central nervous system, liver, kidneys	Colorless liquid with a sweet, pungent, benzene-like odor. BP: 232°F FI.P: 40°F UEL: 7.1% LEL: 1.1% Class IB Flammable Liquid

Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

Compound	CAS#	ACGIH TLV	NIOSH REL	OSHA PEL	IDLH	Routes of Exposure	Toxic Properties	Target Organs	Physical/Chemical Properties
Trichloroethene	79-01-6	TWA 10 ppm STEL 25 ppm	Ca	TWA 100 ppm C 200 ppm 300 ppm (5- minute maximum peak in any 2 hours)	Ca [1000 ppm]	absorption,	Irritation eyes, skin; headache, visual disturbance, lassitude (weakness, exhaustion), dizziness, tremor, drowsiness, nausea, vomiting; dermatitis; cardiac arrhythmias, paresthesia; liver injury; [potential occupational carcinogen]		Colorless liquid (unless dyed blue) with a chloroform-like odor. BP: 189°F UEL(77°F): 10.5% LEL(77°F): 8%
Xylene (m, o & p isomers)	108-38-3, 95-47-6, 106-42-3	TWA 100 ppm (435 mg/m³) STEL 150 ppm	TWA 100 ppm (435 mg/m³)	TWA 100 ppm (435 mg/m $^3$ )	900 ppm	Skin absorption, inhalation, ingestion, skin, and/or eye contact		system, central nervous system, gastrointestinal tract, blood, liver, kidneys	Colorless liquid with an aromatic odor BP: 282°F, 292°F, 281°F Fl. Pt. 82°F, 90°F, 81°F LEL: 1.1%, 0.9%, 1.1% UEL: 7.0%, 6.7%, 7.0% Classs C Flammable Liquid
Zinc	7440-66-6	TWA 10 mg/m3 (Inhalable fraction)	None established	TWA 10 mg/m3 (for zinc oxide fume)	None established	skin and/or eye contact, inhalation, ingestion	Irritation eyes, skin, respiratory tract; gastrointestinal disturbances	Eyes, skin, respiratory system	Bluish gray solid BP: 1664.6°F Flammable

# Table 1. Toxicological, Physical, and Chemical Properties of Compounds Potentially Present at 1022 Old Country Road, Plainview, New York

#### **References:**

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National Institute for Occupational Safety and Health.

#### **Abbreviations:**

ACGIH – American Conference of Governmental Industrial Hygienists

BP – Boiling point at 1 atmosphere, °F

C – Ceiling, is a concentration that should not be exceeded during and part of the working exposure.

Ca – Considered by NIOSH to be a potential occupational carcinogen.

CAS# Chemical Abstracts Service registry number which is unique for each chemical.

Fl. Pt. – Flash point

IDLH – Immediately Dangerous to Life and Health concentrations represent the maximum concentration

from which, in the event of respirator failure, one could escape within 30 minutes without a respirator

and without experiencing any escape-impairing or irreversible health effects.

LEL – Lower explosive (flammable) limit in air, % by volume (at room temperature).

mg/m<sup>3</sup> – Milligrams of substance per cubic meter of air

NIOSH – National Institute for Occupational Safety and Health.

OSHA – Occupational Safety and Health Administration

PEL – OSHA Permissible Exposure Limit (usually) a time weighted average concentration that must not be exceeded during any 8 hour work shift of a 40 hr work week.

ppm - Parts per million

REL - NIOSH Recommended Limit indicated a time weighted average concentration that must not be exceeded during any 10 hour work shift of a 40 hr work week.

STEL – Short-term exposure limit

TLV – ACGIH Threshold Limit Values (usually 8 hour time weighted average concentrations).

TWA – 8-hour, time-weighted average

UEL – Upper explosive (flammable) limit in air, % by volume (at room temperature)

## TABLE 2 ACTION LEVELS FOR WORKER BREATHING ZONE

Instrument	Action Level *	Level of Respiratory Protection/Action
PID	0 to <5 ppm (one minute sustained)	Level D *
PID	>5 to <50 ppm (one minute sustained)	Utilize APR (Level C)
PID	>50 to <100 ppm (one minute sustained)	Level B
PID	>100 ppm	Stop work** (ventilate, apply foam)
CGI/H2S Meter	<5 ppm	Level D
CGI/H2S Meter	>5% to <25 ppm	Level B
CGI/H2S Meter	>25 ppm	Stop work**
CGI/CO Meter	>25 ppm	Level B
CGI/CO Meter	>50 ppm	Stop work** (ventilate area)
CGI/O2 Meter	<10% LEL, in excavation 19.5% oxygen – 23.5%	Level D Level D
CGI/O2 Meter	>10% LEL, in excavation <19.5% or >23.5% oxygen	Allow to vent, apply foam** Stop work, Oxygen Deficient or Enriched ATM**
CGI/CO Meter	>25 to <35 ppm (five minutes sustained) >35 ppm	Allow to vent ** (five minutes sustained) Stop work **

## Note:

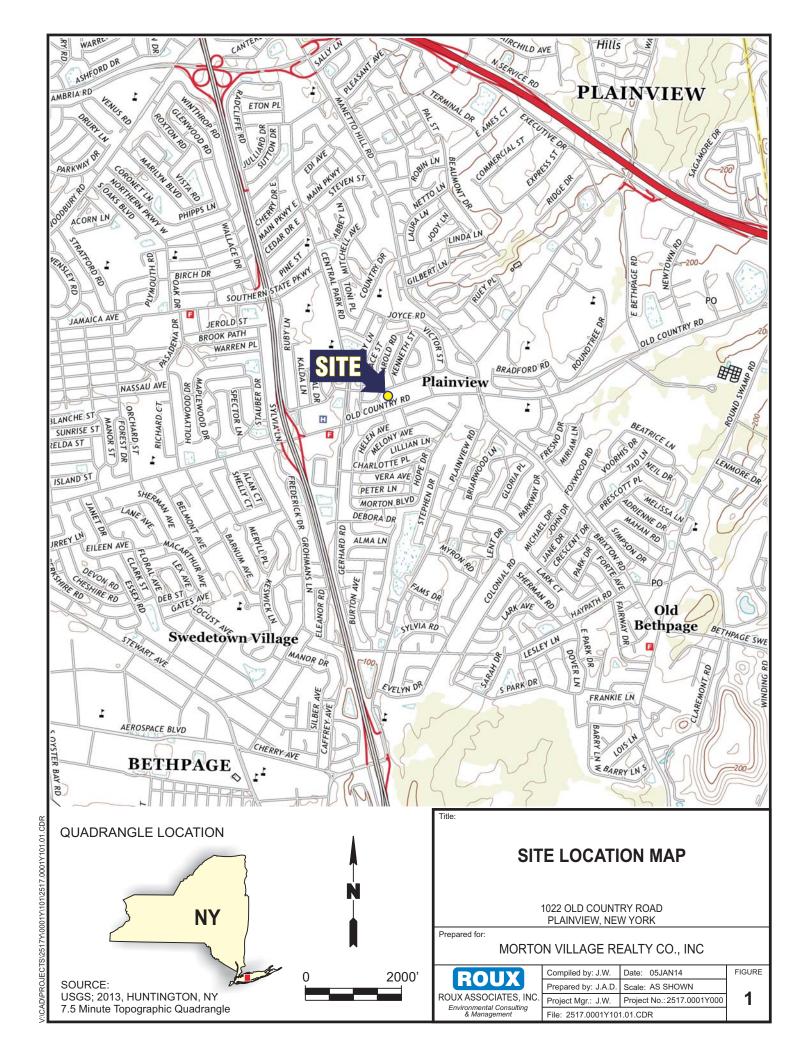
Action levels are based on above background levels.

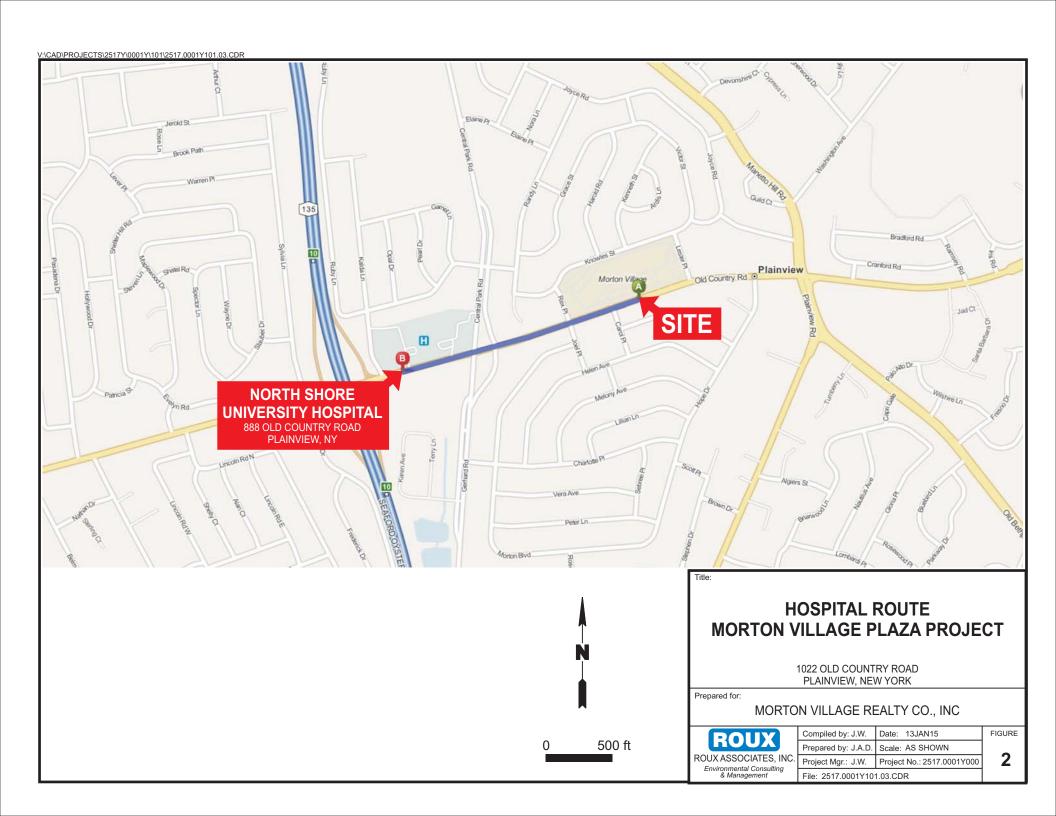
<sup>\*</sup> Instrument readings will be taken in the breathing zone of the workers, unless otherwise indicated.

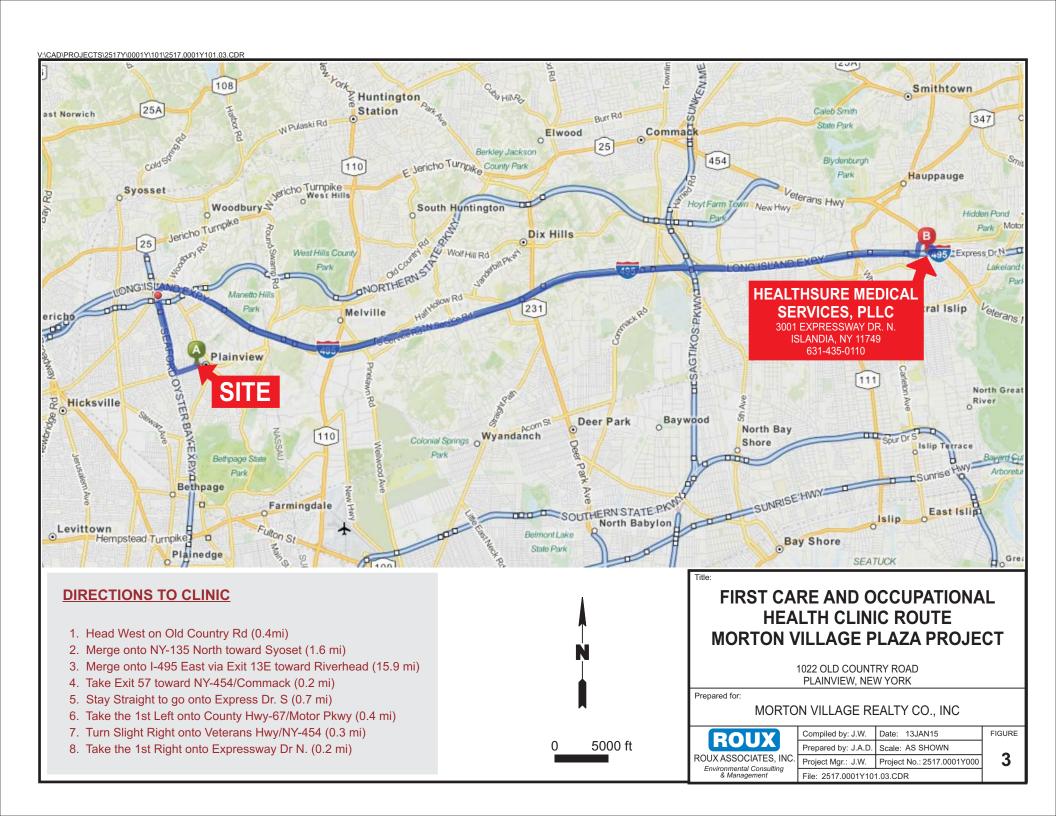
<sup>\*\*</sup> Suspend work in immediate area. Conduct air monitoring periodically to determine when work can continue. Implement mitigative measures.

## **FIGURES**

- 1. Site Location Map
- 2. Hospital Route Map
- 3. Health Clinic Route Map







## **APPENDICES**

- A. Job Safety and Health (OSHA) Poster
- B. Material Safety Data Sheets (MSDS)
- C. Job Safety Analysis
- D. Heat and Cold Stress Guidelines
- E. Health and Safety Briefing/Tailgate Meeting Form
- F. Medical Data Form
- G. Generic Community Air Monitoring Plan
- H. Accident Report and Investigation Form
- I. Acord Automobile Loss Form
- J. Near Loss Reporting Form
- K. OSHA Log of Occupational Injuries and Illnesses

## **APPENDIX A**

Job Safety and Health (OSHA) Poster

## You Have a Right to a Safe and Healthful Workplace.

- You have the right to notify your employer or OSHA about workplace hazards. You may ask OSHA to keep your name confidential.
- You have the right to request an OSHA inspection if you believe that there are unsafe and unhealthful conditions in your workplace. You or your representative may participate in the inspection.
- You can file a complaint with OSHA within 30 days of discrimination by your employer for making safety and health complaints or for exercising your rights under the OSH Act.
- You have a right to see OSHA citations issued to your employer. Your employer must post the citations at or near the place of the alleged violation.
- Your employer must correct workplace hazards by the date indicated on the citation and must certify that these hazards have been reduced or eliminated.
- You have the right to copies of your medical records or records of your exposure to toxic and harmful substances or conditions.
- Your employer must post this notice in your workplace.



The Occupational Safety and Health Act of 1970 (OSH Act), P.L., 91-596, assures safe and healthful working conditions for working men and women throughout the Nation. The Occupational Safety and Health Administration, in the U.S. Department of Labor, has the primary responsibility for administering the OSH Act. The rights listed here may vary depending on the particular circumstances. To file a complaint, report an emergency, or seek OSHA advice, assistance, or products, call 1-800-321-OSHA or your nearest OSHA office: Atlanta (404) 562-2300 • Boston (617) 565-9860 Chicago (312) 353-2220
 Dallas (214) 767-4731
 Denver (303) 844-1600
 Kansas City (816) 426-5861
 New York (212) 337-2378
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> 1-800-321-OSHA www.osha.gov



Health	and	<b>Safety</b>	Plan
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## **APPENDIX B**

Material Safety Data Sheets (MSDS)







## Material Safety Data Sheet Benzene MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Benzene

Catalog Codes: SLB1564, SLB3055, SLB2881

CAS#: 71-43-2

**RTECS:** CY1400000

TSCA: TSCA 8(b) inventory: Benzene

CI#: Not available.

Synonym: Benzol; Benzine

Chemical Name: Benzene

**Chemical Formula:** C6-H6

**Contact Information:** 

Sciencelab.com, Inc. 14025 Smith Rd. Houston. Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400

Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

## **Section 2: Composition and Information on Ingredients**

#### Composition:

Name	CAS#	% by Weight
Benzene	71-43-2	100

**Toxicological Data on Ingredients:** Benzene: ORAL (LD50): Acute: 930 mg/kg [Rat]. 4700 mg/kg [Mouse]. DERMAL (LD50): Acute: >9400 mg/kg [Rabbit]. VAPOR (LC50): Acute: 10000 ppm 7 hours [Rat].

#### Section 3: Hazards Identification

#### **Potential Acute Health Effects:**

Very hazardous in case of eye contact (irritant), of inhalation. Hazardous in case of skin contact (irritant, permeator), of ingestion. Inflammation of the eye is characterized by redness, watering, and itching.

#### **Potential Chronic Health Effects:**

CARCINOGENIC EFFECTS: Classified A1 (Confirmed for human.) by ACGIH, 1 (Proven for human.) by IARC. MUTAGENIC EFFECTS: Classified POSSIBLE for human. Mutagenic for mammalian somatic cells. Mutagenic for bacteria and/or yeast. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Classified Reproductive system/toxin/female [POSSIBLE]. The substance is toxic to blood, bone marrow, central nervous system (CNS). The substance may be toxic to liver, Urinary System. Repeated or prolonged exposure to the substance can produce target organs damage.

#### Section 4: First Aid Measures

#### **Eye Contact:**

Check for and remove any contact lenses. In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Cold water may be used. WARM water MUST be used. Get medical attention immediately.

#### Skin Contact:

In case of contact, immediately flush skin with plenty of water. Cover the irritated skin with an emollient. Remove contaminated clothing and shoes. Wash clothing before reuse. Thoroughly clean shoes before reuse. Get medical attention.

#### **Serious Skin Contact:**

Wash with a disinfectant soap and cover the contaminated skin with an anti-bacterial cream. Seek immediate medical attention.

#### Inhalation:

If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical attention if symptoms appear.

#### Serious Inhalation:

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek medical attention.

#### Ingestion:

Do NOT induce vomiting unless directed to do so by medical personnel. Never give anything by mouth to an unconscious person. If large quantities of this material are swallowed, call a physician immediately. Loosen tight clothing such as a collar, tie, belt or waistband.

Serious Ingestion: Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: Flammable.

Auto-Ignition Temperature: 497.78°C (928°F)

Flash Points: CLOSED CUP: -11.1°C (12°F). (Setaflash)

Flammable Limits: LOWER: 1.2% UPPER: 7.8%

**Products of Combustion:** These products are carbon oxides (CO, CO2).

#### Fire Hazards in Presence of Various Substances:

Highly flammable in presence of open flames and sparks, of heat. Slightly flammable to flammable in presence of oxidizing materials. Non-flammable in presence of shocks.

## **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available. Explosive in presence of oxidizing materials, of acids.

#### **Fire Fighting Media and Instructions:**

Flammable liquid, soluble or dispersed in water. SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use alcohol foam, water spray or fog.

#### **Special Remarks on Fire Hazards:**

Extremely flammable liquid and vapor. Vapor may cause flash fire. Reacts on contact with iodine heptafluoride gas. Dioxygenyl tetrafluoroborate is as very powferful oxidant. The addition of a small particle to small samples of benzene, at ambient temperature, causes ignition. Contact with sodium peroxide with benzene causes ignition. Benzene ignites in contact with powdered chromic anhydride. Virgorous or incandescent reaction with hydrogen + Raney nickel (above 210 C) and bromine trifluoride.

#### **Special Remarks on Explosion Hazards:**

Benzene vapors + chlorine and light causes explosion. Reacts explosively with bromine pentafluoride, chlorine, chlorine trifluoride, diborane, nitric acid, nitryl perchlorate, liquid oxygen, ozone, silver perchlorate. Benzene + pentafluoride and methoxide (from arsenic pentafluoride and potassium methoxide) in trichlorotrifluoroethane causes explosion. Interaction

of nitryl perchlorate with benzene gave a slight explosion and flash. The solution of permanganic acid ( or its explosive anhydride, dimaganese heptoxide) produced by interaction of permanganates and sulfuric acid will explode on contact with benzene. Peroxodisulfuric acid is a very powferful oxidant. Uncontrolled contact with benzene may cause explosion. Mixtures of peroxomonsulfuric acid with benzene explodes.

#### Section 6: Accidental Release Measures

Small Spill: Absorb with an inert material and put the spilled material in an appropriate waste disposal.

#### Large Spill:

Flammable liquid. Keep away from heat. Keep away from sources of ignition. Stop leak if without risk. Absorb with DRY earth, sand or other non-combustible material. Do not touch spilled material. Prevent entry into sewers, basements or confined areas; dike if needed. Be careful that the product is not present at a concentration level above TLV. Check TLV on the MSDS and with local authorities.

## Section 7: Handling and Storage

#### **Precautions:**

Keep locked up.. Keep away from heat. Keep away from sources of ignition. Ground all equipment containing material. Do not ingest. Do not breathe gas/fumes/ vapor/spray. In case of insufficient ventilation, wear suitable respiratory equipment. If ingested, seek medical advice immediately and show the container or the label. Avoid contact with skin and eyes. Keep away from incompatibles such as oxidizing agents, acids.

#### Storage:

Store in a segregated and approved area. Keep container in a cool, well-ventilated area. Keep container tightly closed and sealed until ready for use. Avoid all possible sources of ignition (spark or flame).

## **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value. Ensure that eyewash stations and safety showers are proximal to the work-station location.

#### **Personal Protection:**

Splash goggles. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Vapor respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

## **Exposure Limits:**

TWA: 0.5 STEL: 2.5 (ppm) from ACGIH (TLV) [United States] TWA: 1.6 STEL: 8 (mg/m3) from ACGIH (TLV) [United States] TWA: 0.1 STEL: 1 from NIOSH TWA: 1 STEL: 5 (ppm) from OSHA (PEL) [United States] TWA: 10 (ppm) from OSHA (PEL) [United States] TWA: 3 (ppm) [United Kingdom (UK)] TWA: 1.6 (mg/m3) [United Kingdom (UK)] TWA: 1 (ppm) [Canada] TWA: 3.2 (mg/m3) [Canada] TWA: 0.5 (ppm) [Canada] Consult local authorities for acceptable exposure limits.

## **Section 9: Physical and Chemical Properties**

Physical state and appearance: Liquid.

Odor:

Aromatic. Gasoline-like, rather pleasant. (Strong.)

**Taste:** Not available.

Molecular Weight: 78.11 g/mole

Color: Clear Colorless. Colorless to light yellow.

pH (1% soln/water): Not available.

**Boiling Point:** 80.1 (176.2°F) **Melting Point:** 5.5°C (41.9°F)

Critical Temperature: 288.9°C (552°F)

**Specific Gravity:** 0.8787 @ 15 C (Water = 1)

Vapor Pressure: 10 kPa (@ 20°C)

Vapor Density: 2.8 (Air = 1)

**Volatility:** Not available. **Odor Threshold:** 4.68 ppm

Water/Oil Dist. Coeff.: The product is more soluble in oil; log(oil/water) = 2.1

Ionicity (in Water): Not available.

**Dispersion Properties:** See solubility in water, diethyl ether, acetone.

Solubility:

Miscible in alcohol, chloroform, carbon disulfide oils, carbon tetrachloride, glacial acetic acid, diethyl ether, acetone. Very slightly soluble in cold water.

## **Section 10: Stability and Reactivity Data**

Stability: The product is stable.

**Instability Temperature:** Not available.

Conditions of Instability: Heat, ignition sources, incompatibles.

**Incompatibility with various substances:** Highly reactive with oxidizing agents, acids.

Corrosivity: Non-corrosive in presence of glass.

#### Special Remarks on Reactivity:

Benzene vapors + chlorine and light causes explosion. Reacts explosively with bromine pentafluoride, chlorine, chlorine trifluoride, diborane, nitric acid, nitryl perchlorate, liquid oxygen, ozone, silver perchlorate. Benzene + pentafluoride and methoxide (from arsenic pentafluoride and potassium methoxide) in trichlorotrifluoroethane causes explosion. Interaction of nitryl perchlorate with benzene gave a slight explosion and flash. The solution of permanganic acid ( or its explosive anhydride, dimaganese heptoxide) produced by interaction of permanganates and sulfuric acid will explode on contact with benzene. Peroxodisulfuric acid is a very powferful oxidant. Uncontrolled contact with benzene may cause explosion. Mixtures of peroxomonsulfuric acid with benzene explodes.

Special Remarks on Corrosivity: Not available.

Polymerization: Will not occur.

## **Section 11: Toxicological Information**

Routes of Entry: Absorbed through skin. Dermal contact. Eye contact. Inhalation.

#### **Toxicity to Animals:**

WARNING: THE LC50 VALUES HEREUNDER ARE ESTIMATED ON THE BASIS OF A 4-HOUR EXPOSURE. Acute oral toxicity (LD50): 930 mg/kg [Rat]. Acute dermal toxicity (LD50): >9400 mg/kg [Rabbit]. Acute toxicity of the vapor (LC50): 10000 7 hours [Rat].

**Chronic Effects on Humans:** 

CARCINOGENIC EFFECTS: Classified A1 (Confirmed for human.) by ACGIH, 1 (Proven for human.) by IARC. MUTAGENIC EFFECTS: Classified POSSIBLE for human. Mutagenic for mammalian somatic cells. Mutagenic for bacteria and/or yeast. DEVELOPMENTAL TOXICITY: Classified Reproductive system/toxin/female [POSSIBLE]. Causes damage to the following organs: blood, bone marrow, central nervous system (CNS). May cause damage to the following organs: liver, Urinary System.

#### Other Toxic Effects on Humans:

Very hazardous in case of inhalation. Hazardous in case of skin contact (irritant, permeator), of ingestion.

Special Remarks on Toxicity to Animals: Not available.

#### Special Remarks on Chronic Effects on Humans:

May cause adverse reproductive effects (female fertility, Embryotoxic and/or foetotoxic in animal) and birth defects. May affect genetic material (mutagenic). May cause cancer (tumorigenic, leukemia)) Human: passes the placental barrier, detected in maternal milk.

#### Special Remarks on other Toxic Effects on Humans:

Acute Potential Health Effects: Skin: Causes skin irritation. It can be absorbed through intact skin and affect the liver, blood, metabolism, and urinary system. Eyes: Causes eye irritation. Inhalation: Causes respiratory tract and mucous membrane irritation. Can be absorbed through the lungs. May affect behavior/Central and Peripheral nervous systems (somnolence, muscle weakness, general anesthetic, and other symptoms similar to ingestion), gastrointestinal tract (nausea), blood metabolism, urinary system. Ingestion: May be harmful if swallowed. May cause gastrointestinal tract irritation including vomiting. May affect behavior/Central and Peripheral nervous systems (convulsions, seizures, tremor, irritability, initial CNS stimulation followed by depression, loss of coordination, dizziness, headache, weakness, pallor, flushing), respiration (breathlessness and chest constriction), cardiovascular system, (shallow/rapid pulse), and blood.

## **Section 12: Ecological Information**

Ecotoxicity: Not available.

BOD5 and COD: Not available.

#### **Products of Biodegradation:**

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

Toxicity of the Products of Biodegradation: The products of degradation are less toxic than the product itself.

Special Remarks on the Products of Biodegradation: Not available.

## **Section 13: Disposal Considerations**

#### **Waste Disposal:**

Waste must be disposed of in accordance with federal, state and local environmental control regulations.

## **Section 14: Transport Information**

DOT Classification: CLASS 3: Flammable liquid.
Identification: : Benzene UNNA: 1114 PG: II
Special Provisions for Transport: Not available.

## **Section 15: Other Regulatory Information**

#### Federal and State Regulations:

California prop. 65: This product contains the following ingredients for which the State of California has found to cause cancer, birth defects or other reproductive harm, which would require a warning under the statute: Benzene California prop. 65 (no significant risk level): Benzene: 0.007 mg/day (value) California prop. 65: This product contains the following ingredients

for which the State of California has found to cause cancer which would require a warning under the statute: Benzene Connecticut carcinogen reporting list.: Benzene Connecticut hazardous material survey.: Benzene Illinois toxic substances disclosure to employee act: Benzene Illinois chemical safety act: Benzene New York release reporting list: Benzene Rhode Island RTK hazardous substances: Benzene Pennsylvania RTK: Benzene Minnesota: Benzene Michigan critical material: Benzene Massachusetts RTK: Benzene Massachusetts spill list: Benzene New Jersey: Benzene New Jersey spill list: Benzene Louisiana spill reporting: Benzene California Director's list of Hazardous Substances: Benzene TSCA 8(b) inventory: Benzene SARA 313 toxic chemical notification and release reporting: Benzene CERCLA: Hazardous substances.: Benzene: 10 lbs. (4.536 kg)

## Other Regulations:

OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200). EINECS: This product is on the European Inventory of Existing Commercial Chemical Substances.

#### Other Classifications:

#### WHMIS (Canada):

CLASS B-2: Flammable liquid with a flash point lower than 37.8°C (100°F). CLASS D-2A: Material causing other toxic effects (VERY TOXIC).

#### DSCL (EEC):

R11- Highly flammable. R22- Harmful if swallowed. R38- Irritating to skin. R41- Risk of serious damage to eyes. R45- May cause cancer. R62- Possible risk of impaired fertility. S2- Keep out of the reach of children. S26- In case of contact with eyes, rinse immediately with plenty of water and seek medical advice. S39- Wear eye/face protection. S46- If swallowed, seek medical advice immediately and show this container or label. S53- Avoid exposure - obtain special instructions before use.

## HMIS (U.S.A.):

Health Hazard: 2 Fire Hazard: 3

Reactivity: 0

Personal Protection: h

#### National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 3
Reactivity: 0
Specific hazard:

#### **Protective Equipment:**

Gloves. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Splash goggles.

## **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

Created: 10/10/2005 08:35 PM

Last Updated: 11/06/2008 12:00 PM

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Material Safety Data Sheet cis-1,2-Dichloroethylene, 97%

MSDS# 97773

Section 1 - Chemical Product and Company Identification

MSDS Name: cis-1,2-Dichloroethylene, 97%

Catalog Numbers: AC113380000, AC113380025, AC113380100, AC113380500

Synonyms: cis-Acetylene dichloride.

Acros Organics BVBA

Company Identification: Janssen Pharmaceuticalaan 3a

2440 Geel, Belgium

Acros Organics

Company Identification: (USA)

One Reagent Lane

Fair Lawn, NJ 07410

For information in the US, call:

For information in Europe, call:

Emergency Number, Europe:

+32 14 57 52 11

Emergency Number US:

201-796-7100

CHEMTREC Phone Number, US: 800-424-9300 CHEMTREC Phone Number, Europe: 703-527-3887

Section 2 - Composition, Information on Ingredients

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CAS#: 156-59-2

Chemical Name: cis-1,2-Dichloroethylene

%: 97

EINECS#: 205-859-7

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Hazard Symbols:

×

Risk Phrases:

XN F



11 20 52/53

Section 3 - Hazards Identification

## **EMERGENCY OVERVIEW**

Warning! Flammable liquid and vapor. May cause respiratory tract irritation. Harmful if inhaled. May be harmful if swallowed. Causes eye and skin irritation. Unstabilized substance may polymerize. Target Organs: Central nervous system, respiratory system, eyes, skin.

Potential Health Effects

Eye: Causes moderate eye irritation.

Skin: Causes moderate skin irritation. May cause dermatitis.

Ingestion: May cause gastrointestinal irritation with nausea, vomiting and diarrhea. May be harmful if swallowed. May cause

central nervous system depression.

Inhalation: May cause respiratory tract irritation. May cause narcotic effects in high concentration.

Chronic:

Eyes: In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Get medical aid.

In case of contact, flush skin with plenty of water. Remove contaminated clothing and shoes. Get medical Skin:

aid if irritation develops and persists. Wash clothing before reuse.

If swallowed, do not induce vomiting unless directed to do so by medical personnel. Never give anything by Ingestion:

mouth to an unconscious person. Get medical aid.

If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Inhalation:

Get medical aid.

Notes to Physician:

General Information: Section 5 - Fire Fighting Measures

As in any fire, wear a self-contained breathing apparatus in pressure-demand, MSHA/NIOSH (approved or equivalent), and full protective gear. Vapors may form an explosive mixture with air. Use water spray to keep fire-exposed containers cool. Flammable liquid and vapor. Fire or excessive heat may result in violent rupture of the container due to bulk polymerization. Vapors are heavier than air and may travel to a source

of ignition and flash back. Vapors can spread along the ground and collect in low or confined areas.

Hazardous polymerization may occur under fire conditions.

Extinguishing Media:

Use water fog, dry chemical, carbon dioxide, or regular foam.

Autoignition 440 deg C ( 824.00 deg F) Temperature:

Flash Point: 6 deg C (42.80 deg F)

Explosion 9.70 vol % Limits: Lower:

Explosion 12.80 vol % Limits: Upper:

Handling:

NFPA Rating: health: 2; flammability: 3; instability: 2;

Section 6 - Accidental Release Measures

General Use proper personal protective equipment as indicated in Section 8. Information:

Absorb spill with inert material (e.g. vermiculite, sand or earth), then place in suitable container. Remove Spills/Leaks:

all sources of ignition. Use a spark-proof tool. Provide ventilation.

Section 7 - Handling and Storage

Wash thoroughly after handling. Remove contaminated clothing and wash before reuse. Ground and bond containers when transferring material. Use spark-proof tools and explosion proof equipment. Avoid contact with eyes, skin, and clothing. Empty containers retain product residue, (liquid and/or vapor), and can be dangerous.

Avoid ingestion and inhalation. Do not pressurize, cut, weld, braze, solder, drill, grind, or expose empty containers to heat, sparks or open flames. Use only with adequate ventilation. Pure vapor will be uninhibited and

may polymerize in vents or other confined spaces.

Keep away from sources of ignition. Store in a tightly closed container. Flammables-area. Store protected from Storage:

light and air.

#### Section 8 - Exposure Controls, Personal Protection

++	+	+	++
Chemical Name	ACGIH	NIOSH	OSHA - Final PELs
cis-1,2-Dichloroeth	200 ppm	none listed	none listed
ylene	I		1
++	+		++

## OSHA Vacated PELs: cis-1,2-Dichloroethylene: None listed

## **Engineering Controls:**

Use process enclosure, local exhaust ventilation, or other engineering controls to control airborne levels below recommended exposure limits. Facilities storing or utilizing this material should be equipped with an eyewash facility and a safety shower.

**Exposure Limits** 

Personal Protective Equipment

Eyes: Wear chemical splash goggles.

Skin: Wear appropriate protective gloves to prevent skin exposure.

Clothing: Wear appropriate protective clothing to prevent skin exposure.

Follow the OSHA respirator regulations found in 29 CFR 1910.134 or European Standard EN 149. Use a

Respirators: NIOSH/MSHA or European Standard EN 149 approved respirator if exposure limits are exceeded or if

irritation or other symptoms are experienced.

Section 9 - Physical and Chemical Properties

Physical State: Liquid

Color: Clear

Odor: Pleasant odor pH: Not available

Vapor Pressure: 201 mm Hg @ 25 deg C

Vapor Density: 3.34 (air=1) Evaporation Rate: Not available Viscosity: Not available

Boiling Point: 60 deg C @ 760 mm Hg ( 140.00°F)

Freezing/Melting Point: -80 deg C (-112.00°F)

Decomposition Temperature: Not available Solubility in water: Insoluble

Specific Gravity/Density: 1.2800

Molecular Formula: C2H2Cl2

Molecular Weight: 96.94

Section 10 - Stability and Reactivity

Chemical Stability: Stable under normal temperatures and pressures. This material is a monomer and may

polymerize under certain conditions if the stabilizer is lost.

Conditions to Avoid: Light, ignition sources, exposure to air, excess heat.

Incompatibilities with Other

Materials

Strong oxidizing agents, strong bases, copper.

Hazardous Decomposition

Products

Hydrogen chloride, phosgene, carbon monoxide, carbon dioxide.

Hazardous Polymerization May occur.

Section 11 - Toxicological Information

RTECS#: CAS# 156-59-2: KV9420000

RTECS:

LD50/LC50: **CAS# 156-59-2:** Inhalation, rat: LC50 = 13700 ppm;

.

Carcinogenicity: cis-1,2-Dichloroethylene - Not listed as a carcinogen by ACGIH, IARC, NTP, or CA Prop 65.

Other: See actual entry in RTECS for complete information.

Section 12 - Ecological Information

Not available

Section 13 - Disposal Considerations

Dispose of in a manner consistent with federal, state, and local regulations.

Section 14 - Transport Information

**US DOT** 

Shipping Name: 1,2-DICHLOROETHYLENE

Hazard Class: 3

UN Number: UN1150 Packing Group: II Canada TDG

Shipping Name: Not available

Hazard Class:

UN Number: Packing Group:

## Section 15 - Regulatory Information

## European/International Regulations

European Labeling in Accordance with EC Directives

Hazard Symbols: XN F

Risk Phrases:

R 11 Highly flammable.

R 20 Harmful by inhalation.

R 52/53 Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

#### Safety Phrases:

S 7 Keep container tightly closed.

S 16 Keep away from sources of ignition - No smoking.

S 29 Do not empty into drains.

S 61 Avoid release to the environment. Refer to special instructions/safety data sheets.

## WGK (Water Danger/Protection)

CAS# 156-59-2: Not available

#### Canada

CAS# 156-59-2 is listed on Canada's NDSL List

Canadian WHMIS Classifications: Not available

This product has been classified in accordance with the hazard criteria of the Controlled Products Regulations and the MSDS contains all of the information required by those regulations.

CAS# 156-59-2 is not listed on Canada's Ingredient Disclosure List.

## US Federal

#### **TSCA**

CAS# 156-59-2 is listed on the TSCA Inventory.

Section 16 - Other Information

MSDS Creation Date: 2/09/1998 Revision #6 Date 7/20/2009

The information above is believed to be accurate and represents the best information currently available to us. However, we make no warranty of merchantibility or any other warranty, express or implied, with respect to such information, and we assume no liability resulting from its use. Users should make their own investigations to determine the suitability of the information for their particular purposes. In no event shall the company be liable for any claims, losses, or damages of any third party or for lost profits or any special, indirect, incidental, consequential, or exemplary damages howsoever arising, even if the company has been advised of the possibility of such damages.

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## Material Safety Data Sheet Ethylbenzene MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Ethylbenzene

Catalog Codes: SLE2044

CAS#: 100-41-4

RTECS: DA0700000

TSCA: TSCA 8(b) inventory: Ethylbenzene

CI#: Not available.

Synonym: Ethyl Benzene; Ethylbenzol; Phenylethane

Chemical Name: Ethylbenzene

**Chemical Formula:** C8H10

**Contact Information:** 

Sciencelab.com, Inc. 14025 Smith Rd. Houston. Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400

Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

## **Section 2: Composition and Information on Ingredients**

#### Composition:

Name	CAS#	% by Weight
Ethylbenzene	100-41-4	100

Toxicological Data on Ingredients: Ethylbenzene: ORAL (LD50): Acute: 3500 mg/kg [Rat].

#### Section 3: Hazards Identification

#### **Potential Acute Health Effects:**

Hazardous in case of eye contact (irritant), of ingestion, of inhalation. Slightly hazardous in case of skin contact (irritant, permeator).

#### **Potential Chronic Health Effects:**

Slightly hazardous in case of skin contact (irritant, sensitizer). CARCINOGENIC EFFECTS: Classified 2B (Possible for human.) by IARC. MUTAGENIC EFFECTS: Mutagenic for mammalian somatic cells. Mutagenic for bacteria and/or yeast. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Not available. The substance may be toxic to central nervous system (CNS). Repeated or prolonged exposure to the substance can produce target organs damage.

#### **Section 4: First Aid Measures**

#### **Eye Contact:**

Check for and remove any contact lenses. In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Cold water may be used. WARM water MUST be used. Get medical attention.

Skin Contact: Wash with soap and water. Cover the irritated skin with an emollient. Get medical attention if irritation develops.

Serious Skin Contact: Not available.

#### Inhalation:

If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical attention.

#### Serious Inhalation:

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. WARNING: It may be hazardous to the person providing aid to give mouth-to-mouth resuscitation when the inhaled material is toxic, infectious or corrosive. Seek medical attention.

#### Ingestion:

Do NOT induce vomiting unless directed to do so by medical personnel. Never give anything by mouth to an unconscious person. Loosen tight clothing such as a collar, tie, belt or waistband. Get medical attention if symptoms appear.

Serious Ingestion: Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: Flammable.

**Auto-Ignition Temperature:** 432°C (809.6°F)

Flash Points:

CLOSED CUP: 15°C (59°F). (Tagliabue.) OPEN CUP: 26.667°C (80°F) (Cleveland) (CHRIS, 2001) CLOSED CUP: 12.8 C (55 F) (Bingham et al., 2001; NIOSH, 2001) CLOSED CUP: 21 C (70 F) (NFPA)

Flammable Limits: LOWER: 0.8% - 1.6% UPPER: 6.7% - 7%

**Products of Combustion:** These products are carbon oxides (CO, CO2).

Fire Hazards in Presence of Various Substances: Highly flammable in presence of open flames and sparks, of heat.

#### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available. Slightly explosive in presence of heat.

#### Fire Fighting Media and Instructions:

Flammable liquid, soluble or dispersed in water. SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use alcohol foam, water spray or fog.

#### **Special Remarks on Fire Hazards:**

Vapor may travel considerable distance to source of ignition and flash back. Vapors may form explosive mixtures with air. When heated to decomposition it emits acrid smoke and irritating fumes.

**Special Remarks on Explosion Hazards:** Vapors may form explosive mixtures in air.

#### **Section 6: Accidental Release Measures**

Small Spill: Absorb with an inert material and put the spilled material in an appropriate waste disposal.

#### Large Spill:

Flammable liquid. Keep away from heat. Keep away from sources of ignition. Stop leak if without risk. Absorb with DRY earth, sand or other non-combustible material. Do not touch spilled material. Prevent entry into sewers, basements or confined areas; dike if needed. Be careful that the product is not present at a concentration level above TLV. Check TLV on the MSDS and with local authorities.

## **Section 7: Handling and Storage**

#### Precautions:

Keep away from heat. Keep away from sources of ignition. Ground all equipment containing material. Do not ingest. Do not breathe gas/fumes/ vapor/spray. Avoid contact with eyes. Wear suitable protective clothing. In case of insufficient ventilation, wear suitable respiratory equipment. If ingested, seek medical advice immediately and show the container or the label. Keep away from incompatibles such as oxidizing agents.

## Storage:

Store in a segregated and approved area. Keep container in a cool, well-ventilated area. Keep container tightly closed and sealed until ready for use. Avoid all possible sources of ignition (spark or flame). Sensitive to light. Store in light-resistant containers.

## **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value. Ensure that eyewash stations and safety showers are proximal to the work-station location.

#### **Personal Protection:**

Splash goggles. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Vapor respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

#### **Exposure Limits:**

TWA: 100 STEL: 125 (ppm) from OSHA (PEL) [United States] TWA: 435 STEL: 545 from OSHA (PEL) [United States] TWA: 435 STEL: 545 from OSHA (PEL) [United States] TWA: 435 STEL: 545 (mg/m3) from NIOSH [United States] TWA: 100 STEL: 125 (ppm) from NIOSH [United States] TWA: 100 STEL: 125 (ppm) [United Kingdom (UK)] TWA: 100 STEL: 125 (ppm) [Belgium] TWA: 100 STEL: 125 (ppm) [Finland] TWA: 50 (ppm) [Norway] Consult local authorities for acceptable exposure limits.

## Section 9: Physical and Chemical Properties

Physical state and appearance: Liquid.

Odor: Sweetish. Gasoline-like. Aromatic.

Taste: Not available.

Molecular Weight: 106.16 g/mole

Color: Colorless.

pH (1% soln/water): Not available. Boiling Point: 136°C (276.8°F) Melting Point: -94.9 (-138.8°F)

**Critical Temperature:** 617.15°C (1142.9°F)

Specific Gravity: 0.867 (Water = 1) Vapor Pressure: 0.9 kPa (@ 20°C)

**Vapor Density:** 3.66 (Air = 1)

Volatility: 100% (v/v).
Odor Threshold: 140 ppm

Water/Oil Dist. Coeff.: The product is more soluble in oil; log(oil/water) = 3.1

Ionicity (in Water): Not available.

**Dispersion Properties:** See solubility in water, diethyl ether.

#### Solubility:

Easily soluble in diethyl ether. Very slightly soluble in cold water or practically insoluble in water. Soluble in all proportions in Ethyl alcohol. Soluble in Carbon tetrachloride, Benzene. Insoluble in Ammonia. Slightly soluble in Chloroform. Solubility in Water: 169 mg/l @ 25 deg. C.; 0.014 g/100 ml @ 15 deg. C.

## Section 10: Stability and Reactivity Data

**Stability:** The product is stable.

**Instability Temperature:** Not available.

Conditions of Instability: Heat, ingnition sources (flames, sparks, static), incompatible materials, light

Incompatibility with various substances: Reactive with oxidizing agents.

Corrosivity: Not considered to be corrosive for metals and glass.

Special Remarks on Reactivity:

Can react vigorously with oxidizing materials. Sensitive to light.

Special Remarks on Corrosivity: Not available.

Polymerization: Will not occur.

## **Section 11: Toxicological Information**

Routes of Entry: Absorbed through skin. Inhalation.

**Toxicity to Animals:** Acute oral toxicity (LD50): 3500 mg/kg [Rat].

#### **Chronic Effects on Humans:**

CARCINOGENIC EFFECTS: Classified 2B (Possible for human.) by IARC. MUTAGENIC EFFECTS: Mutagenic for mammalian somatic cells. Mutagenic for bacteria and/or yeast. May cause damage to the following organs: central nervous system (CNS).

#### **Other Toxic Effects on Humans:**

Hazardous in case of ingestion, of inhalation. Slightly hazardous in case of skin contact (irritant, permeator).

#### **Special Remarks on Toxicity to Animals:**

Lethal Dose/Conc 50% Kill: LD50 [Rabbit] - Route: Skin; Dose: 17800 ul/kg Lowest Published Lethal Dose/Conc: LDL[Rat] - Route: Inhalation (vapor); Dose: 4000 ppm/4 H

#### **Special Remarks on Chronic Effects on Humans:**

May cause adverse reproductive effects and birth defects (teratogenic) based on animal test data. May cause cancer based on animals data. IARC evidence for carcinogenicity in animals is sufficient. IARC evidence of carcinogenicity in humans inadequate. May affect genetic material (mutagenic).

## Special Remarks on other Toxic Effects on Humans:

Acute Potential Health Effects: Skin: Can cause mild skin irritation. It can be absorbed through intact skin. Eyes: Contact with vapor or liquid can cause severe eye irritation depending on concentration. It may also cause conjunctivitis. At a vapor exposure level of 85 - 200 ppm, it is mildly and transiently irritating to the eyes; 1000 ppm causes further irritation and tearing; 2000 ppm results in immediate and severe irritation and tearing; 5,000 ppm is intolerable (ACGIH, 1991; Clayton and Clayton, 1994). Standard draize test for eye irritation using 500 mg resulted in severe irritation (RTECS) Inhalation: Exposure to high concentrations can cause nasal, mucous membrane and respiratory tract irritation and can also result in chest constriction and, trouble breathing, respiratory failure, and even death. It can also affect behavior/Central Nervous System. The effective dose for CNS depression in experimental animals was 10,000 ppm (ACGIH, 1991). Symptoms of CNS depression include

headache, nausea, weakness, dizziness, vertigo, irritability, fatigue, lightheadedness, sleepiness, tremor, loss of coordination, judgement and conciousness, coma, and death. It can also cause pulmonary edema. Inhalation of 85 ppm can produce fatigue, insomnia, headache, and mild irritation of the respiratory tract (Haley & Berndt, 1987). Ingestion: Do not drink, pipet or siphon by mouth. May cause gastroinestinal/digestive tract irritation with Abdominal pain, nausea, vomiting. Ethylbenzene is a pulmonary aspiration hazard. Pulmonary aspiration of even small amounts of the liquid may cause fatal pneumonitis. It may also affect behavior/central nervous system with

## **Section 12: Ecological Information**

#### **Ecotoxicity:**

Ecotoxicity in water (LC50): 14 mg/l 96 hours [Fish (Trout)] (static). 12.1 mg/l 96 hours [Fish (Fathead Minnow)] (flow-through)]. 150 mg/l 96 hours [Fish (Blue Gill/Sunfish)] (static). 275 mg/l 96 hours [Fish (Sheepshead Minnow)]. 42.3 mg/l 96 hours [Fish (Fathead Minnow)] (soft water). 87.6 mg/l 96 hours [Shrimp].

BOD5 and COD: Not available.

#### **Products of Biodegradation:**

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

Toxicity of the Products of Biodegradation: The products of degradation are less toxic than the product itself.

Special Remarks on the Products of Biodegradation: Not available.

## **Section 13: Disposal Considerations**

#### Waste Disposal:

Waste must be disposed of in accordance with federal, state and local environmental control regulations.

## **Section 14: Transport Information**

**DOT Classification:** CLASS 3: Flammable liquid. **Identification:** : Ethylbenzene UNNA: 1175 PG: II **Special Provisions for Transport:** Not available.

## **Section 15: Other Regulatory Information**

#### **Federal and State Regulations:**

Connecticut hazardous material survey.: Ethylbenzene Illinois toxic substances disclosure to employee act: Ethylbenzene Illinois chemical safety act: Ethylbenzene New York release reporting list: Ethylbenzene Rhode Island RTK hazardous substances: Ethylbenzene Pennsylvania RTK: Ethylbenzene Minnesota: Ethylbenzene Massachusetts RTK: Ethylbenzene Massachusetts spill list: Ethylbenzene New Jersey: Ethylbenzene New Jersey spill list: Ethylbenzene Louisiana spill reporting: Ethylbenzene California Director's List of Hazardous Substances: Ethylbenzene TSCA 8(b) inventory: Ethylbenzene TSCA 4(a) proposed test rules: Ethylbenzene TSCA 8(d) H and S data reporting: Ethylbenzene: Effective Date: 6/19/87; Sunset Date: 6/19/97 SARA 313 toxic chemical notification and release reporting: Ethylbenzene

### Other Regulations:

OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200). EINECS: This product is on the European Inventory of Existing Commercial Chemical Substances.

## Other Classifications:

#### WHMIS (Canada):

CLASS B-2: Flammable liquid with a flash point lower than 37.8°C (100°F). CLASS D-2A: Material causing other toxic effects (VERY TOXIC). CLASSE D-2B: Material causing other toxic effects (TOXIC).

#### DSCL (EEC):

R11- Highly flammable. R20- Harmful by inhalation. S16- Keep away from sources of ignition - No smoking. S24/25- Avoid contact with skin and eyes. S29- Do not empty into drains.

#### HMIS (U.S.A.):

Health Hazard: 2

Fire Hazard: 3
Reactivity: 0

Personal Protection: h

#### National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 3

Reactivity: 0

Specific hazard:

#### **Protective Equipment:**

Gloves. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Splash goggles.

## **Section 16: Other Information**

#### References:

-Manufacturer's Material Safety Data Sheet. -Fire Protection Guide to Hazardous Materials, 13th ed., Nationial Fire Protection Association (NFPA) -Registry of Toxic Effects of Chemical Substances (RTECS) -Chemical Hazard Response Information System (CHRIS) -Hazardous Substance Data Bank (HSDB) -New Jersey Hazardous Substance Fact Sheet -Ariel Global View -Reprotext System

Other Special Considerations: Not available.

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## Material Safety Data Sheet Cumene MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Cumene

Catalog Codes: SLC3052

CAS#: 98-82-8

RTECS: GR8575000

TSCA: TSCA 8(b) inventory: Cumene

CI#: Not available.

**Synonym:** Isopropyl benzene; Cumol; 2-Phenyl propane;

(1-Methylethyl)benzene

Chemical Name: Isopropylbenzene

Chemical Formula: C6H5CH(CH3)2

**Contact Information:** 

Sciencelab.com, Inc. 14025 Smith Rd. Houston, Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400

Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

## Section 2: Composition and Information on Ingredients

#### Composition:

Name	CAS#	% by Weight
Cumene	98-82-8	100

**Toxicological Data on Ingredients:** Cumene: ORAL (LD50): Acute: 1400 mg/kg [Rat]. 12750 mg/kg [Mouse]. DERMAL (LD50): Acute: 12300 mg/kg [Rabbit].

#### **Section 3: Hazards Identification**

#### **Potential Acute Health Effects:**

Very hazardous in case of skin contact (irritant, permeator), of eye contact (irritant), of ingestion, of inhalation. Inflammation of the eye is characterized by redness, watering, and itching. Skin inflammation is characterized by itching, scaling, reddening, or, occasionally, blistering.

### **Potential Chronic Health Effects:**

Very hazardous in case of skin contact (permeator). CARCINOGENIC EFFECTS: Not available. MUTAGENIC EFFECTS: Not available. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Not available. The substance is toxic to lungs, the nervous system, mucous membranes. Repeated or prolonged exposure to the substance can produce target organs damage.

#### Section 4: First Aid Measures

#### **Eye Contact:**

Check for and remove any contact lenses. Immediately flush eyes with running water for at least 15 minutes, keeping eyelids open. Cold water may be used. Do not use an eye ointment. Seek medical attention.

#### Skin Contact:

After contact with skin, wash immediately with plenty of water. Gently and thoroughly wash the contaminated skin with running water and non-abrasive soap. Be particularly careful to clean folds, crevices, creases and groin. Cover the irritated skin with an emollient. If irritation persists, seek medical attention. Wash contaminated clothing before reusing.

#### Serious Skin Contact:

Wash with a disinfectant soap and cover the contaminated skin with an anti-bacterial cream. Seek immediate medical attention.

Inhalation: Allow the victim to rest in a well ventilated area. Seek immediate medical attention.

#### Serious Inhalation

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek medical attention.

#### Ingestion:

Do not induce vomiting. Examine the lips and mouth to ascertain whether the tissues are damaged, a possible indication that the toxic material was ingested; the absence of such signs, however, is not conclusive. Loosen tight clothing such as a collar, tie, belt or waistband. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek immediate medical attention.

Serious Ingestion: Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: Flammable.

Auto-Ignition Temperature: 424°C (795.2°F)

Flash Points: CLOSED CUP: 36°C (96.8°F). OPEN CUP: 44°C (111.2°F).

Flammable Limits: LOWER: 0.9% UPPER: 6.5%

**Products of Combustion:** These products are carbon oxides (CO, CO2).

Fire Hazards in Presence of Various Substances: Flammable in presence of open flames and sparks.

#### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available.

#### **Fire Fighting Media and Instructions:**

Flammable liquid, soluble or dispersed in water. SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use alcohol foam, water spray or fog. Cool containing vessels with water jet in order to prevent pressure build-up, autoignition or explosion.

Special Remarks on Fire Hazards: Not available.

**Special Remarks on Explosion Hazards:** Not available.

#### **Section 6: Accidental Release Measures**

Small Spill: Absorb with an inert material and put the spilled material in an appropriate waste disposal.

#### Large Spill:

Flammable liquid. Keep away from heat. Keep away from sources of ignition. Stop leak if without risk. Absorb with DRY earth, sand or other non-combustible material. Do not touch spilled material. Prevent entry into sewers, basements or confined areas; dike if needed. Eliminate all ignition sources. Be careful that the product is not present at a concentration level above TLV. Check TLV on the MSDS and with local authorities.

## **Section 7: Handling and Storage**

#### Precautions:

Keep away from heat. Keep away from sources of ignition. Ground all equipment containing material. Do not ingest. Do not breathe gas/fumes/ vapour/spray. In case of insufficient ventilation, wear suitable respiratory equipment If ingested, seek medical advice immediately and show the container or the label. Avoid contact with skin and eyes

#### Storage:

Flammable materials should be stored in a separate safety storage cabinet or room. Keep away from heat. Keep away from sources of ignition. Keep container tightly closed. Keep in a cool, well-ventilated place. Ground all equipment containing material. A refrigerated room would be preferable for materials with a flash point lower than 37.8°C (100°F).

## **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value. Ensure that eyewash stations and safety showers are proximal to the work-station location.

#### **Personal Protection:**

Splash goggles. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Vapor respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

#### **Exposure Limits:**

TWA: 50 CEIL: 75 (ppm) TWA: 245 CEIL: 365 (mg/m3) Consult local authorities for acceptable exposure limits.

## **Section 9: Physical and Chemical Properties**

Physical state and appearance: Liquid.

Odor: Not available.

Taste: Not available.

Molecular Weight: 120.2 g/mole

Color: Clear Colorless.

pH (1% soln/water): Not available. Boiling Point: 152.4°C (306.3°F) Melting Point: -96°C (-140.8°F)

Critical Temperature: Not available.

Specific Gravity: 0.862 (Water = 1)

Vapor Pressure: 8 mm of Hg (@ 20°C)

Vapor Density: 4.14 (Air = 1)

Volatility: Not available.

Odor Threshold: 1.2 ppm

Water/Oil Dist. Coeff.: The product is more soluble in oil; log(oil/water) = 3.7

Ionicity (in Water): Not available.

**Dispersion Properties:** Not available.

**Solubility:** Very slightly soluble in cold water.

Section 10: Stability and Reactivity Data

Stability: The product is stable.

**Instability Temperature:** Not available. **Conditions of Instability:** Not available.

Incompatibility with various substances: Not available.

Corrosivity: Non-corrosive in presence of glass.

Special Remarks on Reactivity: Not available.

Special Remarks on Corrosivity: Not available.

Polymerization: No.

## **Section 11: Toxicological Information**

Routes of Entry: Dermal contact. Eye contact. Inhalation. Ingestion.

**Toxicity to Animals:** 

Acute oral toxicity (LD50): 1400 mg/kg [Rat]. Acute dermal toxicity (LD50): 12300 mg/kg [Rabbit].

Chronic Effects on Humans: The substance is toxic to lungs, the nervous system, mucous membranes.

Other Toxic Effects on Humans: Very hazardous in case of skin contact (irritant, permeator), of ingestion, of inhalation.

**Special Remarks on Toxicity to Animals:** Not available.

Special Remarks on Chronic Effects on Humans: Not available.

Special Remarks on other Toxic Effects on Humans: Not available.

## **Section 12: Ecological Information**

**Ecotoxicity:** Not available.

BOD5 and COD: Not available.

**Products of Biodegradation:** 

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

**Toxicity of the Products of Biodegradation:** The products of degradation are more toxic.

Special Remarks on the Products of Biodegradation: Not available.

## **Section 13: Disposal Considerations**

**Waste Disposal:** 

## **Section 14: Transport Information**

**DOT Classification:** Class 3: Flammable liquid.

Identification: : Isopropylbenzene : UN1918 PG: III

## **Section 15: Other Regulatory Information**

#### Federal and State Regulations:

Pennsylvania RTK: Cumene Massachusetts RTK: Cumene TSCA 8(b) inventory: Cumene SARA 313 toxic chemical notification and release reporting: Cumene CERCLA: Hazardous substances.: Cumene

Other Regulations: OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200).

Other Classifications:

WHMIS (Canada):

CLASS B-3: Combustible liquid with a flash point between 37.8°C (100°F) and 93.3°C (200°F).

DSCL (EEC):

R10- Flammable. R22- Harmful if swallowed. R38- Irritating to skin. R41- Risk of serious damage to eyes.

HMIS (U.S.A.):

Health Hazard: 2

Fire Hazard: 3

Reactivity: 0

Personal Protection: h

National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 3

Reactivity: 1

Specific hazard:

#### **Protective Equipment:**

Gloves. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Splash goggles.

#### **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

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## Material Safety Data Sheet Methyl tert-butyl ether MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Methyl tert-butyl ether

Catalog Codes: SLM2152

CAS#: 1634-04-4

RTECS: KN5250000

TSCA: TSCA 8(b) inventory: Methyl tert-butyl ether

CI#: Not available.

Synonym:

Chemical Name: Methyl tert-Butyl Ether

Chemical Formula: C5-H12-O

**Contact Information:** 

Sciencelab.com, Inc. 14025 Smith Rd. Houston, Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400

Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

## **Section 2: Composition and Information on Ingredients**

#### Composition:

Name	CAS#	% by Weight
Methyl {tert-}butyl ether	1634-04-4	100

**Toxicological Data on Ingredients:** Methyl tert-butyl ether: ORAL (LD50): Acute: 4000 mg/kg [Rat]. 5960 mg/kg [Mouse]. VAPOR (LC50): Acute: 23576 ppm 4 hour(s) [Rat].

#### Section 3: Hazards Identification

#### **Potential Acute Health Effects:**

Extremely hazardous in case of eye contact (irritant), of ingestion. Very hazardous in case of skin contact (irritant), of inhalation. Hazardous in case of skin contact (permeator). Inflammation of the eye is characterized by redness, watering, and itching. Skin inflammation is characterized by itching, scaling, reddening, or, occasionally, blistering.

#### **Potential Chronic Health Effects:**

Extremely hazardous in case of eye contact (irritant), of ingestion. Very hazardous in case of skin contact (irritant), of inhalation. Hazardous in case of skin contact (permeator). CARCINOGENIC EFFECTS: Not available. MUTAGENIC EFFECTS: Not available. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Not available. The substance is toxic to lungs, the nervous system, mucous membranes. Repeated or prolonged exposure to the substance can produce target organs damage. Repeated or prolonged inhalation of vapors may lead to chronic respiratory irritation.

#### **Section 4: First Aid Measures**

#### **Eye Contact:**

Check for and remove any contact lenses. Immediately flush eyes with running water for at least 15 minutes, keeping eyelids open. Cold water may be used. Do not use an eye ointment. Seek medical attention.

#### Skin Contact:

After contact with skin, wash immediately with plenty of water. Gently and thoroughly wash the contaminated skin with running water and non-abrasive soap. Be particularly careful to clean folds, crevices, creases and groin. Cold water may be used. Cover the irritated skin with an emollient. If irritation persists, seek medical attention. Wash contaminated clothing before reusing.

#### **Serious Skin Contact:**

Wash with a disinfectant soap and cover the contaminated skin with an anti-bacterial cream. Seek medical attention.

Inhalation: Allow the victim to rest in a well ventilated area. Seek immediate medical attention.

#### Serious Inhalation:

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek medical attention.

#### Ingestion:

Do not induce vomiting. Loosen tight clothing such as a collar, tie, belt or waistband. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek immediate medical attention.

**Serious Ingestion:** Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: Flammable.

**Auto-Ignition Temperature:** 224°C (435.2°F)

Flash Points: CLOSED CUP: -28°C (-18.4°F).

Flammable Limits: LOWER: 2.5% UPPER: 15.1%

**Products of Combustion:** These products are carbon oxides (CO, CO2).

Fire Hazards in Presence of Various Substances: Flammable in presence of open flames and sparks.

#### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available.

#### **Fire Fighting Media and Instructions:**

Flammable liquid, soluble or dispersed in water. SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use alcohol foam, water spray or fog.

Special Remarks on Fire Hazards: Not available.

Special Remarks on Explosion Hazards: Not available.

#### Section 6: Accidental Release Measures

#### Small Spill:

Dilute with water and mop up, or absorb with an inert dry material and place in an appropriate waste disposal container.

#### Large Spill:

Flammable liquid. Keep away from heat. Keep away from sources of ignition. Stop leak if without risk. Absorb with DRY earth, sand or other non-combustible material. Do not touch spilled material. Prevent entry into sewers, basements or confined areas; dike if needed. Eliminate all ignition sources.

## **Section 7: Handling and Storage**

#### Precautions:

Keep away from heat. Keep away from sources of ignition. Ground all equipment containing material. Do not ingest. Do not breathe gas/fumes/ vapour/spray. In case of insufficient ventilation, wear suitable respiratory equipment If ingested, seek medical advice immediately and show the container or the label. Avoid contact with skin and eyes

#### Storage:

Flammable materials should be stored in a separate safety storage cabinet or room. Keep away from heat. Keep away from sources of ignition. Keep container tightly closed. Keep in a cool, well-ventilated place. Ground all equipment containing material. A refrigerated room would be preferable for materials with a flash point lower than 37.8°C (100°F).

## **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value. Ensure that eyewash stations and safety showers are proximal to the work-station location.

#### **Personal Protection:**

Splash goggles. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Vapor respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

**Exposure Limits:** Not available.

## **Section 9: Physical and Chemical Properties**

Physical state and appearance: Liquid.

**Odor:** Characteristic. (Strong.)

Taste: Not available.

Molecular Weight: 88.15 g/mole

Color: Clear Colorless.

pH (1% soln/water): Not available.

Boiling Point: 55.2°C (131.4°F)

Melting Point: -109°C (-164.2°F)

Critical Temperature: Not available.

Specific Gravity: 0.7405 (Water = 1)

Vapor Pressure: 245 mm of Hg (@ 20°C)

Vapor Density: 3.1 (Air = 1)

**Volatility:** 100% (v/v).

Odor Threshold: Not available.

Water/Oil Dist. Coeff.: Not available. lonicity (in Water): Not available.

**Dispersion Properties:** See solubility in water, methanol, diethyl ether.

Solubility:

Soluble in methanol, diethyl ether. Partially soluble in cold water.

## Section 10: Stability and Reactivity Data

**Stability:** The product is stable.

**Instability Temperature:** Not available. **Conditions of Instability:** Not available.

Incompatibility with various substances: Not available.

Corrosivity: Non-corrosive in presence of glass.

Special Remarks on Reactivity: Not available.

Special Remarks on Corrosivity: Not available.

Polymerization: No.

## Section 11: Toxicological Information

Routes of Entry: Dermal contact. Eye contact. Inhalation. Ingestion.

**Toxicity to Animals:** 

WARNING: THE LC50 VALUES HEREUNDER ARE ESTIMATED ON THE BASIS OF A 4-HOUR EXPOSURE. Acute oral toxicity (LD50): 4000 mg/kg [Rat]. Acute toxicity of the vapor (LC50): 23576 ppm 4 hour(s) [Rat].

Chronic Effects on Humans: The substance is toxic to lungs, the nervous system, mucous membranes.

Other Toxic Effects on Humans:

Extremely hazardous in case of ingestion. Very hazardous in case of skin contact (irritant), of inhalation. Hazardous in case of skin contact (permeator).

Special Remarks on Toxicity to Animals: Not available.

Special Remarks on Chronic Effects on Humans: Not available.

Special Remarks on other Toxic Effects on Humans: Not available.

## **Section 12: Ecological Information**

Ecotoxicity: Not available.

BOD5 and COD: Not available.

### **Products of Biodegradation:**

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

**Toxicity of the Products of Biodegradation:** The products of degradation are more toxic.

Special Remarks on the Products of Biodegradation: Not available.

#### **Section 13: Disposal Considerations**

Waste Disposal:

## **Section 14: Transport Information**

DOT Classification: Class 3: Flammable liquid.

Identification: : Methyl tert-butyl ether : UN2398 PG: II

Special Provisions for Transport: Not available.

## **Section 15: Other Regulatory Information**

#### Federal and State Regulations:

Pennsylvania RTK: Methyl tert-butyl ether Massachusetts RTK: Methyl tert-butyl ether TSCA 8(b) inventory: Methyl tert-butyl ether SARA 313 toxic chemical notification and release reporting: Methyl tert-butyl ether CERCLA: Hazardous substances.: Methyl tert-butyl ether

Other Regulations: OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200).

Other Classifications:

## WHMIS (Canada):

CLASS B-2: Flammable liquid with a flash point lower than 37.8°C (100°F). CLASS D-2A: Material causing other toxic effects (VERY TOXIC).

## DSCL (EEC):

R11- Highly flammable. R38- Irritating to skin. R41- Risk of serious damage to eyes.

#### HMIS (U.S.A.):

Health Hazard: 2 Fire Hazard: 3 Reactivity: 0

Personal Protection: h

#### National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 3
Reactivity: 0
Specific hazard:

#### **Protective Equipment:**

Gloves. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Splash goggles.

#### **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

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# Material Safety Data Sheet Naphthalene MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Naphthalene

Catalog Codes: SLN1789, SLN2401

CAS#: 91-20-3

**RTECS: QJ0525000** 

TSCA: TSCA 8(b) inventory: Naphthalene

CI#: Not available.

Synonym:

Chemical Name: Not available.

**Chemical Formula:** C10H8

**Contact Information:** 

Sciencelab.com, Inc. 14025 Smith Rd. Houston, Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400

Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

## **Section 2: Composition and Information on Ingredients**

#### Composition:

Name	CAS#	% by Weight
Naphthalene	91-20-3	100

**Toxicological Data on Ingredients:** Naphthalene: ORAL (LD50): Acute: 490 mg/kg [Rat]. 533 mg/kg [Mouse]. 1200 mg/kg [Guinea pig]. DERMAL (LD50): Acute: 20001 mg/kg [Rabbit]. VAPOR (LC50): Acute: 170 ppm 4 hour(s) [Rat].

#### Section 3: Hazards Identification

#### **Potential Acute Health Effects:**

Very hazardous in case of ingestion. Hazardous in case of eye contact (irritant), of inhalation. Slightly hazardous in case of skin contact (irritant, permeator). Severe over-exposure can result in death.

#### **Potential Chronic Health Effects:**

CARCINOGENIC EFFECTS: A4 (Not classifiable for human or animal.) by ACGIH. MUTAGENIC EFFECTS: Not available. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Classified Development toxin [POSSIBLE]. The substance is toxic to blood, kidneys, the nervous system, the reproductive system, liver, mucous membranes, gastrointestinal tract, upper respiratory tract, central nervous system (CNS). Repeated or prolonged exposure to the substance can produce target organs damage. Repeated exposure to an highly toxic material may produce general deterioration of health by an accumulation in one or many human organs.

#### Section 4: First Aid Measures

#### Eye Contact:

Check for and remove any contact lenses. Immediately flush eyes with running water for at least 15 minutes, keeping eyelids open. Cold water may be used. Do not use an eye ointment. Seek medical attention.

#### **Skin Contact:**

After contact with skin, wash immediately with plenty of water. Gently and thoroughly wash the contaminated skin with running water and non-abrasive soap. Be particularly careful to clean folds, crevices, creases and groin. Cover the irritated skin with an emollient. If irritation persists, seek medical attention. Wash contaminated clothing before reusing.

Serious Skin Contact: Not available.

**Inhalation:** Allow the victim to rest in a well ventilated area. Seek immediate medical attention.

#### Serious Inhalation:

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. WARNING: It may be hazardous to the person providing aid to give mouth-to-mouth resuscitation when the inhaled material is toxic, infectious or corrosive. Seek immediate medical attention.

### Ingestion:

Do not induce vomiting. Examine the lips and mouth to ascertain whether the tissues are damaged, a possible indication that the toxic material was ingested; the absence of such signs, however, is not conclusive. Loosen tight clothing such as a collar, tie, belt or waistband. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek immediate medical attention.

Serious Ingestion: Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: Flammable.

**Auto-Ignition Temperature:** 567°C (1052.6°F)

Flash Points: CLOSED CUP: 88°C (190.4°F). OPEN CUP: 79°C (174.2°F).

Flammable Limits: LOWER: 0.9% UPPER: 5.9%

**Products of Combustion:** These products are carbon oxides (CO, CO2).

Fire Hazards in Presence of Various Substances: Not available.

#### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available.

## **Fire Fighting Media and Instructions:**

Flammable solid. SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use water spray or fog. Cool containing vessels with water jet in order to prevent pressure build-up, autoignition or explosion.

Special Remarks on Fire Hazards: Not available.

**Special Remarks on Explosion Hazards:** Not available.

## **Section 6: Accidental Release Measures**

**Small Spill:** Use appropriate tools to put the spilled solid in a convenient waste disposal container.

#### Large Spill:

Flammable solid. Stop leak if without risk. Do not touch spilled material. Use water spray curtain to divert vapor drift. Prevent entry into sewers, basements or confined areas; dike if needed. Eliminate all ignition sources. Call for assistance on disposal. Be careful that the product is not present at a concentration level above TLV. Check TLV on the MSDS and with local authorities.

## **Section 7: Handling and Storage**

#### Precautions:

Keep locked up Keep away from heat. Keep away from sources of ignition. Ground all equipment containing material. Do not ingest. Do not breathe dust. Avoid contact with eyes Wear suitable protective clothing In case of insufficient ventilation, wear suitable respiratory equipment If ingested, seek medical advice immediately and show the container or the label. Keep away from incompatibles such as oxidizing agents.

#### Storage:

Flammable materials should be stored in a separate safety storage cabinet or room. Keep away from heat. Keep away from sources of ignition. Keep container tightly closed. Keep in a cool, well-ventilated place. Ground all equipment containing material. Keep container dry. Keep in a cool place.

## **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Use process enclosures, local exhaust ventilation, or other engineering controls to keep airborne levels below recommended exposure limits. If user operations generate dust, fume or mist, use ventilation to keep exposure to airborne contaminants below the exposure limit.

#### **Personal Protection:**

Splash goggles. Lab coat. Dust respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

## Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Dust respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

## **Exposure Limits:**

Israel: TWA: 10 (ppm) TWA: 10 STEL: 15 (ppm) from ACGIH (TLV) [1995] TWA: 52 STEL: 79 (mg/m3) from ACGIH [1995] Australia: STEL: 15 (ppm) Consult local authorities for acceptable exposure limits.

## **Section 9: Physical and Chemical Properties**

Physical state and appearance: Solid. (Crystalline solid.)

Odor: Aromatic.

Taste: Not available.

Molecular Weight: 128.19 g/mole

Color: White.

pH (1% soln/water): Not available. Boiling Point: 218°C (424.4°F) Melting Point: 80.2°C (176.4°F)

**Critical Temperature:** Not available. **Specific Gravity:** 1.162 (Water = 1)

Vapor Pressure: Not applicable.

Vapor Density: 4.4 (Air = 1)

Volatility: Not available.

Odor Threshold: 0.038 ppm

Water/Oil Dist. Coeff.: Not available.

Ionicity (in Water): Not available.

#### **Dispersion Properties:**

Partially dispersed in hot water, methanol, n-octanol. Very slightly dispersed in cold water. See solubility in methanol, n-octanol.

#### Solubility:

Partially soluble in methanol, n-octanol. Very slightly soluble in cold water, hot water.

## Section 10: Stability and Reactivity Data

Stability: The product is stable.

**Instability Temperature:** Not available. **Conditions of Instability:** Not available.

**Incompatibility with various substances:** Highly reactive with oxidizing agents.

**Corrosivity:** Non-corrosive in presence of glass. **Special Remarks on Reactivity:** Not available.

Special Remarks on Corrosivity: May attack some forms of rubber and plastic

Polymerization: No.

## **Section 11: Toxicological Information**

Routes of Entry: Absorbed through skin. Dermal contact. Eye contact. Inhalation. Ingestion.

### **Toxicity to Animals:**

WARNING: THE LC50 VALUES HEREUNDER ARE ESTIMATED ON THE BASIS OF A 4-HOUR EXPOSURE. Acute oral toxicity (LD50): 490 mg/kg [Rat]. Acute dermal toxicity (LD50): 20001 mg/kg [Rabbit]. Acute toxicity of the vapor (LC50): 170 ppm 4 hour(s) [Rat].

#### **Chronic Effects on Humans:**

CARCINOGENIC EFFECTS: A4 (Not classifiable for human or animal.) by ACGIH. DEVELOPMENTAL TOXICITY: Classified Development toxin [POSSIBLE]. The substance is toxic to blood, kidneys, the nervous system, the reproductive system, liver, mucous membranes, gastrointestinal tract, upper respiratory tract, central nervous system (CNS).

#### Other Toxic Effects on Humans:

Very hazardous in case of ingestion. Hazardous in case of inhalation. Slightly hazardous in case of skin contact (irritant, permeator).

Special Remarks on Toxicity to Animals: Not available.

Special Remarks on Chronic Effects on Humans: Not available.

Special Remarks on other Toxic Effects on Humans: Not available.

## Section 12: Ecological Information

**Ecotoxicity:** Ecotoxicity in water (LC50): 305.2 ppm 96 hour(s) [Trout].

BOD5 and COD: Not available.

#### **Products of Biodegradation:**

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

**Toxicity of the Products of Biodegradation:** The products of degradation are more toxic.

Special Remarks on the Products of Biodegradation: Not available.

## **Section 13: Disposal Considerations**

Waste Disposal:

## **Section 14: Transport Information**

**DOT Classification:** CLASS 4.1: Flammable solid. **Identification:** : Naphthalene, refined: UN1334 PG: III **Special Provisions for Transport:** Marine Pollutant

## **Section 15: Other Regulatory Information**

## Federal and State Regulations:

Rhode Island RTK hazardous substances: Naphthalene Pennsylvania RTK: Naphthalene Florida: Naphthalene Minnesota: Naphthalene Massachusetts RTK: Naphthalene TSCA 8(b) inventory: Naphthalene TSCA 8(a) PAIR: Naphthalene TSCA 8(d) H and S data reporting: Naphthalene: 06/01/87 SARA 313 toxic chemical notification and release reporting: Naphthalene: 1% CERCLA: Hazardous substances.: Naphthalene: 100 lbs. (45.36 kg)

## Other Regulations:

OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200). EINECS: This product is on the European Inventory of Existing Commercial Chemical Substances.

#### Other Classifications:

#### WHMIS (Canada):

CLASS B-4: Flammable solid. CLASS D-1B: Material causing immediate and serious toxic effects (TOXIC). CLASS D-2B: Material causing other toxic effects (TOXIC).

## DSCL (EEC):

R36- Irritating to eyes. R40- Possible risks of irreversible effects. R48/22- Harmful: danger of serious damage to health by prolonged exposure if swallowed. R48/23- Toxic: danger of serious damage to health by prolonged exposure through inhalation. R63- Possible risk of harm to the unborn child.

## HMIS (U.S.A.):

Health Hazard: 2

Fire Hazard: 2

Reactivity: 0

Personal Protection: E

#### National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 2

Reactivity: 0

Specific hazard:

## **Protective Equipment:**

Gloves. Lab coat. Dust respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Splash goggles.

## **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

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## MATERIAL SAFETY DATA SHEET

## 1. CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

MATHESON TRI-GAS, INC. Emergency Contact:

150 Allen Road Suite 302 CHEMTREC 1-800-424-9300

Basking Ridge, New Jersey 07920 Calls Originating Outside the US:

**Information: 1-800-416-2505** 703-527-3887 (Collect Calls Accepted)

SUBSTANCE: BUTYL BENZENE

TRADE NAMES/SYNONYMS:

MTG MSDS 139; BUTYLBENZENE; 1-PHENYLBUTANE; N-BUTYLBENZENE; UN 2709;

MAT03530; RTECS CY9070000

**CHEMICAL FAMILY:** hydrocarbons, aromatic

**CREATION DATE:** Jan 24 1989 **REVISION DATE:** Dec 11 2008

## 2. COMPOSITION, INFORMATION ON INGREDIENTS

**COMPONENT:** BUTYL BENZENE

CAS NUMBER: 104-51-8 PERCENTAGE: 100

## 3. HAZARDS IDENTIFICATION

NFPA RATINGS (SCALE 0-4): HEALTH=2 FIRE=2 REACTIVITY=0

## **EMERGENCY OVERVIEW:**

**COLOR:** colorless

PHYSICAL FORM: liquid

**ODOR:** odorless

MAJOR HEALTH HAZARDS: respiratory tract irritation, skin irritation, eye irritation, central nervous

system depression

PHYSICAL HAZARDS: Combustible liquid and vapor.

## POTENTIAL HEALTH EFFECTS:

**INHALATION:** 

**SHORT TERM EXPOSURE:** irritation, vomiting, headache, symptoms of drunkenness, coma







LONG TERM EXPOSURE: lung damage

**SKIN CONTACT:** 

**SHORT TERM EXPOSURE:** irritation, headache, symptoms of drunkenness **LONG TERM EXPOSURE:** same as effects reported in short term exposure

**EYE CONTACT:** 

**SHORT TERM EXPOSURE:** irritation, tearing

LONG TERM EXPOSURE: same as effects reported in short term exposure

**INGESTION:** 

SHORT TERM EXPOSURE: vomiting, headache, symptoms of drunkenness, coma

LONG TERM EXPOSURE: lung damage

## 4. FIRST AID MEASURES

**INHALATION:** If adverse effects occur, remove to uncontaminated area. Give artificial respiration if not breathing. If breathing is difficult, oxygen should be administered by qualified personnel. Get immediate medical attention.

**SKIN CONTACT:** Wash skin with soap and water for at least 15 minutes while removing contaminated clothing and shoes. Get medical attention, if needed. Thoroughly clean and dry contaminated clothing and shoes before reuse.

**EYE CONTACT:** Flush eyes with plenty of water for at least 15 minutes. Then get immediate medical attention.

**INGESTION:** DO NOT induce vomiting. Never make an unconscious person vomit or drink fluids. If vomiting occurs, keep head lower than hips to help prevent aspiration. If person is unconscious, turn head to side. Get medical attention.

**NOTE TO PHYSICIAN:** For inhalation, consider oxygen. For ingestion, consider gastric lavage, catharsis and activated charcoal slurry.

## 5. FIRE FIGHTING MEASURES

**FIRE AND EXPLOSION HAZARDS:** Severe fire hazard. Vapor/air mixtures are explosive above flash point. The vapor is heavier than air. Vapors or gases may ignite at distant ignition sources and flash back.

**EXTINGUISHING MEDIA:** regular dry chemical, carbon dioxide, water, regular foam

Large fires: Use regular foam or flood with fine water spray.

**FIRE FIGHTING:** Move container from fire area if it can be done without risk. Cool containers with water spray until well after the fire is out. Stay away from the ends of tanks. For fires in cargo or storage area: Cool containers with water from unmanned hose holder or monitor nozzles until well after fire is out. If this is impossible then take the following precautions: Keep unnecessary people away, isolate hazard area and deny



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entry. Let the fire burn. Withdraw immediately in case of rising sound from venting safety device or any discoloration of tanks due to fire. For tank, rail car or tank truck: Evacuation radius: 800 meters (1/2 mile). Do not attempt to extinguish fire unless flow of material can be stopped first. Flood with fine water spray. Do not scatter spilled material with high-pressure water streams. Cool containers with water spray until well after the fire is out. Apply water from a protected location or from a safe distance. Avoid inhalation of material or combustion by-products. Stay upwind and keep out of low areas.

FLASH POINT: 160 F (71 C) (OC) LOWER FLAMMABLE LIMIT: 0.8% UPPER FLAMMABLE LIMIT: 5.8% AUTOIGNITION: 770 F (410 C)

FLAMMABILITY CLASS (OSHA): IIIA

## 6. ACCIDENTAL RELEASE MEASURES

#### OCCUPATIONAL RELEASE:

Avoid heat, flames, sparks and other sources of ignition. Stop leak if possible without personal risk. Reduce vapors with water spray. Small spills: Absorb with sand or other non-combustible material. Collect spilled material in appropriate container for disposal. Large spills: Dike for later disposal. Remove sources of ignition. Keep unnecessary people away, isolate hazard area and deny entry.

## 7. HANDLING AND STORAGE

**STORAGE:** Store and handle in accordance with all current regulations and standards.

## 8. EXPOSURE CONTROLS, PERSONAL PROTECTION

## **EXPOSURE LIMITS:**

## **BUTYL BENZENE:**

No occupational exposure limits established.

**VENTILATION:** Provide local exhaust ventilation system. Ensure compliance with applicable exposure limits.

**EYE PROTECTION:** Wear splash resistant safety goggles with a faceshield. Provide an emergency eye wash fountain and quick drench shower in the immediate work area.

**CLOTHING:** Wear appropriate chemical resistant clothing.

**GLOVES:** Wear appropriate chemical resistant gloves.

**RESPIRATOR:** Under conditions of frequent use or heavy exposure, respiratory protection may be needed. Respiratory protection is ranked in order from minimum to maximum. Consider warning properties before





use.

Any supplied-air respirator with a full facepiece that is operated in a pressure-demand or other positive-pressure mode.

Any self-contained breathing apparatus that has a full facepiece and is operated in a pressure-demand or other positive-pressure mode.

## 9. PHYSICAL AND CHEMICAL PROPERTIES

PHYSICAL STATE: liquid

**COLOR:** colorless **ODOR:** odorless

MOLECULAR WEIGHT: 134.21 MOLECULAR FORMULA: C10-H14 BOILING POINT: 356 F (180 C) FREEZING POINT: -116 F (-82 C) VAPOR PRESSURE: 1 mmHg @ 23 C

VAPOR DENSITY (air=1): 4.6

SPECIFIC GRAVITY (water=1): 0.9 WATER SOLUBILITY: insoluble

PH: Not available

**VOLATILITY:** Not available

**ODOR THRESHOLD:** Not available **EVAPORATION RATE:** Not available

**COEFFICIENT OF WATER/OIL DISTRIBUTION:** Not available

**SOLVENT SOLUBILITY: Miscible:** alcohol, ether, benzene

## 10. STABILITY AND REACTIVITY

**REACTIVITY:** Stable at normal temperatures and pressure.

**CONDITIONS TO AVOID:** Avoid heat, flames, sparks and other sources of ignition. Containers may rupture or explode if exposed to heat. Keep out of water supplies and sewers.

**INCOMPATIBILITIES:** oxidizing materials

## **HAZARDOUS DECOMPOSITION:**

Thermal decomposition products: miscellaneous decomposition products

**POLYMERIZATION:** Will not polymerize.

## 11. TOXICOLOGICAL INFORMATION





## BUTYL BENZENE: LOCAL EFFECTS:

Irritant: inhalation, skin, eye

**TARGET ORGANS:** central nervous system

## 12. ECOLOGICAL INFORMATION

#### **ECOTOXICITY DATA:**

**INVERTEBRATE TOXICITY:** 340 ug/L 48 hour(s) EC50 (Immobilization) Water flea (Daphnia magna)

## 13. DISPOSAL CONSIDERATIONS

Dispose in accordance with all applicable regulations.

## 14. TRANSPORT INFORMATION

U.S. DOT 49 CFR 172.101:

**PROPER SHIPPING NAME:** Butyl benzenes

**ID NUMBER:** UN2709

**HAZARD CLASS OR DIVISION: 3** 

**PACKING GROUP: III** 

**LABELING REQUIREMENTS: 3** 

MARINE POLLUTANT: BUTYL BENZENE

#### CANADIAN TRANSPORTATION OF DANGEROUS GOODS:

**SHIPPING NAME:** Butylbenzenes

UN NUMBER: UN2709

CLASS: 3

PACKING GROUP/CATEGORY: III

## 15. REGULATORY INFORMATION

#### **U.S. REGULATIONS:**

CERCLA SECTIONS 102a/103 HAZARDOUS SUBSTANCES (40 CFR 302.4): Not regulated.

**SARA TITLE III SECTION 302 EXTREMELY HAZARDOUS SUBSTANCES (40 CFR 355 Subpart B):** Not regulated.

**SARA TITLE III SECTION 304 EXTREMELY HAZARDOUS SUBSTANCES (40 CFR 355 Subpart C):** Not regulated.

SARA TITLE III SARA SECTIONS 311/312 HAZARDOUS CATEGORIES (40 CFR 370 Subparts B







**and C**):

ACUTE: Yes CHRONIC: No

FIRE: Yes

REACTIVE: No

SUDDEN RELEASE: No

SARA TITLE III SECTION 313 (40 CFR 372.65): Not regulated.

OSHA PROCESS SAFETY (29 CFR 1910.119): Not regulated.

**STATE REGULATIONS:** 

California Proposition 65: Not regulated.

**CANADIAN REGULATIONS:** 

WHMIS CLASSIFICATION: Not determined.

**NATIONAL INVENTORY STATUS:** 

**U.S. INVENTORY (TSCA):** Listed on inventory.

TSCA 12(b) EXPORT NOTIFICATION: Not listed.

CANADA INVENTORY (DSL/NDSL): Not determined.

## 16. OTHER INFORMATION

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# **Material Safety Data Sheet**

HAZARD WARNINGS	RISK PHRASES	PROTECTIVE CLOTHING
	Combustible material; avoid heat and sources of ignition. The health risks of this compound have not been fully determined Exposure may cause irritation of the skin, eyes, and respiratory system.	

Section I.	Chemical Product and Company Id	lentification	
Chemical Name	n-Propylbenzene		
Catalog Number	P0523	Supplier	TCI America 9211 N. Harborgate St.
Synonym	1-Phenylpropane		Portland OR 1-800-423-8616
Chemical Formula	CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> C <sub>6</sub> H <sub>5</sub>		***************************************
CAS Number	103-65-1	In case of Emergency Call	Chemtrec® (800) 424-9300 (U.S.) (703) 527-3887 (International)
		4,,,,,,,	

Section II. Composition and Information on Ingredients					
Chemical Nan	ne	CAS Number	Percent (%)	TLV/PEL	Toxicology Data
n-Propylbenze	ne	103-65-1	Min. 99.0 (GC)		Rat LD <sub>50</sub> (inhalation) 65000ppm/2H Rat LD <sub>50</sub> (oral) 6040mg/kg

Section III.	Hazards identification
Acute Health Effects	No specific information is available in our data base regarding the toxic effects of this material for humans. However, exposure to any chemical should be kept to a minimum. Skin and eye contact may result in irritation. May be harmful if inhaled or ingested. Always follow safe industrial hygiene practices and wear proper protective equipment when handling this compound.
Chronic Health Effect	CARCINOGENIC EFFECTS: Not available. MUTAGENIC EFFECTS: Not available. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITYNot available. Repeated exposure to an highly toxic material may produce general deterioration of health by an accumulation in one or many human organs.

Eye Contact	Check for and remove any contact lenses. IMMEDIATELY flush eyes with runing water for at least 15 minutes. keeping eyelids open. COLD water may be used. DO NOT use an eye oitment. Flush eyes with running water for a minimum of 15 minutes, occasionally lifting the upper eyelids. Seek medical attention. Treat symptomatically and supportively.
Skin Contact	After contact with skin, wash immediately with plenty of water. Gently and thorough wash the contaminated skin with running water and non-abrasive soap. Be particularly careful to clean folds, crevices, creases and groin. COLD water may be used. Cover the irritated skin with an emollient. Seek medical attention. Treat symptomatically and supportively. Wash any contaminated clothing before reusing.
Inhalation	Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform artificial respiration. WARNING: It may be dangerous to the person providing aid to give mouth-to-mouth resuscitation when the inhaled material is toxic, infectious or corrosive. Seek immediate medical attention and, if possible, show the chemical label. Treat symptomatically and supportively.
Ingestion	INDUCE VOMITING by sticking finger in throat. Lower the head so that the vomit will not reenter the mouth and throat. Loosen tight clothing such as a collar, tie, belt, or waistband. If the victim is not breathing, administer artificial respiration. Examine the lips and mouth to ascertain whether the tissues are damaged, a possible indication that the toxic material was ingested; the absence of such signs, however, is not conclusive. Seek immediate medical attention and, if possible, show the chemical label. Treat symptomatically and supportively.

Section V.	Fire and Explosion Data	a	
Flammability	Combustible.	Auto-Ignition	Not available.
Flash Points	47.8°C (118°F).	Flammable Limits	Not available.
Combustion Products	These products are toxic carbon of	oxides (CO, CO <sub>2</sub> ).	
Fire Hazards	No specific information is available	e regarding the flammability of this compo	ound in the presence of various materials.
Explosion Hazards	Risks of explosion of the product i	in presence of mechanical impact: Not avin presence of static discharge: Not available regarding the risks of explosion.	
Continued of	n Next Page	Emergency phone nu	mber (800) 424-9300

Section IV.

First Aid Measures

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Fire Fighting Media and Instructions

SMALL FIRE: Use DRY chemicals, CO<sub>2</sub>, alcohol foam or water spray.

LARGE FIRE: Use alcohol foam, water spray or fog.

#### Section VI. Accidental Release Measures

Spill Cleanup Instructions Combustible liquid.

Keep away from heat and sources of ignition. Mechanical exhaust required. Stop leak if without risk. Absorb with DRY earth, sand or other non-combustible material. DO NOT touch spilled material. Prevent entry into sewers, basements or confined areas; dike if needed. Eliminate all sources of ignition. Consult federal, state, and/or local authorities for assistance on disposal.

## Section VII. Handling and Storage

Handling and Storage Information COMBUSTIBLE. Handle with caution and minimize exposure. DO NOT ingest. Do not breathe gas, fumes, vapor or spray. Wear suitable protective clothing. If ingested, seek medical advice immediately and show the container or the label. Treat symptomatically and supportively.

Always store away from incompatible compounds such as oxidizing agents.

## Section VIII. Exposure Controls/Personal Protection

**Engineering Controls** 

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value. Ensure that eyewash station and safety shower is proximal to the work-station location.

Personal Protection

Splash goggles. Lab coat. Dust respirator. Boots. Gloves. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.



**Exposure Limits** 

Not available

Section IX. Pl	hysical and Chemical	Properties	
Physical state @ 20°C	Liquid.	Solubility	Very slightly soluble in water. Soluble in alcohol, ether, all proportions in
Specific Gravity	0.86		acetone, benzene, and petroleum ether.
Molecular Weight	120.19	Partition Coefficient	Not available.
<b>Boiling Point</b>	159°C (318.2°F)	Vapor Pressure	Not available.
Melting Point	-99°C (-146.2°F)	Vapor Density	Not available.
Refractive Index	1.4920 @ 20°C	Volatility	Not available.
Critical Temperature	Not available.	Odor	Not available.
Viscosity	Not available.	Taste	Not available.

## Section X. Stability and Reactivity Data

Stability This material is stable if stored under proper conditions. (See Section VII for instructions)

Conditions of Instability Avoid excessive heat and light.

Incompatibilities Reactive with strong oxidizing agents.

## Section XI. Toxicological Information

RTECS Number DA8750000

Routes of Exposure Eye contact. Ingestion. Inhalation.

Toxicity Data Rat LD<sub>50</sub> (inhalation) 65000ppm/2H

Rat LD<sub>50</sub> (oral) 6040mg/kg

Chronic Toxic Effects CARCINOGENIC EFFECTS: Not available.

MUTAGENIC EFFECTS: Not available.
TERATOGENIC EFFECTS: Not available.
DEVELOPMENTAL TOXICITYNot available.

Repeated exposure to an highly toxic material may produce general deterioration of health by an accumulation in one or

many human organs.

Acute Toxic Effects No specific information is available in our data base regarding the toxic effects of this material for humans. However,

exposure to any chemical should be kept to a minimum. Skin and eye contact may result in irritation. May be harmful if inhaled or ingested. Always follow safe industrial hygiene practices and wear proper protective equipment when handling

this compound.

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## Section XII. Ecological Information

Ecotoxicity

Not available.

**Environmental Fate** 

Not available.

## Section XIII. Disposal Considerations

Waste Disposal

Recycle to process, if possible. Consult your local or regional authorities. You may be able to dissolve or mix material with a combustible solvent and burn in a chemical incinerator equipped with an afterburner and scrubber system. Observe all federal, state, and local regulations when disposing of the substance.

## Section XIV. Transport Information

DOT Classification

DOT CLASS 3: Flammable liquid.

PIN Number

UN2364

Proper Shipping Name

n-Propylbenzene

Packing Group (PG)

Ш

DOT Pictograms



## Section XV. Other Regulatory Information and Pictograms

TSCA Chemical Inventory

This compound is **ON** the EPA Toxic Substances Control Act (TSCA) inventory list.

(EPA)

WHMIS Classification

)11

WHMIS CLASS B-3: Combustible liquid with a flash point between 35°C (100°F) and 93.3°C (200°F).

(Canada)
EINECS Number (EEC)

203-132-9

**EEC Risk Statements** 

R10- Flammable.

R18- In use, may form flammable/explosive vapor-air mixture.

Japanese Regulatory Data

Not available

## Section XVI. Other Information

Version 1.0

Validated on 10/26/1998.

Printed 3/18/2005.

#### **Notice to Reader**

TCI laboratory chemicals are for research purposes only and are NOT intended for use as drugs, food additives, households, or pesticides. The information herein is believed to be correct, but does not claim to be all inclusive and should be used only as a guide. Neither the above named supplier nor any of its subsidiaries assumes any liability whatsoever for the accuracy or completeness of the information contained herein. Final determination of suitability of any material is the sole responsibility of the user. All chemical reagents must be handled with the recognition that their chemical, physiological, toxicological, and hazardous properties have not been fully investigated or determined. All chemical reagents should be handled only by individuals who are familiar with their potential hazards and who have been fully trained in proper safety, laboratory, and chemical handling procedures. Although certain hazards are described herein, we can not guarantee that these are the only hazards which exist. Our MSDS sheets are based only on data available at the time of shipping and are subject to change without notice as new information is obtained. Avoid long storage periods since the product is subject to degradation with age and may become more dangerous or hazardous. It is the responsibility of the user to request updated MSDS sheets for products that are stored for extended periods. Disposal of unused product must be undertaken by qualified personnel who are knowledgeable in all applicable regulations and follow all pertinent safety precautions including the use of appropriate protective equipment (e.g. protective goggles, protective clothing, breathing equipment, facial mask, fume hood). For proper handling and disposal, always comply with federal, state, and local regulations.

Printed 3/18/2005







# Material Safety Data Sheet Phenanthrene MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Phenanthrene

Catalog Codes: SLP1318

CAS#: 85-01-8

**RTECS:** SF7175000

TSCA: TSCA 8(b) inventory: Phenanthrene

CI#: Not available.

Synonym:

Chemical Name: Not available.

**Chemical Formula:** C14H10

**Contact Information:** 

**Sciencelab.com, Inc.** 14025 Smith Rd. Houston, Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400

Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

## **Section 2: Composition and Information on Ingredients**

## **Composition:**

Name	CAS#	% by Weight
Phenanthrene	85-01-8	100

Toxicological Data on Ingredients: Phenanthrene: ORAL (LD50): Acute: 700 mg/kg [Mouse].

#### **Section 3: Hazards Identification**

#### **Potential Acute Health Effects:**

Hazardous in case of skin contact (irritant, sensitizer), of eye contact (irritant), of ingestion, of inhalation. Slightly hazardous in case of skin contact (permeator).

#### **Potential Chronic Health Effects:**

CARCINOGENIC EFFECTS: Not available. MUTAGENIC EFFECTS: Not available. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Not available. Repeated or prolonged exposure is not known to aggravate medical condition.

## **Section 4: First Aid Measures**

#### **Eye Contact:**

Check for and remove any contact lenses. Immediately flush eyes with running water for at least 15 minutes, keeping eyelids open. Cold water may be used. Do not use an eye ointment. Seek medical attention.

#### Skin Contact:

After contact with skin, wash immediately with plenty of water. Gently and thoroughly wash the contaminated skin with running water and non-abrasive soap. Be particularly careful to clean folds, crevices, creases and groin. Cover the irritated skin with an emollient. If irritation persists, seek medical attention. Wash contaminated clothing before reusing.

#### **Serious Skin Contact:**

Wash with a disinfectant soap and cover the contaminated skin with an anti-bacterial cream. Seek immediate medical attention.

**Inhalation:** Allow the victim to rest in a well ventilated area. Seek immediate medical attention.

Serious Inhalation: Not available.

#### Ingestion:

Do not induce vomiting. Examine the lips and mouth to ascertain whether the tissues are damaged, a possible indication that the toxic material was ingested; the absence of such signs, however, is not conclusive. Loosen tight clothing such as a collar, tie, belt or waistband. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek immediate medical attention.

Serious Ingestion: Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: May be combustible at high temperature.

**Auto-Ignition Temperature:** Not available. **Flash Points:** OPEN CUP: 171°C (339.8°F).

Flammable Limits: Not available.

Products of Combustion: These products are carbon oxides (CO, CO2).

Fire Hazards in Presence of Various Substances: Not available.

### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available.

## Fire Fighting Media and Instructions:

SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use water spray, fog or foam. Do not use water jet.

Special Remarks on Fire Hazards: Not available.

Special Remarks on Explosion Hazards: Not available.

#### **Section 6: Accidental Release Measures**

#### **Small Spill:**

Use appropriate tools to put the spilled solid in a convenient waste disposal container. Finish cleaning by spreading water on the contaminated surface and dispose of according to local and regional authority requirements.

#### Large Spill:

Use a shovel to put the material into a convenient waste disposal container. Finish cleaning by spreading water on the contaminated surface and allow to evacuate through the sanitary system.

## **Section 7: Handling and Storage**

#### Precautions:

Keep away from heat. Keep away from sources of ignition. Empty containers pose a fire risk, evaporate the residue under a fume hood. Ground all equipment containing material. Do not ingest. Do not breathe dust. Wear suitable protective clothing In

case of insufficient ventilation, wear suitable respiratory equipment If ingested, seek medical advice immediately and show the container or the label. Avoid contact with skin and eyes

#### Storage:

Keep container dry. Keep in a cool place. Ground all equipment containing material. Keep container tightly closed. Keep in a cool, well-ventilated place. Combustible materials should be stored away from extreme heat and away from strong oxidizing agents.

## **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Use process enclosures, local exhaust ventilation, or other engineering controls to keep airborne levels below recommended exposure limits. If user operations generate dust, fume or mist, use ventilation to keep exposure to airborne contaminants below the exposure limit.

#### **Personal Protection:**

Splash goggles. Lab coat. Dust respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Dust respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

**Exposure Limits:** Not available.

## **Section 9: Physical and Chemical Properties**

Physical state and appearance: Solid.

Odor: Not available.

Taste: Not available.

Molecular Weight: 178.22 g/mole

Color: Not available.

pH (1% soln/water): Not available. Boiling Point: 340°C (644°F) Melting Point: 101°C (213.8°F)

Critical Temperature: Not available.

Specific Gravity: 1.179 (Water = 1)

Vapor Pressure: Not applicable.

Vapor Density: 6.14 (Air = 1)

Volatility: Not available.

Odor Threshold: Not available.

Water/Oil Dist. Coeff.: Not available.

Ionicity (in Water): Not available.

Dispersion Properties: Not available.

**Solubility:** Very slightly soluble in cold water.

## Section 10: Stability and Reactivity Data

**Stability:** The product is stable.

Instability Temperature: Not available.Conditions of Instability: Not available.

Incompatibility with various substances: Not available.

Corrosivity: Non-corrosive in presence of glass.

Special Remarks on Reactivity: Not available.

Special Remarks on Corrosivity: Not available.

Polymerization: No.

## **Section 11: Toxicological Information**

Routes of Entry: Eye contact. Inhalation. Ingestion.

Toxicity to Animals: Acute oral toxicity (LD50): 700 mg/kg [Mouse].

Chronic Effects on Humans: Not available.

Other Toxic Effects on Humans:

Hazardous in case of skin contact (irritant, sensitizer), of ingestion, of inhalation. Slightly hazardous in case of skin contact

(permeator).

Special Remarks on Toxicity to Animals: Not available.

Special Remarks on Chronic Effects on Humans: Not available.

Special Remarks on other Toxic Effects on Humans: Not available.

## **Section 12: Ecological Information**

Ecotoxicity: Not available.

BOD5 and COD: Not available.

**Products of Biodegradation:** 

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

**Toxicity of the Products of Biodegradation:** The products of degradation are more toxic.

Special Remarks on the Products of Biodegradation: Not available.

## **Section 13: Disposal Considerations**

**Waste Disposal:** 

## **Section 14: Transport Information**

**DOT Classification:** Not a DOT controlled material (United States).

**Identification:** Not applicable.

Special Provisions for Transport: Not applicable.

## **Section 15: Other Regulatory Information**

Federal and State Regulations: TSCA 8(b) inventory: Phenanthrene

Other Regulations: OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200).

Other Classifications:

WHMIS (Canada): Not controlled under WHMIS (Canada).

DSCL (EEC):

R36/38- Irritating to eyes and skin. R43- May cause sensitization by skin contact.

HMIS (U.S.A.):

Health Hazard: 2

Fire Hazard: 1
Reactivity: 0

Personal Protection: E

National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 1

Reactivity: 0

Specific hazard:

## **Protective Equipment:**

Gloves. Lab coat. Dust respirator. Be sure to use an approved/certified respirator or equivalent. Splash goggles.

#### **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

Created: 10/10/2005 11:16 AM

Last Updated: 11/06/2008 12:00 PM

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# Material Safety Data Sheet Tetrachloroethylene MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Tetrachloroethylene

Catalog Codes: SLT3220

CAS#: 127-18-4

RTECS: KX3850000

TSCA: TSCA 8(b) inventory: Tetrachloroethylene

CI#: Not available.

Synonym: Perchloroethylene; 1,1,2,2-

Tetrachloroethylene; Carbon bichloride; Carbon dichloride; Ankilostin; Didakene; Dilatin PT; Ethene, tetrachloro-; Ethylene tetrachloride; Perawin; Perchlor; Perclene; Perclene D; Percosolvel; Tetrachloroethene; Tetraleno;

Tetralex; Tetravec; Tetroguer; Tetropil

Chemical Name: Ethylene, tetrachloro-

Chemical Formula: C2-Cl4

**Contact Information:** 

Sciencelab.com, Inc. 14025 Smith Rd.

Houston, Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400

Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

## **Section 2: Composition and Information on Ingredients**

## **Composition:**

Name	CAS#	% by Weight
Tetrachloroethylene	127-18-4	100

**Toxicological Data on Ingredients:** Tetrachloroethylene: ORAL (LD50): Acute: 2629 mg/kg [Rat]. DERMAL (LD): Acute: >3228 mg/kg [Rabbit]. MIST(LC50): Acute: 34200 mg/m 8 hours [Rat]. VAPOR (LC50): Acute: 5200 ppm 4 hours [Mouse].

## **Section 3: Hazards Identification**

## **Potential Acute Health Effects:**

Hazardous in case of skin contact (irritant), of inhalation. Slightly hazardous in case of skin contact (permeator), of eye contact (irritant), of ingestion.

#### **Potential Chronic Health Effects:**

CARCINOGENIC EFFECTS: Classified A3 (Proven for animal.) by ACGIH. Classified 2A (Probable for human.) by IARC, 2 (anticipated carcinogen) by NTP. MUTAGENIC EFFECTS: Mutagenic for bacteria and/or yeast. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Not available. The substance may be toxic to kidneys, liver, peripheral nervous system, respiratory tract, skin, central nervous system (CNS). Repeated or prolonged exposure to the substance can produce target organs damage.

#### Section 4: First Aid Measures

#### **Eye Contact:**

Check for and remove any contact lenses. In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Get medical attention if irritation occurs.

#### Skin Contact:

In case of contact, immediately flush skin with plenty of water. Cover the irritated skin with an emollient. Remove contaminated clothing and shoes. Wash clothing before reuse. Thoroughly clean shoes before reuse. Get medical attention.

#### **Serious Skin Contact:**

Wash with a disinfectant soap and cover the contaminated skin with an anti-bacterial cream. Seek medical attention.

#### Inhalation

If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical attention if symptoms appear.

#### Serious Inhalation:

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek medical attention.

## Ingestion:

Do NOT induce vomiting unless directed to do so by medical personnel. Never give anything by mouth to an unconscious person. Loosen tight clothing such as a collar, tie, belt or waistband. Get medical attention if symptoms appear.

**Serious Ingestion:** Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: Non-flammable.

Auto-Ignition Temperature: Not applicable.

Flash Points: Not applicable.

Flammable Limits: Not applicable.

Products of Combustion: Not available.

Fire Hazards in Presence of Various Substances: Not applicable.

#### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available.

Fire Fighting Media and Instructions: Not applicable.

Special Remarks on Fire Hazards: Not available.

Special Remarks on Explosion Hazards: Not available.

## **Section 6: Accidental Release Measures**

Small Spill: Absorb with an inert material and put the spilled material in an appropriate waste disposal.

#### Large Spill:

Absorb with an inert material and put the spilled material in an appropriate waste disposal. Be careful that the product is not present at a concentration level above TLV. Check TLV on the MSDS and with local authorities.

## **Section 7: Handling and Storage**

#### Precautions:

Do not ingest. Do not breathe gas/fumes/ vapor/spray. Avoid contact with skin. Wear suitable protective clothing. In case of insufficient ventilation, wear suitable respiratory equipment. If ingested, seek medical advice immediately and show the container or the label. Keep away from incompatibles such as oxidizing agents, metals, acids, alkalis.

**Storage:** Keep container tightly closed. Keep container in a cool, well-ventilated area.

## **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value.

#### **Personal Protection:**

Safety glasses. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Vapor respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

## **Exposure Limits:**

TWA: 25 (ppm) from OSHA (PEL) [United States] TWA: 25 STEL: 100 (ppm) from ACGIH (TLV) [United States] TWA: 170 (mg/m3) from OSHA (PEL) [United States] Consult local authorities for acceptable exposure limits.

## **Section 9: Physical and Chemical Properties**

Physical state and appearance: Liquid.

Odor: Ethereal.

Taste: Not available.

Molecular Weight: 165.83 g/mole

Color: Clear Colorless.

pH (1% soln/water): Not available. Boiling Point: 121.3°C (250.3°F) Melting Point: -22.3°C (-8.1°F)

Critical Temperature: 347.1°C (656.8°F)

Specific Gravity: 1.6227 (Water = 1) Vapor Pressure: 1.7 kPa (@ 20°C)

**Vapor Density:** 5.7 (Air = 1) **Volatility:** Not available.

Odor Threshold: 5 - 50 ppm

Water/Oil Dist. Coeff.: The product is more soluble in oil; log(oil/water) = 3.4

Ionicity (in Water): Not available.Dispersion Properties: Not available.

## Solubility:

Miscible with alcohol, ether, chloroform, benzene, hexane. It dissolves in most of the fixed and volatile oils. Solubility in water: 0.015 g/100 ml @ 25 deg. C It slowly decomposes in water to yield Trichloroacetic and Hydrochloric acids.

## Section 10: Stability and Reactivity Data

Stability: The product is stable.

**Instability Temperature:** Not available.

Conditions of Instability: Incompatible materials

Incompatibility with various substances: Reactive with oxidizing agents, metals, acids, alkalis.

Corrosivity: Non-corrosive in presence of glass.

## **Special Remarks on Reactivity:**

Oxidized by strong oxidizing agents. Incompatible with sodium hydroxide, finely divided or powdered metals such as zinc, aluminum, magnesium, potassium, chemically active metals such as lithium, beryllium, barium. Protect from light.

Special Remarks on Corrosivity: Slowly corrodes aluminum, iron, and zinc.

Polymerization: Will not occur.

## **Section 11: Toxicological Information**

Routes of Entry: Absorbed through skin. Eye contact. Inhalation. Ingestion.

## **Toxicity to Animals:**

WARNING: THE LC50 VALUES HEREUNDER ARE ESTIMATED ON THE BASIS OF A 4-HOUR EXPOSURE. Acute oral toxicity (LD50): 2629 mg/kg [Rat]. Acute dermal toxicity (LD50): >3228 mg/kg [Rabbit]. Acute toxicity of the vapor (LC50): 5200 4 hours [Mouse].

#### **Chronic Effects on Humans:**

CARCINOGENIC EFFECTS: Classified A3 (Proven for animal.) by ACGIH. Classified 2A (Probable for human.) by IARC, 2 (Some evidence.) by NTP. MUTAGENIC EFFECTS: Mutagenic for bacteria and/or yeast. May cause damage to the following organs: kidneys, liver, peripheral nervous system, upper respiratory tract, skin, central nervous system (CNS).

#### Other Toxic Effects on Humans:

Hazardous in case of skin contact (irritant), of inhalation. Slightly hazardous in case of skin contact (permeator), of ingestion.

## **Special Remarks on Toxicity to Animals:**

Lowest Publishe Lethal Dose/Conc: LDL [Rabbit] - Route: Oral; Dose: 5000 mg/kg LDL [Dog] - Route: Oral; Dose: 4000 mg/kg LDL [Cat] - Route: Oral; Dose: 4000 mg/kg

#### Special Remarks on Chronic Effects on Humans:

May cause adverse reproductive effects and birth defects(teratogenic). May affect genetic material (mutagenic). May cause cancer.

#### **Special Remarks on other Toxic Effects on Humans:**

Acute Potential Health Effects: Skin: Causes skin irritation with possible dermal blistering or burns. Symtoms may include redness, itching, pain, and possible dermal blistering or burns. It may be absorbed through the skin with possible systemic effects. A single prolonged skin exposure is not likely to result in the material being absorbed in harmful amounts. Eyes: Contact causes transient eye irritation, lacrimation. Vapors cause eye/conjunctival irritation. Symptoms may include redness and pain. Inhalation: The main route to occupational exposure is by inhalation since it is readily absorbed through the lungs. It causes respiratory tract irritation, . It can affect behavior/central nervous system (CNS depressant and anesthesia ranging from slight inebriation to death, vertigo, somnolence, anxiety, headache, excitement, hallucinations, muscle incoordination, dizziness, lightheadness, disorentiation, seizures, enotional instability, stupor, coma). It may cause pulmonary edema Ingestion: It can cause nausea, vomiting, anorexia, diarrhea, bloody stool. It may affect the liver, urinary system (proteinuria, hematuria, renal failure, renal tubular disorder), heart (arrhythmias). It may affect behavior/central nervous system with symptoms similar to that of inhalation. Chronic Potential Health Effects: Skin: Prolonged or repeated skin contact may result in excessive drying of the skin, and irritation. Ingestion/Inhalation: Chronic exposure can affect the liver(hepatitis,fatty liver degeneration), kidneys, spleen, and heart (irregular heartbeat/arrhythmias, cardiomyopathy, abnormal EEG), brain, behavior/central nervous system/peripheral nervous system (impaired memory, numbness of extremeties, peripheral neuropathy and other

## **Section 12: Ecological Information**

## **Ecotoxicity:**

Ecotoxicity in water (LC50): 18.4 mg/l 96 hours [Fish (Fatthead Minnow)]. 18 mg/l 48 hours [Daphnia (daphnia)]. 5 mg/l 96 hours [Fish (Rainbow Trout)]. 13 mg/l 96 hours [Fish (Bluegill sunfish)].

BOD5 and COD: Not available.

## **Products of Biodegradation:**

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

**Toxicity of the Products of Biodegradation:** The product itself and its products of degradation are not toxic.

Special Remarks on the Products of Biodegradation: Not available.

## **Section 13: Disposal Considerations**

#### Waste Disposal:

Waste must be disposed of in accordance with federal, state and local environmental control regulations.

## **Section 14: Transport Information**

**DOT Classification:** CLASS 6.1: Poisonous material. **Identification:** : Tetrachloroethylene UNNA: 1897 PG: III **Special Provisions for Transport:** Marine Pollutant

## **Section 15: Other Regulatory Information**

#### Federal and State Regulations:

California prop. 65: This product contains the following ingredients for which the State of California has found to cause cancer, birth defects or other reproductive harm, which would require a warning under the statute: Tetrachloroethylene California prop. 65: This product contains the following ingredients for which the State of California has found to cause cancer which would require a warning under the statute: Tetrachloroethylene Connecticut hazardous material survey.: Tetrachloroethylene Illinois toxic substances disclosure to employee act: Tetrachloroethylene Illinois chemical safety act: Tetrachloroethylene New York release reporting list: Tetrachloroethylene Rhode Island RTK hazardous substances: Tetrachloroethylene Pennsylvania RTK: Tetrachloroethylene Minnesota: Tetrachloroethylene Michigan critical material: Tetrachloroethylene Massachusetts RTK: Tetrachloroethylene Massachusetts spill list: Tetrachloroethylene New Jersey: Tetrachloroethylene New Jersey spill list: Tetrachloroethylene Louisiana spill reporting: Tetrachloroethylene California Director's List of Hazardous Substances: Tetrachloroethylene TSCA 8(b) inventory: Tetrachloroethylene TSCA 8(d) H and S data reporting: Tetrachloroethylene: Effective date: 6/1/87; Sunset date: 6/1/97 SARA 313 toxic chemical notification and release reporting: Tetrachloroethylene CERCLA: Hazardous substances:: Tetrachloroethylene: 100 lbs. (45.36 kg)

#### Other Regulations:

OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200). EINECS: This product is on the European Inventory of Existing Commercial Chemical Substances.

#### Other Classifications:

#### WHMIS (Canada):

CLASS D-1B: Material causing immediate and serious toxic effects (TOXIC). CLASS D-2A: Material causing other toxic effects (VERY TOXIC).

## DSCL (EEC):

R40- Possible risks of irreversible effects. R51/53- Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment. S23- Do not breathe gas/fumes/vapour/spray S26- In case of contact with eyes, rinse immediately with plenty of water and seek medical advice. S37- Wear suitable gloves. S61- Avoid release to the environment. Refer to special instructions/Safety data sheets.

HMIS (U.S.A.):

Health Hazard: 2

Fire Hazard: 0

Reactivity: 0

Personal Protection: g

National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 0
Reactivity: 0

Specific hazard:

#### **Protective Equipment:**

Gloves. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Safety glasses.

## **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

Created: 10/10/2005 08:29 PM

Last Updated: 11/06/2008 12:00 PM

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# Material Safety Data Sheet Toluene MSDS

## **Section 1: Chemical Product and Company Identification**

Product Name: Toluene

Catalog Codes: SLT2857, SLT3277

CAS#: 108-88-3

RTECS: XS5250000

TSCA: TSCA 8(b) inventory: Toluene

CI#: Not available.

**Synonym:** Toluol, Tolu-Sol; Methylbenzene; Methacide;

Phenylmethane; Methylbenzol

Chemical Name: Toluene

Chemical Formula: C6-H5-CH3 or C7-H8

#### **Contact Information:**

Sciencelab.com, Inc. 14025 Smith Rd. Houston, Texas 77396

US Sales: **1-800-901-7247** 

International Sales: 1-281-441-4400
Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

# Section 2: Composition and Information on Ingredients

## Composition:

	Neight
Toluene 108-88-3 100	

**Toxicological Data on Ingredients:** Toluene: ORAL (LD50): Acute: 636 mg/kg [Rat]. DERMAL (LD50): Acute: 14100 mg/kg [Rabbit]. VAPOR (LC50): Acute: 49000 mg/m 4 hours [Rat]. 440 ppm 24 hours [Mouse].

#### Section 3: Hazards Identification

## **Potential Acute Health Effects:**

Hazardous in case of skin contact (irritant), of eye contact (irritant), of ingestion, of inhalation. Slightly hazardous in case of skin contact (permeator).

#### **Potential Chronic Health Effects:**

CARCINOGENIC EFFECTS: A4 (Not classifiable for human or animal.) by ACGIH, 3 (Not classifiable for human.) by IARC. MUTAGENIC EFFECTS: Not available. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Not available. The substance may be toxic to blood, kidneys, the nervous system, liver, brain, central nervous system (CNS). Repeated or prolonged exposure to the substance can produce target organs damage.

## **Section 4: First Aid Measures**

## **Eye Contact:**

Check for and remove any contact lenses. In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Get medical attention.

#### Skin Contact:

In case of contact, immediately flush skin with plenty of water. Cover the irritated skin with an emollient. Remove contaminated clothing and shoes. Wash clothing before reuse. Thoroughly clean shoes before reuse. Get medical attention.

#### **Serious Skin Contact:**

Wash with a disinfectant soap and cover the contaminated skin with an anti-bacterial cream. Seek immediate medical attention.

#### Inhalation:

If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical attention.

#### Serious Inhalation:

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. WARNING: It may be hazardous to the person providing aid to give mouth-to-mouth resuscitation when the inhaled material is toxic, infectious or corrosive. Seek medical attention.

#### Ingestion:

Do NOT induce vomiting unless directed to do so by medical personnel. Never give anything by mouth to an unconscious person. If large quantities of this material are swallowed, call a physician immediately. Loosen tight clothing such as a collar, tie, belt or waistband.

Serious Ingestion: Not available.

## **Section 5: Fire and Explosion Data**

Flammability of the Product: Flammable.

**Auto-Ignition Temperature:** 480°C (896°F)

Flash Points: CLOSED CUP: 4.4444°C (40°F). (Setaflash) OPEN CUP: 16°C (60.8°F).

Flammable Limits: LOWER: 1.1% UPPER: 7.1%

**Products of Combustion:** These products are carbon oxides (CO, CO2).

## **Fire Hazards in Presence of Various Substances:**

Flammable in presence of open flames and sparks, of heat. Non-flammable in presence of shocks.

#### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Risks of explosion of the product in presence of static discharge: Not available.

#### **Fire Fighting Media and Instructions:**

Flammable liquid, insoluble in water. SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use water spray or fog.

Special Remarks on Fire Hazards: Not available.

#### Special Remarks on Explosion Hazards:

Toluene forms explosive reaction with 1,3-dichloro-5,5-dimethyl-2,4-imidazolididione; dinitrogen tetraoxide; concentrated nitric acid, sulfuric acid + nitric acid; N2O4; AgClO4; BrF3; Uranium hexafluoride; sulfur dichloride. Also forms an explosive mixture with tetranitromethane.

## **Section 6: Accidental Release Measures**

Small Spill: Absorb with an inert material and put the spilled material in an appropriate waste disposal.

## Large Spill:

Toxic flammable liquid, insoluble or very slightly soluble in water. Keep away from heat. Keep away from sources of ignition. Stop leak if without risk. Absorb with DRY earth, sand or other non-combustible material. Do not get water inside container. Do not touch spilled material. Prevent entry into sewers, basements or confined areas; dike if needed. Call for assistance on disposal. Be careful that the product is not present at a concentration level above TLV. Check TLV on the MSDS and with local authorities.

## Section 7: Handling and Storage

#### **Precautions:**

Keep away from heat. Keep away from sources of ignition. Ground all equipment containing material. Do not ingest. Do not breathe gas/fumes/ vapor/spray. Wear suitable protective clothing. In case of insufficient ventilation, wear suitable respiratory equipment. If ingested, seek medical advice immediately and show the container or the label. Avoid contact with skin and eyes. Keep away from incompatibles such as oxidizing agents.

## Storage:

Store in a segregated and approved area. Keep container in a cool, well-ventilated area. Keep container tightly closed and sealed until ready for use. Avoid all possible sources of ignition (spark or flame).

## **Section 8: Exposure Controls/Personal Protection**

## **Engineering Controls:**

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value. Ensure that eyewash stations and safety showers are proximal to the work-station location.

## **Personal Protection:**

Splash goggles. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Vapor respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

#### **Exposure Limits:**

TWA: 200 STEL: 500 CEIL: 300 (ppm) from OSHA (PEL) [United States] TWA: 50 (ppm) from ACGIH (TLV) [United States] SKIN TWA: 100 STEL: 150 from NIOSH [United States] TWA: 375 STEL: 560 (mg/m3) from NIOSH [United States] Consult local authorities for acceptable exposure limits.

## **Section 9: Physical and Chemical Properties**

Physical state and appearance: Liquid.

Odor: Sweet, pungent, Benzene-like.

Taste: Not available.

Molecular Weight: 92.14 g/mole

Color: Colorless.

**pH (1% soln/water):** Not applicable. **Boiling Point:** 110.6°C (231.1°F)

Melting Point: -95°C (-139°F)

Critical Temperature: 318.6°C (605.5°F)

Specific Gravity: 0.8636 (Water = 1)

Vapor Pressure: 3.8 kPa (@ 25°C)

Vapor Density: 3.1 (Air = 1)

Volatility: Not available.

Odor Threshold: 1.6 ppm

Water/Oil Dist. Coeff.: The product is more soluble in oil; log(oil/water) = 2.7

Ionicity (in Water): Not available.

**Dispersion Properties:** See solubility in water, diethyl ether, acetone.

Solubility:

Soluble in diethyl ether, acetone. Practically insoluble in cold water. Soluble in ethanol, benzene, chloroform, glacial acetic acid, carbon disulfide. Solubility in water: 0.561 q/l @ 25 deg. C.

## Section 10: Stability and Reactivity Data

Stability: The product is stable.

**Instability Temperature:** Not available.

Conditions of Instability: Heat, ignition sources (flames, sparks, static), incompatible materials

**Incompatibility with various substances:** Reactive with oxidizing agents.

Corrosivity: Non-corrosive in presence of glass.

#### Special Remarks on Reactivity:

Incompatible with strong oxidizers, silver perchlorate, sodium difluoride, Tetranitromethane, Uranium Hexafluoride. Frozen Bromine Trifluoride reacts violently with Toluene at -80 deg. C. Reacts chemically with nitrogen oxides, or halogens to form nitrotoluene, nitrobenzene, and nitrophenol and halogenated products, respectively.

Special Remarks on Corrosivity: Not available.

Polymerization: Will not occur.

## **Section 11: Toxicological Information**

Routes of Entry: Absorbed through skin. Dermal contact. Eye contact. Inhalation. Ingestion.

#### **Toxicity to Animals:**

WARNING: THE LC50 VALUES HEREUNDER ARE ESTIMATED ON THE BASIS OF A 4-HOUR EXPOSURE. Acute oral toxicity (LD50): 636 mg/kg [Rat]. Acute dermal toxicity (LD50): 14100 mg/kg [Rabbit]. Acute toxicity of the vapor (LC50): 440 24 hours [Mouse].

## **Chronic Effects on Humans:**

CARCINOGENIC EFFECTS: A4 (Not classifiable for human or animal.) by ACGIH, 3 (Not classifiable for human.) by IARC. May cause damage to the following organs: blood, kidneys, the nervous system, liver, brain, central nervous system (CNS).

## Other Toxic Effects on Humans:

Hazardous in case of skin contact (irritant), of ingestion, of inhalation. Slightly hazardous in case of skin contact (permeator).

## **Special Remarks on Toxicity to Animals:**

Lowest Published Lethal Dose: LDL [Human] - Route: Oral; Dose: 50 mg/kg LCL [Rabbit] - Route: Inhalation; Dose: 55000 ppm/40min

## **Special Remarks on Chronic Effects on Humans:**

Detected in maternal milk in human. Passes through the placental barrier in human. Embryotoxic and/or foetotoxic in animal. May cause adverse reproductive effects and birth defects (teratogenic). May affect genetic material (mutagenic)

## **Special Remarks on other Toxic Effects on Humans:**

Acute Potential Health Effects: Skin: Causes mild to moderate skin irritation. It can be absorbed to some extent through the skin. Eyes: Cauess mild to moderate eye irritation with a burning sensation. Splash contact with eyes also causes conjunctivitis, blepharospasm, corneal edema, corneal abraisons. This usually resolves in 2 days. Inhalation: Inhalation of vapor may cause respiratory tract irritation causing coughing and wheezing, and nasal discharge. Inhalation of high concentrations may affect behavior and cause central nervous system effects characterized by nausea, headache, dizziness, tremors, restlessness, lightheadedness, exhilaration, memory loss, insomnia, impaired reaction time, drowsiness, ataxia, hallucinations, somnolence, muscle contraction or spasticity, unconsciousness and coma. Inhalation of high concentration of vapor may also affect the cardiovascular system (rapid heart beat, heart palpitations, increased or decreased blood pressure, dysrhythmia, ), respiration (acute pulmonary edema, respiratory depression, apnea, asphyxia), cause vision disturbances and dilated pupils, and cause loss of appetite. Ingestion: Aspiration hazard. Aspiration of Toluene into the lungs may cause chemical pneumonitis. May cause irritation of the digestive tract with nausea, vomiting, pain. May have effects similar to that of acute inhalation. Chronic Potential Health Effects: Inhalation and Ingestion: Prolonged or repeated exposure via inhalation may cause central nervous system and cardiovascular symptoms similar to that of acute inhalation and ingestion as well liver damage/failure, kidney damage/failure (with hematuria, proteinuria, oliguria, renal tubular acidosis), brain damage, weight loss, blood (pigmented or nucleated red blood cells, changes in white blood cell count), bone marrow changes, electrolyte imbalances (Hypokalemia, Hypophostatemia), severe, muscle weakness and Rhabdomyolysis. Skin: Repeated or prolonged skin contact may cause defatting dermatitis.

## Section 12: Ecological Information

## **Ecotoxicity:**

Ecotoxicity in water (LC50): 313 mg/l 48 hours [Daphnia (daphnia)]. 17 mg/l 24 hours [Fish (Blue Gill)]. 13 mg/l 96 hours [Fish (Blue Gill)]. 56 mg/l 24 hours [Fish (Fathead minnow)]. 34 mg/l 96 hours [Fish (Fathead minnow)]. 56.8 ppm any hours [Fish (Goldfish)].

BOD5 and COD: Not available.

## **Products of Biodegradation:**

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

**Toxicity of the Products of Biodegradation:** The products of degradation are less toxic than the product itself.

Special Remarks on the Products of Biodegradation: Not available.

## **Section 13: Disposal Considerations**

## Waste Disposal:

Waste must be disposed of in accordance with federal, state and local environmental control regulations.

## **Section 14: Transport Information**

**DOT Classification:** CLASS 3: Flammable liquid.

Identification: : Toluene UNNA: 1294 PG: II

**Special Provisions for Transport:** Not available.

## **Section 15: Other Regulatory Information**

#### Federal and State Regulations:

California prop. 65: This product contains the following ingredients for which the State of California has found to cause cancer, birth defects or other reproductive harm, which would require a warning under the statute: Toluene California prop. 65 (no significant risk level): Toluene: 7 mg/day (value) California prop. 65 (acceptable daily intake level): Toluene: 7 mg/day (value) California prop. 65: This product contains the following ingredients for which the State of California has found to cause birth defects which would require a warning under the statute: Toluene Connecticut hazardous material survey.: Toluene Illinois

toxic substances disclosure to employee act: Toluene Illinois chemical safety act: Toluene New York release reporting list: Toluene Rhode Island RTK hazardous substances: Toluene Pennsylvania RTK: Toluene Florida: Toluene Minnesota: Toluene Michigan critical material: Toluene Massachusetts RTK: Toluene Massachusetts spill list: Toluene New Jersey: Toluene New Jersey spill list: Toluene Louisiana spill reporting: Toluene California Director's List of Hazardous Substances.: Toluene TSCA 8(b) inventory: Toluene TSCA 8(d) H and S data reporting: Toluene: Effective date: 10/04/82; Sunset Date: 10/0/92 SARA 313 toxic chemical notification and release reporting: Toluene CERCLA: Hazardous substances.: Toluene: 1000 lbs. (453.6 kg)

## Other Regulations:

OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200). EINECS: This product is on the European Inventory of Existing Commercial Chemical Substances.

#### Other Classifications:

#### WHMIS (Canada):

CLASS B-2: Flammable liquid with a flash point lower than 37.8°C (100°F). CLASS D-2A: Material causing other toxic effects (VERY TOXIC).

## DSCL (EEC):

R11- Highly flammable. R20- Harmful by inhalation. S16- Keep away from sources of ignition - No smoking. S25- Avoid contact with eyes. S29- Do not empty into drains. S33- Take precautionary measures against static discharges.

## HMIS (U.S.A.):

Health Hazard: 2 Fire Hazard: 3 Reactivity: 0

Personal Protection: h

## National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 3
Reactivity: 0
Specific hazard:

#### **Protective Equipment:**

Gloves. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Splash goggles.

## **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

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# Material Safety Data Sheet

## Trichloroethylene, stabilized

ACC# 23850

# Section 1 - Chemical Product and Company Identification

MSDS Name: Trichloroethylene, stabilized

Product Grade: SQ, ExcelaR, EL

Catalog Numbers: 28455, 28456, 28457, 14715, 41957

**Synonyms:** Trichloroethylene **Company Identification:** 

Fisher Scientific

Part of Thermo Fisher Scientific

THERMO ELECTRON LLS INDIA PVT.LTD.

Godrej Coliseum, 101A-101B, Somaiya Hospital Road,

Off Eastern Express Highway, Sion (East), Mumbai-400 022, India

For information, call: 022 – 6680 3001/2, Call India Toll Free – 1800 209 7001

Emergency Number: 022-66803004/14

**For CHEMTREC assistance**, call: 800-424-9300 [International]

**For International CHEMTREC assistance**, call: 703-527-3887 [International]

## Section 2 - Composition, Information on Ingredients

CAS#	Chemical Name	Percent	EINECS/ELINCS
79-01-6	Trichloroethylene	>99	201-167-4

## Section 3 - Hazards Identification

## **EMERGENCY OVERVIEW**

Appearance: clear, colorless liquid.

**Warning!** Breathing vapors may cause drowsiness and dizziness. Causes eye and skin irritation. Aspiration hazard if swallowed. Can enter lungs and cause damage. May cause cancer based on animal studies. May cause liver damage.

Target Organs: Central nervous system, liver, eyes, skin.

## **Potential Health Effects**





**Eye:** Causes moderate eye irritation. May result in corneal injury. Contact produces irritation, tearing, and burning pain. Contact with trichloroethylene causes pain but no permanent injury to the eyes. (Doc of TLV)

**Skin:** Causes mild skin irritation. Prolonged and/or repeated contact may cause defatting of the skin and dermatitis. May cause peripheral nervous system function impairment including persistent neuritis, and temporary loss of touch. Damage to the liver and other organs has been observed in workers who have been overexposed. **Ingestion:** May cause irritation of the digestive tract. Aspiration of material into the lungs may cause chemical pneumonitis, which may be fatal.

**Inhalation:** May cause respiratory tract irritation. May cause liver abnormalities. May cause cardiac abnormalities. May cause peripheral nervous system effects. Inhalation overexposure may lead to central nervous system depression, producing effects such as dizziness, headache, confusion, incoordination, nausea, weakness, and loss of consciousness. Extreme exposures may cause other CNS effects including death. The chief symptoms of TCE exposure were found to be abnormal fatigue, irritability, headache, gastric disturbances, and intolerance to alcohol. (Doc to TLV)

**Chronic:** Possible cancer hazard based on tests with laboratory animals. Chronic inhalation may cause effects similar to those of acute inhalation. Prolonged or repeated skin contact may cause defatting and dermatitis. May cause peripheral nervous system function impairment including persistent neuritis, and temporary loss of touch. Damage to the liver and other organs has been observed in workers who have been overexposed.

## Section 4 - First Aid Measures

**Eyes:** Immediately flush eyes with plenty of water for at least 15 minutes, occasionally lifting the upper and lower eyelids. Get medical aid imme diately.

**Skin:** Get medical aid if irritation develops or persists. Flush skin with plenty of soap and water.

**Ingestion:** If victim is conscious and alert, give 2-4 cupfuls of milk or water. Never give anything by mouth to an unconscious person. Possible aspiration hazard. Get medical aid immediately.

**Inhalation:** Get medical aid immediately. Remove from exposure and move to fresh air immediately. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Do NOT use mouth-to-mouth resuscitation.

**Notes to Physician:** Treat symptomatically and supportively.

# Section 5 - Fire Fighting Measures

**General Information:** As in any fire, wear a self-contained breathing apparatus in pressure-demand, MSHA/NIOSH (approved or equivalent), and full protective gear. During a fire, irritating and highly toxic gases may be generated by thermal decomposition or combustion. Use water spray to keep fire-exposed containers cool.





**Extinguishing Media:** Use extinguishing media most appropriate for the surrounding

fire.

Flash Point: None

**Autoignition Temperature:** 420 deg C ( 788.00 deg F)

**Explosion Limits, Lower:**8

**Upper:** 10.5

NFPA Rating: (estimated) Health: 2; Flammability: 1; Instability: 0

## Section 6 - Accidental Release Measures

**General Information:** Use proper personal protective equipment as indicated in Section 8.

**Spills/Leaks:** Absorb spill with inert material (e.g. vermiculite, sand or earth), then place in suitable container. Provide ventilation. Approach spill from upwind. Control runoff and isolate discharged material for proper disposal.

# Section 7 - Handling and Storage

**Handling:** Wash thoroughly after handling. Remove contaminated clothing and wash before reuse. Use with adequate ventilation. Avoid contact with eyes, skin, and clothing. Avoid breathing vapor.

**Storage:** Store in a tightly closed container. Store in a cool, dry, well-ventilated area away from incompatible substances.

# Section 8 - Exposure Controls, Personal Protection

**Engineering Controls:** Facilities storing or utilizing this material should be equipped with an eyewash facility and a safety shower. Use adequate general or local exhaust ventilation to keep airborne concentrations below the permissible exposure limits. **Exposure Limits** 

Chemical Name	ACGIH	NIOSH	OSHA - Final PELs
Trichloroethylene	50 ppm TWA; 100 ppm STEL	1000 ppm IDLH	100 ppm TWA; 200 ppm Ceiling

OSHA Vacated PELs: Trichloroethylene: 50 ppm TWA; 270 mg/m3 TWA





Personal Protective Equipment

Eyes: Wear chemical splash goggles.

**Skin:** Wear appropriate protective gloves to prevent skin exposure. **Clothing:** Wear appropriate protective clothing to prevent skin exposure.

**Respirators:** Follow the OSHA respirator regulations found in 29 CFR 1910.134 or

European Standard EN 149. Use a NIOSH/MSHA or European Standard EN 149 approved respirator if exposure limits are exceeded or if irritation or other symptoms are

experienced.

# Section 9 - Physical and Chemical Properties

Physical State: Liquid

**Appearance:** clear, colorless

**Odor:** chloroform-like **pH:** Not available.

Vapor Pressure: 58 mm Hg @ 20 deg C

Vapor Density: 4.5 (air=1) Evaporation Rate:0.69 (CCl4=1)

**Viscosity:** 0.0055 poise **Boiling Point:** 87 deg C

Freezing/Melting Point:-86 deg C

**Decomposition Temperature:** Not available.

Solubility: Slightly soluble. Specific Gravity/Density:1.46 Molecular Formula:C2HCl3 Molecular Weight:131.39

# Section 10 - Stability and Reactivity

**Chemical Stability:** Stable under normal temperatures and pressures.

Conditions to Avoid: Light, confined spaces.

**Incompatibilities with Other Materials:** Active metals.

Hazardous Decomposition Products: Hydrogen chloride, phosgene, carbon

monoxide, carbon dioxide.

Hazardous Polymerization: May occur.

# Section 11 - Toxicological Information





RTECS#:

**CAS#** 79-01-6: KX4550000

LD50/LC50: CAS# 79-01-6:

> Draize test, rabbit, eye: 20 mg/24H Moderate; Draize test, rabbit, skin: 2 mg/24H Severe; Inhalation, mouse: LC50 = 8450 ppm/4H;

Inhalation, mouse: LC50 = 220000 mg/m3/20M; Inhalation, mouse: LC50 = 262000 mg/m3/30M; Inhalation, mouse: LC50 = 40000 mg/m3/4H; Inhalation, rat: LC50 = 140700 mg/m3/1H;

Oral, mouse: LD50 = 2402 mg/kg; Oral, mouse: LD50 = 2400 mg/kg; Oral, rat: LD50 = 4920 mg/kg; Skin, rabbit: LD50 = >20 gm/kg; Skin, rabbit: LD50 = 20 mL/kg;

Carcinogenicity:

CAS# 79-01-6:

**ACGIH:** Not listed.

• California: carcinogen, initial date 4/1/88

 NTP: Suspect carcinogen • **IARC:** Group 2A carcinogen

**Epidemiology:** In six epidemiological studies completed, there was no evidence to suggest that trichloroethylene has increased the incidence of cancer in humans. (Documentation of the TLV, 7th edition)

**Teratogenicity:** No information available.

**Reproductive Effects:** Experimental reproductive effects have been observed. Mutagenicity: Human mutation data has been reported. IARC and the National Toxicology Program (NTP) stated that variability in the mutagencity test results with trichloroethylene may be due to the presence of various stabilizers used in TCEwhich are mutagens (e.g.epoxybutane, epichlorohydrin). See actual entry in RTECS for complete infomation.R68 Mutagen Category 3 (CHIP 2002, UK).

**Neurotoxicity:** No information available.

Other Studies:

## Section 12 - Ecological Information

Ecotoxicity: Fish: Fathead Minnow: 41-67 mg/L; 96 hrs.; LC50Daphnia: Daphnia: 2.2-100 mg/L; 48 hrs.; LC50Mollusk Shrimp: 2 mg/L; 96 hrs.; LC50 Bluegill sunfish, LD50= 44,700 ug/L/96Hr. Fathead minnow, LC50=40.7 mg/L/96Hr.

**Environmental:** In air, substance is photooxidized and is reported to form phosgene, dichloroacetyl chloride, and formyl chloride. In water, it evaporates rapidly. Potential for





mobility in soil is high.

Physical: No information available.

Other: Bioconcentration potential is low (BCF less than 100).

# Section 13 - Disposal Considerations

Chemical waste generators must determine whether a discarded chemical is classified as a hazardous waste. US EPA guidelines for the classification determination are listed in 40 CFR Parts 261.3. Additionally, waste generators must consult state and local hazardous waste regulations to ensure complete and accurate classification.

RCRA P-Series: None listed.

**RCRA U-Series:** 

CAS# 79-01-6: waste number U228.

# Section 14 - Transport Information

	US DOT	Canada TDG		
<b>Shipping Name:</b>	TRICHLOROETHYLENE	TRICHLOROETHYLENE		
Hazard Class:	6.1	6.1		
UN Number:	UN1710	UN1710		
Packing Group:	III	III		

# Section 15 - Regulatory Information

#### **US FEDERAL**

#### **TSCA**

CAS# 79-01-6 is listed on the TSCA inventory.

#### **Health & Safety Reporting List**

None of the chemicals are on the Health & Safety Reporting List.

#### **Chemical Test Rules**

None of the chemicals in this product are under a Chemical Test Rule.

#### Section 12b

None of the chemicals are listed under TSCA Section 12b.

#### **TSCA Significant New Use Rule**

None of the chemicals in this material have a SNUR under TSCA.

#### **CERCLA Hazardous Substances and corresponding RQs**





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CAS# 79-01-6: 100 lb final RQ; 45.4 kg final RQ

#### SARA Section 302 Extremely Hazardous Substances

None of the chemicals in this product have a TPO.

#### **SARA Codes**

CAS # 79-01-6: immediate, delayed, reactive.

#### Section 313

This material contains Trichloroethylene (CAS# 79-01-6, >99%), which is subject to the reporting requirements of Section 313 of SARA Title III and 40 CFR

#### Clean Air Act:

CAS# 79-01-6 is listed as a hazardous air pollutant (HAP).

This material does not contain any Class 1 Ozone depletors.

This material does not contain any Class 2 Ozone depletors.

#### **Clean Water Act:**

CAS# 79-01-6 is listed as a Hazardous Substance under the CWA. CAS# 79-01-6 is listed as a Priority Pollutant under the Clean Water Act. CAS# 79-01-6 is listed as a Toxic Pollutant under the Clean Water Act.

#### OSHA:

None of the chemicals in this product are considered highly hazardous by OSHA.

#### STATE

CAS# 79-01-6 can be found on the following state right to know lists: California, New Jersey, Pennsylvania, Minnesota, Massachusetts.

#### California Prop 65

#### The following statement(s) is(are) made in order to comply with the California Safe Drinking Water Act:

WARNING: This product contains Trichloroethylene, a chemical known to the state of California to cause cancer.

California No Significant Risk Level: CAS# 79-01-6: 50 æg/day NSRL (oral); 80 æg/day NSRL (inhalation)

#### **European/International Regulations European Labeling in Accordance with EC Directives Hazard Symbols:**

#### **Risk Phrases:**

R 36/38 Irritating to eyes and skin.

R 45 May cause cancer.

R 52/53 Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

R 67 Vapours may cause drowsiness and dizziness.

#### Safety Phrases:

S 45 In case of accident or if you feel unwell, seek medical advice immediately (show the label where possible).

S 53 Avoid exposure - obtain special instructions before use.

S 61 Avoid release to the environment. Refer to special instructions /safety data sheets.

#### WGK (Water Danger/Protection)

CAS# 79-01-6: 3





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#### Canada - DSL/NDSL

CAS# 79-01-6 is listed on Canada's DSL List.

#### Canada - WHMIS

This product has a WHMIS classification of D1B, D2B.

This product has been classified in accordance with the hazard criteria of the Controlled Products Regulations and the MSDS contains all of the information required by those regulations.

#### **Canadian Ingredient Disclosure List**

CAS# 79-01-6 is listed on the Canadian Ingredient Disclosure List.

### Section 16 - Additional Information

MSDS Creation Date: 2/01/1999 **Revision #7 Date:** 12/27/2006

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Off Eastern Express Highway,

Sion (E), Mumbai - 400 022,

INDIA



# MATERIAL SAFETY DATA SHEET

Section 1 - Chemical Product and Company Identification

**MSDS Name:** 1,2,4-Trimethylbenzene

Catalog Numbers: AC140090000, AC140090010, AC140090025, AC140095000

**Synonyms:** Pseudocumene.

Company Identification: Acros Organics BVBA

Janssen Pharmaceuticalaan 3a

2440 Geel, Belgium

Company Identification: (USA) Acros Organics

One Reagent Lane Fair Lawn, NJ 07410

For information in the US, call:

For information in Europe, call:

Emergency Number, Europe:

Emergency Number US:

CHEMTREC Phone Number, US:

CHEMTREC Phone Number, Europe:

800-ACROS-01

+32 14 57 52 11

+32 14 57 52 99

201-796-7100

800-424-9300

CHEMTREC Phone Number, US:

703-527-3887

Section 2 - Composition, Information on Ingredients

CAS#: 95-63-6

Chemical Name: 1,2,4-Trimethylbenzene

%: 98

EINECS#: 202-436-9

Hazard Symbols:

XN N







10 20 36/37/38 51/53

Section 3 - Hazards Identification

#### **EMERGENCY OVERVIEW**

Warning! Flammable liquid and vapor. Harmful if inhaled. Causes eye, skin, and respiratory tract irritation. Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment. Target Organs: Blood, central nervous system, respiratory system, eyes, skin.

#### **Potential Health Effects**

**Eye:** Causes eye irritation. Causes redness and pain.

**Skin:** Causes skin irritation. Causes redness and pain. May be harmful if absorbed through the skin.

Ingestion: May cause irritation of the digestive tract. Aspiration of material into the lungs may cause chemical

pneumonitis, which may be fatal. May be harmful if swallowed. May cause central nervous system

depression.

Inhalation: Harmful if inhaled. Causes respiratory tract irritation. May cause drowsiness, unconsciousness, and central

nervous system depression.

Chronic: Prolonged or repeated skin contact may cause dermatitis. May cause anemia and other blood cell

abnormalities. Prolonged exposure may produce a narcotic effect. Prolonged or repeated exposure may

cause nausea, dizziness, and headache.

#### Section 4 - First Aid Measures

Immediately flush eyes with plenty of water for at least 15 minutes, occasionally lifting the upper and Eyes:

lower eyelids. Get medical aid.

Skin: Get medical aid. Immediately flush skin with plenty of water for at least 15 minutes while removing

contaminated clothing and shoes.

Do not induce vomiting. Possible aspiration hazard. Get medical aid immediately. Call a poison control Ingestion:

Inhalation: Get medical aid immediately. Remove from exposure and move to fresh air immediately. If breathing is

difficult, give oxygen. Possible aspiration hazard. Do not use mouth-to-mouth resuscitation if victim ingested or inhaled the substance; induce artificial respiration with the aid of a pocket mask equipped with

a one-way valve or other proper respiratory medical device.

Notes to Physician:

#### Section 5 - Fire Fighting Measures

General As in any fire, wear a self-contained breathing apparatus in pressure-demand, MSHA/NIOSH

(approved or equivalent), and full protective gear. Vapors may form an explosive mixture with air. Information:

Vapors can travel to a source of ignition and flash back. Will burn if involved in a fire. Containers may

explode in the heat of a fire. Flammable liquid and vapor.

**Extinguishing** Use water spray to cool fire-exposed containers. Use water spray, dry chemical, carbon dioxide, or

Media: chemical foam.

Autoignition 500 deg C (932.00 deg F)

Temperature:

Flash Point: 48 deg C (118.40 deg F)

Explosion 0.9 vol %

Limits: Lower:

Explosion 6.4 vol %

Limits: Upper:

Spills/Leaks:

NFPA Rating: health: 2; flammability: 2; instability: 0;

#### Section 6 - Accidental Release Measures

General Use proper personal protective equipment as indicated in Section 8.

Information:

Absorb spill with inert material (e.g. vermiculite, sand or earth), then place in suitable container. Wear a self contained breathing apparatus and appropriate personal protection. (See Exposure Controls, Personal Protection section). Remove all sources of ignition. Use a spark-proof tool. Do not let this

chemical enter the environment.

#### Section 7 - Handling and Storage

Handling: Use spark-proof tools and explosion proof equipment. Do not get in eyes, on skin, or on clothing. Do not ingest or inhale. Use only in a chemical fume hood. Keep away from heat, sparks and flame.

**Storage:** Keep away from sources of ignition. Store in a cool, dry place. Store in a tightly closed container. Flammables-area.

Section 8 - Exposure Controls, Personal Protection

Chemical Name	+	+	++
	ACGIH	NIOSH	OSHA - Final PELs
1,2,4-Trimethylbenz ene		  25 ppm TWA; 125  mg/m3 TWA	none listed

OSHA Vacated PELs: 1,2,4-Trimethylbenzene: 25 ppm TWA; 125 mg/m3 TWA (listed under Trimethyl benzene)

#### **Engineering Controls:**

Use explosion-proof ventilation equipment. Facilities storing or utilizing this material should be equipped with an eyewash facility and a safety shower. Use only under a chemical fume hood.

#### **Exposure Limits**

#### **Personal Protective Equipment**

Wear appropriate protective eyeglasses or chemical safety goggles as described by OSHA's eye and face Eyes:

protection regulations in 29 CFR 1910.133 or European Standard EN166.

Skin: Wear appropriate protective gloves to prevent skin exposure. Clothing: Wear appropriate protective clothing to prevent skin exposure.

Respirators: A respiratory protection program that meets OSHA's 29 CFR 1910.134 and ANSI Z88.2 requirements or

European Standard EN 149 must be followed whenever workplace conditions warrant respirator use.

#### Section 9 - Physical and Chemical Properties

Physical State: Clear liquid Color: colorless Odor: aromatic odor

pH: Not available

Vapor Pressure: 7 mm Hg @ 44.4 deg C

Vapor Density: 4.15 (air=1) **Evaporation Rate:** Not available Viscosity: Not available

**Boiling Point:** 168 deg C @ 760 mmHg ( 334.40°F)

Freezing/Melting Point: -44 deg C (-47.20°F)

**Decomposition Temperature:** Not available Solubility in water: Insoluble Specific Gravity/Density: 0.880 g/cm3 Molecular Formula: C9H12

Molecular Weight: 120.19

#### Section 10 - Stability and Reactivity

**Chemical Stability:** Stable under normal temperatures and pressures. **Conditions to Avoid:** Incompatible materials, ignition sources, excess heat.

**Incompatibilities with Other Materials** Strong oxidizing agents.

**Hazardous Decomposition Products** Carbon monoxide, carbon dioxide.

**Hazardous Polymerization** Will not occur.

#### **Section 11 - Toxicological Information**

RTECS#: CAS# 95-63-6: DC3325000

LD50/LC50: RTECS:

**CAS# 95-63-6:** Inhalation, rat: LC50 = 18000 mg/m3/4H;

Oral, mouse: LD50 = 6900 mg/kg; Oral, rat: LD50 = 5 gm/kg;

Carcinogenicity: 1,2,4-Trimethylbenzene - Not listed as a carcinogen by ACGIH, IARC, NTP, or CA Prop 65.

Other: See actual entry in RTECS for complete information.

**Section 12 - Ecological Information** 

Fish: Fathead Minnow: LC50 = 77.2 mg/L; 96 Hr; Flow-through at 25 C (pH 7.24) **Ecotoxicity:** 

Other: Do not empty into drains.

Section 13 - Disposal Considerations

Dispose of in a manner consistent with federal, state, and local regulations.

**Section 14 - Transport Information** 

**US DOT** 

Shipping Name: FLAMMABLE LIQUIDS, N.O.S. (1,2,4-Trimethylbenzene)

Hazard Class: 3 UN Number: UN1993 Packing Group: III Canada TDG

Shipping Name: Not available

Hazard Class: UN Number: Packing Group:

#### Section 15 - Regulatory Information

#### **European/International Regulations**

European Labeling in Accordance with EC Directives

Hazard Symbols: XN N

Risk Phrases:

R 10 Flammable.

R 20 Harmful by inhalation.

R 36/37/38 Irritating to eyes, respiratory system and skin.

R 51/53 Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

#### Safety Phrases:

S 26 In case of contact with eyes, rinse immediately with plenty of water and seek medical advice.

S 61 Avoid release to the environment. Refer to special instructions/safety data sheets.

WGK (Water Danger/Protection)

CAS# 95-63-6: 3

#### Canada

CAS# 95-63-6 is listed on Canada's DSL List Canadian WHMIS Classifications: B3, D1B, D2B

This product has been classified in accordance with the hazard criteria of the Controlled Products Regulations and the MSDS contains all of the information required by those regulations.

CAS# 95-63-6 is listed on Canada's Ingredient Disclosure List

#### **US Federal**

**TSCA** 

CAS# 95-63-6 is listed on the TSCA Inventory.

#### Section 16 - Other Information

**MSDS Creation Date:** 5/19/1999 **Revision #5 Date** 8/30/2007

**Revisions were made in Sections:** 3, 4, 5, 6, 7, 8, 9, 10, 11, 1

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# Material Safety Data Sheet Xylenes MSDS

#### **Section 1: Chemical Product and Company Identification**

**Product Name:** Xylenes

Catalog Codes: SLX1075, SLX1129, SLX1042, SLX1096

**CAS#:** 1330-20-7

RTECS: ZE2100000

TSCA: TSCA 8(b) inventory: Xylenes

CI#: Not available.

**Synonym:** Xylenes; Dimethylbenzene; xylol;

methyltoluene

Chemical Name: Xylenes (o-, m-, p- isomers)

Chemical Formula: C6H4(CH3)2

#### **Contact Information:**

Sciencelab.com, Inc. 14025 Smith Rd. Houston, Texas 77396

US Sales: 1-800-901-7247

International Sales: 1-281-441-4400
Order Online: ScienceLab.com

CHEMTREC (24HR Emergency Telephone), call:

1-800-424-9300

International CHEMTREC, call: 1-703-527-3887

For non-emergency assistance, call: 1-281-441-4400

# **Section 2: Composition and Information on Ingredients**

#### Composition:

Name	CAS#	% by Weight
Xylenes	1330-20-7	100

**Toxicological Data on Ingredients:** Xylenes: ORAL (LD50): Acute: 4300 mg/kg [Rat]. 2119 mg/kg [Mouse]. DERMAL (LD50): Acute: >1700 mg/kg [Rabbit].

#### Section 3: Hazards Identification

Potential Acute Health Effects: Hazardous in case of skin contact (irritant, permeator), of eye contact (irritant), of ingestion, of inhalation.

#### **Potential Chronic Health Effects:**

CARCINOGENIC EFFECTS: 3 (Not classifiable for human.) by IARC. MUTAGENIC EFFECTS: Not available. TERATOGENIC EFFECTS: Not available. DEVELOPMENTAL TOXICITY: Not available. The substance may be toxic to blood, kidneys, liver, mucous membranes, bone marrow, central nervous system (CNS). Repeated or prolonged exposure to the substance can produce target organs damage.

#### **Section 4: First Aid Measures**

#### **Eye Contact:**

Check for and remove any contact lenses. In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Get medical attention.

#### **Skin Contact:**

In case of contact, immediately flush skin with plenty of water. Cover the irritated skin with an emollient. Remove contaminated clothing and shoes. Wash clothing before reuse. Thoroughly clean shoes before reuse. Get medical attention.

#### Serious Skin Contact:

Wash with a disinfectant soap and cover the contaminated skin with an anti-bacterial cream. Seek immediate medical attention.

#### Inhalation:

If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical attention if symptoms appear.

#### Serious Inhalation:

Evacuate the victim to a safe area as soon as possible. Loosen tight clothing such as a collar, tie, belt or waistband. If breathing is difficult, administer oxygen. If the victim is not breathing, perform mouth-to-mouth resuscitation. Seek medical attention.

#### Ingestion:

Do NOT induce vomiting unless directed to do so by medical personnel. Never give anything by mouth to an unconscious person. Loosen tight clothing such as a collar, tie, belt or waistband. Get medical attention if symptoms appear.

Serious Ingestion: Not available.

#### **Section 5: Fire and Explosion Data**

Flammability of the Product: Flammable.

**Auto-Ignition Temperature:** 464°C (867.2°F)

Flash Points: CLOSED CUP: 24°C (75.2°F). (Tagliabue.) OPEN CUP: 37.8°C (100°F).

Flammable Limits: LOWER: 1% UPPER: 7%

Products of Combustion: These products are carbon oxides (CO, CO2).

#### Fire Hazards in Presence of Various Substances:

Highly flammable in presence of open flames and sparks, of heat. Non-flammable in presence of shocks.

#### **Explosion Hazards in Presence of Various Substances:**

Risks of explosion of the product in presence of mechanical impact: Not available. Slightly explosive in presence of open flames and sparks, of heat.

#### **Fire Fighting Media and Instructions:**

Flammable liquid, soluble or dispersed in water. SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use alcohol foam, water spray or fog. Cool containing vessels with water jet in order to prevent pressure build-up, autoignition or explosion.

Special Remarks on Fire Hazards: Vapors may travel to source of ignition and flash back.

#### **Special Remarks on Explosion Hazards:**

Vapors may form explosive mixtures with air. Containers may explode when heated. May polymerize explosively when heated. An attempt to chlorinate xylene with 1,3-Dichloro-5,5-dimethyl-2,4-imidazolidindione (dichlorohydrantoin) caused a violent explosion

#### Section 6: Accidental Release Measures

Small Spill: Absorb with an inert material and put the spilled material in an appropriate waste disposal.

#### Large Spill:

Flammable liquid. Keep away from heat. Keep away from sources of ignition. Stop leak if without risk. Absorb with DRY earth, sand or other non-combustible material. Do not touch spilled material. Prevent entry into sewers, basements or confined

areas; dike if needed. Be careful that the product is not present at a concentration level above TLV. Check TLV on the MSDS and with local authorities.

# **Section 7: Handling and Storage**

#### Precautions:

Keep away from heat. Keep away from sources of ignition. Ground all equipment containing material. Do not ingest. Do not breathe gas/fumes/ vapor/spray. Wear suitable protective clothing. In case of insufficient ventilation, wear suitable respiratory equipment. If ingested, seek medical advice immediately and show the container or the label. Avoid contact with skin and eyes. Keep away from incompatibles such as oxidizing agents, acids.

#### Storage:

Store in a segregated and approved area. Keep container in a cool, well-ventilated area. Keep container tightly closed and sealed until ready for use. Avoid all possible sources of ignition (spark or flame).

#### **Section 8: Exposure Controls/Personal Protection**

#### **Engineering Controls:**

Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value. Ensure that eyewash stations and safety showers are proximal to the work-station location.

#### **Personal Protection:**

Splash goggles. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Gloves.

#### Personal Protection in Case of a Large Spill:

Splash goggles. Full suit. Vapor respirator. Boots. Gloves. A self contained breathing apparatus should be used to avoid inhalation of the product. Suggested protective clothing might not be sufficient; consult a specialist BEFORE handling this product.

#### **Exposure Limits:**

TWA: 100 (ppm) [Canada] TWA: 435 (mg/m3) [Canada] TWA: 434 STEL: 651 (mg/m3) from ACGIH (TLV) [United States] TWA: 100 STEL: 150 (ppm) from ACGIH (TLV) [United States] Consult local authorities for acceptable exposure limits.

#### **Section 9: Physical and Chemical Properties**

Physical state and appearance: Liquid.

Odor: Sweetish.

Taste: Not available.

Molecular Weight: 106.17 g/mole

Color: Colorless. Clear

pH (1% soln/water): Not available.

Boiling Point: 138.5°C (281.3°F)

Melting Point: -47.4°C (-53.3°F)

Critical Temperature: Not available.

Specific Gravity: 0.864 (Water = 1)

Vapor Pressure: 0.9 kPa (@ 20°C)

Vapor Density: 3.7 (Air = 1)

Volatility: Not available.

Odor Threshold: 1 ppm

p. 3

Water/Oil Dist. Coeff.: The product is more soluble in oil; log(oil/water) = 3.1

Ionicity (in Water): Not available.Dispersion Properties: Not available.

Solubility:

Insoluble in cold water, hot water. Miscible with absolute alcohol, ether, and many other organic liquids.

#### Section 10: Stability and Reactivity Data

Stability: The product is stable.

Instability Temperature: Not available.

Conditions of Instability: Heat, ignition sources, incompatibles

Incompatibility with various substances: Reactive with oxidizing agents, acids.

Corrosivity: Non-corrosive in presence of glass.

Special Remarks on Reactivity: Store away from acetic acid, nitric acid, chlorine, bromine, and fluorine.

Special Remarks on Corrosivity: Not available.

Polymerization: Will not occur.

#### **Section 11: Toxicological Information**

Routes of Entry: Absorbed through skin. Dermal contact. Eye contact. Inhalation.

#### **Toxicity to Animals:**

WARNING: THE LC50 VALUES HEREUNDER ARE ESTIMATED ON THE BASIS OF A 4-HOUR EXPOSURE. Acute oral toxicity (LD50): >1700 mg/kg [Rabbit]. Acute toxicity of the vapor (LC50): 5000 4 hours [Rat].

#### **Chronic Effects on Humans:**

CARCINOGENIC EFFECTS: 3 (Not classifiable for human.) by IARC. May cause damage to the following organs: blood, kidneys, liver, mucous membranes, bone marrow, central nervous system (CNS).

Other Toxic Effects on Humans: Hazardous in case of skin contact (irritant, permeator), of ingestion, of inhalation.

#### **Special Remarks on Toxicity to Animals:**

Lowest Lethal Dose: LDL [Human] - Route: Oral; Dose: 50 mg/kg LCL [Man] - Route: Oral; Dose: 10000 ppm/6H

#### **Special Remarks on Chronic Effects on Humans:**

Detected in maternal milk in human. Passes through the placental barrier in animal. Embryotoxic and/or foetotoxic in animal. May cause adverse reproductive effects (male and femael fertility (spontaneous abortion and fetotoxicity)) and birth defects based animal data.

#### **Special Remarks on other Toxic Effects on Humans:**

Acute Potential Health Effects: Skin: Causes skin irritation. Can be absorbed through skin. Eyes: Causes eye irritation. Inhalation: Vapor causes respiratory tract and mucous membrane irritation. May affect central nervous system and behavior (General anesthetic/CNS depressant with effects including headache, weakness, memory loss, irritability, dizziness, giddiness, loss of coordination and judgement, respiratory depression/arrest or difficulty breathing, loss of appetite, nausea, vomiting, shivering, and possible coma and death). May also affects blood, sense organs, liver, and peripheral nerves. Ingestion: May cause gastrointestinal irritation including abdominal pain, vomiting, and nausea. May also affect liver and urinary system/kidneys. May cause effects similar to those of acute inhalation. Chronic Potential Health Effects: Chronic inhalation may affect the urinary system (kidneys) blood (anemia), bone marrow (hyperplasia of bone marrow) brain/behavior/Central Nervous system. Chronic inhalation may alsocause mucosal bleeding. Chronic ingestion may affect the liver and metabolism (loss of appetite) and may affect urinary system (kidney damage)

#### Section 12: Ecological Information

**Ecotoxicity:** Not available.

BOD5 and COD: Not available.

#### **Products of Biodegradation:**

Possibly hazardous short term degradation products are not likely. However, long term degradation products may arise.

Toxicity of the Products of Biodegradation: The products of degradation are less toxic than the product itself.

Special Remarks on the Products of Biodegradation: Not available.

#### **Section 13: Disposal Considerations**

#### Waste Disposal:

Waste must be disposed of in accordance with federal, state and local environmental control regulations.

#### **Section 14: Transport Information**

**DOT Classification:** CLASS 3: Flammable liquid. **Identification:** : Xylenes UNNA: 1307 PG: III

Special Provisions for Transport: Not available.

#### **Section 15: Other Regulatory Information**

#### **Federal and State Regulations:**

Connecticut hazardous material survey.: Xylenes Illinois chemical safety act: Xylenes New York acutely hazardous substances: Xylenes Rhode Island RTK hazardous substances: Xylenes Pennsylvania RTK: Xylenes Minnesota: Xylenes Michigan critical material: Xylenes Massachusetts RTK: Xylenes Massachusetts spill list: Xylenes New Jersey: Xylenes New Jersey spill list: Xylenes Louisiana spill reporting: Xylenes California Director's List of Hazardous Substances: Xylenes TSCA 8(b) inventory: Xylenes SARA 302/304/311/312 hazardous chemicals: Xylenes SARA 313 toxic chemical notification and release reporting: Xylenes CERCLA: Hazardous substances.: Xylenes: 100 lbs. (45.36 kg)

#### Other Regulations:

OSHA: Hazardous by definition of Hazard Communication Standard (29 CFR 1910.1200). EINECS: This product is on the European Inventory of Existing Commercial Chemical Substances.

#### Other Classifications:

#### WHMIS (Canada):

CLASS B-2: Flammable liquid with a flash point lower than 37.8°C (100°F). CLASS D-2A: Material causing other toxic effects (VERY TOXIC).

#### DSCL (EEC):

R10- Flammable. R21- Harmful in contact with skin. R36/38- Irritating to eyes and skin. S2- Keep out of the reach of children. S36/37- Wear suitable protective clothing and gloves. S46- If swallowed, seek medical advice immediately and show this container or label.

#### HMIS (U.S.A.):

**Health Hazard: 2** 

Fire Hazard: 3

Reactivity: 0

Personal Protection: h

National Fire Protection Association (U.S.A.):

Health: 2

Flammability: 3
Reactivity: 0

Specific hazard:

#### **Protective Equipment:**

Gloves. Lab coat. Vapor respirator. Be sure to use an approved/certified respirator or equivalent. Wear appropriate respirator when ventilation is inadequate. Splash goggles.

#### **Section 16: Other Information**

References: Not available.

Other Special Considerations: Not available.

Created: 10/11/2005 12:54 PM

Last Updated: 11/06/2008 12:00 PM

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# **APPENDIX C**

Job Safety Analysis

JOB SAFETY ANALYSIS	Ctrl. No. generic	DATE: 9/14	/2015	☐ NEW ☑ REVISED	PAGE 1 of 2
JSA TYPE CATEGORY <b>Generic</b>	WORK TYPE: Drilling		WORK ACTIVIT	Y (Description): Point Installation k	ov Hand
DEVELOPMENT TEAM	POSITION / TITLE			WED BY:	POSITION / TITLE
Jeffrey Wills	Project Hydrogeologist		Daniel Abberton		Site Health and Safety Mgr.
Chris Migliorie	Driller (ADT)		Dennis Mayer		Operations Mgr. (ADT)
	JIRED AND / OR RECOM	MENDED P		ECTIVE EQUIPMENT	· · · · · · · · · · · · · · · · · · ·
☐ LIFE VEST ☐ HARD HAT ☐ LIFELINE / BODY HARNESS ☐ SAFETY GLASSES	☐ GOGGLES ☐ FACE SHIELD: (while air knifing and jack hammering) ☐ HEARING PROTECTION: (as needed) ☐ SAFETY SHOES steel or composite toe		AIR PURIFY SUPPLIED R PPE CLOTH sleeve shirt of reflective saf	ING RESPIRATOR RESPIRATOR ING: fluorescent long or long sleeve shirt and ety vest.	GLOVES: Leather. Nitrile and Cut Resistant ANSI Level 2 OTHER: Insect Repellant, sunscreen (as needed)
Vac-Truck or Vac Drum, Jack Ham		, Photoioniza	ation Detector, M	ulti Gas Meter, 42 inch	
Fire Extinguisher, "Work Area" Sign					
EXCLUSION ZONE POLICY: All n			OUR HANDS"	e during any invasive w	OFK
Driller and	اه I helper should show t			controls and movi	ng parts
Assess	Analyze			Act	
<sup>1</sup> JOB STEPS	<sup>2</sup> POTENTIAL HAZAR	DS		3CRITICAL ACT	IONS
Verify pre-clearance protocol and Subsurface Clearance Procedure Checklist	CONTACT: Undergroutility damage; prope damage; personal inj	rty	drilling; hav  1a. Walk the Si  Site Walk I  1a. Review pre location mu ground surf	re DigSafe number avai ite to evaluate utility ma Inspection JSA). Inclearing checklist form Inst be conducted to a magace using hand tools all Iling soil vapor point dee	s were contacted prior to lable at time of work. rkings and review maps (see  . Pre-clearing of the drilling inimum of 5 vertical feet below and/or an air knife/vacuum, eper than 5 feet below ground
Mobilize/demobilize and establish Exclusion Zone	FALL: tripping/falling uneven terrain, weath conditions, and materials/equipment the Site  2b. CONTACT: with traff (including any uninter movement of the wor Contact / Interference Other Site Activities)	ner stored at ic nded k truck),	hazards (i.e. mobilizing et amobilizing et amobilizing et al. Do not clim Practice go 2a. Use establi 2b. When first a space and/vehicles an 2b. Check in with other set al. Use a spott operations; 2b. Maintain 1 personnel at 2b. Use a spott operations; 2b. Set-up the eliminates of 2b. Delineate wother barries 2b. Position lar where apple 2b. Face traffic spotter, and 2b. Chock whee	equipment.  b over stored materials, and housekeeping.  shed pathways and wale arriving onsite park vehice or out of the way location of the way l	letc.), and obstructions prior to dequipment; walk around.  Ilk on stable, secure ground.  Il
Assess ¹JOB STEPS	Analyze	ns		Act	IONS
JUD STEPS	2POTENTIAL HAZAR  2c. EXERTION: during n     equipment (cones an     signage) into work ar	noving of d	never reach	n with a load. t loads are balanced to	eep load close to body, and reduce the potential for

- <sup>1</sup> Each Job or Operation consists of a set of tasks / steps. Be sure to list all the steps needed to perform job.
- A hazard is a potential danger. Break hazards into six types: Contact victim is struck by or strikes an object; Caught - victim is caught on, caught in or caught between objects; Fall - victim falls to ground or lower level (includes slips and trips); Exertion - excessive strain or stress / ergonomics / lifting techniques; Exposure - inhalation/skin hazards; Energy Source - electricity, pressure, compression/tension.
- <sup>3</sup> Using the first two columns as a guide, decide what actions or procedures are necessary to eliminate or minimize the risk. List the recommended safe operating procedures. Say exactly what needs to be done such as "use two persons to lift". Avoid general statements such as, "be careful".

	2d. <b>EXPOSURE:</b> To biological hazards: ticks, bees/wasps, poison ivy, insects, etc. (ticks are most active any time the temperature is above freezing from March to November)	<ul> <li>2d. Inspect area to avoid contact with biological hazards.</li> <li>2d. Wear long sleeved clothing (mandatory PPE) to protect skin and apply insect repellant containing DEET when working in overgrown areas of the Site.</li> <li>2d. Personnel shall examine themselves for ticks.</li> <li>2d. If skin comes in contact with poison ivy, wash skin thoroughly with soap and water as soon as possible.</li> </ul>
	Exposure:     Sun possibly causing sunburn or cold possibly causing cold stress	<ul> <li>2e. Wear sunscreen with an SPF of at least 15 whenever 30 minutes or more of exposure is expected.</li> <li>2e. Wear weather appropriate attire</li> <li>2e. Take frequent heat breaks to avoid cold stress.</li> <li>2e. Drink fluids to remain hydrated.</li> </ul>
	2f. <b>EXPOSURE:</b> Noise hazards (certain sites operate heavy machinery)	Wear hearing protection if necessary.     When possible, position body away from noise origins to reduce exposure.
Concrete saw cutting, jack hammer and hand clearance (air knife) (review completed Subsurface Clearance Form; Review Geophysical Survey notes)	3a. See Clearing, Vactron and Air Knife JSAs	3a. See Clearing, Vactron and Air Knife JSA.
Soil vapor point installation	4a. CAUGHT: Pinch points associated with equipment and installation	<ul> <li>4a. Always wear leather gloves when making connections and installing sample screen/tubing; wear cut-proof (i.e., Kevlar) gloves when handling cutting tools (no fix blade knives).</li> <li>4a. Inspect the equipment prior to use for potential pinch points.</li> <li>4a. Inspect all hand tools for damage and wear prior to use. Remove any damaged tool from service and replace.</li> </ul>
	4b. EXERTION: Muscle strain	4b. See 2c.
Move drum to staging area using drum cart	5a. EXPOSURE/CONTACT: Contamination (e.g., Separate Phase Hydrocarbons (SPH), contaminated groundwater, soil)	<ul> <li>5a. Do not overfill drums. Wear Nitrile gloves and long sleeved shirts to avoid contact with skin.</li> <li>5a. Dispose of used impacted materials/PPE in designated containers.</li> </ul>
	5b. <b>EXERTION:</b> Muscle strain	5b. See 2c.
	5c. CAUGHT: Pinch points associated with handling drum lid	5c. Ensure that fingers are not placed under the lid of the drum. Wear leather or cut-proof gloves while sealing drum lid.
	5d. <b>FALL</b> : Slipping on spilled materials	5d. Clean up any spills using absorbent pads.
6. Decontaminate equipment	6a. EXPOSURE/CONTACT: Contamination (e.g., Separate Phase Hydrocarbons (SPH), contaminated groundwater, vapors)	<ul><li>6a. Wear chemical-resistant disposable gloves and safety glasses.</li><li>6a. See 5a.</li></ul>
	6b. <b>EXPOSURE:</b> Chemicals in cleaning solution including ammonia	6b. See 5a. 6b. See 2a.

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A hazard is a potential danger. Break hazards into six types: Contact - victim is struck by or strikes an object;
 Caught - victim is caught on, caught in or caught between objects; Fall - victim falls to ground or lower level (includes slips and trips); Exertion - excessive strain or stress / ergonomics / lifting techniques; Exposure - inhalation/skin hazards; Energy Source - electricity, pressure, compression/tension.

<sup>3</sup> Using the first two columns as a guide, decide what actions or procedures are necessary to eliminate or minimize the risk. List the recommended safe operating procedures. Say exactly what needs to be done - such as "use two persons to lift". Avoid general statements such as, "be careful".

JOB SAFETY				□ NEW	
ANALYSIS			/2015	REVISED	PAGE 1 of 2
JSA TYPE CATEGORY:	WORK TYPE:			TY (Description):	
Generic	Drilling				Well Installation
DEVELOPMENT TEAM	POSITION / TITL		REVIEW		POSITION / TITLE
Jeffrey Wills	Project Hydrogeologis		Ray Fitzpatric	k	OHSO
Thalassa Sodre	Staff Assistant Engine	er			
DE(	QUIRED AND / OR RECO	MANGENDED D	EDSONAL DOO	FECTIVE EQUIDME	NIT
☐ LIFE VEST	GOGGLES	MINIENDED		ING RESPIRATOR	☐ GLOVES: <u>Leather, Nitrile and cut</u>
	☐ FACE SHIELD		☐ SUPPLIED I	RESPIRATOR	<u>resistant</u>
☐ LIFELINE / BODY HARNESS ☐ SAFETY GLASSES	HEARING PROTECTION (as needed)	ON:		HING: Fluorescent st or high visibility	
M SALETT GLAGGES	SAFETY SHOES: Cor	mposite-toe or	clothing, Lor	ng Sleeve Shirt	Sunscieen (as needed)
	steel toe boots	'D / OD DECC	THE PER EQU	UDATAIT	
Gooprobe or Truck-Mounted Direc			MMENDED EQU		r (or equivalent), Macrocore liners,
Liner Opening Tool, 42" Cones & F		JIS, FIIUIUIUI 112	.allon Detector an	U/OI IVIUILI-GAS IVICIO	(Of equivalent), Macrocore inters,
Exclusion Zone Policy – All non-		aintain a dista	nce of 10' feet fro	m drilling equipmen	t while moving/engaged.
	"0	SHOW ME Y	OUR HANDS"		
Driller an	nd helper should show			n controls and m	oving parts
Assess	Analyze			Act	
<sup>1</sup> JOB STEPS	<sup>2</sup> POTENTIAL HAZAI			3CRITICAL A	
Mobilization of drilling rig	1a. <b>CONTACT</b> :			tower/derrick will be	lowered and secured prior to
(ensure the Subsurface Clearance Protocol and Drill	Equipment/proper damage.		mobilization.	auld he utilized while	moving the drill rig. If personnel
Rig Checklist are completed)	uamay <del>e</del> .	[ ]			the drill rig will be stopped until the
rug onosimet are compress,			path is again	clear. Use a spotter	for all required backing operations.
		[ 1			n equipment in a manner that
				reduces the need for	or backing of support trucks and
			trailers. 1a When backin	o up truck rig with a	n attached trailer use a second
					simultaneously on multiple sides of
			the equipmer	nt or if turning angles	s limit driver visibility.
					en terrain. Level or avoid if needed.
		[ ]			exclusion zone of 10 feet for non- elper, geologist) when the rig is
			moving/ in op		elper, geologist) when the hy is
	1	1			errain, weather-related hazards (i.e.,
	1b. FALL:	_l_	ice, puddles, equipment.	snow, etc.), and obs	structions prior to mobilizing
	Slip/trip/fall hazard		- 1 - 1	over stored materia	s/equipment; walk around. Practice
			good housek	eeping.	
		1			alk on stable, secure ground.
Raising tower/derrick of drill	2a. CONTACT:	<del></del>	20 Prior to raisir	as the towar/derrick	the area above the drilling rig will be
rig	Overhead hazards		inspected for	wires. tree limbs, pi	ping, or other structures, that could
1.9			come in conta	act with the rig's tow	er and/or drilling rods or tools.
		2	2a. Maintain a sa	afe distance from ove	erhead structures.
	Ch CONTACT.		OL Incorpor the e		avaid ningh nainta
	2b. CONTACT: Pinch Points when				e and avoid pinch points. e stability prior to raising rig
	the rig and instabil	0	tower/derrick		s stability prior to raising ng
	]		2b. If the rig need	ds to be mounted, be	e sure to use three points of contact.
Advancement of drilling	3a. CONTACT:	- ;	3a. Be aware of a	and avoid potential I	ines of fire and wear required PPE
equipment and well	Flying debris			ear, and hand prote	
installation					

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Advancement of drilling	3b.	EXPOSURE:		Wet borehole area with sprayer to minimize dust.
equipment and well installation (Continued)		Noise and dust.		Stand upwind and keep body away from rig.  Dust mask should be worn if conditions warrant.
motanation (Continued)				Wear hearing protection when the drill rig is in operation.
	3c.	CAUGHT: Limb/extremity pinching; abrasion/crushing.	3c. 3c. 3c. 3c. 3c.	Always wear leather gloves when making connections and using hand tools; wear cut-resistant (i.e., Kevlar) gloves when handling cutting tools.  Inspect the equipment prior to use for potential pinch points. Keep hands away from being between pinch points and use of tools is preferable compared to fingers and hands.  Inspect drill head for worn surface or missing teeth; replace if damaged or blunt.  Ensure all jewelry is removed, loose clothing is secured, and PPE is secured close to the body.  All non-essential personnel should stay away from the immediate work area; position body out of the line-of-fire of equipment.  Drillers and helpers will understand and use the "Show Me Your Hands" Policy.  Spinning rods/casing have an exclusion zone of 10 feet while in operation.
	3d.	CONTACT: Equipment imbalance during advancement of drill equipment.	3d.	Drillers will advance the borehole with caution to avoid causing the rig to become imbalanced and/or tip.  The blocking and leveling devices used to secure the rig will be inspected by drillers and Roux personnel regularly to see if shifting has occurred.  In addition, personnel and equipment that are non-essential to the advancement of the borehole will be positioned away from the rig at a distance that is at least as far as the boom is high (minimum exclusion zone of 10 feet).
	3e.	EXPOSURE: Inhalation of contamination/vapors.		Air monitoring using a calibrated photoionization detector (PID) will be used to periodically to monitor the breathing zone of the work area. If a reading of >5ppm is recorded, the Roux field personnel must temporarily cease work, instruct all Site personnel to step away from the area of elevated readings and inform the Roux PM of the condition. The Roux PM will then recommend additional precautions in accordance with the site specific health and safety plan.
	3f.	FALL: Slip/trip/fall hazards.		Contain drill cuttings and drilling water to prevent fall hazards from developing in work area. See 1b.
	3g.	EXERTION: Potential for muscle strain/injury while lifting and installing well casings, lifting sand bags, and/or lifting rods.	3g.	Keep back straight and bend at the knees. Utilize team lifting for objects over 50lbs. Use mechanical lifting device for odd shaped objects.
Decontaminate equipment.	4a.	EXPOSURE/CONTACT: To contamination (e.g., Separate Phase Hydrocarbons (SPH), contaminated groundwater, vapors).	4a. 4a.	Wear chemical-resistant disposable gloves and safety glasses. Contain decontamination water so that it does not spill. Use an absorbent pad to clean spills, if necessary. See 3b.
	4b.	<b>EXPOSURE:</b> To chemicals in cleaning solution including ammonia.	4b.	See 4a. Review MSDS to ensure appropriate precautions are taken and understood.

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JOB SAFETY ANALYSIS Ctrl. No. GEN-007 DATE: 1/5/2015				5		☐ NEW ☑ REVISED	F	PAGE 1 of 1	
	TYPE CATEGORY	WORK TYPE			WORK ACTIVITY (Description)				
GEI	GENERIC		Drilling		Мо	Movement of 55-gallon Drums/Drum Handling			
The	DEVELOPMENT TEAM	POSITION / TITLE		1-44	REVIEWED BY:  Jeff Wills		Desia	POSITION / TITLE	
ina	assa Sodre	Starr	Assistant Engineer		Jett	VVIIIS		Proje	ect Manager
	R	EQUIR	ED AND / OR RECOM	MENDED PERSO	NAL P	ROTECTIVE E	EQUIPMENT		
	LIFE VEST HARD HAT LIFELINE / BODY HARNESS SAFETY GLASSES		GOGGLES FACE SHIELD HEARING PROTECT SAFETY SHOES: <u>Ste</u>	ION		AIR PURIFY SUPPLIED PPE CLOTH reflective ve clothing	/ING RESPIRATOR RESPIRATOR HING: <u>Fluorescent</u> st or high visibility		GLOVES: <u>Cut-resistant</u> gloves OTHER:
Rea	uired Equipment: Drum Cart and/	or fork							
	LUSION ZONE: A 10' exclusion								
	Assess		Analyze				Act		
	<sup>1</sup> JOB STEPS		<sup>2</sup> POTENTIAL HAZA	ARDS			3CRITICAL AC	CTION	S
1.	Secure Work Area, Inspect 55- gal drums for proper condition, labeling, check drum ring and bolts.	rea, Inspect 55- roper condition, Tripping/falling due to uneven		<ul> <li>1a. Inspect walking path for uneven terrain, weather-related hazards (i.e., tree debris, puddles, etc.), and obstructions prior to accessing work area.</li> <li>1a. Use established pathways and walk on stable, secure ground.</li> <li>1a. Secure work area and coordinate and communicate the planned work activities with other personnel working in the area.</li> <li>1a.</li> </ul>			c.), and obstructions prior on stable, secure ground. d communicate the sonnel working in the		
				1b. 1b.	<ul> <li>1b. When inspecting drums, don nitrile gloves under cut resistant glove. If drum is not properly labeled, do not open and cease all drum transport related activities. Immediately contact project manager and inform him/her of drum situation.</li> <li>1b. Do not continue drum transport activities until further actions are determined by the project manager.</li> <li>1b. If the drum is properly labeled, but leaking, improperly sealed, or in poor condition, place drum in an over-pack drum.</li> <li>1c. Keep back straight and secure grip on drum ratchet.</li> </ul>				
2.	When using a forklift, position drum clamp in between drum ribs. When using a drum dolly, secure fastening hook on top of drum.	2a. 2b.	Hazards between drum/forklift clamp or dolly fastener/drum.		2b.	<ul> <li>2a. Position drum clamp between the ribs on the drum to preve possible slipping. Do not place hands between drum clam and drum; wear cut resistant gloves.</li> <li>2b. Do not jerk body. Wear cut-resistant gloves and steel toed boots.</li> <li>2b. Ensure that drums are not over-filled.</li> </ul>		s between drum clamp	
3.	Transport drums to designated location and disengage drum clamp.	3a. EXPOSURE/ CONTACT: Hazards associated with drum transport; skin contact and vapors.  3b. CAUGHT:		<ul> <li>3a. Maintain a 10' EZ around forklift. Ensure drum clamp is secure on drum before beginning to move.</li> <li>3a. Ensure that drum is sealed and lid is tight before beginning to move.</li> </ul>			move.		
		3c.	Pinching hazards a maneuvering drum: FALL: Tripping/ falling due obstructions and ur	s. e to	3c. 3c.	See 2b. If path is too		art, utili	

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JOB SAFETY ANALYSIS	Ctrl. No. GEN-006	DATE 1/15/	/2015	☐ NEW ☐ REVISED	PAGE 1 of 2			
JSA TYPE CATEGORY	WORK TYPE		WORK ACTIVITY					
Generic	Surveying		Elevation Su	urveying				
DEVELOPMENT TEAM	POSITION / TITLE		REVIEW	ED BY:	POSITION / TITLE			
Bjorn Wespestad	Project Engineer		Jeff Wills		Project Manager			
Thalassa Sodre	Staff Assistant Engineer							
	REQUIRED AND / OR RECOM	MENDED PER	SONAL PROTECTI	VE EQUIPMENT				
☐ LIFE VEST ☑ HARD HAT ☐ LIFELINE / BODY HARNESS ☑ SAFETY GLASSES	GOGGLES     FACE SHIELD     HEARING PROTECTION     SAFETY SHOES: Steel-t		SUPPLIED F  PPE CLOTH	ING RESPIRATOR RESPIRATOR ING: <u>Fluorescent</u> st or high visibility	GLOVES: Cut-resistant or leather OTHER			
	REQUIRED AND / OR RECOMMENDED EQUIPMENT							
Surveying equipment (i.e., leveling	rod/measuring ruler, tripod an	d scope).						

, , , , ,		Analyza	Act			
Assess 1JOB STEPS		Analyze  2POTENTIAL HAZARDS	3CRITICAL ACTIONS			
Locate surveying position for instrument and rod and set-up work area	1a.	FALL: Slip/trip hazards.	Inspect area for ice, puddles, sno at the survey loc Be aware of onco	uneven terrain, weather-related hazards (i.e., ow, etc.), and obstructions prior to setting up eation.  oming traffic. Utilize a flagman / spotter for		
	CONTACT:     Traffic (surveying locations could potentially be located in parking areas and sidewalks).		Place 42 inch co work zone with c Wear appropriate reflective safety	ntain eye contact with oncoming vehicles,		
	1c.	OVEREXERTION: Hazard due to carrying, lifting, and bending while transporting equipment.	straight, lift with leach with a load Avoid carrying to	y positioning and lifting techniques; keep back legs, keep load close to body, and never d. to much equipment at one time and team-lift is more than 50lb.		
	1d.	CAUGHT/CONTACT: Pinch Points / sharp edges associated with setting up the tripod.	carry tripod by th			
Open / close manhole cover to well that is being surveyed (if necessary).	2a.	OVEREXERTION: Muscle strain		ees when reaching to open well. Use ook or pry bar to avoid bending.		
ourroyou (ii nooossary).	2b.	CAUGHT: Pinch points associated with removing / replacing manholes and working with hand tools.	well cover and h	(ratchet and crowbar or pry bar for well ect before use.		
	2c.	<b>EXPOSURE:</b> To potentially hazardous vapors.		osure to vapors allow well to vent after efore survey activities begin.		
	2d.	CONTACT: With traffic.	See 1b.			

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<sup>&</sup>lt;sup>3</sup> Using the first two columns as a guide, decide what actions or procedures are necessary to eliminate or minimize the risk. List the recommended safe operating procedures. Say exactly what needs to be done - such as "use two persons to lift". Avoid general statements such as, "be careful".

	Assess 1JOB STEPS		Analyze  2POTENTIAL HAZARDS		Act  3CRITICAL ACTIONS
3.	Perform survey.	3a.	FALL: Slip/trip hazards	3a.	See 1a.
		3b.	CONTACT: Traffic (surveying locations could be potentially located in parking areas and sidewalks)	3b. 3b.	See 1b.  Personnel using the scope will be devoting most of their attention to the surveying activity. Personnel holding the measuring stick should be extra vigilant of survey personnel and communicate any potential hazards to the instrument person via handheld radio or similar means. Ensure reflective safety vest is worn.
4.	Break down work area.	4a.	CONTACT: Traffic (surveying locations can potentially be located in parking areas and sidewalks).	4a.	See 1b.
		4b.	EXERTION: Hazard due to carrying, lifting, and bending while transporting equipment	4b.	See 1c.

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JOE	B SAFETY ANALYSIS	Ctrl. No. GEN-005	DATE 1	/5/2015	☐ NEW ☐ REVISED	PAGE 1 of 2		
	TYPE CATEGORY Neric	WORK TYPE:		WORK ACTIVITY (I				
		Gauging and Sampling		Gauging and S		DOCITION / TITLE		
	VELOPMENT TEAM	POSITION / TITLE		REVIEWED	BY:	POSITION / TITLE		
	a Masciello	Project Scientist		Jeff Wills		Project Manager		
rna	lassa Sodre	Staff Assistant Engineer						
		DECLURED AND LOD DECOME	AENIDED DEI	DOONAL BROTEOT	NE FOLUDATA	-		
	LIFE VEST	REQUIRED AND / OR RECOMN GOGGLES	IENDED PER	AIR PURIFYING		I ☐ GLOVES: Leather, Nitrile	and cut	
$\boxtimes$	HARD HAT	FACE SHIELD		☐ SUPPLIED RES		resistant		
	LIFELINE / BODY HARNESS	☐ HEARING PROTECTION				OTHER: Knee pads, Inse		
$\boxtimes$	SAFETY GLASSES	SAFETY SHOES: Composite toe boots	-toe or steel	reflective vest or clothing	high visibility	Repellant, sunscreen (as	needed)	
			OR RECOM	MENDED EQUIPME	NT			
42 ir	nch Safety Cones, Caution T	ape, Interface Probe and/or Water				ools as needed: Socket Wre	nch,	
	w Driver, Crow Bar, Mallet, a			,	,		ŕ	
	Assess	Analyze			Act			
	<sup>1</sup> JOB STEPS	<sup>2</sup> POTENTIAL HAZARDS			3CRITICAL A			
	Mobilization to monitoring	1a. <b>FALL:</b> Personal injury fro				ost suitable designated pathy	vay	
	well(s).	slip/trip/fall due to uneven	terrain	prior to mobiliz		condition on atable cons		
		and/or obstructions.			oid steep hills or	c and/or drive on stable, secu uneven terrain	ie,	
				ground and av	old oldop fillio Of	anovon tonam.		
		1b. CONTACT: With traffic/th	nird	1b. Identify potenti	al traffic sources	and delineate work area with	42 inch	
		parties.				ehicle to protect against onco		
						vide a more visible delineation	n of the	
				work area if ne		g high visibility clothing or ref	octivo	
				vest.	ate i i L ilicidulii	g riigir visibility clottiling of ren	CCIIVE	
					aintain eye conta	act with oncoming vehicles, ar	nd	
				establish a sat	e exit route.	-		
				A. Lancettoni				
		1c. EXPOSURE:		<ol> <li>Inspect work a</li> <li>Use insect/tick</li> </ol>	rea for bees and			
		To biological hazards.		ic. Ose insect/tick	repellent as neo	cssary.		
2.	Open/close well.	2a. OVEREXERTION: Muscl	e strain.	2a. Use proper lift	ing techniques; k	eep back straight, lift with leg	s and	
	•				hen reaching to c			
		Oh OAHOUT Bissh assists		Oh Maarlaathar		stant alougo when working wi	المبيطة	
		2b. CAUGHT: Pinch points a with removing/replacing m		<ol> <li>Wear leather gloves or cut resistant gloves when working with well cover and hand tools.</li> </ol>				
		and working with hand too				ry bar for well cover) and insp	ect	
		and noming mannand to		before use.		,		
				2b. Do not put fing	jers under well co	over.		
		On EVENCUES. To make the		2c No open flame	oc/boot cources			
		<ol> <li>EXPOSURE: To potential hazardous vapors.</li> </ol>	'	<ul><li>2c. No open flames/heat sources.</li><li>2c. To minimize exposure to vapors allow well to vent a</li></ul>			ina it	
		nazardous vapors.			mpling activities b		mig it	
				2c. Stand up-wind	, if possible, to a	void vapors.		
3.	Gauge well.	3a. <b>CONTACT:</b> With contar			l-resistant dispos	able gloves and safety glasse	es when	
		(e.g. contaminated ground	iwater).	gauging well. 3a. Insert and rem	ove probe slowly	to avoid eplaching		
					ent pad to clean			
		3b. CONTACT:			pau 10 0.0u	p. 656.		
		With traffic.		3b. See 1b.				
4.	Purge and sample well.	4a. EXPOSURE/CONTACT:			ample jars slowly	to avoid splashing and conta	act with	
		contamination (e.g., SPH, contaminated groundwate		preservatives.	tant aloves and a	hemical-resistant disposable	alovos	
	and/or sample preservatives.			when sampling		anomioai-resistant disposable	gioves	
					ge container to avoid spilling v	water		
				onto the groun	ıd.	, ,		
				4a. Use an absorb	ent pad to clean	spills.		
	Assess	Analyze			Act			
	1JOB STEPS	<sup>2</sup> POTENTIAL HAZARDS			3CRITICAL A			

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4.	Purge and sample well (Continued).	4b.	<b>CONTACT:</b> Personal injury from cuts, abrasions, or punctures by glassware or sharp objects.	4b. 4b. 4b.	To avoid spills or breakage, place sample ware on even surface. Do not over tighten caps on glass sample ware.  Wear cut-resistant (i.e., Kevlar) gloves and chemical-resistant disposable gloves when sampling and handling glassware (i.e., VOA vials) or when using cutting tools.
		4c.	<b>EXERTION:</b> Muscle strain while carrying equipment.	4c. 4c. 4c.	Use proper lifting techniques when handling/moving equipment; bend knees and keep back straight. Use mechanical assistance or team lifting techniques when equipment is 50lbs or heavier. Make multiple trips to carry equipment.
		4d.	CONTACT: With traffic.	4d.	See 1b.
5.	Management of purge water.	5a.	<b>EXPOSURE/CONTACT:</b> To contamination (e.g., SPH, contaminated groundwater, vapors).	5a. 5a.	Do not overfill container and pour liquids in such a manner that they do not splash.  Properly dispose of used materials/PPE in appropriate container in designated storage area.
		5b.	<b>EXERTION:</b> Muscle strain from lifting/carrying and moving containers.	5b. 5b.	Use proper lifting techniques when lifting / carrying or moving container(s) (see 4c.).  Do not overfill container(s).
6.	Decontaminate equipment.	6a.	<b>EXPOSURE/CONTACT:</b> To contamination (e.g., SPH, contaminated groundwater, vapors).	6a.	Work on the upwind side, where possible, of decon area. Wear chemical-resistant disposable gloves and safety glasses. Use an absorbent pad to clean spills.

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		Г						
			☐ NEW					
JSA TYPE CATEGORY:	WORK TYPE:			WORK ACTIVITY (Description):				
Site Specific	Drilling		Hollow Stem Auger Soil Borings /Well Installation					
Site: Morton Village	DOOLTION / TITLE		DE\#E\#ED E	DOCUTION (TITLE				
DEVELOPMENT TEAM	POSITION / TITLE		REVIEWED E	8Y:	POSITION / TITLE			
Dennis Mayer Gina Vanderlin	Operations Manager		Jeff Wills		Project Manager			
Gina vandenin	Project Scientist							
REC	UIRED AND / OR RECOMM	IENDED P	ERSONAL PROTECTIV	/F FOLLIPMENT				
☐ LIFE VEST	GOGGLES	ILINDED I	☐ AIR PURIFYING		☐ GLOVES: Leather, Nitrile			
☐ HARD HAT	☐ FACE SHIELD		☐ SUPPLIED RESP	PIRATOR	and cut resistant			
☐ LIFELINE / BODY HARNESS		TION:	PPE CLOTHING: flu		OTHER: Insect Repellant,			
	(as needed)  ☐ SAFETY SHOES <u>ste</u>	ما مه	sleeve shirt or long s reflective safety vest		sunscreen (as needed)			
	composite toe	ei oi	ionidente dalet, red	<u></u>				
	REQUIRED AND /		MMENDED EQUIPMEN		<u> </u>			
Truck-Mounted Drilling Rig or Track		oionization	Detector, Multi-Gas Met	er (or equivalent)	), Interface Probe, 20 lb. Fire			
Extinguisher, 42" Cones & Flags, "W	Vork Area" Signs POLICY – All non-essential pe		all maintain a 20 fact of		ile deill ein in nannand			
EXCLUSION ZONE P			OUR HANDS"	ciusion zone wn	ille drill rig is engaged			
Driller an	nd helper should show th			rols and movi	ng parts			
Assess	Analyze	rat mamae		Act	g parto			
<sup>1</sup> JOB STEPS	<sup>2</sup> POTENTIAL HAZARD	S		3CRITICAL AC				
Mobilization / demobilization	1a. See Mobilization/		1a. See Mobilization/	Demobilization J	JSA			
and establish a work area	Demobilization							
Raising tower/derrick of	2a. CONTACT: Overhead	hazards.	2a. Prior to raising the	e tower/derrick, a	area above the drilling rig will be			
drilling rig					vires, tree limbs, piping, or other			
					by the rig's tower or drilling rods.			
					ng a tower/derrick in the area of			
			overhead wires m 2a. The tower/derrick		eed beneath overhead power lines			
			unless approved b	by both the Roux	CPM.			
			2a. Maintain at a mini 2a. Do not move the i		om overhead structures.			
			Za. Do not move the i	ig wrille the towe	er/deffick is faised.			
	2b. CONTACT: Amputation				and avoid amputation points.			
	points when raising the	rig and			stability prior to raising rig tower			
	instability of rig.		riggers	t and body out of	f the line of fire when lowering out-			
				be mounted, be	sure to use three points of contact.			
				,	·			
	0.0017407.51: /							
<ol><li>Advancement of augers for soil borings, steel casing</li></ol>	<ol> <li>CONTACT: Flying / sp debris.</li> </ol>	orayıng			nd, eye, ear protection). e when operating to avoid lines of			
modification, and well	debiis.		fire of possible fly		e when operating to avoid lines of			
material installation.			ĺ	J				
	OF EXPOSURE Notice	and allower			minimize dust. Stand upwind and			
	3b. <b>EXPOSURE</b> : Noise ar	nd aust.	keep body away f		rig is operating/or the noise levels			
			exceed 85dBA.	COUCH WITHE UITH	ing is operating or the hoise levels			
	Ì							

<sup>&</sup>lt;sup>1</sup> Each Job or Operation consists of a set of tasks / steps. Be sure to list all the steps needed to perform job.

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<sup>&</sup>lt;sup>3</sup> Using the first two columns as a guide, decide what actions or procedures are necessary to eliminate or minimize the risk. List the recommended safe operating procedures. Say exactly what needs to be done - such as "use two persons to lift". Avoid general statements such as, "be careful".

Advancement of augers for soil borings, steel casing      The soil borings and wall are soil borings.	4c. <b>CAUGHT</b> : Limb/extremity amputation; abrasion/crushing.	4c. Always wear leather gloves when making connections and using hand tools; wear cut-resistant (i.e., Kevlar) gloves when handling authors tools.
modification, and well material installation (Continued).		cutting tools.  4c. Inspect the equipment prior to use for potential pinch points.  4c. Test all emergency shutdown devices prior to drilling.  4c. Inspect drill head for worn surface or missing teeth; replace if
		damaged or blunt.  4c. Inspect augers, do not use if auger flight if damaged or bent.  4c. Ensure all jewelry is removed, loose clothing is secured, and PPE is
		secured close to the body.  4c. All non-essential personnel should stay away from the immediate work area; position body out of the line-of-fire of equipment particularly when installing auger flights and steel override casings.
		Drillers and helpers will understand and use the "Show Me Your Hands" Policy.
		Spinning augers should have an exclusion zone of 20 feet when in operation.
	4d. <b>CONTACT</b> : Equipment imbalance during advancement	4d. Drillers will advance the borehole with caution to avoid causing the rig to become imbalanced and/or tip.
	of drill equipment, sparks/heat generated during steel casing modification with chop saw, and	The blocking and leveling devices used to secure the rig will be inspected by drillers and Roux personnel regularly to see if shifting has occurred.
	installing steel override casing.	4d. In addition, personnel and equipment that are non-essential to the advancement of the borehole will be positioned away from the rig at a distance that is at least as far as the boom is high (minimum oxclusion zone of 20 foot)
		exclusion zone of 20 feet).  4d. When lifting and lowering steel override casing into borehole, personnel that are non-essential will be positioned away from direct line of fire incase casing falls over or swings during advancement.
		4d. Flame retardant clothing must be worn while sparks are generated.
	4e. <b>EXPOSURE</b> : Inhalation of contamination/vapors.	<ul> <li>4e. Air monitoring using a calibrated photoionization detector (PID) will be used to periodically monitor the breathing zone of the work area.</li> <li>4e. The Action Level for breathing zone air is five parts per million</li> </ul>
		<ul> <li>(sustained) as detected by the PID.</li> <li>4e. If a reading of &gt;5ppm is recorded, the Roux field personnel must temporarily cease work, instruct all Site personnel to step away from the area of elevated readings and inform the Roux PM of the condition. The Roux PM will then recommend additional appropriate precautions in accordance with the site specific health and safety</li> </ul>
	4f. <b>FALL</b> : Slip/trip/fall hazards.	plan.
		<ul><li>4f. See 1b.</li><li>4f. Remove soil cuttings to avoid a tripping hazard from developing near augers.</li></ul>
	4g. <b>EXERTION</b> : Installing well casings and lifting augers.	<ul> <li>4g. Keep back straight and bend at the knees.</li> <li>4g. Utilize team lifting for objects over 50lbs.</li> <li>4g. Use mechanical lifting device for odd shaped objects.</li> </ul>
5. Cleaning the auger flights	5a. CONTACT:	5a. Follow "No Hands" Procedure and make sure auger is out of gear
	Cuts/scrapes or puncture wound from hand tool contacting auger.	before contacting auger with tool or hand.  5a. Pull cleaning tool across your body with handle away from body; do not push toward the auger.
		5a. Do not clean more than ¾ turn around the auger at a time. 5a. Wear cut resistant and leather gloves.
		<ul><li>5a. Always use two hands to operate cleaning tool.</li><li>5a. Inspect tool before use and remove from service if handle or metal are cracked/fatigued.</li></ul>
6 Decenteminate acciment	62 EVENCUER/CONTACT:	5a. Stand out of the line of fire.
Decontaminate equipment.	6a. EXPOSURE/CONTACT: To contamination (e.g., contaminated groundwater, vapors).	<ul> <li>5a. Wear chemical-resistant disposable gloves and safety glasses.</li> <li>5a. Contain decontamination water so that it does not spill.</li> <li>5a. Use an absorbent pad to clean spills, if necessary.</li> </ul>
	6b. EXPOSURE:	
	To chemicals in cleaning solution including ammonia	5b. See 5a.

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JOB SAFETY ANALYSIS Cntrl. No. GEN-010 DATE: 1/5/			2015	□NEW ☑REVISED	PAGE 1 of 2		
JSA TYPE CATEGORY  GENERIC	WORK TYPE		WORK ACTIVITY (Description)				
	Site Recon POSITION / TITLE		Mobilization/Demobilization  REVIEWED BY: POSITION / TITLE				
Jared Lefkowitz	Staff Assistant Scientist		Ray Fitzpatrick	IED BY:	OSHO POSITION / TITLE		
Jeff Wills	Project Hydrogeologist		Kay Fitzpatilick		03110		
OCH WINS	1 Toject Trydrogeologist						
F	EQUIRED AND / OR RECOMMEND	DED PERSON	IAL PROTECTIVE	EQUIPMENT			
☐ LIFE VEST ☑ HARD HAT ☐ LIFELINE / BODY HARNESS ☑ SAFETY GLASSES	GOGGLES FACE SHIELD HEARING PROTECTION (a needed) SAFETY SHOES: Steel Too composite toe		PPE CLOTE	TOR D RESPIRATOR	GLOVES: Leather, nitrile, and cut resistant (as needed) OTHER		
	REQUIRED AND / OR	RECOMMEND	DED EQUIPMENT				
Required Equipment:							
EXCLUSION ZONE: A minimumexo	lusion zone of 10' will be mair	ntained arou	und moving equ	ipment (if necessar	y)		
Assess	Analyze			Act	TIONS		
1JOB STEPS	2POTENTIAL HAZARDS		10 Hea 2 nois	3CRITICAL AC			
1. Mobilize/demobilize and establish work area	1a. FALL: Slip/trips/falls from obstructions, uneven terrain, we conditions, heavy loads, and/or housekeeping.  1b. CONTACT: Personal injurproperty damage caused by bistruck by Site traffic or equipment in Site activities.  1c. CAUGHT: Personal injurprinch points and being in line-	reather r poor	and exiting 1a. Inspect wa obstruction and puddle pathways. 1a. Do not clir Practice ge 1a. Wear bood 1a. Delineate flagging. 1b. Observe and vehicles and vehicles and vehicles and the identify po 1b. Wear PPE 1b. Use a spot backing wi 1b. Maintain a motion. Wi second sp multiple si visibility. 1b. Delineate other barri 1b. Position "V either side 1b. Face traffit a spotter, a  1c. Make sure chocks in	g vehicle.  alking path for unever ns, and/or weather-re es) prior to mobilizing Walk on stable/securing on the stored material to do housekeeping. It is with adequate treasunsafe areas with 42' and maintain the poster arriving onsite, park of the very local of the way local of the works are with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area. The work area with 42" colors. Work Area" signs at Sof work area with 42" colors. Work Area" signs at Sof work area with 42" colors. Work Area" signs at Sof work area with 42" colors. Work Area" signs at Sof work area with 42" colors. Work Area" signs at Sof work area with 42" colors. Work Area" signs at Sof work area with 42" colors.	lated hazards (i.e., ice, snow, equipment. Use established erground. als/equipment; walk around. ds. ds. ds. ds. ds. ds. ds. ds. ds. d		
	vehicle and/or equipment.		<ul> <li>is parked in front/down gradient of work area.</li> <li>1c. Wear leather gloves when handling any tools or equipment. Avoid wearing loose clothing. Wear cut-resistant gloves (Kevlor similar) when handling sharp objects/cutting tools.</li> <li>1c. Keep body parts away from line-of-fire of equipment.</li> <li>1c. Always carry tools by the handles and/or designated carrier. Ensure sharp-edged tools are sheathed/secure.</li> <li>1c. Remove any loose jewelry. Ensure loose clothing is secure.</li> </ul>				

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PL-2204		
Assess	Analyze	Act
1JOB STEPS	<sup>2</sup> POTENTIAL HAZARDS	<sup>3</sup> CRITICAL ACTIONS
	1d. OVEREXERTION: Muscle strains while lifting/carrying equipment.	<ul> <li>1d. Use body positioning and lifting techniques that avoid muscle strain; keep back straight, lift with legs, keep load close to body, and never reach with a load.</li> <li>1d. Ensure that loads are balanced. Use assistance (mechanical or additional person) to carry equipment that is either awkward to carry or over 50 lbs.</li> </ul>
	1e. EXPOSURE: Personal injury from exposure to biological and environmental hazards.	<ul> <li>1e. Inspect area to avoid contact with biological hazards (i.e. poisonous plants, stinging insects, ticks, etc.).</li> <li>1e. Wear long sleeved clothes, apply insect repellant containing DEET, and inspect clothes and skin for ticks during and after work.</li> <li>1e. Apply sunscreen (SPF 15+) if exposure to sun for 30 minutes or more is expected.</li> </ul>
	1f. EXPOSURE: Heat and cold related injuries.	<ol> <li>Watch for heat stress symptoms (muscle cramping, exhaustion, dizziness, rapid and shallow breathing). Take breaks as needed.</li> <li>Watch for cold stress symptoms (severe shivering, slowing of body movement, weakness, stumbling or inability to walk, collapse). Take breaks as needed.</li> <li>Wear clothing appropriate for weather and temperature conditions (e.g., rain jackets, snow pants, multiple layers).</li> <li>If lightning is observed, wait 30 minutes in a sheltered location (car is acceptable) before resuming work.</li> </ol>
	<b>1g. EXPOSURE:</b> Personal injury from noise hazards.	1g. Wear hearing protection if sound levels exceed 85 dBA.

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JOB :	JOB SAFETY						☐ NEW		DAGE 4. ( 0
ANALYSIS Cntrl. No. GEN-009 DATE: 1/5/2		2015		□ REVISED		PAGE 1 of 2			
JSA TY	PE CATEGORY	WOF	RK TYPE		WOF	RK ACTIVITY	(Description)		
GEN	ERIC	Ha	nd Tools		Pre	e-Clearin	g activities, i	ncl	luding Air
							Soil Vacuun		
DEV	ELOPMENT TEAM		POSITION / TITLE			REVIEWE			POSITION / TITLE
Jeff Wi	ills	Proje	ect Hydrogeologist		Ray	Fitzpatrick		OF	HSO
Dennis	Mayer	Man	ager (ADT)			-			
	EE \ /EOT		UIRED AND / OR RECOMI	MENDED PER				Is-a	01.01/50.11/5
	FE VEST ARD HAT		GOGGLES FACE SHIELD (while air k	(nifing)		AIR PURIFY		$\boxtimes$	GLOVES: Nitrile and cut resistant
	FELINE / BODY		HEARING PROTECTION				RESPIRATOR		<u>roototarit</u>
	ARNESS	N 7	needed)	_	$\boxtimes$	PPE CLOTI		$\boxtimes$	OTHER: <u>Dust mask (as</u>
⊠ S/	AFETY GLASSES		SAFETY SHOES: Steel of composite toed	<u>or</u>		or high visib	reflective vest		needed)
			REQUIRED AND	OR RECOMM	IENDE	D EQUIPMEN	NT .		
			ctor Truck (Vac Truck), (			d Tools, Phot	oionization Detecto	or, M	lulti-Gas Meter, Traffic
Cones	, 20 lb. Fire Extinguisher,	"Work	Area" and/or "Exclusion	Zone" Signs					
EXCL	JSION ZONE: A 10 foot	exclu	sion zone will be main	tained aroun	d air	knife and/or	r soil vacuum ope	ratio	ons.
	Assess		Analyze				Act		
	JOB STEPS		POTENTIAL HAZARI	os			3CRITICAL A	CTIC	ONS
	rify pre-clearance	1a.	CONTACT:		1a.	Confirm tha			were contacted prior to
pro	tocol.		Underground utility dam			drilling.			
			property damage; perso	nal injury.	1a.				rkings and review maps or critical actions).
		See	Site Walk Inspection JS	A for	1a				and sub-surface clearance
			ntial hazards.	(101	ıu.				es that clearance must be
									cal feet below ground
								gro gro	und surface in the critical
0 14-	b 92 - 7 d b 92 d	0-	See Mobilization / Den	1. 111	0 -	zone using		- 4 7	. 10 4 (
	bilize/demobilize and ablish work area.	∠a.	JSA for potential haza		∠a.	See WODIIIZ	zation / Demobiliz	atio	n JSA for critical actions.
	e-clear with air knife and	3a.	CONTACT:		3a.	Maintain 10	) foot exclusion z	one.	Only (air knife/vac truck)
soi	I vacuum, and/or		Flying debris.						all remain within exclusion
cle	arance with hand tools								tive. Use the required PPE,
							t a minimum), cut r iields, and long slee		tant gloves, safety glasses
					3a.				from flying debris when
						using air kn			, 3
					3a.			and	others, so to avoid line-of-
					0 -	fire hazards	· -		
					за.	Use anti-wn	ip devices on comp	oress	sor noses.
		3b.	EXPOSURE/ENERGY	SOURCE:	3b.	Monitor brea	athing zone with a	calib	orated PID and multi-gas
			Inhalation/exposure to h	nazardous					ppm, the Roux field
			vapors; inhalation/expos	sure to dust;					work, instruct all Site
			electrocution.						rea of elevated readings and of the condition. The Roux
									end additional precautions.
					3b.		nasks as needed.		γ
					3b.		open flames/heat se	ourc	es are present within the
					O.L.	work area.			ded wiente
							truck is properly gr		glass or equivalent.
					00.	Do not doo	motar any bar, acc		grade or equivalent.
			CONTACT:		3c.				rith the high pressure
			Damage to unknown/kno	own			and using the air k	nife	tip as a physical digging
			utility with air knife.		30	tool. Keep the a	ir knife tin constant	tlv m	oving to reduce direct
					50.		n a potential utility.	•	oving to roddoo diroot
					3c.	increase th	e distance betwee		knife tip and
					0 -	soil/utility.		£	a la alla suddh sua assurer
		3d <b>O</b>	VEREXERTION:		JC.				n hole with vacuum, ct on utility casings.
			body positioning when I	nandlina		windiniay	nave an abrasive	OHEC	or on annly casings.
			pment and materials.	3	3d.				ting techniques that
							nuscle strain; keep o body, and never		k straight, lift with legs, keep

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#### PL-2204

0 0 1 1 11 11 11 11	0.1 <b>0</b> 1/ED	EVEDTION ( .: I)	0.1	
Pre-clearing with air knife and soil vacuum, and/or clearance with hand tools (continued)	Poor b	EXERTION: (continued) ody positioning when ng equipment and materials.		Ensure that loads are balanced to reduce the potential for muscle strain.  Two people or a mechanical lifting aid are required when lifting objects over 50 lb. or when the shape makes the object difficult to lift.
	terrain,	g/falling due to uneven , weather conditions, and als/equipment stored at the	3e. 3e. 3e. 3e.	Inspect walking path for uneven terrain, weather-related hazards (e.g., ice, puddles, snow, etc.), and obstructions prior to mobilizing equipment.  Walk around any stored materials/equipment; do not climb over. Practice good housekeeping.  Use established pathways and walk on stable, secure ground. Equipment and tools will be stored at the lowest point of potential energy and out of the walkway and immediate work area (i.e., tools should not be propped against walls or nearby equipment or vehicles).  Equipment and tools that are not anticipated to be used will be returned to a storage area that is out of the immediate work area.  Ensure power cords/hoses are grouped when used within the work area. Mark out cords/hoses that cross pathways with traffic cones.  Ensure all Site personnel and equipment stay a minimum of 2 feet from an open hole. Mark out open holes with traffic cones/caution tape, etc.  Pre-cleared location will be finished flush to grade as to prevent a slip/trip hazard.
		HT: points associated with the nent and vacuum hose.	3f. 3f. 3f. 3f.	Always wear cut-resistant gloves when making connections and using hand tools. Inspect the equipment prior to use for potential pinch points. Test all emergency shutdown devices prior to using equipment. Ensure all jewelry is removed, loose clothing is secured, and PPE is secured close to the body. All non-essential personnel shall maintain a 10 foot exclusion zone; position body out of the line-of-fire of equipment. Drillers and helpers will understand and use the "Show Me Your Hands Policy".
	3g. <b>EXPOS</b> Noise f	from vac truck and/or air	3g.	Wear hearing protection when vac truck and air compressor are in operation. Otherwise, if sound levels exceed 85 dB, don hearing protection.
Move drum to staging area using drum cart.	Contar Phase contar	SURE/CONTACT: mination (e.g., Separate Hydrocarbons (SPH), minated groundwater, soil).	4a.	Wear chemically resistant gloves (i.e., Nitrile; worn in addition to cut resistant gloves).  Do not overfill drums. Ensure that the drum lids are attached securely.  Stage all drums in the designated storage area (per Roux Project Manager) and ensure they are labeled.
		FION: e strain while maneuvering with drum cart/lift gate.	4b.	See 3d. Do not overfill drums. Use lift gate on back of truck to load and unload drums or drum cart to transport drums.
5. Davidson	handlir	points associated withing drum lid.		Ensure that fingers are not placed under the lid of the drum.  Wear cut-resistant gloves. Use 15/16" ratchet while sealing drum lid.
Decontaminate equipment and tools.	To con Phase	SURE/CONTACT:  Itamination (e.g., Separate Hydrocarbons (SPH),  ninated groundwater, ).	5a. 5a.	See 4a. Contain decontamination water (closed lid) so that it does not spill. Use an absorbent pad to clean spills, if necessary. Store all impacted materials/PPE in a designated storage container (per Roux Project Manager) and ensure the container is labeled.
	5b. <b>EXPOS</b> To che	SURE: emicals in cleaning solution.	5b.	See 4a.

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JOB SAFETY ANALYSIS Cntrl. No. GEN-		Cntrl. No. GEN-012	DAT	E: 1/5/20	15	□NEW ⊠REVISED		PAGE 1 of 2
JSA TYPE CATEGORY:		RK TYPE:	D/(I	WORK ACTIVITY (Description):				
GENERIC Gauging & Sampling			Soil Sampling					
DEVELOPMENT TEAM	Juu	POSITION / TITLE			POSITION / TITLE			
Jeff Wills	Proie	ect Hydrogeologist		Ray Fitz	REVIEWED B patrick	•••	OHS	
	, .							
REG	QUIRE	D AND / OR RECOMME	NDED	PERSON	AL PROTEC	TIVE EQUIPME	NT	
☐ LIFE VEST		GOGGLES			PURIFYING RE		$\boxtimes$	GLOVES: Leather, Nitrile and
<ul> <li>☐ HARD HAT</li> <li>☐ LIFELINE / BODY HARNESS</li> <li>☐ SAFETY GLASSES</li> <li>☐ FLAME RESISTANT</li> </ul>		FACE SHIELD: HEARING PROTECTION: <u>(</u>	20		PLIED RESPIR CLOTHING: F		$\boxtimes$	cut resistant OTHER: Insect Repellant,
☐ SAFETY GLASSES		needed)	<u>uo</u>	_	ctive vest or hig			sunscreen (as needed)
		SAFETY SHOES: Composi	te-toe	cloth	ing			
CLOTHING (as needed)		or steel toe boots  REQUIRED AND / O	RRF	COMMENI	DED FOLLIPM	FNT		
Recommended Equipment; 42" t	raffic c			JOININE IN	DED EQUIT IV	I-IV I		
EXCLUSION ZONE: A minir		•		ntained a	ound movin	n equinment if	nres	ant
Assess	ilulii i	Analyze	e iliai	illailleu ai	ourid movin	Act	pres	ant.
JOB STEPS	2 <b>P</b> (	OTENTIAL HAZARDS				3CRITICAL A	CTION	NS
Secure location	1a.	CONTACT:		1a. If in	an area with			delineate the work area with
		Personnel and vehicula	ar					prevent exposure to traffic
		traffic may enter the wo	ork	and	inform others	of work activity		
		area.				est and/or fluore		
					e the direction ker from traffic		ar tranı	ic. Position vehicle to protect
						rk activity with a	ıdiaceı	nt work areas
							.ajacc.	nom arous
	1b.	FALL:						even terrain, weather-related
		Tripping/falling due to	, .,					and obstructions.
		uneven terrain or entry/ from excavations.	exit					stable, secure ground. venient, stable, and orderly
		nom excavations.				ent at lowest po		
								n in-progress excavations
								vation be appropriate (when
				stab	ilization is co	mplete), ladders	must	be employed for steep
				emb	ankments, ex	cavations, pits,	and tr	encnes.
	1c.	EXPOSURE:		1c. Wea	ar sunscreen v	with an SPF 15	or gre	ater whenever 30 minutes or
		Exposure to sun and			e of exposure		J	
		excessive heat, possible	ly					direct sunlight particularly
		causing sunburn, heat	1			eratures are als		
		exhaustion or heat stro	ĸe,			ocation of all Sit		le cramping, exhaustion,
		Exposure to cold				nd shallow brea		
		temperatures possibly		1c. Wat	ch for cold str	ess symptoms	(sever	e shivering, slowing of body
		causing cold stress.		mov	ement, weak	ness, stumbling	or ina	bility to walk, collapse).
				1c. Tak	e breaks for re	est and water as	s nece	ssary. Move to an area that is ning (i.e., car, site trailer,
						area that is war		rining (i.e., car, site trailer,
		Skin burn as a result of	fire		pen flames/he			
		if occurred. Exposure to						en specified by Site policy.
		explosive vapors due to	)					pecified by Site policy.
		tank farm operations,			treat field clot and insects.	ning with Perme	etnrin	prior to site visit to kill/repel
		Biological hazards - ticl	ks.			d shirts and tucl	k in (o	r tape) pant legs into socks or
		bees/wasps, poison ivy		boo	s to prevent t	icks from reachi	ing ski	n.
		thorns, insects, etc.						on exposed skin when
						own areas of the		nical hazarda
						oid contact with t aloves when h		gical nazards.  Ig branches, shrubs, etc. that
						walking path.		5
							ves ar	nd co-worker's outer clothing
						lly when onsite.	an !	wood olde the recent be with
					in comes in c and water.	ontact with pols	on ivy,	wash skin thoroughly with

<sup>&</sup>lt;sup>1</sup> Each Job or Operation consists of a set of tasks / steps. Be sure to list all the steps needed to perform job.

A hazard is a potential danger. Break hazards into five types: Contact - victim is struck by or strikes an object; Caught - victim is caught on, caught in or caught between objects; Fall - victim falls to ground or lower level (includes slips and trips); Exertion - excessive strain or stress / ergonomics / lifting techniques; Exposure - inhalation/skin hazards; Energy Source - electricity, pressure, compression/tension.

<sup>&</sup>lt;sup>3</sup> Using the first two columns as a guide, decide what actions or procedures are necessary to eliminate or minimize the risk. List the recommended safe operating procedures. Say exactly what needs to be done - such as "use two persons to lift". Avoid general statements such as, "be careful".

Assess 1JOB STEPS	Analyze  2POTENTIAL HAZARDS	Act 3CRITICAL ACTIONS
2. Collect Soil Sample	2a. CONTACT:  Personal injury from pinch points, cuts, and abrasions from sampling equipment tools, and material within soil sample.  Personal injury from contact with moving equipment while sampling.	<ul> <li>2a. Wear cut-resistant (i.e., Kevlar) gloves under chemical-resistant disposable gloves when handling soil samples and sampling jars.</li> <li>2a. Where possible, use trowel or equivalent tool to avoid contact with soil.</li> <li>2a. If sampling from bucket of heavy equipment, ensure all equipment is off and operator utilizes the "show me your hands" policy.</li> <li>2a. See 1a.</li> </ul>
	2b. EXPOSURE: Exposure to contamination (impacted soil) and/or lab preservatives.	Wear chemical-resistant disposable gloves over cut resistant gloves to protect hands when handling samples; use containment material or plastic sheeting to protect surrounding areas.      When collecting soil sample from hand auger, put large zip lock bag over entire auger to prevent spillage of soil on to the ground.      Open sample jars slowly and fill carefully to avoid contact with preservatives.
3. Decontaminate equipment	3a. EXPOSURE/CONTACT: Contamination (e.g., Separate Phase Hydrocarbons (SPH), contaminated vapors and/or soil).	<ul> <li>3a. Wear chemical-resistant disposable gloves and safety glasses.</li> <li>3a. Use an absorbent pad to clean spills.</li> <li>3a. Properly dispose of used materials/PPE in provided drums in designated drum storage area.</li> </ul>
	3b. <b>EXPOSURE:</b> Chemicals in cleaning solution including ammonia.	<ul> <li>3b. Wear chemical-resistant disposable gloves and safety glasses.</li> <li>3b. Work on the upwind side of decon area.</li> <li>3b. Use an absorbent pad to clean spills.</li> <li>3b. Properly dispose of used materials/PPE in provided drums in designated drum storage area.</li> </ul>

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Using the first two columns as a guide, decide what actions or procedures are necessary to eliminate or minimize the risk. List the recommended safe operating procedures. Say exactly what needs to be done - such as "use two persons to lift". Avoid general statements such as, "be careful".

JOB SAFETY ANALYSIS Cntrl#: GEN-015 DATE			/15	□NEW ⊠REVISED		PAGE 1 of 2	
JSA TYPE CATEGORY:				otion):			
GENERIC DEVELOPMENT TEAM	Drilling POSITION / TITLE	VV	ell Developmen REVIEWED BY		OSITION / TITLE		
Amy Hoffman	Staff Geologist	.le	ff Wills	•	Project Manager		
7 tilly Florifical	Otali Ocologist	- 00	11 771113		1 TOJOUL	Manager	
	QUIRED AND / OR RECOMM	IENDED PERS					
☐ LIFE VEST ☑ HARD HAT ☐ LIFELINE / BODY HARNESS ☑ SAFETY GLASSES	☐ GOGGLES ☐ FACE SHIELD ☑ HEARING PROTECTION (aneeded) ☑ SAFETY SHOES: Compostoe or steel toe boots  ■ REQUIRED AND /	<u>iite-</u>	AIR PURIFYING RESUPPLIED RESPIRATED RESPIRATED FOR CLOTHING: For Englishing Fig. 1 and 1 a	ATOR uorescent n visibility	cut i ☑ OTH	OVES: <u>Leather, Nitrile and</u> resistant HER: <u>Insect repellant,</u> screen (as needed)	
Required Equipment as needed: T Submercible Pump, Surge Block/P Driver, Pry Bar, Ratchet, Vault Key	ruck Rig or support truck, Traile lunger,  20 lb. Fire Extinguishe	er, 42 inch Saf	ety cones and flags,	Caution Tape			
·	Maintain a 20 Foot Excl			Activities			
		OW ME YOU					
	d helper should show that	t hands are	clear from contro			S	
Assess ¹JOB STEPS	Analyze  2POTENTIAL HAZARDS	S		Act 3CRITICAL A		3	
Mobilization /     Demobilization (Review Mobilization and Demobilization JSA)	<ul> <li>1a. CONTACT: Equipment/property dam</li> <li>1b. FALL: Slip/trip/fall hazards.</li> </ul>	1a. 1a. 1a. 1b.	mobilization.  Set-up the work are eliminates or reduce. All non-essential particles are been born twice by the backing up clearance simultar turning angles limited inspect the driving linspect walking particles, ice, puddles, equipment.	ea / position e ces the need for ersonnel should efore backing with an attach- neously on mult driver visibilit path for uneven show, etc.), ar stored materia	equipment for backing uld mainta up. ed trailer eltiple side ty. Stay aven terrain terrain, wend obstruct	ed and secured prior to t in a manner that g of trucks and trailers. ain an exclusion zone of use a spotter if there is tight as of the equipment or if way from the line-of-fire. b. Level or avoid if needed. reather-related hazards ctions prior to mobilizing ment; walk around. Store	
2. Open/close well.	2a. <b>OVEREXERTION:</b> Muscle strain (some well large vault covers).	lls have	reach with a load. potential for muscl objects over 50 lbs lift.	Ensure that loe strain. Two performs or when the s	pads are t people are shape ma	d close to body, and never calanced to reduce the e required when lifting likes the object difficult to	
	<ol> <li>CAUGHT:         Pinch points associated removing/replacing man and working with hand to     </li> </ol>	with holes 2b.	tools. Do not put fi	ngers under w	ell vault/c	ell vault/cover and hand cover. I inspect before use.	
	2c. <b>EXPOSURE:</b> Potentially hazardous va		activities to minimi	after opening i ze exposure to set up and du	o vapors.	ore starting development Air monitoring must be vell development activities.	
	2d. <b>CONTACT</b> : Traffic.	2d. 2d.	Delineate work are Position vehicle to	ea with 42" saf protect agains naintain eye c	ety cones st oncomi	y clothing or reflective vest. s and/or other barriers. ng traffic. th oncoming vehicles, and	

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	Assess ¹JOB STEPS	Analyze  2POTENTIAL HAZARDS	Act 3CRITICAL ACTIONS
3.	Develop well (mechanical surging).	3a. CAUGHT: Cut hazards and finger pinch points.	<ul> <li>3a See 2b.</li> <li>3a. Use required PPE including leather/cut-resistant gloves when handling development equipment. Identify finger/hand pinch points. Keep hands away from active surge equipment.</li> <li>3a. All non-essential personnel should maintain an exclusion zone of 20 feet.</li> </ul>
		3b. CONTACT/EXPOSURE: Contamination (e.g., SPH, contaminated groundwater, vapors).	<ul> <li>3b. See 2c.</li> <li>3b. Wear Nitrile gloves and safety glasses. Insert and remove surge block/plunger and line/cable slowly to avoid splashing at the surface.</li> <li>3b. Use an absorbent pad to clean any spills.</li> </ul>
		3c. OVEREXERTION: Muscle strain from lifting equipment.	<ul><li>3c. See 2a.</li><li>3c. Use mechanical device to insert and remove surge block/plunger if greater than 50lb.</li></ul>
		3d. CONTACT: Injury while handling wench line/cable, or with active surging equipment	<ul> <li>3d. If using a drill rig, inspect all wench lines/cables for any kinks or if frayed prior to use. Replace any damaged lines/cables. Review Drill Rig checklist prior to development activities.</li> <li>3d. See 3a.</li> </ul>
4.	Purging well (pumping water to holding tanks/drums/buckets).	4a. CAUGHT: Pinch points associated with connecting hose to tank. Pinch points associated with handling pump and hoses.	<ul> <li>4a. See 3a.</li> <li>4a. Ensure that fingers are not placed near coupling when attaching and securing hose(s). Do not place fingers under pump/hoses. Wear leather or cut-resistant gloves when handling pump/hose(s).</li> <li>4a. Keep hands clear from any line of fire.</li> </ul>
		4b. FALL: Using side mounted ladder when attaching hose to tank. Slip, trip, fall from lines/hoses	<ul> <li>4b. Inspect ladder steps make sure steps are not bent/damaged and free of debris/fluid.</li> <li>4b. Use three points of contact at all times when using ladder.</li> <li>4b. Utilize anti-whip cords on all compressed hoses. Keep hoses and lines coiled and organized out of designated walking paths around the work zone.</li> </ul>
		4c. CONTACT: Contamination (e.g., SPH, contaminated groundwater).	<ul> <li>4c. Secure water hose.</li> <li>4c. Do not overfill tanks, and purge/transfer liquids in such a manner that they do not splash. (See 3b).</li> <li>4c. Dispose of used materials/PPE in the designated impacted PPE container.</li> </ul>
		4d. <b>EXERTION:</b> Muscle strain from lifting/carrying equipment.	<ul> <li>4d. Use lifting techniques to minimize muscle strain when carrying equipment. When possible, use mechanic means to lift equipment.</li> <li>4d. Use two people to lift any equipment or material that is over 50 lbs.</li> </ul>
		4e. <b>FALL:</b> Spilled purge water.	4e. Clean up any spills using absorbent pads or spill kits.
5.	Decontaminate equipment	5a. CONTACT/EXPOSURE: Contamination (e.g., SPH, contaminated groundwater, vapors).	5a. See 3b.
		5b. EXPOSURE/CONTACT: Chemicals in cleaning solution	<ol> <li>Decontaminate equipment in well-ventilated area. Wear nitrile gloves to avoid skin contact with cleaning solutions.</li> </ol>

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# **APPENDIX D**

Heat and Cold Stress Guidelines

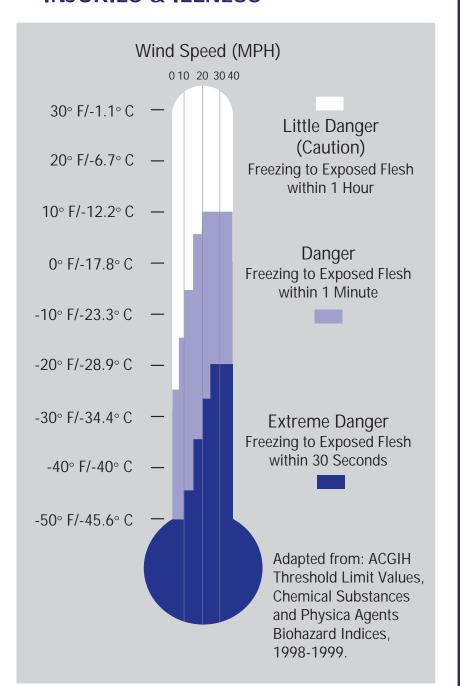
# J.S. Department of Labor Occupational Safety and Health Administration

# THE COLD STRESS EQUATION

# LOW TEMPERATURE + WIND SPEED + WETNESS = INJURIES & ILLNESS

When the body is unable to warm itself, serious cold-related illnesses and injuries may occur, and permanent tissue damage and death may result.

Hypothermia can occur when land temperatures are above freezing or water temperatures are below 98.6°F/ 37°C. Coldrelated illnesses can slowly overcome a person who has been chilled by low temperatures, brisk winds, or wet clothing.



OSHA 3156 1998

# **FROST BITE**

# What Happens to the Body:

FREEZING IN DEEP LAYERS OF SKIN AND TISSUE; PALE, WAXY-WHITE SKIN COLOR; SKIN BECOMES HARD and NUMB; USUALLY AFFECTS THE FINGERS, HANDS, TOES, FEET, EARS, and NOSE.

# What Should Be Done: (land temperatures)

- Move the person to a warm dry area. Don't leave the person alone.
- Remove any wet or tight clothing that may cut off blood flow to the affected area.
- DO NOT rub the affected area, because rubbing causes damage to the skin and tissue.
- **Gently** place the affected area in a warm (105°F) water bath and monitor the water temperature to **slowly** warm the tissue. Don't pour warm water directly on the affected area because it will warm the tissue too fast causing tissue damage. Warming takes about 25-40 minutes.
- After the affected area has been warmed, it may become puffy and blister. The affected area may have a burning feeling or numbness. When normal feeling, movement, and skin color have returned, the affected area should be dried and wrapped to keep it warm. **Note:** If there is a chance the affected area may get cold again, do not warm the skin. If the skin is warmed and then becomes cold again, it will cause severe tissue damage.
- Seek medical attention as soon as possible.

## **HYPOTHERMIA - (Medical Emergency)**

## What Happens to the Body:

NORMAL BODY TEMPERATURE (98.6° F/37°C) DROPS TO OR BELOW 95°F (35°C); FATIGUE OR DROWSINESS; UNCONTROLLED SHIVERING; COOL BLUISH SKIN; SLURRED SPEECH; CLUMSY MOVEMENTS; IRRITABLE, IRRATIONAL OR CONFUSED BEHAVIOR.

## What Should Be Done: (land temperatures)

- Call for emergency help (i.e., Ambulance or Call 911).
- Move the person to a warm, dry area. Don't leave the person alone. Remove any wet clothing and replace with warm, dry clothing or wrap the person in blankets.
- Have the person drink warm, sweet drinks (sugar water or sports-type drinks) if they
  are alert. Avoid drinks with caffeine (coffee, tea, or hot chocolate) or alcohol.
- Have the person move their arms and legs to create muscle heat. If they are unable
  to do this, place warm bottles or hot packs in the arm pits, groin, neck, and head
  areas. DO NOT rub the person's body or place them in warm water bath. This may
  stop their heart.

## What Should Be Done: (water temperatures)

- Call for emergency help (Ambulance or Call 911). Body heat is lost up to 25 times faster in water.
- **DO NOT** remove any clothing. Button, buckle, zip, and tighten any collars, cuffs, shoes, and hoods because the layer of trapped water closest to the body provides a layer of insulation that slows the loss of heat. Keep the head out of the water and put on a hat or hood.
- Get out of the water as quickly as possible or climb on anything floating. DO NOT
  attempt to swim unless a floating object or another person can be reached because
  swimming or other physical activity uses the body's heat and reduces survival time
  by about 50 percent.
- If getting out of the water is not possible, wait quietly and conserve body heat by folding arms across the chest, keeping thighs together, bending knees, and crossing ankles. If another person is in the water, huddle together with chests held closely.

## How to Protect Workers

- Recognize the environmental and workplace conditions that lead to potential cold-induced illnesses and injuries.
- Learn the signs and symptoms of cold-induced illnesses/injuries and what to do to help the worker.
- Train the workforce about cold-induced illnesses and injuries.
- Select proper clothing for cold, wet, and windy conditions. Layer clothing to adjust to changing environmental temperatures. Wear a hat and gloves, in addition to underwear that will keep water away from the skin (polypropylene).
- Take frequent short breaks in warm dry shelters to allow the body to warm up.
- Perform work during the warmest part of the day.
- Avoid exhaustion or fatigue because energy is needed to keep muscles warm.
- Use the buddy system (work in pairs).
- Drink warm, sweet beverages (sugar water, sports-type drinks). Avoid drinks with caffeine (coffee, tea, or hot chocolate) or alcohol.
- Eat warm, high-calorie foods like hot pasta dishes.

## Workers Are at Increased Risk When...

- They have predisposing health conditions such as cardiovascular disease, diabetes, and hypertension.
- They take certain medication (check with your doctor, nurse, or pharmacy and ask if any medicines you are taking affect you while working in cold environments).
- They are in poor physical condition, have a poor diet, or are older.



# **Protecting Workers from Heat Stress**

## **Heat Illness**

Exposure to heat can cause illness and death. The most serious heat illness is heat stroke. Other heat illnesses, such as heat exhaustion, heat cramps and heat rash, should also be avoided.

There are precautions your employer should take any time temperatures are high and the job involves physical work.

## **Risk Factors for Heat Illness**

- High temperature and humidity, direct sun exposure, no breeze or wind
- · Low liquid intake; previous heat illnesses
- Heavy physical labor
- Waterproof clothing
- No recent exposure to hot workplaces

## **Symptoms of Heat Exhaustion**

- · Headache, dizziness, or fainting
- Weakness and wet skin
- Irritability or confusion
- Thirst, nausea, or vomiting

## **Symptoms of Heat Stroke**

- May be confused, unable to think clearly, pass out, collapse, or have seizures (fits)
- May stop sweating

## **To Prevent Heat Illness, Your Employer Should**

- Provide training about the hazards leading to heat stress and how to prevent them.
- Provide a lot of cool water to workers close to the work area. At least one pint of water per hour is needed.



For more complete information:





- Schedule frequent rest periods with water breaks in shaded or airconditioned areas.
- Routinely check workers who are at risk of heat stress due to protective clothing and high temperature.



Consider protective clothing that provides cooling.

## How You Can Protect Yourself and Others

- Know signs/symptoms of heat illnesses; monitor yourself; use a buddy system.
- Block out direct sun and other heat sources.
- Drink plenty of fluids. Drink often and BEFORE you are thirsty.
- Avoid beverages containing alcohol or caffeine.
- Wear lightweight, light colored, loosefitting clothes.
- Be aware that poor physical condition, some health problems (such as high blood pressure or diabetes), pregnancy, colds and flu, and some medications can increase your personal risk. If you are under treatment, ask your healthcare provider.





## What to Do When a Worker is III from the Heat

- Call a supervisor for help. If the supervisor is not available, call 911.
- Have someone stay with the worker until help arrives.
- · Move the worker to a cooler/shaded area.
- Remove outer clothing.
- Fan and mist the worker with water; apply ice (ice bags or ice towels).
- Provide cool drinking water, if able to drink.

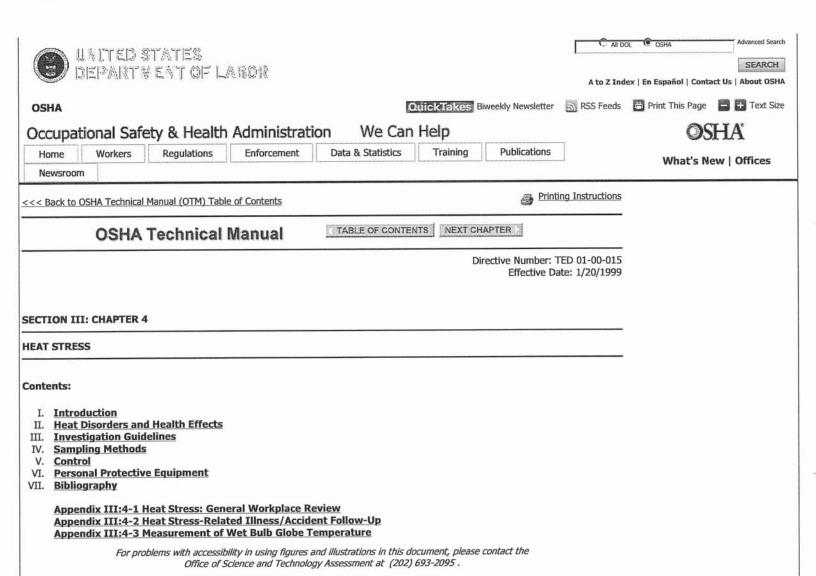
IF THE WORKER IS NOT ALERT or seems confused, this may be a heat stroke. CALL 911 IMMEDIATELY and apply ice as soon as possible.

If you have any questions or concerns, call OSHA at 1-800-321-OSHA.

For more complete information:



www.osha.gov (800) 321-OSHA



#### I. INTRODUCTION.

Operations involving high air temperatures, radiant heat sources, high humidity, direct physical contact with hot objects, or strenuous physical activities have a high potential for inducing heat stress in employees engaged in such operations. Such places include: iron and steel foundries, nonferrous foundries, brick-firing and ceramic plants, glass products facilities, rubber products factories, electrical utilities (particularly boiler rooms), bakeries, confectioneries, commercial kitchens, laundries, food canneries, chemical plants, mining sites, smelters, and steam tunnels.

Outdoor operations conducted in hot weather, such as construction, refining, asbestos removal, and hazardous waste site activities, especially those that require workers to wear semipermeable or impermeable protective clothing, are also likely to cause heat stress among exposed workers.

#### A. CAUSAL FACTORS.

- Age, weight, degree of physical fitness, degree of acclimatization, metabolism, use of alcohol or drugs, and a
  variety of medical conditions such as hypertension all affect a person's sensitivity to heat. However, even the type
  of clothing worn must be considered. Prior heat injury predisposes an individual to additional injury.
- It is difficult to predict just who will be affected and when, because individual susceptibility varies. In addition, environmental factors include more than the ambient air temperature. Radiant heat, air movement, conduction, and relative humidity all affect an individual's response to heat.

## B. **DEFINITIONS**.

- The American Conference of Governmental Industrial Hygienists (1992) states that workers should not be permitted to work when their deep body temperature exceeds 38°C (100.4°F).
- 2. Heat is a measure of energy in terms of quantity.

- A calorie is the amount of heat required to raise 1 gram of water 1°C (based on a standard temperature of 16.5 to 17.5°C).
- 4. Conduction is the transfer of heat between materials that contact each other. Heat passes from the warmer material to the cooler material. For example, a worker's skin can transfer heat to a contacting surface if that surface is cooler, and vice versa.
- Convection is the transfer of heat in a moving fluid. Air flowing past the body can cool the body if the air temperature is cool. On the other hand, air that exceeds 35°C (95°F) can increase the heat load on the body.
- Evaporative cooling takes place when sweat evaporates from the skin. High humidity reduces the rate of evaporation and thus reduces the effectiveness of the body's primary cooling mechanism.
- 7. Radiation is the transfer of heat energy through space. A worker whose body temperature is greater than the temperature of the surrounding surfaces radiates heat to these surfaces. Hot surfaces and infrared light sources radiate heat that can increase the body's heat load.
- 8. Globe temperature is the temperature inside a blackened, hollow, thin copper globe.
- 9. Metabolic heat is a by-product of the body's activity.
- 10. Natural wet bulb (NWB) temperature is measured by exposing a wet sensor, such as a wet cotton wick fitted over the bulb of a thermometer, to the effects of evaporation and convection. The term natural refers to the movement of air around the sensor.
- Dry bulb (DB) temperature is measured by a thermal sensor, such as an ordinary mercury-in-glass thermometer, that is shielded from direct radiant energy sources.

## II. HEAT DISORDERS AND HEALTH EFFECTS.

A. HEAT STROKE occurs when the body's system of temperature regulation fails and body 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 fitness and the timing and effectiveness of first aid treatment.

Regardless of the worker's protests, no employee suspected of being ill from heat stroke should be sent home or left unattended unless a physician has specifically approved such an order.

B. HEAT EXHAUSTION. The signs and symptoms of heat exhaustion are headache, nausea, 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, a medical emergency.

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

C. 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 by both too much and too little salt. Cramps appear to be caused by the lack of water replenishment. Because sweat is a hypotonic solution (±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. Recent studies have shown that drinking commercially available carbohydrate-electrolyte replacement liquids is effective in minimizing physiological disturbances during recovery.

D. HEAT COLLAPSE ("Fainting"). In heat collapse, the brain does not receive enough oxygen because blood pools in the extremities. As a result, the exposed individual may lose consciousness. This reaction is similar to that of heat exhaustion and does not affect the body's heat balance. However, the onset of heat collapse is rapid and unpredictable. To prevent heat collapse, the worker should gradually become acclimatized to the hot environment.

- E. HEAT RASHES are the most common problem in hot work environments. Prickly heat 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.
- F. HEAT FATIGUE. A factor that predisposes an individual to heat fatigue is lack of acclimatization. The use of a program of acclimatization and training for work in hot environments is advisable. The signs and symptoms of heat fatigue include impaired performance of skilled sensorimotor, mental, or vigilance jobs. There is no treatment for heat fatigue except to remove the heat stress before a more serious heat-related condition develops.

#### III. INVESTIGATION GUIDELINES.

These guidelines for evaluating employee heat stress approximate those found in the 1992-1993 ACGIH publication, *Threshold Limit Values for Chemical Substances and Physical Agents and Biological Exposure Indices.* 

#### A. EMPLOYER AND EMPLOYEE INTERVIEWS.

- The inspector will review the OSHA 200 Log and, if possible, the OSHA 101 forms for indications of prior heat stress problems.
- Following are some questions for employer interviews: What type of action, if any, has the employer taken to prevent heat stress problems? What are the potential sources of heat? What employee complaints have been made?
- 3. Following are some questions for employee interviews: What heat stress problems have been experienced? What type of action has the employee taken to minimize heat stress? What is the employer's involvement, i.e., does employee training include information on heat stress? (<u>Appendix III:4-1</u> lists factors to be evaluated when reviewing a heat stress situation, and <u>Appendix III:4-2</u> contains a follow-up checklist.)
- B. WALKAROUND INSPECTION. During the walkaround inspection, the investigator will: determine building and operation characteristics; determine whether engineering controls are functioning properly; verify information obtained from the employer and employee interviews; and perform temperature measurements and make other determinations to identify potential sources of heat stress. Investigators may wish to discuss any operations that have the potential to cause heat stress with engineers and other knowledgeable personnel. The walkaround inspection should cover all affected areas. Heat sources, such as furnaces, ovens, and boilers, and relative heat load per employee should be noted.

## C. WORK-LOAD ASSESSMENT.

 Under conditions of high temperature and heavy workload, the CSHO should determine the work-load category of each job (Table III:4-1 and Figure III:4-1). Work-load category is determined by averaging metabolic rates for the tasks and then ranking them:

Light work: up to 200 kcal/hour
 Medium work: 200-350 kcal/hour

3. Heavy work: 350-500 kcal/hour

Cool Rest Area: Where heat conditions in the rest area are different from those in the work area, the metabolic rate (M) should be calculated using a time-weighted average, as follows:

Equation III: 4-1. Average Metabolic Rate

Average<sub>M</sub> = 
$$\frac{(M_1)(t_1) + (M_2)(t_2) + ... + (M_n)(t_n)}{(t_1) + (t_2) + ... (t_n)}$$

where: M = metabolic rate

t = time in minutes

In some cases, a videotape is helpful in evaluating work practices and metabolic load.

## FIGURE III:4-1. ACTIVITY EXAMPLES

- Light hand work: writing, hand knitting
- Heavy hand work: typewriting
- Heavy work with one arm: hammering in nails (shoemaker, upholsterer)
- Light work with two arms: filing metal, planing wood, raking the garden
- Moderate work with the body: cleaning a floor, beating a carpet
- Heavy work with the body: railroad track laying, digging, barking trees

Sample Calculation: Assembly line work using a heavy hand tool

Walking along
Intermediate value between heavy work with two arms and light work with the body
Add for basal metabolism

1.0 kcal/min
Total: 6.0 kcal/min

Source: ACGIH 1992.

TABLE III:4-1. ASSESSMENT OF WORK

Body position ar	nd movement	kcal/min*
Sitting		0.3
Standing		0.6
Walking		2.0-3.0
Walking uphill		add 0.8 for every meter (yard) rise
Type of work	Average kcal/min	Range kcal/min
Hand work		
Light	0.4	0.2-1.2
Heavy	0.9	
Work: One arm		
Light	1.0	0.7-2.5
Heavy	1.7	
Work: Both arms		
Light	1.5	1.0-3.5
Heavy 2.5		
Work: Whole body	y	
Light	3.5	2.5-15.0
Moderate	5.0	
Heavy	7.0	
Very hear	vv 9.0	

Source: ACGIH 1992.

## IV. SAMPLING METHODS.

- A. BODY TEMPERATURE MEASUREMENTS. Although instruments are available to estimate deep body temperature by measuring the temperature in the ear canal or on the skin, these instruments are not sufficiently reliable to use in compliance evaluations.
- B. ENVIRONMENTAL MEASUREMENTS. Environmental heat measurements should be made at, or as close as possible to, the specific work area where the worker is exposed. When a worker is not continuously exposed in a single hot area but moves between two or more areas having different levels of environmental heat, or when the environmental heat varies substantially at a single hot area, environmental heat exposures should be measured for each area and for each level of environmental heat to which employees are exposed.
- C. WET BULB GLOBE TEMPERATURE INDEX.
  - 1. Wet Bulb Globe Temperature (WBGT) should be calculated using the appropriate formula in Appendix III:4-2. The

WBGT for continuous all-day or several hour exposures should be averaged over a 60-minute period. Intermittent exposures should be averaged over a 120-minute period. These averages should be calculated using the following

Equation III:4-2. Average Web Bulb Globe Temperature (WBGT)

Average<sub>WBGT</sub> = 
$$\frac{(WBGT_1)(t_1) + (WBGT_2)(t_2) + ... + (WBGT_n)(t_n)}{(t_1) + (t_2) + ... (t_n)}$$

For indoor and outdoor conditions with no solar load, WBGT is calculated as:

$$WBGT = 0.7NWB + 0.3GT$$

For outdoors with a solar load, WBGT is calculated as

$$WBGT = 0.7 NWB + 0.2 GT + 0.1 DB$$

WBGT = Wet Bulb Globe Temperature Index NWB = Nature Wet-Bulb Temperature

= Dry-Bulb Temperature DB GT Globe Temperature

- 2. The exposure limits in Table III:4-2 are valid for employees wearing light clothing. They must be adjusted for the insulation from clothing that impedes sweat evaporation and other body cooling mechanisms. Use Table III:4-3 to correct Table III:4-2 for various kinds of clothing.
- 3. Use of Table III:4-2 requires knowledge of the WBGT and approximate workload. Workload can be estimated using the data in Table III:4-1, and sample calculations are presented in Figure III:4-1.
- D. MEASUREMENT. Portable heat stress meters or monitors are used to measure heat conditions. These instruments can calculate both the indoor and outdoor WBGT index according to established ACGIH Threshold Limit Value equations. With this information and information on the type of work being performed, heat stress meters can determine how long a person can safely work or remain in a particular hot environment. See Appendix III:4-2 for an alternate method of

TABLE III:4-2. PERMISSIBLE HEAT EXPOSURE THRESHOLD LIMIT VALUE

		Work Load*	
Work/rest regimen	Light	Moderate	Heavy
Continuous work	30.0°C (86°F)	26.7°C (80°F)	25.0°C (77°F)
75% Work, 25% rest, each hour	30.6°C (87°F)	28.0°C (82°F)	25.9°C (78°F)
50% Work, 50% rest, each hour	31.4°C (89°F)	29.4°C (85°F)	27.9°C (82°F)
25% Work, 75% rest, each hour	32.2°C (90°F)	31.1°C (88°F)	30.0°C (86°F)
*Values are in °C and °F. WBGT.			

These TLV's are based on the assumption that nearly all acclimatized, fully clothed workers with adequate water and salt intake should be able to function effectively under the given working conditions without exceeding a deep body temperature of 38°C (100.4° F). They are also based on the assumption that the WBGT of the resting place is the same or very close to that of the workplace. Where the WBGT of the work area is different from that of the rest area, a time-weighted average should be used (consult the ACGIH 1992-1993 Threshold Limit Values for Chemical Substances and Physical Agents and Biological Exposure Indices (1992).

These TLV's apply to physically fit and acclimatized individuals wearing light summer clothing. If heavier clothing that impedes sweat or has a higher insulation value is required, the permissible heat exposure TLV's in Table III:4-2 must be reduced by the corrections shown in Table III:4-3.

Source: ACGIH 1992.

## E. OTHER THERMAL STRESS INDICES.

- The Effective Temperature index (ET) combines the temperature, the humidity of the air, and air velocity.
  This index has been used extensively in the field of comfort ventilation and air-conditioning. ET remains a
  useful measurement technique in mines and other places where humidity is high and radiant heat is low.
- The Heat-Stress Index (HSI) was developed by Belding and Hatch in 1965. Although the HSI considers all environmental factors and work rate, it is not completely satisfactory for determining an individual worker's heat stress and is also difficult to use.

TABLE III:4-3. WBGT CORRECTION FACTORS IN °C

Clothing type	Clo* value	WBGT correction
Summer lightweight working clothing	0.6	0
Cotton coveralls	1.0	-2
Winter work clothing	1.4	-4
Water barrier, permeable	1.2	-6

\*Clo: Insulation value of clothing. One clo = 5.55 kcal/m²/hr of heat exchange by radiation and convection for each degree °C difference in temperature between the skin and the adjusted dry bulb temperature.

Note: Deleted from the previous version are trade names and "fully encapsulating suit, gloves, boots and hood" including its clo value of 1.2 and WBGT correction of -10.

Source: ACGIH 1992.

#### V. CONTROL.

Ventilation, air cooling, fans, shielding, and insulation are the five major types of engineering controls used to reduce heat stress in hot work environments. Heat reduction can also be achieved by using power assists and tools that reduce the physical demands placed on a worker.

However, for this approach to be successful, the metabolic effort required for the worker to use or operate these devices must be less than the effort required without them. Another method is to reduce the effort necessary to operate power assists. The worker should be allowed to take frequent rest breaks in a cooler environment.

## A. ACCLIMATIZATION.

- The human body can adapt to heat exposure to some extent. This physiological adaptation is called
  acclimatization. After a period of acclimatization, the same activity will produce fewer cardiovascular demands. The
  worker will sweat more efficiently (causing better evaporative cooling), and thus will more easily be able to
  maintain normal body temperatures.
- 2. A properly designed and applied acclimatization program decreases the risk of heat-related illnesses. Such a program basically involves exposing employees to work in a hot environment for progressively longer periods. NIOSH (1986) says that, for workers who have had previous experience in jobs where heat levels are high enough to produce heat stress, the regimen should be 50% exposure on day one, 60% on day two, 80% on day three, and 100% on day four. For new workers who will be similarly exposed, the regimen should be 20% on day one, with a 20% increase in exposure each additional day.
- B. FLUID REPLACEMENT. Cool (50°-60°F) water or any cool liquid (except alcoholic beverages) should be made available to workers to encourage them to drink small amounts frequently, e.g., one cup every 20 minutes. Ample supplies of liquids should be placed close to the work area. Although some commercial replacement drinks contain salt, this is not necessary for acclimatized individuals because most people add enough salt to their summer diets.

## C. ENGINEERING CONTROLS.

- General ventilation is used to dilute hot air with cooler air (generally cooler air that is brought in from the
  outside). This technique clearly works better in cooler climates than in hot ones. A permanently installed
  ventilation system usually handles large areas or entire buildings. Portable or local exhaust systems may be more
  effective or practical in smaller areas.
- Air treatment/air cooling differs from ventilation because it reduces the temperature of the air by removing heat (and sometimes humidity) from the air.
- 3. Air conditioning is a method of air cooling, but it is expensive to install and operate. An alternative to air conditioning is the use of chillers to circulate cool water through heat exchangers over which air from the ventilation system is then passed; chillers are more efficient in cooler climates or in dry climates where

evaporative cooling can be used.

- 4. Local air cooling can be effective in reducing air temperature in specific areas. Two methods have been used successfully in industrial settings. One type, cool rooms, can be used to enclose a specific workplace or to offer a recovery area near hot jobs. The second type is a portable blower with built-in air chiller. The main advantage of a blower, aside from portability, is minimal set-up time.
- 5. Another way to reduce heat stress is to increase the air flow or convection using fans, etc. in the work area (as long as the air temperature is less than the worker's skin temperature). Changes in air speed can help workers stay cooler by increasing both the convective heat exchange (the exchange between the skin surface and the surrounding air) and the rate of evaporation. Because this method does not actually cool the air, any increases in air speed must impact the worker directly to be effective.

If the dry bulb temperature is higher than 35°C (95°F), the hot air passing over the skin can actually make the worker hotter. When the temperature is more than 35°C and the air is dry, evaporative cooling may be improved by air movement, although this improvement will be offset by the convective heat. When the temperature exceeds 35°C and the relative humidity is 100%, air movement will make the worker hotter. Increases in air speed have no effect on the body temperature of workers wearing vapor-barrier clothing.

- Heat conduction methods include insulating the hot surface that generates the heat and changing the surface itself.
- 7. Simple engineering controls, such as shields, can be used to reduce radiant heat, i.e. heat coming from hot surfaces within the worker's line of sight. Surfaces that exceed 35°C (95°F) are sources of infrared radiation that can add to the worker's heat load. Flat black surfaces absorb heat more than smooth, polished ones. Having cooler surfaces surrounding the worker assists in cooling because the worker's body radiates heat toward them.

With some sources of radiation, such as heating pipes, it is possible to use both insulation and surface modifications to achieve a substantial reduction in radiant heat. Instead of reducing radiation from the source, shielding can be used to interrupt the path between the source and the worker. Polished surfaces make the best barriers, although special glass or metal mesh surfaces can be used if visibility is a problem.

Shields should be located so that they do not interfere with air flow, unless they are also being used to reduce convective heating. The reflective surface of the shield should be kept clean to maintain its effectiveness.

#### D. ADMINISTRATIVE CONTROLS AND WORK PRACTICES.

- Training is the key to good work practices. Unless all employees understand the reasons for using new, or changing old, work practices, the chances of such a program succeeding are greatly reduced.
- 2. NIOSH (1986) states that a good heat stress training program should include at least the following components:
  - Knowledge of the hazards of heat stress;
  - Recognition of predisposing factors, danger signs, and symptoms;
  - Awareness of first-aid procedures for, and the potential health effects of, heat stroke;
  - Employee responsibilities in avoiding heat stress;
  - Dangers of using drugs, including therapeutic ones, and alcohol in hot work environments;
  - Use of protective clothing and equipment; and
  - Purpose and coverage of environmental and medical surveillance programs and the advantages of worker participation in such programs.
- 3. Hot jobs should be scheduled for the cooler part of the day, and routine maintenance and repair work in hot areas should be scheduled for the cooler seasons of the year.

## E. WORKER MONITORING PROGRAMS.

- Every worker who works in extraordinary conditions that increase the risk of heat stress should be personally
  monitored. These conditions include wearing semipermeable or impermeable clothing when the temperature
  exceeds 21°C (69.8°F), working at extreme metabolic loads (greater than 500 kcal/hour), etc.
- Personal monitoring can be done by checking the heart rate, recovery heart rate, oral temperature, or extent of body water loss.
- 3. 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.
- 4. The recovery heart rate can be checked by comparing the pulse rate taken at 30 seconds (P<sub>1</sub>) with the pulse rate taken at 2.5 minutes (P<sub>3</sub>) after the rest break starts. The two pulse rates can be interpreted using Table III:4-4.
- Oral temperature can be checked with a clinical thermometer after work but before the employee drinks water. If the oral temperature taken under the tongue exceeds 37.6°C, shorten the next work cycle by one third.
- 6. Body water loss can be measured by weighing the worker on a scale at the beginning and end of each work day.

The worker's weight loss should not exceed 1.5% of total body weight in a work day. If a weight loss exceeding this amount is observed, fluid intake should increase.

- F. OTHER ADMINISTRATIVE CONTROLS. The following administrative controls can be used to reduce heat stress:
  - Reduce the physical demands of work, e.g., excessive lifting or digging with heavy objects;
  - Provide recovery areas, e.g., air-conditioned enclosures and rooms;
  - Use shifts, e.g., early morning, cool part of the day, or night work;
  - Use intermittent rest periods with water breaks;
  - Use relief workers;
  - Use worker pacing; and
  - Assign extra workers and limit worker occupancy, or the number of workers present, especially in confined or enclosed spaces.

TABLE III:4-4. HEART RATE RECOVERY CRITERIA

Heart rate recovery pattern	P <sub>3</sub>	Difference between P <sub>1</sub> and P <sub>3</sub>
Satisfactory recovery	<90	=
High recovery (Conditions may require further study)	90	10
No recovery (May indicate too much stress)	90	<10

#### VI. PERSONAL PROTECTIVE EQUIPMENT.

A. REFLECTIVE CLOTHING, which can vary from aprons and jackets to suits that completely enclose the worker from neck to feet, can stop the skin from absorbing radiant heat. However, since most reflective clothing does not allow air exchange through the garment, the reduction of radiant heat must more than offset the corresponding loss in evaporative cooling. For this reason, reflective clothing should be worn as loosely as possible. In situations where radiant heat is high, auxiliary cooling systems can be used under the reflective clothing.

#### B. AUXILIARY BODY COOLING.

- Commercially available ice vests, though heavy, may accommodate as many as 72 ice packets, which are usually
  filled with water. Carbon dioxide (dry ice) can also be used as a coolant. The cooling offered by ice packets lasts
  only 2 to 4 hours at moderate to heavy heat loads, and frequent replacement is necessary. However, ice vests do
  not encumber the worker and thus permit maximum mobility. Cooling with ice is also relatively inexpensive.
- 2. Wetted clothing is another simple and inexpensive personal cooling technique. It is effective when reflective or other impermeable protective clothing is worn. The clothing may be wetted terry cloth coveralls or wetted two-piece, whole-body cotton suits. This approach to auxiliary cooling can be quite effective under conditions of high temperature and low humidity, where evaporation from the wetted garment is not restricted.
- Water-cooled garments range from a hood, which cools only the head, to vests and "long johns," which offer
  partial or complete body cooling. Use of this equipment requires a battery-driven circulating pump, liquid-ice
  coolant, and a container.

Although this system has the advantage of allowing wearer mobility, the weight of the components limits the amount of ice that can be carried and thus reduces the effective use time. The heat transfer rate in liquid cooling systems may limit their use to low-activity jobs; even in such jobs, their service time is only about 20 minutes per pound of cooling ice. To keep outside heat from melting the ice, an outer insulating jacket should be an integral part of these systems.

4. Circulating air is the most highly effective, as well as the most complicated, personal cooling system. By directing compressed air around the body from a supplied air system, both evaporative and convective cooling are improved. The greatest advantage occurs when circulating air is used with impermeable garments or double cotton overalls.

One type, used when respiratory protection is also necessary, forces exhaust air from a supplied-air hood ("bubble hood") around the neck and down inside an impermeable suit. The air then escapes through openings in the suit. Air can also be supplied directly to the suit without using a hood in three ways:

- by a single inlet;
- by a distribution tree; or
- by a perforated vest.

In addition, a vortex tube can be used to reduce the temperature of circulating air. The cooled air from this tube can be introduced either under the clothing or into a bubble hood. The use of a vortex tube separates the air stream into a hot and cold stream; these tubes also can be used to supply heat in cold climates. Circulating air, however, is noisy and requires a constant source of compressed air supplied through an attached air hose.

One problem with this system is the limited mobility of workers whose suits are attached to an air hose. Another is that of getting air to the work area itself. These systems should therefore be used in work areas where workers are not required to move around much or to climb. Another concern with these systems is that they can lead to dehydration. The cool, dry air feels comfortable and the worker may not realize that it is important to drink liquids frequently.

C. RESPIRATOR USAGE. The weight of a self-contained breathing apparatus (SCBA) increases stress on a worker, and this stress contributes to overall heat stress. Chemical protective clothing such as totally encapsulating chemical protection suits will also add to the heat stress problem.

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#### APPENDIX III:4-1. HEAT STRESS: GENERAL WORKPLACE REVIEW.

**NOTE:** Listed below are sample questions that the Compliance Officer may wish to consider when investigating heat stress in the workplace.

#### WORKPLACE DESCRIPTION.

- A. Type of business
- B. Heat-producing equipment or processes used
- C. Previous history (if any) of heat-related problems
- D. At "hot" spots:
  - Is the heat steady or intermittent?
  - Number of employees exposed?
  - For how many hours per day?
  - Is potable water available?
  - Are supervisors trained to detect/evaluate heat stress symptoms?

#### ARE EXPOSURES TYPICAL FOR A WORKPLACE IN THIS INDUSTRY?

- A. Weather at Time of Review
- B. Temperature
- C. Humidity
- D. Air velocity
- E. Is Day Typical of Recent Weather Conditions? (Get information from the Weather Bureau)
- F. Heat-Reducing Engineering Controls

- G. Ventilation in place?
- H. Ventilation operating?
- I. Air conditioning in place?
- J. Air conditioning operating?
- K. Fans in place?
- L. Fans operating?
- M. Shields or insulation between sources and employees?
- N. Are reflective faces of shields clean?

#### WORK PRACTICES TO DETECT, EVALUATE, AND PREVENT OR REDUCE HEAT STRESS.

- A. Training program?
- B. Content?
- C. Where given?
- D. For whom?
- E. Liquid replacement program?
- F. Acclimatization program?
- G. Work/rest schedule?
- H. Scheduling of work (during cooler parts of shift, cleaning and maintenance during shut-downs, etc.)
- I. Cool rest areas (including shelter at outdoor work sites)?
- J. Heat monitoring program?
- K. Personal Protective Equipment
- L. Reflective clothing in use?
- M. Ice and/or water-cooled garments in use?
- N. Wetted undergarments (used with reflective or impermeable clothing) in use?
- O. Circulating air systems in use?
- P. First Aid Program
- Q. Trained personnel?
- R. Provision for rapid cool-down?
- S. Procedures for getting medical attention?
- T. Transportation to medical facilities readily available for heat stroke victims?
- U. Medical Screening and Surveillance Program
- V. Content?
- W. Who manages program?
- X. Additional Comments

(Use additional pages as needed.)

## APPENDIX III: 4-2. HEAT STRESS-RELATED ILLNESS OR ACCIDENT FOLLOW-UP.

- A. Describe events leading up to the episode.
- B. Evaluation/comments by other workers at the scene.
- C. Work at time of episode (heavy, medium, light)?
- D. How long was affected employee working at site prior to episode?
- E. Medical history of affected worker, if known.
- F. Appropriate engineering controls in place?
- G. Appropriate engineering controls in operation?
- H. Appropriate work practices used by affected employee(s)?
- I. Appropriate personal protective equipment available?
- J. Appropriate personal protective equipment in use?
- K. Medical screening for heat stress and continued surveillance for signs of heat stress given other employees?
- L. Additional comments regarding specific episode(s): (Use additional pages as needed.)

## APPENDIX III: 4-3. MEASUREMENT OF WET BULB GLOBE TEMPERATURE.

Measurement is often required of those environmental factors that most nearly correlate with deep body temperature and other physiological responses to heat. At the present time, the Wet Bulb Globe Temperature Index (WBGT) is the most used technique to measure these environmental factors. WBGT values are calculated by the following equations:

Equation III:4-4. Indoor or Outdoor Wet Bulb Globe Temperature Indexes (WBGI) Indoor or outdoors with no solar load

WBGT = 0.7 NWB + 0.3 GT

Outdoors with solar load

WBGT = 0.7NWB + 0.2GT + 0.1DB

where: WBGT = Wet Bulb Globe Temperature Index

NWB = Natural Wet-Bulb Temperature

DB = Dry-Bulb (air) Temperature

GT = Globe Thermometer Temperature

The determination of WBGT requires the use of a black globe thermometer, a natural (static) wet-bulb thermometer, and a dry-bulb thermometer. The measurement of environmental factors shall be performed as follows:

- 1. The range of the dry and the natural wet-bulb thermometers should be  $-5^{\circ}$ C to  $+50^{\circ}$ C, with an accuracy of  $\pm 0.5^{\circ}$ C. The dry bulb thermometer must be shielded from the sun and the other radiant surfaces of the environment without restricting the airflow around the bulb. The wick of the natural wet bulb thermometer should be kept wet with distilled water for at least one-half hour before the temperature reading is made. It is not enough to immerse the other end of the wick into a reservoir of distilled water and wait until the whole wick becomes wet by capillarity. The wick must be wetted by direct application of water from a syringe one-half hour before each reading. The wick must cover the bulb of the thermometer and an equal length of additional wick must cover the stem above the bulb. The wick should always be clean, and new wicks should be washed before using.
- 2. A globe thermometer, consisting of a 15 cm (6-inch) in diameter hollow copper sphere painted on the outside with a matte black finish, or equivalent, must be used. The bulb or sensor of a thermometer (range -5°C to  $\pm 100$ °C with an accuracy of  $\pm 0.5$ °C) must be fixed in the center of the sphere. The globe thermometer should be exposed at least 25 minutes before it is read.
- A stand should be used to suspend the three thermometers so that they do not restrict free air flow around the bulbs and the wet-bulb and globe thermometer are not shaded.
- It is permissible to use any other type of temperature sensor that gives a reading similar to that of a mercury thermometer under the same conditions.
- 5. The thermometers must be placed so that the readings are representative of the employee's work or rest areas, as appropriate.

Once the WBGT has been estimated, employers can estimate workers' metabolic heat load (see Tables III:4-1 and III:4-2) and use the ACGIH method to determine the appropriate work/rest regimen, clothing, and equipment to use to control the heat exposures of workers in their facilities.

Freedom of Information Act | Privacy & Security Statement | Disclaimers | Customer Survey | Important Web Site Notices | International | Contact Us

U.S. Department of Labor | Occupational Safety & Health Administration | 200 Constitution Ave., NW, Washington, DC 20210 Telephone: 800-321-OSHA (6742) | TTY: 877-889-5627

www.OSHA.gov

## **APPENDIX E**

Health and Safety Briefing/ Tailgate Meeting Form

# HEALTH AND SAFETY BRIEFING / TAILGATE MEETING FORM

Site Name / Location	
Date:	Weather Forecast:
Names of Personnel Attending Briefing	
Planned Work	
<u>Items Discussed</u>	
Work Permit Type and Applicable Restrictions:	
Signatures of Attending Personnel	

## APPENDIX F

Medical Data Form



(Patient Must Present Photo ID at Time of Service)

## **Authorization for Examination or Treatment**

Patient Name:	Social Security Number:
Employer:	Date of Birth:
Street Address:	Location Number:
Temporary Staffing Agency:	
Work Related	Physical Examination
□ Injury □ Illness	☐ Preplacement ☐ Baseline ☐ Annual ☐ Exit
Date of Injury	DOT Physical Examination
Substance Abuse Testing <sup>★</sup> (check all that apply)	☐ Preplacement ☐ Recertification
☐ Regulated drug screen ☐ Breath alcohol	Special Examination
☐ Collection only ☐ Hair collect	□ Asbestos □ Respirator □ Audiogram
☐ Non-regulated drug screen ☐ Rapid drug screen	☐ Human Performance Evaluation*
☐ Other	☐ HAZMAT ☐ Medical Surveillance
Type of Substance Abuse Testing	☐ Other
☐ Preplacement ☐ Reasonable cause	Billing (check if applicable)
☐ Post-accident ☐ Random	☐ Employee to pay charges
☐ Follow-up	
Special instructions/comments:	patient and staff are allowed in the testing/treatment area. Please alert your employee so that they can make
Authorized by: Please print	Title:
Phone: ()	

Concentra now offers urgent care services for non-work related illness and injury. We accept many insurance plans.

(Copies of this form are available at www.concentra.com)

# **Concentra**°

Patient Information

Improving America's health, one patient at a time.

Last name: _	First name:		MI.
	Date of birth (MM/DD/Y)		
	Apt. #		
Contact phone (home or cell):	Work phon	e:	🗅 Female 🗀 Male
Occupation			☐ Single ☐ Married
Employer Requestin	3 Services		
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Contact name;		Contact phon	£(
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s your employment arranged throu	igh a temporary hire agency? $\square$ Yes $\square$ No Name of agenc	cy:	Agency phone:
	The information provided is correct to the best of my known responsible for any errors or omissions that I may have mo		
	employer to verify the purpose of my visit, if necessary.		
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You may be contacted by Westgate Research, acting on behalf of Concentra, to participate in a satisfaction survey about this visit. We rely on your feedback to help us improve.

# **Concentra**°

## Información del Paciente

Mejorando la salud de los Estados Unidos, un paciente a la vez.

pellido:	Norm	bre:		Inicio	al Seg. Nombre:	
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cupación:					☐ Soltero(a)	☐ Casado
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*		Ubia	ación/Tienda Número	,		
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	Nombre (letra imprenta por fo Firma:  Fecha y hora de recibida la no  Si ustes esté aquí por una lesión.	Nificación:				
Fecho de la lesión:	Hora de la lesión:					<b>\</b>
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G. T. C.						
¿Cómo ocurrió la lesión?						
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consulta Nosotros contamos con esta información, la cual nos ayuda a mejorar.

## **Patient Information**



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	Apt #			<u>.</u>	71P-
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How did you learn about Concentra? (Check one, please.)	□Billboard □Direct mail □Doctor referral □Insurance company □Internet □Movie the □School □Apartment Complex	* ' ' '			
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msurare	I certify that the information provided is correct to				
	employees responsible for any errors or omissions	o man i min'i mase minas mi Cambu	eresg me mommunem		
	You may be contacted by Westgate Research, ac We rely on your feedback to help us improve.		_		

## Información Paciente



Gracias por confiarnos sus cuidados de hoy. \_\_\_\_\_\_ Nombre: \_\_\_\_\_ \_\_\_\_\_ Inicial Segundo Nombre: \_\_\_ #SS del paciente: Fecho de nacimiento (MM/DD/AAAA): Casodo(a) Teléfono en casa: Teléfono celular: ■Soltero(a) ☐ Hombre ☐ Mujer Motivo de la consulta: Correo electrónico del paciente: Dirección: \_\_\_\_\_\_ Estado: \_\_\_\_ Cód. Postal: \_\_\_\_\_ Teléfono: Nombre del médico de atención primaria: Nombre del empleador: Dirección del empleador: \_\_\_\_\_\_ Ciudad: \_\_\_\_\_ Estado: \_\_\_\_ Cód. Postal: \_\_\_\_\_\_ Contacto de Emergencia: Teléphono de Contacto de Emergencia: □Vallo □Correo Directo □Remitido por un doctor □Pasábamos por aquí □Empleador □ Paciente existente Cómo se enteró de Concentra? □Amigo/pariente □Compañía de seguro □Internet □Teatro de cine □Periódico □Radio □Farmacia □Escuela (Por favor maraue una) □Compleio de Apartamentos El pago de hay la va a hacer. Pago de hoy 🗆 El paciente — Yo pagaré la cuenta total usando: 🗆 Efectivo 🖵 Cheque 🗀 VISA 🗀 AtasterCard 🗀 Discover 🖵 Tanjeta Débito Cómo va a pagar la cuento de hoy? 🗖 La Compañía paga - Estay participando en un programa que es paçado por la Compañía E seguro — Yo presentaré mi tarieta de seguro y una forma de identificación aprobada (Por fovor complete las siguientes dos secciones). Empleodor de la persona osegurado: Información del seguro Compañía de seguio: Si usted está usando Identificación del Mierribro: # de Grupo: seguro para pagar Dirección de redamas: \_\_\_\_\_\_Cód. Postal: \_\_\_\_\_\_\_Estado: \_\_\_\_\_Cód. Postal: \_\_\_\_\_ la cuento de hoy, por favor provédnos Tiene securo con mas de un plan de salud? 🔲 Si 🔲 No con la siguiente Si si, nombre el otro seguro: (Par favor presente ambas tarjetas de identificación al registrarse) \_\_\_\_\_\_ Nombre: \_\_\_\_\_\_ Inicial Seg. Nombre: \_\_\_\_\_ Información de la cuenta # de SS en lo Cuenta: \_\_\_\_\_\_\_ Fecha de Nocimiento: (MM/DD/AAAA) \_\_\_\_\_\_ Si usted está usando Teléfono celular: seguro, esta es Cindod: Estado: Cód. Postol: información acerca de la persona que tiene el seguro... Relación con el paciente: 🔲 Usted mismo 🗀 Cónyuge 🗀 Padre/Guardián 🗖 Otro: \_\_\_\_\_\_\_ (Por fovor marque una) Yo certifico que la información provista es correcta hasta donde yo sé. Yo no haré responsable a Concentra, sus proveedores de la salud, o sus empleados por cualquier error u omisión que yo haya hecho al llenar la información en este formulario. & Firma: \_\_\_

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Version 2.6

02/13

CONPIF-0911

Health and Safety Plan	Health	and	Safety	Plan
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## **APPENDIX G**

Generic Community Air Monitoring Plan

## **APPENDIX G**

## New York State Department of Health Generic Community Air Monitoring Plan

## Overview

A Community Air Monitoring Plan (CAMP) requires real-time monitoring for volatile organic compounds (VOCs) and particulates (i.e., dust) at the downwind perimeter of each designated work area when certain activities are in progress at contaminated sites. The CAMP is not intended for use in establishing action levels for worker respiratory protection. Rather, its intent is to provide a measure of protection for the downwind community (i.e., off-site receptors including residences and businesses and on-site workers not directly involved with the subject work activities) from potential airborne contaminant releases as a direct result of investigative and remedial work activities. The action levels specified herein require increased monitoring, corrective actions to abate emissions, and/or work shutdown. Additionally, the CAMP helps to confirm that work activities did not spread contamination off-site through the air.

The generic CAMP presented below will be sufficient to cover many, if not most sites. Specific requirements should be reviewed for each situation in consultation with NYSDOH to ensure proper applicability. In some cases, a separate site-specific CAMP or supplement may be required. Depending upon the nature of contamination, chemical-specific monitoring with appropriately sensitive methods may be required. Depending upon the proximity of potentially exposed individuals, more stringent monitoring or response levels than those presented below may be required. Special requirements will be necessary for work within 20 feet of potentially exposed individuals or structures and for indoor work with co-located residences or facilities. These requirements should be determined in consultation with NYSDOH.

Reliance on the CAMP should not preclude simple, common-sense measures to keep VOCs, dust, and odors at a minimum around the work areas.

## **Community Air Monitoring Plan**

Depending upon the nature of known or potential contaminants at each site, real-time air monitoring for VOCs and/or particulate levels at the perimeter of the exclusion zone or work area will be necessary. Most sites will involve VOC and particulate monitoring; sites known to be contaminated with heavy metals alone may only require particulate monitoring. If radiological contamination is a concern, additional monitoring requirements may be necessary per consultation with appropriate DEC/NYSDOH staff.

**Continuous monitoring** will be required for all ground intrusive activities and during the demolition of contaminated or potentially contaminated structures. Ground intrusive activities include, but are not limited to, soil/waste excavation and handling, test pitting or trenching, and the installation of soil borings or monitoring wells.

**Periodic monitoring** for VOCs will be required during non-intrusive activities such as the collection of soil and sediment samples or the collection of groundwater samples from existing

monitoring wells. "Periodic" monitoring during sample collection might reasonably consist of taking a reading upon arrival at a sample location, monitoring while opening a well cap or overturning soil, monitoring during well baling/purging, and taking a reading prior to leaving a sample location. In some instances, depending upon the proximity of potentially exposed individuals, continuous monitoring may be required during sampling activities. Examples of such situations include groundwater sampling at wells on the curb of a busy urban street, in the midst of a public park, or adjacent to a school or residence.

## **VOC Monitoring, Response Levels, and Actions**

Volatile organic compounds (VOCs) must be monitored at the downwind perimeter of the immediate work area (i.e., the exclusion zone) on a continuous basis or as otherwise specified. Upwind concentrations should be measured at the start of each workday and periodically thereafter to establish background conditions, particularly if wind direction changes. The monitoring work should be performed using equipment appropriate to measure the types of contaminants known or suspected to be present. The equipment should be calibrated at least daily for the contaminant(s) of concern or for an appropriate surrogate. The equipment should be capable of calculating 15-minute running average concentrations, which will be compared to the levels specified below.

- 1. If the ambient air concentration of total organic vapors at the downwind perimeter of the work area or exclusion zone exceeds 5 parts per million (ppm) above background for the 15-minute average, work activities must be temporarily halted and monitoring continued. If the total organic vapor level readily decreases (per instantaneous readings) below 5 ppm over background, work activities can resume with continued monitoring.
- 2. If total organic vapor levels at the downwind perimeter of the work area or exclusion zone persist at levels in excess of 5 ppm over background but less than 25 ppm, work activities must be halted, the source of vapors identified, corrective actions taken to abate emissions, and monitoring continued. After these steps, work activities can resume provided that the total organic vapor level 200 feet downwind of the exclusion zone or half the distance to the nearest potential receptor or residential/commercial structure, whichever is less but in no case less than 20 feet, is below 5 ppm over background for the 15-minute average.
- 3. If the organic vapor level is above 25 ppm at the perimeter of the work area, activities must be shutdown.
- 4. All 15-minute readings must be recorded and be available for State (DEC and NYSDOH) personnel to review. Instantaneous readings, if any, used for decision purposes should also be recorded.

## Particulate Monitoring, Response Levels, and Actions

Particulate concentrations should be monitored continuously at the upwind and downwind perimeters of the exclusion zone at temporary particulate monitoring stations. The particulate monitoring should be performed using real-time monitoring equipment capable of measuring

particulate matter less than 10 micrometers in size (PM-10) and capable of integrating over a period of 15 minutes (or less) for comparison to the airborne particulate action level. The equipment must be equipped with an audible alarm to indicate exceedance of the action level. In addition, fugitive dust migration should be visually assessed during all work activities.

- 1. If the downwind PM-10 particulate level is 100 micrograms per cubic meter (mcg/m³) greater than background (upwind perimeter) for the 15-minute period or if airborne dust is observed leaving the work area, then dust suppression techniques must be employed. Work may continue with dust suppression techniques provided that downwind PM-10 particulate levels do not exceed 150 mcg/m³ above the upwind level and provided that no visible dust is migrating from the work area.
- 2. If, after implementation of dust suppression techniques, downwind PM-10 particulate levels are greater than 150 mcg/m³ above the upwind level, work must be stopped and a reevaluation of activities initiated. Work can resume provided that dust suppression measures and other controls are successful in reducing the downwind PM-10 particulate concentration to within 150 mcg/m³ of the upwind level and in preventing visible dust migration.
- 3. All readings must be recorded and be available for State (DEC and NYSDOH) and County Health personnel to review.

Health	and	<b>Safety</b>	Plan
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## **APPENDIX H**

Accident Report and Investigation Form

☐ Roux Associates, Inc.	☐ Remedial Engineering, P.C
(Check applica	able company name)

## ACCIDENT REPORT

Joe Gentile, Corporate Health and Safety Manager
Cell: (610) 844-6911; Office: (856) 423-8800; Office FAX: (856) 423-3220; Home: (484) 373-0953

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Involved in Incident:	Roux/Rem	nedial Subco		Yrs in Cu	urrent Occupation;	Address; an					Phone		
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## Accident Report - Page 2

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2)									
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Authority/Agency Notified:		Name Notifie		Fax # of Person	Address of Person Notified:	Da	te & Time of Notificat	ion: Exact Inform Reported/Pr	
II. PUBLIC RESPONS	ES TO INC	CIDENT	(if applic	able)					
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Roux/Remedial Response:									
(Check all that apply.) (ATTACHED INFORMA		tos, drav □Pr		to help illustrate		orm	☐Police Rep	oort 🗆	ther
Name(s) of person(s) v	who prepa	ared Init	ial and	Title(s):			Phone num	ber(s):	
- 4			PAR	T 3: INVE	STIGATION TEAM	ANA	LYSIS		
(Root Causes: Lack of know	ledge or skill rocedures or	l, Doing the	ne task acco	ording to procedures did not result in an a	CAUSAL FACTORS AN or acceptable practices takes more tin accident, Lack of or inadequate proced	ne or effe	ort, Short-cuts or not fe	ollowing acceptable p	ractices is reinforced
ROOT (	CAUSE	(S) AN	ND SO	LUTION(S)	: HOW TO PREVEN	T INC	IDENT FRO	M RECURRI	NG
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	NAME				JOB POSITION		DATE	SIG	NATURE
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## **APPENDIX I**

Acord Automobile Loss Form

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REMAR	KS (AC	ORD .	101, Additional Ro	marks S	chedule, maj	y be altach	ed if more space i	s required	)	·			
						•							

## APPLICABLE IN ALABAMA

Any person who knowingly presents a false or fraudulent claim for payment of a loss or benefit or who knowingly presents false information in an application for insurance is guilty of a crime and may be subject to restitution fines or confinement in prison, or any combination thereof.

#### **APPLICABLE IN ALASKA**

A person who knowingly and with intent to injure, defraud, or deceive an insurance company files a claim containing false, incomplete, or misleading information may be prosecuted under state law.

## APPLICABLE IN ARIZONA

For your protection, Arizona law requires the following statement to appear on this form. Any person who knowingly presents a false or fraudulent claim for payment of a loss is subject to criminal and civil penalties.

# APPLICABLE IN ARKANSAS, DELAWARE, KENTUCKY, LOUISIANA, MAINE, MICHIGAN, NEW JERSEY, NEW MEXICO, NORTH DAKOTA, PENNSYLVANIA, RHODE ISLAND, SOUTH DAKOTA, TENNESSEE, TEXAS, VIRGINIA, AND WEST VIRGINIA

Any person who knowingly and with intent to defraud any insurance company or another person, files a statement of claim containing any materially false information, or conceals for the purpose of misleading, information concerning any fact, material thereto, commits a fraudulent insurance act, which is a crime, subject to criminal prosecution and civil penalties. In LA, ME, TN, and VA, insurance benefits may also be denied.

## APPLICABLE IN CALIFORNIA

For your protection, California law requires the following to appear on this form: Any person who knowingly presents a false or fraudulent claim for payment of a loss is guilty of a crime and may be subject to fines and confinement in state prison.

## APPLICABLE IN COLORADO

It is unlawful to knowingly provide false, incomplete, or misleading facts or information to an insurance company for the purpose of defrauding or attempting to defraud the company. Penalties may include imprisonment, fines, denial of insurance, and civil damages. Any insurance company or agent of an insurance company who knowingly provides false, incomplete, or misleading facts or information to a policy holder or claimant for the purpose of defrauding or attempting to defraud the policy holder or claimant with regard to a settlement or award payable from insurance proceeds shall be reported to the Colorado Division of insurance within the Department of Regulatory Agencies.

## APPLICABLE IN THE DISTRICT OF COLUMBIA

Warning: It is a crime to provide false or misleading information to an insurer for the purpose of defrauding the insurer or any other person. Penalties include imprisonment and/or fines. In addition, an insurer may deny insurance benefits, if false information materially related to a claim was provided by the applicant.

## APPLICABLE IN FLORIDA

Pursuant to S. 817.234, Florida Statutes, any person who, with the intent to Injure, defraud, or deceive any insurer or insured, prepares, presents, or causes to be presented a proof of loss or estimate of cost or repair of damaged property in support of a claim under an insurance policy knowing that the proof of loss or estimate of claim or repairs contains any false, incomplete, or misleading information concerning any fact or thing material to the claim commits a felony of the third degree, punishable as provided in S. 775.083, or S. 775.084, Florida Statutes.

## APPLICABLE IN HAWAII

For your protection, Hawaii law requires you to be informed that presenting a fraudulent daim for payment of a loss or benefit is a crime punishable by fines or imprisonment, or both.

## **APPLICABLE IN IDAHO**

Any person who knowingly and with the intent to injure, defraud, or deceive any insurance company files a statement of claim containing any false, incomplete or misleading information is guilty of a felony.

## APPLICABLE IN INDIANA

A person who knowingly and with Intent to defraud an insurer files a statement of claim containing any false, incomplete, or misleading information commits a felony.

AGEN	av ci	INTO	Men	in:
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#### **APPLICABLE IN KANSAS**

Any person who, knowledge and with intent to defraud, presents, causes to be presented or prepares with knowledge or belief that it will be presented to or by an insurer, purported insurer, broker or any agent thereof, any written statement as part of, or in support of, an application for the issuance of, or the rating of an insurance policy for personal or commercial insurance, or a claim for payment or other benefit pursuant to an insurance policy for commercial or personal insurance which such person knows to contain materially false information concerning any fact material thereto; or conceals, for the purpose of misleading, information concerning any fact material thereto commits a fraudulent insurance act.

#### APPLICABLE IN MARYLAND

Any person who knowingly or willfully presents a false or fraudulent claim for payment of a loss or benefit or who knowingly or willfully presents false information in an application for insurance is guilty of a crime and may be subject to fines and confinement in prison.

## APPLICABLE IN MINNESOTA

A person who files a claim with intent to defraud or helps commit a fraud against an insurer is guilty of a crime.

## **APPLICABLE IN NEVADA**

Pursuant to NRS 686A.291, any person who knowingly and willfully files a statement of claim that contains any false, incomplete or misleading information concerning a material fact is guilty of a felony.

#### APPLICABLE IN NEW HAMPSHIRE

Any person who, with purpose to injure, defraud or deceive any insurance company, files a statement of claim containing any false, incomplete or misleading information is subject to prosecution and punishment for insurance fraud, as provided in RSA 638:20.

## APPLICABLE IN NEW YORK

Any person who knowingly and with intent to defraud any insurance company or other person files an application for commercial insurance or a statement of claim for any commercial or personal insurance benefits containing any materially false information, or conceals for the purpose of misleading, information concerning any fact material thereto, and any person who in connection with such application or claim knowlingly makes or knowingly assists, abets, solicits or conspires with another to make a false report of the theft, destruction, damage or conversion of any motor vehicle to a law enforcement agency, the Department of Motor Vehicles or an insurance company, commits a fraudulent insurance act, which is a crime, and shall also be subject to a civil penalty not to exceed five thousand dollars and the value of the subject motor vehicle or stated claim for each violation.

### APPLICABLE IN OHIO

Any person who, with intent to defraud or knowing that he/she is facilitating a fraud against an insurer, submits an application or files a claim containing a false or deceptive statement is guilty of insurance fraud.

## APPLICABLE IN OKLAHOMA

WARNING: Any person who knowingly and with intent to injure, defraud or deceive any insurer, makes any claim for the proceeds of an insurance policy containing any false, incomplete or misleading information is guilty of a felony.

#### APPLICABLE IN WASHINGTON

It is a crime to knowingly provide false, incomplete, or misleading information to an insurance company for the purpose of defrauding the company. Penalties include imprisonment, fines and denial of insurance benefits.

## **APPENDIX J**

Near Loss Reporting Form

	Remedial Engineering, P.C. ble company name)									
PART 1: ADMINISTRATIVE INFORMATION										
Office: ☐ New York ☐ Massachusetts ☐ New Jersey [	☐ Illinois ☐ CA - Los Angeles ☐ CA - San Francisco									
	ject Principal:									
Project Name: Pro	ject Location:									
PART 2: LESSONS LEARNED INCIDENT DETAILS										
	e\Time Submitted (MM/DD/YYYY HH:MM):									
	? - Select all that apply (1-7) al (spill, permit exceedance, etc.) on of personnel (vehicle accident)  6. ☐ Property/Equipment Damage 7. ☐ Business Interruption									
Event Leading to Potential Injury/Illness:    Continue   Fourth										
Job Task*: Equ	uipment Involved*:									
WHAT HAPPENED? Do not include individuals' names. Ensure photos been obtained.  Summary (1-2 sentences. Provide brief description of the incident. Provided in the incident of the incident.	s, sketches, etc. are not personally identifiable unless written consent has vide facts only, no speculation or opinion):									
Incident Details (Brief factual details of what, where, when; include pho	tos, sketches, etc. as attachments):									
SERIOUS INJURY OR FATALITY (SIF): IF AN ACTUAL	SIF, USE EXISTING ROUX ACCIDENT REPORTING FORM									
Could this have resulted in a SIF?  Yes  No										
A potential SIF is defined as likely to have caused an injury resulting in significant physical body damage with probable long term and/or life altering complications.										
	significant physical body damage with probable long term and/or life									
altering complications.  INCIDENT INVOLVED:										
altering complications.  INCIDENT INVOLVED:  Roux Employee:  Yes  No Subcontractor Compar										
altering complications.  INCIDENT INVOLVED:  Roux Employee:  Yes  No Subcontractor Compar	ny Name:									
INCIDENT INVOLVED:  Roux Employee:  Yes  No Subcontractor Compar  INVESTIGATION FINDINGS AND FOR Date Investigation Team Assigned (mm/dd/yyyy):	NY Name: ATION TEAM ME JOB TITLE  REPORT QUALITY REVIEW									
INCIDENT INVOLVED:  Roux Employee:  Yes  No Subcontractor Compar  INVESTIGATION FINDINGS AND F	ATION TEAM  ME JOB TITLE  REPORT QUALITY REVIEW  rs and/or conditions may have contributed to the H&S Lessons Learned haviors/conditions; provide a narrative for each that explains how the									
INCIDENT INVOLVED:  Roux Employee: Yes No Subcontractor Compare INVESTIGATION FINDINGS AND FOR INVESTIGATION FINDINGS AND FOR INVESTIGATION SUMMARY: Determine from list below what behavior Incident. Then, use the "Multiple-Why Technique" for each of these behassociated Root Cause(s) was determined. Do not include individuals'	TATION TEAM  ME  JOB TITLE  REPORT QUALITY REVIEW  Tes and/or conditions may have contributed to the H&S Lessons Learned haviors/conditions; provide a narrative for each that explains how the mames.									
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INCIDENT INVOLVED:  Roux Employee: Yes No Subcontractor Compar  INVESTIGATION FINDINGS AND R  Date Investigation Team Assigned (mm/dd/yyyy):  INVESTIGATION SUMMARY: Determine from list below what behavior Incident. Then, use the "Multiple-Why Technique" for each of these behassociated Root Cause(s) was determined. Do not include individuals'  ROOT CAUSES: HOW TO REDUCE POSSIBILITY OF INCIDENT REG Selection of RCs and solutions reflects the analysis of investigation team. It is not	TON TEAM  ME  JOB TITLE  REPORT QUALITY REVIEW  To and/or conditions may have contributed to the H&S Lessons Learned haviors/conditions; provide a narrative for each that explains how the names.  CURRING  The meant as a legally binding conclusion as to causal factors and/or solutions.									
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Altering complications.  INCIDENT INVOLVED:  Roux Employee: Yes No Subcontractor Compar INVESTIGATION FINDINGS AND FINDING	TON TEAM  ME  JOB TITLE  REPORT QUALITY REVIEW  To and/or conditions may have contributed to the H&S Lessons Learned naviors/conditions; provide a narrative for each that explains how the names.  CURRING The meant as a legally binding conclusion as to causal factors and/or solutions.  JOB FACTORS:  E. LACK OF OR INADEQUATE PROCEDURES  F. INADEQUATE COMMUNICATION OF EXPECTATIONS REGARDING									

September 2013 00101Y1026/FORM

Person Responsible for Completion Target Date Completion Actual Date

Solution(s)
(Must Match Root Cause)

Behavior /

Condition

Root

Cause

	Completion	

QUALITY REVIEW Correct root cause(s) identified? Do root cause(s) and solution(s) match? Are solution(s) feasible / maintainable?

Name:

Job Title:

PART 4: Date Solutions were Implemented & Validated (Were Solutions Effective?)										
Solution Verifier / Validator Name and Job Title Details (of I & V performed)										
		Marifford (Mallidates Names and Jak Tida								

JOB TASK - Select the most appropriate one (primary job associated with incident-related work activity, avoid "Other" if possible) 7. Gauging 8. Geoprobe / Direct Push 18. System Startup 19. UST Removal (includes 1. Carbon Change 12. Pavement Cutting 2. Construction 13. Pump Test exposure and backfill) 9. Mobil Remediation (includes 3. Demolition 14. Sampling vacuum event and chemical injection) 4. Dewatering 15. Site Visit / Survey 20. Waste Management 5. Drilling (well install) 21. Well Abandonment 10. NAPL Recovery 16. Subsurface Clearance 6. Excavation / Trenching 11. O&M (remediation system) 17. System Install 22. Other:

EQUIPMENT INVOLVED	THAT CONTRIBUTED TO	H&S LESSONS LEARNED -	Select all that apply	
1. Air Stripper	25. Fire Extinguisher	51. Maintenance Tool, General	77. Safety Shoes / Boots	98. Vapor Extraction System
2. API Separator	26. Forklift	52. Manifold	78. Safety Vest / Clothing	99. Vapor-Phase Treatment
3. Automobile	27. Front End Loader	53, Manlift/Basket/Cherry Picker	79. Rope	System
Boom Material	28. Grader	54. Motor, Electric	80. Bailer	100. Other System, Type:
5. Bulldozer	29. Hammer	55. Oxidizer	81. Geoprobe	101. Surge Tank
6. Cable	30. Knife	56. Pallet	82. Hand Auger	102. Underground Tank
7. Carbon Drum / Vessel	31. Non-Powered Equipment	57. Piping	83. PID	103. Telemetry System
8. Chain Block	32. Powered Equipment	58. Piping, Hose	84. Multi-Gas Meter	104. Testing Devices
9. Compressor, Air	33. Drill	59. Piping, Injection/Mixing Point	85. Sample Container	105. Tractor Trailer
10. Control Panel (local)	34. Grinder	60. Hydrojet	86. Split-Spoon Sampler	106. Truck, Flatbed
11. Crane (mobile)	35. Hydraulic Torque Wrench	61. Centrifugal Pump	87. Sling	107. Truck, Pickup
12. Drill Rig	36. Powered Saw	62. Diaphragm Pump	88. Snow Blower	108. Truck, Tank Truck
13. Drilling Equipment, Vacuum	37. Impact Wrench	63. Reciprocating Pump	89. Snow Plow	109. Truck, Vacuum
14. Drum, Vertical	38. Saw	64. Regenerative Pump	90. Space Heater	110. Safety Valve
15. Dump Truck	39. Screwdriver	65. Rotary Pump	91. Air Sparging System	111. Block Valve
16. Electric Heater	40. Shears	66. Transfer Pump	92. Carbon Treatment System	112. Extraction Well
17. Electrical Power Supply	41. Shovel	67. Submersible Pump	93. Chemical Oxidation System	113. Monitoring Well
18. Engine, Combustion	42. Snip	68. Face Shield	94. Dual Phase Product	114. Recovery Well
19. Equipment Safety Grounding	43. Wrench	69. Fall Protection	Recovery System	115. Winch
20. Excavator / Power Shovel	44. Hoist	70. Gloves	95. Groundwater Pump	116. Wire Rope
21. Exclusion Zone Equipment	45, Hook/Clamp/Buckle, etc.	71. Hard Hat / Helmet	and Treat System	117. No Equipment Involved
22 Fan / Blower	46. Jack	72. Hearing Protection	96. POET System	118. MPT - Traffic Control
23 Fencing	47. Ladder, Extension	73. Respiratory PPE (Chemical)	97. Shed or Trailer	Devices
24 Filter	48. Ladder, Platform	74. Respiratory PPE (Particulate)		118. Not in List (describe):
	49. Ladder, Step	75. Safety Glasses		
	50. Lock Out / Tag Out	76. Safety Goggles		

SEPTEMBER 2013 00101Y1026/FORM

## **APPENDIX K**

## OSHA Log of Occupational Injuries and Illnesses

## OSHA's Form 300 (Rev. 01/2004)

## Log of Work-Related Injuries and Illnesses

**Attention:** This form contains information relating to employee health and must be used in a manner that protects the confidentiality of employees to the extent possible while the information is being used for occupational safety and health purposes.



U.S. Department of Labor
Occupational Safety and Health Administration

Establishment name

Form approved OMB no. 1218-0176

You must record information about every work-related death and about every work-related injury or illness that involves loss of consciousness, restricted work activity or job transfer,
days away from work, or medical treatment beyond first aid. You must also record significant work-related injuries and illnesses that are diagnosed by a physician or licensed health
care professional. You must also record work-related injuries and illnesses that meet any of the specific recording criteria listed in 29 CFR Part 1904.8 through 1904.12. Feel free to
use two lines for a single case if you need to. You must complete an Injury and Illness Incident Report (OSHA Form 301) or equivalent form for each injury or illness recorded on this
form. If you're not sure whether a case is recordable, call your local OSHA office for help.

m. If yo	ou're not sure whether a case	e is recordable, call your i	local OSHA office f	or help.						City			Sta	ate		
dent	ify the person		Describe t	he case			ify the c									
	(B) Employee's name	(C) Job title	(D)  Date of injury	(E) Where the event occurred	(F) Describe injury or illness, parts of body affected,		on the mos	E box for eac t serious out		Enter to days to ill wor	the number of he injured or ker was:	Chec	ck the ose on	"Injur e type	y" colu of illn	umn d iess:
10.		(e.g., Welder)	or onset of illness	(e.g., Loading dock north end)	and object/substance that directly injured or made person ill (e.g., Second degree burns on			Remaine	ed at Work	Away	On job	(M)	order	ory n	90 -1	in i
					right forearm from acetylene torch)	Death		Job transfer or restriction	Other record- able cases	Away from work	transfer or restriction	Injury	Skin dise	Respirat conditio	Poisonin	NI other
						(G)	(H)	(I)	(J)	(K)	(L)	(1)			(4) (5	5) (6
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Public reporting burden for this collection of information is estimated to average 14 minutes per response, including time to review the instructions, search and gather the data needed, and complete and review the collection of information. Persons are not required to respond to the collection of information unless it displays a currently valid OMB control number. If you have any comments about these estimates or any other aspects of this data collection, contact: US Department of Labor, OSHA Office of Statistical Analysis, Room N-3644, 200 Constitution Avenue, NW, Washington, DC 20210. Do not send the completed forms to this office.

Be sure to transfer these totals to the Summary page (Form 300A) before you post it.

Injury	Skin disorder	Respiratory condition	Poisoning	Hearing loss	All other Illnesses
(1)	(2)	(3)	(4)	(5)	(6)

## OSHA's Form 300A (Rev. 01/2004)

# Summary of Work-Related Injuries and Illnesses



U.S. Department of Labor
Occupational Safety and Health Administration

Form approved OMB no. 1218-0176

All establishments covered by Part 1904 must complete this Summary page, even if no work-related injuries or illnesses occurred during the year. Remember to review the Log to verify that the entries are complete and accurate before completing this summary.

Using the Log, count the individual entries you made for each category. Then write the totals below, making sure you've added the entries from every page of the Log. If you had no cases, write "0."

Employees, former employees, and their representatives have the right to review the OSHA Form 300 in its entirety. They also have limited access to the OSHA Form 301 or its equivalent. See 29 CFR Part 1904.35, in OSHA's recordkeeping rule, for further details on the access provisions for these forms.

Number of C	ases		
Total number of deaths	Total number of cases with days away from work	Total number of cases with job transfer or restriction	Total number of other recordable cases
(G)	(H)	(1)	(J)
Number of D	)ays		
Total number of da from work		otal number of days of job ansfer or restriction	
(K)	_	(L)	
Injury and II	Iness Types		
Total number of (M)			
) Injuries		(4) Poisonings	
		(5) Hearing loss	
) Skin disorders		(6) All other illness	es
Respiratory condit	ions		

#### Post this Summary page from February 1 to April 30 of the year following the year covered by the form.

Public reporting burden for this collection of information is estimated to average 58 minutes per response, including time to review the instructions, search and gather the data needed, and complete and review the collection of information. Persons are not required to respond to the collection of information unless it displays a currently valid OMB control number. If you have any comments about these estimates or any other aspects of this data collection, contact: US Department of Labor, OSHA Office of Statistical Analysis, Room N-3644, 200 Constitution Avenue, NW, Washington, DC 20210. Do not send the completed forms to this office.

Your establishment name _	
Street	
City	State ZIP
Industry description (e.g., M	Ianufacture of motor truck trailers)
Standard Industrial Classific	cation (SIC), if known (e.g., 3715)
OR	
North American Industrial	Classification (NAICS), if known (e.g., 336212)
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